

## **Technical Information**

# Flamma<sup>®</sup> PS Bead series

#### Overview

Flamma® PS bead series are fluorescent spherical particles in the colloidal size made up of high-grade polystyrene (PS) doped with our proprietary dyes. Polystyrene beads are not cytotoxic, and they diffuse minimally from the injection site and persist for long periods in nerve cells. Our fluorescent beads display excellent optical properties yet negligible photobleaching under the intense excitation for fluorescence microscopy. The intense fluorescence and spherical shape of Flamma® PS bead series enable to generate stronger signals with small amount of probes in many tracing experiments. To decrease nonspecific binding and provide additional functional groups for conjugation, exteriors of PS bead are equipped with carboxylic acids. These pendent carboxylic acids allow them suitable for covalent coupling of proteins and other aminecontaining biomolecules by using water-soluble coupling reagents. BioActs provides various sizes of Flamma® PS bead series  $(0.1 - 0.5 \,\mu\text{m})$  in order to meet the diverse needs of our customer, and sizes specified in the product names are nominal bead diameters. These effective fluorescent PS beads can be utilized in a variety of applications such as neuronal tracers, cellular antigen markers, cell tracers, and the standardization reagents for flow cytometry. Due to their high fluorescence intensity, Flamma<sup>®</sup> PS bead series are suitable for the many nanoparticle based diagnostic tests, including tracking particles and cells, tracing fluid dynamics, discriminating the size dependence of uptake or transport in vivo. Although limited sizes of our fluorescent beads are available, we can prepare custom-sized particles, custom-dyed or custom-surface modifications orders upon request. Contact our custom service for further information.

Cat. No.	Product name	Ex*	Em*	Bead size	Functionality	Solids
PCS7001	Flamma <sup>®</sup> Deep Red PS Bead 0.2 µm	645	665	0.2 µm	СООН	2%
PCS7004	Flamma <sup>®</sup> Deep Red PS Bead 0.3 µm	645	665	0.3 µm	COOH	2%
PCS7002	Flamma <sup>®</sup> Deep Red PS Bead 0.5 μm	645	665	0.5 µm	СООН	2%

#### Table 1. Flamma<sup>®</sup> PS Bead series list

\*maxima of excitation and emission in nm

#### Handling & Storage

All Flamma<sup>®</sup> Fluors PS beads are stable under sonication, vortexing, or shaking process. All Flamma<sup>®</sup> Fluors PS beads should be stored at 2–6°C, protected from light, and **do not freeze.** Our Flamma<sup>®</sup> Fluors PS beads are supplied as suspensions (2% solids) in water containing 0.05% sodium azide. The PS beads are stable for at least one year, provided recommended storage conditions are strictly observed. Before sampling, mix well by sonication, vigorous shaking, or vortex mixing.

#### Activation of surface carboxylic acids to NHS ester

EDCI-mediated NHS ester formation is a common scheme to activate a carboxylic acid. NHS ester is readily reactive with primary amines of biomolecules. Primary amines such as lysine can react with NHS ester under the physiological pH in aqueous condition.

#### **Protocols for NHS ester activation**

- 1. Dilute 2% PS bead 10-fold by mixing in 50 mM MES buffer (pH 6.4)
- 2. Add N-hydroxysuccinimide and EDCI into the diluted bead solution, and mixed the mixture for 30 min at rt.
- 3. Centrifuge the mixture at 15,000 Xg for10 min, and then remove supernatant.
- 4. Suspend the bead pellet in 50 mM MES buffer (pH 6.4), and repeat the step 3 twice

### Protocols for antibody (Goat-anti-mouse IgG) conjugation

- 1. Suspend the NHS ester-activated bead pellet in 50 mM MES buffer (pH 6.4)
- 2. Add antibody solution to bead suspension solution, and stand the mixture for 4 h at rt (bead 1 mg per antibody 100  $\mu g)$
- 3. Centrifuge the mixture and remove supernatant, and wash bead pellet three times with 0.01% Tween-20 in PBS buffer

#### **Blocking/Quenching**

If you want to block carboxylic acids on the surface of PS beads with BSA or other amines such as glycine, add excess amount of 1x BSA or 1 mM amine solution in PBS after step 2 in conjugation protocol and perform step 3.

#### **Custom Labeling Service**

Based on accumulated know-how and technologies, BioActs provide a wide range of custom services such as protein fluorescence labeling, organic synthesis, oligonucleotide synthesis upon customers' request. Our reliable technology has acknowledged by our clients from domestic and overseas universities, institutions, in vitro diagnostic and pharmaceutical companies and has enabled to steadily conduct their requirements. In addition, we can introduce fluorescent materials to many other compounds such as organic and inorganic compounds, drugs, hormones, polymer, peptides, proteins, antibodies, etc. We also can provide chemical and optical analytical data, along with cell and animal experiments.



Nucleic acid



Peptide/Protein



Antibody



Small molecules /Polymer

#### **Technical Support**

#### ADDRESS

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WEBSITE http://www.bioacts.com/

#### MAILS

SDS (Safety Data Sheets) You can find SDS at www.bioacts.com, the official website of BioActs.

CoA (Certificate of Analysis) provides detailed quality information of each product. To see CoA, check the lot number written on each product's page at www.bioacts.com, when having trouble with check, contact to our technical support team

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