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.