

FAR-PO-29 Novel Bifunctional Compounds Targeting Nicotine and Dopamine Receptor Subtypes: Synthesis and Pharmacological Investigation

Clelia Dallanoce,^a Carlo Matera,^a Luca Pucci,^b Cecilia Gotti,^b Marco De Amici,^a Carlo De Micheli^a

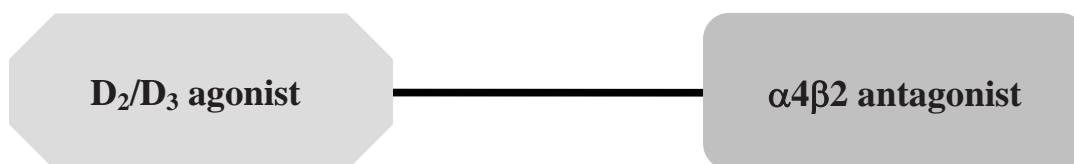
^aDipartimento di Scienze Farmaceutiche “Pietro Pratesi” dell’Università degli Studi di Milano, Via L. Mangiagalli 25, 20133 Milan, Italy

^bIstituto di Neuroscienze, Farmacologia Cellulare e Molecolare - CNR and Dipartimento Farmacologia, Chemioterapia e Tossicologia Medica dell’Università degli Studi di Milano, Via Vanvitelli 32, 20129 Milan, Italy

clelia.dallanoce@unimi.it

Future therapies for diseases associated with altered dopaminergic signaling, including Parkinson’s disease, schizophrenia and drug addiction or drug dependence, may be substantially built on the existence of intramembrane receptor-receptor interactions within receptor mosaics where it is believed that the D₂ receptor may operate as the “hub receptor” [1]. In particular, it has been proposed that striatal dopaminergic neurotransmission could be under the control of receptor heteromers containing D₂ autoreceptors and non-α7 nicotinic acetylcholine heteroreceptors [2].

In an attempt to investigate the biochemical and functional interactions between dopaminergic autoreceptors and nAChRs containing the β2 subunit, we designed and prepared a group of potential bifunctional derivatives incorporating a D₂/D₃ agonist moiety and a nicotinic α4β2 antagonist fragment, linked by polymethylene spacers of different length.



The new compounds have been biologically characterized for their affinity/specificity/functional profile at the target nACh and D₂ receptor subtypes. The synthesis of the designed derivatives and the results of their pharmacological investigation will be presented and discussed.

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- [2] D.Quarta, F.Ciruela, K.Patkar, J.Borycz, M.Solinas, C.Lluis, R.Franco, R.A.Wise, S.R.Goldberg, B.T.Hope, A.Woods, S.Ferré, *Neuropsychopharmacol.*, 32, **2007**, 35-42.