# **Everspark**

#### The long-lasting blinking buffer for dSTORM microscopy



**Everspark** is a super-resolution microscopy buffer that offers prolonged fluorophore blinking in dSTORM microscopy: mount your sample, and image repeatedly for months!

Everspark 1.0 | Everspark 2.0

	Everspark i.o	LVCISPAIR L.O
Compatible fluorophores	yellow to far-red	green, yellow & far-red Optimized for <b>3-colour</b> dSTORM imaging
Blinking stability (after mounting)	2 months	3.5 months
Number of blinking events	+	+++

#### **Key features**

- 1. Simplify your imaging workflow by imaging as many times as you want for 2.5months without replacing the buffer
- **2. Multicolor (including green)**Compatible with green, red & far-red dyes (i.e. AF488, JF549, CF568, JF646, AF647 & Cy5)
- **3. Ready to use**Use it straight from the vial

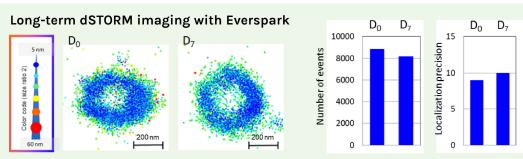
#### 4. Up to 6 months performance

Individualized packaging for optimal longevity

## Compatible microscopy techniques:

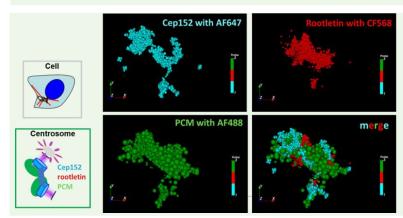
- PALM
- MINFLUX
- dSTORM
- HiLo & TIRF modalities

#### Results



Left: Two centrosomes imaged the same Day (D0) and 7 days after mounting (D7) on the same slide stored in the dark at 4°C and visualised using a colour-coded scale encoded with the IGOR software, where each point appears as a function of its localisation precision (5 to 60 nm; inverted rainbow colour scale). Labeling: Distal-appendages detected by immunofluorescence with AF647 in RPE-1 cells. Right: The number of blinking events per centrosome and the median of the localization precision in nm are presented for each serie of 50,000 images recorded at D0 and D7.

Credits: Camille Fourneaux, Karine Monier, CRCL, Lyon



### Co-localization studies using three-colour 3D dSTORM imaging with Everspark 2.0

Centrosomal proteins Cep152, rootletin and PCM labelled with AF647, CF568 & AF488 respectively were imaged on a Vutara VXL (Bruker) in Everspark 2.0 buffer before 3D dSTORM reconstruction

Credits: Karine Monier, INMG-PGNM, Lyon

