



# Basic EnVision Training

Getting you up to speed on the EnVision

Daniel Cardillo Field Application Scientist

# Training Cirriculum



#### Classroom

- Software Familiarization (classroom setting is preferred)
  - EnVision options overview and definition of what is on your system
  - Light source options
  - Filter, Beamsplitter, Dichroic, and Monochromator overview
  - o Inventory tab review
  - Reagent technologies as a guide to why you would select certain options, light sources, filters, and mirrors
  - Protocol review (Measurement technology, Plate, and well selection/data analysis)
  - Preset protocol tab
  - Building protocols by using copy feature
  - Building your own protocols.
  - Basic Protocol optimization overview

#### Lab Training

- Hardware overview (in front of machine)
  - Proper power up sequence, and starting of software
  - Identify all of the key pieces of hardware
  - Demonstrate filter, mirror(top and bottom), and aperture changing
  - Show plate orientation, and identify the A1 location on plate holder
  - Using stackers (If applicable)
  - Dispenser Overview (If applicable)
- Basic programming of Protocols
- Optimization of 2-3 protocols User creates the protocol
- Demonstration of applying basic calculations and exporting data after assay has been run.
- Cover what is available in an Advanced Training session





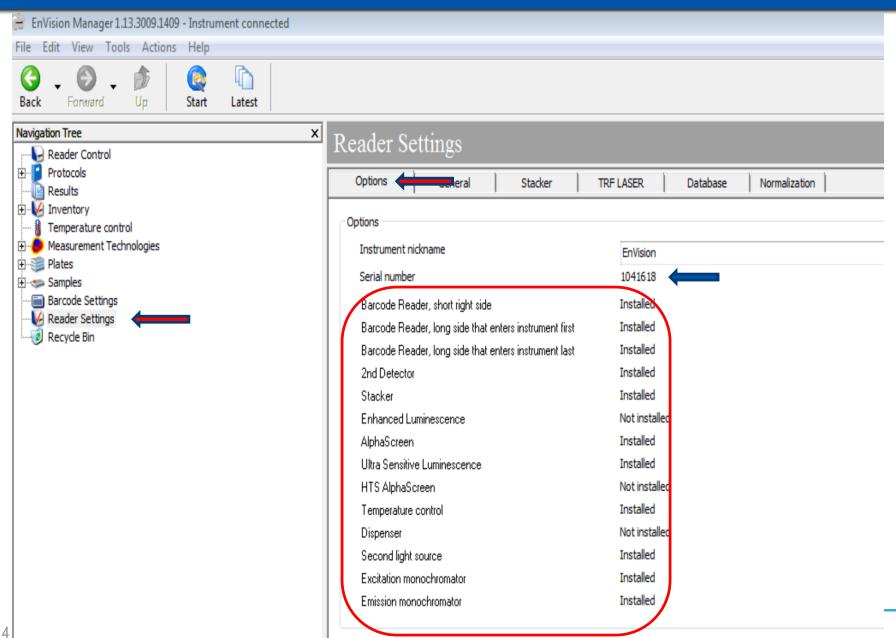
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Top 3 reasons to pay attention to this course...

- 1) You will learn how to use your EnVision
- 2) Once I leave...you're the EnVision expert
- 3) The Basic refresher course is \$3,000!

## **EnVision Options**







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	Emission monochromator

# **Light Sources**





- 1. Xenon Flash Lamp
  - FI,
  - Abs,
  - FP,
  - FRET
  - TRF
  - TR-FRET
- 2. 337 Laser
  - TRF
  - TR-FRET

# 3. 680 Laser

- AlphaScreen
- AlphaLISA
- 4. Luminescent sample
  - Glow
  - Flash





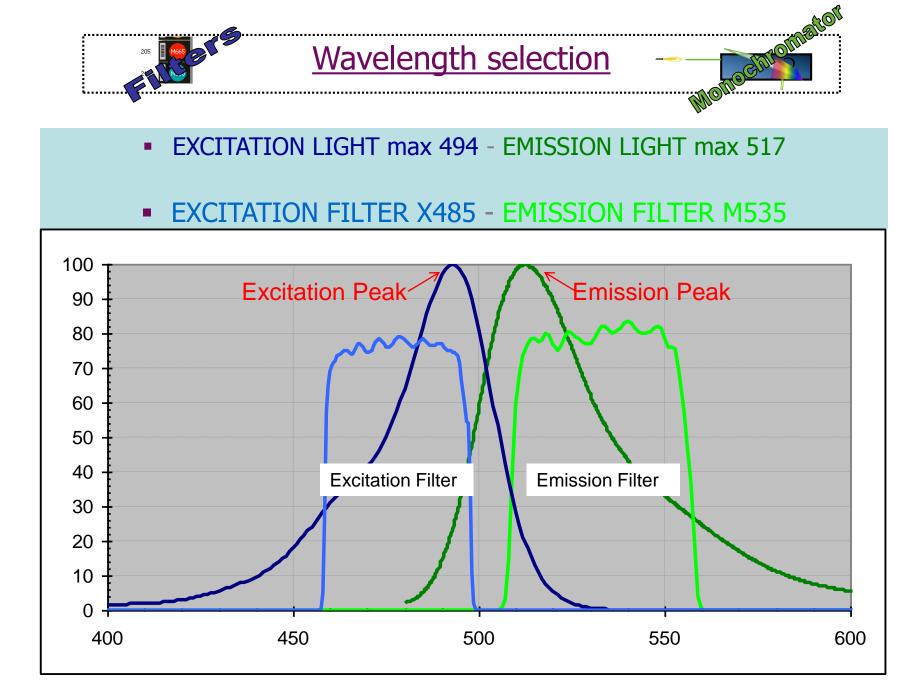


## Filter/Mirror Review





- EnVision can have up to 16 filters and 5 mirrors in the system at any time.
- All Filters and Mirrors are Barcoded
- Additional Filter slides allow for easy swapping of filters

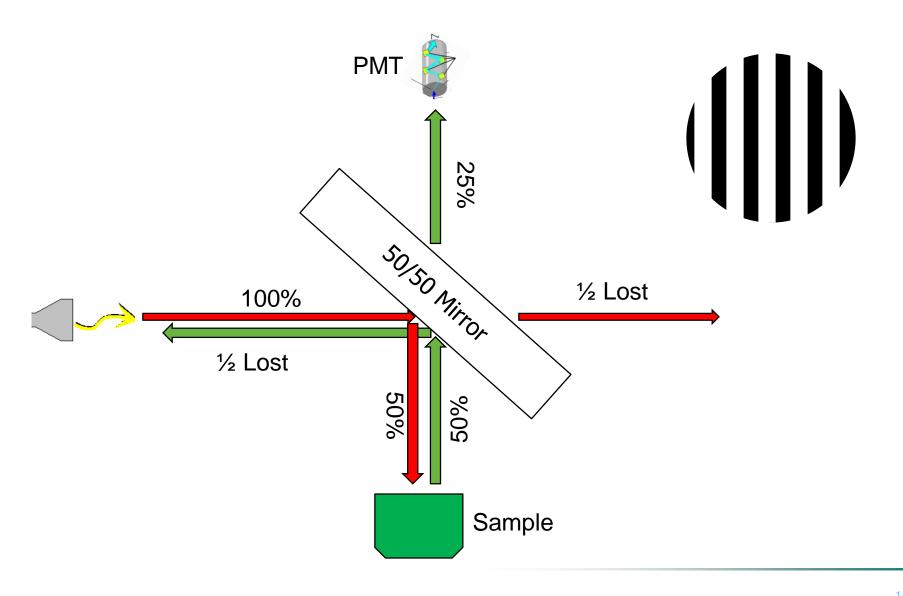






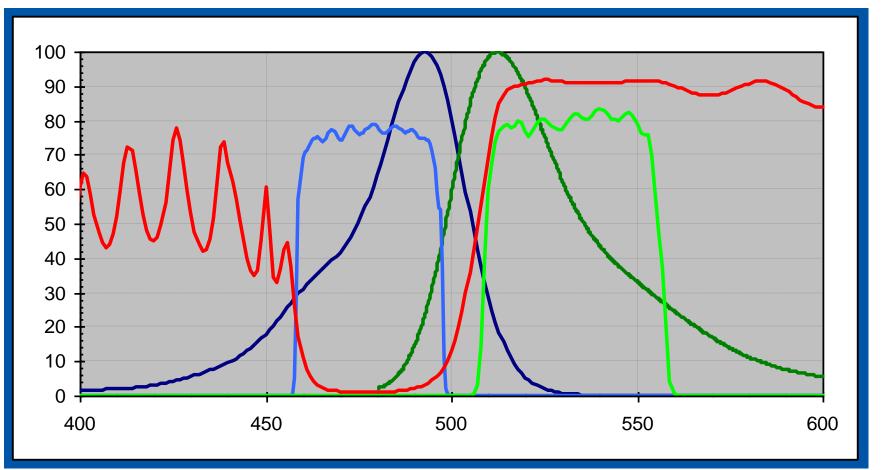
How do you get filtered light in and out of a well with something that is physically in the light path between the sample and the PMT







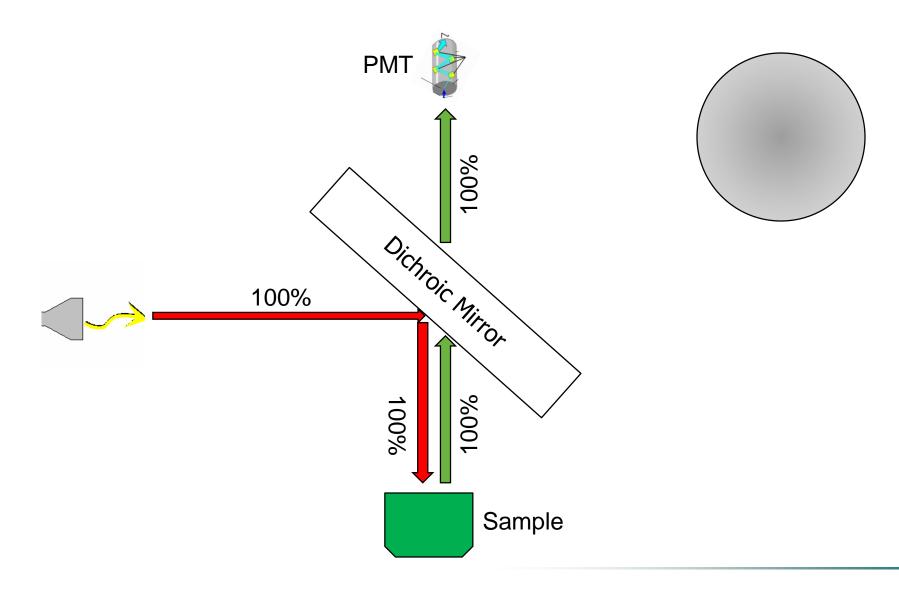
 EXCITATION FILTER X485 - DICHROIC MIRROR D505 - EMISSION FILTER M535



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### **Principle of the Dichroic Mirror**

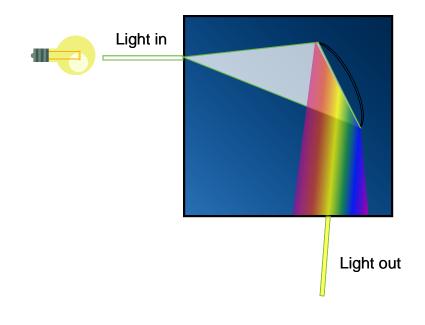




How does a Monochromator work?

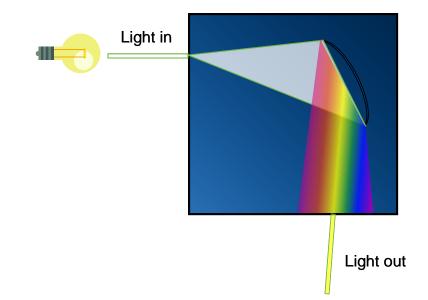


#### DIFFRACTION GRATINGS



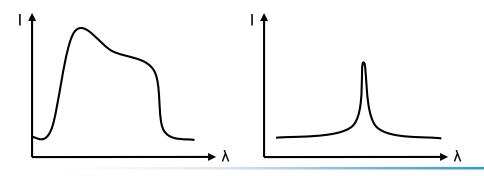


#### Single Monochromator



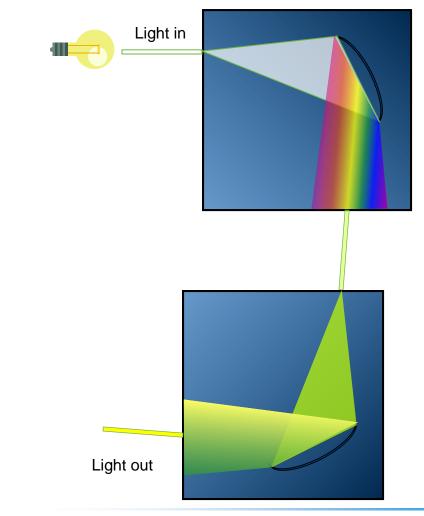
Monochromators is an optical module that splits the the incoming light into partial components

Monochromators grate is mounted to a motorized axis, allowing it to turn. The Spectrum then moves across a slit, which passes the desired wavelength light.





#### Dual Monochromator module



The filtering capacity of a monochromator is limited.

 To address this a multiple of monochromators are sequentially coupled (Dual).

#### In a Dual Monochromator

- The wavelength is selected once, and after that another time in the sequentially connected second grate.
- The filtered light from the second monochromator is then clean.

## Inventory



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	605 Eu-Tb-Sm sequential single mirrri No No	
Beta-Lactamase	658 Sm/Eu simultaneous duplex No Yes	
Beta-Lactamase Dual	🚰 401 General - No BS	50
Beta-Lactamase Dual Enh	- No D5	
BODIPY TMR	🚰 404 Luminescence - No LU	Μ
BODIPY TMR FP	GA 405 BODIPY TMR - No DS	55
🥳 BODIPY TMR FP Dual	408 Dichroic 385 - No D3	85
🚰 BODIPY TMR FP Dual Enh		60
🚰 BRET2	E 🚰 418 Beta-Lactamase - No D4	125
آ BRET2 Enh		58
- GEP/YEP		i05fp
CFP/YFP Dual Enh	Ga 432 BODIPY TMR FP - No DS	55fp
Cy5	Ga 433 Umbelliferone FP - No D4	юбр
	Ga 434 Texas Red FP - No D5	i95fp
Cys FP Dual Enh	🚰 435 General FP - No BS	50fp
G Dichroic 385	Ga 437 Cy5 FP - No D6	558fp
FITC		600
FITC FP	🚰 446 LANCE/DELFIA Dual / Bias - Yes D4	400/D630
🥁 FITC FP Dual		50/BS50
🚰 FITC FP Dual Enh		400/D630
🚰 General		M/BS50
🥻 General Dual		M/D475
General Dual Enh		i25/D490
General FP		400/D630
General FP Dual		i05fp/D535
ANCE Dual		555p/D595
		00fp/8550
LANCE/DELFIA Bias		19510155
LANCE/DELFIA Dual		50fp/BS50
LANCE/DELFIA Dual / Bias		58fp/D688
		M/BS50
🥻 Luminescence		M/D475
🚰 Luminescence Dual	2	125/D490
🥻 Luminescence Dual Enh		100/D630
Texas Red FP		05fp/D535

### Reader Control



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ile Edit View Tools Actions Help	
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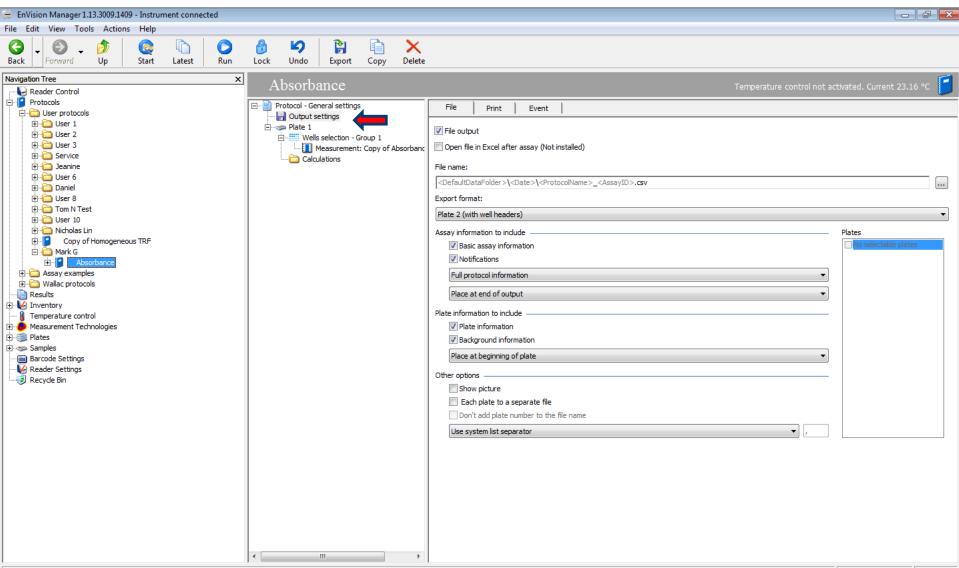
## **Reader Settings**



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#### **Protocol General Settings**

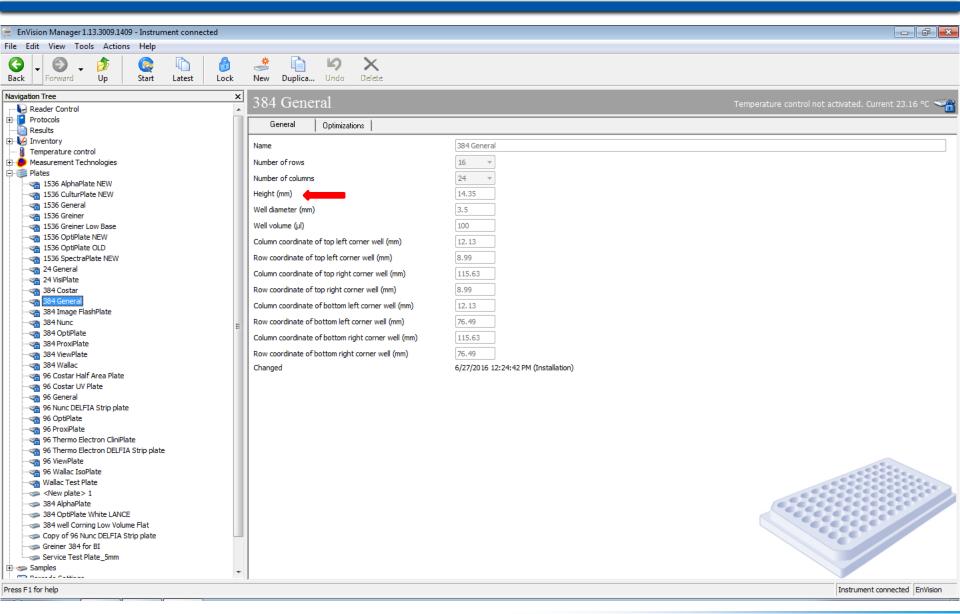




Press F1 for help

#### Plates General Tab





## **Temperature Control**



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File Edit View Tools Actions Help		
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	Temperature control         Image: Plate heating adjustment         Set measurement chamber temperature to 200°C         Condensation prevention for sealed plates         Upper heater temperature is 000°C (warmer in than lower heater temperature)         Set plate heating off after next assay         Image: Plate temperature offset         Plate temperature is 000°C (warmer in than ambient temperature)	Temperature control not activated. Current 23.36 °C
I Press F1 for help	1	Instrument connected EnVision

### Barcode Reader



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Results     Results     Inventory     Temperature control     Measurement Technologies     Plates     Samples     Samples     Reader Settings     Recycle Bin	Barcode reading       Protocol starting         Read barcode from the <ul> <li>short right side and use it as</li> <li>barcode</li> <li>long side that enters instrument first and use it as</li> <li>barcode</li> <li>long side that enters instrument last and use it as</li> <li>barcode</li> </ul> Protocol definition by barcodes <ul> <li>w Use barcodes as plate ID only</li> <li>w Data and the protocol definition by barcodes</li> <li>w Data and the protocol definition by barcodes</li> </ul>	
	<ul> <li>Define the protocol using</li></ul>	
Press E1 for help		Instrument connected EnVision

## **Reagent Technologies**

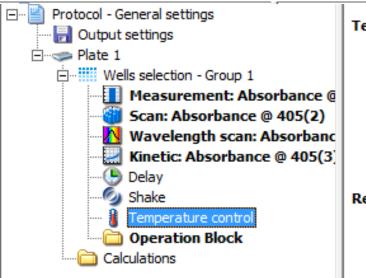


	Absorbance	Fluorescence	Luminescence	FP	DELFIA	LANCE	ALPHASCREEN
<u>LIGHT</u> <u>SOURCE</u>	Ex Xenon Flash + Filter/Mono	Ex Xenon Flash + Filter/mono	Reagent is light source	Ex Xenon Flash + Filter	Ex Xenon Flash + Filter or TRF Laser	Ex Xenon Flash + Filter or TRF Laser	Alpha Laser
MIRROR REQUIRED		Dichroic Mirror or Beamsplitter	Std Lum requires a Lumi mirror block	Dichroic Mirror or Beamsplitt er w/polarize r	Dichroic Mirror	Dichroic Mirror	Std Alpha uses Alpha mirror block
Emission Wavelength Selection	Reagent is Em filter	Em Filter/Mono	-Std Lum uses single or dual Filter -US Lum uses Aperture	Em Filters with S and P polarizers	Em Filter(s)	Em Filter(s)	Std Alpha uses single or dual Em Filters HTS uses Aperture
DETECTION DEVICE(s)	Read on Photodiode	Read on Single or dual PMT with filters Read on Dedicated PMT with Mono	-Std Lum uses single or dual PMT -US Lum uses Dedicated PMT	Single or dual PMTs	Single or dual PMTs	Single or dual PMTs	Std Alpha uses single or dual PMT HTS Alpha uses Aperture and dedicated PMT



DNA Luminescence ATPlite TR-FRET **Protein-Protein** Kinase Surefire-Ultra FP AlphaLISA DELFIA LANCE BrightLite TRF AlphaScreen LANCE-Ultra AlphaPLEX Absorbance Surefire NeoLite SteadyLite Fluorescence





#### Temperature control

Measurement chamber temperature (°C)

Fast start

First assay/plate repeat affected

Last assay/plate repeat affected

#### Repeats

Number of assay repeats

Start assay repeat each

Number of plate repeats

Start plate repeat each



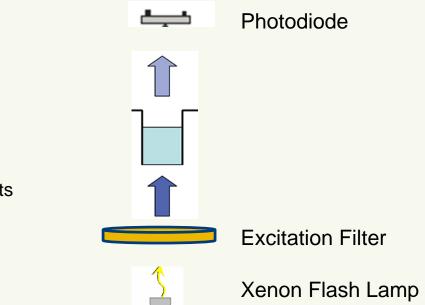




# Absorbance Assays



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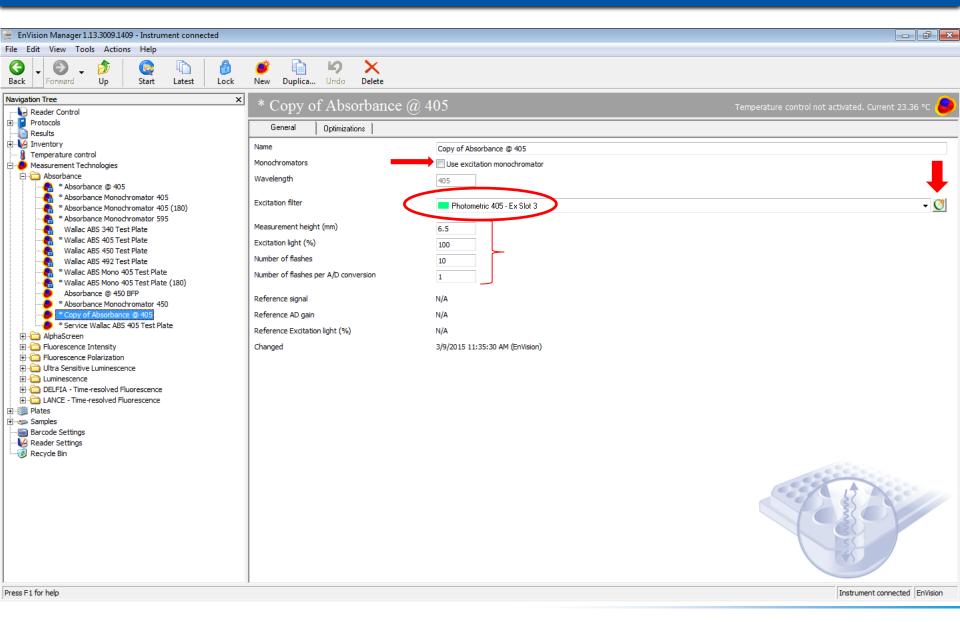
- Inexpensive
- Easy to build
- Large # of commercial kits
- Wash Assay
- Limited Window





## **Optical Density**





# Fluorescence Assays

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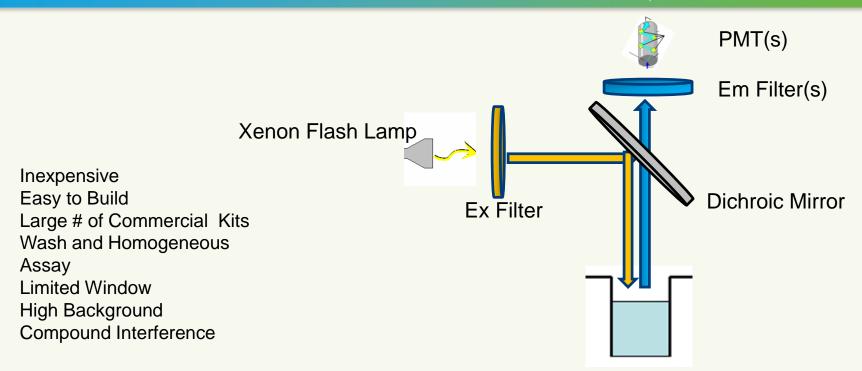
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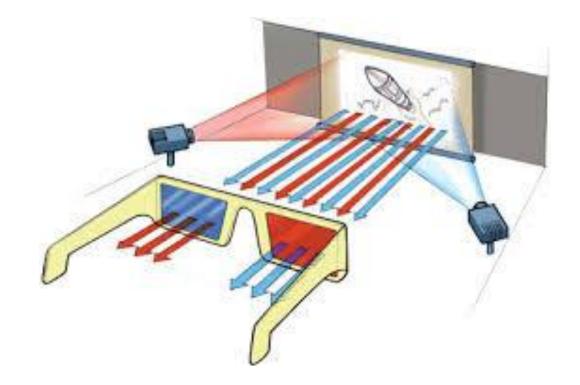
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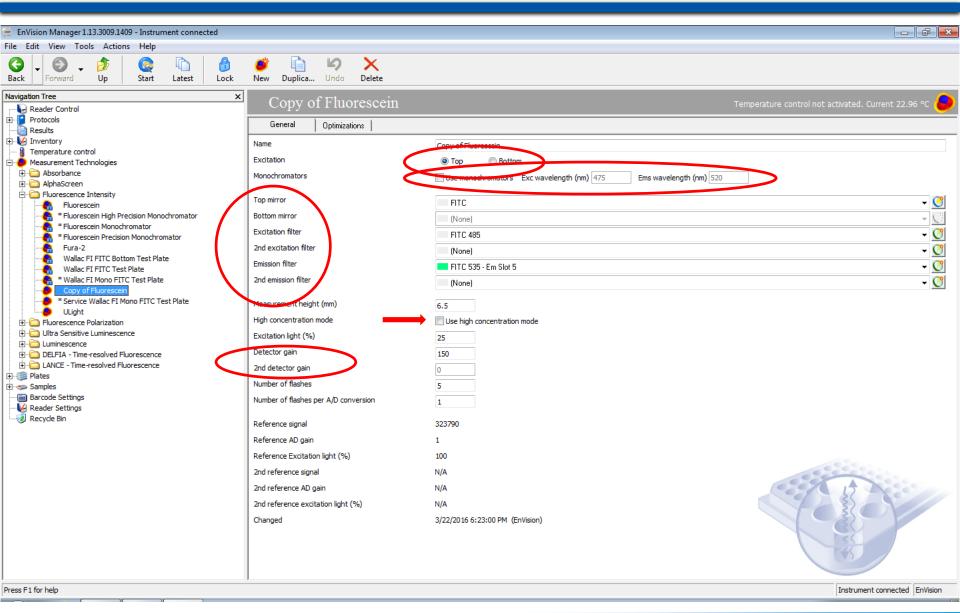




### Filtered Light In- Filtered Light Read

#### Fluorescence Protocol





# Luminescence Assays



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PMT(s)

Emission Filter(s)

Low background •

- Variety of kits and reagents • available
- Good assay window •
- Can be inject and read •
- Signal Diminishes over time .

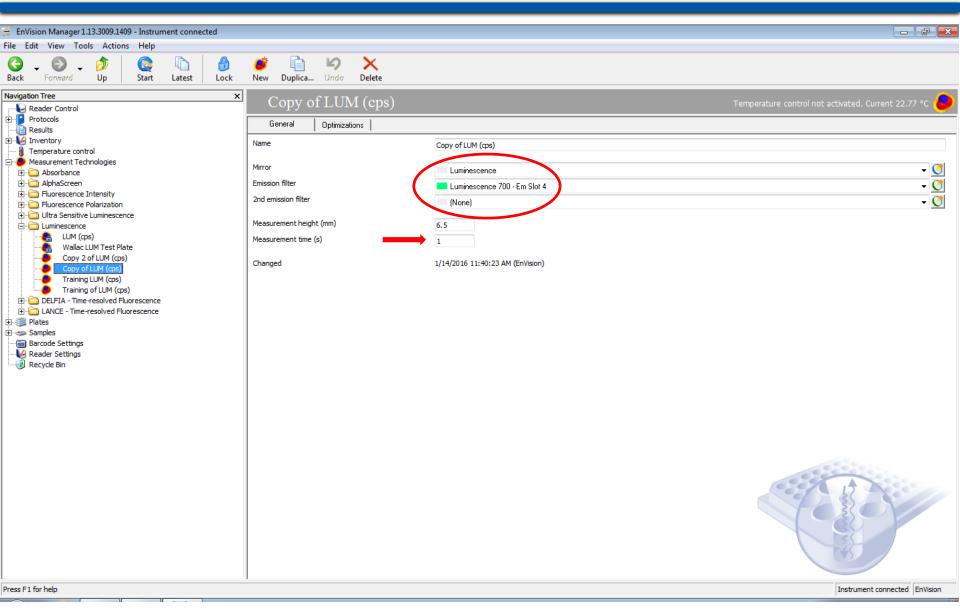


	LUMINESC	ENCE ASSA	Y SYSTEM	S AND INST	RUMENTS	
	AequoScreen AequoZen PhotoScreen	brite <b>lite</b> plus	steady <b>lite</b> plus	neo <mark>lite</mark>	ATP <b>lite</b> 1step	ATPlite
Applications	<ul> <li>Calcium Flux</li> <li>GPCR</li> <li>Ion Channels</li> </ul>	<ul> <li>Reporter Gene Assay</li> <li>GPCR</li> </ul>	<ul> <li>Reporter Gene Assay</li> <li>GPCR</li> </ul>	<ul> <li>Reporter Gene Assay</li> <li>GPCR</li> </ul>	<ul> <li>Cytotoxicity</li> <li>Cell Proliferation Assay</li> </ul>	<ul> <li>Cytotoxicity</li> <li>Cell Proliferation Assay</li> </ul>
Half-life (hours)	Flash	0.5	4 to 5	2.5	0.5	5
Sensitivity	High	Very High	Moderate	High	As few as 5 cells/well	As few as 5 cells/well
Plate Format (wells)	96 384 1536	96 384 1536	96 384 1536	96 384 1536	96 384 1536	96 384 1536
Instrumentation	— MicroBeta <sup>2</sup> LumiJET	TopCount MicroBeta <sup>2</sup> EnSpire	TopCount MicroBeta <sup>2</sup> EnSpire	TopCount MicroBeta <sup>2</sup> EnSpire	TopCount MicroBeta <sup>2</sup> EnSpire	TopCount MicroBeta <sup>2</sup> EnSpire
	EnVision VICTOR X VICTOR X Light —	EnVision VICTOR X VICTOR X Light ViewLux	EnVision VICTOR X VICTOR X Light ViewLux			

### Sample Generated Light- Filtered Light Read

## Standard Luminescence

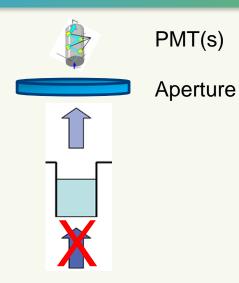




# Ultra Sensitive Luminescence Assays



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## Ultra Sensitive Luminescence

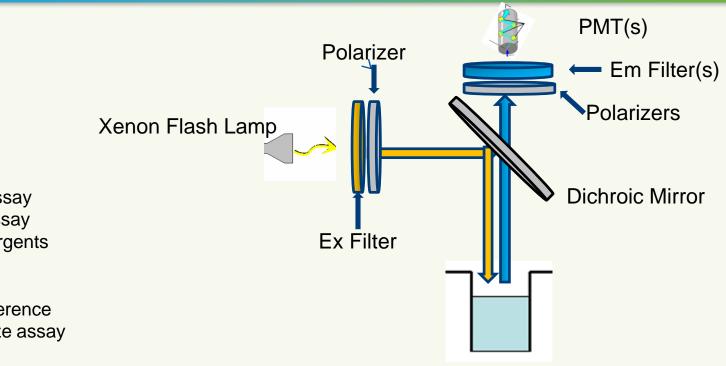


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Protocols     Results	General Optimizations		
🗈 🥳 Inventory	Name	Copy of US LUM 384 (cps)	
Temperature control     Measurement Technologies			
	Aperture	384 Plate US Luminescence aperture	- 🔇
🕀 🛅 AlphaScreen	Distance between eleterand determine (res)		
	Distance between plate and detector (mm)	0	
	Measurement time (s)	0.5	
	Glow (CT2) correction factor (%)	0	
	Changed	7/10/2015 10:59:54 AM (EnVision)	
Wallac US LUM Test Plate	Changed	715/2010 10:00:0 (Envision)	
BMS US LUM 96 (cps)			
Copy of US LUM 384 (cps)			
Training US LUM 384 (cps)			
Herrich - Time-resolved Fluorescence     LANCE - Time-resolved Fluorescence			
Plates			
🗈 🥌 Samples			
Barcode Settings      Keader Settings			
Recycle Bin			
			1
Press F1 for help			Instrument connected EnVision

# **Fluorescence Polarization**



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- Inexpensive
- Easy to Build
- Homogeneous Assay
- Single Binding Assay
- Sensitive to Detergents
- Limited Window
- High Background
- Compound Interference
- Difficult to optimize assay conditions





Without Polarizer

With Polarizer

Filtered + Polarized Light In- Filtered + Polarized Light Read

## Fluorescence Polarization



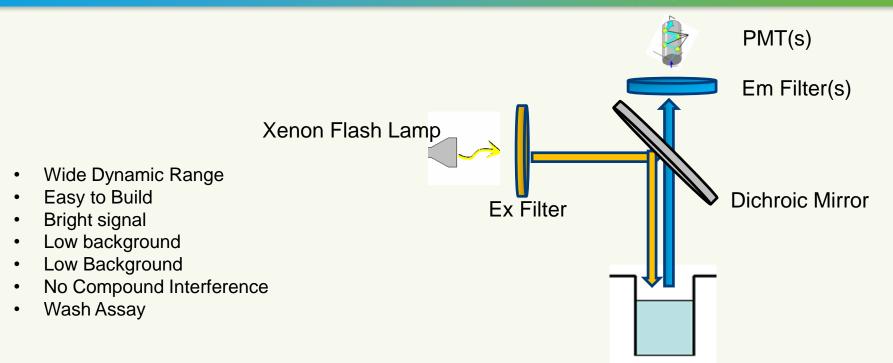
🚊 EnVision Manager 1.13.3009.1409 - Instrument connected - 6 **- X** File Edit View Tools Actions Help P × S G 0 Back Delete Forward Up Start Latest Lock New Duplica... Navigation Tree x Copy of Copy of FP Fluorescein single  $\bigcirc$ + Protocols General Optimizations Results 🗄 😡 Inventory Name Copy of Copy of FP Fluorescein single Temperature control - Measurement Technologies Mirror FITC FP Dual Enh - 0 🗄 🛅 Absorbance 🗄 🛅 AlphaScreen Excitation filter - 🗸 FITC FP 480 E Fluorescence Intensity Emission filter - 🔮 FITC FP P-pol 535 E Fluorescence Polarization FP Fluorescein Dual 2nd emission filter - 0 FITC FP S-pol 535 FP Fluorescein Dual 681 Wallac FP FITC Test Plate HTS Measurement height (mm) 6.5 Wallac FP FITC Test Plate HTS 681 Copy of Copy of FP Fluorescein single High concentration mode Use high concentration mode Copy of FP Fluorescein single Excitation light (%) 100 🗄 🧰 Ultra Sensitive Luminescence G - factor 🗄 🛅 Luminescence 1 E DELFIA - Time-resolved Fluorescence Detector gain 500 E-Cance - Time-resolved Fluorescence 2nd detector gain 700 🗄 🧐 Plates Number of flashes Barcode Settings Number of flashes per A/D conversion W Reader Settings 1 🔞 Recycle Bin Reference signal N/A Reference AD gain N/A Reference Excitation light (%) N/A Changed 6/22/2016 3:44:00 PM (EnVision) Instrument connected EnVision

Press F1 for help

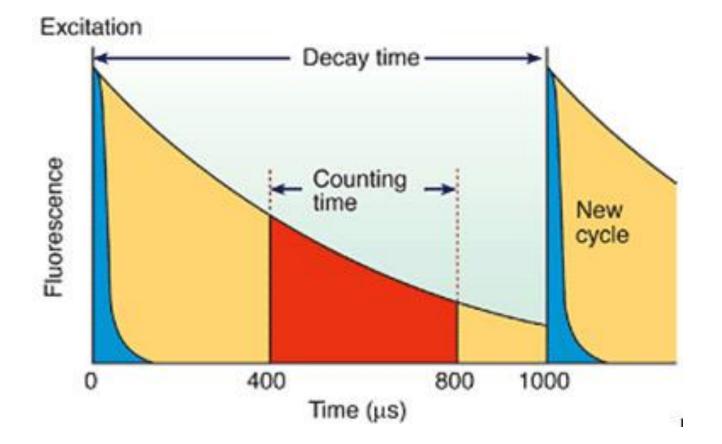


# DELFIA (TRF)

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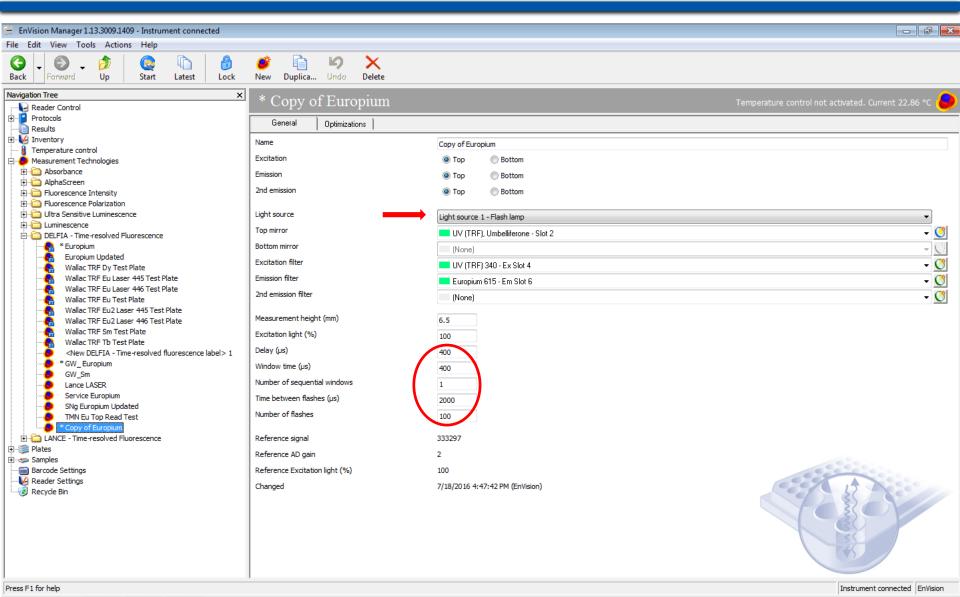




Filtered Light In - Time resolve - Filtered Light Read

# **DELFIA** Flash Lamp





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# DELFIA TRF Laser



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🕀 💼 Absorbance	Emission	Top OBottom	
⊕- 🔂 AlphaScreen ⊕- 🔂 Fluorescence Intensity	2nd emission	Top     O     Bottom	
Horescence Polarization			
🗄 🛗 Ultra Sensitive Luminescence	Light source	Light source 2 - Laser	▼
⊕- 🔂 Luminescence ⊟- 🔂 DELFIA - Time-resolved Fluorescence	Top mirror	LANCE/DELFIA Bias	✓
	Bottom mirror	(None)	
Europium Updated     Wallac TRF Dy Test Plate	Emission filter	Europium 615 - Em Slot 6	- 💟
Wallac TRF Eu Laser 445 Test Plate	2nd emission filter	(None)	- 0
Wallac TRF Eu Laser 446 Test Plate			
Wallac TRF Eu Test Plate Wallac TRF Eu Laser 445 Test Plate	Measurement height (mm)	6.5	
Wallac TRF Eu2 Laser 446 Test Plate	Excitation light (%)	100	
Wallac TRF Sm Test Plate Wallac TRF Tb Test Plate	Delay (µs)	400	
New DELFIA - Time-resolved fluorescence label> 1	Window time (µs)	400	
* GW_Europium	Number of sequential windows	1	
GW_Sm Lance LASER	Time between flashes (µs)	16600	
Service Europium	Number of flashes	100	
SNg Europium Updated TMN Eu Top Read Test			
* Copy of Europium	Reference signal	N/A	
Copy of Wallac TRF Eu Laser 445 Test Plate	Reference AD gain	N/A	
E-C LANCE - Time-resolved Fluorescence ⊡-S Plates	Reference Excitation light (%)	N/A	
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Recycle Bin			
Press F1 for help			Instrument connected EnVision



# LANCE (TR-FRET)

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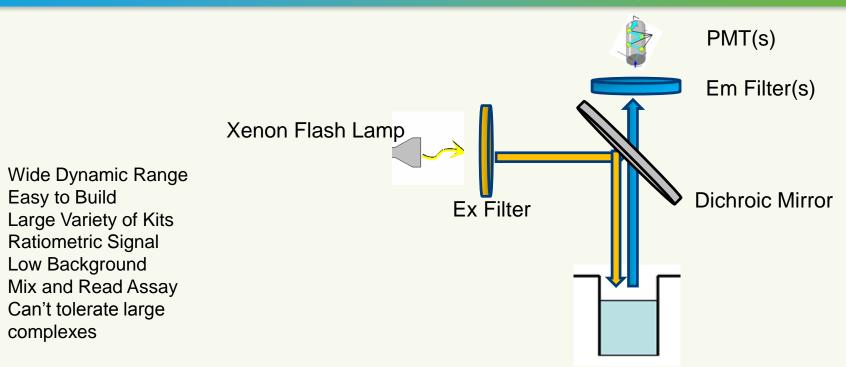
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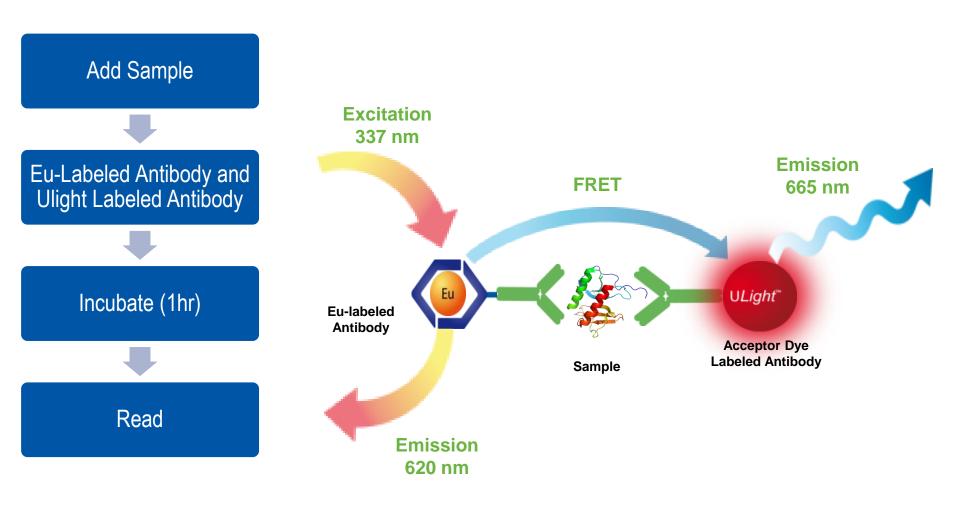
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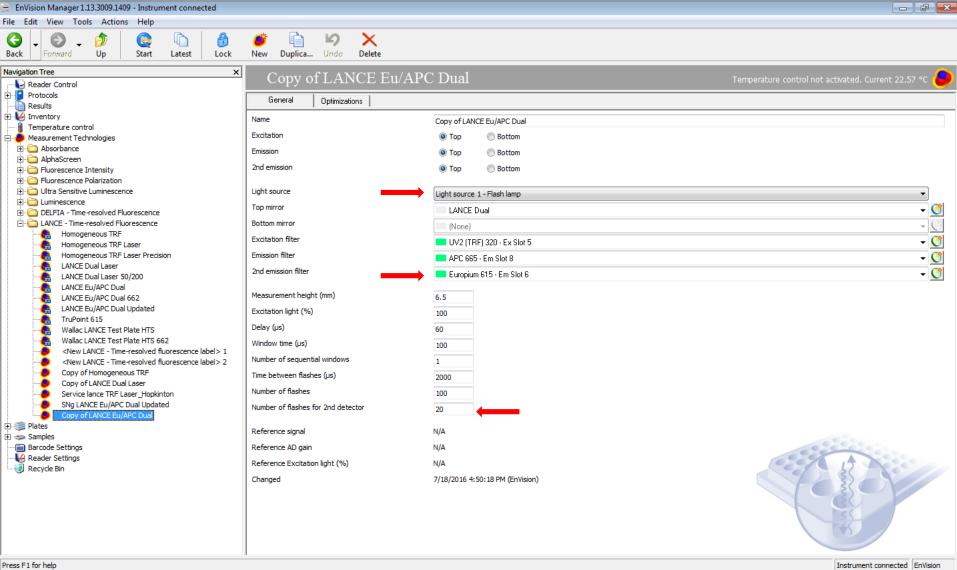


### Filtered Light In - Time resolve - Filtered Light Read

# LANCE/ TR-FRET Flash Lamp



😑 EnVision Manager 1.13.3009.1409 - Instrument connected



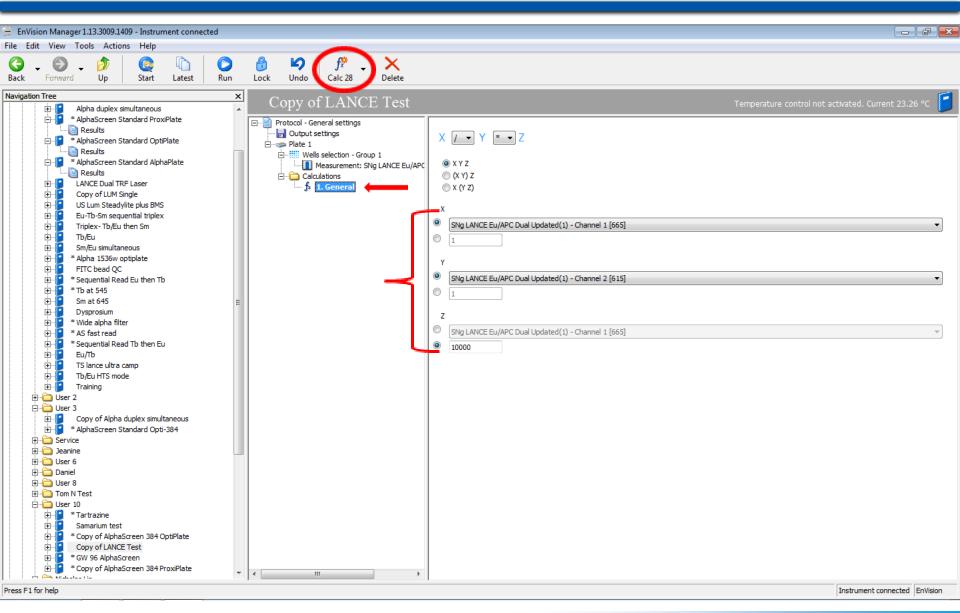
## LANCE/ TR-FRET TRF Laser



😑 EnVision Manager 1.13.3009.1409 - Instrument connected - 6 <u>- x</u> File Edit View Tools Actions Help r × S G θ Ð 0 Back Up Start Latest Lock New Duplica... Undo Delete Forward Navigation Tree × Copy of LANCE Dual Laser - Reader Control Protocols General Optimizations Results + M Inventory Name Copy of LANCE Dual Laser Temperature control Excitation Top Bottom - Measurement Technologies 🗄 🛅 Absorbance Emission Top Bottom 🗄 🫅 AlphaScreen 2nd emission Top Bottom 🗄 🛅 Fluorescence Intensity Fluorescence Polarization Light source Light source 2 - Laser Ŧ 🗄 🧰 Luminescence Top mirror - 🕑 LANCE/DELFIA Dual / Bias E-Contraction - Time-resolved Fluorescence E-Call LANCE - Time-resolved Fluorescence Bottom mirror (None) - 1 Homogeneous TRF Emission filter - 🔮 APC 665 - Em Slot 8 Homogeneous TRF Laser - 0 Homogeneous TRF Laser Precision 2nd emission filter Europium 615 - Em Slot 6 LANCE Dual Laser LANCE Dual Laser 50/200 Measurement height (mm) 6.5 LANCE Eu/APC Dual Delay (µs) LANCE Eu/APC Dual 662 20 LANCE Eu/APC Dual Updated Window time (µs) 200 TruPoint 615 Number of sequential windows 1 Wallac LANCE Test Plate HTS Wallac LANCE Test Plate HTS 662 Time between flashes (µs) 16600 <New LANCE - Time-resolved fluorescence label> 1 Number of flashes 20 <New LANCE - Time-resolved fluorescence label > 2 Copy of Homogeneous TRF Number of flashes for 2nd detector 5 Copy of LANCE Dual Laser Service lance TRF Laser\_Hopkinton Reference signal 283766 SNg LANCE Eu/APC Dual Updated Copy of LANCE Eu/APC Dual Reference AD gain 2 🗄 🧐 Plates Changed 7/7/2015 2:29:24 PM (EnVision) + Samples Barcode Settings Weader Settings 🥡 Recycle Bin Press F1 for help Instrument connected EnVision

### PerkinElmer<sup>\*</sup> For the Better

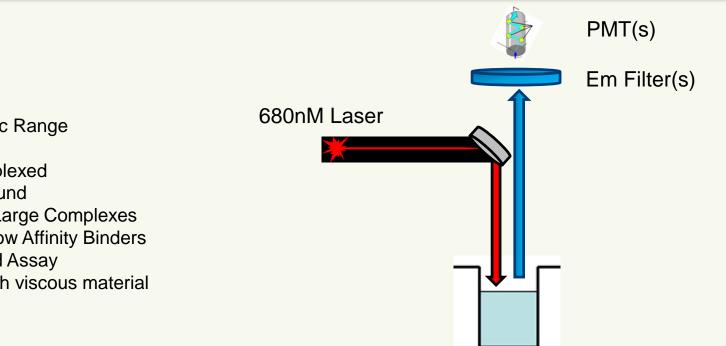
# Applying a Ratio Calculation to TR-FRET assays



## AlphaScreen (AlphaLISA, AlphaPLEX, Surefire)



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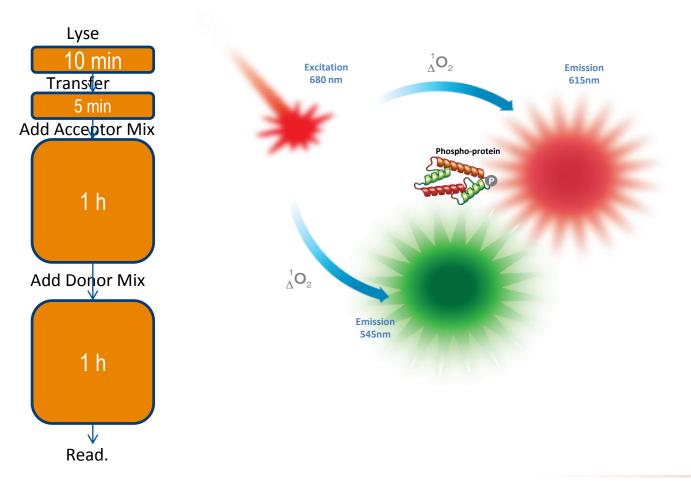


- Wide Dynamic Range •
- Easy to Build .
- Can be Multiplexed ٠
- Low Background •
- Can Handle Large Complexes •
- Works with Low Affinity Binders •
- Mix and Read Assay •
- Cant work with viscous material .

# Alpha SureFire Ultra Multiplex

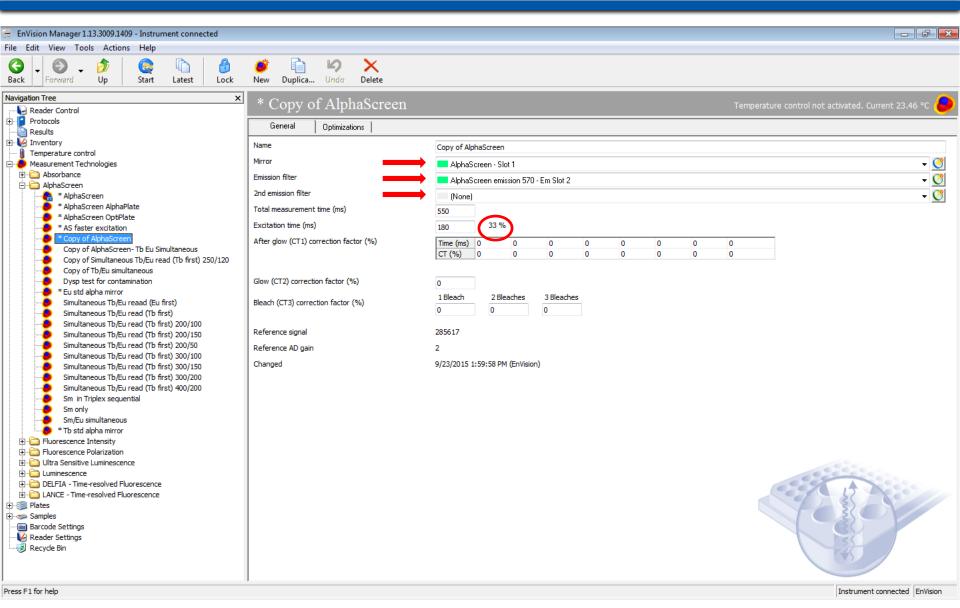


- Alpha SureFire Ultra Multiplex, dual-targets kits:
  - Phospho-ERK (Eu) + total-ERK (Tb): MPSU-PTERK-K500/10K/50K/HV
  - Phospho-AKT1/2/3 (Eu) + total-AKT1 (Tb): мрзи-ртакт-к500/10к/50к/ну



# AlphaScreen/AlphaLISA/AlphaPLEX



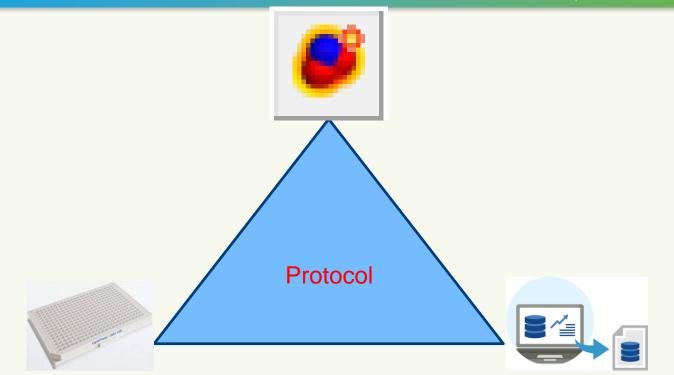


51

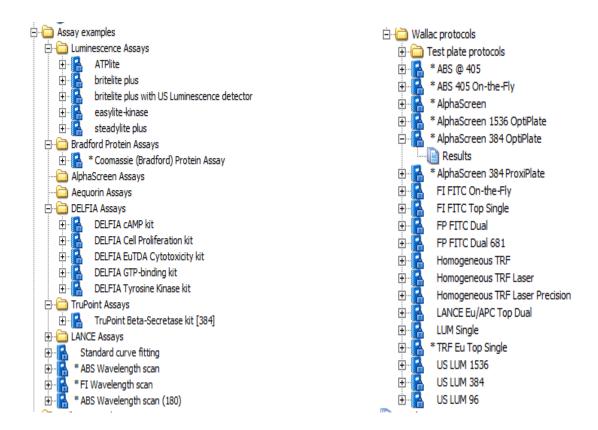
# Creating your Protocol



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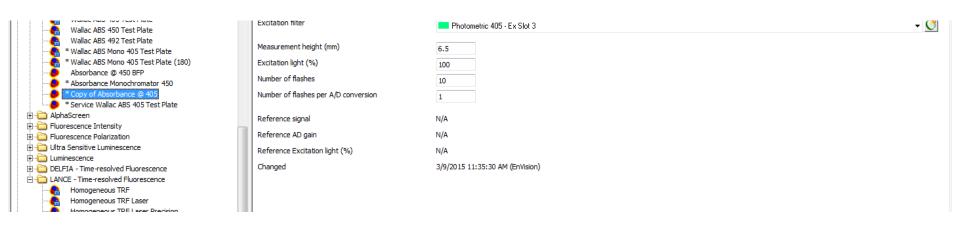
## Creating a Protocol

PerkinElmer<sup>\*</sup> For the Better

EnVision Manager 1.13.3009.1409 - Instrument connected File Edit View Tools Actions Help Training Temperature control not activated. Current 23.46 °C ⊡.... Protocol - General settings Measurement 🗟 Output settings Label (valid labels marked with \*): Measurement: 🛅 Cali Vations Optimizations 1 Invalid operation Repeats Check parameters Number of assay repeats n Start assay repeat each s -Number of plate repeats Start plate repeat each 0 s -· Barcode Settings Statistics Weader Settings Created by EnVision 🔞 Recycle Bin Created 7/18/2016 5:03:25 PM Changed by EnVision Changed 7/18/2016 5:03:57 PM Last run N/A Number of runs 0 Protocol edited Changes will be saved automatically when leaving editor. Click Undo to restore values ₹. ш Press F1 for help Instrument connected EnVision



General	Optimizatio	ns						
Plate	Meas Height	Height Optimized	Delay	Window time	TRF Optimized	Flashes	Z' Optimized	Flatfield Optimized
384 OptiPlate	N/A	N/A	30	300	4/29/2015 3:3	N/A	N/A	N/A
384 AlphaPlate	N/A	N/A	30	300	4/29/2015 3:3	N/A	N/A	N/A
384 OptiPlate White	N/A	N/A	30	300	4/29/2015 3:3	N/A	N/A	N/A
Greiner 384 for BI	N/A	N/A	30	300	4/29/2015 3:3	N/A	N/A	N/A
384 well Corning Lo	N/A	N/A	30	300	4/29/2015 3:3	N/A	N/A	N/A

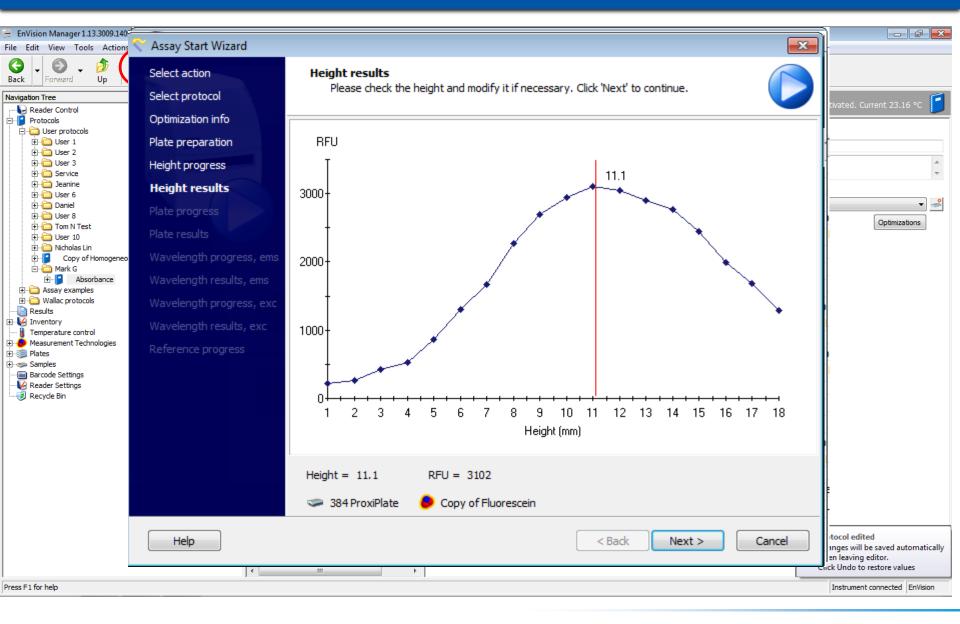


# **Existing Optimizations**



otocol - General settings Output settings Plate 1 Wells selection - G Measurement Scan: Absorb	roup 1 : Absorbance @ 405	Easurement Label (valid labels ma ABS - Absorba	-		Optimiza
Optimizations				×	
Select the label either Return back to protoc	double clicking it or choose of without any selection by (	e a label and accept it clicking Cancel button	oy clicking OK button		
Plate	Label	Wizard	Optimized	<b></b>	
96 General	Absorbance @ 405 Absorbance @ 450 Training US LUM 3	Plate ) BFP Plate	10/14/2015 11:13:11 AM (EnVision ) 10/14/2015 11:13:11 AM (EnVision ) 6/22/2016 5:15:02 AM (EnVision ) 6/22/2016 5:15:02 AM (EnVision )	Y	
			OK Cance	el	









## Results



EnVision Manager 1.13.3009.1409 - Instrument connec	cted			
le Edit View Tools Actions Help	.eu			
3   0 . 1 🔊   🙆 🖓	🔎 🖻 🗙			
Back Forward Up Start Latest	Filter Open Delete			
avigation Tree	× Deculto			
	* Results		Temperature control no	ot activated. Current 23.46 °C 👖
Protocols	Protocol:			
⊡ - Construction Description Descripti Description Description Description Description De	Protocon.	ABS @ 405		
⊞…⊡ Assay examples ⊞…⊡ Wallac protocols	Amount:	Show last 500 results		
Results				
Inventory	Time: F	From: 7 /11/2016 T 12:00:00 AM		
🔋 Temperature control	Refresh	To: 7 /18/2016 4:28:09 PM		
Measurement Technologies		7710/2010		
Plates	Assay ID Simulat		Started Finished	
	1884		7/18/2016 4:20:03 PM 7/18/2016 4:21:24 PM	
Reader Settings	1883		7/18/2016 4:18:32 PM 7/18/2016 4:19:52 PM	
Recycle Bin	1882		7/14/2016 11:03:46 AM 7/14/2016 11:08:17 AM	
Cover bin	1881		7/14/2016 8:53:19 AM 7/14/2016 8:54:40 AM	
	1880		7/14/2016 8:51:52 AM 7/14/2016 8:53:13 AM	
	1879		7/12/2016 3:23:40 PM 7/12/2016 3:25:01 PM	
	1878		7/12/2016 3:22:08 PM 7/12/2016 3:23:28 PM	
	1877 □ 1876 □		7/12/2016 3:21:18 PM 7/12/2016 3:21:47 PM	
	1876 1875		7/12/2016 1:40:34 PM 7/12/2016 1:44:00 PM	
	1875		7/11/2016 2:15:12 PM 7/11/2016 2:19:46 PM 7/11/2016 2:10:13 PM 7/11/2016 2:14:49 PM	
	1874		7/11/2016 2:10:13 PM //11/2016 2:14:49 PM 7/11/2016 2:05:12 PM 7/11/2016 2:09:47 PM	
	1873		7/11/2016 2:05:12 PM 7/11/2016 2:09:47 PM 7/11/2016 1:12:26 PM 7/11/2016 1:14:53 PM	
	1872		7/5/2016 1:12:26 PM 7/11/2016 1:14:53 PM 7/5/2016 1:02:46 PM 7/5/2016 1:07:18 PM	
	1871		7/5/2016 1:02:46 PM 7/5/2016 1:07:18 PM	
	1869		7/5/2016 12:53:54 PM 7/5/2016 12:53:12 PM	
	1869		7/5/2016 12:52:13 PM 7/5/2016 12:52:33 PM	
	1867		7/5/2016 12:48:37 PM 7/5/2016 12:48:52 PM	
	1866		7/5/2016 12:47:46 PM 7/5/2016 12:48:23 PM	
	1865		7/5/2016 11:45:20 AM 7/5/2016 11:45:38 AM	
	1864		6/29/2016 2:52:35 PM 6/29/2016 2:53:08 PM	
	1863		6/29/2016 2:49:05 PM 6/29/2016 2:49:46 PM	
	1862		6/28/2016 9:08:59 AM 6/28/2016 9:10:28 AM	
	1861		6/23/2016 5:07:02 PM 6/23/2016 5:08:11 PM	
	1860		6/23/2016 4:18:36 PM 6/23/2016 4:19:37 PM	
	1859		6/23/2016 3:07:35 PM 6/23/2016 3:10:00 PM	
	1858	Sequential Read Tb then Eu	6/23/2016 2:59:11 PM 6/23/2016 3:02:47 PM	
	1857		6/23/2016 2:56:45 PM 6/23/2016 2:58:40 PM	
	1856		6/23/2016 2:42:00 PM 6/23/2016 2:43:01 PM	
	1855		6/23/2016 2:12:16 PM 6/23/2016 2:13:37 PM	
	1854		6/23/2016 2:02:04 PM 6/23/2016 2:03:24 PM	
	1853	AlphaScreen Corning plate	6/23/2016 1:46:20 PM 6/23/2016 1:48:26 PM	



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# Advanced Training Course



#### Advanced topics for assay optimization

- Techniques for getting maximum sensitivity
- Techniques for getting maximum speed
- Optimizing assays with high energy or saturating levels of energy
- Advanced topics using the Optimization Wizard
- Cover advanced topics for each Measurement technology

#### Advanced topics for optics

- Filter and mirror training
- Understanding the light path
- PMT concepts

#### **Trouble shooting concepts**

- Concepts to determine if an error/plate pattern is caused by the EnVision or not
- Common user errors, and how to remedy
- Techniques for isolating optimization and system
   issues
- Removing passwords from locked protocols and labels

#### Advanced concepts in real-time Data Analysis

- Options with sample maps
- How to generate standard curve and apply it to data
- Blank corrected assay data
- General calculations
- Using advanced calculations

#### Concepts and programming for Liquid Dispenser

#### Hands on assay optimization

- Optimizing a problematic assays in FI, FP, and TR-FRET
- Create a new protocol for each technology to illustrate how all of the settings are used
- Create protocols that use the TRF Laser vs standard Xenon to illustrate why and when you would select the appropriate light source (If TRF Laser option is available)
- Create a inject and read assay (If dispenser is available)

#### **Advanced User Certification**



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# Hands on Training



- Hardware overview (in front of machine)
  - Proper power up sequence, and starting of software
  - o Identify all of the key pieces of hardware
  - Demonstrate filter, mirror(top and bottom), and aperture changing
  - Show plate orientation, and identify the A1 location on plate holder
  - Using stackers (If applicable)
  - Dispenser Overview (If applicable)
- Basic programming of Protocols
- Optimization of 2-3 protocols User creates the protocol
- Demonstration of applying basic calculations and exporting data after assay has been run.
- Cover what is available in an Advanced Training session