# Instructions for Labeling Proteins with Gadolinium

# Reagents. Store all reagents at 4°C.

Reagent 1	Proprietary chelate (7.3x10 <sup>-6</sup> mol) ~5 mg in a 2 ml plastic screw cap vial sealed
	with parafilm
Reagent 2	Reconstitution solution (2 ml) 2.04 ml 1 M sodium acetate and 0.112 ml 1 M
	sodium hydroxido in a 2 ml plastic cal vial soaled with parafilm

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Reagent 3 1 M GdCl<sub>2</sub> (0.2 ml) 2 ml glass vial with gray stoper and silver crimp

Reagent 4 0.2 M carbonate buffer pH 8.9 (10 ml) 10 ml vial with red stopper and silver crimp

### **Generic conjugation procedure**

## 1.1 Protein preparation.

Transfer protein to a suitable solution free of nucleophiles, such as amine and sulfhydryl groups. We recommend that the protein conjugation be done in 0.2 M carbonate buffer pH 8.5 to 8.9. If possible, start by dissolving the solid protein in the reaction buffer.

# 1.2 Reconstitution of proprietary chelate and chelation of gadolinium.

To the container of Reagent 1, add in sequence 393  $\mu$ l of Reagent 2 and 6.66  $\mu$ l (6.66x10<sup>-6</sup> mol) of Reagent 3. Cap container and vortex approximately 2 minutes until solid dissolves. Let the resulting solution sit for 5 minutes.

#### 1.3 Protein conjugation.

At room temperature and with gentle mixing of your protein, add approximately one, two, three or more mols of chelate per mol of protein. Continue mixing for 2 hours.

#### 1.4 Purification.

Use a suitable size separation procedure for purification of your protein, such as gel filtration or dialysis against a suitable buffer for your protein.

## **Example conjugation with albumin**

#### 2.1 Protein preparation.

Dissolve 50 mg of albumin (7.3x10<sup>-7</sup> mol) in 10 ml of Reagent 4.

## 2.2 Reconstitution of proprietary chelate and chelation of gadolinium.

To the container of Reagent 1, add in sequence 393  $\mu$ l of Reagent 2 and 6.66  $\mu$ l (6.66x10<sup>-6</sup> mol) of Reagent 3. Cap container and vortex approximately 2 minutes until solid dissolves. Let the resulting solution sit for 5 minutes.

## 2.3 Protein conjugation.

At room temperature and with gentle mixing of the albumin, add all of the reconstituted chelate (7.3 x 10<sup>-6</sup> mol) to the albumin. Continue mixing for 2 hours.

## 2.4 Purification.

Dialyze the conjugated albumin against a 10K MWCO membrane. Store at 4°C.

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