



## WHEN SHOULD YOU CONTACT TRC?

<b>Long Site Initiation Process</b>	<ul style="list-style-type: none"> <li>• When a study has multiple contracting or legal issues across countries and sites, how will you keep sites enthusiastic about your study?</li> <li>• When do you communicate and train as the sites are slowly initiated into the study?</li> </ul>
<b>Difficult to Recruit Population</b>	<ul style="list-style-type: none"> <li>• When a site's database will not be sufficient, how will you help sites look at alternative ways to find patients?</li> <li>• How will you help sites explain to patients why they should participate in a clinical trial?</li> </ul>
<b>Complex Protocol</b>	<ul style="list-style-type: none"> <li>• What delivery methods do you use to communicate complicated and multiple protocols?</li> <li>• How do you present difficult protocols to effectively train Coordinators? How do you ensure Coordinators are ready to implement your complex study?</li> <li>• When there are multiple vendors involved within patient processes, how do you efficiently educate sites on these processes?</li> </ul>
<b>Long Screening Period/High Screen Failure</b>	<ul style="list-style-type: none"> <li>• How do you train sites to keep patients engaged during a long screening period?</li> <li>• When screen failure is high, how do you work with sites to keep them engaged?</li> </ul>
<b>Intense Market Competition</b>	<ul style="list-style-type: none"> <li>• When there is a lot of competition within a market, how do you differentiate your study?</li> <li>• How do you ensure sites keep your program top of mind?</li> <li>• When there are drugs already on the market for this disease, why does your drug matter and how should sites talk about it for recruitment?</li> </ul>
<b>Rare Disease</b>	<p>With rare diseases, the site may only see one appropriate patient per year.</p> <ul style="list-style-type: none"> <li>• How do you maintain quality when sites received training at the Investigator Meeting 12 to 18 months prior?</li> <li>• How do you maintain interest at the site level?</li> </ul>
<b>Rescue Programs</b>	<ul style="list-style-type: none"> <li>• How do you help sites accelerate patient recruitment when a study is behind schedule?</li> <li>• How do you improve site relations if there has been a history of starts and stops with the study and frustrating road blocks and changes?</li> <li>• What techniques do you use to regain commitment to your study?</li> </ul>
<b>Limited Budget/No F2F Investigator Meeting</b>	<ul style="list-style-type: none"> <li>• How do you efficiently train and motivate sites on your study when Face to Face is not an option?</li> <li>• How will you incorporate accelerated learning techniques into web meetings and online modules?</li> <li>• How will you measure information retention?</li> </ul>
<b>Repetitive Training</b>	<ul style="list-style-type: none"> <li>• How much time do you waste on repetitive training?</li> <li>• Can you trust your CRA to deliver accurate and consistent training across all sites and countries?</li> <li>• Is there a way for your content experts to train all sites concisely?</li> </ul>

*Creating partnerships for success. Enhancing programs for the future.*