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*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

## Pomalidomide

January 26, 2021

### **Therapeutic category**

Other antitumor agents

### **Non-proprietary name**

Pomalidomide

### **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: [safety.info@pmda.go.jp](mailto:safety.info@pmda.go.jp)

Revision in line with the Instructions for Package Inserts of Prescription Drugs, PAB Notification No. 606 by the Director General of Pharmaceutical Affairs Bureau, MHW, dated April 25, 1997 (Old instructions): Revised language is underlined.

Current	Revision
<p>Adverse Reactions</p> <p>Clinically Significant adverse Reactions</p> <p>(N/A)</p>	<p>Adverse Reactions</p> <p>Clinically Significant adverse Reactions</p> <p><u>Progressive multifocal leukoencephalopathy (PML):</u></p> <p><u>Progressive multifocal leukoencephalopathy (PML) may occur.</u></p> <p><u>Patients should be carefully monitored during, and after, the treatment with this drug, and if symptoms such as disturbed consciousness, cognitive disorder, paralytic symptoms (hemiplegia or quadriplegia), dyslalia, or speech loss are observed, diagnostic imaging through MRI and cerebrospinal fluid test should be performed, administration should be discontinued, and appropriate measures should be taken.</u></p>

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 E-mail: [safety.info@pmda.go.jp](mailto:safety.info@pmda.go.jp)

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
<p>11. ADVERSE REACTIONS</p> <p>11.1 Clinically Significant adverse Reactions</p> <p>(N/A)</p>	<p>11. ADVERSE REACTIONS</p> <p>11.1 Clinically Significant adverse Reactions</p> <p><u>Progressive multifocal leukoencephalopathy (PML)</u></p> <p><u>Patients should be carefully monitored during, and after, the treatment with this drug, and if symptoms such as disturbed consciousness, cognitive disorder, paralytic symptoms (hemiplegia or quadriplegia), dyslalia, or speech loss are observed, diagnostic imaging through MRI and cerebrospinal fluid test should be performed, administration should be discontinued, and appropriate measures should be taken.</u></p>

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

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