MINISTERIO
DE SANIDAD, SERVICIOS SOCIALES
E IGUALDAD

agencia española de
medicamentos y
productos sanitarios

DEPARTMENT OF
MEDICINAL PRODUCTS
FOR HUMAN USE

Annex IV

Suitability of the facilities

Version 23rd June 2017
Date of Publication: 15th March 2018

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

Code: XXXXXX
EudraCT: XXXXXX
Department to which Principal Investigator belongs: XXXXX
Site: XXXXXXX

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 _____ on __ (day/month/year)_____

Signed: _____

Dr.: _____

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Site head / Delegated person.

MINISTRY OF HEALTH, SOCIAL SERVICES
AND EQUALITY
Spanish Agency for Medicinal Products and
Medical Devices