

Antibodies for In Vivo Mouse Platelet Labeling

X649

Product Information

Catalog Number: X649

Isotype: Rat IgG (Wistar), derivatised

Contents: 100 µg DyLight649-labelled immunoglobulin derivative in phosphate buffered

saline containing 0.2% BSA

Concentration: 0.1 mg/ml

For research use only, not for diagnostic or therapeutic use. This product is no medical device.

Specificity: This antibody preparation contains a rat IgG derivative against the GPIbβ subunit of the murine platelet/megakaryocyte-specific GPIb-V-IX complex. The modified antibody has been optimized for the easy and stable *in vivo* labelling of circulating platelets in mice. At the recommended concentration (0.1 μg/g body weight), X-649 is non-cytotoxic and does not interfere with platelet adhesion and aggregation *in vivo*. Also, X-649 at the recommended concentration does not alter platelet adhesion on collagen/von Willebrand factor *in vitro*. ¹ *In vivo* platelet labeling has been used for intravital microscopical analysis of platelet involvement in pathological processes, such as thrombosis. ^{2,3}

Preparation and Storage: The antibodies were purified from hybridoma cell culture supernatant by Protein G-Sepharose chromatography and biochemically modified. Stable for six months from date of shipment when stored at 4°C in the dark. KEEP STERILE, the preparation contains no preservative.

Usage: This preparation is optimized for rapid and stable *in vivo* labelling of mouse platelets. Use 0.1 μ g (1 μ l) X-649 per gram body weight in an appropriate volume (50 - 200 μ l) of sterile PBS for i.v. injection. Platelet can be visualized using appropriate filter sets and a light-sensitive camera. Recommended exposure time: 200 – 400 ms.

References

- 1. emfret Analytics, unpublished
- 2. Falati *et al.* (2002) Real-time in vivo imaging of platelets, tissue factor and fibrin during arterial thrombus formation in the mouse. Nature Medicine **8**, 1175-1181
- 3. Grosse *et al.*, (2007): An EF hand mutation in Stim1 causes premature platelet activation and bleeding in mice. J Clin Invest. **117**, 3540-3550