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62

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 Iopamidol

July 20, 2020

**Therapeutic category** 

X-ray contrast media

## Non-proprietary name

lopamidol

## Safety measure

Precautions should be revised in the package insert.

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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
Adverse Reactions	Adverse Reactions
Clinically Significant Adverse Reactions	Clinically Significant Adverse Reactions
(N/A)	Contrast-induced encephalopathy:
	Disturbed consciousness, paralysis, aphasia, cortical blindness, or
	other central nervous system symptoms may occur as a result of
	extracerebrovascular leakage of this drug in cerebral angiography,
	angiocardiography (including pulmonary artery angiography), and
	aortography. Minimum effective dosages should be administered
	and appropriate measures should be taken if any abnormalities are
	observed.

Revision in line with the Instructions for Package Inserts of Prescription Drugs, etc. PSEHB Notification No. 0608-1 by the Director of

Pharmaceutical Safety and Environmental Health Bureau, MHLW, dated June 8, 2017 (New instructions): Revised language is underlined.

Current	Revision
11. ADVERSE REACTIONS	11. ADVERSE REACTIONS
11.1 Clinically Significant Adverse Reactions	11.1 Clinically Significant Adverse Reactions
(N/A)	(Cerebral angiography, angiocardiography (including pulmonary
	artery angiography), and aortography)
	Contrast-induced encephalopathy
	Disturbed consciousness, paralysis, aphasia, cortical blindness, or
	other central nervous system symptoms may occur as a result of

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extracerebrovascular leakage of this drug. Minimum effective
dosages should be administered and appropriate measures should
be taken if any abnormalities are observed.

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

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