

# Data Collection through MDT's

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# Overview

- Purpose of MDT's
- The Cancer Reform Strategy
- The North West
- Electronic Data Collection
  - Why?
  - Challenges
- The Way Forward
- Questions



# Purpose Of MDT's

- A team made up of different professions
- Need to draw a distinction between MDTs and MDT meetings
- MDT meetings are typically weekly or fortnightly local or specialist
- Could discuss up to 50 cases per meeting
- But MDTs exist 24/7 and team members are team members outside of the meeting!



# Purpose Of MDT's

- To ensure multidisciplinary input into all decisions affecting patient care
- To ensure that care is given according to recognised guidelines
- To support entry of eligible patients into clinical trials

Manual of Cancer Services Standards, 2000



#### **MDT Data Collection**

- MDT meeting is pivotal point in patient journey
- BUT MDT meeting needs to be well-structured, wellsupported and well-resourced
- Lots of information is collated and collected at MDT meetings
  - Diagnostic tests
  - Staging
  - Treatment plans (surgery, radiotherapy, chemo, watch and wait, palliative care, hormones)
  - Treatment intentions (palliative, curative)



# Impact of the Cancer Reform Strategy

- New targets have been set, covering a much wider number of patients
  - 31 Day Target extend to cover all cancer treatments
  - Bowel Screening extended to include ages 70-75 by 2010
  - All screening detected cancers will be on 62 day pathway
  - Consultants can upgrade patients to 62 day pathway



# Impact of the Cancer Reform Strategy

- Streamlined submission of MDT data within the National Cancer Registration Dataset
  - Electronic submission of datasets to Cancer Registries by 31<sup>st</sup> March 09
- Compliance with National Cancer Audits
  - DAHNO
  - LUCADA
  - NBOCAP
- .... And subsequently
  - NMBRA
  - Oesophageal
- Improvement in the quality of cancer staging data



#### The "North West" Position

- Recognised to be approx 34,000 cancer registrations across the North West
- Retrospective analysis (2007) would highlight the scale of the problem
- Staging data only collected for
  - Breast
  - Colorectal
  - Melanoma
  - Cervix
- Completeness of staging data variable



### The "North West" Position

Table No. & Parameters	Target / Expected Value UK Average		England Average	NWCIS 2005					
Table 3C Completeness of the dataset - specific staging information including grade									
Stage - Breast cancer - % with known Bloom and Richardson grade	>78%	76.4	77.6	79.3					
Stage - Breast cancer - % with known number of positive nodes	>65%	47.2	43.4						
Stage - Breast cancer - % with known invasive size	>65%	63.4	61.3	54.6					
Stage - % with known NPI score	>65%	39.9	37.8	0.8					
Stage - Colorectal cancer - % with known pathological Dukes stage	>74%	51.4	48.5	46.1					
Stage - Cervical cancer - % with complete FIGO stage	>70%	37.3	41.2	17.0					
Stage - Melanoma of the skin cancer - % with known Breslow thickness	>80%	57.0	54.7	23.8					
Stage - Melanoma of the skin cancer - % with known Clark's level	>80%	47.7	42.9	70.4					



#### The "North West" Position

- The retrospective data analysis presents a number of challenges
  - Purpose of the exercise just staging data??
  - Interpretation and abstraction from notes (expertise required??)
  - Pathology reports may miss clinical staging detail
  - Potential staffing of the exercise
  - Costs of the exercise
- Prospective cancer data collection



#### Live Electronic Cancer Data Collection

#### • Why?

- Live data collection via MDT instant validation
- Less duplication of data
- Encourages clinical engagement at MDT meetings
- Data sharing across organisational boundaries
- More efficient use of time and resources
- Legal issues
- Meets the challenges of CRS head on
  - Electronic feed to Cancer Registries with Path data
  - More informative health outcome data clin staging and path staging



#### Live Electronic Cancer Data Collection

# Challenges?

- Complete culture change within MDT
- Multiple data collection tools for tumour sites
- Finding a coordinated solution for each tumour site
- Links with existing hospital systems i.e.
   PAS, Path, Radio
- Ensure that the coordinated solution meets the CRS informatics agenda



# The Way Forward

- Mandation of data
  - NCASP audits (Healthcare Comm)
  - Utilisation of datasets e.g CWT to include staging data (more timely validation)
  - Performance monitoring within network clinical groups
- Real time health professional involvement
- Strong leadership within MDT



#### Where is the "North West"

- Somerset Cancer Register (SCR)
  - Web based clinical data collection
  - Originally in house database (ASW NHS Trust)
  - Initially designed to support CWT collection
  - ASWCS Network supported further development
  - To create a single database that allows "real time" collection across a number of datasets
  - 35 Trusts either using or in the process of implementing SCR across the UK



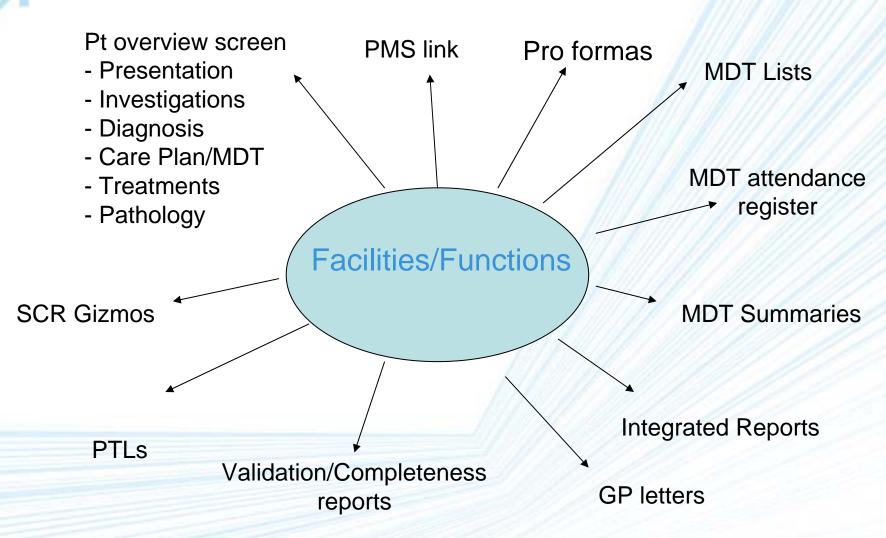
#### Where is the "North West"

Datasets SCR can collect

- National Cancer Waiting Times
- National Cancer Registry
- NCASP
- Royal College
- CSV export facilities for all the above



#### Where is the "North West"





		Mult	idisciplinary Color	ectal Cancer Meetin	<b>d</b>	7,0777				
Patients Name: PARK	ER, Peter D		14/06/1964 (43)	Hospital Number:	-	NHS Number:0000000008				
Consultant:	▼ D	ate First Appointment:	15/03/2008	Recurrence	No 💌	Key Worker:				
Date of Meeting: 05			MPH 💌	Pre/Post Treatmen	it: Pre 💌	Same of the second seco				
Annas Croten German Carl			Park to the later		Days Remaining	1. 9				
Definitive Treatment Target Date: 15/05/2008 Days Remaining: 9  View Referral Symptoms										
Presentation Presentation										
Investigation: Colonos	conv [	Date: 15/03/2008	Investigations	(Abnormal)			-			
View All Investigations										
			Diagno	sis						
Date of Diagnosis:		Diagnosis:				_				
Synchronous Cancer:	<u>*</u>									
Full Colonic Imaging:	No 💌 F	Polyps: No 💌								
Liver Imaging:	No 💌	Cysts: No 💌	Haemangi	oma: No 💌						
Liver Mets:	None 💌 E	Bone Mets: None	Lung Mets	: None 💌	Other Mets: None	<b>*</b>				
Treatment										
Clinical Trial Status:			Donton	✓						
Date of Reporting:		THE CO	Patholo aging: pT  pl							
Chicago de Cara de Car		THAN SE	aging. pr	al Elbat El						
Dukes Stage?	<u> </u>		•							
Local Invasion: Extramural Vascular										
Invasion:										
Nodes Examined:	Positive Nod	es:								
							(A)			
			MDT Decisio	n/Care Plan						
Care Plan Intent:			No Treatment	Reason:		<b>V</b>				
First treatment:			Second treatm	nent:		•				
Referred to:	*		Who Referred	to:	Action By:					
			, , , , , , , , , , , , , , , , , , , ,	1 -						
MDT Comments:							*			
							*			
View All MDTs   Print □   Save   Close										



#### Pre-Treatment Multidisciplinary Colorectal Cancer Meeting

Somerset Health Informatics Service

Date of Meeting:05/05/2008

This management plan relates to the original primary cancer

HHS Humber: 0000000008 Surname: PARKER Hospital Humber: Forename: PETER Consultant: Dr Archer
Date of Decision to Refer: 14/03/2008

Date of Birth: 14/06/1964

Date of First Appt: 15/03/2008

Referral Priority: 2ww Referral

Address:

GP Hame: DR ADAMSON

GP Address: THE OLD SCHOOL SURGERY, MANOR ROAD, FISHPONDS, BS16 2JD

Abnormal

Definitive Treatment Target Date: 15/05/2008

Days Remaining: 9

Presentation

Investigation

Date Investigation Site Outcome Comments

Diagnosis

Date of Diagnosis:

15/03/2008 Colonoscopy

10/05/2008

Diagnosis: Malignant neoplasm of colon

Patient Informed: Ho
Relative/Carer Informed: Ho
Synchronous Cancer: Ho
Full Colonic Imaging: Ho
Liver Imaging: Ho
Liver Mets: Hone
Bone Mets: Hone
Lung Mets: Hone

Treatment

Pathology

THM Staging:

Other Mets:

pT2pHtpM0

Dukes Stage: Extramural B Yes

Hone

Vascular Invasion:

MDT Decision / Care Plan

Care Plan Intent:

CURATIVE

First Treatment: Second Treatment: SURGERY TELETHERAPY (RADIOTHERAPY)

Referred to:

Prof Angelini for Surgery



# "If you are in the business of treating cancer, you are in the business of collecting cancer data"

- Prof Mike Richards



#### Questions?

**Contact Details** 

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