National Research Ethics Committee for Clinical Trials (NREC-CT)

Investigator Curriculum Vitae

This template must be used by Sponsors of clinical trials as part of the application dossier. A separate document should be completed and submitted for each site.

This template was originally developed and endorsed by the EU Clinical Trials Expert Group to comply with Regulation (EU) No. 536/2014 Clinical Trials on Medicinal Products for Human Use and adapted by the National Office for Research Ethics Committees.

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| PERSONAL INFORMATION |
| Name and title: | Click or tap here to enter text. |
| Profession: | Click or tap here to enter text. |
| Current Position: | Click or tap here to enter text. |

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| PROFESSIONAL REGISTRATION |
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| Registration state/province (if applicable): | Click or tap here to enter text. |

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| EDUCATION AND QUALIFICATIONS |
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| E-mail address: | Click or tap here to enter text. |
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| RELEVANT PROFESSIONAL EXPERIENCE |
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| RELEVANT CLINICAL TRIAL/STUDY EXPERIENCE |
| Investigator role | Therapeutic area | Type of trial | Year started | Phase  | Ongoing (Y/N) |
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| TRAINING |
| Research training (including GCP) | Institution name | Year obtained |
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