



SpectraMax XPS & SpectraMax L Microplate Reader

金萬林企業股份有限公司
產品專員 朱家均 Mavis

 MOLECULAR
DEVICES

Introduction of KimForest



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點數破萬
見樹又見林



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Customized**

applied biosystems
by Thermo Fisher Scientific

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DEVICES

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BECKMAN
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DNA TESTING TECHNOLOGIES

SARTORIUS

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DNASTAR®

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the new standard in enzyme production

gen O way

protein simple
a bioactive world



BUSINESS DIRECTORY

Looking for the world's most innovative
science and technology companies? You'll find
them here.

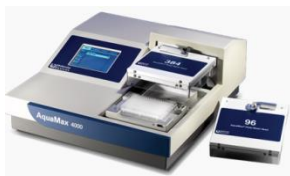


Product Portfolio in Global Market

BioResearch

- Leader in detection
- Focus on gov't and acad. research
- Modular and integrated systems

Microplate reader & Washer



Axon® Conventional Electrophysiology



GenePix® Microarray Analysis



Drug Discovery

- Leaders in niche markets
- Focus on pharma and biotech
- Complete automated solutions

ImageXpress® HCS screening



Reagents & Consumables



FLIPR® HTS screening



IonWorks® Automated Electrophysiology



Biotherapeutics

- Industry standard for cell line, protein and antibody development
- Complete automated solutions

ClonePix™ 2 Cell Line Development



CloneSelect™ Imager



QPix™ - Ab & Protein Development



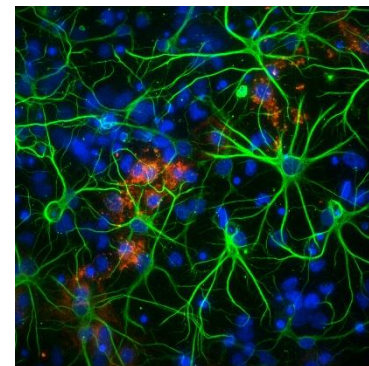
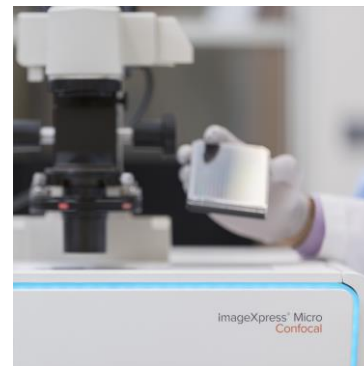
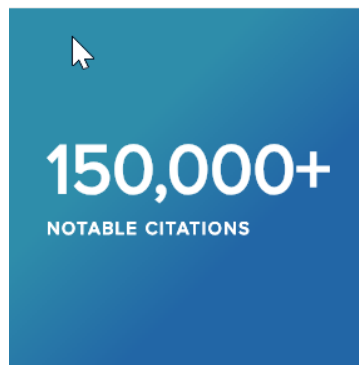
DispenCell™ Single-Cell Dispenser



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About Us

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Latest Citations

Basic Research

Find Your Own Microplate Reader



FlexStation 3



i3x



**iD3s (w/o screen)
/iD5e**



Mini



FilterMax



M2-M5

SpectraMax XPS Multi Mode Reader



Microplate

- ✓ Fluorescence Intensity
- ✓ Time-Resolved Fluorescence
- ✓ Luminescence

SpectraMax XPS Specifications

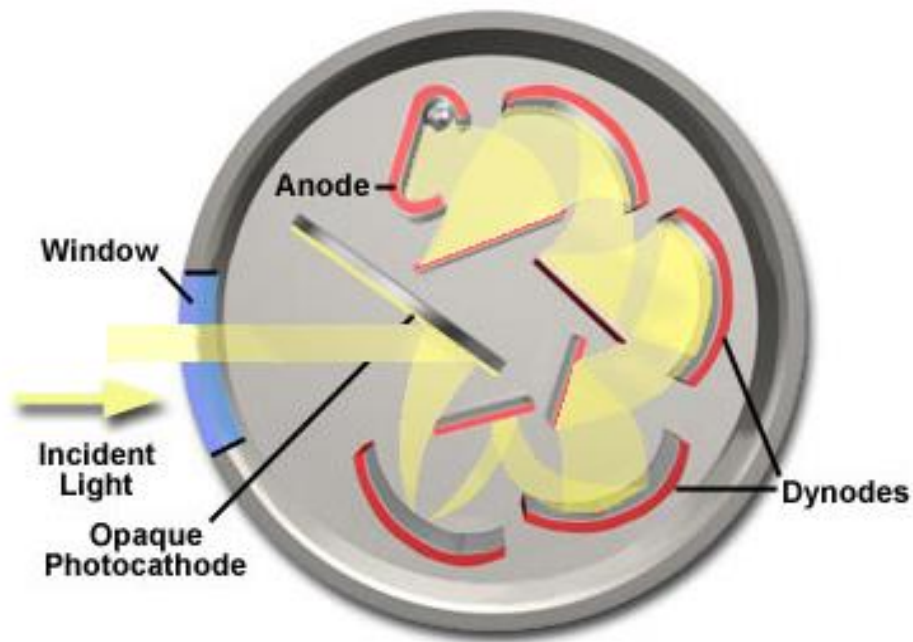
- **Microplates:** 6, 12, 24, 48, 96, and 384 well
- **Light source:**
 - ✓ **Xenon flash lamp** (1 joule/ flash)
 - ✓ Lifetime of **1 billion flashes** ~ 1 million endpoint microplates
- **Wavelengths Supported**
 - ✓ Fluorescence Intensity: EX: **250 – 850** nm
EM: **360 – 850** nm
 - ✓ Time-Resolved Fluorescence: **360 – 850** nm
 - ✓ Luminescence: **360 – 850** nm

SpectraMax XPS Specifications

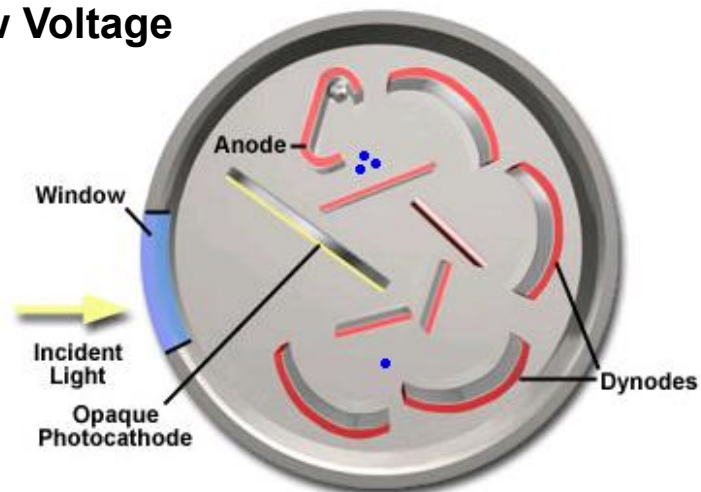
- **Dual monochromators**
 - ✓ Two scanning monochromators determine optimal excitation and emission settings
 - ✓ No need for expensive filters
- **Detectors**
 - ✓ Photomultiplier (PMT)
- **Shaker time:** 1 to 999 seconds
- **Temperature control:**
 - ✓ 4°C above ambient to 45°C
 - ✓ Temperature regulation: 4 sensors

Patented AutoPMT Optimization System

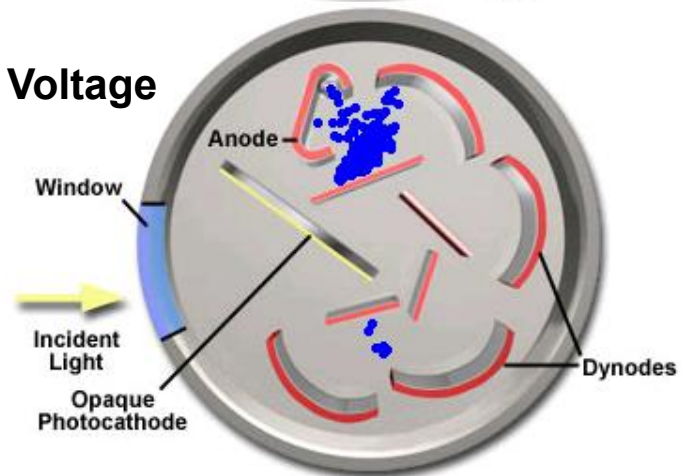
Photoelectric Effect



Low Voltage



High Voltage



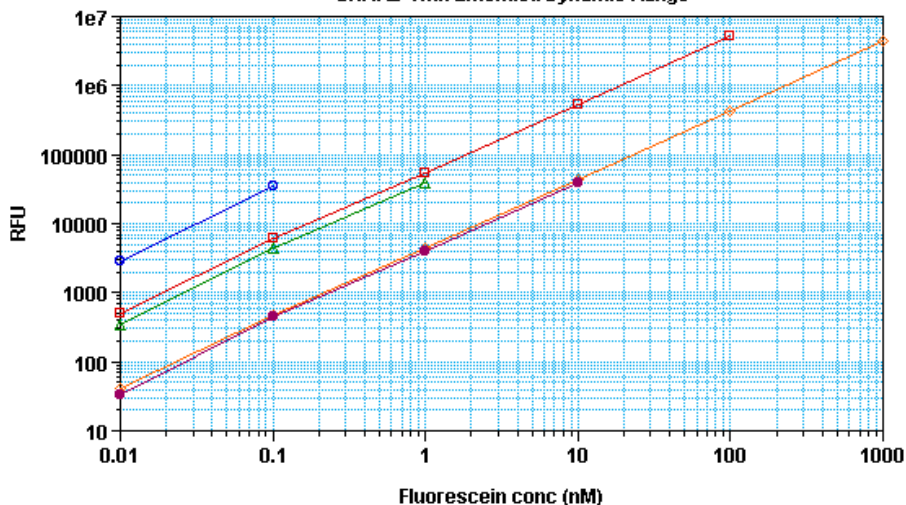
Stop

Gain

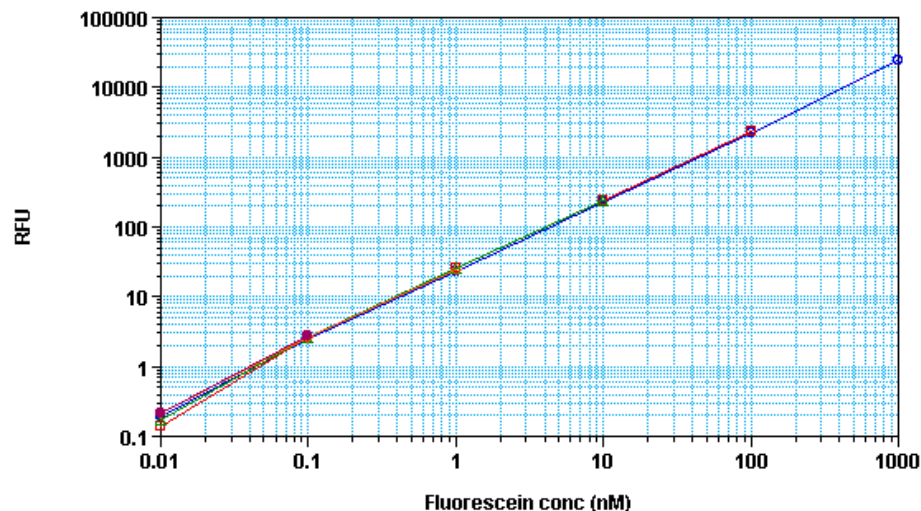


Patented AutoPMT Optimization System

SAFIRE with Extended Dynamic Range



- Plot#1 (0.1 nM Max@SAFIRE ExDyRa: Concentration vs MeanValue)
- Plot#2 (100 nM Max@SAFIRE ExDyRa: Concentration vs MeanValue)
- Plot#3 (1.0 nM Max@SAFIRE ExDyRa: Concentration vs MeanValue)
- Plot#4 (1000 nM Max@SAFIRE ExDyRa: Concentration vs MeanValue)
- Plot#5 (10 nM Max@SAFIRE ExDyRa: Concentration vs MeanValue)



- Plot#1 (1000Max: Concentration vs MeanValue)
- Plot#2 (100Max: Concentration vs MeanValue)
- Plot#3 (10Max: Concentration vs MeanValue)
- Plot#4 (1.0Max: Concentration vs MeanValue)
- Plot#5 (0.1Max: Concentration vs MeanValue)

✓自動切換最適電壓

-允許最大濃度範圍樣品在同一盤檢測中

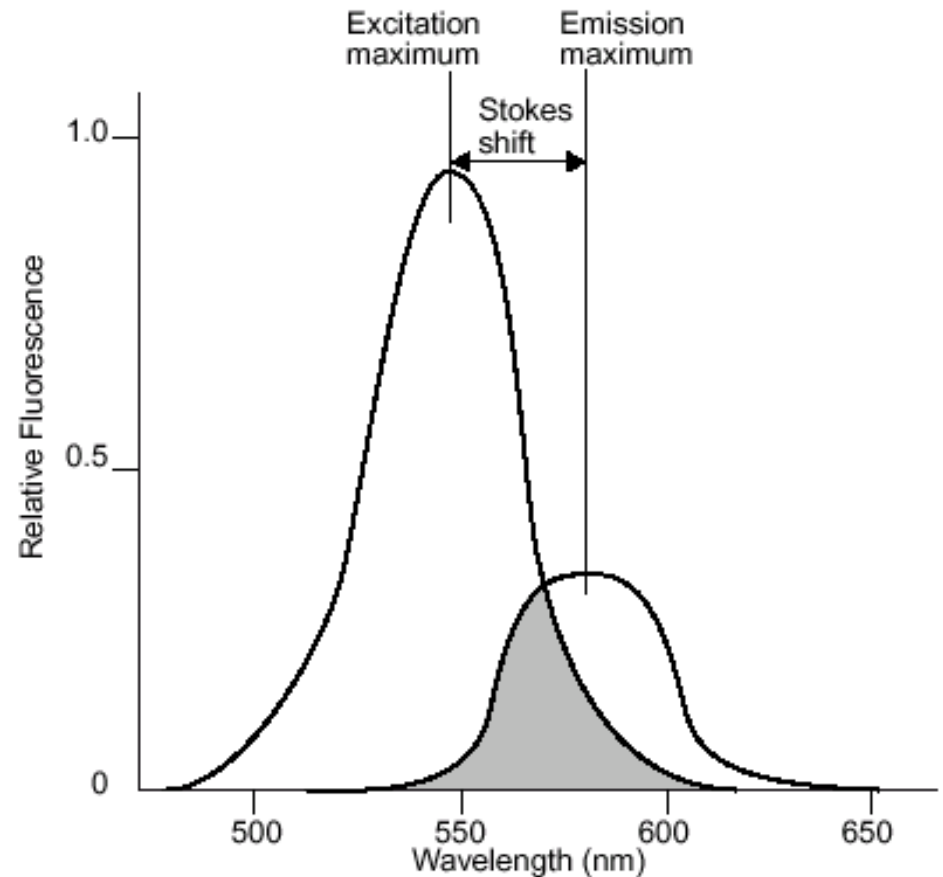
✓自動換算數據

-無須帶入不同曲線換算gain值，避免繁瑣工作

Auto Cutoff - Emission Cutoff Filter

Why Cut-Off Filters?

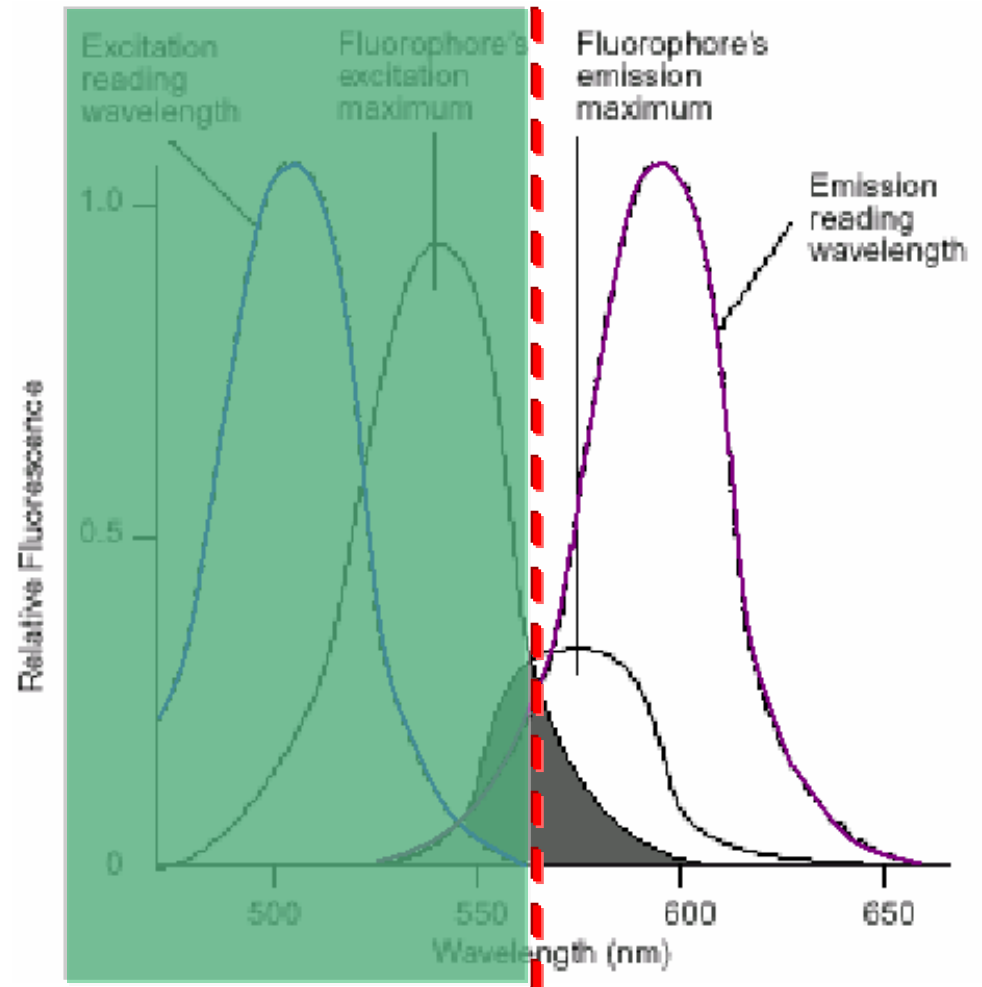
- Excitation light is **10,000 X** brighter than Emission light.
- Spectral separation needed to reduce interference of Ex with Em light.



Auto Cutoff - Emission Cutoff Filter

Optimizing Wavelength Selection

- Using Ex wavelength below Ex max and Em wavelength above Em max may provide best signal/noise ratio.
- Cutoffs cause less Ex light mixing with Em light at the PMT.



Cutoff filter

Applications of SpectraMax XPS

- **Fluorescence assays:**

- ✓ Nucleic acid quantitation
- ✓ Enzyme assay
- ✓ ELISA
- ✓ Cell proliferation & viability
- ✓ GFP
- ✓ SNP
- ✓ FRET assay

SpectraMax L Microplate Reader



Wavelength:
380 – 630 nm

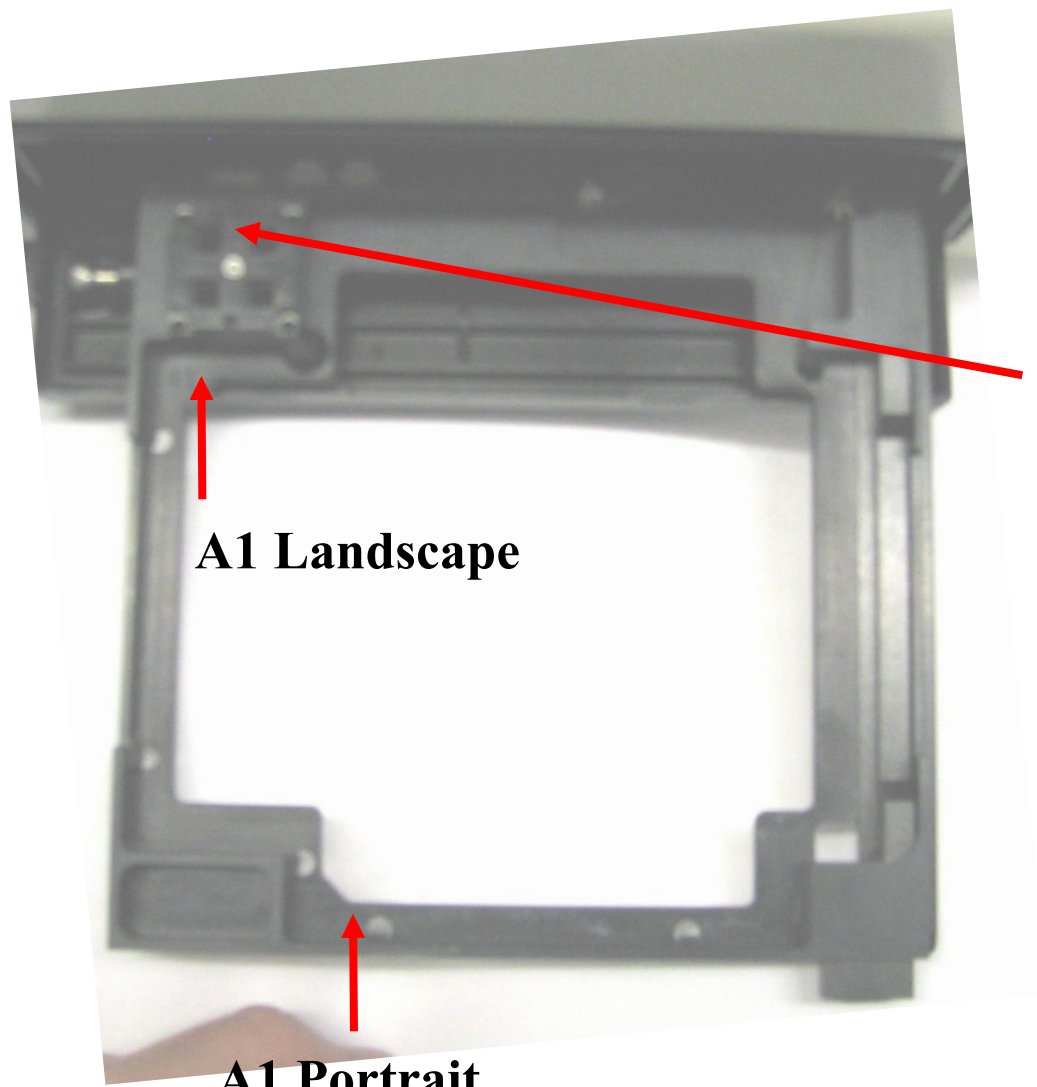
Microplate formats:
96 and 384 well

Read mode:
Top read - Luminescence

Read type:

- Endpoint**
- Dual-Read**
- Kinetic**
- Fast Kinetic**

Plate Drawer



1 Calibration Sources

A1 Landscape

A1 Portrait

Features of SpectraMax L



(1) Auto-PMT :

- High sensitivity luminescence detection
<0.2 fg firefly luciferase per well
- Simultaneous **photon counting** and **analog detection** for extended dynamic range

(2) Aperture Design :

Low Background and crosstalk

- **3×10^{-5}** (white plate)

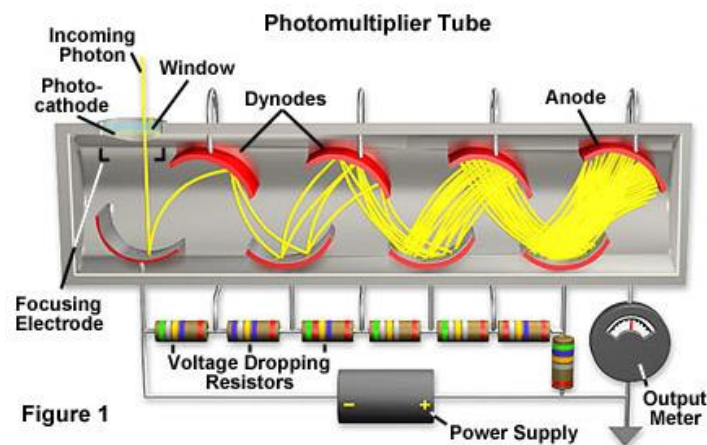
(3) Upgradeable :

- Dual injectors for flash luminescence
- Multi-detector configurations for higher throughput



Luminescence Detection by PMT

- Measuring Luminescence :
 - Photomultiplier tubes (PMT) convert incoming photons to electrons
 - Incoming photon strikes photocathode → generates electron
 - Electron flows through a series of electron multipliers (dynodes) to the anode
 - Current flowing from the anode is proportional to the number of photons at the photocathode
 - Amount of amplification a PMT can produce depends on 1) the number of dynodes and 2) voltage applied to it



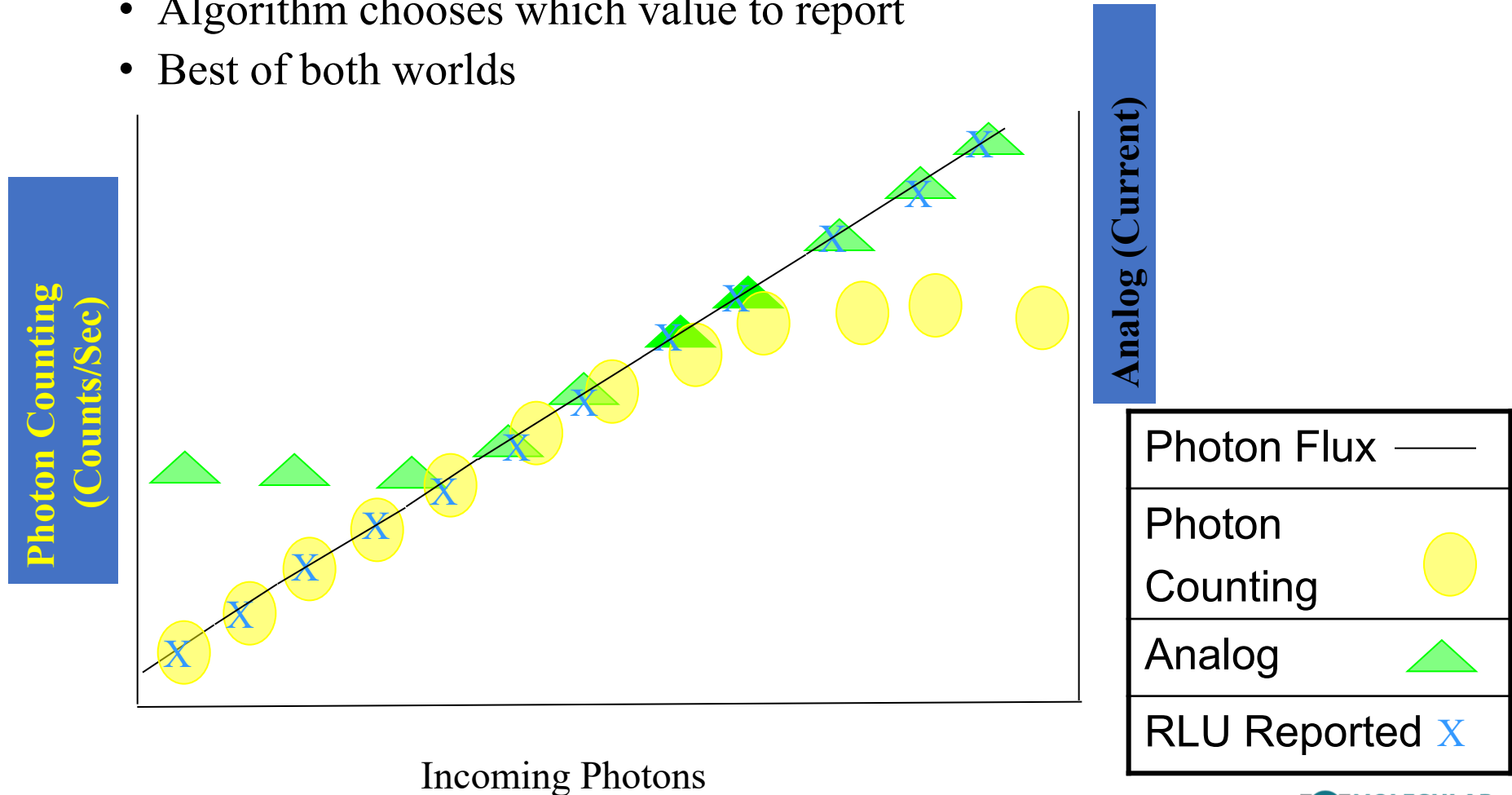
PMT Sensitivity:

How PMT Current Is Measured

- **Analog**
 - Wide dynamic range achieved by converting PMT pulses into current
 - Downside – Loss of sensitivity because of difficulties in separating signal from noise
- **Photon Counting**
 - Excellent sensitivity achieved by converting PMT signal into digital pulses, pulses are then digitally filtered, only pulses larger than a threshold are counted.
 - Digital filtering of pulses does a great job of discriminating between signal and noise, resulting is high sensitivity
 - Downside – Photon counting has a limited linear range compared to analog detection

PMT Sensitivity: MaxRange PMT Setting

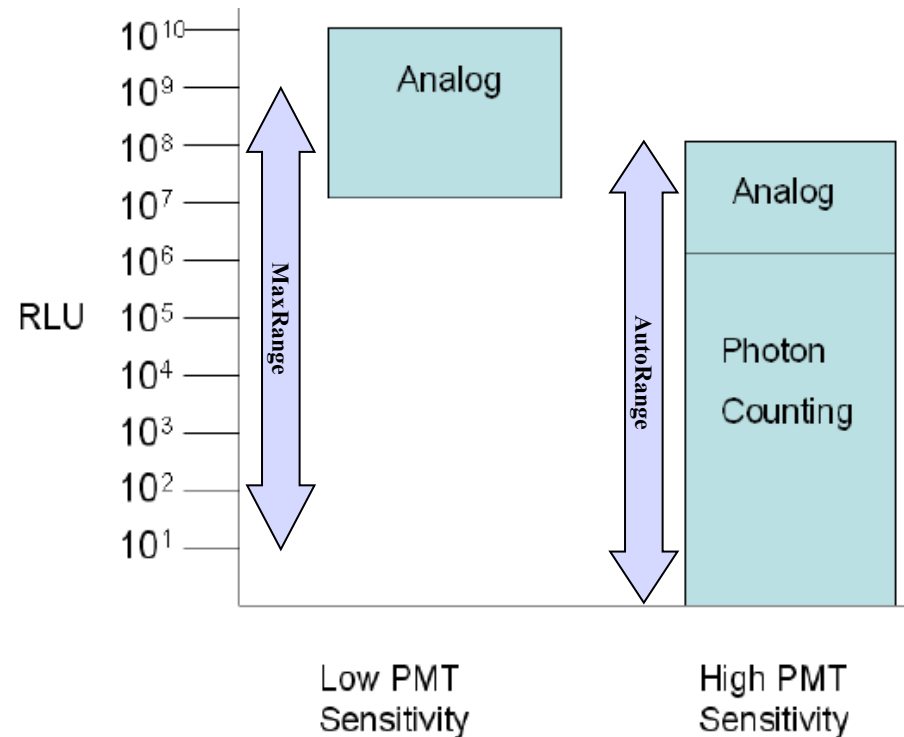
- Simultaneous Photon Counting and Analog Detection
 - Algorithm chooses which value to report
 - Best of both worlds



Extended Dynamic Range

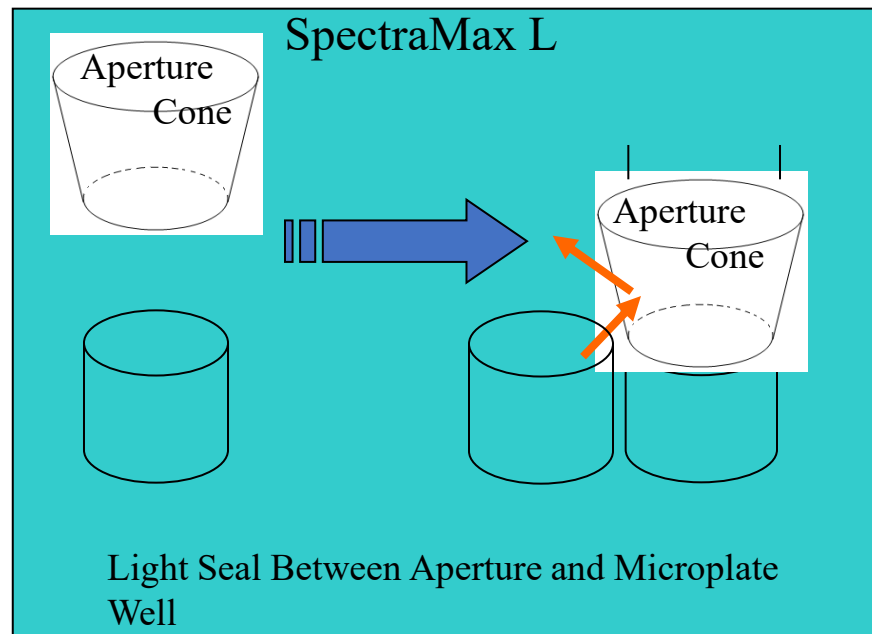
4 PMT Settings

- **Analog Only:** Ideal for very bright signals above 1.6×10^7 RLU (low PMT voltage)
- **Photon Counting:** Ideal for very dim and medium signals below 2.5×10^6 RLU (digital)
- **AutoRange:** Extends range of Photon Counting mode by adding Analog (high PMT voltage)
- **MaxRange:** Combines AutoRange and Analog Only modes
 - Captures entire dynamic range
 - **10 to 10^9 RLU**



SpectraMax L Aperture Design

- By design, the SpectraMax[®] L luminometer offers **higher sensitivity** and **lower crosstalk** by capturing the maximum amount of light from a test well and physically limiting light from adjacent wells.
- The aperture design does limit the types of plates that can be used on the system. (see known issues)



A Configurable System

Available in different configurations
and depot-upgradeable for
changing assay requirements

- Single detector (PMT):
 - 0 or 2 injectors
- 2-detectors:
 - 0, 2, or 4 injectors
- 6-detectors:
 - 0 or 12 injectors
- With injectors → flash
- Without injectors → glow

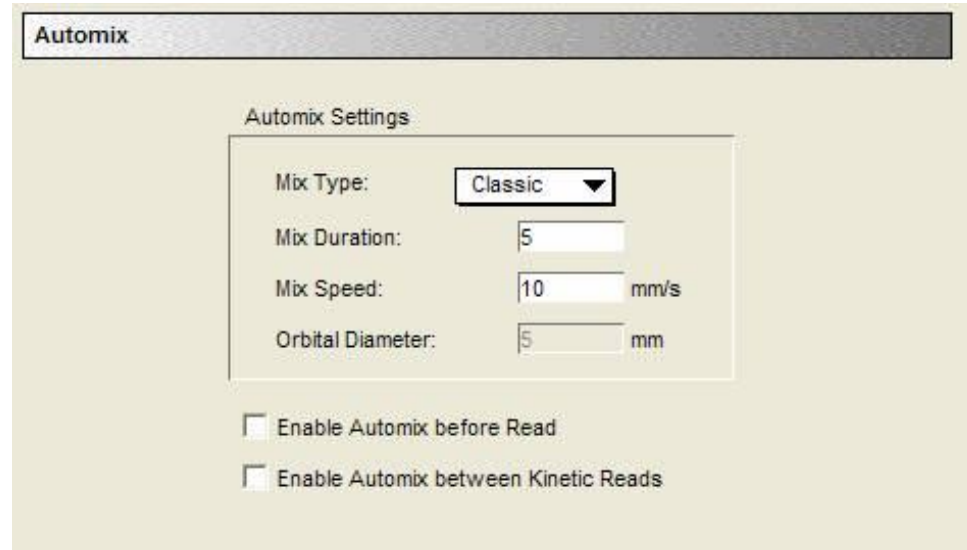


Types of Luminescence Assays

- **Glow**
 - Reaction kinetics are slow
 - Reagents can be added by hand
 - Plate will emit light for 20 minutes up to several hours
- **Flash**
 - Reaction kinetics fast enough to require an injector
 - Slow enough for injection followed by readhead movement and then read
- **“Extremely Fast Flash”**
 - Requires an injector in same position as readhead
 - Acridinium ester

Four Mode Mixing

- **Classic**
 - Same as AutoMix on most SpectraMax readers
 - Shakes along axis from front of instrument to back
- **Single Axis**
 - Shakes along same axis as Classic, but frequency doesn't change
- **Dual Axis**
 - Shakes In “L” pattern
- **Orbital**
 - Shakes in Circle pattern



The screenshot shows a software window titled "Automix". Inside, there is a section labeled "Automix Settings". Within this section, there is a "Mix Type:" label followed by a dropdown menu currently set to "Classic". Below this are three input fields: "Mix Duration:" with the value "5", "Mix Speed:" with the value "10" and the unit "mm/s", and "Orbital Diameter:" with the value "5" and the unit "mm". At the bottom of the settings section, there are two checkboxes: "Enable Automix before Read" and "Enable Automix between Kinetic Reads", both of which are currently unchecked.

SpectraMax® L Applications

- ELISAs and Immunoassays
- Reporter Gene Assays
- Bioluminescence Resonance Energy Transfer (BRET)
- Enzyme Assays
- Cell Viability, Proliferation, and Cytotoxicity
- Aequorin Assays
- ATP Detection
- Mycoplasma monitoring



SoftMax Pro 7 *Features*

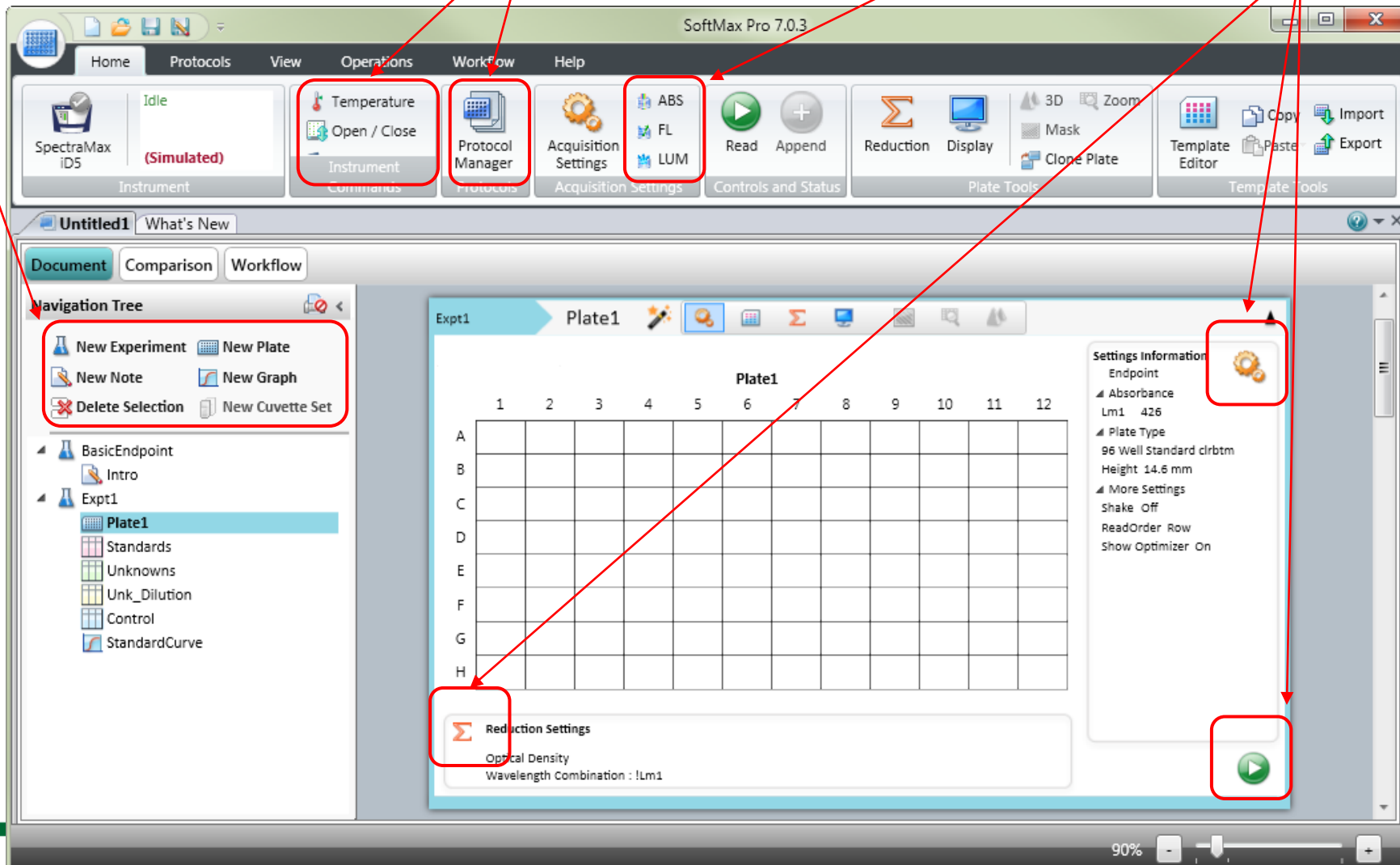
Software Interface

The New Experiment icons moved into the Navigation Tree

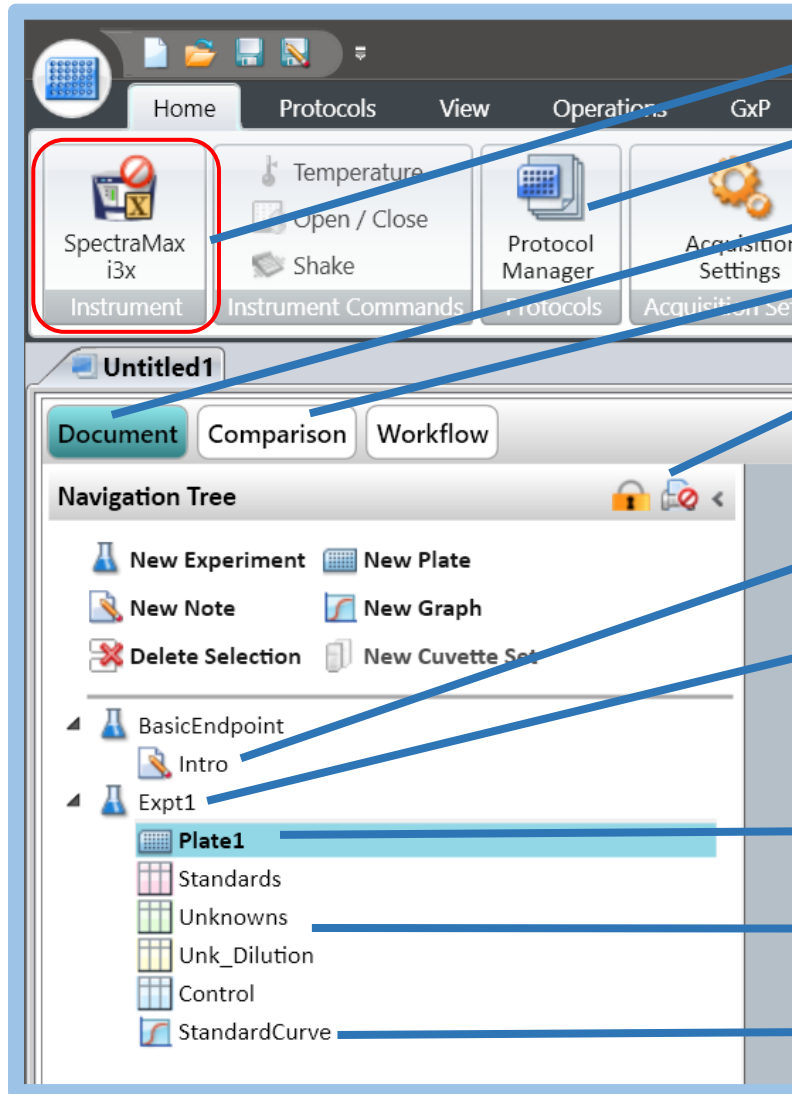
Frequently used Icons moved from the operations tab into the Home tab

New Quick Start reads

Easy access buttons



Project Oriented Interface



軟體連線情況

檔案管理

分頁模式

比較模式

可獨立設定”編輯性” or ”
是否列印”

註解/摘要報告

不同實驗混編

原始資料(Raw Data)

分析資料(Meta Data)

曲線擬合/統計

Advanced Acquisition

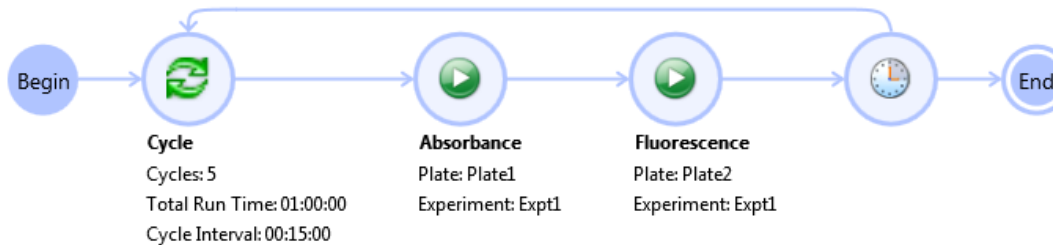


Multitask Kinetics Benefits

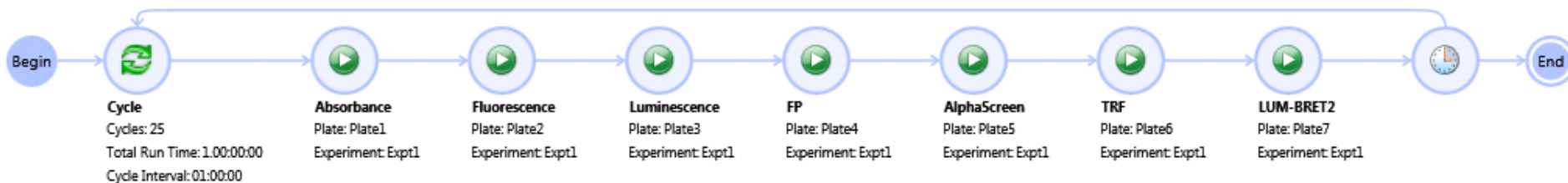
- Customize kinetic cycle tasks using the new Workflow Editor
 - **Say goodbye to Excel Workflows!**
- Run multi-day kinetic assays
- Works with all supported readers that have kinetic mode

Multitask Kinetics Benefits

- **Set up multi-mode kinetic reads**
 - Alternate between absorbance and fluorescence

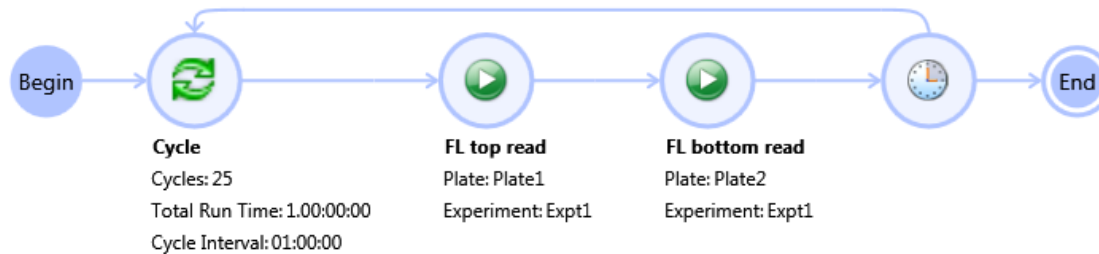


- Alternate between absorbance, fluorescence, luminescence, FP, TRF, cartridges, etc

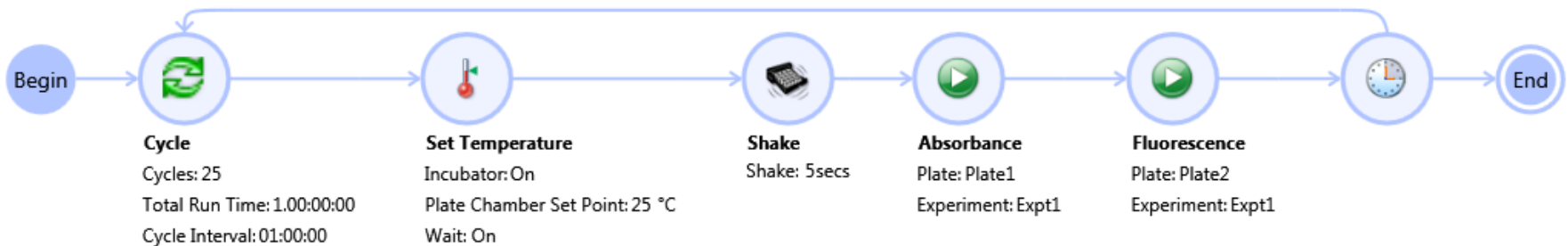


Multitask Kinetics Benefits

- **Can't decide if top read is better than bottom read? Try both!**



- **Add in tasks: delay, shake, open drawer**



Advanced Analysis



Powerful Formula system

SoftMax Pro 7.1.2

Home Protocols View Operations Workflow Help

SpectraMax i3x (Simulated) Instrument

Temperature Open / Close Shake Instrument Commands

Protocol Manager Protocols

Acquisition Settings Acquisition Settings

Read Append Controls and Status

Reduction Display Mask Clone Plate Plate Tools

Template Editor Copy Import Paste Export Template Tools

Plate1 Plate1-1 Plate1-2

Plate1

	1	2	3	4	5	6	7
A	0.087	0.089	3.562	0.566	2.884	2.188	
B	0.238	0.243	3.264	0.667	2.866	2.186	
C	0.636	0.649	3.525	3.569	0.848	2.941	
D	1.475	1.490	3.888	3.234	0.777	2.445	
E	2.603	2.629	0.590	1.668	0.232	3.303	
F	3.399	3.365	0.618	1.998	0.282	3.732	
G	3.702	3.665	2.867	2.899	0.931	0.868	
H	3.779	3.817	2.767	2.812	0.777	0.352	

Reduction Settings

Optical Density
Wavelength Combination : !Lm1

Plate1 Plate1-1 Plate1-2

Plate1-1

	1	2	3	4	5	6	7
A	Blue	Blue	Red	Blue	Red	Blue	
B	Blue	Blue	Red	Blue	Red	Blue	
C	Blue	Blue	Red	Red	Blue	Red	
D	Blue	Blue	Red	Red	Blue	Blue	
E	Red	Red	Blue	Blue	Blue	Red	
F	Red	Red	Blue	Blue	Blue	Red	
G	Red	Red	Red	Red	Blue	Blue	
H	Red	Red	Red	Red	Blue	Blue	

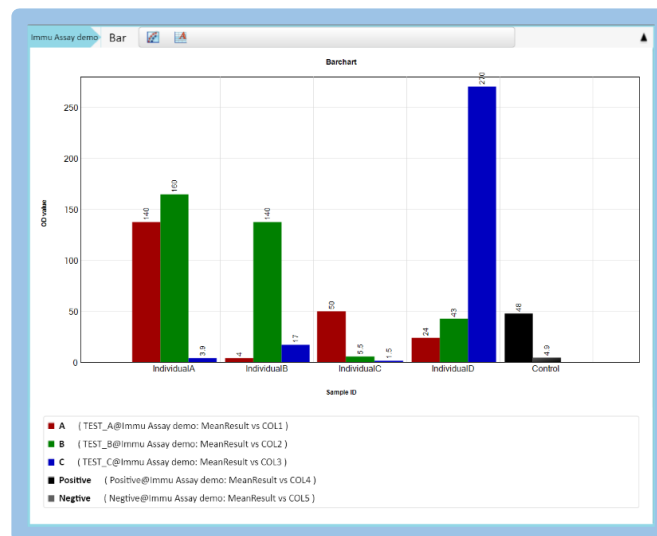
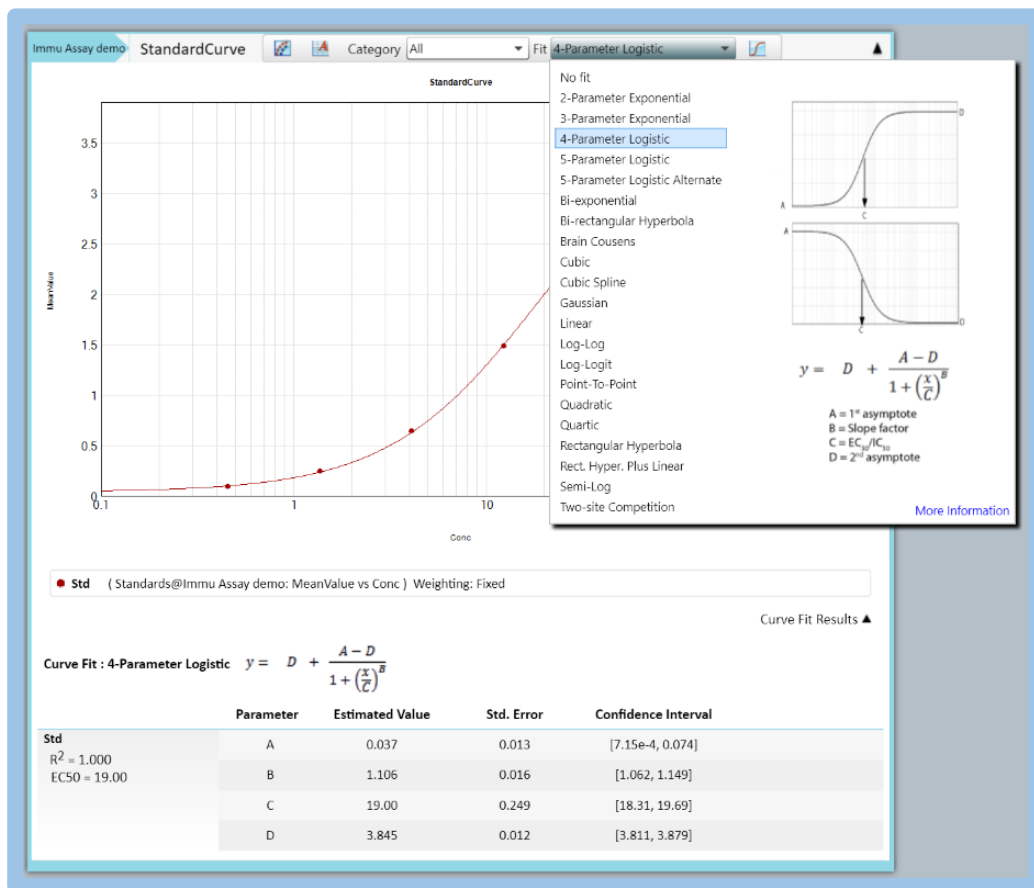
Reduction Settings

Optical Density
Wavelength Combination : If (Lm1>2.5,2,0)

✓ RAW data

✓ QC Dashboard

Adaptable Graph system



Suspend Calculations

Recalculate Now

Function Editor

Calculations

$y = ()$

21 Predefined
Curve Fitting

Bar Chart

New Function
Editor

Powerful Formula system

Immu Assay demo
TEST_A

Sample	Wells	COL1	Value	MeanValue	Risk Class 風險評比
01	A3	1	3.562	3.413	危險
	B3		3.264		
02	A4	2	0.566	0.617	未檢出
	B4		0.667		
03	A5	3	2.884	2.875	偏高
	B5		2.866		
04	A6	4	2.188	2.187	正常
	B6		2.186		

judgement criteria description (公式寫法)
//巢狀條件式，與Excel寫法極為相近

MeanValue>DUL@Positive,"危險",If (MeanValue>DLL@Positive,If ("偏高",If (MeanValue<NDL@Negative,"未檢出","正常")))

- 檢出值 > Pos上限者 ==> 危險
- 檢出值 > Pos下限者 ==> 偏高
- 檢出值 < Pos下限者 ==> 正常
- 檢出值 < Neg限者 ==> 未檢出

Formula
☒ Syntax Helper

MeanValue>DUL@Positive,"危險",If (MeanValue>DLL@Positive,If ("偏高",If (MeanValue<NDL@Negative,"未檢出","正常"))

Mathematical

Statistical

Comparison

Condition

Peak analysis

Array analysis

Sample	Wells	Sample#	Lm_run	OD_read	PEAK_Judge
01	B1	1	401	2.527	-
			431	3.174	PEAK@2
			461	0.726	
			491	0.090	
			521	2.352	
			551	3.661	PEAK@6
			581	0.441	PEAK@7
			611	2.039	
			641	3.771	PEAK@9
			671	3.093	
			701	2.783	
			731	2.488	-

Number of read point 12

Thank you for your attention

產品專員 朱家均 Mavis

服務電話：0955-586-009

Email: mavis.chu@kimforest.com

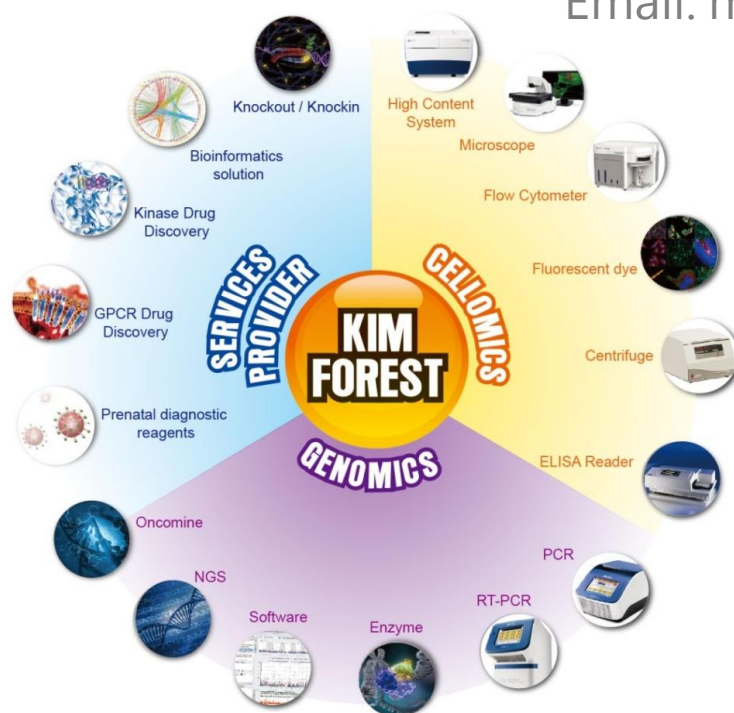


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