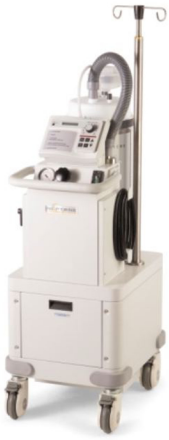
Neptune® Gold & Neptune Bronze



0700-001-000 Neptune Gold

PLEASE REFER TO THE RECALL NOTIFICATION LETTER FOR COMPLETE DETAILS

Required Actions

0700-007-000 Neptune Bronze

1	Ensure all users of the Neptune device, including surgeons, surgical residents and health profession students or OR assignments, are adequately trained, and are aware of the risks associated with the device as detailed in the device recall notification.	☐
2	It is recommended that all facilities keep a master list of all personnel that have been trained on the use of the Neptune. This list should include all users, such as surgeons, anesthesiologists, residents, nurses and technicians.	☐
3	Inform all users of the Neptune device that additional adverse events have been reported.	☐
4	Ensure warning labels, previously supplied by Stryker, are present on all Neptune device(s). If you require additional labels, please contact Stryker at 855-458-7441 or 269-389-2316.	☐
5	Complete the business reply form, acknowledging you completed the above actions.	☐