Fact Sheet

IQVIA Vigilance Platform

Purpose-built PV technology

Enterprise grade pharmacovigilance, but right-sized for emerging biopharma and small pharma teams

Data ownership | Regulatory readiness | AI-ready platform | Right-sized investment

Taking control of your safety data

Emerging biopharma and small companies have long relied on CRO-hosted PV systems for cost-effective access. That model works early — but it limits the data that sponsors can access: the breadth of reports, the frequency of insights, and control over the full information chain. In today's environment, not having the data you need, when you need it, in the format you want is a material disadvantage for time to market and clinical and post-marketing safety.

As organizations grow and markets become more complex, these limitations drive rising operational cost and risk. More sponsors are investing in their own PV platform — not just a system that runs processes, but a modern platform that supports AI, transparency, and high-quality data access. IQVIA Vigilance Platform, for emerging biopharma and small pharma, lowers the barrier to entry with a standardized, right-sized approach that lets lean teams establish an audit-ready PV foundation early — without the burden of traditional enterprise implementations.

Build a strong pharmacovigilance foundation from day one

- **Own the data:** Pharma-owned safety database with full, direct access to your safety data — real-time visibility, not CRO-gated reports.
- **Deploy with focus:** Pre-configured platform and standardized operating model cut implementation complexity from months of custom build to weeks.
- **Operate with readiness:** Out-of-the-box reporting rules, E2B gateway support, and a validation package get you audit-ready from day one.
- **Scale as you grow**: Built for lean teams today on the same enterprise-grade IQVIA Vigilance Platform that scales with portfolio and case volume.

What's included

- **Platform setup:** Products, licenses, studies, SSO, security model, standard workflows.
- **PV operations:** Case intake and processing, duplicate search, tasking, dashboards, worklists.
- **Regulatory submissions:** E2B gateway to FDA, EMA and MHRA; line listings and tabulations.
- **Validation and support:** Validation package, gateway configuration support, 2-week hypercare.

Fast implementation

$49K implementation fee

12-15 weeks + 2 weeks hypercare

SaaS Subscription

$65K annually

3-year term up to 1500 cases/yr*
includes 10 users

Built for:

Sponsors ready to own their safety data, simplify their first PV system, and build a growth-aligned foundation — without overbuying enterprise complexity

IQVIA PV technology expertise

30+	20	12 million	220	400+	200+
Vigilance platform customers SaaS solutions	**Customers in top 50** 6 HQ in Japan 8-9 global HQ	**Production cases hosted** Largest: 600K+/yr Smallest: 5/mo	**Support technicians** Global and Japan support 35 acct managers	**PV tech personnel** Dev and delivery across multiple regions	**Technology customers** 47 single-tenant 149 multi-tenant 8 on-prem

*We offer additional tiers for higher case volumes

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