



EMPOWERING DRUG SAFETY THROUGH END-TO-END PHARMACOVIGILANCE SERVICES



Lambda Therapeutic Research provides comprehensive drug safety & pharmacovigilance services, encompassing all clinical and post-marketing phases.

Our well-established PV framework ensures thorough safety monitoring and regulatory compliance for both clinical and commercial products, offering **end-to-end support across the entire life cycle of medicinal products.**



THE **LAMBDA** ADVANTAGE

Experience & expertise in major Therapeutic Areas & Products:

- **Drugs**
- **Vaccines**
- **Medical Devices**
- **Biologics & Biosimilars**
- **Advanced therapy medicinal products**
- **Combination Products**

Year 2022 at a Glance

86000+

Cases

1200+

Signal
Management
Reports

684,000+

Literature
Screening

Over **99%**

Submission
Compliance





PvEDGE

- Advanced and largely automated **end-to-end PV solution**.
- **Regulatory-compliant** web-based customizable safety database.
- **21 CFR Part 11** compliant & based on ICH E2B specification.
- Supports **MedDRA** and **WHO-DD** browsers.
- **E2B R2** and **R3** compliant database.
- Enabled with **literature automation** and **signal detection** modules.



Ensuring safety & compliance worldwide with team of 150+ PV Professionals



UK-HQ
(UK QPPV & MI Services)

POLAND
(EU QPPV, LRP & MI Services)

INDIA
(PV Delivery & MI Services)

CANADA
(MI Services)

Unlock seamless collaboration with Lambda's expert team and gain access to a comprehensive range of cutting-edge drug safety and pharmacovigilance services.

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