



Antifungal Activity of Resveratrol Derivatives against *Candida* Species

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Auteur	Houillé, Benjamin [1], Papon, Nicolas [2], Boudesocque, Leslie [3], Bourdeaud, Eric [4], Besseau, Sébastien [5], Courdavault, Vincent [6], Enguehard-Gueiffier, Cécile [7], Delanoue, Guillaume [8], Guérin, Laurence [9], Bouchara, Jean-Philippe [10], Clastre, Marc [11], Giglioli-Guivarc'h, Nathalie [12], Guillard, Jérôme [13], Lanoue, Arnaud [14]
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Résumé en anglais	<p><i>trans</i>-Resveratrol (1a) is a phytoalexin produced by plants in response to infections by pathogens. Its potential activity against clinically relevant opportunistic fungal pathogens has previously been poorly investigated. Evaluated herein are the candidacidal activities of oligomers (2a, 3-5) of 1a purified from <i>Vitis vinifera</i> grape canes and several analogues (1b-1j) of 1a obtained through semisynthesis using methylation and acetylation. Moreover, <i>trans</i>-ϵ-viniferin (2a), a dimer of 1a, was also subjected to methylation (2b) and acetylation (2c) under nonselective conditions. Neither the natural oligomers of 1a (2a, 3-5) nor the derivatives of 2a were active against <i>Candida albicans</i> SC5314. However, the dimethoxy resveratrol derivatives 1d and 1e exhibited antifungal activity against <i>C. albicans</i> with minimum inhibitory concentration (MIC) values of 29–37 $\mu\text{g/mL}$ and against 11 other <i>Candida</i> species. Compound 1e inhibited the yeast-to-hyphae morphogenetic transition of <i>C. albicans</i> at 14 $\mu\text{g/mL}$.</p>
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