Supplementary Materials: Passive Mixing Capabilities of Micro- and Nanofibres When Used in Microfluidic Systems

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Figure S1. A typical electrospinning apparatus consisting of a syringe filled with a polymer spinning dope, a high voltage power source, a grounded collector plate, and a syringe pump.

Figure S2. (A) Hot Embossing—A copper template with a raised channel structure is used to emboss the channel design in a piece of PMMA. The PMMA is sandwiched between the copper template and a blank piece of copper and placed on a Carver Laboratory Hot Press at 130 °C and 10,000 lbs (44,482 N) of force. Inlet at outlet holes are drilled at the ends of the channels; (B) UVO Treatment—The embossed PMMA and another piece of PMMA modified with electrospun fibers are both placed in a UV-ozone oven and treated to allow for UVO-assisted thermal bonding; (C) Bonding — The UVO-treated PMMA pieces are sandwiched together (with the fibers facing the embossed channels) and placed between two blank pieces of copper. The assembly is placed on the hot press at bonded together at a lower temperature and pressure than hot embossing; (D) Completed Device—Polyvinyl chloride tubing is glued into the inlet and outlet holes of the completed device.
Data Analysis

We have added a step-by-step discussion of the calculation of the mixing index into the SI:

**Data Analysis**

The mixing index is calculated using the following formula:

\[
\text{mixing index} = \frac{1}{\sqrt{N}} \sum \left( \frac{I_k - I_o}{I_o} \right)^2
\]

- \(I_k\) = intensity of an individual pixel in a column;
- \(I_o\) = average intensity of all pixels in a column;
- \(N\) = number of pixels in a column.

The mixing index for each channel is a value between 0 and 1, where a value of 0 indicates complete mixing within the channel. The mixing index was chosen to quantify the degree of mixing because it allowed us to compare our mixing results with the results from other labs.

When calculating the mixing index, a fluorescent microscopy image of the channel was taken during fluid flow. Images were taken either at the inlet (before the fiber mat) or at the outlet (after the fiber mat):
In order to calculate the mixing index, we first needed to get the intensity of each individual pixel in a column (blue arrow in above image) that extends along the width of the channel (these are our $I_k$ values) as well as the average intensity of all the pixels in that column (this is our $I_0$ value). The length of the column in pixels is our $N$ value. This was done for 50 consecutive columns in each photo taken using the following ImageJ command:

```java
run("ROI Manager...");
for (i=Xn; i<Xn+50;i++) {
    makeLine (i,Y1,i,Y2);
    roiManager("Add");
}
roiManager("Multi Plot");
```

where $X_n$ is the x location of the first column being measured (where x is its location along the length of the channel) and $Y_1$ and $Y_2$ are the y coordinates of the top and bottom of the channel (along the width of the channel) that indicate where the column being measured:

Using this code, ImageJ produced a plot of the pixel intensity along each column as well as a list of the intensities for all the columns measured. An example of the pixel intensity plot for a single column is seen below:
The average intensity ($I_o$) of the pixels in the column was calculated from the ImageJ data and then the mixing index was calculated for each column. The mixing index values for each of the 50 columns were then averaged to get the average mixing index for that channel. Mixing indices are provided here with two decimal places to provide a more refined analyses of difference between the channels and mats analyzed.

**Figure S4.** (A) Examples of flow profiles observed in thick nanofiber mat samples; (B) Variation in nanofiber porosity and distribution along thickness of nanofiber mats. Confocal microscopy.

**Figure S5.** Outlet mixing index observed in 5 mm long two-layer mats at different flow rates. 3 channels were tested at each flow rate.
Table S1. Multiple linear regression variables and their outcome. The outlet mixing index values for each PVA mat morphology was compared to the outlet mixing index for empty control channels using linear regression. The linear regression controlled for the effect of flow rate on the mixing value. It was determined that mixing was significantly increased in all of the fiber mats, with the greatest mixing increase seen with the two-layer mats.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Significant within Family of Comparisons?</th>
<th>Average Change to Mixing Value</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 mm, 1 Layer</td>
<td>Yes</td>
<td>−0.182</td>
<td>1.7234 x 10^{-5}</td>
</tr>
<tr>
<td>3 mm, 2 Layer</td>
<td>Yes</td>
<td>−0.285</td>
<td>2.3516 x 10^{-10}</td>
</tr>
<tr>
<td>5 mm, 1 Layer</td>
<td>Yes</td>
<td>−0.209</td>
<td>3.5847 x 10^{-7}</td>
</tr>
<tr>
<td>5 mm, 2 Layer</td>
<td>Yes</td>
<td>−0.344</td>
<td>4.4726 x 10^{-12}</td>
</tr>
<tr>
<td>10 mm, 1 Layer</td>
<td>Yes</td>
<td>−0.136</td>
<td>2.3049 x 10^{-3}</td>
</tr>
<tr>
<td>10 mm, 2 Layer</td>
<td>Yes</td>
<td>−0.340</td>
<td>6.6718 x 10^{-12}</td>
</tr>
</tbody>
</table>

Table S2. Multiple linear regression variables and their outcome for analyzing the effect of poly(vinyl alcohol) mat morphology. There were three families for comparison: (1) the different fiber mat lengths, (2) having two layers, and (3) the flow rate. Having two layers and flow rate were each considered to be significantly increased. However, the different fiber mat lengths did not appear to have an effect on mixing index.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Significant Within Family of Comparisons?</th>
<th>Average Change to Mixing Index Value</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 mm</td>
<td>No</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>5 mm</td>
<td>No</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>10 mm</td>
<td>No</td>
<td>-</td>
<td>Over all p = 0.24</td>
</tr>
<tr>
<td>2 Layers</td>
<td>Yes</td>
<td>−0.142</td>
<td>5.15 x 10^{-8}</td>
</tr>
<tr>
<td>Flow Rate</td>
<td>Yes</td>
<td>0.03 per (µL/s)</td>
<td>0.0045</td>
</tr>
</tbody>
</table>

Figure S6. Flow profiles observed in channels containing PS fiber mats. Flow at 1 µL/min. Black lines indicate location of nanofiber mats within the channels. Leica Fluorescent Microscope. 5× objective.
Figure S7. Flow profiles at inlets of channels containing different PS fiber morphologies. Flow rate was 1 µL/min.

Figure S8. Flow profiles at outlets of channels containing different PS fiber morphologies. Flow rate was 1 µL/min.
Table S3. Multiple linear regression variables and their outcome. The different fiber mat types were considered as one family for comparison, and the flow rate another using linear regression. The linear regression controlled for the effect of flow rate on fluid mixing. 1 Layer 17.5%, 2 Layer 15%, and 2 Layer 17.5%, along with flow rate, were all considered to be significantly increased.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Significant within Family of Comparisons?</th>
<th>Average Change to Mixing Index Value</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Layer, 12.5%</td>
<td>No</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>1 Layer, 15%</td>
<td>No</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>1 Layer, 17.5%</td>
<td>Yes</td>
<td>-0.128</td>
<td>0.0018</td>
</tr>
<tr>
<td>2 Layer, 12.5%</td>
<td>No</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>2 Layer, 15%</td>
<td>Yes</td>
<td>-0.146</td>
<td>0.00046</td>
</tr>
<tr>
<td>2 Layer, 17.5%</td>
<td>Yes</td>
<td>-0.127</td>
<td>0.0021</td>
</tr>
</tbody>
</table>

Table S4. Multiple linear regression variables and their outcomes for analyzing effects of polystyrene mat morphology on mixing index. The effect of fiber diameter, fiber mat height, and flow rate on outlet mixing index was determined using multiple linear regression. The mixing increased with increasing fiber diameter, fiber mat height, and decreased with increasing flow rate.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Compared To</th>
<th>Significant within Family of Comparisons?</th>
<th>Average Change to Mixing Value</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>15%</td>
<td>12.5%</td>
<td>No</td>
<td>-0.078</td>
<td>0.0077</td>
</tr>
<tr>
<td>17.5%</td>
<td>12.5%</td>
<td>Yes</td>
<td>-0.077</td>
<td>0.0090</td>
</tr>
<tr>
<td>17.5%</td>
<td>15%</td>
<td>Yes</td>
<td>-0.077</td>
<td>0.0090</td>
</tr>
<tr>
<td>2 Layers</td>
<td>1 Layer</td>
<td>Yes</td>
<td>-0.084</td>
<td>5.17 × 10⁻⁴</td>
</tr>
<tr>
<td>Flow Rate</td>
<td>-</td>
<td>Yes</td>
<td>0.017 per (µL/min)</td>
<td>0.0481</td>
</tr>
</tbody>
</table>


**Code S1. PVA Analysis-Effect of Length and Number of Fiber Mat Layers**

```matlab
%Multiple Regression on PVA Data Separated by Length and Layer
%Author: Lauren Colangelo and Nicholas Colangelo
%Last Edit: 03-23-2015

%As long as data input into the Table appropriately, everything should be taken care of between data sets except for the following,
%which should be done for each new data set:
%To do:
%1) Make sure boxplots and other plots represent the data to be analyzed.
%2) Make sure labels for plots are appropriate.
%3) Make sure Holm Test is using the appropriate data from the SortData Table.
%Clear memory of variables that may be used here:
clear
%Load the data:
Data = readtable('C:\MatlabPVASep.xlsx');
SortData = sortrows(Data,'Flow');
SortData.Type = categorical(SortData.Type,'Ordinal',false);
SortData.Length = categorical(SortData.Length,'Ordinal',false);
SortData.Layer = categorical(SortData.Layer,'Ordinal',false);
SortData.Channel = categorical(SortData.Channel,'Ordinal',false);

%Box Plot:
figure(1)
subplot(2,2,1)
```
boxplot(SortData.Mixing,SortData.Length)
xh = xlabel('Length');
yh = ylabel('Mixing Value');
set([xh,yh], 'fontweight', 'bold');
set(gca,'XTick', 1:3)
set(gca,'XTickLabel', {'3 mm', '5 mm', '10 mm'});

subplot(2,2,2)
boxplot(SortData.Mixing,SortData.Layer)
xh = xlabel('Layer');
yh = ylabel('Mixing Value');
set([xh,yh], 'fontweight', 'bold');

subplot(2,2,3)
boxplot(SortData.Mixing,SortData.Flow)
xh = xlabel('Flow (\muL/sec)');
yh = ylabel('Mixing Value');
set([xh,yh], 'fontweight', 'bold');

% Regression:
modelspec = 'Mixing ~ Length + Layer + Flow';
lm = fitlm(table2dataset(SortData), modelspec);
disp(lm);

% Table of ANOVA Results:
disp(anova(lm));

% Show Number of Outliers as Determined by Cook’s Distance >= 3*MeanCooksDistance:
Cook = sum(lm.Diagnostics.CooksDistance./mean(lm.Diagnostics.CooksDistance) >= 3);
fprintf('Number of Values above Cook’s Distance %f, Cook

% Obtain Parameters for Holm Test from the Regression
MSres = double(lm.MSE);
DF = double(lm.DFE);

% Number of each length:
for i = 1:max(double(SortData.Length))
    n(i) = sum(double(SortData.Length) == i);
end

% Input the number of types, less one (So it doesn’t compare values against themselves)
for i = 1:(max(double(SortData.Length)-1))
    for j = (i+1):max(double(SortData.Length))
        % Comparison to Control
        t(i,j) = (lm.Coefficients.Estimate(j) - 0)/ sqrt(MSres*(1/n(j)+1/n(i)));
        % Between types comparison
        else
            t(i,j) = (lm.Coefficients.Estimate(j) - lm.Coefficients.Estimate(i))/ sqrt(MSres*(1/n(j)+1/n(i)));
    end %if i
end %j
end %i

% Now we take the t values to p values
p1 = 2*(1-tcdf(abs(t),DF));
%Organize p-values
Sorted1 = sort(reshape(p1,1,size(t,1)*size(t,2)));

%Holm test p-value for comparison with uncorrected p-value of \( p = 0.05 \):
%Calculate number of comparisons:
Comparisons = 0;
i = 0;
while i < max(double(SortData.Length))
    for i = 1:max(double(SortData.Length))
        Comparisons = Comparisons + (i-1);
    end
end

%Determine Significant Terms
detect1 = 0;
for i = 1:Comparisons
    pcomp = 0.05/(Comparisons-i+1);
    if Sorted1(i) <= pcomp
        Sig1(i) = 1;
detect1 = 1;
    else
        break
    end
end

%Display p values in p matrix that are significant
if detect1 == 1
    fprintf(‘
\( p \) matrix is:
’)
    format short e
    disp(p1.*(p1<=Sorted1(length(Sig1))))
else
    fprintf(‘
\( p \) matrix has no significant values.
’)
end

%Layer
%A lot of this code is unnecessary as there is only two layers to
%compare.
%Number of each Layer:
for i = 1:max(double(SortData.Layer)) \%different types
    n2(i) = sum(double(SortData.Layer) == i);
end

%Input the number of types, less one (So it doesn’t compare values against themselves)
for i = 1:(max(double(SortData.Layer))-1))
    %Input the total number of types, so it can make all the comparisons
    for j = (i+1):max(double(SortData.Layer))
        %Comparison to Control
        %Note: Accessing the coefficient array must account for Length variables
        if i == 1
            t2(i,j) = (lm.Coefficients.Estimate(max(double(SortData.Length))-1+j) - 0)/ sqrt(MSres*(1/n2(j)+1/n2(i)));
        elseif else
            t2(i,j) = (lm.Coefficients.Estimate(max(double(SortData.Length))-1+j)) - lm.Coefficients.Estimate(i))/ sqrt(MSres*(1/n2(j)+1/n2(i)));
        end
    end
end
end %i
%Now we take the t values to p values
p2 = 2*(1-tcdf(abs(t2),DF));
%Organize p-values
Sorted2 = sort(reshape(p2,1,size(t2,1)*size(t2,2)));  
%Holm test p-value for comparison with uncorrected p-value of p=0.05:  
%Calculate number of comparisons:
Comparisons2 = 0;
i = 0;
while i < max(double(SortData.Layer))
    for i = 1:max(double(SortData.Layer))
        Comparisons2 = Comparisons2 + (i-1);
    end %for i
end %while i
%Determine Significant Terms
detect2 = 0;
for i = 1:Comparisons2
    pcomp = 0.05/(Comparisons2-i+1);
    if Sorted2(i) <= pcomp
        Sig2(i) = 1;
        detect2 = 1;
    else
        break
    end %p(i)
end %i
%Display p values in p matrix that are significant
if detect2 == 1
    fprintf('np2 matrix is:
')
    format short e
    disp(p2.*(p2<=Sorted2(length(Sig2))))
else
    fprintf('np2 matrix has no significant values.
')
end %if

%Residuals
figure(2)
gscatter(double(SortData.Type),lm.Residuals.Raw,SortData.Flow)
yh = ylabel('Residuals');
xh = xlabel('Types');
legh = legend;
htitle = get(legh,'Title');
set(htitle,'String','Flow (\muL/sec)')
set([xh,yh], 'fontweight', 'bold');
set(gca,'XTick', 1:6)
set(gca,'XTickLabel',{'3 mm 1 Layer', '3 mm 2 Layer', '5 mm 1 Layer', '5 mm 2 Layer', '10 mm 1 Layer', '10 mm 2 Layer'})

%Test Normality
figure(3)
normplot(lm.Residuals.Standardized)
xh = xlabel('Standardized Residuals');
yh = ylabel('Normal Probability Scale');
set([xh,yh], 'fontweight', 'bold');
Code S2. PVA Analysis—Comparison of each PVA mat type

% Multiple Regression on PVA by Type
% Author: Lauren Colangelo and Nicholas Colangelo
% Last Edit: 03-23-2015
% As long as data input into the Table appropriately, everything should be taken care of between data sets except for the following,
% which should be done for each new data set:
% To do:
% 1) Make sure boxplots and other plots represent the data to be analyzed.
% 2) Make sure labels for plots are appropriate.
% 3) Make sure Holm Test is using the appropriate data from the SortData Table.
% Clear memory of variables that may be used here:
clear

% Load the data:
Data = readtable('C:\MatlabPVAType.xlsx');
SortData = sortrows(Data,'Flow');
SortData.Type = categorical(SortData.Type,'Ordinal',false);
SortData.Channel = categorical(SortData.Channel,'Ordinal',false);

% Box Plot:
figure(1)
subplot(2,1,1)
boxplot(SortData.Mixing,SortData.Type)
xh = xlabel('Type');
yh = ylabel('Mixing Value');
set([xh,yh],’fontweight’,’bold’);
set(gca,’XTick’, 1:7)
set(gca,’XTickLabel’,{'Control’, ‘3 mm 1 Layer’, ‘3 mm 2 Layer’, ‘5 mm 1 Layer’, ‘5 mm 2 Layer’,
’10 mm 1 Layer’, ’10 mm 2 Layer’});
subplot(2,1,2)
boxplot(SortData.Mixing,SortData.Flow)
xh = xlabel(‘Flow (\muL/min)’);
yh = ylabel(‘Mixing Value’);
set([xh,yh],’fontweight’,’bold’);

% Regression:
modelspec = ‘Mixing ~ Type + Flow’;
lm = fitlm(table2dataset(SortData),modelspec);
disp(lm);
% Table of ANOVA Results:
disp(anova(lm));
% Show Number of Outliers as Determined by Cook’s Distance >= 3*MeanCooksDistance:
Cook = sum(lm.Diagnostics.CooksDistance./mean(lm.Diagnostics.CooksDistance) >= 3);
fprintf(’Number of Values above Cook’s Distance %f’, Cook)
% Obtain Parameters for Holm Test from the Regression
% TBL(3,3) = MSres
MSres = double(lm.MSE);
% TBL(3,2) = Degrees of Freedom
DF = double(lm.DFE);
% Number of each type:
for i = 1:max(double(SortData.Type)) % different types
n(i) = sum(double(SortData.Type) == i);
end %
%Holm Test
%Input the number of types, less one (So it doesn't compare values against themselves)
for i = 1:(max(double(SortData.Type)-1))
    %Input the total number of types, so it can make all the comparisons
    for j = (i+1):max(double(SortData.Type))
        %Comparison to Control
        if i == 1
            t(i,j) = (lm.Coefficients.Estimate(j) - 0)/ sqrt(MSres*(1/n(j)+1/n(i)));
        %Between types comparison
        else
            t(i,j) = (lm.Coefficients.Estimate(j) - lm.Coefficients.Estimate(i))/
                sqrt(MSres*(1/n(j)+1/n(i)));
        end %if i
    end %j
end %i
%Now we take the t values to p values
p = 2*(1-tcdf(abs(t),DF));
%Organize p-values
Sorted = sort(reshape(p,1,size(t,1)*size(t,2)));
%Holm test p-value for comparison with uncorrected p-value of p=0.05:
%Calculate number of comparisons:
Comparisons = 0;
i = 0;
while i < max(double(SortData.Type))
    for i = 1:max(double(SortData.Type))
        Comparisons = Comparisons + (i-1);
    end %for i
end %while i
%Determine Significant Terms
detect = 0;
for i = 1:Comparisons
    pcomp = 0.05/(Comparisons-i+1);
    if Sorted(i) <= pcomp
        Sig(i) = 1;
detect = 1;
    else
        break
    end %if Sorted
end %if i
%Display p values in p matrix that are significant
if detect == 1
    fprintf('
np matrix is: 
')
    format short e
    disp(p.*(p<=Sorted(length(Sig))));
else
    fprintf('
np1 matrix has no significant values.
')
end %if
%Residuals
figure(2)
gscatter(double(SortData.Type),lm.Residuals.Raw,SortData.Flow)
yh = ylabel('Residuals');
xh = xlabel('Types');
legh = legend;
htitle = get(legh,'Title');
set(htitle,'String','Flow (\muL/min)')
set([xh,yh], 'fontweight', 'bold');
set(gca, 'XTick', 1:7)
set(gca, 'XTickLabel', {'Control', '3 mm 1 Layer', '3 mm 2 Layer', '5 mm 1 Layer', '5 mm 2 Layer', '10 mm 1 Layer', '10 mm 2 Layer'})

% Test Normality with a Normal Probability Plot
figure(3)
normplot(lm.Residuals.Standardized)
xh = xlabel('Standardized Residuals');
yh = ylabel('Normal Probability Scale');
set([xh,yh], 'fontweight', 'bold');

Code S3. PS Analysis- Effect of polymer weight percent and number of layers

% Multiple Regression on Polystyrene Data Separated by Weight and Layer
% Author: Lauren Colangelo and Nicholas Colangelo
% Last Edit: 03-23-2015
% As long as data input into the Table appropriately, everything should be taken care of between data sets except for the following,
% which should be done for each new data set:
% To do:
% 1) Make sure boxplots and other plots represent the data to be analyzed.
% 2) Make sure labels for plots are appropriate.
% 3) Make sure Holm Test is using the appropriate data from the SortData Table.

% Clear memory of variables that may be used here:
clear

% Load the data:
Data = readtable('C:\MatlabPolyStySep.xlsx');
SortData = sortrows(Data,'Flow');
SortData.Type = categorical(SortData.Type,'Ordinal',false);
SortData.Weight = categorical(SortData.Weight,'Ordinal',false);
SortData.Layer = categorical(SortData.Layer,'Ordinal',false);
SortData.Channel = categorical(SortData.Channel,'Ordinal',false);

% Box Plot the data:
figure(1)
subplot(2,2,1)
boxplot(SortData.Mixing,SortData.Weight)
xh = xlabel('Weight');
yh = ylabel('Mixing Value');
set([xh,yh], 'fontweight', 'bold');
set(gca, 'XTick', 1:3)
set(gca, 'XTickLabel', {'12.5%', '15%', '17.5%'});
subplot(2,2,2)
boxplot(SortData.Mixing,SortData.Layer)
xh = xlabel('Layer');
yh = ylabel('Mixing Value');
set([xh,yh], 'fontweight', 'bold');
subplot(2,2,3)
boxplot(SortData.Mixing,SortData.Flow)
xh = xlabel('Flow (\muL/sec);
yh = ylabel('Mixing Value');
set([xh,yh],'fontweight','bold');

%Regression:
modelspec = 'Mixing ~ Weight + Layer + Flow';
lm = fitlm(table2dataset(SortData),modelspec);
disp(lm);
%Table of ANOVA Results:
disp(anova(lm));
%Show Number of Outliers as Determined by Cook's Distance >= 3*MeanCooksDistance:
Cook = sum(lm.Diagnostics.CooksDistance./mean(lm.Diagnostics.CooksDistance) >= 3);
fprintf('Number of Values above Cook’s Distance %f', Cook)

%Obtain Parameters for Holm Test from the Regression
%MSres
MSres = double(lm.MSE);
%Degrees of Freedom
DF = double(lm.DFE);

%Weight Percentage
%Number of each weight percentage:
for i = 1:max(double(SortData.Weight)) %different types
    n(i) = sum(double(SortData.Weight) == i);
end %i

%Input the number of types, less one (So it doesn’t compare values against themselves)
for i = 1:(max(double(SortData.Weight)-1))
    %Input the total number of types, so it can make all the comparisons
    for j = (i+1):max(double(SortData.Weight))
        %Comparison to Control
        if i == 1
            t(i,j) = (lm.Coefficients.Estimate(j) - 0)/ sqrt(MSres*(1/n(j)+1/n(i)));
        %Between types comparison
        else
            t(i,j) = (lm.Coefficients.Estimate(j) - lm.Coefficients.Estimate(i))/
            sqrt(MSres*(1/n(j)+1/n(i)));
        end %if i
    end %j
end %i

%Now we take the t values to p values
p1 = 2*(1-tcdf(abs(t),DF));
%Organize p-values
Sorted1 = sort(reshape(p1,1,size(t,1)*size(t,2)));

%Holm test p-value for comparison with uncorrected p-value of p=0.05:
%Calculate number of comparisons:
Comparisons = 0;
i = 0;
while i < max(double(SortData.Weight))
    for i = 1:max(double(SortData.Weight))
        Comparisons = Comparisons + (i-1);
    end %for i
end %while i

%determine significant terms
detect1 = 0;
for i = 1:Comparisons
    pcomp = 0.05/(Comparisons-i+1);
    if Sorted1(i) <= pcomp
        Sig1(i) = 1;
        detect1 = 1;
    else
        break
    end
end %i
%Display p values in p matrix that are significant
if detect1 == 1
    fprintf('
    p1 matrix is:
    
    p1.*((p1<=Sorted1(length(Sig1))))
    format short e
    else
    fprintf('
    p1 matrix has no significant values.
    
    end %if

%Layer
%A lot of this code is unnecessary as there is only two layers to 
%compare.
%Number of each Layer:
for i = 1:max(double(SortData.Layer)) %different types
    n2(i) = sum(double(SortData.Layer) == i);
end %i
%Input the number of types, less one (So it doesn’t compare values against themselves)
for i = 1:(max(double(SortData.Layer)-1))
    %Input the total number of types, so it can make all the comparisons
    for j = (i+1):max(double(SortData.Layer))
        %Comparison to Control
        %Note: Accessing the coefficient array must account for weight variables
        if i  == 1
            t2(i,j) = (lm.Coeffici ents.Estimate(max(double(SortData.Weight))-1+j) - 0)/
            sqrt(MSres*(1/n2(j)+1/n2(i))));
        %Between types comparison
        else
            t2(i,j) = (lm.Coefficients.Estimate(max(double(SortData.Weight))-1+j) -
            lm.Coefficients.Estimate(i))/ sqrt(MSres*(1/n2(j)+1/n2(i))));
        end %if i
    end %j
end %i
%Now we take the t values to p values
p2 = 2*(1-tcdf(abs(t2),DF));
%Organize p-values
Sorted2 = sort(reshape(p2,1,size(t2,1)*size(t2,2)));
%Holm test p-value for comparison with uncorrected p-value of p=0.05:
%Calculate number of comparisons:
Comparisons2 = 0;
i = 0;
while i < max(double(SortData.Layer))
    for i = 1:max(double(SortData.Layer))
        Comparisons2 = Comparisons2 + (i-1);
    end %for i
end %while i
% Determine Significant Terms
detect2 = 0;
for i = 1:Comparisons2
    pcomp = 0.05/(Comparisons2-i+1);
    if Sorted2(i) <= pcomp
        Sig2(i) = 1;
        detect2 = 1;
    else
        break
    end %p(i)
end %i
% Display p values in p matrix that are significant
if detect2 == 1
    fprintf('
p2 matrix is:
')
    format short e
    disp(p2.*(p2<=Sorted2(length(Sig2))))
else
    fprintf('
p2 matrix has no significant values.
')
end %if
% Residuals
figure(2)
gscatter(double(SortData.Type),lm.Residuals.Raw,SortData.Flow)
yh = ylabel('Residuals');
xh = xlabel('Types');
legh = legend;
htitle = get(legh,'Title');
set(htitle,'String','Flow (\muL/sec)')
set([xh,yh],'fontweight','bold');
set(gca,'XTick', 1:6)
set(gca,'XTickLabel',{'3 mm 1 Layer', '3 mm 2 Layer', '5 mm 1 Layer', '5 mm 2 Layer', '10 mm 1 Layer', '10 mm 2 Layer'})
% Test Normality
figure(3)
normplot(lm.Residuals.Standardized)
xh = xlabel('Standardized Residuals');
yh = ylabel('Normal Probability Scale');
set([xh,yh],'fontweight','bold');

**Code S4. PS Analysis- Comparison of each type of PS fiber mat**

% Multiple Regression on Polystyrene Data by Type
% Author: Lauren Colangelo and Nicholas Colangelo
% Last Edit: 03-23-2015

% As long as data input into the Table appropriately, everything should be taken care of between data sets except for the following,
% which should be done for each new data set:
% To do:
% 1) Make sure boxplots and other plots represent the data to be analyzed.
% 2) Make sure labels for plots are appropriate.
% 3) Make sure Holm Test is using the appropriate data from the SortData Table.
%Clear memory of variables that may be used here:
clear
%Load the data:
Data = readtable('C:\MatlabPolyStyType.xlsx');
SortData = sortrows(Data,'Flow');
SortData.Type = categorical(SortData.Type,'Ordinal',false);
SortData.Channel = categorical(SortData.Channel,'Ordinal',false);

%Box Plot:
figure(1)
subplot(2,1,1)
boxplot(SortData.Mixing,SortData.Type)
xh = xlabel('Type');
yh = ylabel('Mixing Value');
set([xh,yh],'fontweight','bold');
set(gca,'XTick', 1:7)
set(gca,'XTickLabel',{'Control', '1 Layer 12.5%','1 Layer 15%', '1 Layer 17.5%','2 Layer 12.5%','2 Layer 15%','2 Layer 17.5%'});
subplot(2,1,2)
boxplot(SortData.Mixing,SortData.Flow)
xh = xlabel('Flow (\muL/min)');
yh = ylabel('Mixing Value');
set([xh,yh],'fontweight','bold');
%Regression:
modelspec = 'Mixing ~ Type + Flow';
lm = fitlm(table2dataset(SortData),modelspec);
disp(lm);
%Table of ANOVA Results:
disp(anova(lm));
%Show Number of Outliers as Determined by Cook's Distance >= 3*MeanCooksDistance:
Cook = sum(lm.Diagnostics.CooksDistance./mean(lm.Diagnostics.CooksDistance) >= 3);
fprintf('Number of Values above Cook's Distance %f', Cook);

%Obtain Parameters for Holm Test from the Regression
%TBL(3,3) = MSres
MSres = double(lm.MSE);
%TBL(3,2) = Degrees of Freedom
DF = double(lm.DFE);
%Number of each type:
for i = 1:max(double(SortData.Type)) %different types
    n(i) = sum(double(SortData.Type) == i);
end

%Holm Test
%Input the number of types, less one (So it doesn't compare values against themselves)
for i = 1:(max(double(SortData.Type)-1))
    %Input the total number of types, so it can make all the comparisons
    for j = (i+1):max(double(SortData.Type))
        %Comparison to Control
        if i == 1
            t(i,j) = (lm.Coefficients.Estimate(j) - 0)/ sqrt(MSres*(1/n(j)+1/n(i)));
        else
        %Between types comparison
        end
    end
end
\[
t(i, j) = \frac{(\text{lm.Coefficients.Estimate}(j) - \text{lm.Coefficients.Estimate}(i))}{\sqrt{\text{MSres} \times (1/n(j) + 1/n(i))}};
\]
end \%if i
end \%j
end \%i

% Now we take the t values to p values
p = 2*(1-tcdf(abs(t),DF));

% Organize p-values
Sorted = sort(reshape(p,1,size(t,1)*size(t,2)));

% Holm test p-value for comparison with uncorrected p-value of p=0.05:
% Calculate number of comparisons:
Comparisons = 0;
i = 0;
while i < max(double(SortData.Type))
  for i = 1:max(double(SortData.Type))
    Comparisons = Comparisons + (i-1);
  end \%for i
end \%while i

% Determine Significant Terms
detect = 0;
for i = 1:Comparisons
  pcomp = 0.05/(Comparisons-i+1);
  if Sorted(i) <= pcomp
    Sig(i) = 1;
detect = 1;
  else
    break
  end \%if Sorted
end \%if i

% Display p values in p matrix that are significant
if detect == 1
  fprintf(\n matrix is:\n')
  format short e
  disp(p.*(p<=Sorted(length(Sig))))
else
  fprintf(\n1 matrix has no significant values.\n')
end \%if

% Residuals
figure(2)
gscatter(double(SortData.Type),lm.Residuals.Raw,SortData.Flow)
yh = ylabel('Residuals');
xh = xlabel('Types');
legh = legend;
htitle = get(legh,'Title');
set(htitle,'String','Flow (\muL/min)')
set([xh,yh],'fontweight','bold');
set(gca,'XTick', 1:7)
set(gca,'XTickLabel',{'Control', '1 Layer 12.5%', '1 Layer 15%', '1 Layer 17.5%', '2 Layer 12.5%', '2 Layer 15%', '2 Layer 17.5%'})

% Test Normality with a Normal Probability Plot
figure(3)
normplot(lm.Residuals.Standardized)
xh = xlabel('Standardized Residuals');
yh = ylabel('Normal Probability Scale');
set([xh,yh],’fontweight’,’bold’);