



Thursday, October 13th, 2016

Dear Sponsor, Clinical Team, and/or Clinical Research Organization:

It was our pleasure to meet you at the *2016 Global Site Solutions Summit*. Please allow me to formally introduce you to *In Situ Global Clinical Trials Network, LLC*, as its Founder and Chief Executive Officer.

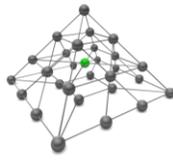
Established in 2016, and based in San Juan, Puerto Rico, *In Situ Global Clinical Trials Network, LLC* is currently a collective of twenty-four (24) United States Board-Certified Specialist and Subspecialist Physicians from ten (10) clinical areas, supported by a central Executive Core, and organized to conduct industry-sponsored Clinical Trials. Our operations are based on collaborations among the best emerging talents; comprehensive logistics to ensure the integrity and continuity of trial processes; and the utmost administrative efficiency to produce quality and timely results.

Puerto Rico currently provides clinical, administrative, and regulatory advantages over other world jurisdictions, as follows:

- Significant incidences and prevalence of some of the most challenging diseases and conditions, thus providing substantial pools of potential Clinical Trials Participants.
- Commercial and manufacturing presence of multiple global pharmaceutical and medical device companies for over the past four decades, providing an ideal corporative background for the conduction of Clinical Trials operations.
- Local corporate economic and tax incentive programs tailored to scientific research operations, and in accordance with the United States of America's Federal *Internal Revenue Service (IRS)*.
- Federal standards and *Food and Drug Administration (FDA)* oversight on Clinical Trials processes as part of the Government's longstanding political status as a Commonwealth of the United States of America.

At an institutional level, we strive to provide added levels of assurances to our Sponsors through six (6) main operational keys:

1. *100% United States Board-Certified Specialist and Subspecialist Physicians.* We trust that our Investigators' medical credentials serve as a testament of our commitment to Sponsors and Clinical Research Organizations, and their Clinical Trials.
2. *Back-Up Investigator.* In addition to the standard appointment of a Principal Investigator and a Sub-Investigator, we designate a *Back-Up Investigator* per Trial as a security clause in the eventuality of an extraordinary occurrence that interferes with the duties and responsibilities of any of these first two officials. We are confident that this tripartite arrangement provides, not only an added safety feature, but a supplementary element of expertise on a specific clinical area that benefits Sponsors, Clinical Research Organizations, and Participants.



3. *Clinical Mentors Board.* Recognizing the clinical perspectives that must be taken into account during the evaluation and conduction of a Clinical Trial, we have developed a *Clinical Mentors Board* comprised of local Key Opinion Leaders that provide their periodic input on our operations.
4. *Interpretation, Translation, and Linguistic Localization.* Understanding that language is an essential element of human interaction that can pose challenges during Clinical Trials at Hispanic Sites, we have partnered with RITA ("*Red de Intérpretes y Traductores Asociados*" or "*Network of Associated Translators and Interpreters*"), a fully certified and accredited Translation and Interpretation Company, to ease these linguistic processes for Participants, Clinical Research Organizations, and Sponsors (www.ritapr.com).
5. *Global Administrative and Logistics Support.* We provide continuous, 24/7/365, administrative and logistics support to our Investigators, thus eliminating unnecessary disruptions, and helping them focus on their clinical affairs for each trial.
6. *Professional Liability Insurance.* All our Clinical Trials operations are backed by a *Professional Liability Insurance* that protects our Sponsors' investments in the eventuality of an extraordinary occurrence that impedes the fulfillment of our duties, as per the pertaining contract agreement.

We would treasure an opportunity to further discuss our professional offerings, and candidly put *In Situ Global Clinical Trials Network, LLC* at your service for any Clinical Trial needs, trusting that our operational model and human resource will satisfy your expectations in the event of a professional endeavor. Please visit our Official Webpage and *LinkedIn* Page through the links provided below for more information on our current and upcoming clinical affairs.

Sincerely,

Dr. Oscar A. Colón-Acosta, MD

Founder/CEO

In Situ Global Clinical Trials Network, LLC

Metro Office Park

Metro Parque 7, Street 1, Suite 204

Guaynabo, PR 00968

founder@insitu.global

Office: 787-200-2702

Mobile: 234-207-4243

Fax: 787-793-4495

www.insitu.global

www.linkedin.com/company/in-situ-global-clinical-trials-network-llc