Certificate of Analysis

Client

E&O Laboratories Ltd Burnhouse Bonnybridge Scotland FK4 2HH



Burnhouse, Bonnybridge Scotland, FK4 2HH Telephone: 01324 840404 Fax:01324 841314 Email: info@eolabs.com

Sample: BM0760 - Max. Recovery Diluent

Batch Number: 04096857 **Expiry Date:** 2022-05-11

Date Received: 2021-11-11

Date Tested: 2021-11-11

Date of Issue: 2021-11-15

Sample Condition: Satisfactory

Sample Number: 412868

The log difference between inocula pre and post a 45 minute retention in the diluent should be ≤ 0.3. Accredited Test Method: See table below			
Productivity	Result	Specification	
Escherichia coli NCTC 12241	Conforms	≤ 0.3	
Staphylococcus aureus NCTC 12981	Conforms	≤ 0.3	

Physical	Result	Specifications	Accredited Test Method
Sterility	Conforms	Within acceptable limits	ED/SOP/005 Visual check and growth assessment following incubation for 3 days at 15-25°C and 37°C
	•	n accordance with ISO 2859-1:1 ling on batch size.	999*. The inspection level is \geq 0.4% of the batch and
рН	7.2	7.0 +/- 0.2	ED/SOP/003 measurement by pH meter
Colour	Conforms	Colourless	ED/SOP/009 by visual observation. Range measured using Pantone® 4 colour process guide
Fill Quantity	Conforms	5000ml +/- 0.3	ED/SOP/054 by gravimetric determination

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Sample Condition: Satisfactory

Accredited Test Method for Solid Media

ED/SOP/008 Quantitative evaluation

using spread inoculum technique

Sample Number: 412868

ED/SOP/008 Semi quantitative inoculation with growth assessment from subculture

Accredited Test Method for Diluents

ED/SOP/008 Semi quantitative evaluation of viability maintenance after a holding time of 45 minutes

All of the results on this certificate of analysis relate only to the samples submitted.

Test specifications are based on ISO 11133:2014/Amd/:2020 and internal product specifications

*Sterility sampling is outwith scope of accreditation.



Douglas Cameron

Technical Manager, E&O Laboratories Ltd