

# DualView- $\Lambda$ ™

## Dual-Channel Simultaneous-Imaging System

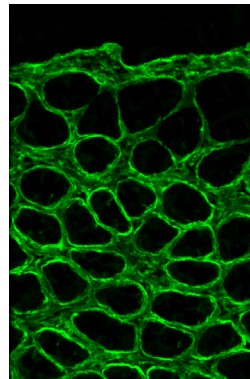
The DV $\Lambda$  is an emission splitting system enabling a user to acquire two spatially identical but spectrally distinct images simultaneously. What distinguishes the DV $\Lambda$  from the DV2 is that DV $\Lambda$  can accommodate image sensors up to a 22 mm diagonal. Simultaneous, multichannel imaging is essential for colocalization, ratiometric analysis, or polarization studies of a single emission wavelength channel.



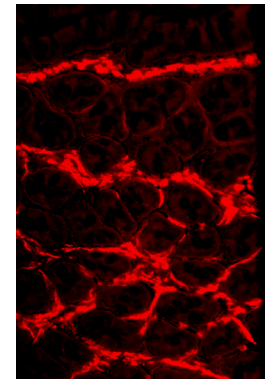
### Features

- Accommodates image sensors up to 22 mm diagonal including most sCMOS sensors
- Simultaneous acquisition of two full-field emission images
- Emission can be separated by wavelength, polarization, or amplitude
- Exchangeable filter cube use standard 25 mm diameter filters and polarizers
- Filter cubes can be configured by the user for different filters and beam splitters
- Easily mounts to most microscope side ports
- Incorporates rectangular aperture to define a region of interest
- Image processing of DualView data is available in leading microscopy packages

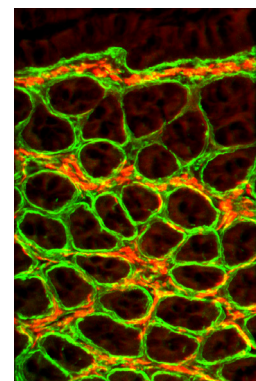
Green



Red



Green/Red Overlay



## DVΛ Specifications

Wavelength range	450 to 900 nm
Transmission per image	>88% (does not include filter)
Operation temperature	10°C to 37°C
Detector attachment	C-mount (male)
Front attachment	C-mount (female)
External mounting option	¼-20 tapped hole on back of unit
Dimensions	320 mm long, 110 mm wide, 79 mm tall
Weight	1.5 kg
Filters	
Emission/Barrier Dichroic	25 mm diameter 25-26 mm wide, 36-40 mm long, 1-3 mm deep
	Recommend Chroma Ultra Flat range in size 26x38x2 mm for best results
Supports LightSpeed™ mode of Evolve™ 512 Delta with firmware newer than 12.2014	

## Applications

- Fluorescence resonance energy transfer (FRET) imaging
- Polarized FRET analysis
- Calcium imaging with fluo-3/Fura Red™ (Molecular Probes) or dual-emission indo-1 imaging
- Fluorescence polarization/anisotropy imaging
- Simultaneous fluorescence/DIC imaging
- Drug discovery with Cy3/Cy5
- Single-molecule fluorescence (SMF) imaging
- pH imaging with SNARF
- Multiwavelength total internal reflection fluorescence (TIRF) imaging
- Voltage sensing with di-4-ANEPPS
- Fluorescence *in situ* hybridization (FISH) imaging
- cAMP imaging with FICRhR
- Multichannel confocal microscopy when used in conjunction with a spinning-disk confocal

Note: All specifications are typical and subject to change.

