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## Review article

# Recent developments in biological activities of indanones



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#### ABSTRACT

Indanone is one of the privileged structures in medicinal chemistry and it's commonly associated with various pharmacologically active compounds. The indanone moiety is found in several natural compounds and also, it can be used as intermediate in the synthesis of many different types of medicinally important molecules. Among the medicinally important indanones, the most significant drug probably is donepezil (IV), an acetylcholinesterase (AChE) inhibitor, which has been approved by the US Food and Drug Administration for the treatment of Alzheimer's disease (AD). Along with donepezil, the indanone moiety can be seen in a number of other pre-clinical and clinical candidates which belong to different categories with diverse therapeutic activities. In summary, the present review article encompasses the recent biological applications such as antialzheimer, anticancer, antimicrobial and antiviral activity of various indanone derivatives.

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## 1. Introduction

Indanone is one of the privileged structures in medicinal chemistry and is commonly associated with various

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pharmacologically active compounds. Indanones have demonstrated a broad spectrum of biological activity. Moreover, they are very useful synthons for the synthesis of various carbocyclic and heterocyclic molecules as synthetic intermediates for several drugs and natural products [1–8]. Their important biological activities include antiinflammatory [9–13], analgesic [14], antimicrobial [15,16], anticholinergic [17,18], dopaminergic [19], anticancer [20] and antimalarial [21] activities. The indanone moiety is present in various bioactive natural products. The important natural products

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Abbreviations		S. aureus Staphylococcus aureus	
		E. coli Escherichia coli	
AD	Alzheimer's disease	K. pneumonia Klebsiella pneumonia	
AChE	Acetylcholinesterase	A. baumannii Acinetobacter baumannii	
BChE	Butylcholinesterase	P.aeruginosa Pseudomonas aeruginosa	
BACE1	β-Site amyloid precursor protein cleaving enzyme 1	C. albicans Candida albicans	
SAR	Structure-activity relationship	C. neoformans Cryptococcus neoformans	
HDACs	Histone deacetylases	A. clavatus Aspergillus clavatus	
ACh	Acetylcholine	B. subtilis Bacillus subtilis	
IC <sub>50</sub>	Half maximal inhibitory concentration	M. luteusgenes Micrococcus luteusgenes	
MAO-B	Monoamine oxidases B	L. Monocytogenes Listeria monocytogenes	
BBB	Blood brain barrier	A. Niger Aspergillus niger	
μΜ	Micromolar	T. mentagrophites Trichophyton mentagrophites	
nM	Nanomolar	RBV Ribavirin	
GI <sub>50</sub>	The concentration of drug to cause 50% reduction in proliferation		

such as 1-methoxy-6-methyl-3-oxo-2,3-dihydro-1*H*-indene-4carbaldehyde (I) and Pterosin B (II) were isolated from marine cyanobacterium and Pteris ensiformisburm (Fig. 1) [22,23]. Another important natural product having the indanone ring system is a potential anti-viral polyphenolic derivative Paucifloral F (III) (Fig. 1) and its isomer isopaucifloral F is a potential anti-osteoporosis agent [24,25]. Because of the biological importance of the indanone core, past several years researchers have produced library of pharmacologically active indanones. Very recently, our laboratory reported the novel indonone analogs as antiviral and antimicrobial agents [26,27]. Among them the most significant drug that bears an indanone moiety is probably donepezil (IV), an acetylcholinesterase (AChE) inhibitor, which has been approved by the US Food and Drug Administration for the treatment of Alzheimer's disease (AD). Along with donepezil, the indanone moiety can be noticed in a number of other pharmacologically active molecules that belong to different categories with diverse therapeutic activities [28–32] (Fig. 1).

H<sub>3</sub>CO 
$$H_3$$
CO  $H_3$ C

Fig. 2. 1-Indanone thiosemicarbazone derivatives.

Fig. 1. Pharmacologically active important natural and unnatural products bearing indanone moiety.

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