PROMPT
PONV RESCUE OUTCOMES AFTER
AMISULPRIDE TREATMENT

PROMPT Study - Site Enrollment Details

Thank you for completing the initial enrollment form for the PROMPT Study via the PACE Registry. In order to route your information correctly through the onboarding process, we have a few more questions.

As a reminder, sites interested in participating should be willing to meet these and other minimum criteria:

- Estimated surgical volume of approximately 1,000 cases per month
- Routine pre/peri-operative antiemetic prophylaxis
- Standard collection EMR-appropriate data points (ICD, CPT, SNOMED, others)
- Adequate IT resources to permit timely retrospective data collection
- Adequate research infrastructure to complete the necessary contracting for initiation of study within 3 to 6 months.

Key Contacts

Anesthesia Department Chair
Name:
Email:
Phone:

Local Technical Support
Name:
Organization (if different from site):
Email:
Phone:

EMR Vendor Technical Support
Name:
EMR Organization:
Email:
Phone:

Site Details

What type of Hospital or Facility best describes your site?

- ❑ Inpatient
- ❑ Outpatient
- ❑ Acute Care
- ❑ Long-Term Care
- ❑ Psychiatric Care
- ❑ VA
- ❑ Ambulatory Surgery Center

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- ❑ Community
- ❑ Rural
- ❑ Other, please specify _____

What EMR system does your site use?

- ❑ EPIC
- ❑ Cerner
- ❑ Greenway
- ❑ NextGen
- ❑ Unknown
- ❑ Other, please specify _____

How many surgical cases does your site see per month?

How many cases of PONV do you treat in the PACU per month?