



# Certificate of Analysis

Becton Dickinson Caribe LTD.  
BD Diagnostics Systems  
6 Vicks Drive, Lot # 6  
Cayey PR 00737-2860 PR

Page: 1 of 2

**Product Name** : Bottle Plastic Bactec Plus Aerob/F 50/Pk  
**Catalog Number** : 442023 **Manufacture Date**: 2024/02/12  
**Batch Number** : 4032494  
**Expiration Date** : 2024/11/08

This is to certify that representative samples of  
BACTEC PLUS AEROBIC F MEDIUM  
were tested in the Quality Control Laboratory by procedures  
conventionally utilized for this type of product, including  
methodology  
and control ATCC cultures specified in the CLSI standard, Quality  
Assurance for Commercially Prepared Microbiological Culture Media\*,  
and  
met the following test parameters:

pH: 7.3 ± 0.2

#### Autoclaving:

The product was exposed to a moist heat sterilization process  
(previously validated following an ISO Standard\*\*).

#### Vacuum draw:

greater than or equal to 8 mL at time of manufacture.

#### Biological Performance:

Satisfactory growth:

| CULTURE                     | ATCC No. |
|-----------------------------|----------|
| Alcaligenes faecalis        | 8750     |
| Candida glabrata            | 66032    |
| Escherichia coli            | 25922    |
| Haemophilus influenzae      | 19418    |
| Neisseria meningitidis      | 13090    |
| Pseudomonas aeruginosa      | 27853    |
| Staphylococcus aureus       | 25923    |
| ***Streptococcus pneumoniae | 6305     |
| S. pyogenes Group A         | 19615    |

Antimicrobial Removal: Satisfactory

ATCC is a trademark of the American Type Culture Collection.

\* Clinical and Laboratory Standards Institute. 2004. Approved  
Standard,  
M22-A3. Quality assurance for commercially prepared microbiological  
culture media, 3rd ed. CLSI, Wayne, PA.

\*\* ISO 11134, Sterilization of health care products - Requirements for  
validation and routine control - Industrial moist heat sterilization,  
1994.

Creation Date: 2024/03/04 19:48:42



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\*\*\*CLSI Strain

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

BD Diagnostic Systems is an ISO 13485:2016 Registered facility. BD Diagnostics - Diagnostic Systems 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 BD Diagnostics - Diagnostic Systems 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.

Manufacturer is Becton Dickinson and Company, 7 Loveton Circle, Sparks, MD 21152 USA. To determine location of manufacturing for this product, please see [www.bd.com/en-us/support/bd-life-sciences-diagnostic-systems-customer-regulatory-support-information](http://www.bd.com/en-us/support/bd-life-sciences-diagnostic-systems-customer-regulatory-support-information).

Jose E. Mojica  
Senior Quality Manager  
Signature Date:2024/03/04

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