



DATE December 17, 2021

FROM Bhanu Kannan, Bioresearch Monitoring Branch (BMB)
Division of Inspections and Surveillance (DIS)
Office of Compliance and Biologics Quality (OCBQ)

THROUGH Dennis Cato, Chief, BMB

THROUGH Carrie Mampilly, M.P.H., Director, DIS

TO Michael Kennedy, Ph.D., Chair, STN 125743/0
Vijay Kumar, M.D., Clinical Reviewer
Nancy Skeeter, RPM

SUBJECT Bioresearch Monitoring Final Review Memo
SPONSOR Greencross Corporation
PRODUCT Immune Globulin Intravenous (Human) 10% Liquid
BLA STN 125743/0

FINAL SUMMARY STATEMENT

Bioresearch Monitoring (BIMO) inspections were conducted at three domestic clinical investigator sites participating in the conduct of study protocol GC5107B_P3. The inspections did not reveal substantive problems impacting the data submitted in support of this Biologics License Application (BLA).

Background

Three clinical investigators (CI) were inspected in support of this BLA. The inspection was conducted in accordance with FDA's Compliance Program (CP) 7348.811, Inspection Program for Clinical Investigators. The inspection assignments were issued for the following study protocol:

GC5107B_P3: An Open-Label, Single-Arm, Historically Controlled, Prospective, Multicenter Phase III Study to Evaluate the Safety, Efficacy and Pharmacokinetics of Immune Globulin Intravenous (Human) GC5107 in Subjects with Primary Humoral Immunodeficiency.

The sites were selected based on previous inspectional history, geographic location, and the data submitted in the BLA. The inspection assignment included specific questions concerning the study protocol, and information submitted in the BLA was compared to source documents at the site. Study GC5107B_P3 was conducted at 17 study centers enrolling a total of 49 subjects aged ≥ 2 to ≤ 70 years with Primary Humoral Immunodeficiency requiring Immune Globulin Intravenous (IGIV) treatment. The domestic CI sites inspected in support of this BLA covered all subjects enrolled in the study.

Inspection Outcome

Site ID	Number of subjects enrolled	Location	Form FDA 483 issued	Final Inspection Classification
21	3	Oklahoma Institute of Allergy and Asthma Clinical Research, LLC Oklahoma City, OK 73131	No	No Action Indicated
27	5	Allergy Associates of The Palm Beaches, P.A. North Palm Beach, FL 33408	No	No Action Indicated
30	5	Pediatric Pulmonary Associates of North Texas, P.A. Frisco, TX 75043	No	No Action Indicated

The inspections verified the data reported in the BLA, including but not limited to subject eligibility, protocol deviations, IGIV infusion including any interruptions, serious bacterial infections, and adverse events for the subjects enrolled at the inspected clinical sites. The inspections further evaluated the adequacy of the study and site monitoring by the sponsor. No Form FDA 483s were issued for the four inspected study sites.

Noteworthy inspectional findings

None.

Sponsor Issues

No significant sponsor issues were noted.

Financial Disclosure

The Clinical Investigator Compliance Program directs the FDA investigators to ask the clinical investigator if and when he/she disclosed information about his/her financial interests to the sponsor and/or interests of any sub-investigators, spouses and dependent children, and if and when the information was updated. The information submitted to the BLA was verified at the inspected clinical sites and found no deviations in the submitted data.

Administrative follow-up

No administrative follow-up is warranted at this time from BIMO for the inspected clinical investigators. Should you have any questions about the contents of this memo or any aspect of BIMO, please contact Dennis Cato at 301-741-7326.

 Bhanu Kannan, M.S.
 Consumer Safety Officer

Electronic Copies

CBER Connect BLA STN125743/0

Michael Kennedy, Chair, STN 125743/0

Vijay Kumar, Clinical Reviewer

Nancy Skeeter, RPM

Carrie Mampilly, Division Director

Dennis Cato, Branch Chief

Bhanu Kannan

cberbimonotification@fda.hhs.gov, Chron file

ORA BIMOE Correspondence

ORA BIMOW Correspondence

Sheri Stephenson, FDA Investigator

Corrine Carter, FDA Investigator

Joanne Schlossin, FDA Investigator

History:

Kannan draft: 11/09/2021