Product Datasheet

Lightning-Link (R) Rapid Atto565 Antibody Labeling Kit 351-0010

Unit Size: 3 x 100ug Reaction Store at -20C.

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351-0010

Lightning-Link (R) Rapid Atto565 Antibody Labeling Kit

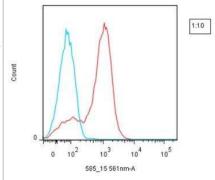
Lighthing-Link (R) Rapid Attoobs	Antibody Labeling Kit	
Product Information		
Unit Size	3 x 100ug Reaction	
Concentration	Concentration is not relevant for this product. Please see the protocols for proper use of this product.	
Storage	Store at -20C.	
Conjugate	Atto565	
Product Description		
Description	Lightning-Link Rapid is an innovative technology that enables direct labeling of proteins, peptides or other biomolecules for use in R&D applications, drug discovery and the development of diagnostic kits (See protocol for further information). The easy-to-use, one step procedure allows researchers to covalently label biomolecules with only 30 seconds hands-on time; furthermore conjugates are ready to use in less than twenty minutes. The researcher simply pipettes the biomolecule into a vial of lyophilized mixture containing the label of interest and incubates (for more details please watch the video below). FeaturesBenefitsQuick and easy to useSave time, no special knowledge requiredNo separation steps100% recovery - no antibody/protein lossCan be used in a wide range of applicationsFlexibleFreeze driedShips at ambient temperature, long shelf-lifeFully scalable (10 ug to 1 g or more)Easy transfer from R&D to manufacturingStringently QC testedConsistent high quality, excellent batch-to-batch reproducibilityLarge number of labels available Experimental flexibilityReliable: nearly 300 referencesSuccessfully used in many fields of research Atto565 is one of a new generation of fluorescent labels. It has a strong absorption at 570nm, high fluorescence at 598nm (extinction coefficient 1.2 x105 cm-1M-1) and quantum yield. Learn more about Lightning-Link™ Conjugation Kits by reading FAQs For more information please check out these useful links! Antibody Labeling Guide Antibody Conjugation Illustrated Assay	
Kit Components	1 or 3 glass vial(s) of Lightning-Link Rapid mix, 1 vial of LL-Rapid Modifier reagent, 1 vial of LL-Rapid Quencher reagent	
Notes	This product is manufactured by Abcam and distributed by Novus Biologicals. This product is for research use only and is not approved for use in humans or in clinical diagnosis. This product is guaranteed for 1 year from date of receipt and this statement overrides any mentioned guarantee period on the limitations section of this products datasheet. Please contact technical@novusbio.com with questions.	
Product Application Details		
Applications	Flow Cytometry	



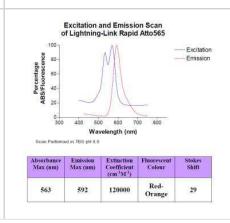
Recommended Dilutions	Flow Cytometry
Application Notes	By circumventing the desalting or dialysis steps that commonly interrupt traditional antibody conjugation procedures, LightningLink technology can be used to label both small (e.g. 10 ug) and large quantities of primary antibodies with ease. Batch-to-batch variation upon scale up is minimal as the process is so simple, and recoveries are always 100%.

Images

Flow Cytometry: Lightning-Link Rapid Atto565 Antibody Labeling Kit [351 -0010] - Mouse anti-human CD3 was conjugated with Atto565 using Lightning-Link® Rapid kit. The conjugated antibody was then used to stain human peripheral blood lymphocytes, followed by analysis with flow cytometry. (Blue line - negative control; red line - positive staining).



Lightning-Link Rapid Atto565 Antibody Labeling Kit [351-0010]



Publications

Robison P, Caporizzo MA, Ahmadzadeh H et al. Detyrosinated microtubules buckle and bear load in contracting cardiomyocytes Science 2016-04-22 [PMID: 27102488] (ICC/IF)

Diesenberg K, Beerbaum M, Fink U et al. SEPT9 negatively regulates ubiquitin-dependent downregulation of EGFR. J Cell Sci 2015-01-15 [PMID: 25472714]

Babusyte A, Kotthoff M, Fiedler J, Krautwurst D. Biogenic amines activate blood leukocytes via trace amine-associated receptors TAAR1 and TAAR2. J Leukoc Biol 2014-01-01 [PMID: 23315425] (IB)

Oki S, Kitajima K, Meno C. Dissecting the role of Fgf signaling during gastrulation and left-right axis formation in mouse embryos using chemical inhibitors. Dev Dyn 2010-01-01 [PMID: 20503372]





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