



Cancer Outcomes and Services Dataset

What is COSD?

Implications for Providers and Networks Sarcoma Workshop January 2013







... Cancer registration and careful monitoring of treatment and outcomes are essential...

Calman-Hine 1995

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

Improving Outcomes: a Strategy for Cancer, 2011

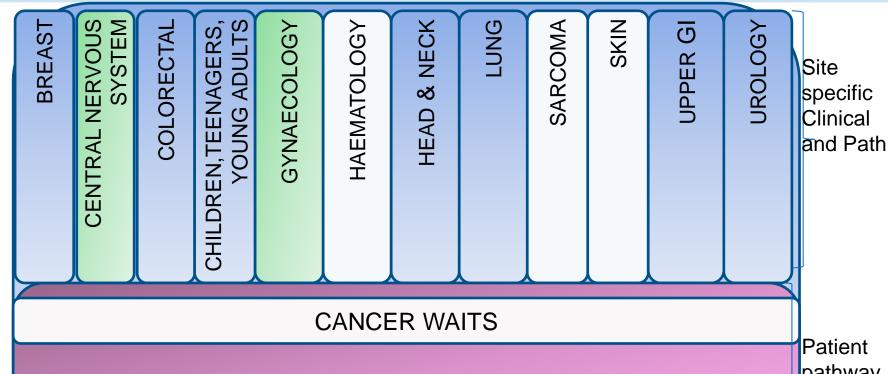
What is COSD?



- The new national cancer dataset
- Incorporates previous cancer registration dataset
- Cancer Outcomes and Services
- Aligned with patient management
- Proposed and supported by clinicians
- Updated and aligned with other datasets
- Clarified definitions of data items, codes and values
- Specifies Provider submissions
- Compiled by registries from Providers and other sources

COSD - Structure





CORE - CANCER REGISTRATION

Demographics/Referral/Diagnostics/Diagnosis/Care Plan/Treatment

Patient pathway referral to treatment

What's different about COSD? (1)



- Complete patient pathway
 - Referral details for all cases
 - All treatments
 - Includes palliative and supportive care
- New core data items including
 - TNM Edition Number
 - Involvement of Clinical Nurse Specialist
 - Duration of symptoms
 - Mandatory for Children, Teenagers, Young Adults (CTYA), Optional for others
 - Year/Month/Day as appropriate or available
- All registerable conditions including
 - in situ bladder, in situ melanoma, benign brain tumours

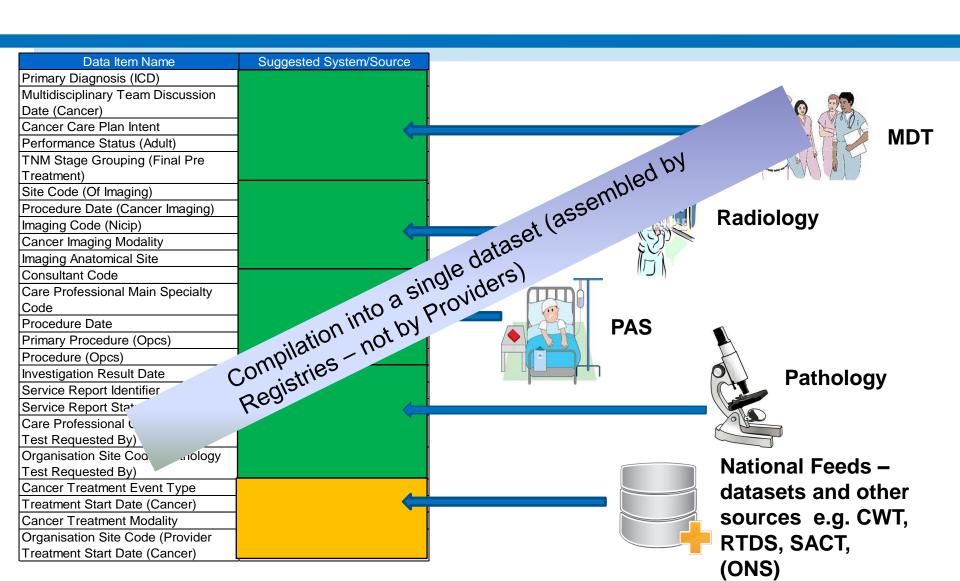
What's different about COSD? (2)



- Site specific data
 - Key site specific clinical items patient management
 - Site specific stage
 - Stage components of RCPath datasets
- Includes recurrences
 - Breast cancers to start with
 - New record including referral information

COSD Dataset

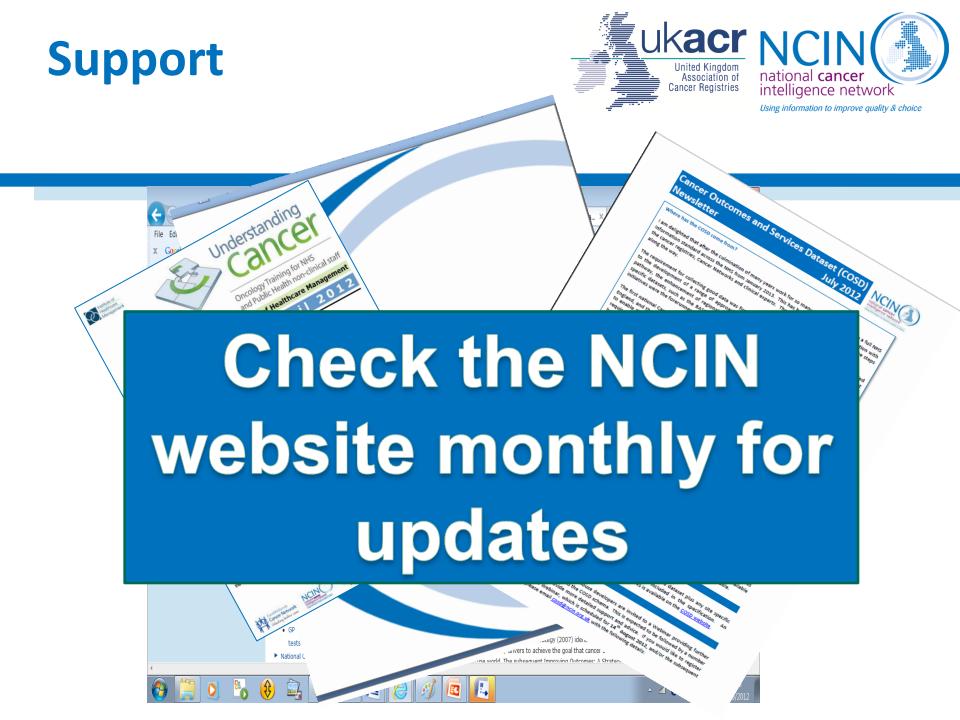




What does this mean for you?

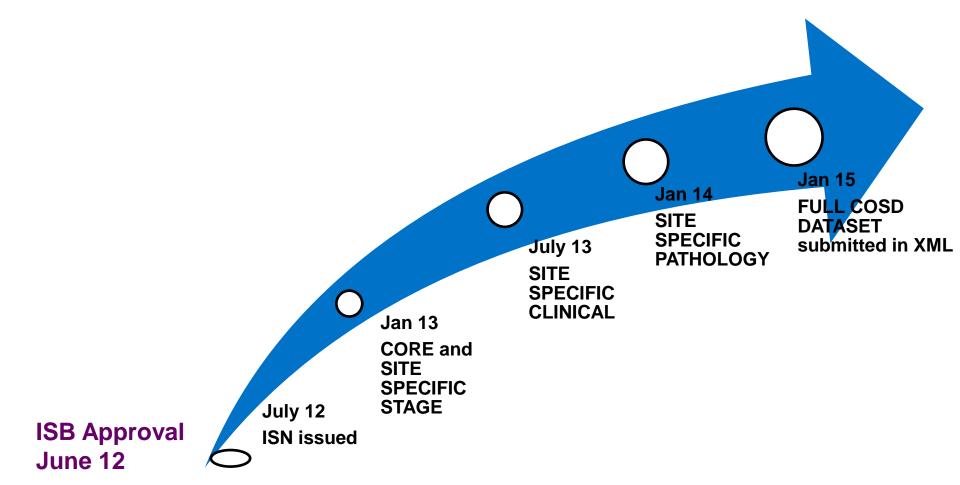


- Monthly submission
 - Current cancer registry feeds expanded to include COSD items
 - 25 working days after diagnosis or treatment
 - Send updates as applicable
 - Aim for three months to complete initial record (to first treatment)
 - Final updates to first treatment within 6 months
 - Further treatments submit 25 working days after treatment
- How to collect in "real time"?
- Clinical ownership/sign off for
 - MDT extract
 - PAS extract
 - Path extract
 - RIS extract



Implementation Timetable





Conformance



- Included in National Contract
 - Information Standard
 - Possible financial penalty
- Simple criteria
 - Monthly feedback to Providers (raw data)
 - e.g. data submitted on time?
 - Staging data completeness
 - Quarterly and annual feedback to follow (processed data)
- Potential Escalation process
 - Informal discussions
 - Notification to CEO
 - Formal notification to commissioners
 - NHS Commissioning Board

Monthly feedback for MDTs- records submitted



- By end of month submitted
 - Number diagnosed
 - Number treated
 - Breakdown of basis of diagnosis
 - % Performance status completed
 - Stage breakdown (at diagnosis/final)
 - % Overall stage grouping completion

Monthly feedback for MDTs records compiled



- Six months after submission month
 - Number diagnosed
 - Number treated
 - Breakdown of basis of diagnosis
 - % Performance status completed (0-4)
 - % Stage grouping completion (at diagnosis/final)
 - % Registry derived stage completion
 - % key pathology items completed
 - Other site specific items -to be agreed in consultation with clinical teams

Quarterly feedback for MDTs records compiled



- Six months after submission quarter
 - Number diagnosed
 - Number treated
 - Breakdown of basis of diagnosis
 - % Performance status completed (0-4)
 - % Stage grouping completion (at diagnosis/final)
 - % Registry derived stage completion
 - % key pathology items completed
 - Other site specific items -to be agreed in consultation with clinical teams
 - Providers to audit progress

Annual feedback for MDTs records compiled



- Six months after final submission
 - Number diagnosed
 - Number treated
 - Breakdown of basis of diagnosis
 - % Performance status completed (0-4)
 - % Stage grouping completion (at diagnosis/final)
 - % Registry derived stage completion
 - % key pathology items completed
 - Other site specific items -to be agreed in consultation with clinical teams
 - 80% expected annual cases (agreed at start of year)



Thank you

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Future Direction for Cancer Registries

What is happening at the cancer registries and how it will support Trusts?

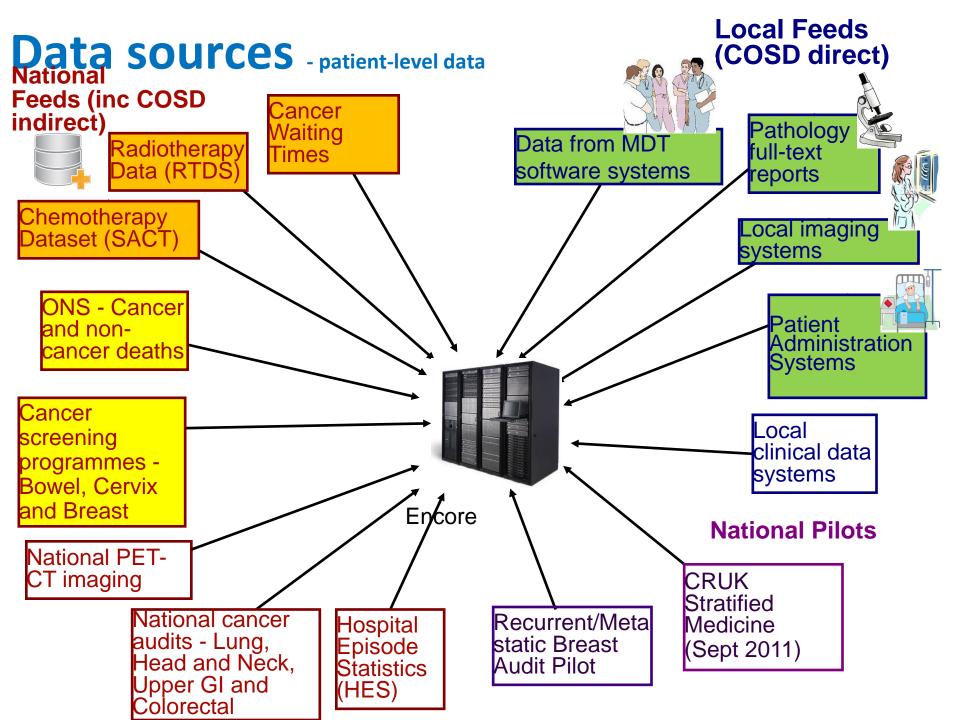


National Cancer Registration Service



- Single England wide system
- Cancer Outcomes and Services Dataset compliant
- Facilitates rapid processing of multiple local and all national data sources
- Common standards and processes including data quality
- Provides rapid and direct feedback of data to clinical teams to enhance data quality
- Support for (near) real-time surveillance, cancer audit and analysis
- Datasets available for external analysis and research

Local data collection at MDTs is key to success





S251 NHS Act exemption

- All cancer registries covered one annual application
- Annual renewal through National Information Governance Board (NIGB) and corresponding statutory instrument

Caldicott Guardian

- sign-off required to allow transfer of data to and from organisations
- existing signoff covers COSD

Information Governance (2)



- Data Protection Act
 - All cancer registries are registered under the DPA.
 - Fair processing notices, including rights to withdraw consent, are provided through the UKACR Patient Information Leaflet (under revision)
 - Latest version of leaflet available from:

http://www.ukacr.org/content/patient-information

About cancer registration

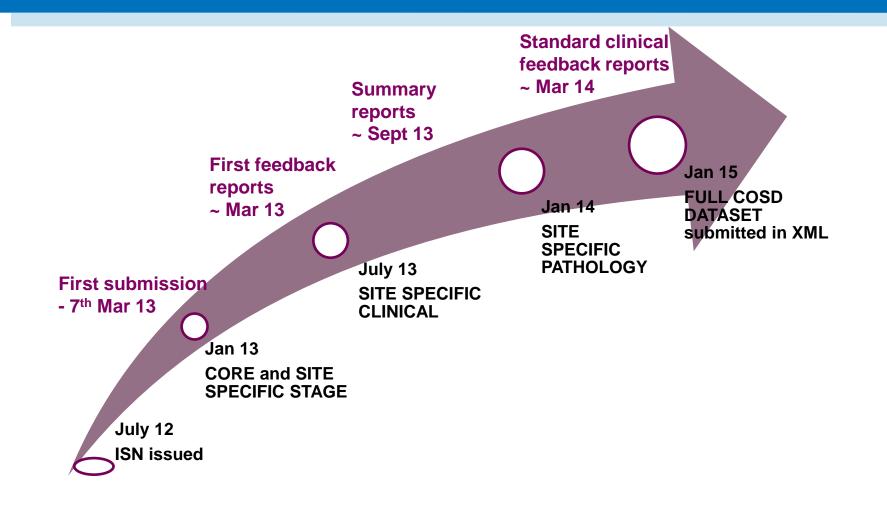
Feedback for Providers and Clinical Teams



- Monthly Conformance Report
 - Extract files received, number of cases, timeliness etc
- Data Quality
 - Initial completeness eg NHS number, stage, performance status, diagnosis
- Summary Information Indicators
 - Eg % histologically verified, RCPath data item completeness
- Clinical and Performance Indicators
 - NICE Quality Standards etc
 - Eg % lung resections
 - % breast conserving surgery with no Radiotherapy

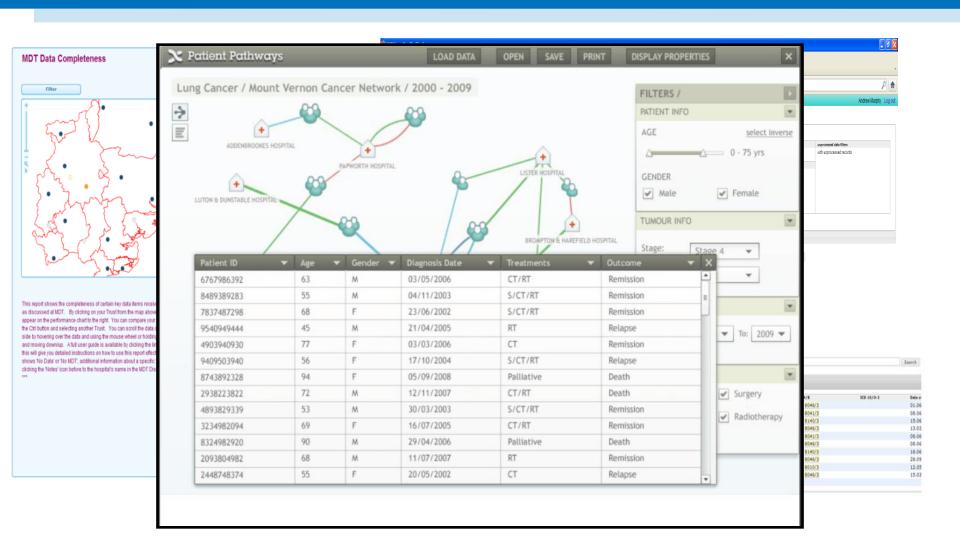
Proposed Feedback Timetable











The Future



- A single cancer registration system (ENCORE) for England
- COSD compliant
- Routine, timely feedback
- Improved registry/clinical partnership working
- Provide powerful data for local, national and international analysis