

UPPER GASTROINTESTINAL SURVEILLANCE IN LYNCH SYNDROME

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Poster

Oesophagus

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Retrospective study of 349 Lynch syndrome patients found that push enteroscopy detected jejunal neoplasia not visible on standard upper endoscopy, with clinically relevant findings in 50% of patients overall.

KEY POINTS

- In 349 Lynch syndrome patients undergoing 960 endoscopic procedures, push enteroscopy exclusively detected jejunal adenomas in 2% of patients and jejunal cancer in 0.6%, while standard esophagogastroduodenoscopy detected none.
- Clinically relevant findings occurred in 50% of patients, including gastric adenomas in 3.7%, gastric cancer in 1.1%, duodenal adenomas in 3.7%, duodenal cancer in 0.9%, and intestinal metaplasia in 28%.
- In a separate cohort of 2859 Lynch syndrome patients from the German Consortium for Familial Intestinal Cancer, gastric cancer occurred in 2.2%, duodenal cancer in 2.1%, and jejunal cancer in 1.9%, predominantly in high-risk pathogenic variant carriers of MLH1, MSH2, and EPCAM.
- The authors conclude that push enteroscopy should be considered in high-risk Lynch syndrome surveillance protocols due to its diagnostic value beyond conventional esophagogastroduodenoscopy.

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