BP LOGIX Product Sheet

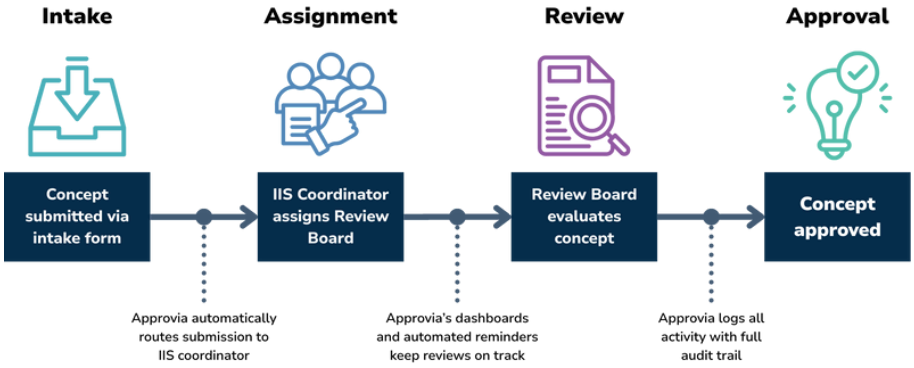
Approvia for Investigator-Initiated Studies

Too many organizations still rely on spreadsheets, email chains, and shared drives to manage IIS submissions. These manual methods slow down approvals, frustrate stakeholders, and increase risk.

As real-world evidence becomes more valuable and regulatory expectations grow, medical affairs teams need a better way to review and approve IIS concepts.

Accelerate investigator-led research with **Approvia**

Approvia streamlines the review and approval of IIS concepts while cutting administrative burden and reducing compliance risk.



Key features

No-code configuration
Customize review & approval workflows and user roles to suit your unique SOPs, no IT intervention required.

Always-on audit tracking
Get full visibility into every step of the approval process, ensuring a full audit trail to prove regulatory compliance.

Seamless integration
Connect your existing tech stack and forms to reduce manual data entry and ensure continuity across systems.

Real-time dashboards
Monitor review progress in real-time, allowing stakeholders to maintain transparency and accountability.

Automated reviewer assignments
IIS concept submissions automatically routed to the appropriate reviewers based on predefined parameters.

Manual IIS approvals slow down research, increase risk, and waste time. Approvia changes that.

Request a demo today.