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Certificate of Analysis

Becton Dickinson and Company BD Diagnostic Systems PO Box 999 Sparks MD 21152-0999 US

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Product Name: BOTTLECatalog Number: 211825Batch Number: 1199470Expiration Date: 2016/04		MED 500G cture Date : 2	011/06/30
 Dehydrated Medium Appendous Solubility: 3% solution Solution Appearance: 1 Medium was tested per (USP) Growth Promotion 100 CFUs. Tubes were to 5 days for (*) orgatindicated. 	on, soluble in Light amber, cl European (EP) n requirements. re incubated ae	distilled or ear and United S Tubes were robically fo	deionized water States Pharmacopeia inoculated with or 3 days and up
	6633 growth 10231 growth 8739 growth	20-25°C 30-35°C,20- 20-25°C 30-35°C 30-35°C 30-35°C	Up to 5 days 25°C Up to 3 days Up to 5 days Up to 3 days Up to 3 days Up to 3 days Up to 3 days
05. Cultural Response: Med Tubes were inoculated temperatures specified necessary.	with the test	organisms an	d incubated at the
TEST ORGANISMS Neisseria meningitidis Staphylococcus epider Streptococcus pneumon Streptococcus pyogenes	s 13090 nidis 12228 Lae 6305	EMPERATURE 30-35°C f 30-35°C 30-35°C 30-35°C 30-35°C	RECOVERY air to good good good good
06. Residual Solvents (CPI Soy Broth indicates th No other solvents were	nat there is le	ss than 5000	ppm of Acetone.
Characteristic Unit	Value	LowLimit	HighLimit
Loss on Drying : % pH at 25°C : Bulk Lot Number : -	2 7.4 1172568	0 7.1	5 7.5



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	Country of	Tissue (Tissue Category		
Animal Source	Origin	BIC	SIC	ABC	
Bovine	Australia	IV	IV	MLK	
Porcine	USA	III	III	IB	
Bovine	New Zealand	IV	IV	MLK	
Porcine	Canada	III	III	IB	

The Batch Number on this certificate is synonymous with the Lot Number shown on the product label.

BD Diagnostic Systems (BDDS) is an ISO 13485:2003 and ISO 9001:2008 Registered facility. BDDS products are manufactured in facilities registered with the United States Food and Drug Administration (FDA), and are regulated by the FDA's Quality System Regulations (QSRs). This product met BDDS stringent quality standards at time of batch/lot release. Any test results reported on this certificate were obtained at time of release.

BD Diagnostic Systems' Certificates of Analysis (COA) typically are set up to contain animal origin information for finished products manufactured using materials of animal origin. The animal origin information may be contained in the animal source table and/or in one or more of the additional paragraphs found on the COA. This information is a compilation of animal origin data from the individual lots of raw materials used to manufacture the batch of BD Diagnostic Systems (BDDS) finished product for which the COA was created.

At the time the BDDS Certificate of Analysis is created and sent to the Internet website address at http://www.bd.com/regdocs/, the animal origin information as provided to BDDS by its suppliers is pulled into the certificate as it is created by the BDDS automated certificate system.

At times, suppliers notify BDDS of new and/or additional information they have received from their raw material suppliers that modifies the animal origin information for lots previously provided to BDDS. When this situation occurs, BDDS updates the animal origin information in the automated certificate system, recreates the affected finished product COAs for batches within expiration date, and sends them to the Internet website where they replace the prior certificate and are immediately available to customers.

Customers enrolled in BD Diagnostic Systems' Automated Change Notification Program will be notified of the changes described above.



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For complete details refer to "BD Position Statement - BD Diagnostic-Diagnostic Systems, COA Animal Origin Information Position Statement", located on the Internet website address at http://www.bd.com/regdocs/.

Juth

John Gerlich Vice President, Quality Management and Regulatory Compliance Signature Date: 2011/08/05