

### **DNA-PAINT KIT**

# MASSIVE-TAG-Q anti-GFP

Expiration after 6 months

(For research use only)

#### **CONTENT**

#### SINGLE DOMAIN ANTIBODY

- FluoTagQ® anti-GFP (clone: 1H1) + Docking site 3 (To be measured with Imager 3)
- Concentration: 5 μM Protein, 5 μM DNA (1 DNA strand per protein)
- Volume: 100 µL
- Storage buffer: PBS, 50 % glycerol + 0.05 % NaN<sub>3</sub>
- Storage: -20 °C
- Recommended dilution: 1:200 1:500 (for optimal results the dilution needs to be optimized depending on the target accessibility and expression level)

### **IMAGERS**

- Imager 3 Cy3B or ATTO 655
- Concentration: 1 μM in TE buffer (10 mM Tris, 1 mM EDTA, pH 8)
- Volume: 300 μL
- Storage: 20 °C (1 μM imager solutions are stable for multiple freeze-and-thaw cycles)

Optional: Prepare 50  $\mu$ L aliquots and store at -20 °C. Working aliquots can be stored at 4 °C for short-term or -20 °C for long term

Note: Further dilutions should be prepared fresh before use. Low imager concentrations are not stable in plastic tubes.

### **BUFFERS**

- Antibody incubation buffer, 50 mL, store at 2-8 °C
  Note: For longer-term storage we recommend to store aliquots at -20 °C.
- Washing buffer (10x), 20 mL, store at room temperature (to be diluted 1:10 in water before use)
- Imaging buffer, 50 mL, store at room temperature

## SAMPLE PREP. PROTOCOL

- 1. Prepare sample using a protocol optimized for your GFP-tagged protein target.
- 2. Block cells in Antibody incubation buffer for ~30 minutes.
- 3. Dilute sdAbs in Antibody incubation buffer.
- 4. Incubate for 30 60 min at room temperature
- 5. Wash three times with washing buffer (1×).
- 6. Optional: Incubate fiducial markers.
- 7. Wash once with imaging buffer before adding the final imaging solution with imager strands.
- 8. Before imaging: Add imager strands diluted in imaging buffer. We recommend a starting concentration of 1 nM. However, the optimum imager concentration strongly depends on the target and labeling density. Thus, the imager concentration should be adjusted such that distinct single molecule blinking events can be observed
- 9. After imaging, exchange buffer to washing buffer (1×) for storage at 4 °C.

## IMAGING PARAMETERS

- Exposure time: 100 150 ms
- Laser-Intensity: ~200 W/cm² (561 nm) and ~300 W/cm² (640 nm). This intensity might vary due to different illumination depths/modes (TIRF/HILO). For dense targets we recommend increasing the laser power to enhance blinking.
- Total imaging time/target: 30 min (Depends on target density and applied imager concentration)
- Temperature: The kit is optimized for image acquisition at 21 25 °C. At higher temperatures shorter exposure times and higher laser powers are required.