MINISTERIO DE SANIDAD
agencia española de medicamentos y productos sanitarios

DEPARTAMENT
OF MEDICINES
FOR HUMAN USE

Clinical Trials Division

Annex IV

Site Suitability

Version 27th March 2023

<<Insert name of site>>
With regard to the clinical trial entitled:
Study title:

Code:
EudraCT / EUCT Number:
Department to which Principal Investigator belongs:
Site:
Planned number of trial participants at the site:

It is hereby stated that, based on the nature and use of the investigational medicinal product, this site has the necessary human resources, equipment and facilities to carry out this study.

In addition, the collaboration of the following departments is acknowledged, which have been informed about their involvement in the study and have given their agreement in this regard.

Other departments involved (specify which and if none indicate "No"):

In *<<City>>* on *<<Day>>* of *<<Month>>* of *<<Year>>*

Signed: _____

Dr.: _____

Site Head / Delegated person (indicate position if that is the case).

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