CURRICULUM VITAE	
First Name	Colin
Family Name	McGregor
Middle Name (if applicable)	Edward
Position	Project Coordinator
Location (City, State, Country)	Chicago, IL, United States of America

EDUCATIONAL BACKGROUND, PROFESSIONAL TRAININGS		
Dates	Degree / Diploma / Certificate	Institution Name, Location
22-Mar-2024	Certificate: Certified Clinical Research Coordinator	Association of Clinical Research Professionals (ACRP)
22-Sep-2022	Certificate: Good Clinical Practice Course (US FDA focus)	Collaborative Institutional Training Initiative (CITI Program)
2020	Bachelor of arts: Cognitive science	Johns Hopkins University

CURRENT AND PREVIOUS POSITIONS (including Academic Appointments)		
Dates	Position	Organization, Location
2024 - Present	Project Coordinator (PC)	CONFIDENCE Pharmaceutical Research, LLC
Main Responsibilities:	• Act as a direct contact for the Sponsor for non-medical issues as directed and applicable • Report project status to the Sponsor • Control and report the project progress to the PM as per contract, budget, and the Company's Quality Management System Documentation • Perform project-related financial reporting • Interact with other departments of the Company on study-related issues • Organize and facilitate the study team meetings (STM) • Write the minutes of the STM • Maintain documentation and perform quality control to ensure the most recent revisions of documents are on portals and shared drives • Assist the project team (Clinical Research Associate, PM, etc.) with gathering the documents required for submission to Regulatory Authorities and Ethics Committees • Assure that all the required financial documents have been received from the sites • Maintain the Trial Master File (TMF) as appropriate • Provide data for invoices of investigators and sites to the financial department • Assist in conducting the Investigators' Meeting • Translate the study documents or arrange them for translation in accordance with local needs as required	

CURRENT AND PREVIOUS POSITIONS (including Academic Appointments)		
	• Train the study team members on how to prevent and resolve the identified study issues • Keep the project-specific training documentation updated	
Dates	Position	Organization, Location
2022 – 2024	Lead clinical oncology research coordinator	Roswell Park Comprehensive Cancer Center
Main Responsibilities:	• Supervisor and primary support for 15 solid-tumor protocols; combined 20-30 active patients and 60-70 on follow-up • Successfully opened 4 pharmaceutical-sponsored trials and ran 7 others, including 4 phase 1 trials with PKs for as many as 3 simultaneous reference drugs • Assessed protocols for efficacy, feasibility, and subject safety; reviewed inclusion/exclusion criteria; facilitated open discussions between sponsor and investigator • Managed high-accruing, multi-phase IIT w/initial dose escalation • Ran two National Cooperative Group studies on site level • Completed in-house monitoring SDV tasks for an investigator-initiated trial (total of 21 patients over roughly 1 year) and shadowed in-house CRAs • Assists in updating and rewriting existing standard operating procedures to adapt to evolving IRB, FDA, and NIH regulations, as well as FDA audit findings • Maintains open communication between sponsor, CRO, investigator, processing labs, treatment center, site data management, and home clinic • Provides oversight and training of new clinical research coordinators, assistants, and laboratory employees • Ensured timely and consistent long-term follow-up of 60-70 patients while consistently recruiting at high-accruing site • Manages 4 data-entry employees and delegated tasks appropriately • Worked in multiple EDC databases (including REDCap, iMedidata RAVE, Advarra, Veeva) and electronic medical records (Epic, Allscripts EHR) • Obtains informed consent, demographics, medical history, concomitant medications, and documented appropriately in source • Maintains accurate case report forms, source binders, and electronic source • Serves as point-of-contact during 6 site initiation visits and hundreds of investigator meetings • Ensured compliance with 1572 and DOA • Coordinates sponsor-supplied equipment approval and testing • Completed and compiled continuing review documentation • Reports, documents, and communicates routine AEs, SAEs, AESIs, and DLTs as applicable and ensures subject safety in accordance with IRB guidelines • Verifies compliance with GCP, FDA, ICH, and any local regulations • Documents protocol deviations and reportable problems; assisted sponsors with study startup/closeout • Provides timely, efficient, and friendly assistance to monitors	
Dates	Position	Organization, Location
2020 – 2022	Clinical neurology research coordinator	Johns Hopkins Medical Institute

CURRENT AND PREVIOUS POSITIONS (including Academic Appointments)		
2017 – 2020	Research assistant	
Main Responsibilities:	• Coordinator of 55 patients on treatment or follow-up • Served as coordinator on 6 protocols involving neurodegenerative movement disorder patients • Designed and built pilot study aiming to test investigational walking device (see Publications) • Attended and participated in investigator/staff meetings. Provided guidance on progress and participant timelines • Obtained informed consent, baseline information, financial disclosure statements • Ensured compliance with ICH, GCP, FDA, and local guidelines • Coordinated meetings and data reviews with sister sites and maintained rapport with colleagues across the world • Administered tests and questionnaires including the MoCA, Purdue Pegboard Assessment, Box and Blocks Test, Hopkins Verbal Learning Assessment, Trail-Making Tests, and Stroop Tests • Completed and compiled continuing review documentation • Actively attended several physicians' clinic days to distribute IRB-approved recruitment materials to prospective patients • Participated in journal club meetings and suggested papers for discussion	

CLINICAL RESEARCH EXPERIENCE			
Dates	Therapeutic Area	Phase	Role in the Study
2022 – 2024	Thoracic medical oncology	I-III	Lead CRC
2018 – 2022	Parkinson's Disease/Atypical Parkinsonism	II	CRC

ADDITIONAL INFORMATION

Languages:

- English

Publications:

- Bastepe-Gray S, Wainwright L, Lanham DC, Gomez G, Kim JS, Forshee Z, Kaas B, McCoy A, McGregor C, Moukheiber E, Rajan S, Suarez-Cedeno G, Wang J, Brennan S, Coughlin S, Kang K, Pantelyat A. GuitarPD: A Randomized Pilot Study on the Impact of Nontraditional Guitar Instruction on Functional Movement and Well-Being in Parkinson's Disease. Parkinsons Dis. 2022 Jun 25;2022:1061045. doi: 10.1155/2022/1061045. PMID: 35795456; PMCID: PMC9252755.
- Zajac JA, Porciuncula F, Cavanaugh JT, McGregor C, Harris BA, Smayda KE, Awad LN, Pantelyat A, Ellis TD. Feasibility and Proof-of-Concept of Delivering an Autonomous Music-Based Digital Walking Intervention to Persons with Parkinson's Disease in a Naturalistic Setting. J Parkinsons Dis. 2023;13(7):1253-1265. doi: 10.3233/JPD-230169. PMID: 37840504; PMCID: PMC10657706.

Signature		Date (DD-MON-YYYY)	3-Oct-2024

Colin McGregor CV 2024_10_02

Final Audit Report 2024-10-03

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