BP LOGIX Product Sheet

PubPro for Global Clearance Processes

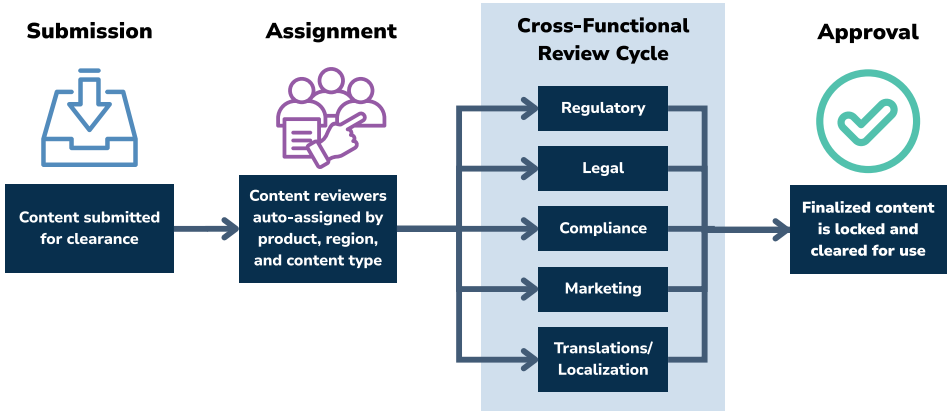
The clearance process is critical — and chronically broken. Medical affairs teams at large pharma and med device companies are stuck managing cross-functional reviews with spreadsheets, siloed tools, and endless email chains.

The result? Delays, duplicated work, compliance risk, and zero visibility into where things stand. Bottlenecks multiply when legal, regulatory, brand, and compliance teams are spread across time zones and systems with no shared source of truth.

PubPro Clearance was built to fix that.

Simplify the global clearance process with **PubPro**

PubPro Clearance Edition centralizes clearance workflows so companies can speed up reviews and stay audit-ready.



Key features

Automated reviewer assignments
Tasks are routed to the appropriate reviewers and stakeholders based on product, region, or content type.

Cross-departmental collaboration
Legal, regulatory, medical, and brand reviewers provide feedback in one shared environment.

Role-specific dashboards
Reviewers only see the tasks, deadlines, and statuses relevant to them. No clutter, just clear next steps.

Built-in reporting and analytics
Monitor timelines, identify delays, and demonstrate compliance with full audit trails and on-demand reporting.

No-code configuration
Admins can adapt clearance workflows and user roles to fit SOPs — no IT involvement or custom dev required.

Learn more about automating your clearance system.

PubPro

Request a demo today.