University of Central Florida Institutional Review Board
Office of Research & Commercialization
12201 Research Parkway, Suite 501
Orlando, Florida 32826-3246
Telephone: 407-823-2901 or 407-882-2276
www.research.ucf.edu/compliance/irb.html

Approval of Human Research

From: **UCF Institutional Review Board #1**
FWA00000351, IRB00001138

To: **Saleh A Naser** and Co-PI **Shazia Ashraf Beg**

Date: **June 09, 2017**

Dear Researcher:

On 06/09/2017 the IRB approved the following modifications to human participant research until 06/08/2018 inclusive:

Type of Review:	IRB Continuing Review Application Form
Modification Type:	Removed External Collaborators: Dr. Srinivas Seela, Dr. Seela Ramesh and Dr. Harinath Sheela
Project Title:	The Role of Mycobacterium avium subspecies paratuberculosis (MAP) in Crohn's disease and Diabetes
Investigator:	Saleh A Naser
IRB Number:	SBE-16-12193
Funding Agency:	
Grant Title:	Crohn's disease Research 250-10-10-2 THe Crohn's research project is funded by the Florida Governor Office as a budget item.
Research ID:	n/a

The scientific merit of the research was considered during the IRB review. **NOTE: Because this study was not approved before the IRB expiration date, there was a lapse in IRB approval from 4/18/2017 to the new approval date above.** The Continuing Review Application must be submitted 30days prior to the expiration date for studies that were previously expedited, and 60 days prior to the expiration date for research that was previously reviewed at a convened meeting. Do not make changes to the study (i.e., protocol, methodology, consent form, personnel, site, etc.) before obtaining IRB approval. A Modification Form **cannot** be used to extend the approval period of a study. All forms may be completed and submitted online at https://iris.research.ucf.edu .

If continuing review approval is not granted before the expiration date of 06/08/2018, approval of this research expires on that date. When you have completed your research, please submit a Study Closure request in iRIS so that IRB records will be accurate.

Use of the approved, stamped consent document(s) is required. The new form supersedes all previous versions, which are now invalid for further use. Only approved investigators (or other approved key study personnel) may solicit consent for research participation. Participants or their representatives must receive a signed and dated copy of the consent form(s).

All data, including signed consent forms if applicable, must be retained and secured per protocol for a minimum of five years (six if HIPAA applies) past the completion of this research. Any links to the identification of participants

should be maintained and secured per protocol. Additional requirements may be imposed by your funding agency, your department, or other entities. Access to data is limited to authorized individuals listed as key study personnel.

In the conduct of this research, you are responsible to follow the requirements of the Investigator Manual.

On behalf of Sophia Dziegielewski, Ph.D., L.C.S.W., UCF IRB Chair, this letter is signed by:

Signature applied by Renea C Carver on 06/09/2017 02:18:04 PM EDT

IRB Coordinator