

PRODUCT INFORMATION SHEET

Version: 1.0 Revision Date: 08/08/2020

LINK NHS

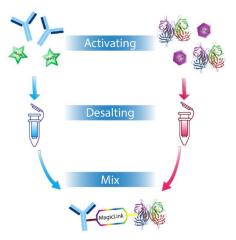
Component	Product size	Storage
	BP-50064	
LINK NHS	2 μmol	-20 °C

Overview

LINK NHS ester contains NHS ester and LINK functional group with a hydrophilic spacer. This compound can react with amine containing biomolecules such as antibody, protein, enzyme, amine-modified oligonucleotides, etc. at neutral or slightly basic pH to form LINK-activated biomolecules. The spacer helps with hydrophilic and bonding properties of the new biomolecules in bioassay.

Scientist can activate their biomolecules with LINK NHS to get LINK-activated biomolecules which will instantly react with a MAGIC-activated biomolecule to form a new biomolecule by MagicLink[™] chemistry. The new linked biomolecule is ready to use.

LINK NHS is one component of BroadPharm proprietary MagicLink[™] crosslinking chemistry. LINK NHS and MAGIC NHS (catalog # BP-50063) are always used together for linking the two biomolecules.



Scheme 1. a typical flow chart of LINK NHS activating protein to form LINK-activated protein (red part)

Technical Considerations

Pre-conjugation considerations for the protein.

The protein should be purified and amine, glycine, BSA, gelatin free. Glycine can be removed by dialyzing against 1X PBS, pH 7.2-7.4. Alternatively, use Amicon Ultra-0.5, Ultracel-10 Membrane, 10K MWCO (Cat # UFC501008 from Millipore). Impure protein or protein stabilized with bovine serum albumin (BSA) or gelatin will not be labeled well.



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- For optimal labeling efficiency a final protein concentration range of 1-5 mg/ml is recommended. The conjugation efficiency is significantly reduced if protein concentration is less than 1 mg/ml.
- Activated protein should be used right away for protein coupling via MagicLink[™] chemistry.

Sample Experimental Protocol

- 1. Reconstitute LINK NHS vial with 100 µl DMSO. Use immediately after reconstitution.
- 2. Activation

Suggested activation condition: 20X LINK NHS to protein ratio. Protein concentration between 1-5 mg/ml in 1X PBS pH 7.2 – 7.5.

Sample calculation of protein amount to use for reaction with every 10 μl of LINK NHS in DMSO at 20X LINK to protein ratio. Protein volume (ml) = protein molecular wt. / protein conc. / 100,000 Unit of molecular weight is g/mol, unit of concentration is mg/ml.

Example calculation for a solution of antibody 150 kD (or 150,000 g/mol) at 5 mg/ml concentration. ml of antibody to use with 10 μl of LINK NHS (at 20 fold ratio) = 150000 / 5 / 100000 = 0.3 ml or 300 μl If a total of 100 μl LINK NHS is to be used up (at 20 fold ratio), then it would require 3 ml of antibody.

Protein activation Protein should be 1-5 mg/ml in 1X PBS (buffer exchange, see note in technical consideration). Add LINK NHS volume according to the calculation above. Mix gently and incubate for 1 hour at room temperature.

Desalt of the Activated Protein

Example of desalting step by Amicon Ultra – 0.5 ml concentrator (not supplied), using a microcentrifuge. Researcher to select appropriate molecular weight cutoff of concentrator, and or different size concentrator; follow manufacturer's direction.

- 1. Hydrate concentrator membrane 'filter device' with 400 to 500 μl of 1X PBS pH 7.2 7.5, or DI water, and microcentrifuge 14,000 x g, for 3 minutes. Discard, liquid from the filter device and collection tube.
- Spin down by adding activated protein to the concentrator/filter device up to 500 μl. Microcentrifuge at 14,000 x g, 8 minutes, or to minimum volume ~ 50 μl left in the filter device. Discard waste from the collection tube.
- 3. Desalt by adding 1X PBS to the filter device up to 500 μ l. Microcentrifuge at 14,000 x g, 8 minutes, or to minimum volume ~ 50 μ l left in the filter device. Discard waste from the collection tube.
- 4. Repeat step 3, twice.
- 5. Collect activated protein from the filter device into a microcentrifuge tube.
- 6. Optional for maximum recovery, add 1X PBS, volume determined by the user, to the filter devices to rinse out residual protein. Microcentrifuge pulse spin, collect proteins/PBS from filter device, and add to the microcentrifuge tube from step 5, mix.

Determine the ratio of the number of LINK group: antibody

Here is a reference for the number of LINK groups after LINK NHS modification reaction:



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antibody conc. (mg/ml)	Mixing ratio (MR)	N _{magic} /N IgG
2	20:1	5~6
10	3:1	1.0~1.5
10	5:1	2.0~2.5
10	10:1	3.0~4.0

The number of LINK groups can be determined with 650-LINK(Cat# BP-50066).

Storage

Activated protein should be used right away for protein coupling via MagicLink[™] chemistry. It is not recommended for longtime storage.