Phosphodiesterase Inhibition and Regulation of Dopaminergic Frontal and Striatal Functioning: Clinical Implications

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Abstract

Background: The fronto-striatal circuits are the common neurobiological basis for neuropsychiatric disorders, including schizophrenia, Parkinson's disease, Huntington's disease, attention deficit hyperactivity disorder, obsessive-compulsive disorder, and Tourette's syndrome. Fronto-striatal circuits consist of motor circuits, associative circuits, and limbic circuits. All circuits share 2 common features. First, all fronto-striatal circuits consist of hyper direct, direct, and indirect pathways. Second, all fronto-striatal circuits are modulated by dopamine. Intracellularly, the effect of dopamine is largely mediated through the cyclic adenosine monophosphate/protein kinase A signaling cascade with an additional role for the cyclic guanosine monophosphate/protein kinase G pathway, both of which can be regulated by phosphodiesterases. Phosphodiesterases are thus a potential target for pharmacological intervention in neuropsychiatric disorders related to dopaminergic regulation of fronto-striatal circuits.

Methods: Clinical studies of the effects of different phosphodiesterase inhibitors on cognition, affect, and motor function in relation to the fronto-striatal circuits are reviewed.

Results: Several selective phosphodiesterase inhibitors have positive effects on cognition, affect, and motor function in relation to the fronto-striatal circuits.

Conclusion: Increased understanding of the subcellular localization and unraveling of the signalosome concept of phosphodiesterases including its function and dysfunction in the fronto-striatal circuits will contribute to the design of new specific inhibitors and enhance the potential of phosphodiesterase inhibitors as therapeutics in fronto-striatal circuits.

Keywords: fronto-striatal circuits, dopamine, phosphodiesterase, phosphodiesterase inhibitors, cyclic adenosine monophosphate

Introduction
Several neuropsychiatric disorders, including Parkinson’s disease, Huntington’s disease, attention-deficit hyperactivity disorder (ADHD), Tourette’s syndrome, schizophrenia, and obsessive-compulsive disorder, share the fronto-striatal circuits, also known as cortico-striatal-thalamic circuits, as their neurobiological basis. The fronto-striatal circuits comprise motor, cognitive, and limbic circuits (Alexander et al., 1986, 1990; Alexander and Crutcher, 1990). These circuits operate in a very complex manner that is extensively described elsewhere (Surmeier et al., 2007; Haber and Rauch, 2010; Gerfen and Surmeier, 2011; Surmeier et al., 2011; Calabresi et al., 2014). Dysfunction of these circuits produces the wide range of motor, cognitive, and affective symptoms observed in related neuropsychiatric disorders. One prominent feature of the complex functioning of the fronto-striatal circuits is their modulation by dopamine, both at the level of the frontal cortex as well as the striatum. As a result, dopaminergic receptors are strongly expressed throughout all fronto-striatal circuits (Gerfen and Surmeier, 2011; Nishi et al., 2011; Kuroiwa et al., 2012). Unsurprisingly, dopaminergic medication has been the first-line therapy for several disorders related to dysfunctional fronto-striatal circuits; however, efficacy is often moderate at best and accompanied by severe side effects (e.g., ADHD, schizophrenia, and Parkinson’s disease).

Dopamine originating from substantia nigra pars compacta (SNc) and/or ventral segmental area (VTA) (nigrostriatal and mesolimbic pathways) binds to both dopamine type1 (D1) receptors and dopamine type2 (D2) receptors on medium spiny neurons (MSNs) in the striatum (Gerfen and Surmeier, 2011). D1 receptors are mainly found on MSNs of the direct pathway, and D2 receptors are mainly found on MSNs of the indirect pathway where they establish antagonistic interactions with adenosine A2A receptors (Gerfen et al., 1990; Ferre et al., 2011). Additionally, dopamine released from VTA (mesocortical pathway) also binds to D1 receptors in the frontal cortex (Kuroiwa et al., 2012). D1 receptors activate the Gi/Go family of G proteins to stimulate cyclic adenosine monophosphate (cAMP) production and thereby striatonigral and frontal signaling (Sibley et al., 1993; Beaulieu and Gainetdinov, 2011). In contrast, the D2 receptors couple to the Gs family of G proteins and thus induce inhibition of cAMP production, thereby inhibiting striatopallidal signaling that eventually leads to disinhibition of the frontal cortex (Figure 1). Actions of the dopamine receptors in both pathways can be viewed as synergistically or complementary.

Intracellularly, the effect of dopamine on striatonigral, striatopallidal, and frontal neurons is largely mediated through the cAMP-activated cascade (Nishi et al., 2008, 2011; Nishi and Snyder, 2010; Kuroiwa et al., 2012). cAMP is synthesized from adenosine triphosphate by adenylyl cyclase, which is activated directly by activated G-protein coupled receptors or by calmodulin (CaM)-dependent protein kinase (PK)/after Ca2+ influx. cAMP affects synaptic plasticity through both presynaptic neurotransmitter release and postsynaptic intracellular pathways (Figure 1). The former might be mediated via a presynaptic calcium (Ca2+)/CaM-dependent PK/cAMP/cAMP-dependent PKA cascade and elevation of cAMP has been found to result in the synthesis and/or release of several neurotransmitters, including 2 main players in the fronto-striatal circuits: glutamate and dopamine (Schoppelmeier et al., 1985; Imanishi et al., 1997; Rodriguez-Moreno and Sihra, 2013).

The influence on postsynaptic intracellular pathways occurs through activation of postsynaptic PKA by cAMP produced by adenylyl cyclase stimulated by either glutamatergic-induced Ca2+ influx or dopamine signaling-stimulated Gz. PKA exerts several effects related to neuroplasticity and neuroprotection. The fastest postsynaptic response in relation to neuroplasticity mediated by cyclic nucleotides is the activation and insertion of stored receptors by PKA through phosphorylation of Glur1 subunits promoting α-3-hydroxy-5-methyl-4-isoxazolopropionic acid (AMPA) receptor trafficking into the postsynaptic membrane for potentiation of glutamatergic transmission (Song et al., 2013). In addition to the mobilization and membrane insertion of stored receptors, the process of protein synthesis (e.g., AMPA receptors) further increases neuroplasticity (Carew and Sutton, 2001; Izquierdo et al., 2006).

PKA also phosphorylates cAMP response element-binding protein (CREB) (Mayr and Montminy, 2001) and Dopamine and cAMP-Regulated Phosphoprotein MR 32kd (DARPP-32) (Greenberg, 2001; Svenningsson et al., 2004). Phosphorylated CREB is also involved in neuroplasticity (e.g., synthesis of other proteins) (Impay et al., 1996; Lu et al., 1999; Sakamoto et al., 2011) and neuroprotection (e.g., neuronal arborization, synaptogenesis, and neurogenesis) (Mantamadiotis et al., 2002; Bruel-Jungerman et al., 2006; Sakamoto et al., 2011). One of the genes transcribed by phosphorylated CREB is bdnf (Scott Bitner, 2012). After release, the protein BDNF binds to the tropomyosin-related kinase B receptor, which is the receptor with the highest affinity for BDNF. BDNF is involved in the proliferation, survival, and differentiation of new neurons (i.e., neurogenesis in the brain) (Minichiello, 2009).

In addition, the activity-dependent release of BDNF and subsequent tropomyosin-related kinase B-mediated activation of CREB is also an important mechanism of enhancing neuronal communication, specifically in active neurons of the brain. For instance, BDNF increases synaptic strength with adjacent neurons by processes like long-term potentiation (LTP), thus ameliorating their connectivity (Lu et al., 2008; Minichiello, 2009). Interestingly, LTP itself has been linked to both synaptogenesis and neurogenesis (Bruel-Jungerman et al., 2006).

DARPP-32 is phosphorylated at Thr24 in both striatal and frontal neurons. DARPP-32 thereby converts into a potent inhibitor of protein phosphatase-1 (PP-1). DARPP-32 is also phosphorylated at Thr75 by Cdk5 and this converts DARPP-32 into an inhibitor of PKA. Thus, DARPP-32 has the unique property of being a dual-function protein, acting either as an inhibitor of PP-1 or of PKA influencing neuroplasticity (Svenningsson et al., 2004). The inhibition of PP-1 controls the phosphorylation state and activity of many downstream physiological effectors, including various neurotransmitter receptors (e.g., AMPA receptor Glur1 subunit, N-methyl-D-aspartate receptor NR1 subunit), ion channels and pumps (e.g., N/P-type Ca2+ channels, Na+ channel, Na+, K+-ATPase), and transcription factors (e.g., CREB, c-Fos, ΔFosB) (Greenberg et al., 1999). Striatal LTP and long-term depression are dependent on cAMP and DARPP-32 phosphorylation (Calabresi et al., 2000).

The cAMP/PKA cascade is thus a potential target for pharmacological intervention in neuropsychiatric disorders related to dopaminergic frontal and striatal dysfunction. cAMP is degraded by cAMP-specific phosphodiesterases (PDEs) and dual substrate PDEs. Eleven PDE families have been described, distinguished by molecular properties, substrate specificity, and regulation (Bender and Beavo, 2006). These enzymes are expressed in unique and overlapping patterns throughout the body and central nervous system (Lakics et al., 2010; Table 1). Selective PDE inhibitors (PDE-Is) prevent the degradation of cyclic nucleotides leading to increased concentrations of cAMP. Due to the differential expression of PDE subtypes in one or more of the frontal and striatal pathways or dopaminergic terminals, different subtype-specific PDE-Is enable stimulation of dopamine synthesis, inhibition of D2 receptor signaling or stimulation of D1 receptor...
Output neurons in the striatum are medium spiny neurons (MSNs), which consist of direct pathway and indirect pathway neurons. The direct pathway neurons inhibit tonically active neurons in globus pallidus interna (GPi)/substantia nigra pars reticulata (SNr). The indirect pathway neurons activate neurons in GPi/SNr via inhibition of the globus pallidus externa (GPe) and activation of the subthalamic nucleus (STN). Direct and indirect pathway neurons induce opposing effects on the output neurons in GPi/SNr, resulting in disinhibition and proinhibition of output, respectively. Within the basal ganglia all projections are GABAergic except those from the STN. Main phosphodiesterases (PDEs) expressed in fronto-striatal circuits are PDE1B, PDE4, and PDE10A. PDE1B is generally colocalized with dopamine (DA) D1 receptors in the brain and thought to represent a major inactivation mechanism of D1 receptors. By acting like a DA D1 agonist, PDE1B-Is can enhance phosphorylation of cAMP response element binding protein (CREB) as well as Dopamine- and cAMP-Regulated Phosphoprotein MR 32kDa (DARPP-32) enhancing synaptic transmission, e.g., AMPA receptors, neuron excitability, and neurogenesis, resulting in neuroplasticity and neuroprotective effects at glutamatergic frontal and fronto-striatal synapses. Regarding fronto-striatal signaling, the effect of PDE4 inhibition on cAMP/protein kinase A (PKA) signaling, is linked to indirect pathway adenosine A2a receptor signaling and has no major role in D1 receptor direct pathway signaling. An opposite situation is observed at frontal dopaminergic signaling. In the frontal cortex, PDE4 is localized at DARPP-32 expressing neurons. In contrast to the striatum, PDE4 inhibition enhances D1 receptor-induced phosphorylation of DARPP-32 in the frontal cortex, indicating a prominent role of PDE4 in frontal DA receptor signaling. Finally, DA release from DAergic midbrain terminals can be influenced with a PDE4 inhibitor as DA is expressed at DAergic terminals in neurons of the SNc in which CAMP has been reported to be a strong inducer of tyrosine hydroxylase (TH) gene transcription rate and mRNA affecting DA synthesis and release. In direct pathway neurons, PDE10A inhibition activates cAMP/PKA signaling related to D1 receptor signaling, whereas in indirect pathway neurons PDE10A inhibition activates cAMP/PKA signaling by simultaneous potentiation of adenosine A2A receptor signaling and inhibition of D2 receptor signaling. Effects of PDE10A inhibition are suggested to predominate the indirect pathway. In contrast to PDE4 inhibition, PDE10A inhibition does not increase TH phosphorylation and therefore has no effects on DA synthesis and release. Nevertheless, it cannot be ruled out that selective PDE inhibitors (PDE-Is) might influence both the direct and indirect pathway via enhancing the release of DA from frontal DAergic projections depending on the presence of PDEs in these terminals.

In striatal interneurons containing nitric oxide synthase (NOS), nitric oxide (NO) is produced and diffuses into dendrites of MSNs which contain high levels of guanylate cyclase (GC), which, when activated, lead to the synthesis of cyclic guanosine monophosphate (cGMP). In the striatum, transient elevations in intracellular cGMP, next to cAMP, primarily act to increase neuronal excitability and to facilitate glutamatergic fronto-striatal transmission. Thus, inhibition of selective PDE subtypes can also target the cGMP/protein kinase G (PKG) pathway and have an effect on fronto-striatal functioning.
Table 1. Localization of the Different PDEs in the Brain of Rodents and Humans in Adulthood

<table>
<thead>
<tr>
<th>PDE</th>
<th>Localization in the Body</th>
<th>Localization in the Brain</th>
</tr>
</thead>
<tbody>
<tr>
<td>PDE1A-C</td>
<td>Heart, smooth muscles, lungs</td>
<td>Hippocampus, cortex, olfactory bulb, striatum (highest expression levels), thalamus, amygdala, cerebellum; expression levels are in general highest for 1A and lowest for 1C</td>
</tr>
<tr>
<td>PDE2A</td>
<td>Heart, adrenal cortex, platelets</td>
<td>Hippocampus, cortex, striatum, hypothalamus, amygdala, midbrain</td>
</tr>
<tr>
<td>PDE3A-B</td>
<td>Heart, smooth muscles, kidneys, platelets</td>
<td>Throughout the brain low expression levels</td>
</tr>
<tr>
<td>PDE4A-D</td>
<td>Wide variety of tissues: e.g., smooth muscles, lungs, kidneys, testes</td>
<td>Hippocampus, cortex, olfactory bulb, striatum, thalamus, hypothalamus, amygdala, midbrain, cerebellum; expression levels in general are for 4A-4D (differs per brain structure) and lowest for 4C</td>
</tr>
<tr>
<td>PDE5A</td>
<td>Smooth muscles, skeletal muscles, lungs, kidneys, platelets</td>
<td>Hippocampus, cortex, cerebellum</td>
</tr>
<tr>
<td>PDE6A-C</td>
<td>Rod and cone cells in retina</td>
<td>Pineal gland</td>
</tr>
<tr>
<td>PDE7A-B</td>
<td>Heart, skeletal muscles, liver, kidneys, testes, pancreas</td>
<td>Hippocampus, cortex, olfactory bulb, striatum, thalamus, hypothalamus, midbrain; expression levels are in general highest for 7B</td>
</tr>
<tr>
<td>PDE8A-B</td>
<td>Heart, liver, kidneys, lungs, testes, thyroid</td>
<td>Hippocampus, cortex, olfactory bulb, striatum, thalamus, hypothalamus, midbrain; expression levels are in general highest for 8B</td>
</tr>
<tr>
<td>PDE9A</td>
<td>Kidneys, spleen, prostate, various gastrointestinal tissues</td>
<td>Hippocampus, cortex, olfactory bulb, striatum, thalamus, hypothalamus, amygdala, midbrain, cerebellum</td>
</tr>
<tr>
<td>PDE10A</td>
<td>Heart, skeletal muscles, lungs, liver, kidneys, testes, pancreas, thyroid</td>
<td>Hippocampus, cortex, striatum (highest expression levels), midbrain, cerebellum</td>
</tr>
<tr>
<td>PDE11A</td>
<td>Skeletal muscles, liver, kidneys, testes, prostate, thyroid</td>
<td>Throughout the brain low expression levels</td>
</tr>
</tbody>
</table>

Abbreviation: PDE, phosphodiesterase.

Note that this table does not provide information with respect to the level of expression (protein or mRNA) of the different PDEs. Adapted from Prickaerts, 2015, based on Lakics et al, 2010; Pérez-Torres et al, 2010.

signaling (Nishi et al., 2011). However, the level of expression of different PDE family members in these fronto-striatal circuits in both physiological and pathological conditions is incompletely understood and a subject of intense investigation. In the fronto-striatal circuits, the main therapeutic mechanism of PDE inhibition is enhanced neuroplasticity and neuroprotection through previously discussed CREB and DARP3-32 signaling cascades (Figure 1). However, known effects of PDEs on neuroinflammation and cytokine-mediated responses may play additional roles (Hebb and Robertson, 2008; Wilson and Brandon, 2015).

The integration of individual PDEs into specific signalosomes within different functional compartments has revealed the functional roles of these PDEs and linked the large number of PDE isoforms to the compartmentalized regulation of specific cyclic nucleotide signaling pathways and biological responses. This compartmentalization therefore contributes to both the fine-tuning and specificity of cyclic nucleotide signaling (Jurevicius and Fischmeister, 1996; Zaccolo et al., 2000; Mongillo et al., 2006; Mauric, 2011; Stangherlin et al., 2011; Stangherlin and Zaccolo, 2012). More in detail, compartmentalization provides spatially distinct pools of PKA and PKG to be activated in different ways. This idea was confirmed by the observation of accumulation of cAMP in localized pools (Houslay, 1995). These pools are created by physical interactions between different components of signaling cascades and structural elements of the cell. Localization and activation of both cyclases and PDEs are important determinants in the process of cyclic nucleotide homeostasis by modulating fluctuations in the compartments. For PDEs, sequestration and anchoring is the principal mechanism to create cyclic nucleotide gradients (Houslay and Milligan, 1997; Houslay and Adams, 2003). Subsequently, different PKA isoforms are anchored at specific intracellular sites by A-kinase anchoring proteins (AKAPs) (Rubin, 1994). AKAPs control the gradients of cAMP in the cell and modify localized target proteins, thereby causing sequestration of PKA into distinct cellular compartments. This also applies to the fronto-striatal circuits and is considered the main determinant of the target PDE isoform within a specific neuropsychiatric disorder. In addition, different PDE isoforms can integrate multiple distinct cellular inputs and allow crosstalk between cyclic nucleotides and other signaling networks and systems (Dodge-Kafka et al., 2005; Mongillo et al., 2006; Houslay et al., 2007; Stangherlin et al., 2011; Wilson et al., 2011; Kritzer et al., 2012).

Since the focus of this review is on frontal and striatal dopaminergic regulation, PDE1B, PDE2A, PDE4, PDE7B, PDE9A, and PDE10A in particular are of special interest (Lakics et al., 2010). PDE1B, PDE7B, and PDE10A are highly enriched in striatum and/or frontal cortex. PDE2A, PDE4 (A, B, D), and PDE9A are more widely distributed but are also expressed in striatum and/or frontal cortex. There are only very limited preclinical data on PDE2A, PDE7A, and PDE9A inhibition (e.g., Duinen et al., 2015). To date, most research has been devoted to the potential of PDE1B, PDE4, and PDE10A for regulation of dopaminergic frontal and striatal signaling, and therefore these subtypes will be discussed below.

**PDE1**

**PDE1 and Dopamine Signaling**

PDE1 hydrolyzes both cAMP and cGMP. PDE1, unlike any other class of PDE, is uniquely activated by the binding of a complex of Ca²⁺ and CaM. PDE1 is encoded by 3 separate genes: PDE1A, PDE1B, and PDE1C. PDE1B is highly colocalized with D1 receptors in the brain and is particularly rich in the striatum, hippocampus, and prefrontal cortex (Lakics et al., 2010). PDE1 activity was first described as cytosolic; however, it now appears that PDE1A is not restricted to the cytosol but is also present in the nucleus where it contributes to the regulation of transcription factors (Nagel et al., 2006). This opens a new field of research in transcriptional regulation. Changes in PDE1 location associated with cell differentiation might contribute to compartmental signaling (Nagel et al., 2006).
Since PDE1B is strongly and selectively expressed in the striatum and frontal cortex, it could be a relevant target for modulating fronto-striatal behaviors. However, only a few studies have been published with PDE1-Is (Medina, 2011; Nunes et al., 2011). Indeed, as noted by the authors, in these studies no potent and selective inhibitors of PDE1 isoforms were available for research. Vinpocetine, often referred to as a PDE1-I, has substantial other activities, including inhibition of Na+ channels and iKβ kinase. Vinpocetine therefore should not be considered as a selective PDE1-I. Since PDE1B is activated by Ca2+ and CaM, it provides a mechanism for crosstalk between Ca2+ and cyclic nucleotide signaling (Nishi et al., 2008, 2011; Nishi and Snyder, 2010). PDE1B was localized to all DARPP-32-positive MSNs indicating expression in both striatal pathways (Nishi et al., 2011). Behavioral profiles of PDE1B knockout mice showed rather mild behavioral effects. The authors show an increased spontaneous locomotor activity in the presence of methamphetamine administration. Cognitive aspects are reported as similar to wild type. Overall, the data suggest predominant effects of PDE1B-Is would be seen in the striatongrall direct pathway (Reed et al., 2002; Ehrman et al., 2006; Siuciak et al., 2007; Zhang, 2010). However, regarding effects on frontal and striatal dopamine release or effects on cAMP/PKA signaling in the frontal cortex, these areas have so far been understudied.

Implications and Clinical Overview of PDE1B-Is

PDE1B is generally colocalized with dopamine D1 receptors in the brain and thought to represent a major inactivation mechanism of D1 receptors. PDE1B is not membrane bound but contained mainly in a soluble intracellular compartment (Fusco and Giampa, 2015). Targeting this subfamily of PDEs is therefore considered a promising therapeutic strategy in disorders characterized by frontal cognitive dysfunction, like schizophrenia and ADHD. Negative and cognitive symptoms of schizophrenia are associated with reduced dopamine (D1) function in the prefrontal cortex, also referred to as hypofrontality (Liemburg et al., 2012; Arnsten, 2013). The reduced prefrontal dopamine function disturbs the balance of excitatory to inhibitory synaptic interactions in this area (Winterer, 2006). Thus, the decreased ratio of D1/D2 signaling in schizophrenia would favor unstable cortical representation of internal and external stimuli (Winterer and Weinberger, 2004) and as such, affect cognition. Likewise, dopaminergic hypofrontality is observed in ADHD patients (Pliszka, 2005; Saygölden et al., 2005; Arnsten and Pliszka, 2011) and linked to inattentiveness, hyperactivity, and impulsivity. By acting supposedly like a dopamine D1 agonist, a PDE1B-I can enhance phosphorylation of GluR1 subunits to potentiate glutamatergic fronto-striatal signaling. In addition, potentiation of dopamine receptors will increase phosphorylation of DARPP-32 and CREB, subsequently inducing gene expression in the prefrontal cortex and benefiting clinical symptoms by activation of neuroplasticity at prefrontal synapses.

Regarding striatal disorders like movement disorders and positive symptoms in schizophrenia, not much data is available. Assuming effects of PDE1B are indeed preferentially induced in the dopamine D1 direct pathway, subsequent DARPP-32 and CREB phosphorylation will enhance synaptic transmission, neuron excitability, and synapto-neurogenesis inducing neuroplasticity and neuroprotection at glutamatergic fronto-striatal synapses. In theory, neuropsychiatric disorders related to striatal hypofunction (like hypokinetic movement disorders such as Parkinson’s disease) would benefit from stimulated plasticity in striatonigral neurons (Nishino et al., 1993; Heckman et al., 2015). This is in contrast to the desired mechanism of action of treatment for hyperkinetic movement disorders (like Huntington’s disease) and antipsychotic treatment, as both preferably target the D2 receptor striatopallidal pathway (Strange, 1998; Walker, 2007). Support for this hypothesis comes from studies that found impaired cyclic nucleotide signaling mechanisms to occur in human Parkinson’s disease as well as in experimental animals (Belmaker et al., 1978; Volcker et al., 1986; Nishino et al., 1993; Sancesario et al., 2004). Additional upregulation in PDE1B activity has also been observed following 6-hydroxydopamine (6-OHDA) lesions (as a Parkinson’s model) in rats (Sancesario et al., 2004). The opposite has been observed in Huntington’s disease (Luthi-Carter et al., 2000; Nuñifora et al., 2001). These studies found decreased cyclic nucleotide (cAMP) levels in the deafferented striatum accompanied by decreased PDE1B activity. The decreased PDE1B activity may be compensatory to the decrease in cyclic nucleotide levels. CREB-mediated transcriptional dysregulation has also been reported to occur during Parkinson’s disease pathology. PDE1-Is have been tested in an animal model of haloperidol-induced catalepsy. As haloperidol is a potent D2 dopamine receptor antagonist, interference with D2 dopamine receptors causes significant motor disturbances seen frequently with schizophrenics treated with antipsychotic medicines. The haloperidol-induced catalepsy model is capable of testing agents for exacerbation or lessening of these motoric effects. The potent and selective PDE1-I, ITI-214 (see below), reversed the haloperidol-induced catalepsy, indicating the potential use of this mechanism to reverse such motoric effects (Wenngöle et al., 2010).

BDNF has been demonstrated to exert protective actions on nigral dopaminergic neurons in vivo and in vitro models of Parkinson’s disease (Hyman et al., 1991; Levivié et al., 1995; Shults et al., 1995; Hung and Lee, 1996; Feng et al., 1999; Mohapel et al., 2005; Sun et al., 2005), whereas inhibition of nigral BDNF expression has been reported to cause dopaminergic neuronal loss (Porritt et al., 2005). Postmortem studies have demonstrated reduced levels of BDNF within the SNs in Parkinson’s disease patients (Mogi et al., 1999; Parain et al., 1999; Howells et al., 2000; Chauhan et al., 2001). Furthermore, during in vitro experiments, BDNF has been demonstrated to promote the survival and differentiation of mesencephalic dopaminergic neurons (Hyman et al., 1991; Feng et al., 1999). The enhancement of cerebral cyclic nucleotide levels by PDE1B inhibition would improve CREB-mediated signaling mechanisms providing therapeutic effects in Parkinson’s disease.

Recently, a set of 4 clinical studies were performed with a truly selective and potent PDE1-I, ITI-214 (Li et al., 2016a). With the exception of work performed with vinpocetine, a nonselective agent as noted above, ITI-214 is the first selective PDE1-I studied in humans. Clinical evaluations included a series of Phase I single- and multiple ascending-dose studies performed in the US and Japan. ITI-214 was given orally to healthy volunteers and patients using once-a-day dosing and was shown to be safe and well tolerated, with a linear pharmacokinetic profile. This study has been reported in a press release (Intra-Cellular Therapies, 2014), where the company concludes that “these studies represent a significant milestone as the first demonstration of the safety of a potent and highly specific PDE1-I in humans.”

PDE4

PDE4 and Dopamine Signaling

PDE4, which is cAMP specific, is encoded by 4 distinct genes in mammals, PDE4A, PDE4B, PDE4C, and PDE4D, and is expressed as at least 25 splice variants. Each of these variants has a modular structure consisting of a variant-specific N-terminal domain,
regulatory domains (upstream conserved region 1 and 2 [UCR1 and UCR2]), a conserved catalytic domain, and an isoform-specific C-terminal domain (McCaffil et al., 2008; Gurney et al., 2011; Richter et al., 2015). Transcription of a number of PDE4 genes is activated by the cAMP/PKA/CREB cascade (D’Sa et al., 2002; Le Jeune et al., 2002), and PKA induction of PDE4 genes serves as a long-term feedback mechanism. The N-terminal domain and UCR1/2 interact with variant-specific binding proteins to direct the subcellular targeting of PDE4 variants (McCaffil et al., 2008). Various targeting proteins have been identified, including arrestin, AKAP5, receptor for activated C kinase 1, disrupted in schizophrenia 1, Src, and extracellular receptor kinase (ERK) (see Nishi and Snyder, 2010).

Nishi and colleagues (2008) showed that the inhibition of PDE4 by rolipram weakly enhanced cAMP/PKA signaling both in neostriatal slices and in vivo (Nishi et al., 2008). Rolipram increased the phosphorylation of DARPP-32 but only at high concentrations. Rolipram treatment enhanced adenosine A$_2$ receptor-mediated phosphorylation of DARPP-32 but had no effect on D1 receptor/cAMP/PKA-mediated phosphorylation at the level of DARPP-32. Enhanced adenosine A$_2$ receptor-mediated signaling is expected to oppose actions of the dopamine D2 receptor in striatopallidal neurons. These findings may suggest that PDE4 is exclusively expressed in indirect pathway neurons. However, immunohistochemical analysis of previously mentioned neostriatal slices revealed that PDE4B expression can be found in both pathways but with a higher expression in indirect pathway neurons. Regarding striatal dopaminergic signaling, it seems that the effect of PDE4 inhibition on cAMP/PKA signaling is linked to adenosine A$_2$ receptor signaling and has no major role in striatal dopamine signaling. An opposite situation is observed at frontal dopaminergic signaling. In the frontal cortex, several PDE isoforms are expressed in cortical neurons (Cherry and Davis, 1999; Pérez-Torres et al., 2000). For the mouse frontal cortex, it has been described that PDE4B is localized at DARPP-32-expressing neurons (Nishi and Snyder, 2010). In contrast to the striatum, rolipram enhanced dopamine D1 receptor-induced phosphorylation of DARPP-32 in the frontal cortex, indicating a prominent role of PDE4 in frontal dopamine receptor signaling. Finally, dopamine is known to be expressed at dopaminergic terminals in neurons of the SNc (Cherry and Davis, 1999), where cAMP has been reported to be a strong inducer of tyrosine hydroxylase (TH) gene transcription rate and mRNA affecting dopamine synthesis (Kumer and Vrana, 1996; Chen et al., 2008). Rolipram enhanced haloperidol-induced phosphorylation of TH at Ser40 in presynaptic dopamine terminals with a proportional increase in dopamine synthesis though failed to do so in the absence of haloperidol. Also, rolipram enhanced levels of 3,4-Dihydroxyphenylacetic acid and the 3,4-Dihydroxyphenylacetic acid/dopamine ratio, indicating an increased dopamine metabolism. However, no increase in the level of dopamine itself was found, indicating the absence of a direct effect on dopamine release (Nishi et al., 2008).

**Implications and Clinical Overview of PDE4-Is**

Compared with PDE1, much more is known regarding the role of PDE4 in frontal and striatal dopaminergic functioning. However, by no means have all the effects been unraveled and all the questions been resolved for PDE4. PDE4 inhibition has been shown to increase dopaminergic tone in striatal neurons by increasing both synthesis and metabolism, though lacking a direct effect on release. It is known that basal ganglia functioning depends on specific amounts of dopamine in order to function at peak performance. Low levels of dopamine cause movement difficulties, while excessive dopamine causes involuntary movements. Even if PDE4 inhibition itself may not be involved in the process of releasing dopamine, the enhanced production of the dopamine precursor levodopa by enhanced cAMP-stimulated TH gene transcription may result in enhanced stimulus-driven dopamine release. PDE4-Is may therefore constitute an interesting treatment for neuropsychiatric disorders involving hypo-functioning striatal dopamine systems, like Parkinson’s disease. Indeed, rolipram has been reported to attenuate MPTP-induced dopamine depletion in the striatum and reduce the loss of nigral TH-positive neurons in vitro (Hulley et al., 1995a; Yamashita et al., 1997a, 1997b) and in vivo (Hulley et al., 1995b; Yang et al., 2008). Next, it remains to be seen if these effects on synthesis and metabolism also apply to frontal dopaminergic terminals arriving from the VTA. If so, PDE4 would be an interesting target for disorders characterized by ventral dopaminergic dysfunction, like ADHD or schizophrenia (cognitive and negative symptoms). Frontal dopaminergic hypo-functioning can not only be opposed by PDE4 inhibition via augmented release of neurotransmitters at dopaminergic terminals but also, like previously discussed for PDE1B, by means of increased DA D1 receptor/cAMP/PKA signaling inducing DARPP-32 and CREB phosphorylation, leading to concomitant gene transcription related to neuronal plasticity. Because PDE4 inhibition affects both these mechanisms, PDE4-Is are particularly interesting as a potential treatment for ADHD and schizophrenia. Finally, in the striatum, PDE4 inhibition regulates adenosine A$_2$ signaling and is therefore often viewed as exerting dopamine D2 antagonistic effects, although it may also be viewed as mimicking the effect of an adenosine A$_3$ agonist. Both would indicate antipsychotic potential of PDE4-Is by counteracting hyperdopaminergia, which has been confirmed by several studies over the years, including early clinical trials (Casacchia et al., 1983; Parkes et al., 1984; Siuciak, 2008). Based on results of rolipram, this latter mechanism is also applicable to Huntington’s disease. Rolipram exerted neuroprotective effects in 2 rodent Huntington’s disease models via increased CREB phosphorylation and subsequent targets like BDNF (DeMarch et al., 2007, 2008; Fusco and Giampa, 2015). Neuroprotective effects of rolipram were induced by sparing of striatal neurons, prevention of intranuclear inclusion formation, and attenuation of microglial reactivity (DeMarch et al., 2008). Furthermore, rolipram was effective in preventing CREB binding protein sequestration into striatal neuronal intranuclear inclusions, sparing interneurons of R6/2 mice and rescuing motor coordination and activity deficits (Giampa et al., 2009b; but see Hannan, 2009). However, there are a range of molecular and cellular mechanisms implicated in the pathogenesis of Huntington’s disease (Gil and Rego, 2008).

When examining actual clinical data, the first clinical trials into PDE4 were done in the field of depression research (Esposito et al., 2009). These first clinical studies showed a good antidepressant response to rolipram treatment (Zeller et al., 1984; Fleischhacker et al., 1992). However, rolipram produces severe dose-limiting side effects, including emesis, headache, gastric hypersecretion, nausea, and vomiting. This has put a serious hold on the further development of rolipram and other related PDE4-Is. It also prevented rolipram from reaching the market. Yet, a clinical Phase II trial started in 2006 to reevaluate the antidepressant properties of rolipram (estimated study completion date: December 2013). No details are yet available to the scientific community. Another PDE4-I, ND1251, was reported to improve memory in a group of 8 depressed subjects (Outsourcing Pharma, 2014).

Although rolipram was primarily developed for treating depression (Zeller et al., 1984; Fleischhacker et al., 1992),
rolipram has also been investigated in early clinical trials as a treatment for Parkinson’s disease (Casaccia et al., 1983; Parkes et al., 1984). Some positive effects of rolipram were observed, however not exceeding the efficacy of levodopa or other dopaminergic drugs. At the moment, second-generation PDE4-Is are being developed, which are supposed to have less-emetic side effects and are being studied for other disorders besides depression. As a result, rolipram was approved by the Food and Drug Administration (FDA) in 2011 as an antiinflammatory drug for the treatment of Chronic Obstructive Pulmonary Disease exacerbations.

PDE4-Is were also tested in clinical trials as treatment for schizophrenia. Takeda has recently finished a proof of mechanism Phase 1 clinical study with the PDE4-I roflumilast in combination with second-generation antipsychotics in schizophrenia patients (ClinicalTrials.gov Identifier: NCT02079844).

PDE4-Is were also examined as a treatment for Huntington’s disease. The new experimental PDE4-I GSK356278 was tested by GlaxoSmithKline as a new treatment for Huntington’s disease in 2 subsequent Phase I studies. In 2012, the first Phase I study was completed investigating the safety, tolerability, pharmacokinetics, and pharmacodynamics of GSK356278 (ClinicalTrials.gov Identifier: NCT01573819). GSK356278 was well tolerated when it was given as a single dose to healthy people, and in this study the objective was to observe effects of GSK356278 after daily intake. Subsequently, a second Phase I positron emission tomography brain occupancy study of GSK356278 was conducted in male healthy volunteers (ClinicalTrials.gov Identifier: NCT01602900; no results are disclosed). It is currently unclear whether this PDE4-I treatment is aimed at the motor or cognitive symptoms observed in Huntington’s disease.

Of note, ibudilast (or AV-411) is another PDE4-I in development as an antiinflammatory drug to treat, for instance, Amyotrophic Lateral Sclerosis (ClinicalTrials.gov Identifier: NCT02238626). However, this compound not only inhibits PDE4 but also serves as a glial activator. Central nervous system applications of AV-411 are being explored in clinical Phase II studies, that is, pain and drug abuse (ClinicalTrials.gov Identifier: NCT00723177, NCT01217970, NCT02025998, NCT01860807).

Additionally, different genetic studies have shown a positive relationship between PDE4B polymorphisms and schizophrenia, which likely results in significantly decreased PDE4B levels as detected in postmortem brain tissue (Fatemi et al., 2008a; Guan et al., 2012). Low PDE4B levels, which might be considered as a compensatory mechanism, do not necessarily result in increased cAMP levels, as several mechanisms can also be activated that counteract the decreased degradation of cAMP by PDE4B. Another genetic link is related to the gene Disrupted-in Schizophrenia-1 (DISC1; Harrison and Weinberger, 2005). A chromosomal translocation of this gene increases susceptibility for schizophrenia (Millar et al., 2000; Sachs et al., 2005), and, interestingly, binding of DISC1 to PDE4B is disrupted, which might result in an overactivity of the latter (Millar et al., 2005; Murdoch et al., 2007).

**PDE10**

**PDE10 and Dopamine Signaling**

PDE10, which is encoded by PDE10A, is a dual substrate PDE, hydrolyzing both cAMP and cGMP. PDE10A is present both in striatonigral direct and striatopallidal indirect pathway MSNs (Xie et al., 2006; Nishi et al., 2008). Additionally, PDE10A regulates cAMP/PKA signaling (Nishi et al., 2008) and gene expression (Strick et al., 2010) in the MSNs of both pathways. Interestingly, PDE10A hydrolyzes both cAMP and cGMP, but it has an approximate 20-fold higher affinity for cAMP (Bender and Beavo, 2006), making it an interesting target for disorders involving the fronto-striatal circuits. In the striatum, PDE10A is expressed in both direct and indirect pathway MSNs, but not in interneurons (Xie et al., 2006; Nishi et al., 2008; Sano et al., 2008). Of the 3 splice variants, PDE10A2 is associated with the membrane, whereas PDE10A1 and PDE10A3 are found in the cytosol (Kotera et al., 2004). In the striatum, mainly PDE10A2 is expressed, and it is found at membranes in dendrites and spines of MSNs (Xie et al., 2006). PDE10A2 is phosphorylated by PKA at Thr16 within the N-terminal region (Kotera et al., 2004). Nishi and colleagues argue that this seems to induce the translocation of PDE10A2 from membrane to cytosol, thereby controlling cAMP/PKA signaling within the spines (Nishi and Snyder, 2010; see also Wilson and Brandon, 2015).

Through this effect on cAMP/PKA signaling, PDE10A inhibition by papaverine showed enhanced phosphorylation of CREB and ERK (Rodefer et al., 2005; Suciak et al., 2006; Becker and Grecksch, 2008) and of their downstream targets DARPP-32 and GluR1 (Nishi et al., 2008) at PKA sites in striatal MSNs both in vitro and in vivo. More specifically, in both direct and indirect pathway neurons, PDE10A shows equal expression patterns (Xie et al., 2006; Nishi et al., 2008; Sano et al., 2008), regulation of cAMP/PKA signaling (Nishi et al., 2008), and gene expression (Strick et al., 2010). However, distinguishing between both pathways, in direct pathway neurons, PDE10A inhibition activates cAMP/PKA signaling related to D1 receptor signaling (Nishi et al., 2008), whereas in indirect pathway neurons, PDE10A inhibition activates cAMP/PKA signaling by simultaneous potentiation of adenosine A1 receptor signaling and inhibition of D2 receptor signaling. A study of neuronal type-specific regulation of DARPP-32 phosphorylation at Thr34 using neostriatal slices showed that papaverine increased DARPP-32 phosphorylation by 6-fold in indirect pathway neurons, whereas it increased DARPP-32 phosphorylation by only 2-fold in direct pathway neurons, indicating that effects of PDE10A inhibition predominate the indirect pathway (Rateup et al., 2008; Nishi et al., 2008). Recent electrophysiological results support this conclusion (Threlfell et al., 2009). More support is provided by recent behavioral studies published by different groups; however, these latter studies also show substantial D1 direct pathway effects of PDE10A-Is (Megens et al., 2014a, 2014b; Gentzel et al., 2015; Suzuki et al., 2016).

In vivo, PDE10A-Is are studied mostly for effects on spontaneous or stimulus behaviors providing evidence for spontaneous indirect pathway effects (similar to effects of D2 receptor blockers). This would include inhibition of spontaneous or stimulant-induced behavior, inhibition of conditioned avoidance behavior, reversal of stimulant-induced sensory gating deficits, and preferential activity against apomorphine-induced climbing (Schmidt et al., 2008; Grauer et al., 2009; Kehler and Nielsen, 2011; Gresack et al., 2013; Megens et al., 2014b). However, concomitant D1 receptor stimulation causes reduced efficiency against behavioral stimulants via direct pathway activation (Menniti et al., 2007; Sotty et al., 2009; Gresack et al., 2013; Megens et al., 2014b). D1 receptor stimulation is also responsible for the cognition-enhancing effects (Rodefer et al., 2005; Grauer et al., 2009) and socializing effects (Grauer et al., 2009) of PDE10A-Is. So indeed, there is compelling evidence for substantial direct pathway activation of PDE10A-Is. This latter notion is supported by recent work from Megens and coworkers (2014a) in suppressed behavior via D1 receptor blockade, D2 receptor blockade or dopamine depletion. Their results indicate...
that PDE10A-Is reverse behavioral suppression after D1 receptor blockade (hypolocomotion) via direct pathway activation (next to suppressing stimulant behavior via indirect pathway activation). These effects are indicative of substantial D1 agonistic effects of PDE10A-Is (next to their D2 antagonistic effects). Still, the main effects of PDE10A-Is are suggested to be exerted through the indirect pathway. By this route, PDE10A-Is can cause extrapyramidal side effects, resembling D2 receptor blockers. The latter may explain why PDE10A-Is have not yet reached the market as antipsychotic treatment. This notion is supported by the recent failure of the Pfizer PDE10A-1 MP-10 (or PF-02545920) in a Phase II clinical trial as antipsychotic treatment, where it showed no efficacy on positive and negative symptoms and produced motor side effects (akathisia and dystonia) in patients with schizophrenia (DeMartinis et al., 2012).

Finally, in contrast to PDE4 inhibition by rolipram, PDE10A inhibition by papaverine showed no increases on TH phosphorylation at Ser40 (PKA site), suggesting no effects of PDE10A inhibitors on dopamine synthesis. Of note, only at high concentrations did papaverine show an effect on TH phosphorylation. Also, results for papaverine should be confirmed by using the more potent PDE10A-Is TP-10 and MP-10. Additionally, PDE10A inhibition showed no effects on dopamine metabolism (Nishi et al., 2008). Therefore, in contrast to PDE4, it is assumed that PDE10A does not play a major role at dopaminergic terminals.

**Implications and Clinical Overview of PDE10A-Is**

PDE10A is even more extensively studied in relation to the frontal-striatal circuits than the previously discussed PDE4 and PDE18 subtypes. Due to the hypothesis of a higher expression in indirect pathway neurons, PDE10A-Is have received much attention as potential dopamine D2 antagonists and as such for their antipsychotic properties. Historically, positive symptoms in schizophrenia have been linked to overstimulation of dopamine receptors in the striatum (Raemder and Francis, 2002), which is attenuated by (PDE10A inhibition-induced) dopamine D2 receptor antagonism. Because of the expected predominant effects in the indirect pathway, PDE10A is also hypothesized as a therapeutic target in Huntington’s disease. Increases in cAMP are expected to drive CREB-dependent signaling pathways, known to be dysregulated in Huntington’s disease mouse models (Choi et al., 2009). In line, like rolipram, TP-10 was shown to be neuroprotective in the quinolinic acid model of Huntington’s disease through CREB-mediated neuroprotection (Giampa et al., 2009a). In a follow-up study, PDE10A-1 treatment of R6/2 mice showed significant delays in development of the motor deficits measured in this model accompanied by reduced striatal and cortical cell loss (Giampa et al., 2010). This was accompanied by increased CREB phosphorylation, suggesting that increased cAMP signaling in these brain regions could slow progression of neurodegeneration. Additionally, gene expression studies have implicated PDE10A and cAMP signaling as a therapeutic strategy for Huntington’s disease (Hebb et al., 2004). Chronic treatment of wild-type mice with TP-10 resulted in an increase in gene expression of members of the ERK and PKA signaling pathways as well as an increase in ERK and MSK phosphorylation (Roze et al., 2008; Kleiman et al., 2011; Martin et al., 2011). Both these effects have proven to be neuroprotective in models of Huntington’s disease. Hypothetically, in Parkinson’s disease, PDE10A-Is could be used in the same way to treat dopamine agonist- or levodopa-induced dyskinesias. Chronic treatment with both classes of drugs leads to improvement in symptoms but causes unwanted side effects. These unwanted symptoms are thought to be due to D1 receptor functional supersensitivity, abnormal cAMP signaling, and enhanced ERK signaling (Bezard et al., 2001; Aubert et al., 2005; Santini et al., 2007). Cyclic nucleotide levels were found to be decreased in the brains of rats treated with a combination of levodopa and 6-OHDA (Giorgi et al., 2008). Consistent with this finding, treatment of levodopa-induced dyskinesias with TP-10 reduced the severity of dyskinesias observed in 6-OHDA rats. In this way, PDE10A-Is rescue decreases in cyclic nucleotide levels and prolong the use of levodopa (Wilson and Brandon, 2015).

Preclinical antipsychotic effects of PDE10A-Is may have initiated fronto-striatal disorder-related research, though lack of clinical efficacy and possible extrapyramidal side effects are hampering PDE10A-Is in reaching the market as antipsychotic treatment. An example of the latter is provided by the failure of the Phase II clinical trial of the Pfizer PDE10A-1 MP-10 (or PF-02545920). MP-10 showed no efficacy and produced motor side effects. Despite the serious challenges, there remains interest in PDE10A-Is as an antipsychotic treatment. For instance, Takeda is currently recruiting participants for a clinical Phase II study to evaluate the efficacy, safety, and tolerability of TAK-063 compared with placebo in treatment of acutely exacerbated schizophrenia. Efficacy was explained as determining whether cognitive impairment associated with schizophrenia would be attenuated (ClinicalTrials.gov Identifier: NCT02477020). Also, a Phase I study by Hoffmann-La Roche has just been completed in which the safety, tolerability, and pharmacokinetics of RO5545965 in patients with schizophrenia on risperidone was tested (no results have been posted; ClinicalTrials.gov Identifier: NCT02019329). Of note, in 2012 Amgen started and terminated a Phase I study to assess the safety and tolerability of their PDE10A-1 AMG 579 following a single oral dose administration in healthy subjects and patients with schizophrenia or stable schizoaffective disorder (ClinicalTrials.gov Identifier: NCT01568203).

A recent study found no difference in PDE10A mRNA expression between schizophrenia patients and comparison subjects in any of the brain regions studied (thalamus, caudate, putamen, nucleus accumbens, globus pallidus, and substantia nigra). This is the first in vivo assessment of PDE10A expression in patients with schizophrenia. However, this should not be interpreted as a case against developing PDE10A drugs in schizophrenia. The study of intracellular signaling pathways makes a persuasive case for how PDE10A-Is could influence the overall signaling in a therapeutic direction, regardless of whether there is an intrinsic change in PDE10A in schizophrenia (Marques et al., 2016).

Pharmaceutical companies have also started to redesignate their PDE10A-Is to Huntington’s disease. A Phase II proof-of-concept trial is now being initiated in which Pfizer’s PDE10A-1 MP-10 will be tested for safety and efficacy in subjects with Huntington’s disease (ClinicalTrials.gov Identifier: NCT02197130). Omeros initiated a Phase II clinical trial in Huntington’s disease patients with OMS824 after an earlier Phase II trial in schizophrenia patients (no results disclosed; ClinicalTrials.gov Identifier: NCT01952132). The Huntington’s disease trial is a sequential-cohort dose escalation study that evaluates the safety and tolerability of OMS824 over 4 weeks (ClinicalTrials.gov Identifier: NCT02074410). In parallel with the clinical OMS824 trial, Omeros is conducting preclinical rat studies to support clinical trials of longer duration. However, based on that data, there might be a safety issue and based on follow-up communications with the FDA, Omeros has suspended the ongoing Huntington’s disease trial. The FDA has requested that Omeros further evaluates the preclinical data in order to characterize the compound more fully prior to reinitiating the clinical trial (Omeros, 2014). Additional support for the
use of PDE10A-Is in Huntington’s disease comes from a recent study that shows that PDE10A levels are lowered early before symptom onset in Huntington’s disease (Niccolini et al., 2015b). Whether this is cause or consequence remains to be determined; however, it most likely resembles a consequence of the degeneration of striatal cells and therefore the PDE10A enzymes within. These results were recently confirmed by a study with the radioligand [18F] MNI-659A (Russell et al., 2016). A comparable large-scale Phase 0 study is currently recruiting new participants. The aim of this study is to measure the availability of the PDE10A enzyme in Huntington’s disease gene expansion carriers using the recently developed radioligand [18F] MNI-659. The study will be cross-sectional, examining Huntington’s disease gene expansion carriers at different stages of the disease (premanifest, stage 1, and stage 2) compared with healthy controls (ClinicalTrials.gov Identifier: NCT02061722).

Of note, Niccolini et al. (2015a) also demonstrated striatal and pallidal loss of PDE10A expression in Parkinson’s disease patients, which is associated with Parkinson’s disease duration and severity of motor symptoms and complications. These results suggest that dopaminergic nigrostriatal degeneration affects the expression of PDE10A in striatum and pallidum. Hypothesizing, it most likely resembles a compensatory mechanism. Less dopaminergic input from the SNc equals less cAMP activation in striatal and pallidal areas decreasing the required levels of PDE10A. In another, more plausible scenario, the decrease is causative. The decrease in PDE10A levels reflects the overall expression of PDE10A in these brain areas not specified for the direct and indirect pathway. Because of the stronger expression of PDE10A in the indirect pathway compared with the direct pathway, PDE10A degeneration will affect the indirect pathway more strongly. Subsequently, reduced PDE10A expression results in enhanced activation of the indirect pathway, resulting in increased inhibition of movement. In both Parkinson’s disease and Huntington’s disease, altered PDE10A levels are likely compensatory/consequential instead of causative. In hyperkinetic movement disorders like Huntington’s disease, PDE10A may thus be a promising target for pharmacological agents (PDE10A-Is enhance the little cAMP signaling that is left in the indirect pathway).

Conclusion

Clinical trials investigating the effects of PDE-Is in neuropsychiatric disorders are overall very sparse, and the wealth of positive preclinical data could not yet be translated into clinical efficacy. As a result, no definitive conclusions can be drawn merely based on clinical trial outcomes. Therefore, the current review provides a discussion of the role of PDEs in dopaminergic frontal and striatal signaling and the potential of their associated inhibitors in specific disorders of the fronto-striatal circuits. Subsequently, an overview is provided of the current clinical status.

The fronto-striatal circuits compose the neurobiological basis for several neuropsychiatric disorders, including Parkinson’s disease, Huntington’s disease, ADHD, Tourette’s syndrome, schizophrenia, and obsessive-compulsive disorder. The fronto-striatal circuits constitute a plurality of parallel segregated circuits, which can be clustered together in motor circuits, associative/cognitive circuits, and limbic circuits (Krack et al., 2010). Together, dysfunctions in these circuits produce the wide range of symptoms observed in related neuropsychiatric disorders.

Intracellularly, direct and indirect pathway signaling in the striatum is largely mediated through the cAMP/PKA cascade (Nishi et al., 2008, 2011; Nishi and Snyder, 2010). Cyclic nucleotide cascades are involved in synaptic transmission, neuron excitability, neuroplasticity, and neuroprotection in all types of fronto-striatal circuits (Figure 1). Additionally, all fronto-striatal circuits are modulated by dopamine. Next to the effects of cAMP/PKA pathways on glutamatergic and GABAergic signaling in the fronto-striatal circuits, these cyclic nucleotide pathways also play a major role in the dopaminergic modulation of the circuits. The intracellular effect of dopamine is mediated through dopamine receptor-regulated activation of cAMP/PKA and subsequent DARPP-32 and CREB phosphorylation in both striatal and frontal neurons.

In the last decades, PDEs have therefore received increased attention for their possible role in disorders involving the fronto-striatal circuits. Based on overall expression patterns in frontal and striatal dopaminergic terminals, indirect pathway neurons, and direct pathway neurons, PDE1B, PDE2A, PDE4, PDE7B, PDE9A, and PDE10A seem to be the most interesting targets (Lakics et al., 2010), although most attention and resources have thus far been devoted to the potential of PDE1B, PDE4, and PDE10A due to their role in dopaminergic signaling. The main site of action and expression of PDE1B, PDE4, and PDE10A as discussed in this clinical review is inferred from biochemical analyses of striatal cAMP/PKA effectors, behavioral phenotypes of knockout mice, and the observation of effects of subtype-specific PDE-Is on dopamine-related behavior. The different PDE subtypes, and more specifically their splice variants, can be related to different disorders due to their differential expression in one or more of the frontal and striatal pathways or dopaminergic terminals inducing stimulation of dopamine synthesis, the inhibition of D2 receptor signaling or the stimulation of D1 receptor signaling. The different PDE isoforms contain a multiplicity of structural and biochemical properties and are located in specific subcellular compartments, with specific transcriptional and posttranscriptional regulation (Keravis and Lugnier, 2012). Therefore, expression of a PDE subtype in a brain area does not make it an interesting target per se. Their particular involvement in dopaminergic modulation of fronto-striatal signaling is what makes them an interesting target for related disorders. Preferably, the targeted cyclic nucleotide signaling cascade is involved in the pathology of the disorder or contributes to the reduction of the pathology. However, even if this is not the case, PDE inhibition could still influence the overall signaling in a therapeutic direction. Currently, researchers are just beginning to unravel the precise subcellular localization and the role of functional compartmentalization in physiological and pathological conditions of the fronto-striatal circuits (e.g., PDE10A: Russ wurm et al., 2015; Li et al., 2016b; MacMullen et al., 2016). Another important consideration is that, in general, PDE-I research involving fronto-striatal disorders is based on the classical view of basal ganglia direct and indirect pathway functioning. Considerable evidence is accumulating to challenge this classical view (Cui et al., 2013; Calabresi et al., 2014; Keeler et al., 2014).

From a therapeutic perspective, inhibition of PDEs with increased expression appears most promising. This way, cognition and plasticity deficits resulting from impaired cAMP/PKA signaling might be improved by inhibiting specific PDE isoforms. However, PDE inhibition might have negative effects on cognition and plasticity when PDEs are already downregulated and AMP levels and PKA activity are high. In this scenario, elevated AMP levels might go over a physiological level and disrupt signaling. Along this line, high doses of rolipram impaired prefrontal cognitive function in aged but not young monkeys, likely due to overstimulation of the already disinhibited cAMP/PKA...
signaling pathway in the aged prefrontal cortex (Ramos et al., 2003; Arnsten et al., 2005). This argues to specifically target PDEs that are overexpressed (Table 2).

In addition to the backbone formed by frontal neurons, MSNs and their dopaminergic modulation, the importance of interneurons in physiological and pathological fronto-striatal functioning is becoming increasingly apparent. Several types of interneurons can be found in the striatum, like cholinergic and different GABAergic interneurons (Gerfen and Surmeier, 2011). In particular, nitric oxide synthase containing GABAergic interneurons we would like to highlight. These nitric oxide-producing interneurons play an important role in fronto-striatal functioning (West and Tseng, 2011). Nitric oxide diffuses from these interneurons into dendrites of MSNs that contain high levels of guanylate cyclase, which, when activated, lead to the synthesis of cGMP (Figure 1). In the intact striatum, transient elevations in intracellular cGMP primarily act to increase neuronal excitability and to facilitate glutamatergic fronto-striatal transmission (West and Tseng, 2011; Threlfell and West, 2013).

Although the main focus in the fronto-striatal system has been on cAMP signaling, several PDE-Is (also) target cGMP (e.g., PDE1-Is and PDE10-Is) and may exert their effects (additionally) on the cGMP signaling cascade (Padovan-Neto et al., 2015).

Summarizing, increased understanding of the subcellular localization and unraveling of the signalosome concept of PDEs including its function and dysfunction in the fronto-striatal circuits will contribute to the design of new specific inhibitors and enhance the potential of PDE-Is as therapeutics in fronto-striatal circuits.

Acknowledgments

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Statement of Interest

L.W. is an employee of Intra-Cellular Therapies, which has a financial interest in the PDE1 inhibitor ITI-214. M.A.v.D., A.B., and J.P. have a proprietary interest in the PDE4 inhibitor roflumilast.

Table 2. Changes in Human Phosphodiesterase mRNA Levels in Several Fronto-Striatal Disorders

<table>
<thead>
<tr>
<th>Phosphodiesterase</th>
<th>Schizophrenia</th>
<th>Parkinson’s Disease</th>
<th>Huntington’s Disease</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cerebellum</td>
<td>Striatum</td>
<td>Pallidum</td>
</tr>
<tr>
<td>PDE1</td>
<td>+ (1C)</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>PDE2</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>PDE3</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>PDE4</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td>(4A1, 4A5, 4A8)</td>
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<td>NS</td>
</tr>
<tr>
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<td>(4B1, 4B2, 4B3, 4B4)</td>
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<tr>
<td></td>
<td>(10A)</td>
<td>(10A)</td>
<td>(10A)</td>
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<tr>
<td>PDE11</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
</tbody>
</table>

+, increased; -, decreased; =, no change; NS, not studied; (Fatemi et al., 2008a, 2008b, 2010; Niccolini et al., 2015a, 2015b; Marques et al., 2016; Russell et al., 2016).

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