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The dopamine D1 receptor (D1R) is essential for neurotransmission in various brain pathways where it modulates key functions including voluntary movement, memory, attention and reward.

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The D-1 receptors are linked to dopamine-sensitive adenylate cyclase, while the D 2 receptors are those that are not linked to this enzyme. It appears necessary to modify this terminology because there is no biochemically practical method for determining the proportion of linked and nonlinked dopamine receptors and there is no relation between the behavioral potencies of dopamine agonists or antagonists and their potencies on this enzyme.

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presence of at least two types of central dopamine da receptors the d 1 da receptors are positively linked to the da sensitive adenylate cyclase system whereas the d 2 da receptors are probably negatively linked

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the d 2 da receptors have been generally recognized as mediating the action of

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D1 or D1A. The D1 receptor is the most abundant dopamine receptor in the brain. This receptor is linked to stimulatory G-proteins that activate adenylate cyclase. The D1 receptors are found in high concentration in the substantia nigra pars reticulata, caudate, putamen, nucleus accumbens, olfactory tubercle, and frontal and temporal cortex.

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Dopamine Receptors in the Human Brain | Psychiatric Times

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Five subtypes of mammalian dopamine receptors have been identified that are divided into D1-like (D1 and D5) or D2-like (D2, D3, and D4) groups. The D1-like receptors couple primarily to the G<sub>s</sub> family of G proteins (G<sub>s</sub> and G<sub>olf</sub>), whereas the D2-like receptors couple primarily to the G<sub>i/o</sub> family.

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The development of a selective D1 dopamine (DA) receptor antagonist SCH 23390 stimulated a number of studies on the functions mediated by central DA receptor subtypes. It was generally assumed that the central D1 DA receptor is a molecular entity whose function awaits further discovery.

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Central D1 Dopamine Receptors | M. J. Goldstein | Springer

Antipsychotic response has been demonstrated to be associated with blockade of dopamine D2 receptors, and antidepressant response has now been linked to blockade of serotonin transporter receptors.

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