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# The effect of enhanced feedback and brief educational reminder messages on laboratory test requesting in primary care: A cluster randomised trial (ISRCTN06490422). Authors: Ruth E Thomas, PhD Research Fellow Health Services Research Unit University of Aberdeen Aberdeen, UK. Bernard Lewis Croal, MD Consultant/Senior Lecturer Department of Clinical Biochemistry NHS Grampian Aberdeen, UK Craig Ramsay, PhD Senior Statistician Health Services Research Unit University of Aberdeen Aberdeen, UK.

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#### Summary

#### Background:

Laboratory services play an important role in screening, diagnosing and managing patients within primary care. Use of laboratory tests is increasing and evidence suggests that overuse is a feature of this increase. Our aim was to assess the effect of two interventions on the number of laboratory tests requested by primary care physicians.

#### Methods:

The study was a cluster randomised controlled trial using a 2x2 factorial design involving 85 primary care practices, (370 general practitioners) who request all laboratory tests from one regional centre. The interventions were: quarterly feedback of practice requesting rates for nine laboratory tests, enhanced with educational messages; brief educational reminder messages added to the test result reports for nine laboratory tests. The primary outcome was the number of targeted tests requested by primary care practices during the 12 months of the intervention.

#### Findings:

Practices that received either or both the enhanced feedback and the reminder messages were significantly less likely than the control group to request the targeted tests in total. (Main effects analyses: enhanced feedback, odds ratio 0.87, 95% CI 0.81 to 0.94; reminder messages, odds ratio 0.89, 95% CI 0.83 to 0.93). The effect of the interventions varied across the targeted tests individually, although the general pattern showed a reduction in the number of tests requested for both interventions. Neither intervention was consistently better than the other.

#### Interpretation:

Enhanced feedback of requesting rates and brief educational reminder messages, alone and in combination are effective strategies for reducing test requesting in primary care. Both strategies are feasible within most laboratory settings.

#### Keywords

Randomised controlled trial, reminder systems, audit and feedback, primary care, laboratory medicine, test requests

#### Introduction

Laboratory services play an important role in screening, diagnosing and managing patients within primary care. Use of laboratory tests has increased substantially in recent years(1, 2) and a survey(3) of United Kingdom (UK) laboratories showed an 83% increase in primary care test requesting between 2000 and 2004.

Potentially there are many reasons for this increase including the development of new useful tests, and the impact of new guidelines and contracts(4, 5). However, previous studies(6-8) provide grounds for thinking that unnecessary test ordering may be a component of this increase.

Unnecessary test requesting is not only a burden on laboratory resources, but may also be detrimental for patients as it can lead to subsequent unnecessary investigation and treatment of healthy individuals following false positive results(9-12). In addition, it is an inappropriate use of the finite resources available for healthcare provision as a whole (1).

The effectiveness of strategies to change health professional practice in general and test requesting in particular is varied(1, 13-18). Previous reviews(10, 13-15) have suggested that audit and feedback of test ordering rates, educational messages, test request form changes, reminders and computer decision support are all potentially effective methods of changing test ordering behaviour. In a systematic review (13) focusing on studies evaluating methods to improve diagnostic test requesting, including 49 studies with a control group, the majority of interventions evaluated claimed effectiveness, however, conclusions are limited by methodological flaws such as lack of a randomised comparison group (in 41 of the 49 studies). In addition, few of the studies evaluated the effectiveness of these interventions within a primary care setting. In a recent systematic review, which included 85 randomised

trials, audit and feedback (19) has been shown to have small to moderate effects on health professional practice, however, the evidence within primary care test requesting is sparse, as only two of the trials evaluated the effect of feedback on laboratory test requesting within primary care. The authors concluded that their review does not provide support for unevaluated use of audit and feedback. Finally, current systematic reviews suggest that "single" intervention strategies may be as effective as multiple complex interventions in changing health professional practice(17, 19).

At the time of planning the current study, we had recently completed a cluster randomised trial of two strategies to reduce requests for lumbar spine and knee x-rays from primary care(20). We found that whilst simple, comparative audit and feedback of request rates had no discernable effects, the provision of educational reminder messages led to a 20% relative reduction in requests. These interventions were considered relatively easy for radiology departments to implement. In view of the limitations in the current evidence we undertook a cluster trial (randomising by practice to minimise contamination across practitioners within practices) using similar interventions to those used in our previous study to determine their effect on laboratory tests requests from primary care.

#### Methods

#### Setting and Population

The study took place in the north east of Scotland, UK, and involved all 85 primary care practices (approximately 370 general practitioners) in the area covered by NHS Grampian. The clinical laboratories based within Aberdeen Royal Infirmary provided all laboratory services for these primary care practices.

#### Interventions

Nine laboratory tests were chosen as targets for intervention (Table 1) due to their perceived limited value for some patient subgroups within primary care, their impact on laboratory workload, and to reflect the range of laboratory services. Educational messages describing inappropriate use were developed for each of the targeted tests (Table 2 shows examples of these educational messages).

The tests were chosen and messages developed during a series of meetings with senior clinicians from four laboratory disciplines within NHS Grampian laboratories (Clinical Biochemistry, Haematology, Immunopathology; and Microbiology). The content and format of the educational messages and feedback were reviewed and commented on by primary care physicians and other laboratory clinicians.

In general, the evidence base for the appropriate and inappropriate use of laboratory tests is poor. Therefore the majority of the target tests were chosen and the content of the educational messages were developed using consensus expert opinion. However there is evidence for some of the targeted tests and their associated educational messages, for example: the use of the tumour markers Carcino-Embryonic Antigen (CEA) and Carbohydrate antigen – 125 (CA125) are not appropriate for diagnostic or screening use, and

should be reserved for following up known patients with relevant cancers(21). In addition, the tests are not licensed for these diagnostic or screening indications(22). Similarly, Follicle Stimulating Hormone (FSH) has been shown to be of limited value and is potentially confusing when used to assess menopausal status(23).

#### Enhanced feedback:

The feedback consisted of a six-sided colour booklet presenting graphs of practice level data for each of the nine targeted tests and for each laboratory discipline as a whole. Each graph showed tests requesting rates over the previous three years (number per 10,000 patients per six months) for the practice compared to the regional rates.

The feedback was enhanced with the educational messages (examples shown in Table 2) which were included alongside the graphs for each of the targeted tests. The booklets were posted to each general practitioner within each intervention group practice on four occasions (updated every three months from the start of the intervention period). Examples of the feedback booklet are available from the corresponding author.

#### Reminder messages:

The brief educational messages (Table 1) were added as reminders to the test result reports sent to the requesting practice. The laboratory information system was programmed to recognise the relevant cues for each of the targeted tests (see Table 1) and automatically add the brief educational reminder messages to the relevant printed and electronic test result reports. The messages were activated every time the cue occurred and were presented at the same time as the test result. The reminders messages were intended to influence future requests for the targeted tests.

The interventions were introduced in February 2002 for 12 months.

#### Study Design

The study design was a cluster randomised controlled trial using a 2 x 2 factorial design(24). All practices (85) were allocated to one of four groups: control, enhanced feedback alone, reminder messages alone or both enhanced feedback and reminder messages, using minimisation(25) based on practice list size (the number of people registered with the practice: 0 to 3000, 3001 to 8000 and over 8000), practice setting (rural or urban) and training status, (training or non training practice). Practice list size was predictive of the number of test requests. Practice setting and training status were chosen as factors that may influence the effect of the interventions. They potentially reflect differing populations, access to health care services, and receptiveness to practice innovation. Allocation was restrained to the enhanced feedback groups for one practice that had taken part in the piloting of the feedback booklet.

A statistician independent of the research team and blinded to the identity of the practices performed the minimization at the end of the pre intervention period.

#### Data collection

The numbers of each targeted test requested per practice (accurate individual physician level requesting data was not available) were collected for the 12 months before (pre intervention) and the 12 months of the intervention period. The data were downloaded from the NHS Grampian laboratory information system. The laboratory personnel who processed the requests were blinded to intervention group status.

#### Sample size

The sample size was based upon a cluster level analysis adjusted for the test ordering patterns in the 12 months prior to the intervention using analysis of covariance. Data on the number of tests requested per practice in Grampian over two years (1998 to 1999) showed a high correlation between the number of requests per practice per year for Thyroid Stimulating Hormone (TSH) and Follicle Stimulating Hormone (FSH) (> 0.9). Using this information, a study with 90 practices had 80% power at 5% significance to detect a relative reduction of 12% in the number of TSH requests assuming a pre intervention level of 129 requests per 1000 of the population (obtained from the 1999 figures). This sample size was for the main effects only and assumed no interaction between the two interventions. Similar detectable differences were obtained for FSH. We did not have pre trial data for the other targeted tests.

#### Statistical analysis

A main effects analysis of covariance, adjusting for pre intervention yearly test requests and minimisation factors was performed on the natural logarithm of the number of test requests per year (sum of the targeted tests). The model fitting strategy was to test the minimisation variables (design) followed by the logarithm of the pre intervention test requests (covariate) and lastly two binary variables (intervention) corresponding to the enhanced feedback and reminder messages main effects were included. Any design or covariate variables failing to achieve statistical significance at the 5% level were excluded. The resultant intervention main effects were transformed onto an odds ratio (OR) scale with corresponding 95% confidence intervals. Analyses of each targeted test were performed similarly. Possible intervention interaction effects (i.e., non-additive effect of both interventions) were also investigated by including an interaction term in the above models.

Subgroup analysis

It was decided a priori to investigate whether larger effects of the interventions were observed for practices reporting a higher pre intervention level of test requests. The models tested were identical to those used for the primary outcomes except for the addition of an interaction term between pre intervention request level and the intervention group. In addition, to test for a reduction in the spread of test requests, Levene's test for equality of variances was applied between the pre intervention and intervention phases.

The study was reviewed by the Local Research Ethics Committee and Grampian Area Medical Committee, General Practice Sub-Committee.

Role of funding source

The funders of this study, other than the initial peer review process prior to funding, did not have any involvement in study design; in the collection, analysis, and interpretation of data; in the writing of the report or in the decision to submit the paper for publication.

#### Results

It was planned to recruit the ninety practices within NHS Grampian, however, four of these practices merged with others prior to the start of the study. Two practices merged with each other during the pre intervention period. Eighty-five practices were thus allocated to the interventions and included in the trial (Figure 1). No practices refused to take part and every practice contributed data to both 12 month pre intervention and intervention periods (Figure 1). Table 3 shows that the practice characteristics were similar across the intervention groups.

Across the total targeted tests, general practices receiving either the enhanced feedback (OR 0.87, 95% CI 0.81 to 0.94) or the brief educational reminder messages (OR 0.89, 95% CI 0.83 to 0.83) were significantly less likely to request targeted tests (Table 4). There was no evidence of an interaction between interventions (OR 0.98, 95% CI 0.84 to 1.14) so the combined effect was equivalent to a 22% reduction (OR 0.78, 95% CI 0.71 to 0.85) in total targeted test requests.

The effect of the interventions varied across each of the targeted tests individually, although the general pattern showed a reduction in the number of tests requested in the intervention groups (Table 4). Practices that received enhanced feedback reduced test ordering for all nine tests, which reached statistical significance for four tests (AAS, FSH, TSH, B12). Practices receiving the brief educational reminder messages also demonstrated reductions in test requests across eight tests, which reached statistical significance for three tests (CEA, TSH, B12).

For eight of the tests, there was no statistical evidence of an interaction effect (median [IQR] size of interaction OR was 0.99 [0.87, 1.23]). For Ferritin, the interaction effect was significant

(OR 0.60, 95% CI 0.42 to 0.87), neither intervention had a significant effect alone, but in combination there was a reduction in test requests (OR 0.58, 95% CI 0.36 to 0.92).

Subgroup analyses exploring the possibility of effect moderation by pre intervention number of test requests provided no evidence that the effect was increased or decreased by the number of pre intervention test requests (enhanced feedback OR 1.05, 95% CI 0.95 to 1.15; reminder messages OR 0.96, 95% CI 0.87 to 1.05). There was no indication that the variability of test requests was changed in the intervention period (Levene's Test, F=0.032, p-value=0.859).

#### Discussion

This study evaluated the effect of enhanced feedback and brief educational reminder messages on nine tests requested from primary care for one year. Enhanced feedback alone and brief educational reminder messages alone each achieved moderate(19) reductions of around 10% in the number of requests in total, whilst the effect of the interventions in combination showed larger reductions of >20 % in the total targeted tests requested.

The effects of the interventions varied across the individual targeted tests but showed a general pattern of reduction in requesting which was particularly consistent when both enhanced feedback and brief educational reminder messages were used together. Neither intervention alone appeared consistently better.

Our study was a rigorous design with all allocated practices included in the analysis. It involved all primary care practices in one region of Scotland (a previous study(20) does not give us any reason to suspect that these practices' requesting of investigations differs from other UK regions).

The targeted tests, although only representing a small proportion of test requesting in primary care, when used inappropriately, potentially have important implications for further diagnostic testing and specialist referral. TSH accounted for the majority of the targeted tests requested, and therefore had a major influence on the overall combined result. However, there were statistically significant reductions in requesting across the range of laboratory tests allowing confidence in the generalisability of the results(26). Our results are particularly impressive when considered in the context of the year on year increases in requesting of the targeted tests observed(2, 3) prior to the intervention period.

Factorial designs are efficient for evaluating two or more interventions when they act independently of each other(24, 27). Our results, except for ferritin, do not suggest an interaction between the feedback and reminders, that is, the combined effect of the feedback and reminders was additive (the sum of the effect for feedback plus the effect for reminders) rather than synergistic. However, as the study was not powered to detect interactions we cannot be sure that they were not present; to fully investigate whether these interventions act independently a much larger trial would be needed.

Due to mergers of primary care practices the original sample size of 90 was not achieved with subsequent loss of power. However, adjusting for pre intervention minimisation factors increased the power of the study and we were able to identify our pre defined clinically important differences in TSH and FSH requesting.

We measured all requests for each test over the study period and we assume that the effects observed are due to reductions in unnecessary requesting. Ideally, we would have included a measure of the proportion of requests that were unnecessary, however data to determine this were not available. An economic analysis was not conducted as this would require further data on the appropriateness of the tests requested and subsequent effects on patient wellbeing for example health effects, reassurance gained or lost.

Whilst the general pattern across the individual targeted tests is a reduction in requesting our results suggest that the effect of the two interventions alone may vary within and across the individual tests. The reductions observed do not all reach statistical significance possibly reflecting the higher variability, low number and lack of power for some of the tests, for instance, CEA. However, across and within tests of similar volume (e.g. FSH, Ferritin and HPS) effectiveness seems to vary. Our outcome measures (all requests for a targeted test)

may be too general to detect changes related specifically to the message for some of the tests. It is also possible that the potential for changes in requesting related to the educational message differs across the individual tests. Requesting a test is a result of complex considerations, motivations and behaviours(28, 29) that may be conflicting(30) and may differ for each of the targeted tests and conditions. The brief educational reminder messages and enhanced feedback may address different considerations and motivations and thus their effectiveness may vary across the individual targeted tests. Further understanding of these motivations and behaviours and how they can be changed is needed to inform and optimise the choice and development of interventions for future trials.

Our study has shown effects that are as large as those in a recent robust study of test requesting using a multifaceted strategy, arguably more complex to deliver and maintain(31). This lends further support to emerging evidence that complex multifaceted interventions may not be superior in effectiveness(17, 19).

Across two similar trials, reminder messages have consistently shown reductions in targeted radiology(20) and total targeted laboratory tests. The reminder messages in both these studies influenced future requests. The advent of computerised order communication systems with interactive real-time interrogation means there will be the opportunity to influence laboratory test requests as they are made. It has been suggested previously (10, 14) that the type of reminder influences their effectiveness and therefore it would be important to investigate the effectiveness of these real time reminders compared to those designed to influence future requests.

The six monthly feedback tested in the radiology requests trial(20) was shown to be ineffective whilst the quarterly feedback enhanced with educational messages tested in this

study showed moderate effects for the total tests, generally consistent with those observed for audit and feedback interventions elsewhere(19). It seems that differences in the presentation of feedback, such as enhancement with educational messages or frequency of the feedback and again, the targeted condition may influence the effectiveness of audit and feedback interventions demonstrating the need for further research testing appropriate permutations head-to-head to inform design of effective audit and feedback interventions.

At the time of planning this study, a control group or usual practice group was necessary as there was insufficient evidence of the effectiveness of reminders or audit and feedback on diagnostic behaviour. Current evidence(17) suggests a need for head to head comparisons of interventions, however, taking into account the primary research question and sample size requirements we would still consider incorporating a usual practice group within a study as it gives an estimate of the absolute effect size.

We studied the effects of these interventions over one year. Using a non-experimental design Winkens et al(32) suggest a persistent effect of feedback on test requesting over nine years in routine practice and Tierney et al(33) in their trial of reminders about test charges suggests the effects do not persist after stopping reminders. To inform the use of these interventions in routine practice their longer term effects on test requesting need to be rigorously evaluated. In addition, the consequences of increasing the number of targeted tests needs to be investigated as this may influence the potency and sustainability of the interventions.

In conclusion, we have identified and rigorously evaluated two strategies that seem feasible within most laboratory settings and together appear to be consistently effective across a range of laboratory tests.

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#### Contributorship

Bernard Croal and Jeremy Grimshaw conceived the project. All authors contributed to the design of the study. Ruth Thomas and Bernard Croal were responsible for running the project. Craig Ramsay was responsible for the statistical analyses. All authors interpreted the data and findings. Ruth Thomas wrote the first draft of the manuscript, all authors commented on it and all further revisions. Bernard Croal and Ruth Thomas are guarantors for the paper.

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## Conflicts of interest statement

None of the authors have any conflicts of interest

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reports

Targeted laboratory tests, brief educational messages and cues for activating addition of messages as reminders to results Table 1

Targeted Test	Brief educational message	Added to test	Cue to activate reminder
		result report for:	message
Autoantibody screen (AAS)	Autoantibody 'screen' requesting is inappropriate for AAS	AAS	AAS request
	investigation of non-specific illness. Requests should be		
	test-specific.		
Carbohydrate antigen - 125	CA125 should not be used to screen, diagnose or exclude CA-125	CA-125	CA-125 request
(CA125)	malignancy.		
Carcino Embryonic	CEA should not be used to screen, diagnose or exclude CEA	CEA	CEA request
Antigen (CEA)	malignancy.		
Ferritin	Ferritin measurement is generally unnecessary in	in Full Blood Count	FBC request result with a mean
	caucasians with hypochromic microcytic anaemia as the	(FBC)	cell volume (MCV) < 80 fl
	underlying cause is almost always iron deficiency.		
Follicle stimulating	In general, FSH testing is of limited value in the	FSH	FSH request
hormone (FSH)	assessment of menopausal status in women over 40 years		

test requests.	
care	
primary care	
for	
reminders	
k and	
Feedback	

HPS request			IgE request			TSH request		B12 request		FBC result with a MCV > 95 fl		
HPS			IgE			TSH		B12		FBC		
Helicobacter Pylori serology should not be used to assess	the efficacy of antibiotic eradication therapy as antibody	levels may remain positive for some time after eradication	General allergen 'screening' is unhelpful. Allergen testing	requests should instead be specific as directed by the	history.	Thyroid Function tests are not indicated as a screening	procedure in young, clinically euthyroid patients.	B12 levels are of no value and should therefore not be	requested in patients undergoing parenteral B12 therapy.	Macrocytosis without anaemia is unlikely to be due to B12	deficiency (thus should not be requested). Thyroid or liver	function tests may be helpful.
Helicobacter pylori serum	(HPS)		Imunoglobulin E (IgE)			Thyroid stimulating	hormone (TSH)	Vitamin B12 (B12)				

### Table 2 - Examples of educational messages used in the feedback reports

Carcino-Embryonic Antigen (CEA) has been found to be raised in the serum of some patients with gastrointestinal malignancy, notably colonic cancer. It is therefore a useful marker for use in monitoring treatment and in detecting recurrence. CEA should not however be used to screen, diagnose or exclude malignancy. It can also be found to be raised in many other benign conditions, especially gastrointestinal disorders. False positive and false negative results can therefore lead to unnecessary further investigation or false reassurance.

Carbohydrate antigen - 125 (CA125) has been found to be raised in the serum of some patients with malignancy, notably ovarian cancer. It is therefore a useful marker for use in monitoring treatment and in detecting recurrence. CA-125 should not however be used to screen, diagnose or exclude malignancy. It can also be found to be raised in many other malignancies and benign conditions, notably gastrointestinal disorders. False positive and false negative results can therefore lead to unnecessary further investigation or false reassurance.

Follicle Stimulating Hormone (FSH) is released by the pituitary gland and acts to stimulate sex hormone production and reproductive processes. In general, FSH testing is of limited value in the assessment of menopausal status in women over 40 years of age, and so should not be requested for this purpose. Menopausal/Peri-menopausal status is best confirmed retrospectively based on clinical symptoms, signs and frequency or absence of menstruation. Biochemical measurement adds little to this classification, and may mislead.

Ferritin measurement is generally unnecessary in caucasians with hypochromic microcytic anaemia as the underlying cause is almost always iron deficiency. While a source of potential blood loss should always be sought in cases of suspected iron deficiency, treatment with iron replacement is best monitored by clinical response and periodic haemoglobin measurement rather than resorting to iron studies - including ferritin assessment.

Imunoglobulin E (IgE) Raised total IgE and positive specific IgE tests are found in around 40% of the population but are only clinically relevant in around 20% (high false positive rate). General allergen "screening" is unhelpful. Allergen testing requests should instead be specific as directed by the history. IgE tests are not a substitute for adequate

history taking. The significance of any result requires careful consideration in the context of the clinical problem.

Helicobacter Pylori serology (HPS) and breath tests are highly sensitive markers of infection associated with symptomatic peptic ulcer disease. HPS should not be used to screen asymptomatic individuals, even if they have a family history of peptic ulcer disease. HPS should not be used to assess the efficacy of antibiotic eradication therapy as antibody levels may remain positive for some time after eradication. A Helicobacter pylori breath test should be instigated if peptic ulcer symptoms persist, so as to confirm/exclude continued infection.

Table 3 Characteristics of study practices

	,							
	Control	trol	Feed	Feedback* only	Rem	Reminders* only	Both	
	N=20	07	N=22	2	N=22	2	N=21	1
	Ħ	(%)	u	%	Z	%	ц	%
Characteristics								
Training practice	8	(40%)	10	(45%)	10	(45%)	6	(43%)
Non training practice	12	(%09)	12	(25%)	12	(25%)	12	(%25)
Location of practice								
Rural	10	(%05)	11	(%09)	10	(45%)	10	(48%)
Urban	10	(20%)	11	(%09)	12	(25%)	11	(52%)
Practice list size								
<3000 patients	4	(50%)	7	(32%)	^	(32%)	9	(29%)
3001-8000 patients	6	(45%)	6	(41%)	8	(36%)	∞	(38%)
>8000 patients	^	(35%)	9	(27%)	7	(32%)	^	(33%)

\*Feedback = quarterly feedback of practice requesting rates enhanced with educational messages, Reminders = brief educational reminder messages added to test result reports

Table 4 Effect of the interventions on total and individual targeted tests

4	P-value	
Effect of interventions*	Odds Ratio (95% CI) P-value	
Intervention period	Median number [IQR]† per	10,000 patients per practice
Pre intervention period	Median number [IQR] † per	10,000 patients per practice

Downstandound		Main effects Feedback <sup>‡</sup> (0.81, 0.94) <0.001	Fotal	argeted tests	<0.001	(0.81, 0.94)	0.87	[726, 2057] [575, 1818] [719, 1590] [362, 1515]	1226 1079 1317 1041	[783, 1804] [601, 1954] [688, 1726] [492, 1749]	1071 1233 1329 1166	roup tonly ston
Feedback <sup>‡</sup> (0.81, 0.94)	Feedback <sup>‡</sup> (0.81, 0.94)		up       1071       [783, 1804]       1226         nly group       1233       [601, 1954]       1079         only group       1329       [688, 1726]       1317	Control group       1071       [783, 1804]       1226         Feedback* only group       1233       [601, 1954]       1079         Reminders* only group       1329       [688, 1726]       1317				[362, 1515]	1041	[492, 1749]	1166	Both group
p	P 1166 [492, 1749] 1041 [362, 1515] 0.87 (0.81, 0.94)	1166 [492, 1749] 1041	1071     [783, 1804]     1226       1233     [601, 1954]     1079	Control group       1071       [783, 1804]       1226         Feedback* only group       1233       [601, 1954]       1079				[719, 1590]	1317	[688, 1726]	1329	Reminders <sup>‡</sup> only group
st only group       1329 [688, 1726]       1317 [719, 1590]         ip       1166 [492, 1749]       1041 [362, 1515]         Feedbackt       0.87 (0.81, 0.94)	st only group 1329 [688, 1726] 1317 [719, 1590]  ip 1166 [492, 1749] 1041 [362, 1515] 0.87 (0.81, 0.94)	only group 1329 [688, 1726] 1317 1166 [492, 1749] 1041	1071 [783, 1804] 1226	Control group 1071 [783, 1804] 1226				[575, 1818]	1079	[601, 1954]	1233	Feedback <sup>‡</sup> only group
tonly group 1233 [601, 1954] 1079 [575, 1818] 1329 [688, 1726] 1317 [719, 1590] 1317 [719, 1590] 1317 [719, 1591] 1317 [719, 1515] 1317 [719,	tonly group 1233 [601, 1954] 1079 [575, 1818] 1329 [688, 1726] 1317 [719, 1590] 1317 [719, 1590] 1329 [492, 1749] 1041 [362, 1515] 0.87 (0.81, 0.94)	nly group       1233 [601, 1954]       1079         only group       1329 [688, 1726]       1317         1166 [492, 1749]       1041		otal				[726, 2057]	1226	[783, 1804]	1071	Control group

Individual

Autoantibody screen (AAS)

Feedback and reminders for primary care test requests.

				0.002	0.599						0.726	0.537
				(0.67, 0.91)	(0.82, 1.12)						(0.65, 1.36)	(0.61, 1.30)
				0.78	96.0						0.94	0.89
[13, 64]	[20, 49]	[18, 63]	[10, 66]				[9, 25]	[3, 19]	[4, 23]	[4, 19]		
41	33	36	31				16	11	12	11		
[13, 68]	[26, 75]	[25, 77]	[18, 97]				[6, 22]	[3, 22]	[4, 30]	[1, 24]		
41	49	38	29				13	∞	6	6		
Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders	Carbohydrate antigen - 125 (CA125)	Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders

Carcino Embryonic Antigen (CEA)

Feedback and reminders for primary care test requests.

				0.177	0.041						0.489	0.746
				(0.52, 1.13)	(0.44, 0.98)						(0.71, 1.18)	(0.81, 1.34)
				0.76	99.0						0.91	1.04
[4, 33]	[2, 15]	[3, 25]	[2, 19]				[49, 137]	[23, 106]	[45, 132]	[16, 87]		
11	6	10	9				79	09	85	28		
[1, 16]	[3, 16]	[5, 26]	[2, 33]				[34, 102]	[43, 95]	[48, 115]	[35, 119]		
10	10	18	13				72	72	98	73		
Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders	Ferritin	Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback <sup>§</sup>	Reminders§

Follicle stimulating hormone (FSH)

Feedback and reminders for primary care test requests.

				0.020	0.559						0.589	0.293
				(0.75, 0.98)	(0.85, 1.09)						(0.74, 1.14)	(0.76, 1.09)
				98.0	96.0						0.95	0.91
[27, 122]	[23, 96]	[30, 92]	[30, 85]				[36, 98]	[21, 104]	[38, 98]	[20, 117]		
77	27	22	49				56	99	92	63		
[41, 121]	[33, 128]	[42, 105]	[43, 132]				[30, 107]	[26, 120]	[38, 99]	[40, 117]		
62	62	77	81				74	29	70	75		
Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders	Helicobacter pylori serum (HPS)	Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders

Imunoglobulin E (IgE)

Feedback and reminders for primary care test requests.

			0.471	0.909						0.005	0.001
			(0.73, 1.16)	(0.79, 1.24)						(0.84, 0.97)	(0.83, 0.95)
			0.92	0.99						06.0	0.82
[9, 34] [10, 36]	[13, 25]	[7, 38]				[552, 1466]	[432, 1359]	[490, 1250]	[287, 1077]		
24	21	23				795	802	891	800		
[7,34] [11,44]	[11, 39]	[18, 46]				[515, 1329]	[476, 1412]	[476, 1338]	[392, 1277]		
20	18	30				750	829	961	891		
Control group Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders	Thyroid stimulating hormone (TSH)	Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders

Vitamin B12 (B12)

Feedback and reminders for primary care test requests.

				0.041	0.043
				(0.66, 0.99)	(0.66, 0.99)
				0.81	0.81
[13, 52]	[15, 48]	[15, 45]	[10, 40]		
34	23	29	19		
[17, 40]	[18, 53]	[21, 49]	[10, 52]		
25	32	37	31		
Control group	Feedback only group	Reminders only group	Both group	Main effects Feedback	Reminders

\*Based upon analysis of covariance adjusting for practice list size and pre intervention request level

# Feedback = quarterly feedback of practice requesting rates enhanced with educational messages, Reminders = brief educational reminder messages added to test result reports

§ Interaction term was statistically significant (p=0.007), effects reported are for main effects and interaction in model

<sup>†</sup> IQR = interquartile range

Figure 1 Trial profile - see separate file