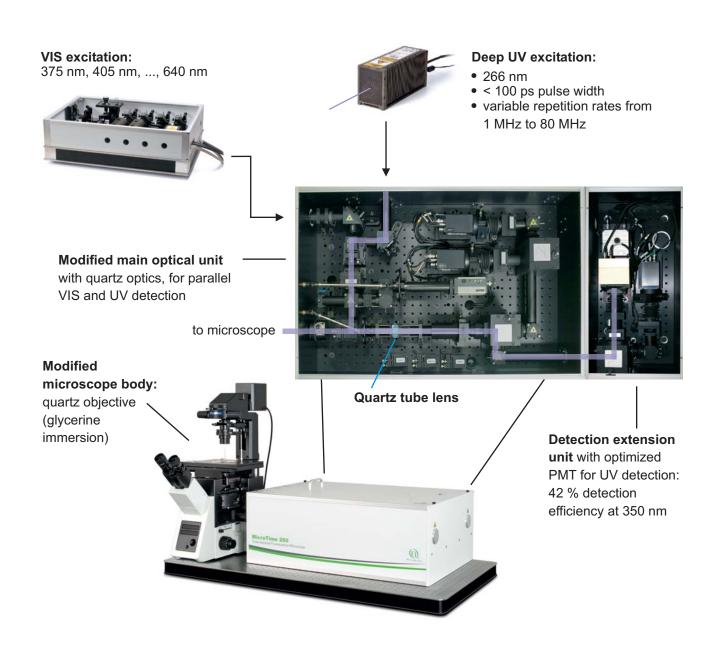
Deep UV Extension for the MicroTime 200



- Excitation at 266 nm with compact diode based laser
- Quartz lenses and UV compatible optics
- UV objective and UV detector for optimal performance
- Parallel UV and VIS detection
- Label-free Fluorescence Lifetime Imaging Microscopy (FLIM)
- Label-free protein detection through intrinsic fluorescence



Deep UV Extension for the MicroTime 200

Applications Dyes with high quantum efficiency FLIM with streptavidin-coated beads in the UV: laser dyes as single (Ø 500 nm, 1 streptavidin = 24 tryptophans) emitters immobilized on quartz coverslips para-Terphenyl para-Terphenyl in water, 33 nM Normalized Autocorrelation 1.2 1,0 0,8 1.5 x 1.5 µm 0,6 0,4 0.2 0,0 10 Lag Time [ms] • diffusion time: $t(diff) = 57 \mu s$ • molecular brightness: 510 Hz per • data acquisition time: approx. 10 min 13 x 13 µm Label-free FLIM Benchmark: in the deep UV FCS in the deep UV Label-free FLIM with biological cells Streptavidin-coated beads → aromatic amino acids within the proteins (Ø 120 nm) (mainly tryptophans) become visible approx. 1000 streptavidin per bead nucleus (1 streptavidin = 24 tryptophans) nucleoli cytoplasm Streptavidin beads, Normalized Autocorrelation in water, Ø 120 nm 3000 0,8 0,6 0,4 0,2 3T3-L1 cells, fixed (methanol) 80 x 80 µm 10⁻² 10⁻¹ 10° 10¹ Lag Time [ms] HEK293 cells, fixed (paraformaldehyde) • diffusion time: t(diff) = 16 ms 73 x 79 µm • molecular brightness: 1.2 kHz per Sample courtesy of Astrid Tannert, • data acquisition time: approx. 5 min University of Leipzig, Germany • excitation: 266 nm (PicoQuant), 20 MHz, < 100 ps pulse width • quartz objective, 40x, NA 0.6, glycerine • optical filters: Z266RDC, 300 nm longpass detection: PMT, 42 % detection efficiency at 350 nm

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