Working with CRI International
Database Inclusion Agreement

☐ Yes, I want to be listed in CRI International's database

We are a Trial Management Organization (TMO) that specializes in investigator identification, study start-up, and project management. As a TMO, we work with sites throughout the world that are capable of conducting studies in a variety of therapeutic areas. We do not own the sites with which we work nor do we have exclusive contracts with them. We do not charge a fee for inclusion in our database.

Our success depends on the quality and commitment of the sites with which we work. Our goal is to build a partnership with our member sites to help meet your goals as clinical research providers. In addition to providing an entirely new opportunity, we also assist you with rapid start-up and patient recruitment. Our success is dependent on your success! Our team is focused on providing solutions customized to your site and location!

At this stage, we are reaching out to you to identify investigators for sponsor trials. Upon receipt of a new study opportunity, we search our database for qualified investigators based on the sponsor's criteria and the investigator's interest and experience. When available, a study synopsis and site information questionnaire will be forwarded to you. (Please note that turnaround deadlines are set by CROs and sponsors. CRI International makes every effort to extend the most time to you for completion). All qualified responses are then submitted for sponsor consideration. You will receive a confirmation of your submission that will contain additional study information.

Once a CRI International affiliated investigator is selected, we will act as the point of contact for critical documents and administrative services for the site, CRO, sponsor and IRB. We will work with you to compile and submit regulatory start-up and maintenance documents per sponsor requirements. CRI International will also negotiate a study budget with you and the sponsor, and will work to complete your Clinical Trial Agreement. CRI International is then responsible for payment of study grant reimbursement to your site.

All clinical aspects of the trial will be managed directly between the CRO/sponsor and your site.

Please contact us if you have any questions about CRI International's business services at cri-indiainfo@criww.com. We look forward to working with you!

By Signing below I indicate that I understand the above statements and wish to be listed in CRI International's database.

_____

Signature Date

_____

Printed Name