

## EC Declaration of Conformity – Issue 4

E&O Laboratories Ltd  
Burnhouse  
Bonnybridge  
Scotland  
FK4 2HH  
info@eolabs.com

Declares that its IVD products have been classified and registered in accordance with Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017.

Each of the products meets the relevant provisions of the Regulation and have been classified as Class A within Annex VIII of the regulation.

Products have been registered with the Health Products Regulatory Authority (HPRA) via E&O's EU Authorised Representative

Syntec Scientific Ltd  
Unit 2, The Business Park  
Northwest Business Park  
Phase 3, Ballycoolin  
Dublin 15  
Ireland  
D15 PY00  
IE-AR-000036690

E&O Laboratories Ltd has developed, implemented, and maintains a documented post-production monitoring process, including the notification of reportable events in compliance with Article 82 Reporting of serious incidents and field safety corrective actions.

Eddie Scott (Quality Director)

Date



14/09/2023



**EO Labs**  
Microbiological Culture Media

E & O Laboratories Ltd  
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Scotland, FK4 2HH  
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## EC Declaration of Conformity Schedule

### Class A Products

#### GMDN

- 58685 Basic /simple nutrient agar culture medium IVD
- 58586 Basic/simple nutrient broth culture medium IVD
- 58585 Differential/selective agar culture medium IVD
- 58684 Differential/selective broth culture medium IVD
- 58688 Enriched agar culture medium IVD
- 58687 Enriched broth culture medium IVD
- 62106 Multiple formulation agar culture medium IVD
- 62392 Oral/respiratory tract specimen container IVD, additive/medium
- 58605 General -microorganism antimicrobial susceptibility culture medium IVD
- 58636 Motility agar culture medium IVD
- 62819 Blood IVD
- 58216 Oral/upper respiratory tract specimen collection kit IVD, clinical
- 62773 General specimen collection kit IVD, clinical
- 62707 Culture Medium Base Ingredient IVD