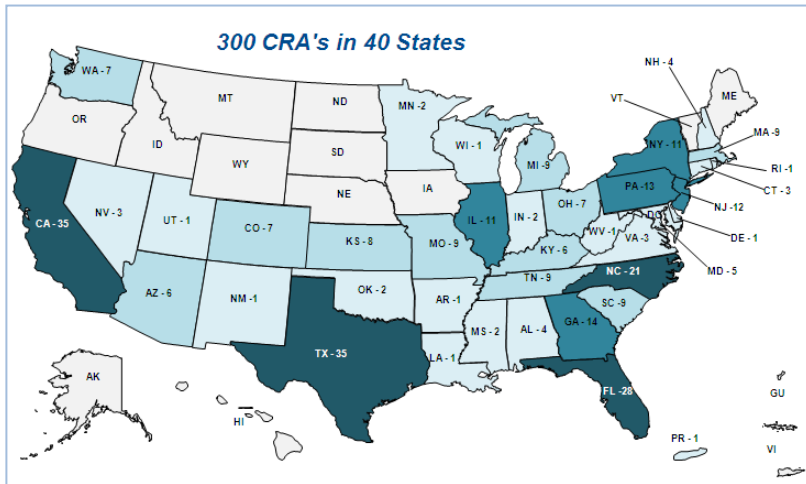


# LARGEST SINGLE CLIENT MONITORING PROGRAM

A Unique Redeployment Opportunity

## A GOLDEN OPPORTUNITY TO RESOURCE YOUR PROGRAMS

As the pioneer in functional outsourcing of regional monitoring, KCR created and managed the largest program in history for a single biopharmaceutical company in North America. More than 350 dedicated clinical research professionals managed 175 studies at 4,500 sites and involving more than 11,000 patients – approximately two-thirds of one of the largest pharmaceutical company's entire portfolio in North America.

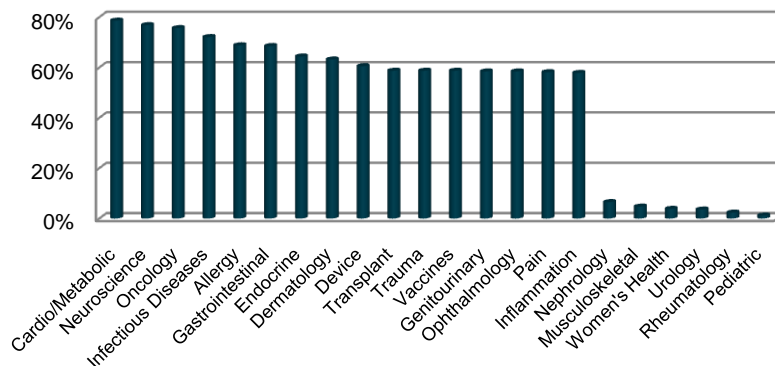


For more than 15 years, we have attracted and retained some of industries' best talent. Due to global vendor consolidation, there is now a unique opportunity to access more than 300 highly experienced clinical monitors and nearly 50 individuals trained in study and line management across the U.S., Canada and Puerto Rico.

## UNPARALLELED ONCOLOGY EXPERIENCE

Oncology monitors complete rigorous training on quality data review for tumor assessment evaluations, adverse event reporting and dosing compliance calculations. Our proprietary training program equips monitors with a realistic understanding of oncology nuances and details that impact clinical, regulatory and research efforts

## BROAD THERAPEUTIC EXPERTISE



## Our Candidates

- Average 15 years of clinical trial and 8 years monitoring experience
- 20% have advanced degrees
- Experience spans all phases and therapeutic areas
- More than 75% have been employees of KCR for at least three years

## Experienced Managers and Directors

- 23 Managers averaging 15 years
- 5 Directors averaging 20 years

## Proven Excellence

- Won client's Supplier Excellence Award in 2009
- During past two years, met all study-start milestones an average of 17 days early and reduced site selection cycle time 50%

## Oncology Expertise

- 65% of all monitors have oncology experience; 30% have Phase I oncology experience
- Oncology monitors average 14 years experience

## About PharmaNet/i3

PharmaNet/i3, the inVentiv Health clinical segment, is recognized as a leading provider of global drug development services to pharmaceutical, biotechnology, generic drug, and medical device companies, including therapeutically-specialized capabilities for Phase I-IV clinical development, bioanalytical services, and staffing from a single clinical professional to an entire functional team. For intelligent solutions needed to accelerate high quality drug development programs of all sizes around the world, *PharmaNet/i3 works for you*. For more information, visit [www.pharmanet-i3.com](http://www.pharmanet-i3.com).

**PharmaNet/i3**  
Strategic Resourcing

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