iEnvision

Communications solutions – Clear

iEnvision Clear®

Does this sound familiar...

You are frustrated by the lack of global visibility of your local affiliate publications?

With no formal system for the release of approved scientific documents, you're concerned about compliance issues?

Your publication managers are overloaded with the manual tracking of non-globally driven publications?

The solution for intuitive, system-managed clearance and approvals

If the compliant, timely, and appropriate communication of evidence is a key responsibility for you, you will no doubt appreciate the need to have a system that provides you with consistency across multiple business areas.

We understand that the automatic processing of clearance and disclosure requests significantly reduces administrative burden, improves efficiency, and ensures compliance.

Envision Pharma Group has provided purpose-built solutions to support the clearance and approval process since 2012. Our technology solutions are used by 19 out of the top 20 global pharma companies every day.

Adding value to your organization

A consistent process for approval of multiple document types

System-managed approval of corporate, promotional, and scientific materials and standard operating procedure (SOP) exceptions

Understand the volume and nature of locally-managed publications

Portfolio view summaries of publication types, origins, and volumes

Capture needs assessments for publications before their initiation

Configurable workflows for specific SOPs

Ease-of-use for submitters, reviewers, and approvers to manage their workloads

Rules-based workflows, and task-based workbenches, ensuring the right people are involved and informed at the right times

Auditable transparency on the approval of materials for external use

Full documentation of approval and clearance for release by medical, legal, and/or regulatory stakeholders

Maximize available time for senior executives

Auto-notifications and deep links to required actions simplify the review and approval process

Minimize administrative time for submitters, reviewers, and approvers

Intuitive, system-led process and interface with minimal training requirements

Adherence to company SOPs

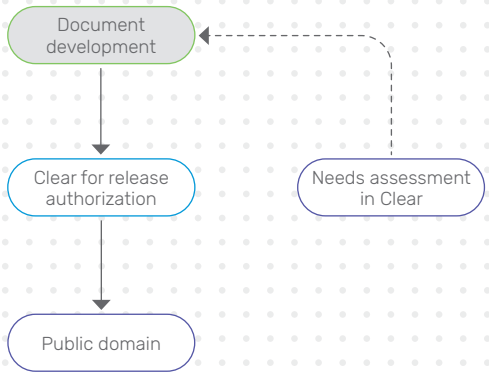
Workflow validation rules ensure required information is entered and appropriate review/approval processes are followed

Reduce approval cycle times and streamline reviewer workload

Single location for resources to perform review/approval tasks

Clear workflow

Helping your business achieve its clearance and approval 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