Resolution

of the Federal Joint Committee on an Amendment of the Pharmaceuticals Directive:
Annex XII – Benefit Assessment of Medicinal Products with New Active Ingredients according to Section 35a (SGB V)
Belantamab mafodotin (repeal of the resolution of 5 October 2023)

of 4 April 2024

At its session on 4 April 2024, the Federal Joint Committee (G-BA) resolved to amend the Pharmaceuticals Directive (AM-RL) in the version dated 18 December 2008 / 22 January 2009 (Federal Gazette, BAnz. No. 49a of 31 March 2009), as last amended by the publication of the resolution of D Month YYYY (Federal Gazette, BAnz AT DD.MM.YYYY BX), as follows:

I. **The findings on the benefit assessment of the active ingredient Belantamab mafodotin in Annex XII of the Pharmaceuticals Directive in the version of the resolution of 5 October 2023 (BAnz AT 21.12.2023 B5) are repealed.**

II. **The resolution will enter into force on the day of its publication on the website of the G-BA on 4 April 2024.**

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

Berlin, 4 April 2024

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

Prof. Hecken