Supplementary table 1

Table 1: Protocol for MRE image acquisition

	Plane	**Slice Width (mm)**	**FOV**	**TR/TE**
3-PI SSFSE		8	256x256	**544.72/89.41**
T2 SSFSE BH	Coronal	5	512x512	**2000/59.33**
T2 SSFSE BH + F	Coronal	5	512x512	**4000/63.94**
FIESTA	Coronal	5	512x512	**4.97/200**
FIESTA	Axial	6	512x512	**5.39/200**
FIESTA	Sagittal	6	512x512	**4.81/200**
FIESTA Dyn	Coronal	8	512x512	**3.10/200**
LAVA	Coronal	6	512x512	**4.96/7**
LAVA	Axial	6	512x512	**4.16/7**
LAVA + C	Coronal	6	512x512	**4.96/7**
LAVA + C	Axial	6	512x512	**4.16/7**
DWI MULT	Axial	6	256x256	**12000/60.50**
ADC	Axial			
Cal Body 48 AA	Axial	16	64x64	**1.43/0.50**
Cal TL Spine 36 4	Axial	16	64x64	**1.43/0.50**
MOD T2 FSE+FAT	Coronal	3	512x512	**5757/67.26**
Cal Body 24 AA2	Axial	15.8	64x64	**1.44/0.50**
MOD T1	Coronal	3	256x256	**460/9.34**
Cal Body 36 AA2	Axial	16	64x64	**1.43/0.50**
Cal Body 24 AA3	Axial	15.8	64x64	**1.44/0.50**
Cal Body 24 AA1	Axial	15.8	64x64	**1.44/0.50**

SSFSE, Single Shot Fast Spin Echo; BH, Breath Hold; FIESTA, Fast Imaging Employing Steady State Acquisition; LAVA, Liver Acquisition with Volume Acquisition; DWI, Diffusion

Weighted Imaging; ADC, Apparent Diffusion Coefficient; FOV, Field Of View; TR, repetition time; TE, Echo Time

STARD checklist for reporting of studies of diagnostic accuracy

(version January 2003)

Section and Topic	**Item #**		**On page #**
TITLE/ABSTRACT/ KEYWORDS	1	Identify the article as a study of diagnostic accuracy (recommend MeSH heading 'sensitivity and specificity').	1
INTRODUCTION	2	State the research questions or study aims, such as estimating diagnostic accuracy or comparing accuracy between tests or across participant groups.	3
METHODS			
Participants	3	The study population: The inclusion and exclusion criteria, setting and locations where data were collected.	6
	4	Participant recruitment: Was recruitment based on presenting symptoms, results from previous tests, or the fact that the participants had received the index tests or the reference standard?	6
	5	Participant sampling: Was the study population a consecutive series of participants defined by the selection criteria in item 3 and 4? If not, specify how participants were further selected.	6
	6	Data collection: Was data collection planned before the index test and reference standard were performed (prospective study) or after (retrospective study)?	6
Test methods	7	The reference standard and its rationale.	7
	8	Technical specifications of material and methods involved including how and when measurements were taken, and/or cite references for index tests and reference standard.	7-9
	9	Definition of and rationale for the units, cut-offs and/or categories of the results of the index tests and the reference standard.	7-9

	10	The number, training and expertise of the persons executing and reading the index tests and the reference standard.	7
	11	Whether or not the readers of the index tests and reference standard were blind (masked) to the results of the other test and describe any other clinical information available to the readers.	7
Statistical methods	12	Methods for calculating or comparing measures of diagnostic accuracy, and the statistical methods used to quantify uncertainty (e.g. 95% confidence intervals).	8
	13	Methods for calculating test reproducibility, if done.	
RESULTS			
Participants	14	When study was performed, including beginning and end dates of recruitment.	9
	15	Clinical and demographic characteristics of the study population (at least information on age, gender, spectrum of presenting symptoms).	9
	16	The number of participants satisfying the criteria for inclusion who did or did not undergo the index tests and/or the reference standard; describe why participants failed to undergo either test (a flow diagram is strongly recommended).	9
Test results	17	Time-interval between the index tests and the reference standard, and any treatment administered in between.	9
	18	Distribution of severity of disease (define criteria) in those with the target condition; other diagnoses in participants without the target condition.	9-10
	19	A cross tabulation of the results of the index tests (including indeterminate and missing results) by the results of the reference standard; for continuous results, the distribution of the test results by the results of the reference standard.	10-11
	20	Any adverse events from performing the index tests or the reference standard.	11
Estimates	21	Estimates of diagnostic accuracy and measures of statistical uncertainty (e.g. 95% confidence intervals).	11

	22	How indeterminate results, missing data and outliers of the index tests were handled.	11
	23	Estimates of variability of diagnostic accuracy between subgroups of participants, readers or centers, if done.	9-11
	24	Estimates of test reproducibility, if done.	
DISCUSSION	25	Discuss the clinical applicability of the study findings.	12-15