

How does the need for information 'From Mpactomation'

Di Riley



CRS, December 2007



.....Better information on cancer services and outcomes will enhance patient choice, drive up service quality and underpin stronger commissioning; [Chapter 8]

High quality data on:

- · Clinical outcomes, including survival
- adjustments for co-morbidity and stage of disease.

Collection of defined datasets

- all cancer patients
- · mandated through the National Contract.
- PCTs responsible for ensuring delivery

NHS Priorities



patients at the heart of everything we do
focus on improving outcomes for patients –
not inputs or processes, but results.
empower clinicians to deliver improvements.

prioritise prevention and create a public health service.

closer integration in how services are commissioned and provided.

Improving outcomes: our ambition

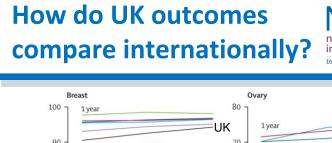


"Our aspiration is that England should achieve cancer outcomes which are comparable with the best in the world"

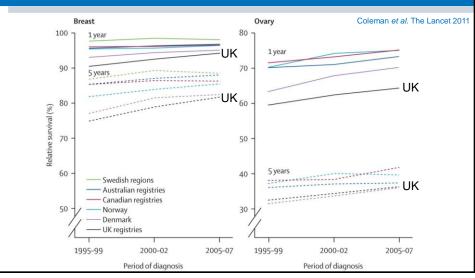


"Quite simply, we want to have the best cancer information service in the world by 2012"

Mike Richards Britain against Cancer Dec 2007

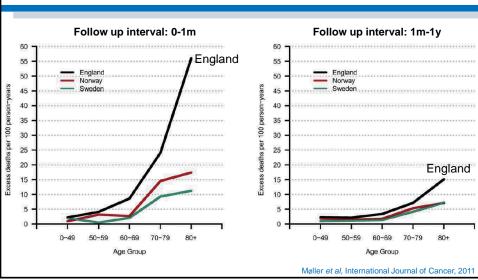






Why is survival poorer in the UK?

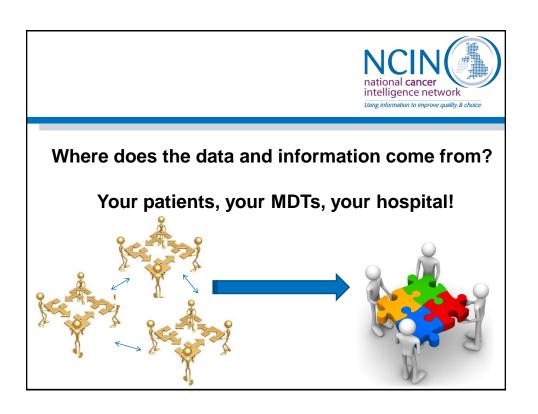


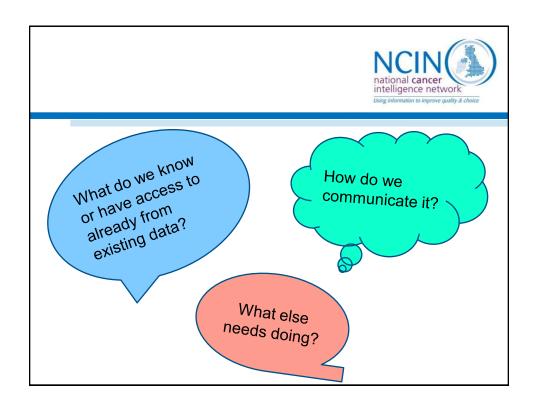


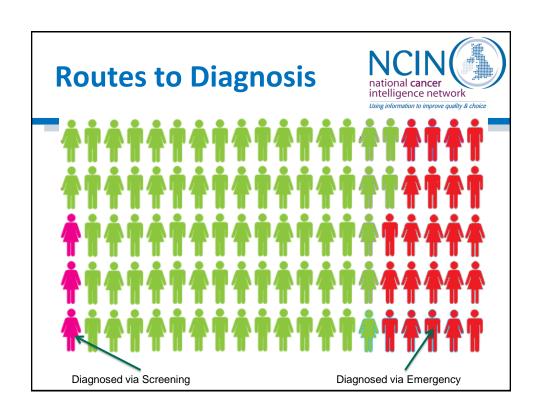
To match the Best in the World

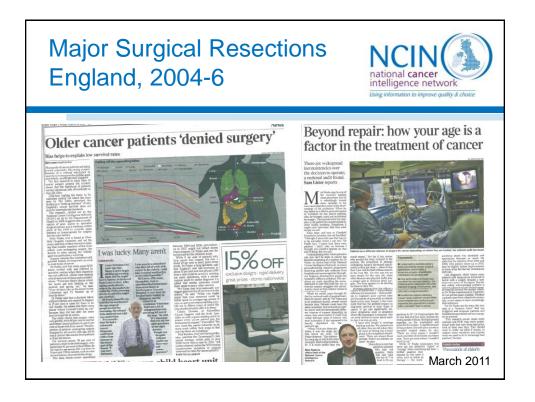


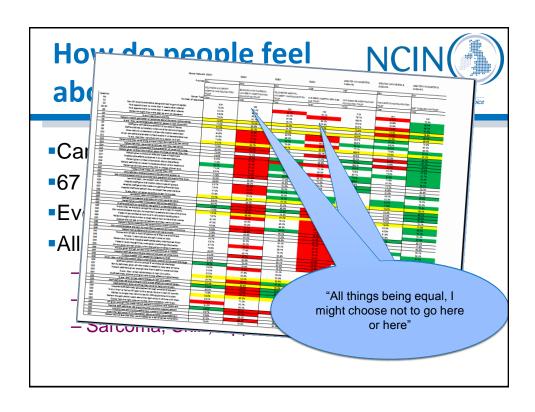
Avoidable Deaths			
Breast	~ 2000	Myeloma	250
Colorectal	~1700	Endometrial	250
Lung	~1300	Leukaemia	240
Oesophagogastric	~ 950	Brain	225
Kidney	~ 700	Melanoma	190
Ovary	~ 500	Cervix	180
NHL/HD	370	Oral/Larynx	170
Bladder	290	Pancreas	75

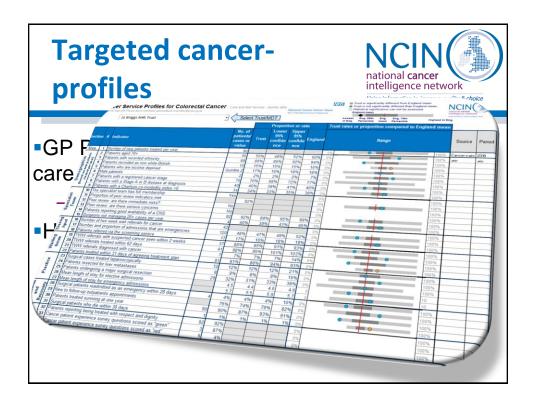


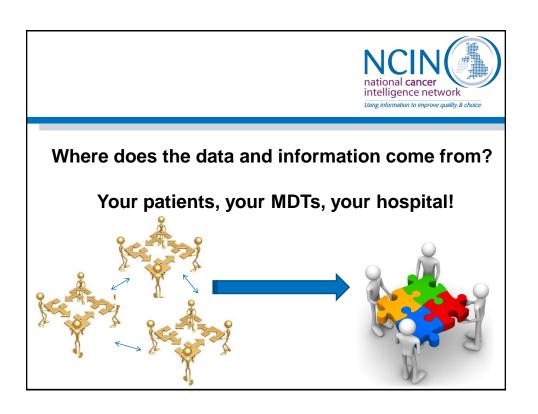


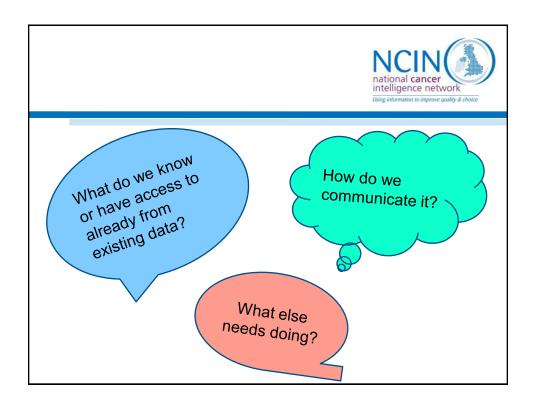








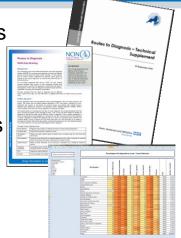




A range of information NCI formats...

- national cancer intelligence network

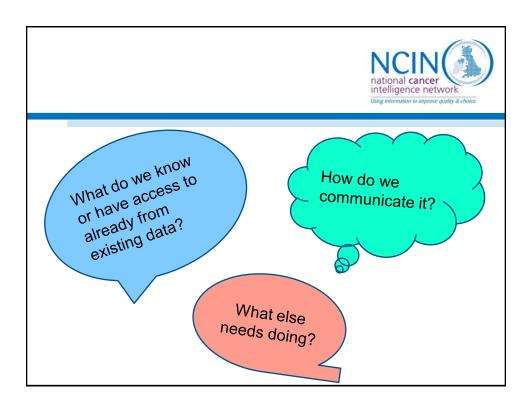
 Using information to improve quality & choice
- Technical/Statistical Reports
- National Data Briefings
- Detailed profiles
- Peer Reviewed Publications
- NHS Toolkits
- Publicly available e-Tools



How to Use Indicators and Data?



- Support Clinical Commissioning Groups to:
 - Understand 'cancer burden' (GP Profile)
 - Understand local services (Service Profile)
- Provide benchmarked information
 - · Understand local services and variation
- Support Patients
 - Understand local services for ME!



Issues?

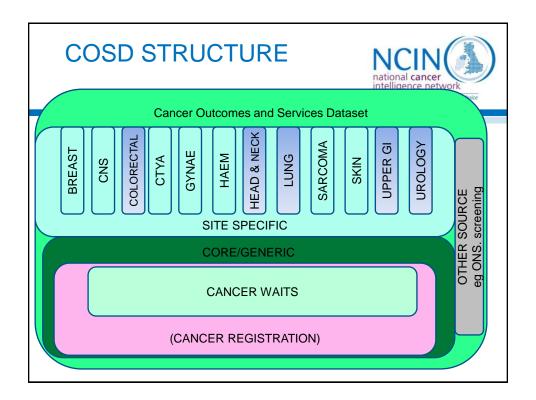


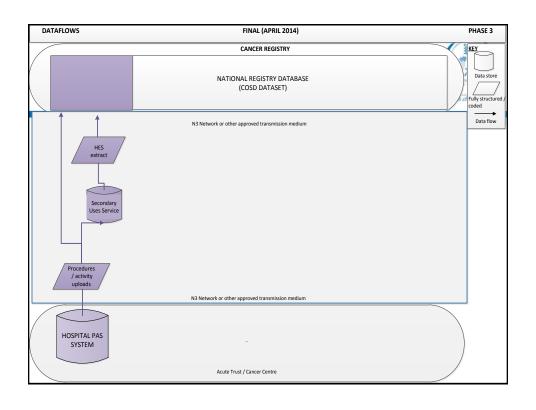
- Are we collecting the correct data
 - Is it specific enough?
 - Can we support the requirements in IOSC?
- Timeliness and quality of data
 - Publish & polish OR
 - Polish & publish
 - Two years old TOO OLD!

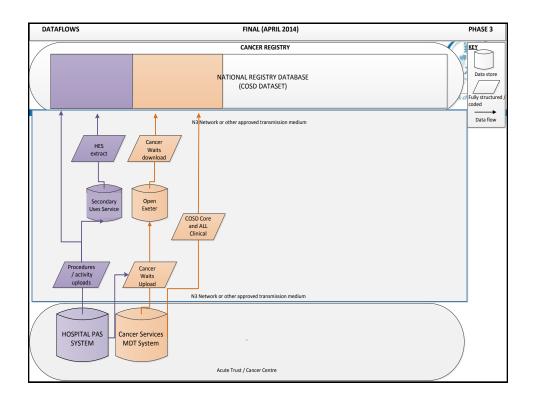
Cancer Outcomes and Services Dataset (COSD)

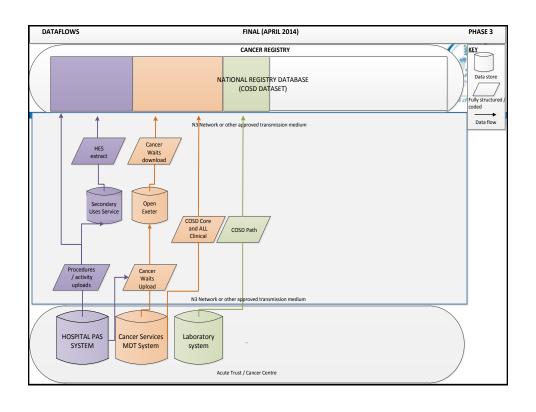


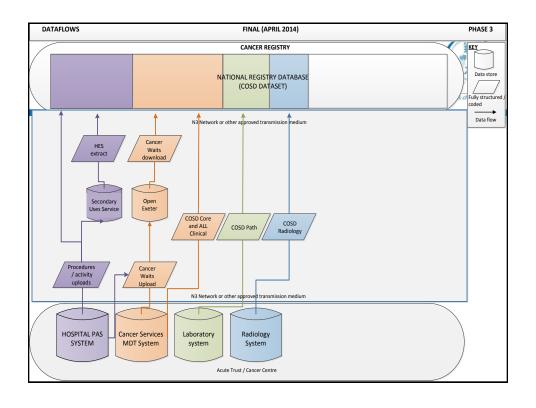
- Replaces 'National Cancer' & 'Cancer Registration' datasets
 - Compiled dataset (data linked and assembled by Registries)
 - · Defines data for direct submission by Trusts
 - · All registerable conditions
 - Wider than CWT
- Components
 - Core (Registration and Cancer Waits)
 - Site specific clinical & pathological

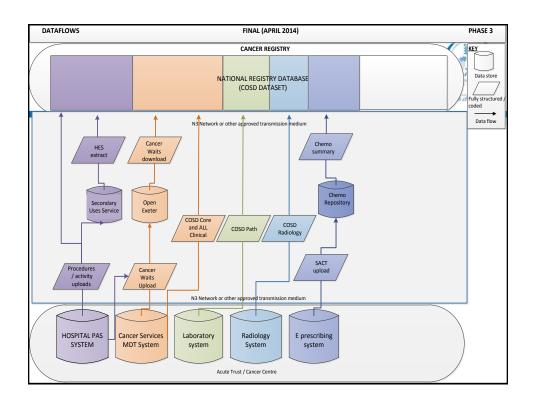


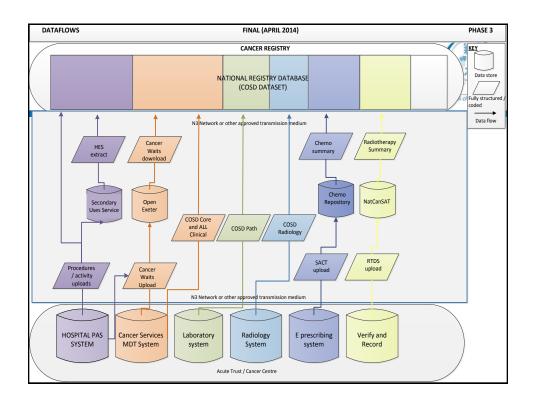


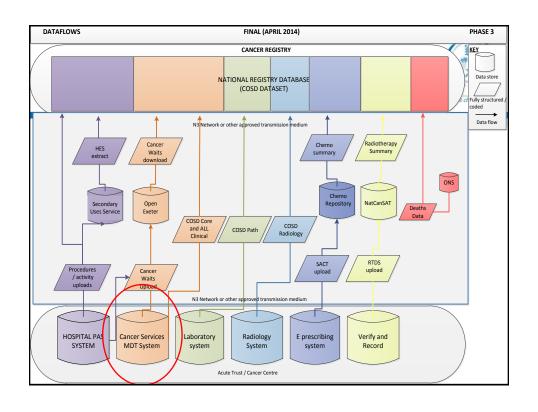


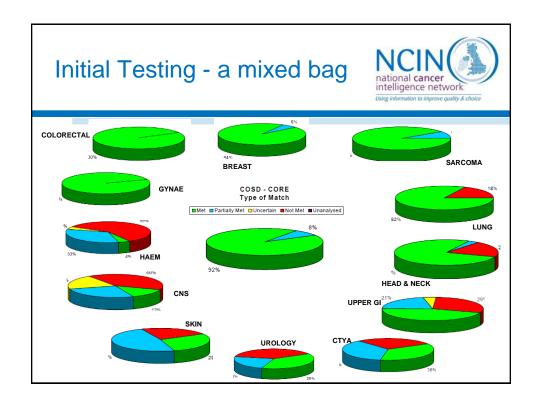








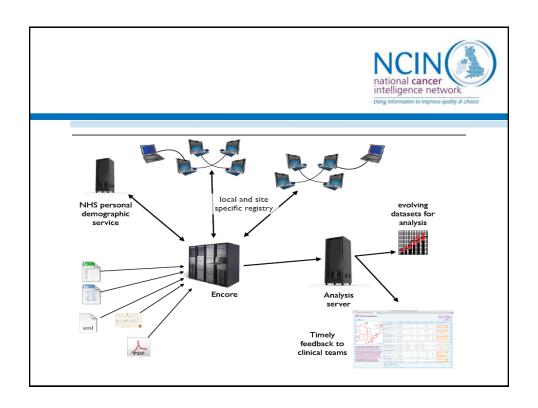


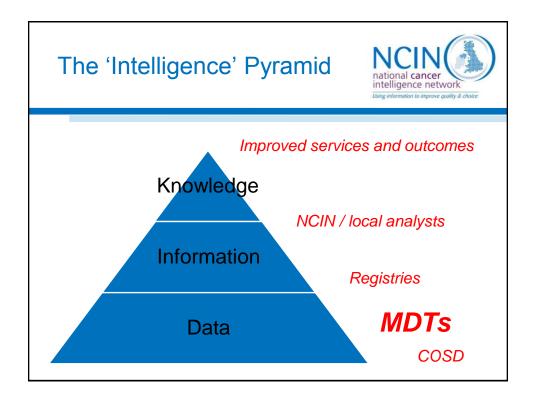


A single Cancer Registration Service for England



- Now need more timely, coordinated data collection
 - By 2013 ONE single Registry for England
 - Central processing, local links
 - More timely, increased quality
 - Closer relationships with local MDTs
 - More rapid feedback processes





In conclusion



- The MDT is central for supporting services and patient care, but also for good data
 - · Patient management & care
 - Understanding services
 - Understanding outcomes
 - Helping patients to choose......

