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REVIEW

Low levels of HIV test coverage in clinical settings in the UK: a systematic review of adherence to 2008 guidelines

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ABSTRACT

Objectives To quantify the extent to which guideline recommendations for routine testing for HIV are adhered to outside of genitourinary medicine (GUM), sexual health (SH) and antenatal clinics.

Methods A systematic review of published data on testing levels following publication of 2008 guidelines was undertaken. Medline, Embase and conference abstracts were searched according to a predefined protocol. We included studies reporting the number of HIV tests administered in those eligible for guideline recommended testing. We excluded reports of testing in settings with established testing surveillance (GUM/SH and antenatal clinics). A random effects meta-analysis was carried out to summarise level of HIV testing across the studies identified.

Results Thirty studies were identified, most of which were retrospective studies or audits of testing practice. Results were heterogeneous. The overall pooled estimate of HIV test coverage was 27.2% (95% CI 22.4% to 32%). Test coverage was marginally higher in patients tested in settings where routine testing is recommended (29.5%) than in those with clinical indicator diseases (22.4%). Provider test offer was found to be lower (40.4%) than patient acceptance of testing (71.5%).

Conclusions Adherence to 2008 national guidelines for HIV testing in the UK is poor outside of GUM/SH and antenatal clinics. Low levels of provider test offer appear to be a major contributor to this. Failure to adhere to testing guidelines is likely to be contributing to late diagnosis with implications for poorer clinical outcomes and continued onwards transmission of HIV. Improved surveillance of HIV testing outside of specialist settings may be useful in increasing adherence testing guidelines.

INTRODUCTION

In 2011, there were an estimated 96 000 people living with HIV in the UK with almost one in four thought to be unaware of their infection.¹ In the same year, 6280 individuals were newly diagnosed and 47% of these cases were at a late stage of infection (CD4 cell count <350 cells/mm³).¹ Patients diagnosed late experience a higher risk of developing AIDS and a tenfold increased risk of death within a year of diagnosis.^{2–4} Timely identification of those who are HIV-positive and appropriate referral into care services is essential not only for the reduction of HIV-associated morbidity and mortality but also for the prevention of onwards transmission of the virus. Reducing late HIV

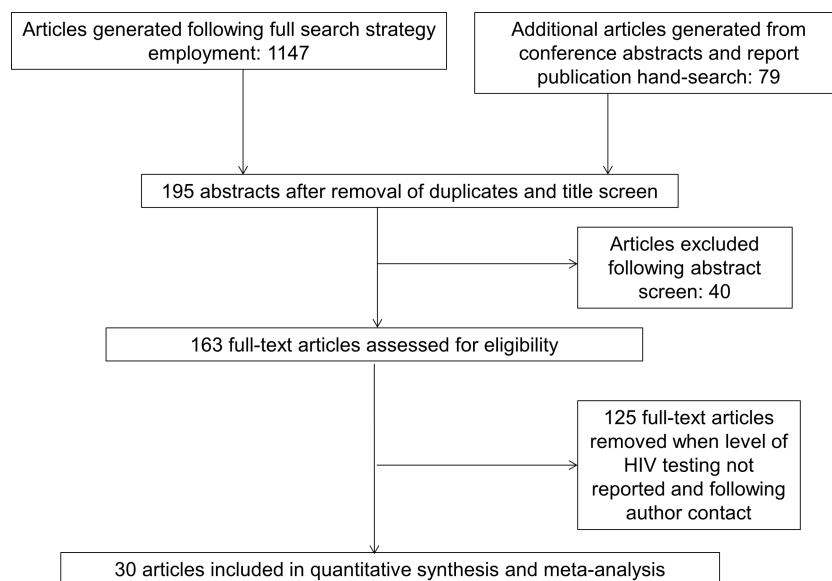
diagnosis is a key indicator of the Public Health Outcomes Framework set by the Department of Health (DoH),⁵ and the primary means of achieving this is widespread testing in populations and settings at increased risk of HIV infection.

The latest national guidelines on HIV testing were published in October 2008. The guidelines were published by the British HIV Association (BHIVA) and written in collaboration with the British Infection Society (BIS) and the British Association for Sexual Health and HIV (BASHH). These guidelines were intended to promote an increase in HIV testing in all healthcare settings to reduce the proportion of individuals with undiagnosed HIV infection. The authors of the guidelines state the reason for the need of their publication as being (a) misconceptions regarding HIV testing remaining a hindrance to increased testing; (b) the importance of both the individual patient and public health benefits of increased testing and (c) the need for up-to-date guidance that would enable any clinician to perform an HIV test within good clinical practice, thereby encouraging the 'normalisation' of HIV testing.⁶ These guidelines recommended HIV testing in a wider range of clinical settings and populations including those with indicator diseases, all medical admissions and new registrants in primary care in areas with a diagnosed adult HIV prevalence of greater than 2 per 1000 population (please see online appendix a, supplementary data). The guidelines have additionally been endorsed by the National Institute of Health and Care Excellence and incorporated into their pathway for strategy, policy and commissioning on HIV testing and prevention.^{7 8}

There have been improvements in earlier diagnosis for HIV in the UK. According to Public Health England figures, there has been a gradual decrease in the proportion of people diagnosed late with HIV in the UK, from 60% in 2002 to 47% in 2011¹; however, this remains high, suggesting that testing programmes continue to miss people at an early stage in their infection. HIV testing is routinely monitored in genitourinary medicine (GUM), sexual health (SH) and antenatal clinics (ANC) where uptake is high, with 70% of GUM and 97% of ANC attendees being tested for HIV in 2010; these locations account for 47% and 31% of total HIV tests in the UK, respectively. However, there is no routine monitoring of testing in other (non-specialist) clinical settings or populations, and

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Figure 1 Flowchart of search results and selection of papers.



therefore adherence to national guidelines is unknown. To inform decision making about future HIV-testing initiatives, we reviewed evidence of adherence to national guidelines in settings not covered by existing surveillance.

METHODS

Search strategy and inclusion/exclusion criteria

We carried out a systematic review and meta-analysis on levels of adherence to national guideline recommended HIV testing in non-specialist settings. A predefined protocol (available as online supplementary file 'Review Protocol') detailing inclusion and exclusion criteria was developed; two authors (RE and SMG) independently used a set combination of terms (HIV, human immunodeficiency virus test*, screen*, diagnos*, United Kingdom, UK, England, Northern Ireland, Scotland, Wales, Britain, British, English, Scottish, Welsh, Northern Irish) to search MEDLINE, Embase, Maternity and Infant Care and PsychINFO databases via the search engine Ovid. The final search was run on 28 February 2013. In addition, bibliographies from eligible papers, conference abstracts and grey literature (including relevant reports^{7 8}) were hand searched. Studies were included if they measured HIV test coverage in a defined, eligible population. Studies were excluded if they related to testing in GUM/SH or ANC (specialist) clinics, included data from before September 2008 or were conducted outside the UK. Studies not measuring HIV testing levels as an outcome were also excluded as were those measuring HIV testing in community settings as, although testing in these settings is encouraged, it is not explicitly recommended in UK national guidelines. In order to identify as wide a range of studies measuring HIV testing levels as possible, all quantitative study designs and methodologies were included. Where key information for article inclusion was missing, an online search for conference presentations/posters was performed and authors were contacted for additional data. Articles were only excluded after the deadline period for author reply had passed.

Data extraction

Data extraction was undertaken separately by two authors, and information on the following variables was retrieved: author(s), exposure status or risk group (if applicable), primary HIV testing outcome (how receipt of HIV test was confirmed),

exclusion criteria, time period and duration of data collection, population, setting (and Public Health England estimations for diagnosed HIV prevalence per 1000 population 15–59 year olds), type and number of centres, study design and methods, measure or reporting method, type of test used, method of service delivery, opt-in/opt-out model, number of patients eligible for testing, number offered testing, number tested and number with positive test result.

Data analysis

Studies were classified into two groups according to patient population or setting where testing took place: persons diagnosed with a disease indicative of HIV infection and persons attending a setting where routine HIV screening should be undertaken (excluding GUM/SH and ANC settings) (see online appendix a, supplementary data).⁵ Test coverage, defined as the percentage of those eligible for HIV testing who were offered and accepted an HIV test, was calculated for each study identified. Additional outcomes including (a) test offer level defined as the percentage of those eligible for testing who were offered a test, (b) test acceptance level (defined as the percentage of those offered an HIV test who were tested) and (c) seroprevalence level (defined as the percentage of those testing positive for HIV) were calculated where this information was available. Using a random effects model, stratified analyses were performed by group. Clopper–Pearson 95% CIs were calculated for each study input. CIs for these results were capped at 0% and 100% for presentation of pooled estimates as percentages.⁸ Cochran's test of heterogeneity (Q statistic) and I^2 statistic was used to assess the presence of and quantify the extent of between-study heterogeneity in testing prevalence estimates.⁹

Univariate meta-regression was used to investigate heterogeneity in overall testing coverage. There were too few studies to explore this for the other outcomes. Proportions were transformed to logits using a continuity correction of 0.1% where the number of patients tested for HIV was either equal to zero or the number of eligible patients.¹⁰ If a covariate was significantly associated with the prevalence estimates, the percentage of between-study variability explained by the covariate (R^2) was calculated as $100 \times (1 - (\tau^2 \text{ regression model with covariate} / \tau^2 \text{ regression without covariate}))$. Analyses were completed in STATA v.11.0 (StataCorp, College station, Texas, USA).

Table 1 Percentage of eligible patients who received HIV tests, plus percentages offered, accepted and HIV prevalence in those tested: summary results from random effects model meta-analysis patient group

Patient group	Percentage of those eligible who received HIV test (95% CI)	N studies	Percentage of those eligible who were offered HIV test (95% CI)	N studies	Percentage of those offered HIV test who accepted (95% CI)	N studies	Percentage of those tested who were HIV-positive (95% CI)	N studies
Patients diagnosed with indicator disease	22.4% (13.9% to 30.9%)	10	9.3% (1.2% to 17.3%)	2	87.4% (57.7% to 100%)	2	2.7% (1.1% to 4.4%)	6
Persons attending screening settings	29.5% (23.6% to 35.4%)	20	45.5% (28% to 63%)	12	69.2% (52.7% to 85.6%)	12	0.4% (0.2% to 0.6%)	17
Overall	27.2% (22.4% to 32%) $I^2=99.9\%$	30	40.4% (24.3% to 56.7%) $I^2=100\%$	14	71.5% (50% to 86.9%) $I^2=99.8\%$	14	0.5% (0.3% to 0.7%) $I^2=51.5\%$	23

95% CIs were bounded to between 0.00% and 100.00% as data are presented as a percentage. For test strategy and type of HIV test, some studies were excluded from the subgroup analyses due to lack of data.

RESULTS

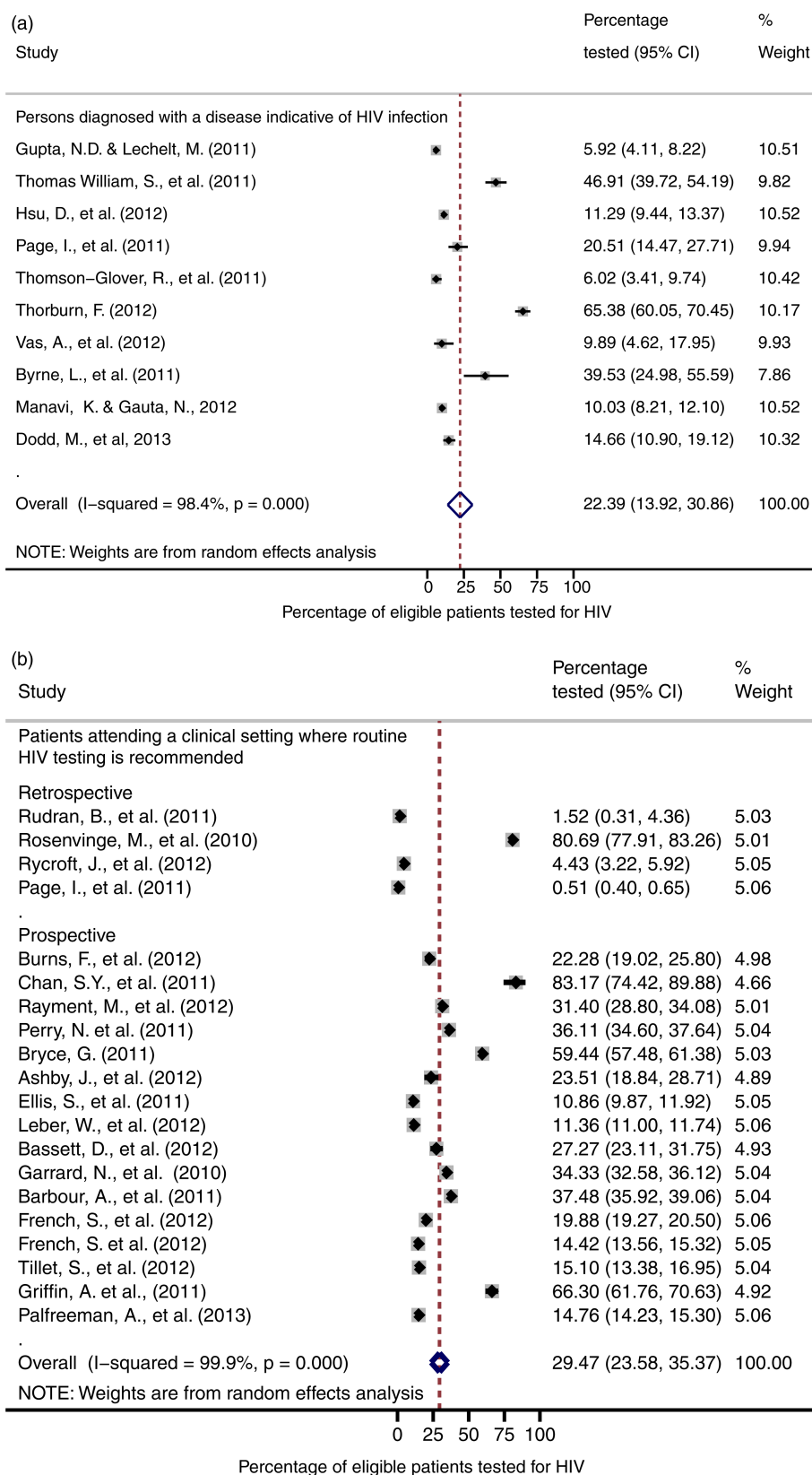
The search identified 1226 references that were screened; after exclusion of duplicates and undertaking a title and abstract screen, 163 full-text articles were evaluated for full inclusion. Of these, 30 reports that measured levels of HIV testing in a range of recommended settings were identified (figure 1). Fourteen were cross-sectional studies or retrospective studies (audits) from hospital settings using either case note review or extraction of data from electronic or paper records. Data from 12 were in journal publications, and data from the remaining 18 studies were extracted from published reports or conference abstracts. Ten studies were in patients diagnosed with an indicator disease and 20 in people attending services where routine HIV testing was recommended due to diagnosed prevalence in the local population. Information for all 30 studies identified can be found in online supplementary data file appendix b: characteristics of studies included: methods, measures and testing levels and appendix c: supplementary data tables for studies identified by group.

Table 1 shows the pooled estimates for the percentage of eligible patients who received HIV tests in the two groups, the percentage who were offered and accepted testing, and HIV prevalence from those studies that reported these outcomes. There was considerable heterogeneity within and between the patient groups, with an overall pooled estimate of 27.2% (95% CI 22.4% to 32.0%) of those eligible being tested. This level of heterogeneity is illustrated in the forest plots (figure 2A,B) for both groups. The higher pooled estimate of the two was 29.5% (95% CI 23.6% to 35.4%), with individual results ranging from 0.5% (95% CI 0.4% to 0.7%) in Page *et al*¹¹ to 83.2% (95% CI 74.4% to 89.9%) in Chan *et al*¹² (figure 2B). Testing was less likely in patients with diseases indicative of HIV infection, with a pooled estimate of 22.4% (95% CI 13.9% to 30.9%) and result ranging from 5.9% (95% CI 4.1% to 8.2%) in Gupta and Lechelt to 65.4% (95% CI 60.1% to 70.5%) in Thorburn *et al*¹³. There was considerable between-study heterogeneity across studies as illustrated by the I^2 statistic value, which was consistently over 97%.

A meta-regression was undertaken to explore other study characteristics that may contribute to the heterogeneity, and the results are presented in table 2. Covariates assessed as contributors to heterogeneity include location of testing (London or non-London), type of test administered (laboratory serological or point-of-care testing), testing strategy (opt-in or opt-out), service model (standard practice, staff training or GUM specialist testing) and study type (retrospective or prospective). None of these factors appeared to contribute significantly to the level of heterogeneity, and the results here should be interpreted with caution as many of the variables had small sample sizes, for example, there were only three studies in the 'GUM specialist testing' group of the service model. In a separate meta-regression model looking at contribution of study type in test coverage level exclusively in persons attending screening settings (the only group to have both retrospective and prospective study types), study type was found to be a significant contributor to the level of heterogeneity seen in test coverage in these studies (OR 6.3, 95% CI 1 to 38.4). The adjusted R^2 for this meta-regression indicates that 15.1% of between-study variance in the pooled estimate for this testing across studies could be explained by study type in this group. The result here should again be carefully interpreted as only four studies were included in the 'Retrospective' group of this model.

Fourteen studies reported both the number of tests being offered to those eligible and the number of those offered tests who were tested. The pooled estimate for HIV test offer level is

Figure 2 (A, B) Forest plots of percentage of eligible patients tested by group (A). Patients with an indicator disease (B). Patients attending a clinical setting where routine HIV testing is recommended (excluding genitourinary medicine/sexual health and antenatal clinics).



at 40.4% (95% CI 24.3% to 56.7%) and the pooled estimate for HIV uptake levels is 71.5% (95% CI 56.0% to 86.9%). These results are presented in table 1 and show that the pooled estimate of eligible people offered an HIV test was 9.3% (95% CI 1.2% to 17.3%) in patients diagnosed with an indicator

disease, which was lower than that seen in persons attending screening settings at 29.5% (95% CI 23.6% to 35.4%). Uptake, that is, the percentage of those offered testing who accepted was 69.2% (95% CI 52.8% to 85.6%) in persons attending screening settings and 87.4% (95% CI 57.7% to 100.0%) in

Table 2 Predictors of HIV testing rates among eligible patients: meta-regression of results from studies identified

Covariate	N studies	OR (95% CI)	p Value
Patient group			
Patients presenting with indicator disease conditions	10	0.8 (0.2 to 2.6)	0.67
Persons attending screening settings	20	1 (ref)	
Location of study			
London	14	1 (ref)	
Non-London	16	0.5 (0.2 to 1.6)	0.26
Type of HIV test			
Laboratory	7	1 (ref)	
Point-of-care	6	0.8 (0.2 to 4)	0.75
Service model*			
Usual practice	18	1 (ref)	
Added staff training	9	1.3 (0.4 to 4.8)	0.62
GUM specialist testing	3	3.5 (0.5 to 24)	0.2
Testing strategy			
Opt-out	12	1 (ref)	
Opt-in	9	1 (0.3 to 3.2)	0.99
Study type			
Retrospective	14	1 (ref)	
Prospective	16	2.6 (0.9 to 7.7)	0.08

*With only three studies in one of the categories, this result should be interpreted with caution due to lack of power.
GUM, genitourinary medicine.

patients diagnosed with indicator diseases, indicating that test offer is lower for patients with indicator diseases despite a higher test acceptance level in this group.

Of the 30 studies, 23 reported the number of those patients who tested positive for HIV, and the meta-analysis results for the seroprevalence observed in these studies are also presented in table 1. The pooled seroprevalence was 0.5% (95% CI 0.3% to 0.7%), with a higher seroprevalence seen in patients diagnosed with an indicator disease (2.7%, 95% CI 1.1% to 4.4%) than those tested in screening settings (0.4%, 95% CI 0.2% to 0.6%).

DISCUSSION

In this review, we found that the estimated percentage of patients eligible for HIV testing who receive a test is 27.2% (95% CI 22.4% to 32%). This low level of testing suggests that adherence to the 2008 UK guidelines for HIV testing is poor in recommended populations and settings. Analysis of test offer and acceptance levels suggests that the low overall level of testing is likely to be due to low levels of provider test offer and not patient acceptance. Provider test offer to those eligible was estimated to be only 40.4% (95% CI 24.3% to 56.7%) while patient acceptance of testing was 71.5% (95% CI 56% to 86.9%). This trend of low provider test offer and high-patient test acceptance has previously been seen in other countries in Europe and in the USA,^{14 15} where it has been suggested that it indicates that health providers assess risk differently, are more likely to offer testing to patients they perceive to be at high risk or more likely to accept testing. Aside from this, operational and training barriers such as inadequate training for routine test offer, lack of time or difficulty in ordering an HIV test have also been

cited as reasons contributing to low levels of health provider test offer.^{16–18}

The highest level of testing (83.2%) was reported by Chan *et al*, who assessed the uptake and acceptability during a study of consecutive HIV test offer in medical admissions in Croydon. A previous audit of HIV testing in this hospital had showed a very low coverage of less than 1% prior to the prospective study. This indicates that consecutive test offer as undertaken in the prospective study can yield a much higher level of coverage. Clearly implementing a study of HIV offer is an intervention, and this may explain the significant difference in coverage in screening settings between retrospective audits and prospective studies. However, some retrospective studies also report high levels of testing, such as in Rosenvinge *et al*¹⁹ with 80.7%, indicating that good coverage can be achieved in the absence of a prospective study.

A higher HIV seroprevalence was found in patients tested who presented with a disease indicative of HIV infection at 2.7% (95% CI 1.1% to 4.4%) than found in those tested in settings where routine HIV testing should be undertaken 0.4% (95% CI 0.2% to 0.6%), and the overall pooled seroprevalence from studies was found to be 0.5% (95% CI 0.3% to 0.7%). These seroprevalence estimates exceed the threshold level 0.1% seropositivity of total tests administered deemed as cost-effective by CDC,²⁰ indicating that HIV testing in these settings and populations is cost-effective and is likely to continue to be so with increased test coverage.

Of those presenting with indicator disease conditions (including tuberculosis, glandular fever and other blood borne viruses), an estimated 22.4% (95% CI 13.9% to 30.9%) received an HIV test compared with an estimated 29.5% (95% CI 23.6% to 35.4%) of those attending screening settings where routine HIV testing should be undertaken. Although the odds of being tested for HIV if diagnosed with an indicator disease condition do not appear to significantly contribute to the difference seen in test coverage (0.8, $p=0.67$), this group represents a particularly high-risk population who are easily identified. Testing in this group is a long-standing recommendation of guidelines prior to 2008, so these results are very disappointing. HIV testing in patients with indicator diseases has previously been explored. Read *et al*²¹ found that 37% of patients newly diagnosed with HIV in their secondary care hospital had presented to health-care services with an HIV indicator condition in the preceding 12 months but had not been tested at the time. In a recent prospective study looking at the effectiveness of indicator condition-guided testing for HIV, Sullivan *et al*²² found an HIV prevalence of 1.8% (95% CI 1.42% to 2.34%) across European centres, similar to our estimate of 2.7% (95% CI 1.1% to 4.4%). Furthermore, findings from an analysis from 13 counselling and testing sites in Italy indicated that those presenting late with HIV were probably already infected at the time their initial indicator disease was diagnosed, but that there was a median lapse of 22.6 months between indicator disease diagnosis and HIV diagnosis.²³ The lack of adherence to guidelines in this group therefore is likely to be hindering timely identification of HIV greatly.

There are a number of limitations to this study, primarily the lack of a comparable routine data set with relevant information. Due to this we have been reliant on a relatively small number of reports from local audits and studies that included a wide variety of populations, settings, duration and methods used for measuring HIV testing. However, as guideline recommendations are broad in their description of settings and populations, further restriction in inclusion criteria was not possible. The

studies were of varied quality, and this could not be systematically assessed as many were published as reports or conference abstracts rather than peer-reviewed papers. Data quality was also variable, with some dependent on patient self-report of previous tests to define eligibility. Several studies were interventional in nature, offering consecutive HIV tests in recommended settings and this may have contributed to an overestimate of testing in routine conditions. However, these limitations could only be overcome through the implementation of standards for reporting in the context of some surveillance system such as those that already exist in established testing settings.

There was a great deal of heterogeneity in the data with some I^2 statistic values at 100%, and as a result caution should be taken in interpreting the summary statistics presented for illustration as an average proportion. True study percentages are likely to vary greatly around the estimate points presented. We do not claim to present a true level of overall test coverage level but rather an estimate from the data collected and we have tried to understand some of the variation that was associated with this. Meta-regression did not identify any factor as the majority a contributor to the between-study variance seen, and it is likely that much more of the heterogeneity could be explained by factors that could not be measured in the meta-regression either due to insufficient study numbers or the fact that potential explanatory variables were not reported for all studies.

CONCLUSION

The results of this review and meta-analysis indicate adherence to 2008 national guidelines for HIV testing in the UK is poor and that low levels of provider test offer appear to be a major contributor to this, particularly in patients presenting with an indicator disease. Failure to adhere to testing guidelines is likely to be contributing to late diagnosis with implications for poorer clinical outcomes and continued onwards transmission of HIV. Improved surveillance of HIV testing outside of specialist settings may be useful in increasing adherence testing guidelines.

Key messages

- Findings indicate HIV test coverage in the UK is poor and low levels of provider test offer appear to be the main contributor to this.
- Seroprevalence estimates show that HIV testing is cost-effective and increasing HIV testing in line with national guidelines would also be cost-effective.
- Further exploration of effective methods for the routine offer of HIV testing in recommended settings.
- Better methods for the clear dissemination of routine HIV testing messages to non-specialist clinicians are required.

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Contributors HW and RE were responsible for study concept and design. RE and SMG conducted the searches and extracted the data. Statistical analysis was carried out by SMG, GGG and RE. All authors contributed to the interpretation of the data.

RE produced the first draft of the manuscript, and all authors contributed to revisions and approved the final version. HW is the guarantor.

Ethics approval Ethical approval was not obtained for this study as it was based on secondary analysis of published data only.

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Appendix a: Classification of settings and populations for routine HIV testing (adapted from national testing guidelines, BHIVA 2008) [6]

<p>Persons diagnosed with a disease indicative of HIV infection</p>	<p>Tuberculosis Pneumocystis Cerebral toxoplasmosis Primary Cerebral Lymphoma Cryptococcal meningitis Progressive multifocal leucoencephalopathy Kaposi's sarcoma Persistent cryptosporidiosis Non-Hodgkin's Lymphoma Cervical Cancer Cytomeglovirus retinitis Bacterial pneumonia Aspergillosis Aseptic meningitis/encephalitis Cerebral abscess Space occupying lesion of unknown cause Guillain-Barré syndrome Transverse myelitis Peripheral neuropathy Dementia Leucoencephalopathy Severe or recalcitrant seborrhoeic dermatitis Severe or recalcitrant psoriasis Multidermatomal or recurrent herpes zoster Oral candidiasis Oral hairy leukoplakia Chronic diarrhoea of unknown cause Weight loss of unknown cause Salmonella, shigella or campylobacter Hepatitis B infection Hepatitis C infection Anal cancer or anal intraepithelial dysplasia Lung cancer Seminoma Head and neck cancer Hodgkin's lymphoma Castleman's disease Vaginal intraepithelial neoplasia Cervical intraepithelial neoplasia (Grade 2 or above) Any unexplained blood dyscrasia including: Thrombocytopenia Neutropenia Lymphopenia Infective retinal disease including herpesviruses and toxoplasma Any unexplained retinopathy Lymphadenopathy of unknown cause Chronic parotitis Lymphoepithelial parotid cysts Mononucleosis-like syndrome (primary HIV infection) Pyrexia of unknown origin Any lymphadenopathy of unknown cause Any sexually transmitted infection</p>
<p>Persons attending a service where routine HIV screening is undertaken</p>	<p>Termination of pregnancy services Drug dependency programmes All patients presenting for healthcare where HIV prevalence</p>

(excluding GUM and antenatal services)	All men and women registering in general practice where diagnosed HIV prevalence in the local population exceeds 2 in 1000 population All general medical admissions where diagnosed HIV prevalence in the local population exceeds 2 in 1000 population
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Appendix b: Characteristics of Studies Included: Methods, Measures and Testing Levels

Author	Risk group	Setting (diagnosed HIV prevalence per 1,000 population 15-59 year olds) *	Methods	Number eligible to test	Number offered test	Number tested	Number testing positive
Persons diagnosed with a disease indicative of HIV infection							
Gupta, N.D. & Lechelt, M. (2011) [1]	Inpatients with indicator diseases (tuberculosis, hepatitis B, hepatitis C, cervical intraepithelial neoplasia (grade I/II), lymphoma, anal cancer, seminoma, aspergillosis or Castleman's disease)	South-west Essex (1.28)	Electronic record audit of attendees attending one secondary care hospital	557	33	33	Not reported
Thomas William, S., <i>et al.</i> (2011) [2]	Patients with indicator disease (tuberculosis)	Birmingham and Solihull (Birmingham East & North; 1.5, Heart of Birmingham; 3.29, South Birmingham; 1.66; Solihull; 0.58)	Retrospective audit	194	Not reported	91	Not reported
Hsu, D., <i>et al.</i> (2012) [3]	Primary care patients presenting with glandular fever-like illness	South London (Lambeth 13.28, Southwark; 10.29)	Retrospective audit of patients attending 72 primary care clinics	1045	Not reported	118	3
Page, I., <i>et al.</i> (2011) [4]	Patients with indicator disease (tuberculosis, hepatitis B, hepatitis C, lymphoma)	Blackpool (3.41)	Retrospective audit of patients attending one secondary care hospital	156	Not reported	32	Not reported
Thomson-Glover, R., <i>et al.</i> (2011) [5]	Patients with indicator disease (hepatitis B, hepatitis C, candida)	Warrington (0.6)	Case-note audit of patients attending two secondary care hospitals	249	Not reported	15	0

	stomatitis)						
Thorburn, F. (2012) [6]	Patients with indicator disease (diagnosed with tuberculosis)	Glasgow (1.7)	Retrospective case-note review of TB patients attending one tertiary care clinic	338	Not reported	221	9
Vas, A., <i>et al.</i> (2012) [7]	Patients with indicator disease (tuberculosis, hepatitis B, hepatitis C)	Manchester (5.22)	Retrospective case-note review of patients attending one secondary care hospital	91	13	9*	Not reported
Byrne, L., <i>et al.</i> (2011) [8]	Patients admitted to acute medical unit with community-acquired pneumonia	London (Newham; 8.12, Tower Hamlets; 5.94)	Retrospective case-note review of patients attending one acute medical admissions unit	43	Not reported	17	2
Manavi, K., Gautam, N. (2012) [9]	Patients diagnosed with clinical indicator conditions as specified in UK HIV testing guidelines	Birmingham (3.29)	Retrospective case note review of patients attending one secondary care hospital	967	Not reported	97	1
Dodd, M. <i>et al</i> (2013) [10]	Patients with an HIV indicator illness in the presenting complaint or past medical history.	Sheffield (1.4)	Retrospective case note review of patients in one General Intensive Care Unit	307	Not reported	45	3
Persons attending recommended testing settings in high prevalence areas							
Burns, F., <i>et al.</i> (2012) [11]	Acute medical admissions	London (5.24)	Prospective, consecutive HIV test offer to patients attending one acute medical admissions unit	606	282	135	3
Chan, S.Y., <i>et al.</i> (2011) [12]	Acute medical admissions	Croydon (4.45)	Prospective offer of HIV test to patients attending one acute medical admissions unit	101	101	84	0
Rayment, M., <i>et al.</i> (2012) [13]	Acute Care unit and Dermatology outpatient clinic	London (City and Hackney (8.25)	Prospective study of patients attending one acute care unit	1223	548	384	4
Perry, N., <i>et al.</i> (2011) [14]	Acute medical admissions	Brighton & Hove PCT (7.57)	Prospective HIV test offer to patients attending one acute medical admissions unit	3913	1553	1413	2
Bryce, G., (2009) [15]	Patients newly registering with GP	Brighton & Hove PCT (7.57)	Prospective HIV test offer to patients attending nine primary	2478	Not reported	1473	2

			care clinics				
Ashby, J., <i>et al.</i> (2012) [16]	Polyclinic attendees in high prevalence area	West London (Kensington & Chelsea; 8.3, Hammersmith & Fulham; 8.5, Westminster; 7.01)	Prospective study of patients attending one polyclinic	302	93	71	0
Ellis, S., <i>et al.</i> (2011) [17]	Acute medical admissions	Newcastle Upon Tyne (1.61)	Prospective audit of patients attending one acute medical admissions unit	3645	478	396	2
Rudran, B., <i>et al.</i> (2011) [18]	Acute medical admissions	Bournemouth and Poole (2.32)	Retrospective case-note review of patients attending one acute medical admissions unit	198	3	3	Not reported
Leber, W., <i>et al.</i> (2012) [19]	Patients newly registering with GP	Hackney (8.25)	Cluster randomised control trial of patients attending 40 primary care units	28274	6607	3213	7
Bassett, D., <i>et al.</i> (2012) [20]	Acute medical admissions	Manchester (5.22)	Prospective case-note review of patients attending one acute medical admissions	429	134	117	Not reported
Rosenvinge, M., <i>et al.</i> (2010) [21]	Women attending termination of pregnancy services	Wandsworth (4.91)	Retrospective review of HIV testing of patients who attended two termination of pregnancy clinic	870	844	702	1
Garrard, N., <i>et al.</i> (2010) [22]	Women attending termination of pregnancy service	Southwark (10.39) and Lambeth (13.28)	Prospective, consecutive test offer to patients attending one termination of pregnancy clinic	2,831	Not reported	972	5
Barbour, A., <i>et al.</i> (2012) [23]	Patients admitted to acute medical admissions	Croydon (4.45)	Prospective intervention at one acute medical unit	3709	Not reported**	1390	7
Rycroft, J., <i>et al.</i> (2012) [24]	Acute medical admissions	Greenwich (5.58)	Retrospective audit of patients who attended one acute medical admissions	970	Not reported	43	3
Page, I., <i>et al.</i> (2011) [4]	Acute medical admissions	Blackpool (3.41)	Retrospective audit of patients who attended one secondary care hospital	13,999	Not reported	72	Not reported

French, S., <i>et al.</i> (2012) [25]	Patients newly registering with GP	Southwark (10.39), Lewisham (7.03), Lambeth (13.28)	Prospective study of patients attending 13 primary care clinics	16,241	6405	3229	12
French, S., <i>et al.</i> (2012) [25]	Patients newly registering with GP	Southwark (10.39), Lewisham (7.03), Lambeth (13.28)	Prospective study of patients attending 5 primary care clinics	6275	4925	905	11
Tillet, S., <i>et al.</i> (2012) [26]	Acute medical admissions	Tower Hamlets (5.94)	Prospective study of patients attending one secondary care hospital	1596	Not reported	241	2
Griffin, A., <i>et al.</i> (2011) [27]	Patient newly registering with GP	Manchester (5.22)	Prospective study of patients attending one primary care clinic	457	Not reported	303	2
Palfreeman, A., <i>et al.</i> (2013) [28]	Patients attending admitted to AMU	Leicester (3.22)	Prospective study of patients admitted to AMU	17226	Not reported	2542	29

* reported for patients diagnosed with TB only. ** 77 test refusals were reported. Additional data regarding testing strategy (opt-in vs. opt-out), service model (standard care vs. specific staff training vs. GUM/Health advisor-led testing), and type of HIV test (POCT vs. 4th generation serology).

Appendix c: Supplementary data tables for studies identified by group

Study	Title	Article	Primary testing outcome	Exclusions	Time period (duration)	Population	Number of centres	Type of centre	Measure/reporting method
Persons diagnosed with a disease indicative of possible HIV infection									
Gupta, N.D. & Lechelt, M. (2011)	Assessment of the implementation and knowledge of the UK national guidelines for HIV testing (2008) in key conditions at	Yes - Audit Report	Electronic departmental record or HIV testing and Electronic pathology records	Non-verifiable data	August 2009 – June 2012 (11 months)	Inpatients at Basildon & Thurrock Hospital	1	Secondary care hospital	Electronic record of HIV test

	a UK district general hospital								
Thom as William, S., et al. (2011)	Changes in HIV testing rates among patients with tuberculosis in a large multi-ethnic city in the UK	Yes - Audit Report	Laboratory database record of HIV test	<18 years, private patients, chemoprophylaxis patients, non-tuberculosis mycobacteria, diagnosis outside catchment area	September 2008 – March 2009 (6 months)	Patients registered on the Birmingham Tuberculosis aftercare register	>1	Various	Laboratory record of HIV test
Hsu, D., et al. (2012)	Diagnosing HIV infection in patients presenting with glandular fever-like illness in primary care: are we missing primary HIV infection?	Yes - Short Communication	Record of HIV test	Not reported	April 2009 - June 2010 (14 months)	Primary care patients in Lambeth and Southwark	72	Primary care clinics	Laboratory record of HIV test request
Page, I., et al. (2011)	The impact of new national HIV testing guidelines at a district general hospital in an area of high HIV seroprevalence	Yes - Paper	Laboratory record of HIV test	HIV test requests from GUM clinics	October 2008 – September 2009	Inpatients in Blackpool	1	Secondary care hospital	Laboratory record of HIV test

Thomson-Glover, R., et al. (2011)	Diagnosing HIV in non-GUM secondary care settings	No - Abstract	Record of HIV test	not reported	November 2009 - April 2010 (6 months)	inpatients Warrington & Halton hospitals	2	Secondary care hospitals	Record of HIV test
Thorburn, F. (2012)	The impact of a multi-disciplinary meeting on the rates of HIV in testing in TB patients	No - Abstract	Record of HIV test offer	not reported	2010 - 2011 (duration not reported)	Tuberculosis patients attending virology centre in Glasgow	1	Tertiary care clinic	Laboratory record of HIV test administration
Vas, A., et al. (2012)	HIV testing and in TB and Hepatitis services in a district general hospital	No - Abstract	Record of HIV test	not reported	2009 (duration not reported)	Indicator disease patients in a Manchester hospital	1	Secondary care hospital	Record of HIV test administration
Byrne, L., et al. (2011)	HIV specialists must lead the way to make HIV testing truly routine	No-Abstract	Record of HIV test	, <18, non-medical specialty, underlying chronic lung disease, hospital-acquired pneumonia	February - April 2010 (3 months)	Patients admitted with community-acquired pneumonia	1	Acute medical admissions unit	Case-note record of HIV test administration
Persons attending services where routine HIV screening should be undertaken									
Chan, S.Y., et al. (2011)	Acceptance of HIV testing in medical inpatients: A local acceptability study	Miscellaneous	Consent to have an HIV test	<15 and >59 years, total time admitted <24 hours, assessed as unable to consent, known to be HIV-positive	September 2009 (2 weeks)	Acute medical admissions in Croydon	1	Acute medical admissions unit	Offer and acceptance of HIV test

Rayment, M., et al. (2012)	HIV Testing in Non-Traditional Settings - the hints study: A multi-centre observational study of feasibility and acceptability.	Article	Offer of HIV test to eligible individual	<16, >65 years, known HIV positive, not accessing healthcare for the first time in testing period, not able to consent	January – September 2010 (12 weeks each site)	Patients attending primary and secondary healthcare services in 4 London centres	4	Acute care units, Dermatology OPD	Administration of HIV oral fluid or 4 th generation HIV serology
Perry, N. et al. (2011)	HIV testing in acute general medical admissions must be universally offered to reduce undiagnosed HIV	Abstract	Record of HIV test	<16 and >79 years, known HIV positive	August 2009 – January 2010 (5 months)	Acute medical admissions in Brighton	1	Acute medical admissions unit	HIV test result
Bryce, G. (2009)	A study to assess the acceptability, feasibility and cost-effectiveness of universal HIV testing with newly registering patients (aged 16-59) in primary care	Abstract	Acceptance of HIV test offer	<16 and >59 years	May - November 2010 (4 months)	Patients attending primary care services in Brighton	9	Primary care clinics	HIV POCT test
Ashby, J., et al. (2012)	HIV testing uptake and acceptability in an inner city polyclinic	Article	Acceptance of HIV test offer	<16, >65 years, unable to consent	2011 dates not specified (random 4-hour duration over a 4 week period)	Polyclinic attendees in west London	1	Polyclinic	Rapid point-of-care HIV test

Ellis, S., et al. (2011)	Offering HIV testing in an acute medical admissions unit in Newcastle upon Tyne	Clinical Medicine research	Record of HIV test	<18 years, no capacity for consent	September - October 2009 (11 weeks) and January - March 2010 (6 weeks)	Acute medical admissions in Newcastle	1	Acute medical admissions unit	HIV test offer and administration
Rudran, B., et al. (2011)	HIV testing in acute medical admissions	Abstract	Record of HIV test	not reported	2010 - exact date not specified (1 week)	Acute medical admissions in Bournemouth	1	Acute medical admissions unit	Hospital or laboratory database record of HIV test
Leber, W., et al. (2012)	Can point-of-care HIV testing in primary care increase identification of HIV? The RHIVA 2 Cluster randomised control trial - update	Abstract	Offer of rapid point-of-care HIV test	<16 years,	May 2010 end date not specified	Patients attending primary care services in London	40	Primary care units	Administration of rapid point-of-care HIV test
Bassett, D., et al. (2012)	Practical challenges implementing national HIV testing guidelines in general medical admissions	Abstract	Record of HIV test	not reported	July 2011 (two weeks)	Acute medical admissions in central Manchester	1	Acute medical admissions	Record of HIV test administration

Rosenvinge, M., et al. (2010)	A successful uptake of HIV testing in south London termination of pregnancy services	Abstract	Record of consent for an HIV test	Known HIV positive, recent (< 6 months) HIV negative test, repeat attendance	April - December 2009 (9 months)	Women attending termination of pregnancy services in south London	2	Termination of pregnancy clinics	Paper and electronic record of HIV test administration
Garrard, N., et al. (2010)	Opt-out HIV testing pilot in termination of pregnancy services - 11-month service evaluation	Abstract	HIV test recommendation	not reported	November 2008 - September 2009 (11 months)	Women attending termination of Pregnancy services north London	1	Termination of pregnancy clinic	Documentation of HIV test result
Barbour, A., et al. (2011)	Opt-out HIV testing policy implemented as routine standard of care for acute medical admissions in a high prevalence area	Abstract	Record of HIV test	<16 and >79 years,	July 2011 – December 2011 (6 months)	Patients attending acute medical admissions in Croydon	1	Acute medical admissions	HIV testing
Rycroft, J., et al. (2012)	HIV testing in the acute medical unit - setting the scene for universal opt-out testing	Abstract	Laboratory record of HIV test	Not reported	June & November 2011 (audited 2 weeks for each admissions cycle)	Patients attending acute medical admissions in south east London	1	Acute medical admissions	Record of HIV test in laboratory database
Page, I., et al. (2011)	The impact of new national HIV testing guidelines at a district general hospital in an area of high	Paper	Laboratory record of HIV test	HIV test requests from GUM clinics	October 2008 – September 2009	Inpatients in Blackpool	1	Secondary care hospital	Laboratory record of HIV test

	HIV seroprevalence								
Palfreman, A., et al. (2013)	HIV testing for acute medical admissions: evaluation of a pilot study in Leicester, England	Paper	Laboratory record of HIV test	Not reported	September 2008 – August 2011 (36 months)	Patients admitted to AMU in Leicester	1	Secondary care hospital	Laboratory record of HIV test

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