



Test form for the External Quality Assessment (EQA) for laboratories participating in the European Antimicrobial Resistance Surveillance Network (EARS-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 DTU and the original providers of the isolate:

- 1. Strains received for the 2025 EARS-Net EQA cannot be re-distributed further by the recipient laboratories.
- 2. It is not possible for DTU or the National EARS-Net EQA Coordinator 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 EARS-Net EQA webtool (https://earsnet.eqa.dtu.dk/). It is based on the test form for 2024. In 2021–2024, the EARS-Net EQA was coordinated by DTU Food, with ECDC, through a framework contract. From 2025, the EARS-Net 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, gentamicin 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.







TEST FORM METHODS

1. Please select a type of laboratory	
☐ National reference laboratory (or laborat	ory with similar functions)
☐ Regional reference laboratory (or laborat	ory with similar functions)
☐ Local laboratory	
2. Which methodology did you mainly use for antin	nicrobial susceptibility testing (AST) of this strain?
☐ Automated system	, , , ,
☐ Disk/Tablet diffusion	
☐ Broth microdilution	
☐ Gradient test	
☐ Macro broth dilution (tubes)	
☐ Agar dilution	
☐ Other – specify:	
🗆 Other – specify.	
	ntibiotics, please change the method for the relevant
antimicrobial(s) below.	
In the webtool, the type of method will by default be	e set to the information already provided above, and be
aware that the settings below will change, if you late	er change method above.
Antimicrobial	Method
Amikacin	
Cefepime	
Ceftazidime	
Ciprofloxacin	
Colistin	
Imipenem	
Levofloxacin	
Meropenem	
Meropenem Piperacillin	
·	
Piperacillin	
Piperacillin Piperacillin-tazobactam (fixed 4)	
Piperacillin Piperacillin-tazobactam (fixed 4) Tobramycin	

If you selected Automated system in the table above, please specify the instrument

- Microscan Walkaway
- o Phoenix
- o VITEK

Imipenem-relebactam (fixed 4)
Meropenem-vaborbactam (fixed 8)

DTU





0	Other – specify:
If you selected	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 selected	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 selected	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 selected	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 selected	Gradient test in the table above, please specify the test and origin
0	E-test (bioMérieux)
0	MIC strip (Liofilchem)
0	Other – specify:
If you selected	Gradient test in the table above, please specify the origin of the agar
0	BD BBL MH II Agar (Becton Dickinson)

- o Biolife MH Agar II (Biolife Italiana)
- o bioMerieux MHE Agar (bioMérieux)
- o Bio-Rad MH Agar (Bio-Rad Laboratories)
- o E&O Laboratories MH Agar (E&O Laboratories)
- Hardy Diagnostics MH Agar (Hardy Diagnostics)
- o HiMedia MH Agar (HiMedia)







- o HiMedia MH Agar no. 2 (HiMedia)
- o Liofilchem MH II Agar (Liofilchem)
- o Oxoid MH Agar (Thermo Scientific)
- Other specify: _____

. Which standard/guideline did you use when performing AST?	
☐ EUCAST – specify breakpoint table version:	
☐ Other – specify:	
. Would you normally send this (invasive!) strain to a reference or other laboratory? (Please note tha	t
ne EQA strains cannot actually be redistributed further).	
□ Yes	
□No	





TEST FORM RESULTS

	≤/=/>	2010 1 / /1)	1
	-7 7	MIC value (mg/L) or zone diameter (mm)	S/I/R/ NA
Amikacin			
Cefepime			
Ceftazidime			
Ciprofloxacin			
Colistin			
Imipenem			
Levofloxacin			
Meropenem			
Piperacillin			
Piperacillin-tazobactam (fixed 4)			
Tobramycin			
Cefiderocol			
Ceftazidime-avibactam (fixed 4)			
Ceftolozane-tazobactam (fixed 4)			
Imipenem-relebactam (fixed 4)			
Meropenem-vaborbactam (fixed 8)			