Commission Delegated Regulation (EU) 2015/1011 of 24 April 2015 supplementing Regulation (EC) No 273/2004 of the European Parliament and of the Council on drug precursors and Council Regulation (EC) No 111/2005 laying down rules for the monitoring of trade between the Union and third countries in drug precursors, and repealing Commission Regulation (EC) No 1277/2005 (Text with EEA relevance)

Article 13

Conditions and requirements concerning the information to be provided on the implementation of the monitoring measures

1 Member States shall submit the communications referred to in Article 32(1) of Regulation (EC) No 111/2005 and Article 13(1) of Regulation (EC) No 273/2004 to the Commission, in the month following each calendar quarter. The communications shall contain the information on all cases where the release of scheduled and non-scheduled substances was suspended or the scheduled and non-scheduled substances were detained.

2 That information shall include the following:

a the name of the substances;

b if known, the origin, provenance and destination of the substances;

c the quantity of the substances, their customs status and the means of transport used.

3 At the end of every calendar year, the Commission shall communicate to all Member States the information received pursuant to paragraph 1.

Changes to legislation:
There are outstanding changes not yet made to Commission Delegated Regulation (EU) 2015/1011. Any changes that have already been made to the legislation appear in the content and are referenced with annotations.
View outstanding changes

Changes and effects yet to be applied to :

– Art. 13 omitted by S.I. 2019/742 reg. 15(7)

Changes and effects yet to be applied to the whole legislation item and associated provisions

– Signature words omitted by S.I. 2019/742 reg. 15(8)
– Art. 9(2)(b) words substituted by S.I. 2019/742 reg. 15(4)
– Art. 10(a) words substituted by S.I. 2019/742 reg. 15(5)(a)