

Diagnostic Accuracy of Fecal Calprotectin for Pediatric Inflammatory Bowel Disease in Primary Care: A Prospective Cohort Study

Gea A. Holtman, MSc¹

Yvonne Lisman-van Leeuwen, PhD¹

Boudewijn J. Kollen, PhD¹

Obbe F. Norbruis, MD, PhD²

Johanna C. Escher, MD, PhD³

Angelika Kindermann, MD, PhD⁴

Yolanda B. de Rijke, MD, PhD⁵

Patrick F. van Rbeenen, MD, PhD⁶

Marjolein Y. Berger, MD, PhD¹

¹Department of General Practice, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

²Department of Pediatrics, Isala Hospital, Zwolle, The Netherlands

³Department of Pediatric Gastroenterology, Erasmus University Medical Centre-Sophia Children's Hospital, Rotterdam, The Netherlands

⁴Department of Pediatric Gastroenterology, Emma Children's Hospital/Academic Medical Center, Amsterdam, The Netherlands

⁵Department of Clinical Chemistry, Erasmus University Medical Centre, Rotterdam, The Netherlands

⁶Department of Pediatric Gastroenterology, Beatrix Children's Hospital/University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

Conflicts of interest: authors report none.

CORRESPONDING AUTHOR

Marjolein Y. Berger, MD, PhD
Department of General Practice, FA21
University Medical Center Groningen
University of Groningen
PO Box 196, 9700 AD
Groningen, The Netherlands
m.y.berger@umcg.nl

ABSTRACT

PURPOSE In specialist care, fecal calprotectin (FCal) is a commonly used noninvasive diagnostic test for ruling out inflammatory bowel disease (IBD) in children with chronic gastrointestinal symptoms. The aim of this study was to evaluate the diagnostic accuracy of FCal for IBD in symptomatic children in primary care.

METHODS We studied 2 prospective cohorts of children with chronic diarrhea, recurrent abdominal pain, or both: children initially seen in primary care (primary care cohort) and children referred to specialist care (referred cohort). FCal (index test) was measured at baseline and compared with 1 of the 2 reference standards for IBD: endoscopic assessment or 1-year follow-up. Physicians were blinded to FCal results, and values greater than 50 $\mu\text{g/g}$ feces were considered positive. We determined specificity in the primary care cohort and sensitivity in the referred cohort.

RESULTS None of the 114 children in the primary care cohort ultimately received a diagnosis of IBD. The specificity of FCal in the primary care cohort was 0.87 (95% CI, 0.80-0.92). Among the 90 children in the referred cohort, 17 (19%) ultimately received a diagnosis of IBD. The sensitivity of FCal in the referred cohort was 0.99 (95% CI, 0.81-1.00).

CONCLUSIONS The findings of this study suggest that a positive FCal result in children with chronic gastrointestinal symptoms seen in primary care is not likely to be indicative of IBD. A negative FCal result is likely to be a true negative, which safely rules out IBD in children in whom a primary care physician considers referral to specialist care.

Ann Fam Med 2016;14:437-445. doi: 10.1370/afm.1949.

INTRODUCTION

Primary care physicians frequently manage recurrent abdominal pain or diarrhea in children. These symptoms account for approximately 2% to 5% of all childhood consultations.¹⁻³ Although they are typically functional in origin, it is essential that organic disease be ruled out. Inflammatory bowel disease (IBD), that is, Crohn disease and ulcerative colitis, is an organic disease that primary care physicians should not miss. Delay in diagnosing IBD, and the resultant delay in receipt of appropriate treatment, may prolong suffering and can lead to complications such as anemia, irreversible growth failure, and delayed sexual maturation.^{4,5}

According to guidelines, primary care physicians should refer children with chronic diarrhea, recurrent abdominal pain, or both for specialist care if red flags are present.^{6,7} The red flags are nonspecific and discriminate poorly between functional and organic gastrointestinal diseases,⁸⁻¹⁰ often leading to referral and extensive diagnostic testing. For children with functional disorders, referral or extensive testing may delay appropriate interventions and further decrease well-being.^{11,12}

Calprotectin is a calcium-binding protein released from neutrophils during intestinal inflammation that can be easily measured in feces.^{13,14} In specialist care, evidence shows it to be a useful, simple, noninvasive test

that can rule out IBD in children with gastrointestinal symptoms.¹⁵⁻¹⁷ The diagnostic accuracy of fecal calprotectin (FCal) has not been assessed in children evaluated in primary care, however.^{10,18,19} Primary and specialist care often have different populations, case mixes, and disease severity, which can affect the pre-test probability of IBD and the sensitivity and specificity of the FCal test. The diagnostic accuracy of FCal in the primary care setting should therefore be clarified before this test is recommended for routine use in primary care. We set out to study the diagnostic accuracy of FCal for identifying IBD in children with chronic gastrointestinal symptoms in primary care.

METHODS

Study Design

This was a prospective cohort study with a delayed-type cross-sectional design.²⁰ Children in the Netherlands with chronic gastrointestinal symptoms were included from July 2011 to July 2013 and had 12 months of follow-up. We studied 2 cohorts: (1) the primary care cohort consisted of consecutive children who were seen by any of 64 primary care physicians (38 practices); (2) the referred cohort consisted of consecutive children who were referred for diagnostic workup by primary care physicians and general pediatricians to any of 4 general hospitals or 3 academic centers, as well

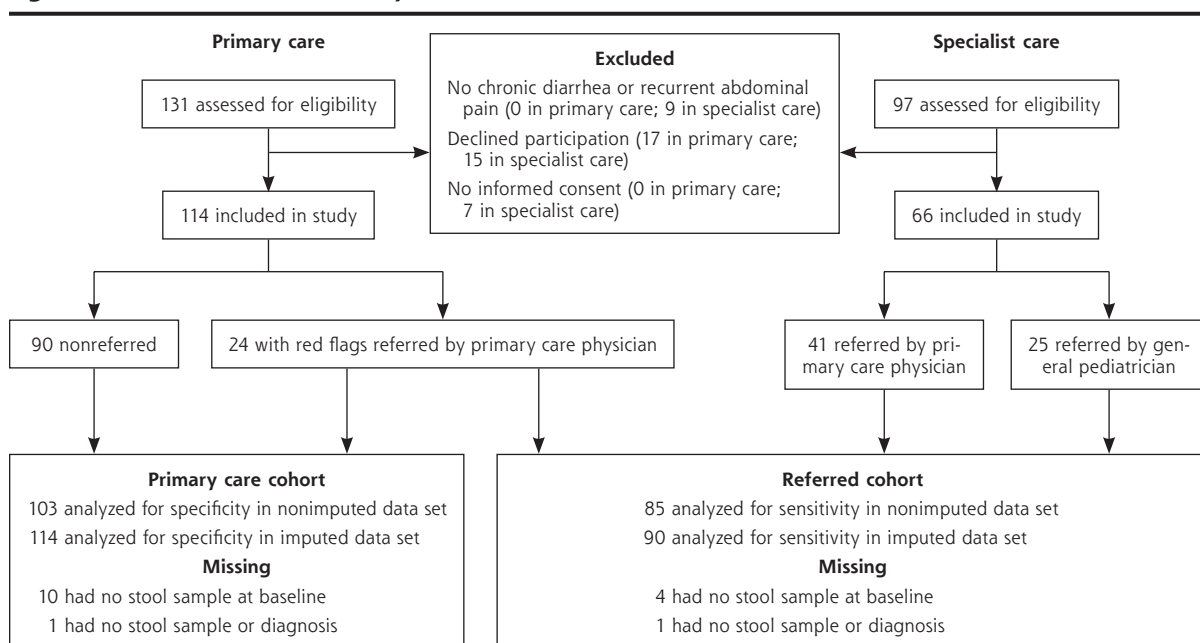
as children selected from the primary care cohort based on the presence of at least 1 red flag (Figure 1).

The medical ethics committee of the University Medical Centre Groningen approved the study. Written informed consent was provided by the parents of all children and by all children aged 12 years or older. The study design has been described in more detail elsewhere.²¹

Participants

Children aged 4 to 18 years who sought care for chronic diarrhea, recurrent abdominal pain, or both were eligible. Chronic diarrhea was defined as soft to watery stool (score of 5, 6, or 7 on the Bristol stool chart²²) for at least 2 weeks or at least 2 episodes in the past 6 months. Recurrent abdominal pain was defined as at least 2 episodes of abdominal pain or discomfort in the past 6 months. Children were excluded if they had a previous diagnosis of chronic organic gastrointestinal disease; an evaluation with endoscopy or FCal for gastrointestinal symptoms in 6 months before this study; or difficulty in understanding questionnaires. Furthermore, we excluded children with long-term use (>3 months) of antibiotics, nonsteroidal anti-inflammatory drugs, or oral corticosteroids in the past 6 months, as well as those aged younger than 4 years, because previous studies have demonstrated elevated calprotectin concentrations in these groups.^{23,24}

Figure 1. Patient flow in the study.



Note: The group of children who were mainly seen in primary care and selected for referral to specialist care based on ≥ 1 red flags were evaluated in both analyses. Fecal calprotectin was not measured in 14 children because no stool sample was collected (9 children), the sample was not stored (2 children), or the sample of feces was too small (3 children). Two children with no stool sample were evaluated in both analyses.

Baseline Evaluation

All participating physicians assessed children for the presence of 10 red flags suggestive of IBD using a structured evaluation form. These red flags consisted of 6 alarm symptoms and 4 blood markers (Table 1).²⁵⁻²⁷ Feces were tested for pathogens—*Salmonella enterica*, *Campylobacter jejuni*, *Shigella* spp/enteroinvasive *Escherichia coli* (EIEC), and Shiga toxin–producing *Escherichia coli* (STEC)—and for parasites—*Giardia lamblia*, *Cryptosporidium* spp, *Dientamoeba fragilis*, and *Entamoeba histolytica*—using real-time polymerase chain reaction.

Patient Flow

All children in the primary care cohort were evaluated by their primary care physician. Children were selected for further diagnostic workup based on the presence of at least 1 red flag. Children in the referred cohort were evaluated by a pediatric gastroenterologist, who decided whether the child required endoscopic evaluation. All children were followed up for 1 year, during which time the attending physician was free to refer a child for further diagnostic workup. At 1 year, the physician did a structured physical examination of all children with persisting gastrointestinal symptoms who had not received a diagnosis of IBD. Children with at least 1 red flag at this time were seen by a pediatric gastroenterologist for further evaluation. The attending physicians of children lost to follow-up were contacted after 1 year to provide updated information on persisting symptoms and additional diagnoses.

Confirmatory Diagnosis

IBD was diagnosed by esophagogastroduodenoscopy, ileocolonoscopy, and histopathologic examination

according to the Porto criteria.⁶ A negative endoscopy was defined as the absence of endoscopic and histopathologic findings of IBD. Children were considered not to have IBD if, after 12 months, the attending physician found no red flags or the pediatric gastroenterologist decided that red flags were not related to IBD.

Index Test

We used as the index test an FCal test with a lowest sensitivity of 19.5 µg/g feces and a cutoff point of greater than 50 µg/g feces according to the manufacturer. Stool samples collected at baseline were stored at –80°C and analyzed by a commercially available quantitative enzyme-linked immunosorbent assay (PhiCal test, Calpro AS) at the end of the data collection period (September 2014) in the department of clinical chemistry at Erasmus Medical Center. All physicians, pathologists, researchers, and patients were blinded to the outcome of the FCal test and were not allowed to apply for another FCal test during the data collection period. The laboratory staff evaluating FCal were blinded to the children's clinical characteristics and diagnoses.

Statistical Analysis

Very few children seen in primary care with chronic gastrointestinal symptoms have IBD. When introducing new tests in these children, it is important that false-positive outcomes are minimized to avoid referrals for endoscopies. We therefore estimated specificity with adequate precision of FCal measurements in the primary care cohort. Given an expected specificity of 93%,^{18,28-30} a 95% CI and precision of 5%, an IBD prevalence of 5%, and a loss to follow-up of 10%, we needed to include 118 children in the primary care cohort.

Table 1. Definitions of Red Flags in Inflammatory Bowel Disease

Red Flag	Method of Ascertainment	Definition of Positive Finding
Alarm symptoms		
Involuntary weight loss	History	Involuntary decrease in weight of >1 kg
Rectal blood loss	History	Rectal blood loss with defecation without constipation according to Rome III criteria
Family history of IBD	History	Affected first-degree relative(s)
Growth failure	History and physical examination	Target height range > –1 standard deviation score
Extraintestinal symptoms	Physical examination	Eyes (episcleritis, scleritis, uveitis), skin (erythema nodosum, pyoderma gangrenosum, psoriasis), mouth ulcers, finger clubbing, arthritis
Perianal lesions	Physical examination	Skin tags, hemorrhoids, fissures, fistulas, abscesses
Blood markers		
Hemoglobin	Local laboratory	Age 4-12 y: <7.1 mmol/L; age 12-18 y: boys <8.1 mmol/L, girls <7.4 mmol/L ²⁵
C-reactive protein	Local laboratory	>10 mg/L ²⁶
Erythrocyte sedimentation rate	Local laboratory	>20 mm/h ²⁶
Platelet count	Local laboratory	>450 × 10 ⁹ /L ²⁷

IBD = inflammatory bowel disease.

The likelihood of IBD increases in children in whom the primary care physician considers a referral. In this group, it is important not to miss a case of IBD, and a low false-negative rate is preferred. We therefore estimated sensitivity with adequate precision of FCal measurement in the referred cohort. Given an expected sensitivity of 95%,^{18,28,30} a 95% CI and precision of $\pm 10\%$, an IBD prevalence of 20%, and a loss to follow-up of 10%, we needed to include 100 children in the referred cohort.²¹

For secondary outcomes, we calculated specificity, posttest probability, and area under the receiver operating characteristic curve (AUC) with the 95% CI in the referred cohort. We also determined the effect of applying different FCal cutoff values ($>50 \mu\text{g/g}$, $>100 \mu\text{g/g}$, $>250 \mu\text{g/g}$)^{16,31} on the test characteristics, number of referrals, and missed diagnoses of IBD.

We assumed that children included in specialist care were comparable to those with at least 1 red flag from the primary care cohort. To test this assumption, we compared the characteristics of these groups. In order to evaluate spectrum bias (whereby the test setting affects the test performance), we performed subgroup analyses in the referred cohort, comparing children who were referred by a primary care physician with those referred by a general pediatrician. To evaluate the likelihood of differential verification bias,³² we separately calculated the test characteristics for both endoscopy and 12-month follow-up.

We conducted a missing value analysis to rule out missing not at random (MNAR) as a possible explanation for missing data from the variables of FCal and diagnosis. The missing data were replaced based on a multiple imputation procedure (conditional specification, predictive mean matching,³³ 20 iterations, and 20 data sets). The patient characteristics, all symptoms, all diagnostic tests, setting, endoscopy performed, and whether IBD was diagnosed were included as predictors. We used the Rubin rule to calculate the pooled AUC.³⁴ Results of the analyses on the nonimputed and imputed data sets were compared to assess the effect of multiple imputations on diagnostic accuracy. Statistical analyses were performed with IBM SPSS for Windows, version 20.0 (IBM Corp).

RESULTS

Participants

The primary care cohort had 114 children, and the referred cohort had 90 children (24 with red flags from primary care plus 66 from specialist care) (Figure 1). Table 2 shows that the 66 children from specialist care more often had weight loss, extra-intestinal symptoms, and decreased hemoglobin levels compared with the

24 children with red flags from primary care. The 25 children who were referred to the pediatric gastroenterologist by a general pediatrician showed a higher IBD prevalence than the 65 children who were referred by their primary care physician (12 vs 5 cases).

Diagnoses

None of the children in the primary care cohort received a diagnosis of IBD (Table 3). The final diagnosis was based on 12 months' follow-up in 111 children and endoscopy in 2 children. Of the 90 children in the referred cohort, 29 (32%) underwent endoscopic evaluation, and 17 (19%) ultimately received a diagnosis of IBD.

Fecal Calprotectin

The median intervals from stool collection to endoscopy were 4 days and 8 days for children with IBD and without IBD, respectively; however, 11 of the 27 children (2 had missing samples) who underwent endoscopy experienced a delay of more than 1 month. Two children in the primary care cohort had similar calprotectin values in samples collected at baseline and shortly before endoscopy. In total, values were missing for 11 children (9.6%) in the primary care cohort and 5 (5.5%) in the referred cohort.

The contingency tables and test characteristics for FCal by follow-up or endoscopy are presented in Figure 2 for the nonimputed data set. The outcomes were comparable for all parameters in the imputed and nonimputed data sets. In the imputed data set, the specificity of FCal for IBD in the primary care cohort was 0.87 (95% CI, 0.80-0.92). Six of the 24 children with red flags and 9 of the 90 children without red flags had a positive calprotectin value. In the referred cohort, the sensitivity was 0.99 (95% CI, 0.81-1.00). The AUC of FCal in referred children cohort was 0.98 (95% CI, 0.96-1.00).

In subgroup analyses, the sensitivities of the test in children referred to specialist care by their primary care physician and by a general pediatrician were 0.98 (95% CI, 0.58-1.00) and 1.00 (95% CI, 0.76-1.00), respectively. An increase in the cutoff value for FCal from $50 \mu\text{g/g}$ to $250 \mu\text{g/g}$ feces in the referred cohort would lead to an extra 14% reduction in referrals for diagnostic workup for IBD, but would also increase the percentage of missed IBD diagnoses from 0% to 18% (Table 4).

DISCUSSION

This is the first study to evaluate the test characteristics of FCal as a marker for IBD in children seen for chronic gastrointestinal symptoms in primary care.

None of the children in the primary care cohort ultimately received a diagnosis of IBD, suggesting that children with chronic gastrointestinal symptoms should not be referred directly for evaluation of IBD. Current guidelines recommend referral for diagnostic workup based on the presence of red flags. In our primary care cohort, referrals based on red flags would have resulted in a higher false-positive referral rate for IBD compared with referrals based on FCal exceeding 50 µg/g feces. To date, however, there is insufficient evidence to recommend FCal as a tool to guide decisions about referral for diagnostic workup of all children with chronic gastrointestinal symptoms seen in primary care.

FCal showed high sensitivity (0.99; 95% CI, 0.81-1.00) in the referred cohort. Therefore, a negative FCal result may safely rule out IBD and thereby reduce the number of referrals for evaluation of IBD in children

whom the primary care physician considers a referral. Nevertheless, the 95% CIs of the false-negative rates are large because of the relatively small numbers of children with IBD included in our study and should be interpreted with caution. In this study we focused on IBD, but from the point of view of the primary care physician, it is important to determine whether the symptoms are related to any organic disease. In addition to 17 children with IBD, 3 children were found to have other organic diseases (1 had celiac disease, 1 had reflux esophagitis, and 1 had a solitary rectal ulcer). Children with celiac disease and reflux esophagitis had normal FCal levels and would have been missed if a referral had been solely based on the results of this test.

Cutoff points might need to be higher in primary care to maintain a high negative predictive value.³⁵ Although we found that an increase of the threshold from 50 µg/g to 250 µg/g feces reduced referrals by

Table 2. Baseline Characteristics of the Primary Care Cohort and the Referred Cohort

Characteristic	Main Analysis		Referred Cohort by Origin		Referred Cohort by Referral	
	Primary Care Cohort (n = 114)	Referred Cohort (n = 90)	Primary Care Patients With Red Flag(s) (n = 24) ^a	Specialist Care Patients (n = 66)	Referred by Primary Care Physician (n = 65) ^b	Referred by General Pediatrician (n = 25)
Male, No. (%)	38 (33)	37 (41)	8 (33)	29 (44)	29 (45)	8 (32)
Age, median (IQR), y	9 (6-12)	11 (7-15)	9 (6-14)	12 (7-15)	10 (7-14)	14 (10-15.5)
Presenting symptoms						
Recurrent abdominal pain, No. (%)	88 (77)	58 (64)	16 (67)	42 (64)	38 (59)	20 (80)
Chronic diarrhea, No. (%)	74 (65)	62 (69)	17 (71)	45 (68)	40 (62)	22 (88)
Involuntary weight loss, No. (%)	5 (4) ^c	23 (26)	1 (4)	22 (33)	10 (15)	13 (52)
Rectal blood loss, No. (%)	7 (6)	27 (30)	6 (25)	21 (32)	13 (20)	14 (56)
Family history of IBD, No. (%)	5 (4)	11 (12)	5 (21)	6 (9)	9 (14) ^d	2 (8)
Growth failure, No. (%)	4 (3)	6 (7)	3 (13)	3 (5)	6 (9)	0 (0)
Extraintestinal symptoms, No. (%)	0 (0)	13 (14)	0 (0)	13 (20)	4 (6)	9 (36)
Perianal lesions, No. (%)	7 (6)	13 (14)	7 (29)	6 (9)	9 (14)	4 (17) ^e
Positive blood markers, n/N (%)						
Hemoglobin ^f	1/111 (1)	11/86 (13)	1/24 (4)	10/62 (16)	5/61 (8)	6/25 (24)
CRP (>10 mg/L)	2/110 (2)	10/76 (13)	2/24 (8)	8/52 (15)	5/56 (9)	5/20 (25)
ESR (>20 mm/h)	5/111 (5)	16/83 (19)	5/24 (21)	11/59 (19)	8/59 (14)	8/24 (33)
Platelet count (>450 × 10 ⁹ /L)	4/111 (4)	7/86 (8)	2/24 (8)	5/62 (8)	4/61 (7)	3/25 (12)
Anti-tissue transglutaminase, ^g n/N	0/100	0/72	0/21	0/51	0/55	0/18
≥1 Red flag, ^h No. (%)	29 (25)	68 (76)	24 (100)	44 (67)	43 (66)	25 (100)
Endoscopy, No. (%)	2 (2)	29 (32)	2 (8)	27 (41)	9 (14)	20 (80)
IBD, No. (%)	0 (0)	17 (19)	0 (0)	17 (26)	5/64 (8)	12 (48)

CRP = C-reactive protein; ESR = erythrocyte sedimentation rate; IBD = inflammatory bowel disease; IQR = interquartile range.

^a Five children with red flags were ultimately not seen by a pediatric gastroenterologist: 3 declined because of reduced symptoms, 2 were lost to follow-up.

^b Including primary care physicians who did not participate in this study.

^c Three children had no further weight loss after 3 weeks.

^d Denominator is 64.

^e Denominator is 24.

^f Age and sex specific: aged 4-12 years <7.1 mmol/L; aged 12-18 years: boys <8.1 mmol/L, girls <7.4 mmol/L.

^g Twenty-five children had IgA deficiency.

^h Growth failure, involuntary weight loss, rectal blood loss, family history of IBD, extraintestinal symptoms, perianal lesions, positive blood markers (hemoglobin, CRP, ESR, platelet count).

Table 3. Prevalence of Symptoms, Blood Marker Positivity, and FCal Positivity by Final Diagnosis

Diagnosis	No. (%)	Symptom Positive, ^a No.	Blood Marker Positive, ^b No.	FCal >50 µg/g, No.	Range of FCal, µg/g
Primary care cohort					
Functional gastrointestinal disorder	108 (95)	24	9	12	20-257
Gastroenteritis ^c	5 (45)	0	0	1	20-88
Declined endoscopy	1 (1)	1	0	–	–
Referred cohort					
IBD					
Crohn disease	7 (8)	7	7	6	152-2,823
Ulcerative colitis	8 (9)	7	4	8	53-916
IBD unclassified	2 (2)	2	1	2	79-778
Non-IBD					
Functional gastrointestinal disorder	66 (73)	40	12	10	20-185
Gastroenteritis ^c	3 (3)	1	0	0	20-45
Reflux esophagitis	1 (1)	0	0	0	22
Celiac disease	1 (1)	1	0	0	20
Solitary rectal ulcer	1 (1)	1	0	1	299

FCal = fecal calprotectin; IBD = inflammatory bowel disease.

^a Presence of 1 or more of the following: growth failure, involuntary weight loss, rectal blood loss, extraintestinal symptoms, perianal lesions, family history of IBD.

^b Hemoglobin (4-12 years old <7.1 mmol/L; 12-18 years old: boys <8.1 mmol/L, girls <7.4 mmol/L), C-reactive protein (>10 mg/L), erythrocyte sedimentation rate (>20 mm/h), platelet count (>450 × 10⁹/L).

^c Due to *Salmonella enterica* (0 cases included by primary care physician; 2 cases included by pediatrician), Shiga toxin-producing *Escherichia coli* (STEC) (1 and 0), and *Giardia lamblia* (4 and 1).

Note: One child declined endoscopy and evaluation of red flags at 12 months' follow-up, so the diagnosis was unknown. Nine children without IBD, including 1 child with a solitary rectal ulcer, underwent upper and lower endoscopy, including ileal intubation. The remaining 3 children did not undergo complete endoscopic evaluation for various reasons: the colonoscopy was prematurely terminated because of mucosal bleeding in 1 child with a functional gastrointestinal disorder, but was not repeated because symptoms subsided; 1 child with a functional gastrointestinal disorder underwent colonoscopy only, but not esophagogastroduodenoscopy; and 1 child received a diagnosis of celiac disease by esophagogastroduodenoscopy only.

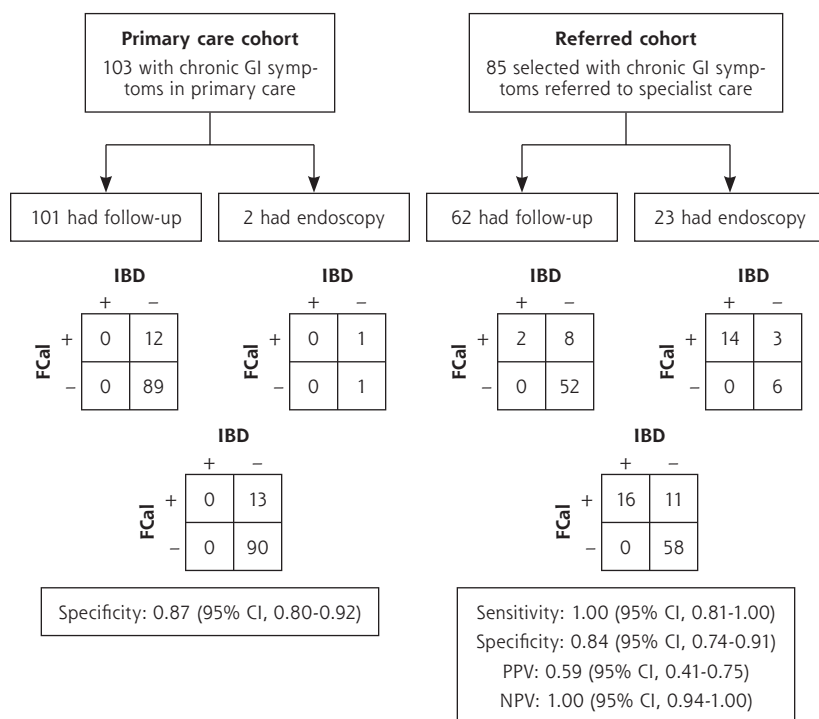
14% (with a drop from 32% to 18% referred), this threshold also led to false-negative results and missed cases of IBD (with an increase from 0% to 18% cases missed) in the referred cohort. A pragmatic approach may be to monitor children with an initial calprotectin value between 50 µg/g and 250 µg/g feces. Children whose symptoms persist and whose calprotectin values remain high can still be referred later. A similar approach has been suggested for adults in primary care, where it was suggested that patients with irritable bowel syndrome and an initial FCal value between 50 µg/g and 150 µg/g feces who have persistent symptoms without treatment should be retested after 3 months.³¹

The specificity of 0.87 that we identified in the primary care cohort is higher than that reported in studies performed in specialist care, where the pooled specificity ranged between 0.68 and 0.76.¹⁵⁻¹⁷ We expected lower specificities because the patient mix was thought to be more diverse in primary care and because calprotectin concentrations increase in conditions such as gastroenteritis.³⁶ Moreover, in the referred cohort, the specificity of FCal was lower in children who underwent endoscopy (0.67) than in those who received clinical follow-up (0.87). The higher specificity might be explained by higher num-

bers of children with functional disease in the primary care cohort and in children with clinical follow-up in the referred cohort. Consequently, the test setting (primary vs specialist care) might affect the specificity of FCal (spectrum bias).

We found a sensitivity of 0.99 in the referred cohort, which is comparable to that reported in other studies performed in specialist care (IBD prevalence of about 60%), where the pooled sensitivity ranged between 0.92 and 0.98.¹⁵⁻¹⁷ The reported sensitivity of FCal in this study might be an overestimation of that in children in whom a primary care physician considers a referral for diagnostic workup. In the subgroup of children referred by a primary care physician (IBD prevalence of 8%), the sensitivity was comparable to that in children referred by a general pediatrician (IBD prevalence of 48%) (0.98 vs 1.00). We therefore assume that spectrum bias did not substantially affect the sensitivity of FCal. Lack of spectrum bias might be explained by the fact that the intestinal inflammation caused by IBD is usually severe enough to increase the calprotectin value to more than 50 µg/g feces, even in the early stages of the disease. A meta-analysis showed that sensitivity remains stable over a range of prevalences and is not substantially influenced by spectrum

Figure 2. Flow charts and contingency tables for the calculation of diagnostic accuracy in the primary care cohort and referred cohort, using the nonimputed data set.



FCal = fecal calprotectin; GI = gastrointestinal; IBD = inflammatory bowel disease; PPV = positive predictive value; NPV = negative predictive value.

Note: The left flow chart shows the specificity of FCal (>50 µg/g) for IBD in the primary care cohort (11 missing values). Specificity of standard follow-up and endoscopy were 0.88 (95% CI, 0.80-0.93) and 0.50 (95% CI, 0.09-0.91), respectively. The right flow chart shows the test characteristics of FCal (>50 µg/g) for IBD in the referred cohort (5 missing values). Sensitivity of the reference standards of follow-up and endoscopy were 1.00 (95% CI, 0.34-1.00) and 1.00 (95% CI, 0.78-1.00), respectively; values for specificity were 0.87 (95% CI, 0.76-0.93) and 0.67 (95% CI, 0.35-0.88), respectively.

Table 4. Test Characteristics at Increasing Calprotectin Cutoff Levels in the Referred Cohort Using the Imputed Data Set (n = 90)

Test Characteristic	Fecal Calprotectin Cutoff		
	>50 µg/g	>100 µg/g	>250 µg/g
Sensitivity (95% CI)	0.99 (0.81-1.00)	0.87 (0.65-0.96)	0.81 (0.58-0.93)
Specificity (95% CI)	0.84 (0.74-0.91)	0.93 (0.84-0.97)	0.98 (0.92-0.99)
PPV (95% CI)	0.60 (0.42-0.76)	0.74 (0.53-0.88)	0.92 (0.69-0.98)
NPV (95% CI)	1.00 (0.94-1.00)	0.97 (0.89-0.99)	0.96 (0.88-0.98)
Referrals avoided, No. (%) ^a	61 (68)	69 (77)	74 (82)
Missed cases of IBD, No. (%) ^b	0 (0)	2 (12)	3 (18)

NPV = negative predictive value; PPV = positive predictive value; IBD = inflammatory bowel disease.

^a Denominator is the 90 children in the referred cohort.

^b Denominator is the 17 children in the referred cohort ultimately given a diagnosis of IBD.

Note: Pretest probability of IBD in this sample was 19%.

Ideally, a study of FCal would include consecutive children with symptoms suggestive of IBD initially evaluated in primary care. This design would be extremely time consuming and costly, however. In order to reflect daily, real-world practice, we included a cohort of children in whom the primary care physician considered a referral for diagnostic workup for chronic gastrointestinal symptoms. We assumed that the children first seen in specialist care were comparable to the selected children with red flags seen in the primary care cohort. We are confident that this assumption is valid as there were only a few differences in characteristics between these groups. Children referred by a primary care physician had a lower probability of IBD (8%) than those referred by a general pediatrician (48%), however. These findings are consistent with what one might expect in the Dutch health care system, where children can consult a pediatrician only after obtaining a referral from their primary care physician, and a pediatric gastroenterologist can be consulted only after a referral from a primary care physician or general pediatrician. Comparable health care systems exist in the United Kingdom, Scandinavia, Canada, New Zealand, and Australia.³⁸

Not all patients received the same reference standard test, which may cause differential verification bias.³² As it is unethical to perform endoscopy in children with a low likelihood of organic gastrointestinal disease, these children received follow-up evaluations over 1 year. An

important aspect for deciding whether the verification leads to biased estimates of accuracy is the length of the follow-up period. We are confident that we identi-

bias.³⁷ A normal FCal level thus could be used to prevent a referral of children with functional symptoms to specialist care.

important aspect for deciding whether the verification leads to biased estimates of accuracy is the length of the follow-up period. We are confident that we identi-

fied all children with IBD, because of the extremely low probability that a child without red flags or indication for endoscopy during 1 year of follow-up has the disease.³⁹ We did not identify new cases of IBD at the end of the follow-up period, even among children who developed red flags. Only 2 children ultimately received a diagnosis of IBD within the follow-up period (both at 6 months). The use of follow-up in children in whom endoscopy is not considered ethical does not correspond very well with the ideal situation that arises when the diagnosis of all cases is determined by endoscopy. Following children during 1 year is the best option given the reality of clinical care, however.²⁰

To recommend a test in a new setting, the diagnostic value of that test needs to be investigated in that setting.⁴⁰ We found that, in selected children in whom a primary care physician considers a referral, FCal has satisfactory discriminatory power between children with and without IBD. Of greater clinical relevance, however, is whether FCal can add to the diagnostic information that is readily available from a thorough history and physical examination.⁴¹ Moreover, the added value of commonly used blood markers should be compared with the added value of FCal. Further research is therefore needed to determine whether FCal should be incorporated into the routine diagnostic evaluation of pediatric patients with chronic gastrointestinal symptoms and red flags in primary care. In addition, research should be performed to evaluate the cost-effectiveness of FCal in primary care.

To read or post commentaries in response to this article, see it online at <http://www.annfammed.org/content/14/5/437>.

Key words: inflammatory bowel disease; calprotectin; child; sensitivity and specificity; primary health care; practice-based research

Submitted October 5, 2015; submitted, revised, March 7, 2016; accepted March 16, 2016.

Funding support: This study was funded by a grant from the Netherlands *Fonds NutsOhra* (grant number 0802-099) and the University Medical Centre Groningen *doelmatigheidsonderzoek* (grant number 689 302).

Disclaimer: The funders did not influence the design, analysis, interpretation of data, writing of the report, or decision to submit the article for publication.

Previous presentations: The results of this study were presented at the European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN), May 6-9, 2015, Amsterdam, the Netherlands, and at NHG-wetenschapsdag, June 19, 2015, Rotterdam, the Netherlands.

Acknowledgment: We would like to thank the primary care physicians and pediatricians for recruiting patients for this study, and we thank all participants for their contribution. The analytical support of Sascha Smit, Erasmus, MC, is highly appreciated. The editorial assistance by Dr Robert Sykes is appreciated.

References

- Gieteling MJ, Lisman-van Leeuwen Y, van der Wouden JC, Schellevis FG, Berger MY. Childhood nonspecific abdominal pain in family practice: incidence, associated factors, and management. *Ann Fam Med*. 2011;9(4):337-343.
- BEACH Program, AIHW General Practice Statistics and Classification Unit. Presentations of abdominal pain in Australian general practice. *Aust Fam Physician*. 2004;33(12):968-969.
- Starfield B, Hoekelman RA, McCormick M, et al. Who provides health care to children and adolescents in the United States? *Pediatrics*. 1984;74(6):991-997.
- Spray C, DeBelle GD, Murphy MS. Current diagnosis, management and morbidity in paediatric inflammatory bowel disease. *Acta Paediatr*. 2001;90(4):400-405.
- Kim SC, Ferry GD. Inflammatory bowel diseases in pediatric and adolescent patients: clinical, therapeutic, and psychosocial considerations. *Gastroenterology*. 2004;126(6):1550-1560.
- Levine A, Koletzko S, Turner D, et al; European Society of Pediatric Gastroenterology, Hepatology, and Nutrition. ESPGHAN revised porto criteria for the diagnosis of inflammatory bowel disease in children and adolescents. *J Pediatr Gastroenterol Nutr*. 2014;58(6):795-806.
- de Ridder L, Rings EH, Escher JC; CBO-werkgroep 'IBD bij kinderen'. [Guideline 'Diagnosis and treatment of inflammatory bowel disease in children']. *Ned Tijdschr Geneesk*. 2010;154:A1898.
- Cabrera-Abreu JC, Davies P, Matek Z, Murphy MS. Performance of blood tests in diagnosis of inflammatory bowel disease in a specialist clinic. *Arch Dis Child*. 2004;89(1):69-71.
- Quail MA, Russell RK, Van Limbergen JE, et al. Fecal calprotectin complements routine laboratory investigations in diagnosing childhood inflammatory bowel disease. *Inflamm Bowel Dis*. 2009;15(5):756-759.
- Henderson P, Casey A, Lawrence SJ, et al. The diagnostic accuracy of fecal calprotectin during the investigation of suspected pediatric inflammatory bowel disease. *Am J Gastroenterol*. 2012;107(6):941-949.
- Gieteling MJ, Bierma-Zeinstra SM, Passchier J, Berger MY. Prognosis of chronic or recurrent abdominal pain in children. *J Pediatr Gastroenterol Nutr*. 2008;47(3):316-326.
- van Tilburg MA, Venepalli N, Ulshen M, Freeman KL, Levy R, Whitehead WE. Parents' worries about recurrent abdominal pain in children. *Gastroenterol Nurs*. 2006;29(1):50-55, quiz 56-57.
- Fagerhol MK, Dale I, Anderson T. Release and quantitation of a leucocyte derived protein (L1). *Scand J Haematol*. 1980;24(5):393-398.
- Røseth AG, Fagerhol MK, Aadland E, Schjønsby H. Assessment of the neutrophil dominating protein calprotectin in feces. A methodologic study. *Scand J Gastroenterol*. 1992;27(9):793-798.
- van Rheenen PF, Van de Vijver E, Fidler V. Faecal calprotectin for screening of patients with suspected inflammatory bowel disease: diagnostic meta-analysis. *BMJ*. 2010;341:c3369.
- Henderson P, Anderson NH, Wilson DC. The diagnostic accuracy of fecal calprotectin during the investigation of suspected pediatric inflammatory bowel disease: a systematic review and meta-analysis. *Am J Gastroenterol*. 2014;109(5):637-645.
- Degraeuwe PL, Beld MP, Ashorn M, et al. Faecal calprotectin in suspected paediatric inflammatory bowel disease. *J Pediatr Gastroenterol Nutr*. 2015;60(3):339-346.
- Fagerberg UL, Lööf L, Myrdal U, Hansson LO, Finkel Y. Colorectal inflammation is well predicted by fecal calprotectin in children with gastrointestinal symptoms. *J Pediatr Gastroenterol Nutr*. 2005;40(4):450-455.
- Holtman GA, Lisman-van Leeuwen Y, Reitsma JB, Berger MY. Noninvasive tests for inflammatory bowel disease: a meta-analysis. *Pediatrics*. 2016;137(1):1-11.
- Knottnerus JA, Dinant GJ. Medicine based evidence, a prerequisite for evidence based medicine. *BMJ*. 1997;315(7116):1109-1110.

21. Holtman GA, Lisman-van Leeuwen Y, Kollen BJ, et al. Challenges in diagnostic accuracy studies in primary care: The fecal calprotectin example. *BMC Fam Pract*. 2013;14:179.
22. Lewis SJ, Heaton KW. Stool form scale as a useful guide to intestinal transit time. *Scand J Gastroenterol*. 1997;32(9):920-924.
23. Oord T, Hornung N. Fecal calprotectin in healthy children. *Scand J Clin Lab Invest*. 2014;74(3):254-258.
24. Gisbert JP, McNicholl AG. Questions and answers on the role of faecal calprotectin as a biological marker in inflammatory bowel disease. *Dig Liver Dis*. 2009;41(1):56-66.
25. Pels LP, Van de Vijver E, Waalkens HJ, et al. Slow hematological recovery in children with IBD-associated anemia in cases of "expectant management." *J Pediatr Gastroenterol Nutr*. 2010;51(6):708-713.
26. Van de Vijver E, Schreuder AB, Cnossen WR, Muller Kobold AC, van Rheeën PF, North Netherlands Pediatric IBD; North Netherlands Pediatric IBD Consortium. Safely ruling out inflammatory bowel disease in children and teenagers without referral for endoscopy. *Arch Dis Child*. 2012;97(12):1014-1018.
27. Mack DR, Langton C, Markowitz J, et al; Pediatric Inflammatory Bowel Disease Collaborative Research Group. Laboratory values for children with newly diagnosed inflammatory bowel disease. *Pediatrics*. 2007;119(6):1113-1119.
28. Canani RB, de Horatio LT, Terrin G, et al. Combined use of noninvasive tests is useful in the initial diagnostic approach to a child with suspected inflammatory bowel disease. *J Pediatr Gastroenterol Nutr*. 2006;42(1):9-15.
29. Bremner A, Roked S, Robinson R, Phillips I, Beattie M. Faecal calprotectin in children with chronic gastrointestinal symptoms. *Acta Paediatr*. 2005;94(12):1855-1858.
30. Carroccio A, Iacono G, Cottone M, et al. Diagnostic accuracy of fecal calprotectin assay in distinguishing organic causes of chronic diarrhea from irritable bowel syndrome: a prospective study in adults and children. *Clin Chem*. 2003;49(6 Pt 1):861-867.
31. Pavlidis P, Chedgy FJ, Tibble JA. Diagnostic accuracy and clinical application of faecal calprotectin in adult patients presenting with gastrointestinal symptoms in primary care. *Scand J Gastroenterol*. 2013;48(9):1048-1054.
32. Naaktgeboren CA, de Groot JA, van Smeden M, Moons KG, Reitsma JB. Evaluating diagnostic accuracy in the face of multiple reference standards. *Ann Intern Med*. 2013;159(3):195-202.
33. Vink G, Frank LE, Pannekoek J, Buuren S. Predictive mean matching imputation of semicontinuous variables. *Stat Neerl*. 2014;68(1):61-90.
34. Marshall A, Altman DG, Holder RL, Royston P. Combining estimates of interest in prognostic modelling studies after multiple imputation: Current practice and guidelines. *BMC Med Res Methodol*. 2009;9:57.
35. Logan R. Faecal calprotectin for the diagnosis of inflammatory bowel disease. *BMJ*. 2010;341:c3636.
36. Chen CC, Huang JL, Chang CJ, Kong MS. Fecal calprotectin as a correlative marker in clinical severity of infectious diarrhea and usefulness in evaluating bacterial or viral pathogens in children. *J Pediatr Gastroenterol Nutr*. 2012;55(5):541-547.
37. Leeftang MM, Rutjes AW, Reitsma JB, Hooft L, Bossuyt PM. Variation of a test's sensitivity and specificity with disease prevalence. *CMAJ*. 2013;185(11):E537-E544.
38. Thomson S, Osborn R, Squires D, Reed SJ. International profiles of health care systems 2011. The Commonwealth Fund. <http://www.commonwealthfund.org/publications/fund-reports/2011/nov/international-profiles-of-health-care-systems-2011>. Published Nov 2011. Accessed 2011.
39. Jellema P, van Tulder MW, van der Horst HE, Florie J, Mulder CJ, van der Windt DA. Inflammatory bowel disease: a systematic review on the value of diagnostic testing in primary care. *Colorectal Dis*. 2011;13(3):239-254.
40. Moons KG, Biesheuvel CJ, Grobbee DE. Test research versus diagnostic research. *Clin Chem*. 2004;50(3):473-476.
41. Moons KG, de Groot JA, Linnet K, Reitsma JB, Bossuyt PM. Quantifying the added value of a diagnostic test or marker. *Clin Chem*. 2012;58(10):1408-1417.