EDGAR LOPEZ PACHECO

CONTACT INFORMATION:

- Mobile phone number: 956-478-3600
- e-mail: edgar_lopezp@hotmail.com

EDUCATION

- **MASTER´S DEGREE OF SCIENCE IN IMMUNOBIOLOGY**. PROFESSIONAL ID: 10052012.
  University of the State of Nuevo Leon (Jan/2014 -May/2016)
- **BACHELOR´S DEGREE AS CHEMIST, BACTERIOLOGIST AND PARASITOLOGIST.** PROFESSIONAL ID: 8659666.
  University of the State of Nuevo Leon (Jan/2009 -Feb/2014)

EXPERIENCE

- **POSITION: LEAD CLINICAL RESEARCH COORDINATOR (ONCOLOGY)**
  FROM: MARCH 2022 TO: CURRENTLY
  COMPANY: DHR HEALTH INSTITUTE FOR RESEARCH AND DEVELOPMENT
  DUTIES: Run study feasibility reports, maintain electronic and hard copy documents related to all protocols, process liquid biopsy samples as per protocol guidelines, order supplies that pertain to a specific protocol, analyze questions/actions brought forth by administrators, employees, board and determine research requirements, must be able to collect and analyze data in relation to expected outcomes, prioritize and handle the diverse workload of projects, correspondences, and project/report tracking, draft, review, and edit reports, opinions, correspondence, articles, and other documents related to Oncology clinical trials.

- **POSITION: STUDY COORDINATOR – CLINICAL RESEARCH ASSOCIATE**
  FROM: SEPTEMBER 2019 TO: FEBRUARY 2022
  COMPANY: NEUROCYTONIX, INC.
  EXPERTISE: MEDICAL DEVICE CLINICAL TRIALS WITH CEREBRAL PALSY PATIENTS (PHASE II CLINICAL TRIALS)
  DUTIES: Responsible for monitoring medical device clinical trials in patients with cerebral palsy. Responsible for performing site initiation visits (SMV), Interim monitoring visits (IMV), and close-out visits. Source data verification and review of the clinical trial EDC System (e-CRF). Responsible for monitoring adverse events (AE), serious adverse events (SAE), and adverse incidents to ensure that these events are recorded, managed, and reported according to FDA regulation, applicable ICH/GCP guidelines and standard operating procedures. Preparation of monitoring plans and monitoring reports. Translation of monitoring reports and procedures from English to Spanish and vice versa.

- **POSITION: STUDY COORDINATOR – CLINICAL RESEARCH ASSOCIATE**
  FROM: APRIL 2016 TO: NOVEMBER 2019
  COMPANY: AVANT SANTE RESEARCH CENTER
  EXPERTISE: CLINICAL TRIALS WITH INVESTIGATIONAL DRUGS IN CANCER PATIENTS AND HEALTHY VOLUNTEERS (PHASE II AND III CLINICAL TRIALS)
  DUTIES: Responsible for Monitoring and coordinating bioavailability/bioequivalence clinical studies in patients with prostate cancer and healthy volunteers. Preparation, and submission of essential documents (Protocol, informed consent form and investigator´s brochure) to institutional review boards and institutional committees. Responsible for preparing and submitting pharmacovigilance and safety reports to regulatory authorities. Responsible for attending audits and inspections from regulatory entities such as COFEPRIS (Mexico), FDA (United States), and ANVISA (Brazil). Research master file management and coordination of along with principal investigator. Responsible for performing drug accountability. Screening of potential clinical trial subjects (consent form, inclusion, and exclusion criteria assessment).

- **POSITION: QUALITY ASSURANCE AND QUALITY CONTROL (QC/QA)**
  FROM: JUNE 2013 TO: JANUARY 2014
  COMPANY: SAN MATEO.
  DUTIES: Preparation, translation (Spanish to English), and implementation of procedures. responsible for quality assurance in the microbiology laboratory.

- **POSITION: CLINICAL LABORATORY TECHNICIAN AND PHARMACY ASSISTANT**
  FROM: JANUARY 2011 TO: JANUARY 2012
  COMPANY: HOSPITAL NOVA. SERVICIOS INTEGRALES NOVA DE MONTERREY.
  DUTIES: Experience performing laboratory blood tests based on immunology, serology, biochemistry, and hematology. Performance of EKG and Phlebotomy procedures. Responsible for handling and safeguarding clinical files. Responsible for retention and dispensing of medications for patients.

LANGUAGES:

- **SIMULTANEOUS TRANSLATOR AND INTERPRETER: ENGLISH AND SPANISH.** From January 2020 – Currently
- **TEST OF ENGLISH AS A FOREIGN LANGUAGE (TOEFL).** Score: 570
- **EFSET EF STANDARD ENGLISH TEST:** C1 Advanced.
- **SPANISH:** Native

CLINICAL RESEARCH CERTIFICATIONS:

- **NIDA CLINICAL TRIAL NETWORK: GOOD CLINICAL PRACTICES** (GCP) by the National Institute on Drug Abuse, National Institute of Health, U.S.A. expiration date: April 20, 2023.
- **GOOD CLINICAL PRACTICES FOR CLINICAL TRIALS WITH INVESTIGATIONAL DRUGS AND MEDICAL DEVICES (U.S. FDA FOCUS).** CITI program. Record ID: 40834887. Expiration date: October 14, 2024.
- **GOOD CLINICAL PRACTICE (GCP).** CITI program. Record ID: 47482349. expiration date: February 15, 2025.
- **CLINICIANS – PRIVACY AND SECURITY (IPS).** Citi program. Record ID: 45441612.
- **HUMAN RESEARCH** – BIOMEDICAL RESEARCHERS. CITI program. Record ID: 37285779. Expiration date: October 20, 2024.
- **CITI CONFLICTS OF INTEREST.** CITI program. Record ID: 47482350. Expiration date: February 16, 2024.
- **CLINICAL RESEARCH MONITORING.** Intercontinental university. Record ID: 003338.
- **BIOETHICS.** UNESCO and TEC SALUD school of medicine. August 2017.
- **INTRODUCTION TO THE PRINCIPLES AND PRACTICE OF CLINICAL RESEARCH (IPPCR).** The National Institutes of Health.U.S.A. From: September 2016 to: May 2017.
- **GOOD LABORATORY PRACTICES (GDP).** University of the State of Nuevo Leon. October 2016.

PUBLISHED CLINICAL RESEARCH PAPERS ON INTERNATIONAL JOURNALS:

- Franco-Molina, Moises & Evangelina Coronado-Cerda, Erika & **Lopez-Pacheco, Edgar** & Zarate, Diana & Galindo, Sergio & del Carmen Salazar-Rodriguez, Maria & Ramos Zayas, Yareellys & Tamez-Guerra, Reyes & Rodríguez-Padilla, Cristina. (2018). **Chitosan nanoparticles plus KLH adjuvant as an alternative for human dendritic cell differentiation.** Current Nanoscience. 14. 10.2174/1573413714666181008110627.

EDGAR LOPEZ PACHECO Edgar Lopez P. MAR/18/2022 (SIGNATURE AND DATE)