Investigator Name:
Site Name:
Site Address:
Phone Number:

Emeka M. Eziri, MD, MS, FCCP Carolina Clinical Research 9040 Nations Ford Road, Charlotte, NC 28273 704-290-4587

**Education: Medical School** and Post-Graduate Studies

New York University Robert F. Wagner Graduate School of Public Service Master of Science in Management May 2001 New York, NY 10012

Albert Einstein College of Medicine Montefiore Medical Center, The University Hospital of AECOM Fellowship Training in Critical Care Medicine August 2001 Bronx, NY 10467

Albert Einstein College of Medicine Bronx Lebanon Hospital Center Fellowship Training in Pulmonary Disease Medicine July 1999 Bronx, NY 10457

Mount Sinai School of Medicine North General Hospital Internship and Residency in Internal Medicine July 1997 New York, NY 10035

University of Calabar College of Medicine Degree: MD Specialization: Medicine, Surgery 1988 Calabar, Nigeria

# Clinical Research Experience

Clinical Investigator, Carolina Clinical Research (CCR)

Charlotte, NC 28273

July 2019 - Present

Clinical Investigator, Vitalink Research - Gastonia

(Effective 01 SEP 2016, Gastonia Pharmaceutical Research is d/b/a Vitalink Research-

Gastonia)

Gastonia, NC 28054

Jan. 2014 - May 2018

**PCC Clinical Research** 

Huntersville, NC 28078

Oct. 2008 - December 2012

# **Board Certification**

American Board of Internal Medicine-1997, 2010

American board of Internal Medicine-Pulmonary-2004, 2012

American Board of Quality Assurance & Utilization Review Physicians-1997

# Fellowships and Licensure

Fellow, American College of Chest Physicians

Fellow, American Institute for Healthcare Quality

Fellow, American Board of Quality Assurance &

**Utilization Review Physicians** 

**Good Clinical Practice** 

Medical and Dental Council of Nigeria

1988-Present

North Carolina Medical Board

2001-Present

South Carolina Medical Board

2015-Present

Florida Medical Board

2016 - Present

# **Professional Experience**

Englewood Hospital and Medical Center House Physician, Department of Medicine Englewood, NJ 1999-2001

Medcare of East Rutherford Urgent Care Physician East Rutherford, NJ 1999-2000

Pulmonary Clinic of the Carolina's PC Medical Director Lincolnton, NC 2001-Present

Sleep Medix Consultant Lekki, Lagos, Nigera 2012-2013

# **Hospital Affiliations**

Atrium Health, Lincolnton NC Caromont Regional Medical Center, Gastonia NC Kings Mountain Hospital, Kings Mountain, NC Novant Health Huntersville Medical Center, Huntersville, NC

# **Sponsors and Studies**

# Chiesi

A phase II, 8 week, randomized, double-blind, placebo and active controlled, parallel group, dose ranging study to evaluate the efficacy and safety of 3 doses of CHF 718 PMDI (beclomethasone dipropionate) in asthmatic subjects.

Protocol: CCD-05993AA3-01

July 2017 - Ongoing (Study ongoing but no longer serving as PI

Expected Enrollment: 6 subjects randomized

Actual Enrollment: 8 subjects randomized prior to departure from PI role

#### GlaxoSmithKline

A phase III, 52 week, randomized, double-blinded, 3-arm parallel group study, comparing the efficacy, safety and tolerability of the fixed dose triple combination FF/UMEC/VI with the fixed dose dual combinations of FF/VI and UMEC/VI, all administered once-daily in the morning via a dry powder inhaler in subjects with chronic obstructive pulmonary disease

Protocol: CTT116855 Jul 2014-Aug 2016 Expected Enrollment: 12 Actual Enrollment: 29

### **Novartis**

A 26 week, randomized, active-controlled safety study of double-blind Formoterol fumarate in free combination with an inhaled corticosteroid Steroid an inhaled corticosteroid in adolescent and adult patients with persistent asthma

Protocol: CFOR258D2416

Nov 2014-Jul 2016

Expected Enrollment: 10 Actual Enrollment: 21

# **Sunovion Therapeutics**

A comparative effectiveness and safety study of Arformoterol Tartrate Inhalation solution and Tiotropiom bromide on re-hospitalization in Chronic obstructive pulmonary disease (COPD) subjects (a phase IIII study) **Protocol: SEP091-402** 

Mar 2015-Dec 2016

**Expected Enrollment: Open enrollment** 

**Actual Enrollment: 17** 

# **Pearl Therapeutics**

A randomized, double-blinded, chronic dosing (24 weeks), placebo-Controlled. Parallel group, multi-center study to assess the efficacy And safety of PT003, PT005, and PT001 in subjects with moderate To very severe COPD, compared with placebo

Protocol: PT003014 Apr 2015-Jul 2016

Expected Enrollment: 10 Actual Enrollment: 15

# GlaxoSmithKline

Mepolizumab vs. Placebo as add-on treatment for frequently exacer-Bating COPD patients characterized by eosinophil level

Protocol: MEA117113 Aug 2015-Jan 2017 Expected Enrollment: 6 Actual Enrollment: 10

# Theravance

A phase 3, 12-week, randomized, double-blinded placebo-controlled Parallel-group study of nebulized TD-4208 in subjects with Chronic Obstructive Pulmonary Disease

Protocol: TD-4208-0126 Oct 2015-Nov 2016 Expected Enrollment: 10 Actual Enrollment: 21

Signature and date: