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What is This?
A Promising New Device for the Prevention of Parastomal Hernia

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Abstract
Parastomal hernia (PSH) is the most frequent long-term stoma complication with serious negative effects on quality of life. Surgical revision is often required and has a substantial morbidity and recurrence rate. The development of PSH requires revisional surgery with a substantial perioperative morbidity and high failure rate in the long-term follow-up. Prophylactic parastomal mesh insertion during stoma creation has the potential to reduce the rate of PSH, but carries the risk of early and late mesh-related complications such as infection, fibrosis, mesh shrinkage, and/or bowel erosion. We developed a new stomaplasty ring (KORING), which is easy to implant, avoids potential mesh-related complications, and has a high potential of long-term prevention of PSH. Here we describe the technique and the first use.

Keywords
biomedical engineering, colorectal surgery, hernias, surgical oncology

Introduction
Parastomal hernia (PSH) is the most frequent long-term complication with serious negative effects on quality of life following stoma formation. PSH appears most commonly within the first 2 years after stoma formation with a prevalence of 35% to 50%. PSH often requires surgical repair with high morbidity and recurrence rates. Preventing PSH by prophylactic mesh insertion during stoma creation remains a challenging procedure, including tissue dissection, carrying the risk of serious mesh-related complications. Therefore, we developed a new stomaplasty ring (KORING), which is easy to implant, requiring limited tissue dissection and avoids potential mesh-related complications providing a high potential of long-term prevention of PSH.

Description of the Device
The new stomaplasty ring (KORING) consists of a non-absorbable and immunologically inert material (polyethylene terephthalate) with a flexible but not dilatable ring (Figure 1). The device is available in different diameters (2.5 cm, 3.5 cm, and 4.5 cm) enabling precise fitting to the stoma site. The KORING is sutured to the anterior rectus fascia, which distributes axial forces evenly to the ring preventing weakness of the fascia (Figure 1). The lip of the ring is inserted between the rectus muscle and the peritoneum to prevent direct contact of the ring with other bowel loops, avoiding bowel entrapment and adhesions (Figure 2).

The Operation
At the designated stoma site a circular skin incision is performed followed by blunt dissection through the subcutaneous tissue. Then a second circular incision is made in the external rectus fascia, the rectus muscle is preserved by blunt separation. Subsequently, the posterior rectus sheath (applicable cranial to the linea arcuata) and the peritoneum are incised. After choosing the correct diameter the ring is fixed to the external rectus fascia with 8 to 10 interrupted, nonabsorbable sutures to prevent constriction of the ring (purse-string effect).

The lip seal of the ring is then fitted between the peritoneum and the muscle (Figure 2). The intestine is then pulled through the ring and fixed according to common recommended practices to form the external orifice.

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The device was first used in a 76-year-old man undergoing abdominoperineal resection of the rectum with formation of an end-colostomy. The implantation of the ring took 10 minutes. No additional tissue mobilization was necessary. The braided structure of the ring made the fixation easy. After implantation the ring had no contact to the other bowel loops in the abdominal cavity and minimal contact with surrounding tissue (Figure 2). The postoperative course was uneventful.

**Summary**

Parastomal hernia is a serious stoma-related complication in terms of morbidity and socioeconomic impact. Prophylactic mesh insertion has shown promising results, remains a demanding procedure, and therefore has not yet become standard during stoma formation. The new stomaplasty ring (KORING) offers a simple and intuitive implantation avoiding large tissue dissection. It has no direct contact with the abdominal content and therefore potentially reduces the risk of adhesions and mesh infection. It may have a high impact on prevention of PSH in the future, improving quality of life of stoma patients as well as decreasing health care costs by preventing further surgical interventions due to PSH. Further clinical trials are necessary.

**Declaration of Conflicting Interests**

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