DECISION OF THE EEA JOINT COMMITTEE

No 227/2017

of 15 December 2017

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2019/1635]

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) 2017/1376 of 25 July 2017 renewing the approval of warfarin as an active substance for use in biocidal products of product-type 14 ([1]) is to be incorporated into the EEA Agreement.

(2) Commission Implementing Regulation (EU) 2017/1377 of 25 July 2017 renewing the approval of chlorophacinone as an active substance for use in biocidal products of product-type 14 ([2]) is to be incorporated into the EEA Agreement.

(3) Commission Implementing Regulation (EU) 2017/1378 of 25 July 2017 renewing the approval of coumatetralyl as an active substance for use in biocidal products of product-type 14 ([3]) is to be incorporated into the EEA Agreement.

(4) Commission Implementing Regulation (EU) 2017/1379 of 25 July 2017 renewing the approval of difenacoum as an active substance for use in biocidal products of product-type 14 ([4]) is to be incorporated into the EEA Agreement.

(5) Commission Implementing Regulation (EU) 2017/1380 of 25 July 2017 renewing the approval of bromadiolone as an active substance for use in biocidal products of product-type 14 ([5]) is to be incorporated into the EEA Agreement.

(6) Commission Implementing Regulation (EU) 2017/1381 of 25 July 2017 renewing the approval of brodifacoum as an active substance for use in biocidal products of product-type 14 ([6]) is to be incorporated into the EEA Agreement.

(7) Commission Implementing Regulation (EU) 2017/1382 of 25 July 2017 renewing the approval of difethialone as an active substance for use in biocidal products of product-type 14 ([7]) is to be incorporated into the EEA Agreement.

(8) Commission Implementing Regulation (EU) 2017/1383 of 25 July 2017 renewing the approval of flocoumafen as an active substance for use in biocidal products of product-type 14 ([8]) is to be incorporated into the EEA Agreement.

([1]) OJ L 194, 26.7.2017, p. 9.
([2]) OJ L 194, 26.7.2017, p. 15.
([3]) OJ L 194, 26.7.2017, p. 21.
([4]) OJ L 194, 26.7.2017, p. 27.
([5]) OJ L 194, 26.7.2017, p. 33.
([6]) OJ L 194, 26.7.2017, p. 39.
([7]) OJ L 194, 26.7.2017, p. 45.
([8]) OJ L 194, 26.7.2017, p. 51.

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

HAS ADOPTED THIS DECISION:

Article 1

The following points are inserted after point 12zzzzw (Commission Implementing Decision (EU) 2017/1282) of Chapter XV of Annex II to the EEA Agreement:

'12zzzzx. **32017 R 1376**: Commission Implementing Regulation (EU) 2017/1376 of 25 July 2017 renewing the approval of warfarin as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 9).

12zzzzy. **32017 R 1377**: Commission Implementing Regulation (EU) 2017/1377 of 25 July 2017 renewing the approval of chlorophacinone as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 15).

12zzzzz. **32017 R 1378**: Commission Implementing Regulation (EU) 2017/1378 of 25 July 2017 renewing the approval of coumatetralyl as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 21).

12zzzzza. **32017 R 1379**: Commission Implementing Regulation (EU) 2017/1379 of 25 July 2017 renewing the approval of difenacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 27).

12zzzzzb. **32017 R 1380**: Commission Implementing Regulation (EU) 2017/1380 of 25 July 2017 renewing the approval of bromadiolone as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 33).

12zzzzzc. **32017 R 1381**: Commission Implementing Regulation (EU) 2017/1381 of 25 July 2017 renewing the approval of brodifacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 39).

12zzzzzd. **32017 R 1382**: Commission Implementing Regulation (EU) 2017/1382 of 25 July 2017 renewing the approval of difethialone as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 45).

12zzzzze. **32017 R 1383**: Commission Implementing Regulation (EU) 2017/1383 of 25 July 2017 renewing the approval of flocoumafen as an active substance for use in biocidal products of product-type 14 (OJ L 194, 26.7.2017, p. 51).'

Article 2

The texts of Implementing Regulations (EU) 2017/1376, (EU) 2017/1377, (EU) 2017/1378, (EU) 2017/1379, (EU) 2017/1380, (EU) 2017/1381, (EU) 2017/1382 and (EU) 2017/1383 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 16 December 2017, 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, 15 December 2017.

For the EEA Joint Committee
The President
Sabine MONAUNI

(*) No constitutional requirements indicated.