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### Patent Search

Invention Title	SUBSTITUTED PHENOXY PHENYLETHYLIDENE DERIVATIVE AS ANTIMICROBIAL AGENT
Publication Number	05/2021
Publication Date	29/01/2021
Publication Type	INA
Application Number	202111002121
Application Filing Date	16/01/2021
Priority Number	
Priority Country	
Priority Date	
Field Of Invention	MICRO BIOLOGY
Classification (IPC)	C12R 1/125 A61K 31/18 A61K 31/5377

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**Abstract:**

Substituted phenoxy phenylethylidene derivatives of Formula I or its pharmaceutically acceptable salt (HG-1 to HG-4) were obtained and screened for antimicrobial activity. Structures of these compounds were established on the basis of M.P, TLC, IR, <sup>1</sup>H-NMR spectral analysis. The target compounds were found to be tested against Bacillus Su (Gram positive), E.Coli (Gram negative) bacteria and Candida albicans, when compared with the reference drugs ampicillin and clotrimazole. Among the compounds tested showed maximum zone of inhibition 20 mm against B.subtilis.

**Complete Specification**

The present invention is in technical field related to synthesis of organic molecules having functional group that contain a carbon-nitrogen double bond with the nitrogen atom connected to aryl group used as an antimicrobial agent. More particularly, the present invention relates to substituted phenoxy phenylethylidene derivatives used to kill or inhibits the growth of microbes such as bacteria, fungi, or viruses commonly used as antimicrobial agent. Further the present invention also relates to process for producing substituted phenoxy phenylethylidene derivatives, its acid addition salt thereof, and to the pharmaceutical composition comprising such compounds and a pharmaceutically acceptable salts thereof.

**BACKGROUND OF THE INVENTION**

Bacteria threatening life and affecting the health status of people living in the 15 world by both Gram-negative and Gram-positive bacteria. The uptake or the ingestion of water and/or food contaminated with bacteria such as Salmonella typhi, Escherichia coli including Clostridium, Pseudomonas and Bacillus species

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Page last updated on: 26/06/2019