

DECISION OF THE EEA JOINT COMMITTEE

No 191/2013

of 8 November 2013

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement

THE EEA JOINT COMMITTEE,

as amended by:

Having regard to the Agreement on the European Economic Area ("the EEA Agreement"), and in particular Article 98 thereof,

— **32013 D 0262**: Commission Implementing Decision 2013/262/EU of 4 June 2013 (OJ L 152, 5.6.2013, p. 52).'

Article 2

Whereas:

The text of Implementing Decision 2013/262/EU in the Icelandic and Norwegian languages, to be published in the EEA Supplement to the *Official Journal of the European Union*, shall be authentic.

(1) Commission Implementing Decision 2013/262/EU of 4 June 2013 amending Implementing Decision 2012/715/EU 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 ⁽¹⁾ is to be incorporated into the EEA Agreement.

Article 3

This Decision shall enter into force on 9 November 2013, provided that all the notifications under Article 103(1) of the EEA Agreement have been made ^(*), or on the day of the entry into force of Decision of the EEA Joint Committee No 162/2013 of 8 October 2013 ⁽²⁾, whichever is the later.

(2) Annex II to the EEA Agreement should therefore be amended accordingly,

Article 4

This Decision shall be published in the EEA Section of, and in the EEA Supplement to, the *Official Journal of the European Union*.

HAS ADOPTED THIS DECISION:

Done at Brussels, 8 November 2013.

Article 1

The following is added in point 15qb (Commission Implementing Decision 2012/715/EU) of Chapter XIII of Annex II to the EEA Agreement:

*For the EEA Joint Committee**The President*

Thórir IBSEN

⁽¹⁾ OJ L 152, 5.6.2013, p. 52.

^(*) No constitutional requirements indicated.

⁽²⁾ OJ L 58, 27.2.2014, p. 15.