

Bowel cancer screening

Risk of colon cancer

The risk of colon cancer in the population in the Czech Republic is constantly increasing. The incidence reached 80 per 100 000 inhabitants in 2007 and is the highest in European comparison. Screening programs include methods of detection of occult bleeding – FOBT, detection of genetic markers and imaging colonoscopic, reps. sigmoidoscopic examination.

Development of molecular biology and applications of type PCR in routine diagnosis, they also open up completely new trends in GIT tumour screening. The latest screening methods are based on the detection of specific mutations by PCR or biochip technology in DNA isolated from a stool sample. For the detection of tumors colon mutations of APC, K-ras, p53, microsatellite instability (MSI) genes and other genetic markers can be determined from a stool sample.

Occult haemorrhage screening

Screening of occult haemorrhage for the detection and early diagnosis of colorectal tumours by the Haemoccult test is carried out in detail, the interval screening, age range and subsequent colloscopic examination programme with positive evidence of occult haemorrhage are established.

Hemoglobin concentration in faeces is an crucial question of setting a 'cut-off' value for screening. The physiological process defines daily blood losses of faust in the volume of 0.5-2.5 ml. If we recalculate this amount of blood by the concentration of haemoglobin in the blood (120-150 mg/ml) and the amount of stool in 24 hours (300-450 g), then we can consider the values of 0.3-1.3 mg hemoglobin per 1 g of stool as a physiological range. A healthy/physiological population with a concentration of 0.3-1.3 mg Hb/g stools and an Hb concentration curve in colorectal cancer can be viewed schematically. The detection cut-off for the standard Haemoccult test (gFOBT) is approximately 5 mg Hb/g stool, so it does not catch all KRCA, but should not detect any healthy individual positively. The level of the detection limit is discussed, e.g. for immunochemical tests - iFOBT.

Studies in recent years have tested several immunochemical analyzers for quantitative determination of haemoglobin in faeces (qi-FOBT), most of which are Japanese-made. ROC curves demonstrate a specificity for advanced adenomas 95,3 % at a sensitivity of 100 ng Hb/ml.

DNA test

Newly introduced molecular-genetic diagnostic methods have a higher sensitivity than the demonstration of haemoglobin in faeces. In the stool sample, the aberrantly methylated DNA of promoter regions *BMP3* and *NDRG4* and mutations in the *KRAS* gene are shown. The sensitivity of the method for carcinoma is 92%, for advanced adenoma as a significant precancerosis 42%^[1].

Links

Source

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Bibliography

- CALISTRI, D, et al. Quantitative fluorescence determination of long-fragment DNA in stool as a marker for the early detection of colorectal cancer. *Cell Oncol.* 2009, vol. 31, no. 1, s. 11-7, ISSN 1570-5870 (Print), 1875-8606 (Electronic). PMID: 19096146 (<https://pubmed.ncbi.nlm.nih.gov/19096146/>).
- AHLQUIST, DA, et al. Stool DNA and occult blood testing for screen detection of colorectal neoplasia. *Ann Intern Med.* 2008, vol. 149, no. 7, s. 441-50, ISSN 0003-4819 (Print), 1539-3704 (Electronic). PMID: 18838724 (<https://pubmed.ncbi.nlm.nih.gov/18838724/>).
- VAN ROSSUM, LG, et al. Random comparison of guaiac and immunochemical fecal occult blood tests for colorectal cancer in a screening population. *Gastroenterology.* 2008, vol. 135, no. 1, s. 82-90, ISSN 0016-5085 (Print), 1528-0012 (Electronic). PMID: 18482589 (<https://pubmed.ncbi.nlm.nih.gov/18482589/>).
- RENNERT, G, et al. Detecting K-ras mutations in stool from fecal occult blood test cards in multiphasic screening for colorectal cancer. *Cancer Lett.* 2007, vol. 253, no. 2, s. 258-64, ISSN 0304-3835 (Print), 1872-7980 (Electronic). PMID: 17349741 (<https://pubmed.ncbi.nlm.nih.gov/17349741/>).
- HAUG, U, et al. Mutant-enriched PCR and allele-specific hybridization reaction to detect K-ras mutations in stool DNA: high prevalence in a large sample of older adults. *Clin Chem.* 2007, vol. 53, no. 4, s. 787-90, ISSN : 0009-9147 (Print), 1530-8561 (Electronic). PMID: 17317884 (<https://pubmed.ncbi.nlm.nih.gov/17317884/>

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- MATSUSHITA, H, et al. A new method for isolating colonocytes from naturally evacuated feces and its clinical application to colorectal cancer diagnosis. *Gastroenterology*. 2005, vol. 129, no. 6, s. 1918-27, ISSN 0016-5085 (Print), 1528-0012 (Electronic). PMID: 16344060 (<https://pubmed.ncbi.nlm.nih.gov/16344060/>).
- GREENWALD, B. The stool DNA test: an emerging technology in colorectal cancer screening. *Gastroenterol Nurs.* 2005, vol. 28, no. 1, s. 28-32, ISSN 1042-895X (Print), 1538-9766 (Electronic). PMID: 15738729 (<https://pubmed.ncbi.nlm.nih.gov/15738729/>).
- OUYANG, DL. Noninvasive testing for colorectal cancer: a review. *Am J Gastroenterol.* 2005, vol. 100, no. 6, s. 1393-403, ISSN 0002-9270 (Print), 1572-0241 (Electronic). PMID: 15929776 (<https://pubmed.ncbi.nlm.nih.gov/15929776/>).
- WHITNEY, D, et al. Enhanced retrieval of DNA from human fecal samples results in improved performance of colorectal cancer screening test. *J Mol Diagn.* 2004, vol. 6, no. 4, s. 386-95, ISSN 1525-1578 (Print), 1943-7811 (Electronic). PMID: 15507679 (<https://pubmed.ncbi.nlm.nih.gov/15507679/>).
- IMPERIALE, TF, et al. Fecal DNA versus fecal occult blood for colorectal-cancer screening in an average-risk population. *N Engl J Med.* 2004, vol. 351, no. 26, s. 2704-14, ISSN 0028-4793 (Print), 1533-4406 (Electronic). PMID: 15616205 (<https://pubmed.ncbi.nlm.nih.gov/15616205/>).
- BRAND, RE, et al. Reproducibility of a multitarget stool-based DNA assay for colorectal cancer detection. *Am J Gastroenterol.* 2004, vol. 99, no. 7, s. 1338-41, ISSN 0002-9270 (Print), 1572-0241 (Electronic). PMID: 15233675 (<https://pubmed.ncbi.nlm.nih.gov/15233675/>).
- SONG, K, et al. Fecal DNA testing compared with conventional colorectal cancer screening methods: a decision analysis. *Gastroenterology.* 2004, vol. 126, no. 5, s. 1270-9, ISSN 0016-5085 (Print), 1528-0012 (Electronic). PMID: 15131787 (<https://pubmed.ncbi.nlm.nih.gov/15131787/>).
- LEVIN, B, et al. Emerging technologies in screening for colorectal cancer: CT colonography, immunochemical fecal occult blood tests, and stool screening using molecular markers. *CA Cancers J Clin.* 2003, vol. 53, no. 1, s. 44-55, ISSN 0007-9235 (Print), 1542-4863 (Electronic). PMID: 12568443 (<https://pubmed.ncbi.nlm.nih.gov/12568443/>).
- NISHIKAWA, T, et al. A simple method of detecting K-ras point mutations in stool samples for colorectal cancer screening using one-step polymerase chain reaction/restriction fragment length polymorphism analysis. *Clin Chim Acta.* 2002, vol. 318, no. 1-2, s. 107-12, ISSN 0009-8981 (Print), 1873-3492 (Electronic). PMID: 11880119 (<https://pubmed.ncbi.nlm.nih.gov/11880119/>).
- PRIK, L, et al. Diagnostic biochip array for fast and sensitive detection of K-ras mutations in stool. *Clin Chem.* 2002, vol. 48, no. 3, s. 428-35, ISSN 0009-9147 (Print), 1530-8561 (Electronic). PMID: 11861435 (<https://pubmed.ncbi.nlm.nih.gov/11861435/>).

Reference

1. IMPERIALE, Thomas F, David F RANSOHOFF a Steven H ITZKOWITZ. Multitarget stool DNA testing for colorectal-cancer screening. *N Engl J Med* [online]. 2014, vol. 371, no. 2, s. 187-8, dostupné také z <<https://www.ncbi.nlm.nih.gov/pubmed/25006736>>. ISSN 0028-4793 (print), 1533-4406.