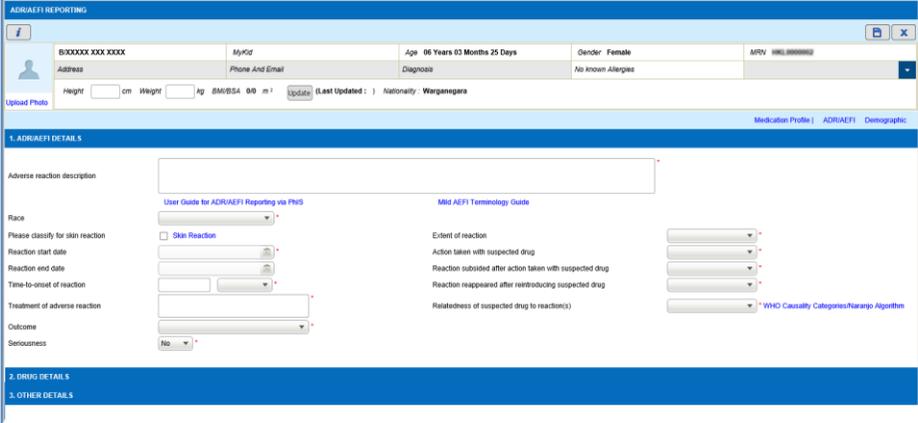
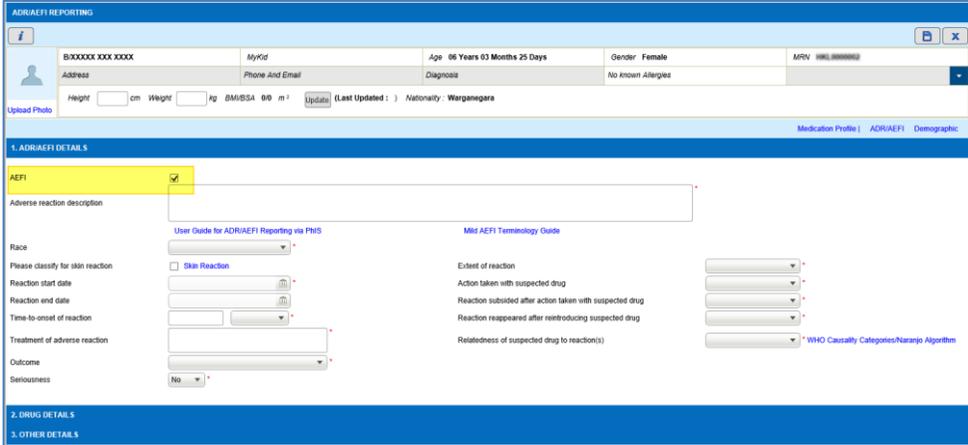


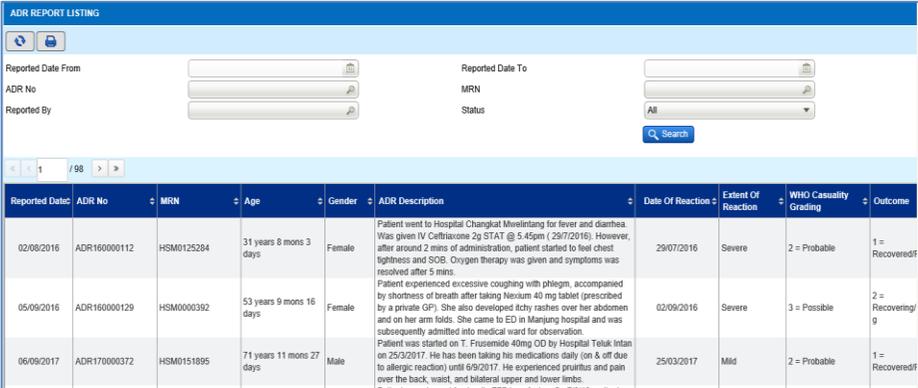
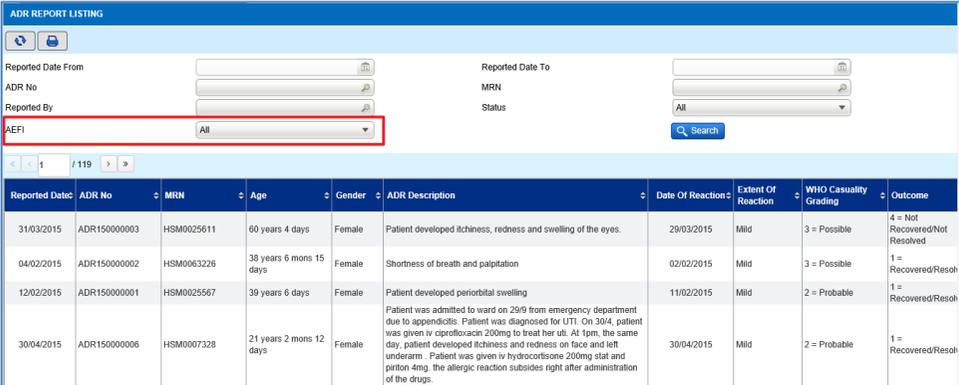
Summary of Ticket Number and Request for ADR/ AEFI Reporting Screen

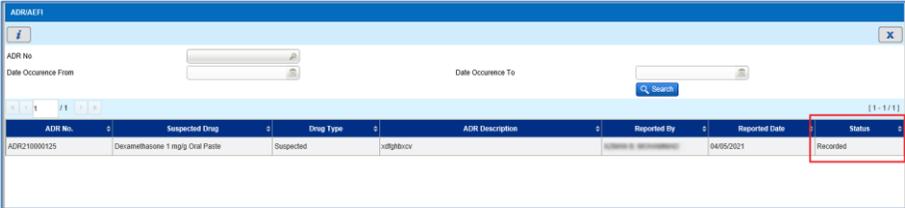
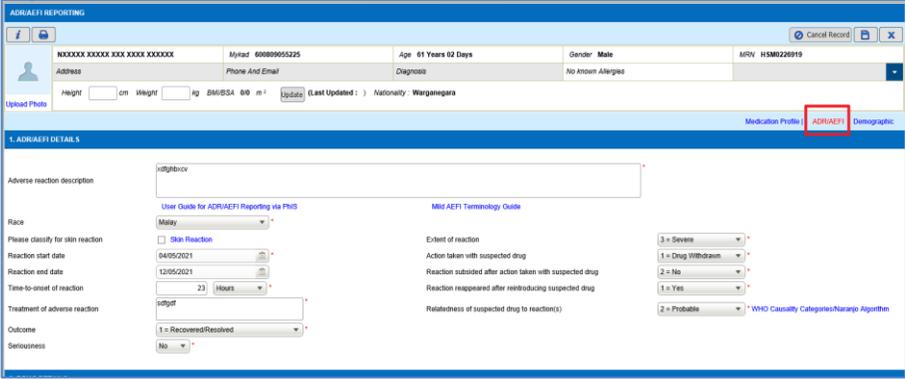
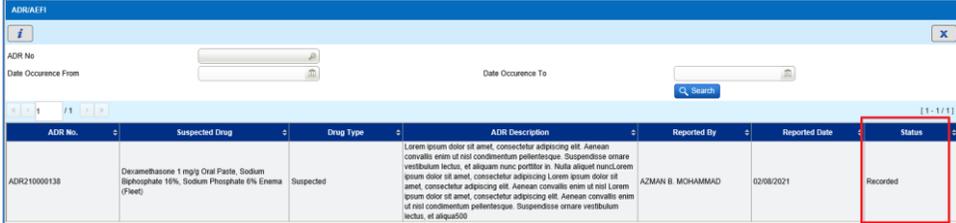
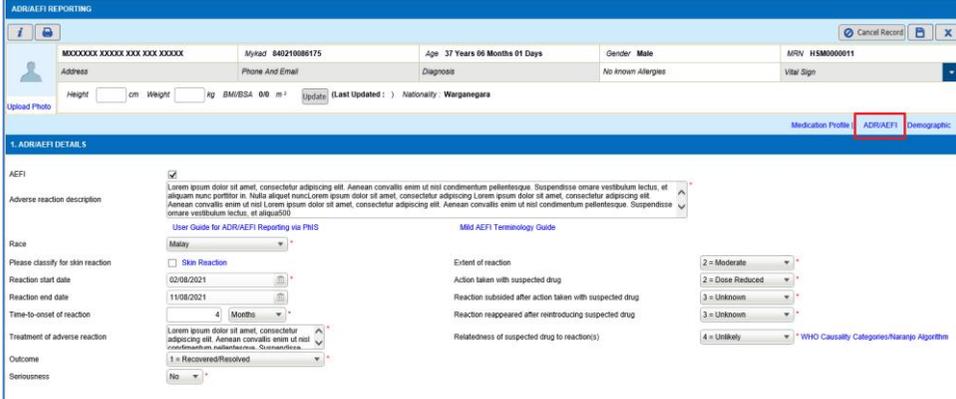
No.	Ticket #	Description	Requests	Solution
1.	200001140E/ 19655033C	User request to add 'click for AEFI' in the ADR module in PhIS so that she can capture the AEFI report from ADR. User are required to report for AEFI (adverse event following immunisation) to BPF but from PhIS, User informed she unable to trace whether the ADR for AEFI has been reported. It will be great if there is a column to summarise no of AEFI from ADR main page.	To add button (indicator) to differentiate between ADR or AEFI	<u>ADR/ AEFI Reporting Screen</u> 1) To add button (indicator) to differentiate between ADR or AEFI 2) To add search filter and column for AEFI at report 'ADR report listing
2.	200001357E/ 19820900C	I would like to request to add 'click for AEFI' in the ADR module in PhIS so that we can capture the AEFI report from ADR. We are required to report for AEFI (adverse event following immunisation) to BPF but from PhIS, we are unable to trace whether the ADR for AEFI has been reported. It will be great if there is a column to summarise no of AEFI from ADR main page.	To add button (indicator) to differentiate between ADR or AEFI	
3.	18564676C	User request ADR Button change into red colour once status change to verified in ADR Reporting screen. Currently, system shows red colour button for ADR in medication order once user create new ADR Reporting.	Request ADR hyperlink changed into red once status change to verified	<u>Patient Banner (ADR Hyperlink)</u> Request ADR hyperlink changed into red once status change to verified - affected to all ADR hyperlink at patient banner
4.	200001360E/ 19820909C	1. Reaction Subsided after stopping drug or Reducing Dose: Yes/No/Unknown in the current setting. However, the new form has N/A (drug continued). 2. Reaction reappeared after reintroducing drug: Yes/No/Unknown in the current setting. However, the new form has N/A (not reintroduced). 3. Seriousness: Yes (results in death is not present in the new ADR form) 4. Additional information: Yes, Others (does not allow for additional input). Please create a column to add remarks for others) 5. Extent of reaction: Mild/moderate/severe/unknown (the new form does not have unknown) 6. Action Taken With Suspected Drug: category 1-6 (drug	1. Reaction Subsided after stopping drug or Reducing Dose: Yes/No/Unknown in the current setting. However, the new form has N/A (drug continued). 2. Reaction reappeared after reintroducing drug: Yes/No/Unknown in the current setting. However, the new form has N/A (not reintroduced)."	<u>ADR Reporting Screen</u> 1. Reaction Subsided after stopping drug or Reducing Dose: Yes/No/Unknown in the current setting. However, the new form has N/A (drug continued) 2. Reaction reappeared after reintroducing drug: Yes/No/Unknown in the current setting. However, the new form has N/A (not reintroduced)." 3. To display both data (if selected) in ADR report printing

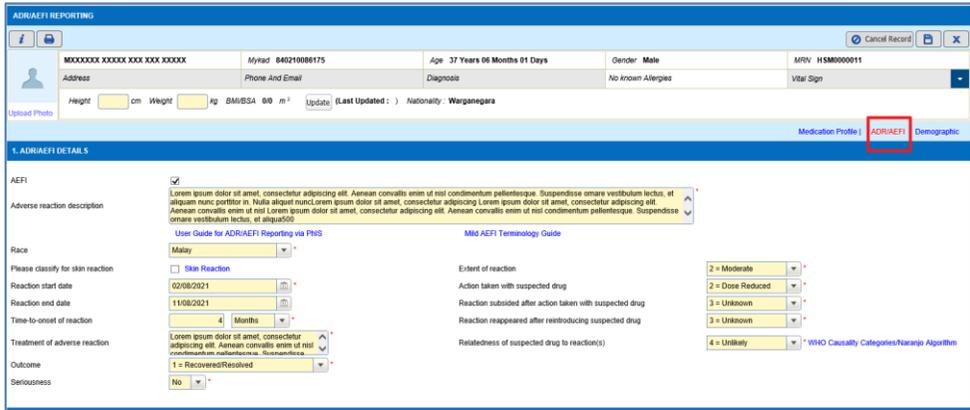
		<p>withdrawn, drug reduced etc). Request to add additional column to type.</p> <p>7. Outcome: 1= Recovered fully, 2 = Recovering 3 = Recovered/Resolved With Sequelae, 4 = Not Recovered/Not Resolved, 5 = Unknown (The new ADR form is Recovered fully, Recovering, Not Recovered, Unknown & Fatal with date and cause of death)</p>		<p><u>Cancel Remarks</u></p> <ol style="list-style-type: none"> 1. To add 'Remarks/Note' when cancelled ADR record 2. To display the 'Remarks'
5.	19694021C	No remark/note indicate 'cancelled record' or the report was cancelled in ADR Report screen		

Function Flow	ADR/ AEFI Reporting Screen in current Version 2.4	ADR/ AEFI Reporting Screen in new Version 2.5
<p>1. ADR/ AEFI Reporting Screen – AEFI Indicator</p>	<p>ADR/ AEFI → ADR/ AEFI Reporting</p> <p>1) Current version there is no indicator to differentiate between ADR and AEFI record</p>  <p>2) Current ADR/ AEFI listing screen, there is No filter option for AEFI</p>	<p>ADR/ AEFI → ADR/ AEFI Reporting</p> <p>1) Added check box for AEFI at ADR/ AEFI reporting screen</p>  <ul style="list-style-type: none"> - Tick on the 'AEFI' check box if user want to record for AEFI reporting - Fill in all the required information and click on SAVE button <p>2) Open ADR/ AEFI listing screen, there is new filter for AEFI and column to differentiate between ADR or AEFI record</p> <ul style="list-style-type: none"> - By default, field will be select 'All' means will display both ADR and AEFI record - User can select 'Yes' to filter record for AEFI - User can select 'No' to filter record for ADR

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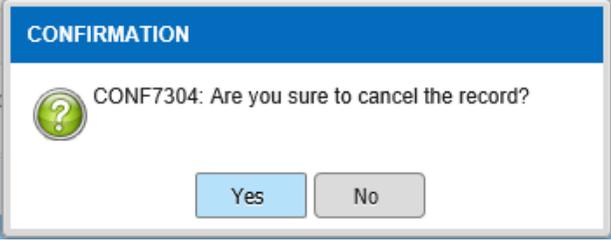
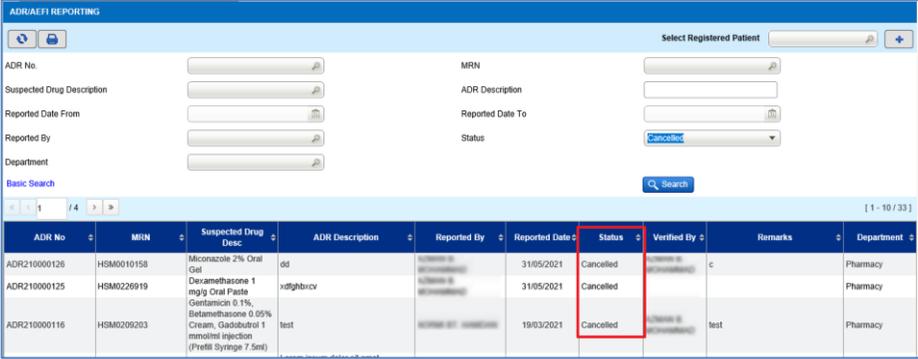
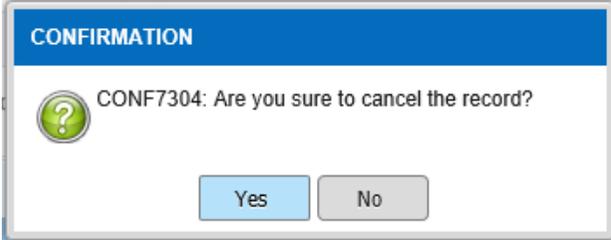
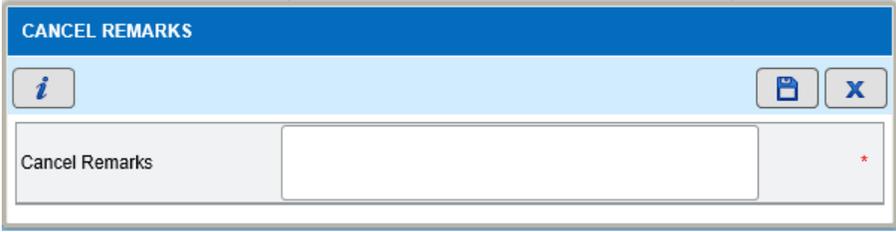
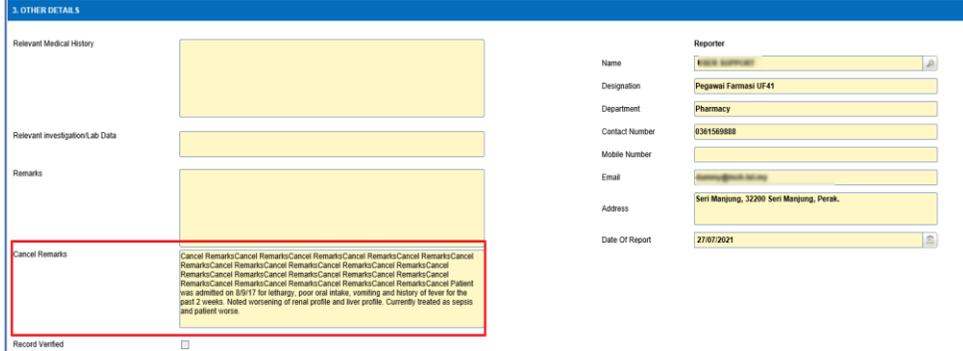
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04/02/2015	ADR150000002	HSM0063226	38 years 6 mons 15 days	Female	Shortness of breath and palpitation	02/02/2015	Mild	3 = Possible	1 = Recovered/Resol																																																																																			
12/02/2015	ADR150000001	HSM0025567	39 years 6 days	Female	Patient developed periorbital swelling	11/02/2015	Mild	2 = Probable	1 = Recovered/Resol																																																																																			
30/04/2015	ADR150000006	HSM0007328	21 years 2 mons 12 days	Female	Patient was admitted to ward on 29/9 from emergency department due to appendicitis. Patient was diagnosed for UTI. On 30/4, patient was given iv ciprofloxacin 200mg to treat her UTI. At 1pm, the same day, patient developed itchiness and redness on face and left underarm. Patient was given iv hydrocortisone 200mg stat and pirron 4mg. the allergic reaction subsides right after administration of the drugs.	30/04/2015	Mild	2 = Probable	1 = Recovered/Resol																																																																																			

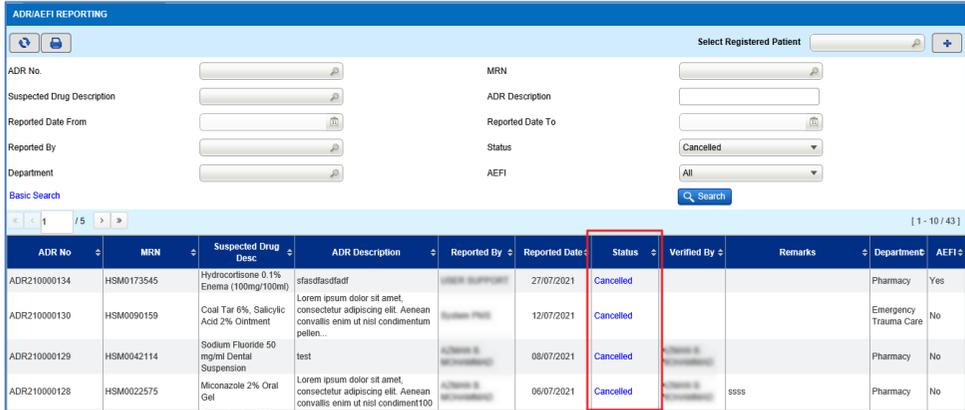
Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5
<p>3. ADR/AEFI Hyperlink</p>	<p>ADR/ AEFI → Patient Banner → ADR/ AEFI Hyperlink</p> <p>Current ADR/ AEFI Hyperlink will be in red color if patient ADR had been recorded with any status</p> <p>Example: ADR record with status = Recorded</p>  <p>ADR Hyperlink automatically in red color once ADR been recorded</p> 	<p>ADR/ AEFI → Patient Banner → ADR/ AEFI Hyperlink</p> <p>New version 2.5, ADR/AEFI hyperlink will only changed into red color once ADR status is “verified”</p> <p>Example: ADR record with status = Recorded</p>  <p>ADR Hyperlink still in blue color once ADR been recorded</p> 

Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5
		<p>Verified the ADR record</p>  <p>ADR Hyperlink automatically in red color once ADR record has been verified</p> 

Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5
<p>4. ADR/AEFI Reporting</p>	<p>ADR/ AEFI → Reporting</p> <p>Current ADR/ AEFI reporting No option to select 'N/A (drug continued)' and 'N/A (not reintroduced)' at:</p> <ol style="list-style-type: none"> Reaction Subsided after stopping drug or Reducing Dose <div data-bbox="264 472 1182 783" style="border: 1px solid blue; padding: 5px;"> <p>Extent of reaction 3 = Severe ▼ *</p> <p>Action taken with suspected drug 1 = Drug Withdrawn ▼ *</p> <p>Reaction subsided after action taken with suspected drug 2 = No ▼ *</p> <div style="border: 1px solid red; padding: 2px; margin-left: 20px;"> <p>1 = Yes</p> <p>2 = No</p> <p>3 = Unknown</p> </div> <p>Relatedness of suspected drug to reaction(s) 2 = Probable ▼ *</p> </div> <ol style="list-style-type: none"> Reaction reappeared after reintroducing drug <div data-bbox="264 858 1182 1161" style="border: 1px solid blue; padding: 5px;"> <p>Extent of reaction 3 = Severe ▼ *</p> <p>Action taken with suspected drug 1 = Drug Withdrawn ▼ *</p> <p>Reaction subsided after action taken with suspected drug 2 = No ▼ *</p> <p>Reaction reappeared after reintroducing suspected drug 1 = Yes ▼ *</p> <div style="border: 1px solid red; padding: 2px; margin-left: 20px;"> <p>1 = Yes</p> <p>2 = No</p> <p>3 = Unknown</p> </div> </div>	<p>ADR/ AEFI → Reporting</p> <p>New version 2.5, added option as below:</p> <ol style="list-style-type: none"> Added drop down value for 'N/A (drug continued)' at "Reaction Subsided after stopping drug or Reducing Dose" field <div data-bbox="1211 472 2175 762" style="border: 1px solid blue; padding: 5px;"> <p>Extent of reaction ▼ *</p> <p>Action taken with suspected drug ▼ *</p> <p>Reaction subsided after action taken with suspected drug ▼ *</p> <div style="border: 1px solid red; padding: 2px; margin-left: 20px;"> <p>1 = Yes</p> <p>2 = No</p> <p>3 = Unknown</p> <p>4 = N/A (drug continued)</p> </div> </div> <ol style="list-style-type: none"> Added drop down 'N/A (not reintroduced)' at "Reaction reappeared after reintroducing drug" <div data-bbox="1211 930 2175 1257" style="border: 1px solid blue; padding: 5px;"> <p>Extent of reaction ▼ *</p> <p>Action taken with suspected drug ▼ *</p> <p>Reaction subsided after action taken with suspected drug ▼ *</p> <p>Reaction reappeared after reintroducing suspected drug ▼ *</p> <div style="border: 1px solid red; padding: 2px; margin-left: 20px;"> <p>1 = Yes</p> <p>2 = No</p> <p>3 = Unknown</p> <p>4 = N/A (not reintroduced)</p> </div> </div>

Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5												
		<p>3. To display both data (if selected) in ADR report printing</p> <div data-bbox="1205 316 2168 1098" style="border: 1px solid black; padding: 5px;"> <p style="text-align: center;">REPORT ON SUSPECTED ADVERSE DRUG REACTION/AEFI NATIONAL CENTRE FOR ADVERSE DRUG REACTION MONITORING</p> <p>ADR NO : ADR210000134 www.npra.gov.my</p> <p><small>(Please report all suspected drug reaction including those for vaccines and traditional medicines. Do not hesitate to report if some details are not known. Identities of Reporter, Patient and Institution will remain Confidential.)</small> <small>REPORT No (for office use only)</small></p> <hr/> <p>PATIENT INFORMATION</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th>Patient NRIC</th> <th>Age</th> <th>Sex</th> <th>Wt (kg)</th> <th>Ethnic Group</th> <th>Institution</th> </tr> </thead> <tbody> <tr> <td>930805075081</td> <td>28 Years 06 Days</td> <td>Male</td> <td>65</td> <td>Berawan</td> <td>Hospital Seri Manjung</td> </tr> </tbody> </table> <hr/> <p>ADVERSE REACTION DESCRIPTION</p> <p>sfasdfsdf</p> <p>Skin Reaction :</p> <p>Please specify Part of Body Affected :</p> <p>Time-to-onset of reaction : <input type="text" value="3 Months"/> Reaction start date : <input type="text" value="07/07/2021"/> Reaction end date : <input type="text" value="22/07/2021"/></p> <p>Action taken with suspected drug : <input type="text" value="6 = Not Applicable"/></p> <p>Reaction subsided after action taken with suspected drug : Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A (drug continued) <input checked="" type="checkbox"/></p> <p>Reaction reappeared after reintroducing suspected drug : Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A (not reintroduced) <input checked="" type="checkbox"/></p> <p>Extent of reaction : Mild <input type="checkbox"/> Moderate <input checked="" type="checkbox"/> Severe <input type="checkbox"/> Unknown <input type="checkbox"/></p> <p>Treatment of adverse reaction & action taken : <input type="text" value="sdfasdf"/></p> <p>Outcome : Recovered <input checked="" type="checkbox"/> Recovering <input type="checkbox"/> Recovered With Sequelae <input type="checkbox"/> Not Recovered <input type="checkbox"/> Unknown <input type="checkbox"/></p> <p>Seriousness: Life Threatening <input type="checkbox"/> Hospitalization/Prolonged Hospitalization <input type="checkbox"/> Disability / Incapacity <input type="checkbox"/> Birth Defect <input type="checkbox"/></p> <p style="text-align: right;">Results In Death <input type="checkbox"/> Date of death <input type="text"/></p> <p>Relatedness of suspected drug to reaction(s) : Certain <input type="checkbox"/> Probable <input type="checkbox"/> Possible <input type="checkbox"/> Unlikely <input type="checkbox"/> Unclassifiable <input checked="" type="checkbox"/></p> </div>	Patient NRIC	Age	Sex	Wt (kg)	Ethnic Group	Institution	930805075081	28 Years 06 Days	Male	65	Berawan	Hospital Seri Manjung
Patient NRIC	Age	Sex	Wt (kg)	Ethnic Group	Institution									
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Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5																																								
<p>5. Cancel Remarks</p>	<p>ADR/ AEFI → Reporting</p> <p>Current version there is No remarks field available when cancelled ADR records</p> <ol style="list-style-type: none"> Click on cancel button, will appear confirmation message as below:  <ol style="list-style-type: none"> Click on 'Yes' to confirm cancel the record, record will be cancelled.  <table border="1" data-bbox="264 981 1182 1117"> <thead> <tr> <th>ADR No</th> <th>MRN</th> <th>Suspected Drug Desc</th> <th>ADR Description</th> <th>Reported By</th> <th>Reported Date</th> <th>Status</th> <th>Verified By</th> <th>Remarks</th> <th>Department</th> </tr> </thead> <tbody> <tr> <td>ADR210000126</td> <td>HSM0010158</td> <td>Miconazole 2% Oral Gel</td> <td>dd</td> <td></td> <td>31/05/2021</td> <td>Cancelled</td> <td></td> <td></td> <td>Pharmacy</td> </tr> <tr> <td>ADR210000125</td> <td>HSM0226919</td> <td>Dexamethasone 1 mg/ly Oral Paste Gentamicin 0.1%, Betamethasone 0.05% Cream, Clostrid 1 mmol/ml injection (Prefill Syringe 7.5ml)</td> <td>vdghbvcv</td> <td></td> <td>31/05/2021</td> <td>Cancelled</td> <td></td> <td></td> <td>Pharmacy</td> </tr> <tr> <td>ADR210000116</td> <td>HSM0209203</td> <td></td> <td>test</td> <td></td> <td>19/03/2021</td> <td>Cancelled</td> <td></td> <td>test</td> <td>Pharmacy</td> </tr> </tbody> </table>	ADR No	MRN	Suspected Drug Desc	ADR Description	Reported By	Reported Date	Status	Verified By	Remarks	Department	ADR210000126	HSM0010158	Miconazole 2% Oral Gel	dd		31/05/2021	Cancelled			Pharmacy	ADR210000125	HSM0226919	Dexamethasone 1 mg/ly Oral Paste Gentamicin 0.1%, Betamethasone 0.05% Cream, Clostrid 1 mmol/ml injection (Prefill Syringe 7.5ml)	vdghbvcv		31/05/2021	Cancelled			Pharmacy	ADR210000116	HSM0209203		test		19/03/2021	Cancelled		test	Pharmacy	<p>ADR/ AEFI → Reporting</p> <p>Added remarks/ note field when cancelled ADR record</p> <ol style="list-style-type: none"> Click on cancel button, will appear confirmation message as below:  <ol style="list-style-type: none"> Click on 'Yes' to confirm cancel the record, another popup message will appear to fill in cancel remarks with maximum 500 characters  <ol style="list-style-type: none"> Click on save button and remarks cancel will be display at 'Other Details' section 
ADR No	MRN	Suspected Drug Desc	ADR Description	Reported By	Reported Date	Status	Verified By	Remarks	Department																																	
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Function Flow	ADR/ AEFI Report Printing in current Version 2.4	ADR/ AEFI Report Printing in new Version 2.5
		<p>4. 'Cancel Remarks' also can be view from ADR/ AEFI report listing screen</p>  <p>- Click on the Cancelled hyperlink will display popup screen as below:</p> 