Nordion Inc. Event Reporting

For the Quarter (October – December 2015)

Event Number	Event	Status/Outcome
15-24*	Vials incorrectly dispensed.	Two product vials were incorrectly dispensed resulting in incorrect amount of activity in each vial. This further resulted in incorrect package labeling and shipping documentation. Currently under investigation
15-26*	Packages returned from a third party site contained a sealed source even though the package was labeled as empty.	Third party has implemented a visual check prior to return shipment. Nordion will be provided with a photograph of the empty package prior to shipment. No action required by Nordion.
15-27*	Shipment received from supplier included an unsealed capsule.	Enhanced manufacturing controls have been put in place at the supplier. No action required by Nordion.
15-28*	Shipment received with contents that exceed allowable activity limit of Type B certification.	Supplier to assess measurement of source process. No action required by Nordion.
15-30*	Damaged Type A package.	Carrier repackaged into another box. No action required by Nordion

**** There was no impact to health, safety, environment or security as a result of these events***