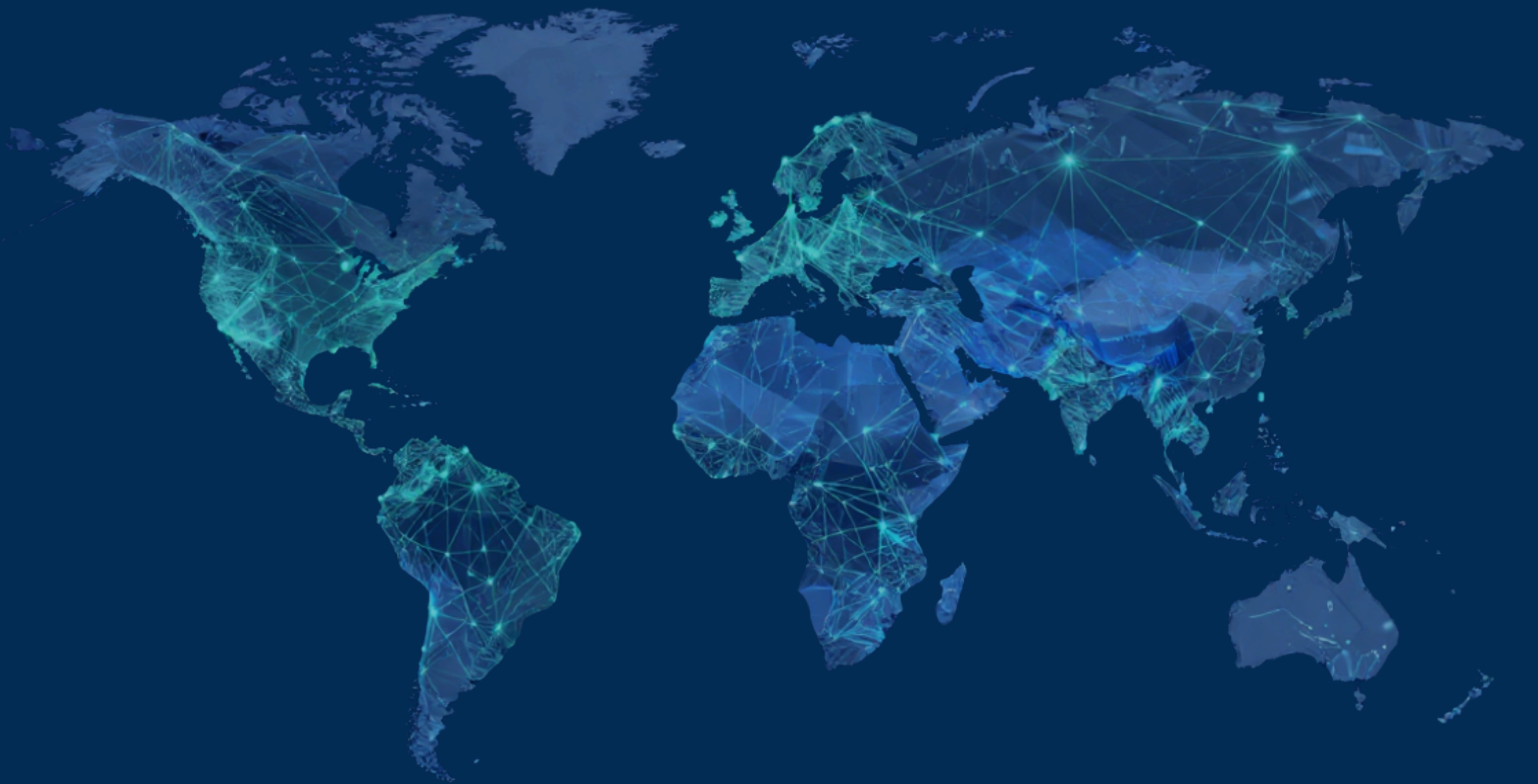


HEALTHCARE INTELLIGENCE

Why APAC Evidence Generation Is Falling Behind

The structural delays costing pharmaceutical organizations 18+ months in regional RWE delivery



Evidence generation in APAC is structurally broken

18+

months average delay
from protocol to publication

67%

of APAC RWE studies miss
regulatory submission windows

3.2x

longer cycle times vs.
US/EU evidence programs



The delay isn't scientific. It's operational.

- Fragmented data partnerships across 12+ APAC markets
- No standardized protocol-to-output infrastructure
- Manual coordination between Medical Affairs, HEOR, and vendors
- Regulatory heterogeneity treated as blockers, not design inputs
- Evidence plans disconnected from market access timelines

"Organizations are solving a systems problem with project-level thinking."



From fragmented projects to connected evidence infrastructure

01 DESIGN

Regulatory-aware protocol architecture that embeds market access from day one

02 CONNECT

Unified data partnerships across APAC with standardized governance

03 GENERATE

Accelerated evidence production through operational infrastructure

04 DELIVER

Synchronized output aligned to submission and launch timelines



What an integrated evidence engine actually looks like

- End-to-end RWE program design aligned to APAC regulatory landscapes
- Pre-built data source connectivity across 12+ markets
- Automated protocol-to-output workflows reducing cycle time by 40%
- Integrated Medical Affairs and Market Access evidence planning
- Real-time program visibility for regional and global stakeholders
- Scalable infrastructure that compounds with each new study



The cost of waiting is compounding

Delayed market access

Every month of evidence delay = lost formulary positioning and competitive ground in key APAC markets

Regulatory risk

Incomplete or late evidence packages trigger additional requirements, extending timelines by 6-12 months

Strategic misalignment

Disconnected evidence and commercial planning creates organizational friction at scale

"The organizations that build evidence infrastructure now will define APAC market leadership for the next decade."



Ready to accelerate your APAC evidence strategy?

Aurumetrics partners with pharmaceutical organizations to design, build, and operate integrated evidence infrastructure across APAC.

- Schedule an evidence infrastructure assessment
- Request the full APAC RWE landscape report
- Connect with our Medical Affairs team

aurumetrics.com



The challenge is no longer whether healthcare data exists.

It is whether the right evidence can be generated fast enough?

Yes, it can.

Aurumetrics supports pharmaceutical and life sciences teams with real-world data intelligence, strategic commercial insight and evidence generation.

GLOBAL MARKET FOOTPRINT

APAC-focused with evidence and data partnerships spanning:

China • Australia • Japan • South Korea • Taiwan • New Zealand

Additional support across the UK, EU, US, and other global markets on an ad hoc basis.

CAPABILITIES

Sourcing & Integrating
Real-World Data Globally

Advancing Analytical
Solutions

Delivering Strategic
Data Insights & Advisory

Building Powerful
Data Collaborations

Claims • EMR • Registry • GP • Lab • Genomics • Linked Data

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