

Episode 53 – The Stiell Sessions: Clinical Decision Rules & Risk Scales

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Clinical Decision Rules (CDRs):

There are hundreds of clinical decision rules and risk scales published in the medical literature, some more widely adopted than others. Ian Stiell shares with us his views and experiences gained from co-creating some of the most influential CDRs to date.

Criteria For Developing a CDR

In order to develop a useful clinical decision tool, a number of criteria must be met. Firstly, the condition needs to be relatively common. Rare conditions will not have the necessary volume of

data to generate high quality decision rules. In addition to being a common complaint or illness, there must also be a perceived inefficiency or clinical variability in practice with regards to the workup of the patient. For example, the inefficiency can be an overuse or under-use of a particular resource (imaging, blood tests etc.) which, given a lack of evidence, physicians have different approaches to. Lastly, the clinical question that leads to the inefficiency needs to be answerable with only a handful of clinical variables.

CDRs vs Risk Stratification Scales

More complex conditions such as CHF and COPD have a spectrum of severity and acuity. Their management cannot be reduced to a binary question, since multiple factors need to be taken into consideration. This is in contradistinction to the Ottawa Ankle rule, in which a binary question is asked – does this patient require an x-ray or not? As a result, these complex conditions require risk stratification scales (rather than CDRs), which estimate the risk of a bad outcome. These scales can help physicians decide what the appropriate management and disposition for the patient would be.

The Development of a CDR

Before CDRs can be safely applied in clinical practice they undergo a rigorous development process. The *four phases of development* include derivation, validation, implementation and studying the barriers to adoption. Although the rule is derived and published in the literature, it should not be used clinically until it has been prospectively validated and shown to be effective.

How CDRs & Risk Scales Can Be Applied to Practice

Despite the rigorous development process that is involved in refining the CDRs, there are some clinicians who are hesitant to apply them in a clinical setting. Some clinicians feel that the rules may be too complex or that they take too long to apply. However, compared to the time it takes to organize and follow up on imaging results, the use of CDRs can be more time efficient for both the physician and the patient, and can improve the flow within a department. Different institutions also have different cultures or habits around investigating and treating a particular condition. This variation is also seen between different health care systems, especially when there are regional differences in the medico-legal environment or funding model. There are some clinicians who believe that their clinical experience is more accurate than clinical decision rules and might place more value on gestalt. Gestalt and experience are valuable and should be used to think critically about our patients. However, what makes emergency medicine challenging is that we rarely get feedback on the patient's clinical course after they have visited the ED. Therefore, no matter how much clinical experience one has, the diagnostic loop is rarely closed as the patient will either follow up with their primary care provider or seen another emergency physician if the problem does not resolve

When considering whether to apply a clinical decision rule, it is important to consider several factors. Most importantly, you must evaluate whether the rule you want to use applies to your patient. To answer this question it's important to know *the inclusion and exclusion criteria* that were used to develop this rule (inclusion and exclusion criteria are provided below along with their respective

CDRs). Secondly, you must consider what information the rule will give you. There are both one-way rules and two-way rules. The PERC rule is an example of a one-way rule. It is designed to rule out a pulmonary embolism in your patient. If the PERC rule is positive it simply means you cannot confidently rule out a PE, but a positive result should not guide your clinical decision-making. The Ottawa Ankle Rule is an example of a two-way rule. Applying it will either direct you to order an x-ray or not order an x-ray.

Below you will find common CDRs and risk stratification tools developed by the research group in Ottawa. We encourage you to become familiar with the inclusion and exclusion criteria. Next time you see a patient, ask yourself whether the rule you are about to apply has been studied in the same population.

The Ottawa Ankle Rules:



Application: A two-way rule to help the physician determine whether a patient with an ankle or foot injury requires an x-ray.

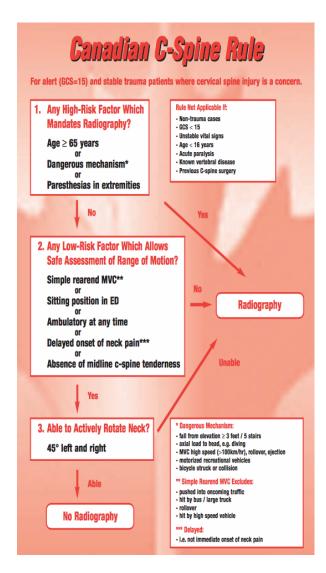
Sensitivity for malleolar fractures: 100% Sensitivity for midfoot fractures: 100%

Inclusion Criteria:

- 1. Adult patient *note that the Ottawa Ankle rules have been validated in Pediatrics
- 2. Any mechanism of blunt ankle injury (including twisting, falls and direct blows)

- 1. Age <18
- 2. Pregnant
- 3. Isolated skin injury
- 4. Injury older than 10 days
- 5. Returning for reassessment of same injury
- 6. Referred from outpatient clinic with existing radiographs

Canadian C-Spine Rule:



Application: A two-way rule to help the physician determine whether a patient with blunt head or neck trauma requires imaging of the C-Spine.

Sensitivity for clinically important C-Spine injury: 100% Specificity for clinically important C-Spine injury: 42.5%

Inclusion Criteria:

- 1. Adult patient
- 2. Acute blunt trauma to the head or neck
- 3. Alert (GCS 15)
- 4. Stable (normal vital signs with SBP>90 mmHg and RR 10-24/min)
- 5. Neck pain
- 6. If no neck then all the following:
 - a. Visible injury above clavicle
 - b. Had not been ambulatory
 - c. Dangerous mechanism of injury

- 1. Age <16
- 2. GCS < 15
- 3. Isolated minor injuries (laceration)
- 4. Grossly abnormal vital signs
- 5. Injury > 48hr ago
- 6. Penetrating trauma
- 7. Presenting with acute paralysis
- 8. Known vertebral disease
- 9. Return for reassessment of same injury
- 10. Pregnant

Canadian CT Head Rule:

Canadian CT Head Rule

CT head is only required for minor head injury patients with any one of these findings:

High Risk (for Neurological Intervention)

- 1. GCS score < 15 at 2 hrs after injury
- 2. Suspected open or depressed skull fracture
- 3. Any sign of basal skull fracture*
- 4. Vomiting ≥ 2 episodes
- 5. Age \geq 65 years

Medium Risk (for Brain Injury on CT)

- 6. Amnesia before impact ≥ 30 min
- 7. Dangerous mechanism ** (pedestrian, occupant ejected, fall from elevation)

*Signs of Basal Skull Fracture

- hemotympanum, 'racoon' eyes, CSF otorrhea/ rhinorrhea, Battle's sign
- ** Dangerous Mechanism
- pedestrian struck by vehicle
- occupant ejected from motor vehicle
- fall from elevation ≥ 3 feet or 5 stairs

Rule Not Applicable If:

- Non-trauma cases
- GCS < 13
- Age < 16 yearsCoumadin or bleeding disorder
- Obvious open skull fracture

Stiell IG, et al. The Canadian CT Head Rule for Patients with Minor Head Injury. Lancet 2001;357:1391-96.

Application: A two- way rule to help physicians determine whether a patient presenting with a minor head injury requires CT imaging of the head

Sensitivity: The five high risk factors have 100% sensitivity for predicting neurological intervention. All seven factors in the rule have a sensitivity of 98.4% for predicting clinically important brain injury.

Specificity: The high risk factors have 68.7% specificity for predicting neurological intervention All seven factors in the rule have a specificity of 49.6% for predicting clinically important brain injury.

Inclusion Criteria:

- 1. Blunt trauma to the head resulting in witnessed loss of consciousness, definite amnesia or witnessed disorientation
- 2. Initial ED GCS >13
- 3. Injury occurred within 24 hours

- 1. Age <16 years old
- 2. Minimal head injury (no LOC, amnesia or disorientation)
- 3. No clear history of trauma as primary event
- 4. Obvious penetrating skull injury or depressed skull fracture
- 5. Acute focal neurological deficit
- 6. Unstable vital signs associated with major trauma
- 7. Seizure before assessment in ED
- 8. Bleeding disorder or use of oral anticoagulants
- 9. Return for assessment of same head injury
- 10. Pregnant

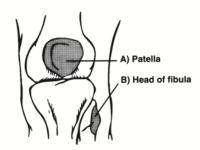
Ottawa Knee Rule:



OTTAWA KNEE RULE



For Knee Injury Radiography



Stiell IG, Greenberg GH, Wells GA, McDowell I, Cwinn AA, Smith NA, Cacciotti TF, Sivilotti MLA.
Prospective validation of a decision rule for the use of radiography in acute knee injuries. JAMA 1996; 275:611-615.

A knee x-ray series is only required for knee injury patients with any of these findings:

- 1. age 55 or older
- 2. isolated tenderness of patella (no bone tenderness of knee other than patella) OR
- 3. tenderness of head of fibula OR
- 4. inability to flex to 90°
- 5. inability to bear weight both immediately and in the emergency department for 4 steps (unable to transfer weight twice onto each lower limb regardless of limping)

Application: A two- way rule to help physicians determine whether a patient presenting with an acute knee injury requires radiography.

Sensitivity for knee fractures: 100% Specificity for knee fractures: 54%

Inclusion Criteria:

- 1. Adult patients
- 2. Blunt knee injury of any mechanism
- 3. Knee defined as: patella, head and neck of the fibula, proximal 8cm of the tibia and distal 8cm of the femur

- 1. Age < 18
- 2. Pregnant
- 3. Isolated injury of the skin without underlying soft-tissue or bone involvement
- 4. Referred with radiographs
- 5. Sustained injury > 7 days ago
- 6. Returned for reassessment of the same injury
- 7. ALOC
- 8. Paraplegic
- 9. Multiple trauma

The Ottawa COPD Risk Scale:

Table 7: Clinical variables contributing to preliminary Ottawa COPD Risk Scale to identify patients with COPD seen in the emergency department who are at high risk of a serious adverse event Variable Points History Coronary bypass graft Peripheral vascular disease intervention 2 Intubation for respiratory distress **Examination** Heart rate on arrival in ED > 110/min Too ill to do walk test after treatment in ED (SaO₂ < 90% or heart rate ≥ 120/min) Investigations Acute ischemic changes on ECG Pulmonary congestion evident on chest radiography Hemoglobin < 100 g/L Urea ≥ 12 mmol/L 1 Serum CO₂ ≥ 35 mmol/L Total score (possible range 0-16) Note: COPD = chronic obstructive pulmonary disease, ECG = electrocardiogram, ED = emergency department, Sao₂ = arterial oxygen saturation.

Clinical variables in the Ottawa COPD Risk Scale

Total score	Risk of adverse event, %	Risk category
0	2.2	Low
1	4.0	Medium
2	7.2	Medium
3	12.5	High
4	20.9	High
5	32.9	Very high
6	47.5	Very high
7	62.6	Very high
8	75.6	Very high
9	NA	Very high
10	91.4	Very high
> 10	NA	Very high

Risk categories for patient in acute COPD exacerbation

Application: This is a risk stratification tool which provides physicians with a risk estimate of short-term serious adverse events for patients presenting to the ED with a COPD exacerbation. It can help the physician make an evidence-based decision regarding admission or discharge of the patient.

Inclusion Criteria:

- 1. Age≥ 50 years
- 2. Presenting with COPD exacerbation, defined as:
 - a. Increase in at least 2 of 3 specified symptoms (breathlessness, sputum volume, sputum purulence)
- 3. COPD has been previously diagnosed or diagnosed in the ED on the basis of a 1year history of chronic dyspnea or cough with sputum production.
- 4. ≥ 15 pack year smoking history
- 5. Prior or current evidence of airflow obstruction

Exclusion Criteria: (Patients too unwell to be discharged)

- 1. Resting O2 Sat <85%
- 2. Heart Rate ≥ 130bpm/min
- 3. SBP <85 mm Hg
- 4. Confusion, disorientation or severe dementia
- 5. Ischemic chest pain requiring treatment on arrival
- 6. Acute ST elevation on ECG on arrival
- 7. Arrival from nursing home or chronic care facility
- 8. Death from chronic illness expected within weeks

The Ottawa Heart Failure Risk Scale:

Ottawa Heart Failure Risk Scale

<u>Items</u>	Points
1. History	
a) Stroke or TIA	1
b) Intubation for respiratory distress	2
2. Examination	
a) Heart rate on ED arrival ≥ 110	2
b) SaO ₂ < 90% on arrival	1
c) Heart rate ≥ 110 during 3-minute walk test	1
(or too ill to perform walk test)	
3. Investigations	
a) ECG has acute ischemic changes	2
b) Urea ≥ 12 mmol/L	1
c) Serum CO₂ ≥ 35 mmol/L	2
d) Troponin I or T elevated to MI level	2
e) NT-proBNP ≥ 5,000 ng/L	1

Total Score (0 - 15):

Heart Failure Risk Categories for Serious Adverse Events

Total Score	Risk	Category
0	2.8%	Low
1	5.1%	Medium
2	9.2%	Medium
3	15.9%	High
4	26.1%	High
5	39.8%	Very High
6	55.3%	Very High
7	69.8%	Very High
8	81.2%	Very High
9	89.0%	Very High

Application: This is a risk stratification tool which provides physicians with a risk estimate of short-term serious adverse events for patients presenting to the ED with acute shortness of breath secondary to heart failure. It can help the physician make an evidence-based decision regarding admission or discharge of the patient.

The score above was developed in the derivation phase. During the validation phase the score was reduced to 5 variables. Although the risk stratification for the variables is not yet published our experts have shared the variables with our audience.

Validated CHF variables (unpublished as of November 2014):

- 1. IV Nitrate use
- 2. Troponin at 5x upper reference
- 3. High PCO2
- 4. High Urea or Cr
- 5. Failing walk test
 - a. O2<90 or HR>110 at rest
 - b. Too sick to walk
 - c. Unable to walk for 3min after standing

Inclusion Criteria:

- 1. Age ≥ 50
- 2. Presenting with acute SOB secondary to an exacerbation of chronic heart failure or new-onset heart failure
- 3. The diagnosis of heart failure was defined as:
 - a. Appropriate symptoms (shortness of breath or fatigue)
 - b. Clinical signs of fluid retention (pulmonary or peripheral
 - c. Presence of an underlying abnormality of cardiac structure or function
 - d. If there was doubt about etiology, a beneficial response to treatment (ie, diuresis) was included.

Exclusion Criteria: (Patients too unwell to be discharged)

- 1. Resting O2 sat <85% on room air or after being on the usual home O2 setting for 20min.
- 2. HR ≥ 120bpm/min on arrival
- 3. SBP <85 mm Hg on arrival
- 4. Confusion, disorientation or dementia
- 5. Ischemic chest pain requiring nitrates on arrival
- 6. Acute ST-segment elevation on ECG on arrival
- 7. Death from chronic illness expected within weeks
- 8. Arrival from nursing home or chronic care facility

The Ottawa TIA Risk Score

Items	Points
Clinical findings	
First TIA (in lifetime)	2
Symptoms ≥10 min	2
History of carotid stenosis	2
Already on antiplatelet therapy	3
History of gait disturbance	1
History of unilateral weakness	1
History of vertigo	-3
Initial triage diastolic blood pressure ≥110 mm Hg	3
Dysarthria or aphasia (history or examination)	1
Investigations in emergency department	
Atrial fibrillation on ECG	2
Infarction (new or old) on CT	1
Platelet count ≥400×10 ⁹ /L	2
Glucose ≥15 mmol/L	3
Total score (-3 to 23)	

CT indicates computed tomography; and TIA, transient ischemic attack.

Clinical Variables

Application: This is a risk stratification tool, which provides physicians with an estimate of the risk that a patient presenting to the ED with TIA will suffer a stroke within 7 days.

Score	No. of Patients (n=3899)	Sensitivity (95% CI)	Specificity (95% CI)	Estimated Probability of Having Outcome, %
-3	1	1.0	0.0	0.01
-1	3	1.0	0.0 (0.0-0.0)	0.03
0	15	1.0	0.0 (0.0-0.0)	0.04
1	20	1.0	0.01 (0.0-0.01)	0.07
2	65	1.0	0.01 (0.01-0.01)	0.11
3	127	1.0	0.03 (0.02-0.03)	0.2
4	226	1.0	0.06 (0.05-0.07)	0.3
5	449	0.99 (0.97-1.0)	0.12 (0.11-0.13)	0.5
6	473	0.98 (0.94-1.0)	0.24 (0.22-0.25)	0.8
7	742	0.94 (0.89-0.99)	0.36 (0.35-0.38)	1.2
8	772	0.84 (0.76-0.92)	0.55 (0.54-0.57)	2.0
9	544	0.63 (0.53-0.73)	0.75 (0.74-0.76)	3.2
10	253	0.41 (0.30-0.51)	0.89 (0.88-0.90)	5.1
11	114	0.28 (0.18-0.37)	0.95 (0.95-0.96)	8.1
12	62	0.20 (0.11-0.28)	0.98 (0.98-0.98)	12.6
13	24	0.05 (0.0-0.09)	0.99 (0.99-1.0)	19.0
14	9	0.01 (0.0-0.03)	1.0 (1.0-1.0)	27.6

CI indicates confidence interval; and TIA, transient ischemic attack.

Probability of TIA based on score

Inclusion Criteria:

- 1. Age > 18
- 2. Diagnosed with a TIA in the ED by either the ED physician or neurologist

- 1. Patients who were diagnosed with a confirmed stroke (neurological deficit present >24 hours)
- 2. Decreased LOC (GCS <15)
- 3. Documented other cause for deficit which was not a TIA
- 4. Presented >7 days following onset of most recent TIA and treated with TPA for an acute stroke.

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