

SecureCHEK AI Streamlines MLR Review to Unlock Data Communications Compliantly, Efficiently

OPPORTUNITY:

A Pivotal Inflection Point to Scale Scientific Communication

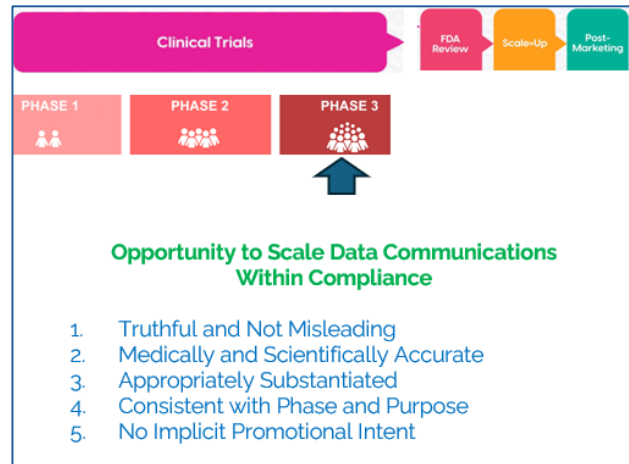
Successful completion of Phase 3 marks a critical transition from clinical development to pre-commercial readiness

- Scientific exchange supports awareness, education, and stakeholder engagement
- More content for medical and investor audiences will need to be reviewed and approved
- Cross-functional stakeholders play a more active role in shaping and approving messaging
- Reliance on external consultants may increase

CHALLENGES:

Manual MLR Processes Cannot Support Required Scale and Speed at Pivotal Stage

- Strict adherence to scientific exchange principles is required to decrease risk around pre-commercial communications
- Lean teams are forced to balance compliance rigor with speed, with many companies outsourcing reviews
- Without modernization, constraints increase rework and consulting costs, limit organizational agility



Deviations That Will Be Automatically Found in Materials

Red Flags

- Implications that the product is safe/effective before FDA approval, or forward-looking efficacy or safety claims; overly promotional tone or language used
- Inconsistent or not supported by reference documents and/or claims library
- Not aligned with clinical trial endpoints, data on file without adequate substantiation, or draft labeling (not available yet)
- Overemphasizing efficacy with limited discussion of risk/safety
- Overstating statistical significance, outcomes

General Deviations

- Corporate signoff incomplete
- Error in chart/table
- Grammar issues
- Incorrect product/generic name
- Lacking citations/support
- Misuse of logos, trademarks, or legal disclaimers
- Missing/wrong reference and footnote
- Missing disclaimers like "investigational" or "not FDA approved"
- Spelling and grammar

SOLUTION

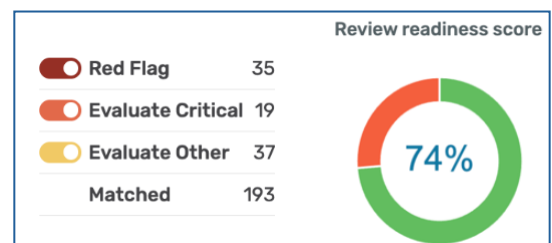
AI-Powered Intelligence for Fast, Scalable, MLR Review With Lower Consulting Costs in 3 Steps

1. Centralized Library establishes single source of truth for approved scientific and investor messaging, enabling consistent and reusable content
2. Automated Pre-Check of Content identifies compliance risks before review, reducing MLR time and external consulting costs (Review Readiness Score)
3. Focused Review to ensures higher-quality submissions, reducing cycles of revision and accelerating approvals

BENEFIT

Faster, Confident Dissemination of Scientific Data for Credibility, Readiness and Momentum

- Speeds up review ready submission packages by 70%
- Focuses reviewers entirely on content requiring adjudication and risk decisions
- Decreases MLR rework, increases MLR productivity
- Enable teams to focus less on compliance mechanics—and more on strategic communication



SecureCHEK AI Increases Efficiency and Productivity, Frees Up Time for Strategy and Creativity

