Fabrication of a microfluidic platform for impedance analysis of cultured endothelial cell monolayers.

Stuart J. Williams 1981-

University of Louisville

Follow this and additional works at: https://ir.library.louisville.edu/etd

Recommended Citation
https://doi.org/10.18297/etd/1577

This Master's Thesis is brought to you for free and open access by ThinkIR: The University of Louisville's Institutional Repository. It has been accepted for inclusion in Electronic Theses and Dissertations by an authorized administrator of ThinkIR: The University of Louisville's Institutional Repository. This title appears here courtesy of the author, who has retained all other copyrights. For more information, please contact thinkir@louisville.edu.
FABRICATION OF A MICROFLUIDIC PLATFORM FOR IMPEDANCE ANALYSIS OF CULTURED ENDOTHELIAL CELL MONOLAYERS

By

Stuart J. Williams
B. S., University of Louisville, 2005

A Thesis
Submitted to the Faculty of the
University of Louisville
Speed School of Engineering
as Partial Fulfillment of the Requirements
for the Professional Degree

MASTER OF ENGINEERING

Department of Mechanical Engineering

December 2005
FABRICATION OF A MICROFLUIDIC PLATFORM FOR IMPEDANCE ANALYSIS OF CULTURED ENDOTHELIAL CELL MONOLAYERS

Submitted by:__________________________________________
Stuart J. Williams

A Thesis Approved on

August 9th, 2005

by the Following Reading and Examination Committee:

____________________________________
Robert Keynton, Ph.D., Thesis Director

____________________________________
Ellen G. Brehob, Ph.D.

____________________________________
Kevin M. Walsh, Ph.D.
ACKNOWLEDGEMENTS

I would like to thank Dr. Robert Keynton for his patience, guidance, and tutelage throughout graduate school. I couldn’t have created the devices without Mark Crain’s cleanroom instruction, training, and fabrication opinions. Dave Morgan, fellow classmate and friend, showed me how to culture the studied cells. Thanks to Scott Cambron, Thomas Roussel, and others in the BioMEMS and Cardiovascular group for their support. Sara Clemons accompanied me many late nights in the clean room and laboratory. Last, but not least, my parents, Joe and Ellen Williams. I couldn’t have made it through this without their continued support (and proofreading tips).
ABSTRACT

Fabrication of a Microfluidic Platform For Impedance Analysis of Cultured Endothelial Cell Monolayers

Stuart J. Williams

August 9th, 2005

Assessing the functionality of the endothelium can provide insight into the initiation and formation of arterial diseases. One of the most important functions of the endothelial layer is its permeability. The integrity of the cell monolayer and its ability to transport molecules can be assessed in vitro by investigating the electrical impedance.

In this study, a microfluidic platform was created using an electrode-patterned glass substrate and microfluidic poly(dimethyl siloxane) (PDMS) substrate. The electrode glass base of the structure was fabricated with platinum square electrodes of various sizes ranging from 10x10 \( \mu \text{m}^2 \) to 160x160 \( \mu \text{m}^2 \) and a larger, common counter electrode. Master microfluidic molds for PDMS casting were created by micro-milling Lexan® and photolithographically patterning SU-8. The microfluidic PDMS substrates reversibly and conformally bonded to the glass-electrode substrate. The microfluidic platforms were characterized by loading the microchannels with cell growth media alone, cell growth media and fibronectin, and cell growth media, fibronectin and human
umbilical vein endothelial cells and obtaining impedance spectra. The experiments were performed under both no flow and flow conditions.

Fibronectin did not significantly alter the collected impedance spectrum compared to media alone under no flow conditions. Under no flow conditions, impedance spectra measurements were able to detect the presence of cultured cells on the electrodes. The presence of fibronectin and various tested flow rates did not alter the impedance spectrum compared to media alone under static conditions. After further investigations, the microfluidic platform will become a versatile means of characterizing endothelial cell layer behavior.
# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>ACKNOWLEDGEMENTS</th>
<th>iv</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABSTRACT</td>
<td>v</td>
</tr>
<tr>
<td>LIST OF TABLES</td>
<td>x</td>
</tr>
<tr>
<td>LIST OF FIGURES</td>
<td>xi</td>
</tr>
</tbody>
</table>

## CHAPTERS

1. INTRODUCTION ................................................................. 1  
   A. Background ........................................................................ 1  
   B. Purpose of Study ............................................................ 3  
   C. Significance of Study ..................................................... 4  

2. LITERATURE REVIEW .......................................................... 5  
   A. The Endothelium.............................................................. 5  
      1. Anatomy of the Blood Vessel ........................................ 5  
      2. Endothelium Function ................................................. 5  
      3. Applied Mechanical Forces on Endothelium ................... 7  
   B. Endothelial Function Characterization .............................. 8  
      1. Transendothelial Electrical Resistance ........................ 9  
      2. Modeling of Measuring Electrodes ............................... 10  
      3. Existing Electrical Systems ....................................... 12  
   C. Shear Stress Investigations ........................................... 13  
      1. Shear Stress Analysis of Endothelial Cells .................... 14  
      2. Electrical Analysis of Endothelial Cells Under Applied Shear ....... 16  
      3. Comparison of Methodologies .................................... 18  

3. INSTRUMENTATION AND MATERIALS ..................................... 20  
   A. Facilities .......................................................................... 20  
   B. Device Fabrication .......................................................... 21  
      1. Materials Used .......................................................... 21  
      2. Chemicals Used ......................................................... 23  
      3. Instrumentation Used .................................................. 24  
   C. Cell Culture and Device Testing ...................................... 28  
      1. Materials Used .......................................................... 28
2. Chemicals Used ................................................................................................................................. 29
3. Instrumentation Used ............................................................................................................................ 29
4. Software .................................................................................................................................................. 32

4. PROCEDURE ............................................................................................................................................... 33
   A. Device Fabrication ................................................................................................................................. 33
       1. Electrode Substrate Design ................................................................................................................. 33
       2. Electrode Substrate Fabrication .......................................................................................................... 35
       3. Microfluidic Substrate Design .............................................................................................................. 38
       4. Microfluidic Substrate Fabrication ....................................................................................................... 39
       5. Topography Measurements .................................................................................................................. 44
       6. PDMS Bonding ..................................................................................................................................... 45
       7. Inlet and Outlet Ports ............................................................................................................................ 47
       8. Wiring .................................................................................................................................................... 49
   B. Cell Culture ............................................................................................................................................. 50
       1. Preparations of Cell Suspensions ......................................................................................................... 50
       2. Cell Culturing in Microchannels ......................................................................................................... 51
   C. Impedance Analysis ................................................................................................................................ 54
       1. Impedance Analyzer Settings ................................................................................................................ 54
       2. Impedance Analysis Tests .................................................................................................................... 54
   D. Analysis .................................................................................................................................................. 56
       1. Impedance Model ................................................................................................................................. 56
       2. Impedance Analysis ............................................................................................................................. 57
       3. Flow Theory ........................................................................................................................................ 59

5. RESULTS .................................................................................................................................................. 61
   A. Fabricated Platform ............................................................................................................................... 61
       1. Glass-Electrode Substrate ...................................................................................................................... 61
       2. Microfluidic PDMS Substrate .............................................................................................................. 64
       3. Complete Assembly ............................................................................................................................ 66
       4. Fluid Testing and Analysis .................................................................................................................... 67
   B. Cell Culture ............................................................................................................................................. 69
   C. Impedance Results ................................................................................................................................ 70
       1. Impedance Measurements in Microchannels with No Flow ............................................................... 70
       2. Impedance Measurements in Microchannels with Flow ................................................................. 77

6. DISCUSSION .......................................................................................................................................... 80
   A. Fabricated Platform ............................................................................................................................... 80
       1. Glass-Electrode Substrate Topography ................................................................................................. 80
       2. A Microfluidic PDMS Substrate ........................................................................................................... 81
       3. Fluid Testing and Analysis ....................................................................................................................... 83
   B. Cell Culture .......................................................................................................................................... 84
   C. Impedance Analysis ................................................................................................................................. 85
       1. Impedance Measurements in Microchannels ....................................................................................... 86

7. CONCLUSIONS ...................................................................................................................................... 88
# LIST OF TABLES

<table>
<thead>
<tr>
<th>TABLE</th>
<th>PAGE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Table 1. Trials for SU-8 fabricated molds.</td>
<td>43</td>
</tr>
<tr>
<td>Table 2. Topography measurements at different locations of the device.</td>
<td>63</td>
</tr>
<tr>
<td>Table 3. Expected and actual dimensions of electrodes and traces for each tested device.</td>
<td>63</td>
</tr>
<tr>
<td>Table 4. Percent error of dimensions of electrodes and traces for all three devices</td>
<td>64</td>
</tr>
<tr>
<td>Table 5. Summary of results for the first four SU-8 fabricated molds.</td>
<td>64</td>
</tr>
<tr>
<td>Table 6. Summary of results for the no-bake SU-8 fabricated molds.</td>
<td>65</td>
</tr>
<tr>
<td>Table 7. Showing calculated fluid flow parameters for 1rpm, 5rpm, 10rpm</td>
<td>69</td>
</tr>
<tr>
<td>Table 8. Standard deviations for the measured electrodes for with medium alone</td>
<td>72</td>
</tr>
<tr>
<td>Table 9. Mathematical model fit with medium alone</td>
<td>72</td>
</tr>
<tr>
<td>Table 10. Standard deviations for the measured electrodes for with medium and fibronectin</td>
<td>74</td>
</tr>
<tr>
<td>Table 11. Mathematical model fit with medium and fibronectin</td>
<td>74</td>
</tr>
</tbody>
</table>
# LIST OF FIGURES

<table>
<thead>
<tr>
<th>FIGURE</th>
<th>PAGE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Figure 1. Electrical model used for impedance analysis of cultured monolayers.</td>
<td>11</td>
</tr>
<tr>
<td>Figure 2. Port assembly, adaptor, and adhesive rings (Upchurch Scientific).</td>
<td>23</td>
</tr>
<tr>
<td>Figure 3. Heidelberg Laser Pattern Generator, DWL 66.</td>
<td>24</td>
</tr>
<tr>
<td>Figure 4. AB-M IR Mask Aligner.</td>
<td>25</td>
</tr>
<tr>
<td>Figure 5. Technics Sputtering System Series 4604.</td>
<td>25</td>
</tr>
<tr>
<td>Figure 6. Veeco Dektak 8 Advanced Development Profiler.</td>
<td>26</td>
</tr>
<tr>
<td>Figure 7. Ultra-High-Precision Micro Milling Machine.</td>
<td>27</td>
</tr>
<tr>
<td>Figure 8. March RIE CS 1701.</td>
<td>27</td>
</tr>
<tr>
<td>Figure 9. Environmental chamber for shear testing.</td>
<td>30</td>
</tr>
<tr>
<td>Figure 10. Ismatec rotary pump.</td>
<td>31</td>
</tr>
<tr>
<td>Figure 11. Agilent 4294A impedance analyzer.</td>
<td>31</td>
</tr>
<tr>
<td>Figure 12. Electrode pattern of the substrate (not to scale).</td>
<td>34</td>
</tr>
<tr>
<td>Figure 13. Electrode pattern of the 80 and 40 ( \mu m ) square electrodes (not to scale).</td>
<td>35</td>
</tr>
<tr>
<td>Figure 14. Electrode photomask.</td>
<td>36</td>
</tr>
<tr>
<td>Figure 15. Summation of electrode fabrication (Keynton 2004).</td>
<td>38</td>
</tr>
<tr>
<td>Figure 16. Superimposed drawings of microchannels and patterned electrodes</td>
<td>39</td>
</tr>
<tr>
<td>Figure 17. The micro-milled Lexan\textsuperscript{\textregistered} mold for the PDMS microfluidic piece.</td>
<td>40</td>
</tr>
<tr>
<td>Figure 18. Photomask used for SU-8 microfabricated mold.</td>
<td>41</td>
</tr>
<tr>
<td>Figure 19. Final SU-8 microfabricated mold.</td>
<td>44</td>
</tr>
<tr>
<td>Figure 20. Schematic of the environment for fluid flow testing.</td>
<td>56</td>
</tr>
<tr>
<td>Figure 21. An example of the Excel worksheet and Premium Solver package.</td>
<td>59</td>
</tr>
<tr>
<td>Figure 22. The resulting fabricated glass-electrode substrate.</td>
<td>61</td>
</tr>
<tr>
<td>Figure 23. Dektak 8 trace of glass-electrode substrate.</td>
<td>62</td>
</tr>
<tr>
<td>Figure 24. Profile results at different Y-coordinates (mm) of the fourth trial.</td>
<td>65</td>
</tr>
<tr>
<td>Figure 25. Profile results at different Y-coordinates (mm) of the fifth trial.</td>
<td>66</td>
</tr>
<tr>
<td>Figure 26. Top and bottom pictures of the assembled device.</td>
<td>67</td>
</tr>
<tr>
<td>Figure 27. Normalized impedance spectrum for each electrode geometry with medium alone.</td>
<td>71</td>
</tr>
<tr>
<td>Figure 28. Normalized impedance spectrum for each electrode geometry with medium and fibronectin.</td>
<td>73</td>
</tr>
<tr>
<td>Figure 29. Device 3 impedance spectrum across two measuring electrodes.</td>
<td>75</td>
</tr>
<tr>
<td>Figure 30. Device 2 impedance spectrum across measuring electrodes over time.</td>
<td>76</td>
</tr>
<tr>
<td>Figure 31. Impedance spectrum of cultured 40x40 ( \mu m^2 ) electrodes with a referenced measurement.</td>
<td>77</td>
</tr>
</tbody>
</table>
Figure 32. Impedance spectra under fluid flow for Device 1 with media alone............ 78
Figure 33. Impedance spectra under fluid flow for Device 1 with media and fibronectin.
........................................................................................................................................... 78
Figure 34. Impedance spectra under fluid flow for 40x40 µm$^2$ electrodes (Device 1) with
media alone. ................................................................................................................................... 79
Figure 35. Impedance spectra under fluid flow for 40x40 µm$^2$ electrodes (Device 1) with
media and fibronectin. .......................................................................................................................... 79
CHAPTER 1

INTRODUCTION

A. Background

In the last half century, investigations have been conducted to better understand the role of cardiovascular disease. As a result, death rates from cardiovascular disease have declined and American’s are living longer, healthier lives. However, cardiovascular disease continues to be the leading cause of morbidity and mortality in the United States. Coronary heart disease, congestive heart failure, stroke, hypertension, and arterial diseases are just a few of the leading types of cardiovascular disease, claiming more than 900 thousand lives in 2001 (Morbidity and Mortality: 2004 Chart Book). A number of investigators have theorized that transport across the endothelium is the primary cause of cardiovascular disease (Davies 1989, Grander 1986, Ross 1993). Thus, the role of endothelium permeability has been investigated in the function of cardiovascular diseases.

The endothelium is a confluent layer of flattened endothelial cells that line arteries, veins, and capillaries. This monolayer provides a multifunctional interface between blood and the vessel wall. Common endothelial activities include intracellular signaling, interaction between the surface and the underlying smooth muscle layer, and the exchange of molecular substances (Davies 1989, Grander 1986, Ross 1993). One of
the most important ways to characterize the endothelial cell monolayer is by investigating its barrier function, defined as the behavior of the transportation of molecules through the cell monolayer.

The permeability of an endothelial cell monolayer is determined by the diffusion of molecules through the individual cells themselves or between their clefts. Investigations of intracellular junctions were used to describe the access of medium to the underlying layers. This characteristic of the barrier function is important for the study of cardiovascular diseases when foreign substances diffuse through the endothelium. A number of factors affect the permeability of the endothelium, including shear induced forces from blood flow. A number of methods have been developed to investigate the behavior of the permeability of a cell monolayer (DePaola 1994, Gojova 2005, James 1995, Jo 1991, Takahashi 1997, Traub 1998). Unlike these methods, which can take an extended period of time to obtain results, electrical measurement techniques to analyze the cell monolayer provide “real time” data collection for barrier function characterization.

Initial endothelial electrical measurement techniques investigated cells cultured on permeable, filtered dishes (Cereijido 1978, Erben 1995, Hein 1992). The measured resistances across the cultured filter were directly related to the transcellular permeability of the endothelial cell layer, as lower resistance values were associated with more permeable monolayers. This technique would later be adapted to cell-culture well plates with gold electrodes (Giaever 1991, Keese 2002, Janshoff 1996, Wegener 1996, Wegener 2000, Seebach 2000). Experiments in a static environment were valid techniques but did
not provide physiologically relevant information since the cells were not exposed to fluid shear stress.

Two types of transendothelial electrical resistance apparatii were developed to investigate endothelial cells under applied fluid shear (DePaola 2001, Seebach 2000). The method developed by Seebach, et al. (2000) incorporated an applied shear force using a cone-and-plate device, whereby the plate contained the cultured cells and electrodes. An impedance analyzer recorded the transendothelial resistance, which was incorporated in a mathematical model to characterize the barrier function across the endothelial lining. Though the mathematical model is a sensible method of describing monolayer permeability, the cone-and-plate apparatus is not a reliable system and does not mimic physiological flow conditions. DePaola, et al. (2001) integrated an impedance system, called Electrical Cell Impedance Sensing (ECIS) (Giaever 1984, 1991, Keese 1990, 2002), into a parallel plate flow device. A major disadvantage of the ECIS analysis system is that it collects data at a predetermined frequency. A frequency spectrum of the measured impedance is needed to apply a comprehensive model describing the barrier function. Although both of these systems are viable, none of them are capable of determining endothelial cell permeability under physiological flow conditions.

B. Purpose of Study

The purpose of this study is to develop a microfluidic platform that uses impedance spectrum analysis to determine the permeability of cultured endothelial monolayers.
C. Significance of Study

The impedance analysis of a cultured endothelial monolayer directly gives insight to its behavior, including the permeability of the endothelial cells (Giaever 1984, 1991, Janshoff 1996, Keese 1990, 2002, Wegener 1996, 2000). Endothelium permeability is an important characteristic associated with the initiation and progression of some cardiovascular diseases (Davies 1989). The method of electrical analysis is advantageous over other permeability characterization studies since this method offers a “real-time” data collection and analysis of the monolayer. The proposed platform can also expose cultured cells to applied shear stress under fluid flow, which more closely mimics physiological conditions. Though impedance analysis studies of endothelial cells under induced shear have been developed by other authors (DePaola 2001, Seebach 2000), this study is the first microfabricated platform to incorporate continual flow conditions. Additionally, the measuring electrodes vary in size in order to analyze local inhomogeneities in cultured endothelial cells.
CHAPTER 2
LITERATURE REVIEW

A. The Endothelium

1. Anatomy of the Blood Vessel

The endothelium is a single layer of flattened, confluent cells that line all arteries and veins, including capillaries. Endothelial cell monolayers are the innermost layer of arteries and are in direct contact with blood. The endothelium is the innermost layer of the tunica intima, one of three main layers of an artery (Langley 1974). The monolayer rests on a layer of tissue connecting it to the underlying elastic membrane (Langley 1974). The next layer is the tunica media, composing of smooth muscle cells and connective tissue. The third, outermost layer is the tunica adventitia that consists mostly of loose collagen fibers (Langley 1974). By understanding the function of the endothelium and its interaction with the underlying layers the relationship between the formation of arterial disease and endothelial cell monolayer behavior can be understood.

2. Endothelium Function

The monolayer of endothelial cells provides a functional barrier between the blood and the underlying blood vessel layers. The endothelium is able to alter the arterial geometry, thus controlling aspects of fluid flow through the vessel. The monolayer
regulates molecular transport through the endothelium to the subendothelium. These functions are executed through a complex process of biological/biochemical signals sent by the endothelial cells in response to their environment.

Endothelial cells have the ability to control vascular tone with the production of vasodilators or vasoconstrictors. Types of vasodilators include nitric oxide, which causes smooth muscles cells to contract or relax and prostacyclin which is released by endothelial cells prevents platelet formation and blood clotting (Verrier, 1996). Vasoconstrictors such as endothelin-1, leukotrienes, and angiotensin II narrow the lumen of blood vessels. Through the production of vasodilators and vasoconstrictors, the endothelial cells control the diameter of the vessel; however, they also control solute interaction across the monolayer itself.

The endothelium is a regulator that mediates the molecular flux between the vessel wall and flowing blood. This permeability function is the most important characteristic of the monolayer. Substances are transported into the subendothelium by permeating through the intercellular junctions (paracellular) or through the cells themselves (transcellular).

Paracellular transport is characterized by the passage of molecules through a number of intercellular junctions. The four main types of intercellular junctions are tight junctions, adherens junctions, gap junctions, and desmosomes (Furuse 1993). Tight junctions prevent the passage of molecules and ions through the intercellular clefts, while adherens junctions provide strong mechanical attachment between cells. Gap junctions, on the other hand, are intercellular channels that provide passage of ions and molecules from cell-to-cell and desmosomes tightly join adjacent cells.
Transcellular transport describes the passage of ions and molecules through the cell itself by passing through the lipid bilayers to the subendothelium layers. The movement of ions through the cell membrane is determined by the size of the ion, its solubility in lipids, its size, and the concentration gradient (Langley 1974). Characteristics of the paracellular and transcellular behavior under different environments provide insight into the permeability function of the endothelium.

In addition, the endothelium controls other circulatory processes including blood coagulation, growth of underlying smooth muscle cells, white blood cell adhesion, transmigration, and lipoprotein uptake (Traub 1998). Thus, it is important to characterize the monolayer’s ability to control and regulate the diffusion of molecules, proteins, and other nutrients into the underlying layers of the artery in order to better understand the initiation and development of cardiovascular disease. In addition to chemical and biological regulation, the endothelium has been found to be highly responsive to mechanical forces which have been identified to potential initiators of cardiovascular disease (Davies 1988, Gojova 2005, Jo 1991, Keynton 2001, Phelps 2000, Traub 1998).

3. Applied Mechanical Forces on Endothelium

Throughout the circulatory system, the flow of blood subjects endothelial cells to two mechanical forces: a normal pressure force and a tangential shear force. The pressurized, pulsating nature of blood exerts a force on the vessel wall, causing the arteries to expand and contract. The vessel wall cushions most of this normal force; however, the endothelial cells absorb all of the shear force (Davies 1988). The tangential shear force is generated from the viscous flowing fluid through the arteries. Typically,
blood flow is laminar in vessels, but turbulence can occur in regions of stenosis or at sites of flow separation, (Davies 1989).

This applied shear force is directly related to the regulation of monolayer functionality, cell morphology, and motility (DePaola 2001). These mechanical forces alter endothelial permeability by modifying the responsive biochemical signaling pathway in endothelial cells. These signaling changes affect the endothelium’s function on both the cellular and molecular levels leading to rapid responses in the underlying tissue (Davies 1993). Furthermore, endothelial cells exposed to shear will rearrange their orientation to align with the direction of flow and elongate. The final shape of the endothelial cell is dependent on the microfilament structure of the cell and its rigidity under shear (Davies 1993, Gray 2002, Levesque 1986, Malek 1996). Permeability between the cell-to-cell junctions in the monolayer is directly related to this morphological adaption to shear stress (Seebach 2000). The monolayer’s reaction to hemodynamic forces is reversible as well as shear and time dependent (Davies 1989, Jo 2001).

B. Endothelial Function Characterization

A number of investigators have studied the effect of disturbed flow and/or applied shear on endothelial cell behavior (DePaola 1994, DePaola 2000, Gojova 2005, James 1995, Jo 1991, Phelps 2000, Seebach 2000, Takahashi 1997, Traub 1998). The function of the monolayer has been monitored by either chemical or electrical methods. Before
these investigations can be explained, it is important to understand the background behind the electrical techniques to describe the endothelial monolayer.

1. Transendothelial Electrical Resistance

The implementation of measured electrical resistances to characterize the behavior of cultured biological substances is not an uncommon practice. For example, transendothelial resistance (TER) provides insight into endothelium permeability. One common method of TER testing requires cells to be cultured on permeable filters, which divide the apparatus into two compartments (Cereijido 1978, Erben 1995, Hein 1992). Resistance measurements are recorded using a series of probes that provide a simple and direct method for real-time data collection and behavior analysis. One pair of electrodes is used to apply a DC voltage across the insert while a second pair detects the corresponding voltage drop. Ohm’s Law is used to determine the resistance across the filter using the resistance across an uncultured container as a correction factor.

Another methodology utilizes impedance analysis of an AC signal across a series of coplanar electrodes fabricated on a base substrate with cultured cells (Giaever 1991, Keese 2002, Janshoff 1996, Wegener 1996, Wegener 2000, Seebach 2000). The AC signal is sent across the electrodes and the impedance spectrum is recorded. This impedance verses frequency spectra is used to determine TER measurements as well as other parameters. The mathematical model used to determine these parameters is described in the next section below. The TER values describe the transcellular permeability of the cultured EC monolayer per unit area of covered electrode. Typical values of TER range from 3 ohm-cm² to 1000 ohm-cm² (Seebach 2000). Additionally,
TER can be monitored over a period of time to characterize cell motility and cell shape over a defined period (Giaever 1991, Keese 2002, Seebach 2000). However, a limitation to this technique is that a mathematical model for the electrical analysis of endothelial cell behavior is needed to perform offline calculations to quantify permeability.

2. Modeling of Measuring Electrodes

The most simplified electrical model that fully describes the characteristics of a cultured cell layer on an electrode using the minimum number of parameters is shown in Figure 1 (Janshoff 1996, Wegener 1996). The capacitor and resistor in parallel characterize or model the behavior of the cultured cell layer, while the resistor and capacitor in series deal with the interaction of the electrode to the electrolytic medium. $R_{\text{cell layer}}$ represents the transendothelial resistance describing the passage of molecules through the junctions between the cells. $C_{\text{cell layer}}$ symbolizes the behavior of transcellular activity through the cytoplasm of the cell itself; this transcellular path is close to $1\mu\text{F/cm}^2$ for most cell types (Seebach 2000). Ionic solutions, like cell culture medium, form an electrical boundary layer at the interface between the measuring electrode surface and the liquid. $C_E$ describes the dielectric properties of this interface and the interaction between the medium and the electrode. Last, $R_{\text{medium}}$ describes the conductivity of the cell culture medium. Note, the measured values of $C_{\text{cell layer}}$ and $R_{\text{cell layer}}$ are interpreted as an average value of the cells resting on the surface area of the electrode.
Parameters of the model are derived from impedance spectra obtained from the input of a low-peak (< 0.5V) AC signal. The range of frequencies for analysis varies but is typically within 1Hz to 1 MHz; the presence of cultured cells noticeably changes the spectrum at 100 Hz to 100 kHz (Janshoff 1996). The impedance spectra collected is inputted into a mathematical model and the variables are determined. Deviations of the model from the experimental data, at points below 1 kHz, may be attributed to the simplified treatment of the electrode/electrolyte interface as an ideal capacitor in series with the bulk conductivity (Janshoff 1996).

The measured electrical impedance of the endothelium is dependent on the input frequency and size of the measuring electrodes. Coplanar electrode analysis of cultured cells has been integrated in the systems of Wegener, et al (1996), Janshoff, et al (1996) and the Electric Cell-Substrate Impedance Sensing (ECIS) system developed by Giaever and Keese (Giaever 1984, 1991, Keese 1990, 2002). Due to the very large size of the counter electrode compared to the smaller, working electrodes the overall impedance of the system is dominated by the small electrodes, even if the larger electrode is covered with cells (Greve 2003). The cell layer cultured on top of the measuring electrodes needs to be confluent in order to monitor accurate impedance behavior. Larger intercellular

Figure 1. Electrical model used for impedance analysis of cultured monolayers.
gaps will influence the impedance spectra, but if the electrodes are uncovered the model is less accurate at predicting the electrical properties (Seebach 2000).

In previous work investigating this electrical analysis technique, the measuring electrodes were substantially larger than the cells they were measuring. Although large measuring electrodes interrogate the mean behavior of a cluster of cells, smaller electrodes provide insight into local cell/monolayer inhomogeneities. Greve, et al. (2003) produced a finite element model depicting the behavior of cells grown on small measuring electrodes. As cellular impedance analysis reaches a smaller scale, considerations for changing the model depicted in Figure 1 must be made in order to accommodate for more precise analysis of cellular characteristics previously ignored, such as focal adhesion sites.

3. Existing Electrical Systems

Two types of systems for impedance analysis of cell monolayers have been developed. The first is the Electronic Cell-Substrate Impedance Sensing (ECIS) system (Giaever 1984, Keese 1990, Giaever 1991) that monitors the behavior of cells at one defined frequency. The second type of system scans over a range of frequencies for impedance analysis (Janshoff 1996, Wegener 1996).

ECIS used one defined frequency (40 kHz) for rapid observation of impedance fluctuations due to cell motility. The investigated culture was grown in well plates that have one gold electrode of 250 µm in diameter and a larger, counter electrode in a coplanar arrangement. One of the well plates was uncultured, which was used to normalize the measured results in the other cultured wells. Impedance measurements
were taken with a lock-in amplifier and data was acquired and processed using a personal computer. By assuming the shape of cultured cells as a flat, circular disc, ECIS techniques can determine morphological parameters including the barrier function, spacing between the cell and the substrate, and cell membrane capacitance.

The other impedance analysis systems (Wegener 1996, Janshoff 1996) were used for the investigation of cultured endothelial cell layers. Electrodes of various sizes were fabricated on the culturing well base using gold films and an impedance analyzer is utilized to sweep through a range of frequencies. The measured values are fit to the model described in Figure 1 using least-squares methods. The disadvantage of these systems, compared to ECIS, is the frequency sweep data collection takes longer, i.e. from seconds to minutes, depending on the number of points and precision.

C. Shear Stress Investigations

Disturbed laminar flow occurs physiologically at sites of sharp curvature, abrupt narrowing and branching of blood flow. At junctions of redirected flow there exists complex flow patterns including regions of relatively high shear, stagnation (zero shear), and low shear varying with the cardiac cycle, resulting in reverse flow (Davies 1988, Keynton 1991). There have been a number of in vitro investigations that examined the endothelial cells’ response to applied shear in an attempt to analyze these regions of disturbed flow (DePaola 1994, 2000, Gojova 2005, James 1995, Jo 1991, Phelps 2000, Seebach 2000, Takahashi 1997, Traub 1998).
The most commonly used devices for investigating, in vitro, the effects of applied shear stress on the endothelial monolayer are parallel plate flow systems (DePaola 1994, 2000, Phelps 2000) and cone-and-plate systems (DePaola 1994, Seebach 2000). Parallel plate flow devices consist of plates arranged in parallel with a top plate being a glass substrate for culturing cells and a bottom substrate which contains the measuring electrodes. The gap between the two plates is small (< 1 mm) in order to produce a laminar, fully developed, two-dimensional flow. Shear forces of cone-and-plate systems are determined by the rotational velocity of the cone, the viscosity of the medium, and the angle between the cone and stationary plate (Davies 1988). The applied shear stress investigations are generally used to determine the relationship between hemodynamic forces, the endothelial layer, and the initiation of arterial disease.

Arterial vessel wall diseases have been linked to damaged or altered endothelium due to the effects of altered flow (Davies 1988, Satcher 1994). Erratic shear stress and disturbed flow hinder barrier function, which alters the permeability of molecules and thus, changes the intercellular signaling (Jo 1991, Phelps 2000). In some cases, endothelial cells do not reorient themselves to the direction of flow due to the disturbance (Traub 1998).

1. Shear Stress Analysis of Endothelial Cells

There are methods, other than electrical analysis, to determine and monitor the activities of endothelial monolayers under induced shear. Jo, et al. (1991) demonstrated that albumin permeability across the endothelial monolayer is acutely sensitive to shear stress. They subjected bovine aortic endothelial cells grown on polycarbonate filters to
shear stresses of 1 dyne/cm² and 10 dyne/cm². Albuminal levels increased by a factor of approximately 4 and 10 for lower and higher shear stresses, respectively, compared to static flow conditions.

James, et al. (1995) characterized the effect of shear on bovine aortic endothelial cell intracellular calcium transport. A shear rate of 15 dyne/cm² was applied to the cells through a confocal cell chamber constructed from polycarbonate. Fluorescence microscopy sensitive to intracellular calcium was used to analyze the cultured endothelial cells. They found that the cells responded to elevated shear levels by increasing intracellular calcium levels.

DePaola, et al. (1994) monitored the effects of shear stress gradients on bovine aortic endothelial cells in both a cone-and-plate system and a parallel plate flow chamber. Flow separation and disturbance was created by placing a rectangular strip in the system. DNA synthesis associated with cell proliferation was monitored by in situ monoclonal antibody detection of Bromodeoxyuridine (BrdU) incorporation. Their results showed that the cell proliferation rate was significantly reduced (only 8-10% of the cells in the disturbed region showed BrdU incorporation compared to 25% of the cells exposed to flow alone) in disturbed flows with shear gradients of 300dynes/cm²-cm.

Takahashi, et al. (1997) studied the mechanisms that induce endothelial cell shear stress response and signaling. They studied various signaling responses of bovine aortic endothelial cells to induced shear including the production of nitric oxide, activation of extracellular-regulated kinases, and tyrosine phosphorylation of focal adhesion kinase. These signaling events in endothelial cells were described as a string of temporary and mechanically distinct events. However, they determined that a more extensive
investigation is needed to compare these results with those induced by other stimuli besides mechanical forces, such as cytokines, growth factors, and hormones.

Traub, et al. (1998) examined data and reviewed investigations that involved stimulation of cellular responses via applied laminar shear stress that are crucial to endothelial cell function and atheroprotective. Coagulation, leukocyte adhesion, migration and proliferation, lipoproteins, endothelial cell survival and other characteristics of the endothelium were reviewed in correlation with past shear investigations. They determined that shear stress influences many of the processes relevant to the development of the atherosclerotic lesion. Additionally, they stated it is unlikely that fluid forces are the sole positive or negative atherogenic stimuli.

Gojova, et al. (2005) visually monitored the wound healing of bovine aortic endothelial cells cultured in parallel plate flow chambers subjected to shear levels of 3 to 19 dyne/cm². Image analysis was used to quantify the cellular response. The presence of shear decreased the rate of cellular spreading and wound closure compared to static conditions. Additionally, the presence of exogenous cholesterol significantly slowed cell migration.

2. Electrical Analysis of Endothelial Cells Under Applied Shear

Seebach, et al. (2000) setup an impedance spectroscopy system to investigate TER under an applied shear using a cone-and-plate apparatus. The rotating cone produced levels of shear stress ranging from 0 to 200 dynes/cm². An impedance spectra was obtained in the frequency range of 10 Hz to 1 MHz. TER values were normalized with TER values collected at the onset of shear ($t = 0$). Confluent cultures of porcine
pulmonary trunk endothelial cells showed a 15% increase in TER within fifteen minutes of applied shear, which was followed by a reduction in TER that continued to decrease by 20% below the initial value. The TER reduction was accompanied by an acceleration in endothelial shape change (monitored visually) and TER levels began to recover as the rate of endothelial shape change decreased.

Research conducted by Phelps, et al. (2000) used two separate parallel plate flow chambers for the analysis of endothelial cells under applied flow. In each apparatus, a rectangular step was introduced to create a disturbance in the flow. The first integrated a porous membrane for dextran extraction at different areas downstream from the site of initial disturbance. The second used electrodes connected to an ECIS device for TER analysis. After 5 hours of flow, areas of high shear stress gradients showed a 5.5-fold increase in transendothelial transport of dextran compared to a 2.9-fold increase in areas of undisturbed, fully developed flow. The TER values collected by ECIS were lowest in the vicinity of the highest shear stress gradients. They concluded these results demonstrate that endothelial permeability is regulated by spatial variation in shear stress, in vitro.

DePaola, et al. (2001) modified the ECIS and incorporated a parallel-flow apparatus to apply a shear force. This parallel plate flow system contained two channels with electrodes, one cultured with cells and the other no cells. Results in the cultured channel were normalized with data collected in the no cells chamber. Periods of uninterrupted flow were followed by time intervals of no-flow. TER values decreased on the onset of flow, started to recover after 15 min and after slightly overshooting its baseline values, decreased again with a prolonged exposure to flow. The observed
changes were reversible and varied with the duration of the preceding flow exposure. The results demonstrated that the impedance changes dramatically with changing endothelial parameters.

3. Comparison of Methodologies

As shown, there are many methodologies to characterize the behavior of endothelial cells under applied shear force. The disadvantage with monitoring endothelial cell behavior chemically or by other substance extractions is that the results are not immediate and the behavior of the system cannot be tracked in “real-time”. However, endothelial biochemical signaling behavior can be monitored. The impedance analysis methodologies are viable, but they too have disadvantages. The ECIS system takes into consideration an assumed disc-shape for a cell in its mathematical model. One of the characteristics of the endothelial cells under shear is their ability to change shape, and hence its permeability. The assumed shape of the cells under investigation would change under applied shear and the derived ECIS mathematical morphological model would be inaccurate. Additionally, the predefined frequency used in ECIS investigations was optimized for static analysis of cell motility and not to characterize the dynamic changes of the monolayer. In order to analyze a comprehensive mathematical model describing the endothelial layer, a frequency spectrum is needed.

The impedance analysis system described by Seebach, et al. (2001) applied shear via a plate-and-cone apparatus. Although it is a viable technique for applying shear to cultured endothelial cells, these systems to not represent the behavior of physiological fluid/endothelial cell interactions. Parallel-plate fluid flow systems are closer at
representing physiological conditions, but it is limited in the assumption of uniform parallel plate flow. Existing parallel-plate flow systems that integrate ECIS for data collection are also limited, with the disadvantages being the same as with the analysis system described above. Therefore, electrical impedance analysis in characterizing the behavior of cultured endothelial cells is preferred. A microfluidic platform would present a more complex fluid profile, thus more closely representing anatomical fluid flow conditions. However, the impedance analysis of cultured endothelial cells on a fabricated microfluidic platform has yet to be investigated.
CHAPTER 3

INSTRUMENTATION AND MATERIALS

A microfluidic platform was created for the purpose of impedance analysis of cultured monolayers. The device consisted of two components, a glass-electrode base for electrical measurements and a microfluidic substrate containing two channels. The fabricated electrode base contains the measuring electrodes of various sizes and a common, larger counter-electrode. The microchannels envelop these measuring electrodes while allowing the culturing of the cells. Before the design and procedures for the fabrication are explained, it is necessary to understand the use and purpose of equipment and materials.

A. Facilities

All fabrication and bonding steps occurred at the University of Louisville’s Lutz Microfabrication Laboratory. The 1500 square foot cleanroom facility is classified as class 100 and 1000. The glass-electrode substrates and SU-8 fabricated microchannel molds were created in the cleanroom environment. Device testing occurred on the University of Louisville’s Health Sciences campus where there was access to endothelial cells, cell culture media and additional testing instruments.
B. Device Fabrication

1. Materials Used

An ultra-flat low reflective chrome soda-lime glass substrate (Nanofilm, Westlake Village, CA) was used for the electrode substrate base. A thin layer of chrome separates the base glass layer and the photoresist surface. The soda-lime glass substrate (4”x4”x0.060”) contained photoresist type AZ 1518 with a thickness of 5300Å. The AZ 1518 photoresist is a positive resist, which means any part of the film exposed to UV light will be washed away in the developing solution. The baking step hardens the photoresist and prevents any contamination from additional solutes in the film.

The material chosen for the micro-milled microfluidic master mold base was Lexan® polycarbonate. A second, microfabricated mold using a negative photoresist for high aspect ratio features was investigated as another possibility for a master mold. A microfabrication process using a viscous, negative photoresist called SU-8-50 (MicroChem, Newton, MA) was developed to create a mold for the PDMS microfluidic substrate. The concentration of solvent (EPON SU-8 resin) in the solution determines the viscosity, which controls the film thickness. Using SU-8-50 produced many microstructures with films of more than 500µm and aspect ratios of more than 50 (Zhang 2001). These characteristics are advantageous for its application in microfluidics and patterning with PDMS. The data sheet for SU-8-50 can be found in Appendix C.

SU-8 is a negative resist, meaning that areas of the photoresist film exposed to UV light will harden and remain after submerging it in developing solution. The
developing solution for SU-8 is Baker BTS 220 (J. T. Baker, Phillipsburg, NJ). The time it takes to develop depends on the thickness of the applied SU-8.

Poly(dimethyl siloxane) (PDMS) was chosen as the material to be used for the microfluidic piece of the system. PDMS is also known as Sylgard 184 (Dow Corning, Midland, MI). PDMS is a low-cost material that has numerous, versatile microsystem applications. PDMS is used for rapid prototyping of microfabricated masters. Resulting molds are sealed directly to the substrate through irreversible and reversible techniques. The versatility of PDMS for microfluidic systems, cellular applications, and fabrication techniques led to its application for the microfluidic platform under investigation. Additionally, it is important to note that none of the previous electrical impedance studies have incorporated PDMS into their systems.

Nanoport assemblies (Upchurch Scientific, Oak Harbor, WA) were applied to the microfluidic platform. The components from Upchurch were made of an inert, biocompatible PEEK polymer and consisted of a nanoport reservoir assembly (part number N-131) that included a nanoport (¼ -28 FB), gasket, insert (10-32/¼-28), and an adhesive ring (N-100). A mechanical clamp, supplied in this assembly, was used to apply the pressure required for successful bonding during adhesive curing. The curing process occurred simultaneously with the curing of the conductive epoxy (H31D, Epoxy Technology, Billerica, MA) applied to the substrate later. This adaptor (P-664) fit into the 10-32 end of the nanoport insert and had a barb on the other end to fit the Pharmed tubing used in experimentation. Teflon tape was applied to the threads of the reservoir and adaptor to ensure a leak-free seal. Each of the parts used from Upchurch is shown in Figure 2.
Figure 2. Port assembly, adaptor, and adhesive rings (Upchurch Scientific).

Copper wires were stripped and attached to the electrode bonding pads with H31D conductive epoxy (Epoxy Technology, Billerica, MA). This is a 100% solids (no solvents) single component, electrically conductive silver epoxy designed for chip bonding. H31D is a soft, smooth, thixotropic paste with high conductivity (<0.005 ohm-cm).

2. Chemicals Used

After exposure, the substrate was placed in a developing bath. For positive resist type AZ 1518, the developing solution was a bath of MF-319 (Rohm Haas, Malborough, MA). The UV-exposed electrode pattern was washed away leaving the chrome surface exposed. The remaining hardened photoresist covered the rest of the substrate.

Nichrome Etchant TFN (Transene Company, Inc., Danvers, MA) was used to dissolve exposed chrome layers from the soda-lime substrate, leaving the underlying layer of glass. The glass was etched with Buffered Oxide Etch (BOE, Honeywell/GEM3, Morristown, NJ). At room temperature and manual agitation, BOE etched the glass at a rate of approximately 100Å /s. Material Safety Data Sheets for these chemicals can be found in Appendix A.
3. Instrumentation Used

Masks used for the fabrication processes were created in-house with a laser pattern generator (DW 66, Heidelberg Instruments, Torrance, CA) (Figure 3). The laser pattern generator is an extremely high-resolution imaging system where over half of a million dpi is achieved using a 40 nanometer writable access grid for exposing chrome plates or wafers. These masks were used to transfer patterns to the photoresist films after exposure to UV light.

![Figure 3. Heidelberg Laser Pattern Generator, DWL 66.]

The patterned masks were exposed to UV light from a mask aligner (Mask Aligner, AB-Manufacturing Inc., San Jose, CA) (Figure 4) and transferred it to the photoresist film. This system was designed to produce improved patterning results due to its ability to precisely control the intensity of UV light. The user-defined intensity/time function was controlled by the system, which reports an accuracy of better than ±2%.
Metal films for the electrodes were sputtered on the patterned layered-glass substrate using an RF/DC sputtering system (Series 4604, Technics Inc., Dublin, CA) (Figure 5). This device was used for the deposition of metallic materials onto the desired substrates. A high-voltage power supply was used to create a potential difference, causing the source material to be bombarded with ions from a plasma gas in the chamber (Argon). This bombardment releases molecules from the source into the chamber, depositing the material onto the substrate.
The topography of the glass-electrode and, later, the microfluidic channel dimensions were tested using a profilometer (Veeco, Dektak 8, Woodbury, New York) (Figure 6). The system has a vertical resolution of 7.5 Å, 1 sigma step height repeatability and a vertical range up to 1mm. This profiler was fitted with a 12.5μm radius stylus and used to measure the height of the microfabricated electrodes.

Figure 6. Veeco Dektak 8 Advanced Development Profiler.

A custom-made ultra-high-precision micro milling machine (Dover Instruments, Westboro, MA) (Figure 7) was used to create a master mold for fabricating the PDMS microfluidic substrates. The milling machine has three-axis resolutions of 1.25 nm, 1.25 nm, and 20 nm in the X-, Y-, and Z-directions, respectively. Micro-endmills and drills were made of micrograin tungsten carbide and can be as small as 10 μm in diameter. Endmills of 1 mm and 330 μm in diameter were used to fabricate the PDMS molds.
Plasma exposure was achieved using the cleanroom’s plasma asher/etcher (CS-1701, March Inc., Concord, CA) (Figure 8). This device is typically used to strip resist or etch patterns in wafers after photolithography. Particles from the substrate are released when high energy ions bombard the surface in the vacuum chamber. It is equipped with a process controller and solid state RF power generator.
An ultrasonic cleaner (8892, Cole Palmer Instruments, Chicago, IL) was used for the degassing of SU-8 photoresist. This instrument is equipped with a heater, digital timer, temperature monitor, and degas feature. This cleaner transforms low-frequency AC current into 42kHz high-frequency sound waves via an electroacoustic transducer.

A hot plate (Dataplate PMC 720 Series, Barnstead International, Dubuque, IA) was used for all fabrication baking steps and PDMS curing. This hotplate is a direct entry, programmable model. A microprocessor accurately controls and monitors the temperature. It has a digital display of all parameters as well as a digital count down timer with an “auto-off” control set.

C. Cell Culture and Device Testing

1. Materials Used

Frozen vials of “passage” 1 Human Umbilical Vein Endothelial Cells (HUVECs) were ordered from Cambrex (Pittsburg, PA) and arrived at “passage” 1. The “passage” of the HUVECs describes the number of times the cells were plated and grown on an artificial surface from the time they were harvested. The cells arrived frozen suspended in a solution of 90% PBS (phosphate buffered saline) and 10% DMSO (dimethyl sulfoxide). When thawed, these cells were plated and cultured with different media solutions.
Pharmed tubing with an inner diameter of 2.79 mm was used for experimentation. When applied with the rotary pump (MCP, ISM 915, Ismatec, Wertheim-Mondfeld, Germany), it is capable of producing a flow rate range of 0.21-52 mL/min. This tubing was also used for the injection of fibronectin and cell suspensions into the PDMS microchannels.

2. Chemicals Used

Trypsin Neutralizing Solution (Cambrex, Pittsburg, PA) was used to detach the cells from their adhered surface. HEPES buffered saline (pH = 7.4, Clonetics, BioWhittaker, Inc., Walkersville, MD) was used as a washing/cleansing solution for the cells, to ensure no traces of the previous medium existed. Endothelial Growth Media-2, EGM-2 (Clonetics, BioWhittaker, Inc., Walkersville, MD) was used as the HUVEC culturing medium. In some tests, a solution of fibronectin (Sigma Aldrich, Inc., St. Louis, MO) was applied. Fibronectin, an extracellular matrix protein, was applied to increase the adhesion of the cultured cells onto the substrate. Each of these solutions were used in sterile environments for the purposes of cell culturing and testing.

3. Instrumentation Used

All cell growth activities were monitored with an inverted, fluorescent microscope (Diaphot Phase Contrast-2, Nikon, Kanagawa, Japan). Impedance measurements and cell growth in the microchannels occurred inside a custom-made environmental chamber (Figure 9). The glove box chamber was constructed out of Lexan and was outfitted with sensors, including a temperature controlled heater and a CO₂
infusion mechanism. The temperature controlled heater (Electro-tech Systems, Inc., Model 513 A, Glenside, PA) kept the temperature near 37°C. The custom-made CO₂ controller was developed by the Bio-MEMS and Cardiovascular Research Group at the University of Louisville. The device consisted of a wall-mounted infrared CO₂ sensor that produced an output voltage that corresponded to the CO₂ concentration in the chamber. This voltage was converted and interpreted by the program in order to keep the CO₂ level of the chamber constant at 5%. A 9”x9” pan of water was placed in the chamber to maintain a constant relative humidity.

![Environmental chamber for shear testing.](image)

A rotary pump (MCP, ISM 915, Ismatec, Wertheim-Mondfeld, Germany) was used for the application of fluid flow through the system (Figure 10). It is capable of generating flow rates of 0.001-3700 ml/min depending on the pump head mounted and tubing diameter used. This pump was equipped with a pump head (MS/CA4-12, ISM 735) capable of flow rates of 0.001-57mL/min depending upon the size of the applied tubing and the rotational speed (1.0-240.0rpm at a resolution of 0.1rpm). With the Pharmed tubing described previously, flow rates of 0.21-52mL/min are possible.
A high-precision impedance analyzer (Agilent 4294A) and impedance probe (Agilent 42941A) were used for data collection (Figure 11). The analyzer has a frequency range of 40 Hz to 100 MHz with the ability to sweep linearly, logarithmically, or run through a list of given frequencies. The analyzer has an impedance range of 3 mΩ to 500 MΩ and it is capable of oscillation levels between 5 mV to 1 Vrms. The 42941A impedance probe enabled in-circuit impedance measurements of electrical components. This probe was used to analyze the measuring electrodes on the microfluidic platform.
4. Software

The Premium Solver Platform (Frontline Systems, Incline Village, NV) expansion for the Excel Solver tool package was used to optimize the fit between the collected data and the mathematical model. It replaced the standard Solver and provided two and a half times the number of decision variables and constraints. The Premium Solver Platform performed better on difficult models, including those involving very large and very small values. It has improved solving conditions for nonlinear problems where the standard Solver stops due to slow improvement in the target cell. This program analyzed collected data on a personal laptop equipped with a 3.20GHz Pentium 4 processor and Windows XP Professional operating system.
CHAPTER 4
PROCEDURE

A. Device Fabrication

1. Electrode Substrate Design

Two similar arrays of different sized square electrodes were used for impedance analysis of the endothelium (Figures 12 & 13). One set was for reference measurements in a microchannel with no cells and the second monitored the behavior of the cultured endothelial cell monolayer. Each array shared a larger, common grounded counter electrode. Five measuring electrodes of sizes 160 µm, 80 µm, 40 µm, 20 µm, and 10 µm square were implemented. With endothelial cells being approximately 10 µm in diameter, these series of electrodes were large enough to determine the behavior of a group of cells within the monolayer and small enough for localized detection.

These square electrodes were arranged in a straight line, with decreasing size, spaced 2.5 mm apart. This set of five decreasing electrodes was repeated within the same channel, for two similar electrodes per channel for a total of ten measuring electrodes. This process was repeated for the second microchannel in parallel for a total of 20 measuring electrodes. Similar sized electrodes from each channel were adjacent to one another, divided by the counter electrode. The counter electrode straddled the two
electrode arrays with a size of 400 µm by 25 mm. The center of each measuring microelectrode is 200 µm from the edge of the counter electrode. A series of twenty-two 1.5 mm-square bonding pads were connected to the electrodes by a series of wire traces. The larger wire trace from the bonding pads is 200 µm wide. The trace sizes from the microelectrodes, going from the largest to the smallest, are 40µm, 20µm, 10µm, 10µm, and 5µm respectively. The bonding pads are spaced 2.5 mm apart. The two sets of parallel bonding pads are spaced 51.4 mm, leaving room for the PDMS microchannel substrate to be placed on top of the electrode substrate.

Figure 12. Electrode pattern of the substrate (not to scale).
2. Electrode Substrate Fabrication

A technique developed at the University of Louisville for the fabrication of capillary electrophoresis devices was utilized for the fabrication of the designed glass-electrode substrate (Baldwin 2002, Keynton 2004). Metal electrodes were recessed in a glass substrate using an existing chrome/photoresist layer as a masking layer. The resulting flat surface permits the resolution and integrity of the fabricated electrodes to be maintained.

The AutoCAD electrode design file was converted to a Caltech Intermediate Format (CIF) file and uploaded on the laser pattern generator. The pattern was transferred to a six-inch, square, low reflective chrome-glass photomask (Nanofilm) with the electrodes and traces being transparent and the chrome remaining on the rest of the mask (Figure 14). The laser pattern generator was equipped with a 40mm head and the mask had G-line exposure at a rate of 150mJ/cm.
An ultra-flat low reflective chrome soda-lime glass substrate (Nanofilm) was used for the base of the electrode substrate. The patterned electrode photomask was placed directly on the chrome/photoresist substrate. It was exposed to UV light for 1.7 s from the ABM IR Mask Aligner to transfer the pattern. After exposure, the substrate was developed in a bath of MF-319 (Rohm Haas, Malborough, MA) for one minute. The UV-exposed electrode pattern was washed away leaving the chrome surface exposed and the hardened photoresist covered the remainder of the substrate.

Next, a DI wash followed with drying by a nitrogen gun before it was submerged in Nichrome Etchant TFN for three minutes. The exposed patterned chrome layer was etched away, leaving exposed glass (Figure 15, a). The substrate was cleaned and dipped in a solution of BOE under mild, manual agitation for one minute. The exposed glass was etched at a rate of approximately 100 Å/s. The resulting recessed pattern provided the space necessary for the subsequent step of the deposition of the near-flush metal electrodes (Figure 15, b). Titanium and platinum were sputtered onto the substrate using
the Technics RF/DC Sputterer. After a ten-minute presputter at 300 W, titanium was RF sputtered for 1.5 minutes at 300 W. The substrate was plasma etched for 30 s before platinum DC sputtering for ten minutes at 120 W. The resulting substrate was covered with the two layers of deposited metal (Figure 15, c).

After deposition, a “lift off” process was performed by placing the substrate in an acetone bath. The remaining photoresist layer dissolved when exposed to acetone, “lifting off” the metal deposited directly on the resist (Figure 15, d). Last, the substrate was placed in a Nichrome Etchant TFN to remove the final layer of chrome. The desired patterned electrodes resulted with the electrodes being nearly-flush with the flat soda lime glass base (Figure 15, e).
3. Microfluidic Substrate Design

The PDMS substrate has two microchannels, which were designed to accommodate the dimensions of the platinum electrode arrays. These channels contained one array of ten microelectrodes and share a portion of the larger counter electrode. The microfluidic PDMS substrate was sized to allow for adherence by conformal contact to the base substrate, leaving sufficient surface area for the insertion of nanoports. An AutoCAD drawing showing the alignment of the PDMS substrate to the electrode base is shown below (Figure 16).
Figure 16. Superimposed drawings of microchannels and patterned electrodes.

The width of each channel is 450 μm and is spaced 250 μm apart from one another. The parallel channels run 30 mm before they veer 45° towards the port insertion sites. Two methods of creating the molds for the microfluidic substrate were considered; one technique utilized micro-milling and the other used microfabrication techniques to create the relief features. The height of the microchannel structures depended on the type of fabrication method used for the mold. Micro-milling yielded better control over the height of the relief structures compared to photolithography.

4. Microfluidic Substrate Fabrication

a. Micro-Milled Mold. The Ultra-High-Precision Micro Milling Machine was used to create a mold constructed out of Lexan® for the PDMS microfluidic substrate
A solid model was developed in Solid Edge and later programmed into EdgeCAM. Two separate files in EdgeCAM were generated (printouts of these files can be found in Appendix B). The first program file was designed to use 1 mm diameter, 4-flute micro-grain tungsten carbide endmill for bulk machining of the substrate, which operated at a spin rate of 4000 rpm. The second program file was designed to machine with a 330 μm (0.013”) 2-flute micro-grain tungsten carbide endmill that operated at a spin rate of 6000 rpm. The smaller endmill was used to “finish” the mold in order to produce more well-defined edges on the microchannel relief structure. The depth of the mold was 0.25” with the microchannel features contained within a 1.5” x 2.5” rectangular perimeter.

b. SU-8 Fabricated Mold. A second mold for the PDMS microfluidic substrate was created using traditional microfabrication methods. The pattern for the microchannels were designed in AutoCAD, converted to CIF, and uploaded into the laser
pattern generator (Figure 18). The settings and procedure for the laser pattern generator were the same as in the creation of the electrode-pattern photomask. The photomask was created with the microfluidic features being transparent to allow for exposure of the negative photoresist (SU-8-50) to the UV light from the mask aligner. Prior to the application of the SU-8, a small quantity of refrigerated SU-8-50 was poured into a small bottle and degassed by placing the bottle in an ultrasonic cleaner (Cole Palmer 8892) for 30 min. This transfer process was performed in order to minimize bubble formation in the SU-8 during the application step.

Figure 18. Photomask used for SU-8 microfabricated mold.

Prior to SU-8 application onto the silicon wafer, an adhesion promoter solution was prepared by combining 95 mL of methanol, 5 mL of DI water, and 1 mL of adhesion promoter VM-651 (HD MicroSystems, Wilmington, DE). The wafers used in this study were single-side polished, n-type <100>, silicon (d = 100 mm, t ≈ 525 µm). The
adhesion promoter was applied to the polished side of the wafer and spun at 3000 rpm for 20 s, followed by the application of SU-8-50. It is important to note that the thickness of the resulting SU-8 film was determined by the amount of resist poured, the spin speed, and the duration of the rotation. For each fabricated mold, SU-8 was carefully poured onto the center of the wafer until the solution covered one-third of the surface. Due to its high viscosity, it was difficult to accurately control the amount of applied photoresist through direct pouring.

For the first trial, the SU-8 coated wafer was spun-on at 500 rpm for 30 s followed by a soft bake at 95°C for 45 min on a hotplate. Due to the SU-8 thickness, a large glass beaker covered the sample on the hotplate to contain the heat. The wafer was then aligned to the photomask and exposed to UV light for 100 s in the ABM mask aligner. The photomask was utilized in the proximity mode rather than contact mode since the photoresist film was not in a solid state after soft-baking.

After exposure, the wafer was post baked at 95°C for 45 min on a covered hotplate and allowed to cool overnight. The next day, the SU-8 was developed in fresh Baker BTS 220 (J. T. Baker, Phillipsburg, NJ) for 45 min followed by a 30 s rinse in Isopropanol. The wafer was washed with DI water and blow dried with a nitrogen gun. The fabricated mold was characterized with the Dektak 8 Profilometer to determine the dimensions of the relief structures at different locations on the wafer. This process was repeated three times with other fabricated, silicon wafers at different spin rates and bake times (Table 1). Unfortunately, each of these wafers produced uneven relief structures and the fabrication process had to be reconsidered.
The absence of precise leveling tools required a new approach to the fabrication process. The alternative approach selected, omitted the use of the hot plates completely, thereby eliminating the pre-exposure or post-exposure bakes on the uneven surfaces. Since the baking steps were primarily used to decrease the amount of foreign solvents in the resist and to improve adhesion of the resist to the wafer, these steps could be eliminated without adversely affecting the final product. Two wafers were fabricated using the non-baking process and a SU-8 spin rate of 500 rpm for 60 s, UV exposure time of 100 s, and developed in Baker BTS 220 for 45 min. Each wafer was profiled with the Dektak 8. The relief structures on these devices were profiled with the Dektak 8 and found to be more uniform than those fabricated previously.

After the wafer was developed, a layer of chrome and gold were sputtered on the substrate. This metallization layer was necessary because, through trial-and-error, it was discovered that a film of cured PDMS on the bare, patterned silicon wafer could not be easily removed and adhered to parts of the wafer. The chrome and gold were RF and DC sputtered at rates of 300 W for 1.5 min and 120 W for 4 min, respectively. With this addition of the gold surfaces, cured PDMS films were easily removed.

<table>
<thead>
<tr>
<th>Trial</th>
<th>Spin Rate</th>
<th>Soft Bake</th>
<th>Hard Bake</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>500rpm for 30s</td>
<td>95C for 15min</td>
<td>95C for 45min</td>
</tr>
<tr>
<td>2</td>
<td>500rpm for 30s</td>
<td>95C for 50min</td>
<td>95C for 60min</td>
</tr>
<tr>
<td>3</td>
<td>300rpm for 60s</td>
<td>95C for 45min</td>
<td>95C for 15min</td>
</tr>
<tr>
<td>4</td>
<td>400rpm for 50s</td>
<td>95C for 45min</td>
<td>95C for 15min</td>
</tr>
</tbody>
</table>

Table 1. Trials for SU-8 fabricated molds.
A retaining barrier for the PDMS was necessary to produce semi-consistent thicknesses for the PDMS microfluidic substrates. Many materials were tried, but the best border material was PDMS itself. A thick layer of PDMS was applied to the finished wafer and the microfluidic substrate was cut with a razor and removed by hand. The remaining PDMS served as a border for future poured molds (Figure 19).

![Figure 19. Final SU-8 microfabricated mold.](image)

5. Topography Measurements

The topography of the fabricated substrates was measured using the Dektak 8 Profilometer. Profiling was necessary to determine whether the patterned electrodes were flush to the glass surface. A near-flush surface would provide a flat base for culturing the endothelial cells and maintain electrode integrity for impedance measurements. Additionally, the laminar profile of the fluid flow would not become disturbed, so a consistent applied shear could be maintained throughout the microchannel. The sample was loaded in the Dektak 8 with settings of a 12.5µm radius stylus, force of 5mg, and
measuring resolution of 65kA. Various electrodes, traces, and points along the counter electrode were profiled.

6. PDMS Bonding

PDMS was formed by combining ten parts of the viscous, liquid silicone rubber base and one part of the curing agent. Once mixed, poured, and degassed it was heated to an elevated temperature of 65°C for four hours. During solidification, the elastomer was cross-linked via the hydrosilylation reaction between the vinyl (SiCH=CH₂) groups and hydrosilane (SiH) groups (Xia 1998). A data sheet for Sylgard 184 can be found in Appendix C.

PDMS has a few advantages over other polymers. Its surface can be oxidized for electro-osmotic flow applications (Ng 2002) and other methods are used to control its surface chemistry after oxidation (Duffy 1999). In a report published by Xia, et al. (1998) PDMS has good chemical stability, does not swell with humidity, is gas permeable, and is thermally stable up to 186°C. In addition, it is optically transparent up to 300 nm, it is isotropic, homogeneous, durable, and has the ability to bend. Disadvantages of PDMS include 1% shrinkage upon curing, its softness limits aspect ratio structures in design considerations, and it is subject to thermal expansion.

Once peeled from the molds, the PDMS microfluidic substrates were visually aligned and bonded to the glass-electrode substrate. Initial bonding protocol consisted of relying on conformal contact between the two, clean surfaces. However, leaking occurred, so an alternative approach was investigated. The alternative approach
consisted of exposing the two surfaces to oxygen plasma then meshing the two substrates together to form an irreversible bond.

The technique to irreversibly bond PDMS to glass, gold films, and other PDMS surfaces has been used in microfluidic applications for the past decade (Ma 1999, Owen 1994, Xia 1998). This bonding process occurs at a much lower temperature than glass fusion bonding, which is a major advantage. Both surfaces of interest were exposed to oxygen plasma which oxidizes the surfaces, creating dangling bonds that allow PDMS to covalently bond to the glass (Owen 1994). This produced a strong, irreversible bond between the PDMS substrate and the glass substrate.

Reversible bonding between the PDMS samples and the glass-electrode substrate was easily achieved with the conformal approach. As the surfaces were peeled and reapplied, the PDMS pieces would collect particles and residue which were difficult to remove from the microfluidic substrate. As the surfaces became more contaminated with particles or residue, the bond between PDMS and the glass surfaces became weaker. Additionally, the smooth surface of the PDMS microfluidic substrate obtained from the SU-8 mold exhibited better adhesion than the micro-milled Lexan® mold since the micro-milling procedure yielded a rougher surface.

PDMS substrates casted via the micro-milled and SU-8 molds and glass microscope slides were cleaned with ethanol then dried with a nitrogen gun. No DI water was used since the presence of water would inhibit the bonding process. The PDMS and glass were placed in the March RIE with the bonding surfaces faced upwards. The oxygen was set at 200 mTorr and the testing power and time settings varied from 1-300 s and 10-150 W, respectively. The variation of exposure was necessary since reports of

7. Inlet and Outlet Ports

The next step was to create an interface to attach the inlet and outlet tubing. Multiple techniques to accomplish this were considered for this device. First, holes were punched through the PDMS microfluidic substrate with different sized punchers (1/4, 3/8, and 1/2-inch diameters). The punched PDMS substrate was visually aligned and bonded with the glass-electrode substrate. Tubing of respective outer diameters was inserted into the PDMS punched holes. The exerted force on the PDMS holes from inserting the tubing caused the PDMS to deform and lift around the ports. Additionally, the movement of the attached tubing itself caused the microfluidic substrate to mechanically bend.

A second approach was explored where a hole-punched PDMS microfluidic substrate was sandwiched between the glass-electrode substrate and a separate piece of glass to prevent bending of the PDMS substrate. Four holes were drilled in the non-electrode piece of glass for the application of adhered nanoports. A water-lubricated drilling process was performed using a standard drill press operated at 3100rpm and fitted with a 5 mm diameter diamond-tipped drill bit. The holes were drilled slowly to prevent cracking and a piece of scrap glass was affixed underneath the sample to prevent additional chipping as the drill bit completely bored through the hole. These samples were placed in an ultrasonic cleaner with DI water and cleaning solution to remove
particles from its surface. Since the top surface of the PDMS was not exposed to oxygen plasma, an irreversible bond between the non-electrode piece of glass and the PDMS substrate was not achieved, and thus, leaking occurred.

The reversibly bonded PDMS microfluidic substrate proved to be sensitive to direct port adherence, thus, an alternative technique was developed to gain fluid access. First, the patterned glass-electrode substrate was drilled (using the same protocol as above) at the desired location for the nanoports. The nanoports, which provide tubing access to the inlet and outlets of the microchannels, were mounted on the bottom surface (non-electrode surface) of the electrode glass substrate to apply fluid flow from underneath the device. The advantage of this technique is that the forces exhibited by the tubing/nanoports were absorbed in the rigid glass base. Second, the PDMS microfluidic substrates were easily replaced and the electrode surfaces were cleaned without having to replace the tubing or ports.

The nanoport reservoir assembly and substrate were cleaned with Isopropanol and dried with nitrogen to aid in adhesion. The gasket was inserted into the nanoport with the adhesive ring applied to the bottom of the nanoport. The nanoport was visually aligned with the machined hole then pressed against the glass-electrode substrate. This procedure was repeated for a total of four port assemblies. Adhesive curing was accomplished simultaneously with the curing of the conductive epoxy, which will be described below. After curing, Teflon tape was applied to the threads and the nanoport insert was screwed into the port. Similarly, the adaptor was trimmed to fit into the device and was inserted into the nanoport assembly until hand-tight.
Leak tests were performed manually with a plastic syringe. Additional leak testing with the SU-8 molded PDMS microfluidic substrate was performed using the rotary pump equipped with a pump head and Pharmed tubing capable of delivering flow rates of 0.21-52 mL/min. After the PDMS microfluidic substrate was aligned and bonded to the glass-electrode substrate, Pharmed tubing was connected to one of the ports. Fluid was injected until leaking or PDMS separation was noticed visually. DI water was the medium used for these initial leak tests. There was no need to color the injected water as fluid flow through the PDMS was easily visualized through the semi-transparent material.

8. Wiring

Copper wires about ¾” long were stripped using a wire stripping tool and attached to the twenty-two electrode bonding pads with H31D conductive epoxy (Epoxy Technology, Billerica, MA). One end of the wire was manually dipped in the epoxy medium and carefully placed on the bonding pads. This wiring process was repeated for the remaining 21 bonding pads. The substrate was placed in an oven at 150°C for 30 min to cure the epoxy. The adhesive rings on the base of the nanoport assembly, which were clamped during heating, also simultaneously cured with the epoxy. After curing, the remaining port assembly was constructed.
B. Cell Culture

1. Preparations of Cell Suspensions

All procedures describing the passaging and injection of cells were performed in a sterile environment under a laminar flow hood to prevent cells from exposure to contaminants. All fluids were first heated to 37°C in a hot water bath. Microfluidic devices, tubing, and reservoir systems were sterilized by ethylene oxide gas at the University of Louisville Research Resource Center.

The cryogenically frozen vials of Human Umbilical Vein Endothelial Cells (HUVECs) were thawed at room temperature. Subsequently, the cells were suspended in 1mL of media (90% PBS, 10% DMSO), extracted with a micropipette then inserted into a sterile centrifuge tube. Trypsin Neutralizing Solution (1mL) was used to wash out the original vial to retrieve any remaining HUVECs and added to the centrifuge tube medium. An additional 0.5 mL of Trypsin was added to equal the 2.5 mL counter-solution used to balance the centrifuge during the 3 min, 3000 rpm spinning process.

Cells accumulated in the form of a pellet at the base of the tube. When the bulk of the medium was aspirated by means of a vacuum line, a pellet of concentrated cells remained. Endothelial Growth Media-2 (EGM-2) was introduced to the centrifuge tube and aspirated similarly to ensure removal of the old medium. Growth media (1 mL) was added to the centrifuge tube by a micropipette. Repeated injection and suction was applied to the tube to break up the cell pellet. This cell-suspended solution was added to a T-75 flask with additional EGM-2 for a total of 10 mL of growth media. The mixture was placed in the incubator at a temperature of 37°C and 5% CO₂.
The endothelial cells in the T-75 flask were grown to approximately 80% confluency. Confluency characterizes the percentage of monolayer coverage of the endothelial cells. Upon achieving this level of confluency, the cell medium in the T-75 flask was aspirated. The cells were washed with ~5 mL of HEPES buffered saline then the solution was aspirated. Trypsin (2.5 mL) was added to the T-75 flask and placed in the incubator for approximately one minute. The T-75 flask was taken out of the incubator and the cell suspended medium was collected and placed in a centrifuge tube. The previously mentioned centrifuge and plating processes were used to divide the larger pellet of cells three separate T-75 flasks. These samples were grown to 80-90% confluency and similarly plated on Petri dishes for testing or placed in vials to be cryogenically frozen. Through this economical growth and division process, the cells have reached passage 3 or 4.

2. Cell Culturing in Microchannels

A number of procedures were attempted to inject and culture the cells inside the microfluidic channels. Before the passaging and injection of the HUVECs, the microchannels were coated with fibronectin to provide an extracellular matrix for the cells to attach and grow. Fibronectin was manually injected with a sterile, one-time use, syringe. The syringe and nanoports were connected with Pharmed tubing that was manufactured to fit the rotary pump. Every three to four minutes, for a total of twenty minutes, the medium was circulated via manual injection. After this period, air was injected into the microchannels and the device was set aside during cell preparation.
The endothelial cells cultured on the Petri dishes were removed and centrifuged in the process described above. During this process, a separate Petri dish containing 10 mL of growth medium was placed in the incubator to acclimate the solution to the proper CO₂ level and temperature (37°C). This medium was then used to break up the concentrated pellet and created a cell-suspended solution that was later injected in the microchannels. The fibronectin coating and cell suspension processes defined above were the only two common procedures executed for culturing the cells inside the microchannels. The cell growth process was monitored visually with an inverted, fluorescent microscope for both the Petri dish and microchannel cell cultures. Upon inspection of the plated Petri dishes, HUVEC cell adherence appeared to occur in the first 1.5 to 2 hours after injection.

Initially, cell culturing inside the channels were performed using a closed system between the microfluidic platform and Pharmed tubing. Detachment and reattachment of Pharmed tubing to components of the system lead to trapping of air bubbles, which resulted in detachment of the cells as the bubbles passed through the channels. To prevent air trapping, clamps were applied to the inlet and outlet Pharmed tubing ports. However, inspection at 2 hours revealed no signs of cell adherence. After these attempts to culture cells inside the channels failed, it was decided to develop an alternative procedure to ensure cell attachment.

First, static cell culturing techniques were implemented by placing suspended HUVEC cells directly on the sterilized glass-electrode substrate. The culture dish containing the glass-electrode substrate was filled with growth medium, placed in an incubation chamber and monitored for attachment and growth. After 1.5 hours
incubating the cell culture dish was covered with a sterile Petri dish lid and visually inspected. The cells showed signs of spreading, which verified that the HUVEC cells were capable of growing on the glass-electrode substrate.

Next, films of either PDMS, Lexan® or Teflon were used in conjunction with the substrate to determine if a reaction between the PDMS substrate and cell culture medium and/or cells was occurring, which may inhibit cell growth. These materials were placed on top of the measuring electrodes and a solution of growth medium with suspended cells was injected on their surface. Openings were created on these films to expose the underlying measuring and counter electrodes to the suspended cells. The PDMS films (less than 3 mm thick) were cured and small slits were cut with a razor or holes were punched for the electrodes. A 1 mm thick slide of Lexan® was machined with a 1.5 mm-wide slit milled the length of the electrode arrays. The Lexan® slide was mechanically clamped to the glass surface, but leaking occurred during cell suspension so this material was abandoned from further testing. Finally, two strips of Teflon tape were visually aligned and pressed onto the substrate. A suspension of cells was applied to the PDMS and Teflon and, upon successful cell adhesion, impedance readings were taken and recorded.

This culturing technique was applied to culture cells inside the PDMS microfluidic channels. First, Teflon tape was aligned on the glass-electrode substrate to completely cover one set of electrodes as these were later used as referenced measurements. Next, another piece of Teflon tape was placed parallel to the first piece, leaving a small slit in between exposing the measuring electrodes. A suspension of cells was applied and left to adhere for 8 hrs. The medium was aspirated and the Teflon tape
was removed, leaving a narrow strip of cultured cells on the measuring electrodes. A sterile PDMS substrate was visually aligned and bonded to the glass-electrode base. EGM-2 growth medium was immediately injected in both channels. The device was transferred to the environmental chamber and impedance readings were recorded.

C. Impedance Analysis

1. Impedance Analyzer Settings

The input signal for the impedance analyzer was set at 10 mV peak-to-peak amplitude for a 100-point logarithmic frequency sweep of 40 Hz to 1 MHz. A bandwidth setting of “3” was selected from a scale of 1 to 5. Lower bandwidth settings decreased collection time at the expense of data precision. The different sized electrodes were manually probed and the impedance spectrum (impedance versus frequency) was collected from each electrode. Immediately following data collection, a spectrum was obtained from the adjacent channel containing the similar sized electrode. All of the saved trials were transferred to a 3.5” floppy disk in ASCII format. Data files were transferred to a laptop and analyzed within Excel.

2. Impedance Analysis Tests

Three different types of analyses were performed to determine the functionality and viability of the platform. First, the PDMS microfluidic substrate was aligned and injected with HUVEC cell growth medium, EGM-2. Measurements were taken and
recorded to compare the impedance spectrums and possible relationships between electrode sizes in each of the three fabricated devices. Additionally, one channel of the sterilized devices was injected with fibronectin in the same procedure explained previously for cell culturing. Next, EGM-2 medium was injected into the channels, being careful not to leave bubbles suspended in the microchannels. An impedance spectrum for each electrode was obtained and analyzed. Electrical analysis was also conducted for the cells cultured inside the PDMS microchannels mentioned above.

Additional experiments were conducted to determine the viability of the devices. The counter electrode should not contribute to the obtained impedance spectra. To test this, the impedance across the two largest electrodes (80x80 and 160x160) within the same channel was obtained. These results were compared to the sum of the impedances of the same electrodes collected across the counter electrode. Tests were also conducted to determine if the dielectric properties of the measuring electrode, or other electrical characteristics of the system, changed over time. In all cases, the impedance spectrums fit the mathematical model discussed previously (Figure 1).

Similar tests were conducted under flow conditions. Three microfluidic devices were tested under rotary pump settings of 1.0 rpm, 5.0 rpm, and 10.0 rpm, equivalent to flow rates of 0.21 mL/min, 1.05 mL/min, and 2.10 mL/min, respectively. Impedance spectra for static conditions (0 rpm) were also collected. Previous to impedance measurements under fluid flow, one microchannel was coated with fibronectin as described in the procedure for cell culturing via injection. The microfluidic platform was placed in the incubation chamber with the rotary pump and a custom-made reservoir assembly. The reservoirs were constructed out of Lexan® and were cylindrical in shape.
with a diameter of 31.75 mm. Four pieces of Pharmed tubing were injected with the gas-
exchanged EGM-2 medium and an additional 20 mL of medium were added to two
reservoirs. The solution-filled tubes were attached to the inlet and outlet connectors to
the reservoir and microfluidic device. The Pharmed tubing was arranged on the rotary
pump and, at different flow rates, impedance spectra were collected. A schematic of the
testing environment is show in Figure 20.

![Figure 20. Schematic of the environment for fluid flow testing.](image)

**D. Analysis**

1. Impedance Model

   Impedance spectra profiles were analyzed with Microsoft Excel and fit to the
simplified model (Figure 1) to determine the resistance and capacitance values. The total
impedance of the circuit, $Z$, in terms of frequency, $R_{cell\ layer}$, $C_{cell\ layer}$, $C_E$, and $R_{medium}$ is
given as:
\[ Z = \sqrt{(Z_1 \cos(\theta_1) + Z_2 \cos(\theta_2))^2 + (Z_1 \sin(\theta_1) + Z_2 \sin(\theta_2))^2} \]

where

\[ Z_1 = \sqrt{1 + \left(\frac{R_{\text{medium}} C_E \omega}{C_E \omega}\right)^2}, \quad \theta_1 = \arctan \left(\frac{-1}{R_{\text{medium}} C_E \omega}\right) \]

\[ Z_2 = \frac{R_{\text{c.l.}}}{\sqrt{1 + \left(\frac{R_{\text{c.l.}} C_{\text{c.l.}} \omega}{C_{\text{c.l.}} \omega}\right)^2}}, \quad \theta_2 = -\arctan \left(R_{\text{c.l.}} C_{\text{c.l.}} \omega\right) \]

\[ \omega = 2\pi f \]

Where \(Z_1\) is the total impedance contributed to the resistance of the electrolyte and the capacitance due to the electrode/electrolyte interaction and \(Z_2\) is the total impedance contributed from the presence of a cell monolayer. Calculations from these derived values can be found in Appendix D.

2. Impedance Analysis

The Premium Solver Platform expansion for the Excel Solver tool package was used to optimize the fit between the collected data and the mathematical model. After data for the investigated trial was imported, Excel cells for the four mathematical model variables \((R_{\text{cell layer}}, C_{\text{cell layer}}, C_E, R_{\text{medium}})\) were established with an initial, arbitrary guess \((R_{\text{cell layer}}, R_{\text{medium}} = 3000 \text{ and } C_{\text{cell layer}}, C_E = 5\text{E-10})\). Next, five columns for calculated values of \(Z_1, \theta_1, Z_2, \theta_2\), and \(Z\) referencing the four variable cells and the respective frequency in each row were created. A column taking the squared difference of measured and calculated values \((Z_{\text{measured}} - Z_{\text{calculated}})^2\) for each row was established. The creation of a cell calculating the summation of these values followed. The Solver was executed to minimize this referenced summation cell. A range of nonnegative values for
the four variables was also established. The range of values for the two resistance and capacitance variables are 0 to 100 MΩ and 0 to 1 μF, respectively; each were well within the expected range of values. The method of optimization for the Solver was chosen as “Standard Evolutionary”. The Solver changed the four variable cells until it reached a specified time, iterations, precision, and/or convergence. Although rare, if the solver calculated a solution that did not correspond to the data (i.e. a flat line or a variable reached its limit), the Solver executed again with new initial guesses for the variables based on previous iterations. This process was repeated and variables were obtained for each recorded impedance spectrum. Similarly, this technique was applied for the analyses of uncultured surfaces where, with the absence of a cell monolayer, $Z_2$, $R_{c.l}$, and $C_{c.l}$ are 0. An example of the Excel Worksheet with the Solver parameters is shown below (Figure 21).
3. Flow Theory

It is important to determine the applied fluid force with respect to the channel geometry and fluid flow for the application of shear stress testing. Additionally, derivations for laminar flow verification and entrance length were calculated. Viscosity and other fluidic parameters for the working medium, EGM-2, could not be found in literature. The fluid was assumed Newtonian and the viscosity was assumed similar to water. The flow rate and cross sectional area were assumed constant throughout the microfluidic channel.

All equations and values for the proceeding calculations were referenced from F. M. White (2002). Shear measurements for fluid flow associated with this investigation is expressed in dyne/cm² (1 dyne = 1x10⁻⁵ N, 10 dyne/cm² = 1 N/m² = 1 Pa). The rotary
pump with the current pump head and Pharmed tubing produces a volumetric flow rate range of 0.21-52mL/min. It is assumed that this flow rate increases at a linear rate proportional with the rotational settings range of 1.0-240.0 rpm. Calculations were recorded at 1.0 rpm at a volumetric flow rate of 210 µL/min, or 3.5x10⁻⁹ m³/s. At 37°C the density, ρ, and viscosity, µ, of EGM-2 are 993.37 kg/m³ and 0.000692 kg/m³, respectively. With the channel width of w and height of h the Reynolds number, Re, was calculated from the hydraulic diameter, DH, and velocity, V, by

\[ DH = \frac{2hw}{h+w} \]

\[ V = \frac{Q}{hw} \]

\[ Re = \frac{\rho V DH}{\mu} \]

The entrance length, Le, for fully developed laminar flow is given as

\[ Le = 0.06 Re DH \]

The shear stress applied to the confluent cell layer was also calculated. Shear was calculated from the average frictional force around the perimeter of a rectangular duct. The product of the frictional factor and Reynolds’s number, fRe, is determined from the aspect ratio of the width and height of the microchannel. The average shear associated with this geometry is calculated by

\[ f = \frac{8 \tau}{\rho V^2} \]

\[ \tau = \frac{f \rho V^2}{8} \]

This applied shear force is linearly proportional to the rotational velocity of the pump head.
CHAPTER 5

RESULTS

A. Fabricated Platform

1. Glass-Electrode Substrate

A total of three glass-electrode substrates were fabricated as described above. Figure 22 represents a typical glass-electrode substrate. Shown is the patterned metal deposited on the glass surface. Twenty-two bonding pads are at the ends of the wire traces. A common, grounding electrode divides the two sets of ten measuring electrodes.

Figure 22. The resulting fabricated glass-electrode substrate.
Topography measurements with the profilometer were performed and a representative graphical result of two 200 µm-wide bonding pad wire traces is depicted in Figure 23. The sputtered metal layers protruded about 90 nm above the surface of the glass. Similar traces were obtained from each of the tested microfabricated glass-electrode substrates and a summary of the results can be found in Table 2.

Figure 23. Dektak 8 trace of glass-electrode substrate.
The surface dimensions of the measuring electrodes were visually inspected under a microscope. The electrode shapes and sizes were different than the dimensions specified in the AutoCAD drawing for mask creation. For example, the expected 10 µm square electrode was nearly elliptical in shape with the major and minor axes being 15 µm and 12 µm, respectively. The wire trace from this electrode was measured at 10 µm instead of the expected width of 5 µm. These actual measured geometries were used in all the impedance analysis calculations. Table 3 summarizes the expected and actual geometries for each substrate. Table 4 shows the percent error for all three devices. Measured values of these surface areas were accurate within 10%.

<table>
<thead>
<tr>
<th>Electrode</th>
<th>Area (µm²)</th>
<th>Trace (µm)</th>
<th>Area (µm²)</th>
<th>Trace (µm)</th>
<th>Area (µm²)</th>
<th>Trace (µm)</th>
<th>Area (µm²)</th>
<th>Trace (µm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>100</td>
<td>5</td>
<td>141</td>
<td>7</td>
<td>165</td>
<td>10</td>
<td>153</td>
<td>9</td>
</tr>
<tr>
<td>20</td>
<td>400</td>
<td>10</td>
<td>660</td>
<td>13</td>
<td>720</td>
<td>15</td>
<td>696</td>
<td>14</td>
</tr>
<tr>
<td>40</td>
<td>1600</td>
<td>10</td>
<td>2230</td>
<td>13</td>
<td>2480</td>
<td>15</td>
<td>2480</td>
<td>14</td>
</tr>
<tr>
<td>80</td>
<td>6400</td>
<td>20</td>
<td>7060</td>
<td>23</td>
<td>7400</td>
<td>25</td>
<td>7400</td>
<td>24</td>
</tr>
<tr>
<td>160</td>
<td>25600</td>
<td>40</td>
<td>27100</td>
<td>43</td>
<td>27900</td>
<td>46</td>
<td>27900</td>
<td>45</td>
</tr>
</tbody>
</table>

Table 3. Expected and actual dimensions of electrodes and traces for each tested device.
Table 4. Percent error of dimensions of electrodes and traces for all three devices

<table>
<thead>
<tr>
<th>Electrode</th>
<th>Percent Error</th>
<th>Trace (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>53.0 ± 12.0</td>
<td>73.3 ± 30.6</td>
</tr>
<tr>
<td>20</td>
<td>73.0 ± 7.5</td>
<td>40.0 ± 10.0</td>
</tr>
<tr>
<td>40</td>
<td>49.8 ± 9.0</td>
<td>40.0 ± 10.0</td>
</tr>
<tr>
<td>80</td>
<td>13.9 ± 3.1</td>
<td>20.0 ± 5.0</td>
</tr>
<tr>
<td>160</td>
<td>7.9 ±1.8</td>
<td>11.7 ± 3.8</td>
</tr>
</tbody>
</table>

2. Microfluidic PDMS Substrate

The dimensions of micro-milled mold were within ±5µm of the design parameters. Table 5 summarizes the first four SU-8 fabricated molds and Figure 24 shows typical profile data obtained on one of the four molds. The fabrication of these molds used baking procedures on an uneven hotplate surface. This fabrication technique led to uneven microchannel profiles with a difference in almost 100µm. Figure 24 continues to exhibit the fabricated uneven surface as shown in the resulting near-linear height profile as it is measured at different locations from its inlet. Note that the Dektak 8 profiler cannot determine the aspect ratio of the SU-8 structures, only their height. The pointed feature between the two relief structures in Figure 24 is the result of the profiler not able to “touch-down” in between the structures.

Table 5. Summary of results for the first four SU-8 fabricated molds.

<table>
<thead>
<tr>
<th>Trial</th>
<th>Spin Rate</th>
<th>Soft Bake</th>
<th>Hard Bake</th>
<th>Highest Profile</th>
<th>Lowest Profile</th>
<th>Channel Width</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>500rpm for 30s</td>
<td>95°C for 15min</td>
<td>95°C for 45min</td>
<td>223μm</td>
<td>130μm</td>
<td>580μm ± 5μm</td>
</tr>
<tr>
<td>2</td>
<td>500rpm for 30s</td>
<td>95°C for 50min</td>
<td>95°C for 60min</td>
<td>&gt;300μm</td>
<td>167μm</td>
<td>515μm ± 5μm</td>
</tr>
<tr>
<td>3</td>
<td>300rpm for 60s</td>
<td>95°C for 45min</td>
<td>95°C for 15min</td>
<td>462μm</td>
<td>366μm</td>
<td>490μm ± 5μm</td>
</tr>
<tr>
<td>4</td>
<td>400rpm for 50s</td>
<td>95°C for 45min</td>
<td>95°C for 15min</td>
<td>353μm</td>
<td>264μm</td>
<td>440μm ± 5μm</td>
</tr>
</tbody>
</table>
Two SU-8 molds were created without the baking steps as described in Table 6. The absence of the uneven baking surfaces allows the relief structures to have a level topography (Figure 25). The difference between the highest and lowest profiles is almost 10µm. The average height and width of the “H” and “I” wafer microchannel relief were 232µm and 441µm, respectively, with a channel division width of 230µm. The length of the channel was within 1mm of designed length.

<table>
<thead>
<tr>
<th>Trial</th>
<th>Spin Rate</th>
<th>Soft Bake</th>
<th>Hard Bake</th>
<th>Highest Profile</th>
<th>Lowest Profile</th>
<th>Width</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>500rpm for 60s</td>
<td>N/A</td>
<td>N/A</td>
<td>227µm</td>
<td>236µm</td>
<td>438µm ± 5µm</td>
</tr>
<tr>
<td>6</td>
<td>500rpm for 60s</td>
<td>N/A</td>
<td>N/A</td>
<td>229µm</td>
<td>237µm</td>
<td>428µm ± 5µm</td>
</tr>
</tbody>
</table>

Table 6. Summary of results for the no-bake SU-8 fabricated molds.
3. Complete Assembly

After the glass-electrode substrate was drilled, wires were bonded, and nanoports adhered, PDMS microfluidic substrates were cured in the SU-8 no-bake molds and aligned visually to the substrate (Figure 26). On the electrode side of the substrate, the PDMS microfluidic piece is aligned such that the ends of the microchannels are at the drilled holes for the nanoports. The PDMS structure dividing the two microchannels straddles the counter-electrode, leaving exposed surface area in each channel. The wires were arranged as to not impede the PDMS adherence and prevent accidental shorting. Ports were adhered on the backside to of the glass-electrode substrate.
4. Fluid Testing and Analysis

The PDMS substrate from the SU-8 mold did not leak during manual injection at low flow rates (< 1 mL/min), however, the PDMS substrates produced from the micro-
milled piece leaked easily. Very thin PDMS microfluidic films (< 5 mm) also leaked under injected flow. The PDMS microfluidic substrate did not leak with the rotary pump connected to the system using a pump rate of 15.0 rpm (equivalent to flow rates of 3.25 mL/min).

With a channel width of 441 µm and height of 232 µm the Reynolds number, $Re$, hydraulic diameter, $D_H$, velocity, $V$, and entrance length, $L_e$, were calculated for a flow rate associated with the 1.0 rpm setting on the rotary pump (0.21 ml/min). Calculated values of $Re$ and $L_e$ are proportional to the applied pump settings.

$$D_H = \frac{2hw}{h+w} = 304 \text{ µm}$$

$$V = \frac{Q}{hw} = 3.42 \text{ cm/s}$$

$$Re = \frac{\rho V D_H}{\mu} = 14.9$$

$$L_e = 0.06 Re D_H = 272 \text{ µm}$$

Under these conditions, the profile is laminar and the flow field was fully developed at 272 µm per applied 1.0 rpm.

With an assumed rectangular cross sectional area, the aspect ratio was $h/w = 0.53$. The product of the frictional factor and Reynolds’s number, $fRe$, was interpolated from a table of values (White 2002) to be 61.74. The average shear associated with this geometry was calculated by

$$fRe = 61.74$$

$$f = \frac{8\tau}{\rho V^2} = \frac{61.74}{Re}$$

$$\tau = \frac{61.74 \rho V^2}{8Re} = \frac{61.74V\mu}{8D_H} = 6.01 \text{ dyne/cm}^2$$
Thus, the applied shear for a flow rate of 0.21 mL/min was found to be 6.01 dyne/cm². This applied shear force was linearly proportional to the rotational velocity of the pump head, so at 15.0 rpm, which is the maximum pump rate before leaking occurred, the applied shear was over 90 dyne/cm². Table 7 shows some of these calculated fluid parameters at 1, 5, and 10 rpm.

<table>
<thead>
<tr>
<th>Pump Setting</th>
<th>Q (mm³/s)</th>
<th>V (mm/s)</th>
<th>Re</th>
<th>f</th>
<th>L_e (mm)</th>
<th>τ (dyne/cm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.0 rpm</td>
<td>3.5</td>
<td>34.2</td>
<td>14.9</td>
<td>4.14</td>
<td>0.272</td>
<td>6.01</td>
</tr>
<tr>
<td>5.0 rpm</td>
<td>17.5</td>
<td>171</td>
<td>74.7</td>
<td>0.83</td>
<td>1.36</td>
<td>30.04</td>
</tr>
<tr>
<td>10.0 rpm</td>
<td>35.0</td>
<td>342</td>
<td>149</td>
<td>0.41</td>
<td>2.72</td>
<td>60.09</td>
</tr>
</tbody>
</table>

Table 7. Showing calculated fluid flow parameters for 1rpm, 5rpm, 10rpm.

B. Cell Culture

After fibronectin coating, attempts were made to seed suspended HUVECs into microchannels; however, endothelial cell attachment/adherence to the fibronectin coated microchannels was limited. During the application of fluid flow, all cells were removed from the microchannel, so alternative HUVEC plating/growth techniques were developed. As described in the procedures chapter, a series of experiments were performed to investigate the effect of the PDMS material on cell attachment and culturing. Specifically, PDMS, Lexan® and Teflon films were applied to the glass-electrode substrate and solutions of suspended HUVECs were placed in the openings of the material onto the glass-electrode substrate.
Cells on PDMS films showed signs of attachment within the first two hours, as expected, but the cells that were suspended in the slit above the measuring electrodes did not attach until after three to four hours. Similar results were observed with the Lexan® slide; however, due to a lack of adhesion between the Lexan® and glass surfaces, the growth medium (and a few cells) between the interface layers. Growth was observed on the device with Teflon tape as well.

Using this technique, cells could be easily patterned and transferred to the PDMS microchannels without the need for injecting cell-suspension media. The cell adhesion was strong enough that, upon the injection of EGM-2 medium, air bubbles flowing through the channels would not detach the adhered HUVECs. Through careful PDMS alignment, only a minimal amount of cells would detach or become trapped at the interface. Additionally, the bulk of the residue left behind by the original cell culturing was removed with the Teflon tape, improving the adhesion between the PDMS substrate and the glass-electrode base. There were no signs that Teflon tape inhibited cell growth or attachment.

C. Impedance Results

1. Impedance Measurements in Microchannels with No Flow

Impedance spectrum for each of the measuring electrodes was obtained for each device for media alone, media and fibronectin, and media, fibronectin and HUVECs under no flow conditions. After fabrication and sterilization, media was injected into the
channels and impedance measurements were obtained for Device 1 (Figure 27), Device 2 (Appendix F) and Device 3 (Appendix F). These impedance values for each of the four similar electrodes were averaged and normalized with respect to the exposed surface area of the electrodes. As anticipated, the normalized impedance values decreased as the area of the electrodes increased. The standard deviations for these values at 1.0kHz, 10kHz, 100kHz, and 1MHz are shown in Table 8. As frequency increased, the standard deviation of each electrode decreased.

![Normalized impedance spectrum for each electrode geometry with medium alone.](image)

**Figure 27.** Normalized impedance spectrum for each electrode geometry with medium alone.
Table 8. Standard deviations for the measured electrodes with medium alone.

<table>
<thead>
<tr>
<th>Frequency</th>
<th>10x10</th>
<th>20x20</th>
<th>40x40</th>
<th>80x80</th>
<th>160x160</th>
</tr>
</thead>
<tbody>
<tr>
<td>1kHz</td>
<td>39.93</td>
<td>6.06</td>
<td>19.39</td>
<td>2.02</td>
<td>3.35</td>
</tr>
<tr>
<td>10kHz</td>
<td>10.00</td>
<td>1.00</td>
<td>3.16</td>
<td>0.26</td>
<td>0.63</td>
</tr>
<tr>
<td>100kHz</td>
<td>3.42</td>
<td>0.17</td>
<td>0.46</td>
<td>0.05</td>
<td>0.11</td>
</tr>
<tr>
<td>1MHz</td>
<td>1.47</td>
<td>0.09</td>
<td>0.08</td>
<td>0.03</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Impedance results from Device 1 were analyzed with the Premium Solver and fit to the mathematical model (Figure 1) with $R_{cell\ layer} = C_{cell\ layer} = 0$. Without the presence of cells, these parameters were omitted. The results of this fit are shown in Table 9. Similar results for Device 2 and Device 3 can be found in Appendix F. Generally, the resistance of the bulk electrolyte decreases and the capacitance increases as the size of the electrode increases.

Table 9. Mathematical model fit with medium alone.

<table>
<thead>
<tr>
<th>Electrode</th>
<th>R (Ω)</th>
<th>C (F)</th>
<th>C (F/cm²)</th>
<th>R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>10x10</td>
<td>28000</td>
<td>2.52E-10</td>
<td>9.99E-06</td>
<td>0.9833</td>
</tr>
<tr>
<td>20x20</td>
<td>12500</td>
<td>3.68E-10</td>
<td>7.29E-06</td>
<td>0.9921</td>
</tr>
<tr>
<td>40x40</td>
<td>12400</td>
<td>4.39E-10</td>
<td>6.78E-06</td>
<td>0.9866</td>
</tr>
<tr>
<td>80x80</td>
<td>6200</td>
<td>9.53E-10</td>
<td>6.46E-06</td>
<td>0.9938</td>
</tr>
<tr>
<td>160x160</td>
<td>6700</td>
<td>9.34E-10</td>
<td>2.24E-06</td>
<td>0.9867</td>
</tr>
</tbody>
</table>

After testing with media alone, fibronectin was injected in the left channel and media was injected in the right channel for a comparative study. Subsequently, impedance measurements were acquired for each electrode. Upon completion of these experiments, the devices were cleaned, fibronectin was loaded in the right channel and
media in the left channel and impedance measurements were again obtained. Figure 28 summarizes the results obtained with the fibronectin-coated channels for Device 1 (Device 2 and Device 3 data can be found in Appendix F). Again, as anticipated, the impedance values increased as electrode size decreased. Comparing the impedance values to the media alone (Figure 27), it was noticed that the presence of fibronectin does not affect the impedance spectrum. The standard deviations at various frequencies and results from the model fit can be found in Table 10 and Table 11, respectively. The observations of the devices with the presence of fibronectin were similar to the previous study of media alone.

![Normalized impedance spectrum for each electrode geometry with medium and fibronectin.](image)

Figure 28. Normalized impedance spectrum for each electrode geometry with medium and fibronectin.
<table>
<thead>
<tr>
<th>Frequency</th>
<th>10x10</th>
<th>20x20</th>
<th>40x40</th>
<th>80x80</th>
<th>160x160</th>
</tr>
</thead>
<tbody>
<tr>
<td>1kHz</td>
<td>20.35</td>
<td>6.01</td>
<td>14.33</td>
<td>0.48</td>
<td>2.50</td>
</tr>
<tr>
<td>10kHz</td>
<td>2.92</td>
<td>1.15</td>
<td>2.17</td>
<td>0.19</td>
<td>0.45</td>
</tr>
<tr>
<td>100kHz</td>
<td>1.90</td>
<td>1.57</td>
<td>0.34</td>
<td>0.04</td>
<td>0.07</td>
</tr>
<tr>
<td>1MHz</td>
<td>0.90</td>
<td>0.39</td>
<td>0.08</td>
<td>0.01</td>
<td>0.01</td>
</tr>
</tbody>
</table>

Table 10. Standard deviations for the measured electrodes for with medium and fibronectin.

<table>
<thead>
<tr>
<th>Electrode</th>
<th>R (Ω)</th>
<th>C (F)</th>
<th>C (F/cm²)</th>
<th>R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>10x10</td>
<td>28600</td>
<td>2.20E-10</td>
<td>8.73E-06</td>
<td>0.9868</td>
</tr>
<tr>
<td>20x20</td>
<td>15800</td>
<td>3.64E-10</td>
<td>7.21E-06</td>
<td>0.9909</td>
</tr>
<tr>
<td>40x40</td>
<td>12800</td>
<td>4.30E-10</td>
<td>6.64E-06</td>
<td>0.9861</td>
</tr>
<tr>
<td>80x80</td>
<td>6470</td>
<td>9.57E-10</td>
<td>6.48E-06</td>
<td>0.9921</td>
</tr>
<tr>
<td>160x160</td>
<td>6180</td>
<td>9.91E-10</td>
<td>2.38E-06</td>
<td>0.9867</td>
</tr>
</tbody>
</table>

Table 11. Mathematical model fit with medium and fibronectin.

In order to determine whether the impedance of the counter electrode contributed to the output obtained from the measuring electrodes, the spectrum across the two largest measuring electrodes were analyzed. Figure 29 illustrates the impedance values across the 80x80 µm² and 160x160 µm² electrodes compared to the sum of their individual impedance values obtained across the counter electrode. The counter electrode did not contribute to the impedance output signal as evidenced by the curve for the sum of the individual impedance values directly overlying on top of the impedance values for the counter electrode.
Additionally, the electrode’s dielectric properties were monitored over time as the electrode was exposed to the cell growth media. Impedance spectra from the five measuring electrodes from Device 2 were measured repeatedly in approximately five minute intervals (Figure 30). The magnitude of the impedance decreases over time as the electrodes are exposed to the cell growth medium (EGM-2). The magnitude of this drop in impedance decreases as the size of the measuring electrode increases. The smaller the electrode, the greater the influence of dielectric properties. The relationship between this trend and the model (Figure 1) without the presence of cells is that the capacitance ($C_E$) and resistance of the bulk electrolyte ($R_{medium}$) increases and decreases with time, respectively.
Figure 30. Device 2 impedance spectrum across measuring electrodes over time.

Impedance spectra were obtained from cells adhered to the channels inside the PDMS substrate. One channel contained the HUVEC culture and the other only growth media, the latter was used as a reference reading to compare to the cell culture data. The following data was obtained from Device 2. Figure 31 shows a set of 40x40 µm$^2$ impedance spectrums with two cultured electrodes having approximately 70% and 30% of their surfaces covered with attached HUVECs. Included with the data is the measured, reference electrode. The magnitude of impedance shifted as the concentration of cells covering the measuring electrode increased. Very little or no cell coverage resulted in a minimal impedance shift.
2. Impedance Measurements in Microchannels with Flow

The impedance spectra for Device 1 were collected at 1.0 rpm, 5.0 rpm, 10.0 rpm and under static conditions (0 rpm) for a channel filled with media alone (Figure 32) and the other coated with fibronectin (Figure 33). The normalized impedance spectra of the 40x40 µm² electrodes in the two channels are shown in Figures 34 & 35. Since the collected impedance spectrum for each electrode was time dependent, the order in which the flow rates were induced were 0 rpm, 5.0 rpm, 1.0 rpm, then 10.0 rpm. As expected, the magnitude of the impedance spectrum decreased as the electrode size increased. As shown in previous static trials, the presence of fibronectin does not alter the spectra. Additionally, the studied volumetric flow rates did not alter the impedance spectra. To verify this trend, impedance spectrum were collected from the 40x40 µm² electrode from Device 2 and the 10x10 µm² and 160x160 µm² electrodes from Device 3 (Appendix F).
Figure 32. Impedance spectra under fluid flow for Device 1 with media alone.

Figure 33. Impedance spectra under fluid flow for Device 1 with media and fibronectin.
Figure 34. Impedance spectra under fluid flow for 40x40 \( \mu \text{m}^2 \) electrodes (Device 1) with media alone.

Figure 35. Impedance spectra under fluid flow for 40x40 \( \mu \text{m}^2 \) electrodes (Device 1) with media and fibronectin.
CHAPTER 6
DISCUSSION

A. Fabricated Platform

1. Glass-Electrode Substrate Topography

This near-flush surface provided a flat culturing base for the endothelial cells and maintained electrode integrity for impedance measurements. Keeping the glass-electrode topography level prevented the fluidic profile in the microchannels from additional disturbance while maintaining the integrity of the cultured cell monolayer. Additionally, a smooth surface aided in the conformal contact bonding of the PDMS. Although the electrodes protrude from the surface around 90 nm, better results are possible with practice and a better operational knowledge of the sputtering machine. Other than sputtering time and RF/DC power input, parameters that affect the amount of deposited metal on the substrate surface are the base chamber pressure and the chamber pressure prior to plasma exposure.

The resulting electrode pattern on the soda-lime glass was observed visually. The shapes of the electrodes did not match the design shape. Specifically, most of the electrodes had rounded corners, but the mask pattern was designed with sharp corners. In some cases, the electrodes produced were elliptical in shape. For example, the 10x10
µm² electrode had an elliptical shape with the major and minor axes being 15 µm and 12 µm, respectively. Although a number of microfabrication techniques could be at fault for this alteration in geometry, the biggest factor was the initial patterning mask. The mask was not printed at the highest quality, meaning resolution of the features would be lost and rounded corners become more prevalent. Upon visual inspection, features of the patterning mask were identical to the fabricated substrate. The mask quality, therefore, was the major contributing factor in the resulting, distorted geometry.

Estimated electrode surface areas were calculated by dividing the structure into smaller geometrical shapes and summing all of its parts. Observations and measurements were initially conducted with the Dektak 8 profiler. However, upon inspection of the curved surfaces, visual characterization was determined to be more efficient. The microscope was equipped with a scale inside the eyepiece and, with 100X magnification, 1-micrometer resolution was possible. These measurements, including the width of the wire traces, were importance for capacitance and other impedance analysis calculations. Cell layer characterization from impedance analysis was dependent on the geometry of the electrode, so precise control over its fabrication was crucial.

2. A Microfluidic PDMS Substrate

The micro-milled Lexan® mold for the PDMS substrate was machined within all specified parameters. The uniformity of the mold was excellent and, with the recessed base, the thickness of the PDMS substrate could be well controlled. However, this machining method produced a rough surface with ridges which were generated during the
machining process. This adversely affected both reversible and irreversible bonding processes since a smooth conformal surface is needed.

Uses of resist SU-8 as means for thick, high aspect-ratio structures have been well established (Zhang 2001). For this reason wafers with PDMS relief structures were created from SU-8 photoresist. Initial trials for fabricating wafers proved troublesome. The profile across the channel relief structure showed a non-uniform structure with a large height disparity ($\approx 100\mu m$). The height disparity increased nearly linearly in the hardened SU-8 thickness from one end of the structure to the other. Three additional wafers were fabricated similarly at different spin rates and resulted in the same characteristic topographical profile.

The cause of the slope in the relief structure was determined to be due to the hotplates not being level. The uneven surface of the hotplate was causing the viscous SU-8 fluid to shift during the baking process. Attempts were made to level the hotplates manually with a handheld bubble-gage, but the difference between the structure’s highest and lowest profiles was improved to approximately 30 $\mu m$. Nevertheless, a more uniform thickness was desired to have stringent control of fluid velocity throughout the length of the channel.

By eliminating the baking step, the amount of error was significantly reduced. The difference between the highest and lowest points on the fabricated relief structures was within 10 $\mu m$. The high point occurred at the center of the relief structure with the lowest points occurring at the inlet and outlet portions of the microchannel. Without the baking step additional fabrication problems were encountered. For example, on closer inspection, the sputtered metal on the SU-8 features had a “cracked” surface. The reason
for the rough surface was not known; however, it could be directly related to the absence of baking steps. Without baking, excess solvents in the SU-8 are present, potentially resulting in the rough metal surface. Initially, this surface did not prove to be a problem since multiple substrates of PDMS were peeled without lifting or releasing of metal particles; however around the fourth or fifth time metal particles were found in the PDMS microfluidic substrates. This problem was easily overcome by fabricating new molds which, without the use of baking steps, could be produced at a faster rate.

Whether or not the pre-exposure bake was conducted, the SU-8 film did not harden for UV exposure. For this reason, proximity patterning was necessary as contact patterning resulted in SU-8 adhering to the mask. The minute separation between the mask and the film caused the dimensions of the exposed features to be altered. This alteration was caused by the diffusion of UV light through the transparent patterns and created larger, exposed patterned areas on the photoresist film.

3. Fluid Testing and Analysis

Both reversible and irreversible bonding was attempted for the PDMS microfluidic substrates. The PDMS substrates were cut, cleaned, and exposed to oxygen plasma for irreversible bonding. After trial and error experiments, it was determined that the optimum exposure to oxygen plasma for ¼” thick samples of PDMS was either 25 W for 20 s or 10 W for 45 s.

Within the first 10 seconds of venting the chamber, the two surfaces needed to come into conformal contact with each other. Also, if the contacted PDMS substrate was lifted off the surface and immediately placed back on the glass surface, adhesion would
not occur. It has been noted that a drop of anhydrous ethanol applied to the surface after plasma exposure delays the bonding between the surfaces, leaving enough time for slight alignment adjustments (Khademhosseini 2004). There were successful trials of smaller samples of delayed PDMS bonding but it proved difficult to provide a thin, uniform layer of anhydrous ethanol to the microfluidic substrates in the amount of time necessary for complete irreversible bonding. Through reversible, conformal contact bonding the PDMS microfluidic substrate was adjusted and realigned multiple times. The irreversible oxygen plasma bonding technique proved difficult to align and bond simultaneously.

Reversible bonding was chosen for a number of reasons. First, it could be easily aligned visually and applied to the glass-electrode substrate. Also, the PDMS microfluidic piece could be peeled away for ease of cleaning or replacing platform components. Additionally, patterning PDMS substrates could be applied immediately before the initiation of microfluidic testing.

B. Cell Culture.

Attempts to culture cells inside microfluidic channels by injection methods were unsuccessful. Through methodical testing it was determined that the presence of PDMS or the glass-electrode substrate were not the causes of the problem. Through static cultures on films of PDMS and Teflon, a new way to pattern and place cells inside the microfluidic PDMS substrate was adapted. The Teflon films were not shown to inhibit cell growth and were sterilized easily. The application and removal of Teflon tape was easier than other methods described. For these reasons, it became a logical choice to use this material to place and pattern a strip of cells. The PDMS placement is crucial.
substrate cannot be misalign so much that the cells crushed and dislodged between the
glass and PDMS interfaces, nor can the time to align the substrate be lengthy since the
cells are without medium during the process. This method proved to be a simple fix to
the problem of culturing cells via injection techniques.

C. Impedance Analysis

As stated before, the size of the measuring electrodes contributed significantly to
the measurement values since the magnitude of impedance decreased with increased
electrode surface area. In addition, the traces leading to the electrodes also played a role
in the impedance spectrum output. Specifically, the PDMS substrate was manually
placed over the glass-electrode substrate and visually aligned. Alignment variations of
the PDMS substrate to the glass-electrode substrate resulted in either an increase or
decrease in the impedance signal dependant upon whether the exposed area was larger or
smaller than the previous alignment. For these reasons, the exposed area for each
electrode was measured for each experiment. Error in the surface measurements could
also lead to error in normalized impedance calculations. Future fabricated platforms
should incorporate a buried trace, leaving it unexposed to media in the channel.

The results in Figure 28 verify that the counter-electrode does not influence the
impedance signal since the amount of area exposed by the counter-electrode is
sufficiently large compared to the measuring electrode. The impedance measured
between the 80-square and 160-square electrodes is given as the sum of impedance across
each electrode. The impedance measured individually across each measuring electrode to
the counter electrode is given as the sum of impedance across each electrode and the
counter electrode. Since the value across the measuring electrodes is equal the sum of those individually (as shown in Figure 28), then the impedance across the counter electrode is zero. The impedance values are influenced heavily on the amount of exposed surface area. As the exposed electrode area decreases, the measured impedance increases. Since the area of the counter-electrode is much greater than the measuring electrodes, the portion of collected impedance due to the presence of the counter-electrode is negligible.

1. Impedance Measurements in Microchannels

There were a number of measurements that, upon observation, were determined to be inconsistent with typical spectra for the measuring electrode. This can be accomplished from a crossing of wires or a loose probing connection. The impedance spectrums from trials without the presence of cells were assumed to be equivalent to a resistor and capacitor in series. Upon inspection, the presence of fibronectin or induced flow rates did not make a significant alteration to the impedance spectrums. However, a few other trends were noticed.

For the microfluidic tests under static conditions, fibronectin was injected in the right channel before growth medium was applied to both. After measurements were recorded, the process was repeated but with the fibronectin in the left channel. In almost every case, the resistance decreases while the capacitance increases, whether or not that electrode had fibronectin previously applied or not. This characteristic is verified in the changing impedance spectrums obtained in Figure 29. The dielectric properties of the electrodes changed the longer they were submerged in the growth medium. One
suggestion for this phenomenon is that the electrode surfaces were originally hydrophobic and as the platinum absorbs the nutrients it becomes more hydrophilic. This changes the dielectric properties of the electrode as well as allow for the attachment of cells. The change of electrode properties from hydrophobic to hydrophilic could also explain why it took longer for the cells to spread on the constructed electrode channel than on other surfaces of the same substrate.

Each of the normalized impedance spectra obtained were comparable between devices with only minor differences. These differences can be attributed to slight exposure differences during fabrication and/or different exposure time to growth medium from previous testing. Another consistent trend from each was that the bulk resistance decreased and overall capacitance increased as the area of exposed, measuring metal electrode increased.

As mentioned previously, the time it takes for HUVECs to plate in the trenches of the films takes longer than on larger, smooth surfaces. Either the topography of the films or the hydrophobic nature of the electrodes could account for this behavior. This technique, though, was necessary to culture cells inside microchannels for data collection.

A lack of confluency or cell coverage will alter the impedance spectrum across the measuring electrode. Even though the electrode was partially covered and the impedance spectrum of the electrode changed slightly over time, there was still a noticeable difference between the uncultured and cultured spectrums. However, because the electrode was not completely covered we cannot use the simplistic mathematical model to describe or characterize the behavior of the cells. Complete cell confluency and electrode behavior need to be examined in-depth before ideal results can be obtained.
CHAPTER 7

CONCLUSIONS

A new microfluidic platform was developed and fabricated, which incorporated impedance-sensing technology together with the use of a poly(dimethyl siloxane) microfluidic substrate to create an apparatus capable of applying shear forces up to 90 dyne/cm² to human umbilical vein endothelial cells. Fibronectin did not significantly alter impedance measurements compared to media alone under no flow and flow conditions. The applied fluidic flow rates of 1.0 rpm, 5.0 rpm, and 10.0 rpm did not alter obtained spectra. Frequency spectrums have detected the presence of cultured cells inside the microchannels under no flow conditions. The degree of impedance shift due to the presence of cells is determined on the amount of confluency and coverage of the measuring electrode. Fabricated measured electrodes display changing dielectric properties over time when submerged in EGM-2 cell growth medium under no-flow conditions. Future investigations need to be conducted to characterize the behavior of suspended cells and determine whether this platform will be a versatile tool for investigating endothelial permeability.
CHAPTER 8
RECOMMENDATIONS

The first and foremost recommendation is electrode characterization on the glass-electrode substrate. The traces from the measuring electrodes need to be buried under a nonconductive layer of material so only the measuring electrodes are exposed to the fluid for every experiment and test, and does not depend on the alignment of the PDMS microchannels. Second, the dielectric properties of the electrodes and their interaction with the EGM-2 medium need to be investigated.

The hard and soft baking steps for the fabrication of the SU-8 microfluidic mold needs to be incorporated. The baking steps would allow contact patterning, preventing alterations caused by proximity patterning. A more precise leveling technique or heating method should be considered. These steps improve SU-8 adhesion and integrity, which may prevent the sputtered metal coating from flaking after multiple uses. If heating cannot be implemented, another coating technique for the SU-8 mold besides a sputtered metal coating could be applied.

From this impedance study, reinvestigations and reevaluation of past experiments or biological phenomena is possible. The custom-made PDMS microchannels can be used and adapted to a more complex measuring electrode base. One example is the investigation of PDMS-created branches to replicate biological arterial arrangements.
Investigation of cells at these regions is important, as flow separation promotes vortex formation which is where arteriosclerotic regions have been found to initiate (Davies 1993).

Additionally, deliberate monolayer injury is possible with an application of a low DC voltage across an individual measuring electrode. The voltage will injure/kill the cells, leaving a portion of the monolayer to be healed. The formation of endothelial cells over the electrode could be monitored. This behavior can be investigated at different shear rates, which is believed to regulate endothelial cell wound closure (Gojova 2005). Furthermore, this deliberate injury compromises the barrier function of the surrounding cells (Davies 1988). The ECIS system has used this voltage-injury technique to investigate this healing function (Keese 2003).

Cell patterning is possible with reversible-bonded PDMS microfluidic channels by hydrodynamic focusing of multiple laminar streams (Takayama 1999) or three-dimensional microfluidic stamps (Chiu 2000). The irreversible-bonding technique utilizing a PDMS patterning stamp and a separate PDMS microfluidic substrate can also be used to apply multiple cells (Khademhosseini 2004, Rhee 2005). Using a number of these techniques it is possible to investigate the impedance analysis of different cell cultures simultaneously under the same fluid flow conditions.

PDMS microfluidic substrates could be altered for more in-depth analysis. First, PDMS irreversible bonding techniques can be investigated and applied as to apply greater shear than possible with conformal bonding (over 90 dyne/cm^2). The user would not be as worried about leaking during cell injection procedures. A PDMS microfluidic system could be incorporated into the measuring platform, complete with branches of differing
cross-sectional areas. The method of patterning metals, organic polymers, ceramics, and inorganic crystals on the inner walls of microchannels could be applied for topography investigations as well (Kenis 1999, Kenis 2000, Takayama 2000).

A more complex measuring microelectrode base could be applied to this study. Localized behavior of the endothelial cells barrier function, including focal adhesion points, can be investigated by impedance analysis with an immense array of tiny electrodes. Also, since shear stress varies over very small distances in major arteries an electrode array paired with a PDMS microfluidic substrate mimicking an artery would be beneficial in the investigation of barrier function. From more in depth experimentation and investigation, a versatile and very useful platform could be produced.
REFERENCES


APPENDIX A

MATERIAL SAFETY DATA SHEETS

BAKER BTS 220

MSDS Number: B0231  Effective Date: 12/16/01  Supercedes: 08/24/00

24 Hour Emergency Telephone: 908-659-2151
CHEMTREC: 1-800-424-8888
National Response in Canada
CANUTEC: 613-995-5665
Outside U.S. And Canada
Chemtrec: 783-527-3887

NOTE: CHEMTREC, CANUTEC and National Response Center emergency numbers to be
used only in the event of chemical emergencies involving a spill, leak, fire,
exposure or accident involving chemicals.

All non-emergency questions should be directed to Customer Service (1-800-620-2027) for assistance.

BTS-220

1. Product Identification

Synonyms: Propylene Glycol Monomethyl Ether Acetate; 1-Methoxy-2-Propanol Acetate
CAS No.: 108-65-6
Molecular Weight: 132.16
Chemical Formula: C6H12O3
Product Codes: 6304, 6314, 6333, 6343

2. Composition/Information on Ingredients

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>CAS No.</th>
<th>Percent</th>
<th>Hazardous</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate</td>
<td>108-65-6</td>
<td>100%</td>
<td>Yes</td>
</tr>
</tbody>
</table>

3. Hazards Identification

Emergency Overview

WARNING! FLAMMABLE LIQUID AND VAPOR. MAY FORM EXPLOSIVE
PEROXIDES. MAY CAUSE IRRITATION TO EYES AND RESPIRATORY TRACT.

SAF-T-DATA™ Ratings (Provided here for your convenience)

- Health Rating: 1 - Slight
- Flammability Rating: 2 - Moderate
- Reactivity Rating: 2 - Moderate
- Contact Rating: 1 - Slight
- Lab Protective Equip: GOGGLES & SHIELD; LAB COAT & APRON; VENT HOOD; PROPER GLOVES; CLASS B EXTINGUISHER
- Storage Color Code: Red (Flammable)

Potential Health Effects

Inhalation:
Inhalation of high concentrations may cause irritation to the respiratory tract.

Ingestion:
Not expected to be a health hazard via ingestion.

Skin Contact:
Not expected to be a health hazard from skin exposure.

Eye Contact:
High vapor concentrations may cause eye irritation.

Chronic Exposure:
No information found.

Aggravation of Pre-existing Conditions:
Persons with pre-existing eye disorders or impaired respiratory function may be more susceptible to the effects of this material.

4. First Aid Measures

Inhalation:
Remove to fresh air. Get medical attention for any breathing difficulty.

Ingestion:
Not expected to require first aid measures. If large amounts were swallowed, give water to drink and get medical advice.

Skin Contact:
Not expected to require first aid measures. Wash exposed area with soap and water. Get medical advice if irritation develops.

Eye Contact:
Immediately flush eyes with plenty of water for at least 15 minutes, lifting upper and lower eyelids occasionally. Get medical attention if irritation persists.

5. Fire Fighting Measures
Fire:
Flash point: 42°C (108°F) CC
Autoignition temperature: 354°C (669°F)
Flammable limits in air % by volume:
LFL: 1.5; UFL: 7.0
Flammable Liquid and Vapor! Contact with strong oxidizers may cause fire.

Explosion:
Sealed containers may rupture when heated. Above the flash point, explosive vapor-air mixtures may be formed. Vapors can flow along surfaces to distant ignition source and flash back. May form unstable peroxides which may explode spontaneously or when heated.

Fire Extinguishing Media:
Water spray, dry chemical, alcohol foam, or carbon dioxide. Water spray may be used to keep fire exposed containers cool.

Special Information:
In the event of a fire, wear full protective clothing and NIOSH-approved self-contained breathing apparatus with full facepiece operated in the pressure demand or other positive pressure mode.

6. Accidental Release Measures

Ventilate area of leak or spill. Remove all sources of ignition. Wear appropriate personal protective equipment as specified in Section 8. Isolate hazard area. Keep unnecessary and unprotected personnel from entering. Contain and recover liquid when possible. Use non-sparking tools and equipment. Collect liquid in an appropriate container or absorb with an inert material (e.g., vermiculite, dry sand, earth), and place in a chemical waste container. Do not use combustible materials, such as saw dust. Do not flush to sewer! If a leak or spill has not ignited, use water spray to disperse the vapors, to protect personnel attempting to stop leak, and to flush spills away from exposures.

7. Handling and Storage

Protect against physical damage. Store in a cool, dry well-ventilated location, away from any area where the fire hazard may be acute. Outside or detached storage is preferred. Separate from incompatibles. Containers should be bonded and grounded for transfers to avoid static sparks. Storage and use areas should be No Smoking areas. Use non-sparking type tools and equipment, including explosion proof ventilation. After opening, purge container with nitrogen before re-closing. Periodically test for peroxide formation on long-term storage. Do not distill to near dryness. If peroxide formation is suspected, do not open or move container. Addition of water or appropriate reducing materials will lessen peroxide formation. Containers of this material may be hazardous when empty since they retain product residues (vapors, liquid); observe all warnings and precautions listed for the product. Do Not attempt to clean empty containers since residue is difficult to remove. Do not pressure, cut, weld, braze, solder, drill, grind or expose such containers to heat, sparks, flame, static electricity or other sources of ignition; they may explode and cause injury or death.
8. Exposure Controls/Personal Protection

Airborne Exposure Limits:
For Propylene Glycol Monomethyl Ether Acetate:
- ACGIH Workplace Environmental Exposure Level (WEEL) -
  100 ppm (TWA).
Ventilation System:
A system of local and/or general exhaust is recommended to keep employee exposures
below the Airborne Exposure Limits. Local exhaust ventilation is generally preferred
because it can control the emissions of the contaminant at its source, preventing dispersion
of it into the general work area. Please refer to the ACGIH document, Industrial
Personal Respirators (NIOSH Approved):
If the exposure limit is exceeded, wear a supplied air, full-facepiece respirator, airlined
hood, or full-facepiece self-contained breathing apparatus.
Skin Protection:
Wear protective gloves and clean body-covering clothing.
Eye Protection:
Use chemical safety goggles and/or a full face shield where splashing is possible. Maintain
eye wash fountain and quick-drench facilities in work area.

9. Physical and Chemical Properties

Appearance:
Clear, colorless liquid.
Odor:
Sweet ether-like odor.
Solubility:
Appreciable in water.
Specific Gravity:
0.969 @ 20C (68°F)
pH:
No information found.
% Volatiles by volume @ 21C (70°F):
No information found.
Boiling Point:
150C (302F)
Melting Point:
No information found.
Vapor Density (Air=1):
4.6
Vapor Pressure (mm Hg):
3.7 @ 20C (68°F)
Evaporation Rate (BuAc=1):
0.39
10. Stability and Reactivity

Stability:
Stable under ordinary conditions of use and storage. Unstable at elevated temperatures. May form explosive peroxides.

Hazardous Decomposition Products:
Carbon dioxide and carbon monoxide may form when heated to decomposition.

Hazardous Polymerization:
Will not occur.

Incompatibilities:
Oxidizers can cause vigorous reaction.

Conditions to Avoid:
Heat, flames, ignition sources and incompatibles.

11. Toxicological Information

For Propylene Glycol Monomethyl Ether Acetate: LD50 oral rat 8532 mg/kg; LD50 skin rabbit > 5 gm/kg.

--- \Cancer List --

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>LD To Carcinogen</th>
<th>Known</th>
<th>Anticipated</th>
<th>IARC Category</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate (108-65-6)</td>
<td>No</td>
<td>No</td>
<td>None</td>
<td></td>
</tr>
</tbody>
</table>

12. Ecological Information

Environmental Fate:
No adverse ecological effects are expected.

Environmental Toxicity:
This material is not expected to be toxic to aquatic life. The LC50/96-hour values for fish are over 100 mg/l. The EC50/48-hour values for daphnia are over 100 mg/l.

13. Disposal Considerations

Whatever cannot be saved for recovery or recycling should be handled as hazardous waste and sent to a RCRA approved waste facility. Processing, use or contamination of this product may change the waste management options. State and local disposal regulations may differ from federal disposal regulations. Dispose of container and unused contents in
14. Transport Information

Domestic (Land, D.O.T.)

Proper Shipping Name: FLAMMABLE LIQUID, N.O.S. (PROPYLENE GLYCOL MONOMETHYL ETHER ACETATE)
Hazard Class: 3
UN/NA: UN1993
Packing Group: III
Information reported for product/size: 425 LB

International (Water, I.M.O.)

Proper Shipping Name: FLAMMABLE LIQUID, N.O.S. (PROPYLENE GLYCOL MONOMETHYL ETHER ACETATE)
Hazard Class: 3
UN/NA: UN1993
Packing Group: III
Information reported for product/size: 425 LB

15. Regulatory Information

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Chemical Inventory Status - Part 1</th>
<th>TSCA</th>
<th>EC</th>
<th>Japan</th>
<th>Australia</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate (108-65-6)</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Chemical Inventory Status - Part 2 - Canada</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate (108-65-6)</td>
<td>Yes</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Federal, State &amp; International Regulations - Part 1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate (108-65-6)</td>
<td>-SARA 302- No</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Federal, State &amp; International Regulations - Part 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Propylene glycol monomethyl ether acetate (108-65-6)</td>
<td>CERCLA 261.53</td>
</tr>
</tbody>
</table>

accordance with federal, state and local requirements.
Australian Hazchem Code: None allocated.
Poison Schedule: None allocated.
WHMIS:
This MSDS has been prepared according to the hazard criteria of the Controlled Products Regulations (CPR) and the MSDS contains all of the information required by the CPR.

16. Other Information

NFPA Ratings: Health: 0 Flammability: 2 Reactivity: 0
Label Hazard Warning:
WARNING! FLAMMABLE LIQUID AND VAPOR. MAY FORM EXPLOSIVE PEROXIDES. MAY CAUSE IRRITATION TO EYES AND RESPIRATORY TRACT.
Label Precautions:
Keep away from heat, sparks and flame.
Keep container closed.
Use only with adequate ventilation.
Avoid contact with eyes, skin and clothing.
Wash thoroughly after handling.
Avoid breathing vapor or mist.
Label First Aid:
If inhaled, remove to fresh air. Get medical attention for any breathing difficulty. In case of eye contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical attention if irritation develops or persists.
Product Use:
Laboratory Reagent.
Revision Information:
MSDS Section(s) changed since last revision of document include: 3.
Disclaimer:
******************************************************************************

Mallinckrodt Baker, Inc. provides the information contained herein in good faith but makes no representation as to its comprehensiveness or accuracy. This document is intended only as a guide to the appropriate precautionary handling of the material by a properly trained person using this product. Individuals receiving the information must exercise their independent judgment in determining its appropriateness for a particular purpose. MALLINCKRODT BAKER, INC. MAKES NO REPRESENTATIONS OR WARRANTIES, EITHER EXPRESS OR IMPLIED, INCLUDING WITHOUT LIMITATION ANY WARRANTIES OF MERCHANTABILITY, FITNESS FOR A PARTICULAR PURPOSE WITH RESPECT TO THE INFORMATION SET FORTH HEREIN OR THE PRODUCT TO WHICH THE INFORMATION REFERS. ACCORDINGLY, MALLINCKRODT
BAKER, INC. WILL NOT BE RESPONSIBLE FOR DAMAGES RESULTING
FROM USE OF OR RELIANCE UPON THIS INFORMATION.

Prepared by: Environmental Health & Safety
Phone Number: (314) 654-1600 (U.S.A.)
Material Safety Data Sheet

1. CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME: BOE
OTHER/GENERIC NAMES: 6:1 BOE
PRODUCT USE: Semiconductor Manufacturing

MANUFACTURER: Honeywell/GEM3
101 Columbia Road
Morristown, New Jersey 07962-1053

FOR MORE INFORMATION CALL: (Monday-Friday, 8:00am-5:00pm)
1-800-278-9998

IN CASE OF EMERGENCY CALL:
(24 Hours/Day, 7 Days/Week)
1-800-707-4555 or Chemtrec 1-800-827-4723
International: 1-701-527-3887

2. COMPOSITION/INFORMATION ON INGREDIENTS

<table>
<thead>
<tr>
<th>INGREDIENT NAME</th>
<th>CAS NUMBER</th>
<th>WEIGHT %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ammonium Fluoride</td>
<td>12125-01-8</td>
<td>&lt; 36%</td>
</tr>
<tr>
<td>Hydrofluoric Acid</td>
<td>7664-39-3</td>
<td>&lt; 10%</td>
</tr>
</tbody>
</table>

Trace impurities and additional material names not listed above may also appear in Section 15 toward the end of the MSDS. These materials may be listed for local 'Right-To-Know' compliance and for other reasons.

3. HAZARDS IDENTIFICATION

EMERGENCY OVERVIEW:
Extremely hazardous liquid and vapor. Causes severe burns that may not be immediately painful or visible. May be fatal if inhaled or swallowed. May cause damage to bones.

POTENTIAL HEALTH HAZARDS

SKIN: Liquid causes severe irritation and burns of eyes and skin
EYES: Both liquid and vapor can cause irritation or corneal burns.
INHALATION: May cause irritation to mucous membranes. Inhalation causes acute systemic poisoning which must receive immediate medical attention.
INGESTION: Ingestion causes vomiting and severe burns of mouth and stomach.

MSDS Number: 6:1 BOE
Current Issue Date: October 10, 2003
MATERIAL SAFETY DATA SHEET

DELAYED EFFECTS: The effects of contact with dilute solutions of hydrofluoric acid or its vapors may be delayed. The potential delay in clinical signs or symptoms for dilute solutions is given below. Symptoms might include pain, redness and possible tissue destruction.

<table>
<thead>
<tr>
<th>HF Concentration</th>
<th>Delay In Symptoms</th>
</tr>
</thead>
<tbody>
<tr>
<td>95%</td>
<td>Immediately</td>
</tr>
<tr>
<td>20%-50%</td>
<td>1-8 Hours</td>
</tr>
<tr>
<td>0%-20%</td>
<td>Up to 24 hours</td>
</tr>
</tbody>
</table>

Can also cause bone and joint changes in humans (Fluorosis).

Ingredients found on one of the OSHA designated carcinogen lists are listed below.

INGREDIENT NAME | NTP STATUS | IARC STATUS | OSHA LIST
--- | --- | --- | ---
No ingredients listed in this section.

4. FIRST AID MEASURES

SKIN: Do not scrub. Remove the victim from the contaminated area and immediately wash the burned area with plenty of water for a minimum of 15 minutes. Limit washing to 5 minutes if treatment specific for HF exposure is available. Remove all contaminated clothing while washing continuously. After thorough washing for at least 5 minutes, the burned area should be immersed in a solution of 0.13% iced aqueous Zephiran® Chloride until pain is relieved. As an alternate first aid treatment, 2.5% calcium gluconate gel may be continuously massaged into the burn area until the pain is relieved. For larger burns or burns treated with calcium gluconate gel (in which pain is present longer than 30 minutes), a physician should inject 5% aqueous calcium gluconate beneath, around and in the burned area. Use of local anesthetics is not recommended, as a reduction in pain is an indicator of effectiveness of treatment.

EYES: Irrigate eyes for at least 15 minutes with copious quantities of water, keeping eyelids apart and away from eyelashes during irrigation. Get competent medical attention immediately, preferably an eye specialist. If a physician is not immediately available, apply one or two drops of 0.5% tetracaine hydrochloride solution, or other aqueous topical cocaine-based anesthetic and continue irrigation. DO NOT use the solution described for skin treatment. Use no oils or greases unless instructed to do so by a physician. Irrigate with 1% calcium gluconate in normal saline for 1 to 2 hours to prevent or lessen corneal damage.

INHALATION: Move to fresh air. Keep the victim lying down, quiet and warm. Get competent medical attention immediately. If breathing has stopped, start artificial respiration at once. An authorized person should administer oxygen to a victim who is having difficulty breathing, until the victim is able to breathe easily by himself. Calcium gluconate, 2.5% in normal saline may be given by nebulizer with oxygen. DO NOT give stimulants unless instructed to do so by a physician. Victim should be examined by a physician and held under observation for at least 24 hours.

INGESTION: Drink large amounts of water to dilute. DO NOT induce vomiting. Several glasses of milk or several ounces of milk of magnesium may be given for their soothing effect. Take the victim to a doctor.

ADVICE TO PHYSICIAN:

Treat symptomatically. For Hydrofluoric Acid:

For burns of large skin areas (greater than 25 square inches), for ingestion and for significant inhalation exposure, severe systemic effects may occur. Monitor and correct for hypocalcemia, cardiac arrhythmias, hypomagnesemia and hyperkalemia. In some cases hemodialysis may be indicated. For certain burns, especially of the digits, use of intra-articular calcium gluconate may be indicated. For inhalation exposures, treat as chemical pneumonia. Monitor for hypocalcemia. 2.5% calcium gluconate in normal saline by nebulizer or by IPPB with 100% oxygen may decrease pulmonary damage. Bronchodilators may also be administered.

A booklet titled “Recommended Medical Treatment for Hydrofluoric Acid Exposure” is available from the Honeywell HF website: http://www.HFacid.com.

MSDS Number: 6190E
Current Issue Date: October 10, 2003
Page 2 of 7
5. FIRE FIGHTING MEASURES

FLAMMABLE PROPERTIES:

FLASH POINT: Not flammable
FLASH POINT METHOD: Closed cup
AUTOIGNITION TEMPERATURE: N/A
UPPER FLAME LIMIT (volume % in air): N/A
LOWER FLAME LIMIT (volume % in air): N/A
FLAME PROPAGATION RATE (solids): N/A
OSHA FLAMMABILITY CLASS: N/A

EXTINGUISHING MEDIA:

Use water or suitable agent for fires adjacent to non-leaking tanks or containers of HF. This substance is not combustible. Use water fog or carbon dioxide. DO NOT use solid water streams near ruptured tanks or spills of HF. Acid reacts with water and can splatter acid onto personnel.

UNUSUAL FIRE AND EXPLOSION HAZARDS:

This product is not flammable. Emits toxic fumes.

SPECIAL FIRE FIGHTING PRECAUTIONS/INSTRUCTIONS:

Use spray to cool exposed containers. Block sewers in the path of spreading spill to prevent entry. Remove containers of this material to cool areas. Prevent the boiling of this formulation. Wear self-contained breathing apparatus approved by NIOSH and full chemical protective clothing. Use water spray to keep containers cool.

6. ACCIDENTAL RELEASE MEASURES

IN CASE OF SPILL OR OTHER RELEASE: (Always wear recommended personal protective equipment.)

Good ventilation is necessary. Discharge will ordinarily be a vapor or a liquid that gives off fumes of HF gas. Those treating spills or repairing leaks must use full protective equipment. Take actions to minimize environmental impact. Try to contain spillage and avoid drainage to areas which cannot be treated. Rapid dilution of the spill with water will reduce the amount of fumes given off. Carefully neutralize the dilute liquid with lime slurry, soda ash, limestone, caustic soda or other alkaline material. (See sections 10 and 13 for more information.)

Spills and releases may have to be reported to Federal and/or local authorities. See Section 15 regarding reporting requirements.

7. HANDLING AND STORAGE

NORMAL HANDLING: (Always wear recommended personal protective equipment.)

DO NOT breathe vapor or mist. Use only adequate ventilation. Avoid all contact with skin, eyes and clothing. Even dilute solutions. DO NOT add water to acid.

STORAGE RECOMMENDATIONS:

Store in approved containers only. Store in cool, well-ventilated area. Flammable hydrogen gas can be generated in contact with metals. Diking of storage tanks is recommended. Store above 70°F.
MATERIAL SAFETY DATA SHEET

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

ENGINEERING CONTROLS:
Material must be handled or transferred in an approved fume hood or with equivalent ventilation sufficient to reduce vapor and acid mists below permissible TLV levels. Packaging and unloading areas and open processing equipment may require mechanical exhaust systems.

PERSONAL PROTECTIVE EQUIPMENT

SKIN PROTECTION:
For routine product use, wear acid-resistant jacket, trousers, boots and gauntlet gloves. For increased protection, use air-supplied totally encapsulating HF resistant protective suit.

EYE PROTECTION:
As a minimum, wear hardhat, chemical safety goggles (plastic lenses), and full-face plastic shield. For increased protection, use air-supplied hydrofluoric acid resistant hood.

RESPIRATORY PROTECTION:
Where required, use a respirator approved by NIOSH for HF gas or mists, as applicable. Some exposures may require a NIOSH-approved, self-contained breathing apparatus or air-supplied respirator.

ADDITIONAL RECOMMENDATIONS:
Eyewash and quick-drench shower facilities, protected from freezing, should be available where HF is stored or handled.

EXPOSURE GUIDELINES

<table>
<thead>
<tr>
<th>INGREDIENT NAME</th>
<th>ACGIH TLV</th>
<th>OSHA PEL</th>
<th>OTHER LIMIT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hydrofluoric Acid</td>
<td>3 ppm C</td>
<td>3 ppm TWA</td>
<td>3 mg/loutine pre-shift</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>10 mg/loutine post-shift**</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ammonium Fluoride</td>
<td>2.5 mg/m³</td>
<td>None</td>
<td></td>
</tr>
</tbody>
</table>

OSHA STEL 6 ppm (15min.)
IDLH 30 ppm

AHA Emergency Response Planning Guideline

<table>
<thead>
<tr>
<th>ERPG-1</th>
<th>ERPG-2</th>
<th>ERPG-3</th>
</tr>
</thead>
<tbody>
<tr>
<td>2 ppm (60min)</td>
<td>20 ppm (60min)</td>
<td>50 ppm (60min)</td>
</tr>
<tr>
<td>2 ppm (10min)</td>
<td>20 ppm (10min)</td>
<td>50 ppm (10min)</td>
</tr>
</tbody>
</table>

* = Limit established by Honeywell International, Inc.
** = Workplace Environmental Exposure Level (AIHA).
*** = Biological Exposure Index (ACGIH).

OTHER EXPOSURE LIMITS FOR POTENTIAL DECOMPOSITION PRODUCTS: None

9. PHYSICAL AND CHEMICAL PROPERTIES

APPEARANCE: Colorless liquid
PHYSICAL STATE: Liquid
MOLECULAR WEIGHT: Mixture
### MATERIAL SAFETY DATA SHEET

**CHEMICAL FORMULA:** NH₄F + HF (Contains a diminutive quantity of surfactant)

**ODOR:** Slight pungent odor

**SPECIFIC GRAVITY (water = 1.0):** 1.114

**SOLUBILITY IN WATER (weight %):** Miscible

**pH:** ~4

**BOILING POINT:** Unknown

**MEETING POINT:** N/A

**VAPOR PRESSURE:** N/A

**VAPOR DENSITY (air = 1.0):** N/A

**EVAPORATION RATE:** N/A

**% VOLATILES:** 99+%  

**FLASH POINT:** N/A  

(Flash point method and additional flammability data are found in Section 5.)

### 10. STABILITY AND REACTIVITY

**NORMALLY STABLE? (CONDITIONS TO AVOID):** Yes.

**INCOMPATIBILITIES:** Chemical is very reactive. Store away from other chemicals

**HAZARDOUS DECOMPOSITION PRODUCTS:** Releases highly corrosive forms of fluorides when heated; oxides of C

**HAZARDOUS POLYMERIZATION:** Data not applicable

### 11. TOXICOLOGICAL INFORMATION

**IMMEDIATE (ACUTE) EFFECTS:**

- Hydrofluoric Acid:
  - inh-hmn LCL: 50 ppm/30M  
  - inh-rat LC50: 1300 ppm/60M  
  - inh-man TLC: 100 mg/m³/1M

- Ammonium Fluoride:
  - oral-rat LD50: 200 mg/kg

**DELAYED (SUBCHRONIC AND CHRONIC) EFFECTS:**

- Prolonged exposure can cause bone and joint changes in humans. (Fluorosis- increased bone density and mottling of teeth)

**OTHER DATA:** Tests on laboratory animals indicate HF in concentrate form may produce adverse mutagenic and reproductive effects. Cited in Registry of Toxic Effects of Substances (RTÉS).

### 12. ECOLOGICAL INFORMATION

N/A

---

**MSDS Number:** 61 BOE  
**Current Issue Date:** October 10, 2003
MATERIAL SAFETY DATA SHEET

STATE RIGHT-TO-KNOW

In addition to the ingredients found in Section 2, the following are listed for state right-to-know purposes.

<table>
<thead>
<tr>
<th>INGREDIENT NAME</th>
<th>WEIGHT %</th>
<th>COMMENT</th>
</tr>
</thead>
<tbody>
<tr>
<td>No ingredients listed in this section.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

ADDITIONAL REGULATORY INFORMATION: None

WHMIS CLASSIFICATION (CANADA): N/A

FOREIGN INVENTORY STATUS: N/A

16. OTHER INFORMATION

CURRENT ISSUE DATE: October 9, 2003
PREVIOUS ISSUE DATE: July 17, 2000

CHANGES TO MSDS FROM PREVIOUS ISSUE DATE ARE DUE TO THE FOLLOWING:
Converted to ANSI 16-section format.

OTHER INFORMATION: Honeywell MAKES NO WARRANTY, EXPRESSED OR IMPLIED, CONCERNING THIS MATERIAL OR THE USE OF THIS PRODUCT OTHER THAN INDICATED ON THE LABEL. BUYER ASSUMES ALL RISK OF USE AND/OR HANDLING OF THIS MATERIAL WHEN SUCH USE AND/OR HANDLING IS CONTRARY TO LABEL INSTRUCTIONS

NFPA Hazard Ratings:
Heath: 4
Flammability: 0
Reactivity: 1
Special Hazards:
Nichrome Etchant TFN

MATERIAL SAFETY DATA SHEET

IDENTITY (As Used on Label and List)
CHROMIUM ETCH 1020
CHROME ETCH 8001N
NICHROME ETCH TFN

Section I
Manufacturers Name
TRANSENE COMPANY INC.
Address
10 ELECTRONICS AVENUE
DANVERS MA 01923

Emergency Telephone Number
CHEMTREC 24HR# 1-800-424-9300

Telephone Number for Information
(978)777-7860 Fax: (978)739-5640 www.transene.com

Date Prepared
REVISED 04-04

Section II – Hazardous Ingredients / Identity Information

<table>
<thead>
<tr>
<th>Hazardous Components</th>
<th>Specifice Chemical Identity Common Name(s)</th>
<th>OSHA PEL</th>
<th>ACGIH TLV</th>
<th>Other Limits Recommended</th>
<th>% (optional)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CERIC AMMONIUM NITRATE</td>
<td>CAS# 16774-21-3 oral rat LD50</td>
<td>244 ppm</td>
<td>10-20</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NITRIC ACID (HNO3)</td>
<td>CAS# 7697-37-2</td>
<td>2 ppm</td>
<td>5-6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NON-HAZARDOUS AQUEOUS SOLUTION</td>
<td></td>
<td>* AS NITRATE</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* THIS PRODUCT CONTAINS NO MORE THAN 6% BY WT. NITRIC ACID, WHICH IS CONSIDERED TO BE A TOXIC CHEMICAL UNDER TITLE III. EMISSIONS DATA MUST BE SUBMITTED ANNUALLY TO THE EPA FOR COMPANIES WHO MANUFACTURE, PROCESS OR OTHERWISE USE THIS PRODUCT IN EXCESS OF CERTAIN QUANTITIES ESTABLISHED BY THE EPA. THIS INFORMATION MUST BE INCLUDED IN ALL MSDS THAT ARE Copied AND DISTRIBUTED FOR THIS MATERIAL.

HMIS Ratings
Health : 3 Reactivity : 0 Flammability : 0

Section III - Physical/Chemical Characteristics

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boiling Point (100°C)</td>
<td>212°F</td>
</tr>
<tr>
<td>Vapor Pressure (mm Hg)</td>
<td>N/A</td>
</tr>
<tr>
<td>Vapor Density (AIR - 1)</td>
<td>N/A</td>
</tr>
<tr>
<td>Specific Gravity (H2O = 1)</td>
<td>1.16</td>
</tr>
<tr>
<td>Melting Point % VOLATILES</td>
<td>85%</td>
</tr>
<tr>
<td>Evaporation Rate (Butyl Acetate = 1)</td>
<td>Similar to water</td>
</tr>
</tbody>
</table>

Solubility in Water: COMPLETE

Appearance and Odor: ORANGE LIQUID WITH PRACTICALLY NO ODOR.

Section IV – Fire and Explosion Hazard Data

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flash Point (Method Used)</td>
<td>NON-FIammable</td>
</tr>
<tr>
<td>Flammable Limits</td>
<td>LEL</td>
</tr>
<tr>
<td>Extinguishing Media: WATER SPRAY IN CASE OF FIRE CAUSED BY SPILLAGE. DO NOT USE ORGANIC FIRE EXTINGUISHING AGENTS.</td>
<td></td>
</tr>
<tr>
<td>Special Fire Fighting Procedures: WEAR NIOSH APPROVED SELF-CONTAINED BREATHING APPARATUS AND APPROPRIATE PROTECTIVE APPAREL</td>
<td></td>
</tr>
<tr>
<td>Unusual Fire and Explosion Hazards: SPILLAGE MAY CAUSE FIRE OR LIBERATE DANGEROUS GAS. INCREASES FLAMMABILITY OF COMBUSTIBLE ORGANICS AND DEADLY OXIDIZABLE MATERIALS. MAY CAUSE IONIZATION.</td>
<td></td>
</tr>
</tbody>
</table>

Section V – Reactivity Data

<table>
<thead>
<tr>
<th>Stability</th>
<th>Conditions to Avoid</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unstable</td>
<td>ORGANIC MATERIALS, STRONG OXIDANTS, HEAT</td>
</tr>
<tr>
<td>Stable</td>
<td>X</td>
</tr>
</tbody>
</table>

PAGE 1
Incompatibility (Materials to Avoid): METALLIC POWDERS, CARBIDES, HYDROGEN SULFIDE AND EASILY OXIDIZED MATERIALS

Hazardous Decomposition or Byproducts:
OXIDES OF NITROGEN, STRONG BASES, AND METALS

Hazardous Polymerization: May Occur
Conditions to Avoid
Will Not Occur X EXCESSIVE HEAT

Section VI - Health Hazard Data
Route(s) of Entry: Inhalation? Skin? Ingestion?
Inhalation? YES YES YES

Health Hazards (Acute and Chronic):
MAY CAUSE IRRITATION TO LUNGS AND RESPIRATORY SYSTEM.
MAY CAUSE IRRITATION AND BURNS. VAPOR OR SPRAYS CAN BE HAZARDOUS TO THE LUNGS AND RESPIRATORY SYSTEM. MATERIAL IS CORROSIVE TO EYES, SKIN, AND MUCOUS MEMBRANES.

Carcinogenicity: NTP? IARC Monographs? OSHA Regulated?
NO NO YES

Signs and Symptoms of Exposure:
MAY CAUSE IRRITATION AND BURNS.
MEDICAL CONDITIONS GENERALLY AGGRAVATED BY EXPOSURE:
MAY CAUSE IRRITATION AND BURNS. VAPOR OR SPRAYS CAN BE HAZARDOUS TO THE LUNGS AND RESPIRATORY SYSTEM. MATERIAL IS CORROSIVE TO EYES, SKIN, AND MUCOUS MEMBRANES.

Emergency and First Aid Procedures:
EYE CONTACT: IMMEDIATELY GET MEDICAL ATTENTION.
SKIN: REMOVE CONTAMINATED CLOTHING AND SHOES. WASH EXPOSED AREA WITH SOAP AND WATER. GET MEDICAL ATTENTION.
INHALATION: REMOVE TO FRESH AIR. IF BREATHING HAS STOPPED, GIVE ARTIFICIAL RESPIRATION.
INGESTION: DO NOT INDUCE VOMITING. GET MEDICAL ATTENTION.

Section VII - Precautions for Safe Handling and Use
Steps to Be Taken in Case Material Is Released or Spilled:
ISOLATE OR ENCLOSE THE AREA OF SPILL OR LEAK. NEUTRALIZE WITH CALCIUM CARBONATE. WASH WITH EXCESS WATER. PICK UP WITH ABSORBENT MATERIAL (SAND OR VERMICULITE) AND DISPOSE IN A RCRA APPROVED WASTE FACILITY.

Waste Disposal Method: CLEAN UP PERSONNEL SHOULD WEAR PROTECTIVE CLOTHING AND RESPIRATORY EQUIPMENT SUITABLE FOR TOXIC FUMES. DISPOSE OF IN ACCORDANCE WITH LOCAL, STATE OR FEDERAL ENVIRONMENTAL REGULATIONS.

Precautions to Be Taken in Handling and Storing: STORE IN WELL-VENTILATED AREA. ROOM TEMPERATURE STORAGE-SEALED BOTTLES.

Other Precautions: STORE AWAY FROM INCOMPATIBLE MATERIALS. PROTECT FROM PHYSICAL DAMAGE AND DIRECT SUNLIGHT.

Section VIII - Control Measures
Respiratory Protection (Specify Type):
IF TLV IS EXCEEDED WEAR A NIOSH APPROVED SELF-CONTAINED BREATHING APPARATUS.

Ventilation: Local Exhaust: MAINTAIN ADEQUATE VENTILATION
Mechanical (general): EXPOSURE TO LIMITS BELOW AIRBORNE EXPOSURE LIMITS

Special: EYE BATH AND SHOWER FACILITIES WORK AREA.

Protective Gloves: RUBBER

Eye Protection: CHEMICAL SAFETY GOGGLES OR FACE SHIELD.

Other Protective Clothing or Equipment:
PROTECTIVE CLOTHING, RUBBER APRON / USE APPROPRIATE INDUSTRIAL PRACTICES FOR HANDLING ACIDS.

Work / Hygienic Practices: USE APPROPRIATE INDUSTRIAL PRACTICES FOR HANDLING ACIDS.
## ADDENDUM TO MATERIAL SAFETY DATA SHEET

### REGULATORY STATUS

This addendum must not be detached from the MSDS. Identification of SARA substances must include a copy of this addendum.

### HAZARD CATEGORIES FOR SARA

<table>
<thead>
<tr>
<th>Product or Components Of Products</th>
<th>SARA EHS Sect. 302</th>
<th>SARA Section 313 Chemicals</th>
<th>CERCLA Sec. 103</th>
<th>RCRA</th>
</tr>
</thead>
<tbody>
<tr>
<td>NITRIC ACID (7697-37-2)</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>No</td>
</tr>
</tbody>
</table>

**Applicable Products:**
- Reagent Semiconductor Etchants (RSE 100 - RSE 1155)
- Nitric Acid 70%, Wright Etchant, TFB, TFD, TFC, TPN / 1020, Chromium Mask Etchant

SARA Section 302 EHS RQ: Reportable Quantity of Extremely Hazardous Substance, listed at 40 CFR 355.

SARA Section 302 EHS TPC: Threshold Planning Quantity of Extremely Hazardous Substance. An asterisk (*) following a Threshold Planning Quantity signifies that if the material is a solid and has a particle size equal to or larger than 100 micrometers, the Threshold Planning Quantity is 10,000 lbs.

SARA Section 313 Chemical Toxic Substances subject to annual release reporting requirements listed at 40 CFR 372.65.

CERCLA Sec. 103: Comprehensive Environmental Response, Compensation, and Liability Act (Superfund) Releases to air, land or water of these hazardous substances which exceed the Reportable Quantity (RQ) must be reported to the National Response Center (800-424-8802).


---

Effective Date: 02-17-87 Supersedes 04-30-86

---

**NITRIC ACID**
Material Safety Data Sheet

1. PRODUCT AND COMPANY IDENTIFICATION

MICROPOSIT(TM) MF(TM) -319 Developer

Supplier: Pohm and Haas Electronic Materials LLC
455 Forest Street
Marlborough, MA 01752 United States of America

Revision date: 01/01/2004

For non-emergency information contact: 508-481-7950

Emergency telephone number
Chemtrec 800-424-9300
Rohm and Haas Emergency 215-592-3000

2. COMPOSITION/INFORMATION ON INGREDIENTS

<table>
<thead>
<tr>
<th>Component</th>
<th>CAS-No.</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>7732-18-5</td>
<td>&gt; 95.0 %</td>
</tr>
<tr>
<td>Surfactant</td>
<td></td>
<td>&lt; 1.0 %</td>
</tr>
<tr>
<td>Tetramethylammonium hydroxide</td>
<td>75-59-2</td>
<td>2.2 %</td>
</tr>
</tbody>
</table>

3. HAZARDS IDENTIFICATION

Emergency Overview

Appearance
Form: liquid
Colour: colourless
Odour: amines

Hazard Summary

WARNING!
Alkaline liquid and vapor. Causes skin, eye, and respiratory tract irritation. Onset of symptoms may be delayed. Prolonged, repeated contact, inhalation, ingestion, or absorption through the skin, may cause toxic effects to internal organ systems (liver, kidney, central nervous system).

Potential Health Effects
Primary Routes of Entry: Inhalation, ingestion, eye and skin contact, absorption.

Eyes: May cause pain, transient irritation and superficial corneal effects.

Skin: Material may cause irritation. Prolonged or repeated exposure may have the following effects:
- central nervous system depression
- drowsiness
- desquamation of skin leading to irritation and dermatitis

Ingestion: Swallowing may have the following effects:
- irritation of mouth, throat and digestive tract
- Repeated doses may have the following effects:
- central nervous system depression
- drowsiness

Inhalation: Inhalation may have the following effects:
- irritation of nose, throat and respiratory tract
- Higher concentrations may have the following effects:
- systemic effects similar to those resulting from ingestion

Target Organs: Eye
Respiratory System
Skin
nervous system

Carcinogenicity
Not considered carcinogenic by NTP, IARC, and OSHA

4. FIRST AID MEASURES

Inhalation: Remove from exposure. If there is difficulty in breathing, give oxygen. Seek medical attention if symptoms persist.

Skin contact: Wash skin with water. Continue washing for at least 15 minutes. Obtain medical attention if blistering occurs or redness persists.

Eye contact: Immediately flush the eye with plenty of water for at least 15 minutes, holding the eye open. Obtain medical attention if soreness or redness persists.

Ingestion: Wash out mouth with water. Have victim drink 1-3 glasses of water to dilute stomach contents. Immediate medical attention is required. Never administer anything by mouth if a victim is losing consciousness, is unconscious or is convulsing.

Notes to physician
Treat symptomatically.

5. FIRE-FIGHTING MEASURES

Flash point: Nonflammable

Suitable extinguishing media: Not readily combustible. Select extinguishing agent appropriate to other materials involved.
Specific hazards during fire fighting: No specific measures necessary.

Special protective equipment for fire-fighters: Wear full protective clothing and self-contained breathing apparatus.

Further Information: This product may give rise to hazardous vapors in a fire.

6. ACCIDENTAL RELEASE MEASURES

Personal precautions
Wear suitable protective clothing.

Environmental precautions
Prevent the material from entering drains or water courses.
Do not discharge directly to a water source.
Advise Authorities if spillage has entered watercourse or sewer or has contaminated soil or vegetation.

Methods for cleaning up
Cover with absorbent or contain. Collect and dispose.

7. HANDLING AND STORAGE

Handling
Use only in well-ventilated areas. Avoid breathing vapor. Avoid contact with eyes, skin and clothing. Keep container tightly closed.

Further Information on storage conditions: No special precautions necessary.

Storage
Storage conditions: Store in original container. Storage area should be: cool, dry, well ventilated out of direct sunlight away from incompatible materials

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Exposure limit(s)
Exposure limits are listed below, if they exist.

Eye protection: goggles

Hand protection: Butyl rubber gloves. Other chemical resistant gloves may be recommended by your safety professional.

Skin and body protection: Normal work wear.
Respiratory protection: Respiratory protection if there is a risk of exposure to high vapor concentrations. The specific respirator selected must be based on the airborne concentration found in the workplace and must not exceed the working limits of the respirator.

Engineering measures: Engineering methods to prevent or control exposure are preferred. Methods include process or personnel enclosure, mechanical ventilation (local exhaust), and control of process conditions.

9. PHYSICAL AND CHEMICAL PROPERTIES

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Appearance</td>
<td>liquid</td>
</tr>
<tr>
<td>Colour</td>
<td>colourless</td>
</tr>
<tr>
<td>Odour</td>
<td>amines</td>
</tr>
<tr>
<td>pH</td>
<td>13</td>
</tr>
<tr>
<td>Boiling point/range</td>
<td>100 °C (212 °F)</td>
</tr>
<tr>
<td>Flash point</td>
<td>Nonflammable</td>
</tr>
<tr>
<td>Component</td>
<td>Tetramethylammonium hydroxide</td>
</tr>
<tr>
<td>Vapour pressure</td>
<td>17.5 mmHg at 20 °C</td>
</tr>
<tr>
<td>Relative vapour density</td>
<td>no data available</td>
</tr>
<tr>
<td>Water solubility</td>
<td>completely soluble</td>
</tr>
<tr>
<td>Relative density</td>
<td>1.00</td>
</tr>
<tr>
<td>Evaporation rate</td>
<td>Slower than ether</td>
</tr>
<tr>
<td>VOC's</td>
<td>not applicable</td>
</tr>
</tbody>
</table>

NOTE: The physical data presented above are typical values and should not be construed as a specification.

10. STABILITY AND REACTIVITY

Hazardous reactions Stable under normal conditions.

Conditions to avoid contact with incompatible materials

Materials to avoid Acids, Oxidizers

Hazardous decomposition Methanol, oxides of carbon, nitrogen oxides (NOx),
products
polymerization Will not occur.

11. TOXICOLOGICAL INFORMATION

Toxicological information on this product or its components appear in this section when such data is available.
Component: Tetramethylammonium hydroxide
Acute dermal toxicity 2.1 % (m)
A single 4h semi-occlusive application to intact rabbit skin produced no signs of dermal irritation.
No clinical signs of toxicity were observed.
DOT Corrosivity testing conducted on stainless steel and laboratory animals determined that this product is not corrosive.

Component: Tetramethylammonium hydroxide
Acute dermal toxicity 3.5 % (m)
A single 4h semi-occlusive application to intact rabbit skin produced minimal signs of irritation (mean scores for erythema or edema less than 2).
No clinical signs of toxicity were observed.

Component: Tetramethylammonium hydroxide
Acute dermal toxicity 5 % (m)
A single 4h semi-occlusive application to intact rabbit skin produced burns (full thickness destruction of skin).
This material is corrosive.
No clinical signs of toxicity were observed.
Corrosive to aluminum per DOT corrosivity testing.

Component: Tetramethylammonium hydroxide
Acute dermal toxicity 7 % (m)
A single 4h semi-occlusive application to intact rabbit skin produced burns (full thickness destruction of skin).
This material is corrosive.
No clinical signs of toxicity were observed.
Corrosive to aluminum per DOT corrosivity testing.

Component: Tetramethylammonium hydroxide
Acute dermal toxicity <5% (w/v):
Repeated application to rat skin for 6 h/d, 5 d/wk for 4 weeks did not produce systemic toxicity.
Test material was applied continuously through a reservoir affixed to shaved animal backs.

Component: Tetramethylammonium hydroxide
Acute dermal toxicity >=5% (w/v):
Repeated application to rat skin for 6 h/d, 5 d/wk for 4 weeks produced rapid toxicity and following effects:
convulsions
death
Effects were noted after 2 hours of initial application.
Test material was applied continuously through a reservoir affixed to shaved animal backs.
12. ECOLOGICAL INFORMATION

Ecotoxicological information on this product or its components appear in this section when such data is available.

**Tetramethylammonium hydroxide**

- **Ecotoxicity effects**
- **Toxicity to aquatic invertebrates**
  - LC50 *Daphnia magna* (water flea) 96 h: 0.07 - 1.2 mg/l
  - A pH neutralized solution has been shown to be toxic to aquatic organisms.

13. DISPOSAL CONSIDERATIONS

**Environmental precautions:** Prevent the material from entering drains or water courses. Do not discharge directly to a water source. Advise Authorities if spillage has entered watercourse or sewer or has contaminated soil or vegetation.

**Disposal**

Dispose in accordance with all local, state (provincial), and federal regulations. Under RCRA, it is the responsibility of the product's user to determine at the time of disposal, whether the product meets RCRA criteria for hazardous waste. This is because the product uses, transformations, mixtures, processes, etc. may render the resulting materials hazardous.

Do not remove label until container is thoroughly cleaned. Empty containers may contain hazardous residues. This material and its container must be disposed of in a safe way.

14. TRANSPORT INFORMATION

**DOT**

- Not regulated for transport

**IMO/IMDG**

- Not regulated (Not dangerous for transport)

15. REGULATORY INFORMATION

**SARA TITLE III: Section 311/312 Categorizations (40CFR370):** Immediate health hazard

**SARA TITLE III: Section 313 Information (40CFR372)**

This product does not contain a chemical which is listed in Section 313 at or above de minimis concentrations.
U.S. Toxic Substances Control Act (TSCA) Section 12(b) Export Notification (40 CFR 707, Subpt D):

This product does not contain any substances subject to Section 12(b) export notification.

U.S. Toxic Substances Control Act (TSCA) All components of this product are in compliance with the inventory listing requirements of the U.S. Toxic Substances Control Act (TSCA) Chemical Substance Inventory.

California (Proposition 65)
This product does not contain materials which the State of California has found to cause cancer, birth defects or other reproductive harm.

16. OTHER INFORMATION

Hazard Rating

<table>
<thead>
<tr>
<th>NFPA</th>
<th>Health</th>
<th>Fire</th>
<th>Reactivity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>3</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

Legend

- ACGIH: American Conference of Governmental Industrial Hygienists
- BAC: Butyl acetate
- OSHA: Occupational Safety and Health Administration
- PEL: Permissible Exposure Limit
- STEL: Short Term Exposure Limit (STEL):
- TLV: Threshold Limit Value
- TWA: Time Weighted Average (TWA):
- —: Bar denotes a revision from prior MSDS.

The information provided in this Safety Data Sheet is correct to the best of our knowledge, information and belief at the date of its publication. The information given is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered a warranty or quality specification. The information relates only to the specific material designated and may not be valid for such material used in combination with any other materials or in any process, unless specified in the text.

Version: 2.0
Print Date: 02/06/2004
Revision date: 01/01/2004
APPENDIX B

FILES USED FOR MICROMILLING MACHINE

STU_COMPLETE_1_ROUGH.txt

;MICROCHANNEL MOLD
;POLYCARBONATE
;1 MM 2 FLUTE ENDMILL

CLOSE
DELETE GATHER
OPEN PROG 1
CLEAR

P4=4000 ;SPINDLE RPM
P2=1 ;SPINDLE ON
I13=10 ;MOVE SEGMENTATION TIME

ABS
F(2.50)
G1 X23.6 Y23.6

Q700=1
WHILE(Q700<14)
INC
F(0.100)
Z(0.200*Q700)
DWELL 100

ABS
F(4.0)
G1 X48.9 DWELL 10
Y23.9 DWELL 10
X23.6 DWELL 10
Y23.6 DWELL 10
Y23.1 DWELL 10
X49.4 DWELL 10
Y24.4 DWELL 10
X23.1 DWELL 10
Y23.1 DWELL 10
X23.6 DWELL 10
Y23.1 DWELL 10
X23.6 DWELL 10
Y22.3 DWELL 10
X50.2 DWELL 10
Y25.2 DWELL 10
X22.3     DWELL 10
Y22.3     DWELL 10
X23.6     DWELL 10
Y21.5     DWELL 10
X51.0     DWELL 10
Y26.0     DWELL 10
X21.5     DWELL 10
Y21.5     DWELL 10
X23.6     DWELL 10
Y20.7     DWELL 10
X51.8     DWELL 10
Y26.8     DWELL 10
X20.7     DWELL 10
Y20.7     DWELL 10
X23.6     DWELL 10
Y19.9     DWELL 10
X52.6     DWELL 10
Y27.6     DWELL 10
X19.9     DWELL 10
Y19.9     DWELL 10
X23.6     DWELL 10
Y19.1     DWELL 10
X53.4     DWELL 10
Y28.4     DWELL 10
X19.1     DWELL 10
Y19.1     DWELL 10
X23.6     DWELL 10
Y18.3     DWELL 10
X54.2     DWELL 10
Y29.2     DWELL 10
X18.3     DWELL 10
Y18.3     DWELL 10
X23.6     DWELL 10
Y17.5     DWELL 10
X55.0     DWELL 10
Y30.0     DWELL 10
X17.5     DWELL 10
Y17.5     DWELL 10
X23.6     DWELL 10
Y16.7     DWELL 10
X55.8     DWELL 10
Y30.8     DWELL 10
X16.7     DWELL 10
Y16.7     DWELL 10
X23.6     DWELL 10
Y15.9     DWELL 10
X56.6     DWELL 10
Y31.6     DWELL 10
X15.9     DWELL 10
Y15.9     DWELL 10
X23.6     DWELL 10
Y15.1     DWELL 10
X57.4     DWELL 10
Y32.4     DWELL 10
X15.1     DWELL 10
Y15.1     DWELL 10

123
X65.4     DWELL 10
Y40.4     DWELL 10
X7.1     DWELL 10
Y7.1     DWELL 10
X23.6     DWELL 10
Y6.3     DWELL 10
X66.2     DWELL 10
Y41.2     DWELL 10
X6.3     DWELL 10
Y6.3     DWELL 10
X23.6     DWELL 10
Y5.5     DWELL 10
X67.0     DWELL 10
Y42.0     DWELL 10
X5.5     DWELL 10
Y5.5     DWELL 10
X23.6     DWELL 10
X23.6 Y23.6    DWELL 10

INC
F(0.100)
Z(-0.200*Q700)
DWELL 100
Q700=Q700+1
END WHILE
;
************************************************************************

ABS
F(2.50)
G1 X60.9 Y23.262

INC
F(0.100)
Z((Q700-1)*.200)
Q701=1
WHILE(Q701<3)
INC
F(0.100)
Z(0.200*Q701)
DWELL 100

ABS
F(4.0)
G1 Y24.238     DWELL 10
X60.412 Y23.75     DWELL 10
X60.9 Y23.262     DWELL 10
X60.546 Y22.909     DWELL 10
X61.4 Y22.055     DWELL 10
X61.471 Y21.56     DWELL 10
X61.7 Y21.331     DWELL 10
Y21.655     DWELL 10
X61.4 Y22.055     DWELL 10
Y25.445     DWELL 10
X61.7  Y25.845  Dwell 10
Y26.169  Dwell 10
X61.471 Y25.94  Dwell 10
X61.4  Y25.445  Dwell 10
X59.705 Y23.75  Dwell 10
X60.546 Y22.909  Dwell 10
X59.981 Y22.343  Dwell 10
X61.376 Y20.948  Dwell 10
G17  G2  X62.2  Y19.882  R5.32  Dwell 10
G1  X62.125 Y19.388  Dwell 10
G2  X62.5  Y18.338  R5.02  Dwell 10
G1  Y19.482  Dwell 10
X62.2  Y19.882  Dwell 10
Y27.618  Dwell 10
X62.5  Y28.018  Dwell 10
Y29.162  Dwell 10
G2  X62.125 Y28.112  R5.02  Dwell 10
G1  X62.2  Y27.618  Dwell 10
G2  X61.376 Y26.552  R5.32  Dwell 10
G1  X58.574 Y23.75  Dwell 10
X59.981 Y22.343  Dwell 10
X61.376 Y20.948  Dwell 10
G2  X62.2  Y19.882  R5.32  Dwell 10
G1  X62.125 Y19.388  Dwell 10
G2  X62.5  Y18.338  R5.02  Dwell 10
G1  X61.7  Y12.642  Dwell 10
X61.513 Y12.465  Dwell 10
X61.4  Y11.978  Dwell 10
G2  X59.877 Y11.1  R6.12  Dwell 10
G1  X61.4  Dwell 10
Y11.978  Dwell 10
X61.7  Y12.378  Dwell 10
Y12.642  Dwell 10
Y12.378  Dwell 10
X61.4  Y11.978  Dwell 10
Y11.1  Dwell 10
X59.877  Dwell 10
X47.644 Y14.0  Dwell 10
X47.514 Y14.13  Dwell 10
X24.986  Dwell 10
X24.856 Y14.0  Dwell 10
X47.644  Dwell 10
Y13.5  Dwell 10
X48.851  Dwell 10
X49.251 Y13.2  Dwell 10
X49.575  Dwell 10
X49.346 Y13.429  Dwell 10
X48.851 Y13.5  Dwell 10
X47.721 Y14.63  Dwell 10
X24.779  Dwell 10
X23.649 Y13.5  Dwell 10
X23.154 Y13.429  Dwell 10
X22.925 Y13.2  Dwell 10
X23.249  Dwell 10
X23.649 Y13.5  Dwell 10
X47.644  Dwell 10
Y12.7 DWELL 10
X50.782 DWELL 10
X51.182 Y12.4 DWELL 10
X51.507 DWELL 10
X51.277 Y12.629 DWELL 10
X50.782 Y12.7 DWELL 10
X48.052 Y15.43 DWELL 10
X24.448 DWELL 10
X21.718 Y12.7 DWELL 10
X21.223 Y12.629 DWELL 10
X20.993 Y12.4 DWELL 10
X21.318 DWELL 10
X21.718 Y12.7 DWELL 10
X47.644 DWELL 10
Y11.9 DWELL 10
X52.714 DWELL 10
X53.114 Y11.6 DWELL 10
X53.499 DWELL 10
X53.21 Y11.843 DWELL 10
X52.714 Y11.9 DWELL 10
X48.384 Y16.23 DWELL 10
X24.116 DWELL 10
X19.786 Y11.9 DWELL 10
X19.29 Y11.843 DWELL 10
X19.001 Y11.6 DWELL 10
X19.386 DWELL 10
X19.786 Y11.9 DWELL 10
X47.644 DWELL 10
Y11.1 DWELL 10
X55.35 DWELL 10
X55.75 Y10.8 DWELL 10
X59.477 DWELL 10
X59.877 Y11.1 DWELL 10
X59.399 Y11.247 DWELL 10
G2 X55.828 R5.82 DWELL 10
G1 X55.35 Y11.1 DWELL 10
G2 X53.286 Y12.459 R6.12 DWELL 10
G1 X48.715 Y17.03 DWELL 10
X23.785 DWELL 10
X19.214 Y12.459 DWELL 10
G2 X17.15 Y11.1 R6.12 DWELL 10
G1 X47.644 DWELL 10
X17.15 DWELL 10
X16.672 Y11.247 DWELL 10
G2 X13.101 R5.82 DWELL 10
G1 X12.623 Y11.1 DWELL 10
G2 X11.1 Y11.978 R6.12 DWELL 10
G1 X10.987 Y12.465 DWELL 10
X10.8 Y12.642 DWELL 10
Y12.378 DWELL 10
X11.1 Y11.978 DWELL 10
Y11.1 DWELL 10
X12.623 DWELL 10
X13.023 Y10.8 DWELL 10
X16.75 DWELL 10
X17.15 Y11.1 DWELL 10
X16.854 Y11.843
G2  X10.3  Y14.09  R5.32
G1  X10.375 Y14.584
G2  X10.0  Y15.634 R5.02
G1  Y14.49
X10.3  Y14.09
Y10.3
X62.2
Y14.09
X62.5  Y14.49
Y15.634
G2  X62.125 Y14.584 R5.02
G1  X62.2  Y14.09
G2  X53.852 Y13.024 R5.32
G1  X49.046 Y17.83
X23.454
X18.648 Y13.024
G2  X16.854 Y11.843 R5.32
X10.3  Y14.09  R5.32
G1  X10.375 Y14.584
G2  X10.0  Y15.634 R5.02
G1  X11.6  Y23.262
X12.088 Y23.75
X11.6  Y24.238
Y23.262
X11.954 Y22.909
X12.795 Y23.75
X11.1  Y25.445
X11.029 Y25.94
X10.8  Y26.169
Y25.845
X11.1  Y25.445
Y22.055
X10.8  Y21.655
Y21.331
X11.029 Y21.56
X11.1  Y22.055
X11.954 Y22.909
X12.519 Y22.343
X13.926 Y23.75
X11.124 Y26.552
G2  X10.3  Y27.618 R5.32
G1  X10.375 Y28.112
G2  X10.0  Y29.162 R5.02
G1  Y28.018
X10.3  Y27.618
Y19.882
X10.0  Y19.482
Y18.338
G2  X10.375 Y19.388 R5.02
G1  X10.3  Y19.882
G2  X11.124 Y20.948 R5.32
G1  X12.519 Y22.343
X11.124 Y20.948
G3  X10.3  Y19.882 R5.32
G1  X10.375 Y19.388
G3  X10.0  Y18.338 R5.02  Dwell 10
G1  Y19.482  Dwell 10
X10.3  Y19.882  Dwell 10
Y22.343  Dwell 10
X9.5  Dwell 10
Y38.0  Dwell 10
X59.877  Dwell 10
Y36.4  Dwell 10
G2  X61.4  Y35.522 R6.12  Dwell 10
G1  X61.513 Y35.035  Dwell 10
X61.7  Y34.858  Dwell 10
Y35.122  Dwell 10
X61.4  Y35.522  Dwell 10
Y36.4  Dwell 10
X59.877  Dwell 10
X47.644 Y33.5  Dwell 10
X24.856  Dwell 10
X24.986 Y33.37  Dwell 10
X47.514  Dwell 10
X47.644 Y33.5  Dwell 10
X47.997 Y33.146  Dwell 10
X48.851 Y34.0  Dwell 10
X49.346 Y34.071  Dwell 10
X49.575 Y34.3  Dwell 10
X49.251  Dwell 10
X48.851 Y34.0  Dwell 10
X23.649  Dwell 10
X23.249 Y34.3  Dwell 10
X22.925  Dwell 10
X23.154 Y34.071  Dwell 10
X23.649 Y34.0  Dwell 10
X24.779 Y32.87  Dwell 10
X47.721  Dwell 10
X47.997 Y33.146  Dwell 10
X48.563 Y32.581  Dwell 10
X50.782 Y34.8  Dwell 10
X51.277 Y34.871  Dwell 10
X51.507 Y35.1  Dwell 10
X51.182  Dwell 10
X50.782 Y34.8  Dwell 10
X21.718  Dwell 10
X21.318 Y35.1  Dwell 10
X20.993  Dwell 10
X21.223 Y34.871  Dwell 10
X21.718 Y34.8  Dwell 10
X24.448 Y32.07  Dwell 10
X48.052  Dwell 10
X48.563 Y32.581  Dwell 10
X49.129 Y32.015  Dwell 10
X52.714 Y35.6  Dwell 10
X53.211 Y35.657  Dwell 10
X53.499 Y35.9  Dwell 10
X53.114  Dwell 10
X52.714 Y35.6  Dwell 10
X19.786  Dwell 10
X19.386 Y35.9  Dwell 10
X19.001     DWELL 10
X19.29 Y35.657     DWELL 10
X19.786 Y35.6     DWELL 10
X24.116 Y31.27     DWELL 10
X48.384     DWELL 10
X49.129 Y32.015     DWELL 10
X49.694 Y31.449     DWELL 10
X53.286 Y35.041     DWELL 10
G2 X55.35 Y36.4 R6.12     DWELL 10
G1 X55.828 Y36.253     DWELL 10
G2 X59.399 R5.82     DWELL 10
G1 X59.877 Y36.4     DWELL 10
X59.477 Y36.7     DWELL 10
X55.75     DWELL 10
X55.35 Y36.4     DWELL 10
X17.15     DWELL 10
G2 X19.214 Y35.041 R6.12     DWELL 10
G1 X23.785 Y30.47     DWELL 10
X48.715     DWELL 10
X49.694 Y31.449     DWELL 10
X48.715 Y30.47     DWELL 10
X23.785     DWELL 10
X19.214 Y35.041     DWELL 10
G3 X17.15 Y36.4 R6.12     DWELL 10
G1 X16.75 Y36.7     DWELL 10
X13.023     DWELL 10
X12.623 Y36.4     DWELL 10
X11.1     DWELL 10
Y35.522     DWELL 10
X10.8 Y35.122     DWELL 10
Y34.858     DWELL 10
X10.987 Y35.035     DWELL 10
X11.1 Y35.522     DWELL 10
G2 X12.623 Y36.4 R6.12     DWELL 10
G1 X13.101 Y36.253     DWELL 10
G2 X16.672 R5.82     DWELL 10
G1 X17.15 Y36.4     DWELL 10
X16.854 Y35.657     DWELL 10
G2 X18.648 Y34.476 R5.32     DWELL 10
G1 X23.454 Y29.67     DWELL 10
X49.046     DWELL 10
X53.852 Y34.476     DWELL 10
G2 X62.2 Y33.41 R5.32     DWELL 10
G1 X62.125 Y32.916     DWELL 10
G2 X62.5 Y31.866 R5.02     DWELL 10
G1 Y33.01     DWELL 10
X62.2 Y33.41     DWELL 10
Y37.2     DWELL 10
X10.3     DWELL 10
Y33.41     DWELL 10
X10.0 Y33.01     DWELL 10
Y31.866     DWELL 10
G2 X10.375 Y32.916 R5.02     DWELL 10
G1 X10.3 Y33.41     DWELL 10
G2 X16.854 Y35.657 R5.32     DWELL 10
G1 X16.558 Y34.913     DWELL 10
X13.387 Y28.815
G2 X12.766 Y30.314 R2.12
G1 Y30.714
G2 X14.889 Y32.834 R2.12
G1 Y40.4
X7.1
Y7.1
X65.4
Y40.4
X14.889
Y30.714
X13.585 Y30.935
G2 X15.819 Y31.647 R1.32
G1 X21.797 Y25.67
X50.703
X56.681 Y31.647
G2 X58.934 Y30.714 R1.32
G1 Y30.314
G2 X58.547 Y29.381 R1.32
G1 X52.917 Y23.75
X58.547 Y18.119
G2 X58.934 Y17.186 R1.32
G1 Y16.786
G2 X56.681 Y15.853 R1.32
G1 X50.703 Y21.83
X21.797
X15.819 Y15.853
G2 X13.566 Y16.786 R1.32
G1 Y17.186
G2 X13.953 Y18.119 R1.32
G1 X19.583 Y23.75
X13.953 Y29.381
G2 X13.566 Y30.314 R1.32
G1 Y30.714
X13.585 Y30.935
Y5.5
X30.935
X5.5
X6.3
Y6.3
X66.2
Y41.2
X6.3
Y30.935
X5.5
Y5.5
X67.0
Y42.0
X5.5
Y30.935
Y30.726
X14.366 Y30.715
G2 X15.254 Y31.082 R0.52
G1 X21.465 Y24.87
X51.035
X57.246 Y31.082
G2 X58.134 Y30.714 R0.52
G1 Y30.314
G2  X57.982 Y29.946 R0.52  DWELL 10
G1  X51.785 Y23.755  DWELL 10
X57.982 Y17.554  DWELL 10
G2  X58.134 Y17.186 R0.52  DWELL 10
G1  Y16.786  DWELL 10
G2  X57.246 Y16.418 R0.52  DWELL 10
G1  X51.035 Y22.63  DWELL 10
X21.465  DWELL 10
X15.254 Y16.418  DWELL 10
G2  X14.366 Y16.786 R0.52  DWELL 10
G1  Y17.186  DWELL 10
G2  X14.518 Y17.554 R0.52  DWELL 10
G1  X20.715 Y23.75  DWELL 10
X14.518 Y29.946  DWELL 10
G2  X14.366 Y30.314 R0.52  DWELL 10
G1  Y30.714  DWELL 10
Y30.715  DWELL 10
X60.9  Y23.262

INC
F(0.100)
Z(-0.200*Q701)
DWELL 100

Q701=Q701+1
END WHILE

P2=0
F(2.5)
Z(-20)
CLOSE
STU COMPLETE 1 FINISH.txt

;MICROCHANNEL MOLD
;POLYCARBONATE
;0.013" (0.330 MM) 2 FLUTE ENDMILL

CLOSE
DELETE GATHER
OPEN PROG 1
CLEAR

P4=6000 ;SPINDLE RPM
P2=1 ;SPINDLE ON
I13=10 ;MOVE SEGMENTATION TIME

ABS
F(2.50)
G1 X51.182 Y24.515

Q700=1
WHILE(Q700<5)
INC
F(0.100)
Z(0.100*Q700)
DWELL 100

ABS
F(1.0)
G1 X57.497 Y30.831 DWELL 10
G17 G2 X57.779 Y30.714 R0.165 DWELL 10
G1 Y30.314 DWELL 10
G2 X57.731 Y30.197 R0.165 DWELL 10
G1 X51.367 Y23.833 DWELL 10
G2 X51.25 Y23.785 R0.165 DWELL 10
G1 X21.25 DWELL 10
G2 X21.133 Y23.833 R0.165 DWELL 10
G1 X14.769 Y30.197 DWELL 10
G2 X14.721 Y30.314 R0.165 DWELL 10
G1 Y30.714 DWELL 10
G2 X15.003 Y30.831 R0.165 DWELL 10
G1 X21.318 Y24.515 DWELL 10
X51.182 DWELL 10

INC
F(1.00)
Z(-1.0) DWELL 10
DWELL 100

ABS
F(2.5)
G1 Y22.985 DWELL 10
INC
F(1.00)
Z(1.0)          DWELL 10
DWELL 100

ABS
F(1.0)
G1 X21.318      DWELL 10
X15.003 Y16.669 DWELL 10
G2 X14.721 Y16.786 R0.165 DWELL 10
G1 Y17.186      DWELL 10
G2 X14.769 Y17.303 R0.165 DWELL 10
G1 X21.133 Y23.667 DWELL 10
G2 X21.25 Y23.715 R0.165 DWELL 10
G1 X51.25       DWELL 10
G2 X51.367 Y23.667 R0.165 DWELL 10
G1 X57.731 Y17.303 DWELL 10
G2 X57.779 Y17.186 R0.165 DWELL 10
G1 Y16.786      DWELL 10
G2 X57.497 Y16.669 R0.165 DWELL 10
G1 X51.182 Y22.985 DWELL 10

INC
F(1.00)
Z(-1.0)         DWELL 10
DWELL 100

ABS
F(2.5)
G1 X51.182 Y24.515 DWELL 10

INC
F(1.00)
Z(1.0)          DWELL 10
DWELL 100

INC
F(0.100)
Z(-0.100*Q700) DWELL 100
Q700=Q700+1
END WHILE
P2=0
F(2.5)
Z(-20)
CLOSE
APPENDIX C

MANUFACTURER DATA SHEETS FOR SYLGARD 184 AND SU-8-50

Information About High Technology Silicone Materials

SYLGARD® 184 SILICONE ELASTOMER, BASE & CURING AGENT

<table>
<thead>
<tr>
<th>Type</th>
<th>Two-part silicone elastomer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physical Form</td>
<td>Transparent</td>
</tr>
<tr>
<td>as supplied</td>
<td>Medium-Viscosity liquid</td>
</tr>
<tr>
<td>as cured</td>
<td>Flexible elastomer</td>
</tr>
<tr>
<td>Cure</td>
<td>25 to 150°C (77 to 302°F)</td>
</tr>
<tr>
<td>Special Properties</td>
<td>Transparent, reversion resistant, stable dielectric, high physical properties</td>
</tr>
<tr>
<td>Primary Uses</td>
<td>Potting and encapsulating</td>
</tr>
<tr>
<td>Benefits</td>
<td>Low toxicity, repairability</td>
</tr>
</tbody>
</table>

DESCRIPTION
SYLGARD® 184 silicone elastomer, base & curing agent, is supplied as a two-part kit comprised of liquid components. When the base and the curing agent are thoroughly mixed in a 10:1 weight ratio, the medium-viscosity liquid mixture has a consistency resembling SAE No. 40 motor oil. The liquid mixture will cure in thick or thin sections to a flexible, transparent elastomer ideally suited for electrical/electronic potting and encapsulating applications.

SYLGARD 184 silicone elastomer offers a flexible cure schedule from 25 to 150°C (77 to 302°F), without an exotherm. SYLGARD 184 silicone elastomer requires no post cure and can be placed in service immediately following the completion of the cure schedule at any operating temperature from -65 to 200°C (-67 to 392°F). See Table 1 for special features and benefits of this product.

USES
Because of its many special features, SYLGARD 184 silicone elastomer is ideal for a wide variety of electrical/electronic potting and encapsulating applications and provides environmental protection for:

- Equipment modules
- Relays, power supplies and magnetic amplifiers
- Transformers, coils and ferrite cores
- Connectors
- Fiber optic waveguide coatings
- Encapsulation of circuit boards

SYLGARD 184 silicone elastomer is especially useful in applications where clarity is desirable, including:

- Encapsulation of solar cells
- Encapsulation of opto-electronic displays

HOW TO USE
Mixing
SYLGARD 184 silicone elastomer is supplied in two parts, a 10:1-matched base and curing agent, mixed in a ratio of 10 parts base to one part curing agent, by weight.

For best curing results, glassware or tinned cans and glass or metal stirring implements should be used. Mix with a smooth action that will minimize the introduction of excess air.

Pot Life – Working Time
Cure reaction of SYLGARD 184 silicone elastomer, base & curing agent, begins with the mixing process. Initially, cure is evidenced by a gradual increase in viscosity, followed by gelation and conversion to a solid elastomer. Pot life is defined as the time required for viscosity to double following addition of curing agent to base. At 25°C (77°F), the pot life of SYLGARD 184 silicone elastomer is 2 hours. Pot life of this product can be extended by refrigeration at 4°C (40°F); however, do not allow moisture from condensation to collect in the silicone elastomer.

SYLGARD 184 silicone elastomer will cure at 25°C (77°F), will become solid in 24 hours, and will reach full cure in 7 days. See “Curing.”

Processing
Thoroughly mix SYLGARD 184 silicone elastomer, base & curing agent, in a ratio of 10 parts base to 1 part curing agent, by weight. Agitate gently to reduce the amount of air introduced. Allow mixture to set 30 minutes before pouring.

Since air bubbles are usually present following mixing, vacuum de-airing is recommended. De-air in a container with at least four times the liquid volume to allow for expansion of material. Air entrapped in the mixture can be removed by using a vacuum of 25 to 29 inches of mercury. Continue the vacuum until the liquid expands and settles to its original volume and bubbling subsides. This may take 15 minutes to 2 hours, depending on...
## Typical Properties

These values are not intended for use in preparing specifications.

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>As Supplied</strong></td>
<td></td>
</tr>
<tr>
<td>CTM 0001A</td>
<td>Specific Gravity at 25 C (77 F)</td>
</tr>
<tr>
<td>CTM 0050</td>
<td>Viscosity at 25 C (77 F), centipoise</td>
</tr>
<tr>
<td><strong>As Cured</strong></td>
<td></td>
</tr>
<tr>
<td>CTM 0050</td>
<td>Viscosity at 25 C (77 F), centipoise</td>
</tr>
<tr>
<td>CTM 0055</td>
<td>Pol Life, minimum, hours</td>
</tr>
<tr>
<td><strong>As Cured - Physical</strong></td>
<td></td>
</tr>
<tr>
<td>CTM 0176</td>
<td>Appearance</td>
</tr>
<tr>
<td>CTM 0099</td>
<td>Durometer Hardness, Shore A</td>
</tr>
<tr>
<td>CTM 0137A</td>
<td>Tensile Strength, MPa (psi)</td>
</tr>
<tr>
<td>CTM 0137A</td>
<td>Elongation, percent</td>
</tr>
<tr>
<td>CTM 0159A</td>
<td>Tear Strength, die B, kN/cm (ipsi)</td>
</tr>
<tr>
<td>CTM 0022</td>
<td>Specific Gravity at 25 C (77 F)</td>
</tr>
<tr>
<td>CTM 0224</td>
<td>Thermal Conductivity, cal/cm²-sec-C/°C</td>
</tr>
<tr>
<td>CTM 0585</td>
<td>Linear Coefficient of Thermal Expansion, cm/cm per °C from -55 to 150 C (-67 to 302 F)</td>
</tr>
<tr>
<td>MIL-I-16923G</td>
<td>Thermal Shock Resistance, from -49 to 88 C (-55 to 155 F), passes 10 cycles</td>
</tr>
<tr>
<td>ASTM D 570</td>
<td>Water Absorption, after 7 days immersion at 25 C (77 F), percent</td>
</tr>
<tr>
<td>ASTM D 449</td>
<td>Internal Resistance, ohm-cm</td>
</tr>
<tr>
<td>UL 94</td>
<td>Flammability Classification</td>
</tr>
<tr>
<td>UL</td>
<td>Temperature Rating, Mechanical, °C (°F)</td>
</tr>
<tr>
<td>Electrical, °C (°F)</td>
<td>130 (265)</td>
</tr>
<tr>
<td><strong>Electrical</strong></td>
<td></td>
</tr>
<tr>
<td>CTM 0114A</td>
<td>Dielectric Strength, volts/mil</td>
</tr>
<tr>
<td>CTM 0112</td>
<td>Dielectric Constant, at 60 Hz</td>
</tr>
<tr>
<td></td>
<td>100 Hz</td>
</tr>
<tr>
<td></td>
<td>100 kHz</td>
</tr>
<tr>
<td>CTM 0112</td>
<td>Dissipation Factor, at 60 Hz</td>
</tr>
<tr>
<td></td>
<td>100 Hz</td>
</tr>
<tr>
<td></td>
<td>100 kHz</td>
</tr>
<tr>
<td>CTM 0249A</td>
<td>Volume Resistivity, ohm-cm</td>
</tr>
<tr>
<td>CTM 0171</td>
<td>Arc Resistance, track, seconds</td>
</tr>
<tr>
<td><strong>Electrical - after heat aging for 1000 hours at 200 C (392 F)</strong></td>
<td></td>
</tr>
<tr>
<td>CTM 0114A</td>
<td>Dielectric Strength, volts/mil</td>
</tr>
<tr>
<td>CTM 0112</td>
<td>Dielectric Constant, at 60 Hz</td>
</tr>
<tr>
<td></td>
<td>100 Hz</td>
</tr>
<tr>
<td>CTM 0249A</td>
<td>Volume Resistivity, ohm-cm</td>
</tr>
</tbody>
</table>

*In most cases, CTMs (Corporate Test Methods) correspond to ASTM standard tests. Copies of CTM procedures are available upon request.

*Storvized Viscosimeter Model LHF, spindle H at 60 rpm.

*Mix rate 10 parts base to 1 part curing agent, by weight.

*Time required to double catalyzed viscosity.

*Based on sample thickness of 125 mils and a minimum cure of 4 hours at 85 C (185 F).

*Tests, claims, representations and descriptions regarding flammability are based on standard small scale laboratory tests.

*Properties obtained on 1.5-mil-film (0.032-inch) specimens, cure 4 hours at 65 C (149 F).

*Measured with 1.4-inch standard ASTM electrode, 500 volts per second rate of rise.

**Specification Writers:** Please contact Dow Corning Corporation, Midland, Michigan, before writing specifications on this product.

Amount of air introduced during curing:

Clean and degrease all application surfaces using a solvent to remove all mold release agent, processing oils and surface contaminant. Dry and remove all solvent before application.

For best adhesion the application substrate should be primed with SYLGARD® primer. Obtain a technical bulletin for proper application instructions. See "Bonning."

**CAUTION:** SYLGARD primer is flammable. Keep away from heat, sparks and open flame. Use only with adequate ventilation.

When pouring SYLGARD 184 silicone elastomer, use a pouring container, care should be taken to minimize air entrapment. When practical, pouring should be done under vacuum, particularly if the component being poured or encapsulated has many fine voids. If this technique cannot be used, the unit should be evacuated after SYLGARD 184 silicone elastomer has been poured.

After allowing time for the material to reach temperature, cure according to the cure time-temperature listed in "Curing."

**Reparability**

In the manufacture of electrical/electronic devices it is often desirable to salvage or reclaim damaged or defective units. With most rigid types of potting and encapsulating materials, removal of entry is often impossible without causing excessive damage to internal circuitry. SYLGARD 184 silicone elastomer should be selectively removed with relative ease, allowing repair or changes to be made and the area to be repotted with additional product. SYLGARD 184 silicone elastomer, base & curing agent, can be removed by cutting with a sharp blade or knife and tearing material away from the area to be repaired. Thin sections of the adhesive elastomer are best removed from substrates and circuitry by mechanical action such as scraping or rubbing; removal can be assisted by the application of isopropyl alcohol.

Before repotting a repaired device, if possible, roughen the exposed surface of the SYLGARD 184 silicone elastomer, base & curing agent, with an abrasive paper. This will enhance...
adhesion and permit the repair material to become an integral matrix with the existing elastomer. Silicone prime coats are not recommended for adhering SYLGARD 184 silicone elastomer, base & curing agent, to itself in rejoicing applications.

Reversion Resistance
SYLGARD 184 silicone elastomer, base & curing agent, provides excellent reversion resistance even when exposed to temperatures in excess of 200 C (392 F) while under confinement or in deep section.

Temperature Stability
SYLGARD 184 silicone elastomer, base & curing agent, cures to a thermostatic material that will not melt or appreciably soften even at elevated temperatures of 250 C (482 F). The material may harden or become brittle after prolonged exposure to these elevated temperatures. SYLGARD 184 silicone elastomer, base & curing agent, has a 190 C (385 F) UL yellow-card temperature index classification for both electrical and mechanical functional use.

Upon exposure to lower temperatures, SYLGARD 184 silicone elastomer, base & curing agent, does not approach a stifferening point until -55 C (-67 F). Overall, the cured elastomer will maintain its basic elastomeric flexibility over an extremely wide range of -55 to 200 C (-67 to 392 F), making it the ideal selection for applications that may experience high or low temperature cycling.

PROCESSING TECHNIQUES
Curing
SYLGARD 184 silicone elastomer, base & curing agent, can be satisfactorily cured either with exposure to air or completely sealed, and at temperatures ranging from 25 to 150 C (77 to 302 F). Curing time can be appreciably decreased by elevating the cure temperature. One of the following cure cycles is suggested:

- 25 C (77 F) for 24 hours
- 65 C (149 F) for 4 hours
- 100 C (212 F) for 1 hour
- 150 C (302 F) for 15 minutes

Larger parts will require additional time in the oven to heat to the selected cure temperature. Satisfactory sources of heat include circulating and non-circulating ovens, infrared heating lamps and heat guns.

Full mechanical strength will not be achieved for 7 days when SYLGARD 184 silicone elastomer, base & curing agent, is cured at 25 C (77 F). The majority of its physical strength, however, is present after 24 hours, although surface cure may not yet be complete.

Bonding
SYLGARD 184 silicone elastomer, base & curing agent, will not normally bond to clean, nonporous surfaces such as metal or glass. A primer coat is required to ensure adhesion to these surfaces. SYLGARD prime coat is recommended to obtain the best adhesion. The prime coat should be applied in a thin layer to clean, dry surfaces where adhesion is desired. It should be air dried 1 to 2 hours before the silicone elastomer is applied.

Lowering Viscosity
The viscosity of SYLGARD 184 elastomer, base & curing agent, can be lowered by the addition of 200% fluid, 50 cSt, from Dow Corning. Quantities of 10 percent or less will have little or no effect on the physical or electrical properties. Quantities of 10 percent or greater may decrease the physical strength and hardness but will have no effect on the electrical properties. At concentrations greater than 10 percent the fluid may possibly bleed from cured SYLGARD 184 silicone elastomer. The addition of thinning fluid does not change the mixing ratio of curing agent to base.

Varying the Hardness
Variations of up to 10 percent in concentration of curing agent in SYLGARD 184 silicone elastomer will have little effect on cure time or on the physical properties of the final cured elastomer. Lowering the curing agent concentration by more than 10 percent will result in a softer and weaker elastomer; increasing the concentration by more than 10 percent will result in over-hardening of the cured elastomer and will tend to degrade the physical and thermal properties. Changes in the curing agent concentration will have little or no effect on the electrical properties, however.

Release Agents
When SYLGARD 184 silicone elastomer, base & curing agent, is cured in a mold, the mold should first be treated with a release agent to prevent sticking. Suitable release agents include DUPONOL® WAX at a 5 percent concentration with isopropyl alcohol. For a 2 percent concentration with CHLOROTHENE® or similar chlorinated solvent; a mild liquid detergent at a 2 to 5 percent concentration with water; or petroleum jelly at a 5 percent solution in a chlorinated solvent.

LIMITATIONS
Inhibition of Cure
In the presence of inhibitors, cure of SYLGARD 184 silicone elastomer is poor. In the inhibited area (usually less than 0.020-inch thick) the silicone elastomer remains in a liquid or tacky state even though the cure schedule has been completed, and despite any subsequent attempts to convert it to a hard, dry, rubbery mass, this material will remain uncured. Extremely minute quantities of inhibitor may be sufficient to produce this effect. The most notable causes of inhibition include:

- Organotin and other organometallic compounds
- Silicone rubbers containing organotin catalyst
- Sulfur, polysulfides, polytetraflouror and other sulfur-containing materials
- Amines, urethanes, amine-containing materials and other nitrogen-containing materials
- Unsaturated hydrocarbon plasticizers

If a substrate or material is considered questionable in respect to potential curing inhibition, a small-scale compatibility test to ascertain suitability in the particular application is recommended. If inhibition is present it may sometimes be overcome by probing the unit at the highest tolerable temperature for approximately 1 to 4 hours to remove volatile chemicals. See bulletin No. 10-022, "How To

"Duponol" is a registered trademark of E. I. du Pont de Nemours & Company.
"Chlorothene" is a registered trademark of The Dow Chemical Company.
Process SYLGARD® Brand Elastomers® for a listing of inhibition-causing materials.

**Thermal Expansion**
SYLGARD 184 silicone elastomer, base & curing agent, has a notable volume coefficient of thermal expansion (9.6 x 10^-4/°C; see Typical Properties chart). The volume of the cured elastomer will increase or decrease approximately 9.6 percent for each 100 °C (212 °F) of temperature differential. When using SYLGARD 184 silicone elastomer, base & curing agent, at higher temperatures in applications that are highly confined or hermetically sealed, allowance should be made to accommodate volume expansion and prevent pressure build-up. Normal thermal expansion and contraction stresses may be minimized by selecting a cure temperature close to the midpoint of the high and low extremes of the thermal cycle.

Temperature limits may be influenced by differences in the thermal expansion values between the silicone elastomer and the encapsulated or potted components, and also by their configurations. Therefore, thermal operating limits should be determined by testing before large scale use.

**SAFETY REQUIREMENTS**

**Handling**
SYLGARD 184 silicone elastomer, base & curing agent, does not contain volatile solvents. Special ventilation is not required in the normal use or storage of this product. Base and curing agent liquid components or their cured mixture do not present any significant toxicological hazard incidental to normal industrial handling. Minimal eye protection, such as standard safety glasses, should be adequate for normal industrial use. Direct eye contact can cause temporary eye discomfort; flush thoroughly with copious amounts of water should contact occur. See Material Safety Data Sheet for more detailed handling instructions.

**TABLE: SPECIAL FEATURES AND BENEFITS**

<table>
<thead>
<tr>
<th>Special Feature</th>
<th>Benefit</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low toxicity</td>
<td>No special precautions required for normal industrial handling</td>
</tr>
<tr>
<td>No solvents or cure by-products; no exotherm during cure</td>
<td>Requires no special venting; will not cause corrosion; low shrinkage during cure</td>
</tr>
<tr>
<td>Cures to a transparent, flexible elastomer</td>
<td>Provides stability and relief from mechanical shock; low transmission of vibration; visual inspection of components; and easy repairability</td>
</tr>
<tr>
<td>Environmental protection</td>
<td>Low water absorption; good radiation resistance; little out-gassing in hard vacuum</td>
</tr>
<tr>
<td>Excellent dielectric properties</td>
<td>Maintains and protects existing electrical insulation requirements</td>
</tr>
<tr>
<td>Stability over a wide temperature range; reversion resistant</td>
<td>Maintains elastomeric flexibility and provides functional stability from -55 to 200 °C (-67 to 392 °F), even in confinement</td>
</tr>
<tr>
<td>Flame resistant</td>
<td>UL flammability classification of 94-V-1 and a temperature rating of 130 °C (265 °F)</td>
</tr>
</tbody>
</table>

**Abnormal Exposures**

**CAUTION:** The liquid curing agent component of SYLGARD 184 silicone elastomer may generate hydrogen gas if contaminated with strong acids, bases or catalytic oxidizing materials. If exposure is suspected, use appropriate caution to relieve hydrogen gas pressure. Keep away from sparks and flame, and supply adequate ventilation to reduce localized build-up of hydrogen gas.

**Spills**
Spills of the liquid base and curing agent components of SYLGARD 184 silicone elastomer can become extremely slippery. Sawdust or other absorbent material should be immediately applied to any liquid spill for temporary relief. The spill should be removed with high flash point mineral spills or other suitable solvent.

**SHIPPING LIMITATIONS**
None.

**STORAGE AND SHELF LIFE**
When stored in original unopened containers at or below 32 °C (90 °F), SYLGARD 184 silicone elastomer, base & curing agent, has a shelf life of 24 months from date of manufacture.

**PACKAGING**
SYLGARD 184 silicone elastomer, base & curing agent, is shipped in kits that contain both the base and curing agent, liquid components in separate containers. Each kit contains the appropriate weight of curing agent for the amount of base. Complete kits are available in 1.1-, 8.8-, 44- and 495-lb (0.5-, 4-, 20- and 225-kg) quantities, net weight.

**MSDS INFORMATION**
ATTENTION: PRODUCT SAFETY INFORMATION REQUIRED FOR SAFE USE IS NOT INCLUDED. BEFORE HANDLING, READ PRODUCT AND MATERIAL SAFETY DATA SHEETS.
AND CONTAINER LABELS FOR SAFE USE. PHYSICAL AND HEALTH HAZARD INFORMATION. THE MATERIAL SAFETY DATA SHEET IS AVAILABLE FROM YOUR DOW CORNING REPRESENTATIVE, OR DISTRIBUTOR, OR BY WRITING TO DOW CORNING CUSTOMER SERVICE, OR BY CALLING (517) 496-9000.

WARRANTY INFORMATION—PLEASE READ CAREFULLY

Dow Corning believes that the information in this publication is an accurate description of the typical characteristics and/or uses of the product or products, but it is your responsibility to thoroughly test the product in your specific application to determine its performance, efficacy and safety.

Unless Dow Corning provides you with a specific written warranty of fitness for a particular use, Dow Corning’s sole warranty is that the product or products will meet Dow Corning’s then current sales specifications. DOW CORNING SPECIFICALLY DISCLAIMS ANY OTHER EXPRESS OR IMPLIED WARRANTIES OF MERCHANTABILITY AND OF FITNESS FOR USE. Your exclusive remedy and Dow Corning’s sole liability for breach of warranty is limited to refund of the purchase price or replacement of any product shown to be other than as warranted, and Dow Corning expressly disclaims any liability for incidental or consequential damages. Suggestions of use should not be taken as inducements to infringe any particular patent.
NANO™ SU-8
Negative Tone Photoresist Formulations 50-100

- High aspect ratio imaging with near vertical side walls
- Near UV (350-400nm) processing
- Film thicknesses from 1 to >200μm with single spin coat processes
- Superb chemical and temperature resistance

SU-8 is a high contrast, epoxy based photoresist designed for micromachining and other microelectronic applications, where a thick chemically and thermally stable image is desired. The exposed and subsequently cross-linked portions of the film are rendered insoluble to liquid developers. SU-8 has very high optical transparency above 360nm, which makes it ideally suited for imaging near vertical sidewalls in very thick films. SU-8 is best suited for permanent applications where it is imaged, cured and left in place.

Process Guidelines
SU-8 is most commonly processed with conventional near UV (350-400nm) radiation, although it may be imaged with e-beam or x-ray. (x-line 365nm) is recommended. Upon exposure, cross-linking proceeds in two-steps (1) formation of a strong acid during the exposure processes, followed by (2) acid-initiated, thermally driven epoxy cross-linking during the post-exposure bake (PEB) step.

A normal process is: spin coat, soft bake, expose, post-exposure bake (PEB) and develop. A controlled hard bake is recommended to further cross-link the imaged SU-8 structures where they will remain as part of the device. The entire process should be optimized for the specific application. A baseline process is given here to be used as a starting point.
**Substrate Pretreatment**

To obtain maximum process reliability, substrates should be clean and dry prior to applying the SU-8 resist. Start with a solvent cleaning, or rinse with dilute acid, followed by DI water rinse. Where applicable, substrates should be subjected to a piranha etch/clean (H₂SO₄ & H₂O₂). To dehydrate the surface, bake at 200°C for 5 minutes on a hotplate. For applications that require electroplating and subsequent removal of SU-8, use MicroChem’s Omnicoat prior to processing SU-8.

**Coat**

SU-8 resists are designed to produce low defect coatings over a very broad range of film thickness. The film thickness versus spin speed data displayed in Table 1, and Figure 1, provide the information required to select the appropriate SU-8 resist and spin conditions, to achieve the desired film thickness.

The recommended coating conditions are:

1. **STATIC Dispense**: Approximately 1 ml of SU-8 per inch of substrate diameter.
2. **Spread Cycle**: Ramp to 500 rpm at 100 rpm/second acceleration. Hold at this speed for 5-10 seconds to allow the resist to cover the entire surface.
3. **Spin Cycle**: Ramp to final spin speed at an acceleration of 300 rpm/second and hold for a total of 30 seconds.

![SU-8 Spin Speed Curve](image)

**Figure 1. Spin speed vs. thickness curves for selected SU-8 resists.**

<table>
<thead>
<tr>
<th>Product Name</th>
<th>Viscosity (cSt)</th>
<th>Thickness (μm)</th>
<th>Spin Speed (rpm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SU-8 50</td>
<td>12250</td>
<td>40</td>
<td>3000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>100</td>
<td>1000</td>
</tr>
<tr>
<td>SU-8 100</td>
<td>51500</td>
<td>750</td>
<td>1000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>250</td>
<td>1000</td>
</tr>
</tbody>
</table>

Table 1. Thickness vs. spin speed data for selected SU-8 resists.

**Soft Bake**

After the resist has been applied to the substrate, it must be soft baked to evaporate the solvent and densify the film. SU-8 is normally baked on a level hot plate, although convection ovens may be used. The following bake times are based on contact hot plate processes. Bake times should be optimized for proximity and convection oven bake processes since solvent evaporation rate is influenced by the rate of heat transfer and ventilation.

For best results, ramping or stepping the soft bake temperature is recommended. Lower initial bake temperatures allow the solvent to evaporate out of the film at a more controlled rate, which results in better coating fidelity, reduced edge bevel and better resist-to-substrate adhesion. Refer to Table 2, for TWO STEP contact hot plate process recommendations.

<table>
<thead>
<tr>
<th>Product Name</th>
<th>Thickness (μm)</th>
<th>Prebake @ 65°C</th>
<th>Softbake @ 95°C</th>
</tr>
</thead>
<tbody>
<tr>
<td>SU-8 50</td>
<td>50</td>
<td>6</td>
<td>20</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>10</td>
<td>30</td>
</tr>
<tr>
<td>SU-8 100</td>
<td>130</td>
<td>20</td>
<td>50</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>30</td>
<td>90</td>
</tr>
</tbody>
</table>

Table 2. Recommended soft bake parameters
Expose

SU-8 is optimized for near-UV (350-400nm) exposure. i-line exposure tools are recommended. SU-8 is virtually transparent and insensitive above 400nm but has high optical absorption below 350nm. This can be seen in Figure 2. Excessive dose below 350nm may, therefore, result in overexposure of the top portion of the resist film, resulting in exaggerated negative sidewall profiles or T-topping. The optimal exposure dose will depend on film thickness (thicker films require higher dosage) and process parameters. The exposure dose recommendations in Table 3, are based on source intensity measurements taken with an i-line (365nm) radiometer and probe.

Exposure tip: When using a broad spectral output source, for best imaging results, i.e., straightest sidewalls, filter out excessive energy below 350nm.

Catastrophic adhesion failure, severely negative sidewalls and excessive cracking often indicate an under-cure condition. To correct the problem, increase the exposure dose and/or increase the post exposure bake (PEB) time.

Post Exposure Bake

Following exposure, a post-exposure bake (PEB) must be performed to selectively cross-link the exposed portions of the film. This bake can be performed either on a hot plate or in a convection oven. Optimum cross-link density is obtained through careful adjustments of the exposure and PEB process conditions. The bake recommendations below are based on results obtained with a convection hot plate.

PEB tip: SU-8 is readily cross-linked and can result in a highly stressed film. To minimize stress, wafer bowing and resist cracking, a slow ramp or TWO STEP contact hot plate process, as shown in Table 4, is recommended. Rapid cooling after PEB should be avoided.

![Graph showing SU-8 absorbance vs. film thickness](Figure 2. SU-8 absorbance vs. film thickness)

![Graph showing SU-8 exposure vs. film thickness](Figure 3. SU-8 2000 exposures vs. film thickness)

<table>
<thead>
<tr>
<th>Product Name</th>
<th>Thickness (μm)</th>
<th>PEB 1 @65°C</th>
<th>PEB 2 @95°C</th>
</tr>
</thead>
<tbody>
<tr>
<td>SU-8 50</td>
<td>50</td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>1</td>
<td>10</td>
</tr>
<tr>
<td>SU-8 100</td>
<td>150</td>
<td>1</td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>1</td>
<td>20</td>
</tr>
</tbody>
</table>

Table 3. Recommended exposure dose processes

Table 4. Recommended post exposure bake parameters
Develop
SU-8 resists have been optimized for use with MicroChem’s SU-8 Developer. Immersion, spray or spray-puddle processes can be used. Other solvent based developers such as ethyl lactate and diacetone alcohol may also be used. Strong agitation is recommended for high aspect ratio and/or thick film structures. Recommended develop times are given in Table 5 for immersion processes. These proposed develop times are approximate, since actual dissolution rates can vary widely as a function of agitation rate, temperature and resist processing parameters.

<table>
<thead>
<tr>
<th>Product Name</th>
<th>Thickness (um)</th>
<th>Development (minutes)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SU-8 50</td>
<td>50</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>SU-8 100</td>
<td>150</td>
<td>15</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>20</td>
</tr>
</tbody>
</table>

Table 5. Recommended develop processes

Rinse and Dry
Following development, the substrate should be rinsed briefly with isopropyl alcohol (IPA), then dried with a gentle stream of air or nitrogen.

Rinse tip: If a white film is produced during rinse, this is an indication that the substrate has been underdeveloped. Simply immerse or spray the substrate with SU-8 developer to remove the film and complete the development process. Repeat the rinse step.

Hard Bake (cure)
SU-8 has good mechanical properties, therefore hard bakes are normally not required. For applications where the imaged resist is to be left as part of the final device, the resist may be ramped hard baked between 150-200°C on a hot plate or in a convection oven to further cross link the material. Bake times vary based on type of bake process and film thickness.

Remove
SU-8, after expose and PEB, is a highly cross-linked epoxy, which makes it extremely difficult to remove with conventional solvent based resist strippers. MicroChem’s Remover PG will swell and lift off minimally cross-linked SU-8 2000.

However, if OmniCoat has been applied immersion in Remover PG should effect a clean and thorough Lift-Off of the SU-8 2000 Material. It will not remove fully cured or hard baked SU-8 2000 without the use of OmniCoat. Alternate removal processes include immersion in oxidizing acid solutions such as piranha etch / clean, plasma ash, RIE, laser ablation and pyrolysis.

To remove minimally cross-linked SU-8 2000, or if using Omnicoat, with Remover PG, heat the bath to 50-80°C and immerse the substrates for 30-90 minutes. Actual strip time will depend on resist thickness and cross-link density.

Formore information on MicroChem Omnicoat and Remover PG please see the relevant product data sheets.

Storage
Store SU-8 resists upright in tightly closed containers in a cool, dry environment away from direct sunlight at a temperature of 40-70°F (4-21°C). Store away from light, acids, heat and sources of ignition. Shelf life is twelve months from date of manufacture.

Disposal
SU-8 resists may be included with other waste containing similar organic solvents to be discarded for destruction or reclaim in accordance with local state and federal regulations. It is the responsibility of the customer to ensure the disposal of SU-8 resists and residues made in observance all federal, state, and local environmental regulations.

Environmental, Health and Safety
Consult product Material Safety Data Sheet before working with SU-8 resists. Handle with care. Wear chemical goggles, chemical gloves and suitable protective clothing when handling SU-8 resists. Do not get into eyes, or onto skin or clothing. Use with adequate ventilation to avoid breathing vapors or mist. In case of contact with skin, wash affected area with soap and water. In case of contact with the eyes, rinse immediately with water and flush for 15 minutes lifting eyelids frequently. Get emergency medical assistance.

The information is based on our experience and is, we believe to be reliable, but may not be complete. We make no guarantee or warranty, expressed or implied, regarding the information, use, handling, storage, or possession of these products, or the application of any process described herein or the results desired, once since the conditions of use and handling of these products are beyond our control.
APPENDIX D

DERIVATIONS OF IMPEDANCE MODEL EQUATIONS

\[ \frac{1}{Z_{RC_b}} = \frac{1}{R_b} \frac{Z_c}{R + Z_c} \rightarrow Z_{RC_b} = \frac{RZ_c}{R + Z_c} = \frac{R_B}{R_B + \left( -\frac{1}{\omega C_B} \right)} = \frac{R_B}{jR_B C_B \omega + 1} \]

\[ Z = \frac{R_A - \frac{1}{j \omega C_A} + Z_{RC_b}}{Z_1 Z_2} \]

\[ |Z_1| = \sqrt{\left( -\frac{1}{j \omega C_A} \right)^2 + R_A^2} = \frac{\sqrt{1 + (R_A C_A \omega)^2}}{C_A \omega}, \quad \Theta_1 = \arctan \left( \frac{-1}{R_A C_A \omega} \right) \]

\[ Z_2 = \frac{R_B \angle 0^\circ}{\sqrt{1 + (R_B C_B \omega)^2}} \quad \angle \Theta_2 = \arctan \left( \frac{R_B C_B \omega}{1} \right) \rightarrow L - \Theta_2 = -\arctan \left( R_B C_B \omega \right) \]
APPENDIX E

SAMPLE WORKSHEET OF COLLECTED IMPEDANCE DATA

4294A REV1.11
DATE: Jul 23 2005

MEASURE PARAMETER: IMPEDANCE MAG PHASE (DEG)
ADAPTER: PROBE
SWEEP TYPE: LOG FREQ
NUMBER of POINTS: 100
POINT DELAY TIME: 0 Sec
SWEEP DELAY TIME: 0 Sec
OSC LEVEL: 10 mVolt
DC BIAS: OFF
BW: 3
SWEEP AVERAGING: OFF
POINT AVERAGING: OFF

TRACE: A
FORMAT: LINEAR

<table>
<thead>
<tr>
<th>Frequency</th>
<th>Data Trace Real</th>
<th>Data Trace Imag</th>
</tr>
</thead>
<tbody>
<tr>
<td>4.00E+01</td>
<td>5.43E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.43E+01</td>
<td>7.07E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.91E+01</td>
<td>6.90E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.44E+01</td>
<td>4.27E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.02E+01</td>
<td>3.98E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.67E+01</td>
<td>4.22E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.39E+01</td>
<td>3.75E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>8.19E+01</td>
<td>3.72E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>9.07E+01</td>
<td>3.23E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.00E+02</td>
<td>2.96E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.11E+02</td>
<td>2.73E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.23E+02</td>
<td>2.40E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.37E+02</td>
<td>2.35E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.51E+02</td>
<td>2.18E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.67E+02</td>
<td>2.07E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.86E+02</td>
<td>1.87E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.06E+02</td>
<td>1.67E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.28E+02</td>
<td>1.61E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.52E+02</td>
<td>1.46E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.79E+02</td>
<td>1.38E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.09E+02</td>
<td>1.18E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>Value</td>
<td>Value</td>
<td>Value</td>
</tr>
<tr>
<td>---------</td>
<td>---------</td>
<td>---------</td>
</tr>
<tr>
<td>3.43E+02</td>
<td>1.12E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.80E+02</td>
<td>1.03E+06</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.21E+02</td>
<td>9.45E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.66E+02</td>
<td>8.72E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.16E+02</td>
<td>8.40E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.72E+02</td>
<td>7.54E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.33E+02</td>
<td>6.97E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.01E+02</td>
<td>6.38E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.77E+02</td>
<td>5.90E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>8.61E+02</td>
<td>5.47E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>9.53E+02</td>
<td>5.00E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.06E+03</td>
<td>4.66E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.17E+03</td>
<td>4.28E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.30E+03</td>
<td>3.91E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.44E+03</td>
<td>3.66E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.59E+03</td>
<td>3.37E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.76E+03</td>
<td>3.12E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.95E+03</td>
<td>2.88E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.16E+03</td>
<td>2.64E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.39E+03</td>
<td>2.46E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.65E+03</td>
<td>2.27E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.94E+03</td>
<td>2.10E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.25E+03</td>
<td>1.94E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.60E+03</td>
<td>1.78E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.99E+03</td>
<td>1.65E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.42E+03</td>
<td>1.53E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.90E+03</td>
<td>1.41E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.42E+03</td>
<td>1.30E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.01E+03</td>
<td>1.20E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.66E+03</td>
<td>1.11E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.37E+03</td>
<td>1.02E+05</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>8.17E+03</td>
<td>9.50E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>9.05E+03</td>
<td>8.77E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.00E+04</td>
<td>8.08E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.11E+04</td>
<td>7.46E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.23E+04</td>
<td>6.91E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.36E+04</td>
<td>6.37E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.51E+04</td>
<td>5.90E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.67E+04</td>
<td>5.46E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.85E+04</td>
<td>5.04E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.05E+04</td>
<td>4.66E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.27E+04</td>
<td>4.30E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.52E+04</td>
<td>3.98E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.79E+04</td>
<td>3.69E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.09E+04</td>
<td>3.41E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.42E+04</td>
<td>3.16E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.79E+04</td>
<td>2.93E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.20E+04</td>
<td>2.72E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.65E+04</td>
<td>2.53E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.15E+04</td>
<td>2.35E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.70E+04</td>
<td>2.18E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.32E+04</td>
<td>2.04E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.00E+04</td>
<td>1.90E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.75E+04</td>
<td>1.77E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>8.59E+04</td>
<td>1.66E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>9.51E+04</td>
<td>1.55E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.05E+05</td>
<td>1.45E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.17E+05</td>
<td>1.37E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.29E+05</td>
<td>1.29E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.43E+05</td>
<td>1.22E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.59E+05</td>
<td>1.15E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.76E+05</td>
<td>1.09E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.95E+05</td>
<td>1.04E+04</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.16E+05</td>
<td>9.85E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.39E+05</td>
<td>9.42E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.65E+05</td>
<td>9.02E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>2.93E+05</td>
<td>8.65E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.25E+05</td>
<td>8.33E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.60E+05</td>
<td>8.03E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>3.98E+05</td>
<td>7.77E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.41E+05</td>
<td>7.54E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>4.89E+05</td>
<td>7.33E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>5.41E+05</td>
<td>7.14E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.00E+05</td>
<td>6.97E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>6.64E+05</td>
<td>6.80E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>7.36E+05</td>
<td>6.65E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>8.15E+05</td>
<td>6.51E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>9.03E+05</td>
<td>6.39E+03</td>
<td>0.00E+00</td>
</tr>
<tr>
<td>1.00E+06</td>
<td>6.27E+03</td>
<td>0.00E+00</td>
</tr>
</tbody>
</table>
APPENDIX F

IMPEDEANCE DATA FOR DEVICE 2 AND DEVICE 3

Results from Device 2 with media only – no flow

<table>
<thead>
<tr>
<th>Frequency</th>
<th>10x10</th>
<th>20x20</th>
<th>40x40</th>
<th>80x80</th>
<th>160x160</th>
</tr>
</thead>
<tbody>
<tr>
<td>1kHz</td>
<td>71.68</td>
<td>32.06</td>
<td>12.19</td>
<td>16.76</td>
<td>0.38</td>
</tr>
<tr>
<td>10kHz</td>
<td>36.59</td>
<td>9.31</td>
<td>4.06</td>
<td>3.85</td>
<td>0.18</td>
</tr>
<tr>
<td>100kHz</td>
<td>15.19</td>
<td>2.80</td>
<td>1.49</td>
<td>0.94</td>
<td>0.07</td>
</tr>
<tr>
<td>1MHz</td>
<td>3.09</td>
<td>0.52</td>
<td>0.30</td>
<td>0.18</td>
<td>0.02</td>
</tr>
<tr>
<td>Electrode</td>
<td>$R$ (Ω)</td>
<td>$C$ (F)</td>
<td>$C$ (F/cm$^2$)</td>
<td>$R^2$</td>
<td></td>
</tr>
<tr>
<td>-----------</td>
<td>---------</td>
<td>---------</td>
<td>----------------</td>
<td>-------</td>
<td></td>
</tr>
<tr>
<td>10x10</td>
<td>84600</td>
<td>1.58E-10</td>
<td>4.88E-06</td>
<td>0.9458</td>
<td></td>
</tr>
<tr>
<td>20x20</td>
<td>34600</td>
<td>2.70E-10</td>
<td>5.05E-06</td>
<td>0.9687</td>
<td></td>
</tr>
<tr>
<td>40x40</td>
<td>25900</td>
<td>4.08E-10</td>
<td>5.64E-06</td>
<td>0.9672</td>
<td></td>
</tr>
<tr>
<td>80x80</td>
<td>21400</td>
<td>5.18E-10</td>
<td>3.33E-06</td>
<td>0.9652</td>
<td></td>
</tr>
<tr>
<td>160x160</td>
<td>6780</td>
<td>2.79E-09</td>
<td>6.43E-06</td>
<td>0.9593</td>
<td></td>
</tr>
</tbody>
</table>
Results from Device 2 with media and fibronectin – no flow

<table>
<thead>
<tr>
<th>Standard Deviation (Ω/µm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frequency 10x10 20x20 40x40 80x80 160x160</td>
</tr>
<tr>
<td>1kHz 56.95 15.11 5.13 1.98 0.20</td>
</tr>
<tr>
<td>10kHz 11.70 2.56 1.42 0.23 0.03</td>
</tr>
<tr>
<td>100kHz 1.50 0.52 0.23 0.04 0.01</td>
</tr>
<tr>
<td>1MHz 0.31 0.21 0.08 0.03 0.01</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Model Fit</th>
</tr>
</thead>
<tbody>
<tr>
<td>Electrode R (Ω) C (F) C (F/cm²) R²</td>
</tr>
<tr>
<td>10x10 47300 1.28E-10 3.96E-06 0.9768</td>
</tr>
<tr>
<td>20x20 23800 2.72E-10 5.08E-06 0.9808</td>
</tr>
<tr>
<td>40x40 17800 4.12E-10 5.70E-06 0.9772</td>
</tr>
<tr>
<td>80x80 9010 9.46E-10 6.08E-06 0.9806</td>
</tr>
<tr>
<td>160x160 4800 3.05E-09 7.03E-06 0.9767</td>
</tr>
</tbody>
</table>
Results from Device 3 with media only – no flow

![Graph showing impedance vs. frequency for different electrode sizes](image)

<table>
<thead>
<tr>
<th>Frequency</th>
<th>10x10</th>
<th>20x20</th>
<th>40x40</th>
<th>80x80</th>
<th>160x160</th>
</tr>
</thead>
<tbody>
<tr>
<td>1kHz</td>
<td>102.07</td>
<td>78.98</td>
<td>46.73</td>
<td>4.47</td>
<td>0.77</td>
</tr>
<tr>
<td>10kHz</td>
<td>19.75</td>
<td>13.09</td>
<td>7.92</td>
<td>1.15</td>
<td>0.10</td>
</tr>
<tr>
<td>100kHz</td>
<td>2.87</td>
<td>2.28</td>
<td>1.09</td>
<td>0.18</td>
<td>0.01</td>
</tr>
<tr>
<td>1MHz</td>
<td>0.50</td>
<td>0.41</td>
<td>0.18</td>
<td>0.04</td>
<td>0.00</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>electrode</th>
<th>R (Ω)</th>
<th>C (F)</th>
<th>C (F/cm²)</th>
<th>R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>10x10</td>
<td>36100</td>
<td>1.42E-10</td>
<td>3.93E-06</td>
<td>0.9790</td>
</tr>
<tr>
<td>20x20</td>
<td>37700</td>
<td>1.39E-10</td>
<td>2.31E-06</td>
<td>0.9767</td>
</tr>
<tr>
<td>40x40</td>
<td>31300</td>
<td>1.67E-10</td>
<td>2.15E-06</td>
<td>0.9783</td>
</tr>
<tr>
<td>80x80</td>
<td>17500</td>
<td>3.98E-10</td>
<td>2.75E-06</td>
<td>0.9682</td>
</tr>
<tr>
<td>160x160</td>
<td>6250</td>
<td>1.14E-09</td>
<td>2.77E-06</td>
<td>0.9787</td>
</tr>
</tbody>
</table>
Results from Device 3 with media and fibronectin – no flow

<table>
<thead>
<tr>
<th>Frequency</th>
<th>10x10</th>
<th>20x20</th>
<th>40x40</th>
<th>80x80</th>
<th>160x160</th>
</tr>
</thead>
<tbody>
<tr>
<td>1kHz</td>
<td>134.37</td>
<td>68.61</td>
<td>35.04</td>
<td>11.45</td>
<td>1.34</td>
</tr>
<tr>
<td>10kHz</td>
<td>23.16</td>
<td>13.05</td>
<td>6.47</td>
<td>2.03</td>
<td>0.23</td>
</tr>
<tr>
<td>100kHz</td>
<td>3.02</td>
<td>2.03</td>
<td>1.07</td>
<td>0.25</td>
<td>0.03</td>
</tr>
<tr>
<td>1MHz</td>
<td>0.54</td>
<td>0.29</td>
<td>0.17</td>
<td>0.02</td>
<td>0.01</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Electrode</th>
<th>R (Ω)</th>
<th>C (F)</th>
<th>C (F/cm²)</th>
<th>R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>10x10</td>
<td>47900</td>
<td>1.11E-10</td>
<td>3.07E-06</td>
<td>0.9773</td>
</tr>
<tr>
<td>20x20</td>
<td>44300</td>
<td>1.10E-10</td>
<td>1.83E-06</td>
<td>0.9763</td>
</tr>
<tr>
<td>40x40</td>
<td>34600</td>
<td>1.47E-10</td>
<td>1.90E-06</td>
<td>0.9776</td>
</tr>
<tr>
<td>80x80</td>
<td>16600</td>
<td>2.85E-10</td>
<td>1.97E-06</td>
<td>0.9800</td>
</tr>
<tr>
<td>160x160</td>
<td>6920</td>
<td>8.47E-10</td>
<td>2.06E-06</td>
<td>0.9807</td>
</tr>
</tbody>
</table>
Results from Device 2 (40x40 $\mu$m$^2$) and Device 3 (10x10 $\mu$m$^2$ and 160x160 $\mu$m$^2$) with medium alone (top) and medium and fibronectin (below) – flow
CURRICULUM VITAE

NAME: Stuart Joseph Williams

ADDRESS: 2535 Dundee Rd
Louisville, KY 40205

DOB: Louisville, KY – October 14, 1981

EDUCATION: B. S., Mechanical Engineering
University of Louisville
Louisville, KY
2000-2005

PROFESSIONAL EXPERIENCE: Sandia National Laboratories
Organization 1764: Micro Analytical Systems
Albuquerque, NM
2002-2003

PROFESSIONAL SOCIETIES: ASME, BMES