

## DECISION OF THE EEA JOINT COMMITTEE

No 163/2013

of 8 October 2013

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

THE EEA JOINT COMMITTEE,

*Article 2*

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

The texts of Regulation (EU) No 1027/2012 in the Icelandic and Norwegian languages, to be published in the EEA Supplement to the *Official Journal of the European Union*, shall be authentic.

Whereas:

*Article 3*

(1) Regulation (EU) No 1027/2012 of the European Parliament and of the Council of 25 October 2012 amending Regulation (EC) No 726/2004 as regards pharmacovigilance<sup>(1)</sup> is to be incorporated into the EEA Agreement.

This Decision shall enter into force on 9 October 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 158/2013 of 8 October 2013<sup>(2)</sup>, whichever is the later.

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

*Article 4*

HAS ADOPTED THIS DECISION:

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

*Article 1*

The following indent is added in point 15zb (Regulation (EC) No 726/2004 of the European Parliament and of the Council) of Chapter XIII of Annex II to the EEA Agreement:

Done at Brussels, 8 October 2013.

— **32012 R 1027**: Regulation (EU) No 1027/2012 of the European Parliament and of the Council of 25 October 2012 (OJ L 316, 14.11.2012, p. 38).'

For the EEA Joint Committee

The President

Thórir IBSEN

<sup>(1)</sup> OJ L 316, 14.11.2012, p. 38.

<sup>(\*)</sup> No constitutional requirements indicated.

<sup>(2)</sup> See page 10 of this Official Journal.