



Becton Dickinson and Company
BD Diagnostic Systems
1475 Athens Highway
Grayson GA 30017 US

Certificate of Analysis

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Product Name : VIAL COAG PLASMA RABBIT EDTA 10X3ML
Catalog Number : 240827 **Manufacture Date** : 2013/09/04
Batch Number : 3255367
Expiration Date : 2016/09/03

Dehydrated Appearance: Off-white to cream, dried button or fluffy material.
Solubility: Soluble in distilled or deionized water.(per package instructions)
Rehydrated Appearance: Off-white to cream, opaque solution without visible fibrin.
Coagulase Test: A fixed amount of Coagulase Plasma was pipetted into one tube per test specimen. 0.05 milliliters of an overnight broth culture was added, mixed and incubated at $37 \pm 1^{\circ}\text{C}$ for six hours. Coagulation was examined after two, four and six hours of incubation.

| TEST ORGANISMS | ATCC® | COAGULASE REACTION |
|----------------------------|-------|--------------------|
| Staphylococcus aureus | 25923 | clot (positive) |
| Staphylococcus aureus | 29213 | clot (positive) |
| Staphylococcus epidermidis | 12228 | no clot (Negative) |
| Enterobacter aerogenes | 13048 | no clot (Negative) |

| Animal Source | Country of Origin | Tissue Category BIC | SIC | ABC |
|---------------|-------------------|---------------------|-----|-----|
| Leporine | USA | IV | IV | IB |

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

BD Diagnostics - Diagnostic Systems products are manufactured in ISO 9001:2008 Registered facilities. In addition, BD Diagnostics - Diagnostic Systems facilities are registered with the United States Food and Drug Administration (FDA), are regulated by the FDA's Quality System Regulations (QSRs), and are also ISO 13485:2003 Registered. 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. This material is not for human or animal consumption.

BD Diagnostics - Diagnostic Systems' Certificates of Analysis (COA) typically contain animal origin information when products are manufactured using materials of animal origin. This information may be contained in the animal source table and/or in one or more of the additional paragraphs found on the COA. Following Quality Control release, the COA is created and published at <http://www.bd.com/regdocs>. For each batch of finished product that contains animal origin raw

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materials, the COA shows the animal origin data from the individual lots of animal origin raw materials used, as provided by the raw material suppliers.

At times, suppliers notify BD Diagnostics - Diagnostic Systems 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 BD. See "COA Animal Origin Information Position Statement" located at <http://www.bd.com/regdocs> under "Position Statements" for the impact that retrospective information has on COAs and on customers enrolled in the BDDS and BDAB Automated Change Notification Program.

For complete details on animal origin information, refer to "BD Position Statement - BD Diagnostic-Diagnostic Systems, COA Animal Origin Information Position Statement", at <http://www.bd.com/regdocs> under "Position Statements".

Paula A. Elliott
WW Quality Manager, Media
BD Diagnostics - Diagnostic Systems
Signature Date: 2014/01/10