

SUSPECTED ADVERSE DRUG REACTION REPORTING FORM

For VOLUNTARY reporting of Adverse Drug Reactions by Healthcare Professionals

<div>INDIAN PHARMACOPOEIA COMMISSION</div> <div>(National Coordination Centre-Pharmacovigilance Programme of India)</div> <div>Ministry of Health &amp; Family Welfare, Government of India</div> <div>Sector-13, Plot-18/19, Connaught Place, New Delhi-110028</div>							FOR AMC/NCC USE ONLY			
A. PATIENT INFORMATION							AMC Report No. _____			
1. Patient Initials _____		2. Age at time of Event or Date of Birth _____		3. M <input type="checkbox"/> F <input type="checkbox"/> Other <input type="checkbox"/>			Worldwide Unique No. _____			
				4. Weight _____ Kgs			12. Relevant tests/ laboratory data with dates _____			
B. SUSPECTED ADVERSE REACTION							13. Relevant medical/ medication history (e.g. allergies, race, pregnancy, smoking, alcohol use, hepatic/renal dysfunction etc.) _____			
5. Date of reaction started (dd/mm/yyyy) _____							14. Seriousness of the reaction (Yes <input type="checkbox"/> No <input type="checkbox"/> <input type="checkbox"/> Death (dd/mm/yyyy) <input type="checkbox"/> Congenital anomaly <input type="checkbox"/> Life threatening <input type="checkbox"/> Required intervention to prevent permanent impairment/damage <input type="checkbox"/> Hospitalization/Prolonged <input type="checkbox"/> Disability <input type="checkbox"/> Other (specify) _____			
6. Date of recovery (dd/mm/yyyy) _____										
7. Describe reaction or problem _____										
C. SUSPECTED MEDICATION(S)							15. Outcomes <input type="checkbox"/> Recovered <input type="checkbox"/> Recovering <input type="checkbox"/> Not recovered <input type="checkbox"/> Fatal <input type="checkbox"/> Recovered with sequelae <input type="checkbox"/> Unknown			
S.No	B. Name (Brand/Generic)	Manufacturer (if known)	Batch No. / Lot No.	Exp. Date (if known)	Dose used	Route used	Frequency (OD, BD etc.)	Therapy dates		Indication
								Date started	Date stopped	
i										
ii										
iii										
iv										
5. No. as per C	9. Action Taken						10. Reaction reappeared after reintroduction			
	Drug withdrawn	Dose increased	Dose reduced	Dose not changed	Not applicable	Unknown	Yes	No	Effect unknown	Dose (if reintroduced)
i										
ii										
iii										
iv										
11. Concomitant medical product including self medication and herbal remedies with therapy dates (Exclude those used to treat reaction)							D. REPORTER DETAILS			
17. Causality Assessment:  Additional information: _____							16. Name and Professional Address: _____ Pin: _____ E-mail: _____ Tel. No. (with STD code): _____ Occupation: _____ Signature: _____			
							18. Date of this report (dd/mm/yyyy): _____			
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# How To Report Adverse Drug Reaction

**Michael Brown**



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## **Table of Contents How To Report Adverse Drug Reaction**

1. Understanding the eBook How To Report Adverse Drug Reaction
  - The Rise of Digital Reading How To Report Adverse Drug Reaction
  - Advantages of eBooks Over Traditional Books
2. Identifying How To Report Adverse Drug Reaction
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an How To Report Adverse Drug Reaction
  - User-Friendly Interface
4. Exploring eBook Recommendations from How To Report Adverse Drug Reaction

- Personalized Recommendations
- How To Report Adverse Drug Reaction User Reviews and Ratings
- How To Report Adverse Drug Reaction and Bestseller Lists
- 5. Accessing How To Report Adverse Drug Reaction Free and Paid eBooks
  - How To Report Adverse Drug Reaction Public Domain eBooks
  - How To Report Adverse Drug Reaction eBook Subscription Services
  - How To Report Adverse Drug Reaction Budget-Friendly Options
- 6. Navigating How To Report Adverse Drug Reaction eBook Formats
  - ePub, PDF, MOBI, and More
  - How To Report Adverse Drug Reaction Compatibility with Devices
  - How To Report Adverse Drug Reaction Enhanced eBook Features
- 7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of How To Report Adverse Drug Reaction
  - Highlighting and Note-Taking How To Report Adverse Drug Reaction
  - Interactive Elements How To Report Adverse Drug Reaction
- 8. Staying Engaged with How To Report Adverse Drug Reaction
  - Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - Following Authors and Publishers How To Report Adverse Drug Reaction
- 9. Balancing eBooks and Physical Books How To Report Adverse Drug Reaction
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection How To Report Adverse Drug Reaction
- 10. Overcoming Reading Challenges
  - Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
- 11. Cultivating a Reading Routine How To Report Adverse Drug Reaction
  - Setting Reading Goals How To Report Adverse Drug Reaction
  - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of How To Report Adverse Drug Reaction

- Fact-Checking eBook Content of How To Report Adverse Drug Reaction
- Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
  - Utilizing eBooks for Skill Development
  - Exploring Educational eBooks
- 14. Embracing eBook Trends
  - Integration of Multimedia Elements
  - Interactive and Gamified eBooks

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