Site Suitability Template - Adopted in Hungary

Site name:

Site address:

Name of principal investigator:

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| 1. Please provide a written statement on the suitability of the site adapted to the nature and use of the investigational medicinal product. |
| Click or tap here to enter text. |
| 1. Please describe the suitability of the facilities |
| Click or tap here to enter text. |
| 1. Please describe the suitability of the equipment |
| Click or tap here to enter text. |
| 1. Please provide a description of all trial procedures which will take place at the site. |
| Click or tap here to enter text. |
| 1. Please provide a description of Human Resources arrangements and expertise at the site |
| Click or tap here to enter text. |
| 1. How will potential participants be identified (e.g. patients of the investigators’ patient pool; patients appeared as a result of recruitment; patients referred by an other specialist)? |
| Click or tap here to enter text. |
| 1. Please provide a description of the medical attendance policy following the end of the trial (i.e. how the continous medical attendance of the participant is ensured?)   Click or tap here to enter text. |
| 1. Please describe how an urgent medical help is organised in an emergency situation |
| Click or tap here to enter text. |

I confirm that the site has the facilities and equipment to be able to conduct the clinical trial and has suitable arrangements in place to ensure that all investigators and other individuals involved in conducting the trial have the suitable qualifications, expertise and training in relation to their role in the clinical trial, in compliance with EU Regulation 536/2014, and all conditions identified, which might influence the impartiality of any investigators, were addressed.

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Signature of Principal Investigator

or other responsible person Date