

P2Y receptors in GtoPdb v.2021.3

Maria-Pia Abbracchio¹, Jean-Marie Boeynaems², José L. Boyer³, Geoffrey Burnstock⁴, Stefania Ceruti¹, Marta Fumagalli¹, Christian Gachet⁵, Rebecca Hills⁶, Robert G. Humphries⁷, Kazu Inoue⁸, Kenneth A. Jacobson⁹, Charles Kennedy¹⁰, Brian F. King¹¹, Davide Lecca¹, Christa E. Müller¹², Maria Teresa Miras-Portugal¹³, Vera Ralevic¹⁴ and Gary A. Weisman¹⁵

1. University of Milan, Italy
2. Université Libre de Bruxelles, Belgium
3. University of North Carolina, USA
4. Autonomic Neuroscience Centre, University College London, UK
5. INSERM, France
6. University of Edinburgh, UK
7. AstraZeneca, UK
8. University of Kyushu, Japan
9. National Institutes of Health, USA
10. University of Strathclyde, UK
11. University College London, UK
12. Universität Bonn, Germany
13. Universidad Complutense de Madrid, Spain
14. University of Nottingham, UK
15. University of Missouri, USA

Abstract

P2Y receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on P2Y Receptors [3, 5, 192]**) are activated by the endogenous ligands **ATP**, **ADP**, **uridine triphosphate**, **uridine diphosphate** and **UDP-glucose**. The relationship of many of the cloned receptors to endogenously expressed receptors is not yet established and so it might be appropriate to use wording such as '**uridine triphosphate**-preferring (or **ATP**-, *etc.*) P2Y receptor' or '**P2Y₁**-like', *etc.*, until further, as yet undefined, corroborative criteria can be applied [47, 110, 190, 383, 396]. Clinically used drugs acting on these receptors include the dinucleoside polyphosphate **diquafosol**, agonist of the P2Y₂ receptor subtype, approved in Japan for the management of dry eye disease [241], and the P2Y₁₂ receptor antagonists **prasugrel**, **ticagrelor** and **cangrelor**, all approved as antiplatelet drugs [53, 323].

Contents

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