The effectiveness of indicator condition based HIV testing across Europe: results from HIDES-2, a prospective multi-centre study

Michael Rayment on behalf of the HIV Indicator Diseases Across Europe (HIDES) Study Group

British HIV Association Annual Conference April 2015

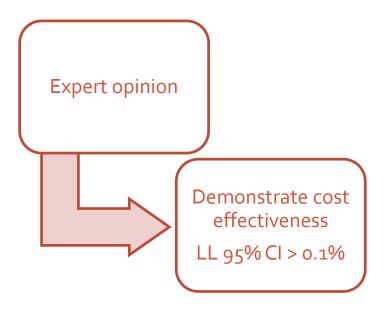


What is indicator condition based HIV testing?

- Indicator conditions are conditions known to be, or believed to be, associated with an excess risk of being HIV-positive
- There is a paucity of evidence of prevalence of previously undiagnosed HIV in indicator conditions
- Indicator condition based testing may be an effective HIV testing strategy
- Routine HIV testing is cost effective when the undiagnosed HIV prevalence in the target group >0.1%

Objectives of HIV Indicator Diseases Across Europe Study

 To implement large scale surveys to prospectively assess prevalence of previously undiagnosed HIV in patients presenting for care of putative, non-AIDS defining indicator conditions





HIV Indicator Diseases Across Europe Study – Phase 2

- Open call to European centres
- Routine offer of HIV test to patients (18-65 yrs) presenting with indicator condition
- Simple demographic data collected; additional data items for those newly diagnosed HIV+
- Primary endpoint:
 demonstration of previously
 undiagnosed HIV infection >0.1%
 in each indicator condition (IC)
- Projected n=11 000
- Open 2012 2014

Disease Area	Indicator Conditions
Malignancies	Lymphoma Cervical dysplasia or cancer (CIN II and above) Anal dysplasia or cancer (AIN II and above) Primary lung cancer
Viral infections	Hepatitis B infection Hepatitis C infection Hepatitis B & C co-infection Ongoing mononucleosis-like illness
Haematological disorders	Leucocytopaenia and / or thrombocytopaenia Lymphadenopathy
Dermatological	Severe psoriasis Seborrhoeic dermatitis
Other	Pneumonia (hospitalised) Peripheral neuropathy

Enrolment

- 150 surveys were performed, across 42 clinical centres in 20 countries across four regions of Europe
- 10 139 patients were enrolled
- Excluded participants: 668
- Total of 9471 participants

Recruitment by region	Number enrolled	%
Total	9471	100
South	500	5.3
Central	942	10.0
North	2297	24.3
East	5732	60.5



Characteristics of participants

- 54% male
- Median age 37 yrs (IQR 29 49 yrs)
- 86.6% white
- 14.4% had previously tested for HIV (median time since last test was 1.3 years [IQR 0.4 3.2 years])
- Setting of test:

	Number enrolled	%
Outpatient	4500	47.5
Inpatient	3564	37.6
Primary Care	270	2.9
Unknown	1137	12.0



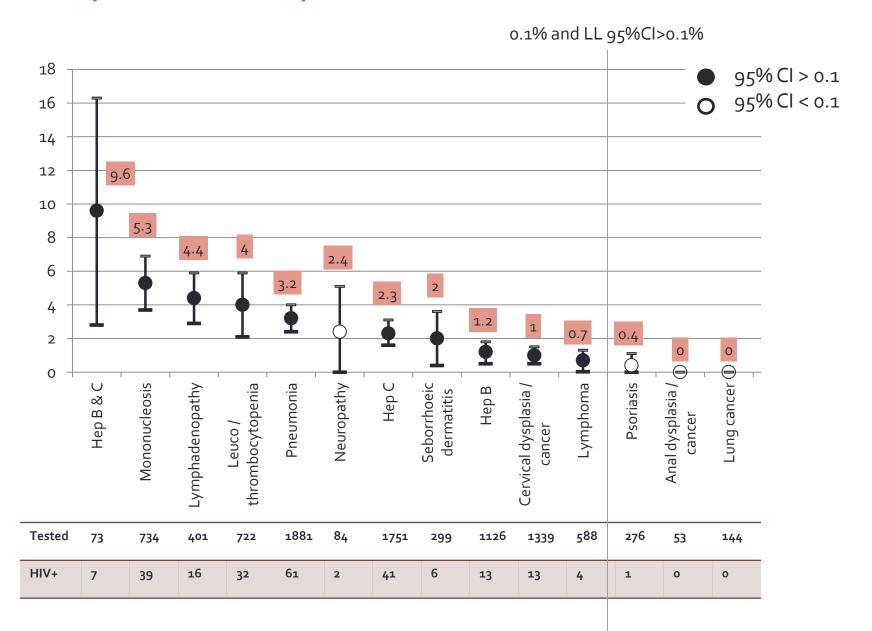
HIV test positivity (overall)

- 235/9471 individuals tested HIV+
- HIV prevalence: 2.5% [95%Cl 2.2 2.8]
- Marked variation by region:

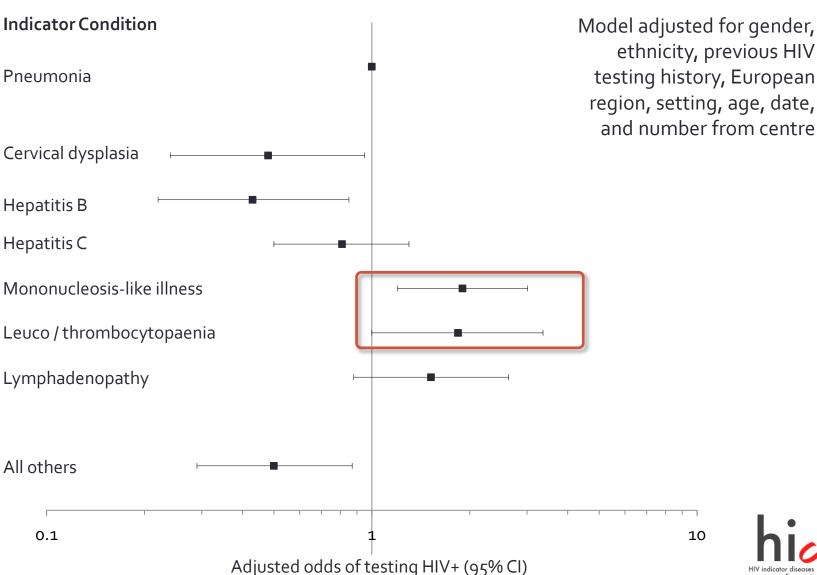
Region	Number enrolled	Number HIV+	%	95%CI
All	9471	235	2.5	2.2 – 2.8
South	500	25	5.0	3.1-6.9
Central	942	10	1.1	0.4 - 1.7
North	2297	31	1.4	0.9 – 1.8
East	5732	169	3.0	2.5 – 3.4



HIV prevalence by indicator condition



Odds of testing HIV+ by indicator condition (adjusted)





Stage of presentation

- Median CD₄ count at diagnosis was 200 cells/μl (IQR 65 390)
- 71.9% (143/235) were late presenters (defined as CD4<350 cells/μl)
- 28.2% (61/216) persons reported prior minor HIV-associated symptoms
- In multivariate analysis, older persons were more likely to be late presenters, as were those seen outside of outpatient departments. Of note, IC and region did not predict late presentation

Conclusions

- Cost effectiveness was established for ten of 14 ICs, in which an HIV prevalence >0.1% was definitively demonstrated
- For the remaining conditions, relatively low numbers of patients were tested, and there were few events



Recommendations and ongoing work

- The conditions with a proven HIV prevalence of >0.1% should be adopted into HIV testing and IC specialty guidelines on both national and European level
- Audits of testing performance in indicator conditions should be performed to evaluate the level of implementation
- An extension of the survey in mononucleosis-like illness is continuing until end of June 2015
- An EU funded project on "Optimising Testing and Linkage to Care for HIV across Europe" (OptTEST) will build on and develop tools for the implementation of IC-guided testing

Acknowledgments

The HIV Indicator Diseases Across Europe Study Group. Centres: Austria: R Zangerle, M Kitchen, University Hospital Innsbruck, Department of Dermatology and Venereology, Innsbruck. Belarus: A Vassilenko, Minsk Municipal Infectious Diseases Hospital, Minsk, VM Mitsura, Gomel State Medical University, Gomel. Belgium: C Necsoi, AF Genotte, Saint-Pierre University Hospital, Brussels. Bosnia: V Hadziosmanovic, Clinical Center, University of Sarajevo, Infectious Diseases Clinic, Sarajevo. Croatia: J Begovac, University Hospital of Infectious Diseases, Zagreb. Denmark: C Pedersen, L Redder, Odense Universitetshospital, UB Dragsted, Roskilde Hospital. France: F Caby, Hôpital de la Pitié-Salpễtriére, Paris, E Bouvet, Hôpital Bichat Claude Bernard, Paris, MA Khuong, Hỗpital Delafontaine, St. Denis, Centre Hospitalier René Dubois, Pontoise, S Caldato, P Morlat, Hôpital Saint-André, Bordeaux, D Coban, CHU de Clermont-Ferrand, Clermont-Ferrand, C Arvieux, Hôpital Pontchaillou, Rennes, F Ajana, Centre Hospitalier de Tourcoing, Tourcoing, A Cabié, CHU Fort-de-France, Fortde-France, Matinique. Georgia: N Chkhartishvili, Infectious Diseases, AIDS and Clinical Immunology Research Center, Tbilisi, Georgia. Germany: J Rockstroh, Immunologische Ambulanz, Department of Medicine, University of Bonn, S Gerdes, Universität-Hautklinik Kiel, Kiel Greece: H Sambatakou, Ippokration General Hospital, Athens. Israel: Z M Sthoeger, D Elbirt, Ben Ari Institute of Clinical Immunology, Rehovot. Italy: A d'Arminio Monforte, L Comi, T Bini, Unit of Infectious Diseases, San Paolo Hospital, Milan, BM Celesia, M Gussio, Unit of Infectious Diseases, University of Catania, Catania, Sicily. Netherlands: K Brinkman, Onze Lieve Vrouwe Gasthuis, Internal Medicine, Amsterdam. **Poland:** A Grzeszczuk, Medical University of Bialystok, Department of Infectious Diseases and Hepatology, Bialystok. Romania: EC Rosca, University of Medicine and Pharmacy Victor Babes, Clinical Emergency County Hospital, Timisoara. Serbia: D Pesut, Clinical Centre of Serbia, Teaching Hospital of Lung Diseases, Belgrade. Spain: MA Goenaga Sánchez, Hospital Donostia, San Sebastian, V P Estrada, Hospital Universitario San Carlos, Madrid, E Ortega Gonzalez, Consorcio Hospital General Univ de Valencia, Valencia, A Ocampo, Complexo Xeral Cies de Vigo, Vigo, M Masiá, Hospital Universitario de Elche, Elche, C Agustí, CEEISCAT, Badalona. Sweden: A Sönnerborg, Department of Infectious Diseases, Karolinska University Hospital, Stockholm. Switzerland: P Vernazza, B Bertisch, Kantonsspital, St Gallen. United Kingdom: A Sullivan, M Rayment, C Rae, Chelsea and Westminster Hospital, London, S Morris, Western General Hospital, Edinburgh, J Anderson, Homerton University Hospital, London, A Palfreeman, Leicester, J Minton, J Calderwood, St James's University Hospital, Leeds, p Farazmand, Huddersfield Royal Infirmary, West Yorkshire, ELC Ong, The Newcastle upon Tyne Hospital, Newcastle, D Mummery, North End Medical Centre, London. Ukraine: G Kutsyna, Luhansk AIDS Center, Luhansk, A Kuznetsova, Kharkov Regional Clinic of Infectious Diseases, Kharkov. Advisory Group: N Clumeck, Saint-Pierre University Hospital, Brussels, Belgium, J Gatell, Hospital Clinic de Barcelona, Barcelona, Spain, B Gazzard, Chelsea and Westminster Hospital, London, England, J Lundgren, University of Copenhagen and Rigshospitalet, Copenhagen, Denmark, A d'Arminio Monforte, Unit of Infectious Diseases, San Paolo Hospital, University of Milan, Milan, Italy, J Rockstroh, Department of Medicine, University of Bonn, Germany, A Mocroft, University College London Medical School, UK, Y Yazdanpanah, Hopital Bichat Claude Bernard, Paris, France. Coordinating Centre Staff: A Sullivan, Chelsea and Westminster Hospital, London, UK, K Champenois, Inserm U738, ATIP/AVENIR Team, Paris, France, P Lopez, D Raben, M L Jakobsen, R S Brandt, CHIP, Rigshospitalet, Copenhagen, Denmark. Statistical Analysis: A Mocroft, University College London, UK.

