

September 7, 2018

To whom it may concern

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**Issuance of Notification on the Extension of the Reexamination Period for the
Hyperuricemia and Gout Treatment FEBURIC® (febuxostat)**

Tokyo, Japan, September 7, 2018 --- Teijin Limited announced today that a notification on the extension of the reexamination period for the hyperuricemia and gout treatment FEBURIC® (febuxostat) of Teijin Pharma Limited, a subsidiary of Teijin, was issued on September 4 by the Director of the Pharmaceutical Evaluation Division, Pharmaceutical Health and Safety Bureau, Ministry of Health, Labour and Welfare. Details are as follows.

Name of pharmaceutical : FEBURIC Tablet 10 mg, FEBURIC Tablet 20 mg, FEBURIC Tablet 40 mg
Extended reexamination period : Until January 20, 2021
Holder of approval : Teijin Pharma Limited
Reasons : It was recognized that there is a need to implement trials to set the dosage and administration for pediatric patients and assess efficacy and safety in the pediatric population.

This matter will have a negligible impact on the consolidated full-term operating results forecasts for the fiscal year ending March 31, 2019.