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► **B** COMMISSION IMPLEMENTING DECISION

of 22 November 2012

establishing a list of third countries with a regulatory framework applicable to active substances for medicinal products for human use and the respective control and enforcement activities ensuring a level of protection of public health equivalent to that in the Union, in accordance with Directive 2001/83/EC of the European Parliament and of the Council

(Text with EEA relevance)

(2012/715/EU)

(OJ L 325, 23.11.2012, p. 15)

Amended by:

		Official Journal		
		No	page	date
► <b><u>M1</u></b>	Commission Implementing Decision 2013/196/EU of 24 April 2013	L 113	22	25.4.2013
► <b><u>M2</u></b>	Commission Implementing Decision 2013/262/EU of 4 June 2013	L 152	52	5.6.2013
► <b><u>M3</u></b>	Commission Implementing Decision 2013/301/EU of 11 June 2013	L 169	71	21.6.2013
► <b><u>M4</u></b>	Commission Implementing Decision (EU) 2015/1057 of 1 July 2015	L 171	23	2.7.2015
► <b><u>M5</u></b>	Commission Implementing Decision (EU) 2019/769 of 14 May 2019	L 126	70	15.5.2019
► <b><u>M6</u></b>	Commission Implementing Decision (EU) 2023/172 of 24 January 2023	L 24	37	26.1.2023
► <b><u>M7</u></b>	Commission Implementing Decision (EU) 2023/2484 of 9 November 2023	L 2484	1	10.11.2023

Corrected by:

► **C1** Corrigendum, OJ L 90103, 17.11.2023, p. 1 (2023/2484)

**▼B**

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**(2012/715/EU)**

*Article 1*

The list of third countries referred to in Article 111b(1) of Directive 2001/83/EC is set out in the Annex to this Decision.

*Article 2*

This Decision shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

