HIV testing in General Practice

Dr Sylvia Bates

HIV – the basics

Acquisition

- Sex
- Vertical transmission
- Injecting drug use
- Blood transfusions with infected blood
- Percutaneous injury needle stick

HIV infection

Acute infection - seroconversion

Asymptomatic

HIV related illnesses

AIDS defining illness

Death

Sero- conversion	CD4 > 500	CD4 500 - 200	CD4 <200
 oFever oMyalgia oArthralgia oAdenopathy oMalaise oRash oMeningo-encephalitis 	oGuillain-Barre syndrome oBell's palsy oPolymyositis oChronic demyelinating neuropathy oldiopathic thrombo- cytopaenia oTinea	oSeborrhoeic dermatitis oGingivitis oWarts oMolluscum oTB oHerpes Zoster/ Simplex oOral candida oKS oCIN oPrimary CNS lymphoma	 oCryptosporidiosis oPCP oToxoplasmosis oCryptococcal meningitis oCMV oMAC

Primary HIV infection

Classical triad of:

- Pharyngitis
- Rash
- Fever

Clinical Manifestations













HIV in the UK

- There are currently an estimated 96,000 people living with HIV in the UK.
- 12% of those infected remain undiagnosed HPA 2016
- 42% of new HIV diagnoses are diagnosed late (CD4 < 350).
- 35% of HIV-related deaths attributable to late diagnosis BHIVA audit 2006
- Effective treatments greatly reduce morbidity
- Better response to treatment if started early
- Fully suppressed viral loads revice PREVENT onward transmission

U=U

Undetectable = un-transmittable

Figure 18: The continuum of HIV care, England excluding London, 2016

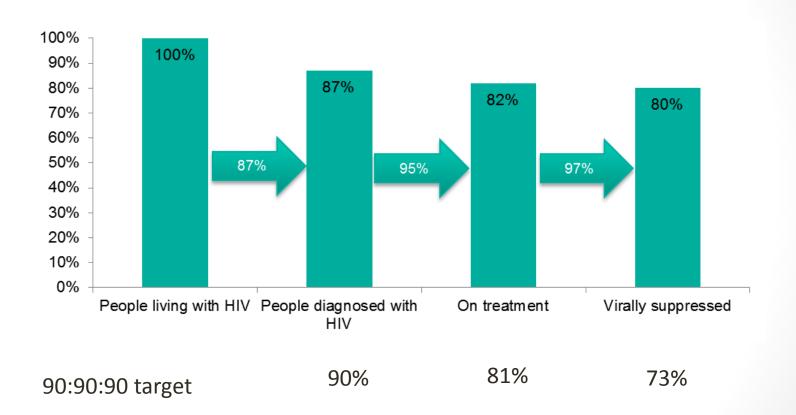
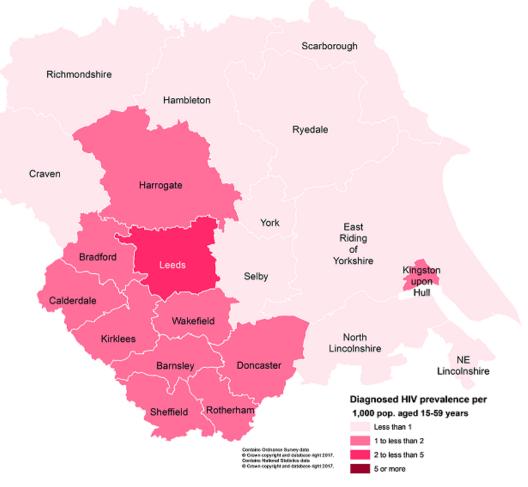


Figure 16: Diagnosed HIV prevalence per 1,000 residents aged 15-59 years by local authority, Yorkshire and the Humber, 2016



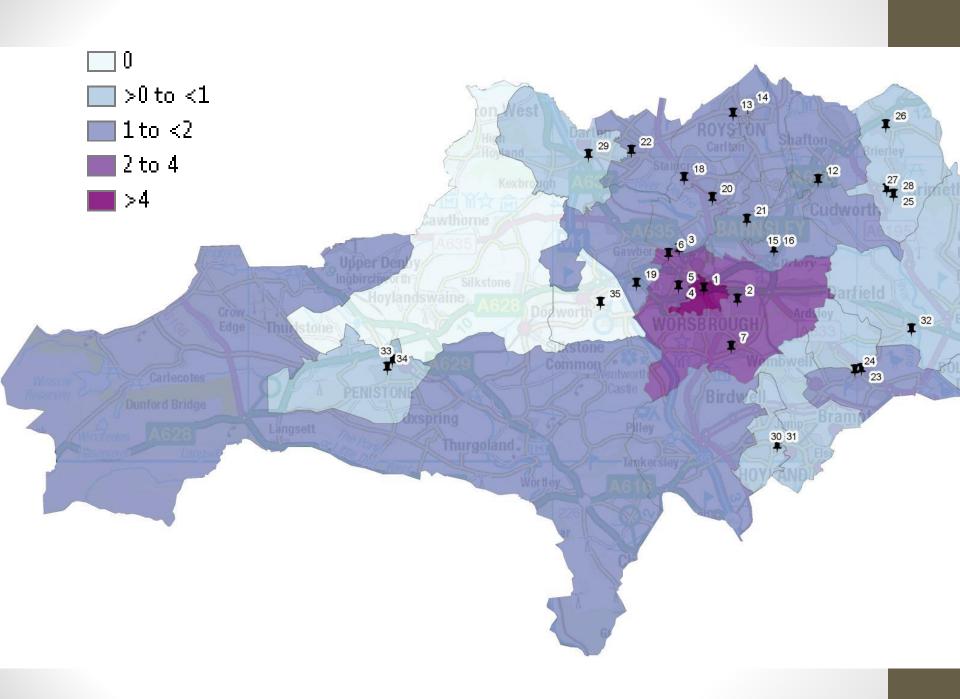
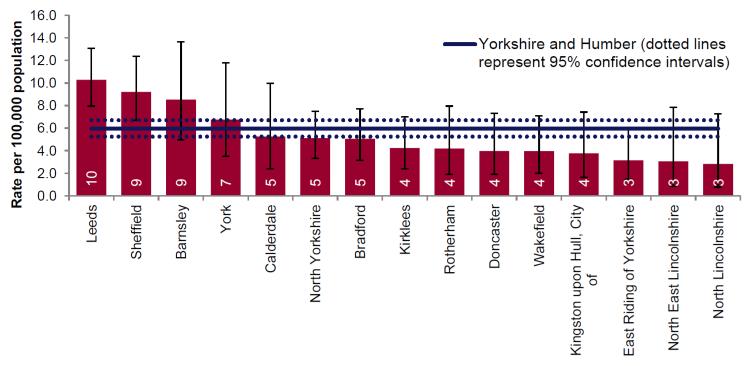


Figure 2: New HIV diagnoses per 100,000 population aged 15 years or older by upper tier local authority of residence, Yorkshire and the Humber residents, 2016

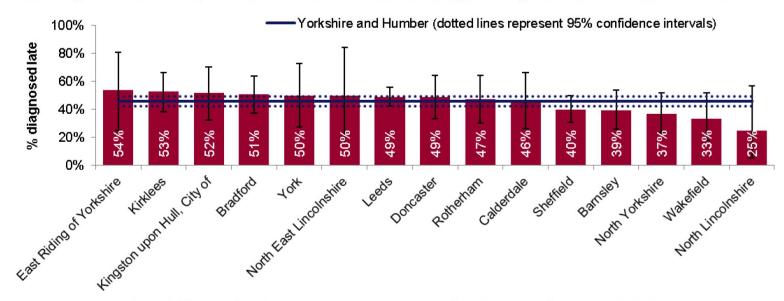


- +

Source: Public Health England, HIV & AIDS New Diagnoses and Deaths (HANDD).

0

The number of new diagnoses will depend on accessibility of testing as well as infection and transmission.



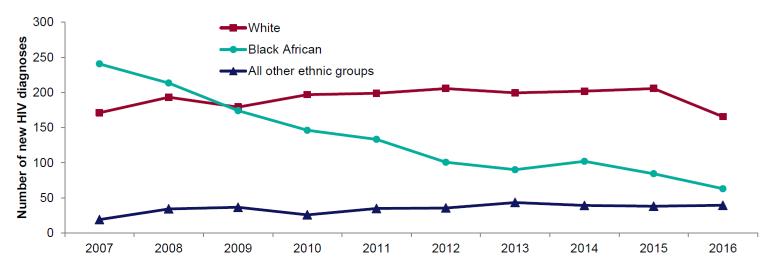
Source: Public Health England, HIV and AIDS New Diagnosis Database/System, HIV & AIDS Reporting System (HARS).

The underlying population will impact on the proportion diagnosed late, e.g. MSM are less likely to be diagnosed late.

0

^{*} Only includes new diagnoses for which CD4 count was reported within 91 days of diagnosis; late diagnosis defined as CD4 count <350 cells/mm³.

Figure 6: Number of new HIV diagnoses by ethnic group (adjusted for missing ethnic group information), Yorkshire and the Humber residents, 2007-2016 (please see footnote on interpreting trends)*



Source: Public Health England, HIV & AIDS New Diagnoses and Deaths (HANDD).

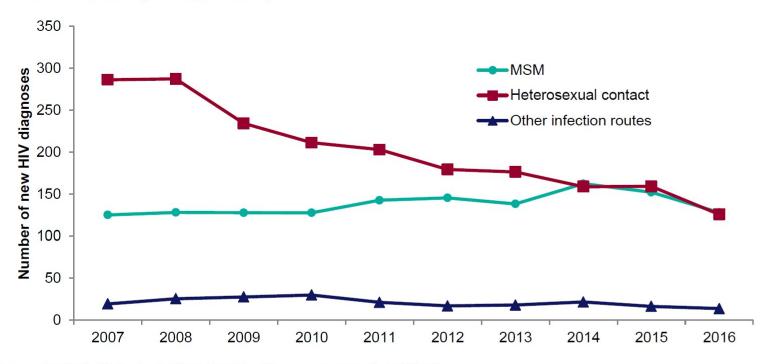
0

[]

The number of new diagnoses will depend on accessibility of testing as well as infection and transmission.

*Numbers may rise as further reports are received and more information is obtained on area of residence of those diagnosed. This is more likely to affect more recent years, particularly **2016**. Please see important note on data earlier in this report. This will impact on interpretation of trends in more recent years.

Figure 4: New HIV diagnoses by probable route of infection (adjusted for missing route of infection information), Yorkshire and the Humber residents, 2007-2016 (please see footnote on interpreting trends)*



Source: Public Health England, HIV & AIDS New Diagnoses and Deaths (HANDD).

12 / 25

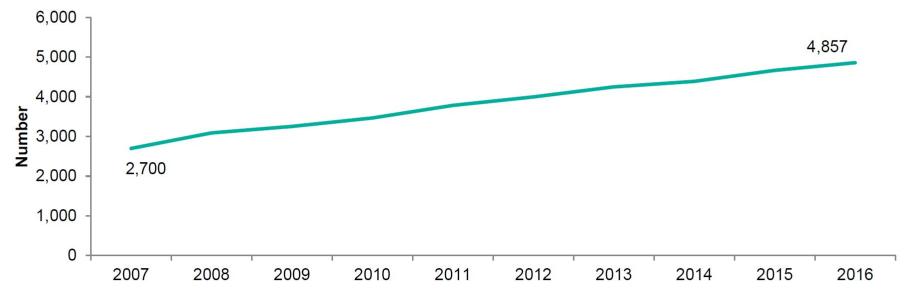
0

The number of new diagnoses will depend on accessibility of testing as well as infection and transmission.

*Numbers may rise as further reports are received and more information is obtained on area of residence of those diagnosed. This is more likely to affect more recent years, particularly **2016**. Please see important note on data earlier in this report. This will impact on interpretation of

Spotlight on HIV in Yorkshire and the Humber

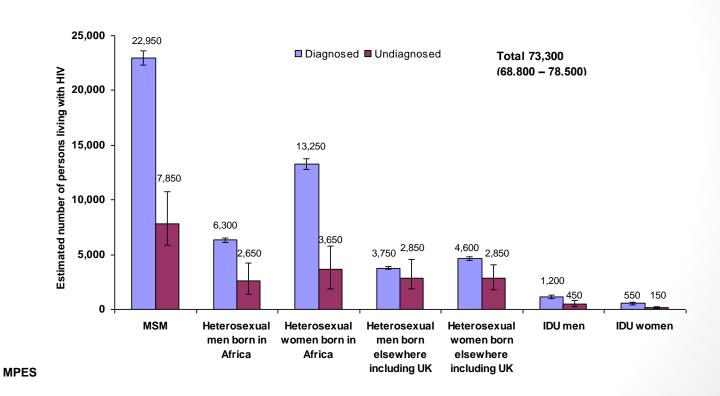
Figure 11: Number of residents living with diagnosed HIV and accessing care, Yorkshire and the Humber, 2007-2016



Source: Public Health England, HIV & AIDS Reporting System (HARS).



Estimated number of adults (15 to 59 years) living with HIV (both diagnosed and undiagnosed) in the UK: 2007



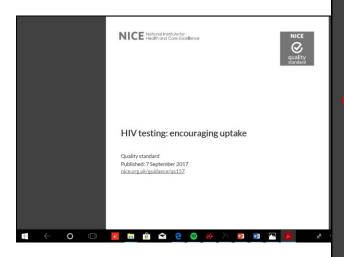
Why test in General practice?

- 1. Recommendations based on prevalence
 - 11 practices in Barnsley have an HIV prevalence of > 2/1000
 - A further 10 Practices have a rate between 1-2 / 1000

NICE recommendations
BASHH recommendations

2. Testing for HIV saves lives - Look back review of late diagnosis

NICE guidance



HIV testing should be seen as routine practice, says NICE in new draft quality standard

Millions of people could be offered HIV tests during routine appointments as outlined by NICE.



Quality statements

Statement 1 Young people and adults are offered an HIV test when admitted to hospital or attending an emergency department in areas of extremely high HIV prevalence, or when having a blood test when admitted to hospital or attending an emergency department in areas of high HIV prevalence.

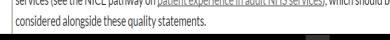
Statement 2 Young people and adults in areas of high or extremely high HIV prevalence are offered an HIV test by their GP practice when registering or when having a blood test if they have not had an HIV test in the past 12 months.

Statement 3 Young people and adults newly diagnosed with an HIV indicator condition are offered an HIV test.

Statement 4 Young people and adults in at-risk groups who test negative for HIV are advised that the test should be repeated at least annually.

Statement 5 People who may have been exposed to HIV by a person newly diagnosed with HIV are offered an HIV test.

NICE has developed guidance and a quality standard on patient experience in adult NHS services (see the NICE pathway on patient experience in adult NHS services), which should be



- Healthcare professionals should emphasise that having an HIV test is a routine procedure but if the test is declined, they should provide information on how to access other local HIV testing services.
- Local authorities with a diagnosed HIV prevalence of between
 - 2 and 5 per 1,000 people aged 15 to 59 years have a high HIV prevalence
 - 5 or more per 1,000 people aged 15 to 59 years have an extremely high HIV prevalence

Strongly recommend test

Neoplasms:

- Cervical cancer
- Non-Hodgkin lymphoma
- Kaposi's sarcoma

Bacterial infections

- · Mycobacterium Tuberculosis, pulmonary or extrapulmunary
- Mycobacterium avium complex (MAC) or Mycobacterium kansasii, disseminated or extrapulmonary
- Mycobacterium, other species or unidentified species, disseminated or extrapulmunary
- · Pneumonia, recurrent (2 or more episodes in 12 months)
- · Salmonella septicaemia, recurrent

Viral infections

- · Cytomegalovirus retinitis
- · Cytomegalovirus, other (except liver, spleen, glands)
- · Herpes simplex, ulcer(s) > I month/bronchitis/pneumonitis
- · Progressive multifocal leucoencephalopathy

Parasitic infections

- Cerebral toxoplasmosis
- · Cryptosporidiosis diarrhoea, > 1 month
- Isosporiasis, > 1 month
- · Atypical disseminated leismaniasis
- Reactivation of American trypanosomiasis (meningoencephalitis or myocarditis)

Fungal infections

- · Pneumocystis carinii pneumonia
- · Candidiasis, oesophageal
- Candidiasis, bronchial/ tracheal/ lungs
- · Cryptococcosis, extra-pulmonary
- · Histoplasmosis, disseminated/ extra pulmonary
- · Coccidiodomycosis, disseminated/ extra pulmonary
- · Penicilliosis, disseminated

3. Conditions where not identifying the presence of HIV infection may have significant adverse implications for the individual's clinical management

Offer testin

- · Conditions requiring aggressive immuno-suppressive therapy:
- Cancer
- Transplantation
- · Auto-immune disease treated with immunosuppressive therapy
- Primary space occupying lesion of the brain.
- · Idiopatic/Thrombotic thrombocytopenic purpura

2a. Conditions associated with an undiagnosed HIV prevalence of ≥0.1

- · Sexually transmitted infections
- · Malignant lymphoma
- · Anal cancer/dysplasia
- · Cervical dysplasia
- · Herpes zoster
- · Hepatitis B or C (acute or chronic)
- · Unexplained lymphadenopathy
- · Mononucleosis-like illness
- · Community-acquired pneumonia
- Unexplained leukocytopenia/thrombocytopenia lasting >4 weeks
- · Seborrheic dermatitis/exanthema
- · Invasive pneumococcal disease
- Unexplained fever
- Candidaemia

Strongly recommend testing

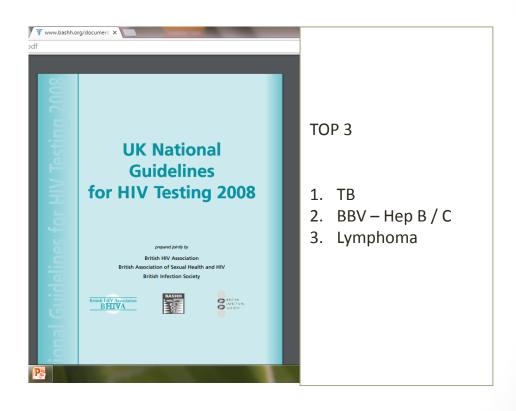
Offer testing:

- Visceral leishmaniasis
- · Pregnancy (implications for the unborn child)

2b. Other conditions considered likely to have an undiagnosed HIV prevalence of >0.1%

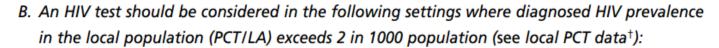
- · Primary lung cancer
- · Lymphocytic meningitis
- · Oral hairy leukoplakia
- · Severe or atypical psoriasis
- · Guillain-Barré syndrome
- Mononeuritis
- Subcortical dementia
- · Multiplesclerosis-like disease
- Peripheral neuropathy
- · Unexplained weightloss
- · Unexplained oral candidiasis
- · Unexplained chronic diarrhoea
- · Unexplained chronic renal impairment
- · Hepatitis A
- Candidiasis

When to test?



cuments/1838.pdf





- all men and women registering in general practice
- all general medical admissions.

The introduction of universal HIV testing in these settings should be thoroughly evaluated for acceptability and feasibility and the resultant data made available to better inform the ongoing implementation of these guidelines.

C. HIV testing should be also routinely offered and recommended to the following patients:

- all patients presenting for healthcare where HIV, including primary HIV infection, enters the differential diagnosis (see table of indicator diseases and section on primary HIV infection)
- all patients diagnosed with a sexually transmitted infection
- 3. all sexual partners of men and women known to be HIV positive
- all men who have disclosed sexual contact with other men
- all female sexual contacts of men who have sex with men
- all patients reporting a history of injecting drug use
- all men and women known to be from a country of high HIV prevalence (>1%*)
- all men and women who report sexual contact abroad or in the UK with individuals from countries of high HIV prevalence.*
 - * for an up to date list see

Xc

Deskton

Testing in General Practice

- In 2016, HIV test positivity rates in general practice in high (0.5%) and extremely high (0.4%) diagnosed HIV prevalence areas, and those tested in A&E and other secondary care settings (0.6%), now exceed those seen in SHS.
- HIV Testing in general practice have increased by 17% between 2014 and 2016.
- In extremely high prevalence areas, 101 per 10,000 of the general practice population were tested for HIV in 2016 (positivity of 0.4%).
- In high prevalence areas, 44 per 10,000 of the general practice population were tested, 0.5% of whom were positive.

Local epidemiology – new cases 2015/2016;

2015

- 21 male, 4 female;
- 17 white British, 14 MSM (82%);
- 3 Black African
- 5 Eastern European (20%)
- Africans in UK since 2015;
- E Europeans since 2003, 2003, 2004, 2007 and 2013;
- GP diagnosed 2 (x 1 MSM), 4 by medical admissions (1 died)
- Mean CD4 White British = 438
- Mean CD4 non-white Br = 305

2016

- 15 male, 4 female;
- 13 white British, 6 MSM (46%);
- 3 Black African
- 3 Eastern European (16%)
- Africans in UK since 1999, 2003,2007; E Europeans 2016;
- GP diagnosed 1 (African), 4 by medical admissions
- Mean CD4 White British = 458
- Mean CD4 non-white Br = 263

- 35 year old male.
- Latvian.
- Previously under drug addiction services in Derby since 2013 methadone programme.
- Migrant.
- Attended GP practice on 6 occasions April to October 2016 with bad skin.
- Diagnosed with psoriasis;
- Clippings showed fungal nails.
- Referred Dermatology confirmed psoriasis and suggested treatment

- October 2016: drug and addiction services performed a POCT:
 - HIV reactive; referred to BISH.
- BISH: "diagnosed with HIV 4 years previously; in UK 8 months; past history of IVDU and Hepatitis B";
- O/E; florid oral candida; seborrhoeic dermatitis; psoriasis; fungal nails;
- HIV confirmed; also HCV genotype 1b, HCV VL 439,672c/ml; ALT 78; HepBcore Ab+ve;
- CD4 = 6 cells/ml; VL 4.04×10^6 /ml;
- Commenced on prophylaxis with septrin, and ARVs
- March 17: CD4 78, VL 54 c/ml; skin & mouth much better;

with clues

- 35 year old male.
- Latvian.
- Previously under drug addiction services in Derby since 2013 methadone programme.
- Migrant.
- Attended GP practice on 6 occasions April to October 2016 with bad skin.
- Diagnosed with psoriasis;
- Clippings showed fungal nails.
- Referred Dermatology confirmed psoriasis and suggested treatment

- October 2016: drug and addiction services performed a POCT:
 - HIV reactive; referred to BISH.
- BISH: "diagnosed with HIV 4 years previously; in UK 8 months; past history of IVDU and Hepatitis B";
- O/E; florid oral candida; seborrheoic dermatitis; psoriasis; fungal nails;
- HIV confirmed; also HCV genotype 1b, HCV VL 439,672c/ml; ALT 78; antiHBcore +ve;
- CD4 = 6 cells/ml; VL 4.04×10^6 /ml;
- Commenced on prophylaxis with septrin, and ARVs
- March 17: CD4 78, VL 54 c/ml; skin & mouth much better;

- 47 year old female.
- Born in Zimbabwe. In Uk since 2007. previously in the Midlands until ~ 2013.
- On ramipril for hypertension.
- Admitted to BHNFT in July 2017 with appendicitis; asked for an HIV test. Believed also had one with GP in previous year.
- HIV detected, CD4 462 cells/mm³, VL 32,000c/ml³, TPPA +ve;
- VL not detected, CD4 588;
- ICE records:
 - bloods 2013 for hypertension;
 - in 2014 "patient requested blood tests";

- with clues

- 47 year old female.
- Born in Zimbabwe. In Uk since 2007. previously in the Midlands until ~ 2013.
- On ramipril for hypertension.
- Admitted to BHNFT in July 2017 with appendicitis; asked for an HIV test. Believed also had one with GP in previous year.
- HIV detected, CD4 462 cells/mm³, VL 32,000c/ml³, TPPA +ve;
- VL not detected, CD4 588;
- ICE records:
 - bloods 2013 for hypertension;
 - in 2014 "patient requested blood tests";

- 37 year old male.
- Admitted to MAU with cough, SOB, fever; CXR showed small area of consolidation right base.
- Married, same partner 10 years, 2 children;
- Weightloss and malaise for 5 months.
- F2 performed an HIV test on admission
- CD4 count 8 cells/mm³
- ICE reports:
 - Nil.

- with clues

- 37 year old male.
- Admitted to MAU with cough, SOB, fever; CXR showed small area of consolidation right base.
- Married, same partner 10 years, 2 children;
- Weightloss and malaise for 5 months.
- F2 performed an HIV test on admission;
- CD4 count 8 cells/mm³; VL 2.9 x 10⁶/L
- ICE reports:
 - Nil.
- CD4 count 180 cells/ml, VL not detected;

Commonly reported clinical signs

- Skin
 - Psoriasis;
- Recurrent fungal infections
- Herpes zoster
- Recurrent oral ulceration
- Weight loss;
- Persistent diarrhoea;
- Seroconversion symptoms;
 - Rash, fever, glands, sore throat, malaise, abdo pain;

Laboratory clues

- Abnormal FBC
 - Low (or high) platelets;
 - Low (or high) neutrophils;
 - High (or low) lymphocytes;
- Hepatitis C or B
- High total protein (esp globulins)

Local epidemiological clues

- Born in Eastern Europe
- Born in Africa;
- IVDU;
- Hepatitis C or B;
- MSM;
- Clinical or laboratory features of HIV, regardless of age/race/gender;

HAART

- Several widely available single tablet regimens
- 'Cleaner' drugs fewer side effects
- Newer drug groups with fewer drug-drug interactions
- Compliance is the key to successful treatment



ART is for life – please always contact the HIV unit before discontinuing a patients antiretroviral therapy

Treatments









Other commonly used combinations:

Darunavair ritonavir Truvada or Kivexa







or

Nevirapine Truvada or Kivexa





Benefits of treatment

- Preservation of immune function
- Improved life expectancy
- Reduced onward transmission
- Reduction of mother to baby transmission
- Fewer hospital admissions

How to do an HIV test

You need one of these....



.....verbal consent and a standard microbiology form – simply request HIV test!

You do not need....

Formal pre test counselling*, written consent, cat 3 stickers (unless other risk), special permission!

* If patient identified as being at high risk of infection GUM health advisors will come and see patient

Resources for General Practice

- Patient information sheet
- Flow chart for testing
- Flow chart for management of positive results
- Practice sessions

Patient information sheet

Testing for HIV

At this practice we offer HIV testing to:

- Everyone wanting to register with us as part of standard medical care.
- ✓ Some people who are already registered and are having blood tests for other reasons.
- Sometimes your doctor may recommend an HIV test, for example if you are at increased risk of HIV or if you have a condition associated with HIV infection.

What is HIV infection?

HIV is a virus that can damage the body's defence system so that it cannot fight off certain infections.

HIV can be found in people who are fit and well, as symptoms can take years to develop.

AIDS is the condition that can occur if someone who is HIV positive goes on to develop certain other serious illnesses. AIDS can be prevented if HIV is diagnosed early enough.

Why do we offer HIV testing?

- ✓ In this area the number of people living with HIV is high enough for National organisations to recommend testing everyone registering with a medical practice
- Excellent medicines have now been developed for treating HIV. This treatment will help to keep you healthy and reduce the risk of you passing the virus on to another person.
- ✓ The earlier you find out about HIV infection, the better the chance of the treatment working.

Who is at risk of HIV?

Anyone who has been in a sexual relationship or has had a needle or blood exposure can have HIV. However, some groups of people are at higher risk. These include:

- ✓ Men who have sex with men
- People from countries such as Africa, South East Asia and parts of Eastern Europe where there are higher rates of HIV
- ✓ People who have injected drug
- ✓ People who have had a blood transfusion abroad
- ✓ Those with other health problems known to be more commonly seen in those with HIV

If you want more information about HIV, please ask us. If you decided that you do not want to test for HIV please tell the doctor or nurse looking after you.

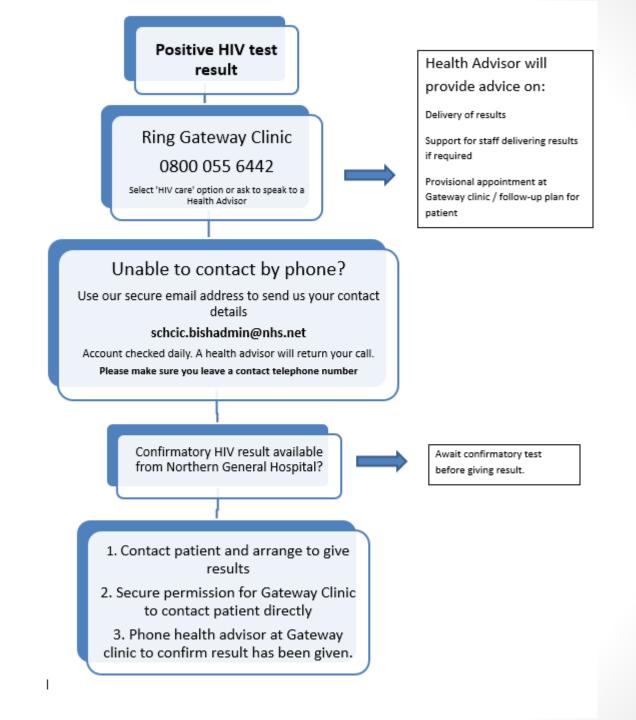
HIV Screening

Who to test:

- 1. All new adult registrants
- 2. Anyone having a blood test who has not been tested within the last 12 months
- 3. People from Africa / South East Asia / Eastern Europe.
- 4. Anyone with a sexually acquired infection
- 5. All men having sex with men
- 6. People who inject drugs
- 7. Anyone with a clinic condition where HIV may be the cause
- 8. Anyone requesting an HIV test

How to test:

- 1. Obtain verbal consent to test
 - Formal pre-test councelling is not required
- Take sample.
 - Brown tube search ICE for HIV and select HIV screening test
- 3. Agree process for getting results with patient
 - Usual practice policy for obtaining other test results is acceptable
- 4. Consider whether a repeat test is required
 - Window period is 4 weeks from last risk



What might you be asked by someone having an HIV test?

- What about insurance?
- How reliable is the test?
- Do I need another test?
- Where can I get more information?
- How do I get my results?
- What happens if I have it?

Prompts to encourage testing

- Add to templates
 - TB templates
 - Hepatitis templates
- New patient forms

Barriers to testing?



Terrence Higgins Trust ♥ @THTorguk · 16m



Prince Harry's message for #HIVTestWeek is clear: get tested and know your HIV status.



Prince Harry plays shopkeeper as he hands out free HIV testing kits

Prince Harry has made a cameo appearance in the job of shopkeeper, as he took a turn handing out free HIV testing kits to the public.

telegraph.co.uk









PEP(SE)
Post Exposure Prophylaxis
(following sexual exposure)

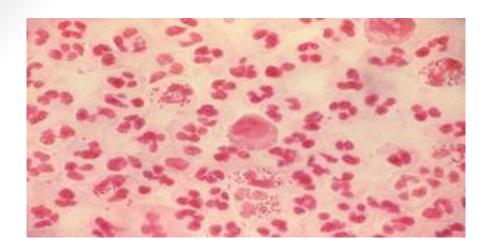
- PEP(SE) is a 4 week course of medication used to treat HIV infection
- Start within 72 hours of exposure risk
- Refer directly to Gateway clinic or A+E if we are closed

Pre Exposure prophylaxis to HIV infection (PrEP)

- PROUD study
- MSM having UPSI were randomised to either start PrEP immediately or delayed start of PrEP for 1 year
- 544 participants (275 in the immediate group, 269 in the deferred group)
- Three HIV infections occurred in the immediate group ($1\cdot2/100$ person-years) versus 20 in the deferred group ($9\cdot0/100$ person-years) despite 174 prescriptions of post-exposure prophylaxis in the deferred group
- Relative reduction 86%,
- 13 men (90% CI 9–23) in a similar population would need access to 1 year of PrEP to avert one HIV infection. (cost about £63 K vs cost HIV £380 k) Also improved broader sexual health and wellbeing

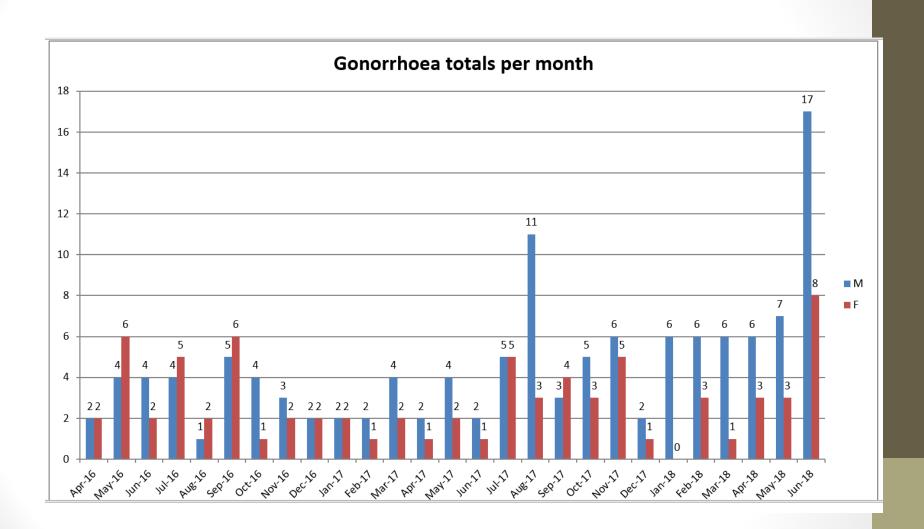
Chem Sex

- Taking drugs and having sex
- Increasing prevalence in the UK
- Current focus on MSM (although HS may also practice Chem Sex)
- Common drugs include GHB, Methamphetamine (Crystal meth, Ice), mephadrone (meow meow, M-CAT)



Gonorrhoea

Testing Management



Background

- Gram negative intracellular diplococci
- 50% cases in < 25 yrs
- 36% cases in MSM
- Asymptomatic / dysuria, discharge, LAP pain, dyspareunia



Tests

Women

Vulvo vaginal swab (CT and GC NAAT)

Men

Urine (1st void) NAAT CT/GC

MSM

- Urine (1st void) NAAT CT/GC
- Pharyngeal swab NAAT CT/GC
- Rectal swab NAAT CT/GC



Genital Gonorrhoea in a male patient

Culture

- 100% specificity
- 100% positive predictive value
- 85-95% sensitivity
- Enriched medium, CO2 enriched environment,>90% humidity.
 - Contains antibiotic/antifungal to prevent overgrowth (combinations of vancomycin / linomycin, colistin/trimethoprim and nystatin/amphoteracin)
- Can do swabs and store at 4 degrees, loss of viability after 48 hours.

NAAT

- Better sensitivity for extra-genital sites
- Sensitivity good for:
 - urine and urethral swabs in men
 - Endocervical and vaginal swabs in women Not good for urine in women.
- False positives with other Neisseria species test on 2 primers

Management of positive results

- Offer confirmed contacts epidemiological treatment at first visit
- Ensure 2 NAAT tests have been performed
- Ensure cultures taken form all relevant sites

- TOC following treatment:
 - 2/52 for NAAT tests
 - 3/7 for cultures
- HA for contact tracing

Treatment

First line:

- Cefixime 400mg po stat with azithromycin 1 g po stat or
- Ceftriaxone 500mg IM stat with azithromycin 1 g po stat

MSM or acquired overseas: ceftriaxone 500mg IM stat with azithromycin 1 g po stat

Alternatives: ciprofloxacin 500mg po stat with azithromycin 1 g po stat or

azithromycin 2g po stat

Pregnancy/breastfeeding:

- Ceftriaxone 500mg IM stat with azithromycin 1 g po stat or
- Cefixime 400mg po stat with azithromycin 1 g po stat

(Manufacturers advise that azithromycin should only be used if no suitable alternatives.)

Penicillin allergy: ciprofloxacin 400mg po stat with azithromycin 1g po stat or

azithromycin 2g po stat

Pharyngeal infection: ceftriaxone 500mg IM stat with azithromycin 1g po stat or discuss with Consultant.