SecureCHEK® AI™

ACCURACY / COMPLIANCE / CONFIDENCE

Get Launch-Ready. Stay Market-Ready.

Turn product approval into market momentum with AI-powered medical, legal, and regulatory (MLR) readiness.

A product launch creates an immediate race against the clock. Teams must translate approved labeling and scientific evidence into a growing volume of promotional, educational, and scientific communications—across brands, audiences, channels, and markets. Every claim must be substantiated, every reference accurate, and every piece of content compliant.

Yet manual MLR processes, fragmented evidence, and repeated rounds of revisions can slow approvals, increase costs, and delay the impact of a launch.

SecureCHEK AI helps life sciences teams build a stronger compliance foundation before launch—and move faster once the product reaches the market.

Prepare the foundation. Accelerate the launch.

Whether preparing for a first commercial launch or managing an established portfolio, SecureCHEK AI helps you turn approved science into compliant, high-impact communications—faster and with greater confidence.

1 Establish a single source of truth

Build a centralized, reusable Claims Library from approved product labeling, scientific references, and existing approved materials. Organize claims and their supporting evidence so teams can maintain consistency, traceability, and visibility into approved content across products, audiences, and markets.

2 Identify issues before formal review

Run new materials through AI-powered precheck before submitting them to MLR. SecureCHEK AI identifies unsupported or inconsistent claims, missing or incorrect references, discrepancies with labeling, safety and fair-balance concerns, and other potential compliance issues. A Review Readiness Score helps teams understand what needs attention before submission.

3 Focus expert review where it matters most

Combine precision, rules-based analysis with GenAI reasoning grounded in structured claims and supporting evidence. Surface new claims, changed language, potential risks, and deviations requiring judgment—so reviewers can focus on scientific accuracy, risk assessment, and final decisions rather than preventable errors.

4 Scale from launch to lifecycle management

As the product expands into new indications, audiences, channels, and markets, reuse and update the claims foundation rather than repeatedly starting from scratch. Support consistent global standards while accommodating local labeling, language, regulatory requirements, and market-specific content.

The result: greater speed, confidence, and control.

Accelerate time to market

Make submission packages review-ready up to 70% faster.

Reduce rework and costs

Catch preventable errors earlier and reduce unnecessary revision cycles and external consulting needs.

Protect scientific integrity

Connect claims to substantiating evidence and identify potential inconsistencies before content reaches formal review.

Increase team capacity

Enable medical, legal, regulatory, and agency teams to spend less time correcting materials and more time on strategic decisions.

Extend value beyond launch

Create a foundation for efficient, consistent review as products, claims, and communication needs evolve.

Ready before launch. Built for what comes next.

SecureCHEK AI combines analytical, rules-based AI, GenAI reasoning, precision claims management, and human oversight. MLR teams retain responsibility for final review and approval.

BUILD THE LIBRARY. PRECHECK THE CONTENT. FOCUS REVIEW ON WHAT MATTERS.
GET TO MARKET WITH GREATER SPEED AND CONFIDENCE.