

## Prostanoid receptors in GtoPdb v.2021.2

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### Abstract

Prostanoid receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Prostanoid Receptors [694]**) are activated by the endogenous ligands prostaglandins [PGD<sub>2</sub>](#), [PGE<sub>1</sub>](#), [PGE<sub>2</sub>](#), [PGF<sub>2α</sub>](#), [PGH<sub>2</sub>](#), prostacyclin [[PGI<sub>2</sub>](#)] and [thromboxane A<sub>2</sub>](#). Differences and similarities between human and rodent prostanoid receptor orthologues, and their specific roles in pathophysiologic conditions are reviewed in [448]. Measurement of the potency of [PGI<sub>2</sub>](#) and [thromboxane A<sub>2</sub>](#) is hampered by their instability in physiological salt solution; they are often replaced by [cicaprost](#) and [U46619](#), respectively, in receptor characterization studies.

### Contents

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### Database links

#### Prostanoid receptors

<https://www.guidetopharmacology.org/GRAC/FamilyDisplayForward?familyId=58>

#### Introduction to Prostanoid receptors

<https://www.guidetopharmacology.org/GRAC/FamilyIntroductionForward?familyId=58>

#### Receptors

##### DP<sub>1</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=338>

##### DP<sub>2</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=339>

##### EP<sub>1</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=340>  
EP<sub>2</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=341>  
EP<sub>3</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=342>  
EP<sub>4</sub> receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=343>  
FP receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=344>  
IP receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=345>  
TP receptor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=346>

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