

**ADVERSE EVENT REPORTING FORM**

<b>Patient Details:</b>								
Patient's Name: _____						Country: _____		
Age/Date of Birth: _____				Weight: _____ Kg.		Height: _____		
Sex: <input type="checkbox"/> Male			<input type="checkbox"/> Female (Pregnancy Status: _____)					
<b>Adverse Event Details:</b>								
Date of Adverse Event: _____					Adverse Event Description:			
Seriousness:								
<input type="checkbox"/> Death			<input type="checkbox"/> Life Threatening					
<input type="checkbox"/> Hospitalization- Initial /Prolonged			<input type="checkbox"/> Disability					
<input type="checkbox"/> Congenital-anomaly			<input type="checkbox"/> Other Medically Important					
<input type="checkbox"/> Required intervention to prevent permanent impairment/ damage								
<b>Adverse Event Outcome:</b>								
<input type="checkbox"/> Recovering			<input type="checkbox"/> Unknown			<input type="checkbox"/> Not Recovered		
<input type="checkbox"/> Recovered with sequelae			<input type="checkbox"/> Recovered (Recovered Date: _____)			<input type="checkbox"/> Fatal (Date of Death: _____)		
<b>Related Tests/ Laboratory data(If available):</b>					<b>Other Medical History(Including Pre-existing Condition):</b>			
<b>Suspected Medication Details:</b>								
S/ No.	Product Name	Batch/Lot No.	Mfg. Date	Exp. Date	Dose and Frequency	Therapy Dates		Reason for Use/ Indication
						Start Date	Stopped Date	
1.								
2.								
Action taken after reaction: <input type="checkbox"/> Drug withdrawn <input type="checkbox"/> Dose reduced <input type="checkbox"/> Dose increased <input type="checkbox"/> Dose continued								
Reaction abated after use stopped or dose reduced : <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not applicable								
Reaction reappeared after reintroduction: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not applicable								
<b>Concomitant Medication Details (Excluding Treatment of Reaction):</b>								
S/ No.	Product Name	Batch/Lot No.	Mfg. Date	Exp. Date	Dose and Frequency	Therapy Dates		Reason for Use/ Indication
						Date Start	Date Start	
1.								
2.								
3.								
4.								
5.								
<b>Reporter Details:</b>					<b>To be Filled by National Healthcare:</b>			
Name: _____					Date of Receipt of Adverse Event: _____			
Profession: _____					Name and sign of Receiver: _____			
Contact No. : _____					Report type: <input type="checkbox"/> Initial <input type="checkbox"/> Follow up			
E-mail: _____					<b>Where to Report:</b>			
Address: _____					<b>Mail to:</b> National Healthcare Pvt. Ltd., Chhatapipra-3, Bara, Nepal.			
Date of Reporting: _____					<b>Phone No.:</b> +977-51-580236, +977-51-580508			
Signature: _____					<b>or Email at:</b> bpm@nationalhealthcare.com.np			
<b>Confidentiality:</b> The identity of patient shall be kept in strict confidence and protected to the fullest extent. No any legal implication on reporter upon submission of ADR reports.								