Training Program Performing systematic reviews of critical values in key laboratory analytes

A collaboration between
Australasian Association of Clinical Biochemists,
Royal College of Pathologists of Australasia
Macquarie University







Welcome and Thank You

RCPA-AACB HIGH RISK RESULTS WORKING PARTY



Andrea Rita Horvath



Andrew Georgiou



Grahame Caldwell



Craig Campbell



Penelope Coates



Hans Schneider



Alan McNeil



Robert Flatman



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Towards a Harmonized Alert Table

- Don't need to produce complete table in first pass.
- Be ready to ready to defend any recommended alert thresholds.
- A systematic, transparent approach is required for wide adoption.

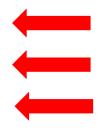
- Review the literature to identify appropriate alert thresholds
- 2. Rate the quality of the evidence on which these thresholds are based

- 3. Perform risk analysis to assess threshold suitability
- 4. Assess transferability and consider the pre- and postanalytical aspects of the alert threshold
- Assess the impact of the selected thresholds on the frequency of critical alerts
 - Seek endorsement for selected thresholds from laboratories and clinical groups

Risk Management Approach

Table 1. Examples of Risk Management for Critical- and Significant-Risk Results²⁰

Stage of Risk Management		Abnormal Laboratory Result	
Risk Analysis	Hazard identification	Neonatal hypoglycemia (< 40 mg/dL; < 2.2 mmol/L)	Unexpected malignancy in routine anatomic pathology specimen
	Potential harm associated with the laboratory or anatomic pathology result	Irreversible neurological injury	Progression of disease that affects therapy, prognosis
	Clinical intervention that can reduce the risk of harm	Correction of hypoglycemia	Referral to specialist for consideration of therapy
Risk Estimation	Probability: Is there reasonable likelihood of harm in absence of intervention?	Yes	Yes
	Severity: Is there reasonable likelihood of severe damage if harm occurs?	Yes	Yes
	Urgency: Is immediate intervention necessary to reduce risk of harm?	Yes	No
	Risk of Process Failure: Is there reasonable likelihood that routine reporting would not permit timely intervention?	Yes	Possible
Risk Evaluation	Is the risk of process failure greater than the clinically acceptable risk, given the estimation of potential harm?	Yes	Yes (unless organization has a process to identify routine reports that are not reviewed by clinician)
Risk Control	Category of abnormal laboratory result (critical-risk vs significant-risk)	Critical-risk result	Significant-risk result
Risk Monitoring	 Are results communicated within intended time frame? Do outcomes support the alert threshold? Are alternative systems available for communicating results available? 	N/A	N/A



Abbreviation: N/A, not applicable.

It would be great if the evidence supports a particular alert threshold

BUT it is just as important to know:

That there is a <u>lack</u> of high quality evidence.