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Abstract (poster session)

**Antimicrobial activity of PTK 0796 (omadacycline) and comparator agents against contemporary pathogens commonly associated with community-acquired respiratory tract infections collected during 2011 from the European Union**

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**Objectives:** To determine the activity of PTK 0796 (omadacycline) and comparator agents against recent (2011) *Streptococcus pneumoniae* (SPN), *Haemophilus influenzae* (HI), and *Moraxella catarrhalis* (MCAT) isolated in the European Union (EU). PTK 0796 is a novel aminomethylcycline which is currently under clinical development for both intravenous and oral formulations. It has excellent activity against pathogens from the respiratory tract and overcomes tetracycline resistance. **Methods:** Susceptibility (S) testing for omadacycline and commonly used antimicrobials was performed by Clinical and Laboratory Standards Institute (CLSI) broth microdilution methodology on a total of 1,025 isolates in 2011 from medical centers in the SENTRY Antimicrobial Surveillance Program platform in the EU. S interpretations were performed using CLSI and EUCAST guidelines. **Results:** PTK 0796 was very active against SPN independent of S to penicillin (PEN; MIC<sub>50/90</sub>, 0.06/0.06 mg/L for PEN-S and -resistant [R] strains). PTK 0796 was 16-fold more active than levofloxacin (MIC<sub>50/90</sub>, 1/1 mg/L) and ceftriaxone (MIC<sub>90</sub>, 1 mg/L) against SPN. SPN showed high R rates to erythromycin (S, 60.7%) and tetracycline (S, 69.2% CLSI/68.9% EUCAST) even though all isolates had PTK 0796 MIC values  $\leq$  0.25 mg/L. PTK 0796 against HI (13.9% beta-lactamase positive) and MCAT (98.5% beta-lactamase positive) exhibited low MIC values (Table) independent of beta-lactamase production. **Conclusions:** PTK 0796 was very active against SPN, regardless of PEN-S status, with MIC<sub>50/90</sub> value of 0.06/0.06 mg/L and no MIC value greater than 0.25 mg/L. PTK 0796 was also very active against *M. catarrhalis* (MIC<sub>50/90</sub>, 0.12/0.12 mg/L) and *H. influenzae* (MIC<sub>50/90</sub>, 0.5/1 mg/L) with activity independent of beta-lactamase status.

	Organism (no. tested)		
	SPN (601)	HI (359)	MCAT (65)
<b>PTK 0796</b>			
<b>MIC<sub>50/90</sub><sup>a</sup></b>	0.06 / 0.06	0.5 / 1	0.12 / 0.12
<b>Ceftriaxone</b>			
<b>MIC<sub>50/90</sub><sup>a</sup></b>	≤0.06 / 1	≤0.06 / ≤0.06	0.25 / 0.5
<b>%S<sup>b</sup></b>	91.9 (73.0)	100.0 (100.0)	100.0 (98.5)
<b>Erythromycin</b>			
<b>MIC<sub>50/90</sub><sup>a</sup></b>	≤0.12 / >16	NT <sup>e</sup>	NT <sup>e</sup>
<b>%S<sup>b</sup></b>	60.7 (60.7)	NT <sup>e</sup>	NT <sup>e</sup>
<b>Levofloxacin</b>			
<b>MIC<sub>50/90</sub><sup>a</sup></b>	1 / 1	≤0.12 / ≤0.12	≤0.12 / ≤0.12
<b>%S<sup>b</sup></b>	98.7 (98.7)	100.0 (100.0)	100.0 (100.0)
<b>Tigecycline</b>			
<b>MIC<sub>50/90</sub><sup>a</sup></b>	≤0.03/0.06	0.25/0.5	0.06/0.06
<b>%S<sup>c</sup></b>	100.0 (IE) <sup>d</sup>	87.2 (IE) <sup>d</sup>	- (IE) <sup>d</sup>

a. MIC values in mg/L; b. Categorized by CLSI breakpoints and in parentheses by EUCAST breakpoints; c. categorized by USA drug package insert breakpoints and in parentheses by EUCAST breakpoints; d. IE= insufficient evidence to determine breakpoint; e. NT= not tested