

University of Dundee

Faecal haemoglobin concentration is related to detection of advanced colorectal neoplasia in the next screening round

Digby, Jayne; Fraser, Callum G.; Carey, Francis A.; Diament, Robert H.; Balsitis, Margaret; Steele, Robert J. C.

Published in:
Journal of Medical Screening

DOI:
[10.1177/0969141316653983](https://doi.org/10.1177/0969141316653983)

Publication date:
2017

Document Version
Peer reviewed version

[Link to publication in Discovery Research Portal](#)

Citation for published version (APA):

Digby, J., Fraser, C. G., Carey, F. A., Diament, R. H., Balsitis, M., & Steele, R. J. C. (2017). Faecal haemoglobin concentration is related to detection of advanced colorectal neoplasia in the next screening round. *Journal of Medical Screening*, 24(2), 62-68. <https://doi.org/10.1177/0969141316653983>

General rights

Copyright and moral rights for the publications made accessible in Discovery Research Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from Discovery Research Portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain.
- You may freely distribute the URL identifying the publication in the public portal.

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Faecal haemoglobin concentration is related to detection of advanced colorectal neoplasia in the next screening round.

Jayne Digby,^{1,2} Callum G Fraser,^{1,2} Francis A Carey,³ Robert H Diamant,⁴
Margaret Balsitis,⁵ Robert JC Steele.^{1,2,6}

¹Scottish Bowel Screening Research Unit, Ninewells Hospital and Medical School, Dundee, Scotland

²Centre for Research into Cancer Prevention and Screening, Ninewells Hospital and Medical School, Dundee, Scotland

³Department of Pathology, Ninewells Hospital and Medical School, Dundee, Scotland

⁴Department of Surgery, Crosshouse Hospital, Kilmarnock, Ayrshire, Scotland

⁵Department of Pathology, Crosshouse Hospital, Kilmarnock, Ayrshire, Scotland

⁶Medical Research Institute, Division of Cancer, Ninewells Hospital and Medical School, Scotland

Correspondence to: Jayne Digby, Scottish Bowel Screening Research Unit, Mailbox 7, Level 7, Ninewells Hospital and Medical School, Dundee DD1 9SY,

Scotland

E-mail: jaynedigby@nhs.net

Tel: 01382383982

Short Title: Faecal haemoglobin and advanced neoplasia detection.

Keywords: advanced neoplasia, colorectal cancer, faecal haemoglobin, faecal immunochemical test, faecal occult blood test, risk, screening

Number of words (abstract): 250

Number of words (text): 2975

Number of Tables: 5

Number of Figures: 1

Number of references: 24

Abstract

Introduction Relationships between faecal haemoglobin concentrations (f-Hb) below the cut-off used in colorectal cancer (CRC) screening and longer-term outcomes are not well established.

Objectives To examine associations between f-Hb reported as negative and outcomes in the next screening round.

Setting Scottish Bowel Screening Programme.

Methods f-Hb and diagnostic outcomes were investigated for participants with a negative result, f-Hb cut-off < 80.0 µg Hb/g faeces, but then positive within the next two years.

Results Of 37,780 with a negative result, at the next screening round 556 (1.5%) had positive and 30,293 (80.2%) negative results. Initial median f-Hb (2.1 µg Hb/g faeces, IQR: 0.0 - 13.2) was higher in those with a subsequently positive result than those who had a negative result at the next round (0.0 µg Hb/g faeces, IQR: 0.0 - 1.4; $p < 0.0001$). Using f-Hb 0.0 - 19.9 µg Hb/g faeces as reference, logistic regression analysis showed high adjusted OR for advanced neoplasia (AN: CRC or higher-risk adenoma) detection at the next round of 14.3 (95% CI: 8.9 - 23.1) in those with initial f-Hb 20.0 - 39.9 µg Hb/g faeces and 38.0 (95% CI: 20.2 - 71.2) with 60.0 - 79.9 µg Hb/g faeces.

Conclusions A higher proportion of participants with f-Hb of ≥ 20 µg Hb/g faeces had AN detected at the next round, compared to those with lower f-Hb. Although most relevant when using high f-Hb cut-offs, studies of f-Hb and

outcomes over screening rounds may provide strategies to direct available colonoscopy towards those at highest risk.

Introduction

Participants in colorectal cancer (CRC) screening programmes with faecal haemoglobin concentrations (f-Hb) that are below the selected cut-off for positivity and referral for follow-up colonoscopy, as determined by a quantitative faecal immunochemical test (FIT) for haemoglobin (Hb), are more likely to be diagnosed with an interval cancer (IC) than those with undetectable f-Hb.¹ Rarely documented, however, has been the relationship between a detectable f-Hb below the programme selected cut-off and diagnostic outcome in the longer term. Such knowledge could assist in determining the role of f-Hb as a predictor of the future risk of advanced neoplasia (AN), defined as CRC plus higher-risk adenoma (HRA).

Only two relevant studies have been performed previously. Firstly, in a longitudinal follow-up of CRC screening participants in Taiwan,² a cohort of 44,324 participants, aged 40 - 69 years, with f-Hb below the cut-off applied of ≥ 20.0 μg Hb/g faeces were followed for a median of 4.39 years. The incidence of AN rose from 1.75/1000 person-years for those with f-Hb of 0.2 – 3.9 μg Hb/g faeces to 7.08/1000 in those with f-Hb of 16.0 – 19.9 μg Hb/g faeces. Moreover, relative to those with f-Hb of 0.2 – 3.9 μg Hb/g faeces, adjusted hazard ratios (HR) for AN were calculated as 3.41 for those with f-Hb in the category lower than but closest to the cut-off f-Hb used (16.0 – 19.9 μg Hb/g faeces). Secondly, a recent study documents results of regression modelling to further establish the value of f-Hb as a predictor for colorectal neoplasia:³ 54,921 participants invited between 2001 and 2007 were followed up in the annual FIT-based CRC screening programme in Taiwan to identify

those diagnosed with IC, screen-detected CRC and adenoma. A trend of increasing HR with increasing baseline f-Hb was demonstrated for colorectal neoplasia, including for those with f-Hb above the traditionally used cut-off of $\geq 20 \mu\text{g Hb/g faeces}$.

Thus, the available limited evidence indicates that the higher the baseline f-Hb, the greater the likelihood of a future finding of AN. If verified, this could have implications for the design and execution of CRC screening programmes; for example, those with a negative screening test result who are at high risk of having AN by virtue of a detectable f-Hb despite being lower than the cut-off f-Hb applied, could be prioritised for future repeat screening at a shorter interval than the current usual annual or biennial invitation.

Conversely, those with undetectable f-Hb, and therefore deemed very unlikely to have a future finding of AN, could perhaps require less frequent invitations to screening. Such a strategy could be an important consideration in countries such as Scotland where, in view of demands on the colonoscopy resource, adoption of a cut-off f-Hb of $\geq 80.0 \mu\text{g Hb/g faeces}$ (equivalent to $\geq 400 \text{ ng Hb/ml buffer}$) for the planned FIT-based programme is likely, given that this was used in the pilot evaluation required to inform future adoption of FIT as a first-line test.⁴ This f-Hb cut-off was selected in order to achieve approximately 2% positivity, to mimic the current screening algorithm based on a two-tier reflex guaiac faecal occult blood test (gFOBT)/qualitative FIT strategy.⁴ Since it is known that IC are common at this f-Hb cut-off,¹ it is reasonable to postulate is that a proportion of participants with f-Hb $< 80.0 \mu\text{g Hb/g faeces}$ will have AN detected at the next screening round. Our aim therefore, was to investigate the relationship between f-Hb reported as a negative screening

test result in our FIT evaluation and diagnostic outcomes identified over the two year period of the next screening round to test the hypothesis that those with f-Hb lower than but closer to the cut-off applied are more likely to be subsequently diagnosed with AN.

Materials and Methods

Outside the setting of the FIT evaluation, the Scottish Bowel Screening Programme (SBoSP) uses a two-tier reflex gFOBT/FIT screening algorithm, which has been described in detail previously.⁴ The pilot evaluation of using FIT as a first-line screening test in Scotland has also been described in detail previously:⁵ all participants with f-Hb ≥ 80.0 $\mu\text{g Hb/g}$ faeces were reported as positive and offered colonoscopy.

Following completion of the FIT evaluation, the SBoSP returned the two NHS Boards to the two-tier reflex gFOBT/FIT screening algorithm. The screening test results for all those eligible to take part in the two years of the next screening round after the FIT evaluation (13 January 2011 to 12 January 2013) and resident in either NHS Tayside or NHS Ayrshire & Arran were examined. The f-Hb from the previous FIT-based pilot evaluation round for those with a positive gFOBT/FIT screening test result was retrieved from the Scottish Bowel Screening IT System (BoSS). Data for colonoscopy outcomes and any subsequent pathology for those with a positive test result were downloaded from the appropriate clinical IT systems in NHS Tayside and NHS Ayrshire & Arran. Data on colonoscopy findings, including number, size,

Dukes' stage, and localisation of CRC and adenomas were collected. As in the SBoSP, HRA were defined as those greater than 10 mm in diameter or when there were three or more adenoma present.⁶ AN was defined as CRC and HRA.

MedCalc (MedCalc Software, Mariakerke, Belgium) statistical software was used for all calculations. The Mann-Whitney U test was used for comparison of median f-Hb between classes. Probability of $p < 0.05$ was considered significant. Logistic regression analysis was performed to calculate odds ratios (OR) for diagnosis of AN at the next screening round amongst those in different f-Hb classes and different demographic groups, both adjusted and unadjusted for age and gender as known confounding variables.

Results

The majority of participants (37,780) in the FIT pilot evaluation had an initial negative test result.⁵ Of those, 92.7% were eligible for an invitation to further screening with the gFOBT/FIT algorithm: the majority of those not invited were above the age limit applied in the SBoSP (75 years), with others having died or being no longer resident in Scotland. In total, 30,849 participants participated in the FIT evaluation and the subsequent screening round with gFOBT/FIT. Table 1 details the outcomes of all participants who had f-Hb < 80 $\mu\text{g Hb/g faeces}$, the cut-off used in the FIT evaluation, by age and gender.

Table 1. Number of participants with a negative FIT screening test result and their screening test results in two years in the next gFOBT/FIT screening round.

	All		Men		Women	
	n	%	n	%	n	%
Total with negative FIT screening test result < 80.0 µg Hb/g faeces	37,780		17,525		20,255	
Test result in next gFOBT/FIT round:						
Positive	556	1.5	339	1.9	217	1.1
Negative	30,293	80.2	13,910	79.4	16,383	80.9
Non-responder	4,165	11.0	1,976	11.3	2,189	10.8
Excluded	2,766	7.3	1,300	7.4	1,466	7.2

556 (1.5%) participants had a positive test result in the subsequent screening round and 30,293 (80.2%) had a negative test result. The initial median f-Hb was statistically significantly higher in those who then had a positive test result in the next round than those who remained negative (2.1 µg Hb/g faeces, IQR 0.0 - 13.2 v 0.0 µg Hb/g faeces, IQR 0.0 - 1.4; $p < 0.0001$).

Table 2 shows the diagnostic outcomes, with median f-Hb, of participants who had a positive test result in the next screening round. The differences in median f-Hb between men and women were only statistically significant in those diagnosed at colonoscopy with either non-neoplastic pathology or normal findings (both $p < 0.05$).

Table 2. Diagnostic outcomes and median faecal haemoglobin concentrations (f-Hb) at the initial FIT screening round in participants then having a positive screening test result at the next gFOBT/FIT screening round.

	All				Men				Women			
	n	%	initial median f-Hb (µg Hb/g faeces)	IQR	n	%	initial median f-Hb (µg Hb/g faeces)	IQR	n	%	initial median f-Hb (µg Hb/g faeces)	IQR
Total positive	556		2.1	0.0 - 13.0	339		1.6	1.2 - 3.0	217		2.4	0 - 12.9
Colorectal cancer (CRC)	26	4.7	16.7	1.2 - 31.6	19	5.6	21.0	0.5 - 36.3	7	3.2	10.2	2.5 - 24.0
Higher risk adenoma (HRA)	85	15.3	13.6	1.2 - 38.5	63	18.6	9.6	0.9 - 38.8	22	10.1	16.6	1.8 - 35.2
Advanced neoplasia (AN: CRC and HRA)	111	20.0	13.6	1.2 - 37.6	82	24.2	14.7	0.8 - 38.8	29	13.4	13.6	1.8 - 35.2
Low risk adenoma	65	11.7	1.7	0.0 - 8.5	42	12.4	1.6	0.0 - 10.9	23	10.6	1.8	0.0 - 7.0
Non-neoplastic pathology	131	23.6	1.6	0.0 - 7.4	72	21.2	0.7	0.0 - 5.8	59	27.2	3.4	0.1 -15.1
Normal findings	169	30.4	1.4	0.0 - 6.2	93	27.4	0.3	0.0 - 5.6	76	35.0	2.0	0.0 - 7.3

Footnote: IQR is the interquartile range (difference between the upper quartile and the lower quartile); non-neoplastic pathology comprises conditions including diverticular disease, haemorrhoids and inflammatory bowel disease and haemorrhoids.

A numerical f-Hb result was available for 30,823 of the 30,849 participants in both screening rounds. The majority of these (96.6%) had f-Hb in the FIT round that was in the lowest class of 0.0 - 19.9 µg Hb/g faeces. 87.4% of participants with a positive gFOBt/FIT result who did not have AN detected were in this low f-Hb class, compared to only 56.8% of those with AN. The proportions of participants with different diagnostic outcomes according to which class of f-Hb they fell into at the initial screening round with FIT are shown in Table 3. A lower proportion of participants with previously low f-Hb had AN detected than less severe outcomes, but this trend reversed with rising f-Hb: this is demonstrated in Figure 1. Since a commonly used f-Hb cut-off for screening programmes outside Scotland is ≥ 20 µg Hb/g faeces, Table 4 shows further analysis of the class with a f-Hb below this concentration, dissected into two f-Hb classes; those with undetectable f-Hb at the initial FIT screening round and those with f-Hb 1.0 - 19.9 µg Hb/g faeces.

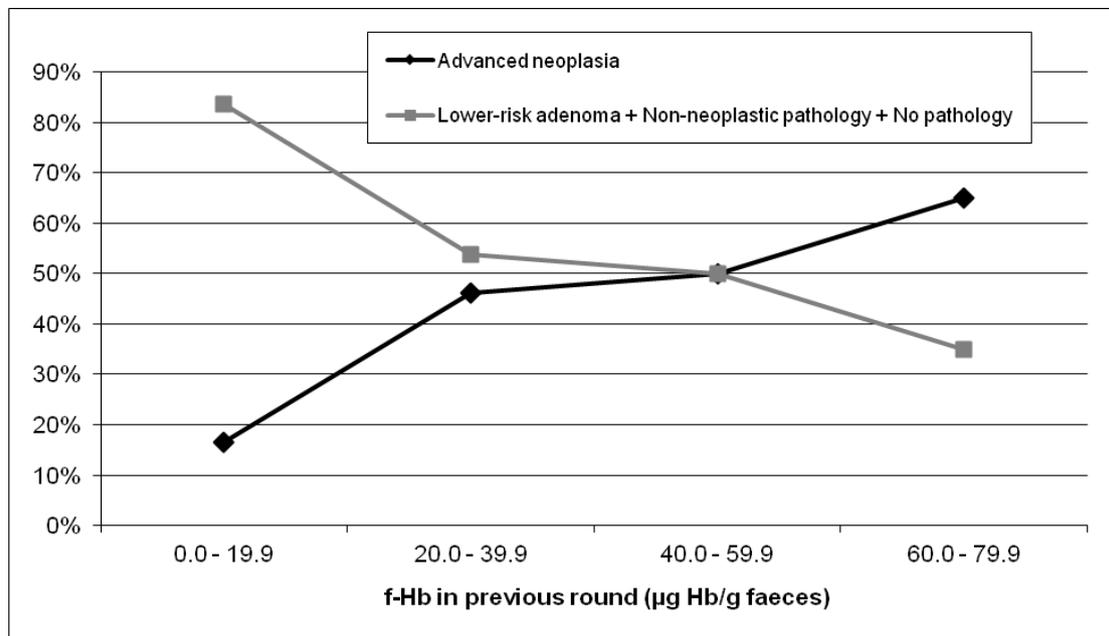
Table 3. Outcomes of participants in both screening rounds according to faecal haemoglobin concentration class at the initial FIT screening round.

Outcome in next FOBt/FIT screening round	Faecal haemoglobin concentration class in the initial FIT screening round (µg Hb/g faeces)							
	0.0 - 19.9		20.0 - 39.9		40.0 - 59.9		60.0 - 79.9	
	n	%	n	%	n	%	n	%
Negative screening test result	29,049	96.6%	661	2.2%	227	0.8%	125	0.4%
Positive screening test result	449	80.8%	59	10.6%	26	4.7%	22	4.0%
Follow-up complete	383		52		22		20	

Colorectal cancer (CRC)	14	3.7%	7	13.5%	1	4.5%	4	20.0%
Higher-risk adenoma (HRA)	49	12.8%	17	32.7%	10	45.5%	9	45.0%
Advanced neoplasia (AN: CRC and HRA)	63	16.4%	24	46.2%	11	50.0%	13	65.0%
Low-risk adenoma	58	15.1%	8	15.4%	2	9.1%	0	0.0%
Non-neoplastic pathology	115	30.0%	6	11.5%	7	31.8%	3	15.0%
No pathology detected	147	38.4%	14	26.9%	2	9.1%	4	20.0%

Footnote: Non-neoplastic pathology comprises conditions including diverticular disease, haemorrhoids and inflammatory bowel disease and haemorrhoids.

Figure 1. Percentages of FIT positive screening participants completing follow-up who had advanced neoplasia (AN) or less severe outcomes in the next gFOBT/FIT screening round, according to faecal haemoglobin concentration (f-Hb) class.



Footnote: Non-neoplastic pathology comprises conditions including diverticular disease, haemorrhoids and inflammatory bowel disease and haemorrhoids.

Table 4. Outcomes of participants of both screening rounds with faecal haemoglobin concentration < 20 µg Hb/g faeces at the initial FIT screening round, dissected into two classes.

Outcome in subsequent screening round	Faecal haemoglobin concentration class in the initial FIT screening round (µg Hb/g faeces)			
	Undetectable		1.0 - 19.9	
	n	%	n	%
Negative screening test result	16,621	56.8	12,633	43.2
Positive screening test result	181	40.3	268	59.7
Follow-up complete	151		232	
Colorectal cancer (CRC)	5	3.3	9	3.8
Higher-risk adenoma (HRA)	14	9.3	35	15.1
Advanced neoplasia (AN: CRC and HRA)	19	12.6	44	19.0
Lower-risk adenoma (LRA)	24	15.9	34	14.7
Non-neoplastic pathology	43	28.5	72	31.0
No pathology detected	65	43.0	82	35.3

Footnote: Non-neoplastic pathology comprises conditions including diverticular disease, haemorrhoids and inflammatory bowel disease and haemorrhoids.

As shown in Table 5, logistic regression analysis showed very high age and gender adjusted OR for AN even in those with f-Hb 20.0 - 39.9 µg Hb/g faeces, using those with f-Hb 0.0 - 19.9 µg Hb/g faeces as the reference group (adjusted OR = 14.3, 95% CI 8.9 - 23.1). Almost one-tenth of all participants who had f-Hb within the highest f-Hb class examined, 60.0 - 79.9 µg Hb/g faeces, and participated in the subsequent round, had AN detected at

the follow-up investigations of a positive test result. AN was over 40-times more prevalent in this class than those with f-Hb previously 0.0 - 19.9 µg/g faeces, of whom 0.21% had AN. The adjusted odds ratio for AN in the highest f-Hb class was calculated to be 38.0 (95% CI 20.2 - 71.2).

Table 5. Odds ratios (both unadjusted and adjusted for age and gender) for advanced neoplasia (AN) found at the next gFOBT/FIT round after a positive screening test result, according to faecal haemoglobin concentration (f-Hb) class in the previous FIT screening round.

f-Hb class (µg Hb/g faeces)	% with AN detected	Odds ratio	
		Unadjusted (95% CI)	Adjusted (95% CI)
0.0 - 19.9	0.2%	1.0	1.0
20.0 - 39.9	3.4%	16.2 (10.1 - 26.2)	14.3 (8.9 - 23.1)
40.0 - 59.9	4.4%	21.5 (11.2 - 41.4)	17.7 (9.2 - 34.2)
60.0 - 79.9	9.0%	45.9 (24.7 - 85.4)	38.0 (20.2 - 71.2)

Footnote: Advanced neoplasia (AN) includes colorectal cancer and higher-risk adenoma.

Discussion

The finding that a higher proportion of participants with a previous f-Hb lower than, but approaching, the cut-off used in our FIT pilot evaluation had AN detected at the next round, compared to those with low f-Hb, provides further evidence that f-Hb is a strong predictor of future risk.

Positivity in those participants who, in the previous round, had f-Hb below the cut-off applied of $\geq 80.0 \mu\text{g Hb/g faeces}$ was 1.5%. This was lower than the overall positivity seen in the FIT pilot evaluation done in Scotland⁷ and also with the current gFOBT/FIT algorithm at ca. 2%.⁴ This suggests that those who have previously had a negative screening test result, even with a relatively high cut-off f-Hb, are less likely to have a positive test result in the subsequent screening round compared with the overall population invited for screening, which would include first-time participants as well as participants who have previously had a positive test result.

Our findings further support the concept of f-Hb being a valuable predictor of risk of AN since f-Hb was statistically significantly elevated, not only in those who would go on to have a positive test result in the next round compared to those who again had a negative result, but also in those who had AN detected at follow-up investigations compared with those with no pathology (both $p < 0.0001$). Furthermore, one in five participants who had f-Hb in the f-Hb range nearest to the cut-off f-Hb applied and were then positive in the next screening round, were diagnosed with CRC. This compares with one in 27 participants in the lowest f-Hb class examined: thus, those with elevated f-Hb

who go on to have a positive screening test result in the next round are over five times more likely to have CRC detected. When also taking into account detection of HRA, 65% of those participants with a positive screening test result who previously had f-Hb in the highest f-Hb class examined had any AN diagnosed, compared to just 16.4% in the lowest. Moreover, a trend exists of a higher proportion of AN found in participants from each of the four classes of increasing f-Hb. From these data, the odds of having AN detected following a positive screening result are even in those with previous f-Hb between 40.0 and 59.9 $\mu\text{g/g}$ faeces. After this point, it is more likely than not that AN will be present at follow-up investigations in those with a positive screening test result in the next round. Almost one-tenth of all participants with f-Hb approaching the cut-off f-Hb applied went on to have AN detected, representing a 40-fold increase in the risk as compared to those in the lowest f-Hb class. This finding is important when considering f-Hb as a predictor of future risk of AN at the time of a negative screening test result. Not only is strong evidence provided for the value of f-Hb as a risk factor for AN being detected at follow-up investigations of participants going on to have a positive test result, but also for predicting future positivity and the subsequent detection of AN. In addition, since it has been suggested that f-Hb is a predictor, not only of CRC mortality, but also all-cause death,⁸ there may be other benefits in the more general application of f-Hb as a predictor of ill health.

If screening programmes collect data such as this, irrespective of the faecal test used, then the effect of cut-off f-Hb on yield of AN can be determined using a graph such as shown in our Figure 1. The f-Hb where the percentage

of participants with positive screening test results completing follow-up who had advanced neoplasia (AN) crosses that of those less severe outcomes can give an indication of the cut-off f-Hb associated with the greatest benefit in terms of yield of advanced neoplasia (AN), although this only includes participants with a positive result at the next screening round and not those who had a negative test result at the subsequent round. CRC screening programmes using FIT have adopted a wide range of f-Hb cut-offs, and the Scottish Bowel Screening Programme FIT evaluation used a much higher f-Hb cut-off than that used in most other countries. Our data allows some comparison of other strategies using lower f-Hb cut-offs. Combining the data presented here with that in our recent publication on IC arising in the cohort participating in the pilot evaluation of screening using FIT,¹ if Scotland used a $\geq 60.0 \mu\text{g Hb/g faeces}$ cut-off rather than the $\geq 80.0 \mu\text{g Hb/g faeces}$ used in the FIT evaluation,⁴ then 25.6% additional colonoscopy would yield 20.0% more CRC, or at least possible pre-cursor lesions, presuming these were present at the time of the FIT-based screening. However, in addition to the reduction in the number of false negative screening test results, the number of false positive results will also increase, requiring assessment of the associated harms such as over-diagnosis of lesions that would never have become symptomatic and the risk of complications of colonoscopy.

Further analysis was performed on clinical outcomes at the subsequent screening round of those with an initial f-Hb below the commonly used cut-off f-Hb $\geq 20 \mu\text{g Hb/g faeces}$. With the prevalence of AN higher in those with f-Hb between 1.0 and 19.9 $\mu\text{g Hb/g faeces}$ than in those with undetectable f-

Hb, evidence is provided that f-Hb is a predictor of future risk of colorectal neoplasia, even at a lower f-Hb cut-off than that used in Scotland.

It is well accepted that men have higher median f-Hb than women.^{9,10}

Therefore, it is surprising that the median initial f-Hb of participants with a positive screening test result at the next round was higher in women than in men, although this did not reach statistical significance. Significantly higher median initial f-Hb was detected in women compared with men in those with non-neoplastic pathology or normal findings at colonoscopy triggered by a positive screening test result at the next screening round. It appears from this analysis that men who had a false positive screening test result were more likely to have had a low f-Hb in the previous screening round before experiencing short term colonic blood loss to trigger their subsequent positive screening test result. Women, however, may have had longer term colonic bleeding, rising over time. The reasons for this are unclear and may be worthy of further study. Although not statistically significant, median f-Hb of participants with CRC detected at the next screening round was higher in men than in women, but the opposite was apparent for HRA. In fact, previous median f-Hb for women with CRC was very similar to previous median f-Hb of men with HRA. This further demonstrates the variation in f-Hb by gender as documented previously,^{9,10} which should be taken into account in setting cut-off f-Hb to be used in any screening programme.

An obvious weakness of this analysis is that different tests were employed in the two rounds screening examined. However, the cut-off employed in the quantitative FIT pilot evaluation was equivalent in terms of positivity to the

current gFOBT/FIT algorithm to which the programme reverted. Of course, it would be of interest to conduct a similar study in which quantitative FIT was used in consecutive screening rounds to allow investigation of the variation in f-Hb over time according to diagnostic outcomes. In addition, the number of CRC cases is low, limiting detailed analysis by sub-groups such as age, gender and lesion site.

Two very recent studies comparing clinical outcomes between two consecutive screening rounds have emerged with very large cohorts allowing sub-analyses by gender and lesion site. Recent work from Spain showed that the second screening round was associated with lower median f-Hb and an increased proportion of proximal CRC.¹¹ The authors suggested various strategies to improve diagnosis of proximal CRC, such as using a lower cut-off f-Hb at the first compared with the second round. Another study from France has demonstrated that subsequent screening was associated with increased detection of proximal AN, particularly in women, and CRC was more often later stage.¹² Here, it was concluded that, since the late stage CRC were likely to be previously missed lesions, the findings stressed the importance of repeated screening after a previous negative test result. These studies demonstrate the scope that exists for further work into the relationship between f-Hb and outcomes in subsequent screening rounds according to various sub-groups, if working with a large enough sample size.

Our data could impact on CRC screening strategies. Risk scoring models are becoming increasingly developed, with an escalation in published studies in recent years.¹³⁻²² We have confirmed that f-Hb does predict the presence of

colorectal disease through analysis of positivity in the next screening round. In consequence, screening programmes using FIT and looking to apply risk-scoring should certainly incorporate f-Hb into such models, along with age and gender, and perhaps deprivation,²³ to improve their predictive power. The fact that f-Hb is related, not only to colorectal disease severity,²⁴ but also to future risk, has important implications for FIT-based CRC screening strategies.

Declaration of interests

CGF undertook consultancy with Immunostics Inc., Ocean, NJ, USA, Mode Diagnostics, Glasgow, Scotland, and Kyowa-Medex, Tokyo, Japan. All other authors had no conflicting interests.

Funding

Funded by the Chief Scientist Office of the Scottish Government Health Department. Project: Establishing a Bowel Screening Research Unit (Grant Ref: CZH/6/4). JD was supported from a grant from Tenovus Tayside. The funders had no role in the study design, data collection, statistical analyses, interpretation of the results, or writing of the manuscript.

Research Ethics

Ethical approval was not required for this study. It was approved by the Programme Board, Scottish Bowel Screening Programme.

References

1. Digby J, Fraser CG, Carey FA, et al. Interval cancers using a quantitative faecal immunochemical test for haemoglobin (FIT) when colonoscopy capacity is limited. *J Med Screen OnlineFirst*, November 19, 2015: doi:10.1177/0969141315609634.
2. Chen LS, Yen AM, Chiu SY, Liao CS, Chen HH. Baseline immunochemical faecal occult blood test (iFOBT) concentration as a predictor of incident colorectal neoplasia: longitudinal follow-up of a Taiwanese population-based colorectal cancer screening cohort. *Lancet Oncol* 2011;**12**:551-8.
3. Yen AM, Chen SL, Chiu SY, et al. A new insight into fecal haemoglobin concentration-dependent predictor for colorectal neoplasia. *Int J Cancer* 2014;**135**:1203-12.
4. Fraser CG, Digby J, McDonald PJ, et al. Experience with a two-tier reflex gFOBT/FIT strategy in a national bowel screening programme. *J Med Screen* 2012;**19**:8-13.
5. Steele RJ, McDonald PJ, Digby J, et al. Clinical outcomes using a faecal immunochemical test for haemoglobin as a first-line test in a national programme constrained by colonoscopy capacity. *United European Gastroenterol J* 2013;**1**:198-205.
6. Atkin WS, Saunders BP. Surveillance guidelines after removal of colorectal adenomatous polyps. *Gut* 2002;**51**(Suppl v):v6–v9.

7. Steele RJ, McClements PL, Libby G, et al. Results from the first three rounds of the Scottish demonstration pilot of FOBT screening for colorectal cancer. *Gut* 2009;**58**:530-5.
8. Chen LS, Yen AM, Fraser CG, et al. Impact of faecal haemoglobin concentration on colorectal cancer mortality and all-cause death. *BMJ Open* 2013;**3**:e003740.
9. McDonald PJ, Strachan JA, Digby J, Steele RJ, Fraser CG. Faecal haemoglobin concentrations by gender and age: implications for population-based screening for colorectal cancer. *Clin Chem Lab Med* 2012;**50**:935-40.
10. Fraser CG, Rubeca T, Rapi S, Chen LS, Chen HH. Faecal haemoglobin concentrations vary with sex and age, but data are not transferable across geography for colorectal cancer screening. *Clin Chem Lab Med* 2014;**52**:1211-6.
11. Bujanda L, Sarasqueta C, Castells A, et al. Colorectal cancer in a second round after a negative faecal immunochemical test. *Eur J Gastroenterol Hepatol* 2015;**27**(7):813-8.
12. Azimoufoussé Assogba GF, Jezewski-Serra D, Lastier D, et al. Impact of subsequent screening episodes on the positive predictive value for advanced neoplasia and on the distribution of anatomic subsites of colorectal cancer: A population-based study on behalf of the French

colorectal cancer screening program. *Cancer Epidemiol* 2015;**39**(6):964-71.

13. Aniwan S, Rerknimitr R, Kongkam P, et al. A combination of clinical risk stratification and fecal immunochemical test results to prioritize colonoscopy screening in asymptomatic participants. *Gastrointest Endosc* 2015;**81**:719-27.
14. Stegeman I, de Wijkerslooth TR, Stoop EM, et al. Combining risk factors with faecal immunochemical test outcome for selecting CRC screenees for colonoscopy. *Gut* 2014;**62**(3):466-71.
15. Auge JM, Pellise M, Escudero JM, et al. Risk stratification for advanced colorectal neoplasia according to fecal haemoglobin concentration in a colorectal cancer screening program. *Gastroenterology* 2014;**147**:628-36.
16. Chen G, Mao B, Pan Q, Liu Q, Xu X, Nung Y. Prediction rule for estimating advanced colorectal neoplasm risk in average-risk populations in Southern Jiangsu Province. *Chin J Cancer Res* 2014;**26**:4-11.
17. Kaminski MF, Polkowski M, Kraszewska E, et al. A score to estimate the likelihood of detecting advanced colorectal neoplasia at colonoscopy. *Gut* 2014;**63**:1112-9.

18. Wong MC, Lam TY, Tsoi KK, et al. A validated tool to predict colorectal neoplasia and inform screening choice for asymptomatic subjects. *Gut* 2014;**63**:1130-6.
19. Chong HY, Roslani AC, Law CW. Colonoscopic prioritization in colorectal carcinoma screening using quantitative immunochemical faecal occult blood test: a pilot study. *Med J Malaysia*.2013;**68**:30-3.
20. Yeoh KG, Ho KY, Chiu HM, et al; Asia-Pacific Working Group on Colorectal Cancer. The Asia-Pacific Colorectal Screening score: a validated tool that stratifies risk for colorectal advanced neoplasia in asymptomatic Asian subjects. *Gut*. 2011;**60**:1236-41.
21. Omata F, Shintani A, Isozaki M, et al. Diagnostic performance of quantitative fecal immunochemical test and multivariate prediction model for colorectal neoplasms in asymptomatic individuals. *Eur J Gastroenterol Hepatol* 2011;**23**:1036-41.
22. Driver JA, Gaziano JM, Gelber RP, Lee IM, Buring JE, Kurth T. Development of a risk score for colorectal cancer in men. *Am J Med* 2007;**120**:257-63.
23. Digby J, McDonald PJ, Strachan JA, et al. Deprivation and faecal haemoglobin: implications for bowel cancer screening. *J Med Screen* 2014;**21**:95-7

24. Digby J, Fraser CG, Carey FA, et al. Faecal haemoglobin concentration is related to severity of colorectal neoplasia. *J Clin Pathol* 2013;**66**:415-9