

## Why is co-morbidity important?

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## Co-morbidity in cancer



#### Definition:-

Co-morbidity is a disease or illness affecting a cancer patient in addition to but not as a result of their index (current) cancer.





# Why is co-morbidity important for cancer patients?

- Clinical decision making
- Risk adjusted outcomes analyses
- Highlighted in the CRS
  - Important but variably collected



# What influences cancer decision making?



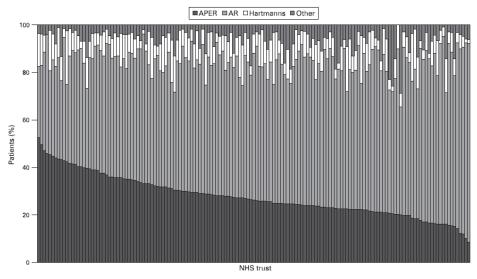
- Tumour factors
  - Stage/grade
- Individual factors
  - Patient preferences
  - Performance status
  - Frailty
  - Fitness
  - CO-MORBIDITY
- To predict outcome personal prognostograms?

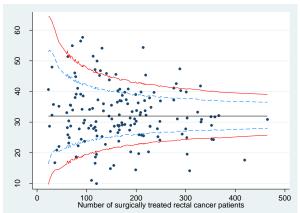


## Unacceptable variation in abdominoperineal excision rates for rectal cancer: time to intervene?

E Morris, 1,2 P Quirke, 2 J D Thomas, 1,2 L Fairley, 4 B Cottier, 3 D Forman 1,4











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#### Main elements



- Selection for treatment
  - Intent, toxicity and mortality
- Late effects:
  - Predicting them
  - Identifying them
- Impact along with stage on overall survival / prognosis
- Is it feasible to expect a single scale to answer all these questions?



# Questionnaire to Site-Specific Clinical Reference Group Chairs



#### In your speciality area, what are:

- the indices/scores are in use?
- the most important ways in which co-morbidity affects treatment and/or outcomes?
- the major C-Ms which impact on treatment decisions and outcomes?
- Do you use 'frailty' as an indicator?
- Other comments



## Site-specific review



	Breast	Colo- rectal	Gynae	Haem	H&N	Lung	Sarcoma	Skin	UGI	TYA
PS	±	+++	+	+++	+	+++	±	++	+++	±
C-M	++	+++	++	+	++	+++	+	+	+++	±
Surgery	+	+++	+	-	++	+++	+	±	+++	±
Chemo	++	++	++	++	++	++	+	+	++	±
RT	++	+	+	±	+	++	±	-	±	±
Peri-op mortality	+	++	+	-	+	+++	+	-	+++	±
Tools	ASA	ASA Possum	UK GOSoc	ACE27 ADL	ACE 27	No (lung function)	No	No	ASA	No
Overall survival	+	++	+	+	++	+			±	±
Late effects	+++	++	+	+++	+	+	+	+	+	+++

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National Cancer Researcl Institute

#### When to record?



## Prospective Recording

- Presence or absence?
- Severity?
- Type of co-morbidity present?
- ACE-27?
- Other scale e.g. ASA?

## Derive retrospectively

- HES favours admitted care
- Accuracy/completeness of coding
- Less timely





## **Adult Co-morbidity Evaluation-27**

prospectively recorded by MDT as part of treatment planning process?



#### ACE-27



- Chart-based comorbidity index for patients with cancer
- Developed through modification of the Kaplan-Feinstein Comorbidity Index (KFI)
- Modifications were made through discussions with clinical experts and a review of the literature
- Validated in study of 19,268 cancer patients treated at Barnes-Jewish Hospital, USA



#### ACE-27



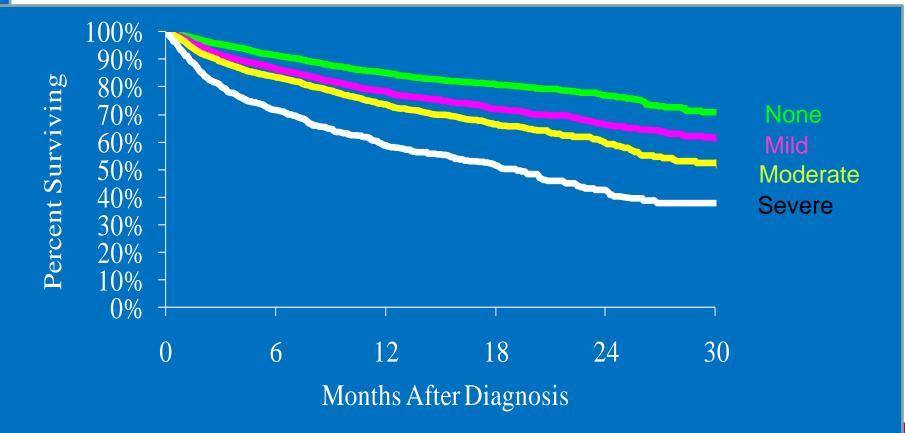
Cogent comorbid ailment	Grade 3 Severe Decompensation	Grade 2 Moderate Decompensation	Grade 1 Mild Decompensation
Cardiovascular Syster			_
Myocardial Infarct	■ MI ≤ 6 months	■ MI > 6 months ago	Old MI by ECG only, age undetermined
Angina / Coronary Artery Disease	■ Unstable angina	<ul> <li>Chronic exertional angina</li> <li>Recent (≤ 6 months) Coronary</li> <li>Artery Bypass Graft (CABG) or</li> <li>Percutaneous Transluminal Coronary</li> <li>Angioplasty (PTCA)</li> <li>Recent (≤ 6 months) coronary stent</li> </ul>	■ ECG or stress test evidence or catheterization evidence of coronary disease without symptoms ■ Angina pectoris not requiring hospitalization ■ CABG or PTCA (>6 mos.) ■ Coronary stent (>6 mos.)
Congestive Heart Failure (CHF)	<ul> <li>Hospitalized for CHF within past 6 months</li> <li>Ejection fraction &lt; 20%</li> </ul>	<ul> <li>Hospitalized for CHF &gt;6 months prior</li> <li>CHF with dyspnea which limits activities</li> </ul>	<ul> <li>CHF with dyspnea which has responded to treatment</li> <li>Exertional dyspnea</li> <li>Paroxysmal Nocturnal Dyspnea (PND)</li> </ul>
Arrhythmias	■ Ventricular arrhythmia ≤ 6 months	<ul> <li>Ventricular arrhythmia &gt; 6 months ago</li> <li>Chronic atrial fibrillation or flutter</li> <li>Pacemaker</li> </ul>	■ Sick Sinus Syndrome
Hypertension	<ul> <li>DBP≥130 mm Hg</li> <li>Severe malignant papilledema or other eye changes</li> <li>Encephalopathy</li> </ul>	<ul> <li>DBP 115-129 mm Hg</li> <li>Secondary cardiovascular symptoms: vertigo, epistaxis, headaches</li> </ul>	■ DBP 90-114 mm Hg ■ DBP <90 mm Hg while taking antihypertensive medications
Venous Disease	<ul> <li>Recent PE (≤ 6 mos.)</li> <li>Use of venous filter for PE's</li> </ul>	<ul> <li>DVT controlled with Coumadin or heparin</li> <li>Old PE &gt; 6 months</li> </ul>	Old DVT no longer treated with Coumadin or Heparin
Peripheral Arterial Disease	■ Bypass or amputation for gangrene or arterial insufficiency < 6 months ago ■ Untreated thoracic or abdominal aneurysm (≥6 cm)	<ul> <li>Bypass or amputation for gangrene or arterial insufficiency &gt; 6 months</li> <li>Chronic insufficiency</li> </ul>	■ Intermittent claudication ■ Untreated thoracic or abdominal aneurysm (< 6 cm) ■ s/p abdominal or thoracic aortic aneurysm repair

http://cancercomorbidity.wustl.edu/ElectronicACE27.aspx Using information to improve quality & choice



# Prognostic Impact of Comorbidity





Log Rank  $\chi^2 = 379.24$ , p < 0.0001





#### **Charlson Score**

derived retrospectively by analysts based on information in notes coded by clinical coders



## **Cancer Diagnosis**



	HES 6	episodes	1 yr previ	us			
				▼			
time							
	<u> </u>	/ 	T		Charlson	0 5	T
HESID DIAG_1 DIAC		DIAG_4	DIAG_5		Group 1	Group Description Acute Myocardial Infarction	Score
5494782 I211 T814 5494782 5494782 0259 -	Y838	1802				Congestive Heart Failure	
5494782 K740 K528	3			₹	3	Peripheral Vascular Disease	1
5494782 C679 - 5494782				•	4	Cerebral Vascular Accident	1
5494782 D171 -					5	Dementia	1
5494782 H332 D569	2853			7		Pulmonary Disease	1
5494782 M720 -						Connective Tissue Disorder	1
-	<del>!</del>	ļ.	-1		8	Peptic Ulcer	1
						Diabetes	
					10	Diabetes Complications	2
					11	Paraplegia	:
Acute Myocardial Infa	rction 1				12	Renal Disease	2
				13	Cancer	2	
Liver Disease	2	——			14	Metastatic Cancer	6
Final Score	3	_					
					15	Severe Liver Disease	3
					16	ПІТ	6
					17	Liver Disease	<del>-</del>



## Complications

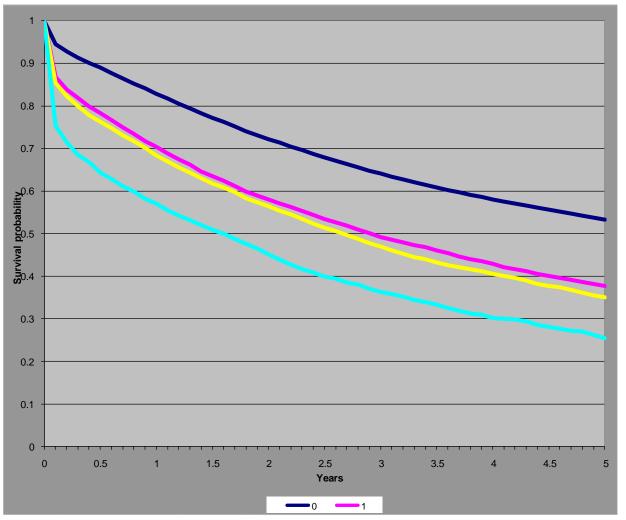


- Score is very dependent on date of cancer diagnosis
  - Differences in registration processes between registries
- Cancer diagnosis is often first in-patient episode
  - Only including episodes prior to diagnosis may miss co-morbidity codes
- Coding of Cancers differ in Registry/HES meaning cancers can be counted twice
  - e.g. an individual colorectal tumour could be coded as C18 in registry and C19 in HES, this could lead to
- Suspected cancer diagnosis coded in HES
  - over-reporting of cancer diagnosis in HES
- Cancers and Metastatic Cancer make up main proportion of scores
  - Should any cancer information be used in the calculation of the score for cancer purposes.
  - Would it be better to use definitive data on multiple tumours/mets



# Colorectal survival by Charlson Score







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## Conclusions



- NCDR has Charlson score available at individual tumour level
- Further analysis required to assess best approach to calculating co-morbidity from available data
- Work with DH/CfH on national co-morbidity project
  - SSCRGs to define pertinent conditions



## Workshop Action Plan



#### Prospective:

- Recommend collection of ACE-27 co-morbidity score is mandated for all adult cancer patients
- Ensure that appropriate training is delivered
- Research different collection methodologies e.g. patient questionnaires
- Identify where supplementary indices or information may be required
- Retrospective:
  - Continue to evaluate co-morbidity scores from HES
- Consider establishing a Co-morbidity 'CRG'





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## Thank you

www.ncin.org.uk

