TITLE: Antifungal activity of silver tungstate against dermatophytes and Candida spp.

AUTHORS: BIANCO, L. M¹; SIQUEIRA, J.P.Z.²; BRIZZOTTI, N. S.²; FERNANDES, J.M.F.²; LEMES, T.H.¹; ALMEIDA, B.G.¹; SILVA, C. A.; LONGO, E. ³; ASSIS, M. ³; ALMEIDA M. T. G.²

INSTITUTION: ¹UNIVERSIDADE ESTADUAL PAULISTA – UNESP. INSTITUTO DE BIOCIÊNCIAS, LETRAS E CIÊNCIAS EXATAS - CÂMPUS SÃO JOSÉ DO RIO PRETO, SP (RUA CRISTOVÃO COLOMBO, 2265, CEP 15054-000, SÃO JOSÉ DO RIO PRETO-SP, BRASIL).

²FACULDADE DE MEDICINA DE SÃO JOSÉ DO RIO PRETO (AV. BRIGADEIRO FARIA LIMA, 5416, CEP 15090-000, SÃO JOSÉ DO RIO- SP, BRASIL).

³UNIVERSIDADE FEDERAL DE SÃO CARLOS – UFSCar. CENTRO DE CIÊNCIAS EXATAS E DE TECNOLOGIA (JARDIM GUANABARA, CEP 13565905, SÃO CARLOS-SP, BRASIL).

ABSTRACT:

Fungal infections represent a global public health problem. The incidence and the etiology of skin disease is variable, including dermatophytes, non-dermatophytes, or yeast fungi. The objective of this study was to evaluate the antifungal activity of a nanostructured molecule of silver tungstate (α -Aq₂WO₄) against fungi responsible for superficial mycoses. The α -Aq₂WO₄ molecule was synthesized by the simple coprecipitation method in 1:1 v/v of water and ethanol. The material was irradiated with electrons using a high resolution scanning electron microscope (SEM) model Supra 35-VP (Carl Zeiss, Germany), with an acceleration voltage of 15 kV. The antifungal susceptibility tests were performed considering the protocols of the Clinical and Laboratory Standards Institute M38-A2, for filamentous fungi, and M27-A3, for yeasts. Microplates were prepared with α -Ag₂WO₄ final concentrations of 0.03 – 1000 µg/ml. The fungi tested included reference strains (Trichophyton rubrum CBS 118892, Candida albicans ATCC 90028, Candida glabrata ATCC 2001, Candida parapsilosis ATCC 22019, and Candida krusei ATCC 40147) and clinical isolates (Trichophyton rubrum, Trichophyton mentagrophytes, Candida albicans, Candida glabrata, Candida parapsilosis, and Candida krusei). Minimum inhibitory concentration (MIC) was defines as the lowest drug concentration capable of prevent any discernible growth. Regarding the results, MIC values observed for dermatophytes were 0.48-1.95 µg/ml (0.48, 0.95, and 1.95 µg/ml for T. rubrum CBS 118892, T. rubrum clinic, and T. mentagrophytes clinic, respectively). For Candida spp., MIC values exhibited a wider range (0.95–7.81 μg/ml). Greater activity of α-Ag₂WO₄ was observed for C. krusei (0.48 μg/ml for both strains), while the lowest activity was observed for C. glabrata (1.95 and 7.81 µg/ml, for ATCC 2001 and clinical isolate, respectively) and C. parapsilosis (3.9 and 1.95 µg/ml, for ATCC 22019 and clinical isolate, respectively). MIC values for Candida albicans were 1.95 µg/ml (ATCC 90028) and 0.95 µg/ml (clinic). These results open possibilities for new therapeutic approaches using silver tungstate for the control of fungal infections, which may have a positive impact on clinical dermatology and infectious diseases management.

KEYWORDS: silver tungstate, fungal infections, antifungal activity.