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**Record 1 of 1****Title:** Efficacy of some synthesized thiazoles against dermatophytes**Author(s):** Ouf, SA (Ouf, S. A.); Abu Taleb, AM (Abu Taleb, A. M.); Tharwat, NA (Tharwat, N. A.); Geweely, NS (Geweely, N. S.)**Source:** JOURNAL DE MYCOLOGIE MEDICALE **Volume:** 23 **Issue:** 4 **Pages:** 230-236 **DOI:** 10.1016/j.mycmed.2013.07.056 **Published:** DEC 2013**Times Cited in Web of Science Core Collection:** 2**Total Times Cited:** 2**Usage Count (Last 180 days):** 0**Usage Count (Since 2013):** 4**Cited Reference Count:** 32

**Abstract:** Twelve thiazoles and their fused derivatives were tested for their antimicrobial activity against *Trichophyton rubrum*, *T terrestris*, *Epidermophyton floccosum*, and *Microsporum gypseum*. Most of the synthesized compounds were inhibitory to the tested fungi. The most effective compound was 5-(4-ethoxybenzylidene)-4,5-dihydro-4-oxothiazol-2-yl)-N,3-diphenylbut-2-namide (3c) followed by 2-(4-oxo-4,5-dihydrothiazol-2-yl)-3-phenyl-but-2-enic acid-(3-cyano-4,5,6,7-tetrahydrobenzo[b]thiophen-2-yl)-amide (2b). These compounds were more efficacious than terbinafine, the reference drug. The tested compounds caused variable reduction in the activity of keratinase of the dermatophytes, depending on the azole derivative and the test fungus. Thiazole derivatives (2b) and (3c) exhibited the highest efficacy in decreasing ergosterol biosynthesis of the tested dermatophytes. The treatment of guinea pigs with compound (3c) induced complete curing in the case of all the test dermatophytes 30 days post-treatment. The percent curing for compounds (3c) and (2b) was better than the reference drug. (C) 2013 Published by Elsevier Masson SAS.

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