Confocal laser endomicroscopy for the diagnosis of diversion colitis

Confocal laser endomicroscopy (CLE) and virtual chromoendoscopy (VCE) are advanced endoscopic imaging techniques. *i*-Scan (Pentax, Japan) is a contrast-enhancing computed program [1], while CLE enables *in vivo* histological examination [2,3]. Studies from our group described the diagnostic utility of CLE and VCE, such as for Crohn’s disease [4–6]. However, data regarding the applicability of VCE and CLE for the diagnosis of diversion colitis have not been provided so far. Here, we describe the case of a 33-year-old woman who underwent an ileocecal resection and colorectal diversion 4 years previously because of a complicated ileocecal and rectal endometriosis with poor postoperative wound healing and the occurrence of a rectovaginal fistula. She reported diarrheic stools, which were still abundant (12–15 occurrences of diarrhea/day) even after a therapeutic attempt with high-dose steroids and antibiotics.

High definition endoscopy (EPK-i, Pentax, Japan) showed a highly inflamed and friable mucosa with large ulcerations, loss of haustration, spontaneous bleeding, and mucus mixed with pus in the lumen (Fig. 1a) of the diverted colorectum. Subsequent examination of the same area by CLE (Pentax EC-3870 CIFK, Japan) upon intravenous administration of fluorescein as a contrast agent, revealed a pronounced crypt rarefaction, hypervascularization with moderate leakage, and typical crypt abscesses (Fig. 1b).

Subsequent histopathological evaluation (hematoxylin and eosin stain) of conven-

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**Fig. 1**  
(a) High definition white-light endoscopy reveals a highly inflamed and friable mucosa with large ulcerations, loss of haustration, and mucus mixed with pus, in the lumen of the sigmoid colon. (b) Confocal laser endomicroscopy of the same colonic area shows rarefied crypts with large lumens which are invaded by a dense polymorph infiltrate (crypt abscesses), as well as hypervascularization with mild leakage in the lamina propria.

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**Fig. 2**  
(a) Conventional hematoxylin and eosin stain of a biopsy sample from the same area as in Fig. 1 shows similar crypt abscess findings (cross-sectioned crypt marked by green arrowheads, and longitudinal sectioned crypt highlighted by yellow arrowheads) confirming the endomicroscopic diagnosis. (b) A high definition virtual chromoendoscopy image (*i*-Scan) of an improved colonic mucosa with deposits (but without ulcerations) in the same patient after 10 weeks of topical salicylate therapy. (The intense erythematous aspect is due also to the contrast enhancement of the *i*-Scan mode.)
tional biopsies taken from the same location confirmed the endoscopic findings (Fig. 2a). There was no evidence of cytomegalovirus with the specific stain. As a therapeutic consequence, the patient received topical therapy with salicylates. In a control colonoscopy with i-Scan, performed approximately 10 weeks later, we saw an endoscopic improvement of the colitis (Fig. 2b), a fact that confirmed our diagnosis and choice of specific therapy.

In conclusion, this article reports the first description and in vivo diagnosis of diversion colitis after surgery, by virtual chromoendoscopy and fluorescein-guided CLE, respectively.

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Competing interests: None

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References
1 Kodashima S, Fujishiro M. Novel image-enhanced endoscopy with i-scan technology. World J Gastroenterol 2010; 16: 1043

Bibliography
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