

Availability of cancer decision-support tools: A cross-sectional survey of UK primary care

Sarah Price, Anne Spencer, Antonieta Medina-Lara, Willie Hamilton on behalf of the Exeter/Bangor/Leeds Cancer Diagnostic Support in Primary Care HTA Group

Contact details:

Sarah Price University of Exeter Medical School, Exeter, EX1 2LU, S.J.Price@exeter.ac.uk

Anne Spencer University of Exeter Medical School, Exeter, EX1 2LU, a.e.spencer@exeter.ac.uk

Antonieta Medina-Lara University of Exeter Medical School, Exeter, EX1 2LU, A.Medina-Lara@exeter.ac.uk

Willie Hamilton University of Exeter Medical School, Exeter, EX1 2LU, W.Hamilton@exeter.ac.uk

Corresponding author: Dr Sarah J. Price, Room 1.20 College House, University of Exeter Medical School, St Luke's campus, Exeter, EX1 2LU; email S. Price S.J.Price@exeter.ac.uk; tel: +44-1392-726347

Word count: 2,858 (excluding tables, figures and references)

Abstract (250/250 words)

Background Decision-support tools quantify the risk of undiagnosed cancer in symptomatic patients, and may help general practitioners (GPs) when making referrals.

Aims: To quantify the availability and use of cancer decision-support tools (QCancer[®] and Risk Assessment Tools). To explore the association between tool availability and two-week-wait referrals for suspected cancer.

Design and setting: Cross-sectional postal survey in UK primary care.

Methods: 4,600 GPs from a random sample of 975 UK general practices were invited to participate. Outcome measures included the proportions of UK general practices where: (1) cancer decision-support tools are available, and (2) at least one GP uses the tool. Weighted least-squares linear regression with robust errors tested the association between tool availability and number of two-week-wait referrals, adjusting for practice size, sex, age and index of multiple deprivation.

Results: 476 GPs in 227 practices responded (response rates: practitioner, 10.3%; practice, 23.3%). Cancer decision-support tools were available in 83/227 (36.6%, 95% confidence interval 30.3% to 43.1%) practices. Tools were available and likely to be used in 38/227 (16.7%, 12.1% to 22.2%) practices. In sub-group analyses of 172 English practices, there was no difference in mean two-week-wait referral rate between practices with tools and those without (mean adjusted difference in referrals per 100,000: 3.1, -5.5 to +11.7).

Conclusions: This is the first survey of cancer decision-support tool availability and use. It suggests that the tools are an underused resource in the UK. Given the cost of cancer investigation, a randomised controlled trial of such clinical decision-support aids would be appropriate.

How this fits in

Clinical decision-support tools quantify the risk of an undiagnosed cancer in symptomatic patients and may help GPs improve their selection of patients for investigation for suspected cancer. The tools are an integral part of the National Cancer Strategy, yet their uptake in general practice is unknown. Our survey - the first of the availability and use of cancer decision-support tools in the UK - reports that the paper-based and electronic tools are available to GPs in approximately one-third

(36.6%, 95% CI 30.3% to 43.1%) of UK practices and likely to be used in 16.7% (95% CI 12.1% to 22.2%).

Introduction

Diagnosing cancer quickly after patients develop symptoms is a UK priority (1). National guidelines help UK general practitioners (GPs) select which patients warrant referral and investigation for suspected cancer (2–4). This “gatekeeper” system may cause diagnostic delay (5); for example, two-week-wait referrals for suspected cancer are less likely when patients present with “low-risk but not no-risk” than when they have “alarm” symptoms (6). Clinical decision-support tools for cancer quantify the risk of an undiagnosed cancer in symptomatic patients (7). Two main types are available: Risk Assessment Tools (RATs) and QCancer® (8,9). RATs are available for 18 specific cancer sites, and use symptoms and test results to estimate the risk of cancer (8,10–17). QCancer® uses symptoms, test results and patient risk factors for six specific cancer sites (9,18–22), plus one for each sex estimating the overall risk of cancer (23,24). RATs were distributed to all 10,000 general practices in England in 2012 as mousemats and flipcharts (1). QCancer® is freely accessible on the internet (<http://www.qcancer.org/>).

In 2013, both RATs and QCancer® were incorporated into GP software systems and renamed collectively as “**electronic clinical decision-support tools for cancer**”. For simplicity, they are hereafter called “cancer tools”. RATs were integrated into the GP software system Vision (INPS, London), and QCancer® into EMIS Web (Egton Medical Information Systems, Leeds). Together, EMIS Web and Vision had 62% of the market share of GP IT systems in 2015 (25).

There is little research on the clinical utility of cancer tools, or on their availability and uptake in UK primary care (7). A recent qualitative study of a convenience sample of 126 GPs aimed to improve the understanding of how GPs use cancer tools. The study reported that 18.3% of GPs used either a RAT or QCancer®, but that overall awareness of these tools was low [Chisnell et al. submitted]. A cohort study compared the numbers of cancer investigations and diagnoses before and after the

introduction of colorectal and lung RATs to 165 general practices in England. The introduction of RATs was associated with increased diagnostic activity and additional diagnoses of lung and colorectal cancer (26). A 2×2 design trial of a GP intervention, which included the colorectal and lung RATs, found no evidence that it was associated with faster time to diagnosis of cancer in rural Australia (27). No studies have investigated the association between use of cancer decision-support tools and use of the UK's urgent referral pathway for suspected cancer. Understanding this association is important for two reasons: (1) the impact of increased referrals on resources; and (2) use of the two-week-wait referral pathway is associated with improved cancer outcomes (28).

Therefore, the primary aims of this study were to identify the proportions of general practices and of GPs with access to cancer tools, and, where there is access to tools, what proportions of practices actually use them. The secondary aim was to investigate any association between a practice's access to cancer tools and referral activity for suspected cancer.

Two main measures of cancer referral activity are available in the Public Health England dataset. The first is a diagnostic process indicator – the age- and sex-standardised number of referrals adjusted for practice size – and is reliable at the practice level (29). This indicator is suitable for assessing whether use of the tools is associated with increased number of referrals and potential impact on resources. The second is a diagnostic outcome indicator – the proportion of patients undergoing a two-week-wait referral who are subsequently diagnosed with cancer (conversion rate). However, the small numbers of cancers diagnosed per practice make this measure unreliable (29), so we decided against using it as an outcome measure for investigating the association between use of tools and cancer outcomes.

Methods

This was a cross-sectional postal survey in UK primary care. The questionnaire was planned using the best practice guidelines for survey design, and was further reviewed and edited by the originators of RATs (Hamilton) and QCancer[®] (Hippisley-Cox and Coupland). Images of the paper-based tools and screenshots of the electronic cancer tools were included to ease their identification. Questionnaires included a general practice identifier, but not the name of the responding GP. The questionnaire was piloted with five GPs for its clarity and design. To measure tool usage, GPs were asked how likely they would be to consult desktop or electronic tools in a patient with symptoms of possible cancer, using a four-point Likert scale: very likely, likely, unlikely and very unlikely (30). Participants were asked to select any aspects of the tools they found helpful, from a list of positive aspects of lung and colorectal cancer RATs reported previously (26,31,32). Participants were also asked to rank in order of usefulness the three main interactive functions of the electronic tools: **Alert/prompt** cancer risk scores appear automatically once a patient's electronic notes are opened, if there is a risk of any individual cancer $\geq 2\%$; **Symptom checker** GPs can request a patient's cancer risk; and **Searches/report** GPs can search records and produce summaries of patients ranked by cancer risk. The questionnaire, covering letter and information sheet are available from the authors. The questionnaire had no free text comments section; however, any written comments, or comments sent by email or phone, were recorded (see Supplementary material).

The survey was administered by a commercial firm, Binley's (www.binleys.com). It was conducted at the practice level, reflecting how practice software decisions are generally made. The invited population was general practices in the UK and clinically active GP partners/principals, sessional GPs (including salaried and locum GPs) and GP registrars. For 5% precision, assuming a population proportion of 50% of practices with access to a tool and adjusting for the clustered design, we estimated a sample size of 392 general practices was required at the 95% confidence level. We estimated a 40% response rate, so obtained a random probability sample of 975 general practices

from Binley's (33). Questionnaires were sent to all GPs (n=4,350) and registrars (n=250) in these practices in July 2017, with a follow-up questionnaire for non-responding practices one month later. The data collection stopped 14 weeks after reminders were issued. To incentivise participation, a charitable donation of £7.50 (to Cancer Research UK and Macmillan Cancer Support equally) was made for the first 400 replies.

Analyses

If any single GP reported that they had access to cancer tools, it was assumed that this was true for all other GPs at that practice. We used simple descriptive statistics for access to and use of cancer tools. For completeness, GP- and practice-level responses are reported. Survey responses from English practices were linked to the Public Health England dataset. Referral activity was measured using the practices' age- and sex-adjusted number of two-week-wait referrals for suspected cancer per 100,000 head of population (29). The association between practice tool availability and two-week-wait referral rate was estimated using weighted least-squares regression with robust errors, adjusted for the practice's index of multiple deprivation (34–37).

Results

Sample characteristics

Responses were received from 473 GPs and 3 GP registrars in 227 practices. The response rate at the practice level was 23.3% and at the practitioner level, 10.3%. Responding practices had a median of 6 GPs (interquartile range (IQR), 4 to 8), of whom a median of 2 (IQR 1 to 3) responded. The mean within-practice response rate was 43.7% (95% confidence interval 39.3% to 48.1%). Unprompted comments indicated that lack of time (n=12) and lack of awareness of the tools (n=6) were the most common reasons for non-response. 294 (61.8%) of responders had been practising for 11 years or more; 299 (62.8%) were working between 5 and 8 sessions per week (Supplementary Table 1). EMIS Web was the most frequently used IT software (96/227, 42.3%), followed by TPP SystemOne (74/227,

32.6%) and INPS Vision (32/227, 14.1%), largely matching the national market share of these software packages (Supplementary Table 2). The distribution of practices by Index of Multiple Deprivation was broadly representative of practices in the UK (data not shown).

Access to a paper-based cancer tool in mousemat or flipchart form was reported by 63 of the 476 (13.2%) GPs. At the practice level, tools were available in 51 of the 227 (22.5%, 95% CI 17.2 to 28.5%) practices (Table 1). The “other” tools are listed in Supplementary Table 3, and consist of national guidelines or summaries thereof, which do not quantify the risk of undiagnosed cancer. Of the 63 GPs with access to a mouse mat or flip chart, 39 (61.9%) reported that they were unlikely or very unlikely to use it during a consultation with a patient with possible symptoms of cancer. The participants’ choices from a selected list of helpful aspects of the paper-based cancer tools are reported in Table 2.

The electronic cancer tool was downloaded or activated on the IT system of 58 of 476 GPs (12.2%) (Table 3), equating to a practice level of 42/227 (19.0%, 14.0% to 24.6%). Practices using EMIS Web and INPS Vision were equally likely to have downloaded/activated the software (n=32/96, 33.3% EMIS Web, n=10/32, 31.3% INPS Vision) (Table 4). Of the 476 GPs, 174 (36.6%) were unaware of electronic tools, and 39 (8.2%) reported that they would like to have them but that they are not available for their system.

Of the 58 GPs with access to the electronic cancer tools, 17 (29.3%) reported having integrated it into their practice, and 9 (15.5%) having received training. Only 5 GPs had both received training and had integrated the tool into their practice. At the practice level, training had been received by at least one GP in 6 (14.3%) practices with access to the tool. The tool was integrated into the practice of at least one GP in 15 (35.7%) practices.

The “alert prompt” and “symptom checker” functions were deemed the most useful by 16 (27.6%) and 14 (24.1%) of the 58 GPs with access to the tool, respectively. Two-thirds (39/58,

67%) reported that they would be unlikely or very unlikely to use an electronic cancer tool to assess a patient whose symptoms may represent cancer. The participants' choices from a selected list of helpful aspects of the electronic cancer tools are reported in Table 2.

Overall, of the 476 GPs, 112 (23.5%, 95% CI 19.7% to 27.6%) had access to a cancer tool in either paper or electronic format, or both. At the practice level, this equates to at least one GP with access in 83 practices (36.6%, 30.3% to 43.1%). Of the 227 general practices, 38 (16.7%, 12.1% to 22.2%) contained at least one GP who had access to the tools and was likely or very likely to use them.

Association between use of tools and two-week-wait referral activity

Of the 172 practices in England with published two-week-wait referral and conversion rates, 68 had access to either a paper or electronic cancer tool. There was no difference in mean two-week-wait referral rate between practices with or without access to either type of tool, after adjusting for index of multiple deprivation (mean difference 3.1 referrals per 100,000, -5.5 to +11.7, per 100,000) (Table 5).

Discussion

Summary

This is the first UK-wide survey of the availability of cancer tools. These tools, in paper or electronic format, are available to GPs in approximately one-third (36.6%, 95% CI 30.3% to 43.1%) of UK practices. The proportion of general practices where at least one GP had access to the tools and was likely or very likely to use them was 16.7% (95% CI 12.1% to 22.2%).

There are no current plans to re-release paper-based tools, with the expectation that the electronic version will become the norm. Therefore, the 19.0% (14.0% to 24.6%) with access to the electronic version may be the more important measure. Currently, the tools are only available via EMIS Web and INPS Vision, and approximately one-third of the practices using these software systems had opted to download or activate them. The software will shortly be

integrated into SystemOne, with approximately 33% of the UK market share. Between them, EMIS Web, SystemOne and INPS Vision represent over 95% of the GP software systems available (25); therefore, in the near future it is reasonable to assume that nearly all GPs could access tools, should they choose to download/activate them.

It could be argued that use of the tools risks overwhelming secondary care resources; however, we found no evidence of an association between tool availability and an increase in the number of two-week-wait referrals at the practice level. The inability to find differences may be because the tools have only been available for a short while and are not yet embedded in clinical practice. To assess the effectiveness of the tools future studies will need to consider the two-week-wait referrals and the impact these have on stage at diagnosis and survival. Our finding that the tools are an underused resource in the UK suggests that there is potential to explore the effectiveness of these tools on appropriate referrals to improve cancer outcomes within a randomised controlled trial.

Strengths and limitations

Our selection of a 40% response rate had seemed reasonable, based on a reported value of 61% (95% confidence interval 59% to 63%) in 2011, and adjusted downward to reflect the current workload crisis in general practice (33,38). However, our achieved sample was smaller than planned, resulting in wide confidence intervals. The low response rate probably reflects high GP workload, as volunteered by practice managers and reported elsewhere (38) (Chisnell et al. submitted). Responder bias is important to consider, given our low response rate. Our study would overestimate tool availability if responders are more likely than non-responders to have access to the tools. However, the proportion of practices with computer systems supporting electronic tools was not overrepresented in our sample: 57% of responding practices had Vision and EMIS systems – very similar to the national picture of 62%. This suggests that the response rate is unrelated to access to tools via the software used

at the practice, and that the effect of responder bias on the estimates of tool availability and use is likely to be small. The possibility remains that responses to questions about use of the tools may have been influenced by GPs' cognitive biases. Furthermore, it could be argued that practices that have chosen to access the cancer tools are more engaged in the early cancer diagnosis framework than practices who have not. This might be expected to lead to overestimates of the association between use of the tools and the number of two-week-wait referrals.

Comparison with existing literature

There is no comparable literature on practice-level availability and use of cancer tools for cancer in the UK or elsewhere. Chisnell et al. (submitted) reported that use of cancer tools was low (18.3% of GPs), but this estimate is at the GP level.

Our finding of low levels of use of cancer tools are supported by qualitative studies reporting that the cancer tool's screen alerts increase the risk of disuse through "prompt fatigue" (31,32), and generally low levels of awareness (Chisnell et al submitted).

Implications for research and/or practice

This study and previous qualitative work suggests that improvements in design and training of tools may increase uptake (26,32,39). Any training should encourage GPs to maximise symptom recording in a patient's medical record, using a code rather than text fields. This is because the algorithms rely on coded data, and omission of data recorded in text fields is associated with bias (40).

As the levels of tool uptake are relatively low, it remains possible to carry out a randomised controlled trial to assess whether these tools are genuinely helpful in improving the selection of patients for investigation and to assess the impact on resource use in a cost-effectiveness framework. The potential benefits of improved patient selection include better targeting of

investigation resources, earlier diagnosis and reduced treatment costs (26,39,41–46). Such a trial should include a study of barriers to use, and ways to overcome them.

Acknowledgements: The authors thanks Dr Ruben Mujica-Mota for advice on how to conduct the regression sub-analyses, and Professors Hippisley-Cox and Coupland for their help with questionnaire development.

Author Contributions: SP wrote the analysis plan, designed the survey, conducted the data analysis and wrote the first draft of the paper. AS contributed to the analysis plan, advised on data analysis and commented on all drafts of the paper. AML commented on all drafts of the paper. WH commented on the design of the survey and commented on all drafts of the paper.

References

1. HM Government. *Improving Outcomes: A Strategy for Cancer*. London: The Department of Health; 2011.
2. National Institute for Health and Care Excellence. *Suspected cancer: recognition and referral* [NG12]. London: NICE, 2015. Available from: <http://www.nice.org.uk/guidance/NG12>
3. NHS Scotland. *Scottish Cancer Referral Guidelines*. 2016. Available from: www.cancerreferral.scot.nhs.uk/
4. Health and Social Care in Northern Ireland (HSCNI). *Northern Ireland Referral Guidance for Suspected Cancer*. 2012. Available from: [http://cancerni.net/files/file/Northern Ireland Referral Guidance for Suspected Cancer _Dec12\(2\).pdf](http://cancerni.net/files/file/Northern%20Ireland%20Referral%20Guidance%20for%20Suspected%20Cancer%20_Dec12(2).pdf)
5. Vedsted P, Olesen F. Are the serious problems in cancer survival partly rooted in gatekeeper principles? An ecologic study. *Br J Gen Pract* 2011;**61**(589):e508-12.
6. Zhou Y, Mendonca SC, Abel GA, et al. Variation in “fast-track” referrals for suspected cancer by patient characteristic and cancer diagnosis: Evidence from 670 000 patients with cancers of 35 different sites. *Br J Cancer* 2018;**118**(1):24–31.
7. Usher-Smith J, Emery J, Hamilton W, Griffin SJ, Walter FM. Risk prediction tools for cancer in primary care. *Br J Cancer* 2015;**113**(12):1645–50.
8. Hamilton W, Peters TJ, Round A, Sharp D. What are the clinical features of lung cancer before the diagnosis is made? A population based case-control study. *Thorax* 2005;**60**(12):1059–65.
9. Hippisley-Cox J, Coupland C. Identifying patients with suspected lung cancer in primary care : derivation and validation of an algorithm. *Br J Gen Pract* 2011;**61**(592):e715-23.

10. Nicholson BD, Mant D, Neal RD, et al. International variation in adherence to referral guidelines for suspected cancer: a secondary analysis of survey data. *Br J Gen Pract* 2016;**66**(643):e106-13.
11. Hamilton W, Round A, Sharp D, Peters TJ. Clinical features of colorectal cancer before diagnosis: a population-based case-control study. *Br J Cancer* 2005;**93**(4):399–405.
12. Hamilton W, Sharp DJ, Peters TJ, Round AP. Clinical features of prostate cancer before diagnosis: a population-based, case-control study. *Br J Gen Pract* 2006;**56**(531):756–62.
13. Shephard E, Neal R, Rose P, Walter F, Hamilton WT. Clinical features of kidney cancer in primary care: A case-control study using primary care records. *Br J Gen Pract* 2013;**63**(609):e250-5.
14. Shephard E, Stapley S, Neal RD, Rose P, Walter FM, Hamilton W. Clinical features of bladder cancer in primary care. *Br J Gen Pract* 2012;**62**:e598–604.
15. Stapley S, Peters TJ, Neal RD, Rose PW, Walter FM, Hamilton W. The risk of pancreatic cancer in symptomatic patients in primary care: a large case-control study using electronic records. *Br J Cancer* 2012;**106**(12):1940–4.
16. Stapley S, Peters TJ, Neal RD, Rose PW, Walter FM, Hamilton W. The risk of oesophago-gastric cancer in symptomatic patients in primary care: a large case-control study using electronic records. *Br J Cancer* 2013;**108**(1):25–31.
17. Walker S, Hyde C, Hamilton W. Risk of uterine cancer in symptomatic women in primary care: Case-control study using electronic records. *Br J Gen Pract* 2013;**63**(614):e643-8.
18. Hippisley-Cox J, Coupland C. Identifying patients with suspected gastro-oesophageal cancer in primary care: derivation and validation of an algorithm. *Br J Gen Pract* 2011;**61**(592):e707-14.
19. Hippisley-Cox J, Coupland C. Identifying women with suspected ovarian cancer in primary care: derivation and validation of algorithm. *BMJ* 2011;**344**:d8009.
20. Hippisley-Cox J, Coupland C. Identifying patients with suspected pancreatic cancer in primary care: derivation and validation of an algorithm. *Br J Gen Pract* 2012;**62**(594):e38-45.
21. Hippisley-Cox J, Coupland C. Identifying patients with suspected renal tract cancer in primary care: derivation and validation of an algorithm. *Br J Gen Pract* 2012;**62**(597):e251-60.
22. Hippisley-Cox J, Coupland C. Identifying patients with suspected colorectal cancer in primary care: derivation and validation of an algorithm. *Br J Gen Pract* 2012 **62**(594):e29-37.
23. Hippisley-Cox J, Coupland C. Symptoms and risk factors to identify women with suspected cancer in primary care: Derivation and validation of an algorithm. *Br J Gen Pract* 2013;**63**(606):e11-21.
24. Hippisley-Cox J, Coupland C. Symptoms and risk factors to identify men with suspected cancer in primary care: derivation and validation of an algorithm. *Br J Gen Pract* 2013;**63**(606):e1-10.
25. NHS Digital. GP Systems of Choice. Practice and System Data: Market Share 2015. Available at <https://digital.nhs.uk/services/gp-systems-of-choice> [Accessed 21/11/2018].
26. Hamilton W, Green T, Martins T, Elliott K, Rubin G, Macleod U. Evaluation of risk assessment tools for suspected cancer in general practice: a cohort study. *Br J Gen Pract* 2013;**63**(606):e30-6.

27. Emery JD, Gray V, Walter FM, et al. The Improving Rural Cancer Outcomes (IRCO) Trial: a factorial cluster-randomised controlled trial of a complex intervention to reduce time to diagnosis in rural patients with cancer in Western Australia: a study protocol. *BMJ Open* 2014;**4**(9):e006156–e006156.
28. Møller H, Gildea C, Meechan D, Rubin G, Round T, Vedsted P. Use of the English urgent referral pathway for suspected cancer and mortality in patients with cancer: cohort study. *BMJ* 2015;**351**:h5102.
29. Abel G, Saunders CL, Mendonca SC, Gildea C, McPhail S, Lyratzopoulos G. Variation and statistical reliability of publicly reported primary care diagnostic activity indicators for cancer: A cross-sectional ecological study of routine data. *BMJ Qual Safety* 2018;**27**(1):21-30.
30. Lozano LM, García-Cueto E, Muñiz J. Effect of the number of response categories on the reliability and validity of rating scales. *Methodology* 2008;**4**:73-79.
31. Dikomitis L, Green T, Macleod U. Embedding electronic decision-support tools for suspected cancer in primary care: a qualitative study of GPs' experiences. *Prim Health Care Res Dev* 2015;**16**(6):548–55.
32. Green T, Martins T, Hamilton W, Rubin G, Elliott K, Macleod U. Exploring GPs' experiences of using diagnostic tools for cancer: a qualitative study in primary care. *Fam Pract* 2015;**32**(1):101–5.
33. Creavin ST, Creavin AL, Mallen CD. Do GPs respond to postal questionnaire surveys? A comprehensive review of primary care literature. *Fam Pract* 2011;**28**(4):461-7.
34. Imbens GW, Kolesár M. Robust standard errors in small samples: Some practical advice. *Rev Econ Stat* 2016;**98**(4):701-12.
35. UCLA Statistical Consulting Group. Stata Analysis Tools: Weighted Least Squares Regression. Available online at <https://stats.idre.ucla.edu/stata/ado/analysis/stata-analysis-toolsweighted-least-squares-regression/> [Accessed 21/11/2018].
36. Abel GA, Barclay ME, Payne RA. Adjusted indices of multiple deprivation to enable comparisons within and between constituent countries of the UK including an illustration using mortality rates. *BMJ Open* 2016;**6**(11):e012750.
37. HM Government. English Indices of Deprivation 2015: technical report. London: Ministry of Housing, Communities and Local Government, 2015. Available online at <https://www.gov.uk/government/statistics/english-indices-of-deprivation-2015> [Accessed 21/11/2018].
38. Hobbs FDR, Bankhead C, Mukhtar T, et al. Clinical workload in UK primary care: a retrospective analysis of 100 million consultations in England, 2007-14. *Lancet* 2016;**387**(10035):2323-2330.
39. Moffat J, Ironmonger L, Green T. Clinical Decision Support Tool for Cancer (CDS) Project: Evaluation Report to the Department of Health. London: CRUK, 2014. Available online at https://www.cancerresearchuk.org/sites/default/files/cds_final_310714.pdf [Accessed 21/11/2018].
40. Price SJSJ, Stapley SASA, Shephard E, Barraclough K, Hamilton WT. Is omission of free text records a possible source of data loss and bias in Clinical Practice Research Datalink studies? A case-control study. *BMJ Open* 2016;**6**(5):e011664.

41. Collins GS, Altman DG. Identifying patients with undetected colorectal cancer: An independent validation of QCancer (Colorectal). *Br J Cancer* 2012;**107**(2):260-5.
42. Collins GS, Altman DG. Identifying patients with undetected pancreatic cancer in primary care: An independent and external validation of QCancer[®] (Pancreas). *Br J Gen Pract* 2013;**63**(614):e636-42.
43. Collins GS, Altman DG. Identifying patients with undetected renal tract cancer in primary care: an independent and external validation of QCancer[®] (Renal) prediction model. *Cancer Epidemiol* 2013;**37**(2):115–20.
44. Collins GS, Altman DG. Identifying patients with undetected gastro-oesophageal cancer in primary care: External validation of QCancer[®] (Gastro-Oesophageal). *Eur J Cancer* 2013;**49**(5):1040-8.
45. Collins GS, Altman DG. Identifying women with undetected ovarian cancer: Independent and external validation of QCancer[®] (Ovarian) prediction model. *Eur J Cancer Care (Engl)* 2013;**22**(4):423-9.
46. Department of Health. *The Likely Impact of Earlier Diagnosis of Cancer on Costs and Benefits to the NHS*. London: Department of Health, 2011.