**PHRC Interrégional GIRCI AuRA 2025**

**CURRICULUM VITAE**

**de l’investigateur coordonnateur**

*Le CV peut être complété en français ou en anglais.*

*Le CV doit être daté et signé de moins d’un an.*

*La formation BPC est obligatoire, validité 3 ans.*

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| **Personal Information** |
| **Name:** | Click or tap here to enter text. |
| **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[[1]](#endnote-1)** |
| **Registration number:** | Click or tap here to enter text. |
| **Registration body:** | Click or tap here to enter text. |
| **Registration expiry date (if applicable):** | Click or tap here to enter text. |
| **Registration state/province (if applicable):** | Click or tap here to enter text. |

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| **Education and Qualifications[[2]](#endnote-2)**  |
| **Institution name** | **Qualification** | **Year** |
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| **Affiliation to a research organization?**(If so, please specify : Inserm, CNRS, CEA, INRA…) |
| **No** [ ]  | Click or tap here to enter text. |
| **Yes** [ ] **Name and address of the organization** | Click or tap here to enter text. |

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| **Current employment** |
| **Institution name:**  | Click or tap here to enter text. |
| **Department:** | Click or tap here to enter text. |
| **Institution address:** | Click or tap here to enter text. |
| **Telephone number:** | Click or tap here to enter text. |
| **E-mail address:** | Click or tap here to enter text. |

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| **Professional experience[[3]](#endnote-3)** |
| **Position** | **Institution name and department** | **Start year** | **End year** |
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| **Relevant clinical trial/study experience[[4]](#endnote-4)** |
| **Investigator role** | **Therapeutic area** | **Type of trial** | **Year started** | **Phase**  | **Ongoing** |
| Choose an item. | Click or tap here to enter text. | Choose an item. | Click or tap here to enter text. | Choose an item. | Choose an item. |
| Choose an item. | Click or tap here to enter text. | Choose an item. | Click or tap here to enter text. | Choose an item. | Choose an item. |
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| **Training** *Formation obligatoire, doit dater de moins de 3 ans (dont les BPC)* |
| **Research training (including GCP)** | **Institution name** | **Year obtained** |
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| **Date completed[[5]](#endnote-5):**  | Click or tap to enter a date. |
| **Signature[[6]](#endnote-6) (if required):** | Click or tap here to enter text. |

1. As per national legislation [↑](#endnote-ref-1)
2. Relevant to be an investigator [↑](#endnote-ref-2)
3. This should cover the preceding 10 years as a maximum [↑](#endnote-ref-3)
4. *Idem* [↑](#endnote-ref-4)
5. The CTR does not require signing individual documents in the clinical trial application – a request for signature could however be subject to national legislation. [↑](#endnote-ref-5)
6. As per national legislation, a signed version of the CV should be included in the trial master file however a signed version may not be required for regulatory review, this should be confirmed nationally. [↑](#endnote-ref-6)