

Supplementary Table S1. Search strategy PubMed and EMBASE

PubMed full search

("nephroblastomatosis"[Title/Abstract] OR "nephrogenic rest*"[Title/Abstract])

AND

("molecular biology"[MeSH Terms] OR "molecular" [Title/Abstract] OR "biochemical"[Title/Abstract]
OR "biomarker*"[Title/Abstract] OR "mutation*"[Title/Abstract])

Last performed: 25th of January 2022

Total of 79 results

EMBASE full search

'nephroblastomatosis'/exp OR 'nephroblastomatosis' OR 'nephroblastomatosis':ti,ab,kw OR

'nephrogenic rest*':ti,ab,kw

AND

'biochemistry'/exp OR 'biochemistry':ti,ab,kw OR 'mutation'/exp OR 'mutation':ti,ab,kw OR 'molecular
biology'/exp OR 'molecular biology':ti,ab,kw OR 'molecular':ti,ab,kw OR 'biomarker'/exp OR

'biomarker':ti,ab,kw

Last performed: 25th of January 2022

Total of 124 results (without MEDLINE)

Supplementary Table S2a. “Standards for Reporting Diagnostic Accuracy 2015” (STARD 2015) checklist

Section & Topic	No	Item	Reported on page #
TITLE OR ABSTRACT			
	1	Identification as a study of diagnostic accuracy using at least one measure of accuracy (such as sensitivity, specificity, predictive values, or AUC)	
ABSTRACT			
	2	Structured summary of study design, methods, results, and conclusions (for specific guidance, see STARD for Abstracts)	
INTRODUCTION			
	3	Scientific and clinical background, including the intended use and clinical role of the index test	
	4	Study objectives and hypotheses	
METHODS			
<i>Study design</i>	5	Whether data collection was planned before the index test and reference standard were performed (prospective study) or after (retrospective study)	
<i>Participants</i>	6	Eligibility criteria	
	7	On what basis potentially eligible participants were identified (such as symptoms, results from previous tests, inclusion in registry)	
	8	Where and when potentially eligible participants were identified (setting, location and dates)	
	9	Whether participants formed a consecutive, random or convenience series	
<i>Test methods</i>	10a	Index test, in sufficient detail to allow replication	
	10b	Reference standard, in sufficient detail to allow replication	
	11	Rationale for choosing the reference standard (if alternatives exist)	
	12a	Definition of and rationale for test positivity cut-offs or result categories of the index test, distinguishing pre-specified from exploratory	
	12b	Definition of and rationale for test positivity cut-offs or result categories of the reference standard, distinguishing pre-specified from exploratory	
	13a	Whether clinical information and reference standard results were available to the performers/readers of the index test	
	13b	Whether clinical information and index test results were available to the assessors of the reference standard	
<i>Analysis</i>	14	Methods for estimating or comparing measures of diagnostic accuracy	
	15	How indeterminate index test or reference standard results were handled	
	16	How missing data on the index test and reference standard were handled	
	17	Any analyses of variability in diagnostic accuracy, distinguishing pre-specified from exploratory	
	18	Intended sample size and how it was determined	
RESULTS			
<i>Participants</i>	19	Flow of participants, using a diagram	
	20	Baseline demographic and clinical characteristics of participants	
	21a	Distribution of severity of disease in those with the target condition	
	21b	Distribution of alternative diagnoses in those without the target condition	

	22	Time interval and any clinical interventions between index test and reference standard	
<i>Test results</i>	23	Cross tabulation of the index test results (or their distribution) by the results of the reference standard	
	24	Estimates of diagnostic accuracy and their precision (such as 95% confidence intervals)	
	25	Any adverse events from performing the index test or the reference standard	
DISCUSSION			
	26	Study limitations, including sources of potential bias, statistical uncertainty, and generalisability	
	27	Implications for practice, including the intended use and clinical role of the index test	
OTHER INFORMATION			
	28	Registration number and name of registry	
	29	Where the full study protocol can be accessed	
	30	Sources of funding and other support; role of funders	

Supplementary Table S2b. Case report guidelines (CARE checklist 2013)



CARE Checklist of information to include when writing a case report



Topic	Item	Checklist item description	Reported on Line
Title	1	The diagnosis or intervention of primary focus followed by the words "case report"	_____
Key Words	2	2 to 5 key words that identify diagnoses or interventions in this case report, including "case report"	_____
Abstract (no references)	3a	Introduction: What is unique about this case and what does it add to the scientific literature?	_____
	3b	Main symptoms and/or important clinical findings	_____
	3c	The main diagnoses, therapeutic interventions, and outcomes	_____
	3d	Conclusion—What is the main "take-away" lesson(s) from this case?	_____
Introduction	4	One or two paragraphs summarizing why this case is unique (may include references)	_____
Patient Information	5a	De-identified patient specific information	_____
	5b	Primary concerns and symptoms of the patient	_____
	5c	Medical, family, and psycho-social history including relevant genetic information	_____
	5d	Relevant past interventions with outcomes	_____
Clinical Findings	6	Describe significant physical examination (PE) and important clinical findings	_____
Timeline	7	Historical and current information from this episode of care organized as a timeline	_____
Diagnostic Assessment	8a	Diagnostic testing (such as PE, laboratory testing, imaging, surveys)	_____
	8b	Diagnostic challenges (such as access to testing, financial, or cultural)	_____
	8c	Diagnosis (including other diagnoses considered)	_____
	8d	Prognosis (such as staging in oncology) where applicable	_____
Therapeutic Intervention	9a	Types of therapeutic intervention (such as pharmacologic, surgical, preventive, self-care)	_____
	9b	Administration of therapeutic intervention (such as dosage, strength, duration)	_____
	9c	Changes in therapeutic intervention (with rationale)	_____
Follow-up and Outcomes	10a	Clinician and patient-assessed outcomes (if available)	_____
	10b	Important follow-up diagnostic and other test results	_____
	10c	Intervention adherence and tolerability (How was this assessed?)	_____
	10d	Adverse and unanticipated events	_____
Discussion	11a	A scientific discussion of the strengths AND limitations associated with this case report	_____
	11b	Discussion of the relevant medical literature with references	_____
	11c	The scientific rationale for any conclusions (including assessment of possible causes)	_____
	11d	The primary "take-away" lessons of this case report (without references) in a one paragraph conclusion	_____
Patient Perspective	12	The patient should share their perspective in one to two paragraphs on the treatment(s) they received	_____
Informed Consent	13	Did the patient give informed consent? Please provide if requested	Yes <input type="checkbox"/> No <input type="checkbox"/>

Supplementary Table S3a. Quality assessment of 18 out of 23 included articles based on the STARD 2015 checklist

STARD-item	1	2	3	4	5	6	7	8	9	10a	10b	11	12a	12b	13a	13b	14	15	16	17	18	19	20	21a	21b	22	23	24	25	26	27	28	29	30
Coorens et al. 2019	N	N	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	N	N	N	N	N	N	N	N	N	N	Y	N	Y	Y	N	Y	Y
Wegert et al. 2018	N	N	Y	Y	Y	Y	Y	U	U	Y	Y	U	Y	N	U	U	Y	N	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	Y	Y
Charlton et al. 2015	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	Y	N	U	U	Y	Y	Y	N	N	N	N	N	N	N	N	N	Y	Y	N	Y	Y	
Grill et al. 2010	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	N	Y	
Fukuzawa et al. 2010	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	Y	
Vuononvirta et al. 2008	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	Y	N	U	U	Y	Y	Y	N	N	N	N	N	N	N	N	Y	N	Y	Y	N	N	Y
Brown et al. 2008	N	N	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	N	N	N	N	N	Y	N	Y	Y	N	N	Y
Chilukamarri et al. 2007	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	N	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	Y	
Hancock et al. 2007	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	Y	Y	
Fukuzawa et al. 2006	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	N	Y	
Ravenel et al. 2001	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	N	N	N	N	N	N	N	N	N	N	Y	N	Y	Y	N	N	Y
Powlesland et al. 1999	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	Y	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	Y	
Charles et al. 1998	N	Y	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	Y	Y	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	N	
Cui et al. 1997	N	N	Y	Y	Y	Y	Y	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	Y	
Steenman et al. 1997	N	Y	Y	Y	Y	N	N	U	U	Y	N	N	Y	N	U	U	N	N	N	N	N	N	Y	N	N	N	N	N	Y	Y	N	N	Y	
Austruy et al. 1995	N	Y	Y	Y	Y	Y	N	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	N	Y	
Yun et al. 1993	N	Y	Y	Y	Y	N	N	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	Y	N	N	N	N	Y	Y	N	N	Y		
Pritchard-Jones et al. 1991	N	Y	Y	Y	Y	Y	N	U	U	Y	N	N	N	N	U	U	N	N	N	N	N	N	N	N	N	N	N	Y	Y	N	N	N		

Y = Present, N = Absent, U = Unclear

Supplementary Table S3b. Quality assessment of 5 out of 23 included articles based on the CARE 2013 checklist

CARE-item	1	2	3a	3b	3c	3d	4	5a	5b	5c	5d	6	7	8a	8b	8c	8d	9a	9b	9c	10a	10b	10c	10d	11a	11b	11c	11d	12	13
Chang et al. 2021	N	N	Y	Y	Y	Y	Y	Y	Y	Y	N	Y	N	Y	N	Y	Y	N	N	N	N	Y	N	N	Y	Y	Y	Y	N	U
Slack et al. 2021	N	N	Y	Y	Y	Y	Y	Y	Y	Y	Y	Y	N	Y	N	Y	Y	N	N	N	N	Y	N	N	Y	Y	Y	Y	N	U
MdZin et al. 2011	N	N	Y	Y	Y	Y	Y	Y	Y	N	Y	N	N	Y	N	Y	Y	Y	N	Y	N	Y	N	N	Y	Y	Y	Y	N	U
Hoban et al. 1995	N	N	Y	Y	Y	N	N	Y	Y	N	N	N	N	Y	N	Y	Y	N	N	N	N	Y	N	N	Y	Y	Y	Y	N	U
Park et al. 1993	N	N	Y	Y	Y	N	N	Y	Y	Y	Y	N	N	Y	N	Y	Y	Y	N	N	N	N	N	N	Y	Y	Y	Y	N	U

Y = Present, N = Absent, U = Unclear