

Resolution

of the Federal Joint Committee on a Finding in the Procedure of Routine Practice Data Collection and Evaluations according to Section 35a, paragraph 3b SGB V:

Fedratinib (myelofibrosis) – Study protocol and statistical analysis plan submission

of 1 June 2023

At its session on 1 June 2023, the Federal Joint Committee (G-BA) decided the following in the procedure of routine practice data collection and evaluations according to Section 35a, paragraph 3b SGB V for the active ingredient fedratinib (myelofibrosis):

- I. The routine practice data collection will not be carried out as that the pharmaceutical company has not fulfilled its obligation - by resolution of 3 November 2022 - to prepare a statistical analysis plan and study protocol prior to the implementation of the routine practice data collection.

- II. The resolution will enter into force on the day of its publication on the website of the G-BA on 1 June 2023.

The justification to this resolution will be published on the website of the G-BA at www.g-ba.de.

Berlin, 1 June 2023

Federal Joint Committee (G-BA)
in accordance with Section 91 SGB V
The Chair

Prof. Hecken