

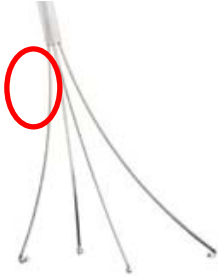
URGENT: TALON[®] GRASPING DEVICE RECALL

Customer Name: [REDACTED]
Regarding: US Endoscopy Talon[®] grasping device
Model No: 00711175

Dear Distributor Partner:

US Endoscopy is conducting a voluntary product recall for certain lots of the Talon[®] grasping device. We received a report of a grasping wire detachment during use (please see picture for reference below). We have determined the product lots listed below may be affected by a condition which could result in a wire detachment. Our records indicate you have purchased devices from the affected lots.

No other Talon[®] device, manufactured before or after those listed below, are affected by this action.

Affected Lot Numbers (manufactured April – June 2014)	Talon [®] grasping device (00711175)																
<table border="1"> <tr><td>1407418</td><td>1410178</td></tr> <tr><td>1407419</td><td>1410179</td></tr> <tr><td>1408069</td><td>1410478</td></tr> <tr><td>1408070</td><td>1410480</td></tr> <tr><td>1408739</td><td>1410990</td></tr> <tr><td>1408740</td><td>1411518</td></tr> <tr><td>1408741</td><td>1423937</td></tr> <tr><td>1408742</td><td>1423938</td></tr> </table>	1407418	1410178	1407419	1410179	1408069	1410478	1408070	1410480	1408739	1410990	1408740	1411518	1408741	1423937	1408742	1423938	
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1408740	1411518																
1408741	1423937																
1408742	1423938																

US Endoscopy will provide a free replacement for each unused Talon[®] grasping device that is returned. US Endoscopy requests that you:

- 1. Quarantine the affected Talon[®] grasping devices in your stock.**
- 2. Contact your customers to arrange for return of the affected devices to you.**
- 3. Complete the attached Customer Response Card and return to the attention of Cindy Perpar, via fax 1-440-639-6256 or email recall@usendoscopy.com.**
- 4. Mark the shipping container with code: RMA 7859833 QIF**
- 5. Return affected stock (yours and customer) via Fed Ex Account #293799547, to:**

**US Endoscopy
6091 Heisley Road
Mentor, Ohio 44060 USA**

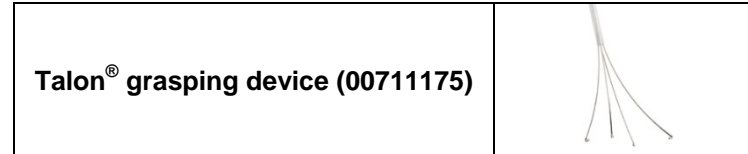
Once the product(s) has been received by US Endoscopy, replacement products will be sent to you.

US Endoscopy is conducting this recall in the continuing interest of ensuring the delivery of quality devices and customer safety.

If you have any questions, please contact Cindy Perpar at 1-440-358-6051.

PLEASE RETAIN THIS LETTER FOR YOUR RECORDS

**CUSTOMER RESPONSE CARD FOR:
DEVICE TECHNOLOGIES**



Please check the appropriate boxes below. If you have no stock of the listed product lots, indicate this below and return the completed card. If you currently have stock, indicate the quantity in units next to the lot number and return the completed card.

- We acknowledge the receipt of this recall notification and understand its content.
- We have no stock of the listed lot numbers.
- We do have stock of the listed lot numbers and have quarantined the units for return to US Endoscopy. Please list below:

Product	Lot Number Received	Quantity Received
00711175	1410480	15 units
	1410478	15 units
	1408740	15 Units
	1408741	10 Units
	1408739	15 Units
	1407419	10 Units

Lot Number Returned	Quantity Returned

- We do have stock of the listed lot numbers and have elected to retain the product for the following reason(s):

Please fax the completed signed form to the attention of Cindy Perpar at 1-440-639-6256, or email to recall@usendoscopy.com.

Return the affected product to US Endoscopy, if applicable.

If you have any questions concerning this recall notification, please contact Cindy Perpar at 1-440-358-6051.

Contact Name (please print)

Title

Signature and date

Facility Name

Address

Phone Number