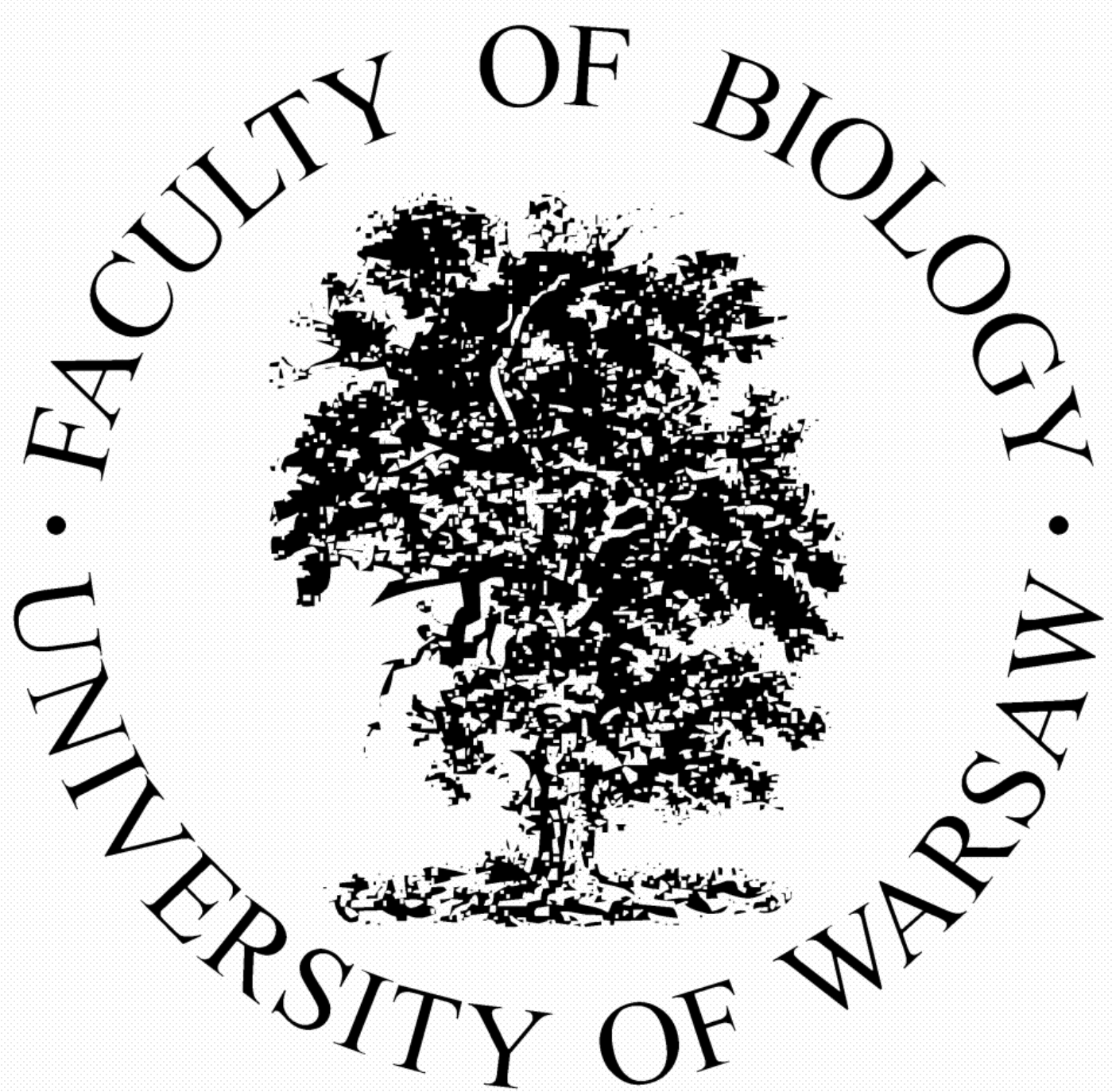


SUSCEPTIBILITY TESTING OF *MYCOBACTERIUM KANSASII* STRAINS ISOLATED FROM PATIENTS BETWEEN 2000 AND 2015 IN POLAND

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INTRODUCTION

Nontuberculous mycobacterial (NTM) infections including those caused by *Mycobacterium kansasii* appear to be increasing worldwide. In Poland, *M. kansasii* accounts for over a third of the total NTM isolations. Reports on drug susceptibility of this species are very scarce. The aim of this study was to determine drug susceptibility profiles of *M. kansasii* strains isolated from patients of a pulmonary clinic (Department of Internal Medicine, Pulmonology, and Allergology, Warsaw Medical University) in Warsaw, Poland.

MATERIALS AND METHODS

The study included 62 *M. kansasii* strains collected from as many patients of the Department of Internal Medicine, Pulmonology, and Allergology, Warsaw Medical University between 2000 and 2015. Patients (40 women and 22 men; age range: 21 to 89 years; median age: 64.5 ± 17.9 years) were classified as having or not having an infection, following the criteria of the American Thoracic Society (ATS). Susceptibility testing to 8 anti-TB drugs: rifampicin (RMP), amikacin (AMK), ethambutol (EMB), isoniazid (INH), streptomycin (STR), clarithromycin (CLM), kanamycin (KAN) and trimethoprim/ sulfamethoxazole (SXT), was performed by the E-test method, strictly according to the manufacturer's instructions (bioMérieux®). As a control, the *M. kansasii* ATCC12478 strain was used. The critical concentrations of tested drugs were based on Clinical and Laboratory Standards Institute guidelines, and in their absence – on literature data.

| Drug | Critical concentration [mg/L] | No of resistant strains (%) | No of susceptible strains (%) | Average MIC (value of MIC on the E-test strip) | MIC ₅₀ | MIC ₉₀ |
|------|-------------------------------|-----------------------------|-------------------------------|--|-------------------|-------------------|
| RMP | 1 | 0 (0%) | 62 (100%) | 0.011 ± 0.008 (0.012) | 0.008 | 0.023 |
| AMK | 32 | 0 (0%) | 62 (100%) | 2.12 ± 1.96 (2) | 1.5 | 4 |
| STR | 10 | 1 (2%) | 61 (98%) | 18.51 ± 129.8 (16) | 1.5 | 4 |
| EMB | 4 | 2 (3%) | 60 (97%) | 1.01 ± 1.78 (1) | 0.5 | 1.5 |
| CLM | 16 | 2 (3%) | 60 (97%) | 8.32 ± 45.6 (8) | 0.047 | 0.094 |
| KAN | 4 | 22 (35%) | 40 (65%) | 4.91 ± 5.73 (4) | 2 | 12 |
| INH | 4 | 53 (85%) | 9 (15%) | 218.84 ± 90.90 (192) | >256 | >256 |
| SXT | 2/38 | 62 (100%) | 0 (0%) | >32/608 | >32/608 | >32/608 |

RESULTS

Of the 62 patients under the study, 38 (61.3%) met ATS criteria for the definition of the *M. kansasii* disease, whereas 24 (38.7%) patients did not have the *M. kansasii* disease.

All strains tested exhibited full susceptibility to RMP (MIC<1 mg/L) and AMK (MIC<32 mg/L). The number of strains showing resistance to INH (MIC >256 mg/L), KAN (MIC≥4 mg/L), EMB (MIC≥8 mg/L), CLR (MICs>256 mg/L), STR (MIC>1024 mg/L) was 53, 22, 2, 2, and one, respectively. All strains were found resistant to SXT (MIC>32/608 mg/L).

Strains from patients who met the ATS case definition criteria for disease had their MIC₉₀s for RMP higher than strains from patients with no *M. kansasii* disease (0.023 vs 0.016 mg/L).

CONCLUSIONS

The results showed a high activity of tested drugs against clinical strains of *M. kansasii*, except for SXT, INH, and KAN.

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