

URGENT: TALON® GRASPING DEVICE RECALL

Customer Name: GULF MEDICAL CO LTD

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.

	ot Numbers pril – June 2014)	Talon [®] grasping device (00711175)	
1407418 1407419 1408069 1408070 1408739 1408740 1408741 1408742	1410178 1410179 1410478 1410480 1410990 1411518 1423937 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 7859827 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.



CUSTOMER RESPONSE CARD FOR:

GULF MEDICAL CO LTD

Talon [®] grasping device (00711175)	
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		riate boxes below. If you have no		ts, indicate this below and return completed card.	the completed card. If you			
	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							
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 Numbe				