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Virtual Ports Receives CE Mark Approval for Selling the EndoGrab™ in Europe

The EndoGrab is currently sold in the U.S. after receiving clearance from the FDA

Virtual Ports (www.virtual-port.com) announced today that it received European CE Mark approval for the EndoGrab™.

The EndoGrab™ reduces the number of incisions (or ports) needed to perform laparoscopic surgical procedures. EndoGrab™'s internal anchoring station replaces the port, the introducer and the retractor traditionally used during laparoscopic procedures; a demo can be viewed at http://www.youtube.com/watch?v=g_BZ0how05U. EndoGrab™ reduces scarring and is likely to minimize some of the complications associated with abdominal incisions – all without lengthening the operating time.

Udi Gordin, the Company's CEO stated, "We are pleased to receive the CE Mark and are preparing for marketing the EndoGrab™ in Europe through leading distributors with whom we are in advanced negotiations."
Sales of the EndoGrab™ were recently initiated in the U.S. through the Company's own sales department.

About Virtual Ports
Virtual Ports (http://www.virtual-ports.com) specializes in developing and marketing medical devices for abdominal and thoracic endoscopic surgeries. The Company is located in the Western Galilee. Two of the Company's products, the EndoGrab™ and the EndoClear™, are cleared for sale by the U.S. Food and Drug Administration (FDA).

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