



Product Informatiion Sheet

Polyclonal Anti- Dopamine receptor D₁, DRD1 (Magnetic Bead Conjugate)

Catalogue No. PA1231-M Immunogen

Lot No. 09E01 A synthetic peptide corresponding to a sequence at the C-terminal of human DRD1,

identical to the related rat and mouse sequence.

Ig type: rabbit IgG1 Purification

Immunogen affinity purified

Size: 100µg/Vial

Contents

Specificity Each vial contains 1mg/ml Magnetic Bead in PBS, pH 7.2, 0.05mg NaN₃.

Human, rat, mouse.

No cross reactivity with other Storage

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proteins. Store at 4°C for frequent use.

Immunoprecipitation(IP) This Antagene antibody is immobilized by the covalent reaction of

hydrazinonicotinamide-modified antibody with formylbenzamide-modified magnetic beads.

It is useful for immunoprecipitation

BACKGROUND

Dopamine receptor D₁, also known as DRD1, is a human gene. It is the most highly expressed DA receptor subtype among the DA receptor family. Receptors for dopamine have been classified into two functional types, D1 and D2. They belong to the family of receptors acting through G (or guanine nucleotide-binding) proteins. D2 receptors inhibit adenylyl cyclase, but D1 receptors stimulate adenylyl cyclase and activate cyclic AMP-dependent protein kinases. Dopamine D1 and D2 receptors are targets of drug therapy in many psychomotor disorders, including Parkinson's disease and schizophrenia, and may also have a role in drug addiction and alcoholism. D1 receptors regulate neuron growth and differentiation, influence behaviour and modify dopamine D2 receptor-mediated events. And the presence of a D1 receptor gene restriction fragment length polymorphism will be helpful for future disease linkage studies. DRD1 also regulates the neurochemical architecture of the striatum and is critical for the normal expression of motor activity.

REFERENCE

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- 3. Xu, M.; Moratalla, R.; Gold, L. H.; Hiroi, N.; Koob, G. F.; Graybiel, A. M.; Tonegawa, S.: Dopamine D1 receptor mutant mice are deficient in striatal expression of dynorphin and in dopamine-mediated behavioral responses. *Cell* 79: 729-742, 1994.

