# Cost-savings Analysis with Use of Omadacycline Among Hospitalized Community-Acquired Pneumonia Patients At Risk of *Clostridium difficile* Infection Being Treated with Moxifloxacin: Budget Impact Model Findings

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## BACKGROUND

- Clostridium difficile (CDI) infection is a substantial cause of morbidity and mortality across the United States (US)
- CDI has been reported with the use of nearly all antibacterial agents and ranges in severity from mild diarrhea to fatal colitis
- CDI places a huge burden on healthcare systems, particularly for hospitalized patients, because it is associated with extended hospital stays, repeat admissions, and significant mortality
- In 2011, *C. difficile* was responsible for nearly 500,000 infections and associated with ~29,000 deaths, in the USA<sup>1</sup>
- A meta-analysis of CDI rates associated with antibiotic use suggests that tetracyclines may be linked to a decreased risk of CDI compared with other classes of antibiotics<sup>2</sup>
- This suggests that it may be appropriate to use tetracyclines to decrease the burden of CDI associated with antibiotic use<sup>2</sup>
- Omadacycline (OMC; Paratek Pharmaceuticals, Inc., King of Prussia, PA) is an aminomethylcycline antibiotic that was recently approved in the US for the treatment of adult patients with community-acquired bacterial pneumonia (CABP)<sup>3</sup>
- In the Phase 3 Omadacycline for Pneumonia Treatment In the **C**ommunity (OPTIC) study, OMC was shown to be non-inferior to moxifloxacin (MOX) in the treatment of patients with CABP (NCT02531438)<sup>3</sup>
- Adverse event rates were comparable between treatment groups. However, C. difficile infections occurred in 8 patients (2%) who received MOX and 0 patients who received OMC<sup>3</sup>
- A budgetary impact model was created to highlight costsaving opportunities for treating patients hospitalized for CABP with OMC compared to MOX, taking the risk and cost of treatment-associated CDI into account

### **Model Structure and Populations**

### **Costs Included**

- Hospital room and board fees were assumed to be the same for OMC and MOX
- Drug acquisition costs (MOX: US\$46/day, wholesale acquisition cost)
- OMC acquisition cost varied between \$150 and \$600/day
- No additional adverse events were considered

### **Sensitivity Analysis**

- Sensitivity analysis with incremental CDI incidence of MOX (0-12%) and cost of CDI treatment (Low: \$1,522; Mean: \$34,149; High: \$122,318) were undertaken to capture incidence uncertainty
- The assumptions were that treatment with OMC has a lower propensity to induce CDI relative to MOX and has the potential to avoid CDI events, leading to a reduction in overall hospital costs

- From the hospital/health system perspective and a CDI treatment cost of \$34,149, for every 100 patients treated with OMC instead of MOX, the incremental cost of reductions in CDI rate with OMC ranged between \$52,000 and \$132,884 (cost saving) depending on the acquisition cost of OMC and CDI incidence for MOX (0-12%) (Fig.1; Table 1)
- Analysis showed that for OMC to become cost saving, the incidence of CDI in MOX-treated patients would need to range between 1.5% and 8.1%, depending on the acquisition cost of OMC (\$150-\$600/day)
- At the lower end of the cost per hospitalized CDI case, OMC would not be cost saving at any price point
- As the cost per case of CDI increases, OMC use becomes cost saving for a hospital compared with MOX at lower CDI incidence rates

# METHODS

• The economic model was developed using a hospital perspective to estimate the budget impact of replacing the current strategy of treating 100 patients hospitalized for CABP with 5 days of MOX, to treating with 5 days of OMC – This time period reflects the initial 5 days of a minimum treatment of 7 days for either OMC or MOX in CABP

- CDI adverse event treatment (average cost-per-case
- attributed to hospital-onset CDI was \$34,1574)

# RESULTS





RESULTS													CONCLUSIONS	
Figure Moxif	Figure 1. Incremental Cost per 100 Patients Treated with Omadacycline Relative to Moxifloxacin at Varying Incidences of Moxifloxacin-associated CDI Event Rates													• The model illustrated the economic impact associated with reductions in CDI rates with omadacycline compared to moxifloxacin, when treating patients hospitalized for CABP
Incremental Cost per 100 Patients (\$34,149) \$150										- \$150	<ul> <li>As part of the analysis, the model demonstrated the incremental increase in CDI rates with moxifloxacin compared to omadacycline that conferred cost savings</li> </ul>			
\$30 \$20 \$10 -\$10 -\$20	0,000. 0,000. 0,000. \$0. 0,000. 0,000.	$ \begin{array}{c} 00 - \\ 0$											\$300 \$450 \$600	Use of omadacycline has the potential to reduce the economic burden associated with patients hospitalized for CABP who are treated with moxifloxacin if it can avoid approximately 2-8 cases of moxifloxacin-associated CDI per 100 patients
-\$3( -\$4( -\$5(	0,000. 0,000. 0 000	00 - 00												<ul> <li>Future studies are required to identify CABP patients at greatest risk of moxifloxacin-associated CDI</li> </ul>
ΨΟ	,	0	0 1 2 3 4 5 6 7 8 9 10 11 12 %						9	12		<ul> <li>Like all studies of this nature, the findings in this analysis and modeling require validation in real-world settings</li> </ul>		
CDI-											REFERENCES			
attributable Cost per Case, \$ <sup>4</sup>			1,522 (Lov	ver Range)		34,149 (Mean) 122,318 (Upper R						oper Range)		<ol> <li>Lessa FC, et al. N Engl J Med. 2015;372(9):825-834.</li> <li>Tariq R. Clin Infect Dis. 2018;66(4):514-522.</li> </ol>
Omadacycline Daily Acquisition												<ol> <li>Stets R, et al. Open Forum Infect Dis. 2017;4:S543-S544.</li> <li>Zhang S, et al. BMC Infect Dis. 2016;16(1):447.</li> </ol>		
Cost, \$		150	300	450	600	150	300	450	600	150	300	450	600	
CDI Incidence, %	0	29,372	109,322	189,272	269,222	29,372	109,322	189,272	269,222	29,372	109,322	189,272	269,222	
	1	27,850	107,800	187,750	267,700	-4,776	75,173	155,123	235,073	-92,945	-12,995	66,954	146,904	Thomas Lodise: Consultant - Paratek Pharmaceuticals, Inc. Kenneth LaPensee: Employee - Paratek Pharmaceuticals, Inc. Rohit Mistry: Consultant - Paratek Pharmaceuticals, Inc. Kate Young: Consultant - Paratek Pharmaceuticals, Inc. This work was funded by Paratek Pharmaceuticals, Inc. Medical editorial assistance, funded by Paratek Pharmaceuticals, Inc., was provided by Innovative Strategic Communications.
	2	26,328	106,278	186,228	266,178	-38,925	41,024	120,974	200,924	-215,263	-135,313	-55,363	24,586	
	3	24,806	104,756	184,706	264,656	-73,074	6,875	86,825	166,775	-337,581	-257,631	-177,681	-97,731	
	4	23,284	103,234	183,184	263,134	-107,223	-27,273	52,676	132,626	-459,899	-379,949	-299,999	-220,049	
	5	21,762	101,712	181,662	261,612	-141,372	-61,422	18,527	98,477	-582,217	-502,267	-422,317	-342,367	
	0	20,240	00 660	170 610	200,090	-175,521	-95,571	-13,021	04,320 20.170	-704,000	-024,303	-544,035	-404,000	
	/ Q	17 106	90,000	177,006	250,500	-209,070	162 260	-49,770	2,060	-020,000	-740,903	-000,900	-307,003	
	0	15 674	97,140	175 574	257,040	-243,019	102,009	112 062	-3,909 20 110	-949,171	-009,221	-709,271	-709,321	ACKNOWLEDGMENTS
	10	14 152	93,024	174 052	254 002	-217,300	-232 167	-152 217	-72 267	-1,071,403	-1 113 857	-1 033 907	-051,009	The authors wish to thank the PAREXEL Access RWE team, who provided real-world data used in this study.
	11	12 630	92 580	172,530	252 480	-346 266	-266,316	-186,366	-106 416	-1 316 125	-1 236 175	-1 156 225	-1 076 275	
	12	11,108	91,058	171,008	250,958	-380,415	-300,465	-220,515	-140,565	-1,438,443	-1,358,493	-1,278,543	-1,198,593	
CDI = Clostr	dium difficile	nfection												

Cost of nospital-onset CDI tested over the range of values reported by Zhang et al Purple highlighted cells show cost savings (negative \$ values) associated with omadacycline in the model, at the given omadacycline costs and CDI incidence rates.