## In Vitro Susceptibilities of Aeromonas hydrophila to 32 Antimicrobial Agents

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Minimal inhibitory concentrations of 32 antimicrobial agents for 20 strains of *Aeromonas hydrophila* were determined by a microdilution method. Moxalactam was the most active drug tested. All strains were also susceptible to clinically achievable concentrations of mecillinam, cefamandole, cefuroxime, cefotaxime, aminoglycosides (except streptomycin), tetracycline, chloramphenicol, and trimethoprim-sulfamethoxazole.

Aeromonas hydrophila is an organism found in natural water sources which occasionally causes human infections. In recent years, it has been recognized as the cause of a variety of serious infections, particularly in compromised hosts, and as a nosocomial pathogen (1, 6, 7). Susceptibilities to antimicrobial agents have varied, but isolates were usually susceptible to chloramphenicol, tetracycline, aminoglycosides, and trimethoprim-sulfamethoxazole and relatively resistant to penicillins, cephalosporins, erythromycin, and polymixins (1, 2, 3-7).

In this study, we tested the susceptibilities of 20 strains of *A. hydrophila* to 32 antimicrobial agents. The isolates were from patients hospitalized in University Hospitals, Columbus, Ohio during the past two years. Minimal inhibitory concentrations were determined by a microdilution method in cation-supplemented Mueller-Hinton broth (Difco Laboratories, Detroit, Mich.), using an inoculum size of approximately  $2 \times 10^5$  colony-forming units per ml (Fass and Barnishan, Rev. Infect. Dis., in press).

Results are shown in Table 1. Among the  $\beta$ lactam antibiotics, moxalactam was most active. Mecillinam, cefuroxime, and cefotaxime inhibited all strains in concentrations of 8  $\mu g$  or less per ml, and cefamandole inhibited all strains in concentrations of 32  $\mu g$  or less per ml. A previous study (2) indicated that cefoperazone was also active; minimal inhibitory concentrations were 0.13 to 8  $\mu$ g/ml. Among the aminoglycoside antibiotics, gentamicin was most active. All aminoglycosides except streptomycin inhibited 100% of strains in clinically achievable concentrations. Among the other antimicrobial agents studied, only tetracycline, chloramphenicol, and trimethoprim-sulfamethoxazole were consistently active.

Antimicrobial agent	Minimal inhibitory concn (µg/ml)		
	Range	50°c inhibition	90% inhibition
Penicillin G	64->128	>128	>128
Ampicillin	16->128	>128	>128
Carbenicillin	8->128	>128	>128
Ticarcillin	8->128	>128	>128
Azlocillin	8->128	32	64
Mezlocillin	4->128	8	8
Piperacillin	1->128	4	8
Mecillinam	0.25-4	1	2
Cephalothin	2->128	>128	>128
Cefazolin	4->128	128	>128
Cefamandole	0.25-32	2	8
Cefoxitin	0.5 - 128	8	32
Cefazaflur	0.5->128	64	>128
Cefuroxime	0.25-8	1	4
Cefotaxime	≤0.06-4	≤0.06	0.25
Moxalactam	≤0.06	≤0.06	≤0.06
Streptomycin	2->64	8	16
Kanamycin	0.5-8	2	8
Gentamicin	0.13-0.5	0.25	0.5
Tobramycin	0.13 - 2	1	2
Amikacin	0.5-4	2	4
Sisomicin	0.13-1	0.5	1
Netilmicin	0.13-1	0.5	1
Colistin	1->64	4	>64
Tetracycline	0.5 - 2	1	2
Minocycline	0.5-4	1	4
Doxycycline	0.5-4	1	2
Chloramphenicol	0.5-8	1	4
Erythromycin	8-64	16	32
Clindamycin	≥64	>64	>64
Vancomycin	≥64	>64	>64
TMP-SMZ <sup>a</sup>	2-32	4	8

 TABLE 1. In vitro susceptibilities of A. hydrophila

 to 32 antimicrobial agents

" TMP-SMZ, Total trimethoprim-sulfamethoxazole in a ratio of 1:19.

## LITERATURE CITED

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