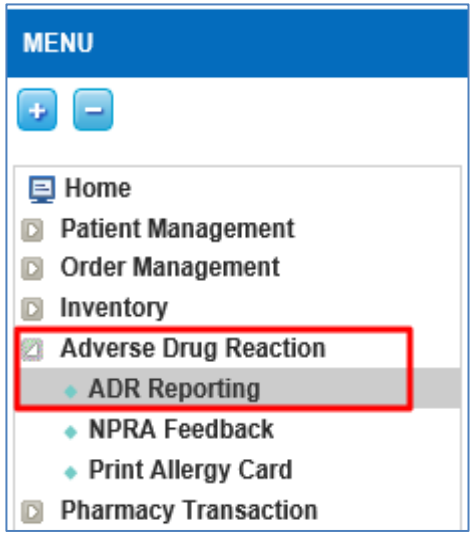
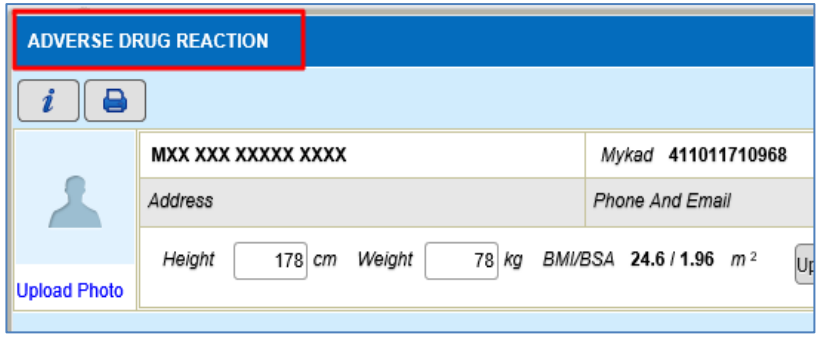
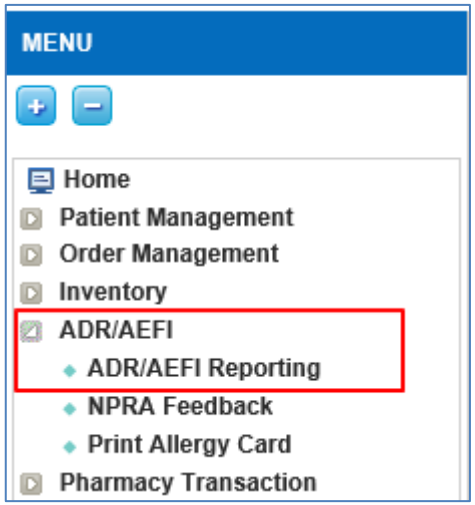
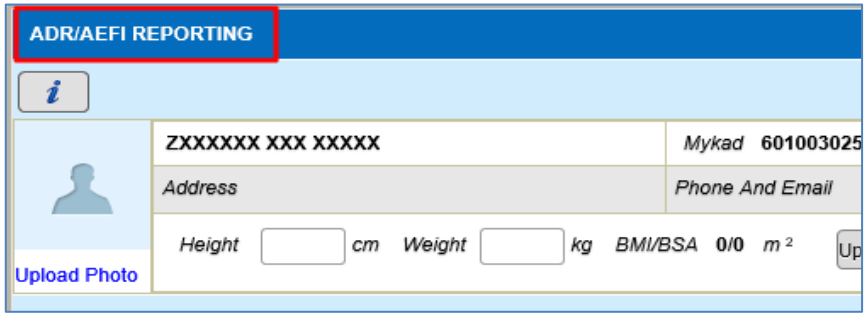
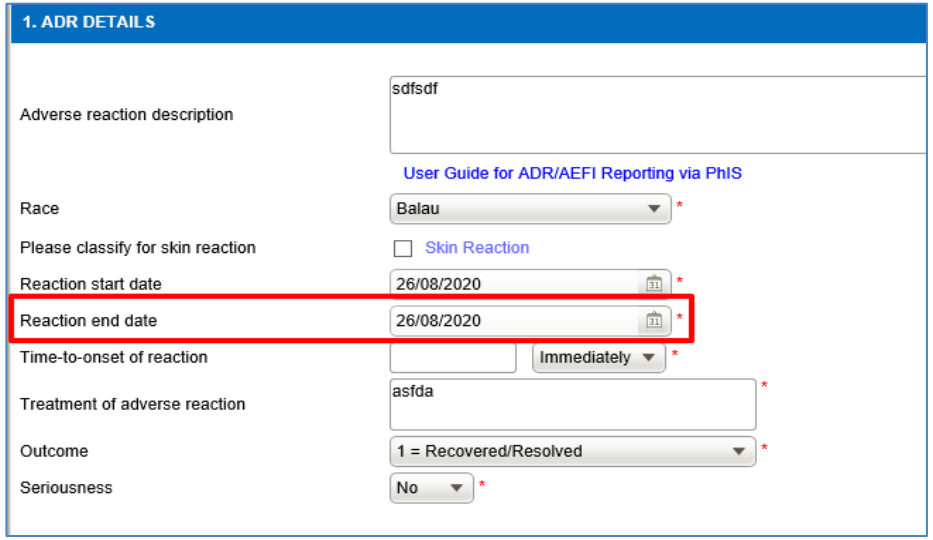
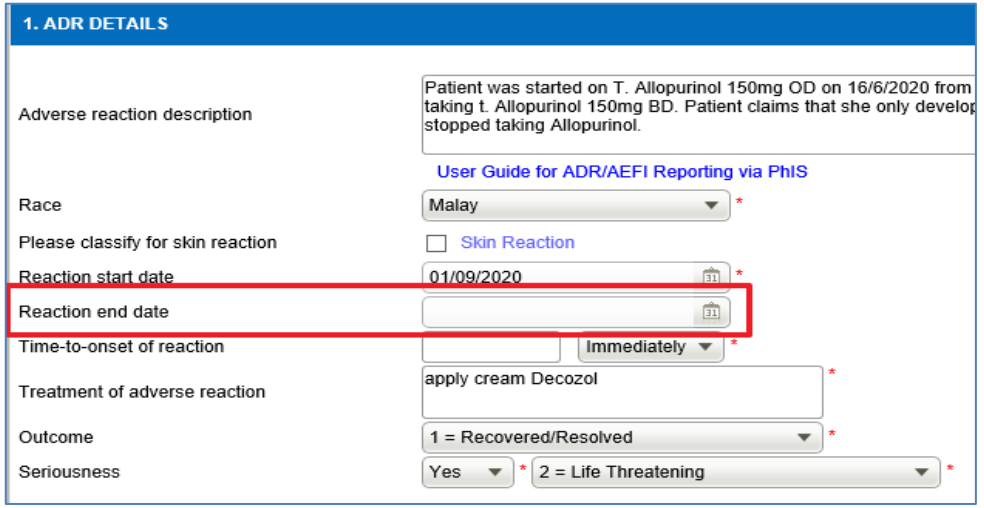
Summary of Ticket Number and Request for ADR Reporting Screen

No.	Ticket #	Description	Requests	Solution
1.	I-PhIS035 781118S	User request date end of reaction not mandatory because sometime patient still got reaction during ADR reporting. User also request got remark column for patient still ongoing the reaction.	Request date end of reaction not mandatory	ADR Reporting Screen 1) To remove mandatory for Date End of Reaction 2) To rename ' Hospitalization/Prolong Hospitalization' to 'Hospitalization/Prolonged Hospitalization' 3) To rename 'ADR Reporting' to 'ADR/AEFI Reporting' 4) To remove this confirmation alert (CONF0003: Please use WHO Classification for ADR description. Proceed to Verify?) 5) To have link to medication profile in ADR reporting section ADR Report Printing 1. To display 'Action taken with suspected drug' in ADR Report printing 2. To rename 'www.bpfk.gov.my' to 'www.npra.gov.my' 3. to display full name 'Hospitalization/Prolonged Hospitalization'"
2.	I-PhIS045 488918S	User request to remove mandatory for Date End of Reaction and add field Date End of Reaction REMARK . As per drug details for Therapy End Date Remark	Request date end of reaction not mandatory	
3.	I-PhIS047 907718S	User request Date End Of Reaction field not compulsory to key in.	Request date end of reaction not mandatory	
4.	I-PhIS049 313619S	To rename ' Hospitalization/Prolong Hospitalization' to 'Hospitalization/Prolonged Hospitalization'	To rename ' Hospitalization/Prolong Hospitalization' to 'Hospitalization/Prolonged Hospitalization'	
5.	I-PhIS049 314419S	To rename 'ADR Reporting' to 'ADR/AEFI Reporting'	To rename 'ADR Reporting' to 'ADR/AEFI Reporting'	
6.	I-PhIS049 355219S	Remove mandatory for 'Date end of Reaction' field	Request date end of reaction not mandatory	
7.	I-PhIS049 355319S	To remove this confirmation alert (CONF0003: Please use WHO Classification for ADR description. Proceed to Verify?)	To remove this confirmation alert (CONF0003: Please use WHO Classification for ADR description. Proceed to Verify?)	
8.	I-PhIS049 355419S	1. To display 'Action taken with suspected drug' in ADR Report printing 2. To rename 'www.bpfk.gov.my' to 'www.npra.gov.my' 3. to display full name 'Hospitalization/Prolonged Hospitalization'	1. To display 'Action taken with suspected drug' in ADR Report printing 2. To rename 'www.bpfk.gov.my' to 'www.npra.gov.my' 3. to display full name 'Hospitalization/Prolonged Hospitalization'	
9.	I-PhIS021 674717S	User request to view all medication in drug details column. Currently at drug details column only appear active medication. User request to view all medication include previous medication.	ADR Reporting Screen To have link to medication profile in ADR report section	

Function Flow	ADR Reporting Screen in current Version 2.2	ADR Reporting Screen in new Version 2.3
1. ADR/ AEFI Reporting Screen	**Adverse Drug Reaction → ADR Reporting** 1) Menu and header name will be displayed as "ADR Reporting" • At the main Menu:  • At the screen header: 	**ADR/ AEFI → ADR/ AEFI Reporting** 1) Menu and header has been renamed form 'ADR Reporting' to 'ADR/AEFI Reporting' • At the main menu:  • At the screen header: 

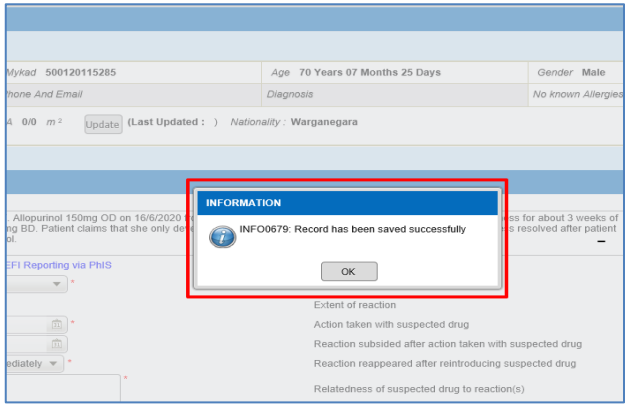
Function Flow	ADR Reporting Screen in current Version 2.2	ADR Reporting Screen in new Version 2.3
	2) 'Reaction end date' field is mandatory to be selected	2) Mandatory rules for 'Reaction End Date' field has been removed

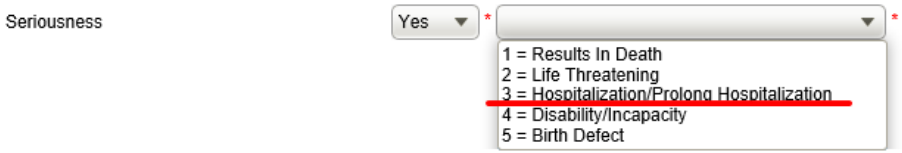
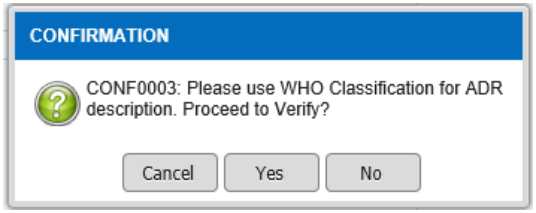
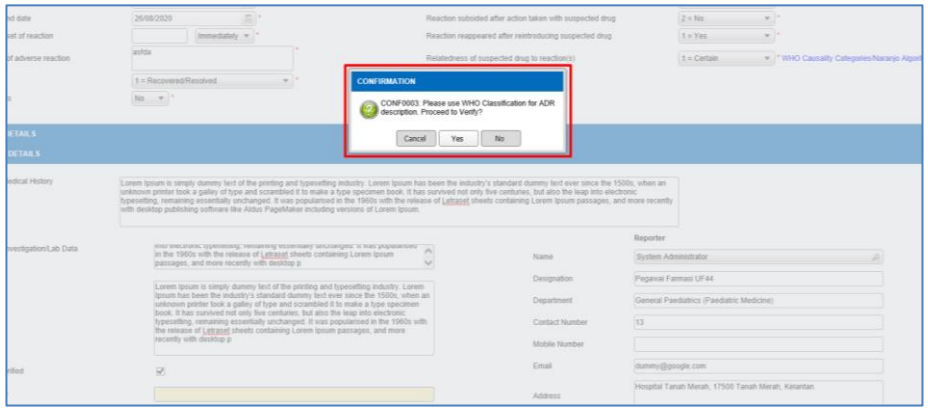
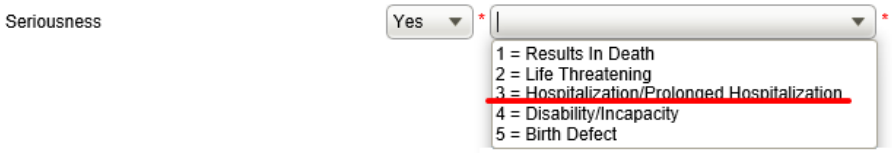
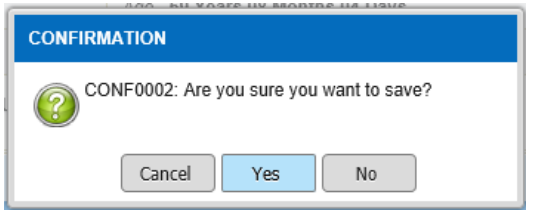
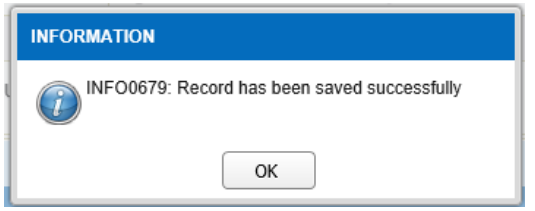




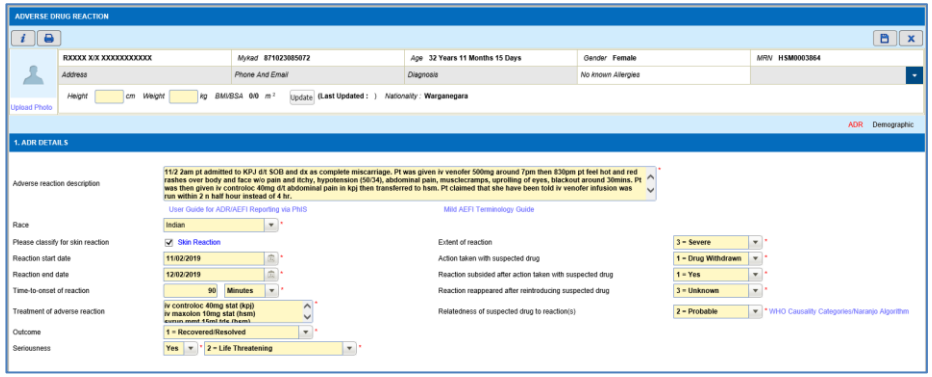
Steps:

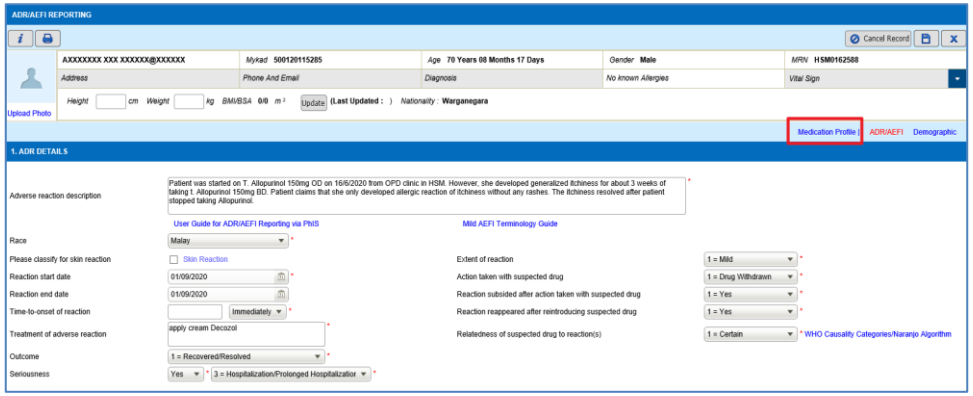
- Create ADR record with empty 'Reaction end date' field
- Click on Save button
- Record saved successfully



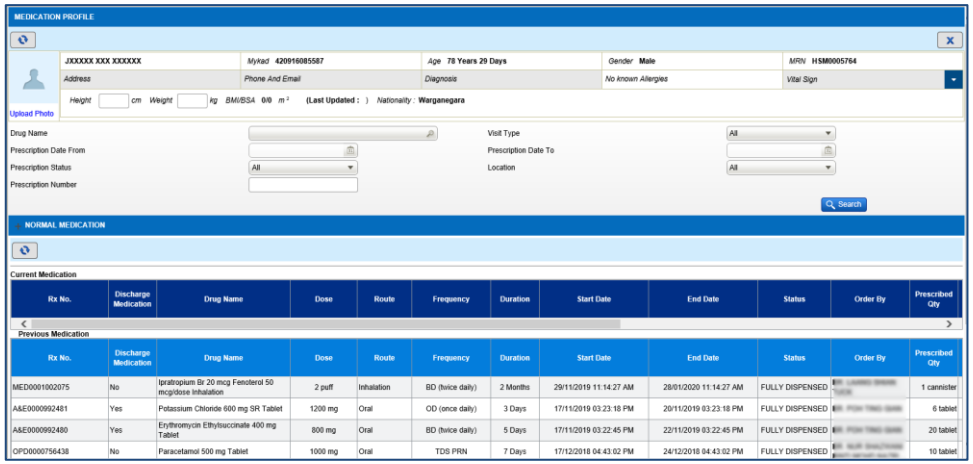
Function Flow	ADR Reporting Screen in current Version 2.2	ADR Reporting Screen in new Version 2.3
	3) In version 2.2, display as 'Hospitalization/Prolong Hospitalization'  4) There is alert message on using WHO classification when click on verify check box. The alert message as below:  Step: - Tick on 'Record Verified' check box, message alert prompted as below: 	3) In new version has been renamed from 'Hospitalization/Prolong Hospitalization' to 'Hospitalization/Prolonged Hospitalization'  4) Alert message on using WHO classification has been removed. Steps: - Tick on 'Record Verified' check box, no more message alert prompted - User may proceed to click on Save button to save the records, message alert shall prompt "Are you sure you want to save?"  - Click 'yes', another message alert prompted to informed record has been saved successfully 

Function Flow	ADR Reporting Screen in current Version 2.2	ADR Reporting Screen in new Version 2.3
	5) Currently there is no place to view patient's medication profile in ADR reporting section	5) Link to view patient's medication profile has been added in ADR reporting screen





- Click on the hyperlink will display patients, medication profile screen



Function Flow	ADR Report Printing in current Version 2.2	ADR/ AEFI Report Printing in new Version 2.3
2. ADR/AEFI Report Printing	**Adverse Drug Reaction → ADR Reporting → Report Printing** Current ADR Report Printing	**ADR/ AEFI → ADR/ AEFI Reporting → Report Printing** New ADR/ AEFI Report Printing 1. Displayed 'Action taken with suspected drug' in the ADR/ AEFI Report printing 2. Renamed 'www.bpfk.gov.my' to 'www.npra.gov.my' 3. Displayed full name of 'Hospitalization/Prolonged Hospitalization'

REPORT ON SUSPECTED ADVERSE DRUG REACTION
NATIONAL CENTRE FOR ADVERSE DRUG REACTION MONITORING

ADR NO : ADR190000035 www.bpfk.gov.my

(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**.)
REPORT No ..... (for office use only)

PATIENT INFORMATION

Patient NRIC	Age	Sex	Wt (kg)	Ethnic Group	Institution
871023085072	32 Years 11 Months 15 Days	Female		Indian	Hospital Seri Manjung

ADVERSE REACTION DESCRIPTION

11/2 2am pt admitted to KPJ d/t SOB and dx as complete miscarriage. Pt was given iv venofer 500mg around 7pm then 830pm pt feel hot and red rashes over body and face w/o pain and itchy, hypotension (50/34), abdominal pain, musclecramps, uprolling of eyes, blackout around 30mins. Pt was then given iv controloc 40mg d/t abdominal pain in kpj then transferred to hsm. Pt claimed that she have been told iv venofer infusion was run within 2 n half hour instead of 4 hr.

Skin Reaction : Urticaria ,

Please specify Part of Body Affected : include wholebody and face

Time-to-onset of reaction : 90 Minutes | Reaction start date : 11/02/2019 | Reaction end date : 12/02/2019

Reaction subsided after action taken with suspected drug : Yes ☑ No ☐ Unknown ☐

Reaction reappeared after reintroducing suspected drug : Yes ☐ No ☐ Not applicable ☑

Extent of reaction : Mild ☐ Moderate ☐ Severe ☑

Treatment of adverse reaction & action taken: iv controloc 40mg stat (kpj); iv maxolon 10mg stat (hsm); syrup mmt 15ml tds (hsm); t.pantoprazole 40mg od (hsm); c.tramal 50mg stat & tds

Outcome : Recovered ☑ Recovering ☐ Recovered With Sequelae ☐ Not Recovered ☐ Unknown ☐

Seriousness: Life Threatening ☑ Hospitalization / Prolong Hospitalization ☐ Disability / Incapacity ☐ Birth Defect ☐ Results In Death ☐ Date of death:

Relatedness of suspected drug to reaction(s) : Certain ☐ Probable ☑ Possible ☐ Unlikely ☐ Unclassifiable ☐

REPORT ON SUSPECTED ADVERSE DRUG REACTION/AEFI
NATIONAL CENTRE FOR ADVERSE DRUG REACTION MONITORING

ADR NO : ADR200000097 www.npra.gov.my (2)

(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**.)
REPORT No ..... (for office use only)

PATIENT INFORMATION

Patient NRIC	Age	Sex	Wt (kg)	Ethnic Group	Institution
500120115285	70 Years 08 Months 18 Days	Male		Malay	Hospital Seri Manjung

ADVERSE REACTION DESCRIPTION

Patient was started on T. Allopurinol 150mg OD on 16/6/2020 from OPD clinic in HSM. However, she developed generalized itchiness for about 3 weeks of taking t. Allopurinol 150mg BD. Patient claims that she only developed allergic reaction of itchiness without any rashes. The itchiness resolved after patient stopped taking Allopurinol.

Skin Reaction :

Please specify Part of Body Affected :

Time-to-onset of reaction : Immediately | Reaction start date : 01/09/2020 | Reaction end date : 01/09/2020

Action taken with suspected drug : 1 = Drug Withdrawn (1)

Reaction subsided after action taken with suspected drug : Yes ☑ No ☐ Unknown ☐

Reaction reappeared after reintroducing suspected drug : Yes ☑ No ☐ Not applicable ☐

Extent of reaction : Mild ☑ Moderate ☐ Severe ☐

Treatment of adverse reaction & action taken: apply cream Decozol

Outcome : Recovered ☑ Recovering ☐ Recovered With Sequelae ☐ Not Recovered ☐ Unknown ☐

Seriousness: Life ☐ (3) Hospitalization/Prolonged Hospitalization ☑ Disability / Incapacity ☐ Birth Defect ☐ Results In Death ☐ Date of death

Relatedness of suspected drug to reaction(s) : Certain ☑ Probable ☐ Possible ☐ Unlikely ☐ Unclassifiable ☐