

## DECISION OF THE EEA JOINT COMMITTEE

No 85/2012

of 30 April 2012

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

THE EEA JOINT COMMITTEE,

HAS ADOPTED THIS DECISION:

*Article 1*

Having regard to the Agreement on the European Economic Area, as amended by the Protocol adjusting the Agreement on the European Economic Area, hereinafter referred to as 'the Agreement', and in particular Article 98 thereof,

The following indents shall be added in point 13 (Commission Regulation (EU) No 37/2010) of Chapter XIII of Annex II to the Agreement:

Whereas:

— **32010 R 0758**: Commission Regulation (EU) No 758/2010 of 24 August 2010 (OJ L 223, 25.8.2010, p. 37),

(1) Annex II to the Agreement was amended by Decision of the EEA Joint Committee No 47/2012 of 30 March 2012 <sup>(1)</sup>.

— **32010 R 0759**: Commission Regulation (EU) No 759/2010 of 24 August 2010 (OJ L 223, 25.8.2010, p. 39),

(2) Commission Regulation (EU) No 758/2010 of 24 August 2010 amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance valnemulin <sup>(2)</sup> is to be incorporated into the Agreement.

— **32010 R 0761**: Commission Regulation (EU) No 761/2010 of 25 August 2010 (OJ L 224, 26.8.2010, p. 1).'

*Article 2*

The texts of Regulations (EU) No 758/2010, (EU) No 759/2010 and (EU) No 761/2010 in the Icelandic and Norwegian languages, to be published in the EEA Supplement to the *Official Journal of the European Union*, shall be authentic.

*Article 3*

(3) Commission Regulation (EU) No 759/2010 of 24 August 2010 amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance tildipirosin <sup>(3)</sup> is to be incorporated into the Agreement.

This Decision shall enter into force on 1 May 2012, provided that all the notifications under Article 103(1) of the Agreement have been made to the EEA Joint Committee (\*).

*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*.

(4) Commission Regulation (EU) No 761/2010 of 25 August 2010 amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance methylprednisolone <sup>(4)</sup> is to be incorporated into the Agreement,

Done at Brussels, 30 April 2012.

For the EEA Joint Committee

The Acting President

Gianluca GRIPPA

<sup>(1)</sup> OJ L 207, 2.8.2012, p. 27.

<sup>(2)</sup> OJ L 223, 25.8.2010, p. 37.

<sup>(3)</sup> OJ L 223, 25.8.2010, p. 39.

<sup>(4)</sup> OJ L 224, 26.8.2010, p. 1.

(\*) No constitutional requirements indicated.