

# Participation in faecal immunochemical testing-based colorectal cancer screening programmes in the northwest of Europe

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## Abstract

**Objective:** This study compared the participation in four faecal immunochemical testing-based screening programmes for colorectal cancer in Flanders, France, Basque country and the Netherlands, to identify factors to further optimize faecal immunochemical testing programmes.

**Method:** Background information and data on performance indicators were collected and compared for the four programmes.

**Results:** Invitation method, reminders, funding, faecal immunochemical testing cut-off and follow-up after positive faecal immunochemical testing differed in the four programmes. In France, only an invitation letter is sent by mail, while the sample kit must be collected from the general practitioner. In the other programmes, an invitation letter including the sample kit is sent by mail. Participation rates vary substantially according to the method of invitation, with the highest participation rates in the Netherlands (73.0%) and Basque country (72.4%), followed by Flanders (54.5%) and France (28.6%). Basque country (92.8%) and France (88.4%), the two programmes with most active involvement of general practitioners in referral for colonoscopy, had the highest participation rates for colonoscopy.

**Conclusions:** Large differences in screening participation observed between programmes according to the invitation method used suggest that changes to the design of the programme, such as including the sample kit with the invitation, or active involvement of GPs, might increase participation.

## Keywords

Colorectal cancer, screening, faecal immunochemical testing, invitation method, population-based, participation

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## Introduction

Many countries and regions have implemented colorectal cancer (CRC) screening by faecal occult blood testing (FOBT), in particular using faecal immunochemical testing (FIT).<sup>1</sup> Screening using FOBT is recommended by European Union CRC screening guidelines.<sup>2</sup> In France, a national population-based CRC screening programme was initiated in 2002 using guaiac FOBT, which was changed to FIT-based screening in April 2015. FIT-based screening was introduced in 2009 in the Basque country (Spain), in 2013 in Flanders (Belgium) and in 2014 in the Netherlands. As these four European programmes are geographically close and connected, and have recently been implemented, similar outcomes with respect to CRC screening may have been expected.

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The effectiveness of population-based screening programmes is driven not only by the sensitivity of the screening method but also by the availability of resources, healthcare infrastructure and population preferences. Population preferences in particular will be reflected in participation rate. To determine the optimal screening method for the population, pilot studies were performed in the Basque country, Flanders and the Netherlands before the initiation of the screening programmes. The Basque Country pilot study in 2009 demonstrated a participation rate with FIT screening of 64.3%.<sup>3</sup> In Flanders, the pilot study compared participation for two invitation strategies: FIT directly sent by mail (52.3%), or invitation only by mail and FIT to be collected from the general practitioner (GP) (24.6%).<sup>4</sup> In the Dutch pilot studies comparing different screening methods, FIT screening resulted in the highest CRC detection per invitee compared with other screening methods, such as gFOBT, colonoscopy and sigmoidoscopy screening.<sup>5-7</sup>

Crucial to optimal screening performance is the organizational structure of the programme, including pre-invitation letters, reminders, and FIT mailing; however, almost all studies of these aspects have been carried out in trials.<sup>4,8,9</sup> Actual performance in routine screening programmes may differ because many other factors are involved, such as government-endorsement, organization of healthcare systems and healthcare insurance. Participation in the trial preceding the implementation of the national Dutch programme, for example, was 60%, which was significantly lower than the 71% achieved in the first year of the national programme. It is unknown how the different organizational aspects will affect individuals residing in different countries, with different sociocultural backgrounds.

This study aimed to identify similarities and differences in the organizational structure of four large population-based CRC screening programmes using FIT in France, Basque country (Spain), Flanders (Belgium) and the Netherlands, and then compare these with important programme performance indicators of participation and detection rates to infer relationships between organizational structure and these programme outcomes.

## Methods

Information on year of initiation, target population, eligible population, screening interval, methods of invitation to FIT screening and to colonoscopy following a positive FIT, funding and executive organization of the four screening programmes was collected. The target population was defined for each population-based CRC screening programme according to programme-specific policies. The eligible population is the target population minus those who are ineligible for screening based on exclusion criteria. The eligible population was all individuals who should have been invited in 2016. This number can deviate from the total target population, because of biennial screening

or phased implementation of the national screening programme.

Data on performance indicators were extracted from each of the national or regional screening databases. In France, data were extracted from the database of French Public Health Agency (Santé Publique France) and Organized screening structure of the Big East region and the Pyrénées. Data from Flanders were extracted from the screening database, the Belgian Cancer Registry and reimbursement data from the Health insurance companies. All data from the Basque country were extracted from the programme (Programa de Cáncer ColoRectal) database, which is linked with medical records, population and hospital cancer registries. All data from the Netherlands were extracted from the national database for screening programmes (ScreenIT). Data on the invitees were collected in France starting in April 2015 until June 2017. Data on the invitees of 2016 were collected in the Basque country until December 2017, and in Flanders and the Netherlands until 30 June 2017.

Data were collected on the main performance indicators: participation rate, FIT positivity rate, participation rate in colonoscopy following positive FIT, detection rate of CRC or advanced neoplasia (AN) per participant and diagnostic yield. The definitions of performance indicators were those recommended in the European Union CRC screening guidelines.<sup>10</sup>

*Participation rate* was the number of persons sending back the FIT sample divided by the number receiving an invitation letter. For Flanders and France, persons were only considered participants if they returned the FIT sample within 12 months after the invitation. In the Basque country, participants were those who returned an assessable stool sample within six months following the invitation. In the Netherlands, individuals were considered participants until the date of the invitation of subsequent screening round. *Positivity rate* was the number of persons with a FIT result at or above the cut-off level, divided by the number of persons with an assessable stool sample. *Participation rate colonoscopy* was the number undergoing a colonoscopy divided by the number with a positive FIT result. *Detection rate* was defined as the number of persons with AN detected during colonoscopy per participant. AN was considered a relevant abnormality within a CRC screening programme, and was defined as CRC or any adenoma with histology showing  $\geq 25\%$  villous component or high-grade dysplasia or adenoma with size  $\geq 10$  mm. In Flanders, only adenomas with any villous component and/or high-grade dysplasia were counted as advanced adenoma, because no data were available on adenoma size or the amount of villous components. In the Basque country, in addition to histology, dysplasia and size, having  $\geq 3$  adenomas was also considered as advanced adenoma. *Diagnostic yield of the programme* was defined as the number of persons with AN detected during colonoscopy divided by all individuals who received an invitation. In Flanders, data of colonoscopy

yield were not linked to the date of invitees of the programme. The denominator can contain individuals invited in previous year.

The organizational structures of the four programmes were compared using thematic analysis to identify similarities or differences. Outcomes of the performance indicators for each of the programmes were then compared. To rule out the possibility that the observed difference may be related to cultural differences between populations rather than organizational differences, the programme of Basque country in Spain was compared with the Basque country in France, as these two regions are very close with respect to geographical location and cultural background. The different subgroups were compared using chi-squared test. This test was performed using R version 3.5.0.

## Results

The age range of the target populations differed, with France and the Basque country having the lowest starting age of 50 and the Basque country having the lowest stopping age of 69 (Table 1). All four programmes used a two-year screening interval. Exclusion criteria prior to invitation differed among the four programmes. In the Netherlands, persons are only excluded based on a positive FIT result of previous screening round (Table 1). France, Flanders and Basque country all excluded individuals with history of CRC, proctocolectomy and recently performed colonoscopy before invitation. France and Flanders also excluded individuals with a recently performed FIT test. Additionally, the Basque country excluded individuals with severe or terminal illness.

Methods of invitation also differed among the four programmes. The eligible population in France received an invitation letter to collect the FIT sample kit at the GP. In Flanders, the invitation included the FIT sample kit. In the Basque country and the Netherlands, a pre-invitation letter was sent, followed by an invitation letter including the FIT sample kit. All four screening programmes used a reminder letter, but all at different time points, ranging from 30 days (Basque country) to six months (France). The programme in France sent two reminder letters. All four programmes used different cut-offs for a positive FIT and referral to colonoscopy: in Flanders, 15 µg Hb/g faeces; the Basque country, 20 µg Hb/g faeces; France, 30 µg Hb/g faeces and the Netherlands, 47 µg Hb/g faeces (Table 1).

Among the four programmes, 18.9 million individuals were invited to participate in FIT screening. The highest participation rate was observed in the Netherlands (73.0%), followed by the Spanish Basque country (72.4%), Flanders (54.5%) and France (28.6%,  $p < 0.001$ ). Because of the different FIT cut-off levels used, the positivity rate differed between the four programmes, from 4.7% in France to 6.7% in Flanders ( $p < 0.001$ ). The participation rate for colonoscopy following a positive FIT result was 92.8% in the Basque

country, 88.4% in France, 81.9% in Flanders and 82.8% in the Netherlands ( $p < 0.001$ ). Detection rate for AN per participant was highest in the Netherlands (2.3%) and lowest in Flanders (1.0%). Diagnostic yield for AN per invitee was highest in the Basque country (1.6%) and lowest in France (0.4%,  $p < 0.001$ ). Figure 1 shows the impact of FIT participation rate, positivity rate and colonoscopy participation rate on the diagnostic yield. This shows that programmes with the highest FIT participation rate have the highest diagnostic yield, but this is less visible for the positivity rate and colonoscopy participation rate.

In the comparison of the French with the Spanish Basque country, despite cultural similarities, differences in screening performance indicators were observed (Table 2). The participation rate in the Spanish part was 72.4%, 2.5 times as high as in the French part, with 24.6% ( $p < 0.001$ ) (Table 3). Participation rate to colonoscopy was high in both the Spanish and the French part (92.8% and 87.4%, respectively;  $p = 0.37$ ).

## Discussion

Large differences in screening participation were observed between programmes, in line with the invitation method used. There are several possible explanations. First, sending the FIT home is more effective than having it collected at the GP. Almost all studies show a large increase in participation when including the FIT sample kit with the invitation.<sup>8,11–13</sup> One Italian study showed only a modest increase in participation, but this study was performed in previously screened individuals (used to another screening strategy).<sup>14</sup> One French study showed low uptake rates when directly mailing the FOBT.<sup>14</sup> This inconsistency may be due to the test modality, gFOBT instead of FIT, resulting in lower participation rates.<sup>15</sup>

A second explanation for a higher FIT participation may be the advanced notification letter, as illustrated by the higher participation rate in the Spanish Basque country and the Netherlands. However, this will only explain a small proportion of the total difference, as studies have shown that sending a pre-invitation letter results in a three percentage point increase.<sup>9,16</sup> Only one study from Australia showed a higher increase, nine percentage points, when a pre-invitation letter was sent.<sup>17</sup> The higher participation with both direct mailing and the pre-invitation letter is in line with a recent systematic review.<sup>18</sup> However, one large difference was that GP involvement improved participation. This has also been shown in France, with an increase of 4% if GPs were notified of the screening status of their patients so that they could actively promote CRC screening to non-participants.<sup>8</sup> We showed the opposite in this study; in a country that sends out the FIT by mail with no involvement of GPs, such as the Netherlands, participation rates were very high, while in a country with active involvement of GPs, such as France, participation rates were substantially

**Table 1.** Background information on screening programmes.

	France	Flanders (Belgium)	Netherlands	Basque country (Spain)
<i>1. General</i>				
Year of introduction	Started in 2009 and switched nationally to the FIT in April 2015.	October 2013	2014	2009
Piloting	Yes	Yes	Yes	No
Public awareness campaigns	Yes. Through dedicated websites: National Cancer Institute (INCa), Health Insurance, some local initiatives.	Yes. Since 2015 short announcements on the Flemish television and since 2017 advertising in public transport. Both only during the month March (international CRC month).	Yes. Information on the website of the national institute of public health and the environment. Only at the start in 2014, there was extra media attention through television and radio. No advertising.	Yes. National yearly campaign by patient association and main results are presented in television, leaflets, radio and social network.
Population-based programme	National	Regional	National	Regional
Age group	50–74	56–74 Per 1 July 2017, it was extended to 55–74, and per July 2018, it was extended to 53–74	55–75	50–69
Screening interval	Two years	Two years	Two years	Two years
Cut-off level (Hb/g faeces)	30 µg	15 µg	47 µg	20 µg
Brand name of the test	OC-Sensor, Eiken, Japan	OC-Sensor, Eiken, Japan	FOB-Gold, Sentinel, Italy	OC-Sensor, Eiken, Japan
<i>2. Methods of invitation</i>				
Pre-invitation letter	No	No	Yes, three weeks in advance	Yes, four weeks in advance
Invitation by mail including FIT	No, test to be collected at GP's office	Yes	Yes	Yes
Reminder letter	12 Weeks and 24 weeks	8 Weeks	6 Weeks	4 Weeks
Exclusion of individuals before invitation	Yes	Yes	No	Yes
Exclusion criteria mentioned in the invitation letter	No	Past or current treatment of colorectal cancer, occult blood in stool and unexplained and persistent change in the bowel movement patterns, colonoscopy in past 10 years, stool test in past two years, higher risk at CRC, having one or more first relatives who have/had CRC.	Past or current treatment of colorectal cancer, occult blood in stool and unexplained and persistent change in the bowel movement patterns.	Colonoscopy performed ≤ 5 years.
<i>3. Methods of invitation to diagnostic colonoscopy</i>				
Invitation to follow-up colonoscopy	By letter. Results are sent to the participant, the general practitioner.	By letter. Results are sent to the participant and the general practitioner.	By letter, including appointment for colonoscopy intake. Results are	By letter. Results are sent to the participant. In that letter,

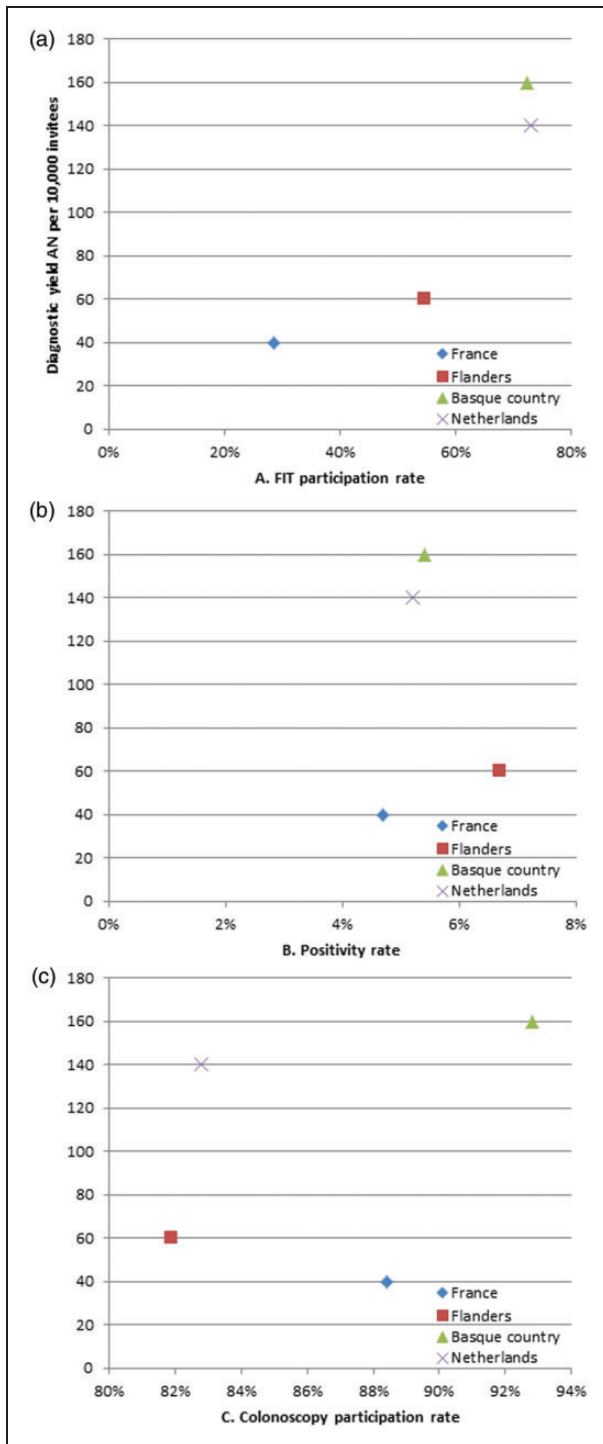
(continued)



Table 1. Continued

	France	Flanders (Belgium)	Netherlands	Basque country (Spain)
Reminder following initial invitation colonoscopy	practitioner and the structure in charge of organized CRC screening. No	No, from 2019, persons receive a new invitation for colonoscopy intake two years after their positive FIT. When not following this second advise, two years later (four years after the positive FIT), they will receive a new FIT.	Yes, persons receive a no show letter. In case of no response, persons received a new invitation for colonoscopy intake after two years after positive FIT (until 2017). From 2018 onwards, persons will receive a new FIT after two years.	participant is recommended to visit GP for colonoscopy referral. Yes, after 30 days, the participant is reminded to make an appointment with the general practitioner for colonoscopy referral. If participants reject to have colonoscopy follow-up, they will receive a new FIT after two years.
4. Organization				
Funding of screen test	Free of charge, but not the GP's encounter to collect the kit that is covered at 70% by national public health insurance. The remaining 30% are being covered by complementary insurance if the participant has one.	Free of charge	Free of charge	Free of charge
Funding of colonoscopy	Standard healthcare insurance covers 70%. The remaining 30% is covered by complementary insurance if any.	Standard healthcare insurance and partially payment of personal funds (out-of-pocket costs).	Standard healthcare insurance. Note: Not in all situations colonoscopy costs are fully covered by the healthcare insurance. Up to 350–850 euro are paid of personal funds (out-of-pocket costs, only once a year a deductible excess of all healthcare costs).	Free of charge
Responsible organization of the screening programme	INCa, National cancer institute, with dedicated screening structure in each of the departments. The latest are in charge of invitations.	Centre for Cancer Detection (CvKO). In Flanders, CvKO consists of five regional screening programmes, but the invitation letter, reminder letter and FIT results are organized by one central organization. The five regional screenings programmes offer free telephone advice and are responsible for administration.	The national institute of public health and the environment is the responsible organization of the programme. Five regional screening organizations are responsible for the execution of the programme. Those five regional screening organizations are brought together in one national cooperation.	The Basque Health Service is responsible for the execution of the programme, according with authorities planning, organizing and monitoring the quality of the process and results. The final responsibility of the programme is the Regional Ministry of Health in the Basque country.

CRC: colorectal cancer; GP: general practitioner; FIT: faecal immunochemical testing.



**Figure 1.** Diagnostic yield of advanced neoplasia by FIT participation rate.

AN: advanced neoplasia; FIT: faecal immunochemical testing.

lower. We hypothesize that GP endorsement can have a positive impact on participation, as long this requires no effort for the participants. This is in line with findings of the CRC screening programme in England, which showed an increase in participation if the invitation letter was added with a GP endorsement banner.<sup>19</sup>

Our analysis of the two Basque regions in France and Spain showed that very similar cultures can have very different rates in screening participation, and that culture may not be the driving factor of performance differences between programmes, although we cannot rule out cultural differences completely. We know from the literature that cultural difference in screening attitude is also observed in the participation rates of other cancer screening programmes. In 2016, for example, participation in breast cancer screening was also lower in Flanders (51.9%) than in the Netherlands (77.6%) and the Basque country (80.1%), with France having the lowest participation rate (50.7%).<sup>20–22</sup> The participation rate for breast cancer screening in France is similar to Flanders, while there is a much larger difference in participation rate for CRC screening. Gender cannot explain this difference, as both men and women show a similar pattern in participation. This again reflects the negative impact of using a different invitation method in France for FIT-based screening.

Participation rate to follow-up colonoscopy was high in all four screening programmes, although slightly below the recommended 85% in the Netherlands and Flanders. We hypothesize that higher participation to follow-up colonoscopy in Basque country and France may be the result of the active involvement of GPs during the screening process. In both countries, GPs play an active role in selecting the population eligible for FIT screening, at collection of the screening test, at follow-up after a negative FIT, or in defining the eligible population by excluding those with severe comorbidity from invitation. Consequently, those participating in FIT screening are all healthy enough to undergo follow-up colonoscopy. Conversely, it may also explain the lower participation to colonoscopy in the Netherlands, as there is no exclusion of individuals based on co-morbidities or medical history. Active involvement of GPs after positive FIT only for referral to colonoscopy, without involvement in the total screening process, will be less effective.<sup>8</sup> Reimbursement differences for colonoscopy do not seem to explain participation differences. Although in the Basque country, the colonoscopy is free of charge, participation in France was only slightly lower, while French individuals may have significant expenses which have to be paid from personal funds. One explanation may be that in France, only the most motivated individuals collect the FIT sample kit at their GP practice, and they may thus also be more motivated to attend colonoscopy in case of a positive FIT.

Positivity rate differed for all four programmes, due to three important reasons: cut-off level of the FIT, target age group and screening round (first or subsequent round).<sup>23</sup> Higher cut-off level and subsequent screening round will result in lower positivity rate, while higher age will result in higher positivity rate. The same explanations may partly hold for the difference in diagnostic yield of the programme. Previous research showed that using a higher FIT cut-off level will have a negative

**Table 2.** Performance indicators for France, Flanders, the Netherlands and Basque country.

	France	Flanders	Netherlands	Basque country	<i>p</i>
Calendar year	2015–2016	2016	2016	2016	
Age (years)	50–74	56–74	59–76	50–69	
Target population	19,043,771	1,447,434 <sup>a</sup>	Unknown	273,084	
Eligible population	16,701,387	830,665	1,543,223	229,380	
Invited	16,701,387	571,034	1,457,976	229,380	
%	100	68.7 <sup>a</sup>	94.5	87.7	
Number of participants	4,779,845	311,453	1,063,651	166,110	<0.001
Participation rate FIT % (95% CI)	28.6 (28.6–28.6)	54.5 (54.4–54.7) <sup>b</sup>	73.0 (72.9–73.0)	72.4 (72.2–72.6)	
Men	27.8	53.1	71.1	70.0	
Women	30.8	56.0	74.8	74.6	
Screen round	Any round	First and second	First and second	First to fourth round	
Cut-off level (Hb/g faeces)	30 µg	15 µg	47 µg	20 µg	
Positivity rate % (95% CI)	4.7 (4.7–4.7)	6.7 (6.7–6.8)	5.4 (5.4–5.5)	5.2 (5.1–5.3)	<0.001
Participation rate colonoscopy % (95% CI)	88.9 (88.8–89.0) <sup>c</sup>	81.9 (81.4–82.4)	82.8 (82.5–83.1)	92.8 (92.3–93.4)	<0.001
Detection rate					
AN% (95% CI)	1.5 (1.5–1.5) <sup>c</sup>	1.0 (1.0–1.1)	2.3 (2.2–2.3)	1.9 (1.9–2.0)	<0.001
CRC% (95% CI)	0.31 (0.30–0.32) <sup>c</sup>	0.28 (0.26–0.30)	0.35 (0.34–0.36)	0.20 (0.18–0.22)	<0.001
Diagnostic yield programme					
AN% (95% CI)	0.4 (0.4–0.4) <sup>c</sup>	0.6 (0.5–0.6)	1.6 (1.6–1.7)	1.4 (1.3–1.4)	<0.001
CRC% (95% CI)	0.09 (0.09–0.09) <sup>c</sup>	0.15 (0.14–0.16)	0.25 (0.25–0.26)	0.15 (0.13–0.16)	<0.001

Note: Advanced neoplasia was defined as CRC or any adenoma with histology showing  $\geq 25\%$  villous component or high-grade dysplasia or adenoma with size  $\geq 10$  mm. In Basque country, also  $\geq 3$  adenomas were considered AN. In Flanders, only adenoma with a villous component and/or high-grade dysplasia was counted as advanced adenoma. There were no data available on the size or the amount of villous components in an adenoma. Detection rate: invitees with CRC or AN per participant. Diagnostic yield: individuals with CRC or AN per invitees.

AN: advanced neoplasia; NA: not available; FIT: faecal immunochemical testing.

<sup>a</sup>Eligible population in Flanders is the total number aged 56–74 for two years minus those excluded for invitation. Eligible population for 2016 only could not be provided.

<sup>b</sup>Coverage by examination, also including opportunistic screening by FIT or colonoscopy, resulted in 65.5% of the target population to be screened.

<sup>c</sup>In France, the participation rate of colonoscopy and number of colorectal cancers and advanced neoplasia was based on data from April 2015 until December 2015.

**Table 3.** Outcome performance indicators Basque region.

	Basque country in France	Basque country in Spain	<i>p</i>
Year	2016	2016	
Age	50–74	50–69	
Invited	45,923	229,380	
Number of participants	11,293	166,110	
Participation rate FIT % (95% CI)	24.6 (24.2–25.0)	72.4 (72.2–72.6)	<0.001
Cut-off level (Hb/g faeces)	30 µg	20 µg	
Positivity rate % (95% CI)	4.8 (4.4–5.2)	5.2 (5.1–5.3)	0.07
Participation rate follow-up colonoscopy % (95% CI)	87.4 (84.4–90.0)	92.8 (92.3–93.4)	0.37
Detection rate			
AN% (95% CI)	1.4 (1.2–1.7)	1.9 (1.9–2.0)	<0.001
CRC% (95% CI)	0.27 (0.19–0.39)	0.20 (0.18–0.22)	
Diagnostic yield			
AN% (95% CI)	0.4 (0.3–0.4)	1.4 (1.3–1.4)	<0.001
CRC% (95% CI)	0.07 (0.05–0.10)	0.15 (0.13–0.16)	

AN: advanced neoplasia; FIT: faecal immunochemical testing.

Note: Advanced neoplasia was defined as CRC or any adenoma with histology showing  $\geq 25\%$  villous component or high-grade dysplasia or adenoma with size  $\geq 10$  mm. In Basque country, also  $\geq 3$  adenomas were considered AN. Detection rate: invitees with CRC or AN per participant. Diagnostic yield: individuals with CRC or AN per invitees.

impact on detection rates.<sup>24</sup> Nevertheless, the impact of positivity rate and colonoscopy participation rate on diagnostic yield was modest. Programmes with the highest FIT participation rate still have the highest diagnostic yield, regardless of their positivity rate or colonoscopy participation rate. This highlights the importance of high participation rates to primary screening, as this will result in the highest detection of AN. However, the difference in detection rates of AN is surprising, with the Netherlands having high detection rates while having the highest cut-off. This might be related to several factors: more incident screens in France and Basque country (lower detection), more prevalent screens in Flanders and the Netherlands, difference in age range (lower age, lower detection) and stricter definition of advanced adenomas in Flanders (lower detection).

Our study has three important strengths. It is the first study that gives detailed information on organizational structure of four large population-based programmes provided by representatives of each country. These details are generally unknown, as key elements of CRC screening programmes are only described in the local language (Flemish, French, Spanish/Basque, Dutch). These details can be used by other countries/regions considering CRC screening and are valuable for policymakers. In addition, our study contains the most recent outcomes of these large population-based programmes, all using the same test modality (FIT). Lastly, our study compared screening programmes of neighbouring countries with cultural similarities and differences, and can thus address the impact of cultural and organizational aspects on the uptake of CRC screening.

The study has also some limitations. Comparing quality indicators was challenging due to different definitions and differences in cut-off level and number of screening rounds. Unfortunately, we could not restrict the comparison to first screen round data only, as not all programmes have detailed information. In addition, data collection may be of concern as, for example, France does not yet have a centralized recording of quality indicators and does not centrally collect data on diagnostic yield.

Our findings suggest that organizational structure, for example, sending out the FIT, pre-invitation letter, involvement of the GP in the screening programme and colonoscopy referral, has an impact on the participation rate to FIT and follow-up colonoscopy. These results could be used to optimize each of the four screening programmes, or as an example for other organized FIT-based CRC screening programmes. Possibilities for optimization can differ for every organized programme, as healthcare systems, funding of the colonoscopy and available resources also differ. Interventions for optimization will have cost implications, and these results can therefore be used to explore the additional benefits and costs for each programme. France has already begun to optimize the screening programme, by deciding to mail the FIT with the first reminder, but only to those individuals who participated in

previous rounds. The study by Giorgi-Rossi et al. suggests that these individuals may not be the best target.<sup>25</sup>

Although posting the FIT and actively approaching FIT positives for the colonoscopy seems to be most effective, this may be considered an infringement of free will.<sup>26</sup> The goal of the screening programmes should not be high participation, but the level of informed choice, although infringement of free will should be considered in the light of the strength of the recommendation. For CRC screening, and in particular for colonoscopy in FIT positives, the evidence for net benefit is considered strong, and we can presume that the vast majority would weigh the balance of benefits and harm in the direction of having the intervention.<sup>27</sup> Moreover, there is no indication that high participation in the Netherlands, for example, results in a lower level of informed choice.<sup>28,29</sup> Besides these ethical considerations, there is also a remaining difference in participation that cannot be explained by the organizational structure and is difficult to unravel. This difference seems to be a difference in attitude towards screening in general between the different regions or countries, and it is unclear how this arises or how it can be solved.

## Conclusion

This study shows that there is large variability in the design of these screening programmes, and that programmes with more evidence-based interventions in place (e.g. mailed-out FIT, pre-invitation and reminder letters) experience significantly higher screening participation than those without. Active involvement of GPs may result in slightly higher participation with colonoscopy after a positive FIT result, but should not come at the expense of extra barriers to collect the FIT, which can dramatically reduce primary screening participation.

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