Pharma & Life Sciences
decision-intelligence done differently.

REGULATION ANCHOR: GMP (ICH Q10) · EU FALSIFIED MEDICINES DIRECTIVE · US DSCSA

API supplier integrity, sanctions, climate events on the single-source upstream.

WHAT THE SECTOR STRUGGLES WITH

- Single-source API suppliers in sanctioned geographies — and the compliance team finds out from a news alert.
- Climate events (flood, typhoon) disrupt upstream manufacturing; dependency isn't modelled in the GMP register.
- Sanctions screening is run on named counterparties, not **ultimate beneficial owners** up the chain.
- Quality-critical events don't have a £ risk framing for the board.

WHAT ERIO SHIPS OUT OF THE BOX

- Live n-tier supplier graph with sanctions + UBO overlays (GLEIF + Companies House + BYO D&B).
- Ingests GDACS flood/earthquake + ReliefWeb humanitarian alerts — maps onto your dependency graph.
- Open FAIR £ curves per material risk — quality excursions get a board-level number.
- Replayable evidence packs for GMP / FMD / DSCSA audits — inputs chained with the output.

"We used to find out from a news alert. Now ERIO tells us the day the ownership changed." Sector brief · Nov 2026