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## Systematic review and meta-analysis: Prognostic tests of paracetamol-induced acute liver failure

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**Systematic review and meta-analysis: Prognostic tests of  
paracetamol-induced acute liver failure**

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3 **Systematic review and meta-analysis: Prognostic tests of paracetamol-**  
4 **induced acute liver failure**  
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46 failure  
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**ABBREVIATIONS**

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8	AFP	Alpha-fetoprotein
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10	ALF	Acute liver failure
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12	ALT	Alanine aminotransferase
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14	APACHE II	Acute Physiology and Chronic Health Evaluation II
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16	AUC	Area under the curve
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18	DOR	Diagnostic odds ratio
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20	HE	Hepatic encephalopathy
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23	IL-6	Interleukin-6
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25	KCC	King's College Criteria
26		
27	KCH	King's College Hospital
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29		
30	MELD	Model for End-Stage Liver Disease
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32	OLT	Orthotopic liver transplantation
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34	ROC	Receiver operating characteristics
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36	SIRS	Systemic inflammatory response syndrome
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39	SOFA	Sequential Organ Failure Assessment
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## ABSTRACT

**Background:** Paracetamol toxicity remains the leading cause of acute liver failure (ALF) in the developed world. In the UK, the recently modified King's College Criteria (KCC) are used to list patients for emergency liver transplantation, but these criteria have been criticised for their low sensitivity and for spectrum bias in their application.

**Aim:** To critically evaluate existing prognostic criteria for predicting death without transplantation in paracetamol-induced ALF.

**Methods:** MEDLINE, EMBASE and CINAHL were searched to identify studies containing adult patients with paracetamol-induced ALF. Selected studies were evaluated, and data were pooled if appropriate, to calculate sensitivity, specificity, and diagnostic odds ratios (DORs) of applied prognostic tests.

**Results:** Of 6507 studies identified, 14 were eligible for inclusion, evaluating 1960 patients. The original KCC had a pooled sensitivity of 58.2% and specificity of 94.6%, with a DOR of 27.7. Addition of arterial lactate to the KCC reduced the DOR to 26.1. **Several other clinical and laboratory variables** had higher DORs than the KCC, but were only evaluated in single studies of limited quality.

**Conclusion:** **The original KCC remain well validated criteria with high prognostic accuracy.** Other potential prognostic variables should be prospectively assessed in multi-centre studies to further refine the criteria.

## INTRODUCTION

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Paracetamol (acetaminophen) toxicity remains the leading cause of acute liver failure (ALF) in the developed world, accounting for over 40% of cases in selected case series.(1,2) Whilst the vast majority of patients recover spontaneously following paracetamol overdose, a small number develop severe acute liver injury, hepatic encephalopathy (HE) and consequently, ALF. Despite significant advances in supportive care, the only effective treatment for the condition remains emergency orthotopic liver transplantation (OLT).(3)

The decision to transplant a patient with paracetamol-induced ALF involves balancing the inherent risks associated with delaying listing for OLT against the potential for spontaneous recovery with medical therapy alone, the risks of surgery in the context of a rapidly evolving critical illness, the scarcity of donor grafts, and the requirement for lifelong immunosuppression. Furthermore, the psychosocial implications of paracetamol overdose cases are considerable, with over 30% of patients that fulfil transplant criteria unsuitable for OLT due to severe psychological illness, or coexistent chronic alcohol or drug dependency.(4) Accurate prognostication in ALF is therefore vital to utilise OLT effectively and prevent unnecessary transplantation, whilst procuring donor organs in a timely fashion for those most likely to benefit.

In 1989 O'Grady and colleagues developed the 'King's College Criteria' (KCC) in an attempt to determine which patients with paracetamol and non-paracetamol-induced ALF have a poor prognosis with medical therapy alone, and will therefore benefit most from OLT.(5) The original KCC for paracetamol-induced ALF were highly specific, but have been criticised for their relatively low negative predictive

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3 value (6) and, since up to 26% of patients with paracetamol toxicity are medically  
4 unfit to undergo surgery at the point they fulfil the KCC,(4) modifications to these  
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6 criteria now utilise arterial lactate (>3.5 mmol/L after early fluid resuscitation or >3.0  
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8 mmol/L after adequate fluid resuscitation) in an attempt to extend the time-window  
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10 for acquisition of a suitable graft.(7)  
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16 Over recent years a plethora of alternative prognostic variables have been  
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18 proposed, in an attempt to improve, or replace, the KCC. These criteria variously  
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20 involve radiological,(8) histological,(9) serological,(10,11) or mathematical (12-14)  
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22 indices, but their assessment is complicated by the use of OLT; due to the inevitably  
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24 imperfect nature of current listing criteria a small proportion of ALF patients are  
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26 transplanted who would have spontaneously survived, invalidating further  
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28 assessment of that particular patient. Studies evaluating the prognostic accuracy of  
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30 a particular variable should therefore exclude transplanted patients from subsequent  
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32 analysis. The purpose of this systematic review was to critically evaluate existing  
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34 prognostic criteria for predicting death without transplantation in paracetamol-  
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36 induced ALF, to update previous studies in this area,(15,16) and to examine the  
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38 accuracy of recently described prognostic variables.  
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## METHODS

### Search Strategy and Study Selection

We conducted a systematic review of the medical literature using MEDLINE (1950 to June 2009), EMBASE (1980 to June 2009), and CINAHL (1982 to June 2009) to identify studies recruiting adult (>age 16 years) patients that evaluated prognostic markers of paracetamol-induced ALF. Potential studies were identified by combining the search terms “*acetaminophen*” and “*paracetamol*” (as both Medical Subject Headings (MeSH) and free text terms) using the set operator OR. Prognostic studies of ALF were identified, and combined using the set operator OR, by using the terms “*Kings adj3 College adj3 Criteria*”, “*Clichy\$*”, “*APACHE\$*”, “*lactate*”, “*Gc-globulin*”, “*alpha-fetoprotein*”, “*phosphate*” and “*MELD*” (all free text terms) and the McMaster expert search strategy for prognostic studies. These studies were combined using the set operator AND with papers evaluating studies on “*liver failure*”, “*hepatic encephalopathy*” or “*liver failure, acute*” (all MeSH). The search was limited to human studies without language restrictions. Abstracts of the studies identified by the initial search were evaluated for appropriateness to the study question by two independent reviewers (DC, KS) and all potentially relevant papers were obtained and evaluated in detail. The bibliographies of these studies and relevant review articles were screened to perform a recursive search of the literature, and abstract books of international liver conferences from the preceding 4 years were hand-searched for potentially relevant studies.

Selected studies were required to report mortality data on cohorts of patients admitted to hospital with acute severe liver injury or ALF secondary to paracetamol overdose (**see Box for study eligibility criteria**). Acute severe liver injury was defined as severe hepatotoxicity (serum alanine aminotransferase (ALT) levels >1000



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3 U/mL and co-existent coagulopathy), whilst ALF required the additional presence of  
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5 HE, in patients without pre-existing liver disease. Unselected cohorts of patients with  
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7 mixed etiologies of acute liver injury were eligible for inclusion if separate data for  
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9 paracetamol overdose patients were available. Case control studies and studies  
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11 comparing ALF patients with chronic liver disease patients were excluded due to  
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13 potential bias in favour of the prognostic variable in question,(17) as were studies  
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15 where transplantation and death were combined as a single end-point. Studies were  
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17 required to include more than 25 patients with paracetamol-induced acute liver  
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19 injury, since the majority of studies in this area include the KCC as a prognostic  
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21 variable in addition to the main variable studied, and smaller studies may invalidate  
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23 analysis.(18) All potential articles were assessed independently by two researchers  
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25 according to eligibility criteria, which were defined prospectively, and disagreements  
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27 resolved by consensus.  
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### 37 **Data extraction and quality assessment**

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40 Data were extracted using predesigned forms by two separate reviewers (DC  
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42 and KS) on to a Microsoft Excel spreadsheet (Microsoft Corp, Redmond, WA, USA),  
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44 and any discrepancies were resolved by consensus. The following data were  
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46 extracted for each study: setting, country and geographical region, year(s)  
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48 conducted, retrospective or prospective design, inclusion criteria, definitions of  
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50 paracetamol-induced liver injury, definitions of ALF, total number of subjects  
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52 included, total number of subjects with paracetamol-induced ALF or acute liver  
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54 injury, total number of subjects with paracetamol-induced ALF transplanted, duration  
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56 of follow-up, prognostic score used, timing of application of the prognostic score, and  
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58 outcome definitions. Study quality was assessed semi-quantitatively according to six  
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potential sources of bias that can be encountered in studies of prognostic variables (19) and a grade (poor, moderate, good or excellent) allocated (**Table 1**).

Study population	Study population adequately represents the population of interest; source population adequately described; sampling timeframe, place and period of recruitment described; inclusion and exclusion criteria described
Follow-up	Adequate study completion rate; loss to follow-up not associated with key characteristics; reasons for loss to follow-up described; no important differences between participants completing/not completing study
Prognostic factor	Prognostic measure clearly defined; continuous data reported and any cut-points described and appropriate; method and setting of measurement of prognostic factor identical for all patients; adequate proportion of study population has prognostic factor measured
Outcome measurement	Clear definition of outcome measure of interest described and recorded, including duration of follow-up; outcome measure valid; outcome measure identical
Confounding	All important confounders measured and accounted for at all stages of study design, performance and analysis
Analysis	Sufficient presentation of data to permit assessment of analytical analysis; appropriate model-building, study design etc.; no selective reporting of results

**Table 1** Quality assessment of included studies (adapted from Hayden 2006 (19))

### Data synthesis and analysis

Prognostic tests evaluating the outcomes of survival without transplantation or death without transplantation following paracetamol-induced liver injury were assessed individually. From each study cohort we extracted total number of subjects dying without OLT, total number of subjects surviving without OLT, and total

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3 numbers of subjects fulfilling or not fulfilling the prognostic variable in question, and  
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5 constructed 2x2 contingency tables. The sensitivity, specificity, and diagnostic odds  
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7 ratio (DOR) with their 95% confidence intervals (CI) were calculated for each test  
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9 using Microsoft Excel (Microsoft Corp, Redmond, Washington, USA) and checked  
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11 using Meta-DiSc version 1.4 (Universidad Complutense, Madrid, Spain). The value  
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13 of a DOR ranges from zero to infinity, with higher values indicating better  
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15 discriminatory test performance, and is calculated from the following formula:  
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17  $\text{sensitivity}/(1 - \text{sensitivity}) / (1 - \text{specificity})/\text{specificity}.$ (20) Where required, a zero cell  
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19 correction of 0.5 was added to all cells to prevent computational problems arising  
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21 where proportions were equal to zero.(21) Where two or more studies analysed a  
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23 particular prognostic marker, data were pooled using a random effects model and  
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25 the pooled DOR calculated. Heterogeneity between studies was assessed using the  
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27  $I^2$  statistic with a cut off of 50%, and the  $\chi^2$  test with a P value < 0.10, used to define  
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29 a statistically significant degree of heterogeneity. We explored study setting and  
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31 study design as potential reasons for heterogeneity.(22) These are exploratory  
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33 analyses only, and the results should therefore be interpreted with caution.  
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## 44 RESULTS

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47 The search strategy identified 6507 studies of which 105 were potentially  
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49 eligible for inclusion and were retrieved for further analysis (**Figure 2**). Of these, 14  
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51 were considered eligible for inclusion,(5,7,10,23-33) including two studies published  
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53 in abstract form only.(32,33) Characteristics of included studies are provided in  
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55 **Table 2**. Cohen's kappa test for inter-observer agreement was excellent at 0.86  
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60 (95% CI 0.75-0.98).

Study	Country	Liver Unit	Study period	Inclusion criteria	Prognostic test(s) evaluated	No paracetamol patients included	Retrospective/prospective cohort	Retrospective/prospective test development/evaluation	Study quality
O'Grady 1989(5)	UK	KCH	1973-1985; 1986-7	FHF as per Trey & Davidson(34) with Grade III-IV HE	pH<7.3 Concurrent PT>100 seconds, serum creatinine> 300 mmol/L, Grade III/IV HE	121 99	Both	Prospective	Moderate
O'Grady 1991(23)	UK	KCH	1988-1990	Severe liver damage	KCC as per O'Grady (5)	60	Prospective	Prospective	Moderate
Izumi 1996 (24)	UK	KCH	Not stated	FHF as per Trey & Davidson (34)	KCC as per O'Grady (5) Factor V ratio <20% Factor V ratio <10%	} 81	Prospective	Prospective (partially)	Poor
Anand 1997(25)	UK	Birmingham	1990-1994	FHF as per Trey & Davidson (34)	pH<7.3 Concurrent PT>100 seconds, serum creatinine> 300 mmol/L, Grade III/IV HE	72 89	Retrospective	Prospective	Poor
Bernal 1998(26)	UK	KCH	1990-1996	Severe hepatotoxicity (KCC); Death whilst not meeting KCC (APACHE III)	KCC as per O'Grady(5) Adapted APACHE III	504 56	Retrospective	Prospective (KCC) Retrospective (APACHE III)	Poor
Mitchell 1998(27)	UK	KCH	1993-1994	Coagulopathy + recent history paracetamol ingestion	KCC as per O'Grady(5) APACHE II>15 at 24 hr post-admission APACHE II>15	} 94	Prospective	Prospective (KCC) Retrospective (APACHE II)	Good
Bernal 2002(7)	UK	KCH	1998-1999 (learning set) 1999-2000 (validation set)	Severe paracetamol-induced hepatotoxicity	KCC as per O'Grady(5) Arterial lactate>3.5 mmol/L at 4 hours Arterial lactate>3.0 mmol/L at 12 hours	99 97 85	Both	Prospective	Moderate

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					KCC + combination of lactate criteria	85			
Bernal 2003(28)	UK	KCH	1998-2000	Acute severe hepatotoxicity	KCC as per O'Grady(5) Phosphate >1.2 mmol/L on admission day +1 or 2	} 170	Retrospective	Prospective	Moderate
Larson 2005(29)	USA	22 academic centres	1998-2003	INR>1.4; HE; jaundice to HE interval<26 weeks	KCC as per O'Grady(5)	252	Prospective	Prospective	Moderate
Schmidt 2005(10)	Denmark	Copenhagen	1999-2002	Peak ALT>1000U/L	KCC as per O'Grady(5) AFP <3.9 at D1 post peak ALT AFP <3.9 & INR>2.4 at D1 post peak ALT	234 188 188	Prospective	Prospective (KCC) Retrospective (AFP)	Moderate
Schmidt 2006(30)	Denmark	Copenhagen	1999-2004	Severe paracetamol-induced FHF	KCC as per O'Grady(5) Arterial lactate Modified KCC SOFA score >8 at admission/ >12 at onset of Grade III/IV HE SIRS at admission/at onset of Grade III/IV HE	95 91 91 95 95	Prospective	Retrospective	Moderate
Zaman 2006(31)	Ireland	Dublin	1994-2005	Paracetamol-induced ALF (jaundice to HE <8 weeks) or rapid bilirubin/ INR/ renal impairment/ hypoglycaemia if no HE	KCC as per O'Grady(5) MELD >30	} 60	Retrospective	Prospective	Poor
Bates 2007(32)	UK	Edinburgh	2004-2007	Acute severe liver injury	KCC as per O'Grady(5) Lactate modifications to KCC as per Bernal(7)	69 32	Retrospective	Prospective	Poor
Bernal 2007(33)	UK	KCH	Not stated	Acute liver failure	IL-6 within 24 hours of admission	31	Prospective	Retrospective	Poor

**Table 2** Characteristics of included studies

PT, prothrombin time; FHF, fulminant hepatic failure

For Peer Review

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3 The eligible studies evaluated a total of 1960 patients with paracetamol-  
4 induced acute liver injury or ALF. Three (10,24,27) studies had complete temporal  
5 overlap with other studies from the same unit; however, these studies were included  
6 for their evaluation of unique prognostic markers separate to the KCC. There was  
7 only one multicentre study.(29) Five (25,26,28,31,32) studies evaluated patient  
8 cohorts retrospectively, whilst five (10,26,27,30,33) studies developed prognostic  
9 test thresholds retrospectively (usually from receiver operator characteristics (ROC)  
10 curve analysis). No studies blinded observers to patient outcome or other potentially  
11 confounding prognostic data; and only two (5,7) validated their prognostic marker in  
12 a separate cohort. Consequently, six (24-26,31-33) of the 14 studies were graded as  
13 poor and seven (5,7,10,23,28-30) as moderate quality (**Table 2**). The 14 eligible  
14 studies analysed a total of 22 different prognostic markers or variations thereof. The  
15 sensitivity, specificity, and DORs of these are reported in **Table 3**.  
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Prognostic test	Study	N/deaths	Test +ve/ deaths	Sensitivity (95% CIs)	Specificity (95% CIs)	Diagnostic Odds Ratio (95% CIs)	Heterogeneity	
							$\chi^2$	$I^2$
pH<7.3	O'Grady 1989(5)	121/43	22/21	48.8 (33.3-65.5)	98.7 (93.1-100)	73.5 (9.4-577.4)	4.91 p=0.027	80%
	Anand 1997(25)	72/39	31/24	61.5 (44.6-76.6)	78.8 (61.1-91.0)	5.9 (2.1-17.1)		
	<b>POOLED</b>			<b>54.9 (43.5-65.9)</b>	<b>92.8 (86.3-96.8)</b>	<b>18.0 (1.1-229.6)</b>		
Concurrent PT>100s, serum creatinine>300 mmol/L, Grade III/IV HE	O'Grady 1989(5)	99/22	15/10	45.5 (24.4-67.8)	93.5 (85.5-97.9)	12.0 (3.5-41.3)	0.8 p=0.374	0%
	Anand 1997(25)	89/45	24/19	42.2 (27.7-57.9)	88.6 (75.4-96.2)	5.7 (1.9-17.2)		
	<b>POOLED</b>			<b>43.3 (31.2-56.0)</b>	<b>91.7 (85.3-96.0)</b>	<b>7.9 (3.5-18.1)</b>		
KCC (combined)†	O'Grady 1989(5)	220/65	37/31	47.7 (35.1-60.5)	96.1 (91.8-98.6)	22.6 (8.8-58.6)	8.4 p=0.038	64%
	O'Grady 1991(23)	60/26	23/19	73.1 (52.2-88.4)	88.2 (72.6-96.7)	20.4 (5.2-79.0)		
	Bernal 1998(26)	504/99	80/71	71.7 (61.8-80.3)	97.8 (95.8-99.0)	111.6 (50.5-246.4)		
	Bernal 2002(7)	99/21	20/16	76.2 (52.8-91.8)	94.9 (87.4-98.6)	59.2 (14.3-245.3)		
	<b>POOLED (KCH)</b>			<b>65.9 (59.0-72.3)</b>	<b>96.1 (94.4-97.5)</b>	<b>43.9 (17.6-109.3)</b>		
	Larson 2005(29)							
	(at admission)	252/74	34/19	25.7 (16.2-37.2)	91.6 (86.5-95.2)	3.8 (1.8-7.9)		
	Schmidt 2006(30)	95/48	36/27	77.1 (62.7-88.0)	83.0 (69.2-92.4)	16.4 (5.9-45.3)		
	Zaman 2006(31)	60/29	21/21	72.4 (52.8-87.3)	100.0 (88.8-100.0)	159.4 (8.7-2908.9)		
Bates 2007(32)	69/17	18/15	88.2 (63.6-98.5)	94.2 (84.1-98.8)	122.5 (18.7-803.1)			
<b>POOLED (non-</b>								

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	<b>KCH)</b>			<b>48.8 (41.0-56.6)</b>	<b>91.2 (87.5-94.1)</b>	<b>16.5 (3.5-77.8)</b>	17.0 p=0.001	82%
	<b>POOLED (overall)</b>			<b>58.2 (53.1-63.3)</b>	<b>94.6 (93.0-95.9)</b>	<b>27.7 (9.2-83.5)</b>	53.0 p<0.001	87%
	Bernal 2002(7)	97/21	18/14	66.7 (43.0-85.4)	94.7 (87.1-98.6)	36.0 (9.3-139.6)		
	Schmidt 2006(30)	91/46	61/39	84.8 (71.1-93.7)	51.1 (35.8-66.3)	5.8 (2.2-15.8)		
	Bates 2007(32)	69/17	37/15	88.2 (63.6-98.5)	57.7 (43.2-71.3)	10.2 (2.1-49.4)		
	<b>POOLED</b>			<b>81.0 (70.9-88.7)</b>	<b>72.3 (64.9-78.8)</b>	<b>12.2 (4.0-37.4)</b>	4.54 p=0.103	56%
	Bernal 2002(7)	85/21	18/16	76.2 (52.8-91.8)	96.9 (89.2-99.6)	99.2 (17.6-559.3)		
	Schmidt 2006*(30)	91/46	55/36	78.3 (63.6-89.1)	57.8 (42.2-72.3)	4.9 (2.0-12.3)		
	Bates 2007(32)	32/12	20/12	100.0 (73.5-100.0)	60.0 (36.1-80.9)	36.8 (1.9-708.0)		
	*at onset of Grade III/IV HE						9.83 p=0.007	80%
	<b>POOLED</b>			<b>81.0 (70.6-89.0)</b>	<b>77.5 (69.3-84.4)</b>	<b>22.8 (2.5-210.0)</b>		
	Arterial lactate >4.0 at admission		51/34	73.9 (58.9-85.7)	62.2 (46.5-76.2)	4.7 (1.9-11.4)		
	Arterial lactate >4.0 at onset of Grade III/IV HE	91/46	40/31	67.4 (52.0-80.5)	80.0 (65.4-90.4)	8.3 (3.2-21.5)	-	-
	Bernal 2002(7)	85/21	24/19	90.5 (69.6-98.8)	92.2 (82.7-97.4)	112.1 (20.1-625.7)		
	Schmidt 2006(30)	91/46	69/42	91.3 (79.2-97.6)	40.0 (25.7-55.7)	7.0 (2.1-22.9)		
	<b>POOLED</b>			<b>91.0 (81.5-96.6)</b>	<b>70.6 (61.2-79.0)</b>	<b>26.1 (1.7-393.7)</b>	8.83 p=0.003	89%

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Factor V ratio <20%	Izumi 1996(24)	81/35	69/34	97.1 (85.1-99.9)	23.9 (12.6-38.8)	10.7 (1.3-87.3)	-	-
Factor V ratio <10%			51/29	63.0 (47.6-76.8)	37.1 (21.5-55.1)	5.3 (1.8-15.1)	-	-
Adapted APACHE III	Bernal 1998(26)	56/28	19/16	57.1 (37.2-75.5)	89.3 (71.8-97.7)	11.1 (2.7-45.6)	-	-
APACHE II>15 at 24 hr post- admission	Mitchell 1998(27)	94/14	13/11	78.6 (49.2-95.3)	97.5 (91.3-99.7)	143.0 (21.4 -953.5)	-	-
APACHE II>15 within 5 days of admission			23/13	92.9 (66.1-99.8)	87.5 (78.2-93.8)	91.0 (10.7-772.8)	-	-
Phosphate >1.2 mmol/L on admission day +1 or 2	Bernal 2003(28)	170/52	55/42	80.8 (67.5-90.4)	89.0 (81.9-94.0)	33.9 (13.8-83.3)	-	-
AFP <3.9 at D1 post peak ALT	Schmidt 2005(10)	188/33	54/33	100.0 (89.4-100.0)	73.6 (65.9-80.3)	184.9 (11.1-3085.6)	-	-
AFP <3.9 & INR>2.4 at D1 post peak ALT			74/33	100.0 (89.4-100.0)	86.5 (80.0-91.4)	419.1 (24.8-7097.5)	-	-
SOFA score >8 at admission	Schmidt 2006(30)	95/48	48/32	66.7 (51.6-79.6)	66.0 (50.7-79.1)	3.9 (1.7-9.1)	-	-
SOFA score >12 at onset of Grade III/IV HE			54/39	81.3 (67.4-91.1)	68.1 (52.9-80.9)	9.2 (3.6-23.9)	-	-

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SIRS at admission	Schmidt 2006(30)	95/48	52/33	68.8 (53.7-81.3)	59.6 (44.3-73.6)	3.2 (1.4-7.5)	-	-
SIRS at onset of Grade III/IV HE			41/34	70.8 (55.9-83.0)	85.1 (71.7-93.8)	13.9 (5.0-38.3)		
MELD >30	Zaman 2006(31)	60/29	38/28	96.6 (82.2-99.9)	67.7 (48.6-83.3)	58.8 (7.0-495.8)	-	-
IL-6 within 24 hours of admission	Bernal 2007(33)	31/8	7/6	75.0 (34.9-96.8)	95.7 (78.1-99.9)	66.0 (5.1-857.7)	-	-

**Table 3** Sensitivity, specificity, and diagnostic odds ratios of individual prognostic markers.

† Anand 1997 (25) excluded as 2x2 table not reconstructable; Mitchell 1998,(27) Bernal 2003,(28) and Schmidt 2005(10) excluded due to complete temporal overlap with previous studies from same unit

## King's College Criteria

A total of 13 studies evaluated the original KCC for paracetamol-induced ALF, either as separate elements (arterial pH<7.3 or concurrent grade III/IV HE, serum creatinine>300µmol/L and prothrombin time (PT)>100 seconds) (5,25) or as a whole, (7,10,23,24,26-32) including a total of 1929 patients. One study (29) evaluated the KCC only on admission. After exclusion of studies with complete temporal overlap with other studies,(10,24,27) pooled specificity of the KCC was high, at 94.6% (95% CI 93.0-95.9), but the pooled sensitivity was relatively poor at 58.2% (95% CI 53.1-63.3). The pooled DOR for the KCC was 27.7 (95% CI 9.2-83.5) (**Table 3**). The summary ROC curve is shown in **Figure 3**. The area under the curve (AUC) was calculated as 0.91 (95% CI 0.79-0.99), suggesting good performance of the KCC overall.(35) The accuracy of the KCC improved when studies originating from KCH were analysed separately from those outside KCH (DOR 43.9 (95% CI 17.6-109.3) for KCH-based studies vs. DOR 16.5 (95% CI 3.5-77.8) for non-KCH-based studies). Statistically significant heterogeneity ( $I^2=87\%$ ) existed between individual study results, though this fell to 79% after exclusion of the single multi-centre study,(29) and to 64% after exclusion of studies from outside KCH.

## Lactate modifications to the KCC

The original study (7) that evaluated arterial lactate in the prognosis of paracetamol-induced ALF reported similar specificity, but improved sensitivity, when compared with the original KCC. However, two (30,32) subsequent studies evaluating arterial lactate alone failed to replicate this high prognostic accuracy (**Table 3**), with pooled DORs of 12.2 (95% CI 4.0-37.4) and 22.8 (95% CI 2.5-210.0)

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3 for the early and post-resuscitation lactate values respectively. This was mainly due  
4 to the reduced specificity seen in the two studies from outside KCH. Combination of  
5 the KCC with a post-resuscitation lactate value  $>3.0$  offered higher prognostic  
6 accuracy in the original lactate study (7) but this was not replicated in a subsequent  
7 evaluation.(30)  
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### 19 **Other prognostic markers**

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22 Several other markers, all evaluated in single studies, appeared to offer  
23 improved prognostic accuracy when compared with the KCC, with low serum alpha-  
24 fetoprotein (AFP) (DOR 419.1),(10) 24-hour post-admission Acute Physiology and  
25 Chronic Health Evaluation II (APACHE II) score (DOR 143.0),(27) serum interleukin-6  
26 (IL-6) levels (DOR 66.0),(33) Model for End-Stage Liver Disease (MELD) score  
27 (DOR 58.8),(31) and serum phosphate (DOR 33.9) (28) all outperforming the KCC  
28 (**Table 3**). However, further studies would be required to replicate these findings to  
29 confirm or refute whether these other markers are indeed superior to the KCC.  
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### 45 **DISCUSSION**

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48 This systematic review and meta-analysis has demonstrated that the original  
49 KCC for paracetamol-induced ALF have high pooled specificity (94.6%), but low  
50 pooled sensitivity (58.2%) in determining prognosis in patients with paracetamol-  
51 induced ALF. Additionally, the benefit of the arterial lactate modifications to the KCC  
52 (7) is questionable according to these data. Other proposed prognostic markers, in  
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3 particular AFP, APACHE II scores, and serum IL-6 levels, showed encouraging  
4 prognostic accuracy but were only evaluated in single studies of variable quality.  
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9 This review is limited by the quality of the included studies, which had  
10 significant heterogeneity and were generally of poor or moderate quality, with only  
11 two (5,7) studies validating their prognostic model prospectively in a separate cohort.  
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13 Acute liver failure is a rare syndrome and, as a result, many studies were small and  
14 retrospective in nature. Continuous variables were frequently analysed  
15 retrospectively using cut-off values designed to maximise AUC values, a method  
16 which assumes constant risk amongst the 'high' and 'low' risk groups.(18) Several  
17 studies attempted to evaluate multiple prognostic markers retrospectively using the  
18 same cohort, an approach which increases the risk of obtaining a statistically  
19 significant result by chance. Another potentially confounding issue is the lack of  
20 international standardisation of laboratory variables, such as prothrombin time and  
21 creatinine, and changes to assay reagents over the time course of these studies.(36)  
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23 We also appreciate that exclusion of studies where transplantation and death were  
24 deemed equivalent may have introduced spectrum bias, since prognostic tests are  
25 usually applied in settings where OLT is available. This problem is difficult to  
26 circumvent given that emergency OLT will never be subjected to a clinical trial, but,  
27 given the reduced quality-of-life seen following transplantation for paracetamol-  
28 induced ALF, the accumulated risks of immunosuppression, and the scarcity of liver  
29 donors,(16) accurate calculation of the specificity of each prognostic test is vital in  
30 order to minimise inappropriate transplantation.  
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57 This study expands upon a recent systematic review (16) evaluating the KCC  
58 for paracetamol-induced ALF by including additional prognostic markers. The former  
59 study found spectrum bias in studies originating from KCH, with increased  
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3 spontaneous survival amongst patients listed for transplantation, but not receiving a  
4 graft, compared with those meeting criteria but never listed, suggesting that a  
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6 'healthier' cohort may be preferentially transplanted in KCH. The current data confirm  
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8 the increased prognostic accuracy of the original KCC in studies originating from  
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10 KCH compared with studies from other units, with a higher pooled DOR in KCH-  
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12 based studies compared with that from studies conducted in other units. This raises  
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14 further questions regarding the overall generalisability of the KCC to patients with  
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16 paracetamol-induced ALF treated outside KCH.  
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23 The heterogeneity of the included studies evaluating the KCC was partly due  
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25 to a multicentre US study (29) which explicitly applied the KCC solely at the time of  
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27 admission, rather than dynamically throughout admission as originally  
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29 intended.(5,37) Whilst early and accurate prognostication in ALF is vital in order to  
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31 permit timely listing for OLT, application of the KCC solely at admission reduces the  
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33 time available for the disease to evolve and may introduce spectrum bias. Given that  
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35 the median time taken to fulfil the KCC following admission to a tertiary liver  
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37 transplant unit is 12 hours,(7) application of the KCC solely at admission may explain  
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39 the low sensitivity of the criteria in the US study, whilst concerns have also been  
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41 expressed about the prophylactic use of fresh frozen plasma in this US study and the  
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43 potential confounding effects upon prothrombin time.(38) Furthermore, 48% of cases  
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45 resulted from unintentional overdoses in this US study, a type of overdose seen less  
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47 frequently in the UK.(39) It may be that repeated ingestion of supratherapeutic  
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49 doses of paracetamol over a protracted time course disrupts liver function  
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51 subacutely, so that patients are less likely to develop the profound acidosis or  
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53 concurrent severe coagulopathy, renal dysfunction, and encephalopathy required to  
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55 fulfil the KCC, but have at least as poor a prognosis as an intentional overdose at a  
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3 single time point. Future prognostic scoring systems may therefore need to take the  
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5 pattern of paracetamol overdose into account in addition to traditional biochemical  
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7 parameters.  
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11 The addition of post-resuscitation arterial lactate to the KCC (7) is questioned  
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13 by this study. Only two additional (30,32) studies, one of which was reported in  
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15 abstract-form, that reported this modification fulfilled eligibility criteria for inclusion,  
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17 but these studies suggest little benefit from this. Furthermore, the reduced specificity  
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19 of the lactate criteria undermines the traditional benefit of the KCC in 'ruling in' a  
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21 hopeless prognosis. The presence of systemic inflammation, with or without sepsis,  
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23 is increasingly recognised as important in ALF,(40) but hyperlactataemia can result  
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25 from numerous other organ sources and therefore, perhaps not surprisingly, arterial  
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27 lactate is a relatively non-specific prognostic indicator in paracetamol-induced ALF.  
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29 Within critical care settings, the use of APACHE II and Sequential Organ Failure  
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31 Assessment (SOFA) scores to monitor organ dysfunction have greater recognition  
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33 than the KCC, and are attractive as early prognostic markers in ALF.(41) APACHE II,  
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35 in particular, showed encouraging prognostic accuracy but was only evaluated in a  
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37 single eligible study.(27) One (29) additional study evaluating APACHE II scores was  
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39 excluded from this particular analysis as the prognostic scoring reported in the study  
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41 could not be reconstructed into a 2x2 table; this study is also notable for the atypical  
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43 nature of the patient cohort as outlined above. Given that APACHE III, systemic  
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45 inflammatory response syndrome (SIRS) and SOFA scores all performed less well  
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47 than the KCC (albeit in single studies), the role of these markers may be to permit  
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49 earlier identification of a high-risk cohort requiring transfer to tertiary centres that  
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51 offer liver transplantation, rather than as definitive transplant listing criteria.  
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4 Persistently elevated serum phosphate or reduced AFP levels may reflect  
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6 failure of hepatic regeneration, and therefore could help predict a poorer outcome in  
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8 paracetamol-induced ALF. Serum AFP showed high prognostic performance in one  
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10 (10) study, whilst serum phosphate showed equivalence with the KCC in one (28)  
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12 retrospective study from KCH. Other authors, (in studies where transplantation was  
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14 equated with death and hence excluded from this study), have demonstrated  
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16 conflicting results with serum phosphate,(42-45) so further evaluation of both serum  
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18 AFP and phosphate in future studies would be worthwhile. The MELD scoring  
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20 system has been widely adopted for organ allocation in chronic liver disease and has  
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22 shown encouraging prognostic accuracy in non-paracetamol ALF.(12,46-48) More  
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24 limited data exist regarding the use of MELD in paracetamol-induced ALF, but one  
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26 additional study (excluded from this analysis as the 2x2 table was not  
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28 reconstructable) suggested that its use may be limited by a high false positive  
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30 rate.(49)  
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38 In summary, this systematic review and meta-analysis has demonstrated that  
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40 the original KCC for paracetamol-induced ALF have high pooled specificity, but low  
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42 pooled sensitivity in determining prognosis in patients with paracetamol-induced  
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44 ALF. The KCC had reduced prognostic accuracy when applied outside of KCH and  
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46 were occasionally applied only at admission. The reduced specificity of the KCC  
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48 following the addition of arterial lactate calls into question the benefit of this  
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50 modification, suggesting that re-evaluation of this as a prognostic marker is required.  
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52 Urgent consideration should be given to the design of a high-quality, prospective  
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54 study evaluating the KCC, APACHE II scores, and markers of hepatic regeneration,  
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56 such as serum AFP and phosphate, in paracetamol-induced ALF. Given the  
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58 relatively rare nature of ALF, such a study is likely to require cooperation between  
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3 several large centres, and argues the need for a collaborative network of tertiary  
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5 hospitals experienced in the management and prognostication of ALF, similar to that  
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7 developed by the Acute Respiratory Distress Syndrome Network program.(50)  
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7 **Figure legends**  
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10 **Figure 1** Eligibility criteria for included studies  
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13 **Figure 2** Flow diagram of assessment of studies identified in the systematic  
14 review  
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18 **Figure 3** Summary receiver operator curve (SROC) of studies evaluating the  
19 original KCC  
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24 AUC: Area under the curve; SE: Standard Error; Q: heterogeneity  
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# PRISMA Checklist

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Section/topic	#	Checklist item	Reported on page #
<b>TITLE</b>			
Title	1	Identify the report as a systematic review, meta-analysis, or both.	1
<b>ABSTRACT</b>			
Structured summary	2	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number.	3
<b>INTRODUCTION</b>			
Rationale	3	Describe the rationale for the review in the context of what is already known.	4-5
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	5,6
<b>METHODS</b>			
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	Figure 1
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	6
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	6
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	6-7
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	7-9

Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	7-9
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	8
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	12, Table 3
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., $I^2$ ) for each meta-analysis.	12, Table 3
Section/topic	#	Checklist item	Reported on page #
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	19
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	17
<b>RESULTS</b>			
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram.	Figure 2
Study characteristics	18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	Table 2
Risk of bias within studies	19	Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12).	Table 2
Results of individual studies	20	For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot.	Table 3
Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	Table 3
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies (see Item 15).	
Additional analysis	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	Table 3
<b>DISCUSSION</b>			

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Summary of evidence	24	Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers).	18
Limitations	25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	19
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	18-21
<b>FUNDING</b>			
Funding	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.	1

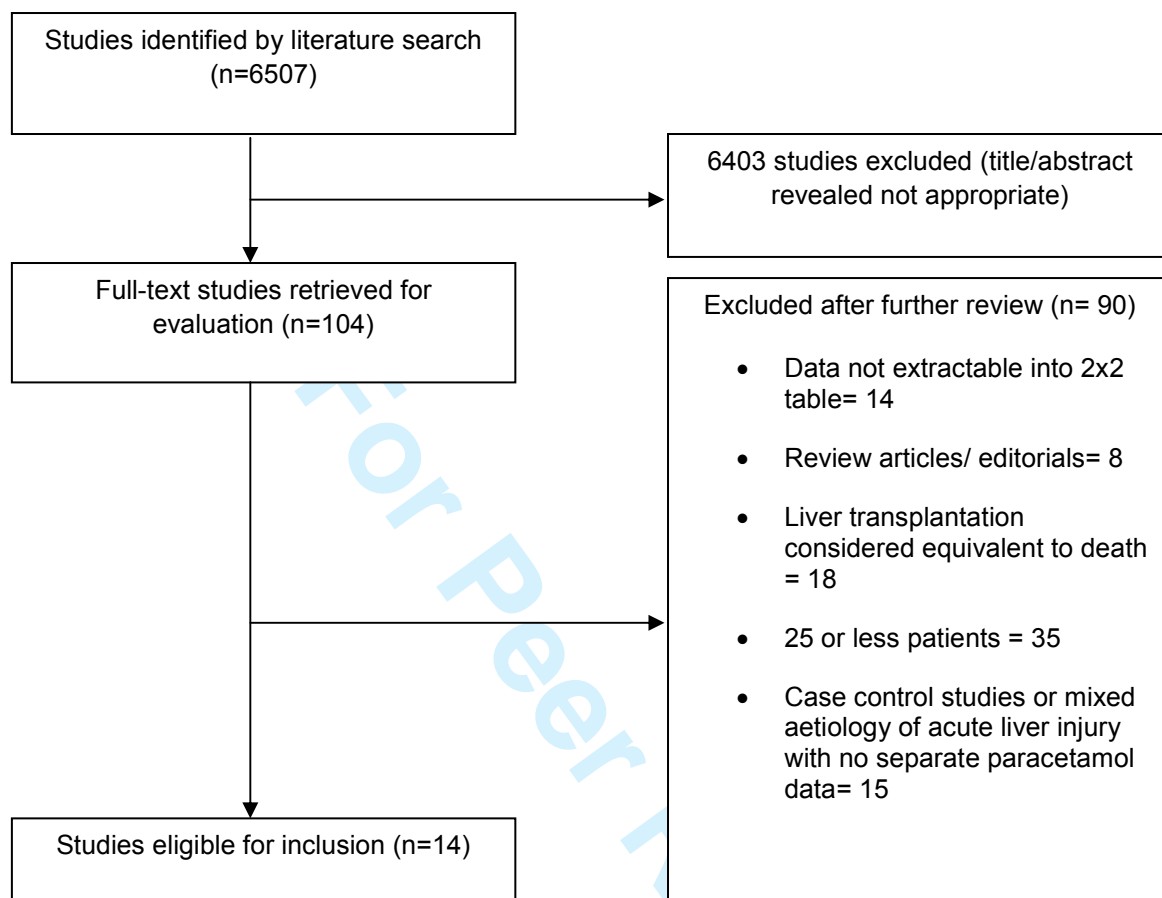
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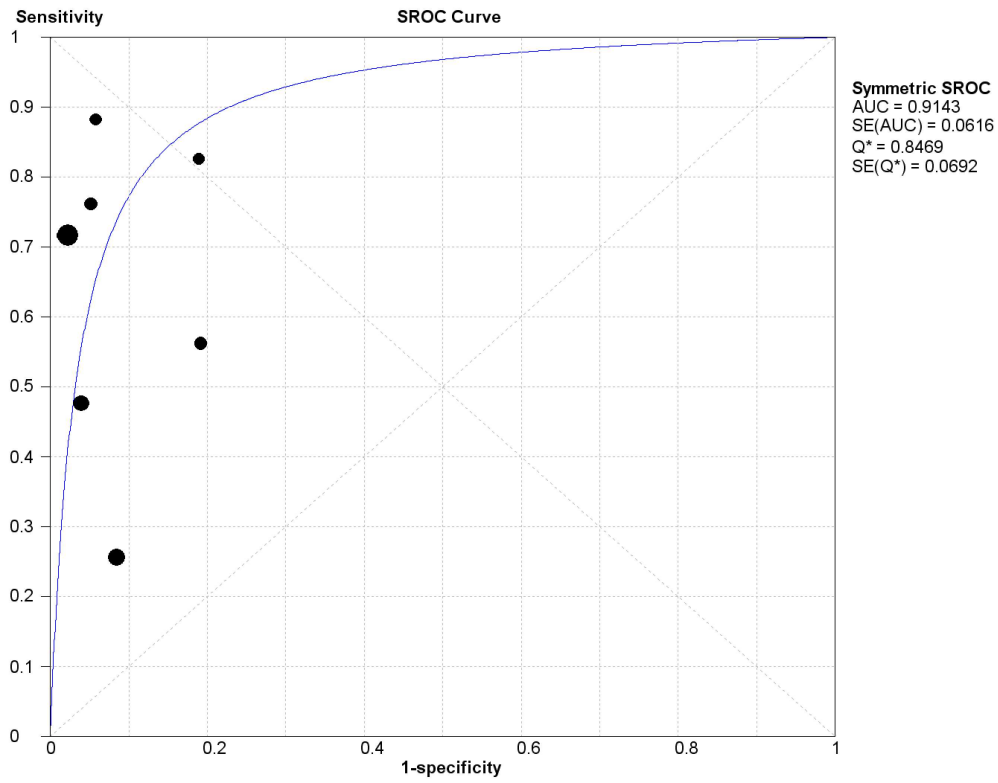
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- Adult patients (aged > 16 years) presenting with acute severe liver injury or ALF attributed to paracetamol hepatotoxicity
- Cohort studies
- Included >25 patients with paracetamol hepatotoxicity
- Mortality data recorded
- Data for death and liver transplantation considered separately

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Figure 3 Summary receiver operator curve (SROC) of studies evaluating the original KCC  
AUC: Area under the curve; SE: Standard Error; Q: heterogeneity

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view