Appendix 1: Imaging Sequences [posted as supplied by author]

The protocol employed a thin 3mm section turbo spin-echo T2-weighted technique using a surface pelvic phased array coil. For all tumours, scans were performed perpendicular to the long axis of the tumour. Coronal imaging was performed for all tumours arising at, or below, the levator muscle origins. Images were stored in DICOM format on CD. Extramural depth of tumour invasion was measured, for each patient, as the maximum depth of penetration beyond the outer edge of the longitudinal muscle layer, measured using the workstation electronic callipers. The closest distance of tumour to the mesorectal fascia was recorded. Potential circumferential resection margin involvement by tumour was defined as tumour, tumour deposit or, involved lymph node abutting or extending through the mesorectal fascia or, extending <1mm to the mesorectal fascia.

No bowel preparation, air insufflation or intravenous anti-spasmodic agents were used. For a 1.5T magnetic resonance imaging scanner, four sequences were used:

- 1. After a coronal localiser, sagittal scans were required from inner pelvic sidewall to sidewall using a 24cm field of view, 5mm contiguous/interleaved slices (no gap), TR>2500 and <5000, TR=85. These acquisitions were used to plan thin section oblique axial images.
- 2. Axial T2FSE acquisitions of the anatomic pelvis by using a 24cm field of view, a 5millimetre contiguous section thickness, 4000/85, 512 x 256 matrix, an echo train length of eight, no fat saturation, a 32kHz bandwidth, and two signals acquisitions (2NEX).
- 3. The sagittal T2 weighted images obtained were then used to plan T2-weighted thin-section axial images through the rectal cancer and adjacent peri-rectal tissues. These images were performed perpendicular to the long-axis of the rectum. These were obtained by using a 16cm field of view, a 3mm section thickness, no intersection gap, 4000/85, a 256 x 256 matrix, an echo train length of eight, no fat saturation, a 32kHz bandwidth and four acquisitions (4 NEX).
- 4. For low tumours these sequences were repeated with imaging in the coronal plane.

For a 1.0T magnetic resonance imaging Scanner, the sequences were similar with a modification of the imaging parameters to obtain an adequate SNR. The high resolution images are obtained with 20cm field of view, 3mm section thickness, no intersection gap, a 256 x 256 matrix, a TR >2500 (<5000), and a TE > 80.

Appendix 2. Example of reporting proforma used in study [posted as supplied by author]

Code No:	MRI Reporting	
	Proforma	Addressograph
D 1: 1 : :		
Radiologist Patient Name:	Date:	/ /200
Date of Birth:	/ / Hosp. No	
Exam performed elsew		No If yes, where
Exam technically satis		No
Image quality		ub-Optimal
Pathology identified Has the patient receive	Yes ed Radiotherapy Yes	No No
Has the patient had a previous rectal MRI Yes No		
If Yes, date of previou		
Gross Morphology		
Polypoidal	Annular ulcerating Annu	ular non ulcerating
Infiltrating margin of	f extramural spread	
Eroding	Pushing Infiltrating	No Extramural spread
Mucinous Tumour	Yes No	
Metastatic spread	Nodes demonstrated not suspicious	Yes No No
	Nodes demonstrated suspicious	Yes No No
	Extramural venous invasion	Yes No No
	Tumour deposits / satellites present	Yes No No
Local invasion	Submucosa (T1)	Muscularis (T2)
Beyond Muscularis <1.00 mm (T3a) Beyond Muscularis 1.01-5.00 mm (T3b)		
Beyond Muscularis 5.01-15.00mm (T3c) Beyond muscularis >15.01mm (T3d)		
Into adjacent organs (T4a) Perforation of visceral peritoneum (T4b)		
Margins		
Distance to mesorectal fa	ascia <1.00 mm Me1 Distance to m	esorectal fascia >1.01 mm Me0
Low tumour (below leva	tor) >T2 MeLev	
Measurements	Maximum extramural spread of tumo	ourmm
Min distance to mesor	ectal fascia/potential CRM from outer e	edge of tumourmm
Please state distance to	CRM for:	
a. Main tumourmm		
b. Suspicious lymph nodec. Extramural venous invasion		mm
c. Extramural venous invasion d. Tumour satellite/deposit		mm mm
u. Tumo	a. Satemie deposit	
Distan	ace to sphincter (Low tumours only)	mm

Members of the MERCURY Study Group [posted as supplied by author] P Toomey (surgeon), M A Raja (surgeon), C D George (radiologist), L Temple (pathologist), S Woodward (nurse specialist), I Swift (surgeon), M Abulafi (surgeon), N Bees (radiologist), H Blake (radiologist), N Jeyadeven (radiologist), A Arnaout (pathologist), G Brown (radiologist), A Wotherspoon (pathologist), A Massey (nurse specialist), R J Heald (surgeon), B J Moran (surgeon), T D Cecil (surgeon), D M Gold (surgeon), P D Peppercorn (radiologist), J Finch (pathologist), I Ilesley (pathologist), A Leppington-Clarke (nurse specialist), P J Finan (surgeon), P Sagar (surgeon), D Burke (surgeon), K Sasapu (research fellow), A Chalmers (radiologist), P Quirke (pathologist), A Cairns (pathologist), N Chauhan-Lall (YCRN trials coordinator), S Ambrose (surgeon), I Botterill (surgeon), D Jayne (surgeon), A Guthrie (radiologist), N Scott (pathologist), C Verbeke (pathologist), J Wiig (surgeon), K Kotanska-Groeholt (pathologist), H Emblemsvaag (radiologist), T Vetrhus (radiologist), Knut Haakon Hole (radiologist), S Larsen (surgeon), M Gudgeon (surgeon), D Edwards (surgeon), S Mellor (surgeon), H Massouh (radiologist), M Elmahallawy (pathologist), K Bundy (nurse specialist), Donaldson (surgeon), H J Scott (surgeon), P Bearn (surgeon), K Galbraith (surgeon), M Creagh (radiologist), S Dodd (pathologist), L De Snoo (nurse specialist), J Strassburg (surgeon), K Ludwig (surgeon), A Weskott (surgeon), A Lewin (surgeon), M Frei (research assistant), P Knuth (radiologist), J Linke (pathologist), V Loy (pathologist), A G Radcliffe (surgeon), J Torkington (surgeon), R Bleehen (radiologist), N S Dallimore (pathologist), Y Perston (nurse specialist), M W Bourne (radiologist), T Holm (surgeon), L Blomqvist (radiologist), J Lindholm (pathologist), M Torkzad (radiologist), Ms B Andersson (nurse specialist), Ms Y Ericsson-Alm (nurse specialist).