Datasheet: BUF034A BATCH NUMBER 168919

Description:	ELISA SYNBLOCK
Name:	ELISA SYNBLOCK
Format:	Ready To Use
Product Type:	Accessory Reagent
Quantity:	100 ml

Product Details

Applications	This product has been reported to work in the following applications. This information is derived from testing within our laboratories, peer-reviewed publications or personal communications from the originators. Please refer to references indicated for further information. For general protocol recommendations, please visit <u>www.bio-rad-antibodies.com/protocols</u> .							
	Yes No Not Determined Suggested Dilution							
	ELISA Where this product has n	• ot been to	sted for u	se in a particular tech	Neat			
	a guide only. It is recommoder system using appropriate	se in such nended tha	procedure at the user	es. Suggested working titrates the product for	g dilutions are given as			
Product Form	Ready to use - liquid							
Buffer Solution	Phosphate buffered saline	e						
Preservative Stabilisers	<0.1% sodium azide (Nal	N ₃)						
Product Information	ELISA Synblock is an El with buffers containing ar signals in ELISA assays v	nimal prote	eins (e.g. E	3SA) and reduce non-	specific background			
Intended Use	ELISA SynBlock is a novel protein-free blocking buffer suitable for use in all ELISA formats requiring maximum blocking strength. With Tween and synthetic blocking agents, the inert nature of this unique buffer, enables maximum reduction of non-specific binding and interference associated particularly with sandwich ELISA assays.							
	Additional molecular stab environment for coating a temperature and stored o	antigen or	capture ar	ntibody. Plates can be	-			

	N.B. SYNBLOCK is not suitable for use on Immunolon-2 plates. Bio-Rad recommends the use of <u>BUF033A</u> for this purpose.
Instructions For Use	1. Coat ELISA plate with antibody or antigen as required.
	2. After incubation, remove the coating solution and wash the plate x2 with wash buffer. <u>BUF031A</u> can be used for this purpose.
	3. Add 300-400ul of BUF034A and incubate for 2-24 hours. Use a volume equal to or greater than the volume of coating solution.
	4. After removal of the blocking buffer continue with the assay or dry the plate for long-term storage at +4°C.
References	 Afrough, B. <i>et al.</i> (2007) Identification and elimination of false-positives in an ELISA-based system for qualitative assessment of glycoconjugate binding using a selection of plant lectins. <u>Biotechniques. 43</u> (4): 458, 460, 462 passim. Dalley, D. <i>et al.</i> (2008) Development and evaluation of a gamma-interferon assay for tuberculosis in badgers (<i>Meles meles</i>). <u>Tuberculosis (Edinb). 88: 235-43.</u> Ahmed, R.R. <i>et al.</i> (2010) BACE1 and BACE2 enzymatic activities in Alzheimer's disease. <u>J Neurochem. 112: 1045-53.</u> Chambers, M.A. <i>et al.</i> (2009) Performance of TB immunodiagnostic tests in Eurasian badgers (<i>Meles meles</i>) of different ages and the influence of duration of infection on serological sensitivity. <u>BMC Vet Res. 5: 42.</u> Thompson, R. <i>et al.</i> (2011) Optimization of the enzyme-linked lectin assay for enhanced glycoprotein and glycoconjugate analysis. <u>Anal Biochem, 413: 114-22.</u> Kuramitz, H. <i>et al.</i> (2012) Multiplexed assay for proteins based on sequestration electrochemistry using the protein binding electroactive magnetic microbeads. <u>Anal Sci. 28 (1): 77.</u> Dwek, M.V. <i>et al.</i> (2010) A sensitive assay to measure biomarker glycosylation demonstrates increased fucosylation of prostate specific antigen (PSA) in patients with prostate cancer compared with benign prostatic hyperplasia. <u>Clin Chim Acta. 411 (23-24): 1935-9.</u> Verhelst, R. <i>et al.</i> (2010) The effects of plant polyphenols on enterotoxigenic <i>Escherichia coli</i> adhesion and toxin binding <u>Livestock Science. 133 (1-3): 101-3</u> Verhelst, R. <i>et al.</i> (2013) <i>E. coli</i> heat labile toxin (LT) inactivation by specific polyphenols is aggregation dependent. <u>Vet Microbiol. 163 (3-4): 319-24.</u> Greenwell P <i>et al.</i> (2008) Purification and analysis of DNases of <i>Tritrichomonas foetus:</i> evidence that these enzymes are glycoproteins. Int J Parasitol. 38 (7): 749-56. Beckett, T.L. <i>et al.</i> (2013) A ketogenic diet improv

	15. Wilcock, D.M. <i>et al.</i> (2015) Down syndrome individuals with Alzheimer's disease have	
	a distinct neuroinflammatory phenotype compared to sporadic Alzheimer's disease. Neurobiol Aging. 36 (9): 2468-74.	
	16. Chinthamani S <i>et al.</i> (2017) Macrophage inducible C-type lectin (Mincle) recognizes	
	glycosylated surface (S)-layer of the periodontal pathogen <i>Tannerella forsythia</i> . <u>PLoS</u> One. 12 (3): e0173394.	
	17. LeVine, H. 3rd. <i>et al.</i> (2017) Down syndrome: age-dependence of PiB binding in	
	postmortem frontal cortex across the lifespan. Neurobiol Aging. 54: 163-9.	
Storage	Store at +4°C.	
	DO NOT FREEZE	
Guarantee	Guaranteed until date of expiry. Please see product label.	
Health And Safety	Material Safety Datasheet documentation #10380 available at:	
Information	https://www.bio-rad-antibodies.com/SDS/BUF034A 10380	
Regulatory	For research purposes only	
Related Produ	cts	
Recommended U	seful Reagents	

5x ELISA COATING BUFFER (BUF030A) 10x ELISA WASH BUFFER (BUF031A)

North & South America	Tel: +1 800 265 7376 Fax: +1 919 878 3751 Email: antibody_sales_us@bio-ra	Worldwide d.com	Tel: +44 (0)1865 852 700 Fax: +44 (0)1865 852 739 Email: antibody_sales_uk@bio-rac	Europe	Tel: +49 (0) 89 8090 95 21 Fax: +49 (0) 89 8090 95 50 Email: antibody_sales_de@bio-rad.com	
To find a batch/lot specific datasheet for this product, please use our online search tool at: bio-rad-antibodies.com/datasheets						

'M417390:230322'

Printed on 10 Sep 2024

© 2024 Bio-Rad Laboratories Inc | Legal | Imprint