# Class 2 Device Recall Stryker FlowPort II Adapter

**Date Posted**: June 03, 2016

**Recall Status**: Terminated on June 03, 2016

**Recall Number**: Z-1910-2016

**Recall Event ID**: 7407023

**Product Classification**: OR - Orthopedic - Product Code NBH

**Product**: Stryker FlowPort II Adapter: Model number: 00CAT00778

The FlowPort II Adapter is intended to connect the FlowPort II Cannulas to commercially available, 4mm arthroscopes in surgical procedures.

**Code Information**: Lot Numbers Affected: 13604

**Recalling Firm/Manufacturer**: Stryker Corporation

5900 Optical Ct
San Jose CA 95138-1400

**For Additional Information Contact**: Michael Hilldoerfer
408-754-2664

**Manufacturer Reason for Recall**: Complaints were received for the Stryker FlowPort II Adapter, and investigation found that the scope sat in the adapter 180 degrees in the wrong direction

**FDA Determined Cause**: Device Design

**Action**: A recall letter was not sent as all affected devices have been returned. A ship hold was placed on the device on October 16, 2015.

**Quantity in Commerce**: 21 units

**Distribution**: GA, UT, MD, NJ, MT

**Total Product Life Cycle**: TPLC Device Report

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1. For details about termination of a recall see [Code of Federal Regulations (CFR) Title 21 §7.55](http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfres/res.cfm?id=146181)
2. Per FDA policy, recall cause determinations are subject to modification up to the point of termination of the recall.

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Links on this page:

4. http://www.fda.gov/MedicalDevices/default.htm
6. /scripts/cdrh/devicesatfda/index.cfm
7. /scripts/cdrh/cfdocs/cfPMN/pmn.cfm

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http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfres/res.cfm?id=146181

6/14/2016