New Product Committee Guide

General Information

**Name of Product:** AccuTrac® Single-Use 200 micron Holmium Laser Fiber and Flexiva® TracTip High Power Single-Use 200 micron Holmium Laser Fiber

**Product Codes:**

<table>
<thead>
<tr>
<th>Product Code</th>
<th>Description</th>
<th>Unit</th>
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<tbody>
<tr>
<td>M0068404110</td>
<td>AccuTrac 200 micron Holmium Laser Fiber</td>
<td>Single</td>
</tr>
<tr>
<td>M0068404112</td>
<td>AccuTrac 200 micron Holmium Laser Fiber</td>
<td>Box 5</td>
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<tr>
<td>M0068403960</td>
<td>Flexiva TracTip 200 micron Holmium Laser Fiber</td>
<td>Single</td>
</tr>
<tr>
<td>M0068403961</td>
<td>Flexiva TracTip 200 micron Holmium Laser Fiber</td>
<td>Box 5</td>
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**Product Description:**
The AccuTrac Laser Fibers are fiber optic laser energy delivery devices consisting of a SMA-905 connector, strain relief, and a silica core fiber jacketed with ethylene tetrafluoroethylene (ETFE). The Flexiva TracTip fibers are equipped with a polished and reinforced ball-shaped TracTip. These fibers may be used in a variety of laser based surgical cases as an integral part of laser systems.

The Flexiva TracTip Laser Fibers are fiber optic laser energy delivery devices consisting of a SMA-905 connector, strain relief, and a silica core fiber jacketed with ethylene tetrafluoroethylene (ETFE). The AccuTrac fibers are equipped with a polished and reinforced ball-shaped TracTip. These fibers may be used in a variety of laser based surgical cases as an integral part of laser systems.

**Manufacturer:** Boston Scientific Corporation

**Distributor Federal Tax ID:** 04 269 5240

**Will this product(s) replace or supplement a current in-house product(s) now performing the same function?**
The AccuTrac and Flexiva TracTip fibers will supplement the product line of AccuMax® and Flexiva® (End-Firing) single use Holmium Laser Fibers currently on the market.

**Laser Compatibility:**
AccuTrac Fibers are recommended for use with Ho:YAG laser systems with a standard SMA-905 connector that have been cleared for surgical use. Recommended Ho:YAG lasers are Dornier® and New Star. Please refer to the laser system User Manual for complete information regarding applications, contraindications, precautions and warnings.

Flexiva TracTip Fibers are recommended for use with Ho:YAG laser systems with a standard SMA-905 connector that have been cleared for surgical use. Recommended for use with Lumenis manufactured Ho:YAG and Nd:YAG lasers. Please refer to the laser system User Manual for complete information regarding applications, contraindications, precautions and warnings.

**Product Usage**

Briefly describe this product and other required components/accessories, specific items and their product numbers. Please include information as to whether in-house support is needed for use of product during OR cases.

**Flat Tip Laser Fibers**

“The fiber (flat) tip can damage the channel within a flexible endoscope. Therefore, they must be passed only when the channel is straight. For example, to reach calculus within the lower pole of the kidney with a flexible ureteroscope, the tip should be straightened within the renal pelvis and the fiber passed. The fiber is advanced into the field of view and then withdrawn until it is just no longer visible. At that point, it will be located approximately ½ mm beyond the tip of the ureteroscope. The tip can then be deflected and placed into the lower pole for treatment. The fiber can then be advanced.”¹

**TracTip Laser Fibers**
The ball shape tip is designed to reduce procedure steps associated with initial advancement of a deflected laser fiber to the treatment site. One step passage eliminates the need to re-access challenging stone locations.²,³

**The fiber should not be re-passed through a deflected scope once laser energy has been applied to the treatment site.**⁴
**Product Usage (cont.)**

**May reduce the risk of Flexible Endoscope Damage:**
According to “Evaluation of a New 240 micron Single Use Holmium:YAG Optical Fiber for Flexible Ureteroscopy” – “No (Flexiva) fibers fractured during 100 consecutive ureteroscopy procedures. The lack of fiber fracture during clinical use may reduce the risk of flexible endoscope damage due to fiber failure.”

**May reduce the risk of Laser Fiber Connector Failure:**
According to “Evaluation of a New 240 micron Single Use Holmium:YAG Optical Fiber (Flexiva) for Flexible Ureteroscopy” – During 100 consecutive ureteroscopy procedures,”No (Flexiva) fibers failed at the connector end.”

**Improved Scope Flexibility:**
With a 242 micron core size, the AccuTrac and Flexiva TracTip Laser Fibers are designed to improve scope flexibility over larger core fibers (273 micron & 365 micron).

According to the “Evaluation of 24 Holmium:YAG Laser Optical Fibers for Flexible Ureteroscopy,”

“We advocate using as flexible a fiber as possible to increase the likelihood that the stone can be reached and promote endoscope longevity.”

**Cost/Utilization**

**Is this item/technology on contract with GPO’s and/or IDN’s?**
Please speak to your Boston Scientific sales representative for the contract status of specific GPO’s & IDN’s.

**Ship Unit:** 1  
**UOM:** Each

**Mode of transportation:** FedEx® Delivery

**Minimum order quantity?** No

**Lead Time in working days?** 1-2 days

**What are the dimensions of the Product Carton?**
The Product Carton for the fiber is 16” x 12”

**What is the list price per each unit or unit of utilization?**
Each = $550.00  
Box 5 = $2,750.00

**Method of Purchase:** The purchase would be an outright purchase

**Does this item require special storage considerations?**
No, store in a cool, dry, dark place
Regulatory

Is this product FDA cleared for this intended use?
The AccuTrac Laser Fibers are intended for use in laser-based surgical applications, including but not limited to, endoscopic, laparoscopic and open surgical procedures involving vaporization, ablation and fragmentation of calculi (urinary and biliary) and surgical procedures involving vaporization, ablation, coagulation, hemostasis, excision, resection and incision of soft and cartilaginous tissue. The AccuTrac Laser Fibers are designed for use with Ho:YAG lasers with a standard SMA-905 connector that have been cleared for surgical use.

The Flexiva TracTip Laser Fibers are intended for use in laser-based surgical applications, including but not limited to, endoscopic, laparoscopic and open surgical procedures involving vaporization, ablation and fragmentation of calculi (urinary and biliary) and surgical procedures involving vaporization, ablation, coagulation, hemostasis, excision, resection and incision of soft and cartilaginous tissue. The Flexiva TracTip Laser Fibers are designed for use with Ho:YAG lasers with a standard SMA-905 connector that have been cleared for surgical use.

What is the FDA classification of this device?
The AccuTrac® Holmium Laser Fiber is marketed in the US in accordance with US 21 Code of Federal Regulations 878.4810 as a Laser Powered Surgical Instrument. Laser Fibers are Class II devices and are subject to the premarket notification (510k) process.

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Is this a dated product? Yes, with 3 year shelf life

Will this product require evaluation by any of the following departments?
• Epidemiology/Infection Control? No
• Bio Engineering Maintenance? No
• Safety and Security? No
• Pathology/Labs? No

Does this product contain metal substances that may affect tests and or procedures performed on patients? No

Does this item and its packaging contain no detectable latex? Yes

Is this product reusable? Single-Use

What additional waste or recycle costs are anticipated? None

Reimbursement

Is this product reimbursable by insurance?
The procedure for which it is used is reimbursable. A billing guide with respective coding and Medicare reimbursement for Ureteroscopy with laser lithotripsy is available upon request. For additional coding and reimbursement information, contact your local Territory Manager or the Urology Reimbursement Help Desk at (508) 683-4022.

What is the Medicare Pass-Through Code (aka C-code or HCPCS)?
Through Code for this product is 52353 (Ureteroscopy with laser lithotripsy - ISWL).

Is this a patient-chargeable product?
Yes. The appropriate Revenue Code is 272 - Medical / Surgical Supplies and Devices – Sterile Supply. Medicare does not dictate a provider’s charge structure or how it itemizes those charges. Section 2202.8 of the Medicare Provider Reimbursement Manual dealing with Ancillary Services (e.g. operating room) does not specifically address which items are part of the basic (routine) charge and which are charged in addition to the basic charge. Medicare is on record that it is up to the individual hospital to determine whether to and how to itemize the charge for a specific supply. However, Medicare does require that charges billed on the CMS-1450 form (aka UB-04) be aggregated under the appropriate Revenue Code.
Health economic and reimbursement information provided by Boston Scientific Corporation is gathered from third-party sources and is subject to change without notice as a result of complex and frequently changing laws, regulations, rules and policies. This information is presented for illustrative purposes only and does not constitute reimbursement or legal advice. Boston Scientific encourages providers to submit accurate and appropriate claims for services. It is always the provider’s responsibility to determine medical necessity, the proper site for delivery of any services and to submit appropriate codes, charges, and modifiers for services that are rendered. Boston Scientific recommends that you consult with your payers, reimbursement specialists and/or legal counsel regarding coding, coverage and reimbursement matters. Boston Scientific does not promote the use of its products outside their FDA-approved label.

1 "Endourologic Use of the Holmium Laser", 2001 Demetrius Bagley, Akhil Das
3 The fiber should not be re-passed through a deflected scope once laser energy has been applied to the treatment site.

VersaPulse PowerSuite is a trademark of Lumenis, Inc.

Caution: Federal Law (USA) restricts this device to sale by or on the order of a physician. Refer to package insert provided with the product for complete Indications for Use, Contraindications, Warnings, Precautions, Adverse Events, and Instructions prior to using this product.