Principles of First Order Optics Applied to Components of the BioTek Absorbance Spectrophotometer Cuvette Assembly

Tutorial Presentation to the BioTek Mechanical Design Team OPTI521

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What is "First Order Optics"?

First Order Optics = Perfect Lenses (No Aberrations) Analysis Methods:

- Gaussian Optics Formulas referencing:
 - Principal Points
 - Focal Points
- Paraxial Optics
 - Rays near the axis (ignore curvature of lens)
 - Small angles (sin $\theta \approx \theta$)

BioTek Cuvette Assembly



Not an imaging system. We put energy into the sample at a particular wavelength and collect what comes out, measuring absorbance. Wavelength range = 200-999nm.

UV GRADE FUSED SILICA

- Because of the 200 380 nm wavelengths, we used fused silica optics.
- Note the variation in Refractive Index
- Refractive Index:
 - n = <u>Speed of light in vacuum</u> Speed of light in the material

HPFS® Fused Silica Standard Grade



Refractive Index and Dispersion

Data in 22ºC in 760mm Hg dry nitrogen gas

Wavelength	Refractive	Thermal	
[air]	Index "2	Coefficient	Ŀ
λ [nm]	n	$\Delta n/\Delta T^{*3}$ (ppm/K)	
1128.64	1.448870	9.6	
1064.00	1.449633	9.6	
1060.00	1.449681	9.6	
1013.98 n _t	1.450245	9.6	
852.11 n _s	1.452469	9.7	
706.52 n _r	1.455149	9.9	
656.27 n _c	1.456370	9.9	
643.85 n _c	1.456707	10.0	
632.80 n _{He-Ne}	1.457021	10.0	
589.29 n _D	1.458406	10,1	
587.56 n _d	1.458467	10.1	
546.07 n _e	1.460082	10.2	
486.13 n _p	1.463132	10,4	
479.99 n _e	1.463509	10.4	
435.83 n,	1.466701	10.6	
404.66 n _a	1.469628	10.8	
365.01 n _i	1.474555	11.2	
334.15	1.479785	11.6	
312.57	1.484514	12.0	
308.00	1.485663	12.1	
248.30	1.508433	14.2	
248.00	1.508601	14,2	
214.44	1.533789	17.0	
206.20	1.542741	18.1	
194.17	1.559012	20,4	
193.40	1.560208	20.5	
193.00	1.560841	20.6	
184.89	1.575131	22.7	

Snell's Law



www. http://hyperphysics.phy-astr.gsu.edu/

 $N_{air} = 1$ $N_{water} = 1.333$ $N_{fused silica} = 1.45$

Optical Fiber Numerical Aperture (NA)





http://www.photonics.com/EDU/Handbook.aspx?AID=25151

- NA = $n_{air}sin\theta = (n_{core}^2 n_{cladding}^2)^{1/2}$
- θ = half angle
- Our fiber is .040 inch diameter and has NA = 0.22. This is a common fiber NA.

Lens 1 images the fiber face into the cuvette sample



Where is the image? How big is it?



- Thin lens imaging equation: 1/z' = 1/z + 1/f (primed coordinates refer to the image)
- Magnification = z'/z = h'/h
- So what is f?

Focal Length of a Surface

- Curved surfaces separating materials of different index have <u>optical power</u>.
- Power of a surface = $\Phi = (n'-n)/R$
- Curvature = C = 1/R
- f = 1/Φ



• But the lens has two surfaces...

Gaussian Optics – Cardinal Points

- Front and Rear Focal Points
- Front and Rear Principal Planes



Reference 1

Focal Length of a Lens



 $\Phi = (n-1)[C_1 - C_2 + (n-1)(C_1)(C_2)T]$

- $\tau = t/n = Reduced Thickness$
- d and d' locate the principal planes from the Vertices V and V'
- Some lenses are so thin that the separation between P and P' approaches 0.

Focal Length of a Lens

- For a thin lens in air, $t \rightarrow 0$.
- Power = $\Phi = (n-1)(C_1 C_2)$
- d = d' = 0, so the principal planes are on the surfaces and the surfaces coincide as t \rightarrow 0.

• So what about our lens?

Thin and Thick Lens focal length for 7092199

7092199	[mm]	wavelength	index								
Thickness t =	3.1	206nm	1.543		ENTER DATA IN TH	E GREEN CELLS					
		248nm	1.508								
		546nm	1.460								
		1013nm	1.450								
							Power				
Power of first surface	Radius			Power of	second surface		of Lens	THICK LENS EQ'N	THICK LENS		
	of surface	Φ = (n' - n)/R		wave	Radius of surface	$\Phi = (n' - n)/R$	wave	$\Phi=(n\text{-}1)[1/\text{R1}-1/\text{R2}+(n\text{-}1)(1/\text{R1})(1/\text{R2})\tau]$	Focal Length	Focal Length	
wavelength	[mm]	[1/mm]		length	[mm]	[1/mm]	length	[1/mm]	[mm]	[inch]	
206nm	9.9	0.054848485		206nm	-9.9	0.054848485	206nm	0.103652962	9.65	0.380	
248nm	9.9	0.051313131		248nm	-9.9	0.051313131	248nm	0.09721352	10.29	0.405	
546nm	9.9	0.046464646		546nm	-9.9	0.046464646	546nm	0.088345193	11.32	0.446	
1013nm	9.9	0.045454545		1013nm	-9.9	0.045454545	1013nm	0.086491878	11.56	0.455	
								THIN LENS EQ'N	THIN LENS		
@206nm:							wave	Φ = (n-1)[1/R1 - 1/R2]	Focal Length	Focal Length	Error
Front Principal Plane:	$d = (\Phi_2/\Phi)\tau =$	1.063111185					length	[1/mm]	[mm]	[inch]	[%]
Rear Principal Plane:	$d' = (-\Phi_1/\Phi)\tau =$	-1.063111185					206nm	0.10969697	9.12	0.359	5.5
							248nm	0.102626263	9.74	0.384	5.3
NOTES:	REV. DESCRIPTIO A RELEASE TO PRODUCTION 8 NOTE 2:3' TO 5'	N BC0 DATE 42199 3/4.08 42299 4.23.08	BY RMP RMP				546nm	0.092929293	10.76	0.424	4.9
1. ALL DIMENSIONS IN MILLIMETE	RS						1013nm	0.090909091	11.00	0.433	4.9

BI-CONVEX LENS MATERIAL: FUSED SILICA, UV GRADE COMMERCIAL POLISH 80-50 NOT COATED. NOT COATED CENTERING: 5' DEVIATION, NOTE : WITH RESPECT TO FLATS AND OUTSIDE DIAMETER. RADIUS TOLERANCE: 10 RINGS REGULARITY TOLERANCE: 2 RINGS 3.1±0.1 -Ø7.90 -0.25 ±0.10 X 0.25 ±0.10 2X → (1.3) ******BioTek AW ANSI Y14.5-82 LENS, OPTICAL, SYM STEP BI-CONVEX HARE THEFT EXPRESS VETTER CORE IN OF A BUTE YORT DESCRIPTION. 7092199-DG SCALE ADVIS PROFETER SALE SALE 1/1

• Note the 5% error.

• Note the wavelength dependence of the focal length. Blue focuses closer than red.

So where is the image?

- Now that we know the focal length, we can locate the image: 1/z' = 1/z + 1/f and m = z'/z z, z', f all measured from the principal planes.
- Let's go with the 5% error thin lens result for f.

			Z'	
	Z		Image	
	Object	Focal	Distance	m
	Distance	Length	$Z' = [(1/Z) + (1/f)]^{-1}$	Magnification
index	[inch]	[inch]	[inch]	m = Z'/Z
206nm	-0.8	0.359	0.651	-0.81
248nm	-0.8	0.384	0.737	-0.92
546nm	-0.8	0.424	0.901	-1.13
1013nm	-0.8	0.433	0.944	-1.18

- Note that the magnification is ~ 1:1. (Negative means inverted.)
- Note that these values are in air. (The cuvette will hold a liquid.)

Next the beam passes thru a tilted plate beam splitter



A ray passing through a **plane parallel plate** is displaced but not deviated; the input and output rays are parallel.



An image formed through a plane parallel plate is longitudinally displaced, but its magnification is unchanged.

Reference 1

- D ≈ tθ(n-1)/n
- This results in an offset of ~.007 inch.

Next, we enter the cuvette



displaced, but its magnification is unchanged. $d = \left(\frac{n-1}{n}\right)t$ $d = \frac{t}{3} \text{ for } n = 1.5$ Actual
Reference 1

An image formed through a plane parallel plate is longitudinally

d = (n-1)t/n

- The cuvette acts as a plane parallel plate of index 1.33 if it is full of water.
- The image displacement due to ½ the cuvette is ~.062 inch.
- The beam splitter on the previous page also has this effect ~.014 inch.
- The two plates together displace the image ~ .076 axially.

Influences on the Image location

- The system is used with many wavelengths, each with its own focal length.
- The cuvette sample is usually aqueous, but its index may vary.
- The sample may be turbid.
- The system is used in air when running its startup routine.

So the location of the image in the cuvette can vary around the general location we determined. The second lens could be analyzed in a similar way using the image in the cuvette as its object.

Conclusion

The name of the game for this system is:

- To use inexpensive optics that can
- Put energy at a wide variety of wavelengths into the sample chamber
- And collect all that comes out the other side regardless of what is in the chamber.

First order optics can help with the initial layout of a system, or to understand in general how it works. To design it to be robust and accurate requires more advanced methods and lots of testing.

References

- Grievenkamp, John E., "Field Guide to Geometrical Optics" SPIE Press, Bellingham WA, 2004
- Prof. John Grievenkamp, class notes and lectures from "Optical Design and Instrumentation I, Fall 2013.