

## Prostanoid receptors in GtoPdb v.2025.3

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### Abstract

Prostanoid receptors (**nomenclature as agreed by the NC-IUPHAR Subcommittee on Prostanoid Receptors [709]**) 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 [457]. 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

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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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