



## Product Information & Manual

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### HyLink™ Biotin Labeling Kit

Cat no. LDG0004RC/LDG0005RC/LDG0006RC

#### Product Overview

##### Package component

Item	LDG0004RC (10 µg x 3)	LDG0005RC (100 µg x 3)	LDG0006RC (1 mg x 1)
Biotin	3 vials	3 vials	1 vial
10X Modifier	1 vial	1 vial	1 vial
10X Quencher	1 vial	1 vial	1 vial

#### Description

Biotin is a widely used and powerful tool for research due to the specific and high affinity with streptavidin/avidin. Antibody or protein conjugated with several biotin molecules can also amplify the detection signal through streptavidin-conjugated molecule such as streptavidin-HRP, streptavidin-FITC, etc. Biotinylated antibody or protein can be used in various applications including ELISA, WB, IHC, IFA and FACS. Leadgene HyLink™ Biotin Labeling Kit is designed for biotinylation of a small quantity (10 µg-1 mg) of antibody or protein. It provides a rapid and easy process with high efficiency to conjugate biotin to antibody or protein.

#### Storage and Stability

- (1) Stored at -20°C. Avoid repeated freeze/thaw cycles.
- (2) Equilibrate kit to room temperature before use.
- (3) Is stable for one year under proper storage conditions.

#### Procedure

##### Biotin conjugation protocol

- (1) Dissolve antibody in PBS or other buffers that do not contain amine, Tris or glycerol. Use **10X Modifier** (e.g. Add 1 µL of 10X Modifier for 9 µL of antibody) or dialysis against PBS if pH value of used buffer is out of 7 to 8.
- (2) Spin down and equilibrate the vial of **Biotin** at room temperature before opening the cap.
- (3) Make sure all buffers are well dissolved before using. If not, please vortex the vial to make salts dissolved.
- (4) Remove the cap of the vial of **Biotin** and pipette antibody into the vial. Mix gently by pipetting several times.
- (5) Cover the cap and incubate in the dark at room temperature for 3 hours.
- (6) After incubating, add **10X Quencher** (e.g. 1 µL of 10X Quencher for 9 µL of antibody-biotin mixture) and mix gently by pipetting. The conjugates can be used after 30 minutes.

\* For protein conjugation, the amount of protein can be calculated by formula below:

$$\text{Quantities of protein} = \text{quantities of kit (e.g. 10 µg)} \times \frac{\text{M.W. of target protein}}{(150 \text{ (M.W. of IgG)})}$$

#### Important notes

- (1) Antibody concentrations of 1-4 mg/mL generally give optimal results. Recommended amount and volume of antibody for optimal results.

Kit size	Antibody amount	Reaction volume
3 x 10 µg	10-20 µg	4-20 µL
3 x 100 µg	100-200 µg	40-200 µL
1 x 1 mg	1-2 mg	400-2000 µL

- (2) Common non-buffering salts (e.g. sodium chloride) have no effect on conjugation efficiency. Avoid buffer component that contains primary amine (e.g. amino acid or ethanolamine) and thiols (e.g. 2-Mercaptoethanol or DTT).

Components that have on effect or little effect on labeling reaction:

- up to 50 mM Tris
- up to 50 mM HEPES
- up to 10% glycerol
- up to 0.02% sodium azide

## Disclaimer

This product is for research use only and is not intended for diagnostic use.

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