Annex III

Suitability of the investigator

Version 13th January 2016
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Sponsor Logo
With regard to the clinical trial entitled:

Study title: XXXXXXX
Code: XXXXXX
EudraCT: XXXXXX

There is enclosed the list of the planned sites for the conduct of the clinical trial, the name and position of the principal investigators, and the planned number of subjects in each site.

Principal investigator	Study site	Planned number of subjects

The updated curriculum vitae accrediting the principal investigator's training in good clinical practice principles, professional experience in clinical trials and patient care is also attached. In this document, or in a ***specific annex to the document (conflict of interest declaration),*** it shall be indicated if there is any circumstance that could influence the impartiality of the investigators, such as economic interests or institutional affiliations.

In _____ on __ (day/month/year) _____

Signed: _____

Mr./Ms representative of the sponsor
PROTOCOL CODE: XXXXXX
EudraCT: XXXXXXXX