

## Proteus mirabilis (LTR62809)

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**Organism**
**Proteus mirabilis**


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**Clinical**

This organism is found in a variety of environmental sources and may be part of the normal flora of the gastrointestinal tract. It is a cause of both community and nosocomial urinary tract infections, usually catheter associated infections or in chronically catheterized patients. It accounts for 90% of *Proteus spp.* infections. Infection with this organism indicates upper urinary tract involvement.

*P. mirabilis* has also been associated with post-operative wound infections, skin/soft tissue infections, and bacteremia (usually secondary to a urinary tract focus).

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**Usual  
susceptibility  
pattern**

This organism does not produce any chromosomal beta-lactamase and is usually susceptible to ampicillin and cephalosporin. However, plasmid mediated beta-lactamases [penicillinase, cephalosporinase, extended spectrum beta-lactamases (ESBL) and carbapenemase (metalloenzyme, KPC or Class D OXA enzyme)] are increasingly being described. ESBL production may be more difficult to detect. Beta-lactam resistance may also be mediated by permeability mutations. *P. mirabilis* is usually susceptible to aminoglycosides, quinolones, TMP-SMX and carbapenems. (Exception: May exhibit decreased susceptibility to imipenem (decreased affinity to PBP2) that does not affect other carbapenems.)

*P. mirabilis* is resistant to nitrofurantoin, tetracycline and colistin.

**Note:** Although generally susceptible to beta-lactam antibiotics, non-beta-lactam antibiotics appear to be more efficacious in treatment of urinary tract infections with this organism.

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**Susceptibility  
method**

VITEK2. Additional tests (Disc diffusion or Etest method) are performed using Mueller-Hinton agar incubated in ambient air at 35°C for 16-20 hours.

**Note:** For Etest use 0.5 McFarland suspension in saline.  
For mucoid strains use 1.0 McFarland.

**Proteus mirabilis, Continued**

**Susceptibility reporting**

	CSF/ Brain	Blood/ Endo- vascular Catheter	Sterile Body Site	Urine	Other	Comments
Amikacin		3	3	3	3	3 <sup>rd</sup> line if gent and tobra I/R Disc diffusion
Amoxicillin/ Clavulanate oral				✓	✓	If Amp S and Amox-Clav I/R – Do Amp and AMC disc diffusion, consult tech 2 with results <b>See Special Considerations</b>
Amoxicillin/ Clavulanate IV		2*	2*		2*	2 <sup>nd</sup> line if ampicillin I/R, ceftazidime I/R and ceftriaxone S *Report 1 <sup>st</sup> line if ampicillin I/R and ceftriaxone S and anaerobes, <i>Enterococcus</i> or <i>S. aureus</i> (MSSA) co-isolated Report same as AMC oral
Ampicillin	*	✓	✓	✓	✓	* Report only in neonates (<1 month)
Cefazolin		✓	✓	✓*	✓	*If MIC ≤16 Do not report (offer); ≥32 report as R. Refer to Beta-Lactam Resistance Detection Charts
Cefixime				✓		
Ceftriaxone	✓*	✓*	2*	2*	2*	Always report if I/R 2 <sup>nd</sup> line if cefazolin I/R *If patient < 1 mo - report cefotaxime instead of ceftriaxone using the same interpretation.
Cephalexin				✓		<b>Add comment:</b> For uncomplicated lower UTI only <b>#clx1</b>
Ciprofloxacin		✓	✓	2*	✓	Do not report in patients < 18 y 2 <sup>nd</sup> line if cefixime and TMP-SMX I/R. Always report if I/R <b>*See Special Considerations</b>
Ertapenem		3	3	3	3	3 <sup>rd</sup> line if ceftriaxone or ceftazidime I/R If S do not report in patients < 3 months <b>See Special Considerations</b>
Gentamicin	*	✓**	✓**	✓	✓**	* Report only in neonates (< 1 month) <b>**See Special Considerations</b>
Meropenem	2	3	3	3	3	2 <sup>nd</sup> or 3 <sup>rd</sup> line if ceftriaxone or ceftazidime I/R <b>See Special Considerations</b>
Nitrofurantoin				R		<b>Add comment:</b> For uncomplicated lower UTI only <b>#f1</b>
Piperacillin/ Tazobactam		3* †	3*		3*	3 <sup>rd</sup> line if AMC IV R and ceftriaxone S * Report if <i>P.aeruginosa</i> co-isolated and ceftriaxone S † For bloods report if I/R and ceftriaxone S <b>See Special Considerations</b>
TMP-SMX	*	✓	✓	✓	✓	*Report only at physician request
Tobramycin		2*	2*	2	2*	2 <sup>nd</sup> line if gent I/R <b>*See Special Considerations</b>

## Proteus mirabilis, Continued

### Special considerations

<p><u>Amoxicillin/Clavulanate oral:</u></p>	<p>A MIC of 8/4 µg/mL is at upper limit of susceptibility. This may be adequate to achieve reasonable pharmacodynamics in urine but may not be optimal for non-urinary sites.</p> <p>For all non-urinary sites if MIC 8/4 µg/mL and interpretation is S add comment:            “This isolate tests at the upper limit of susceptibility for amoxicillin/clavulanate. Clinical failure may occur despite in vitro susceptibility”. <b>#A315</b></p> <p><b>I/R VITEK 2 results are not recommended due to a card limitation. Perform an alternate method prior to reporting of results.</b></p>
<p><u>Ciprofloxacin:</u></p>	<p>For urine cultures add the following comment when <b>not</b> reporting ciprofloxacin (patients ≥ 18 y):</p> <p>“Ciprofloxacin is not routinely reported, given the potential for significant adverse events and increasing antimicrobial resistance.” <b>&amp;3206</b></p>
<p><u>Ertapenem / Meropenem:</u></p>	<p>If reporting ertapenem and meropenem add comment:            “Imipenem has intrinsically low activity against this organism.” <b>#A375</b></p>
<p><u>Gentamicin/tobramycin:</u></p>	<p>Organisms testing at upper limit of susceptibility (4µg/mL) may not achieve optimal pharmacokinetics/pharmacodynamics.</p> <p><b>For non-urine isolates:</b>            If MIC 4.0 µg/mL add comment:            “This isolate tests at the upper limit of susceptibility for gentamicin. Clinical failure may occur despite in vitro susceptibility”. <b>#A312</b>  <b>or</b>            “This isolate tests at the upper limit of susceptibility for tobramycin. Clinical failure may occur despite in vitro susceptibility”. <b>#A313</b>  <b>or</b>            “This isolate tests at the upper limit of susceptibility for both gentamicin and tobramycin. Clinical failure may occur despite in vitro susceptibility”. <b>#A314</b></p>
<p><u>Piperacillin/tazobactam:</u></p>	<p>This antibiotic is frequently used as empiric therapy for polymicrobial infections (i.e. co-infections with <i>S. aureus</i>, <i>Enterococcus</i>, <i>Pseudomonas aeruginosa</i> and/or anaerobes), febrile neutropenia or sepsis syndromes.</p> <p><b>Note:</b> Do not report as S if ceftriaxone or ceftazidime I/R (≥2 µg/mL) as piperacillin/tazobactam is not recommended if either an extended spectrum beta-lactamase (ESBL) and/or a cephalosporinase is present.</p>

## Proteus mirabilis, Continued

**Interpretation** For Etest, report actual MIC result. For interpretation (S, I, and R) report according to nearest higher doubling dilution (**Appendix 1**).

Use **CLSI** interpretive document for **Enterobacterales**.

**For Beta-lactam drugs – Refer to Beta-lactam Resistance Detection Charts.**  
**For Amoxicillin/Clavulanate, gentamicin and tobramycin – Refer to Special Considerations**