

Datasheet: LNK033APC BATCH NUMBER 160415

rad-antibodies.com/protocols. Yes No Not Determined Sugge Conjugation • We recommend that for each conjugation the user determines the best an ratio. Product Information LYNX Rapid APC Antibody Conjugation Kit® enables the rapid conjugation conjugation containing Allophycocyanin (APC) label to protein. Activation of proprietary reagents within the antibody-label solution directional covalent bonding of APC to the antibody. The LYNX Rapid Conjugation kit® can be used to label small quantities of antibody/protein at near neutral pH, allowing a high conjugation efficiency antibody recovery. Reagents In The Kit 1 Vial of 1mg LYNX lyophilized APC mix 1 Vial LYNX Quencher reagent 1 Vial LYNX Quencher reagent Preparing The Antibody The following buffer solutions are recommended for preparing the antibod 10-50mM amine-free buffer (e.g HEPES, MES, MOPS and phosphate) photon							
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If possible, avoid buffers containing nucleophilic components such a		10-50mM amine-free buffer (e.g HEPES, MES, MOPS and phosphate) pH range 6.5-8.5, although moderate concentrations of Tris buffer (<20mM) may be tolerated.					
amines and thiols (e.g. thiomersal/thimerosal) since they may react v		•		-			

chemicals. Azide (0.02-0.1%), EDTA, up to 50% Glycerol and common non-buffering salts and sugars have little or no effect on conjugation efficiency. For optimal results the antibody should be at a concentration of 1mg/ml, with a maximum volume of 1ml and a maximum antibody amount of 1mg. Antibody at a concentration of greater than 1mg/ml requires dilution. Antibody below 1mg/ml can still be used as long as the maximum volume is not exceeded. Using less than the recommended amount of antibody may result in unbound label, but this will be removed during subsequent application wash steps. Antibody below 0.5mg/ml should be concentrated before use with the kit. Instructions For Use 1. To the antibody sample add 1ul of the Modifier reagent for every 10ul of antibody and mix gently. 2. Pipette the mixed antibody-modifier sample directly onto the LYNX lyophilized mix and gently pipette up and down twice to resuspend. 3. Replace cap onto vial and incubate in the dark at room temperature (20-25°C) for 3 hours, or overnight if preferred. 4. After incubation, add 1ul of Quencher reagent for every 10ul of antibody used. Leave to stand for 30 minutes before use. References 1. Wang, Y. et al. (2010) Local host response to chlamydial urethral infection in male guinea pigs. Infect Immun.78: 1670-81. 2. Lacy, H.M. et al. (2011) Essential Role for Neutrophils in Pathogenesis and Adaptive Immunity in Chlamydia caviae Ocular Infections. Infect Immun. 79: 1889-97 3. Paget, C. et al. (2012) Interleukin-22 is produced by invariant natural killer T lymphocytes during influenza A virus infection: potential role in protection against lung epithelial damage. J Biol Chem. 287: 8816-29. 4. Seliger, C. et al. (2011) A rapid high-precision flow cytometry based technique for total white blood cell counting in chickens. Vet Immunol Immunopathol. 145: 86-99. 5. Fu, Y. et al. (2014) Development of a FACS-based assay for evaluating antiviral potency of compound in dengue infected peripheral blood mononuclear cells. J Virol Methods. 196: 18-24. 6. TraxImayr, M.W. et al. (2014) Construction of pH-sensitive Her2-binding IgG1-Fc by directed evolution. Biotechnol J. 9: 1013-22. 7. Wielgosz, M.M. et al. (2015) Generation of a lentiviral vector producer cell clone for human Wiskott-Aldrich syndrome gene therapy. Mol Ther Methods Clin Dev. 2: 14063. 8. Hofer, C.C. et al. (2015) Infection of mice with influenza A/WSN/33 (H1N1) virus alters alveolar type II cell phenotype. Am J Physiol Lung Cell Mol Physiol. 308 (7): L628-38. 9. Poh, C.M. et al. (2014) Damage to the blood-brain barrier during experimental cerebral malaria results from synergistic effects of CD8+ T cells with different specificities. Infect Immun. 82: 4854-64. 10. Hasenhindl, C. et al. (2014) Creating stable stem regions for loop elongation in Fcabs - insights from combining yeast surface display, in silico loop reconstruction and molecular

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Information	https://www.bio-rad-antibodies.com/SDS/LNK033APC					
Health And Saf						
Guarantee	12 months from date of despatch					
	reach room temperature to minimize condensation.					
	undiluted. Avoid repeated freezing and thawing. Before opening, allow the components t					
	Storage in frost-free freezers is not recommended. This product should be stored					
	is shipped under ambient conditions with silica packets to avoid exposure to moisture. O receipt, Bio-Rad recommend that the kit is stored at -20°C and protected from moisture.					
Storage	This kit contains lyophilized hygroscopic components that are moisture-sensitive. This ki					
	and humoral immunity in pigs. <u>Virol J. 20 (1): 181.</u>					
	Cytometry-Based Differential Blood Count. <u>Microbiol Spectr. 11 (4): e0435122.</u> 16. Haach, V. <i>et al.</i> (2023) A polyvalent virosomal influenza vaccine induces broad cellul					
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	<u>10406387221077969.</u>					
	the equine systemic inflammatory response syndrome. <u>J Vet Diagn Invest.</u>					
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	improves schwann cell adhesion and proliferation. <u>J Mater Sci Mater Med. 27 (12): 188.</u>					
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	dynamics simulations. <u>Biochim Biophys Acta.1844: 1530-40.</u>					

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Printed on 18 Jan 2024

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