**Site Suitability**

*Updated according to AEMPS version 27th March 2023 AEMPS*

**COMPLEJO ASISTENCIAL UNIVERSITARIO DE SALAMANCA**

With regard to the clinical trial entitled:

**Study title: XXXX**

**Code:** XXXXX

**EudraCT / EUCT Number::**  XXXXX

**Department to which Principal Investigator belongs:** XXXXX

**Site:** Complejo Asistencial Universitario de Salamanca.

**Planned number of trial participants at the site**: XX

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 Salamanca, on \_\_\_ of of

|  |  |
| --- | --- |
| Signed Dr. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_  Head of Department\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ | Signed Dr.\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_  Site Head / Delegated person |