Sumanirole, a Highly Dopamine D2-Selective Receptor Agonist: In Vitro and in Vivo Pharmacological Characterization and Efficacy in Animal Models of Parkinson’s Disease

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ABSTRACT

The purpose of this study is to demonstrate that sumanirole is a novel dopamine receptor agonist with high in vitro and in vivo selectivity for the D2 receptor subtype. Sumanirole, (R)-5,6-dihydro-5-(methylamino)-4H-imidazo[4,5,1-ij]quinolin-2(1H)-one (Z)-2-butenedioate (1:1), is unique; it has greater than 200-fold selectivity for the D2 receptor subtype versus the other dopamine receptor subtypes in radioligand binding assays. In cell-based assays, sumanirole is a fully efficacious agonist, with EC50 values between 17 and 75 nM. In animals, sumanirole elicits many physiological responses attributed to D2-like receptor function. In rats, sumanirole is a full agonist for elevation of striatal acetylcholine levels (ED50 = 12.1 μmol/kg i.p.). Sumanirole s.c. dose dependently decreased plasma prolactin levels and depressed dopamine neuron firing rates in the substantia nigra pars compacta with an ED50 of 2.3 μmol/kg i.v. This high selectivity for D2 receptors translates into excellent locomotor stimulant activity in animal models of Parkinson’s disease. In reserpinated, α-methyl-para-tyrosine-treated rats, sumanirole caused a significant and sustained increase in horizontal activity at doses ≥12.5 μmol/kg s.c. In unilateral 6-hydroxydopamine-lesioned rats, sumanirole caused profound, sustained rotational behavior and was substantially more efficacious than any other agonist tested. Sumanirole-stimulated rotational behavior was blocked by the dopamine receptor antagonist haloperidol. Sumanirole dose dependently improved disability scores and locomotor activities of two of three 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine-lesioned monkeys. In summary, sumanirole is the first published selective D2 receptor agonist. The compound has activity in animal models of dopamine hypofunction and has a high level of efficacy in animal models of Parkinson’s disease.

Parkinson’s disease results from degeneration of the dopaminergic cells in the substantia nigra and is characterized by bradykinesia, tremors, and muscular rigidity. Several dopamine receptor agonists have been used as antiparkinsonian therapy, including bromocriptine, cabergoline, pergolide, pramipexole, and ropinirole (Hagan et al., 1997). Pramipexole, cabergoline, and ropinirole are effective as monotherapy in early stage Parkinson’s disease and their use significantly delays the initiation of L-dopa therapy (Inzelberg et al., 2003; Bracco et al., 2004). Dopamine agonists are also used as adjuncts to L-dopa. The combined treatment enables lower doses of L-dopa, which decreases the incidence of response fluctuations and dyskinesias. However, currently available dopaminergic therapy, including agonists and L-dopa therapy, can elicit side effects, including psychiatric complications and somnolence (Lieberman, 1995). Thus, there is a need for new compounds with fewer side effects (Hobson et al., 2002; Etminan et al., 2003).

There are at least five subtypes of dopamine receptors, grouped into two subfamilies, D1-like and D2-like, based on pharmacological and amino acid sequence similarities (Civelli et al., 1993). The dopamine agonists bind at D2-like dopamine receptors (Montastruc et al., 1993). The D2-like subfamily includes D2, D3, and D4 receptors. Based on mRNA distributions, it has been suggested that D2 receptors are more abundant in basal ganglia than in mesolimbic/mesocortical areas, whereas the D3 and D4 receptors are relatively more abundant in the limbic/cortical areas than in the striatum.
atum (Bouthenet et al., 1991; Van Tol et al., 1991). Because of the neuroanatomical distribution of these receptors, it has been suggested that D₃ and D₄ receptors may contribute to the psychiatric disturbances that accompany dopamine agonist and L-dopa therapeutics. In fact, clozapine, a dopamine receptor blocker with high affinity for the D₃ subtype, has been used to ameliorate the psychoses accompanying antiparkinsonian treatment (Friedman, 1995) suggesting that this may be a D₃-mediated side effect. Additionally, dopamine agonists such as pramipexole and ropinirole demonstrate daytime hypomellosome in Parkinson’s disease patients, along with other sleep-related problems (Razmy et al., 2004). The receptor subtype involved in this response has not been delineated. Eliminating D₃- and D₄-activating proper-

Materials and Methods

The chemical structure of sumanirole is shown in Fig. 1, and the synthesis was described previously (Heier et al., 1997). The drug has been tested as either of two salt forms: the monohydrochloride salt with a molecular weight of 239.71 g/mol and the maleate salt with a molecular weight of 319.32 g/mol.

Cell Growth. Chinese hamster ovary (CHO) cells expressing human D₃, D₄, or D₅ receptors or transfected with the 3C vector alone were grown in minimal essential medium [α modification (α-MEM)] supplemented with 10% fetal calf serum, 2 mM glutamine, 10 U/ml penicillin, and 100 µg/ml streptomycin. Media contained 1 mg/ml G418 for growth of CHO-3C (mock-transfected), D2-L6, D2-S8, and D3-3 cells. HEK293 cells expressing human D₂ receptors were grown in Dulbecco’s modified Eagle’s medium high glucose supplemented with 10% fetal bovine serum, 4 mM glutamine, 100 U/ml penicillin G, 100 µg/ml streptomycin, and 100 U/ml hygromycin B. D₂-L6 cells express 3.5 pmol/mg human D₂α receptors measured with [³H]spiroperidol, and 1.0 pmol/mg protein measured with [³H]U-86170, an agonist ligand. D₂-S8 cells express 3.0 pmol/mg human D₂α receptors measured with [³H]spiroperidol. D₃-3 cells express D₃ receptors at 2.5 pmol/mg protein measured with [³H]spiroperidol (Chio et al., 1994b). HEK293 cells express D₄ receptors at a density of 0.74 pmol/mg protein measured with [³H]spiroperidol (Chio et al., 1994a).

Receptor Binding Methods. Radioligands used were [³H]SCH23390 (D₁-dopamine, 96 Ci/mmol, 1 nM), [³H]U-86170 (Lahti et al., 1991) (D₂-dopamine, 62 Ci/mmol, 2 nM), and [³H]spiperone (D₃ and D₄-dopamine, 96 Ci/mmol, 0.2 nM). Rat striatal membranes were the source of D₁ receptors. CHO cells and HEK293 cells expressing D₃, D₄, and D₅ receptors were rinsed with ice-cold Cs²⁺/Mg²⁺-free phosphate-buffered saline and harvested in the same buffer. Cells were pelleted (500g, 5 min), resuspended in 25 mM Tris, 5 mM EDTA, and 5 mM EGTA, pH 7.5, and frozen in liquid nitrogen. After thawing, the cells were homogenized and centrifuged at 1000g to remove nuclei and broken cells. The supernatant was centrifuged at 47,000g; the membrane pellet was washed once with Tris, EGTA, EDTA, resuspended in 20 mM HEPES, pH 7.4, 150 mM NaCl, 10 mM MgCl₂, and 1 mM EDTA, and frozen in liquid nitrogen. Membrane aliquots were stored at −70°C. For the receptor binding assays, the membranes were thawed and diluted into 20 mM HEPES, pH 7.4, 150 mM NaCl, 10 mM MgCl₂, or 1 mM EDTA. IC₅₀ values were determined by fitting the data to a one-site model by nonlinear least-squares minimization. Kᵢ values were calculated with the Cheng-Prusoff equation (Cheng and Prusoff, 1973).

CAMP Measurements. cAMP accumulation was measured in intact CHO cells plated at a density of 15,000 cells/well in a 24-well plate 48 h before the experiment. The cells were incubated in serum-free medium 1 h before the experiment. Fresh medium (0.5 ml) containing 100 µM forskolin, 100 µM isobutyl methylxanthine, and varying concentrations of drugs were added to each well, and cAMP was allowed to accumulate for 15 min at room temperature. The reactions were terminated by the removal of the medium and the addition of 100 µl of cold 7.5% trichloroacetic acid. The samples were diluted by the addition of 1.0 ml of 50 mM sodium acetate, pH 6.2, and aliquots were assayed by radioimmunoassay using the Biomed-

cAMP radiommunoassay kit.

Mitogenesis Assays. CHO cells were seeded into 96-well plates at a density of 5000 cells/well and were grown at 37°C in a α-MEM, with 10% fetal calf serum for 48 h. The wells were rinsed three times with serum-free α-MEM. Ninety microliters of fresh α-MEM were added along with 10 µl of drug (diluted in sterile water and filtered through 0.2-µm filters) or sterile water alone. Eight wells of every plate received 100 µl of α-MEM with 10% fetal calf serum. After culture for 16 to 17 h, [³H]thymidine (1 µCi/well) was added for 2 h. The cells were trypsinized and harvested onto filter mats with a Skatron cell harvester (Molecular Devices, Sunnyvale, CA). The filters were counted in a Betaplate counter (PerkinElmer Life and Analytical Sciences, Boston, MA).

Measurement of [³H]Arachidonic Acid Release. Cells were plated in 24-well plates at a density of 15,000 cells/well 48 h before use. Cells were labeled by incubation with [³H]arachidonic acid (210 Ci/mm, 0.4 µCi/ml; Amersham Biosciences Inc., Piscataway, NJ) in α-MEM (1 ml) supplemented with 10 mM HEPES, pH 7.5, and 0.5% fatty acid-free bovine serum albumin for 2 h at 37°C. The cells were then washed twice with 1 ml of the same buffer. Drugs were added in 1 ml of the same buffer, and the cells were incubated at 37°C for 30 min. Samples (0.5 ml) from each well were counted by liquid scintillation spectroscopy.

In Vivo: General Methods, Rat. Unless otherwise stated, the following generalizations apply to all in vivo rat testing. Animals were male Sprague-Dawley rats from Harlan (Indianapolis, IN) and were maintained in a temperature-controlled environment at 22 ± 1°C, with a 12:12-h light/dark cycle (lights on from 6:00 AM to 6:00 PM). They were group-housed and received food (Rodent Laboratory Chow #5001; Purina Mills Inc., Richmond, IN) and water ad libitum. All rodent testing was in compliance with the Animal Welfare Act Regulations, 9CFR Parts 1, 2, and 3, and also with the Guide for the Care and Use of Laboratory Animals (National Academy Press, National Academy of Sciences, 1996).

Electrophysiology. Rats weighing between 280 and 330 g were anesthetized with chloral hydrate (400 mg/kg i.p.). The femoral artery and vein were catheterized for monitoring blood pressure and

Fig. 1. Chemical structure of sumanirole.
administration of drugs, respectively. Glass microelectrodes filled with pontamine sky blue in 2 M NaCl used for extracellular recordings were lowered through a small hole burred through the calvarium by means of a hydraulic microdrive. Stereotaxic coordinates for placement of recording electrodes were AP, −4.8 to −5.0; L, +2.0; and V, −6.8 to −7.8 mm, relative to bregma (Paxinos and Watson, 1986). Substantia nigra pars compacta (SNPC) neurons were identified by waveform and firing patterns (Bunney et al., 1973). Histological localization of iontophoresed pontamine sky blue dye spots verified electrode locations. Drug solutions were made in distilled water with equimolar citric acid added as needed. Drug effects were measured as changes in firing rates monitored by an integrated rate meter.

**Plasma Prolactin.** Male Long Evans rats weighing 200 to 225 g (Harlan) were used in this assay. Sumanireole was dissolved in saline and administered s.c. in a volume of 1 ml/kg at various times. Thirty minutes after drug or vehicle administration, animals were killed by decapitation, and trunk blood was collected and plasma was stored. Plasma prolactin was measured by a double-antibody radioimmunoassay using the reagents and procedures of the National Institute of Diabetes and Digestive and Kidney Diseases assay kit (kindly supplied by Drs. A. F. Parlow and S. Raiti, National Institute of Diabetes and Digestive and Kidney Diseases National Hormone and Pituitary Program, Torrance, CA). National Institute of Diabetes and Digestive and Kidney Diseases rat prolactin (RP-3) was used as the standard. Using a 100-µl aliquot of plasma, the lower limit of sensitivity for prolactin was 0.1 ng/ml. The intra-assay coefficient of variation is usually about 8.6%.

**Striatal Acetylcholine Concentration.** Rats (130–150 g; Charles River Laboratories, Inc., Wilmington, MA) were used in this study. Solutions of sumanirole and ropinirole were prepared in 0.25% carboxymethylcellulose. Drug treatments were given i.p., and control rats received an equal volume of vehicle (2 ml/kg). Thirty minutes after treatment, animals were killed by decapitaiton; the brain was quickly removed from the skull and placed in ice-cold 0.32 M sucrose. Bilateral striata were dissected and homogenized in 0.05 N perchloric acid containing ethylhomocholine as an internal standard. Acetylcholine concentration was determined by high-pressure liquid chromatography. Each point represents five to six animals, and statistical analysis was done by a one-way ANOVA, followed by Student’s t test.

**Locomotor Activity Measurements in Reserpinized Rats.** Rats pretreated with reserpine and the dopamine synthesis inhibitor α-methyl-para-tyrosine (AMPT) are akinetic and cataleptic. These effects can be reversed by l-dopa (Carlsson et al., 1957). Reserpine/AMPT-treated rats have been used as a model of dopamine deplet...ho of dopamine depletion to mimic parkinsonian conditions. Rats weighing 200 to 250 g were used in this assay. Reserpine and haloperidol were purchased from Sigma/RBI (Natick, MA). Sumanireole was dissolved in physiological saline (0.9% NaCl), whereas haloperidol was dissolved in a few drops of glacial acetic acid and further diluted with 5.5% glucose solution. All compounds were administered s.c. in a volume of 5 ml/kg. Animals were pretreated with reserpine (5 mg/kg s.c., 18 h prior) and AMPT (100 mg/kg s.c., 1 h prior) before the experiment. The animals were injected with sumanirole or saline, and locomotor activity was measured using animal activity monitors (Digiscan model RXYM TAO; Omnitrace Inc., Columbus, OH). Data are presented as horizontal counts expressed as percentage of saline-treated controls (mean ± S.E.M.) and comparisons were done at discrete 10-min intervals, comparing vehicle to individual doses of drug. Statistical analysis was done by one-way ANOVA, followed by an unpaired t test; on a point-by-point basis, any data differing from vehicle at p ≤ 0.05 were considered an increase in activity.

**Turning in 6-Hydroxydopamine (6-OHDA)-Lesioned Rats.** Unilateral 6-OHDA injections into the substantia nigra cause selective destruction of dopamine neurons, leading to supersensitivity of the dopamine receptors in the caudate putamen on the injected side. In these animals, dopamine receptor agonists cause contralateral turning (Ungerstedt, 1971).

**Results**

**Studies with Cloned Receptors.** Sumanireole is an agonist selective for the D2 subtype of dopamine receptors (Table 1). To measure high-affinity, guanine nucleotide-sensitive agonist interactions at D2 receptors, an agonist ligand, [3H]U-86170, was used (Lahti et al., 1991). The affinity of sumanirole for D2 receptors is 9.0 ± 1.0 nM. This is similar to the affinity of D2 receptors for ropinirole and slightly higher

<table>
<thead>
<tr>
<th>Receptor</th>
<th>Sumanireole</th>
<th>Ropinirole</th>
<th>Bromocriptine</th>
<th>Pergolide</th>
</tr>
</thead>
<tbody>
<tr>
<td>D2</td>
<td>9.0 ± 1</td>
<td>7.2 ± 0.8</td>
<td>27 ± 9</td>
<td>1 ± 0.2</td>
</tr>
<tr>
<td>D3</td>
<td>1840 ± 142</td>
<td>22 ± 4</td>
<td>18 ± 2</td>
<td>0.4 ± 0.03</td>
</tr>
<tr>
<td>D4</td>
<td>&gt;2190</td>
<td>1450 ± 390</td>
<td>375 ± 15</td>
<td>9.3 ± 1</td>
</tr>
<tr>
<td>D1</td>
<td>&gt;7140</td>
<td>&gt;7140</td>
<td>3420 ± 130</td>
<td>1300 ± 132</td>
</tr>
</tbody>
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and lower than the receptor affinities for bromocriptine and pergolide, respectively, all agents used clinically for the treatment of Parkinson's disease (Eden et al., 1991; De Keyser et al., 1995; Rascol et al., 1996). The affinity of D₃, D₄, and D₁ receptors for sumanirole was calculated to be at least 200-fold lower than at D₂ receptors. In contrast, ropinirole, bromocriptine, and pergolide all showed high affinity for D₃ receptors; pergolide also showed high affinity for the D₄ receptor subtype.

We next examined whether sumanirole is an agonist or antagonist at the D₂ receptor using three measures: cAMP, arachidonic acid release, and mitosis. Forskolin-stimulated cAMP accumulation is inhibited by sumanirole and ropinirole in CHO cells (Fig. 2). Both compounds caused near maximal inhibition of forskolin-stimulated cAMP at the highest concentrations. Sumanirole had no effect on forskolin-stimulated cAMP accumulation, while sumanirole and ropinirole, and dopamine in two assays for activation of D₂ receptors in CHO-L6 cells: cAMP inhibition and potentiation of ATP-stimulated arachidonic acid release (Lajiness et al., 1994). Sumanirole is approximately 3 times less potent than ropinirole at each measurement (Table 2). Both compounds were as efficacious as dopamine at maximal concentrations, indicative of high levels of intrinsic activity. Sumanirole activated D₂ receptor-stimulated mitogenesis with an EC₅₀ of 4.6 nM, but it had no activity at D₃ and D₁ receptors in measurements of receptor-activated mitogenesis at concentrations up to 1 μM; this was consistent with the very low affinity of these receptors for this compound. In comparison, ropinirole potentiates activated D₂ and D₃ receptors in the mitogenesis assay [EC₅₀ values of 12 ± 1.0 nM (n = 5) and 7.5 ± 2.9 nM (n = 5), respectively].

Studies in Animals. Dopamine agonists that activate the D₂-like subfamily of dopamine receptors inhibit the release of prolactin from anterior pituitary cells (Ho and Thorner, 1988). The effects of four doses of sumanirole on plasma prolactin levels in male Sprague-Dawley were tested. Sumanirole decreased plasma prolactin at all doses studied with significant decreases at all doses (Table 3). The effects at 31 μmol/kg lasted at least 120 min postinjection.

Dopamine agonists depress dopamine neuron firing rates in the SNPC by activation of presynaptic dopamine receptors of the D₂-like subfamily. As demonstrated by population dose-response curves, sumanirole resulted in inhibition of SNPC dopamine neuronal firing with an ED₅₀ ± S.E.M. of 2.3 ± 0.9 μmol/kg i.v. (n = 6) (Fig. 3). Complete inhibition of basal firing rate was seen at 16.8 μmol/kg i.v. The inhibition of cell firing was reversed by haloperidol.

Activation of striatal postsynaptic D₂-like receptors decreases acetylcholine release and raises striatal acetylcholine concentrations (Sethy, 1979). Administration of both ropinirole and sumanirole to rats caused a dose-dependent increase in striatal acetylcholine levels (Fig. 4). The maximal effects of both compounds were similar, with a near doubling of striatal acetylcholine concentration. The ED₅₀ values for the effect (calculated as the fitted half-maximal response ± S.E.M. of the fit using the dose-response fit equation of Sigma Plot (SPSS Inc., Chicago, IL) were 6.0 ± 8.2 μmol/kg i.p. for ropinirole and 12.1 ± 4.1 μmol/kg i.p. for sumanirole.

Effects in Animal Models of Parkinson's Disease. Sumanirole increased locomotor activity in reserpinized/AMPT-pretreated rats with pronounced effects at 12.5 μmol/kg s.c. (Fig. 5). The onset of this effect was approximately 30 min postinjection, and the duration of the 12.5 μmol/kg dose was at least 2.5 h. Even greater locomotor activation was observed in reserpinized/AMPT-treated rats at 42 μmol/kg s.c., and the activation was completely blocked by 0.3 mg/kg haloperidol.

TABLE 2
In vitro D₂ potency of sumanirole and other agonists
Data are mean EC₅₀ ± S.E.M. for n shown in parentheses.

<table>
<thead>
<tr>
<th>Assay</th>
<th>Sumanirole</th>
<th>Ropinirole</th>
<th>Dopamine</th>
</tr>
</thead>
<tbody>
<tr>
<td>cAMP Inhibition</td>
<td>17 ± 4.8 (4)</td>
<td>5.6 ± 0.6 (3)</td>
<td>13&quot;</td>
</tr>
<tr>
<td>[³H]Arachidonate release</td>
<td>75 ± 16 (6)</td>
<td>19 ± 4.1 (3)</td>
<td>31 ± 9.3 (5)</td>
</tr>
<tr>
<td>Mitogenesis</td>
<td>32 ± 6.4 (3)</td>
<td>12 ± 1.0 (5)</td>
<td>5.4&quot;</td>
</tr>
</tbody>
</table>

" With permission from Lajiness et al. (1994).

![Fig. 2. Dose-dependent inhibition of forskolin-stimulated cAMP in CHO cells expressing recombinant D₂A receptors by sumanirole and ropinirole. Each point is a mean ± S.E.M. of triplicate determinations expressed as percentage of the 100 μM forskolin response. Concentration of drug given as nanomolar. ○, ropinirole; ●, sumanirole.](image-url)
operidol. These data suggest that sumanirole induces strong behavioral activation through a D₂ dopamine receptor.

Sumanirole was also highly efficacious at causing contralateral turning in unilateral 6-OHDA-lesioned rats after either oral or s.c. administration. A dose-response and time-course study of sumanirole given orally at 4.2, 12.5, or 42 μmol/kg showed turning activity in substantia nigra-lesioned rats, with significant effects seen at the two higher doses (Fig. 6). Maximal turning behavior was reached at 2 to 3 h after oral administration. Additionally, in this assay, sumanirole had a long duration of action as 12.5 and 42 μmol/kg still showed near-maximal efficacy at 4.5 h postinjection.

Ropinirole also caused dose-dependent increases in contralateral turning in the 6-OHDA-lesioned rats (Fig. 7). Significant turning behavior was measured at 19 μmol/kg s.c. A maximally effective dose of ropinirole was not as efficacious as sumanirole, as measured by the degree of turning. In the same set of rats, sumanirole (12.5 μmol/kg s.c.) induced approximately 3 times as many turns per 10-min interval than 19 μmol/kg s.c. ropinirole. Note that the effects of 12.5 μmol/kg sumanirole s.c. (Fig. 6) are equivalent to the effects produced by 42 mol/kg sumanirole administered orally (Fig. 7). Thus, the oral potency is approximately one-third the potency of the parenteral route. The peak response of sumanirole was also greater than the maximally effective doses of the following drugs given s.c.: 0.37 μmol/kg apomorphine, 4.6 μmol/kg bromocriptine, 9.5 μmol/kg pergolide, 13 μmol/kg quinpirole, and 11.7 μmol/kg SKF 38393. The turning activity induced by sumanirole is completely blocked by the coadministration of 0.27 or 2.7 μmol/kg haloperidol (Fig. 8), indicating that turning is mediated by a D₂-like dopamine receptor.

Drug-experienced cynomolgus monkeys rendered parkin-
sonian with MPTP-induced lesions of the substantia nigra were treated with different dosages of sumanirole and tested for improvement in scores using a MPTP monkey disability scale (Fig. 9). Two monkeys (A and B) had dose-dependent reductions in the disability scores after sumanirole, especially at the 12.5 \( \mu \text{mol/kg} \) dose. Monkey A had a dramatic recovery at 2.5 \( \mu \text{mol/kg} \) s.c. lasting for 5 to 6 h after a delay of 0.5 to 1 h. This monkey also had a dyskinesia accompanying the recovery. Monkey B required 12.5 \( \mu \text{mol/kg} \) sumanirole to show a dramatic response, and the response also lasted for 5 to 6 h after a 0.5 to 1 h delay. No dyskinesias were observed in monkey B to either sumanirole or other dopamine agonists. Monkey C did not show dramatic improvement even with the 12.5 \( \mu \text{mol/kg} \) dose; however, this monkey was unresponsive to apomorphine as well. In locomotion measurements, sumanirole at 12.5 \( \mu \text{mol/kg} \) s.c. increased the activity of monkeys A and B more than Prolopa. Monkey C did not increase locomotion with sumanirole or with apomorphine, although this monkey did respond to Prolopa (Fig. 10).

**Discussion**

The striking difference between sumanirole and the other examined dopamine agonists (ropinirole, bromocriptine, and pergolide) lies in sumanirole’s lack of affinity for D3 and D4 receptors, whereas the other dopamine agonists all showed high affinity for D3 and/or D4 receptors. Sumanirole is an imidazoquinolinone and the first compound described as a highly selective agonist for D2 receptors over the other D2-like subtypes, D3 and D4. The selectivity of this compound was demonstrated in radioligand binding assays, where it has at least 200-fold higher affinity for D2 receptors than for the other dopamine receptor subtypes. The other dopamine agonists examined did not demonstrate D2 receptor-selectivity. Sumanirole has been shown to be inactive in more than 80 enzyme and receptor assays. The compound does have moderate affinity at the 5-HT1A receptor (\( K_i = 95 \text{nM} \); unpublished observations).

D2 receptors are abundant in motor areas such as the basal ganglia; therefore, their activation may be associated with therapeutic efficacy of dopamine agonists in Parkinson’s disease. In contrast, D3 and D4 receptors are located in limbic and cortical areas and may contribute to the psychiatric disturbances that accompany dopamine agonist and L-Dopa therapeutics. In support of this hypothesis, clozapine, a dopamine receptor blocker with higher affinity for the D2 subtype, has been used to ameliorate the psychoses accompany-
Antiparkinsonian treatment (Friedman, 1995). D₃ receptor agonists have also been shown to decrease locomotor activity (Lagos et al., 1998). This suggests that sumanirole's unique selectivity may confer beneficial effects in the treatment of Parkinson's disease. Eliminating D₂- and D₃-activating properties of dopamine receptor agonists may potentially reduce nonmotor side effects and improve tolerability. Whether the unique selectivity of sumanirole translates into a clinical advantage over other dopamine agonists remains to be determined.

The unique selectivity of sumanirole allows the compound to be used to characterize D₂-specific activity in the central nervous system that had previously been only inferred by using nonselective D₂-like receptor agonists. Sumanirole inhibits prolactin release. D₂ receptor mRNA but not D₃ receptor mRNA is found in the pituitary gland (Bunzow et al., 1988; Sokoloff et al., 1990), and D₂ receptors inhibit prolactin release from lactotrophs (Missale et al., 1991). Thus, it is not surprising that the D₂-selective agonist sumanirole can mediate the prolactin inhibition response, but these results provide evidence that sumanirole has anticipated in vivo D₂ activity. D₂-like receptors inhibit the firing rates of SNPC dopamine neurons (Pinnock, 1984; Piercey et al., 1996a,b).

Both mRNA for D₂ and D₃ receptors have been localized to these dopaminergic neurons (Meador-Woodruff et al., 1989; Bouthenet et al., 1991). Sumanirole inhibited the firing rate of these cells in a similar dose range as the doses shown to inhibit prolactin release. These findings support a role for the molecularly defined D₂ receptors as autoreceptors that regulate the firing rates of SNPC neurons. Furthermore, activation of D₂-like receptors decreases acetylcholine release (Consolo et al., 1987). Changes in acetylcholine content in the striatum were used as an in vivo measure of D₂-like receptor activation. By this measurement, sumanirole is equally efficacious in vivo as other compounds used as antiparkinsonian agents. In three animal models of Parkinson's disease, sumanirole was shown to have good locomotor stimulant properties, including reserpine/AMPT-treated rats, 6-OHDA-lesioned rats, and MPTP monkeys. Reserpine treatment of rats to deplete catecholamine stores results in a hypokinetic animal that can be activated by L-dopa and provides an animal model for Parkinson's disease (Carlsson et al., 1957). Reserpine-induced akinesia was dose dependently reversed by sumanirole and the activation effect was blocked by haloperidol. The mechanism for this locomotor activation is likely to be direct stimulation of postsynaptic D₂ dopamine receptors.

**Fig. 7.** Comparison of dose-response ropinirole s.c. to a single dose of sumanirole s.c. on turning behavior in unilaterally 6-OHDA-lesioned rats. Turning behavior of rats with unilateral 6-OHDA lesions of the substantia nigra was measured after injections of vehicle (○), n = 13; 3.8 μmol/kg ropinirole s.c. (●), n = 6; 19 μmol/kg ropinirole s.c. (▲), n = 8; and 12.5 μmol/kg sumanirole s.c. (●), n = 6. *p < 0.05 significance compared with controls after F-test for variance of means, followed by one-way ANOVA with t test for individual points.

**Fig. 8.** Haloperidol antagonism of sumanirole-induced turning behavior in unilaterally 6-OHDA-lesioned rats. Turning behavior of rats with unilateral 6-OHDA lesions of the substantia nigra was measured after injections of either vehicle or 12.5 μmol/kg sumanirole and the indicated concentrations of haloperidol. Each bar is the total area under the curve ± S.D. (n ≥ 4) for turning behavior from 0 to 210 min postinjection. *p < 0.05 significance when compared with controls after F-test for variance of means, followed by one-way ANOVA with t test for area under curve.
Sumanirole was highly efficacious at causing contralateral turning in unilateral 6-OHDA-lesioned rats after either oral or s.c. administration, demonstrating superior efficacy over other dopamine agonists tested in this model. Ropinirole was slightly more potent but equally efficacious as sumanirole in assays of D$_2$ receptor activation in cell-based assays, and at D$_2$-like receptor activation in other in vivo responses. However, even maximally effective doses of ropinirole did not produce the same degree of response as sumanirole on rotational behavior. Furthermore, sumanirole was more efficacious than other agonists (bromocriptine, pergolide, and apomorphine) tested at maximally effective doses in this model. The enhanced efficacy of sumanirole in the 6-OHDA turning model could be a result of other undiscovered activities of this compound; however, turning was completely blocked by the dopamine receptor antagonist haloperidol. This suggests the enhanced efficacy in this model resulted from sumanirole’s unique D$_2$-receptor selectivity.

MPTP-lesioned monkeys represent another animal model for Parkinson’s disease. The toxin MPTP causes selective destruction of dopaminergic neurons resulting in behavioral deficits similar to those observed in diseased humans (Burns et al., 1983). L-Dopa and dopamine agonists reverse the behavioral deficits in MPTP-treated monkeys. Chronic L-dopa can also cause dyskinetic symptoms in monkeys as it does in patients with Parkinson’s disease (Bedard et al., 1986). Sumanirole has only been tested to a limited extent in the MPTP-lesioned monkey model and only in monkeys tested extensively with other dopaminergic agents. Sumanirole caused improvement in Parkinson’s disability scores in two of the three monkeys. Sumanirole also increased the locomotion of the two responding monkeys to a greater extent than Prolopa. This is consistent with the high efficacy effects of sumanirole in the rat 6-OHDA turning model. The studies in the MPTP-lesioned monkey suggest that sumanirole is efficacious in this model but are too limited to draw solid conclusions about the relative efficacy or the dyskinetic potential of this compound. The current issue of *Journal of Pharmacology and Experimental Therapeutics* reports results of a subsequent study in MPTP-treated monkeys in which sumanirole shows antiparkinsonian effects comparable with existing dopaminergic therapies without inducing dyskinetic symptoms using both behavioral and pathological assessments (Stephenson et al., 2005).
Conclusions

Sumanire represents the first dopamine agonist with high selectivity for the D2 receptor subtype over the other closely related D3 and D4 receptor subtypes. In vitro studies demonstrate that sumanire is a selective D2 receptor agonist and can therefore be used to evaluate the importance of the D2 receptor in the physiological effects of dopamine agonists. The D2-selective nature of sumanire is assumed to allow for the assessment of D2 agonist effects on locomotor activity in the absence of D3 and D4 receptor coactivation. Studies presented here suggest that, in animals, many of the physiologically responses thought to be mediated via the D2 receptor subtype, namely, prolactin release, regulation of striatal acetylcholine content, and autoreceptor-mediated inhibition of nigrostriatal firing rates, are indeed mediated by the molecularly defined D2 receptor subtype, since sumanire was fully efficacious at in vivo concentrations expected to selectively activate D2 receptors. Sumanire thus represents a potential tool for studying D2 receptor subtype specificity both in vitro and in vivo.

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