

Protecting and improving the nation's health

What does the cancer landscape look like now?

The national context for cancer intelligence

Nicky Coombes, SSCRG Programme Manager National Cancer Intelligence Network (NCIN)

The Health & Social Care Act 2012: two new organisations from April 2013

NHS England

- "The purpose of NHS England is to use the £80bn commissioning budget to secure the best possible outcomes for patients"
- To ensure the whole commissioning architecture is in place; will also commission some services directly

Public Health England (PHE)

- Use information & Intelligence to support local Public Health (PH) organsiations and the public to make healthier choices
- National Leadership to PH, supporting national policy
- Development of PH workforce
- A civil service function, not NHS

Independent Cancer Taskforce

- The NHS Five Year Forward View (FYFV) presents a vision for improving health, including for all those diagnosed with cancer: - better prevention - swifter diagnosis - better treatment, care and aftercare
- Taskforce established in January 2015 to produce a new fiveyear national cancer strategy for England, delivering this vision
- Taskforce recommends that this strategy is adopted by the FYFV Board: Care Quality Commission, Health Education England, Monitor, NHS England, NICE, Public Health England, Trust Development Authority



ACHIEVING WORLD-CLASS CANCER OUTCOMES

A STRATEGY FOR ENGLAND 2015-2020



Summary

Save many thousands more lives

Transform patient experience and quality of life

Invest to save

Slides courtesy of:



Six strategic priorities

Spearhead a radical upgrade in prevention and public health

Drive a national ambition to achieve earlier diagnosis

Establish patient
experience on par with
clinical effectiveness
and safety

Transform our approach to support people living with and beyond cancer

Make the necessary investments required to deliver a modern, high-quality service

Overhaul the processes for commissioning, accountability and provision



Ambitions

Incidence

Discernible fall in agestandardised incidence

Adult smoking rates should fall to 13% by 2020 Survival

Increase in five and tenyear survival, with 57% of patients surviving ten years or more

Increase in one-year survival to 75%, with a reduction in CCG variation

Reduction in survival deficit for older people Patient experience and quality of life

Continuous improvement in patient experience with a reduction in variation

Continuous improvement in longterm quality of life



Implementation

- Independent Taskforce has set out its recommendations
- FYFV board, comprised of ALB CEOs, has welcomed the report
- Government has welcomed
- Spending review will determine phasing of implementation
- ALBs will set out more detailed plans



Data Drivers

- Government
 - A spotlight on the role of data and transparency
- Commissioning
 - NHS Outcomes Framework
- Regulation
 - New regulation framework (CQC & Monitor)
- The 'public', patients and families
 - (e.g. 'NHSE Friends and Family Test')

Providers of information

Health & Social
Care
Information
Centre
HSCIC

Office for National Statistics ONS

UK and Ireland Cancer Registries UKIACR

National Audits

PHE

including Health Intelligence Networks **NHS England**

Business Intelligence Teams

Incisive Health

Macmillan Cancer Support

Cancer Research UK **Dr Foster** and other intermediaries

Who do we produce intelligence for?

- Clinicians & Clinical Teams
- NHS England (e.g. specialist commissioning)
- Clinical Commissioning Groups
- Health Care Providers
- Local Government
- NICE
- CQC
- Research Community
- National Statistics
- International Cancer Benchmarking Partnership
- Pharmaceutical Industry
- Patients and the public
- Cancer charities

Cancer functions in Public Health England

- Prevention (smoking; obesity; HPV vaccination, etc.)
- Screening and its QA
- Environmental aetiology (including cluster analyses)
- Public Awareness Campaigns (Be Clear on Cancer Campaigns) – links with Local Authorities and Health & Well Being Boards
- Cancer Intelligence:
 - Registration
 - Analysis
 - Reporting
 - Support (e.g. e-learning)
- PHE heavily involved in the Cancer Taskforce





Chief Knowledge Officer's Directorate

CKO: John Newton

Research

Data

Information and Intelligence

Research and development
Deputy Director
Bernadette Hannigan

National disease registration Deputy Director Jem Rashbass



Knowledge and intelligence service
Deputy Director
Peter Bradley

Digital

Business

Digital programme
Deputy Director
Diarmaid Crean

Business planning and commercial development
Deputy Director
Sian Nash

National Cancer Registration Service

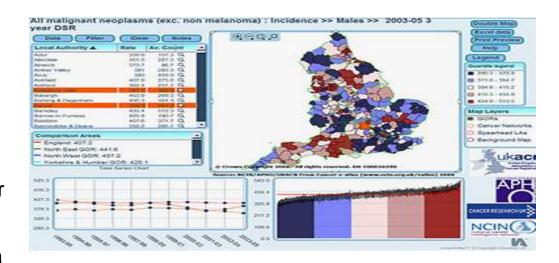
- 8 Regional Cancer Registries now incorporated into a National Cancer Registration Service for England
- 1 Director, 8 local offices
- Complete migration 'normalisation' of 8 legacy systems with 11m cancer registrations to ENCORE
- Same practice and processes, single national system
- Standardised data, consistency, comparability & efficiency

National Cancer Intelligence Network (2007 to date)

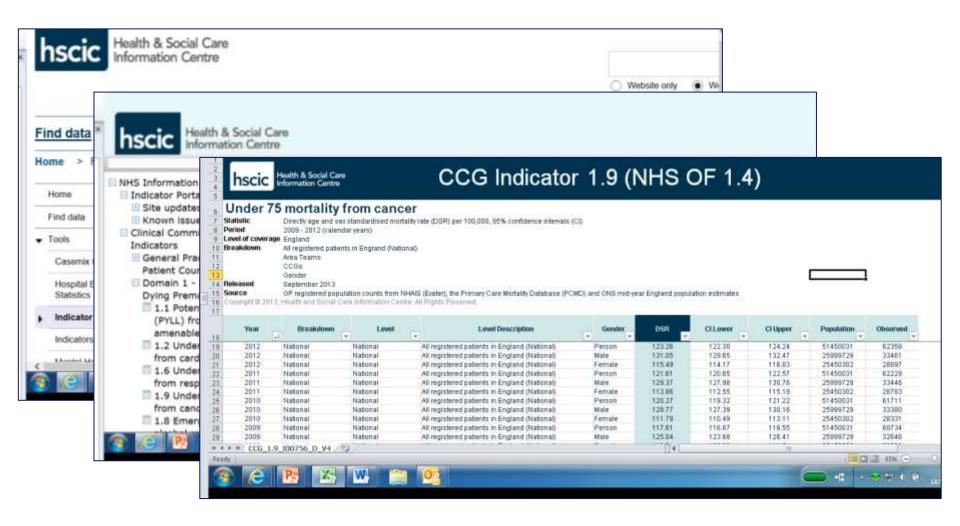
5 aims:

- Enabling use of cancer information to support audit and research programmes
- Exploiting information to drive improvements in cancer care and clinical outcomes
- Providing a common national repository for cancer datasets
- Producing expert analyses to monitor patterns of cancer care
- Promoting efficient and effective data collection

Goal for NCIN: "To develop the best cancer information service of any large country in the world – by 2012"



HSCIC Indicator Portal



CCG Outcomes Indicator Set (cancer)

2013/14

- under 75 mortality rate from cancer
- 1 and 5 year survival from all cancers
- 1 and 5 year survival from breast, lung & colorectal cancers

2014/15 and 2015/16 additional indicators

- cancers diagnosed via emergency routes
- cancer stage at diagnosis
- cancers early detection (stage 1 or 2)
- record of lung cancer stage at diagnosis
- breast cancer mortality

NHS Outcome Framework 2015/16 Dashboard

Outcomes and corresponding indicators used by the Secretary of State to hold NHS England to account for improvements in health outcomes.

Preventing people from dying prematurely

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	Latest data	Indicator value	Unit	
1a.i Potential Years of Life Lost (PYLL) from causes considered amenable to healthcare - Adults	2013	M - 3,167.8 F - 2,470.0	per 100,000 population	
1a.ii - Children and young people	2013	M - 486.5 F - 478.9	per 100,000 population	
1b.i Life expectancy at 75 - Males	2013	11.5	period expectations of	
1b.ii Life expectancy at 75 - Females	2013	13.3	life - years	
1c Neonatal mortality and stillbirths	2013	7.3	per 1,000 births	
Improvement areas				
1.1 Under 75 mortality rate from cardiovascular disease	2013	76.6	per 100,000 population	
1.2 Under 75 mortality rate from respiratory disease	2013	33.2	per 100,000 population	
1.3 Under 75 mortality rate from liver disease	2013	17.7	per 100,000 population	
1.4 Under 75 mortality rate from cancer	2013	141.5	per 100,000 population	
1.4.i One-year survival from all cancers	2011_12	68.2	%	
1.4.ii Five-year survival from all cancers	2007_12	47.9	%	
 1.4.iii One-year survival from breast, lung and colorectal cancer 	2011_12	69.5	%	
1.4.iv Five-year survival from breast, lung and colorectal cancer	2007_12	51.4	%	
1.4.v and vi One- and Five-year survival from cancers diagnosed at stage 1&2	Indicators to be developed			
1.5.i Excess under 75 mortality rate in adults with serious mental illness	2012/13	347.2	SMR percentage	
1.5.ii Excess under 75 mortality rate in adults with common mental illness	Indicator to be developed			
1.5.iii Suicide and mortality from injury of undetermined intent among people with recent contact from NHS services	Indicator to be developed			
1.6.i Infant mortality	2013	3.8	per 1,000 live births	
1.6.ii Five-year survival from all cancers in children	2006_11	81.3	%	
1.7 Excess under 60 mortality rate in adults with a learning disability	Indicator to be developed			

Examples of the clinical value of new data

- Supporting 'intelligent commissioning'
- Demonstration of variation
- Teasing out the causes of variation
- Demonstrating value of specialisation
- Building data into quality improvement
- Adding outcome data into Quality Surveillance Team (Peer Review)
- More meaningful regulation CQC
- Providing robust evidence behind National Guidelines and Quality Standards (NICE)
- Supporting Clinical Trials

Conclusions

- The quality and range of clinically relevant data on cancer is increasing rapidly
- The collection and intelligent use of data are at the heart of good clinical practice and commissioning
- We now have a large and expanding clinical and patient community engaged with cancer data
- Feedback and ongoing interaction with clinicians is an essential part of the process – peer pressure is powerful
- There is a need to improve how information is used at a local level, and we hope this roadshow can help to achieve that.

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