

2018 Upgrade

Manage Oncology Research Adverse Events – Oncology Providers

When research coordinators finish documenting a patient's adverse events for a research study, they send them to you as the principal investigator or treating physician for review. From In Basket, you can review the documentation and mark it as reviewed.

1. From In Basket, select the **Adverse Event Review** folder.
2. Select a message.
3. Use the report to view the patient's new, changed, and unchanged adverse events since the last time you reviewed them.
4. Click Adverse Events to open the Adverse Events activity.
5. Document the Relation to Study and click Mark as Reviewed.
6. Back in your In Basket when you're done reviewing, click Mark As Reviewed.

Salazar, Grace
 Female, 53 yo, 11/7/1964
 Weight: 148 lb (67.1 kg)
 PCP: Seeger, Marty
 MRN: 127539
 MyChart: Inactive
 Next Appt: 11/01/2017

Adverse Event Review Received: Today
 Jensen, Nora → You

New Adverse Events
 These adverse events have been created since the last review.

Constipation	Expected	Current Grade	Relation to Study	Start	Stop
Serious No	No	1	Possible	10/10/2017	--
Grade History ⤴					
Grade			Start		
1			10/10/2017		

Changed Adverse Events
 These adverse events have been changed since the last time they were reviewed.

Fatigue	Expected	Current Grade	Relation to Study	Start	Stop
Serious No	No	3	Possible	10/4/2017	10/5/2017
Grade History ⤴					
Grade			Start		
3			10/4/2017		

Unchanged Adverse Events ⤴
 These adverse events have not been changed since the last time they were reviewed.