



SANDHYASHI NEURO PANCHKARMA

(A unit of Sandhya Pharma and Research Unit)
(BF-45, Shalimar Bagh, Delhi-110088)

SUSPECTED ADVERSE DRUG REACTION REPORTING FORM

For VOLUNTARY reporting
of Adverse Drug Reactions
by health care professionals

Report #

To be filled in by Pharmacovigilance
centres receiving the form.

A. Patient information

1. Patient identifier initials _____	2. Age at time of event: or _____ Date of Birth: _____	3. Sex: <input type="checkbox"/> M <input type="checkbox"/> F
In confidence	4. Weight _____ Kgs	

B. Suspected Adverse Reaction

5. Date of reaction started (dd/mm/yy):
6. Date of recovery (dd/mm/yy):
7. Describe reaction or problem

12. Relevant tests/ laboratory data, including dates

13. Other relevant history, including pre-existing medical conditions
(e.g., allergies, race, pregnancy, smoking alcohol use, hepatic/
renal dysfunction, etc.)

14. Seriousness of the reaction

- | | |
|---|--|
| <input type="checkbox"/> Death (dd/mm/yy) _____ | <input type="checkbox"/> Congenital anomaly |
| <input type="checkbox"/> Life threatening | <input type="checkbox"/> Required intervention |
| <input type="checkbox"/> Hospitalization-initial or prolonged | <input type="checkbox"/> to prevent permanent impairment/ damage |
| <input type="checkbox"/> Disability | <input type="checkbox"/> Other (specify) _____ |

15. Outcomes

- | | | |
|-------------------------------------|-------------------------------------|--|
| <input type="checkbox"/> Fatal | <input type="checkbox"/> Recovering | <input type="checkbox"/> Unknown |
| <input type="checkbox"/> Continuing | <input type="checkbox"/> Recovered | <input type="checkbox"/> Other (specify) _____ |

C. Suspected medication(s)

Sl. No.	8. Name (brand and / or generic name)	Manufacturer (If known)	Batch No. / Lot No. (If known)	Exp. Date (If known)	Dose used	Route used	Frequency	Therapy dates (if unknown, give duration)		Reason for Use or prescribed for
								Date started	Date stopped	
i										
ii										
iii										
iv										

9. Reaction abated after drug stopped or dose reduced

Sl. No. As per C	Yes	No	Unknown	NA	Reduced dose
i					
ii					
iii					
iv					

10. Reaction reappeared after reintroduction

Sl. No. As per C	Yes	No	Unknown	NA	If reintroduced, dose
i					
ii					
iii					
iv					

11. Concomitant medical products and therapy dates including self medication and herbal remedies (exclude those used to treat reaction)

D. Reporter (see confidentiality section in first page)

16. Name and Professional Address: _____

Pin code: _____ E-mail: _____

Cell No. / Tel. No. with STD Code: _____

Speciality: _____ Signature: _____

17. Occupation

18. Date of this report