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## VARIATION IN CERVICAL AND BREAST CANCER SCREENING COVERAGE IN ENGLAND: A CROSS-SECTIONAL ANALYSIS TO CHARACTERISE PRIMARY CARE TRUSTS (PCTS) WITH ATYPICAL BEHAVIOUR

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# Variation in cervical and breast cancer screening coverage in England: a cross-sectional analysis to characterise Primary Care Trusts (PCTs) with atypical behaviour 

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\text { Running title: } \quad \text { Atypical Screening coverage in England }
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## Strengith and limitations of this study

- This study reports on thorough analysis of breast and cervical screening coverage rates to identify area-level factors associated with high and low coverage.
- This is the first study to characterise English PCTs with atypically high or low cervical or breast screening coverage using a risk-adjustment approach.
- At PCT level, high rates of deprivation, urbanisation, and ethnic minority groups other than Asian, Black, or Mixed remain independent predictors of lower coverage for both programmes, and explain most of the lower cervical screening coverage seen in London.
- PCTs with atypically low screening coverage displayed distinct correlation patterns between their population characteristics, in particular distinct correlates of deprivation: these districts may benefit from the development of new approaches to target the low-attending communities living within their boundaries.
- This study deals only with area-level rather than individual-level factors. However, this is often the only data available on participation in public health interventions; the method used is fairly simple and could easily be applied to other settings.


#### Abstract

Objectives. Reducing cancer screening inequalities in England is a major focus of the 2011 Department of Health cancer outcome strategy. Screening coverage requires regular monitoring in order to implement targeted interventions where coverage is low. This study aimed to characterise districts with atypical coverage levels for cervical or breast screening.

Design. Observational study of Primary Care Trust (PCT)-level coverage in the English Cervical and Breast screening programmes in 2012.


Setting. England, UK.

Participants. All English women invited to participate to the Cervical (age group 25-49 and 50-64) and Breast (age group 50-64) screening programmes.

Outcomes. Risk adjustment models for coverage were developed based on PCT-level characteristics. Funnel plots of adjusted coverage were constructed and atypical PCTs examined by correlation analysis.

Results. Variability in coverage was primarily explained by population factors, whereas general practice characteristics had little independent effect. Deprivation and ethnicity other than White, Asian, Black, or Mixed were independently associated with poorer coverage in both screening programmes, with ethnicity having the strongest effect; in comparison the influence of Asian, Black, or Mixed ethnic minority was limited. Deprivation, ethnicity and urbanisation largely accounted for the lower cervical screening coverage in London. However, for breast screening, being located in London remained a strong negative predictor. A subset of PCTs was identified as having atypical coverage across programmes.

Correlates of deprivation in PCTs with relatively low adjusted coverage were substantially different from overall correlates of deprivation.

Discussion. These results inform the continuing drive to reduce avoidable cancer deaths in England, and encourage implementation of targeted interventions in communities residing in districts identified as having atypically low coverage. Sequential implementation to monitor the impact of local interventions would help accrue evidence on 'what works'.

## Introduction

The English National Cervical and Breast Screening Programmes aim either to prevent cancer by treating pre-cancerous changes or diagnose cancer at earlier stages when treatment outcomes are more successful ${ }^{1,2}$. Their success is dependent upon high levels of participation ${ }^{3}$.

Reducing cancer screening inequalities in England is a major focus of the 2011 Department of Health cancer outcome strategy to promote early diagnosis and save lives ${ }^{4,5}$. There is a need to characterise districts that require most support in reducing inequalities or those which could be used as leading examples.

Funnel plots overlapped with control limits have been shown to be a useful tool for comparing proportional outcomes between centres or districts ${ }^{6-8}$. The outcome is plotted against a measure of precision for each district, and control limits are set around the target value. Districts lying outside the limits are subject to 'special-cause variation' and may repay further investigation. Control limits can be adjusted to incorporate sources of variation such as demographic and socio-economic factors in order to identify districts with atypically high or low outcomes, given their known characteristics ${ }^{8,9}$.

Identification of atypical districts might be expected to be a simple matter. It is, however, challenging due to the necessarily incomplete nature of aggregate data, the possible collinearities in such data, and the multiplicity of model choices, even with relatively small numbers of potential risk factors.

Factors associated with variation in screening coverage in England have previously been identified: deprivation, non-Caucasian ethnicity and poorer primary care-level service have
been found linked with lower attendance at both cervical ${ }^{10,11}$ and breast ${ }^{12,13}$ screening. In addition, coverage in London has generally been observed to be lower than the national average ${ }^{1,2}$.

We constructed funnel plots to display the scatter of cervical and breast screening coverage around the national average in areas defined by former English Primary Care Trusts (PCTs). We developed risk adjustment models based on demographic, socio-economic and primary care-level characteristics, and control limits were adjusted accordingly. PCTs with atypically high or low coverage were identified, and associations among PCT characteristics were investigated in an attempt to highlight those districts where further investigation may be beneficial in informing policy to improve coverage.

## Methods

## Data source

Coverage data were available in geographical areas defined by former English PCTs. Data from April 2011 to March 2012 were sourced from the Health \& Social Care Information Centre (HSCIC) ${ }^{1,2}$. Cervical screening coverage was defined as the percentage of eligible women registered with a general practice, who had an adequate screening test within the last 3.5 years for $25-49$ year-olds, and the last 5 years for 50-64 year-olds. PCT-level data were obtained for the two age groups separately. Breast screening coverage was defined as the percentage of eligible women registered with a general practice, who had an adequate screening mammogram within the last 3 years. Data for 50-64 year-olds were obtained to match the older cervical screening group.

The percentage urbanisation within each PCT was derived from the urban-rural classification ${ }^{14}$. For two PCTs with missing data (Stockton-on-Tees, Isle of Wight), the Local Authority urbanisation score was used instead.

The income deprivation domain score from the English Indices of Multiple Deprivation (IMD) 2010 was obtained and the percentage deprivation calculated as a population-weighted average of Lower Super Output Area (LSOA) income deprivation score ${ }^{15}$.

Ethnicity data and the percentage of the total population without any higher education were sourced from the Office of National Statistics (ONS) 2011 Census ${ }^{16,17}$. For ethnicity, two explanatory variables were derived: the percentage of Asian, Black, or Mixed ethnic minority groups, and the percentage of other ethnic minority groups, which includes Asian and African Arabs and any other minor ethnic groups (e.g. Polynesians, Melanesians and Micronesians).

General practice characteristics data were sourced from the $\mathrm{HSCIC}^{18}$, and included average list size, percentage of single-handed practices (only 1 working provider or salaried/other general practitioner (GP) with possible additional GP registrar/retainer), practitioner headcount (excluding retainers and registrars) per $10^{5}$ population, practice staff (excluding GPs and registrars) full-time equivalent (FTE), and percentage of GPs who attained their primary medical qualification outside the UK.

## Statistical analysis

Grouped logistic regression was applied to coverage data aggregated at PCT level ${ }^{19}$. A generalized linear model with quasibinomial error distribution was used to account for within-PCT extra-binomial variation ${ }^{20}$. For the purpose of the analysis, variables were


#### Abstract

classified as "population" and "general practice" risk factors (Table 1). Continuous covariates were mean-centred. Covariates found to be significant at the $1 \%$ level using Wald tests in univariate analyses ${ }^{21}$ were considered for inclusion in two multiple regression sub-models, the first including population factors only and the second including general practice factors only. Correlation and collinearity were evaluated based on Pearson correlation coefficients (Supplementary file Table A1 \& Figure 3a) and generalized variance-inflation factors (GVIF) for covariate coefficients, respectively ${ }^{22}$. Differences between correlation coefficients in two independent groups were assessed for significance by applying Fisher's z test on ztransformed correlations ${ }^{23}$.

The full regression model was built by including both population and general practice factors that were significant at the $5 \%$ level in the sub-models. Percent of deviance (-2 loglikelihood statistic) explained by the adjusted model compared to the null (unadjusted) model was used as a descriptive measure of attribution of variation ${ }^{19}$.

Funnel plots of coverage against eligible population in each PCT were constructed ${ }^{9}$. The covariate-adjusted coverage proportion for each PCT was calculated as the product of the national average by the ratio of observed to expected values from the full regression model. The national average for coverage was used as a target value, and the 95\% and 99.8\% control limits were plotted around it using the asymptotic normal approximation, with a variance inflation factor for extra-binomial variation ( ${ }^{24}$ details available from NJM). All statistical analyses were performed in $R$ version 3.0.2.


## Results

## Data description

PCT-level data on cervical (age groups 25-49 and 50-64) and breast (age group 50-64) screening coverage are summarized in Table 1; overall, and separately for London and the rest of England. Between-PCT variability was more pronounced for breast screening (median 76.9, IQR 6.5) and the younger cervical screening group aged 25-49 (median 74.6, IQR 5.9) than for the cervical screening group aged 50-64 (median 77.5, IQR 3.5, Table 1). The difference in coverage level between London and the rest of England was also larger for the breast and younger cervical screenings groups; with median coverage 7-8\% lower in London.
[Table 1 here]

Table 1. PCT-level summary of population factors, general practice factors, and screening coverage in England in 2012 ( $n=151$ )

| Population factors | Min-Max | Mean (SD) | Median (IQR) |
| :---: | :---: | :---: | :---: |
| \% Urbanisation | 31.0-100.0 | 81.2 (21.5) | 91.0 (35.03) |
| \% Deprivation | 6.8-33.8 | 16.2 (5.8) | 15.3 (8.4) |
| \% Asian, Black, or Mixed ethnicity | 1.3-67.6 | 15.1 (15.4) | 8.9 (20.5) |
| \% Other minor ethnicity | 0.1-11.1 | 1.2 (1.6) | 0.6 (1.3) |
| \% No higher education | 10.1-35.2 | 23.0 (5.1) | 23.0 (6.8) |
| \% Registered women aged 25-29 | 12.2-32.2 | 19.5 (4.2) | 18.3 (5.2) |
| General practice factors | Min-Max | Mean (SD) | Median (IQR) |
| Average practice list size | 4026.4-9566.2 | 6656.2 (1371.2) | 6537.1 (2236.0) |
| \% Single-handed practices | 0.0-41.0 | 13.45 (10.2) | 11.0 (16.0) |
| Practitioner headcount per $10^{5}$ population | 50.9-95.3 | 68.7 (8.3) | 67.7 (10.8) |
| Practice staff FTE | 146.3-1884.2 | 513.7 (296.7) | 424.0 (283.7) |
| \% Practitioners qualified outside UK | 3.0-70.0 | 26.4 (14.7) | 25.0 (19.2) |
| Screening coverage (\%) | Min-Max | Mean (SD) | Median (IQR) |
| Cervical group aged 25-49 |  |  |  |
| Overall | 58.7-80.4 | 73.4 (4.4) | 74.6 (5.9) |
| London SHA (Q36) | 58.7-77.7 | 67.8 (4.6) | 67.8 (5.7) |
| Rest of England | 67.4-80.4 | 74.8 (3.0) | 75.4 (3.8) |
| Cervical group aged 50-64 |  |  |  |
| Overall | 69.1-82.0 | 77.2 (2.5) | 77.5 (3.5) |
| London SHA (Q36) | 69.1-80.9 | 75.7 (2.8) | 75.6 (3.1) |
| Rest of England | 70.1-82.0 | 77.6 (2.3) | 77.9 (2.8) |
| Breast group aged 50-64 |  |  |  |
| Overall | 59.5-84.7 | 75.6 (5.1) | 76.9 (6.5) |
| London SHA (Q36) | 59.5-78.8 | 69.0 (4.9) | 68.8 (8.6) |
| Rest of England | 64.6-84.7 | 77.3 (3.6) | 78.1 (5.5) |

FTE, Full-Time Equivalent; IQR: Inter Quartile Range; SD, Standard Deviation; SHA, Strategic Health Authority

## Relationships between population and general practice factors, and coverage

Tables 2.1, 2.2 and 2.3 show the unadjusted and adjusted odds ratios of the associations between population and general practice risk factors with coverage. Each factor was found to be univariately associated with coverage in all screening groups, except for the percentage of population with no higher education and the practitioner headcount, which were only significant for the cervical screening group aged 25-49.

Variability in coverage was primarily explained by population factors with general practice characteristics only accounting for a small fraction of the residual variability (< $2 \%$ of total deviance after adjustment for population factors). Population covariates explained a lesser percentage of the total deviance among the cervical screening group aged 50-64 (45\%, Table 2.2) than the cervical screening group aged 25-49 (78\%, Table 2.1) or the breast screening group (72\%, Table 2.3); overall variability was also lowest among the former group (IQR 3.5 versus IQR 5.9 and 6.5, respectively, Table 1).

With regard to general practice factors, only staff FTE remained positively associated with cervical screening coverage after accounting for population factors (Table 2.2).

After adjusting for deprivation, ethnicity and education, residing in London and urbanisation were no longer significantly associated with lower cervical screening coverage, but both remained associated with lower breast screening coverage.

Deprivation remained inversely associated with coverage in all screening groups, but displayed some collinearity with other factors for the cervical screening group aged 25-49
(Tables 2.1).

Absence of higher education remained associated with higher coverage in the cervical screening group aged 25-49 after adjusting for other population factors (Table 2.1). In this latter group, the effect of deprivation and education were no longer significant when the model accounted for the percentage of registered women aged 25-29 (Supplementary file Table A2.1).

After adjusting for other population factors, the percentage of other ethnic minority groups remained negatively correlated with coverage in all screening groups, whereas the percentage of Asian, Black, or Mixed ethnic minority groups was no longer associated with lower breast screening coverage (Tables 2.2-2.3).
[Tables 2.1-2.3 here]

Table 2.1 Regression modelling for cervical screening coverage among women aged 25-49


CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority
${ }^{\text {\$ }}$ The variance of the coefficient estimate is being inflated by multicollinearity with other factors ( $\sqrt{\text { GVIF }}=2.7$ ).

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Table 2.2 Regression modelling for cervical screening coverage among women aged 50-64

| Model | Univariate |  |  | Population |  | General practice |  | Population \& General practice 45.3\% |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  |  |  |
| Population factors | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $\begin{gathered} \text { p-value } \\ \text { (Wald, } \chi 2 \text { ) } \end{gathered}$ | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $\begin{gathered} \text { p-value } \\ \text { (Wald, } \chi^{2} \text { ) } \end{gathered}$ | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value (Wald, ${ }^{2}$ ) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value (Wald, $\mathrm{x}^{2}$ ) |
| \% Urbanisation | $\begin{gathered} 0.997 \\ (0.996,0.998) \end{gathered}$ | <0.001 | 25.5\% | $\begin{gathered} 0.9986 \\ (0.9976-0.9995) \end{gathered}$ | 0.004 | - |  | $\begin{gathered} 0.999 \\ (0.9978-0.9998) \end{gathered}$ | 0.02 |
| London SHA (Q36) | $\begin{gathered} 0.886 \\ (0.837,0.937) \end{gathered}$ | $<0.001$ | 10.6\% | $\begin{gathered} 0.940 \\ (0.875,1.010) \end{gathered}$ | NS (0.09) | - |  | - |  |
| \% Deprivation | $\begin{gathered} 0.987 \\ (0.984,0.990) \end{gathered}$ | < 0.001 | 31.1\% | $\begin{gathered} 0.989 \\ (0.985,0.992) \end{gathered}$ | $<0.001$ | - |  | $\begin{gathered} 0.990 \\ (0.985,0.994) \end{gathered}$ | $<0.001$ |
| \% Asian, Black, or Mixed ethnicity | $\begin{gathered} 0.997 \\ (0.996,0.998) \end{gathered}$ | < 0.001 | 9.9\% | $\begin{gathered} 1.005 \\ (1.003,1.007) \end{gathered}$ | < 0.001 | - |  | $\begin{gathered} 1.004 \\ (1.002,1.006) \end{gathered}$ | < 0.001 |
| \% Other minor ethnicity | $\begin{gathered} 0.959 \\ (0.947,0.972) \end{gathered}$ | < 0.001 | 19.6\% | $\begin{gathered} 0.970 \\ (0.952,0.988) \end{gathered}$ | 0.001 | - |  | $\begin{gathered} 0.963 \\ (0.946,0.980) \end{gathered}$ | $<0.001$ |
| \% No higher education | $\begin{gathered} 0.997 \\ (0.993,1.002) \\ \hline \end{gathered}$ | NS (0.3) | 0.9\% | ()- |  | - |  | - |  |
| General practice factors | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | p-value <br> (Wald, x2) | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | p-value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ \text { (95\% CI) } \end{gathered}$ | $p$-value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, x2) |
| Average practice list size | $\begin{gathered} \hline 1.00004 \\ (1.00003,1.00006) \end{gathered}$ | < 0.001 | 20.2\% | - |  | $\begin{gathered} \hline 1.000025 \\ (1.000003-1.000047) \end{gathered}$ | 0.02 | $\begin{gathered} \hline 0.999996 \\ (0.999979,1.000012) \end{gathered}$ | 0.6 |
| \% Single-handed practices | $\begin{gathered} 0.995 \\ (0.993,0.997) \end{gathered}$ | $<0.001$ | 13.1\% | - |  | $\begin{gathered} 0.999 \\ (0.995,1.002) \end{gathered}$ | NS (0.4) | - |  |
| Practitioner headcount per $10^{5}$ population | $\begin{gathered} 0.998 \\ (0.996,1.001) \end{gathered}$ | NS (0.2) | 1.2\% | - |  | (-) |  | - |  |
| Practice staff FTE | $\begin{gathered} 1.00015 \\ (1.00010,1.00020) \end{gathered}$ | < 0.001 | 19.5\% | - |  | $\begin{gathered} 1.00010 \\ (1.00005,1.00016) \end{gathered}$ | < 0.001 | $\begin{gathered} 1.000058 \\ (1.000007,1.000109) \end{gathered}$ | 0.03 |
| \% Practitioners qualified outside UK | $\begin{gathered} 0.997 \\ (0.996,0.999) \end{gathered}$ | $<0.001$ | 7.8\% | - |  | $\begin{gathered} 1.001 \\ (0.998,1.002) \end{gathered}$ | NS (0.5) | - |  |

CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority

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Table 2.3 Regression modelling for breast screening coverage among women aged 50-64

| Model <br> Deviance explained by model | Univariate |  |  | Population General practice |  |  |  | Population \& General practice 70.6\% |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | - |  |  |  |  |  |  |  |  |
| Population factors | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | p -value <br> (Wald, x ) | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | p -value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{CI}) \end{gathered}$ | p-value <br> (Wald, x2) |
| \% Urbanisation | $\begin{gathered} 0.992 \\ (0.991,0.993) \end{gathered}$ | < 0.001 | 50.5\% | $\begin{gathered} 0.996 \\ (0.995,0.998) \end{gathered}$ | < 0.001 | - |  | $\begin{gathered} 0.996 \\ (0.995,0.998) \end{gathered}$ | < 0.001 |
| London SHA (Q36) | $\begin{gathered} 0.642 \\ (0.587,0.703) \end{gathered}$ | $<0.001$ | 37.7\% | $\begin{gathered} 0.896 \\ (0.811,0.990) \end{gathered}$ | 0.03 | - |  | $\begin{gathered} 0.885 \\ (0.806,0.970) \end{gathered}$ | 0.009 |
| \% Deprivation | $\begin{gathered} 0.972 \\ (0.967,0.978) \end{gathered}$ | $<0.001$ | 38.8\% | $\begin{gathered} 0.991 \\ (0.986,0.997) \end{gathered}$ | 0.002 | - |  | $\begin{gathered} 0.991 \\ (0.985,0.997) \end{gathered}$ | 0.004 |
| \% Asian, Black, or Mixed ethnicity | $\begin{gathered} 0.987 \\ (0.985,0.989) \end{gathered}$ | < 0.001 | 49.1\% | $\begin{gathered} 0.999 \\ (0.996,1.002) \end{gathered}$ | NS (0.5) | - |  | - |  |
| \% Other minor ethnicity | $\begin{gathered} 0.880 \\ (0.863,0.898) \end{gathered}$ | $<0.001$ | 50.8\% | $\begin{gathered} 0.948 \\ (0.923,0.973) \end{gathered}$ | < 0.001 | - |  | $\begin{gathered} 0.945 \\ (0.922,0.969) \end{gathered}$ | $<0.001$ |
| \% No higher education | $\begin{gathered} 1.010 \\ (1.001,1.019) \end{gathered}$ | NS (0.03) | 3.1\% | - _ |  | - |  | - |  |
| General practice factors | OR (95\% CI) | p-value <br> (Wald, x2) | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $\begin{gathered} \text { p-value } \\ \text { (Wald, } \chi^{2} \text { ) } \end{gathered}$ | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $\begin{gathered} \text { p-value } \\ \text { (Wald, }{ }^{2} \text { ) } \end{gathered}$ | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, x2) |
| Average practice list size | $\begin{gathered} 1.00010 \\ (1.00007,1.00012) \end{gathered}$ | < 0.001 | 26.5\% |  |  | 1.000046 $(1.000006,1.000087)$ | 0.03 | $\begin{gathered} 1.00001 \\ (0.99998,1.00003) \end{gathered}$ | 0.6 |
| \% Single-handed practices | $\begin{gathered} 0.988 \\ (0.984,0.991) \end{gathered}$ | < 0.001 | 24.2\% | - |  | $\begin{gathered} 0.9945 \\ (0.9886,1.0004) \end{gathered}$ | NS (0.07) | - |  |
| Practitioner headcount per $10^{5}$ population | $\begin{gathered} 0.996 \\ (0.991,1.001) \end{gathered}$ | NS (0.1) | 1.7\% | - |  | - |  | - |  |
| Practice staff FTE | $\begin{gathered} 1.00025 \\ (1.00015,1.00035) \end{gathered}$ | $<0.001$ | 14.1\% | - |  | $\begin{gathered} 1.000099 \\ (0.999990,1.000209) \end{gathered}$ | NS (0.07) | - |  |
| \% Practitioners qualified outside UK | $\begin{gathered} 0.993 \\ (0.990,0.995) \end{gathered}$ | $<0.001$ | 16.0\% | - |  | $\begin{gathered} 0.9992 \\ (0.9957,1.0027) \end{gathered}$ | NS (0.6) | - |  |

CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority

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## Identification of PCTs with atypical coverage

Figure 1 illustrates the PCTs with coverage estimates lying outside the control limits prior to (Figure 1a-c) and after full covariate adjustment (Figure $1 a^{\prime}-c^{\prime}$ ). The geographical location of PCTs with atypical coverage is shown in Figure 2.

Over two-thirds of the PCTs initially lying below limits for cervical screening - for most, located within London - no longer lay below limits after adjustment. For the breast screening group, only one out of the four initial outliers (Kensington \& Chelsea in London data not shown) was found to lie within limits after adjustment, while a new London PCT was uncovered as atypically low (Wandsworth, London). For two London PCTs, the adjusted coverage remained below the $99.8 \%$ lower limit for the cervical screening group aged 25-49, and ranked among the 15 lowest PCTs for the other two screening groups (Hammersmith and Fulham, and Camden, Figure 2).

In contrast to what was observed for the PCTs lying below limits, the PCTs lying above the 95\% upper limits after adjustment were mostly different from those identified prior to adjustment: only 1 in 2 PCTs for the cervical screening group aged 25-49, 1 in 5 for the cervical screening group aged 50-64, and 2 in 5 for the breast screening group would have been identified as atypically high performers without adjustment (Figure 1 \& data not shown). Two PCTs displayed atypically high coverage in all screening groups irrespective of age (Enfield, London and Nottinghamshire County Teaching, East Midlands).

## Characteristics of PCTs with relatively high and low adjusted coverage

PCTs were ranked according to their adjusted coverage values (Supplementary file Tables A3.1 \& A3.2). Associations between population factors were investigated among the 15 lowest- (Figure 3b) and the 15 highest-ranking PCTs (Figure 3c).

For all screening groups, we noted strong positive associations between deprivation and non-white ethnicities among the highest-ranking PCTs, which differed significantly from the associations seen among lowest-ranking PCTs (Fisher's z test p<0.05 for cervical screening and $\mathrm{p}=0.05$ for breast screening group among minor ethnicity groups only, Figure 3d).

For cervical screening, a strong positive correlation between deprivation and absence of higher education was observed among lowest-ranking PCTs ( $\rho=0.77$ and 0.68 for age group 25-49 and 50-64, respectively), which tended to not be as strong overall or among highestranking PCTs, in particular for the younger age group (Fisher's $z$ test $p=0.04$ ).

Lowest-ranking PCTs tended to have populations of other minor ethnicity with a higher level of education ( $\rho=-0.88,-0.77$ and -0.70 for cervical age groups $25-49$ and $50-64$, and breast age group 50-64 respectively) compared with overall or high-ranking PCTs, in particular for cervical screening (Fisher's z test p=0.1 for both cervical age groups).

## DISCUSSION

This aim of this analysis was to identify and characterise PCTs that displayed atypically high or low cervical or breast screening coverage given population and general practice PCT-level risk factors. We found that a subset of PCTs with atypical coverage levels was common to both programmes, while other sets were more specific to the programme or age group. Our risk adjustment results confirm the importance of demographic and socio-economic characteristics for coverage levels, and highlight the comparatively minor impact of various aspects of primary care. This suggests that strategies targeted at raising awareness or addressing barriers among socially-and culturally-diverse populations are likely to be the most effective at increasing coverage.

The number of practice staff FTE remained positively associated with cervical screening coverage but not breast screening coverage after adjusting for population factors. The finding that cervical screening coverage is more likely to be influenced by general practice factors is unsurprising since many women are screened at their local practice ${ }^{25}$, and previous studies have shown the number of nurses per practice to be associated with cervical screening coverage in deprived areas ${ }^{10}$.

Coverage in London has generally been observed to be lower than the national average ${ }^{1,2}$, in spite of some other public health features (for example obesity rates) being better in London ${ }^{26}$. We found that urbanisation, ethnicity, and deprivation, largely accounted for the lower cervical screening uptake in London. For breast screening however, being located in London, remained a strong independent negative risk factor, which warrants further investigation.

Deprivation was an independent negative risk factor for all screening groups, as also found for cervical screening by Bang and coll. ${ }^{27}$. In the cervical screening group aged 25-49, this effect was in part explained by numbers of women under 30 , as was the positive impact of lack of higher education on coverage. Cervical screening coverage has been reported to be lower in younger women ${ }^{28}$, but younger women of lower socio-economic status or with fewer educational qualifications, regardless of ethnicity, have also been shown to be positively influenced by the 2009 Jade Goody's story with respect to cervical screening behaviour ${ }^{29}$, giving hints as to potential strategies for improving uptake.

The impact of Asian, Black, or Mixed ethnic minority groups on coverage differed between programmes after controlling for other population factors. For breast screening, it was no longer significant. For cervical screening, we found it negatively influenced coverage in the age group 25-49, but was associated with greater coverage in the age group 50-64. Previously, only an overall negative overall association after adjustment for other population factors had been reported for cervical screening in women aged 25-64 ${ }^{27}$. For both programmes, and regardless of age, other ethnic minority groups were still associated with poorer coverage after accounting for deprivation and urbanisation, with a particularly strong effect in breast screening. In addition, our results suggest that women of other ethnic minority background, who may be well educated and living in areas with smaller Asian, Black, or Mixed ethnic minority populations, are less likely to go for screening. Arabs communities account for a moderately large subset of the 'other' ethnic minority groups (40\%), and uptake of cervical and breast screening has been shown to be low in these populations for a number of reasons, including religious beliefs, emotional barriers (embarrassment/fear), language barriers or taboos surrounding sexual activity (for cervical
screening) ${ }^{30-32}$. These populations may therefore require newly targeted interventions to promote screening.

Our correlation analyses suggest that PCTs with atypical coverage levels differ from one another not only in respect of a number of population- and general practice-level characteristics, but also in how these characteristics relate to each other. Correlates of deprivation in PCTs with relatively low adjusted coverage were substantially different from the general results, and even more so for cervical screening. In particular, the nature of the relationship between deprivation and non-White ethnicity differed, with an inverse relationship between deprivation and non-White ethnic groups among lowest-ranking PCTs.

Using funnel plots based on crude performance data to assess quality of care at area level may overestimate the number of "underperforming" districts, and overdispersion needs to be addressed a priori. We chose a risk adjustment approach to uncover PCTs with atypically high or low coverage given particular population and general practice characteristics. PCTs with adjusted coverage values lying outside control limits display a behaviour which cannot solely be explained by the area-level risk factors investigated.

PCTs with atypically high coverage were singled out and could be investigated to identify any local health interventions and policies that might help improve coverage in districts with similar characteristics but lower performance. Unfortunately, there is a general lack of reporting in the research literature across PCTs on the impact of local interventions that have been implemented to improve screening uptake (ED, unpublished PhD thesis), so identifying 'what works' is challenging.

Simultaneously, PCTs with atypically low coverage were distinguished from those lying within bounds after accounting for urbanisation, deprivation and ethnicity, in particular for the London region. These districts may benefit from further investigation to uncover the features driving their atypically low coverage and help design population-specific strategies. Additional risk factors that may explain low coverage, as well as differences in PCT performance between programmes, include the percentage of women who are disabled ${ }^{33}$, incarcerated ${ }^{34}$, have greater difficulty in accessing services as indexed by time to screening centre ${ }^{13}$, and differential utilization behaviour as a result of socio-cultural factors, such as marital status ${ }^{35}$, occupation ${ }^{36}$, sexual orientation ${ }^{37}$, and overseas birthplace or religious beliefs ${ }^{11,38}$ that might apply to particular programmes.

Our results are limited by the aggregated nature of the data, which may conceal ecological associations within districts. This could account for the weak association seen between coverage and general practice characteristics after adjustment for population factors. However, similar trends were observed when analysing general practice-level data for cervical screening coverage ${ }^{27}$. Another limitation is that PCT no longer exist, but the findings can easily be applied to the newly defined English Clinical Commissioning Group level (CCG) by direct mapping ${ }^{39}$.

The strength of the approach of combining risk adjustment modelling with funnel plots was to allow us to identify districts with unusual level of screening coverage after accounting for some of the important demographic and socio-economic characteristics of their populations and their primary care settings, known to affect coverage level. Such an approach could be implemented sequentially to monitor the impact of local interventions in a centralised fashion. This method could also be adapted for use with other health indicators.

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$$

Our results demonstrate that population factors largely explain the lower coverage in London. In addition, PCTs in London and other urban centres with specific population characteristics such as non-deprived ethnic minority groups were identified as requiring targeted intervention to improve coverage levels. Bilingual outreach and community-based advocacy, such as support from family and community leaders including GPs, has been found to be valuable in increasing uptake of cancer screening in ethnic minorities ${ }^{40}$.

We hope these results will inform the continued drive to reduce inequalities in cancer screening and avoidable deaths, and encourage implementation of targeted interventions in communities residing within districts identified as having atypically low coverage.

## Abbreviations

| CI | Confidence Interval |
| :--- | :--- |
| FTE | Full-Time Equivalent |
| GP | General Practitioner |
| GVIF | Generalized Variance-Inflation Factor |
| HSCIC | Health and Social Care Information Centre |
| IMD | Indices of Multiple Deprivation |
| IQR | Interquartile Range |
| LSOA | Lower Super Output Area |
| PCT | Primary Care Trust |
| SD | Standard Deviation |
| SHA | Strategic Health Authority |

## Acknowledgements

We would like to thank the Knowledge and Intelligence Team (KIT) - East for supplying the PCT digital boundary information and creating contrast maps according to our results, in particular Aphrodite Niggebrugge, and acknowledge all who contributed to the collection and maintenance of HSCIC, ONS and APHO data.

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## Competing interests

None

## AUTHOR'S CONTRIBUTION

NJM performed the statistical analysis, interpreted the results and wrote the manuscript. ED carried out the data informatics and data checks, and co-wrote the manuscript. JW \& JW provided general expert guidance. SWD provided general statistical guidance. All authors reviewed and approved the final manuscript.

## Data sharing statement

Data are freely available from the HSCIC:

Cervical screening: http://www.hscic.gov.uk/catalogue/PUB10339/bres-scre-prog-eng-2011-12-tab.xls Breast screening: http://www.hscic.gov.uk/catalogue/PUB07990/cerv-scre-prog-eng-2011-12-tab.xls No additional data available.

## ETHICS APPROVAL

None

## List of Figures

Figure 1. Funnel plots of screening coverage and list of PCTs lying outside the 95\% control limits prior to and after risk adjustment
Top left panel. Funnel plots of screening coverage prior to any adjustment
(a) Cervical screening in women aged 25-49.
(b) Cervical screening in women aged 50-64.
(c) Breast screening in women aged 50-64.

Top right panel. Funnel plots of screening coverage after adjustment for population and general practice factors
( $a^{\prime}$ ) Cervical screening in women aged 25-49.
(b') Cervical screening in women aged 50-64.
(c') Breast screening in women aged 50-64.
_ - _ - - - 95.0\% control limits
99.8\% control limits

SHA, Strategic Health Authority; Q30, North East; Q31, North West ;Q33, East Midlands; Q34, West Midlands; Q35, East of England; Q36, London; Q37, South East Coast; Q38, South Central; Q39, South West.

Table. Number of PCTS lying outside the $95 \%$ control limits prior to and after risk adjustment. The number of PCTs within London SHA (Q36) is shown in brackets.

## Figure 2. Geographical location of atypical PCTs

Map. Map of PCT 2006 boundaries with PCTs lying below the $95 \%$ lower control limits after risk adjustment coloured in red and PCTs lying above the $95 \%$ upper control limits after risk adjustment coloured in green.

Table. PCTs lying outside the control limits are listed with corresponding percentile given in brackets. PCTs with coverage ranking among the 15 lowest- (rank $\leq 15$ ) or 15 highest (rank $\geq 137$ ) are specified. All PCTs lying outside the control limits had relative coverage rankings $\leq 15$ for lower $95 \%$ limit and $\geq 137$ for upper $95 \%$ limit.

SHA, Strategic Health Authority; Q30, North East; Q31, North West ;Q33, East Midlands; Q34, West Midlands; Q35, East of England; Q36, London; Q37, South East Coast; Q38, South Central; Q39, South West.

## Figure 3. Correlations between population factors overall, and among the 15 highest-

 and 15 lowest-ranking PCTs after risk adjustmenta-c. Correlation coefficients are displayed in each cell. a, All PCTs; b, 15-lowest ranking PCTs; c, 15 highest ranking PCTs.
For the 15 lowest and 15 highest-ranking PCTs, correlation coefficients which are significantly different from zero at the $1 \%$ level are highlighted in green for positive correlations, and in red for negative correlations.

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## References

1. Health \& Social Care Information Centre (HSCIC). Cervical Screening Programme, England Statistics for 2012-13. 2014;Available from http://www.hscic.gov.uk/catalogue/PUB07990/cerv_scre_prog_eng_201112_rep_v3.pdf (Accessed 14/11/14).
2. Health \& Social Care Information Centre (HSCIC). Breast Screening Programme, England Statistics for 2011-12. 2013;Available from http://www.hscic.gov.uk/catalogue/PUB10339/bres-scre-prog-eng-2011-12rep.pdf (Accessed 14/11/14).
3. Stead MJ, Wallis MG, Wheaton ME. Improving uptake in non-attenders of breast screening: selective use of second appointment. Journal of Medical Screening 1998;5(2):69-72.
4. Weller DP, Campbell C. Uptake in cancer screening programmes: a priority in cancer control. Br J Cancer 2009;101 Suppl 2:S55-9.
5. Department of Health. Improving Outcomes: a strategy for cancer. 2011;https://www.gov.uk/government/publications/the-national-cancer-strategy (Accessed 14/11/14).
6. Spiegelhalter DJ. Funnel plots for comparing institutional performance. Statistics in Medicine 2005a;24(8):1185-202.
7. Spiegelhalter DJ. Handling over-dispersion of performance indicators. Quality and Safety in Health Care 2005b;14(5):347-51.
8. Association of Public Health Observatories (APHO). Statistical process control methods in public health intelligence. Technical briefing 2: 2008;Available from http://www.apho.org.uk/resource/item.aspx?RID=39445 (Accessed 01/12/14).
9. Dover DC, Schopflocher DP. Using funnel plots in public health surveillance. Popul Health Metr 2011;9(1):58.
10. Baker D, Middleton E. Cervical screening and health inequality in England in the 1990s. J Epidemiol Community Health 2003;57(6):417-23.
11. Webb R, Richardson J, Pickles A. A population-based study of primary care predictors of non-attendance for cervical screening. Journal of Medical Screening 2004;11(3):13540.
12. Eilbert KW, Carroll K, Peach J, et al. Approaches to improving breast screening uptake: evidence and experience from Tower Hamlets. Br J Cancer 2009;101 Suppl 2:S64-7.
13. Chen C, Yu W, Huabing W, et al. Determinations of low breast screening uptake using geographically weighted regression model. GEOINFORMATICS 2012 (20th International Conference on Geoinformatics) 15-17 June 2012, Hong Kong, China;http://ieeexplore.ieee.org/xpls/icp.jsp?arnumber=6270323:1-6.
14. Association of Public Health Observatories (APHO). Urban-rural classification of PCTs (post October 2006 boundaries). 2008;Available from http://www.apho.org.uk/resource/item.aspx?RID=53312 (Accessed 14/11/14).
15. UK Department for Communities and Local Government. English indices of deprivation 2010. Statistics 2011;Available from https://www.gov.uk/government/statistics/english-indices-of-deprivation-2010 (Accessed 02/12/14).

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml
16. Office of National Statistics (ONS). 2011 Census: KS201UK Ethnic group, local authorities in the UK - NOMIS. 2011;Available from http://www.nomisweb.co.uk/census/2011/ks201ew (Accessed 14/11/14).
17. Office for National Statistics (ONS). 2011 Census: Highest level of qualification - NOMIS. 2011;Available from http://www.nomisweb.co.uk/census/2011/qs501ew (Accessed 14/11/14).
18. Health \& Social Care Information Centre (HSCIC). NHS Staff 2002-2012, General Practice. 2013;Available from http://www.hscic.gov.uk/catalogue/PUB09536 (Accessed 14/11/14).
19. Hilbe JM, Robinson AP. Generalized Linear Models. Methods of Statistical Model Estimation: Chapman and Hall/CRC, 2013:99-120.
20. vanEngelsdorp D, Lengerich E, Spleen A, et al. Standard epidemiological methods to understand and improve Apis mellifera health. Journal of Apicultural Research 2013;52(4):1-16.
21. McCullagh P, Nelder, JA. Generalized Linear Models. 2nd ed. London: Chapman \& Hall, 1989.
22. Kabacoff RI. R in Action: Data Analysis and Graphics with R. Shelter Island, NY, USA: Manning Publications Co., 2011.
23. Diedenhofen B. cocor: Comparing correlations (Version 1.0-0). Compare two correlations based on independent groups 2013;Available from http://r.birkdiedenhofen.de/pckg/cocor/ (Accessed 14/11/14):18-21.
24. Gelman A, Hill J. Generalized linear models. Data Analysis Using Regression and Multilevel/Hierarchical Models: Cambridge University Press, 2006:117.
25. Public Health England. NHS Cervical Screening Programme. 2014;Available from http://www.cancerscreening.nhs.uk/cervical/index.html (Accessed 14/11/14).
26. Scholes S, Prescott P, Bajekal M. Health \& lifestyle indicators for Strategic Health Authorities 1994-2002. Mean diastolic blood pressure. Health Survey for England 2014;Available from http://www.dh.gov.uk/prod_consum_dh/idcplg?IdcService=GET_FILE\&dID=1084\& Rendition=Web (Accessed 14/11/14).
27. Bang JY, Yadegarfar G, Soljak M, et al. Primary care factors associated with cervical screening coverage in England. J Public Health (Oxf) 2012;34(4):532-8.
28. Waller J, Jackowska M, Marlow L, et al. Exploring age differences in reasons for nonattendance for cervical screening: a qualitative study. BJOG 2012;119(1):26-32.
29. Marlow LAV, Sangha A, Patnick J, et al. The Jade Goody Effect: whose cervical screening decisions were influenced by her story? Journal of Medical Screening 2012;19(4):184-88.
30. Szarewski A, Cadman L, Ashdown-Barr L, et al. Exploring the acceptability of two selfsampling devices for human papillomavirus testing in the cervical screening context: a qualitative study of Muslim women in London. Journal of Medical Screening 2009;16(4):193-98.
31. Donnelly TT, Khater AH, Al-Bader SB, et al. Arab women's breast cancer screening practices: a literature review. Asian Pac J Cancer Prev 2013;14(8):4519-28.
32. Padela AI, Murrar S, Adviento B, et al. Associations Between Religion-Related Factors and Breast Cancer Screening Among American Muslims. Journal of immigrant and minority health / Center for Minority Public Health 2014.
33. Horner-Johnson W, Dobbertin K, Andresen EM, et al. Breast and Cervical Cancer Screening Disparities Associated with Disability Severity. Women's Health Issues 2014;24(1):e147-e53.
34. Plugge E, Fitzpatrick R. Factors affecting cervical screening uptake in prisoners. J Med Screen 2004;11(1):48-9.
35. Lo SH, Waller J, Wardle J, et al. Comparing barriers to colorectal cancer screening with barriers to breast and cervical screening: a population-based survey of screening-age women in Great Britain. Journal of Medical Screening 2013;20(2):73-79.
36. Moser K, Patnick J, Beral V. Inequalities in reported use of breast and cervical screening in Great Britain: analysis of cross sectional survey data. BMJ 2009;338:b2025.
37. Reiter PL, McRee AL. Cervical cancer screening (Pap testing) behaviours and acceptability of human papillomavirus self-testing among lesbian and bisexual women aged 21-26 years in the USA. The journal of family planning and reproductive health care / Faculty of Family Planning \& Reproductive Health Care, Royal College of Obstetricians \& Gynaecologists 2014.
38. Szczepura A, Price C, Gumber A. Breast and bowel cancer screening uptake patterns over 15 years for UK south Asian ethnic minority populations, corrected for differences in socio-demographic characteristics. BMC Public Health 2008;8(1):346.
39. Allies computing. Translate PCTs to CCGs conversion tool. 2014;Available from http://www.alliescomputing.com/innovation/pct-ccg-mapping (Accessed 14/11/14).
40. Shankleman J, Massat NJ, Khagram L, et al. Evaluation of a service intervention to improve awareness and uptake of bowel cancer screening in ethnically-diverse areas. Br J Cancer 2014;111(7):1440-47.

Figure1
$190 \times 254 \mathrm{~mm}$ ( $96 \times 96$ DPI)


Below lower
95\% control limit

| 5H1 / Hammersmith and Fulham | Q36 | Below 99.8\% (0.01\%) | Rank $\geq 137$ | Rank $\geq 137$ |
| :---: | :---: | :---: | :---: | :---: |
| 5k6 / Harrow | Q36 | Below 95\% (0.2\%) | Rank $\geq 137$ |  |
| 517 / Camden | Q36 | Below 99.8\% (0.001\%) | Rank $\geq 137$ | Below 95\% (0.02\%) |
| $5 \mathrm{NJ} / \mathrm{Sefton}$ | Q31 | Rank $\geq 137$ | Below 95\% (0.3\%) | - |
| 5PG / Birmingham East \& North | Q34 | Rank $\geq 137$ | Below 95\% (1.2\%) |  |
| 5LD / Lambeth | Q36 | - | Rank $\geq 137$ | Below 95\% (0.01\%) |
| 5LG/Wandsworth | Q36 | - | Rank $\geq 137$ | Below 95\% (0.4\%) |
| 5NT / Manchester | Q 31 | - | Rank $\geq 137$ | Below 95\% (0.2\%) |
| Above upper |  |  |  |  |
| 95\% control limit |  |  |  |  |
| $5 \mathrm{C1}$ / Enfield | 036 | Above 95\% (99.8\%) | Above 95\% (99.9\%) | Above 95\% (99.9\%) |
| 5N8 / Nottinghamshire County Teaching | Q33 | Above 95\% (93.9\%) | Above 95\% (93.9\%) | Above 95\% (93.5\%) |
| 5 KL / Sunderland Teaching | Q30 | Rank $\leq 15$ | Above 95\% (98.6\%) | Rank $\leq 15$ |
| 5NC/Waltham Forest | Q36 | - | Above 95\% (99.3\%) | Rank $\leq 15$ |
| TAN / North East Lincolnshire Care Trust Plus | 032 | Rank $\leq 15$ | Above 95\% (99.8\%) | - |
| $5 \mathrm{N4}$ / Sheffield | 032 | - |  | Above 95\% (98.5\%) |
| 5N7 / Derby city | Q33 | - | Rank $\leq 15$ | Above 95\% (99.7\%) |
| 5PA / Ledestershire County \& Rutland | 033 | - | - | Above 95\% (96.5\%) |

Figure2
$190 \times 254 \mathrm{~mm}$ ( $96 \times 96$ DPI)


Figure3
$190 \times 254 \mathrm{~mm}$ ( $96 \times 96$ DPI)

## SUPPLEMENTARY FILE (MASSAT, Douglas et AL.)

Table A1. Correlations between population and general practice factors, and screening coverage (all PCTS)
Upper diagonal: Correlation coefficient; Lower diagonal: p-value of test for significant correlation between paired samples

|  | Population factors |  |  |  |  |  | General practice factors |  |  |  |  | Coverage |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | \% <br> Urbanization | \% <br> Deprivation | \% Asian, Black, or Mixed ethnicity | $\begin{aligned} & \text { \% Other } \\ & \text { minor } \\ & \text { ethnicity } \end{aligned}$ | \% No higher education | \% Registered women aged 25-29 | Average practice list size | \% Singlehanded practices | Practitioner headcount per $10^{5}$ population | Practice staff FTE | \% <br> Practitioners qualified outside UK | Cervical group aged 25-49 | Cervical group aged 50-64 | Breast group aged 50-64 |
| Population factors |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| \% Urbanization | 1 | 0.61 | 0.56 | 0.48 | -0.002 | 0.63 | -0.50 | 0.49 | 0.02 | -0.51 | 0.42 | -0.60 | -0.49 | -0.66 |
| \% Deprivation | <0.001 | 1 | 0.58 | 0.39 | 0.41 | 0.64 | -0.58 | 0.46 | 0.23 | -0.36 | 0.54 | -0.56 | -0.47 | -0.58 |
| \% Asian, Black, or Mixed ethnicity | < 0.001 | < 0.001 | 1 | 0.70 | -0.27 | 0.62 | -0.35 | 0.39 | 0.14 | -0.25 | 0.43 | -0.78 | -0.24 | -0.68 |
| \% Other minor ethnicity | < 0.001 | < 0.001 | < 0.001 | 1 | -0.45 | 0.60 | -0.40 | 0.38 | 0.14 | -0.25 | 0.22 | -0.78 | -0.45 | -0.74 |
| \% No higher education | NS (0.9) | < 0.001 | 0.001 | $<0.001$ | 1 | -0.11 | -0.26 | 0.21 | -0.25 | -0.04 | 0.37 | 0.31 | 0.02 | 0.31 |
| \% Registered women $\qquad$ <br> aged 25-29 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | NS (0.2) | 1 | -0.33 | 0.30 | 0.26 | -0.24 | 0.21 | -0.71 | -0.43 | -0.69 |
| General practice factors |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Average practice list size | <0.001 | < 0.001 | < 0.001 | < 0.001 | 0.001 | < 0.001 | 1 | -0.75 | 0.06 | 0.36 | -0.52 | 0.42 | 0.39 | 0.49 |
| \% Single-handed practices | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.01 | < 0.001 | $<0.001$ | 1 | -0.35 | -0.27 | 0.61 | -0.41 | -0.28 | -0.43 |
| Practitioner headcount per $10^{5}$ population | NS (0.8) | 0.006 | NS (0.1) | NS (0.08) | 0.002 | 0.001 | NS (0.5) | $<0.001$ | 1 | 0.06 | -0.35 | -0.19 | -0.14 | -0.22 |
| Practice staff FTE | < 0.001 | $<0.001$ | 0.002 | 0.002 | NS (0.7) | 0.004 | < 0.001 | 0.001 | NS (0.4) | 1 | -0.36 | 0.34 | 0.37 | 0.35 |
| \% Practitioners qualified outside UK | < 0.001 | < 0.001 | < 0.001 | 0.006 | < 0.001 | 0.009 | < 0.001 | < 0.001 | < 0.001 | $<0.001$ | 1 | -0.29 | -0.18 | -0.32 |
| Coverage |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Cervical group aged 25-49 | < 0.001 | < 0.001 | <0.001 | <0.001 | <0.001 | $<0.001$ | <0.001 | <0.001 | 0.02 | $<0.001$ | <0.001 | 1 | 0.68 | 0.84 |
| Cervical group aged 50-64 | < 0.001 | < 0.001 | 0.004 | < 0.001 | NS (0.8) | < 0.001 | < 0.001 | < 0.001 | NS (0.09) | < 0.001 | 0.03 | < 0.001 | 1 | 0.65 |
| Breast group aged 50-64 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.006 | < 0.001 | < 0.001 | <0.001 | <0.001 | 1 |

NS: not significant at the 5\% level; FTE: Full Time Equivalent
$1 \mid P a g e$

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Table A2.1. Regression modelling for cervical screening coverage among women aged 25-49, including \% registered women aged 25-29.


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Table A3.1. 15 lowest-ranking PCTs prior to and after adjustment for population and general practice factors

| Rank | No adjustment | Adjustment for Population |
| :--- | :---: | :---: | :---: |
| (percentile) | \& General practice factors |  |
| (percentile) |  |  |

PCT: Primary Care Trust; SHA: Strategic Health Authority
Dark red: PCTs lying below the $95 \%$ lower control limits using full model (atypical PCTs).
Orange: Atypical PCTs found among PCTs with lowest relative coverage prior to adjustment.
Percentile is given for those PCTs lying below the $95 \%$ lower control limits prior to and after full adjustment.

* PCT in London SHA (Q36)
\$PCT lying between the $95 \%$ and $99.8 \%$ lower control limits
\$ PCT lying below the 99.8\% lower control limits

Table A3.2. 15 highest-ranking PCTs prior to and after adjustment for population and general practice factors

|  | Rank | No adjustment | Adjustment for Population <br> (percentile) |
| :---: | :---: | :---: | :---: |
| (/151) | General practice factors |  |  |
| (percentile) |  |  |  |

[^2]STROBE Statement-checklist of items that should be included in reports of observational studies

|  | $\begin{gathered} \text { Item } \\ \text { No } \\ \hline \end{gathered}$ | Recommendation |
| :---: | :---: | :---: |
| Title and abstract | 1 | (a) Indicate the study's design with a commonly used term in the title or the abstract |
|  |  | (b) Provide in the abstract an informative and balanced summary of what was done and what was found |
| Introduction |  |  |
| Background/rationale | 2 | Explain the scientific background and rationale for the investigation being reported |
| Objectives | 3 | State specific objectives, including any prespecified hypotheses |
| Methods |  |  |
| Study design | 4 | Present key elements of study design early in the paper |
| Setting | 5 | Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection |
| Participants | 6 | (a) Cohort study-Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up <br> Case-control study-Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controls <br> Cross-sectional study-Give the eligibility criteria, and the sources and methods of selection of participants |
|  |  | (b) Cohort study-For matched studies, give matching criteria and number of exposed and unexposed <br> Case-control study-For matched studies, give matching criteria and the number of controls per case |
| Variables | 7 | Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable |
| Data sources/ measurement | 8* | For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group |
| Bias | 9 | Describe any efforts to address potential sources of bias |
| Study size | 10 | Explain how the study size was arrived at |
| Quantitative variables | 11 | Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why |
| Statistical methods | 12 | (a) Describe all statistical methods, including those used to control for confounding |
|  |  | (b) Describe any methods used to examine subgroups and interactions |
|  |  | (c) Explain how missing data were addressed |
|  |  | (d) Cohort study-If applicable, explain how loss to follow-up was addressed <br> Case-control study-If applicable, explain how matching of cases and controls was addressed <br> Cross-sectional study-If applicable, describe analytical methods taking account of sampling strategy |

(e) Describe any sensitivity analyses

Continued on next page

| Results |  |  |
| :---: | :---: | :---: |
| Participants | 13* | (a) Report numbers of individuals at each stage of study-eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed |
|  |  | (b) Give reasons for non-participation at each stage |
|  |  | (c) Consider use of a flow diagram |
| Descriptive data | 14* | (a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders |
|  |  | (b) Indicate number of participants with missing data for each variable of interest |
|  |  | (c) Cohort study-Summarise follow-up time (eg, average and total amount) |
| Outcome data | 15* | Cohort study-Report numbers of outcome events or summary measures over time |
|  |  | Case-control study-Report numbers in each exposure category, or summary measures of exposure |
|  |  | Cross-sectional study-Report numbers of outcome events or summary measures |
| Main results | 16 | (a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, $95 \%$ confidence interval). Make clear which confounders were adjusted for and why they were included |
|  |  | (b) Report category boundaries when continuous variables were categorized |
|  |  | (c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period |
| Other analyses | 17 | Report other analyses done-eg analyses of subgroups and interactions, and sensitivity analyses |
| Discussion |  |  |
| Key results | 18 | Summarise key results with reference to study objectives |
| Limitations | 19 | Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias |
| Interpretation | 20 | Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence |
| Generalisability | 21 | Discuss the generalisability (external validity) of the study results |
| Other information |  |  |
| Funding | 22 | Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based |

*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

## BMJ Open

## VARIATION IN CERVICAL AND BREAST CANCER SCREENING COVERAGE IN ENGLAND: A CROSS-SECTIONAL ANALYSIS TO CHARACTERISE DISTRICTS WITH ATYPICAL BEHAVIOUR

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SCHOLARONE ${ }^{\mathrm{m}}$
Manuscripts

## Variation in cervical and breast cancer screening coverage in England: a cross-sectional ANALYSIS TO CHARACTERISE DISTRICTS WITH ATYPICAL BEHAVIOUR

Running title: Atypical Screening coverage in England

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## Keywords

Screening; Cervical cancer; Breast cancer; Health inequalities; Public health

## Word count

Abstract: 278; Text: 3,086

## Strengths and limitations of this study

- This study reports on an analysis of breast and cervical screening coverage rates to identify area-level factors associated with high and low coverage.
- This is the first study to characterise English districts with atypically high or low cervical or breast screening coverage using a risk-adjustment approach.
- At district level, high rates of deprivation, urbanisation, and ethnic minority groups other than Asian, Black, or Mixed remain independent predictors of lower coverage for both programmes, and explain most of the lower cervical screening coverage seen in London.
- Districts with atypically low screening coverage displayed distinct correlation patterns between their population characteristics, in particular with regard to deprivation: these districts may benefit from the development of new approaches to target the low-attending communities living within their boundaries.
- This study deals only with area-level rather than individual-level factors. However, this is often the only data available on participation in public health interventions; the method used is fairly simple and could easily be applied to other settings.


#### Abstract

Objectives. Reducing cancer screening inequalities in England is a major focus of the 2011 Department of Health cancer outcome strategy. Screening coverage requires regular monitoring in order to implement targeted interventions where coverage is low. This study aimed to characterise districts with atypical coverage levels for cervical or breast screening.

Design. Observational study of district-level coverage in the English Cervical and Breast screening programmes in 2012.


Setting. England, UK.

Participants. All English women invited to participate to the Cervical (age group 25-49 and 50-64) and Breast (age group 50-64) screening programmes.

Outcomes. Risk adjustment models for coverage were developed based on district-level characteristics. Funnel plots of adjusted coverage were constructed and atypical districts examined by correlation analysis.

Results. Variability in coverage was primarily explained by population factors, whereas general practice characteristics had little independent effect. Deprivation and ethnicity other than White, Asian, Black, or Mixed were independently associated with poorer coverage in both screening programmes, with ethnicity having the strongest effect; in comparison the influence of Asian, Black, or Mixed ethnic minority was limited. Deprivation, ethnicity and urbanisation largely accounted for the lower cervical screening coverage in London. However, for breast screening, being located in London remained a strong negative predictor. A subset of districts was identified as having atypical coverage across
programmes. Correlates of deprivation in districts with relatively low adjusted coverage were substantially different from overall correlates of deprivation.

Discussion. These results inform the continuing drive to reduce avoidable cancer deaths in England, and encourage implementation of targeted interventions in communities residing in districts identified as having atypically low coverage. Sequential implementation to monitor the impact of local interventions would help accrue evidence on 'what works'.

## Introduction

The English National Cervical and Breast Screening Programmes aim either to prevent cancer by treating pre-cancerous changes or diagnose cancer at earlier stages when treatment outcomes are more successful ${ }^{1,2}$. Their success is dependent upon high levels of participation ${ }^{3}$.

Reducing cancer screening inequalities in England is a major focus of the 2011 Department of Health cancer outcome strategy to promote early diagnosis and save lives ${ }^{4,5}$. There is a need to characterise districts that require most support in reducing inequalities or those which could be used as leading examples.

Funnel plots overlapped with control limits have been shown to be a useful tool for comparing proportional outcomes between centres or districts ${ }^{6-8}$. The outcome is plotted against a measure of precision for each district, and control limits are set around the target value. Districts lying outside the limits are subject to 'special-cause variation' and may repay further investigation. Control limits can be adjusted to incorporate sources of variation such as demographic and socio-economic factors in order to identify districts with atypically high or low outcomes, given their known characteristics ${ }^{8,9}$.

Identification of atypical districts might be expected to be a simple matter. It is, however, challenging due to the necessarily incomplete nature of aggregate data, the possible collinearities in such data, and the multiplicity of model choices, even with relatively small numbers of potential risk factors.

Factors associated with variation in screening coverage in England have previously been identified: deprivation, non-Caucasian ethnicity and poorer primary care-level service have
been found linked with lower attendance at both cervical ${ }^{10,11}$ and breast ${ }^{12,13}$ screening. In addition, coverage in London has generally been observed to be lower than the national average ${ }^{1,2}$.

We constructed funnel plots to display the scatter of cervical and breast screening coverage around the national average in areas defined by former English Primary Care Trusts (PCTs), the commissioning groups for GPs at the time of data collection. We developed risk adjustment models based on demographic, socio-economic and primary care-level characteristics, and control limits were adjusted accordingly. Districts with atypically high or low coverage were identified, and associations among district characteristics were investigated in an attempt to highlight those districts where further investigation may be beneficial in informing policy to improve coverage.

## Methods

## Data source

Coverage data were available in 152 geographical areas (referred to in this paper as districts) defined by the commissioning groups for GPs at the time the data were collected, i.e. the English Primary Care Trusts (PCTs). Data from April 2011 to March 2012 were sourced from the Health \& Social Care Information Centre (HSCIC) ${ }^{1,2}$. Cervical screening coverage was defined as the percentage of eligible women registered with a general practice, who had an adequate screening test within the last 3.5 years for $25-49$ year-olds, and the last 5 years for 50-64 year-olds. District-level data were obtained for the two age groups separately. Breast screening coverage was defined as the percentage of eligible
$6 \mid P a g e$
women registered with a general practice, who had an adequate screening mammogram within the last 3 years. Data for 50-64 year-olds were obtained to match the older cervical screening group.

The percentage urbanisation within each PCT was derived from the urban-rural classification ${ }^{14}$. For two PCTs with missing data (Stockton-on-Tees, Isle of Wight), the Local Authority urbanisation score was used instead.

The income deprivation domain score from the English Indices of Multiple Deprivation (IMD)
2010 was obtained and the percentage deprivation calculated as a population-weighted average of Lower Super Output Area (LSOA) income deprivation score ${ }^{15}$.

Ethnicity data and the percentage of the total population without any higher education were sourced from the Office of National Statistics (ONS) 2011 Census ${ }^{16,17}$. For ethnicity, two explanatory variables were derived: the percentage of Asian, Black, or Mixed ethnic minority groups, and the percentage of 'other' ethnic minority groups, which includes Asian and African Arabs and any other ethnic minority groups (e.g. Polynesians, Melanesians and Micronesians).

General practice characteristics data were sourced from the $\mathrm{HSCIC}^{18}$, and included average list size, percentage of single-handed practices (only 1 working provider or salaried/other general practitioner (GP) with possible additional GP registrar/retainer), practitioner headcount (excluding retainers and registrars) per $10^{5}$ population, practice staff (excluding GPs and registrars) full-time equivalent (FTE), and percentage of GPs who attained their primary medical qualification outside the UK.

## Statistical analysis

Grouped logistic regression was applied to coverage data aggregated at district level ${ }^{19}$. A generalized linear model with quasibinomial error distribution was used to account for within-district extra-binomial variation ${ }^{20}$. For the purpose of the analysis, variables were classified as "population" and "general practice" risk factors (Table 1). Continuous covariates were mean-centred. Covariates found to be significant at the $1 \%$ level using Wald tests in univariate analyses ${ }^{21}$ were considered for inclusion in two multiple regression sub-models, the first including population factors only and the second including general practice factors only. Correlation and collinearity were evaluated based on Pearson correlation coefficients (Supplementary file Table A1 \& Figure 3a) and generalized variance-inflation factors (GVIF) for covariate coefficients, respectively ${ }^{22}$. Differences between correlation coefficients in two independent groups were assessed for significance by applying Fisher's $z$ test on ztransformed correlations ${ }^{23}$.

The full regression model was built by including both population and general practice factors that were significant at the $5 \%$ level in the sub-models. Percent of deviance ( -2 loglikelihood statistic) explained by the adjusted model compared to the null (unadjusted) model was used as a descriptive measure of attribution of variation ${ }^{19}$.

Funnel plots of coverage against eligible population in each district were constructed ${ }^{9}$. The covariate-adjusted coverage proportion for each district was calculated as the product of the national average by the ratio of observed to expected values from the full regression model. The national average for coverage was used as a target value, and the $95 \%$ and $99.8 \%$ control limits were plotted around it using the asymptotic normal approximation, with a variance inflation factor for extra-binomial variation ( ${ }^{24}$ details available from NJM). All statistical analyses were performed in $R$ version 3.0.2.

## Results

## Data description

District-level data on cervical (age groups 25-49 and 50-64) and breast (age group 50-64) screening coverage are summarized in Table 1; overall, and separately for London and the rest of England. Between-district variability was more pronounced for breast screening (median 76.9, IQR 6.5) and the younger cervical screening group aged 25-49 (median 74.6, IQR 5.9) than for the cervical screening group aged 50-64 (median 77.5, IQR 3.5, Table 1). The difference in coverage level between London and the rest of England was also larger for the breast and younger cervical screenings groups; with median coverage 7-8\% lower in London.
[Table 1 here]

Table 1. District-level summary of population factors, general practice factors, and screening coverage in England in 2012 ( $\mathrm{n}=151$ )

| Population factors | Min-Max | Mean (SD) | Median (IQR) |
| :---: | :---: | :---: | :---: |
| \% Urbanisation | 31.0-100.0 | 81.2 (21.5) | 91.0 (35.03) |
| \% Deprivation | 6.8-33.8 | 16.2 (5.8) | 15.3 (8.4) |
| \% Asian, Black, or Mixed ethnicity | 1.3-67.6 | 15.1 (15.4) | 8.9 (20.5) |
| \% 'Other' ethnicity | 0.1-11.1 | 1.2 (1.6) | 0.6 (1.3) |
| \% No higher education | 10.1-35.2 | 23.0 (5.1) | 23.0 (6.8) |
| \% Registered women aged 25-29 | 12.2-32.2 | 19.5 (4.2) | 18.3 (5.2) |
| General practice factors | Min-Max | Mean (SD) | Median (IQR) |
| Average practice list size | 4026.4-9566.2 | 6656.2 (1371.2) | 6537.1 (2236.0) |
| \% Single-handed practices | 0.0-41.0 | 13.45 (10.2) | 11.0 (16.0) |
| Practitioner headcount per $10^{5}$ population | 50.9-95.3 | 68.7 (8.3) | 67.7 (10.8) |
| Practice staff FTE | 146.3-1884.2 | 513.7 (296.7) | 424.0 (283.7) |
| \% Practitioners qualified outside UK | 3.0-70.0 | 26.4 (14.7) | 25.0 (19.2) |
| Screening coverage (\%) | Min-Max | Mean (SD) | Median (IQR) |
| Cervical group aged 25-49 |  |  |  |
| Overall | 58.7-80.4 | 73.4 (4.4) | 74.6 (5.9) |
| London SHA (Q36) | 58.7-77.7 | 67.8 (4.6) | 67.8 (5.7) |
| Rest of England | 67.4-80.4 | 74.8 (3.0) | 75.4 (3.8) |
| Cervical group aged 50-64 |  |  |  |
| Overall | 69.1-82.0 | 77.2 (2.5) | 77.5 (3.5) |
| London SHA (Q36) | 69.1-80.9 | 75.7 (2.8) | 75.6 (3.1) |
| Rest of England | 70.1-82.0 | 77.6 (2.3) | 77.9 (2.8) |
| Breast group aged 50-64 |  |  |  |
| Overall | 59.5-84.7 | 75.6 (5.1) | 76.9 (6.5) |
| London SHA (Q36) | 59.5-78.8 | 69.0 (4.9) | 68.8 (8.6) |
| Rest of England | 64.6-84.7 | 77.3 (3.6) | 78.1 (5.5) |

FTE, Full-Time Equivalent; IQR: Inter Quartile Range; SD, Standard Deviation; SHA, Strategic Health Authority

## Relationships between population, general practice factors, and coverage

Tables 2.1, 2.2 and 2.3 show the unadjusted and adjusted odds ratios of the associations between population and general practice risk factors with coverage. Each factor was found to be univariately associated with coverage in all screening groups, except for the percentage of population with no higher education and the practitioner headcount, which were only significant for the cervical screening group aged 25-49.

Variability in coverage was primarily explained by population factors with general practice characteristics only accounting for a small fraction of the residual variability (< $2 \%$ of total deviance after adjustment for population factors). Population covariates explained a lesser percentage of the total deviance among the cervical screening group aged 50-64 (45\%, Table 2.2) than the cervical screening group aged 25-49 (78\%, Table 2.1) or the breast screening group (72\%, Table 2.3); overall variability was also lowest among the former group (IQR 3.5 versus IQR 5.9 and 6.5, respectively, Table 1).

With regard to general practice factors, only staff FTE remained positively associated with cervical screening coverage after accounting for population factors (Table 2.2).

After adjusting for deprivation, ethnicity and education, residing in London and urbanisation were no longer significantly associated with lower cervical screening coverage, but both remained associated with lower breast screening coverage.

Deprivation remained inversely associated with coverage in all screening groups, but displayed some collinearity with other factors for the cervical screening group aged 25-49
(Tables 2.1).

Absence of higher education remained associated with higher coverage in the cervical screening group aged 25-49 after adjusting for other population factors (Table 2.1). In this latter group, the effect of deprivation and education were no longer significant when the model accounted for the percentage of registered women aged 25-29 (Supplementary file Table A2.1).

After adjusting for other population factors, the percentage of 'other' ethnic minority groups remained negatively correlated with coverage in all screening groups, whereas the percentage of Asian, Black, or Mixed ethnic minority groups was no longer associated with lower breast screening coverage (Tables 2.2-2.3).
[Tables 2.1-2.3 here]

Table 2.1 Regression modelling for cervical screening coverage among women aged 25-49


CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority
${ }^{\text {\$ }}$ The variance of the coefficient estimate is being inflated by multicollinearity with other factors ( $\sqrt{\text { GVIF }}=2.7$ ).

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Table 2.2 Regression modelling for cervical screening coverage among women aged 50-64


CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority

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Table 2.3 Regression modelling for breast screening coverage among women aged 50-64

| Model <br> Deviance explained by model | Univariate |  |  | Population General practice |  |  |  | Population \& General practice 70.6\% |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | - |  |  |  |  |  |  |  |  |
| Population factors | OR (95\% CI) | p -value <br> (Wald, x ) | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | p -value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{CI}) \end{gathered}$ | p-value <br> (Wald, x2) |
| \% Urbanisation | $\begin{gathered} 0.992 \\ (0.991,0.993) \end{gathered}$ | < 0.001 | 50.5\% | $\begin{gathered} 0.996 \\ (0.995,0.998) \end{gathered}$ | < 0.001 | - |  | $\begin{gathered} 0.996 \\ (0.995,0.998) \end{gathered}$ | < 0.001 |
| London SHA (Q36) | $\begin{gathered} 0.642 \\ (0.587,0.703) \end{gathered}$ | $<0.001$ | 37.7\% | $\begin{gathered} 0.896 \\ (0.811,0.990) \end{gathered}$ | 0.03 | - |  | $\begin{gathered} 0.885 \\ (0.806,0.970) \end{gathered}$ | 0.009 |
| \% Deprivation | $\begin{gathered} 0.972 \\ (0.967,0.978) \end{gathered}$ | $<0.001$ | 38.8\% | $\begin{gathered} 0.991 \\ (0.986,0.997) \end{gathered}$ | 0.002 | - |  | $\begin{gathered} 0.991 \\ (0.985,0.997) \end{gathered}$ | 0.004 |
| \% Asian, Black, or Mixed ethnicity | $\begin{gathered} 0.987 \\ (0.985,0.989) \end{gathered}$ | < 0.001 | 49.1\% | $\begin{gathered} 0.999 \\ (0.996,1.002) \end{gathered}$ | NS (0.5) | - |  | - |  |
| \% 'Other' ethnicity | $\begin{gathered} 0.880 \\ (0.863,0.898) \end{gathered}$ | $<0.001$ | 50.8\% | $\begin{gathered} 0.948 \\ (0.923,0.973) \end{gathered}$ | < 0.001 | - |  | $\begin{gathered} 0.945 \\ (0.922,0.969) \end{gathered}$ | $<0.001$ |
| \% No higher education | $\begin{gathered} 1.010 \\ (1.001,1.019) \end{gathered}$ | NS (0.03) | 3.1\% | -) - |  | - |  | - |  |
| General practice factors | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, $\mathrm{x}^{2}$ ) | Deviance explained | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $\begin{gathered} \text { p-value } \\ \text { (Wald, }{ }^{2} \text { ) } \end{gathered}$ | $\begin{gathered} \text { OR } \\ (95 \% \mathrm{Cl}) \end{gathered}$ | $p$-value <br> (Wald, x2) | $\begin{gathered} \text { OR } \\ \text { (95\% CI) } \end{gathered}$ | p-value <br> (Wald, x2) |
| Average practice list size | $\begin{gathered} 1.00010 \\ (1.00007,1.00012) \end{gathered}$ | < 0.001 | 26.5\% |  |  | 1.000046 $(1.000006,1.000087)$ | 0.03 | $\begin{gathered} 1.00001 \\ (0.99998,1.00003) \end{gathered}$ | 0.6 |
| \% Single-handed practices | $\begin{gathered} 0.988 \\ (0.984,0.991) \end{gathered}$ | < 0.001 | 24.2\% | - |  | $\begin{gathered} 0.9945 \\ (0.9886,1.0004) \end{gathered}$ | NS (0.07) | - |  |
| Practitioner headcount per $10^{5}$ population | $\begin{gathered} 0.996 \\ (0.991,1.001) \end{gathered}$ | NS (0.1) | 1.7\% | - |  | - |  | - |  |
| Practice staff FTE | $\begin{gathered} 1.00025 \\ (1.00015,1.00035) \end{gathered}$ | $<0.001$ | 14.1\% | - |  | $\begin{gathered} 1.000099 \\ (0.999990,1.000209) \end{gathered}$ | NS (0.07) | - |  |
| \% Practitioners qualified outside UK | $\begin{gathered} 0.993 \\ (0.990,0.995) \end{gathered}$ | $<0.001$ | 16.0\% | - |  | $\begin{gathered} 0.9992 \\ (0.9957,1.0027) \end{gathered}$ | NS (0.6) | - |  |

CI, Confidence Interval; FTE, Full-Time Equivalent; NS, Considered non-significant (see Methods for details); SHA, Strategic Health Authority

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## Identification of districts with atypical coverage

Figure 1 illustrates the districts with coverage estimates lying outside the control limits prior to (Figure 1a-c) and after full covariate adjustment (Figure 1a'-c'). The geographical location of districts with atypical coverage is shown in Figure 2.

Over two-thirds of the districts initially lying below limits for cervical screening - for most, located within London - no longer lay below limits after adjustment. For the breast screening group, only one out of the four initial outliers (Kensington \& Chelsea in London data not shown) was found to lie within limits after adjustment, while a new London district was uncovered as atypically low (Wandsworth, London). For two London districts, the adjusted coverage remained below the $99.8 \%$ lower limit for the cervical screening group aged 25-49, and ranked among the 15 lowest districts for the other two screening groups (Hammersmith and Fulham, and Camden, Figure 2).

In contrast to what was observed for the districts lying below limits, the districts lying above the $95 \%$ upper limits after adjustment were mostly different from those identified prior to adjustment: only 1 in 2 districts for the cervical screening group aged $25-49,1$ in 5 for the cervical screening group aged 50-64, and 2 in 5 for the breast screening group would have been identified as atypically high performers without adjustment (Figure 1 \& data not shown). Two districts displayed atypically high coverage in all screening groups irrespective of age (Enfield, London and Nottinghamshire County Teaching, East Midlands).

## Characteristics of districts with relatively high and low adjusted coverage

Districts were ranked according to their adjusted coverage values (Supplementary file Tables A3.1 \& A3.2). Associations between population factors were investigated among the 15 lowest- (Figure 3b) and the 15 highest-ranking districts (Figure 3c).

For all screening groups, we noted strong positive associations between deprivation and non-white ethnicities among the highest-ranking districts, which differed significantly from the associations seen among lowest-ranking districts (Fisher's z test $p<0.05$ for cervical screening and $p=0.05$ for breast screening group among ethnic minorities groups only, Figure 3d).

For cervical screening, a strong positive correlation between deprivation and absence of higher education was observed among lowest-ranking districts ( $\rho=0.77$ and 0.68 for age group 25-49 and 50-64, respectively), which tended to not be as strong overall or among highest-ranking districts, in particular for the younger age group (Fisher's z test p=0.04). Lowest-ranking districts tended to have populations of ethnicity other than Asian, Black, or Mixed with a higher level of education ( $\rho=-0.88,-0.77$ and -0.70 for cervical age groups $25-$ 49 and 50-64, and breast age group 50-64 respectively) compared with overall or highranking districts, in particular for cervical screening (Fisher's z test p=0.1 for both cervical age groups).

## DISCUSSION

This aim of this analysis was to identify and characterise districts that displayed atypically high or low cervical or breast screening coverage given population and general practice risk factors at district level. We found that a subset of districts with atypical coverage levels was common to both programmes, while other sets were more specific to the programme or age group.

Our risk adjustment results confirm the importance of demographic and socio-economic characteristics for coverage levels, and highlight the comparatively minor impact of various aspects of primary care. This suggests that strategies targeted at raising awareness or addressing barriers among socially- and culturally-diverse populations are likely to be the most effective at increasing coverage.

The number of practice staff FTE remained positively associated with cervical screening coverage but not breast screening coverage after adjusting for population factors. The finding that cervical screening coverage is more likely to be influenced by general practice factors is unsurprising since many women are screened at their local practice ${ }^{25}$, and previous studies have shown the number of nurses per practice to be associated with cervical screening coverage in deprived areas ${ }^{10}$.

Coverage in London has generally been observed to be lower than the national average ${ }^{1,2}$, in spite of some other public health features (for example obesity rates) being better in London ${ }^{26}$. We found that urbanisation, ethnicity, and deprivation, largely accounted for the lower cervical screening uptake in London. For breast screening however, being located in

London, remained a strong independent negative risk factor, which warrants further investigation.

Deprivation was an independent negative risk factor for all screening groups, as also found for cervical screening by Bang and coll. ${ }^{27}$. In the cervical screening group aged 25-49, this effect was in part explained by numbers of women under 30 , as was the positive impact of lack of higher education on coverage. Cervical screening coverage has been reported to be lower in younger women ${ }^{28}$, but younger women of lower socio-economic status or with fewer educational qualifications, regardless of ethnicity, have also been shown to be positively influenced by the 2009 Jade Goody's story with respect to cervical screening behaviour ${ }^{29}$, giving hints as to potential strategies for improving uptake.

The impact of Asian, Black, or Mixed ethnic minority groups on coverage differed between programmes after controlling for other population factors. For breast screening, it was no longer significant. For cervical screening, we found it negatively influenced coverage in the age group 25-49, but was associated with greater coverage in the age group 50-64. Previously, only an overall negative overall association after adjustment for other population factors had been reported for cervical screening in women aged 25-64 ${ }^{27}$. For both programmes, and regardless of age, 'other' ethnic minority groups were still associated with poorer coverage after accounting for deprivation and urbanisation, with a particularly strong effect in breast screening. In addition, our results suggest that women of 'other' ethnic minority background, who may be well educated and living in areas with smaller Asian, Black, or Mixed ethnic minority populations, are less likely to go for screening. Arabs communities account for a moderately large subset of the 'other' ethnic minority groups (40\%), and uptake of cervical and breast screening has been shown to be low in
these populations for a number of reasons, including religious beliefs, emotional barriers (embarrassment/fear), language barriers or taboos surrounding sexual activity (for cervical screening) ${ }^{30-32}$. These communities may therefore require newly targeted interventions to promote screening.

Our correlation analyses suggest that districts with atypical coverage levels differ from one another not only in respect of a number of population- and general practice-level characteristics, but also in how these characteristics relate to each other. Correlates of deprivation in districts with relatively low adjusted coverage were substantially different from the general results, and even more so for cervical screening. In particular, the nature of the relationship between deprivation and non-White ethnicity differed, with an inverse relationship between deprivation and non-White ethnic groups among lowest-ranking districts.

Using funnel plots based on crude performance data to assess quality of care at area level may overestimate the number of "underperforming" districts, and overdispersion needs to be addressed a priori. We chose a risk adjustment approach to uncover districts with atypically high or low coverage given particular population and general practice characteristics. Districts with adjusted coverage values lying outside control limits display a behaviour which cannot solely be explained by the area-level risk factors investigated. Districts with atypically high coverage were singled out and could be investigated to identify any local health interventions and policies that might help improve coverage in districts with similar characteristics but lower performance. Unfortunately, there is a general lack of
reporting in the research literature across districts on the impact of local interventions that have been implemented to improve screening uptake (ED, unpublished PhD thesis), so identifying 'what works' is challenging.

Simultaneously, districts with atypically low coverage were distinguished from those lying within bounds after accounting for urbanisation, deprivation and ethnicity, in particular for the London region. These districts may benefit from further investigation to uncover the features driving their atypically low coverage and help design population-specific strategies. Additional risk factors that may explain low coverage, as well as differences in district performance between programmes, include the percentage of women who are disabled ${ }^{33}$, incarcerated ${ }^{34}$, have greater difficulty in accessing services as indexed by time to screening centre ${ }^{13}$, and differential utilization behaviour as a result of socio-cultural factors, such as marital status ${ }^{35}$, occupation ${ }^{36}$, sexual orientation ${ }^{37}$, and overseas birthplace or religious beliefs ${ }^{11,38}$ that might apply to particular programmes.

Our results are limited by the aggregated nature of the data, which may conceal ecological associations within districts. This could account for the weak association seen between coverage and general practice characteristics after adjustment for population factors. However, similar trends were observed when analysing general practice-level data for cervical screening coverage ${ }^{27}$.

The districts boundaries used in this study (152 PCTs) are no longer in place; however, the findings may be applied to the newly defined boundaries (210 Clinical Commissioning Groups (CCGs)) by direct mapping ${ }^{39}$.

The strength of the approach of combining risk adjustment modelling with funnel plots was to allow us to identify districts with unusual level of screening coverage after accounting for some of the important demographic and socio-economic characteristics of their populations and their primary care settings, known to affect coverage level. Such an approach could be implemented sequentially to monitor the impact of local interventions in a centralised fashion. This method could also be adapted for use with other health indicators.

Our results demonstrate that population factors largely explain the lower coverage in London. In addition, districts in London and other urban centres with specific population characteristics such as non-deprived ethnic minority groups were identified as requiring targeted intervention to improve coverage levels. Bilingual outreach and community-based advocacy, such as support from family and community leaders including GPs, has been found to be valuable in increasing uptake of cancer screening in ethnic minorities ${ }^{40}$.

We hope these results will inform the continued drive to reduce inequalities in cancer screening and avoidable deaths, and encourage implementation of targeted interventions in communities residing within districts identified as having atypically low coverage.

## Abbreviations

| CCG | Clinical Commissioning Group |
| :--- | :--- |
| CI | Confidence Interval |
| FTE | Full-Time Equivalent |
| GP | General Practitioner |
| GVIF | Generalized Variance-Inflation Factor |
| HSCIC | Health and Social Care Information Centre |
| IMD | Indices of Multiple Deprivation |
| IQR | Interquartile Range |
| LSOA | Lower Super Output Area |
| PCT | Primary Care Trust |
| SD | Standard Deviation |
| SHA | Strategic Health Authority |

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## Competing interests

None

## AUTHOR'S CONTRIBUTION

NJM performed the statistical analysis, interpreted the results and wrote the manuscript. ED carried out the data informatics and data checks, and co-wrote the manuscript. JW \& JW provided general expert guidance. SWD provided general statistical guidance. All authors reviewed and approved the final manuscript.

## Data sharing statement

Data are freely available from the HSCIC:

Cervical screening: http://www.hscic.gov.uk/catalogue/PUB10339/bres-scre-prog-eng-2011-12-tab.xls Breast screening: http://www.hscic.gov.uk/catalogue/PUB07990/cerv-scre-prog-eng-2011-12-tab.xls No additional data available.

## ETHICS APPROVAL

None

## List of Figures

Figure 1. Funnel plots of screening coverage and list of districts lying outside the 95\% control limits prior to and after risk adjustment
Top left panel. Funnel plots of screening coverage prior to any adjustment
(a) Cervical screening in women aged 25-49.
(b) Cervical screening in women aged 50-64.
(c) Breast screening in women aged 50-64.

Top right panel. Funnel plots of screening coverage after adjustment for population and general practice factors
( $a^{\prime}$ ) Cervical screening in women aged 25-49.
(b') Cervical screening in women aged 50-64.
(c') Breast screening in women aged 50-64.
_ - _ - - - 95.0\% control limits
99.8\% control limits

SHA, Strategic Health Authority; Q30, North East; Q31, North West ;Q33, East Midlands; Q34, West Midlands; Q35, East of England; Q36, London; Q37, South East Coast; Q38, South Central; Q39, South West.

Table. Number of districts lying outside the $95 \%$ control limits prior to and after risk adjustment. The number of districts within London SHA (Q36) is shown in brackets.

## Figure 2. Geographical location of atypical districts

Map. Map of PCT 2006 boundaries with districts lying below the $95 \%$ lower control limits after risk adjustment coloured in red and districts lying above the $95 \%$ upper control limits after risk adjustment coloured in green.

Table. Districts lying outside the control limits are listed with corresponding percentile given in brackets. Districts with coverage ranking among the 15 lowest- (rank $\leq 15$ ) or 15 highest (rank $\geq 137$ ) are specified. All districts lying outside the control limits had relative coverage rankings $\leq 15$ for lower $95 \%$ limit and $\geq 137$ for upper $95 \%$ limit.

SHA, Strategic Health Authority; Q30, North East; Q31, North West ;Q33, East Midlands; Q34, West Midlands; Q35, East of England; Q36, London; Q37, South East Coast; Q38, South Central; Q39, South West.

## Figure 3. Correlations between population factors overall, and among the 15 highest-

 and 15 lowest-ranking districts after risk adjustmenta-c. Correlation coefficients are displayed in each cell. a, All districts; b, 15-lowest ranking districts; c, 15 highest ranking districts.
For the 15 lowest and 15 highest-ranking districts, correlation coefficients which are significantly different from zero at the $1 \%$ level are highlighted in green for positive correlations, and in red for negative correlations.

> d. Fisher's z test for significant differences in correlation coefficients between two independent groups.
> Bold, p-values < 0.05. Italic, p-values not significant at the $10 \%$ level.
> 1, \% Deprivation; 2, \% Urbanisation; 3, \% Asian, Black or Mixed ethnic mirority groups; 4, \% 'Other' ethnic minority groups; 5, \% No higher education.

## References

1. Health \& Social Care Information Centre (HSCIC). Cervical Screening Programme, England Statistics for 2012-13. 2014;Available from http://www.hscic.gov.uk/catalogue/PUB07990/cerv_scre_prog_eng_201112_rep_v3.pdf (Accessed 14/11/14).
2. Health \& Social Care Information Centre (HSCIC). Breast Screening Programme, England Statistics for 2011-12. 2014;Available from http://www.hscic.gov.uk/catalogue/PUB13567/bres-scre-prog-eng-2012-13rep.pdf (Accessed 14/11/14).
3. Stead MJ, Wallis MG, Wheaton ME. Improving uptake in non-attenders of breast screening: selective use of second appointment. Journal of Medical Screening 1998;5(2):69-72.
4. Weller DP, Campbell C. Uptake in cancer screening programmes: a priority in cancer control. Br J Cancer 2009;101 Suppl 2:S55-9.
5. Department of Health. Improving Outcomes: a strategy for cancer. 2011;https://www.gov.uk/government/publications/the-national-cancer-strategy (Accessed 14/11/14).
6. Spiegelhalter DJ. Funnel plots for comparing institutional performance. Statistics in Medicine 2005a;24(8):1185-202.
7. Spiegelhalter DJ. Handling over-dispersion of performance indicators. Quality and Safety in Health Care 2005b;14(5):347-51.
8. Association of Public Health Observatories (APHO). Statistical process control methods in public health intelligence. Technical briefing 2: 2008;Available from http://www.apho.org.uk/resource/item.aspx?RID=39445 (Accessed 01/12/14).
9. Dover DC, Schopflocher DP. Using funnel plots in public health surveillance. Popul Health Metr 2011;9(1):58.
10. Baker D, Middleton E. Cervical screening and health inequality in England in the 1990s. J Epidemiol Community Health 2003;57(6):417-23.
11. Webb R, Richardson J, Pickles A. A population-based study of primary care predictors of non-attendance for cervical screening. Journal of Medical Screening 2004;11(3):13540.
12. Eilbert KW, Carroll K, Peach J, et al. Approaches to improving breast screening uptake: evidence and experience from Tower Hamlets. Br J Cancer 2009;101 Suppl 2:S64-7.
13. Chen C, Yu W, Huabing W, et al. Determinations of low breast screening uptake using geographically weighted regression model. GEOINFORMATICS 2012 (20th International Conference on Geoinformatics) 15-17 June 2012, Hong Kong, China;http://ieeexplore.ieee.org/xpls/icp.jsp?arnumber=6270323:1-6.
14. Association of Public Health Observatories (APHO). Urban-rural classification of PCTs (post October 2006 boundaries). 2008;Available from http://www.apho.org.uk/resource/item.aspx?RID=53312 (Accessed 14/11/14).
15. UK Department for Communities and Local Government. English indices of deprivation 2010. Statistics 2011;Available from https://www.gov.uk/government/statistics/english-indices-of-deprivation-2010 (Accessed 02/12/14).

For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml
16. Office of National Statistics (ONS). 2011 Census: KS201UK Ethnic group, local authorities in the UK - NOMIS. 2011;Available from http://www.nomisweb.co.uk/census/2011/ks201ew (Accessed 14/11/14).
17. Office for National Statistics (ONS). 2011 Census: Highest level of qualification - NOMIS. 2011;Available from http://www.nomisweb.co.uk/census/2011/qs501ew (Accessed 14/11/14).
18. Health \& Social Care Information Centre (HSCIC). NHS Staff 2002-2012, General Practice. 2013;Available from http://www.hscic.gov.uk/catalogue/PUB09536 (Accessed 14/11/14).
19. Hilbe JM, Robinson AP. Generalized Linear Models. Methods of Statistical Model Estimation: Chapman and Hall/CRC, 2013:99-120.
20. vanEngelsdorp D, Lengerich E, Spleen A, et al. Standard epidemiological methods to understand and improve Apis mellifera health. Journal of Apicultural Research 2013;52(4):1-16.
21. McCullagh P, Nelder, JA. Generalized Linear Models. 2nd ed. London: Chapman \& Hall, 1989.
22. Kabacoff RI. R in Action: Data Analysis and Graphics with R. Shelter Island, NY, USA: Manning Publications Co., 2011.
23. Diedenhofen B. cocor: Comparing correlations (Version 1.0-0). Compare two correlations based on independent groups 2013;Available from http://r.birkdiedenhofen.de/pckg/cocor/ (Accessed 14/11/14):18-21.
24. Gelman A, Hill J. Generalized linear models. Data Analysis Using Regression and Multilevel/Hierarchical Models: Cambridge University Press, 2006:117.
25. Public Health England. NHS Cervical Screening Programme. 2014;Available from http://www.cancerscreening.nhs.uk/cervical/index.html (Accessed 14/11/14).
26. Scholes S, Prescott P, Bajekal M. Health \& lifestyle indicators for Strategic Health Authorities 1994-2002. Mean diastolic blood pressure. Health Survey for England 2014;Available from http://www.dh.gov.uk/prod_consum_dh/idcplg?IdcService=GET_FILE\&dID=1084\& Rendition=Web (Accessed 14/11/14).
27. Bang JY, Yadegarfar G, Soljak M, et al. Primary care factors associated with cervical screening coverage in England. J Public Health (Oxf) 2012;34(4):532-8.
28. Waller J, Jackowska M, Marlow L, et al. Exploring age differences in reasons for nonattendance for cervical screening: a qualitative study. BJOG 2012;119(1):26-32.
29. Marlow LAV, Sangha A, Patnick J, et al. The Jade Goody Effect: whose cervical screening decisions were influenced by her story? Journal of Medical Screening 2012;19(4):184-88.
30. Szarewski A, Cadman L, Ashdown-Barr L, et al. Exploring the acceptability of two selfsampling devices for human papillomavirus testing in the cervical screening context: a qualitative study of Muslim women in London. Journal of Medical Screening 2009;16(4):193-98.
31. Donnelly TT, Khater AH, Al-Bader SB, et al. Arab women's breast cancer screening practices: a literature review. Asian Pac J Cancer Prev 2013;14(8):4519-28.
32. Padela AI, Murrar S, Adviento B, et al. Associations Between Religion-Related Factors and Breast Cancer Screening Among American Muslims. Journal of immigrant and minority health / Center for Minority Public Health 2014.
33. Horner-Johnson W, Dobbertin K, Andresen EM, et al. Breast and Cervical Cancer Screening Disparities Associated with Disability Severity. Women's Health Issues 2014;24(1):e147-e53.
34. Plugge E, Fitzpatrick R. Factors affecting cervical screening uptake in prisoners. J Med Screen 2004;11(1):48-9.
35. Lo SH, Waller J, Wardle J, et al. Comparing barriers to colorectal cancer screening with barriers to breast and cervical screening: a population-based survey of screening-age women in Great Britain. Journal of Medical Screening 2013;20(2):73-79.
36. Moser K, Patnick J, Beral V. Inequalities in reported use of breast and cervical screening in Great Britain: analysis of cross sectional survey data. BMJ 2009;338:b2025.
37. Reiter PL, McRee AL. Cervical cancer screening (Pap testing) behaviours and acceptability of human papillomavirus self-testing among lesbian and bisexual women aged 21-26 years in the USA. The journal of family planning and reproductive health care / Faculty of Family Planning \& Reproductive Health Care, Royal College of Obstetricians \& Gynaecologists 2014.
38. Szczepura A, Price C, Gumber A. Breast and bowel cancer screening uptake patterns over 15 years for UK south Asian ethnic minority populations, corrected for differences in socio-demographic characteristics. BMC Public Health 2008;8(1):346.
39. Allies computing. Translate PCTs to CCGs conversion tool. 2014;Available from http://www.alliescomputing.com/innovation/pct-ccg-mapping (Accessed 14/11/14).
40. Shankleman J, Massat NJ, Khagram L, et al. Evaluation of a service intervention to improve awareness and uptake of bowel cancer screening in ethnically-diverse areas. Br J Cancer 2014;111(7):1440-47.


Figure1
$190 \times 254 \mathrm{~mm}$ ( $300 \times 300$ DPI)


Figure2
$190 \times 254 \mathrm{~mm}$ ( $300 \times 300$ DPI)

b. 15 lowest-ranking districts

c. 15 highest-ranking districts


Cervical age group 50-64


d. Differences in correlates between the 15 lowest- and the 15 highest-ranking districts Cervical age group 25-49 Cervical age group 50-64 Breast age group 50-64


Figure 3 revised - TIFF version $190 \times 254 \mathrm{~mm}$ ( $300 \times 300$ DPI)

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## Supplementary file (Massat, Douglas et al.)

Table A1. Correlations between population and general practice factors, and screening coverage (all districts)
Upper diagonal: Correlation coefficient; Lower diagonal: $p$-value of test for significant correlation between pairés samples

|  | Population factors |  |  |  |  |  | General practice factors? |  |  |  |  | Coverage |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | \% <br> Urbanization | \% Deprivation | \% Asian, Black, or Mixed ethnicity | \% 'Other' ethnicity | \% No higher education | \% Registered women aged 25-29 | Average practice list size | \% Singlehanded practices | Practitioner headcount per $10^{5}$ population |  | \% <br> Practitioners qualified outside UK | Cervical group aged 25-49 | Cervical group aged 50-64 | Breast group aged 50-64 |
| Population factors |  |  |  |  |  |  |  |  |  | $\stackrel{8}{2}$ |  |  |  |  |
| \% Urbanization | 1 | 0.61 | 0.56 | 0.48 | -0.002 | 0.63 | -0.50 | 0.49 | 0.02 | $\overrightarrow{\hat{\mathrm{o}}}-0.51$ | 0.42 | -0.60 | -0.49 | -0.66 |
| \% Deprivation | < 0.001 | 1 | 0.58 | 0.39 | 0.41 | 0.64 | -0.58 | 0.46 | 0.23 | $\underset{\underset{y}{3}}{\substack{3}}-0.36$ | 0.54 | -0.56 | -0.47 | -0.58 |
| \% Asian, Black, or Mixed ethnicity | < 0.001 | < 0.001 | 1 $<0001$ | 0.70 | $-0.27$ | 0.62 | -0.35 | 0.39 | 0.14 | $\text { 華 }-0.25$ | 0.43 | -0.78 | -0.24 | -0.68 |
| \% 'Other' ethnicity | < 0.001 | < 0.001 | < 0.001 | 1 | -0.45 | 0.60 | -0.40 | 0.38 | 0.14 | 3-0.25 | 0.22 | -0.78 | -0.45 | -0.74 |
| \% No higher education | NS (0.9) | < 0.001 | 0.001 | < 0.001 | 1 | -0.11 | -0.26 | 0.21 | -0.25 | $\text { 울 }^{-0.04}$ | 0.37 | 0.31 | 0.02 | 0.31 |
| \% Registered women aged 25-29 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | NS (0.2) | 1 | -0.33 | 0.30 | 0.26 | $\begin{aligned} & \text { خे } \\ & \ddot{⿳ ㇒}^{-0.24} \end{aligned}$ | 0.21 | -0.71 | -0.43 | -0.69 |
| General practice factors |  |  |  |  |  |  |  |  |  | ¢ |  |  |  |  |
| Average practice list size | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.001 | < 0.001 | 1 | -0.75 | 0.06 | $\bigcirc 0.36$ | -0.52 | 0.42 | 0.39 | 0.49 |
| \% Single-handed practices | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.01 | < 0.001 | < 0.001 | 1 | -0.35 | $\begin{aligned} & \text { 윽 -0.27 } \\ & D \end{aligned}$ | 0.61 | -0.41 | -0.28 | -0.43 |
| Practitioner headcount per $10^{5}$ population | NS (0.8) | 0.006 | NS (0.1) | NS (0.08) | 0.002 | 0.001 | NS (0.5) | < 0.001 | 1 | 을.06 | -0.35 | -0.19 | -0.14 | -0.22 |
| Practice staff FTE | < 0.001 | < 0.001 | 0.002 | 0.002 | NS (0.7) | 0.004 | < 0.001 | 0.001 | NS (0.4) | $\bigcirc 1$ | -0.36 | 0.34 | 0.37 | 0.35 |
| \% Practitioners qualified outside UK | < 0.001 | < 0.001 | < 0.001 | 0.006 | < 0.001 | 0.009 | < 0.001 | < 0.001 | < 0.001 | $\begin{aligned} & \mathrm{N}_{2} 0.001 \\ & \mathrm{~N} \\ & \hline \end{aligned}$ | 1 | -0.29 | -0.18 | -0.32 |
| Coverage |  |  |  |  |  |  |  |  |  | ¢ |  |  |  |  |
| Cervical group aged 25-49 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.02 | ${\stackrel{\underset{\widetilde{D}}{\mid}}{\stackrel{Q}{\widetilde{D}}}}_{<0.001}$ | < 0.001 | 1 | 0.68 | 0.84 |
| Cervical group aged 50-64 | < 0.001 | < 0.001 | 0.004 | < 0.001 | NS (0.8) | < 0.001 | < 0.001 | < 0.001 | NS (0.09) |  | 0.03 | < 0.001 | 1 | 0.65 |
| Breast group aged 50-64 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | 0.006 | $\begin{aligned} & \text { प्र }^{2} 0.001 \\ & \stackrel{\rightharpoonup}{\oplus} \\ & \stackrel{\rightharpoonup}{D} \end{aligned}$ | < 0.001 | < 0.001 | < 0.001 | 1 |
| NS: not significant at the 5\% level; FTE: Full Time Equivalent |  |  |  |  |  |  |  |  |  |  |  |  | a g e |  |

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Table A2.1. Regression modelling for cervical screening coverage among women aged 25-49, including \% registered women aged 25-29.


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Table A3.1. 15 lowest-ranking PCTs prior to and after adjustment for population and general practice factors

|  | $\begin{aligned} & \text { Rank } \\ & (/ 151) \end{aligned}$ | No adjustment (percentile) | Adjustment for Population \& General practice factors (percentile) |
| :---: | :---: | :---: | :---: |
| Cervical group aged 25-49 | 137 | 5MX | 5A9* |
|  | 138 | 5LD* | 5LA* |
|  | 139 | 5AT* | 5HY* |
|  | 140 | 5C3* | 5PN |
|  | 141 | 5K5* | 5PG |
|  | 142 | 5C4* | 5NL |
|  | 143 | 5C5* ${ }^{\text {(2.79\%) }}$ | TAP |
|  | 144 | 5A9* ${ }^{\text {( }}$ (2.65\%) | 5FE |
|  | 145 | 5HY* \$ (2.04\%) | 5AT* |
|  | 146 | $5 L C^{*}(1.56 \%)$ | 5 KM |
|  | 147 | 5HX* ${ }^{\text {(1.11\%) }}$ | 5NJ |
|  | 148 | 5K6* ${ }^{\text {( }}$ (0.86\%) | 5HP |
|  | 149 | 5LA* ${ }^{\text {( }}$ (0.69\%) | $5 K 6 * \text { * }(0.22 \%)$ |
|  | 150 | 5H1* \$\$ (0.04\%) | $5 \mathrm{H} 1^{* \$(0.01 \%)}$ |
|  | 151 | 5K7* \$ ${ }^{\text {(0.04\%) }}$ | 5K7* \$ ${ }^{\text {(0.001\%) }}$ |
| Cervical group aged 50-64 | 137 | 5LG* | 5LD* |
|  | 138 | 5 J 4 | 5M7* |
|  | 139 | 5M1 | 5HY* |
|  | 140 | 5C3* | 5NT |
|  | 141 | 5HP | 5K6* |
|  | 142 | 5PG ${ }^{\text {\$ }}$ (2.70\%) | 5M1 |
|  | 143 | 5NT ${ }^{\text {\$ }}$ (2.09\%) | 5LG |
|  | 144 | 5NL ${ }^{\text {\$ }}$ (1.92\%) | TAM |
|  | 145 | 5K7* | 5KM |
|  | 146 | 5 NJ \$ $0.98 \%$ ) | 5K7* |
|  | 147 | 5 KM | 5LA* |
|  | 148 | 5LC* ${ }^{\text {( }}$ ( $0.31 \%$ ) | $5 \mathrm{PG}^{\$}(1.20 \%)$ |
|  | 149 | 5D9 | $5 \mathrm{NJ} \text { \$ }(0.30 \%)$ |
|  | 150 | 5LA* ${ }^{\text {( }}$ (0.08\%) | 5H1* |
|  | 151 | 5H1* (0.02\%) | 5D9 |
| Breast group aged 50-64 | 137 | 5K5* | 5F5 |
|  | 138 | 5A8* | 5FL |
|  | 139 | 5HP | 5NH |
|  | 140 | 5C4* | 5NG |
|  | 141 | 5LF* | 5LF* |
|  | 142 | 5LG* | 5LQ |
|  | 143 | 5C3* | 5H1* |
|  | 144 | $5 \mathrm{NT}^{\text {\$ }}$ (0.89\%) | TAP |
|  | 145 | 5C9* | 5LE* |
|  | 146 | 5LE* | 5LG* ${ }^{\text {( }} 0.42 \%$ ) |
|  | 147 | 5H1* | 5HP |
|  | 148 | 5LC* | $5 \mathrm{NT}^{\text {\$ }}$ (0.22\%) |
|  | 149 | 5LD* ${ }^{\text {( }}$ (0.17\%) | 5LA* |
|  | 150 | 5K7* ${ }^{\text {( }}$ (0.09\%) | 5K7* ${ }^{\text {( }}$ (0.02\%) |
|  | 151 | 5LA* ${ }^{\text {( }} 0.03 \%$ ) | 5LD* ${ }^{\text {(0.01\% }}$ ) |

[^3]Table A3.2. 15 highest-ranking PCTs prior to and after adjustment for population and general practice factors

|  | $\begin{aligned} & \text { Rank } \\ & (/ 151) \end{aligned}$ | No adjustment (percentile) | Adjustment for Population \& General practice factors (percentile) |
| :---: | :---: | :---: | :---: |
| Cervical group aged 25-49 | 1 | 5N8 ${ }^{\text {s }}$ (94.4\%) | 5C1* ${ }^{\text {(99.8\%) }}$ |
|  | 2 | 5N6 \$ (93.9\%) | TAN |
|  | 3 | TAC | 5C9* |
|  | 4 | 5A3 | 5MX |
|  | 5 | 5NW | 5D8 |
|  | 6 | TAN | 5K5* |
|  | 7 | 5QM | TAK* |
|  | 8 | 5D8 | 5A3 |
|  | 9 | 5 J 6 | 5N8 \$ (93.9\%) |
|  | 10 | 5 ET | 5A7* |
|  | 11 | 5QH | 5 J 6 |
|  | 12 | 5PA | 5C5* |
|  | 13 | 5 EF | 5 J 9 |
|  | 14 | 5JE | 5LC* |
|  | 15 | 5P9 | 5 KL |
| Cervical group aged 50-64 | 1 | 5 A 3 | 5C1* ${ }^{\text {\% }}$ (99.9\%) |
|  | 2 | 5N8 \$ ${ }^{\text {( }} 96.4 \%$ ) | TAN \$ $(99.8 \%)$ |
|  | 3 | 5PA \$ (95.8\%) | 5NC* ${ }^{\text {( }}$ (99.3\%) |
|  | 4 | TAN | 5A3 |
|  | 5 | 5NA* | $5 \mathrm{KL}{ }^{\text {\$ }}$ (98.6\%) |
|  | 6 | 5P2 | 5MX |
|  | 7 | 5QF | 5JE |
|  | 8 | 5N2 | 5C9* |
|  | 9 | 5 QH | 5NA* |
|  | 10 | 5N6 \$ (91.8\%) | 5N7 |
|  | 11 | 5QM | 5 J 6 |
|  | 12 | 5 J 6 | 5N8 \$ (93.9\%) |
|  | $13$ | $5 P 9$ | 5QN |
|  | $14$ | $5 \text { QV }^{\$}(89.8 \%)$ | 5N2 |
|  | $15^{\#}$ | 5FL | 5F1 |
| Breast group aged 50-64 | 1 | 5PA ${ }^{\text {s }}$ (97.4\%) |  |
|  | 2 | 5M2 | $5 N 7 \text { (99.7\%) }$ |
|  | 3 | TAC | 5 N 4 \$ $(98.5 \%)$ |
|  | 4 | 5PL \$ (93.1\%) | 5A4* |
|  | 5 | 5 N 8 \$ $(93.0 \%)$ | 5F1 |
|  | 6 | 5N6 \$ (92.9\%) | 5PC |
|  | 7 | 5 CN | 5NC* |
|  | 8 | 5 NV \$ (92.5\%) | 5JE |
|  | 9 | 5JE | 5H8 |
|  | 10 | 5QD | 5PA \$ ${ }^{\text {(96.5\%) }}$ |
|  | 11 | 5PX | 5KL |
|  | 12 | 5N7 | 5 PJ |
|  | 13 | 5PT | 5N8 \$ (93.5\%) |
|  | 14 | 5PW | TAK* |
|  | 15 | 5H8 | 5MK |

PCT: Primary Care Trust; SHA: Strategic Health Authority
Dark green: PCTs lying above the $95 \%$ upper control limits using full model (atypical PCTs).
Light green: Atypical PCTs found among PCTs with highest relative coverage prior to adjustment.
Percentile is given for those PCTs lying above the $95 \%$ upper control limits prior to and after full adjustment.

* PCT in London SHA (Q36)
${ }^{\$}$ PCT lying between the $95 \%$ and $99.8 \%$ upper control limits
\$\$ PCT lying above the $99.8 \%$ upper control limits
\# The $17^{\text {th }}$ highest-ranking PCT (5QC) also lay above the $95 \%$ upper control limit (percentile 89.1\%)

STROBE Statement-checklist of items that should be included in reports of observational studies

|  | $\begin{gathered} \text { Item } \\ \text { No } \\ \hline \end{gathered}$ | Recommendation | Checklist |
| :---: | :---: | :---: | :---: |
| Title and abstract | 1 | (a) Indicate the study's design with a commonly used term in the title or the abstract | Yes |
|  |  | (b) Provide in the abstract an informative and balanced summary of what was done and what was found | Yes |
| Introduction |  |  |  |
| Background/rationale | 2 | Explain the scientific background and rationale for the investigation being reported | p.5-6 |
| Objectives | 3 | State specific objectives, including any pre-specified hypotheses | p. 6 |
| Methods |  |  |  |
| Study design | 4 | Present key elements of study design early in the paper | p. 6 |
| Setting | 5 | Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection | p. 6 |
| Participants | 6 | (a) Cohort study-Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up <br> Case-control study-Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controls Cross-sectional study-Give the eligibility criteria, and the sources and methods of selection of participants <br> (b) Cohort study-For matched studies, give matching criteria and number of exposed and unexposed <br> Case-control study-For matched studies, give matching criteria and the number of controls per case | p. 6 |
| Variables | 7 | Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. <br> Give diagnostic criteria, if applicable | p. 6 |
| Data sources/ measurement | 8* | For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group | p. 6 |
| Bias | 9 | Describe any efforts to address potential sources of bias | p. 8 |
| Study size | 10 | Explain how the study size was arrived at | p. 6 |
| Quantitative variables | 11 | Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why | p. 8 |
| Statistical methods | 12 | (a) Describe all statistical methods, including those used to control for confounding | p. 8 |
|  |  | (b) Describe any methods used to examine subgroups and interactions | p. 8 |
|  |  | (c) Explain how missing data were addressed |  |
|  |  | (d) Cohort study-If applicable, explain how loss to follow-up was addressed <br> Case-control study-If applicable, explain how matching of cases and controls was addressed Cross-sectional study-If applicable, describe analytical methods taking account of sampling strategy | p.6-7 |
|  |  | (e) Describe any sensitivity analyses | p. 12 |

Continued on next page
*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.


[^0]:    d. Fisher's z test for significant differences in correlation coefficients between two independent groups.
    Bold, p-values < 0.05. Italic, p-values not significant at the $10 \%$ level.
    1, \% Deprivation; 2, \% Urbanisation; 3, \% Asian, Black or Mixed ethnic mirority groups; 4, \% Other ethnic minority groups; 5, \% No higher education.

[^1]:    Cl: Confidence Interval; FTE: Full-Time Equivalent; NS: Considered non-significant (see Methods for details); SHA: Strategic Health Authority
    \$ The variance of the coefficient estimate is being inflated by multicollinearity with other factors ( $\sqrt{\text { GVIF }}=2.8$ ).

[^2]:    PCT: Primary Care Trust; SHA: Strategic Health Authority
    Dark green: PCTs lying above the $95 \%$ upper control limits using full model (atypical PCTs).
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    Percentile is given for those PCTs lying above the $95 \%$ upper control limits prior to and after full adjustment.

    * PCT in London SHA (Q36)
    \$ PCT lying between the $95 \%$ and $99.8 \%$ upper control limits
    \$ PCT lying above the 99.8\% upper control limits
    \# The $17^{\text {th }}$ highest-ranking PCT (5QC) also laid above the $95 \%$ upper control limit (percentile 89.1\%)

[^3]:    PCT: Primary Care Trust; SHA: Strategic Health Authority
    Dark red: PCTs lying below the $95 \%$ lower control limits using full model (atypical PCTs).
    Orange: Atypical PCTs found among PCTs with lowest relative coverage prior to adjustment.
    Percentile is given for those PCTs lying below the $95 \%$ lower control limits prior to and after full adjustment.

    * PCT in London SHA (Q36)
    \$ PCT lying between the $95 \%$ and $99.8 \%$ lower control limits
    ${ }^{\$ \$}$ PCT lying below the $99.8 \%$ lower control limits

