DECISION OF THE EEA JOINT COMMITTEE

No 265/2014

of 12 December 2014

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2015/2132]

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 658/2014 of the European Parliament and of the Council of 15 May 2014 on fees payable to the European Medicines Agency for the conduct of pharmacovigilance activities in respect of medicinal products for human use [1] is to be incorporated into the EEA Agreement.

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

HAS ADOPTED THIS DECISION:

Article 1

The following point is inserted after point 15zp (Commission Implementing Decision 2012/707/EU) of Chapter XIII of Annex II to the EEA Agreement:

'16. **32014 R 0658**: Regulation (EU) No 658/2014 of the European Parliament and of the Council of 15 May 2014 on fees payable to the European Medicines Agency for the conduct of pharmacovigilance activities in respect of medicinal products for human use (OJ L 189, 27.6.2014, p. 112).'

Article 2

The texts of Regulation (EU) No 658/2014 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 13 December 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, 12 December 2014.

For the EEA Joint Committee
The President
Kurt JÄGER

[1] OJ L 189, 27.6.2014, p. 112.
(*) No constitutional requirements indicated.