Editorial

The behavioural neurology of basal ganglia disorders

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Basal ganglia disorders are common clinical conditions characterised by specific motor dysfunction and associated behavioral problems. The underlying pathophysiology primarily consists in alterations of the cortico-striato-thalamo-cortical pathways, which can be related to both neurodegenerative pathology (e.g. Parkinson disease, Huntington disease) and neurodevelopmental processes (e.g. Tourette syndrome, dystonia). Modulation of dopaminergic neurotransmission within the basal ganglia circuitries represents the mainstay of pharmacological treatment for these conditions. This thematic issue on “The behavioural neurology of basal ganglia disorders” explores the pathophysiological mechanisms underlying the most common basal ganglia disorders, with special focus on the neurochemical targets for pharmacotherapy.

The introductory article (Ward et al.) presents the functional neuroanatomy and behavioral correlates of basal ganglia as demonstrated by case studies of patients with selective lesions affecting the cortico-basal pathways. Lesions involving the caudate nucleus, putamen and globus pallidus have been found to be associated with a spectrum of behavioral and cognitive symptoms, including abulia, poor working memory and

deficits in emotional recognition. The paper by Alobaidi and Pall focuses on the pharmacotherapy of Parkinson disease, an area which is often challenging as clinicians have to find a delicate balance between the efficacy on motor symptoms and the side effects profiles of different dopaminergic medications aimed at restoring basal ganglia activity. This article assesses the available evidence on the role of dopamine agonist monotherapy as an alternative to Levodopa in the treatment of the motor symptoms of Parkinson disease, along with the role of dopamine antagonists in the treatment of Parkinson disease-related psychosis. A related area of high interest over the last few years has been the pathophysiology of impulse control disorders in patients with Parkinson disease: this topic is covered by the following article (Balarajah and Cavanna), along with the implications of these behavioral symptoms for the pharmacological management of Parkinson disease. Another neurodegenerative disorder at the interface between neurology and psychiatry is Huntington disease. The neuropathological process of this condition also involves the basal ganglia, and has been solidly anchored to genetic bases. The paper by Pidgeon and Rickards covers the current evidence supporting the pharmacological treatment for both motor and non-motor symptoms of Huntington disease, whilst Mittal and Eddy's contribution focuses on the pharmacological modulation of dopaminergic and glutamatergic pathways, which are crucially affected in this condition. The article by Waldon et

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al. offers an overview of the evidence-based treatment of tic symptoms in Tourette syndrome, a quintessentially neuropsychiatric basal ganglia disorder, in which challenging behavioural problems often accompany the motor and vocal manifestations. The paper by Patel and Martino presents a systematic review on the evidence-based pharmacotherapy for cervical dystonia, whilst the closing article by Jones and Cavanna focuses on the pathophysiology and treatment of a common and often under-recognized condition, restless legs syndrome.

It is the editor's hope that this timely issue on basal ganglia disorders will prove to be a useful resource for both basic and clinical researchers, plus neuropsychiatrists with a special interest in movement disorders. Finally, this issue is proposed as an up-to-date reference for clinicians opting for the pharmacological modulation of what Ivan Pavlov intuitively called "the blind force of the subcortex".