

ACUTE RHEUMATIC FEVER and RHEUMATIC HEART DISEASE NOTIFICATION

DEFINITE and PROBABLE ARF (including recurrences) and RHD ARE NOTIFIABLE CONDITIONS IN THE NT
(Use this form also to notify possible ARF and borderline RHD cases)

RHD AND NTNDS - OFFICIAL USE ONLY:	
Notification date - date notified by Clinician:	Bed-list flag (DNSOT) and CWS alert (OTHAL) required: Yes <input type="checkbox"/> No <input type="checkbox"/> DNSOT required for: RDH <input type="checkbox"/> PRH <input type="checkbox"/> GDH <input type="checkbox"/> KH <input type="checkbox"/> TCH <input type="checkbox"/> ASH <input type="checkbox"/>
Notification received date - date received by NTNDS:	DSNOT and OTHAL flags added to CARESYS: Yes <input type="checkbox"/> No <input type="checkbox"/> DSNOT completed for: RDH <input type="checkbox"/> PRH <input type="checkbox"/> GDH <input type="checkbox"/> KH <input type="checkbox"/> TCH <input type="checkbox"/> ASH <input type="checkbox"/>

Notification Type
Notification Type: ARF only <input type="checkbox"/> RHD only <input type="checkbox"/> ARF and RHD <input type="checkbox"/>

Biographical Information			
HRN:	First Name:	Surname:	
Consent Obtained: Yes <input type="checkbox"/> No <input type="checkbox"/>	Date of birth:	Gender: Female <input type="checkbox"/> Male <input type="checkbox"/> Not Stated <input type="checkbox"/>	
Ethnicity: Aboriginal only <input type="checkbox"/> Torres Strait Islander only <input type="checkbox"/> Aboriginal and TSI <input type="checkbox"/> Non Aboriginal or TSI (other) <input type="checkbox"/> Unknown <input type="checkbox"/>			
Address:		Primary Clinic:	
Primary clinic notified: <input type="checkbox"/>	Date:	Secondary clinic:	

Diagnosis Information		
DATE DIAGNOSED (ARF):	DIAGNOSIS: Definite ARF (meets criteria) <input type="checkbox"/> Probable ARF <input type="checkbox"/> Possible ARF <input type="checkbox"/>	
DATE DIAGNOSED (RHD):	RHD <input type="checkbox"/> Borderline RHD (≤ 20 years of age) <input type="checkbox"/> Resolved RHD <input type="checkbox"/>	
ARF STATUS (ARF only): First known episode <input type="checkbox"/> Recurrence <input type="checkbox"/> Unknown <input type="checkbox"/>	LIKELY ONSET DATE: (ARF only)	
Clinic at onset (ARF):		
Clinic at presentation (onset) RHD (if different):		
Presentation: Clinical presentation - opportunistic <input type="checkbox"/> Clinical presentation - symptomatic <input type="checkbox"/> Screening <input type="checkbox"/> Other _____		
Hospital Admission: <input type="checkbox"/>	Echo performed: Yes <input type="checkbox"/> No <input type="checkbox"/>	Echo date:
Echo Location:		
Valve Severity at diagnosis: ARF only/no RHD <input type="checkbox"/> Borderline RHD <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe <input type="checkbox"/>		
Echo report consistent with rheumatic heart disease as defined in the World Heart Federation case definition - RHD only (see page 5 for criteria) <input type="checkbox"/>		
Risk Status: High Risk <input type="checkbox"/> Low Risk <input type="checkbox"/>	Priority for recall: 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/>	
Priority Indication: (if different from valve severity)		
Deceased*: Yes <input type="checkbox"/> No <input type="checkbox"/>	Date of death:	

*If patient is deceased, please also complete "Deceased Notification Form"

Notification Information	
Clinician/s who notified the ARF:	
Clinician/s who notified the RHD (if different):	
This form completed by:	Date:

ARF CRITERIA (The 2020 Australian Guideline for prevention, diagnosis and management of acute rheumatic fever and rheumatic heart disease – 3rd edition) – use the ARF diagnosis calculator at: <https://www.rhdaustralia.org.au/apps>

EVIDENCE OF GROUP A STREP (GAS) INFECTION			Meets criteria
Skin swab culture:	Date:	Result:	
Throat swab culture:	Date:	Result:	<input type="checkbox"/>
ASOT:	Date:	Result:	<input type="checkbox"/>
Anti-DNase B:	Date:	Result:	<input type="checkbox"/>

Diagnosis is determined by the number of major and minor criteria and counting rules, however all clinical values available must be recorded on this form

ARF Criteria		Comments:	Meets criteria
Major Manifestation	Aseptic mono-arthritis (high risk)		<input type="checkbox"/>
	Poly-arthralgia (high risk)		<input type="checkbox"/>
	Polyarthritis		<input type="checkbox"/>
	Carditis (includes subclinical evidence of rheumatic valvulitis on echo)		<input type="checkbox"/>
	Chorea (can meet ARF criteria on its own)		<input type="checkbox"/>
	Subcutaneous nodules		<input type="checkbox"/>
Minor manifestation	Erythema marginatum		<input type="checkbox"/>
	Mono-arthralgia (high risk)		<input type="checkbox"/>
	Aseptic mono-arthritis (low risk)		<input type="checkbox"/>
	Poly-arthralgia (low risk)		<input type="checkbox"/>
	ESR ≥ 30mm/hr (high risk)		<input type="checkbox"/>
	ESR ≥ 60mm/hr (low risk)		<input type="checkbox"/>
	Or CRP ≥ 30mg/L		<input type="checkbox"/>
	Fever ≥ 38°C (high risk)		<input type="checkbox"/>
	Fever ≥ 38.5°C (low risk)		<input type="checkbox"/>
	Where documented or reported recent history		
Normal P-R interval on ECG	<input type="checkbox"/>		<input type="checkbox"/>
Prolonged P-R interval on ECG (age specific)	<input type="checkbox"/>		<input type="checkbox"/>
2 nd degree heart block	<input type="checkbox"/>		<input type="checkbox"/>
Complete heart block (3 rd degree)	<input type="checkbox"/>		<input type="checkbox"/>
A/V conduction node abnormality	<input type="checkbox"/>		<input type="checkbox"/>
No ECG	<input type="checkbox"/>		
Evidence of GAS infection?: Yes <input type="checkbox"/> No <input type="checkbox"/>		# Major Manifestations:	# Minor Manifestations:

Prophylaxis Information	
Prophylaxis type: BPG 3w <input type="checkbox"/> BPG 4w <input type="checkbox"/> None <input type="checkbox"/> Oral (Erythromycin) <input type="checkbox"/> Oral (Penicillin) <input type="checkbox"/>	
Other <input type="checkbox"/> _____ Unknown <input type="checkbox"/>	
Start prophylaxis:	Proposed Ceased date:
Date BPG given:	Location BPG given:

Cessation comments: (Please tick either one of the following boxes or write the appropriate comments regarding cease date - as per Cessation comments document)

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16

Education Information	
Education Date:	Education Type: Family <input type="checkbox"/> Initial <input type="checkbox"/> Ongoing <input type="checkbox"/> Surgical <input type="checkbox"/>
Education Clinic:	Education Comments:

Recall Details (Only tick the required reviews, add approximate date review required and any specific comments)		
Review by:	Review due:	Review comments:
Cardiologist <input type="checkbox"/>	Date:	Comment:
Echocardiogram <input type="checkbox"/>	Date:	Comment:
Paediatrician <input type="checkbox"/>	Date:	Comment:
Paediatric cardiologist <input type="checkbox"/>	Date:	Comment:
Physician <input type="checkbox"/>	Date:	Comment:
RHD team internal use for data entry only:		
Discharge summary to be entered: Yes <input type="checkbox"/> Date:		

General Comments:

Please send form to the NT RHD Control Program: RHDDarwin.THS@nt.gov.au or RHDAliceSprings.THS@nt.gov.au

Testing considerations in suspected ARF (Decision Support tool)

(The 2020 Australian Guideline for prevention, diagnosis and management of acute rheumatic fever and rheumatic heart disease – 3rd edition, p 111 – Table 7.3)

Test	Complete	Comments
ECG (if prolonged P-R interval or other rhythm abnormality, repeat in 2 weeks and again at 2 months, if still abnormal)	<input type="checkbox"/>	
Echocardiogram (consider repeating after 1 month, if normal) All cases of suspected or confirmed ARF should undergo early echocardiography to confirm or refute the presence of rheumatic carditis or valvulitis	<input type="checkbox"/>	
FBE <input type="checkbox"/> ESR <input type="checkbox"/> CRP <input type="checkbox"/> ASOT <input type="checkbox"/> Anti-DNase B <input type="checkbox"/>	<input type="checkbox"/>	
In relevant situations:		
Throat swab <input type="checkbox"/> Skin sore swab <input type="checkbox"/>	<input type="checkbox"/>	
Blood cultures, if febrile (plus repeated blood cultures, if possible endocarditis)	<input type="checkbox"/>	
Synovial Fluid aspirate – ensure sample does not clot by using correct tubes, which have been well mixed and transporting them promptly to the laboratory. Include request for cell count, microscopy, culture and gonococcal polymerase chain reaction (PCR)	<input type="checkbox"/>	
Pregnancy test	<input type="checkbox"/>	
Creatinine test (UEC [urea, electrolytes, creatinine]) since NSAIDS can affect renal function	<input type="checkbox"/>	
Differential diagnosis:		
Joint aspirate (microscopy and culture) for possible septic arthritis	<input type="checkbox"/>	
Copper, ceruloplasmin, antinuclear antibody, drug screen for choreiform movements	<input type="checkbox"/>	
Serology and autoimmune markers for autoimmune or reactive arthritis	<input type="checkbox"/>	
Autoantibodies <input type="checkbox"/> Double-stranded DNA <input type="checkbox"/> Anti-cyclic citrullinated peptide (anti-CCP) antibodies <input type="checkbox"/>	<input type="checkbox"/>	
Urine for Neisseria gonorrhoea molecular test	<input type="checkbox"/>	
Urine for Chlamydia trachomatis molecular test	<input type="checkbox"/>	
Serological or other testing for: Viral hepatitis <input type="checkbox"/> Yersinia spp <input type="checkbox"/> Cytomegalovirus (CMV) <input type="checkbox"/> Parvovirus B19 <input type="checkbox"/> Respiratory viruses <input type="checkbox"/> Ross River virus <input type="checkbox"/> Barmah Forest virus <input type="checkbox"/>	<input type="checkbox"/>	

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World Heart Federation criteria for echocardiographic diagnosis of RHD (2012)

(The 2020 Australian Guideline for prevention, diagnosis and management of acute rheumatic fever and rheumatic heart disease – 3rd edition, p 135 – Table 8.5)

Echocardiographic criteria for individuals aged ≤ 20 years

Definite RHD (either A, B, C or D):

- A) Pathological MR and at least two morphological features of RHD of the MV
- B) MS mean gradient ≥ 4 mmHg (note: congenital mitral valve anomalies must be excluded)
- C) Pathological AR and at least two morphological features of RHD of the AV
- D) Borderline disease of both the AV and the MV†

Borderline RHD (either A, B, or C):

- A) At least two morphological features of RHD of the MV without pathological MR or MS
- B) Pathological MR
- C) Pathological AR

Normal echocardiographic findings (all of A, B C, and D):

- A) MR that does not meet all four Doppler echocardiographic criteria (physiological MR)
- B) AR that does not meet all four Doppler echocardiographic criteria (physiological AR)
- C) An isolated morphological feature of RHD of the MV (for example, valvular thickening) without any associated pathological stenosis or regurgitation
- D) Morphological feature of RHD of the AV (for example, valvular thickening) without any associated pathological stenosis or regurgitation

Echocardiographic criteria for individuals aged >20 years

Definite RHD (either A, B, C or D):

- A) Pathological MR and at least two morphological features of RHD of the MV
- B) MS mean gradient, ≥ 4 mmHg
- C) Pathological AR and at least two morphological features of RHD of the AV, only in individuals aged < 35 years
- D) Pathological AR and at least two morphological features of RHD of the MV

† Combined AR and MR in high-prevalence regions and in the absence of congenital heart disease is regarded as rheumatic.

AR, Aortic regurgitation; MR, Mitral regurgitation; MS, mitral stenosis.

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