DECISION OF THE EEA JOINT COMMITTEE

No 218/2014

of 24 October 2014

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2015/1442]

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) Regulation (EU) No 334/2014 of the European Parliament and of the Council of 11 March 2014 amending Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products, with regard to certain conditions for access to the market ([1]) is to be incorporated into the EEA Agreement.

(2) Commission Implementing Regulation (EU) No 405/2014 of 23 April 2014 approving lauric acid as an existing active substance for use in biocidal products for product-type 19 ([2]) is to be incorporated into the EEA Agreement.

(3) Commission Implementing Regulation (EU) No 406/2014 of 23 April 2014 to approve ethyl butylacetylaminopropionate as an existing active substance for use in biocidal products for product-type 19 ([3]) is to be incorporated into the EEA Agreement.

(4) Commission Implementing Regulation (EU) No 407/2014 of 23 April 2014 approving transfluthrin as an existing active substance for use in biocidal products for product-type 18 ([4]) is to be incorporated into the EEA Agreement.

(5) Commission Implementing Regulation (EU) No 408/2014 of 23 April 2014 approving synthetic amorphous silicon dioxide as an existing active substance for use in biocidal products for product-type 18 ([5]) is to be incorporated into the EEA Agreement.

(6) Commission Implementing Regulation (EU) No 437/2014 of 29 April 2014 approving 4,5-Dichloro-2-octyl-2H-isothiazol-3-one as an existing active substance for use in biocidal products for product-type 21 ([6]) is to be incorporated into the EEA Agreement.

(7) Commission Implementing Regulation (EU) No 438/2014 of 29 April 2014 approving cyproconazole as an existing active substance for use in biocidal products for product-type 8 ([7]) is to be incorporated into the EEA Agreement.

(8) Commission Implementing Decision 2014/227/EU of 24 April 2014 on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council ([8]) is to be incorporated into the EEA Agreement.

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

HAS ADOPTED THIS DECISION:

Article 1

Chapter XV of Annex II to the EEA Agreement shall be amended as follows:

(1) the following indent is added in point 12n (Regulation (EU) No 528/2012 of the European Parliament and of the Council):

'— **32014 R 0334**: Regulation (EU) No 334/2014 of the European Parliament and of the Council of 11 March 2014 (OJ L 103, 5.4.2014, p. 22).';

([1]) OJ L 103, 5.4.2014, p. 22.
([2]) OJ L 121, 24.4.2014, p. 8.
([3]) OJ L 121, 24.4.2014, p. 11.
([4]) OJ L 121, 24.4.2014, p. 14.
([5]) OJ L 121, 24.4.2014, p. 17.
([6]) OJ L 128, 30.4.2014, p. 64.
([7]) OJ L 128, 30.4.2014, p. 68.
([8]) OJ L 124, 25.4.2014, p. 27.

(2) the following points are inserted after point 12ns (Commission Implementing Regulation (EU) No 94/2014):

'12nt. **32014 R 0405**: Commission Implementing Regulation (EU) No 405/2014 of 23 April 2014 approving lauric acid as an existing active substance for use in biocidal products for product-type 19 (OJ L 121, 24.4.2014, p. 8).

12nu. **32014 R 0406**: Commission Implementing Regulation (EU) No 406/2014 of 23 April 2014 to approve ethyl butylacetylaminopropionate as an existing active substance for use in biocidal products for product-type 19 (OJ L 121, 24.4.2014, p. 11).

12nv. **32014 R 0407**: Commission Implementing Regulation (EU) No 407/2014 of 23 April 2014 approving transfluthrin as an existing active substance for use in biocidal products for product-type 18 (OJ L 121, 24.4.2014, p. 14).

12nw. **32014 R 0408**: Commission Implementing Regulation (EU) No 408/2014 of 23 April 2014 approving synthetic amorphous silicon dioxide as an existing active substance for use in biocidal products for product-type 18 (OJ L 121, 24.4.2014, p. 17).

12nx. **32014 D 0227**: Commission Implementing Decision 2014/227/EU of 24 April 2014 on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 124, 25.4.2014, p. 27).

12ny. **32014 R 0437**: Commission Implementing Regulation (EU) No 437/2014 of 29 April 2014 approving 4,5-Dichloro-2-octyl-2H-isothiazol-3-one as an existing active substance for use in biocidal products for product-type 21 (OJ L 128, 30.4.2014, p. 64).

12nz. **32014 R 0438**: Commission Implementing Regulation (EU) No 438/2014 of 29 April 2014 approving cyproconazole as an existing active substance for use in biocidal products for product-type 8 (OJ L 128, 30.4.2014, p. 68).'.

Article 2

The texts of Regulation (EU) No 334/2014 and Implementing Regulations (EU) No 405/2014, (EU) No 406/2014, (EU) No 407/2014, (EU) No 408/2014, (EU) No 437/2014 and (EU) No 438/2014 and Implementing Decision 2014/227/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.

Article 3

This Decision shall enter into force on 1 November 2014, 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, 24 October 2014.

For the EEA Joint Committee
The President
Kurt JÄGER

(*) No constitutional requirements indicated.