```
In[@]:= ClearAll
Out[@]=
ClearAll
```

biological parameters

```
ln[ \circ ] := C = 299792458;
         hb = 1.0545715964207855 * 10^-34;
         \epsilon 0 = 8.85 * 10^{-12};
         re = 2.82 * 10^{-13} * 10^{-2};
         q = 1.602 * 10^{-19};
 In[ \circ ] := Wsig = 9 * 10^3 * q / hb;
 In[*]:= λsig = 2 * Pi * c / wsig;
 ln[-]:= f1C = 6.01; f1N = 7.02; f1H = 1; f1O = 8.04; f1S = 16.27; f1CI = 17.31;
 ln[a]:=AC=12.011; AN=14.007; AH=1.008; AO=15.999; AS=32.06; ACI=35.45;
         NA = 6.022 * 10^23;
         cmCubeToMeterCube = 100^3;
 ln[a]:= phase[sH_, sC_, sN_, sO_, sS_, sCI_, thickness_, materialDensity_, \lambda_{-}] :=
           \left(\frac{2\pi}{\lambda}\right) * thickness * \left(\frac{\text{re}}{2\pi}\right) * materialDensity * NA *
             \left(\frac{\bot}{\mathsf{SH} * \mathsf{AH} + \mathsf{SC} * \mathsf{AC} + \mathsf{SN} * \mathsf{AN} + \mathsf{SO} * \mathsf{AO} + \mathsf{SS} * \mathsf{AS} + \mathsf{SCI} * \mathsf{ACI}}\right) *
             (SH * f1H + SC * f1C + SN * f1N + SO * f1O + SS * f1S + SCI * f1CI) * \lambda^2 * cmCubeToMeterCube
 ln[e]:= phase [48.6, 32.9, 8.9, 8.9, 0.6, 0, 5 * 10^-9, 1.35, \lambdasig]
Out[0]=
         0.000843499
```

cropping functions

```
In[@]:= Clear[croping]
     croping[biosample_] := ImageData[
       ImageTake[Image[biosample], {Dimensions[biosample] [1]] - upperight[biosample] [1]] + 1,
          Dimensions[biosample] [1] - lowerleft[biosample] [1] + 1},
         {lowerleft[biosample] [2], upperight[biosample] [2]}}]
```

uploading the biological features and assigning them the biological data

upload the folded protein pattern

```
In[@]:= Clear[foldedProtein]
       foldedProtein =
          Import["G:\\Haim\\SU 11 project\\papers\\Imaging\\My paper\\codes\\biological
             imaging\\biological features\\foldedProtein.JPG"];
 In[*]:= temp = foldedProtein;
       foldedProtein = ImageResize[temp, 150] (*ImageResize[temp, 150] *)
Out[\circ] =
 In[@]:= Clear[temp]
       temp = foldedProtein;
       foldedProtein = ImageData[ColorConvert[temp, "Grayscale"]];
 In[*]:= Image[foldedProtein]
Out[0]=
```

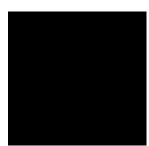


```
In[@]:= Clear[temp]
       temp = croping[foldedProtein];
       Clear[foldedProtein]
       foldedProtein =
          Table [temp[i, j]], \{i, 1, Dimensions [temp][1]]\}, \{j, 1, Dimensions [temp][2]]\}];\\
 In[*]:= Image[foldedProtein]
Out[0]=
```



$assign corresponding {\it refractive} coefficient {\it s} to the folded protein$ pattern

```
In[@]:= Clear[temp]
       temp = foldedProtein;
       biosample01 = temp * phase [48.6, 32.9, 8.9, 8.9, 0.6, 0, (5 * 5 / 2) * 10^-9, 1.35, \lambda sig];
       Image[biosample01]
       Image[biosample01 / Max[biosample01]]
Out[@]=
```



Out[0]=



```
(*1. Import, Resize, and Convert to Grayscale*)
      ClearAll[foldedProtein];
      SetDirectory["G:\\Haim\\SU 11 project\\papers\\Imaging\\My
           paper\\codes\\biological imaging\\biological features"];
       (*Import the image and resize it to 150 pixels width*)
       foldedProtein = Import["foldedProtein.JPG"] // ImageResize[#, 150] &;
       (*Convert to Grayscale and extract numerical ImageData for mathematical processing*)
      foldedProteinData = ImageData[ColorConvert[foldedProtein, "Grayscale"]];
       (*2. Dimension Checks (if needed for subsequent steps)*)
       Print["Dimensions: ", Dimensions[foldedProteinData]];
       (*Example:{143,150}*)
       (*3. Cropping*)
       (*Apply the pre-defined'croping' function and update the main variable*)
      foldedProteinData = croping[foldedProteinData];
       (*Print["Cropped Dimensions: ",Dimensions[foldedProteinData]];
      Image[foldedProteinData] (*Display the cropped image*)
       (*4. Create the Bio Sample*)
       (*Multiply the image data by the pre-defined'phase' function*)
       biosample01 =
         foldedProteinData * phase [48.6, 32.9, 8.9, 8.9, 0.6, 0, (5 * 5 / 2) * 10^-9, 1.35, \lambda sig];
       (*Display the processed sample image*)
      Image[biosample01]
       (*Display the normalized image for better visualization*)
       Image[biosample01 / Max[biosample01]]
 upload the small protein pattern
 In[*]:= Clear[twoSmallProteins]
      twoSmallProteins =
         Import["G:\\Haim\\SU 11 project\\papers\\Imaging\\My paper\\codes\\biological
            imaging\\biological features\\twoSmallProteins.JPG"];
 In[*]:= Clear[temp]
      temp = twoSmallProteins;
      twoSmallProteins = ImageResize[temp, 150] (*ImageResize[temp,150]*)
Out[0]=
```

```
In[@]:= Clear[temp]
       temp = twoSmallProteins;
       twoSmallProteins = ImageData[ColorConvert[temp, "Grayscale"]];
 In[@]:= Image[twoSmallProteins]
Out[0]=
 In[@]:= Clear[temp]
       temp = croping[twoSmallProteins];
       Clear[twoSmallProteins]
       twoSmallProteins =
          Table [temp [\![i,j]\!], \{i,1,Dimensions [temp] [\![1]\!]\}, \{j,1,Dimensions [temp] [\![2]\!]\}];
 In[@]:= Image[twoSmallProteins]
Out[0]=
```

$assign corresponding {\it refractive} coefficient {\it stothesmall} protein$ pattern

```
In[*]:= Clear[temp]
       temp = twoSmallProteins;
       biosample02 = temp * phase [48.6, 32.9, 8.9, 8.9, 0.6, 0, (3 * 5 / 2) * 10^-9, 1.35, \lambda sig];
       Image[biosample02]
       Image[biosample02 / Max[biosample02]]
Out[0]=
Out[0]=
 upload the large protein pattern
 In[*]:= Clear[twoLargeProteins]
       twoLargeProteins =
         Import["G:\\Haim\\SU 11 project\\papers\\Imaging\\My paper\\codes\\biological
             imaging\\biological features\\twoLargeProteins.JPG"];
 In[@]:= Clear[temp]
       temp = twoLargeProteins;
       twoLargeProteins = ImageResize[temp, 150] (*ImageResize[temp,150]*)
Out[0]=
```

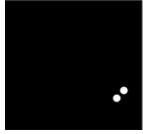
```
In[@]:= Clear[temp]
     temp = twoLargeProteins;
     twoLargeProteins = ImageData[ColorConvert[temp, "Grayscale"]];
```

```
In[*]:= Image[twoLargeProteins]
```

Out[@]=



```
In[*]:= Clear[temp]
       temp = croping[twoLargeProteins];
       Clear[twoLargeProteins]
       twoLargeProteins =
         Table [temp[i, j]], \{i, 1, Dimensions [temp][1]]\}, \{j, 1, Dimensions [temp][2]]\}]; \\
 In[@]:= Image[twoLargeProteins]
Out[0]=
```



assign corresponding refractive coefficients to the large protein pattern

```
In[@]:= Clear[temp]
       temp = twoLargeProteins;
       biosample03 = temp * phase [48.6, 32.9, 8.9, 8.9, 0.6, 0, (5 * 5 / 2) * 10^-9, 1.35, \lambda sig];
       Image[biosample03]
       Image[biosample03 / Max[biosample03]]
Out[0]=
Out[0]=
```

$assign corresponding {\it refractive} coefficient {\it stothelipid layer}$

```
In[ • ]: = lipid =
           Table[phase[62.5, 31.5, 0, 6.3, 0, 0, 45 * 10^-9, 1, \lambda sig], {i, 1, 139}, {j, 1, 144}];
 In[*]:= Image[lipid]
Out[0]=
```

 $assign corresponding {\it refractive} coefficient {\it s} to the lice cube$

```
ln[*]:= water = Table[phase[2, 0, 0, 1, 0, 0, 5 * 10^-6, 0.92, \lambdasig], {i, 1, 139}, {j, 1, 144}];
```

In[@]:= Image[water] Out[0]=

creating the total biological sample

```
In[@]:= Clear[totalbiosample]
       totalbiosample = biosample01 + biosample02 + biosample03 + lipid(*+water*);
 In[@]:= Clear[totalbiosample]
       totalbiosample = ImageData[%292];
       Image[totalbiosample / Max[totalbiosample]]
Out[0]=
 In[@]:= Image[1 - (totalbiosample / Max[totalbiosample])]
Out[0]=
 In[*]:= Clear[totalimageBG]
       totalimageBG = Table[phase[2, 0, 0, 1, 0, 0, 5 * 10^-6, 0.92, \lambda sig],
           {i, 1, Dimensions[totalbiosample] [1]]}, {j, 1, Dimensions[totalbiosample] [2]}}
        (* this is for 5 micron water *);
 In[@]:= Clear[temp]
       temp = totalbiosample;
       totalbiosample = temp + totalimageBG;
```

In[*]:= Image[totalbiosample]

Out[•]=



In[@]:= totalbiosample

Out[@]=

```
\{ \{ 0.605691, 0.605691, 0.605691, 0.605691, \cdots 126\cdots \}
  0.605691, 0.605691, 0.605691, 0.605691}, ....132...., \{ \dots 1 \dots 1 \} 
large output
              show less
                           show more
                                         show all
                                                    set size limit...
```

In[*]:= Image[totalbiosample / Max[totalbiosample]]

Out[@]=

In[@]:= Image[1 - (totalbiosample / Max[totalbiosample])]

Out[@]=



creating a matrix for the signal ("S") angular deviations

```
the object pixel size must be smaller than the imaging resolution →
  \frac{\text{FOV\_sig}}{\text{minimal resolution}} = \frac{10^{-6}}{10^{-9}} = 10^{3} < \text{number of object 1D pixels}
  ln[\circ] := \sigma\Theta p0 = 4.13 * 10^-6
          (* \sigma\Thetap should be of the order of 0.77*objXpixel to achieve the Rayleigh limit *);
        \Thetap0 = -1.1556175944769262`;
        ks = 4.559401012488846`*^10;
        kp = 4.560414567875652 *^10;
        ki = 2.446282748793892 *^7;
 In[.] = fs = 0.003;
 In[*]:= res = _____
Out[0]=
        \textbf{4.99866}\times\textbf{10}^{-9}
 ln[*]:= pixelsize = 0.5 * res (* the smallest separation in my image is 2 pixels than
            we need to multiply by 0.5 for that separation to correspond to 5 nm *)
Out[0]=
        2.49933 \times 10^{-9}
 in[*]:= angularPixel = 0.5 * res / fs
Out[0]=
        8.3311 \times 10^{-7}
 In[*]:= 2 * angularPixel / σθρ0
Out[0]=
        0.403443
 In[@]:= angularPixel * (Dimensions[totalbiosample] [1])
Out[0]=
        0.000111637
 In[\bullet]:=\delta S\Theta xmin = -angularPixel * (Dimensions[totalbiosample][1]) / 2
        \delta s \theta x max = angular Pixel * (Dimensions [totalbiosample] [1]) / 2
Out[0]=
        -0.0000558184
Out[0]=
        0.0000558184
 In[*]:= ScientificForm[δsθxmax]
Out[@]//ScientificForm=
        5.58184 \times 10^{-5}
```

```
In[\bullet]:=\delta S\Theta ymin = -angularPixel * (Dimensions[totalbiosample][2]) / 2
          δsθymax = angularPixel * (Dimensions[totalbiosample] [2]) / 2
Out[0]=
          -0.0000558184
Out[0]=
          0.0000558184
  in[*]:= jmax = Dimensions[totalbiosample] [2];
          imax = Dimensions[totalbiosample] [2];
          imin = 1;
  In[\bullet]:= (\delta s \theta x max - \delta s \theta x min) / (imax - imin + 1)
Out[0]=
          8.3311 \times 10^{-7}
  ln[\bullet]:= objXpixel = (\delta s \theta x max - \delta s \theta x min) / (imax - imin + 1)
             (* this should TURN OUT TO BE EQUAL to angularPixel *)
Out[0]=
          \textbf{8.3311}\times\textbf{10}^{-7}
  In[@]:= N[objXpixel]
Out[0]=
          \textbf{8.3311}\times\textbf{10}^{-7}
  In[\sigma]:= objYpixel = (\delta s \theta y max - \delta s \theta y min) / (jmax - jmin + 1)
             (* this should TURN OUT TO BE EQUAL to angularPixel *)
Out[0]=
          \textbf{8.3311}\times\textbf{10}^{-7}
  In[@]:= N[objYpixel]
Out[0]=
          8.3311 \times 10^{-7}
  ln[*]:= \delta s\theta x = Range[\delta s\theta xmin * 0.995, \delta s\theta xmax * 0.995, objXpixel];
         Dimensions [\delta s \theta x]
Out[0]=
          {134}
  In[\bullet]:=\delta s\theta y = Range[\delta s\theta ymin * 0.995, \delta s\theta ymax * 0.995, objYpixel];
         Dimensions [\delta s \theta y]
Out[0]=
          {134}
  ln[a] := N[129 * (5 * 10^-9) / 2] (* this is the FOV of this image in meters *)
Out[0]=
          3.225 \times 10^{-7}
```

creating a matrix for the idelr ("I") angular

deviations

```
In[*]:= δiθxmin = δsθxmin * ks / ki
           \delta i\theta x max = \delta s\theta x max * ks / ki
           \delta i \theta y min = \delta s \theta y min * ks / ki
           \delta i\Theta y max = \delta s\Theta y max * ks / ki
Out[0]=
           -0.104035
Out[0]=
           0.104035
Out[0]=
           -0.104035
Out[0]=
           0.104035
  In[\bullet]:= imageXpixel = (\delta i\theta x max - \delta i\theta x min) / (imax - imin + 1)
Out[0]=
           0.00155276
  ln[\bullet]:=\delta i\theta x = Range[\delta i\theta x min * 0.995, \delta i\theta x max * 0.995, imageXpixel];
           Dimensions [\delta i\theta x]
Out[0]=
           {134}
  In[\bullet]:= imageYpixel = (\delta i\theta ymax - \delta i\theta ymin) / (jmax - jmin + 1)
Out[0]=
           0.00155276
  ln[\bullet]:=\delta i\theta y = Range[\delta i\theta ymin * 0.995, \delta i\theta ymax * 0.995, imageYpixel];
           Dimensions [\delta i\theta y]
Out[0]=
           {134}
  In[\bullet]:= Dimensions [\delta i\theta y] [1]
Out[0]=
           134
```

writing the input pump function

writing the matrix of the idler angular deviations based on the pump beam width

```
In[@]:= Clear[func]
                                                                                                                                                                                                                                                                                         \underline{\left(\left(\texttt{ki}\,\delta \mathsf{i}\theta \mathsf{x}\, [\![m]\!] + \mathsf{ks}\,\delta \mathsf{s}\theta \mathsf{x}\, [\![i]\!]\right)^2 + \left(\texttt{ki}\,\delta \mathsf{i}\theta \mathsf{y}\, [\![n]\!] + \mathsf{ks}\,\delta \mathsf{s}\theta \mathsf{y}\, [\![j]\!]\right)^2\right)\,\mathsf{Sec}\left[\theta \mathsf{p}\theta\right]^2}
                             func [\sigma \Theta p_{j}, i_{j}, m_{j}, m_{j}] := (*\frac{1}{\sqrt{2 \times \pi} \sigma \Theta p} **) e^{i}
```

writing the matrix of the idler image

```
part 1:
```

```
In[@]:= (* nij = (#nPDC per unit area)*(area of the ENTIRE object) *)
  In[\bullet]:= Dimensions [\delta i\theta y]
Out[0]=
            {134}
  In[@]:= Clear[normalization]
            normalization[\sigma\Thetap_] := ParallelTable[
                 Sum[func[\sigma\theta p, i, j, m, n], \{i, 1, Dimensions[\delta s\theta x][1]]\}, \{j, 1, Dimensions[\delta s\theta y][1]]\}], \{i, 1, Dimensions[\delta s\theta y][1]]\}], \{i, 1, Dimensions[\delta s\theta y][1]]\}
                  {m, 1, Dimensions [\delta i\theta x] [1]}, {n, 1, Dimensions [\delta i\theta y] [1]}];
  In[*]:= Clear[sf]
            sf[\sigma\theta p_{-}] := (1 / normalization[\sigma\theta p_{-}]) *
                 ParallelTable \left[ \text{Sum} \left[ (*2*\frac{\text{nPDC}}{\text{objarea}} **) \text{Sin} \left[ \text{totalbiosample} \right] \right] * \text{func} \left[ \sigma \theta \text{p, i, j, m, n} \right] \right]
                     {i, 1, Dimensions [\delta s \theta x] [1]}, {j, 1, Dimensions [\delta s \theta y] [1]},
                    {m, 1, Dimensions [\delta i \theta x] [1]}, {n, 1, Dimensions [\delta i \theta y] [1]};
  In[@]:= Image[Abs[1 - totalimageBG]]
            Image[Abs[totalimageBG]]
            Clear[sbg]
            sbg[\sigma\theta p_{]} := (1 / normalization[\sigma\theta p]) *
                 ParallelTable \left[ \text{Sum} \left[ (*2*\frac{\text{nPDC}}{\text{objarea}} **) \text{Sin} \left[ \text{totalimageBG[[i, j]]} \right] * \text{func} \left[ \sigma \theta \text{p, i, j, m, n} \right] \right] \right]
                     {i, 1, Dimensions [\delta s \theta x] [1]}, {j, 1, Dimensions [\delta s \theta y] [1]},
                    {m, 1, Dimensions [\delta i \theta x] [1]}, {n, 1, Dimensions [\delta i \theta y] [1]};
Out[0]=
Out[0]=
```

```
In[*]:= Clear[add\total]
          add∆total[q_, n1_, objarea_] :=
             ParallelTable [RandomVariate [NormalDistribution [0, Sqrt [2*(1+q)*\frac{n1}{objarea}]]]]
                {m, 1, Dimensions [\delta i\theta x] [1]}, {n, 1, Dimensions [\delta i\theta y] [1]} \Big| (* e^{-(x-\mu)^2/(2\sigma^2)},
            \textbf{x} \rightarrow \textbf{noise}_{sf}[[\texttt{m},\texttt{n}]], \ \mu[[\texttt{m},\texttt{n}]] = \textbf{0}, \ \sigma^2 = \Delta 2sf[[\texttt{m},\texttt{n}]] + \Delta 2sbg[[\texttt{m},\texttt{n}]] = 2*\frac{nPDC}{objarea} \ + 2*\frac{nPDC}{objarea} \ *); 
part 2:
In[*]:= Clear[Sidler]
          Sidler[\sigma\Thetap_] := sf[\sigma\Thetap] - sbg[\sigma\Thetap];
```

choosing Region Of Interest (ROI):

```
In[\bullet]:= SidlerN\sigma\Thetap0 = Sidler[\sigma\Thetap0];
 In[*]:= Image[SidlerNσθp0 / Max[Image[SidlerNσθp0]]]
Out[0]=
 In[*]:= Clear[SidlerROIn]
         SidlerROIn[i1_, i2_, j1_, j2_] :=
          ImageData[ImageTake[Image[SidlerN\sigma\Thetap0(*/Max[SidlerN\sigma\Thetap0]*)],
             {Dimensions [\delta i\theta x] [1] - i2 + 1, Dimensions [\delta i\theta x] [1] - i1 + 1},
             {Dimensions [\delta i\theta y] [1] - j2 + 1, Dimensions [\delta i\theta y] [1] - j1 + 1}]
 In[@]:= Clear[add\totalROIn]
         add∆totalROIn[q_, n1_, i1_, i2_, j1_, j2_] :=
          ImageData[ImageTake[Image[add∆total[q, n1, 1]]],
             {Dimensions [\delta i\theta x] [1] - i2 + 1, Dimensions [\delta i\theta x] [1] - i1 + 1},
             {Dimensions [\delta i\theta y] [1] - j2 + 1, Dimensions [\delta i\theta y] [1] - j1 + 1}]
 Indes: Image [ (totalbiosample - totalimageBG) / Max[totalbiosample - totalimageBG] ]
Out[0]=
```

```
In[@]:= Clear[i1Folded, i2Folded, j1Folded, j2Folded, rowMinFolded, rowMaxFolded]
 In[@]:= rowMinFolded = 69;
       rowMaxFolded = 134;
       i1Folded = Dimensions [\delta i\theta x] [1] - rowMaxFolded + 1
       i2Folded = Dimensions [\delta i\theta x] [1] - rowMinFolded + 1
       j1Folded = 24; (* the most left column *)
 In[*]:= Clear[foldedProteinData]
       foldedProteinData = ImageData[ImageTake[Image[(totalbiosample - totalimageBG)],
            {i1Folded, i2Folded}, {j1Folded, j2Folded}]];
       Image[foldedProteinData / Max[foldedProteinData]]
Out[0]=
```

Fourier transforming the folded protein

```
In[@]:= Clear[SidlerROInFolded]
       SidlerROInFolded = SidlerROIn[i1Folded, i2Folded, j1Folded, j2Folded];
       Clear[qj]
       qj = Table[q, {q, {1, 5, 25}}];
       Clear[n1]
       n1 = Table[NPDC, {NPDC, 10^5, 1 * 10^8, 1.999 * 10^6}];
 In[*]:= Clear[a1folded]
       alfolded = ParallelTable[Fourier[2 * n1[j]] * Sqrt[qj[i]]] * SidlerROInFolded +
            add∆totalROIn[qj[i], n1[j], i1Folded, i2Folded, j1Folded, j2Folded],
           FourierParameters \rightarrow {1, 1}], {i, 1, Dimensions[qj][1]}, {j, 1, Dimensions[n1][1]}]
Out[0]=
                                                     set size limit...
                     show less
                                            show all
         large output
                                show more
```

```
in[o]:= a1folded = ParallelTable[RotateRight[a1folded[i]][j]], Floor[
             ({Dimensions[foldedProteinData][2], Dimensions[foldedProteinData][1]} - 1) / 2]],
          \{i, 1, Dimensions[qj][1]\}, \{j, 1, Dimensions[n1][1]\}\}
Out[\circ] =
                                              show all
                                                        set size limit...
          large output
                       show less
                                  show more
 In[*]:= Clear[a2folded]
        a2folded = ParallelTable[Fourier[2 * n1[j]] * Sqrt[qj[i]]] * SidlerROInFolded +
             add∆totalROIn[qj[i], n1[j], i1Folded, i2Folded, j1Folded, j2Folded],
            FourierParameters \rightarrow {1, 1}], {i, 1, Dimensions[qj][1]}, {j, 1, Dimensions[n1][1]}]
Out[\circ] =
          large output
                       show less
                                  show more
                                              show all
                                                        set size limit..
 In[@]:= a2folded = ParallelTable[RotateRight[a2folded[i]][j]], Floor[
             ({Dimensions[foldedProteinData][2], Dimensions[foldedProteinData][1]} - 1) / 2]],
          {i, 1, Dimensions[qj][1]}, {j, 1, Dimensions[n1][1]}}
Out[0]=
                       show less
                                  show more
                                              show all
                                                        set size limit...
          large output
```

creating a search ring filter in the Fourier space

```
In[@]:= Clear[sign]
     sign[i_, j_, min_, max_] := If[(i - (Floor[Dimensions[foldedProteinData][1] / 2]))^2 +
            (j - (Floor[Dimensions[foldedProteinData][2] / 2]))^2 ≥ min^2&&
         (i - (Floor[Dimensions[foldedProteinData][2] / 2]))^2 +
            (j - (Floor[Dimensions[foldedProteinData][2]/2]))^2 \le max^2, 1, 0]
In[*]:= Clear[pp]
     pp = Table[0, {i, 1, Dimensions[foldedProteinData][1]]},
         {j, 1, Dimensions[foldedProteinData][2]]}];
     For[i = 1, i ≤ Dimensions[foldedProteinData][1],
      i++, For [j = 1, j \le Dimensions [foldedProteinData] [2],
        j++, pp[[i, j]] = (*ppp[[i,j]]**)sign[i, j, 1, 4]]]
```

Out[0]=

Out[0]=

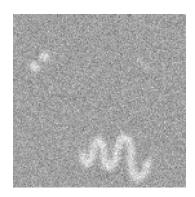
```
In[@]:= Image[pp]
Out[0]=
 In[*]:= Clear[numpix, halfBit]
 In[@]:= numpix[min_, max_] := Floor[Pi * (max^2 - min^2)];
        halfBit[min_, max_] :=
           (0.2071 + 1.9102 / (\sqrt{\text{numpix}[\min, max]})) / (1.2071 + 0.97102 / (\sqrt{\text{numpix}[\min, max]}));
```

plotting at different exposures

{min, 1, Floor[Dimensions[foldedProteinData][2] / 2]}, $\{\Delta r, 1, 4\}$];

 $halfBitTable = Table[halfBit[min, min + \Delta r],$

```
In[*]:= NPDCs = Table[NPDC, {NPDC, 5 * 10^6, 5 * 10^8, 7.07143 * 10^7}];
In[\sigma]:= image1 = NPDCs[1] * SidlerN\sigma\Thetap0 + add\DeltatotalROIn[NPDCs[1]],
            1, Dimensions [SidlerN\sigma\Thetap0] [1], 1, Dimensions [SidlerN\sigma\Thetap0] [2]];
      Image[image1 / Max[image1]]
```



```
In[\bullet]:= image4 = NPDCs[4] * SidlerN\sigma\Thetap0 + add\DeltatotalROIn[NPDCs[4]],
             1, Dimensions [SidlerN\sigma\thetap0] [1], 1, Dimensions [SidlerN\sigma\thetap0] [2]];
       Image[image4 / Max[image4]]
```

```
In[\bullet]:= image7 = NPDCs[[7]] * SidlerN\sigma\Thetap0 + add\DeltatotalROIn[NPDCs[[7]],
               1, Dimensions [SidlerN\sigma\Thetap0] [1], 1, Dimensions [SidlerN\sigma\Thetap0] [2]];
         Image[image7 / Max[image7]]
Out[0]=
```

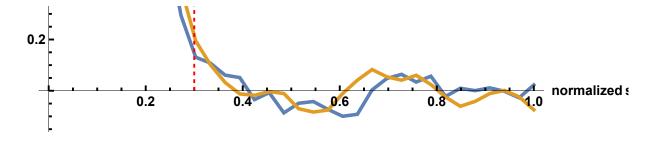
showing the ROI image of the the proteins:

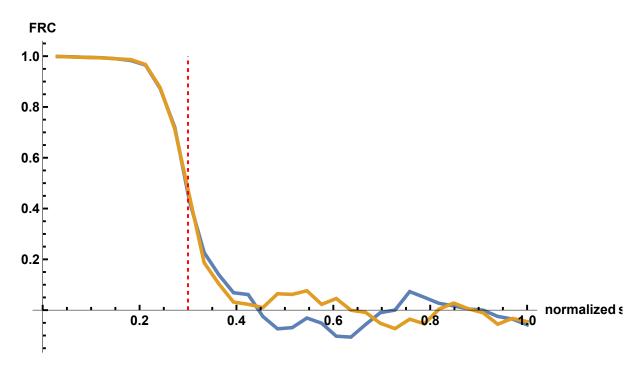
```
in[o]:= Image[SidlerROIn[i1Folded, i2Folded, j1Folded, j2Folded] /
         Max[SidlerROIn[i1Folded, i2Folded, j1Folded, j2Folded]]]
Out[0]=
```

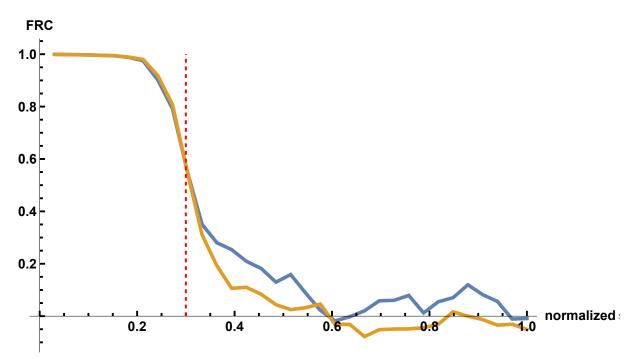
FRC of the folded protein:

```
Clear[p1, p2]
p1 = \{50, 29, 8\};
p2 = {30, 18, 5};
Clear[∆r]
\Delta r = 3;
```

```
Table Clear[frcDatan1, frcDatanq];
 frcDatan1 = ParallelTable[
    (Sum[Re[a1folded[1][p1[k1]][i, j] * Conjugate[a2folded[1][p1[k1]][i, j]] *
           sign[i, j, min, min + \Delta r]], \{i, 1, Dimensions[foldedProteinData][1]]\},
        {j, 1, Dimensions[foldedProteinData][2]]}]) /
      (\mathsf{Sqrt}[(\mathsf{Sum}[\mathsf{sign}[\mathsf{i},\mathsf{j},\mathsf{min},\mathsf{min}+\Delta r]*(\mathsf{Abs}[\mathsf{alfolded}[\![1]\!][\![\mathsf{pl}[\![k1]\!]]\![\![\mathsf{i},\mathsf{j}]\!])^2, \\
            {i, 1, Dimensions[foldedProteinData] [1]]},
            {j, 1, Dimensions[foldedProteinData][2]]}) *
         (Sum[sign[i, j, min, min + \Delta r] * (Abs[a2folded[1][p1[k1]][i, j]])^2,
            {i, 1, Dimensions[foldedProteinData] [1]]},
            {j, 1, Dimensions[foldedProteinData][2]}])]),
    {min, 1, Floor[Dimensions[foldedProteinData] [2] / 2] }];
 frcDatanq = ParallelTable[
    (Sum[Re[a1folded[2][p2[k2]][i, j] * Conjugate[a2folded[2][p2[k2]][i, j]] *
           sign[i, j, min, min + \Delta r]], \{i, 1, Dimensions[foldedProteinData][1]\},
        {j, 1, Dimensions[foldedProteinData] [2]}}]) /
      (\mathsf{Sqrt}[(\mathsf{Sum}[\mathsf{sign}[\mathsf{i},\mathsf{j},\mathsf{min},\mathsf{min}+\Delta r]*(\mathsf{Abs}[\mathsf{alfolded}[\![2]\!][\![\mathsf{p2}[\![k2]\!]]\![\![\mathsf{i},\mathsf{j}]\!])^2, \\
            {i, 1, Dimensions[foldedProteinData] [1] },
            {j, 1, Dimensions[foldedProteinData][2]}]) *
         (Sum[sign[i, j, min, min + \Delta r] * (Abs[a2folded[2][p2[k2]][i, j]])^2,
            {i, 1, Dimensions[foldedProteinData] [1]]},
            {j, 1, Dimensions[foldedProteinData][2]]}))),
    {min, 1, Floor[Dimensions[foldedProteinData] [2] / 2] }];
 Show
  ListPlot[{Table[{i / (Floor[Dimensions[foldedProteinData][2]] / 2]), frcDatan1[i]]},
       {i, 1, Floor[Dimensions[foldedProteinData][2] / 2]}],
     Table[{i / (Floor[Dimensions[foldedProteinData][2]] / 2]), frcDatanq[i]]},
       {i, 1, Floor [Dimensions [foldedProteinData] [2] / 2] } ] }, Joined → True,
   PlotStyle → Thickness[0.007], LabelStyle → {13, GrayLevel[0], Bold},
   TicksStyle → Thick],
  AxesLabel \rightarrow {HoldForm[normalized spatial frequency \left[\frac{1}{2.5 \text{ nm}}\right]], HoldForm[FRC]},
  PlotLabel → None, LabelStyle → {14, GrayLevel[0], Bold},
  Epilog \rightarrow {Dashed, Red, AbsoluteThickness[2], Line[{{0.3, 0}, {0.3, 1}}]}
 , {k1, 1, Length[p1]}, {k2, 1, Length[p2]}
 FRC
1.0
8.0
0.6
```







In[@]:= **minF** Out[0]= $\{13, 14, 15, 16, 18\}$

0.12

```
ln[*]:= bla = {1 / 13, 1 / 14, 1 / 15, 1 / 16, 1 / 18}
Out[0]=
 In[*]:= bla2 = Table[NPDC, {NPDC, 30, 140, 24}]
Out[0]=
        {30, 54, 78, 102, 126}
 In[@]:= ListPlot[
         ParallelTable[\{bla2[i]\}, (5(**10^-9*)) / (1/30) * bla[i]\}, \{i, 1, 5\}], Joined <math>\rightarrow True]
        11.0
        10.5
        10.0
 In[@]:=
         9.5
         9.0
         8.5
                                                      100
 In[\circ]:= Show[%111, AxesLabel \rightarrow {HoldForm[num of photons per pixel], HoldForm[resolution[nm]]},
         PlotLabel → None, LabelStyle → {GrayLevel[0], Bold}]
Out[0]=
        resolution(nm)
          11.5
          11.0
          10.5
          10.0
           9.5
           9.0
           8.5
                                                     num of photons per pixel
                                                120
 ln[*]:= 1/((1/18)*(5)/(1/30))
Out[0]=
         3
 In[*]:= (1 / 18) * (5) / (1 / 30)
Out[0]=
         25
         3
 In[ \circ ] := N[1 / ((1 / 18) * (5) / (1 / 30))]
Out[0]=
```

```
In[*]:= 0.12 * 10^9
Out[0]=
        1.2 \times 10^8
 In[*]:= N[(1/18)*(5)/(1/30)]
Out[•]=
        8.33333
 In[\circ]:= N[(1/18)*(5*10^-9)/(1/30)]
Out[0]=
        8.33333 \times 10^{-9}
 In[@]:= frcMinusBit =
         Table[(Re[(Sum[a1[5][i, j] * Conjugate[a2[5][i, j]] * sign[i, j, min, min + \Delta r],
                 {i, 1, 61}, {j, 1, 61}])]/(Sqrt[
                 (Sum[sign[i, j, min, min + \Delta r] * (Abs[a1[5][i, j]])^2, \{i, 1, 61\}, \{j, 1, 61\}]) * \\
                 (Sum[sign[i, j, min, min + \Delta r] * (Abs[a2[5][i, j]])^2, \{i, 1, 61\}, \{j, 1, 61\}])))
            (*-0.5*halfBit[min,min+△r]*), {min, 1, 30}];
        \label{listPlotTable[i, frcMinusBit[i]}, \{i, 1, 30\}], \ \mbox{Joined} \rightarrow \mbox{True}]
Out[0]=
        1.0
        0.8
        0.6
        0.2
                   5
                             10
                                      15
```