

Test form for the External Quality Assessment (EQA) for laboratories participating in the European Antimicrobial Resistance Surveillance Network (EURGen-Net), 2025 – Pseudomonas aeruginosa

Participating laboratories can only submit results online via the webtool. This form cannot be submitted.

Kindly note, due to the Material Transfer Agreements (MTAs) between SSI and the original providers of the isolate:

- 1. Strains received for the 2025 EURGen-Net EQA cannot be re-distributed further by the recipient laboratories.
- 2. It is not possible for DTU to distribute strains to laboratories after the EQA exercise, e.g. for confirmatory, training, or reference purposes.

It is recommended to store the strain in your strain collection (e.g. in a -80°C freezer), at least until you have reviewed your results from this EQA exercise. This will allow for repetition of species identification and AST, if needed, in light of your individual performance.

This form is designed to help participating laboratories prepare their results before submission in the EURGen-Net EQA webtool (https://EURL-PH-AMR.eqa.dtu.dk). The EURGen-Net phenotypic EQA is coordinated by DTU Food, in their capacity as a consortium member of the EURL-PH-AMR, in consultation with ECDC.

When submitting the results online in the webtool, participants will be asked for the following information:

Pseudomonas aeruginosa - strain no

The isolate should be considered as being obtained from a patient with a bloodstream infection.

For colistin and aminoglycosides (amikacin and tobramycin), it should be assumed that the antimicrobials will be administered in combination with other agents.

Breakpoints currently based on ECOFF values can be used for interpretation of results, when applicable, if no other relevant EUCAST clinical breakpoints exist.

Currently, EUCAST recommends using disk diffusion for testing of cefiderocol, but only after consulting the EUCAST Warnings! page (Warning 12) about certain media and disks (https://www.eucast.org/ast-of-bacteria/warnings).

Non-reported results will not be scored, but the antimicrobials and respective empty result sections will still be visible in the individual evaluation reports.





INFORMATION ABOUT THE METHOD

1. Which methodology did you use for s	species identification of this strain?
☐ Biochemical tests	
☐ MALDI-TOF	
☐ Whole-genome sequencing	
☐ Other sequencing (e.g. 16S rRNA)	
□ PCR	
☐ Other – Specify:	
2. Which mothodology did you mainly u	use for antimicrobial susceptibility testing (AST) of this
strain?	ise for antimicrobial susceptibility testing (AST) of this
☐ Automated system	
☐ Disk/Tablet diffusion	
☐ Broth microdilution	
☐ Gradient test	
☐ Macro broth dilution (tubes)	
☐ Agar dilution	
☐ Other – specify:	
In the webtool, the type of method will by d aware that the settings below will change, if	efault be set to the information already provided above, and be you later change method above.
Antimicrobial	Method
Amikacin	
Aztreonam	
Cefiderocol	
Ceftazidime	
Ceftazidime-avibactam fixed 4 mg/L	
Ceftolozane-tazobactam fixed 4 mg/L	
Ciprofloxacin	
Colistin	
Imipenem	
Imipenem-relebactam fixed 4 mg/L	
Levofloxacin	
Meropenem	
Meropenem-vaborbactam fixed 8 mg/L Piperacillin-tazobactam fixed 4 mg/L	



Tobramycin



o Selecte	ed Automated system in the table above, please specify the instrument Microscan Walkaway
0	Phoenix
0	VITEK
0	Other – specify:
If you selecte	ed Disk/Tablet diffusion in the table above, please specify the origin of the
disks/tablets	
0	BD/BBL sensi disc
0	Liofilchem
0	MAST
0	Neo sensitabs
0	Oxoid
0	Other– specify:
If you selecte	ed Disk/Tablet diffusion in the table above, please specify the origin of the agar
0	BD BBL MH II Agar (Becton Dickinson)
0	Biolife MH Agar II (Biolife Italiana)
0	bioMerieux MHE Agar (bioMérieux)
0	Bio-Rad MH Agar (Bio-Rad Laboratories)
0	E&O Laboratories MH Agar (E&O Laboratories)
0	Hardy Diagnostics MH Agar (Hardy Diagnostics)
0	HiMedia MH Agar (HiMedia)
0	HiMedia MH Agar no. 2 (HiMedia)
0	Liofilchem MH II Agar (Liofilchem)
0	Oxoid MH Agar (Thermo Scientific)
0	Other – specify:
If you selecte	ed Broth microdilution in the table above, please specify the test and origin
0	ComASP
0	Liofilchem
0	Sensititre MIC plates
0	UMIC (Bruker)
0	MIC plates prepared in-house
0	Other– specify:
If you selecte	ed Broth microdilution in the table above, please specify the origin of the broth
0	BD BBL
0	Oxoid
0	Sensititre
0	Sigma-Aldrich
0	Other – specify:
If you selecte	ed Gradient test in the table above, please specify the test and origin

- o E-test (bioMérieux)
- o MIC strip (Liofilchem)
- o Other specify: _____

If you selected Gradient test in the table above, please specify the origin of the agar

o BD BBL MH II Agar (Becton Dickinson)



o Bio	olife MH Agar II (Biolife Italiana)
o bio	oMerieux MHE Agar (bioMérieux)
o Bio	o-Rad MH Agar (Bio-Rad Laboratories)
o E8	&O Laboratories MH Agar (E&O Laboratories)
о На	ardy Diagnostics MH Agar (Hardy Diagnostics)
o Hi	Media MH Agar (HiMedia)
o Hi	Media MH Agar no. 2 (HiMedia)
o Lic	ofilchem MH II Agar (Liofilchem)
o Ox	xoid MH Agar (Thermo Scientific)
o Ot	ther – specify:
3. Which standar	rd/guideline did you use when performing AST?
☐ EUCAST	– specify breakpoint table version:
☐ Other –	- specify:
4. Would you no	rmally send this (invasive!) strain to a reference or other laboratory? (Please
note that the EQ	A strains cannot actually be redistributed further).

RESULTS

☐ Yes \square No

Strain ID	Antimicrobial	Results and interpretation			
		≤/=/>	MIC value (mg/L) or zone diameter (mm)	S/I/R/NA	
	Amikacin				
	Aztreonam				
	Cefiderocol				
	Ceftazidime				
	Ceftazidime-avibactam fixed 4 mg/L				
	Ceftolozane-tazobactam fixed 4 mg/L				
	Ciprofloxacin				
	Colistin				
	Imipenem				
	Imipenem-relebactam fixed 4 mg/L				
	Levofloxacin				
	Meropenem				
	Meropenem-vaborbactam fixed 8 mg/L				
	Piperacillin-tazobactam fixed 4 mg/L				
	Tobramycin				