



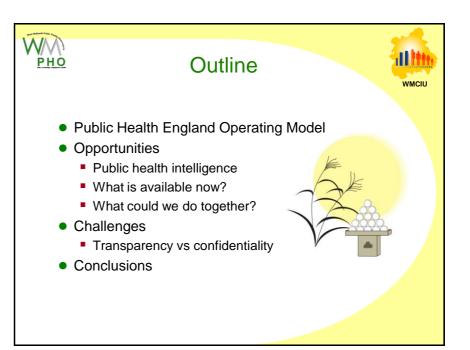
Information to Improve Public Health

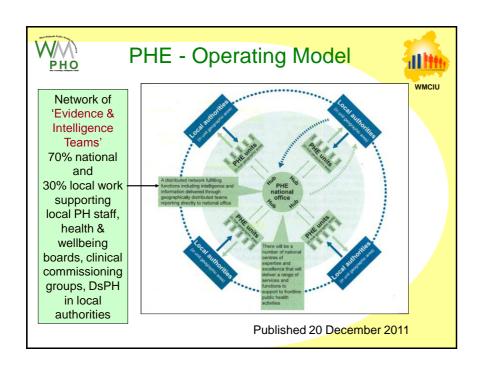
Cancer Outcomes Conference 2012 NEC, Birmingham 14 June 2012

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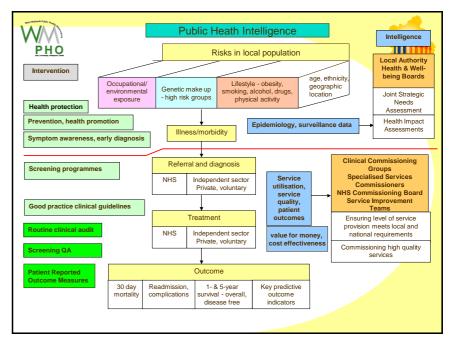


Network Teams

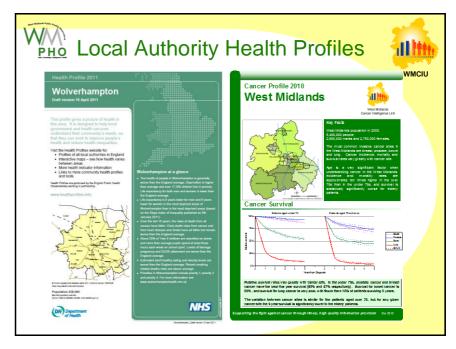


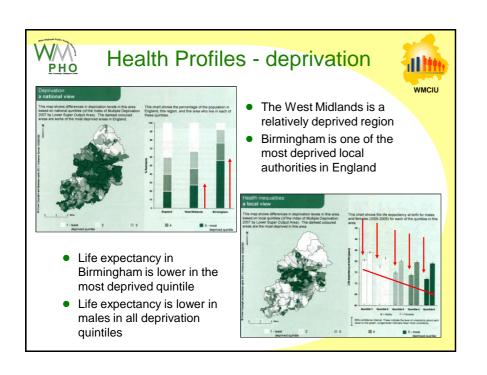
- Will be accountable to 'PHE National Office'
- Will carry out
 - Replicated functions same function performed according to a single service specification by each team agreed with 'PHE National Office' e.g.
 - cancer registration,
 - screening QA visits
 - Distributed lead roles
 - * Roles carried out for whole of England by a single team e.g.
 - NCIN cancer lead roles
 - National Obesity Observatory
 - Child Health and Maternity Information
 - * Integrated work programmes, which could be
 - within a single function NCIN and cancer registry work programme, move to single national cancer registration database
 - across functions PH Outcomes Framework Indicators, joint projects with e.g. PHO, HPA, NTA, cancer screening QARC

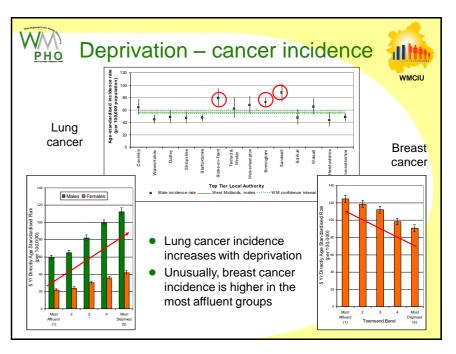


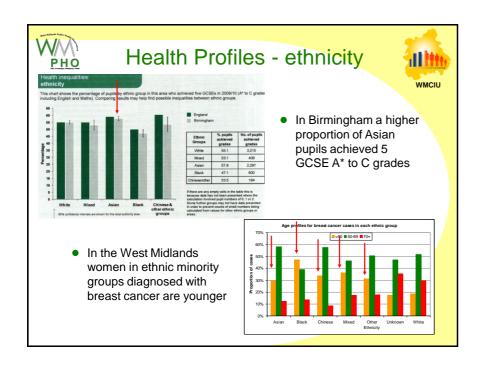


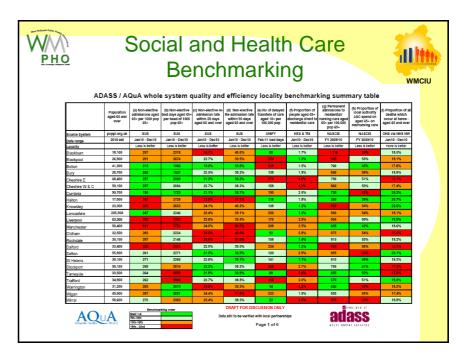




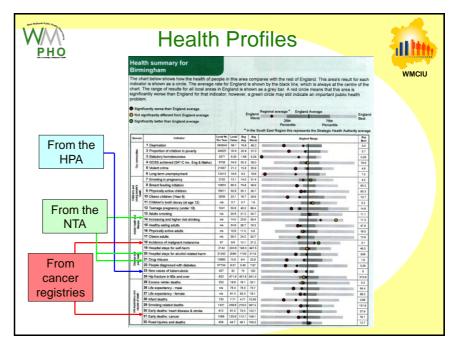


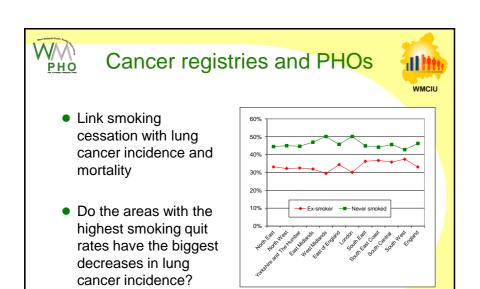


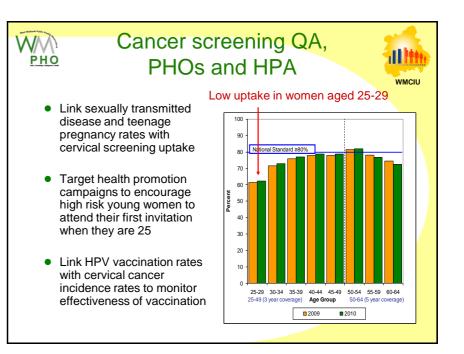












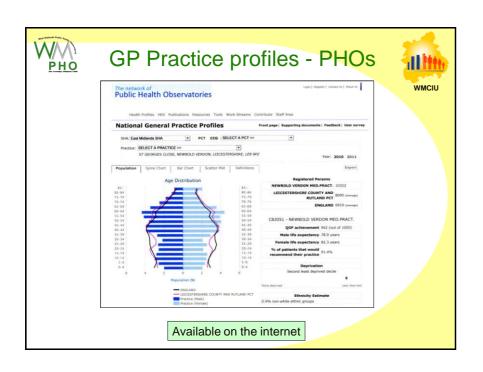




Data Transparency



- Transparency and release of open data throughout the health and social care sector are to be driven forward in line with the policies set out in the Coalition agreement and the Public Data Principles agreed by the Public Sector Transparency Board
- Information Strategy includes plans to:
 - *release large routine public service datasets to analysts
 - provide support for individuals and organisations to obtain fast, easy access to health and social care data
 - ❖ give access for service users to their own data and records
 - provide user feedback to services



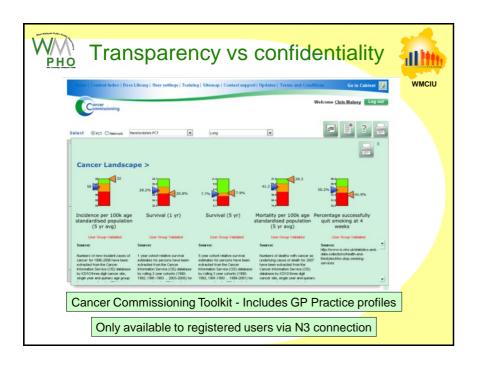


UKACR Guidelines



Guidelines governing the release or publication of potentially identifiable data

- As a general rule, the following categories of data should be regarded as potentially identifiable:
 - Individual records even if they do not include variables, such as names, full postcodes, and dates of birth which would make them obviously identifiable
 - Tabular data, based on small geographic areas, with cell counts of fewer than five cases/events (or where counts of less than five can be inferred by simple arithmetic) – hereafter referred to as "sparse cells"
 - Tabular data containing cells that have underlying population denominators of less than approximately 1,000
 - Tabular data relating to diagnoses/ treatments at provider level, with cells containing counts of fewer than five cases/events





Conclusions



- There is much to gain from the 'sender organisations' working more closely together within Public Health England to provide evidence and intelligence
- Information Governance issues will need to be resolved if equivalent data are to be made available for all types of disease to PH staff working in local authorities and within PHE
- The release of large routine public service data sets as part of the Transparency Agenda means that commercial organisations and community interest groups may be 'in competition' with PHE to provide the intelligence required by local authorities and commissioners to improve the Public's Health

