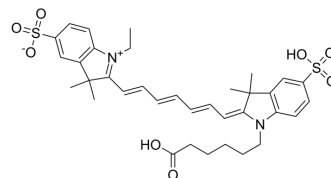


## CY7

<b>Cat. No.:</b>	HY-D0825
<b>CAS No.:</b>	943298-08-6
<b>Molecular Formula:</b>	C <sub>35</sub> H <sub>42</sub> N <sub>2</sub> O <sub>8</sub> S <sub>2</sub>
<b>Molecular Weight:</b>	682.85
<b>Target:</b>	Fluorescent Dye
<b>Pathway:</b>	Others
<b>Storage:</b>	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : ≥ 33 mg/mL (48.33 mM)  
\* "≥" means soluble, but saturation unknown.

Solvent	Mass	Concentration		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.4645 mL	7.3223 mL	14.6445 mL
	5 mM	0.2929 mL	1.4645 mL	2.9289 mL
	10 mM	0.1464 mL	0.7322 mL	1.4645 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

CY7 is a CY dye. CY, short for Cyanine, is a compound consisting of two nitrogen atoms connected by an odd number of methyl units. Cyanine compounds have the characteristics of long wavelength, adjustable absorption and emission, high extinction coefficient, good water solubility and relatively simple synthesis<sup>[1]</sup>. CY dyes are often used for the labeling of proteins, antibodies and small molecular compounds. For the labeling of protein antibodies, the combination can be completed through a simple mixing reaction. Below, we introduce the labeling method of protein antibody labeling, which has certain reference significance<sup>[2]</sup>.

#### In Vitro

##### Protocol

##### 1. Protein Preparation

- In order to obtain the best labeling effect, please prepare the protein (antibody) concentration as 2 mg/mL.
- The pH value of protein solution shall be 8.5±0.5. If the pH is lower than 8.0, 1 M sodium bicarbonate shall be used for adjustment.
- If the protein concentration is lower than 2 mg/mL, the labeling efficiency will be greatly reduced. In order to obtain the best labeling efficiency, it is recommended that the final protein concentration range is 2-10 mg/mL.
- The protein must be in the buffer without primary amine (such as Tris or glycine) and ammonium ion, otherwise the labeling efficiency will be affected.

## 2. Dye Preparation (Example for CY3-NHS ester)

Add anhydrous DMSO into the vial of CY3-NHS ester to make a 10 mM stock solution. Mix well by pipetting or vortex. Before use, it must be activated with condensation solution (500 µg/mL) (HY-D0178) before subsequent labeling experiments can be performed.

## 3. Calculation of dye dosage

The amount of CY3-NHS ester required for reaction depends on the amount of protein to be labeled, and the optimal molar ratio of CY3-NHS ester to protein is about 10.

Example: assuming the required marker protein is 500 µL 2 mg/mL IgG (MW=150,000), use 100 µL DMSO dissolve 1 mg CY3-NHS ester, the required CY3-NHS ester volume is 5.05 µL, and the detailed calculation process is as follows:

$$1) \text{ mmol (IgG)} = \text{mg/mL (IgG)} \times \text{mL (IgG)} / \text{MW (IgG)} = 2 \text{ mg/mL} \times 0.5 \text{ mL} / 150,000 \text{ mg/mmol} = 6.7 \times 10^{-6} \text{ mmol}$$

$$2) \text{ mmol (CY3-NHS ester)} = \text{mmol (IgG)} \times 10 = 6.7 \times 10^{-6} \text{ mmol} \times 10 = 6.7 \times 10^{-5} \text{ mmol}$$

$$3) \mu\text{L (CY3-NHS ester)} = \text{mmol (CY3-NHS ester)} \times \text{MW (CY3-NHS ester)} / \text{mg}/\mu\text{L (CY3-NHS ester)} = 6.7 \times 10^{-5} \text{ mmol} \times 753.88 \text{ mg/mmol} / 0.01 \text{ mg}/\mu\text{L} = 5.05 \mu\text{L (CY3-NHS ester)}$$

## 4. Run conjugation reaction

1) A good volume of freshly prepared 10 mg/mL CY3-NHS ester is slowly added to 0.5 mL protein sample

In solution, gently shake to mix, then centrifuge briefly to collect the sample at the bottom of the reaction tube. Don't mix well to prevent protein samples from denaturation and inactivation.

2) The reaction tubes were placed in a dark place and incubated gently at room temperature for 60 minutes at intervals. For 10-15 minutes, gently reverse the reaction tubes several times to fully mix the two reactants and raise the bar efficiency.

## 5. Purify the conjugation

The following protocol is an example of dye-protein conjugate purification by using a SepHadex G-25 column.

1) Prepare SepHadex G-25 column according to the manufacture instruction.

2) Load the reaction mixture (From "Run conjugation reaction") to the top of the SepHadex G-25 column.

3) Add PBS (pH 7.2-7.4) as soon as the sample runs just below the top resin surface.

4) Add more PBS (pH 7.2-7.4) to the desired sample to complete the column purification. Combine the fractions that contain the desired dye-protein conjugate.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## CUSTOMER VALIDATION

- Chem Eng J. 2021, 129284.
- J Control Release. 2021 Apr 10;332:194-209.
- J Cachexia Sarcopenia Muscle. 2023 Jan 27.
- Cell Mol Biol Lett. 2022 Dec 6;27(1):107.
- Cell Mol Biol Lett. 2022 Oct 12;27(1):92.

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## REFERENCES

- [1]. Ptaszek M. Rational design of fluorophores for in vivo applications. Prog Mol Biol Transl Sci. 2013;113:59-108.
- [2]. Shindy, H. A. (2017). Fundamentals in the chemistry of cyanine dyes: A review. Dyes and Pigments, 145, 505–513. doi:10.1016/j.dyepig.2017.06.029
- [3]. Lim B, et al. A Unique Recombinant Fluoroprobe Targeting Activated Platelets Allows In Vivo Detection of Arterial Thrombosis and Pulmonary Embolism Using a Novel Three-Dimensional Fluorescence Emission Computed Tomography (FLECT) Technology. Theranostics. 2017 Feb 26;7(5):1047-1061.

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**Caution: Product has not been fully validated for medical applications. For research use only.**

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