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Using resource modelling to inform decision making and service planning: the case of colorectal cancer screening in Ireland

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Abstract

Background: Organised colorectal cancer screening is likely to be cost-effective, but cost-effectiveness results alone may not help policy makers to make decisions about programme feasibility or service providers to plan programme delivery. For these purposes, estimates of the impact on the health services of actually introducing screening in the target population would be helpful. However, these types of analyses are rarely reported. As an illustration of such an approach, we estimated annual health service resource requirements and health outcomes over the first decade of a population-based colorectal cancer screening programme in Ireland.

Methods: A Markov state-transition model of colorectal neoplasia natural history was used. Three core screening scenarios were considered: (a) flexible sigmoidoscopy (FSIG) once at age 60, (b) biennial guaiac-based faecal occult blood tests (gFOBT) at 55–74 years, and (c) biennial faecal immunochemical tests (FIT) at 55–74 years. Three alternative FIT roll-out scenarios were also investigated relating to age-restricted screening (55–64 years) and staggered age-based roll-out across the 55–74 age group. Parameter estimates were derived from literature review, existing screening programmes, and expert opinion. Results were expressed in relation to the 2008 population (4.4 million people, of whom 700,800 were aged 55–74).

Results: FIT-based screening would deliver the greatest health benefits, averting 164 colorectal cancer cases and 272 deaths in year 10 of the programme. Capacity would be required for 11,095–14,820 diagnostic and surveillance colonoscopies annually, compared to 381–1,053 with FSIG-based, and 967–1,300 with gFOBT-based, screening. With FIT, in year 10, these colonoscopies would result in 62 hospital admissions for abdominal bleeding, 27 bowel perforations and one death. Resource requirements for pathology, diagnostic radiology, radiotherapy and colorectal resection were highest for FIT. Estimates depended on screening uptake. Alternative FIT roll-out scenarios had lower resource requirements.

Conclusions: While FIT-based screening would quite quickly generate attractive health outcomes, it has heavy resource requirements. These could impact on the feasibility of a programme based on this screening modality. Staggered age-based roll-out would allow time to increase endoscopy capacity to meet programme requirements. Resource modelling of this type complements conventional cost-effectiveness analyses and can help inform policy making and service planning.

Keywords: Colorectal cancer, Adenomatous polyps, Mass screening, Resource utilization, Markov chains

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Background

Various international organisations have recommended screening for colorectal cancer [1,2]. While screening programmes are in place, or being piloted, in several European countries [3], many other countries have not yet implemented screening. Several screening tests are available. These include endoscopy-based tests (such as colonoscopy and flexible sigmoidoscopy (FSIG)), radiology-based tests (such as barium enema and CT colonography) and faecal tests (such as the guaiac-based faecal occult blood (gFOBT) and faecal immunochemical test (FIT)). Each test has its own strengths and limitations [4]. In addition, while most cost-effectiveness analyses suggest that organised screening - by most of the available tests - is likely to be considered cost-effective, none of the tests appears optimal across all settings [5]. This creates considerable uncertainty for those tasked with making policy and providing services.

Cost-effectiveness is only one of the issues which needs to be considered when taking decisions on screening implementation. Long-established criteria from the World Health Organisation (WHO) recommend also considering the safety of the screening test and associated diagnostic investigations, the acceptability of the test, and availability of diagnosis and treatment facilities [6]. The challenge for policy-makers is, therefore, how to take all of these issues into account in weighing screening options against one another to determine the "best" strategy in a given setting. The challenge for health service planners is slightly different: it concerns how to deliver the screening tests and associated health services required for diagnosis, treatment and follow-up. Both of these challenges are further complicated by demographic changes in western populations, which mean that the population in the screening age-range is projected to rise. Moreover, screening tests which are repeated regularly (e.g. faecal tests) impact on the eligible population, since people diagnosed with cancer or adenoma(s) are no longer eligible to be invited for screening. Thus consideration needs to be given to capacity requirements immediately upon programme implementation and over the coming five, ten or twenty years. Cost-effectiveness analyses, with results expressed in costs per quality adjusted life year gained (QALY) or life years saved (LYS), may not help health service managers address such practical issues [7]. Other types of analyses, which express the real, immediate, impact on the health services of actually introducing a screening programme (such as budget impact [8] or other similar analyses) may be of more use. However, these other types of analyses are rarely reported.

We conducted a cost-effectiveness analysis of three strategies for colorectal cancer screening in Ireland: biennial gFOBT at ages 55–74, biennial FIT at ages 55–74 and once-only FSIG at age 60. All three options were

considered to be highly cost-effective compared to the *status quo* of no screening. The cost per QALY gained was €589 for FSIG, €1,696 for FIT and €4,428 for gFOBT [9]. FIT-based screening was expected to result in the greatest gain in QALYs and largest reductions in cancer incidence and mortality once fully implemented (i.e. the population cohort had completed 10 screening rounds). However, it was associated with major resource requirements over the lifetime of the cohort and higher rates of screening-related harms, including bowel perforations and deaths, than the other options. These results presented a significant challenge for decision-makers and service providers, raising questions about the feasibility, practicality and acceptability of a programme based on screening individuals aged 55–74 by biennial FIT. For example, there were questions about the actual resources required for diagnosis and treatment (for example, the numbers of colonoscopies which would need to be conducted), whether the service could deliver these, and the impact of higher than estimated uptake. Safety was also a concern, with questions about the actual number of people expected to suffer adverse effects of screening each year, and whether this was likely to be within acceptable limits. These questions stimulated queries about whether taking a different approach to screening delivery would make a programme based on FIT more feasible: for example, initially offering screening to a restricted age-group with gradual roll-out to incorporate the full age-range (as has been done in England in relation to the roll-out of the gFOBT-based screening programme). These types of questions cannot be answered by a conventional cost-effectiveness analysis.

In this paper we describe how we used a Markov state-transition model to estimate health service resource requirements and health outcomes, relative to the actual population of Ireland, over the first decade of a population-based colorectal cancer screening programme. These analyses were done with a view to aiding decision-making and service planning, and intended to complement the cost-effectiveness analyses.

Methods

Setting and screening scenarios

The country of Ireland is a republic comprising the largest part of the island of Ireland (the remainder of the island is Northern Ireland). The country has a population of 4.4 million, 700,800 of whom are aged 55–74. The healthcare system involves a mixture of public and private provision. Cancer screening programmes are provided free at the point of delivery, so the costs of provision fall on the healthcare payer, in this instance the Health Service Executive (HSE).

This analysis was part of a health technology assessment (HTA) of population-based colorectal cancer screening in Ireland. The commissioners, the Health

Information & Quality Authority, established an Expert Group to scope the HTA, and recommend which screening modalities and age-groups should be evaluated [10]. These decisions were informed by: the available scientific evidence at the time, age-specific colorectal cancer incidence and mortality in Ireland, and considerations of feasibility and likely acceptability. The three base-case scenarios recommended for evaluation were: (1) biennial gFOBT, with reflex FIT (i.e. those with a weak positive gFOBT would be invited to complete a FIT), in individuals aged 55–74 years; (2) biennial FIT in individuals aged 55–74 years; and (3) once-only FSIG at age 60. Investigation of positive screening tests was assumed to be by colonoscopy or, in those unfit for colonoscopy, by CT colonography. Follow-up of those who had adenomas detected and removed was assumed to follow existing guidelines [11]. For gFOBT and FIT, we assumed that approximately half the population aged 55 or 56 would be invited to participate in year one of the programme and the remainder in year two, and similarly for those aged 57 or 58, and so on. Thus biennial screening in the 55–74 age group involved a maximum of ten screening rounds.

Since FIT-based screening was considered the optimal option in terms of cost-effectiveness, health service requirements associated with alternative scenarios for implementing FIT-screening were also considered. The two main alternative implementation approaches involve either restrictions by area or by age. The former is considered unacceptable by the National Cancer Screening Service (NCSS) in Ireland, so only age-based implementation strategies were considered. The first variant involved restricting screening to the 55–64 age-group (i.e. a maximum of five screening rounds). The second and third variants involved staggered roll-out of biennial screening in the 55–74 age-group. “Medium implementation” involved inviting those aged 55 and 65 in year one; those aged 55, 57, 65 and 67 in year two; those aged 55, 57, 59, 65, 67 and 69 in year three; and so on until the full age range would be included in year five. “Slow implementation” involved inviting those aged 55 in year one; those aged 55 and 57 in year two, those aged 55, 57 and 59 in year three; and so on until year ten when the full age-range would be included. These scenarios were not intended to represent the only or preferred options, but rather were chosen to illustrate the health service impact of different possible approaches to implementation.

Estimating resource use and health outcomes

“Conventional” analysis of a Markov model - for the purposes of cost-effectiveness analysis - follows a single notional cohort, usually comprising 100,000 individuals, over their lifetime accumulating costs and benefits as the cohort ages. The approach adopted for this analysis was similar to that taken in a budget impact analysis [8].

We considered a short time horizon - the first 10 years following screening implementation. We estimated requirements separately for a range of resources (from screening tests to colorectal resections). We expressed our results in relation to the actual target population in Ireland. To do this, we ran our economic model, described below, using a cross-sectional whole population approach. We established 70 different age-cohorts, corresponding to the 2008 Irish population; cohort one comprised individuals aged 30, cohort two comprised individuals aged 31, and so on. We followed these 70 age-cohorts through the model for 10 years. For each cohort, the screening-related resources required (e.g. screening tests, diagnostic tests, etc.) and health outcomes experienced (e.g. numbers diagnosed with adenomas, had bowel perforations) were estimated, then summed across the whole population (i.e. all 70 cohorts) for each year following screening implementation. The predictions incorporated changes in the population age distribution over time and lower cancer prevalence in later screening rounds.

Resources associated with screening were assumed to be provided by the HSE and, within that, the NCSS. The health service resources estimated included those associated with: screening (gFOBT and FIT kits dispatched and returned; FSIG examinations); and diagnosis and treatment of screen-detected lesions (up to and including the point of surgery) and surveillance of individuals with screen-detected adenomas (colonoscopy, CT colonography, pathology, diagnostic radiology, neoadjuvant radiotherapy (with or without chemotherapy) and colorectal resection). The health outcomes considered were the numbers of individuals who were diagnosed with screen-detected adenomas and cancers and the numbers who sustained complications (major abdominal bleeding requiring hospital admission, bowel perforation, death following perforation).

Model

We used a Markov state-transition model which is described in detail elsewhere [9,12]. Briefly, the model included three interlinked components relating to: (a) the natural history of colorectal neoplasia; (b) screening scenarios, and subsequent adenoma surveillance; and (c) mortality. The natural history model simulated the experience of a cohort of individuals over their lifetime (from age 30 to 100) through a finite number of defined health states relating to the progression from having normal colorectal epithelium through the adenoma-carcinoma sequence to death (Additional file 1: Figure S1). The screening intervention model was superimposed upon the natural history model and the impact of the screening and diagnostic tests, and management of adenomas and cancers, modelled by re-distributing the cohort across health states in each Markov cycle. The mortality model allowed

for deaths due to colorectal cancer, endoscopic bowel perforation, and other causes.

Parameters and calibration

To populate the model, comprehensive literature reviews were undertaken. These were augmented with data and results from population-based screening programmes and pilot programmes. When relevant data was unavailable, values were based on expert clinical opinion. Table 1 shows the parameter values and the sources from which they were derived [13-49].

Base-case estimates of screening uptake and colonoscopy compliance came from the UK pilot colorectal cancer screening programmes and the UK FSIG trial [34,35,38]. GFOBT and FIT uptake were assumed to be the same and not to change over screening rounds. Compliance with post-polypectomy surveillance was assumed to be the same as with diagnostic colonoscopy. GFOBT and FIT sensitivity and specificity values were derived from pooled analysis of information from diagnostic cohort studies of “screening populations” (i.e. the diagnosis had not been determined prior to recruitment and all participants underwent the index test and the reference “gold-standard” test) and, for gFOBT, which used Hemocult[®] or Hemocult II[®] [13-32]. While we did not set out to evaluate FIT at different cut-off values (because of lack of high-quality data at the time the model was parameterized) the key studies which informed the parameter estimates for FIT performance characteristics used a cut-off of 100 ng/mL. Three studies were combined to estimate FSIG sensitivity for intermediate/high-risk adenomas (adenomas ≥ 10 mm in size) [18,20,33]. Estimates of sensitivity for low-risk adenomas (<10 mm) and cancers, and specificity, were based on expert clinical opinion; these estimates were based on the assumption that sensitivity for low-risk adenomas would be lower than that for high-risk adenomas and sensitivity for cancer would be higher than that for high-risk adenomas. Colonoscopy performance characteristics were based on “miss rates” from studies of individuals who had undergone tandem colonoscopies [41,42]. CT colonography performance characteristics were based on expert opinion informed by reviews and large-scale studies [43-46]. It was assumed that, on average, 1.9 adenomas would be removed from each person undergoing polypectomy [47]. The probability of perforation following FSIG was derived from the UK FSIG Trial [38]. For colonoscopy, probabilities of perforation (with and without polypectomy) were estimated from a Swedish nationwide colonoscopy audit [49]. Estimates of the probability of death following perforation came from Gatto et al. [48]. Estimates of major bleeding following endoscopy procedures were from the UK FSIG Trial [38].

Several parameters, including natural history transition probabilities, could not be empirically observed and were obtained by calibration: This involved fitting the model to colorectal cancer incidence (by stage) and mortality data for Ireland. Further details of the approach and the transition probabilities are provided elsewhere [9,50]. The colorectal cancer incidence and mortality figures predicted by the model were very close to the observed values [9].

Analysis

The primary analysis concerned absolute screening impact (i.e. annual numbers of each health service resource required, and each health outcome, under each screening scenario). One-way sensitivity analyses explored the impact of varying (a) screening uptake, (b) screening test sensitivity and (c) screening test specificity. Values for lower and higher uptake were based on experience in other countries [36,37,39,40]. For test sensitivity and specificity, the values used were the lower and upper 95% confidence intervals obtained from the pooled data used to derive the base-case values.

The secondary analysis assessed the relative impact of screening. For each core screening scenario, we estimated the additional resources needed for diagnosis and treatment, and the population-level health gains, compared to those required or generated under a policy of no screening.

Results

Primary (absolute) analysis of base-case scenarios

The model predicted that, in year one of a programme based on gFOBT or FIT, 357,812 individuals would be sent test kits, and 189,640 completed kits would be returned for laboratory processing (Table 2). With a FSIG-based programme, in year one, an estimated 18,617 individuals would undergo screening. Assuming uptake remained constant, between years one and 10, the number screened by FIT or gFOBT would increase by 16-17% and by FSIG by 11%, due to demographic changes.

For FIT-based screening, in year one, resources would be required to perform 11,095 diagnostic colonoscopies and 1,441 CT colonographies, rising to 12,414 colonoscopies and 1,614 CT colonographies in year 10. The diagnostic resources required under gFOBT screening would be one-tenth of those required for FIT. With once-only FSIG, only 381-423 individuals would undergo diagnostic colonoscopy each year. Similar patterns were evident in the colonoscopy and CT colonography resources required for surveillance of those with intermediate/high-risk adenomas removed. As a consequence of the numbers of colonoscopies, more individuals would suffer adverse events with

Table 1 Parameter estimates

<i>Model parameter</i>	<i>Base-case value</i>	<i>Lower and upper values in sensitivity analysis</i>	<i>References</i>
<i>Performance of screening tests</i>			
gFOBT sensitivity for adenomas	11%	10%, 12% ¹	[13-21]
gFOBT sensitivity for CRC	36%	31%, 42% ¹	
gFOBT specificity for adenomas and CRC	97%	96%, 98%	
FIT sensitivity for adenomas	21%	19%, 22% ¹	[16,22-32]
FIT sensitivity for CRC	71%	67%, 75% ¹	
FIT specificity for adenomas and CRC	95%	94%, 96%	
FSIG sensitivity for low-risk distal adenomas	65%	60%, 70% ¹	Expert opinion informed by [18,20,22]
FSIG sensitivity for intermediate/high-risk distal adenomas	74%	68%, 78% ¹	
FSIG sensitivity for distal CRC	90%	85%, 95% ¹	
FSIG specificity for distal adenomas and CRC	92%	90%, 95%	
<i>Uptake and compliance with screening, diagnosis and surveillance</i>			
gFOBT uptake	53%	32%, 70%	[34-37]
FIT uptake	53%	32%, 70%	
FSIG uptake	39%	24%, 67%	[38-40]
COL compliance (diagnostic test or adenoma surveillance)	86%	-	[34,35]
<i>Performance of diagnostic tests and related parameters</i>			
COL sensitivity for low-risk adenomas	77%	-	[41,42]
COL sensitivity for intermediate/high-risk adenomas	98%	-	
COL sensitivity for CRC	98%	-	
COL specificity for adenomas and CRC	97%	-	Expert opinion
CTC sensitivity for low-risk adenomas	53%	-	Expert opinion, informed by [43-46]
CTC sensitivity for intermediate/high-risk adenomas	85%	-	
CTC sensitivity for CRC	85%	-	
CTC specificity for adenomas and CRC	86%	-	
Average number of adenomas removed per person	1.9	-	[47]
% of those with intermediate/high-risk adenomas removed in whom the adenoma was high-risk	29%	-	[34]
<i>Harms of screening</i>			
FSIG probability of perforation (with or without polypectomy)	0.002%	-	[38]
FSIG probability of death following perforation	6.452%	-	[48]
Probability of (major) bleeding following FSIG	0.029%	-	[38]
COL probability of perforation (with polypectomy)	0.216%	-	[49]
COL probability of perforation (without polypectomy)	0.107%	-	
COL probability of death following perforation	5.195%	-	[48]
Probability of (major) bleeding following COL	0.379%	-	[38]

COL colonoscopy, CRC colorectal cancer, CTC CT colonography, FIT faecal immunochemical test, gFOBT guaiac faecal occult blood test, FSIG flexible sigmoidoscopy; low-risk adenoma(s), <10 mm; intermediate/high-risk adenoma(s), ≥10 mm. ¹ parameters varied simultaneously in sensitivity analysis; - parameters not varied in sensitivity analyses.

screening based on FIT, than with screening based on gFOBT or FSIG. With FIT, in year 10, an estimated 62 people would suffer major abdominal bleeding resulting in hospital admission, and there would be 27 bowel perforations and one associated death.

The yield of adenomas and cancers was predicted to be higher for FIT-based screening than the other two options, and therefore the resources required to manage these would be much greater. In year one, under FIT-based screening, 7,161 lesions (6,308 adenomas and 853

Table 2 Estimated screening-related resource use and health outcomes by year and screening scenario: biennial gFOBT at 55–74 years, biennial FIT at 55–74 years, and FSIG once at 60 years

Year of programme Screening scenario		Year 1			Year 10		
		gFOBT	FIT	FSIG	gFOBT	FIT	FSIG
Screening-related resource use							
Screening tests	No. of kits sent out	357,812	357,812		420,151	417,464	
	No. of kits processed	189,640	189,640		222,637	220,999	
	No. of FSIG done ¹			18,617			20,625
COL/CTC	No. of diagnostic COL	967	11,095	381	1,103	12,414	423
	No. of diagnostic CTC	126	1,442	50	143	1,614	55
	No. of surveillance COL	0	0	0	297	2,406	620
	No. of surveillance CTC	0	0	0	39	313	81
Pathology	No. of adenomas and CRCs requiring pathology ²	1,004	7,161	1,599	1,356	8,909	2,222
CRC work-up and treatment	No. receiving PET scan	31	85	6	34	69	8
	No. receiving MRI scan	111	307	23	121	247	28
	No. receiving CT scan(s)	309	853	64	336	687	78
	No. receiving TUS	16	43	3	17	35	4
	No. receiving pre-operative radiotherapy ³	71	196	15	75	146	17
	No. undergoing colorectal resection ⁴	281	779	59	307	635	71
Screening-related health outcomes							
Harms⁵	No. with major bleeding following endoscopy	4	48	7	6	62	10
	No. with perforation following endoscopy	2	21	1	2	27	2
	No. of deaths from perforation following endoscopy	0	1	0	0	1	0
Screen-detected lesions	No. with adenoma(s) ⁶ Total	366	3,320	808	537	4,327	1,128
	low-risk	229	2,081	544	333	2,770	740
	intermediate/high-risk	137	1,239	264	204	1,558	388
	No. with CRC ⁷ Total	309	853	64	336	687	78
	stage I	111	308	23	130	303	31
	stage II	105	293	22	115	237	26
	stage III	69	192	14	68	115	16
	stage IV	24	60	4	23	31	5

COL colonoscopy, CRC colorectal cancer, CTC CT colonography, gFOBT guaiac-based faecal occult blood test, FIT faecal immunochemical test, FSIG flexible sigmoidoscopy, TUS ultrasound; intermediate/high-risk = adenoma(s) ≥10 mm; low-risk = adenoma(s) <10 mm;

¹ includes FSIG with and without polypectomy.

² sum of number of adenomas and colorectal cancers requiring pathology, assuming average of 1.9 adenomas per person; includes screen-detected and surveillance-detected adenomas.

³ applies to rectal cancer only; includes radiotherapy given with or without chemotherapy.

⁴ sum of estimates of colon and rectal resections required.

⁵ includes complications from diagnostic and surveillance endoscopy, including FSIG where relevant.

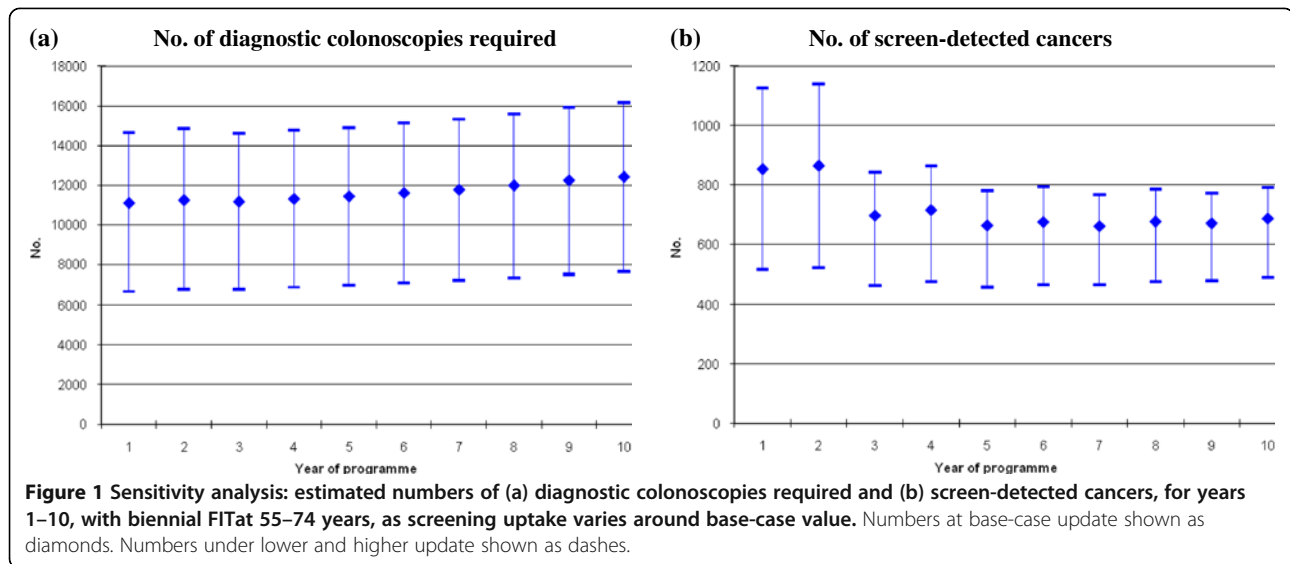
⁶ includes individuals with screen-detected and surveillance-detected adenomas.

⁷ includes individuals with CRC detected at screening and at surveillance.

cancers) would require pathological analysis, compared to 1,599 (1,525 and 64) using FSIG, and 1,004 (695 and 309) using gFOBT. Resources would be required to conduct 774 resections of screen-detected cancers in year one with FIT screening, compared to 280 with gFOBT and 59 with FSIG. In year 10, these numbers would rise to 635, 307 and 71 for FIT, gFOBT and FSIG screening, respectively.

Sensitivity analyses

Numbers of diagnostic colonoscopies and screen-detected cancers estimated from the sensitivity analyses are shown in Additional file 2: Table S1 for all three screening scenarios, and in Figure 1 for biennial FIT at 55–74 years. Varying screening uptake had the greatest impact. For example, for FIT-based screening, compared to base-case uptake (53%), with low uptake (32%) the annual number



of diagnostic colonoscopies required would be around 40% lower; with high uptake (70%) it would be almost one-third higher. Compared to base-case uptake, with low uptake 40% fewer cancers would be detected by screening in year one and 29% fewer in year 10; with high uptake, 32% more cancers would be detected in year one and 14% more in year 10. Varying specificity had a notable effect on colonoscopy requirements; for example, for FIT-based screening, in year one, high specificity (96%) increased the annual number of diagnostic colonoscopies by 13%, while low specificity (94%) reduced colonoscopy requirements by 13%. The impact of varying sensitivity was modest (1–6% variation around base-case values).

Alternative FIT scenarios

Additional file 3: Table S2 shows the resource requirements and health outcomes for the three alternative implementation options for FIT-based screening. Resource requirements under the age-restricted option (FIT at 55–64 years) were approximately 60% of those required for screening the full age-range (e.g. 6,437–7,070 diagnostic colonoscopies annually vs 11,095–12,414 for the base-case). Only around half the number of cancers and adenomas would be detected under this scenario compared to screening the full age-range. Under the medium roll-out option for the full age-group, resource requirements would start low (2,441 diagnostic colonoscopies in year one vs 11,095 for the base-case scenario) rising to the same as those for the base-case by year 10. Numbers of adenomas and cancers detected followed a similar pattern. The slow roll-out option generated the lowest resource requirements in early years, reaching around 60% of the base-case requirement by year 10, but only 317 cancers would be detected in year 10 under this scenario compared to 687 under the base-case.

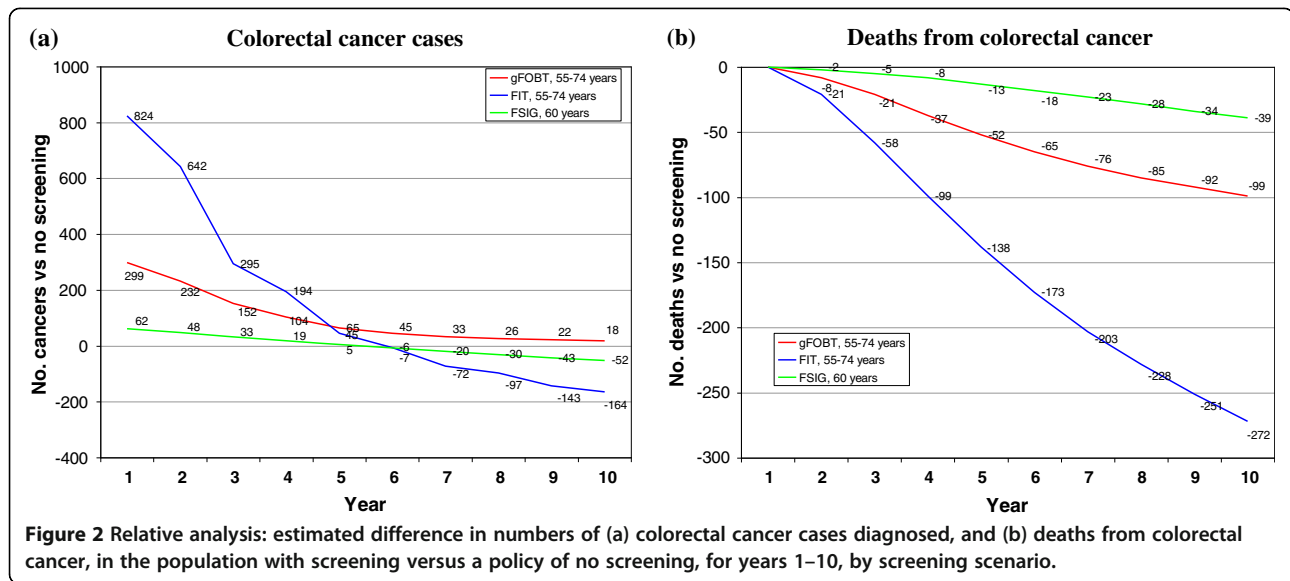
Relative (secondary) analysis of base-case scenarios

Compared to a policy of no screening, screening by gFOBt, FIT and FSIG all had potential to reduce deaths from colorectal cancer in the population within a decade of implementation. This reduction was evident from the second year, and was greatest for FIT-based screening (Figure 2). FIT-based screening at ages 55–74 would result in 272 colorectal cancer deaths avoided in the population in year 10 (26% fewer deaths than under no screening). In the first five years of screening, all three scenarios would generate an overall increase in colorectal cancer cases (Figure 2). After this time, with screening based on FIT or FSIG, the number of cases would fall below the number expected with no screening; this fall would be greatest with FIT (164 cases averted in year 10; 7% reduction compared to no screening).

Discussion

Value of resource modelling

The resource modelling described in this paper could be viewed as a particular type of budget impact analysis, with costs expressed in terms of resources and harms rather than monetary units. As in budget impact analyses [8], we focussed on the short-term health service impact of adopting technologies and generated results specific to the actual target population. Like budget impact analyses, the types of analyses shown here are intended to address the practical needs of both decision-makers and health service planners. They are also intended to complement more conventional cost-effectiveness analyses. In fact, these analyses did inform decision making in Ireland: a population-based colorectal cancer screening programme in 2013 commenced (see below) [51], and the estimates were used in planning the implementation of the programme. Our findings suggested that the



annual endoscopy requirements associated with screening the 55–74 age-group by biennial FIT would be difficult, if not impossible, for the health services to meet in the short-term. However, the magnitude of the health gains that could be achieved within five years of screening implementation meant that FIT-based screening remained the preferred option of policy-makers, despite the lower resource requirements for gFOBT and FSIG. Therefore, the policy decision was to introduce biennial FIT screening; initially the programme will invite individuals aged 60–69 to be screened with the intention of eventually rolling-out screening to the full 55–74 age-range.

Delivering sufficient capacity

The few previous analyses of the health service impact of colorectal cancer screening [12,52,53] all noted that the ability of the health services to deliver sufficient endoscopy services was a crucial determinant of programme feasibility. In England, for example, although screening based on once-only FSIG was the most cost-effective option [12], the associated requirements for endoscopy facilities and staff were considered impossible to meet in the short-term; instead, a programme based on gFOBT was implemented.

In this study, we found that a programme based on FIT the option considered most cost-effective [9] – would require substantially more resources for colonoscopy, CT colonography, and other diagnostic procedures, than one based on gFOBT (with FIT as the second-line test in those who test weakly positive) or FSIG. FIT-based screening will generate demand for 11,000–15,000 additional colonoscopies annually in Ireland. In Ireland, in 2008, 44,000 publicly-funded colonoscopies were performed, 2,000 of these under the National Treatment Purchase Fund. This fund provides for procedures to be undertaken in private

facilities in the event of long-waiting times in the public system. In April 2009 more than 2,300 individuals had been waiting for a colonoscopy for more than three months [54]. These figures suggest there is little spare capacity in the service, at least under current working patterns. They also justify the assumption, inherent in our analysis, that meeting the demand for “screening-generated” colonoscopies would require additional capacity over and above existing services.

If not prepared for, increases in colonoscopy demand can increase waiting times [3]. Studies of screening programmes for other cancers have found that the period between having a positive screening test and waiting for diagnostic follow-up can be a time of considerable anxiety for the individuals concerned (see, for example, [55]). Although this issue does not appear to have been investigated to any great extent for colorectal screening, it is entirely plausible that many people will be distressed or anxious in the time between receiving a positive colorectal screening test result and attending for colonoscopy. Since anxiety and distress can be considered a cost of screening, serious consideration should be given to ensuring that as many people as possible undergo diagnosis and treatment (if required) as quickly as possible. Thus, while recent studies continue to build the clinical case for screening with FIT in preference to other tests [56–59], the implication of our results is that any FIT-based programme – in Ireland or elsewhere – will have to plan carefully how to deliver sufficient services for initial diagnosis and ongoing surveillance. The types of analyses illustrated here, and elsewhere [60], could usefully inform that planning process.

Strategies that might be used to help address endoscopy capacity and other resource challenges include

restricting screening to a narrower age-range, and/or gradually phasing in screening across the full age-range. Although restricting FIT-based screening to those aged 55–64 would reduce resource requirements by around 40%, our cost-effectiveness analyses concluded that, when fully implemented this strategy would be slightly less desirable than screening the 55–74 age group [9]. This suggests that gradual roll-out across the full age-range would be preferable, and indeed this is what has been implemented in Ireland, albeit in a different age-group initially to that considered here [51]. However, it is worth remembering that the short-term health gains achieved with such phased-in screening will be lower than if the full age-group was screened from the outset.

Safety

Safety is a key consideration in the implementation of any screening programme [6]. The chances of bowel perforation and death are, in fact, the major reasons why some – including the NCSS in Ireland – consider colonoscopy unsuitable as a primary screening tool [61]. Colonoscopy is, however, the main option for diagnostic follow-up of those with positive screening tests. Our analysis suggest that the much higher numbers of colonoscopies generated by FIT-based screening than the other options will result in a higher number of adverse events. A higher adverse event rate was also evident in our cost-effectiveness analysis [9]. However, the advantage of the results presented here is that they express the actual numbers of events that might be expected to occur in the real-world eligible population: these values are likely to be more informative for the purposes of assessing and managing risk.

Screening uptake

A screening test must be acceptable to the population [6] and one of the main measures of acceptability is screening uptake. As might be expected, the resource requirements and health gains attainable with screening were highly dependent on estimates of uptake. For all screening tests, if actual uptake was higher than our base-case estimates more cancers would be averted; however, resource requirements would be higher and there would be more colonoscopy-related adverse events. On the other hand, if uptake was lower than the base-case value, resource requirements would be lower and there would be fewer adverse events, but there would also be lower health gains.

Since it is impossible to be entirely certain what uptake might be attained for any screening test in any population, base-case values were chosen to be conservative. For the faecal tests, the base-case estimate of 53% was based on the second round of the UK pilot programmes [34,35]. Although this is lower than

considered desirable by the NCSS in Ireland (60%) [62], it is higher than in some other European programmes and pilot programmes [58,63–65]. In Ireland, 51% of almost 10,000 Dublin residents invited to complete a FIT did so [66], suggesting the base-case estimate may be reasonably realistic. For FSIG, there was no local data to inform what level of uptake might be realistic. Although participation rates of 55% and higher have been achieved in some settings [40,67], generally uptake has been lower [58,63]. We decided to derive our base-case value from the large UK Flexible Sigmoidoscopy Trial [38] to provide some consistency with the source of the estimates of uptake for the faecal tests.

Since we conducted our analysis evidence has accrued that uptake may be higher with FIT than gFOBT [68], although this has not been a universal finding [59]. If FIT is more acceptable than gFOBT, our estimates of the resource requirements for FIT relative to gFOBT may be conservative. However, it is worth noting that the difference in numbers of screening-related harms may also be under-estimated.

Limitations

Although parameter values were based on review of the best available evidence, as with all modelling studies, some uncertainty remains. Even for gFOBT, which has been shown to reduce mortality in several trials [69], performance characteristics remain uncertain. This is particularly true for newer versions of the test [70]. For FIT, the available tests have heterogeneous performance characteristics [70] and for FSIG, information on sensitivity and specificity is limited [18,20,33]. For all tests, assessment of true sensitivity and specificity is further complicated by the fact that the reference (gold) standard, colonoscopy, misses lesions [41]. We addressed this uncertainty through sensitivity analyses, and found that varying specificity, in particular, had a notable impact on requirements for diagnostic investigations. Hence, if the true test specificity differs from the values used here, the resource requirements and health outcomes might differ from those estimated.

We assumed, for gFOBT and FIT, that sensitivity and specificity remained constant across all screening rounds. Recent evidence suggests that the ability of gFOBT-based programmes to identify cancers might decline with repeated screenings [71]. The same may be true for FIT. If so, this would suggest that we may have over-estimated the health gains associated with screening programmes based on FIT and gFOBT, although the extent of this over-estimation is difficult to quantify from currently available evidence.

We evaluated a strategy that combined gFOBT with reflex FIT, instead of the more conventional approach of reflex gFOBT. This was because second-line FIT has

been shown to limit the number of referrals to colonoscopy [72,73]. Finding effective strategies to “adjust” screening when colonoscopy capacity is limited is very topical [60]. Although gFOBT with reflex FIT is currently not widely used, our results indicate that it could be an attractive option for delivering population screening in settings where capacity for diagnostic investigation is expected to be limited.

We did not estimate resources required for adjuvant chemotherapy in screen-detected colon cancers, or for follow-up after resection (which can involve annual colonoscopy or CT colonography for five years) since responsibility for providing these would most likely fall outwith the screening programme/NCSS. Facilities would, however, need to be provided elsewhere in the health services, so this distinction is somewhat arbitrary.

Conclusions

This study illustrates how resource modelling complements cost-effectiveness analyses and can help inform health service decision making and planning. Our results show that the introduction of FIT-based screening would quite quickly generate attractive health outcomes – in terms of reduced numbers of colorectal cancer cases and deaths – but that it has heavy resource requirements. These heavy requirements could impact on the feasibility of any programme based on this screening modality. Staggered age-based roll-out of FIT-based screening would allow time to increase endoscopy capacity to meet requirements. Overall our findings suggest that there are likely to be many challenges ahead in delivering a high-quality, safe, and acceptable screening programme to the population of Ireland.

Additional files

Additional file 1: Figure S1. Simplified diagram of Markov states in natural history mode.

Additional file 2: Table S1. Sensitivity analysis: estimated numbers of diagnostic colonoscopies required and screen-detected cancers, for years one and 10, by screening scenario, as test sensitivity, test specificity and screening uptake vary around base-case values.

Additional file 3: Table S2. Scenario analysis: estimated screening-related resource use and health outcomes by year for alternative FIT implementation options: biennial FIT at 55-64 years, biennial FIT at 55-74 years (medium roll-out), and biennial FIT at 55-74 years (slow roll-out)¹.

Abbreviations

CT: Computed tomography; FIT: Faecal immunochemical test; FSIG: Flexible sigmoidoscopy; gFOBT: guaiac-based Faecal Occult Blood; HSE: Health Service Executive; LYS: Life years saved; MRI: Magnetic resonance imaging; NCSS: National Cancer Screening Service; PET: Positron emission tomography; QALY: Quality adjusted life year; UK: United Kingdom; WHO: World Health Organisation.

Competing interests

None of the authors have any conflicts of interest to declare.

Authors' contributions

All authors were involved in the design of the study, interpretation of data and commenting critically on the manuscript. In addition, LS co-ordinated the study, drafted the initial version of the manuscript, and undertook systematic reviews together with LT, AOC and CU. LT, CU and MB collated cost data. PT, JC and SW developed and modified the Markov model. CW provided mathematical and statistical input. All authors read and approved the final manuscript.

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References

1. Council of the European Union: **Council recommendation of 2 December 2003 on cancer screening (2003/878/EC)**. *Off J Eur Union* 2003, **L327**:34–38.
2. U.S. Preventive Services Task Force: **Screening for colorectal cancer: U.S. Preventive Services Task Force recommendation statement**. *Ann Intern Med* 2008, **149**:627–637.
3. Kavanos P, Schurer W: **Dynamics of colorectal cancer management in 17 countries**. *Eur J Health Econ* 2010, **10**(Suppl 1):S115–S129.
4. Whitlock EP, Lin JS, Liles S, Beil TL, Fu R: **Screening for colorectal cancer: a targeted, updated systematic review for the U.S. Preventive Services Task Force**. *Ann Intern Med* 2008, **149**:638–658.
5. Landsdorp-Vogelaar I, Knudsen AB, Brenner H: **Cost-effectiveness of colorectal cancer screening- an overview**. *Best Pract Res Clin Gastroenterol* 2010, **24**:439–449.
6. Wilson JMG, Junger G: *Principles and practice of screening for disease*. Geneva: WHO; 1968.
7. Neuman P: **Budget impact analyses get some respect**. *Value Health* 2007, **10**:324–325.
8. Mauskopf JA, Sullivan SD, Annemans L, Caro J, Muklins CD, Nuijten M, Orlewska E, Watkins J, Trueman P: **Principles of good practice of budget impact analysis: report of the ISPOR task force on good research practices – budget impact analysis**. *Value Health* 2007, **10**:336–347.
9. Sharp L, Tilson L, Whyte S, O'Ceilleachair A, Walsh C, Usher C, Tappenden P, Chilcott J, Staines A, Barry M, Comber H: **Cost-effectiveness of population-based screening for colorectal cancer: a comparison of guaiac-based faecal occult blood testing, faecal immunochemical testing and flexible sigmoidoscopy**. *Br J Cancer*. in press.
10. Sharp L, Walsh C, Whyte S, Tilson L, O'Ceilleachair A, Walsh C, Usher C, Tappenden P, Chilcott J, Staines A, Barry M, Comber H: **Letter to the editor**. *Br J Cancer*. 2013, **108**:1211–1212.
11. Atkin WS, Saunders BP: **Surveillance guidelines after removal of colorectal adenomatous polyps**. *Gut* 2002, **51**(Supplement 5):v6–v9.
12. Tappenden P, Chilcott J, Eggington S, Patrick J, Sakai M, Karnon J: **Option appraisal of population-based colorectal cancer screening programmes in England**. *Gut* 2007, **56**:677–684.
13. Allison JE, Feldman R, Tekawa IS: **Hemoccult screening in detecting colorectal neoplasm: sensitivity, specificity, and predictive value. Long-term follow-up in a large group practice setting**. *Ann Intern Med* 1990, **112**:328–33.

14. Castiglione G, Grazzini G, Poli A, Bonardi R, Ciatto S: **Hemoccult sensitivity estimate in a screening program for colorectal cancer in the Province of Florence.** *Tumori* 1991, **77**:243–5.
15. Foley DP, Dunne P, Dervan PJ, Callaghan TW, Crowe J, Lennon JR: **Left-sided colonoscopy and haemoccult screening for colorectal neoplasia.** *Eur J Gastroenterol* 1992, **4**:925–36.
16. Allison JE, Tekawa IS, Ransom LJ, Adrain AL: **A comparison of fecal occult-blood tests for colorectal-cancer screening.** *N Engl J Med* 1996, **334**:155–9.
17. Brevinge H, Lindholm E, Buntzen S, Kewenter J: **Screening for colorectal neoplasia with faecal occult blood testing compared with flexible sigmoidoscopy directly in a 55–56 years' old population.** *Int J Colorect Dis* 1997, **12**:291–5.
18. Lieberman DA, Weiss DG, Veterans Affairs Cooperative Study Group 380: **One-time screening for colorectal cancer with combined fecal occult-blood testing and examination of the distal colon.** *N Engl J Med* 2001, **345**:555–560.
19. Niv Y, Lev-El M, Fraser G, Abuksis G, Tamir A: **Protective effect of faecal occult blood test screening for colorectal cancer: worse prognosis for screening refusers.** *Gut* 2002, **50**:33–7.
20. Sung JJ, Chan FK, Leung WK, Wu JC, Lau JY, Ching J, To KF, Lee YT, Luk YW, Kung NN, Kwok SP, Li MK, Chung SC: **Screening for colorectal cancer in Chinese: comparison of fecal occult blood test, flexible sigmoidoscopy, and colonoscopy.** *Gastroenterol* 2003, **124**:608–14.
21. Collins JF, Lieberman DA, Durbin TE, Weiss DG, Veterans Affairs Cooperative Study #380 Group: **Accuracy of screening for fecal occult blood on a single stool sample obtained by digital rectal examination: a comparison with recommended sampling practice.** *Ann Intern Med* 2005, **142**:81–5.
22. Itoh M, Takahashi K, Nishida H, Sakagami K, Okubo T: **Estimation of the optimal cut off point in a new immunological faecal occult blood test in a corporate colorectal cancer screening programme.** *J Med Screen* 1996, **3**:66–71.
23. Chen K, Jiao DA, Zheng S, Zhou S, Yu H: **Diagnostic value of fecal occult blood testing for screening colorectal cancer.** *China Natl J New Gastroenterol* 1997, **3**:166–8.
24. Nakama H, Fattah AS, Zhang B, Kamijo N, Uehara Y: **Association of diverticulosis coli and vascular ectasias and the results of fecal occult blood test.** *Hepatogastroenterol* 2000, **47**:1277–1279.
25. Nakama H, Zhang B, Fattah AA, Kamijo N, Zhang X: **Characteristics of colorectal cancer that produce positive immunochemical occult blood test results on stool obtained by digital rectal examination.** *Can J Gastroenterol* 2001, **15**:227–30.
26. Cheng TI, Wong JM, Hong CF, Cheng SH, Cheng TJ, Shieh MJ, Lin YM, Tso CY, Huang AT: **Colorectal cancer screening in asymptomatic adults: comparison of colonoscopy, sigmoidoscopy and fecal occult blood tests.** *J Formos Med Assoc* 2002, **101**:685–90.
27. Gondal G, Grotmol T, Hofstad B, Bretthauer M, Eide TJ, Hoff G: **The Norwegian colorectal cancer prevention (NORCCAP) screening study: baseline findings and implementations for clinical work-up in age groups 50–64 years.** *Scand J Gastroenterol* 2003, **38**:635–42.
28. Liu HH, Huang TW, Chen HL, Wang TH, Lin JT: **Clinicopathologic significance of immunohistochemical fecal occult blood test in subjects receiving bidirectional endoscopy.** *Hepatogastroenterol* 2003, **50**:1390–1392.
29. Morikawa T, Kato J, Yamaji Y, Wada R, Mitsushima T, Shiratori Y: **A comparison of the immunochemical fecal occult blood test and total colonoscopy in asymptomatic population.** *Gastroenterol* 2005, **129**:422–8.
30. Nakazato M, Yamano H, Matsushita H, Sato K, Fujita K, Yamanaka Y, Imai Y: **Immunologic fecal occult blood test for colorectal cancer screening.** *Jap Med Assoc J* 2006, **49**:203–7.
31. Allison JE, Sakoda LC, Levin TR, Tucker JP, Tekawa IS, Cuff T, Pauly MP, Shlager L, Palitz AM, Zhao WK, Schwartz JS, Ransohoff DF, Selby JV: **Screening for colorectal neoplasms with new fecal occult blood tests: update on performance characteristics.** *J Natl Cancer Inst* 2007, **99**:1462–70.
32. Morikawa T, Kato J, Yamaji Y, Wada R, Mitsushima T, Sakaguchi K, Shiratori Y: **Sensitivity of immunochemical fecal occult blood test to small colorectal adenomas.** *Am J Gastroenterol* 2007, **102**:2259–64.
33. Rozen P, Ron E, Fireman Z, Hallak A, Grossman A, Baratz M, Rattan J, Gilat T: **The relative value of fecal occult blood tests and flexible sigmoidoscopy in screening for large bowel neoplasia.** *Cancer* 1987, **60**:2553–8.
34. Weller D, Moss S, Butler P, Campbell C, Coleman D, Melia J, Robertson R: *English pilot of bowel cancer screening: an evaluation of the second round, Final report to the Department of Health.* Edinburgh: University of Edinburgh; 2006.
35. Information Services Division: *Summary of the key performance indicators (KPIs) used to monitor and evaluate the Scottish bowel screening pilot.* Edinburgh: NHS National Services Scotland; 2008.
36. Malila N, Oivanen T, Hakama M: **Implementation of colorectal cancer screening in Finland: experiences from the first three years of a public health programme.** *Z Gastroenterol* 2007, **45**:1–4.
37. Segnan N, Senore C, Andreoni B, Azzoni A, Bisanti L, Cardelli A, Castiglione G, Crosta C, Ederle A, Fantin A, Ferrari A, Fracchia M, Ferrero F, Gasperoni S, Recchia S, Risio M, Rubeca T, Saracco G, Zappa M, SCORE3 Working Group-Italy: **Comparing attendance and detection rate of colonoscopy with sigmoidoscopy and FIT for colorectal cancer screening.** *Gastroenterol* 2007, **132**:2304–2012.
38. UK Flexible Sigmoidoscopy Screening Trial Investigators: **Single flexible sigmoidoscopy screening to prevent colorectal cancer: baseline findings of a UK multicentre randomised trial.** *Lancet* 2002, **359**:1291–300.
39. Gray M, Pennington CR: **Screening sigmoidoscopy: a randomised trial of invitation style.** *Health Bull (Edinb)* 2000, **58**:137–140.
40. Bretthauer M, Gondal G, Larsen IK, Carlsen E, Eide TJ, Grotmol T, Skovlund E, Tveit KM, Vatn MH, Hoff G: **Design, organization and management of a controlled population screening study for detection of colorectal neoplasia: attendance rates in the NORCCAP study (Norwegian Colorectal Cancer Prevention).** *Scand J Gastroenterol* 2002, **37**:568–573.
41. van Rijn JC, Reitsma JB, Stoker J, Bossuyt PM, van Deventer SJ, Dekker E: **Polyp miss rate determined by tandem colonoscopy: a systematic review.** *Am J Gastroenterol* 2006, **101**:343–50.
42. Bressler B, Paszat LF, Chen Z, Rothwell DM, Vinden C, Rabeneck L: **Rates of new or missed colorectal cancers after colonoscopy and their risk factors: a population-based analysis.** *Gastroenterol* 2007, **132**:96–102.
43. Cotton PB, Durkalski VL, Pineau BC, Palesch YY, Mauldin PD, Hoffman B, Vining DJ, Small WC, Affronti J, Rex D, Kopecky KK, Ackerman S, Burdick JS, Brewington C, Turner MA, Zfass A, Wright AR, Iyer RB, Lynch P, Sivak MV, Butler H: **Computed tomographic colonography (virtual colonoscopy): a multicenter comparison with standard colonoscopy for detection of colorectal neoplasia.** *J Am Med Assoc* 2004, **291**:1713–9.
44. Halligan S, Altman DG, Taylor SA, Mallett S, Deeks JJ, Bartram CI, Atkin W: **CT colonography in the detection of colorectal polyps and cancer: systematic review, meta-analysis, and proposed minimum data set for study level reporting.** *Radiology* 2005, **237**:893–904.
45. Mulhall BP, Veerappan GR, Jackson JL: **Meta-analysis: computed tomographic colonography.** *Ann Intern Med* 2005, **142**:635–50.
46. Nino CD, Chen MH, Toledano AY, Heiken JP, Dachman A, Kuo MD, Menias CO, Siewert B, Cheema JJ, Obregon RG, Fidler JL, Zimmerman P, Horton KM, Coakley K, Iyer RB, Hara AK, Halvorsen RA Jr, Casola G, Yee J, Herman BA, Burgart LJ, Limburg PJ: **Accuracy of CT colonography for detection of large adenomas and cancers.** *N Engl J Med* 2008, **359**:1207–1217.
47. Winawer SJ, Flehinger BJ, Schottenfeld D, Miller DG: **Screening for colorectal cancer with fecal occult blood testing and sigmoidoscopy.** *J Natl Cancer Inst* 1993, **85**:1311–8.
48. Gatto NM, Frucht H, Sundararajan V, Jacobson JS, Grann VR, Neugut AI: **Risk of perforation after colonoscopy and sigmoidoscopy: a population-based study.** *J Natl Cancer Inst* 2003, **95**:230–6.
49. Dafnis G, Ekbohm A, Pahlman L, Blomqvist P: **Complications of diagnostic and therapeutic colonoscopy within a defined population in Sweden.** *Gastrointest Endosc* 2001, **54**:302–9.
50. Whyte S, Walsh C, Chilcott J: **Bayesian calibration of a natural history model with application to a population model for colorectal cancer.** *Med Decis Making* [Epub ahead of print].
51. National Cancer Screening Service: *Update on Progress on the Implementation of the National Colorectal Cancer Screening Programme.* Dublin: NCSS; 2011. Available at <http://www.cancerscreening.ie/bowel-screening.html> [Accessed 21st March 2013].
52. O'Leary BA, Olynyk JK, Neville AM, Platell CF: **Cost-effectiveness of colorectal cancer screening: comparison of community-based flexible sigmoidoscopy with fecal occult blood testing and colonoscopy.** *J Gastroenterol Hepatol* 2004, **19**:38–47.
53. Ho C, Heitman S, Membe SK, Morrison A, Moulton K, Manns B, Au F, Reed M, Hilsden R: *Computed tomographic colonography for colorectal cancer screening in*

- an average risk population: systematic review and economic evaluation. Ottawa: Canadian Agency for Drugs and Technologies in Health; 2008.
54. Health Information and Quality Authority: *Report of the evaluation of use of resources in the national population-based cancer screening programmes and associated services*. Dublin: HIQA; 2009. Available at URL http://www.hiqa.ie/system/files/Colorectal_cancer_evaluation_report_HTA.pdf [Accessed 21st March 2013].
 55. Gray NM, Sharp L, Cotton SC, Masson LF, Little J, Walker LG, Avis M, Philips Z, Russell I, Whynes D, Cruickshank M, Woolley CM, TOMBOLA group: **Psychological effects of a low-grade abnormal cervical smear test result: anxiety and associated factors**. *Br J Cancer* 2006, **94**:1253–62.
 56. Guittet L, Bouvier V, Mariotte N, Vallee JP, Arsène D, Boutreux S, Tichet J, Launoy G: **Comparison of screening for colorectal cancer in a general average immunochemical faecal occult blood test in a general average risk population**. *Gut* 2007, **56**:210–4.
 57. Van Rossum LG, van Rijn AF, Verbeek AL, van Oijen MG, Laheij RJ, Fockens P, Jansen JB, Adang EM, Dekker E: **Colorectal cancer screening comparing no screening, immunochemical and guaiac fecal occult blood tests: a cost-effectiveness analysis**. *Int J Cancer* 2011, **128**:1908–17.
 58. Hol L, van Leerdam ME, van Ballegooijen M, van Vuuren AJ, van Dekken H, Reijerink JC, van der Togt AC, Habbema JD, Kuipers EJ: **Screening for colorectal cancer: randomized trial comparing guaiac-based and immunochemical faecal occult blood testing and flexible sigmoidoscopy**. *Gut* 2010, **59**:62–68.
 59. Levi Z, Birkenfeld S, Vilkin A, Bar-Chana M, Lifshitz I, Chared M, Maoz E, Niv Y: **A higher detection rate for colorectal cancer and advanced adenomatous polyp for screening with immunochemical fecal occult blood test than guaiac fecal occult blood test, despite lower compliance rate. A prospective, controlled, feasibility study**. *Int J Cancer* 2011, **128**:2415–24.
 60. Wilschut JA, Habbema DF, van Leerdam ME, Hol L, Lansdorp-Vogelaar I, Kuipers EJ, Ballegooijen M: **Fecal occult blood testing when colonoscopy capacity is limited**. *J Natl Cancer Inst* 2011, **103**:1741–51.
 61. Ransohoff DF: **Screening colonoscopy in balance. Issues of implementation**. *Gastroenterol Clin North Am* 2002, **31**:1031–44i.
 62. National Cancer Screening Service: *First report of the National Cancer Screening Service Expert Advisory Group on colorectal cancer screening*. Dublin: National Cancer Screening Service; 2007.
 63. Zorzi M, Baracco S, Fedato C, Grazzini G, Naldoni C, Sassoli De Biachi P, Senore C, Visioli CB, Cogo C: **Screening for colorectal cancer in Italy: 2008 survey**. *Epidemiol Prev* 2010, **34**(5–6 Suppl 4):53–72.
 64. Peris M, Espinas JA, Munoz L, Navarro M, Binefa G, Borrás JM, Catalan Colorectal Cancer Screening Pilot Programme Group: **Lesson learnt from a population-based pilot programme for colorectal cancer screening in Catalonia (Spain)**. *J Med Screen* 2007, **14**:81–86.
 65. Kis RK: **National colorectal cancer screening program in the Republic of Croatia – experiences, outcomes and obstacles in the program implementation in the Medimurje County**. *Acta Med Croatica* 2010, **64**:363–74.
 66. McNemara D, Qasim A, Lee N, Condon C, O'Morain C: **Round one of the adelaide and meath hospital/trinity college colorectal cancer screening programme: programme report and analysis based on established international key performance indices**. *Ir J Med Sci* 2011, **180**:549–52.
 67. Brotherstone H, Vance M, Edwards R, Miles A, Robb KA, Evans RE, Wardle J, Atkin W: **Uptake of population-based flexible sigmoidoscopy screening for colorectal cancer: a nurse-led feasibility study**. *J Med Screen* 2007, **14**(2):76–80.
 68. Vart G, Banzi R, Minozzi S: **Comparing participation rates between immunochemical and guaiac faecal occult blood tests: a systematic review and meta-analysis**. *Prev Med* 2012, **55**:87–92.
 69. Hewitson P, Glasziou P, Irwig L, Towler B, Watson E: **Screening for colorectal cancer using the faecal occult blood test**. *Hemoccult Cochrane Database Syst Rev* 2007, **24**:CD001216.
 70. Soares-Weiser K, Burch J, Duffy S, St John J, Smith S, Westwood M, Kleijnen J: **Diagnostic accuracy and cost-effectiveness of faecal occult blood tests used in screening for colorectal cancer: a systematic review**. York: Centre for Reviews and Dissemination; 2007.
 71. Steele RJ, McClements P, Watling C, Libby G, Weller D, Brewster D, Black R, Carey FA, Fraser CG: **Interval cancers in a FOBT-based colorectal cancer population screening programme: implications for stage, gender and tumour site**. *Gut* 2012, **61**:576–61.
 72. Fraser CG, Mathew CM, Mowat NA, Wilson JA, Carey FA, Steele RJ: **Immunochemical testing of individuals positive for guaiac faecal occult blood test in a screening programme for colorectal cancer: an observational study**. *Lancet Oncol* 2006, **7**:127–31.
 73. Fraser CG, Mathew CM, Mowat NA, Wilson JA, Carey FA, Steele RJ: **Evaluation of a card collection-based faecal immunochemical test in screening for colorectal cancer using a two-tier reflex approach**. *Gut* 2007, **56**:1415–8.

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