DECISION OF THE EEA JOINT COMMITTEE
No 87/2018

of 27 April 2018

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement

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 Delegated Regulation (EU) 2018/92 of 18 October 2017 amending Regulation (EU) No 658/2014 of the European Parliament and of the Council as regards the adjustment to the inflation rate of the amounts of the 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 is added in point 16 (Regulation (EU) No 658/2014 of the European Parliament of the Council) of Chapter XIII of Annex II to the EEA Agreement:

', as amended by:

- **32018 R 0092**: Commission Delegated Regulation (EU) 2018/92 of 18 October 2017 (OJ L 17, 23.1.2018, p. 2).'

Article 2

The text of Delegated Regulation (EU) 2018/92 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 28 April 2018, provided that all the notifications under Article 103(1) of the EEA Agreement have been made*.

[1] OJ L 17, 23.1.2018, p. 2.

* [No constitutional requirements indicated.] [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, 27 April 2018.

For the EEA Joint Committee
The President

Claude Maerten

The Secretaries
to the EEA Joint Committee

Dag Wernø Holter *Vaclav Navratil*