

# Magnosphere™ MX200/Carboxyl

# PRODUCT DESCRIPTION

Magnosphere™ MX200/CarboxyI beads are well-designed magnetic microparticles designed for immobilization of ligands through either physical or chemical means. The particle surfaces are covered with a JSR Life Sciences proprietary hydrophobic polymer that has charge density to maximize physical adsorption of proteins and chemical coupling through carboxy groups.

#### **FEATURES**

- Uniform particle size
- Superparamagnetic
- · Rapid magnetic responsiveness
- Maximized for physical adsorption of ligands
- · Maximized for chemical coupling through carboxyl surface groups

# **EXAMPLE APPLICATIONS**

Immunoassay

# **SPECIFICATIONS**

Package volume 5 mL

Solid content in slurry 2 % (2 x 10<sup>9</sup> beads/mL approx.)

Dispersion media 0.05 % Nonionic surfactant + 0.01 % ProClin 950 Al in H<sub>2</sub>O

Bead diameter 2.2 µm (micrometer)

Bead magnetite content 35 % approx.

Surface charge density 5 nmol/mg bead approx. Shelf life Labeled on the bottle

\*Surface charge density = amount of carboxyl groups per 1 mg beads

## **STORAGE**

**Magnosphere™ MX200/CarboxyI** is stable for 24 months when stored at 2-8 °C. Do not freeze the vial. Vortex the vial or pipette gently up and down to obtain a homogeneous dispersion before.

#### RECOMMENDED PROTOCOLS

#### [Protocol I] PHYSICAL COUPLING

## Reagent and equipment requirement

Binding Buffer: 50 mM MES buffer [2-(N-morpholino)ethanesulfonic acid] pH 6.2 (or

other appropriate buffer, if needed)

Washing & Storage Buffer: TBS or PBS buffer

Equipment: Magnetic separator. Vortex tube mixer. Tube rotator.

- Suspend the Magnosphere<sup>™</sup> MX200/CarboxyI beads well using Vortex mixer and put 1 mL of the suspension (i.e., 10 mg beads) into a microtube.
- Place the tube on a magnetic separator for 1 minute (or longer if needed) and remove the supernatant carefully.
- 3. Add 1 mL of Binding Buffer and suspend the beads by vortexing. Then, remove the supernatant as in step 2.
- Repeat step 3 for a total of 3 times.
- 5. Add 1 mL of Binding Buffer and suspend the beads by vortexing.
- Add 100 μg of antibody (100 μL, if antibody was diluted to 1 mg/mL) and suspend the beads by vortexing.

- 7. Keep rotating the tube with Tube rotator for 3 hours at room temperature.
- 8. Remove the supernatant as in step 2.
- 9. Wash the beads using 1 mL of Washing Buffer and suspend the beads by vortexing.
- 10. Remove the supernatant as in step 2.
- Repeat steps 9 & 10 for a total of 3 times.
- Suspend the beads with desired buffer suitable for downstream applications and store at 2-8 °C until needed.

# [Protocol II] CHEMICAL COUPLING

## Reagent and equipment requirement

Binding Buffer: 0.1 M MES\* buffer pH 5.0

(\*MES: 2-(N-morpholino)ethanesulfonic acid)

Washing Buffer: TBS-T (25 mM Tris-HCl, pH 7.2, 0.15 M NaCl, 0.05 % Tween20)

Coupling Reagent 10 mg/mL EDC\*\* in ice-cooled Binding Buffer, prepared just before

the coupling reaction

(\*\*EDC: 1-Ethyl-3-[3-dimethylaminopropyl]carbodiimide

Hydrochloride)

Equipment: Magnetic separator. Vortex tube mixer. Tube rotator.

- Suspend the Magnosphere<sup>™</sup> MX200/CarboxyI beads well using Vortex mixer and put 500 μL of the suspension (i.e., 10 mg beads) into a microtube.
- Place the tube on a magnetic separator for 1 minute (or longer if needed) and remove the supernatant carefully.
- Add 1 mL of Binding Buffer and suspend the beads by vortexing. Then, remove the supernatant as in step 2.
- 4. Add 1 mL of Binding Buffer and suspend the beads by vortexing.
- Add 100 μg of antibody (100 μL, if antibody was diluted to 1 mg/mL) and suspend the beads by vortexing.
- 6. Keep rotating the tube with Tube rotator for 30 minutes at room temperature.
- 7. Add 100 µL of Coupling Reagent and suspend the beads by vortexing.
- 8. Keep rotating the tube with Tube rotator for 3 hours at room temperature.
- 9. Remove the supernatant as in step 2.
- 10. Wash the beads using 1 mL of Washing Buffer and suspend the beads by vortexing.
- 11. Remove the supernatant as in step 2.
- Repeat steps 10 & 11 for a total of 3 times.
- Suspend the beads with a desired buffer suitable for downstream applications and store at 2-8 °C until needed.

## **IMPORTANT NOTICE**

- This product is for research use only and not intended for therapeutic or in vivo diagnostic use.
- The specifications of the product may be changed without a notice.
- JSR Life Sciences Corporation does not guarantee that this product will be continuously available.
- JSR Life Sciences Corporation makes no warranties as to this product including, but not limited to, implied warranties of merchantability or fitness for a particular purpose.

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