

## DECISION OF THE EEA JOINT COMMITTEE

No 159/2013

of 8 October 2013

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

THE EEA JOINT COMMITTEE,

— **32011 L 0062**: Directive 2011/62/EU of the European Parliament and of the Council of 8 June 2011 (OJ L 174, 1.7.2011, p. 74).'

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

*Article 2*

The text of Directive 2011/62/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.

Whereas:

*Article 3*

(1) Directive 2011/62/EU of the European Parliament and of the Council of 8 June 2011 amending Directive 2001/83/EC on the Community code relating to medicinal products for human use, as regards the prevention of the entry into the legal supply chain of falsified medicinal products<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 to the EEA Joint Committee (\*).

(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 October 2013.

*Article 1*

The following indent is added in point 15q (Directive 2001/83/EC of the European Parliament and of the Council) of Chapter XIII of Annex II to the EEA Agreement:

*For the EEA Joint Committee**The President*

Thórir IBSEN

<sup>(1)</sup> OJ L 174, 1.7.2011, p. 74.

<sup>(\*)</sup> Constitutional requirements indicated.