

EU DECLARATION OF CONFORMITY FOR IN VITRO DIAGNOSTIC MEDICAL DEVICES

Manufacturer Name Liofilchem® S.r.l.

Manufacturer Address Via Scozia, 64026 Roseto degli Abruzzi (TE) - Italy

SRN (Single Registration Number) IT-MF-000026495

Basic UDI-DI, Name of the Device(s), Product code(s), Intended Purpose

See Table no.1

Classification in accordance with the

rules set out in Annex VIII

Class C

References to CS used and in relation to

which conformity is declared

Not yet applicable

Description of the conformity assessment

procedure performed:

Annex IX

Identification of the certificate issued: EU Quality Management System Certificate (IVDR) No. V12

071067 0008

Notified Body Name: TÜV SÜD Product Service GmbH

Notified Body Identification Number: 0123

This declaration of conformity is written according to Annex IV of Regulation (EU) 2017/746 and is issued under the sole responsibility of Liofilchem S.r.l.

We hereby declare that the in vitro diagnostic medical device specified above meet the provision of the Regulation (EU) 2017/746 for in vitro diagnostic medical devices.

All supporting documentation is retained at the premises of the manufacturer.

Roseto degli Abruzzi (TE),

07.03.2025

Signature:

LIOFILCHEM s.r.l.
BACTERIOLOGY PRODUCTS
Via Scozia
64028 Roseto degli Abruzzi (TE)
Cod. Fisc. e Partita IVA 00530130673

Technical Director (Dr. Silvio Brocco)



Table no.1

Code	Description	Basic UDI-DI	Intended purpose
Antibiotic Discs			
9253 9253/1 9253/2	Imipenem- relebactam (10+25) I/R 35 μg	805518287ATBDISC253Z3	Imipenem-relebactam (10+25) I/R 35 µg is an <i>in vitro</i> semi-quantitative method for antimicrobial susceptibility of clinical isolates tested on agar media using overnight incubation. The test is used against the following microorganisms: • Gram-negative bacteria • Enterobacterales • Pseudomonas aeruginosa
9270 9270/1 9270/2	Cefepime- enmetazobactam (30+20) FPE 50 µg	805518287ATBDISC270Z3	Cefepime-enmetazobactam (30+20) FPE 50 µg is an <i>in vitro</i> semi- quantitative method for antimicrobial susceptibility of clinical isolates tested on agar media using overnight incubation. The test is used against the following microorganisms: • Gram-negative bacteria • Enterobacterales
9272 9272/1 9272/2	Aztreonam- avibactam (30+20) AZA 50 µg	805518287ATBDISC272Z7	Aztreonam-avibactam (30+20) AZA 50 µg is an in vitro semi-quantitative method for antimicrobial susceptibility of clinical isolates tested on agar media using overnight incubation. The test is used against the following microorganisms: • Gram-negative bacteria • Enterobacterales

DoC Class C Rev.1 Page 2 of 2