Regulation (EU) No 536/2014 of the European Parliament and of the Council of 16 April 2014 on clinical trials on medicinal products for human use, and repealing Directive 2001/20/EC (Text with EEA relevance)

CHAPTER XV

COOPERATION BETWEEN MEMBER STATES

Article 85

Clinical Trials Coordination and Advisory Group

1 A Clinical Trials Coordination and Advisory Group (CTAG), composed of the national contact points referred to in Article 83 is hereby established.

2 The CTAG shall have the following tasks:

- a to support the exchange of information between the Member States and the Commission on the experience acquired with regard to the implementation of this Regulation;
- b to assist the Commission in providing the support referred to in the second paragraph of Article 84;
- c to prepare recommendations on criteria regarding the selection of a reporting Member State.

3 The CTAG shall be chaired by a representative of the Commission.

4 The CTAG shall meet at regular intervals and whenever the situation requires, on a request from the Commission or a Member State. Any item of the agenda of the meeting shall be placed at the request of the Commission or a Member State.

5 The secretariat shall be provided by the Commission.

6 The CTAG shall draw up its rules of procedure. The rules of procedure shall be made public.

Changes to legislation:
There are currently no known outstanding effects for the Regulation (EU) No 536/2014 of the European Parliament and of the Council, Article 85.