DECISION OF THE EEA JOINT COMMITTEE

No 84/2016

of 29 April 2016

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2017/2034]

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) 2016/105 of 27 January 2016 approving biphenyl-2-ol as an existing active substance for use in biocidal products for product-types 1, 2, 4, 6 and 13 ([1]) is to be incorporated into the EEA Agreement.

(2) Commission Implementing Regulation (EU) 2016/124 of 29 January 2016 approving PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-type 4 ([2]) is to be incorporated into the EEA Agreement.

(3) Commission Implementing Regulation (EU) 2016/125 of 29 January 2016 approving PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-types 2, 3, 11 ([3]) is to be incorporated into the EEA Agreement.

(4) Commission Implementing Regulation (EU) 2016/131 of 1 February 2016 approving C(M)IT/MIT (3:1) as an existing active substance for use in biocidal products for product-types 2, 4, 6, 11, 12 and 13 ([4]) is to be incorporated into the EEA Agreement.

(5) Commission Implementing Decision (EU) 2016/107 of 27 January 2016 not approving cybutryne as an existing active substance for use in biocidal products for product-type 21 ([5]) is to be incorporated into the EEA Agreement.

(6) Commission Implementing Decision (EU) 2016/108 of 27 January 2016 not approving 2-Butanone, peroxide as an existing active substance for use in biocidal products for product-types 1 and 2 ([6]) is to be incorporated into the EEA Agreement.

(7) Commission Implementing Decision (EU) 2016/109 of 27 January 2016 not to approve PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-types 1, 6 and 9 ([7]) is to be incorporated into the EEA Agreement.

(8) Commission Implementing Decision (EU) 2016/110 of 27 January 2016 not approving triclosan as an existing active substance for use in biocidal products for product-type 1 ([8]) is to be incorporated into the EEA Agreement.

(9) Commission Implementing Decision (EU) 2016/135 of 29 January 2016 postponing the expiry date of approval of flocoumafen, brodifacoum and warfarin for use in biocidal products for product-type 14 ([9]) is to be incorporated into the EEA Agreement.

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

([1]) OJ L 21, 28.1.2016, p. 74.
([2]) OJ L 24, 30.1.2016, p. 1.
([3]) OJ L 24, 30.1.2016, p. 6.
([4]) OJ L 25, 2.2.2016, p. 48.
([5]) OJ L 21, 28.1.2016, p. 81.
([6]) OJ L 21, 28.1.2016, p. 83.
([7]) OJ L 21, 28.1.2016, p. 84.
([8]) OJ L 21, 28.1.2016, p. 86.
([9]) OJ L 25, 2.2.2016, p. 65.

HAS ADOPTED THIS DECISION:

Article 1

The following points are inserted after point 12zzzb (Commission Implementing Regulation (EU) 2015/985) of Chapter XV of Annex II to the EEA Agreement:

'12zzzc. **32016 R 0105**: Commission Implementing Regulation (EU) 2016/105 of 27 January 2016 approving biphenyl-2-ol as an existing active substance for use in biocidal products for product-types 1, 2, 4, 6 and 13 (OJ L 21, 28.1.2016, p. 74).

12zzzd. **32016 D 0107**: Commission Implementing Decision (EU) 2016/107 of 27 January 2016 not approving cybutryne as an existing active substance for use in biocidal products for product-type 21 (OJ L 21, 28.1.2016, p. 81).

12zzze. **32016 D 0108**: Commission Implementing Decision (EU) 2016/108 of 27 January 2016 not approving 2-Butanone, peroxide as an existing active substance for use in biocidal products for product-types 1 and 2 (OJ L 21, 28.1.2016, p. 83).

12zzzf. **32016 D 0109**: Commission Implementing Decision (EU) 2016/109 of 27 January 2016 not to approve PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-types 1, 6 and 9 (OJ L 21, 28.1.2016, p. 84).

12zzzg. **32016 D 0110**: Commission Implementing Decision (EU) 2016/110 of 27 January 2016 not approving triclosan as an existing active substance for use in biocidal products for product-type 1 (OJ L 21, 28.1.2016, p. 86).

12zzzh. **32016 R 0124**: Commission Implementing Regulation (EU) 2016/124 of 29 January 2016 approving PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-type 4 (OJ L 24, 30.1.2016, p. 1).

12zzzi. **32016 R 0125**: Commission Implementing Regulation (EU) 2016/125 of 29 January 2016 approving PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-types 2, 3, 11 (OJ L 24, 30.1.2016, p. 6).

12zzzj. **32016 R 0131**: Commission Implementing Regulation (EU) 2016/131 of 1 February 2016 approving C(M)IT/MIT (3:1) as an existing active substance for use in biocidal products for product-types 2, 4, 6, 11, 12 and 13 (OJ L 25, 2.2.2016, p. 48).

12zzzk. **32016 D 0135**: Commission Implementing Decision (EU) 2016/135 of 29 January 2016 postponing the expiry date of approval of flocoumafen, brodifacoum and warfarin for use in biocidal products for product-type 14 (OJ L 25, 2.2.2016, p. 65).'

Article 2

The texts of Implementing Regulations (EU) 2016/105, (EU) 2016/124, (EU) 2016/125 and (EU) 2016/131 and Implementing Decisions (EU) 2016/107, (EU) 2016/108, (EU) 2016/109, (EU) 2016/110 and (EU) 2016/135 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 30 April 2016, provided that all the notifications under Article 103(1) of the EEA Agreement have been made (*).

(*) No constitutional requirements indicated.

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, 29 April 2016.

For the EEA Joint Committee
The President
Claude MAERTEN