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Translated by Pharmaceuticals and Medical Devices Agency



This English version is intended to be a reference material to provide convenience for users. In the event of inconsistency between the Japanese original and this

English translation, the former shall prevail.

Revision of Precautions Pramipexole hydrochloride hydrate (conventional tablets, OD tablets)

August 22, 2019

Therapeutic category Antiparkinsonian agents Central nervous system agents-miscellaneous

Non-proprietary name Pramipexole hydrochloride hydrate

Safety measure

Precautions should be revised in the package insert.

Pharmaceuticals and Medical Devices Agency

3-3-2 Kasumigaseki, Chiyoda-ku, Tokyo 100-0013 Japan E-mail: <u>safety.info@pmda.go.jp</u> Revision in line with the Instructions for Package Inserts of Prescription Drugs, PAB Notification No. 606 by the Director General ofPharmaceutical Affairs Bureau, MHW, dated April 25, 1997 (Old instructions):Revised language is underlined.

Current	Revision
Important Precautions	Important Precautions
Rapid dose reduction or discontinuation of this drug in patients with	When dose reduction or discontinuation of this drug is necessary in
Parkinson's disease may cause syndrome malin. When dose	patients with Parkinson's disease, the dose should be gradually
reduction or discontinuation is necessary, the dose should be	reduced. Rapid dose reduction or discontinuation may cause
gradually reduced.	syndrome malin. In addition, rapid dose reduction or
It should be noted that gradual dose reduction is not necessary in	discontinuation of dopamine receptor agonists may cause drug
patients with idiopathic restless legs syndrome because their doses	withdrawal syndrome (characterized by apathy, anxiety, depression,
are lower than for patients with Parkinson's disease.	fatigue, sweating, pain, etc.).
	It should be noted that gradual dose reduction is not necessary in
	patients with idiopathic restless legs syndrome because their doses
	are lower than for patients with Parkinson's disease.
Adverse Reactions	Adverse Reactions
Other Adverse Reactions	Other Adverse Reactions
(N/A)	Drug withdrawal syndrome* (apathy, anxiety, depression, fatigue,
	sweating, pain, etc.)
	*When any abnormalities are observed, appropriate measures
	should be taken such as resuming administration or returning the
	dose to the level prior to reduction.

N/A: Not Applicable, because the section is not included in the current package insert.

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