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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

## Sumatriptan

### Sumatriptan succinate (oral dosage form)

### Sumatriptan succinate (injectable dosage form, ampules)

June 4, 2019

#### **Therapeutic category**

Vasoconstrictors

#### **Non-proprietary name**

Sumatriptan, Sumatriptan succinate

#### **Safety measure**

Precautions should be revised in the package insert.

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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>Important Precautions (N/A)</p>	<p>Important Precautions <u>Triptans including this drug may lead to an exacerbation of headache. If headache does not improve, consider a possibility of “medication overuse headache” and appropriate measures such as discontinuation of administration should be taken.</u></p>
<p>Adverse Reactions Clinically Significant Adverse Reactions (N/A)</p>	<p>Adverse reactions Clinically significant Adverse Reactions <u>Medication overuse headache may occur. If any abnormalities are observed, appropriate measures such as discontinuation of administration should be taken.</u></p>

(Reference) International Headache Society:Cephalalgia 2018;38(1):1-211

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

\* This drug is designated as a drug requiring preparation of a Drug Guide for Patients.

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