Investigator Curriculum Vitae

This template may be used by Sponsors of clinical trials as part of the application dossier.

EU CT number : 2024-519879-25-00

A separate document should be completed and submitted for each site.

This template has been 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.

Personal Information

Name:	Elise BISMUTH-REISMAN
Title:	Doctor
Profession:	Pediatrician
Current position:	Full time Hospital Practitioner

Professional Registration[i]

Registration number:	RPPS: 10004394366
Registration body:	CNOM - France
Registration expiry date (if applicable):	N/A
Registration state/province (if applicable):	N/A

Education and Qualifications[ii]

Institution name	Qualification	Year
Joslin Diabetes Center (Boston)	Fellowship	2006-2007
Paris VI University	Medical School Student	2007
Paris VI University	Qualification in Pediatrics	2007
Paris VI University	Certification in Pediatric Endocrinology and Diabetology	2006
Medical departments in Paris Hospitals	Resident in Pediatrics	2001-2006
Paris VI University	Medical School Student	1995-2006
Paris VI University	Master 2 in Biology, physiology, pathology	2005

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Paris VI University	Master 1 in Biology and Medical Sciences	2001

Current employment

Institution name:	Robert Debré Hospital
Department:	Department of Pediatric, Endocrinology and Diabetes
Institution address:	48 boulevard Sérurier – 75019 Paris - France
Telephone number:	+33 1 40 03 20 67
E-mail address:	Elise.bismuth@aphp.fr

Professional experience[iii]

Position	Institution name and department	Start year	End year
Full time Hospital Practitioner	Department of Pediatric, Endocrinology and Diabetes – Robert Debré Hospital - Paris	2016	On going
Full time Hospital Practitioner	Pediatrics / Department of Endocrinology – Saint Pierre Hospital – La Réunion	2012	2016
Hospital Practitioner	Pediatrics Department – CHR Sud Réunion – La Réunion	2010	2011
Chief Resident	Department of Pediatric Endocrinology – Robert Debré Hospital - Paris	2007	2009

Relevant clinical trial/study experience[iv]

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Principal Investigator	Diabetes Type 1	Clinical Study	2025	Other	Yes
Principal Investigator	Diabetes Type 1	Interventional Trial	2023	Phase III	Yes
Chief Investigator	Diabetes Type 2	Interventional Trial	2022	Phase III	Yes

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Training

Research training (including GCP)	Institution name	Year obtained
ICH GPC 6E R3	Shclinical site education	2025
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Date completed[v]:	05/09/2025

[i] As per national legislation
[ii] Relevant to be an investigator
[iii] This should cover the preceding 10 years as a maximum
[iv] Idem
[v] The CTR does not require signing individual documents in the clinical trial application – a request for signature could however be subject to national legislation.