

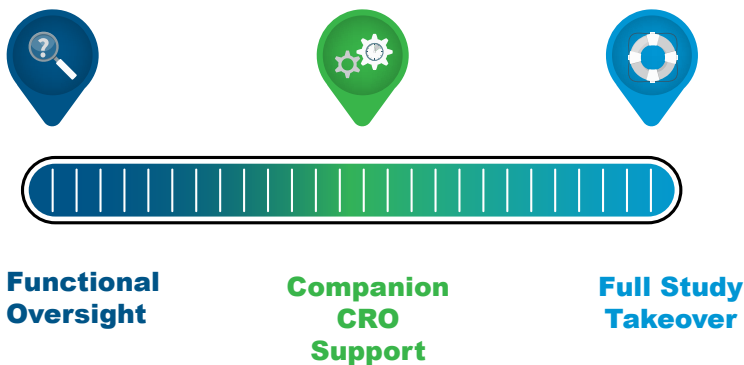
SOLUTION SHEET

Clinical Trial Rescue Services

PharPoint is a trusted partner for rescue studies, with the right experience, innovative solutions, and dedication to quality and on-time delivery to help sponsors get their trials back on track.

Consultative and client-focused, we're obsessed with your success – not your dollars. Rescues are stressful, and the PharPoint team is here to find you the most cost-effective and least stress-inducing fix possible.

Options for study rescues may include:



ABOUT PHARPOINT

PharPoint is an award-winning CRO that offers innovative sponsors of all sizes clinical operations, data management, biostatistics, statistical programming, and strategic clinical trial consulting services. We support studies from Phase 1 to Phase 4, with experience across a range of therapeutic areas.

Our successful rescue experience includes:

30+ Rescues

Since 2007, including biometric and/or clinical support for studies within therapeutic areas such as:



Explore **Rescue Case Studies**

Full-Service Neonatal Rescue

PharPoint took over all services for a Phase 1 cardiovascular study in a neonatal patient population.

Together, PharPoint's experts worked with the sponsor to address issues with their previous CRO. We set and accomplished clear, realistic timeline and budget goals, taking into consideration the complexities involved in enrollment for neonatal patient populations.

ISS/ISE in Rare Disease

PharPoint's biostatisticians were brought in as part of a rare disease rescue, taking on all statistical services.

Since being brought on as a rescue partner for this study, PharPoint has continued a successful relationship with the sponsor. This has included supporting ad hoc and DMC analyses, conducting oversight of other CROs, supporting an NDA submission and approval, and writing ISS/ISE plans and analyses.

Phase 3 Influenza Rescue

PharPoint's data management team handled a mismanaged trial being led by a big box CRO.

PharPoint's team created a comprehensive plan which included predicted timelines and a plan for communication and processes for quality checking that aided in an adequate transition. Our services included desinging a database that went live within 2 weeks of receiving information and over 1,100 forms were entered and cleaned within 6 weeks of the database going live.

[Learn more about our team & capabilities](#)

TRUSTED BY 180+ INNOVATIVE COMPANIES LIKE YOURS

To learn more about how PharPoint can support your study, schedule time to meet with our experts.

Recent Industry Recognitions:

