Tuberculosis Detection by Purified Protein Injection Design Description Document

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1 Revision History

<table>
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<tr>
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<th>Description</th>
<th>Date</th>
<th>Authorization</th>
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<td>Initial Draft DDD</td>
<td>23 Jan 2018</td>
<td>All</td>
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<td>B</td>
<td>Revised Draft DDD</td>
<td>7 Feb 2018</td>
<td>All</td>
</tr>
<tr>
<td>C</td>
<td>Revised Draft DDD</td>
<td>21 Feb 2018</td>
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<tr>
<td>D</td>
<td>Midterm Review</td>
<td>26 Feb 2018</td>
<td>All</td>
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<td>E</td>
<td>Revised Review</td>
<td>9 April 2018</td>
<td>All</td>
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<td>F</td>
<td>Final Review</td>
<td>6 May 2018</td>
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2 Background

Tuberculosis is one of the top 10 causes of death worldwide. It is the leading cause of death in patients with HIV. This epidemic is especially prevalent in developing countries where access to testing and treatment is limited. Testing for tuberculosis is necessary because someone can be a carrier but not show signs, nor be contagious until they have immunosuppression. It is best to catch and treat tuberculosis before this point and this cheap testing is the best way to do that.

Latent TB is diagnosed by measuring the reaction of white cells that have responded to the presence of TB bacteria. One of the method is a skin test with PPD, Purified Protein Derivative. A volume of 0.1 ml PPD is placed in the skin of the patient. The reaction size of the skin area is then read after 48 to 72 hours, usually with a ruler to measure the width. This method dates back to 2000 years and often is dependent on the reader’s judgement. The redness on the reaction spot also makes the measurement harder to see with the naked eye.

This senior design project goal is to improve the existing measuring method with optics and maintaining the cost low.

3 Vision Statement

The ideal system has a source and detector connected to a data processing device. The user would place the detector and source above the spot of skin with PPD injection. The device would be able to calculate the surface topography of the bleb after injection, and provide a volume measurement in milliliters. This includes, cross sectional area and height of the bleb. After the patient returns for a follow up meeting 48 hr to 72 hr later, the device would be able to calculate the cross sectional area, most importantly the width perpendicular to the arm, of the physical reaction on the dermis. We are exploring a profilometry method, an approach that meets the basic criteria requested by the customer. This approach explores the physical traits such as area and volume of the skin reaction to the PPD injection. This solution is similar to that of the current medical procedure, but will eliminate the subjectivity of measuring the spot by hand or simply judging size via the human eye.
4 Project Scope

We are responsible for creating a customized software to obtain data from our imaging device

1. A Matlab program that after analyzing images, provides width and volumetric data as well as a 3D model of the bleb or reaction spot for the technician.

2. A proof of concept setup: consisting mainly of a lens system imaging a line pattern and a camera to capture and record the image for processing

3. Explanation of next steps to take in order to achieve portability and easy to use requirements

4.1 Programming Theory

The equations below are from the paper [9.2.1.] and are the baseline for our program. These equations are very standard across many profilometry papers we read. A Matlab program was written to perform fourier transform, and phase unwrapping on the lab images. Numerical data was extracted from the shifts of the line pattern which is caused by the elevation of the bump. With this data, height and width of the reaction bump were calculated.

4.1.1 Forward Data Acquisition

\[
g_o(x, y) = a(x, y) + b(x, y)\cos[2\pi f_o x + \phi_o(x)] \quad (1)
\]

\[
g_s(x, y) = a(x, y) + b(x, y)\cos[2\pi f_o x + \phi(x, y)] \quad (2)
\]

\[
G_s(u, v) = \text{FFT}(g_s); G_o^*(u, v) = \text{FFT}(g_o^*) \quad (3)
\]

\[
\Delta\phi(x, y) = \Im\{\log(\text{FFT}^{-1}[G_s(u, v)G_o^*(u, v)])\} \quad (4)
\]

\[
h(x, y) = \frac{L_o}{2\pi L_o^2 \cos \theta \frac{\Delta\phi(L_o + x \cos \theta \sin \theta)}{P_o(\Delta\phi(L_o + x \cos \theta \sin \theta) + 1)} - \frac{\Delta\phi(L_o + x \cos \theta \sin \theta) + 1}{P_o \cos \theta}} \quad (5)
\]

\[
P_o = \frac{1}{f_o} = \frac{P_o}{\cos \theta} \quad (6)
\]
Figures 1 through 6 visualize the steps our algorithm follows. See the full code in the appendix.

**Figure 1:** Input image with boundary points (red) selected manually

**Figure 2:** Image in Fourier Transform

**Figure 3:** Limited image in Fourier Transform

**Figure 4:** Filtered and shifted Fourier Transform

**Figure 5:** Wrapped phase map

**Figure 6:** Result- A 3D model of the bump from figure 1
4.1.2 Width Measurement and Volume Calculation

Currently able to obtain the length, width and height of measured bump. This does require using human interaction with program to define boundaries. Next steps towards automation of boundary definition could involve training a neural network.

Though we are able to give fairly accurate measurements for the length, width and height of the reaction bump, we are not yet able to give a reliable value for the volume. This is due to the fact that our program cannot yet detect the boundary around the bump. Since reaction bumps are not perfectly round or symmetric, we need information for the edge of the entire bump to deliver accurate volume measurement. Currently, we are only able to manually select four points on the boundary of the bump and use the height at the center intersection, which is not an accurate representation of the volume of the bump. Once the edge detection algorithm is completed, calculating the accurate volume will be possible.

Figure 7: The technician needs to locate 4 points on the boundary of the bleb before the computer runs any calculations.

Figure 8: Best current results of bleb imaging.

Our current results for physical length measurements (completed with a ruler) versus using the program show promising results. The physical measurements should be measured with a more precise tool like calipers.

<table>
<thead>
<tr>
<th>Physical Measurement</th>
<th>Computer Measurement</th>
<th>Percent Error</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.9 mm</td>
<td>5.4 mm</td>
<td>10.2%</td>
</tr>
<tr>
<td>6.3 mm</td>
<td>6.1 mm</td>
<td>3.2%</td>
</tr>
<tr>
<td>9.8 mm</td>
<td>9.7 mm</td>
<td>1.0%</td>
</tr>
<tr>
<td>9.6 mm</td>
<td>10.2 mm</td>
<td>6.3%</td>
</tr>
</tbody>
</table>

4.2 Experimental

4.2.1 Optical System Overview

The light source is collimated by the first lens, and evenly illuminates the grating that is placed in the path of light. Then the grating is imaged by a second lens at two focal lengths away, resulting in a magnification of 1. The image plane is placed perpendicular to the camera to prevent loss of information due to shadows. Two polarizers, placed after the grating and before the camera, improves images quality from light scattering off of the surface of the skin where the bleb would be measured.
4.2.2 Lab Setup

Our final presented system setup has two parallel polarizers [9.2.2] to reduce scatter; the image plane is perpendicular to the CCD instead of the projection system; the lens mounts were made smaller to reduce the angle between the projection and the CCD to 14.7°; and the system was rotated vertically to allow for patients to rest arm on flat surface under CCD.

<table>
<thead>
<tr>
<th>Illumination</th>
<th>System</th>
</tr>
</thead>
<tbody>
<tr>
<td>Collimator Lens Diameter</td>
<td>30 mm</td>
</tr>
<tr>
<td>EFL</td>
<td>60 mm</td>
</tr>
<tr>
<td>F-Number</td>
<td>2</td>
</tr>
<tr>
<td>MTF</td>
<td>0.9 at 10 lp/mm</td>
</tr>
<tr>
<td>Magnification</td>
<td>1X</td>
</tr>
<tr>
<td>System Length</td>
<td>350 mm</td>
</tr>
<tr>
<td>Grating Spacing</td>
<td>2-4 lp/mm</td>
</tr>
<tr>
<td>Light Source</td>
<td>White LED, cell phone flashlight</td>
</tr>
<tr>
<td>CCD Resolution</td>
<td>1MP at 5 µm per pixel</td>
</tr>
</tbody>
</table>

Figure 9: Schematics of the optical projection and detecting system. Grating is projected onto an image plane that is perpendicular to the camera.

Figure 10: A Zemax stimulation of the design. The quality of the image of the grating was analyzed with CodeV. The MTF of the image is 0.5 at 10 lp/mm.

Figure 11: Final presented lab setup.

Figure 12: System from top view with CCD and projection arm.
4.2.3 Environment

As a device intended for clinics, it needs to operate in the following environment:

- Temperature: 59-77 °F – operation range
- Relative Humidity: 40% room temperature
- Outlet/battery power
- Ambient room light

5 In Vivo Testing

We were able to project the grating pattern onto skin and still maintaining fringe contrast despite scattering. A TB team member had a PPD injection from UHS, and attempted to measure the bleb with the designed system. However, due to an unexpected factor of sharing lab equipments, the system needed realignment. Therefore, the bleb dissipated before a measurement took place. Below are earlier images captured of a raised mole on the skin, this gives the same idea as what the injection would have been but only the same color.

Figure 13: This is a mole on one of the teammates under green LED.

Figure 14: This is the same as Figure 13 but under red LED.

Figure 15: Results from the green light mole. This was early in our code and lab setup so it is not as high definition as we have been achieving. However it shows that skin testing does work.
6 Cost Analysis

This is the current cost analysis for what was presented on Design Day. The fourth column is the cost of the item had we not had the benefit of the teaching lab supplies. [9.2.3-4]

<table>
<thead>
<tr>
<th>Items Needed</th>
<th>Item Found</th>
<th>Cost</th>
</tr>
</thead>
<tbody>
<tr>
<td>Visible Light Source</td>
<td>Cell Phone Flashlight</td>
<td>Free -</td>
</tr>
<tr>
<td>Line Grating</td>
<td>Teaching Lab Equipment</td>
<td>Free $150</td>
</tr>
<tr>
<td>Two Lenses</td>
<td>Edmund Optics (60mm FL, 30 mm Dia.)</td>
<td>$30 ea. -</td>
</tr>
<tr>
<td>Camera</td>
<td>The Imaging Source, S/N:DMK31BF03</td>
<td>Free TBD</td>
</tr>
<tr>
<td>Optical Mounts</td>
<td>Teaching Lab Equipment</td>
<td>Free $30 ea.</td>
</tr>
<tr>
<td>Linear Polarizers</td>
<td>Teaching Lab Equipment</td>
<td>Free $10-20 ea.</td>
</tr>
</tbody>
</table>

7 Design Day

Our plan was to present a system that when a created 3D object is placed in the system we will receive a computer 3D model and the volume, height and width measurements. Ideally we would have liked to use skin or chicken skin, but that proved to be difficult with the current equipment we have so we settled for a demo with our calibration bump.

At design day, we presented a vertically aligned prototype system to show proof of concept. People were able to put their arm in the system with a fake* (maybe a different word here) bump placed on their forearm. We captured that image and manually ran it through our program to produce a 3D model as well as the height, length and width of the bump.

We expanded our demo further by explaining our vision for the future of this prototype. We discussed how we used the cell phone light as the light source because our final vision is to have a small system that can be attach to and be used with a cell phone.

Figure 16: Our poster on Design Day May 4, 2018.

Figure 17: Our setup on Design Day May 4, 2018.

Figure 18: Our setup on Design Day May 4, 2018.
8 Future Design Suggestions

**Truncate the System:** One of the concerns of the customer was to have a design that is easily portable as well as small in size. We have worked towards achieving this goal with the reduction of angle between the projection and imaging arm, thus reducing the width of the system. The length of the design was limited by the focal length of the lens, 30 mm. As part of this project, we also created a truncated version of our design system with a goal of maintaining the field of view of an 25 mm object as well as an projected grating image of 1X magnification.

A ThorLab Achromatic Doublet (focal length of 25 mm and a diameter of 12.7 mm) was used to attempt to replicate the performance of the imaging lens, lens between the grating and the image plane. The collimating lens was replaced by an Edmund Optics double-convex lens with a focal length of 20 mm and a diameter of 15 mm. Although the new design's overall length was potentially truncated to 160.5 mm, almost half of the original design, the magnification was lowered to 0.29 to compensate for the 25 mm object field of view.

**Automation of Program:** Ideally the program would be fully automated after an image is taken. In order to achieve this the program needs to be able to detect the boundaries of the bump without interaction. This can be achieved by training a neural network algorithm.

**Cell Phone Attachment:** One purpose of us using the cell phone light was to take the first step in proving this system could be isolated to a cell phone. Using a clip on projection system (to be designed) on the light source, the camera and an app. This would be the ultimate portability method. There is already a small angle between the light source and camera, however this will require some serious engineering and is possibly a project for another senior design team.

**Reaction Spot Database:** This technology would allow for a creation of a database that could give countless examples of reactions that were positive, negative, false-positive or false-negatives. This could lead to further understanding of the development of the disease.
9 Appendix

9.1 Variables

\( \theta \): angle between light source and camera

\( f_0 \): fundamental frequency of observed grating image

\( d \): distance between light source and camera

\( P \): Projected grating period

\( L_o \): length between reference plane and image/object plane

\( G_0^*, G_o^* \): Fourier transforms of original images

9.2 Citations


3. **Camera**: https://www.theimagingsource.com/products/industrial-cameras/firewire-400-monochrome/dmk31bf03/


9.3 Schedule Spring 2018

1. January
   - Create the Matlab Code
   - Collect lab equipments and make customized fringe patterns

2. February
   - Finish the Matlab Code
   - Design a 3D print Bump
   - Run Code in the lab with the experiment set up

3. March
   - Make Code revision
   - IA Presentation
4. April
   - Continue test trial
   - Expand test bump samples
   - Portability & Design Day

5. May
   - Final Document Submission

9.4 Progression of Image Results

Figure 20: Feb 25. Paper. Grating 4 lp/mm. No Polarizers. Image perpendicular to source

Figure 21: February 25. Mole. No polarizers. Grating: 1 lp/mm. Image Plane perpendicular to source.
Figure 22: March 25, 2018. Model Magic White Bump. 5 mm width on white background. Image Plane perpendicular to source. No polarizers.

Figure 23: May 3. Bump 6 mm width on white background. With Polarizers and image plane perpendicular to camera.

Figure 24: Development of lab setup.

9.5 Matlab Code
Contents

- Initialization
- for each quened pixel, its four neighbors are unwrapped.
- Insert a pixel into AQP
- Split the Queue if it is too long
- Copy pixels from APP to AQP if AQP is empty

```matlab
function p = unwrapping(g)

Initialization

[SX SY] = size(g);
MaxQueueSize = SX + SY;  % maximum Queue Size
HalfMaxQueueSize = fix(MaxQueueSize/2);  % half of MaxQueueSize
MinQualityThresh = 0.01;  % Minimum quality threshold
m = ones(SX, SY);

a = abs(g);  % a = a .* mask;
p = angle(g);  % phase to be unwrapped
L = bwlabel(m);
Lnumber = max(max(L));

Unwrapped = zeros(SX, SY);  % to indicate whether a pixel has been unwrapped

Qa = zeros(MaxQueueSize, 1);  % amplitude (array for quened pixels, AQP)
Qx = zeros(MaxQueueSize, 1);  % x coordinate
Qy = zeros(MaxQueueSize, 1);  % y coordinate
Qn = 0;  % number of queued pixels

Pa = zeros(SX*SY, 1);  % amplitude (array for postponed pixels, APP)
Px = zeros(SX*SY, 1);  % x coordinate
Py = zeros(SX*SY, 1);  % y coordinate
Pn = 0;  % number of postponed pixels
Un = 0;  % number of pixels unwrapped

Not enough input arguments.

Error in unwrapping (line 4)
[SX SY] = size(g);

for i = 1:Lnumber
    % find highest quality
    [start_x start_y] = find(a == max(max(a.*(L==i))), 1, 'first');
    % push into AQP
    [Qx Qy Qa Qn] = InsertQueue(Qx, Qy, Qa, Qn, start_x, start_y, a(start_x, start_y));
    Unwrapped(start_x, start_y) = 1;  % seed is taken as unwrapped
    Un = Un + 1;  % update Un
end
```
for each quened pixel, its four neighbors are unwrapped.

```matlab
while Qn>0
    %step 3.1:
    cx=Qx(1);cy=Qy(1);
    Qx(1:Qn-1)=Qx(2:Qn);%delete
    Qy(1:Qn-1)=Qy(2:Qn);%delete
    Qa(1:Qn-1)=Qa(2:Qn);%delete
    Qn=Qn-1;
    %push the left neighbor into the AQP or APP
    if cx-1>0 && Unwrapped(cx-1,cy)==0 && m(cx-1,cy)==1
        %unwrapping the left neighbor
        p(cx-1,cy)=p(cx-1,cy)-round((p(cx-1,cy)-p(cx,cy))/2/pi)*2*pi;
        if a(cx-1,cy)>MinQualityThresh %push into AQP if quality is high
            [Qx Qy Qa Qn]=InsertQueue(Qx,Qy,Qa,Qn,cx-1,cy,a(cx-1,cy));
            if Qn==MaxQueueSize %if AQP reaches preset size, split it
                [Qn,Px,Py,Pa,Pn,MinQualityThresh]=SplitQueue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize);
            end
        else %push into APP if quality is low
            Pn=Pn+1;Px(Pn)=cx-1;Py(Pn)=cy;Pa(Pn)=a(cx-1,cy);
        end
        Unwrapped(cx-1,cy)=1;%mark this pixel as unwrapped.
        Un=Un+1;
    end
    %push the right neighbor into the AQP or APP
    if cx+1<SX+1 && Unwrapped(cx+1,cy)==0 && m(cx+1,cy)==1
        p(cx+1,cy)=p(cx+1,cy)-round((p(cx+1,cy)-p(cx,cy))/2/pi)*2*pi;
        if a(cx+1,cy)>MinQualityThresh %push into AQP if quality is high
            [Qx Qy Qa Qn]=InsertQueue(Qx,Qy,Qa,Qn,cx+1,cy,a(cx+1,cy));
            if Qn==MaxQueueSize %if AQP reaches preset size, split it
                [Qn,Px,Py,Pa,Pn,MinQualityThresh]=SplitQueue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize);
            end
        else %push into APP if quality is low
            Pn=Pn+1;Px(Pn)=cx+1;Py(Pn)=cy;Pa(Pn)=a(cx+1,cy);
        end
        Unwrapped(cx+1,cy)=1;
        Un=Un+1;
    end
    %push the upper neighbor into the AQP or APP
    if cy-1>0 && Unwrapped(cx,cy-1)==0 && m(cx,cy-1)==1
        p(cx,cy-1)=p(cx,cy-1)-round((p(cx,cy-1)-p(cx,cy))/2/pi)*2*pi;
        if a(cx,cy-1)>MinQualityThresh %push into AQP if quality is high
            [Qx Qy Qa Qn]=InsertQueue(Qx,Qy,Qa,Qn,cx,cy-1,a(cx,cy-1));
            if Qn==MaxQueueSize %if AQP reaches preset size, split it
                [Qn,Px,Py,Pa,Pn,MinQualityThresh]=SplitQueue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize);
            end
        else %push into APP if quality is low
            Pn=Pn+1;Px(Pn)=cx;Py(Pn)=cy-1;Pa(Pn)=a(cx,cy-1);
        end
        Unwrapped(cx,cy-1)=1;
        Un=Un+1;
    end
    %push the lower neighbor into the AQP or APP
end
```
if cy+1<SY+1 && Unwrapped(cx,cy+1)==0 && m(cx,cy+1)==1
  p(cx,cy+1)=p(cx,cy+1)-round((p(cx,cy+1)-p(cx,cy))/2/pi)*2*pi;
if a(cx,cy+1)>MinQualityThresh
  [Qx Qy Qa Qn]=InsertQueue(Qx,Qy,Qa,Qn,cx,cy+1,a(cx,cy+1));
  if Qn==MaxQueueSize
    [Qn,Px,Py,Pa,Pn,MinQualityThresh]= ... 
    SplitQueue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize);
  end
else
  Pn=Pn+1;Px(Pn)=cx;Py(Pn)=cy+1;Pa(Pn)=a(cx,cy+1);
end
Unwrapped(cx,cy+1)=1;
Un=Un+1;

%if AQP is empty, copy data from APP
if Qn==0 && Pn>0
  [Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,MinQualityThresh]= ... 
  Copy2Queue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize);
end
end
p=p.*m+(min(min(p.*m))-2*pi).*(1-m);

Insert a pixel into AQP

function [Qx, Qy, Qa, Qn]=InsertQueue(Qx,Qy,Qa,Qn,x,y,a)
I=find(Qa(1:Qn)<a,1,'first'); %find its proper inserting point
if isempty(I) %put in the end of AQP
  Qx(Qn+1)=x;
  Qy(Qn+1)=y;
  Qa(Qn+1)=a;
else %insert into AQP
  Qx(I+1:Qn+1)=Qx(I:Qn);
  Qx(I)=x;
  Qy(I+1:Qn+1)=Qy(I:Qn);
  Qy(I)=y;
  Qa(I+1:Qn+1)=Qa(I:Qn);
  Qa(I)=a;
end
Qn=Qn+1;%update Qn

Split the Queue if it is too long

function [Qn,Px,Py,Pa,Pn,MinQualityThresh]= ... 
  SplitQueue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize)

%put second half of AQP into APP
Pa(Pn+1:Pn+Qn-HalfMaxQueueSize)=Qa(HalfMaxQueueSize+1:Qn);
Px(Pn+1:Pn+HalfMaxQueueSize)=Qx(HalfMaxQueueSize+1:Qn);
Py(Pn+1:Pn+Qn-HalfMaxQueueSize)=Qy(HalfMaxQueueSize+1:Qn);
Pn=Pn+Qn-HalfMaxQueueSize; %update Pn
Qn=HalfMaxQueueSize; %update Qn
MinQualityThresh=Qa(Qn);%Update MinQualityThresh
Copy pixels from APP to AQP if AQP is empty

function [Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,MinQualityThresh]= Copy2Queue(Qx,Qy,Qa,Qn,Px,Py,Pa,Pn,HalfMaxQueueSize)
  Cn=min(Pn,HalfMaxQueueSize);  % number of pixel to be copied
  [temp I]=sort(Pa(1:Pn),'descend');  % sort APP and store in 'temp'
  Qa(1:Cn)=temp(1:Cn);  % copy to AQP
  Qx(1:Cn)=Px(I(1:Cn));
  Qy(1:Cn)=Py(I(1:Cn));
  Qn=Cn;  % update Qn
  MinQualityThresh=Qa(Qn);  % update MinQualityThresh
  Pa(1:Pn-Cn)=temp(Cn+1:Pn);  % arrange APP
  Px(1:Pn-Cn)=Px(I(Cn+1:Pn));
  Py(1:Pn-Cn)=Py(I(Cn+1:Pn));
  Pn=Pn-Cn;  % update Pn
Contents

- Clean all the setting
- 1. Load the image
- Distance
  - 3. Filtering the spectrum centered around fo and translating to the origin results in
  - 4. Phase Unwrapping using quality guided Flood Filling algorithm
  - 5. 3D calibration
  - Show the highest and lowest point in this 3D model

Clean all the setting

clear; clc;close all;

1. Load the image

```matlab
%image = imread('coin111.tif'); % read an image
%image = imread('sample1.jpeg'); % read an image
%image = imread('gl_2.png'); % read an image
%image = imread('r13.png'); % read an image
%image = imread('b&w.png'); % read an image
%image = imread('MM1.bmp'); % read an image
image = imread('biggerbumponwhite.bmp'); % read an image

%image = imread('1.png'); % read an image
if(size(image,3)>1) % Color to Black&white
    image=rgb2gray(image);
end
image=im2double(image); % Convert Image to Double Precision
imshow(image);title('1.Input Image') % display the original image
```

Distance

```matlab
hold on;
xy = [];
n = 0;
distance = [];
while n < 4
    [xi,yi] = ginput(1);
    plot(xi,yi,'ro')
    n = n+1;
    xy(:,n) = [xi;yi];
end
%disp(xy(1,1));
%disp(xy(1,2));
%disp(xy(2,1));
%disp(xy(2,2));
disp('Pick 2 points for width and then 2 for length');
C_d = 16410;
d1= sqrt((xy(2,2)-xy(1,1))^2+(xy(1,2)-xy(2,1))^2);
d1= d1 / C_d;
d2= sqrt((xy(2,4)-xy(1,3))^2+(xy(2,3)-xy(1,4))^2);
```
Pick 2 points for width and then 2 for length
The width is: (m)
0.0066
The length is: (m)
0.0069


```matlab
image_fft = fftshift(fft2(image));  %% // https://www.mathworks.com/help/matlab/ref/fft2.html
figure; imagesc(log(abs(image_fft))); %%// https://www.mathworks.com/help/matlab/ref/imagesc.html
title('2.Image in Fourier Transform');
index = zeros(size(image_fft));
[x, y] = find(abs(image_fft)==max(max(abs(image_fft))));
image_fft(x-15:x+15,y-18:y+18) = 0; %% select area with most information and least noise
figure; imagesc(log(abs(image_fft)));
title('3.Limited Image in Fourier Transform');
```

3. Filtering the spectrum centered around fo and translating to the origin results in

```matlab
[sax, sy] = find(abs(image_fft)==max(max(abs(image_fft))));
index(sax-15:sax+15,sy-18:sy+18) = 1;
image_fft_bpf=image_fft.*index;
shift_x = x - sax;
shift_y = y - sy;
image_fft_bpf = circshift(image_fft_bpf,[shift_x shift_y]);%% https://www.mathworks.com/help/matlab/ref/circshift.html
figure; imagesc(log(abs(image_fft_bpf))); title('4.Filtered and shifted fourier');
image_rec = ifft2(fftshift(image_fft_bpf));  %% Inverse transformation to the image
phi = atan(imag(image_rec)./real(image_rec));
figure; imagesc(phi);title('5.Wrapped phase map');colormap gray;
```

4. Phase Unwrapping using quality guided Flood Filling algorithm

```matlab
image_unwrp = unwrapping(image_rec);
C = 8.5*10e-6;  %% pixel to meter constant
image_unwrp = image_unwrp * 8.5*10e-6;
```

5. 3D calibration

```matlab
figure;imagesc(image_unwrp);
figure;mesh(image_unwrp);  %% https://www.mathworks.com/help/matlab/ref/mesh.html
ax1 = gca;
ax1.XColor = 'r';
ax1.YColor = 'r';
```
ax1_pos = ax1.Position; % position of first axes
ax2 = axes('Position',ax1_pos,'XAxisLocation','bot','YAxisLocation','left','Color','none');
ax2 = axes('Position',[0.1 0.1 0.8 0.001],'Color','none')
first_axis = gca;
sqz = 0.12; % distance to squeeze the first plot
set(first_axis, 'Position', get(first_axis, 'Position') + [0 sqz 0 -sqz ]);
ax2 = axes('Position', get(first_axis, 'Position') .* [1 1 1 0.001] - [0 sqz 0 0], 'Color','none');
scale_factor = 8.5*10e-6; % change this to your satisfaction
xlim(get(first_axis, 'XLim') * scale_factor);
set(ax2, 'XScale', get(first_axis, 'XScale'));

Show the highest and lowest point in this 3D model

disp(max(max(image_unwrp)));
disp(min(min(image_unwrp)));