

## Prokineticin receptors (version 2019.4) in the IUPHAR/BPS Guide to Pharmacology Database

Rebecca Hills<sup>1</sup>, Philippe Rondard<sup>2</sup>, Oualid Sbai<sup>2</sup> and Qun-Yong Zhou<sup>3</sup>

1. University of Edinburgh, UK
2. Université de Montpellier, France
3. University of California Irvine, USA

### Abstract

Prokineticin receptors, PKR<sub>1</sub> and PKR<sub>2</sub> (**provisional nomenclature as recommended by NC-IUPHAR [23]**) respond to the cysteine-rich 81-86 amino-acid peptides prokineticin-1 (also known as endocrine gland-derived vascular endothelial growth factor, mambakine) and prokineticin-2 (protein Bv8 homologue). An orthologue of PROK1 from black mamba (*Dendroaspis polylepis*) venom, mamba intestinal toxin 1 (MIT1, [65]) is a potent, non-selective agonist at prokineticin receptors [41], while Bv8, an orthologue of PROK2 from amphibians (*Bombina sp.*, [44]), is equipotent at recombinant PKR<sub>1</sub> and PKR<sub>2</sub> [48], and has high potency in macrophage chemotaxis assays, which are lost in PKR<sub>1</sub>-null mice.

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## Receptors

**PKR<sub>1</sub>**

<http://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=335>

**PKR<sub>2</sub>**

<http://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=336>

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