DECISION OF THE EEA JOINT COMMITTEE

No 18/2016

of 5 February 2016

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2017/1301]

THE EEA JOINT COMMITTEE,

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

Whereas:

(1) Commission Implementing Regulation (EU) 2015/1981 of 4 November 2015 approving Formaldehyde released from N,N-Methylenebismorpholine as an existing active substance for use in biocidal products for product-types 6 and 13 ([1]) is to be incorporated into the EEA Agreement.

(2) Commission Implementing Regulation (EU) 2015/1982 of 4 November 2015 approving hexaflumuron as an existing active substance for use in biocidal products for product-type 18 ([2]) is to be incorporated into the EEA Agreement.

(3) Commission Implementing Decision (EU) 2015/1985 of 4 November 2015 pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on an anti-viral tissue impregnated with citric acid ([3]) is to be incorporated into the EEA Agreement.

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

HAS ADOPTED THIS DECISION:

Article 1

The following points are inserted after point 12nns (Commission Implementing Regulation (EU) 2015/1759) of Chapter XV of Annex II to the EEA Agreement:

'12nnt. **32015 R 1981**: Commission Implementing Regulation (EU) 2015/1981 of 4 November 2015 approving Formaldehyde released from N,N-Methylenebismorpholine as an existing active substance for use in biocidal products for product-types 6 and 13 (OJ L 289, 5.11.2015, p. 9).

12nnu. **32015 R 1982**: Commission Implementing Regulation (EU) 2015/1982 of 4 November 2015 approving hexaflumuron as an existing active substance for use in biocidal products for product-type 18 (OJ L 289, 5.11.2015, p. 13).

12nnv. **32015 D 1985**: Commission Implementing Decision (EU) 2015/1985 of 4 November 2015 pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on an anti-viral tissue impregnated with citric acid (OJ L 289, 5.11.2015, p. 26).'

Article 2

The texts of Implementing Regulations (EU) 2015/1981 and (EU) 2015/1982 and Implementing Decision (EU) 2015/1985 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]) OJ L 289, 5.11.2015, p. 9.
([2]) OJ L 289, 5.11.2015, p. 13.
([3]) OJ L 289, 5.11.2015, p. 26.

Article 3

This Decision shall enter into force on 6 February 2016, provided that all the notifications under Article 103(1) of the EEA Agreement have been made (*).

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*.

Done at Brussels, 5 February 2016.

For the EEA Joint Committee
The President
Claude MAERTEN

(*) No constitutional requirements indicated.