Use of hospital resources in the Finnish colorectal cancer screening programme: a randomised health services study

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ABSTRACT

Objective: To estimate the difference in use of hospital resources in the Finnish Colorectal Cancer (CRC) screening programme between those invited and controls, within the year of randomisation and the next year.

Design: CRC screening was implemented in Finland in 2004 as a population-based randomised design using biennial faecal occult blood test (FOBT) for men and women aged 60–69 years. Those randomised to screening and control groups during years 2004–2009 were included in this analysis and use of hospital resources was estimated. Data were collected from the national register on hospital discharges. Outpatient visits, inpatient episodes and colonoscopies were compared between the two groups.

Results: The screening group comprised of 123 149 and control group of 122 930 people. Most people in both groups had not used hospital resources at all. More people in the screening group than in the control group had at least one hospital-based outpatient visit (7.8% vs 7.4%), inpatient episode (3.9% vs 3.8%) and colonoscopy (1.5% vs 1.3%). In total, the screening group had 31 975 and control group 27 061 cumulative outpatient visits, 9260 and 7903 inpatient episodes, and 2686 and 1756 hospital colonoscopies, respectively. The proportion of those with a positive FOBT result with at least one outpatient visit, one inpatient episode or one colonoscopy, was 3.7 times, 2.5 times or 9 times that of those with a negative FOBT result, respectively.

Conclusions: CRC screening using the FOBT slightly increased the volume of hospital outpatient visits, inpatient episodes and hospital colonoscopies in Finland.

INTRODUCTION

Colorectal cancer is the third most common cancer, following breast and prostate cancers, and the second most common cause of cancer deaths after lung cancer, in Europe, including in Finland. Within the 27 Member states of the European Union, 340 000 people were diagnosed with colorectal cancer in 2012 and about 150 000 people died because of it.1 Among 5.4 million Finns, the corresponding figures are approximately 2500 people diagnosed with colorectal cancer and more than 1000 deaths every year. The incidence of colorectal cancer increased for decades while mortality was stable among men and decreased among women since the 1970s.2 3

Colorectal cancer screening aims to reduce the mortality of the disease by discovering the cancer at an early, asymptomatic stage, to enable more effective and less invasive treatment. Screening with a faecal occult blood test (FOBT) has been shown to reduce mortality from colorectal cancer in a number of randomised screening studies.4–7 In a Cochrane review,8 the combined results of these studies showed a 16% reduction in the relative risk of colorectal cancer mortality.
Gradual expansion of the programme as a randomised level within municipality, birth year and in sex specific municipalities was randomised at an individual level. The target population in piloting municipalities has increased from 22 (of 444) in 2004 to 153 (of 336) in 2012. The target population of the Finnish colorectal cancer screening programme for colorectal cancer to the EU Member States.16

In Finland, colorectal cancer screening started in 2004 as a population-based randomised study. The idea was to launch a new screening programme in such a way that it can be reliably evaluated. The target population of screening is men and women aged 60–69 years, based on the fact that the incidence of and mortality from colorectal cancer increase after 60 years of age.3 The municipalities responsible for providing health services in Finland can voluntarily join the programme, and the number of piloting municipalities has increased from 22 (of 444) in 2004 to 153 (of 336) in 2012. The target population in these municipalities was randomised at an individual level within municipality, birth year and in sex specific blocks, to screening and control groups. Owing to the gradual expansion of the programme as a randomised health services study,17 the effects of the ultimate national screening programme on the allocation of health care resources can be evaluated as a part of routine health service.

The aim of this article was to estimate the difference in the use of hospital resources between those invited and the controls in the Finnish colorectal cancer screening programme within the first screening round.

MATERIALS AND METHODS

When the Finnish colorectal cancer screening study was launched, a national screening centre was established in the city of Tampere, at the local cancer society, to deal with the invitations, responses and recommendations for referrals, and to analyse the faecal samples. FOBT kits (Hemoccult; Beckman Coulter, USA) are posted to those offered screening along with a letter of invitation to participate in the programme and advice on how to take the sample. The kits are also returned by post. If any traces of blood are found in the sample, the person is referred to a contact nurse in their home municipality, for further examinations. Diagnostic confirmation, treatment and follow-up of people who screened positive are conducted within the normal health services at the local level and are paid for mostly by the home municipality of the participant being screened. Those in the control arm are identified but not contacted. The control population receives routine health services available in Finland.18

All individuals (n=246 079) who were randomised to the colorectal cancer screening (n=123 149) or control (n=122 930) groups during years 2004–2009 were identified and their data on healthcare resource use were gathered from the Finnish Hospital Discharge Register (FHDR). The follow-up of each individual was from the beginning of the year of randomisation up to the end of the next calendar year (two full years). The study period was 2004–2010.

For those randomised during the years 2004–2009, healthcare utilisation data on all gastrointestinal-related hospital-based outpatient visits and inpatient episodes of care were gathered on the year of randomisation and the following year. The data were restricted by ICD-10 codes that were related to gastrointestinal symptoms: C15-C26; D12 and K00-K93. Colonoscopies performed during those visits and episodes in hospitals were also examined, including procedure codes UJF32, UJF34 and UJF45. Colonoscopies performed in primary healthcare (including the screening colonoscopies) are not included in the analysis for either the screening or the control group. See table 1 for the explanation of the codes used in this article.

The FHDR is one of the oldest individual level hospital discharge registers covering an entire country. The FHDR has total (legislative) coverage of all inpatient care provided at university, general and psychiatric hospitals or primary care health centres, as well as treatment in military and prison wards and private hospitals, since 1969. Each record contains data on several variables, such as personal identity number, age at admission, gender, hospital identifier code, admission and discharge dates, reason for admission (ICD-10 code since 1996), and diagnostic and treatment activities. The register includes data on clinical procedures from 1986 onwards and the current coding system for procedures has been in use since 2004. The data on hospital outpatient visits have been collected since 1998 with some limitations and they are considered to be comparable from 2006 onwards. The completeness and accuracy of the FHDR has been reviewed to be good. Data on outpatient services at the primary care level (general practitioner visits, etc) are available from 1994 onwards at aggregate level.19 20

The percentage of people in the target population who used hospital services and the cumulative number of outpatient visits, hospital inpatient episodes and hospital colonoscopies, were calculated. We compared those randomised to screening to those in the control group. The screening group was also divided according to uptake of screening and test result. The differences between the arms in distributions of the individual variables were tested for statistical significance by the $\chi^2$ test.

RESULTS

The study population included 246 079 people, 123 149 (50%) in the screening group and 122 930 in the...
control group. Based on the randomised setting of the study, the groups were similar in demographic characteristics.

By 2009, the programme covered about 40% of the ultimate target population and 20% were invited to screening. The attendance rate within the first screening round was 68% during the years 2004–2009, with some variation by calendar year. During these years, 2944 individuals (2.4%) had a positive FOBT (table 2).

The number and proportion of people who had at least one gastrointestinal-related hospital outpatient visit, inpatient episode or colonoscopy during the year of randomisation or the next year, is presented in table 3. Most of the study population in both screening and control groups had not had any gastrointestinal-related hospital outpatient visits or inpatient episodes within the 2 years following randomisation. The proportion of people with at least one outpatient visit, inpatient episode or colonoscopy, was higher in the screening group than in the control group, for all the types of resource use evaluated. The difference between the groups was the largest in outpatient visits: there were 523

| Table 1 | Explanation of the diagnose and procedure codes used in the article |
|---------|---------------------------------------------------------------|
| ICD-10 codes | Malignant neoplasms of digestive organs |
| C15-C26 | Malignant neoplasm of oesophagus |
| C15   | Malignant neoplasm of stomach |
| C17   | Malignant neoplasm of small intestine |
| C18   | Malignant neoplasm of colon |
| C19   | Malignant neoplasm of rectosigmoid junction |
| C20   | Malignant neoplasm of rectum |
| C21   | Malignant neoplasm of anus and anal canal |
| C22   | Malignant neoplasm of liver and intrahepatic bile ducts |
| C23   | Malignant neoplasm of gall bladder |
| C24   | Malignant neoplasm of other unspecified parts of biliary tract |
| C25   | Malignant neoplasm of pancreas |
| C26   | Malignant neoplasm of other ill-defined digestive organs |
| D12   | Benign neoplasm of colon, rectum, anus and anal canal |
| K00-K93 | Diseases of the digestive system |
| K00-K14 | Diseases of oral cavity, salivary glands and jaws |
| K20-K31 | Diseases of oesophagus, stomach and duodenum |
| K35-K38 | Diseases of appendix |
| K40-K46 | Hernia |
| K55-K63 | Other diseases of intestines |
| K65-K67 | Diseases of peritoneum |
| K70-K77 | Diseases of liver |
| K80-K87 | Disorders of gallbladder, biliary tract and pancreas |
| K90-K93 | Other diseases of the digestive system |
| National procedure codes* | |
| UJF32 | Colonoscopy |
| UJF34 | Left colonoscopy |
| UJF45 | Flexible sigmoidoscopy with biopsy |

*Based on the Nordic Classification of Surgical Procedures.

| Table 2 | Number and percentage of individuals in the Finnish Colorectal Cancer Screening programme by year of invitation, arm and screening test result |
|---------|----------------------------------------------------------------------------------------------------------------------------------|
| Year of invitation | Screening | Negative test result | Non-attender |
|              | Positive test result | Per cent | n | Per cent | n | Per cent | n | Per cent | n |
| 2004 | 109 | 2.4 | 3283 | 72.3 | 1149 | 25.3 | 4539 | 4539 |
| 2005 | 542 | 2.3 | 16709 | 70.9 | 6311 | 26.8 | 23562 | 23565 |
| 2006 | 585 | 2.3 | 16625 | 66.8 | 7687 | 30.9 | 24897 | 24898 |
| 2007 | 513 | 2.4 | 14303 | 66.2 | 6778 | 31.4 | 21594 | 21375 |
| 2008 | 576 | 2.4 | 14756 | 60.9 | 8899 | 36.7 | 24231 | 24229 |
| 2009 | 621 | 2.6 | 15299 | 62.9 | 8406 | 34.6 | 24326 | 24324 |
| Total | 2944 | 2.4 | 80975 | 65.8 | 39230 | 31.9 | 123149 | 122930 |
people more in the screening group than in the control group with at least one gastrointestinal-related outpatient visit. The corresponding figures for people with at least one inpatient episode or hospital-based colonoscopy were 132 and 315, respectively. The people in the screening group were more likely to undergo more colonoscopies per person than those in the control group (figure 1).

A positive FOBT result increased the probability of using hospital resources. After a positive test result, a quarter of the people had at least one outpatient visit, while 95% of those with a negative test result had no outpatient visits at all. Also, the probability of undergoing colonoscopy while in hospital was 9.6 times higher in those with a positive test result compared to those with a negative test result. Furthermore, those who were invited to screening but did not attend were more likely to use hospital resources for gastrointestinal reasons than those with a negative FOBT result (table 3).

On average, 2.4% of the screening population had only one outpatient visit and 5.5% had two visits or more. The corresponding numbers for the control arm were 3.9% and 3.5%, respectively. Maximum number of visits per person was 86 in the screening group and 141 in the control group.

The cumulative number of outpatient visits, inpatient episodes and hospital colonoscopies over the whole study period is consistently higher in the screening arm than in the control arm (table 4). The difference between the arms was largest in the number of hospital colonoscopies: there were 53% more colonoscopies performed per individual in the screening group than in the control group. For the outpatient visits and inpatient episodes, the number of events was 18% and 17% higher, respectively, in the screening group compared to the control group.

DISCUSSION

According to this study, the vast majority of the study population, that is, 60–69-year-old Finnish men and women, had not used hospital resources at all for any gastrointestinal-related reasons. Slightly more people in the screening group than in the control group had at least one hospital-based outpatient visit (7.8% vs 7.4%) or inpatient episode (3.9% vs 3.8%). In total, the screening group had 4914 more cumulative outpatient visits and 1357 more inpatient episodes than the control group. The proportions of test-positive people with at least one outpatient visit (26.0% vs 7.0%) or with at least one inpatient episode (8.3% vs 3.3%) were clearly higher than those of people with a negative FOBT result.

The difference in use of hospital resources between the screening and control group is reliably comparable. The setting of the Finnish colorectal cancer screening programme makes it possible to evaluate the feasibility and effectiveness of screening in a randomised, yet real-world healthcare setting. Any possible systematic errors within the Finnish Hospital Discharge registry will be equally present in both groups.

| Event | Screening Positive test result | Screening Negative test result | Non-attender | Total | Control | p Value |
|-------|--------------------------------|--------------------------------|--------------|-------|---------|---------|
| Number of individuals | 2944 | 80 975 | 39 230 | 123 149 | 122 930 | <0.001 |
| Outpatient visits* | N | 764 | 5705 | 3190 | 9659 | 9136 | <0.001 |
| | Per cent | 26.0 | 7.0 | 8.1 | 7.8 | 7.4 | <0.001 |
| Inpatient episodes* | N | 243 | 2690 | 1911 | 4844 | 4712 | <0.001 |
| | Per cent | 8.3 | 3.3 | 4.9 | 3.9 | 3.8 | <0.001 |
| Colonoscopies | N | 339 | 942 | 593 | 1874 | 1559 | <0.001 |
| | Per cent | 11.5 | 1.2 | 1.5 | 1.5 | 1.3 | <0.001 |

The data includes the calendar years 2004–2010.
*ICD-10 codes C15-C26, D12 and K00-K93.
One limitation of the study is the lack of primary care data. Resources used at the primary healthcare level, either by the screening or the control groups, were not registered at FHHDR during the study years. Small municipalities with inadequate resources refer patients to hospital for the primary colonoscopy. Other municipalities have their own endoscopy units and their primary colonoscopies are not included in our data. In both cases, if any further examination or treatment (eg, removal of large polyps, suspicious cancer) is necessary, the patient is referred to the secondary healthcare. From earlier work, we know that 90% of the test-positives complied to colonoscopy, which means that approximately 2650 primary colonoscopies should be added to the figures of the screening group reported here. However, the lack of data on primary healthcare concerns both, and no comprehensive data are available on possible colonoscopies or other diagnostics at the primary healthcare level in the control group or non-attenders. According to national care guidelines, the primary investigation for symptomatic patients is endoscopy (not FOBT) and thus controls also have access to primary care endoscopy. The difference in use of primary healthcare services between the two groups can only be speculated, but it is likely that the proportional differences based on FHHDR data should indicate any overall differences.

A major concern in FOBT screening and resource use has been the possible increase in the number of diagnostic procedures, mainly colonoscopies, and our study confirms this. In the UK, Price et al reported an increase of 21–31% in the number of hospital colonoscopies due to FOBT screening, but a simultaneous increase was also seen in the demand for symptom-based colonoscopies. In Australia, it has been estimated that a biennial FOBT would result in five times more colonoscopies than the option of no screening. In our study, a total of 930 more hospital colonoscopies were conducted for the screening group than for the control group, resulting in a 53% increase in hospital colonoscopies. The proportion of people with at least one colonoscopy was slightly higher in the screening group than in the control group (1.5% vs 1.3%). In particular, patients with a positive FOBT result seemed to increase the probability of hospital resource use, including colonoscopies, compared to those with a negative FOBT or to those who did not attend the screening. This implies that an abnormal finding in the screening results in a number of colonoscopies while without screening, the control group will receive only one diagnostic colonoscopy based on the symptoms, without FOBT.

Based on the present study, it is, however, impossible to say whether the increase is only temporary. For example, Thiis-Evensen et al showed that, with a 9-year observation period after a screening colonoscopy, the screening group had 50% fewer usual-care distal endoscopic examinations than the control group. Furthermore, Seeff et al have argued that if all of the potential endoscopic capacity existing in the USA was used for organised colorectal cancer screening, the unscreened population could be screened within 1 year using the FOBT and colonoscopy for those with a positive test result. For example, follow-up after polypectomy is carried out at more frequent intervals than recommended. In comparison, many published estimates of colonoscopies in cost-effectiveness analyses are reported only for different screening strategies, while the alternative of ‘no systematic screening’ is often left out or assumed to be zero.

Our screening interval was 2 years. After that, the confounding effect of the second screen is introduced, presuming that the effects of the first and second round are different. In the Finnish CRC screening programme, everyone in the screening group will receive an invitation to screening also during the second round, regardless of the screening result in the first round. The long term evaluation is thus not that straightforward and may include a source of bias.

In conclusion, colorectal cancer screening using the FOBT seems to slightly increase the volume of hospital outpatient visits, inpatient episodes and hospital colonoscopies in Finland.

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