iEnvision

Research solutions – ISR

iEnvision® ISR

Does this sound familiar...

You are spending an increasing amount of time gathering the information you need to be able to report on your investigator sponsored research (ISR) program, all because your system does not have what you need?

You would love a holistic view of your ISR program, but global and local markets as well as business units are using different systems, or no system at all, so you're unable to truly reflect on what's going on?

Your medical affairs colleagues are increasingly asking for visibility of your ISR program, but providing them with the information they need isn't easy?

The solution for the management of research requests or evidence generation programs

As your ISR program continues to grow, you need a system that can grow with you, and if you've reached a point where you need to streamline your processes so they're not holding you back – you're not alone!

We understand that the identification and generation of evidence beyond clinical trial data is a significant component of any pharmaceutical company's medical affairs activity.

Envision Pharma Group has provided purpose-built solutions to support research requests and the management of evidence generation since 2006. Our technology solutions are used by 19 out of the top 20 global pharmaceutical companies every day.

Adding value to your organization

Save time

Easy-to-use web portal collects all required data, reducing the need to request or find additional information

Efficient and compliant progression to publication/data disclosure

Ability to enter publication plans and data can be automatically pushed into Datavision™ for planning purposes

Effective high-level study program management

Ability to align projects to medical objectives/strategies to determine data gaps and report appropriately

Progress projects efficiently

Auto notifications triggered by status or date (eg, Project Update Notification, institutional review board [IRB] expiration)

Confidence in compliance with the ability to investigate queries

Documentation of all reviews, reconciliation, and close out processes with a full audit trail of all changes and organizational checks

Effective study portfolio performance

Key performance indicators are visible on real-time dashboards (eg, number of submissions, approval rate, number of publications)

Ensure all relevant steps in your project are covered from day one

A standardized workflow in line with industry standards and can be configured to company standard operating procedures

Strategic alignment and elimination of redundancies

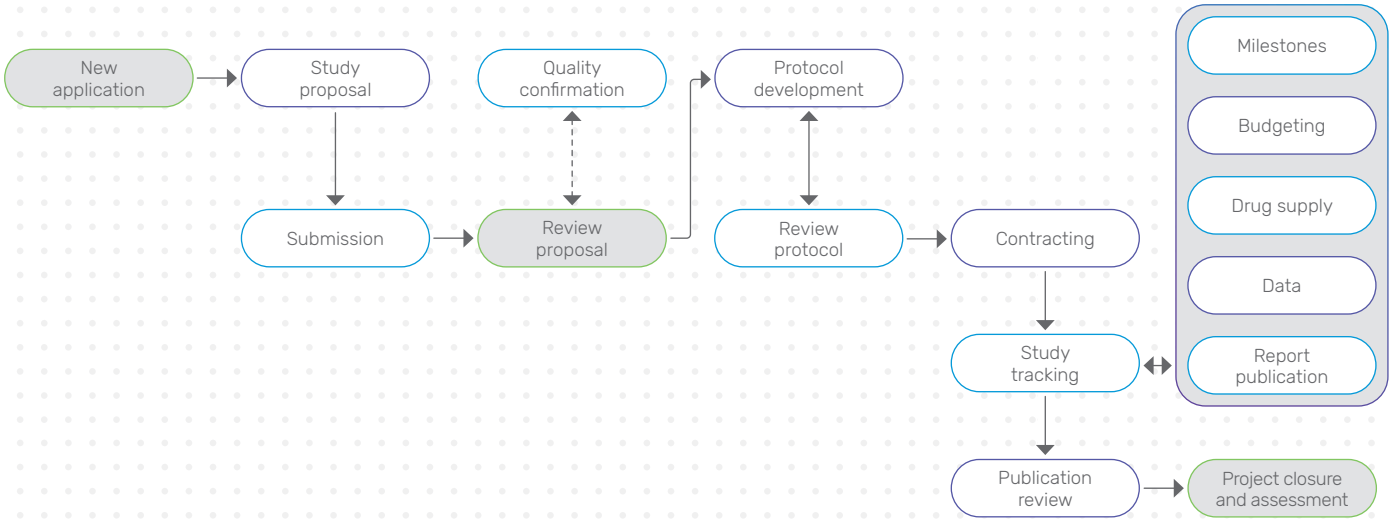
Allows local and global visibility into all studies and outcomes

Ensure the currency and accuracy of your reported information

Ability to filter and develop real-time ad hoc reports

ISR workflow

Helping your business achieve its external evidence goals



iEnvision – The global medical affairs platform

The rapidly evolving medical affairs functional landscape and its growing importance within pharmaceutical companies is accompanied by an increasingly complex world of stakeholder and partner interactions, connections, and compliance requirements.

iEnvision is an advanced software platform supporting medical affairs-led business transformation, operational excellence, and connectivity – purpose-built to help strategically plan and manage medical evidence generation, grant programs, and communication activities.

IMPROVES **EFFICIENCY**

- Reduce internal support and training needs
- Complete processes in a timely manner
- Achieve seamless collaboration

AIDS **COMPLIANCE**

- Align processes with industry standards
- Create auditable project records
- Maintain visibility and governance of both global and local activities

DELIVERS **IMPACT**

- Identify gaps in evidence
- Achieve organizational goals
- Align medical and communication plans with strategy

ENVISION PHARMA GROUP

For further information, or to arrange a demonstration, please contact **enquiries@envisionpharmagroup.com**

envisionpharmagroup.com