

PHARMACOVIGILANCE SERVICES

**Cloud based, E2b(R2)/(R3) complaint database
with integrated
Automated Literature Module and Signal Detection
Using Artificial Intelligence**

This project will be undertaken across five core phases:

Phase 1: PLANNING - Project start up activities

- Establishing an organogram
- Identifying & defining Roles & Responsibilities (Job descriptions)
- Developing Standard Operating Procedures & Work Instructions
- PSMF or Drug Safety System
- Preparation of Risk Management Plan

Phase 2: PHARMACOVIGILANCE SET UP (INITIAL/ANNUAL) COST

The fee for various requirements for Pharmacovigilance set up as per the European Pharmacovigilance guidelines are as follows:

Table 2.1: One time and annual maintenance cost (To be finalized as per client requirement)

Activities	First Time (INR)	Annual Maintenance (INR)
Safety Database Configuration Implementation & Validation	450000	100000
Literature monitoring setup- Configuration of search criteria in literature data base for each set of 10 brands	10000	NA
MedDRA and WHO DD subscription	75000	Depending on new subscription from MedDRA and WHO DD
Preparation of Client specific SOPs & Work Instruction (Per Doc)	8000	5000 for each revision required
Preparation of Pharmacovigilance System Master File (PSMF)	100000	15000
Training of client employees	75000	20000
Record Management	50000	20000
Optional Services		
MICC installation and One time Setup cost	450000	NA
<ul style="list-style-type: none"> • For 100 cases • For 250 cases • For 500 cases 		450000PA/ 45000 PM 600000PA/ 60000 PM 800000PA/ 80000 PM

Activities		First Time (INR)	Annual Maintenance (INR)
Preparation of Risk Management Plan (per document)	For India & ROW	100000	NA
	For EU/USFDA	150000	NA
XEVMPD- For EU region only, if required		35000	NIL
Update for each brand in XEVMPD		2000	NIL
EVWEB registration (Required for EU market)- Optional		50000	NA
Submission Gateway(per gateway)		150000	NA
Data Migration		Depends on amount of data	Data Migration
Clinical Trail Database One Time Setup <ul style="list-style-type: none"> • ECD • CTMS 		450000	50000 Per study per month 60000 Per Study Per month

Phase 3: REAL TIME PV ACTIVITIES

Collection of ADR, Case Processing and Reporting

Collection of ADRs is to be done in the through **MICC** or from patients/relatives/ medical and paramedical staff and Clinical Trials followed by Processing of ICSRs and Expedited Reporting. The cost of each activity is as follows:

Table 3.1: Fee for ADR collection through MICC (If required)

Activities	Fee (INR)
Up to 100 queries/ Month	45000 Per Month
From 101 Up to 250 queries/ Month	60000 Per Month
From 251 queries onwards/ Month	80000 Per Month

Table 3.2: Fee for ADR collection through literature monitoring, **charged on monthly basis Mandatory activities**

Activities	Fee (INR)
1) Weekly Searches through mail alert, random searches, duplicate check, tracking, repository and over all data collation from literature databases	400/articles

Activities	Fee (INR)
2) Literature evaluation of articles, obtained in step 1, for valid and invalid cases	

Note: Articles to be purchased for identified valid cases will be charged separately as per actuals.

Table 3.3: Fee for processing and reporting of ADRs collected from **all applicable sources (Call Centre/ Literature Monitoring/ Clinical Trials/ Health Authority/ Business Partner)**

Activities	Fee (INR)
Clinical Trail Cases	7500/Case
Serious ICSRs - Case processing and reporting from all sources	6000/Case
Non-Serious ICSRs - Case processing and reporting from all sources	5000/Case

Phase 4: AGGREGATE REPORTING

PSURs, PBRER, PADER etc. will be done as per regulatory requirements on annually/biannually/quarterly bases

Table 4.1: Step 1- Screening of literature for ICSRs per **molecule through monthly reconciliation**

S. No.	Activity type	Activities	Fee (INR)
1	Core	1) Screening, collection of cases from literature databases for each Brand 2) Identification of valid and invalid cases for cases collected in step 1	400/articles
2	Core	Serious ICSRs - Case processing from all sources (Cost/case)	6000
3	Core	Non-Serious ICSRs - Case processing from all sources (Cost/case)	5000

Table 4.2: Step 2- Preparation of periodic reports/updates

Sr. No	Activity type	Preparation/compilation of AGGREGATE REPORTS	First Report (INR)	Subsequent Reports (INR)
1	Core	PSUR (per document)* for Indian and ROW markets	70000	50000
2	Core	PBRER/PADER (per document)*	250000	150000

Phase 5: RISK MANAGEMENT ACTIVITIES

Table 5.1 Signal Detection

Activities	Fee (INR)
Signal Management and Reporting per brand annually as and when required	125000

Phase 6: QPPV services

In order to perform the above activities, services of following key personnel will be billed to the sponsor.

Table 6.1: Fee for QPPV personals

Description	Fixed Cost/Month (INR)
QPPV Fee	To be discussed as per requirement

INVOICE SCHEDULE

The below is a proposed invoicing schedule, for which we will be happy to discuss alternative options.

Table 7.1: Invoice schedule for one time cost

Invoice	Percent (%)	Invoice Point	Estimated Invoice Date
1	60%	Upon contract signature	TBD
2	40%	Upon set up of PV system	TBD

Table 7.2: Invoice schedule for monthly variable cost

Invoice	Percent (%)	Invoice Point	Estimated Invoice Date
1	100%	End of the month	8 th of every month

Taxes as applicable would be charged extra.

For Indivirtus Healthcare Services Pvt. Ltd.

Indivirtus Healthcare Services (P) Ltd.

 Director

(Dr. Upendra K Jain)