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<b>1</b> Abdominal - Flank Pain		
Pain between the lower border of the chest and	the pelvis. If pain in the middle of t	the back, see 4-Back Pain.
<ul> <li>Abdominal - Flank Pain</li> <li>Pain between the lower border of the chest and</li> <li>BACKGROUND</li> <li>M • Current medications? <ul> <li>NSAIDs?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>Prior abdominal operations / procedures?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> </ul> </li> <li>HISTORY</li> <li>O • When did the pain start? What were you doing? <ul> <li>Time till max intensity: sec? min? hr?</li> <li>P • Pain location? Size of the painful area?</li> <li>Radiation?</li> <li>Q • Burning, aching, sharp?</li> <li>R • Worse with deep inspiration?</li> <li>Worse with movement?</li> </ul> </li> <li>S • VAS (1-10)? How pain affects daily function?</li> </ul>	the pelvis. If pain in the middle of the pelvis. If pain is the performance of the perfo	<ul> <li>the back, see 4-Back Pain.</li> <li>CONSIDER IF UNCLEAR: The cause of abdominal - flank pain may remain unclear after routine bedside information is obtained. Admission for observation and/or abdominal CT should be considered for these patients in the following situations:</li> <li>1. Abdominal pain + shock</li> <li>2. Severe abdominal pain with sudden onset</li> <li>3. Decreased functional ability</li> <li>4. Generalised peritonitis</li> <li>5. Suspected bowel obstruction</li> <li>6. Inflammed right lower quadrant</li> </ul>
<ul> <li>Constant or intermittent? Increasing?</li> <li>Prior similar painful episodes?</li> <li>PO: nausea, vomiting?</li> <li>PR: diarrhea, constipation?</li> <li>PU: dysuria?</li> <li>PV (for fertile women): last period? Discharge?</li> </ul>	ycars)	

# **1** Abdominal - Flank Pain: Clinical Syndromes & Diagnostic Rules

### **ABDOMINAL PAIN & CHOCK**

Abdominal pain with the following:

- Tachycardia and/or hypotension
- Elevated lactate, base deficit

Potential diagnoses:

- Ruptured abdominal aortic aneurysm
- Ruptured ectopic pregnancy
- Perforation (e.g. ulcer, diverticulus) and sepsis
- Severe pancreatitis, cholangitis

### SEVERE & SUDDEN ABDOMINAL PAIN

- Sudden onset of diffuse abdominal pain
- Severe pain that does not respond to analgesics
- Peritoneal findings are absent

Potential diagnoses:

- Mesenteric ischemia
- Aortic dissection
- Perforated ulcer
- Ovarian torsion, testicular torsion

## DECREASED FUNCTIONAL ABILITY

Patients (often elderly patients) who are sufficiently affected by their abdominal pain that they cannot function at home.

## **GENERALIZED PERITONITIS**

- Pain worsens with movement
- Diffuse tenderness Rigidity or rebound tenderness

Potential diagnoses:

- Perforated ulcer
- Perforated diverticulitis
- Perforated appendicitis
- Cholecystitis, pancreatitis

### **BOWEL OBSTRUCTION**

Pain with several of the following:

- Prior abdominal surgery
- Diffuse, crampy pain, intermittent spikes
- Vomiting, decreased bowel movements, absent flatus
- Swollen abdomen
- Constant, hyperactive, "metallic" abdominal sounds
- The abdomen is diffusely tender in the absence of peritoneal findings

## **RIGHT LOWER QUADRANT**

- Right lower quadrant (RLQ) pain
- RLQ peritonitis OR elevated WBC/CRP

Potential diagnoses:

- Acute appendicitis
- Salpingitis
- Ovarial pathology
- Mesenteric adenitis
- Sigmoiditis

### APPENDICITIS INFLAMMATORY RESPONSE SCORE

Criteria	Points
RLQ pain	1
Vomiting	1
Peritonitis	1, 2 or 3
WBC count	1 (10-14.9), 2 (≥ 15)
% Neutrofils	1 (70-84%), 2 (≥ 85%)
CRP	$1(10-49), 2 \ge 50)$
Temp ≥ 38.5°	1

Probability: 0-4 low, 5-8 indet., 9-12 high

### **APPENDICITIS vs SALPINGITIS**

In fertile women:

Criteria	Salpingitis
Absent pain migration	OR 4.2
Bilateral tenderness	OR 16.7
No nausea or vomiting	OR 8.4
All of the above	99%

2 Allergic Reaction		
Suspected allergic reaction (rash, pruritus, swellin	ig etc).	
BACKGROUND	PHYSICAL	<b>CONSIDER:</b>
<ul> <li>M • Recently taken/terminated medications/substances?</li> <li>• Recent NSAID use?</li> <li>A • Known allergies to medications, food, other?</li> <li>P • Past medical history?</li> <li>• Recent medical test (e.g. with contrast agent)?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> </ul>	<ul> <li>A • Hoarse? Stridor?</li> <li>• Lip- tongue swelling?</li> <li>B • SpO2%</li> <li>• Respiratory rate?</li> <li>• Lung auscultation?</li> <li>• Chest wall examination</li> <li>C • Pulse/blood pressure</li> </ul>	<ol> <li>Anaphylaxis</li> <li>Angioedema</li> </ol>
<ul><li>S • Smoking: amount? Prior smoking?</li><li>HISTOPN</li></ul>	<ul> <li>Heart rate</li> <li>D Level of consciousness?</li> <li>E Front side of the body</li> </ul>	
<ul> <li>O • When did the symptoms start? What were you doing?</li> <li>Time till max intensity: sec? min? hr?</li> <li>P • Which body parts are affected?</li> <li>Q • Rash? Swelling? Itch? Pain?</li> <li>R • Effect of measures if taken (e.g. corticosteroids, antihistamine)?</li> <li>S • Effect on daily function?</li> <li>T • Constant, intermittent, increasing symptoms?</li> <li>Prior similar episodes?</li> <li>+ • Food intake?</li> <li>Insect bite?</li> <li>New soap / washing detergent?</li> </ul>	<ul> <li>Back side of the body</li> <li>Temperature?</li> </ul>	

# **2** Allergic Reaction: Clinical Diagnostic Clues

# ANAPHYLAXIS

Anaphylaxis is a severe, systemic hypersensitivity reaction that affects airway, breathing and/or circulation and is usually associated with skin (e.g. urticarial) and/or mucosal symptoms (Soar 2010). Anaphylaxis is highly likely in any one of the following three contexts (Sampson 2006):

1. Acute onset of an illness (minutes to several hours) with involvement of the skin, mucosal tissue, or both (eg, generalized hives, pruritus or flushing, swollen lips-tongue-uvula) and at least one of the following:

- Respiratory compromise (eg, dyspnea, wheeze-bronchospasm, stridor, reduced PEF, hypoxemia)
- Reduced BP or associated symptoms of end-organ dysfunction (eg, hypotonia [collapse], syncope, incontinence)

2. Two or more of the following that occur rapidly after exposure to a *likely* allergen for that patient (minutes to several hours):

- Involvement of the skin-mucosal tissue (eg, generalized hives, itch-flush, swollen lips-tongue-uvula)
- Respiratory compromise (eg, dyspnea, wheeze-bronchospasm, stridor, reduced PEF, hypoxemia)
- Reduced BP or associated symptoms (eg, hypotonia [collapse], syncope, incontinence)
- Persistent gastrointestinal symptoms (eg, crampy abdominal pain, vomiting)

3. Reduced BP after exposure to *known* allergen for that patient (minutes to several hours):

- Infants and children: low SBP (age specific) or > 30% decrease in SBP
- Adults: SBP of less than 90 mm Hg or > 30% decrease from that person's baseline.

## ANGIOEDEMA

Angioedema results from the fast onset of increased vascular permeability in subcutaneous or submucosal tissue. Symptoms and signs include:

- Swelling of the face (eyelids, lips, tongue), extremities and genitalia
- Swelling of the larynx, resulting in throat tightness, dysphea, dysphonia, dysphagia
- Swelling of the intestine, resulting in abdominal pain, nausea and vomiting
- Urticaria, flushing, generalized pruritus, bronchospasm and/or hypotension are present in the setting of histamineinduced angioedema but absent in the setting of bradykinin-induced angioedema (e.g. ACE-inhibitor induced, hereditary or acquired C1-inhibitor deficiency)

BACKGROUND		PHY	SICAL	TESTS
<ul> <li>M • Current medica</li> <li>Recent changes</li> <li>A • Allergies?</li> <li>P • Past medical his</li> <li>L • Life circumstant</li> <li>E • Alcohol: how of</li> <li>S moking: among</li> </ul>	tions? ? story? ces? often? How much? unt? Prior smoking?	A B	<ul> <li>Trauma to the head?</li> <li>Tongue bite?</li> <li>SpO2%</li> <li>Respiratory rate?</li> <li>Lung auscultation?</li> <li>Chest wall examination</li> <li>Pulse/blood pressure</li> </ul>	<ul> <li>Acid-base: pH, pCO<sub>2</sub>, HCO<sub>3</sub>/BE</li> <li>Electrolytes: Na, K, Ca</li> <li>Hg, WBC, CRP, Trombocytes, INR</li> <li>Creatinin</li> <li>Liver function tests</li> <li>EKG if &gt; 50 years</li> </ul>
<ul> <li>HISTORY</li> <li>When did the priving?</li> <li>Time till max in</li> <li>Decreased or al</li> <li>Effect on daily</li> <li>T Time course? I</li> <li>Prior similar ep</li> <li>Pain?</li> <li>Fever/chills?</li> </ul>	roblem start? Activity at the atensity: sec? min? hr? tered consciousness? function? Diurnal fluctuation? isodes?	D	<ul> <li>Heart rate</li> <li>QRS width, regularity?</li> <li>Level of consciousness?</li> <li>Eye / pupil examination</li> <li>Focal neurological deficits arm/leg?</li> <li>Glucose?</li> <li>Front side of the body</li> <li>Back side of the body</li> <li>Temperature?</li> </ul>	<ul> <li>CONSIDER IF UNCLEAR:</li> <li>1. Stroke including basilar thrombosis</li> <li>2. Sepsis, meningitis</li> <li>3. Herpes encephalitis</li> <li>4. Non-convulsive status</li> <li>5. Wernicke's encephalopathy</li> </ul>

# **3** Altered Consciousness: Clinical Syndromes

## **METABOLIC CAUSE**

The presence of the following three findings suggests a metabolic cause of coma (Sn 96%):

- Age  $\leq 50$  years
- SBT  $\leq$  150 mm Hg
- Lack of focal neurological findings

# **BACTERIAL MENINGITIS**

95% of adults with community-acquired bacterial meningitis had  $\ge 2$  of the following:

- Headache
- Fever
- Neck stiffness
- Change in mental status

# WERNICKE'S ENCEPHALOPATHY

The classic triad of encephalopathy, ocular abnormalities and gait ataxia is present in only 17% of cases. Caine et al recommend the following operational criteria to identify patients with Wernicke's encephalopathy:  $\geq 2$  of:

- **Dietary deficiencies** (e.g. chronic alcohol abuse, anorexia nervosa, gastrointestinal surgery including bariatric surgery, hyperemesis of pregnancy, prolonged intravenous feeding without proper supplementation)
- Altered mental status (e.g. confusion, apathy, inattentiveness, inability to concentrate, disorientation) or mild memory impairment
- **Oculomotor abnormalities** (e.g. nystagmus, symmetrical or asymmetrical palsy of both lateral recti or the other ocular msucles, conjugated-gaze palsies)
- Cerebellar dysfunction (incoordination of gait or truncal ataxia)

4 Back Pain		
Pain in the middle of the back. If lateral/anterior p	oain, see 5-Chest - Thoracic Pa	ain or 1-Abdominal - Flank Pain
BACKGROUND	PHYSICAL	<b>CONSIDER:</b>
<ul> <li>Pain in the middle of the back. If lateral/anterior p</li> <li>BACKGROUND</li> <li>M • Current medications? <ul> <li>Analgesics: amount, frequency?</li> </ul> </li> <li>A • Allergies?</li> <li>P • Past medical history? Prior cancer? <ul> <li>Recent invasive procedures / medical tests?</li> </ul> </li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> <li>HISTORY</li> <li>O • When did the pain start? What were you doing? <ul> <li>Time till max intensity: sec? min? hr?</li> <li>P • Pain location? Size of the painful area?</li> <li>Radiation?</li> <li>Q • Type of pain: aching, sharp/riping?</li> </ul> </li> </ul>	PHYSICAL         Vitals       • RR, SpO2%, HF         BP, Temp?         Back       • Inspection         • Palpation         Neuro       • Leg strength & g         • Leg sensation         • Patella reflex & Babinski         • Romberg	An or 1-Abdominal - Flank Pain CONSIDER: A. 1. Ruptured abdominal aortic aneurysm 2. Aortic dissection 3. Cauda equina / conus medullaris 4. Cancer, osteomyelitis, discitis 5. Fracture
<ul> <li>Decreased pain with analgesia?</li> <li>Decreased pain when lying down?</li> <li>Increased pain upon flexion, extension, walking?</li> <li>S VAS (1-10)? Effect on daily function?</li> <li>T Constant or intermittent? Increasing?</li> <li>Prior similar painful episodes?</li> <li>+ Leg weakness? Decreased perineal/leg sensation?</li> <li>Altered urination / defecation?</li> <li>Fever / chills?</li> </ul>		

ACUTE LOW B	BACK PAIN	LUMBAR SPINAL STENOSIS	
Forseen and Core	y recommend categorizing patients with acute low back	RISK FACTORS	POINTS
pain (< 4 weeks o	of symptoms) into three categories for the sake of further	History	
management, wit	h radiology (e.g. MRI, bone scan) and lab tests restricted to	• Age 60-70 years	1
patients with serie	ous conditions.	• Age > 70 years	2
Serious	Presence of $\geq 1$ "red flag":	Absence of diabetes	1
condition*	• Age $> 50$ years	Neurogenic claudication	3
	• Steroid use	• Exacerbation of symptoms when standing up	2
	Intravenous drug use	• Symptom improvement when bending forward	3
	History of cancer	Physical Examination	
	<ul><li>Osteoporosis</li></ul>	Symptoms induced by having patients bend forward	-1
	<ul> <li>Trauma history</li> <li>Unintentional weight loss</li> <li>Progression of symptoms</li> </ul>	<ul> <li>Symptoms induced by having patients bend backward</li> </ul>	1
	<ul> <li>Flogression of symptoms</li> <li>Focal neurologic deficit</li> </ul>	Good peripheral artery circulation	3
Spinal stenosis	<ul> <li>Spinal stenosis: low back or radicular pain that</li> </ul>	Abnormal Achilles tendon reflex	1
/ radiculopathy	increases with walking and improves with flexion (sitting or propping)	• Straight Leg Raise test positive for reproducing pain	-2
	• Radiculopathy: dysfunction of a nerve root associated with pain, sensory impairment, weakness, or diminished deep tendon reflexes in nerve root distribution (see 13-Neurological deficit)	$\geq$ 7 points: sensitivity 93%, specificity 72%, LR+ 3.3	1, LR- 0.1
Idiopathic /	No red flags		
nonspecific	• No signs / symptoms of spinal stenosis/radiculopathy		

5	Chest - Thoracic Pain		
Pain	or discomfort localized to or under the chest	wall (including the back). If pain in the	e midline of the back, see 4
BAC M A P L E S	<ul> <li>Current medications?</li> <li>Birth control pill, other hormonal treatments?</li> <li>Allergies?</li> <li>Past medical history?</li> <li>Prior heart- or thromboembolic disease?</li> <li>Life circumstances?</li> <li>Alcohol: how often? How much?</li> <li>Smoking: amount? Prior smoking?</li> </ul>	PHYSICALVitalsRR, SpO2%, HR, BP, Temp?HeartS3/S4, murmurs?Elevated JVP?LungRales?Decreased breath sounds?MSKRedness? Rash?Tenderness on palpation?AbdoUpper abdominal tenderness?	<ul><li>CONSIDER:</li><li>1. Acute coronary syndrome</li><li>2. Pulmonary embolism</li><li>3. Aortic dissection</li></ul>
HIST O P Q R S T +	<ul> <li><b>FORY</b></li> <li>When did the pain start? What were you doing?</li> <li>Time till max intensity: sec? min? hr?</li> <li>Pain location? Size of the painful area?</li> <li>Radiation?</li> <li>Cramping, aching, sharp, riping, burning?</li> <li>Worse with deep inspiration?</li> <li>Worse with deep inspiration?</li> <li>Worse with movement?</li> <li>VAS (1-10)?</li> <li>Constant or intermittent? Increasing?</li> <li>Prior similar painful episodes?</li> <li>Wind: shortness of breath?</li> <li>Walk: leg pain/swelling?</li> <li>Warm: fever/chills?</li> </ul>	TESTS <ul> <li>WBC &amp; CRP</li> <li>Troponin if &gt; 40 years</li> <li>EKG</li> </ul>	

# **5** Chest - Thoracic Pain: Clinical Diagnostic Rules

### ACUTE CORONARY SYNDROME

Age	< 40 years	40 - 65 years	> 65 years
ACS Prevalence	0-2%	8-10%	12-19%
0 Risk Factors*	LR 0.17	LR 0.53	LR 0.96
$\geq$ 4 Risk Factors*	LR 7.4	LR 2.1	LR 1.09

\* diabetes, smoking, hypercholesterolemia, hypertension, heredity

**History:** high-risk features include pressure-type pain, radiation to one or both arms, worsening with exertion (but not with inspiration, position), similarity to prior ischemia.

EKG	ST Elevation	ST depression	T wave inversion
LR	22	5.3	1.8

### **Oh-Troponin**

hs-cTnT < 5 ng/L + History not high-risk + EKG non-ischemic rules-out 30-day MACE (acute myocardial infarction, unstable angina, cardiac arrest, cardiogenic shock, death, high-risk arrhythmias) with 99.2% sensitivity and a negative predictive value of 99.7%.

### **0h/1h-Troponin** ( $\Delta$ = difference)

Rule-Out 30-day MACE	Rule-In 30-day MACE
0h hs-cTnT < 12 ng/L AND	$0h hs-cTnT \ge 52 ng/L OR$
$1h\Delta < 3 \text{ ng/L AND}$	$1h\Delta \ge 5 \text{ ng/L OR}$
History not high-risk AND	0h or 1h hs-cTnT > 14 ng/L + either history
EKG non-ischemic	high-risk or ischemic EKG
Patients for whom 30 day MACE neither	rulad in nor rulad out: consider additional transmir

Patients for whom 30-day MACE neither ruled-in nor ruled-out: consider additional troponin testing or stress testing / myocardial imaging (as out-patient?).

### AORTIC DISSECTION DETECTION (ADD) RISK SCORE

**High risk conditions:** Marfan syndrome, family history of aortic disease, known aortic valve disease, recent aortic manipulation, known thoracic aortic aneurysm

**High risk pain features**: abrupt in onset, severe in intensity, ripping or tearing

**High risk examination features**: evidence of perfusion deficit (pulse deficit, systolic BP differential, focal neurologic deficit in conjunction with pain), murmur of aortic insufficiency (new or not known to be old and in conjunction with pain), hypotension or shock state

ADD score: #categories featuring  $\ge 1$  high risk feature/condition. High risk if ADD score  $\ge 2$ .

### **AORTIC DISSECTION & d-dimer**

A negative serum D-dimer (<500 ng/dL) rules out AD if the ADD score is  $\leq 1$ 

### WELLS SCORE FOR PE See 6-Dyspnea

# 6 Diarrhea

### Loose bowel movements.

### BACKGROUND

- **M** Current medications?
  - Recent antibiotic use?
- **A** Allergies?
- **P** Past medical history?
- L Life circumstances?
- **E** Alcohol: how often? How much?
- **S** Smoking: amount? Prior smoking?

## **HISTORY**

- When did the diarrhea start?
  - Travel history? Food prior to diarrhea onset?
- **Q** Watery? Bloody? Tarry?
- **R** Worsened with foot / fluid intake?
- **S** Volume? Frequency?
- **T** Duration?
  - Prior similar episodes?
- + Fever?
  - Abdominal pain?

# PHYSICAL

Abdo

PR

**TESTS** 

• CRP

- Vitals RR, SpO2%, HR, BP, Temp?
  - Inspection
  - Auscultation
  - Palpation
  - Stool colour?

# **CONSIDER:**

- 1. Sepsis
- 2. Gastrointestinal bleeding
- 3. Invasive bacterial syndrome
- 4. Epidemiological features justifying presumptive antimicrobial therapy

# **6** Diarrhea: Clinical Diagnostic Clues

# **BAYESIAN APPROACH TO ACUTE INFECTIOUS DIARRHEA IN ADULTS**

Goodgame recommends categorizing adults with acute infectious diarrhea ( $\geq$  3 loose stools per day for < 14 days) into three categories for the sake of further management:

Category	Features	Infectious agent	Management
Viral or "norovirus-like" diarrhea	<ul> <li>No specific epidemiologic risk factor</li> <li>No clinical feature suggestive of severe bacterial infection</li> </ul>	<ul> <li>Norovirus</li> <li>Bacteria (including e.g. Salmonella) and protozoa producing an uncomplicated gastroenteritis syndrome</li> </ul>	<ul> <li>No specialized diagnostic testing or antimicrobial management</li> <li>Avoid milk products</li> <li>Loperamid 4 mg once and 2 mg with each liquid stool</li> </ul>
Severe bacterial infection	<ul> <li>Fever &gt; 38.5°C</li> <li>Bloody diarrhea</li> <li>Voluminous diarrhea</li> <li>Severe abdominal pain</li> <li>&gt; 6 stools per 24 hours</li> <li>Diarrhea persisting &gt; 7 days</li> </ul>	<ul> <li>Salmonella, Campylobacter, Shigella</li> <li>Shiga-toxin producing E coli</li> <li>Yersinia</li> <li>Vibrio</li> <li>Clostridium difficile</li> </ul>	<ul> <li>Stool testing for bacterial (or amoebic) infection, shiga toxin</li> <li>If the signs and symptoms are severe, presumptive antibiotic therapy is recommended (unless E coli O157:H7 is suspected)</li> </ul>
Epidemiologic risk factors	<ul> <li>Travel</li> <li>Hospitalized &gt; 3 days</li> <li>Antiobiotic use</li> <li>Contact with health care</li> </ul>	<ul> <li>80% probability of bacterial etiology</li> <li>Persistent diarrhea suggests a protozoa</li> <li>Clostridium difficile</li> </ul>	<ul> <li>Presumptive antibiotic therapy combined with clinical observation</li> <li>Stools for Clostridium difficile toxin</li> <li>Presumptive treatment while awaiting test results is appropriate in severely</li> </ul>
	<ul><li>personnel</li><li>Immunocompromised host</li></ul>	Virus, bacteria, mycobacteria, protozoa	ill patients

## HEMOLYTIC-UREMIC SYNDROME

Diarrhea occurring in the setting of hemolysis, thrombocytopenia and uremia suggests hemolytic-uremic syndrome. Most cases are caused by E coli O157:H7.

# 7 Dyspnea

### Shortness of breath.

## BACKGROUND

- **M** Current medications?
  - Birth control pill, other hormonal treatments?
- A Allergies?
- **P** Past medical history?
  - Prior heart- or thromboembolic disease?
- L Life circumstances? (e.g. occupation, pets)?
- **E** Alcohol: how often? How much?
- **S** Smoking: amount? Prior smoking?

## **HISTORY**

- When did the dyspnea start? What were you doing?
  - Time till max intensity: sec? min? hr?
- **P** Worse when lying down?
- **Q** Air hunger? Chest tightness?
- **R** Worse when lying down?
- **S** Effect on daily function?
- **T** Constant or intermittent? Increasing?
  - Prior similar episodes?
- + Chest pain or discomfort?
  - Leg pain or swelling?
  - Fever / chills?
  - Cough (dry or productive-sputum colour?)

# PHYSICAL

- Vitals RR, SpO2%, HR, BP, Temp?
- **Heart** S3/S4, murmurs?
  - Elevated JVP?
- Lung Chest wall movements?
  - Auscultation: rales? ronchi? decreased breath sounds?
- Leg Swelling? Edema?

# TESTS

- Venous blood gas (pH, pCO2, HCO3/BE)
- CRP
- EKG if > 40 years

# **CONSIDER:**

- 1. Upper respiratory tract problem
- 2. Acute coronary syndrome
- 3. Pulmonary embolism
- 4. Pneumonia

# 7 Dyspnea: Clinical Diagnostic Rules & Clues

### PULMONARY EMBOLISM: THE SIMPLIFIED WELLS (CANADIAN) SCORING SYSTEM

Purpose: ruling-out PE with a negative d-dimer

**Inclusion**: clinically suspected PE: sudden onset of dyspnea, sudden deterioration of existing dyspnea, or sudden onset of pleuritic chest pain without another apparent cause

**Exclusion**: therapeutic doses of unfractionated or low-molecular-weight heparin for > 24 hrs, life expectancy < 3 mo, pregnancy, < 18 years, allergy to IV contrast, renal insufficiency (Crea clearance < 30 ml/min),

too ill to undergo CT scanning, hemodynamic instability

RISK FACTORS	POINTS
• Clinical signs and symptoms of deep venous thrombosis*	3
• Alternative diagnosis less likely than pulmonary embolism	3
• Heart rate > 100/min	1.5
• Immobilization (> 3 days) or surgery in previous 4 weeks	1.5
• Previous pulmonary embolism or deep ven thrombosis	1.5
• Hemoptysis	1
• Malignancy (receiving treatment, treated in the last 6 mo or palliative)	1

\* minimum of leg swelling and pain with palpation of the deep veins PE unlikely (score  $\leq 4$ ) + negative d-dimer: 0.5% nonfatal PE/DVT at 3 month follow-up

# HEART FAILURE

Background	<ul> <li>Heart failure</li> </ul>	LR+ 5.8	LR- 0.45
	<ul> <li>Myocardial infarction</li> </ul>	LR+ 3.1	LR- 0.69
Symptoms	• PND*	LR+ 2.6	LR- 0.70
	Orthopnea	LR+ 2.2	LR- 0.65
	<ul> <li>Dyspnea on exertion</li> </ul>	LR+ 1.3	LR- 0.48
Physical	• S3	LR+11	LR- 0.88
	• JVD**	LR+ 5.1	LR- 0.66
	• Rales	LR+ 2.8	LR- 0.51
	• Wheezing	LR+0.5	LR- 1.3
EKG	<ul> <li>Atrial fibrillation</li> </ul>	LR+ 3.8	LR-0.79
	• Any abnormal finding	LR+ 2.2	LR- 0.64
Ultrasound	Reduced EF	LR+ 4.1	LR-0.24
	• IVC ≥ 20.5 mm	SN 90%	SP 73%
	• Pleural effusion(s)	LR+ 2.0	LR- 0.49
	<ul> <li>Positive B-line scan</li> </ul>	LR+ 7.4	LR-0.16
Chest X-ray	Venous congestion	LR+ 12.0	LR-0.48
	Cardiomegaly	LR+ 3.3	LR- 0.33
BNP	• > 100 pg/ml	LR+ 2.2	LR-0.11
NT-proBNP	• > 300 pg/ml	LR+ 1.8	LR- 0.09

\* PND = paroxysmal nocturnal dyspnea.

\*\* JVD = Jugular venous distension

7 Dyspnea: Clinical Diagnostic Rules & Clues						
OTTAWA HEART FAILURE RISK SCALE	CATEGORY	POINTS	SCORE RISK			
<b>Purpose</b> : predict death from any cause within	Initial assessment		0 3%			
30 days or ED visit or serious adverse event	History of stroke or TIA	1	1 5%			
within 14 days of ED visit (regardless of	• History of intubation for respiratory distress	2	2 9%			
whether admitted): admission to critical care or	• Heart rate on ED arrival $\geq 110$	2	3 16%			
acute monitoring unit where the patient is too ill	• Room air SaO2 < 90% on EMS or ED arriva	1 1	4 26%			
to ambulate, endotracheal intubation or NIV,	Investigations		5 40%			
CABG/PCI/cardiac surgery return to ED for any	• EKG has acute ischemic changes	2	6 55%			
related medical problem (e.g. for respiratory	• Urea $\ge 12 \text{ mmol/L}$	1	7 70%			
distress fever sepsis) and admission	• Serum $CO2 \ge 35 \text{ mmol/L}$	2	8 81%			
<b>Inclusion</b> : $\geq$ 50 vr. presenting to ED with	• Troponin I or T elevated to MI level	2	9 89%			
shortness of breath $< 7$ days duration due to	• NT-ProBNP $\geq$ 5,000 ng/L	1	*Patient is asked to walk at			
exacerbation of chronic HF or new-onset HF	Walk Test* after ED treatment		their own pace for 3 minutes			
(pulmonary or peripheral fluid retention +	• One of the following:	1	in the ED, regardless of the			
abnormal cardiac structure or function)	$\circ$ SaO2 < 90% on room air or usual O2		distance covered			
<b>Exclusion</b> : too ill to be discharged after 2-15	○ HR $\ge$ 110 during 3-minute walk test					
hrs of ED management: $SpO2 < 85\%$ or after	$\circ$ Too ill to walk					
being on home oxygen levels > 20 min, heart						
rate $\geq$ 120/min on arrival, SBP < 85 mm Hg on						
arrival, confusion / disorientation / dementia,						
Ischemic chest pain or acute \$1-1 changes,						
STEMI, terminal status, nursing nome or						
chronic care facility, chronic nemodialysis						

7 Dyspnea: Clinical Diag	nostic Rules & Clues			
OTTAWA COPD RISK SCALE	CATEGORY POINTS		SCORE	RISK
<b>Purpose</b> : predict death from any cause within 30	History		0	2%
days or ED visit or serious adverse event within	Coronary bypass graft	1	1	4%
14 days of ED visit (regardless of whether	Peripheral vascular disease intervention	1	2	7%
admitted): admission to critical care or acute	• Intubation for respiratory distress	2	3	13%
monitoring unit where the patient is too ill to	Examination		4	21%
ambulate, endotracheal intubation or NIV,	• Heart rate on arrival in $ED \ge 110$ /min	2	5	33%
myocardial infarction, unplanned	• Too ill to do the Walk Test* after treatment in	ı	6	48%
CABG/PCI/cardiac surgery/new nemodialysis,	ED (SaO2 < 90% or heart rate $\ge$ 120/min)	2	7	63%
for respiratory distress fever sensis) and	Investigations		8	76%
admission	• Acute ischemic changes on ECG	2	9	NA
<b>Inclusion</b> : > 50 years, COPD previously	• Pulmonary congestion evident on chest X-ray	1	10	91%
diagnosed or diagnosed in ED on the basis of 1	• Hemoglobin < 100 g/L	3	*Patient is ask	ed to walk at their
year of chronic dyspnea or cough with sputum	• Urea $\ge 12 \text{ mmol/L}$	1	own pace for 3	minutes in the ED,
production, $\geq 15$ pack year smoking history, prior	• Serum $CO2 \ge 35 \text{ mmol/L}$	1	regardless of th	he distance covered
or current evidence of moderate airflow			1	
obstruction, COPD exacerbation (increase in $\geq 2/3$	PNEUMONIA: CRB-65			
of breathlessness, sputum volume, sputum	RISK FACTOR	F	POINTS	
purulence)	Confusion of new onset	1		
<b>Exclusion</b> : too ill to be discharged: resting SpO2	• Respiratory rate $\geq$ 30 breaths / min	1		
$< 85\%$ ; heart rate $\ge 130/\text{min}$ ; SBP $< 85$ mm Hg;	• SBP < 90 mm Hg or DBP < 60 mm Hg	1		
confusion, disorientation or severe dementia,	• Age $> 65$ years	1		
STEMI on arrival: death from chronic illness	• Score = 0: outpatient therapy: I R 0.15 (0.10.0)	22)	for 30-day morts	ality
expected within weeks: arrival from a nursing	• Score = 1.2: consider hospitalization		101 JO-day monte	uity
home or chronic care facility	• Score $\sim 3$ : hospitalization: LD 4.4 (2.6.5.5) for	30 -	day mortality	
	- Score $\geq$ 5. hospitalization, LK 4.4 (5.0-5.5) 101	30-0	uay monanty	

<ul> <li>levated body temperature not caused by exogenou</li> <li>ACKGROUND</li> <li>A • Current medications? New medications?</li> </ul>	us factors. If other symptoms present (e.g PHYSICAL	. headache) see other checklists.
ACKGROUND • Current medications? New medications?	PHYSICAL	
<ul> <li>Acetaminophen usage?</li> <li>Allergies?</li> <li>Past medical history?</li> <li>Life circumstances? (e.g. travel history?)</li> <li>Alcohol: how often? How much?</li> <li>Smoking: amount? Prior smoking?</li> </ul>	Vitals• RR, SpO2%, HR, BP, Temp?Head• Meningismus?Heart• S3/S4, murmurs?Lung• Rales?Abdo• Inspection• Auscultation• Palpation	<ul> <li>CONSIDER FOR ALL:         <ol> <li>Sepsis</li> <li>Risk for contagion (e.g. influensa)</li> </ol> </li> <li>CONSIDER IF UNCLEAR:         <ol> <li>The list of causes of fever is long.</li> </ol> </li> </ul>
<ul> <li>ISTORY</li> <li>When did the fever begin?</li> <li>Degree of fever?</li> <li>Constant or intermittent? Increasing?</li> <li>Prior similar episodes?</li> </ul>	<ul> <li>Back Inspection</li> <li>Percussion tenderness over the kidneys?</li> <li>Leg Unilateral swelling?</li> <li>Skin Rash on the trunk / extremities?</li> </ul>	If the history and physical examination do not suggest a specific cause, consider the following diagnoses: 1. Pulmonary embolism 2. Cholecystitis 3. Pyelonephritis 4. Appendicitis
<ul> <li>Headache? Neck stiffness?</li> <li>Shortness of breath? Cough? Chest pain?</li> <li>Abdominal pain? Diarrhea?</li> <li>Back pain? Dysuria?</li> <li>Leg pain or swelling?</li> <li>Rash?</li> </ul>	<b>TESTS</b> • WBC (ev Diff), CRP	<ol> <li>5. Diverticulitis</li> <li>6. Infectious endocarditis</li> <li>7. Drug fever</li> <li>8. Malignancy</li> </ol>

# **8** Fever: Clinical Syndromes & Prediction Rule

### SEPSIS

Sepsis is defined as life-threatening organ dysfunction caused by a dysregulated host response to infection (Sepsis-3 definition). The clinical criteria for sepsis are the presence of both:

- Suspected or documented infection
- Acute increase in the Sequential Organ Failure Assessment (SOFA) score ≥ 2 points consequent to infection. The SOFA score assigns 0-4 points depending on the degree of dysfunction in each of six organ systems (respiration, cardiovascular, central nervous system, renal, coagulation, liver). Bilirubin, platelet count, PaO2 and creatinine are necessary to calculate the SOFA score.

# SEPTIC CHOCK

Septic shock is a subset of sepsis associated with substantially increased mortality due to profound circulatory and cellular/metabolic abnormalities. The clinical criteria for severe sepsis (associated with a hospital mortality > 40%) are the presence of both:

- Persisting hypotension requiring vasopressors to maintain MAP ≥ 65 mm Hg
- Serum lactate level > 2 mmol/L despite adequate volume resuscitation (30 ml/kg cystalloid during the first 3 hours; Dellinger 2013; 1000 ml over the first 30 min Gårdlund 2011).

**QUICK SEQUENTIAL ORGAN FAILURE ASSESSMENT (qSOFA)** The qSOFA score uses bedside clinical criteria to identify patients with suspected infection who have an increased risk of mortality or prolonged ICU admission, i.e. those with  $\geq 2$  of the following criteria:

- Respiratory rate  $\geq 22/\min$
- Systolic blood pressure  $\leq 100 \text{ mm Hg}$
- Altered mentation

The qSOFA score had similar predictive validity to the full SOFA score outside the ICU (Seymour 2016). Its purposes are to (Singer 2016):

- help identify adults with infections who are likely to have a poor outcome
- prompt consideration of possible infection if infection is not yet suspected
- prompt testing for biochemical organ dysfunction
- prompt the physician to initiate or escalate therapy
- increase the frequency of monitoring or refer to critical care

## TOXIC CHOCK SYNDROME

Toxic shock syndrome (TSS) is cause by exotoxins synthesize by Staphylococcus aureus or Group A Streptococcus (GAS). These exotoxins act as 'superantingens' and activate large numbers of T cells resulting in massive cytokine production. Staphylococcal toxic shock syndrome is associated with a variety of clinical settings, e.g. menstruation, postpartum and postsurgical states, barrier contraceptive use, staphylococcal pneumonia. The cytokines cause capillary leak and tissue damage, leading to

- Shock
- Diffuse, sunburn-like erythematous rash
- Multiorgan failure

or neck pain: see 16 Throat or NSICALNEUR5• RR, SpO2%, HR, BP, Temp?• Palpationfunction• Redness?Crania• Fundoscopy: papilledema?nerves	OLOGICAL EXAM         • Level of consciousness         al       • Orientation         • Dysphasia / dysarthria         • Visual fields & neglect
SICAL NEUR 5 • RR, SpO2%, HR, BP, Temp? cerebr • Palpation function • Redness? Crania • Fundoscopy: papilledema?	OLOGICAL EXAM• Level of consciousnessal• Orientation• Dysphasia / dysarthria• Visual fields & neglect
<ul> <li>RR, SpO2%, HR, BP, Temp?</li> <li>Palpation</li> <li>Redness?</li> <li>Fundoscopy: papilledema?</li> <li>Higher cerebr function</li> </ul>	<ul> <li>Level of consciousness</li> <li>Orientation</li> <li>Dysphasia / dysarthria</li> <li>Visual fields &amp; neglect</li> </ul>
bleeding?	<ul><li> Pupil size, reactivity</li><li> Eye movements</li><li> Excipl connection</li></ul>
steeding.	<ul><li>Facial sensation</li><li>Facial movement</li></ul>
P if > 50 years G if > 50 years Motor SIDER: arachnoid hemorrhage terial meningitis ous intracranial pathology poral arteritis Reflex Coord nation	<ul> <li>Soft palate and uvula</li> <li>Tongue movement</li> <li>Proximal and distal arm strength</li> <li>Proximal and distal leg strength</li> <li>Y Sensation touch and pinch in the distal arm</li> <li>Sensation touch and pinch in the distal leg</li> <li>Arm</li> <li>Patella</li> <li>Finger-nose</li> <li>Knee-shin</li> <li>Romberg</li> </ul>
	papilledema? bleeding? S P if > 50 years G if > 50 years Motor SIDER: varachnoid hemorrhage terial meningitis ious intracranial pathology nporal arteritis Reflex Coordination

# 9 Headache - Facial Pain: Clinical Diagnostic Rules

## **OTTAWA SUBARACHNOID HEMORRHAGE RULE**

Purpose: ruling-out SAH clinically

**Inclusion**: adults ( $\geq$  16 years); nontraumatic headache reaching max intensity within 1 hour; alert and oriented (GCS 15); no fall or direct head trauma within previous 7 days; presenting to the ED within 14 days of headache onset

**Exclusion**: new neurologic deficits (e.g. isolated cranial nerve palsies, limb weakness); papilledema on fundoscopic examination; previous diagnosis of cerebral aneurysm, SAH, brain neoplasm, or hydrocephalus; history of recurrent headaches ( $\geq 3$  episodes of the same character and intensity over the course of  $\geq 6$  months); returned for reassessment of the same headache if already investigated with both CT and lumbar puncture

Rule: investigate for SAH if  $\geq$  1 high-risk variable present:

Age ≥ 40 y
Onset during exertion
Thunderclap headache\*
Witnessed loss of consciousness
Neck pain or stiffness (subjective)
Limited neck flexion on examination\*\*

\* Instantly peaking

\*\* Inability to touch chin to chest or raise head 8 cm off bed if supine

# SUBARACHNOID HEMORRHAGE & CT HEAD

- CT head (modern, correctly interpreted) within 6 hours of onset of isolated headache (no primary neck pain, no loss of consciousness, normal neuro exam): SN 100%, LR- 0.01
- CT head beyond 6 hours from headache onset: SN 89%, LR- 0.07

# **BACTERIAL MENINGITIS**

95% of adults with community-acquired bacterial meningitis had  $\ge 2$  of the following:

Headache	<ul> <li>Neck stiffness</li> </ul>
• Fever	• Change in mental status

## SERIOUS INTRACRANIAL PATHOLOGY

Among alert (GCS 15) patients > 15 years presenting to the ED with nontraumatic headache, the presence of  $\geq$  1 of the following had a Sn 98.6%, Sp 34.4%, LR+ 1.50, LR- 0.04 for serious IC pathology:

٠	Abnormal findings on neurological examination	n	
•	Sudden onset of the headache	•	Age > 50 years

# **TEMPORAL ARTERITIS**

The presence of the following combination motivates empiric treatment with corticosteroids and temporal artery biopsy:

- New onset headache w/o alternative explanation (e.g. normal CT)
- Elevated CRP w/o alternative explanation Age > 50 years

## MIGRAINE: "POUNDing"

 $\geq$  4/5 had a LR of 24 while  $\leq$  2/5 had a LR of 0.41 for migraine:

- Nausea and vomiting
- Duration 4-72 hOurs Disabling intensity
- Unilateral location

• Pulsatile quality

# **10** Joint Pain

## Joint Pain. If pain in the lower extremity, see also 11 Leg Pain - Swelling.

### BACKGROUND

- **M** Current medications?
- Allergies? Α
- Р • Past medical history?
- Life circumstances? L
- Alcohol: how often? How much? Ε
- Smoking: amount? Prior smoking? S

# **HISTORY**

- When did the pain start? What were you doing?
  - Time till max intensity: sec? min? hr?
- Location of the pain? One or several joints? Р
  - Radiation?
- **Q** Pain? Stiffness?
- **R** Worse with movement? In such case, which?
- VAS (1-10)? Effect on daily function? S
- Constant or intermittent? Increasing? Т
  - Prior similar painful episodes?
- Fever / chills? +
  - Pain somewhere else?

# **PHYSICAL**

- Vitals RR, SpO2%, HR, BP, Temp?
- **Joint** Inspection: red, swollen?
  - Palpation: warm, tender, joint effusion?
- **CONSIDER:**
- 1. Septic arthritis
- 2. If shoulder pain: acute coronary syndrome

• Range of motion?

# **ARTHROCENTESIS**

- WBC + Neutrophils
- Culture
- Crystals
- Glucose

# **10** Joint Pain: Clinical Diagnostic Rules

## SEPTIC ARTHRITIS

**WBC COUNT:** The higher the WBC count in the synovial fluid, the more likely septic arthritis:

- WBC < 25 x  $10^{9}$ /L: LR 0.32 (0.23-0.43)
- WBC  $\ge 25 \times 10^{9}$ /L: LR 2.9 (2.5-3.4)
- WBC > 50 x  $10^{9}$ /L: LR 7.7 (5.7-11.0)
- WBC >  $100 \times 10^{9}$ /L: LR 28.0 (12-66)

However, a low WBC count can occur in early infection, and WBC >  $50 \times 10^9$ /L can occur with rheumatoid arthritis, gout and pseudogout (Adams 2009)

**PMN PERCENTAGE:** Polymorphonuclear cells count > 90% in the synovial fluid suggests septic arthritis LR+ 3.4 (2.8-4.2); LR- 0.34 (0.25-0.47)

**GLUCOSE**: Low synovial fluid glucose (defined as serum/synovial fluid glucose ratio < 0.75 and/or synovial fluid glucose < 1.5 mmmol/ml) is weakly associated with septic arthritis Sn 51% Sp 85% LR+ 3.4 (2.2-5.1); LR- 0.58 (0.44-0.76)

**LDH:** LDH > 250 U/L in the synovial fluid is sensitive but not specific for septic arthritis Sn 100%; Sp 51%; LR+ 1.9 (1.5-2.5); LR- 0.10 (0.00-1.60)

### **ACUTE PRIMARY GOUT**

(American Rheumatism Association) The presence of  $\geq$  7 of the following is required for a diagnosis of acute gout (Sn 74%, Sp 99%, +LR 74, -LR 0.26):

- More than 1 attack of acute arthritis
- Maximum inflammation developed within 1 day
- Attack of monoarthritis
- Redness observed over joints
- First metatarsophalangeal joint painful and swollen
- Unilateral attack of first metatarsophalangeal joint
- Unilateral attack of tarsal joint
- Tophus (proven or suspected)
- Hyperuricemia
- Asymmetric swelling within a joint on radiograph
- Subcortical cysts without erosions on radiograph
- Monosodium urate monohydrate microcrystals in joint fluid during attack
- Culture of joint fluid negative for organisms during attack

### **KNEE OSTEOARTHRITIS**

(American College of Rheumatology) Knee pain  $+ \ge 3$  of the following suggests osteoarthritis (Sn 95%, Sp 69%; LR+ 3.1; LR- 0.07):

- Age > 50 years
- Morning stiffness lasting < 30 min
- Crepitus on active range of motion
- Bony tenderness
- Bony enlargement
- No palpable warmth

Knee pain + osteophytes on radiograph + ≥ 1 of the following suggests osteoarthritis (Sn 91%, Sp 86%; LR+ 6.5; LR- 0.10):

- Age > 50 years
- Morning stiffness lasting < 30 min
- Crepitus on active range of motion

# **11** Leg Pain - Swelling

### Leg pain and/or swelling.

## BACKGROUND

- **M** Current medications?
  - Birth control pill? Hormones?
- **A** Allergies?
- **P** Past medical history?
  - Prior clots in the leg or lung?
- L Life circumstances?
- **E** Alcohol: how often? How much?
- **S** Smoking: amount? Prior smoking?

## **HISTORY**

- **O** When did the pain/swelling start? What were you doing?
  - Time till max intensity: sec? min? hr?
- **P** Location of the pain/swelling? Size of the area?
  - Radiation (if pain is present)?
- **Q** Pain? Swelling? Other symptoms (t ex redness, itch)?
- **R** Is the pain exacerbated by leg/foot movements?
  - Is the pain/swelling affected by position (supine, sitting)?
- **S** VAS (1-10)? Effect on daily function?
- T Constant or intermittent? Increasing?
  - Prior similar painful episodes?
- + Chest pain?
  - Shortness of breath?
  - Fever?

# PHYSICAL

Vitals • RR, SpO2%, HR, BP, Temp?

- Leg Inspection
  - Palpation

# TESTS

• CRP

# **CONSIDER:**

- 1. Deep vein thrombosis
- 2. Arterial insufficiency
- 3. Infection
- 4. Compartment syndrome
- 5. Ruptured Achilles tendon

# **11** Leg Pain - Swelling: Clinical Diagnostic Rules

# SIMPLIFIED CLINICAL MODEL FOR ASSESSMENT OF DEEP VEIN THROMBOSIS

RISK FACTORS	POINTS
• Active cancer (treated within the previous 6	1
months or currently receiving palliative	
treatment)	
<ul> <li>Paralysis, paresis, or recent plaster</li> </ul>	1
immobilization of the lower extremities	
• Recently bedridden for $\geq 3$ days or major	1
surgery within the previous 12 weeks requiring	
general or regional anesthesia	
• Localized tenderness along the distribution of	1
the deep venous system	
Entire leg swollen	1
• Calf swelling at least 3 cm larger than on the	1
asymptomatic side (measured 10 cm below the	
tibial tuberosity)	
• Pitting edema confined to the symptomatic leg	1
Collateral superficial veins (nonvaricose)	1
Previously documented deep-vein thrombosis	1
• Alternative diagnosis at least as likely as deep-	-2
vein thrombosis	

In patients with symptoms in both legs, the more symptomatic leg is used.

## **D-DIMER USE**

Wells et al evaluated the use of routine D-dimer testing in the diagnosis of deep vein thrombosis. D-dimer testing was performed with either the SimpliRED assay (Agen Biomedical) or the IL-Test (Instrumentation Laboratory). For the SimpliRED test, the result was considered negative if no agglutination was seen. For the IL-Test, the result was considered negative if the value was less than 200 µg per liter. According to their study, deep-vein thrombosis can be ruled out in the following situations:

- Score < 2 + negative d-dimer
- Score < 2 + positive d-dimer + negative ultrasound
- Score  $\geq 2$  + negative d-dimer + negative ultrasound
- Score ≥ 2 + positive d-dimer + negative ultrasound + negative repeat (+ 1 week) ultrasound

## STATENS BEREDNING FÖR MEDICINSK UTVÄRDERING

Enligt Statens Beredning för Medicinsk Utvärdering (2004) kan djup ventrombos uteslutas i följande situationer:

- Låg klinisk sannolikhet (< 2 poäng) + negativ d-dimer
- Låg klinisk sannolikhet (< 2 poäng) + negativt proximalt ultraljud
- Hög klinisk sannolikhet + negativ d-dimer + negativt proximalt ultraljud
- Hög klinisk sannolikhet +
  - negativt ultraljud, både proximalt + av underbenets vener
  - negativ flebografi
  - negativt proximalt ultraljud + negativt upprepat ultraljud +1v

<b>12</b> Neurological Deficit		
Weakness and/or paresthesia. If head trauma	, see also 17 Trauma to the Head o	or Neck
<ul> <li>BACKGROUND</li> <li>M • Current medications?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> </ul>	PHYSICAL Vitals • RR, SpO2%, HR, BP, Temp? Heart • S3/S4, murmurs? • Irregular rhythm? TESTS • EKG if > 50 years	NEUROLOGICAL EXAMCortical functions• Orientationfunctions• Dysphasia / dysarthria• Visual fields & neglectCranial nerves• Pupil size, reactivity• Eye movements• Facial sensation• Facial movement
<ul> <li>HISTORY</li> <li>O • When did the deficit start?</li> <li>• What were you doing?</li> <li>• Time till max intensity: sec? min? hr?</li> <li>P • Location of the deficit?</li> <li>Q • Weakness? Paresthesia? Both?</li> <li>S • Degree of deficit? Effect on daily function?</li> <li>T • Constant or intermittent? Increasing?</li> <li>• Prior similar episodes?</li> </ul>	<ul> <li>CRP if &gt; 50 years</li> <li>CONSIDER: <ol> <li>Stroke / TIA within 5 hours?</li> <li>Dissection (aorta, carotid, vertebrobasilar arteries)?</li> <li>Myelopathy</li> <li>Temporal Arteritis</li> </ol> </li> </ul>	<ul> <li>Soft palate and uvula</li> <li>Tongue movement</li> <li>Proximal arm strength</li> <li>Distal arm strength</li> <li>Proximal leg strength</li> <li>Distal leg strength</li> <li>Sensory</li> <li>Sensation touch hand</li> <li>Sensation pinch hand</li> <li>Sensation touch foot</li> </ul>
<ul> <li>Difficulty finding/understanding words?</li> <li>Vision problems?</li> <li>Urinary incontinence/retention?</li> <li>Pain (head, neck, chest, back)?</li> <li>Fever?</li> </ul>		<ul> <li>Sensation pinch foot</li> <li>Arm         <ul> <li>Patella</li> <li>Finger-nose</li> <li>Knee-shin</li> <li>Romberg</li> </ul> </li> </ul>

# **12** Neurological Deficit: Clinical Syndromes

### FOCAL FOREBRAIN LESION

- Unilateral weakness in the face (forehead sparing), arm and/or leg
- Dysphasia, neglect, conjugated eye deviation and/or homonymous hemianopsia are present with cortical involvement

### FOCAL BRAINSTEM and/or CEREBELLAR LESION

- Unilateral cranial nerve dysfunction (no forehead sparing)
- Contralateral weakness and/or decreased sensation with long tract involvement.

### **MYELOPATHY**

Absence of cortical and cranial nerve involvement; a sensory or motor level is present:

- **Total cord syndrome**: bilateral weakness, loss of sensation for all modalities and sphincter dysfunction
- Anterior cord syndrome: bilateral weakness and loss of sensation for pain; preserved touch
- **Posterior cord syndrome**: bilateral loss of touch; preserved strength and pain sensation
- **Central cord syndrome**: bilateralt loss of strength and pain sensation in the arms
- **Brown-Séquard**: ipsilateral weakness and loss of sensation for touch; preserved pain sensation
- **Conus medullaris/cauda equina syndromes**: leg weakness in specific myotomes; saddle anesthesia; incontinence

•	/						
RAD	RADICULOPATHY						
	Paresthesia	Weakness	Hyporeflexia				
C5	Lower lateral shoulder	Arm abduction					
<b>C6</b>	Lateral lower arm	Elbow flexion	Biceps				
<b>C7</b>	Dig 3	Elbow extension	Triceps				
<b>C8</b>	Mediala lower arm	Finger flexion					
<b>T1</b>	Medial side of elbow	Finger abduction					
L3	Medial thigh	Hip adduction					
L4	Medial calf	Knee extension	Patella				
L5	First web space (dig 1-2)	Extension of dig 1					
<b>S1</b>	Sole of the foot	Foot plantar flexion	Achilles				

### PERIPHERAL MONONEUROPATHY

Nerve	Paresthesia*	Weakness*
Axillary	Lower lateral shoulder	Arm abduction
Musculo-cutaneus	Lateral forearm	Elbow flexion
Radial	Radial aspect of the	Elbow extension & flexion
	back of the hand	Wrist & finger extension
Median	Radial aspect of the palm	Thumb opposition
Ulnar	Ulnar aspect of the hand	Finger abduction & adduction
Lateral cutaneous	Lateral thigh	None
Obturator	Medial thigh	Hip adduction
Femoral	Anterior thigh & medial calf	Knee extension
Peroneal, deep	First web space of the foot	Foot & toe dorsiflexion
Peroneal, superficial	Lateral calf and foot	Foot eversion
Tibial	Sole	Foot & toe dorsiflexion
Ischial	Lateral thigh and calf	Knee flexion
	Dorsom and sole of the foot	
* The distribution of the	deficit depends on the level of	iniurv

13 Poisoning		
Suspected poisoning. If altered consciousn	ess, see also 3. If trauma to the head	, see also 17.
Suspected poisoning. If altered consciousn         BACKGROUND         M       Current medications?         A       Allergies?         P       Past medical history? Substance abuse?         Prior poisoning / self-harm?         L       Life circumstances? Children < 18 year?         E       Alcohol: how often? How much?         S       Smoking: amount? Prior smoking?         HISTORY         What?       Which substances? Amounts?         When?       Time of intoxication?         Why?       Reason? Suicide attempt?         Now?       How do you currently feel (somatically)?         How do you currently feel (mentally / psychologically)?	<ul> <li>ess, see also 3. If trauma to the head.</li> <li>PHYSICAL</li> <li>A • Trauma to the head?</li> <li>• Tongue bite?</li> <li>B • SpO2%</li> <li>• Respiratory rate?</li> <li>• Lung auscultation?</li> <li>• Chest wall examination</li> <li>C • Pulse / blood pressure</li> <li>• Heart rate</li> <li>• QRS width, regularity?</li> <li>D • Level of consciousness?</li> <li>• Eye / pupil examination</li> <li>• Focal neurological deficits in the arms/legs?</li> <li>• Glucose?</li> <li>E • Front side of the body</li> <li>• Back side of the body</li> <li>• Temperature?</li> </ul>	<ul> <li>, see also 17.</li> <li>TESTS <ul> <li>Acid-base: pH, pCO<sub>2</sub>, HCO<sub>3</sub>/BE</li> <li>Electrolytes: Na, K, Cl, anion gap</li> <li>Paracetamol (4 hr after consumption)</li> <li>Ethanol</li> <li>Pregnancy test in fertile women</li> </ul> </li> <li>CONSIDER: <ul> <li>Toxidrome?</li> <li>Contact Poison Control Center</li> <li>Specific tests, e.g.:</li> <li>Medication levels, toxic alcohols</li> <li>Urine toxicologic screen</li> <li>CK, myoglobin (rhabdomyolysis?)</li> <li>Lever function tests</li> <li>INR</li> <li>Chronic alcohol abuse (thiamine?)</li> <li>Admission for somatic reasons?</li> <li>Residual risk for suicide / self-harm?</li> <li>Contact with social services (e.g. caretaker of child)?</li> </ul> </li> </ul>

# **13** Poisoning: Toxidromes

### **ABCDE TOXIDROMES**

		NEITHER	DRY	WET
			• Red, warm, dry skin	<ul> <li>Sweaty skin</li> </ul>
			• Dry mouth	<ul> <li>Salivation</li> </ul>
			• Dry eyes	<ul> <li>Increased tearing</li> </ul>
			• Ileus	• Diarrhea
			<ul> <li>Urinary retention</li> </ul>	• Urinary incontinence
Н	B: Tachypnea, normal $O_2\%$	Sympathomimetic	Anticholinergic	Serotonergic
Ι	C: Hypertension, tachycardia	/ Hallucinatory		
G	D: Agitation, mydriasis, seizure			
Η	E: Hyperthermia			
L	B: Bradypnea, low $O_2\%$ , bronchospasm	Sedative-	Opioid	Cholinergic
0	C: Hypotension, bradycardia	Hypnotic		_
W	D: Somnolence, miosis, hyporeflexia			
	E: Hypothermia			

### Sympathomimetic/Hallucinatory Anticholinergic Serotonergic Cocaine, amphetamines, ephedrine, Tricvclic antidepressants. Serotonin reuptake inhibitors. theophyllamine, caffeine, antihistamines, antiparkinson monoamine oxidase inhibitors. digoxin phencyclidine (PCP), ketamine, medications, phenthiazines, tricyclic antidepressives, Llysergsyredietylamid (LSD), scopolamine, muscle relaxants, white tryptophan, ecstasy (MDMA<sup>2</sup>), mescaline, psilocybin angel's trumpet, Jimson weed, deadly cocaine nightshade Sedative-Hypnotic Opioid Cholinergic Morphine, methadone, oxycodone, Bensodiazepines, zopiklon, zolpidem, Acetylcholinesterase inhibitors (e.g. alpha 2 agonists, barbiturates, ethanol, hydromorphone, buprenorphine, neostigmine, donepezil), insecticides ٠ gamma-hydroxibutansyra (GHB), loperamide, diphenoxylate, heroin, (organophosphates, carbamates), gamma-butyrolactone (GBL), certain pesticides, certain mushrooms, ٠ fentanyl butanediol (BD) organophosphorous ("nerve") gases (e.g. sarin)

### **ACID-BASE TOXIDROMES**

### **Respiratory Alkalosis**

Salicylates, theophylline, caffeine, nicotine

### **Increased Anion Gap**

Methanol, metformin, paraldehyde, phenformin, iron, isoniazid, ibuprofen, ethylene glycol, salicylates, cyanide, toluene (glue sniffing), solvents

### **Decreased Anion Gap**

Lithium, iodide, bromide (falsely low), salicylates (falsely low)

### **EKG TOXIDROMES**

AV nodal blocking Beta-blockers, verapamil, diltiazem,

### Na channel blocking (wide QRS), K channel blocking (long QTc)

- Antiarhythmics (Ia & Ic)
- Tricyclic antidepressants
- Antipsychotics
- Antihistamines
- Chloroquine

<b>14</b> Scrotal - Testicular Pain					
Pain localized to the scrotum or testicle. If concu	Pain localized to the scrotum or testicle. If concurrent abdominal pain, see also 1-Abdominal - Flank Pain.				
<ul> <li>BACKGROUND</li> <li>M • Current medications?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>L • Life circumstances? Sexual activity?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> </ul>	Vitals•RR, SpO2%, HR, BP, Temp?Buk•InspectionGenitalia•InspectionGenitalia•Inspection••Palpation••Cremaster reflex	<b>CONSIDER:</b> 1. Testicular torsion 2. Epididymitis			
<ul> <li>HISTORY</li> <li>O When did the pain start? What were you doing? <ul> <li>Time till max intensity: sec? min? hr?</li> </ul> </li> <li>P Pain location? Size of the painful area? <ul> <li>Radiation?</li> </ul> </li> <li>Q Description of pain quality</li> <li>R Worse with movement?</li> <li>S VAS (1-10)?</li> </ul> <li>T Constant or intermittent? Increasing? <ul> <li>Prior similar painful episodes?</li> </ul> </li> <li>+ Dysuria, urgency, discharge? <ul> <li>Fever / chills?</li> <li>Nausea, vomiting?</li> </ul> </li>	<b>TESTS</b> • CRP • Urine dipstick				

# 14 Scrotal - Testicular Pain: Clinical Diagnostic Rule

## **TESTICULAR TORSION**

A prospective cohort study of 228 male patients aged 0-21 years evaluated for acute ( $\leq$  72 hours) scrotal pain in the Emergency Department of an urban children's hospital reported the following features associated with testicular torsion (defined by diminished blood flow on testicular doppler US, ischaemic/infarcted testicle at operative assessment, or presence of testicular atrophy at 1- to 3-month follow-up):

- Horizontal or inguinal lie OR 18.17 (6.2-53.2)
- Nausea or vomiting OR 5.63 (2.08-15.22)
- Age 11-21 years OR 3.9 (1.27-11.97)

The authors propose the following clinical decision tool to rule out testicular torsion clinically:

- Normal testicular lie
- Lack of nausea and vomiting
- Age 0-10 years

The presence of all three criteria ruled-out testicular torsion with a sensitivity of 100% and negative predictive value of 100%. The tool has not been externally validated.

<b>15</b> Syncope - Seizure			
Transient loss of consciousness with rapid ons	set & complete recovery. If res	sidual alte	red consciousness see 3.
BACKGROUND M • Current medications?	<b>PHYSICAL</b> Vitals • RR. SpO2%, HR.	EKG Rate	<ul> <li>Tachy- bradycardia?</li> </ul>
<ul> <li>Recent additions, dosage changes?</li> </ul>	BP, Temp?	Rhythm	<ul> <li>AV block?</li> <li>Atrial fibrillation?</li> </ul>
<ul> <li>P • Past medical history?</li> </ul>	Head • Head trauma?	P	<ul><li> Left atrial hypertrophy?</li></ul>
Prior episodes with transient loss of consciousness?	• Elevated JVP?	PR Q	<ul><li>Short PR segment?</li><li>Deep, narrow in lateral leads?</li></ul>
<ul><li>L • Life circumstances?</li><li>E • Alcohol: how often? How much?</li></ul>	Leg • Swelling?	R/S	<ul><li>Signs of prior infarction?</li><li>Tall precordial R waves?</li></ul>
<b>S</b> • Smoking: amount? Prior smoking?	<b>CONSIDER:</b> 1. Syncope or seizure?	QRS	<ul><li>Bundle branch block?</li><li>Delta wave?</li></ul>
HISTORY	2. Unclear syncope: cardiogenic?	CTT.	• Epsilon wave?
<ul><li>Prior</li><li>Circumstances (activity? Standing/sitting/supine?)</li></ul>		51	<ul><li>Ischemia?</li><li>Brugada pattern?</li></ul>
<ul><li>Prodrome? Pain? Palpitations?</li><li>Transmontations and a second seco</li></ul>		T OTe	<ul><li>Ischemia?</li><li>Drolog and 2 Short?</li></ul>
<b>During (if witnessed)</b>		QIC	• Prolonged? Short?
<ul><li>Shaking?</li><li>Skin colour?</li></ul>			
<ul> <li>Duration of loss of consciousness?</li> </ul>			
<ul><li>Atter</li><li>Confusion? If so, duration?</li></ul>			
• Pain (muscular, head, chest, back, abdomen, leg)?			

# **15** Syncope - Seizure: Clinical Diagnostic Rules

# CANADIAN SYNCOPE ARRHYTHMIA RISK SCORE

**Purpose**: predict death, arthyhmia or procedural interventions to treat arrhythmias within 30 days of ED evaluation among patients for whom arrythmia and non-arrhythmic serious conditions were not identified during the ED evaluation

**Inclusion**: adults ( $\geq 16$  yr) with syncope presentin within 24 hours after the event

**Exclusion**: prolonged loss of consciousness (> 5 min), change in mental status from baseline after the syncope, obvious witnessed seizure or head trauma causing loss of consciousness, major trauma requiring hospital admission, intoxication with alcohol or illicit drugs, language barrier

CATEGORY	POINTS	SCORE	RISK
Clinical Evaluation		-2	0.2%
<ul> <li>Vasovagal predisposition*</li> </ul>	-1	-1	0.5%
<ul> <li>History of heart disease÷</li> </ul>	+1	0	0.9%
• Any ED SBP < 90 or > 180 mm Hg	g‡ +1	1	1.9%
Investigations		2	3.8%
• Troponin > 99%ile	+1	3	7.5%
• QRS duration > 130 ms	+2	4	14.3%
• QTc interval > 480 ms	+1	5	25.4%
Diagnosis in Emergency Departme	nt	6	41.1%
• ED diagnosis of vasovagal syncope	e -1	7	58.8%
• ED diagnosis of cardiac syncope	+2	8	74.5%

Score of  $\geq 0$  had SN 97% and SP 53% for death/arrhythmia/intervention within 30 days.

\*Warm-crowded place, prolonged standing, fear, emotion or pain ÷ Includes history of coronary or valvular heart disease, cardiomyopathy, congestive heart failure or non-sinus rhythm (ECG evidence during the index visit or documented history of ventricular or atrial arrhythmias, or device implantation) ‡ Includes blood pressure values from triage until ED disposition

### SYNCOPE VERSUS SEIZURE

S

QUESTIONS	POINTS
• At times do you sweat before your spells?	-2
• At times is emotional stress associated with losing consciousness?	1
• At times do you have a sense of deja vu or jamais vu before your spells?	1
• Have you ever had lightheaded spells?	-2
• Is prolonged sitting or standing associated with your spells?	-2
• Unresponsive, unusual posturing, jerking limbs during spells or no memory of spells afterwards?	1
• Has anyone ever noted your head turning during a spell?	1
• At times do you wake with a cut tongue after your spells?	2
• Has anyone ever noted that you are confused after a spall?	1

16 Throat - Neck Pain				
<ul> <li>Pain in the throat or neck. If concurren</li> <li>BACKGROUND</li> <li>M • Current medications?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> </ul>	t headache see 9 Headache - Facial Pain. PHYSICAL Vitals • RR, SpO2%, HR, BP, Temp? Throat • Redness? Swelling? Exudate? Neck • Swelling (e.g. lymph nodes)? • Tenderness?	<ul> <li>If post-traumatic see 17.</li> <li>CONSIDER: <ol> <li>Epiglottitis</li> <li>Serious infection, e.g. retropharyngeal abscess, Ludwig's angina, Lemierre's syndrome</li> <li>Dissection (carotid, vertebro-basilar)</li> </ol> </li> </ul>		
<ul> <li>S • Smoking: amount? Prior smoking??</li> <li>HISTORY</li> <li>O • When did the pain start?</li> </ul>	<b>TESTS</b> <ul> <li>CRP</li> <li>EKG if &gt; 50 years</li> </ul>	4. Acute coronary syndrome		

- Time till max intensity: sec? min? hr?
- **P** Pain location?
  - Radiation?
- **Q** Pain quality?
- **R** Worse with swallowing?
- **S** VAS (1-10)?
- **T** Constant or intermittent? Increasing?
  - Prior similar painful episodes?
- + Fever / chills?
  - Cough?
  - Trauma to the head / throat / neck?

# 16 Throat - Neck Pain: Clinical Syndromes & Decision Rule

### **EPIGLOTTITIS**

Fever + the 4 D's:

- Dypnea
- Dysphagia (odynophagia)
- Dysphonia
- Drooling

### **DEEP NECK SPACE INFECTIONS**

Description, pathophysiology.

- Peritonsillar abscess (quinsy), Parotitis
- Infection in the submandibular space (Ludwig's angina)
- Infection in the parapharyngeal space
- Infection in the retropharyngeal space
- Symptoms that may occur:
- Sore throat
- Trismus (the inability to open the jaw)
- Purulent oral discharge, pooling of saliva in the mouth, asymmetry of the oropharynx
- Lymphadenopathy is usually present.
- Dysphagia and odynophagia are secondary to inflammation of the cricoarytenoid joints.
- Dysphonia and hoarseness are late findings in neck infections and may indicate involvement of the tenth cranial nerve
- Unilateral tongue paresis indicates involvement of the twelfth cranial nerve.
- Stridor and dyspnea signify airway obstruction and may be manifestations of local pressure or spread of infection to the mediastinum.

### **MODIFIED CENTOR CRITERIA**

Criteria	Points
Temperature > 38.0	1
Tonsillar swelling or exudate	1
Swollen tender anterior cervical nodes	1
Absence of cough	1
3-14 years	1
$\geq$ 45 years	-1

Points	Likelihood of positive throat culture for Group A Streptococcal Pharyngitis
< 0	1-2.5%
1	5-10%
2	11-17%
3	28-35%
≥4	51-53%

There are different thresholds for performing a throat culture or rapid antingen-detection test (RADT):

•  $\geq 2$  points

Trauma to the head of neck. If altered	consciousness	, see also 3. If loss of conscious	sness prior to trauma, see also
<ul> <li>BACKGROUND</li> <li>M • Current medications?</li> <li>Platelet inhibitors? Anticoagulant?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> </ul>	PHYSICA Vitals Head C-spine Face	<ul> <li>RR, SpO2%, HR, BP, Temp?</li> <li>Inspection</li> <li>Palpation</li> <li>Visual acuity</li> <li>Swinging flashlight test</li> <li>Eye movements</li> <li>Palpation of the orbital rims</li> <li>Palpation of the nasal bridge</li> <li>Examination of the nasal</li> </ul>	<ul> <li>CONSIDER:</li> <li>1. Poisoning, arrhythmia, seizure, assault etc.</li> <li>2. Intracranial bleed</li> <li>3. C-spine fracture</li> <li>4. Facial fracture</li> <li>5. Admission for observation</li> </ul>
<ul> <li>Circumstances?</li> <li>Prior symptoms (e.g. palpitations?) Trauma</li> <li>Mechanism of injury?</li> <li>Loss of consciousness? After</li> <li>Amnesia (retrograde, anterograde)?</li> <li>Vomiting?</li> </ul>	Neuro	<ul> <li>Inspection of the oral cavity</li> <li>Examintion of jaw movement</li> <li>Otoscopy</li> <li>Level of consciousness</li> <li>Gross sensation and strength in the extremities</li> </ul>	
<ul> <li>Headache? Neck pain?</li> <li>Seizure?</li> <li>Paresthesia?</li> <li>Vision disturbance?</li> <li>Altered bite?</li> </ul>	<ul> <li>TESTS</li> <li>EKG if &gt; 5</li> <li>INR and the an anticoand</li> </ul>	50 years prombocytes if the patient is taking gulant	

<b>17</b> Trauma to the	Head or Neck: Clinic	al Decision Rules
SCANDINAVIAN NEUROTRAUM Applies to all adults with minimal, mi RLS 1-3) within 24 hrs of injury	IA COMMITTEE GUIDELINES ld and moderate head injury (GCS 9-15 /	<ul> <li>NEXUS LOW-RISK CRITERIA</li> <li>No cervical spine x-ray is required if all 5 are present:</li> <li>Normal level of alertness</li> <li>No evidence of intoxication</li> </ul>
GCS 9-13 / RLS 3GCS 14-15 / RLS 1-2 + any of:• posttraumatic seizures• focal neurological deficits• clinical signs of depressed or basal skull fracture• shunt-treated hydrocephalus• therapeutic anticoagulation or coagulation disordersGCS 14-15 / RLS 1-2 + both of:• age $\geq 65$ years• anti-platelet medicationGCS 14 / RLS 2 or GCS 15 / RLS 1 and any of:• suspected/confirmed loss of consciousness• repeated vomiting ( $\geq 2$ episodes)	<ul> <li>CT head and admission for observation &gt; 24 hrs</li> <li>CT head and admission for observation &gt; 24 hrs</li> <li>CT head or admission for observation ≥ 12 hours; discharge* if CT normal</li> <li>S100B if &lt; 6 hrs since injury; discharge* if &lt; 0.1 ug/L</li> <li>CT head or admission for observation ≥ 12 hrs if &gt; 6 hrs or S100B not available or S100B &gt; 0.1 ug/L : discharge* if CT normal</li> </ul>	<ul> <li>No evidence of intoxication</li> <li>No painful distracting injuries</li> <li>No focal neurologic deficit</li> <li>No posterior cervical-spine tenderness</li> <li>CANADIAN C-SPINE RULE</li> <li>No cervical spine x-ray is required if all 4 are present:</li> <li>Fulfills the inclusion criteria</li> <li>0 high risk factors</li> <li>≥ 1 low risk factor</li> <li>Able to rotate the neck actively &gt; 45° left and right</li> <li>Inclusion criteria: &gt; 15 years, no history of back or vertebral disease, normal level of consciousness, trauma &lt; 48 hrs old</li> <li>High risk factors: age ≥ 65 years, paresthesias in the extremities, dangerous mechanism of injury (fall from ≥ 1 m or 5 stairs, axial load on the head, motor vehicle collision at high speed (&gt; 100 km/h) or with rollover or ejection, a collision involving a motorized recreational vehicle, a bicycle collision)</li> </ul>
GCS 15 / RLS 1 and none of the risk factors listed above	• Discharge*	<b>Low risk factors:</b> simple rear-end motor vehicle collision, sitting position in the ED, ambulatory at any time, delayed (not immediate) onset of neck pain, absence of midline
with oral and written instructions		cervical-spine tenderness

18	8 Vertigo							
Tra	Transient or permanent illusion of motion or unsteadiness. If feeling of impending faint, see 15 Syncope - Seizure.							
BA	CKGROUND	PHYSICAL	NEUROL	OGICAL EXAM				
Μ	• Current medications?	• RR, SpO2%, HR, BP, Temp?	Higher	<ul> <li>Level of consciousness</li> </ul>				
Α	• Allergies?		cerebral	Orientation				
Р	• Past medical history?	<b>CONSIDER:</b>	functions	<ul> <li>Dysphasia / dysarthria</li> </ul>				
L	• Life circumstances?	1. Stroke, including dissection	Cranial	<ul> <li>Visual fields &amp; neglect</li> </ul>				
Ε	• Alcohol: how often? How much?	2. Vestibular neuritis	nerves	• Pupil size, reactivity				
S	<ul> <li>Smoking: amount? Prior smoking?</li> </ul>	3. Bacterial labyrinthitis		• Eye movements				
				<ul> <li>Facial sensation</li> </ul>				
HIS	TORY			Facial movement				
0	• When did vertigo start? Activity at the time?			• Soft palate and uvula				
	• Time till max intensity: sec? min? hr?			• Tongue movement				
Q	• Illusion of motion? Faintness?		Motor	• Proximal and distal arm				
R	• Worse with movement of the head?			strength				
S	• Effect on daily function?			• Proximal and distal leg				
Т	• Duration: sec, min, hr, days?		<b>C</b>	strength				
	• Prior similar episodes?		Sensory	• Sensation touch and				
+	• Diplopia?			pinch in the distal arm				
	• Dysarthria?			• Sensation touch and pipeh in the distal log				
	• Dysphagia?		Refley	• Arm				
	<ul> <li>Decreased hearing / tinnitus?</li> </ul>		KURA	Ann     Datella				
	• Decreased strength or sensation?		Coordi-	<ul> <li>Finger_nose</li> </ul>				
	• Dysmetria?		nation	<ul> <li>Knee_shin</li> </ul>				
	Headache / neck pain?			Romberg				
	• Trauma to the head / neck recently?							

# 18 Vertigo: Clinical Diagnostic Rules & Tests

### ACUTE VESTIBULAR SYNDROME (AVS)

AVS consists of dizziness with the following:

- rapid onset (over seconds to hours)
- duration  $\geq 1$  day
- nystagmus
- gait unsteadiness
- nausea/vomiting
- intolerance to head motion

### HINTS

HINTS (Horizontal head Impulse test, Nystagmus and Test of Skew) is a clinical decision rule to identify stroke among patients with AVS.

A stroke can be rule out in a patient with AVS if **all of the following** are present:

- Positive impulse test
- No change in direction of the nystagmus
- No skew deviation

A patient with AVS is likely to have a stroke if **any of the following** are present (acronym INFARCT):

- Impulse Normal
- Fast-phase Alternating
- Refixation on Cover Test

The HINTS examination has the following test characteristics for stroke: SN 98%, SP 85%, LR- 0.02.

## BENIGN PAROXYSMAL POSITIONAL VERTIGO (BPPV)

Affirmative answers to both of the following questions yielded a LR of 6.81 (5.11-9.10) for diagnosis of DHT (Dix-Hallpike test positive) + BPPV, while negative answers to both had a LR of 0.19 (0.08-0.47):

- Duration of dizziness  $\leq 15$  seconds
- Onset when turning over in bed

The **Dix-Hallpike Test** can help diagnose BPPV affecting the posterior semicircular canal. A structured critical appraisal of the litterature suggests that the Dix-Hallpike has the following test characteristics: Sn 79% (65-94); Sp 75% (33-100); LR+ 3.17 (0.58-17.50); LR- 0.28 (0.11-0.69) (Halker 2008).

The **<u>Pagnini-McClure (Head-Roll) Test</u>** can help identify BPPV affecting the horizontal (lateral) semicircular canal.

BACKGROUND	PHYSICAL	NEUROLOGICAL EXAM
<ul> <li>M • Current medications?</li> <li>A • Allergies?</li> <li>P • Past medical history?</li> <li>L • Life circumstances?</li> <li>E • Alcohol: how often? How much?</li> <li>S • Smoking: amount? Prior smoking?</li> </ul>	<ul> <li>Vitals • RR, SpO2%, HR, BP, Temp?</li> <li>Eye • Inspection of the eyelids, conjunctiva, cornea</li> <li>• Visual acuity</li> <li>• Visual fields</li> <li>• Pupil size, reaction to light</li> </ul>	Higher cerebral functionsLevel of consciousnessOrientationOrientationfunctionsDysphasia / dysarthriaCranial nervesVisual fields & neglectPupil size, reactivityEye movements
<ul> <li>HISTORY</li> <li>O • When did the vision disturbance start? What were you doing?</li> <li>• Time till max intensity: sec? min? hr?</li> </ul>	<ul> <li>Swinging flashlight test</li> <li>Fundoscopy</li> </ul> <b>TEST</b> <ul> <li>CRP if &gt; 50 years</li> </ul>	<ul> <li>Facial sensation</li> <li>Facial movement</li> <li>Soft palate and uvula</li> <li>Tongue movement</li> <li>Proximal and distal arm</li> </ul>
<ul> <li>P • Does the disturbance affect vision from one or both eyes?</li> <li>• Which part of the visual field is affected?</li> <li>Q • Description of the distrubance:</li> </ul>	<ul><li>CONSIDER:</li><li>1. Central retinal artery occlusion?</li><li>2. Stroke?</li><li>3. Temporal arteritis?</li></ul>	<ul> <li>strength</li> <li>Proximal and distal leg strength</li> <li>Sensory</li> <li>Sensation touch and pinch in the distal arm</li> </ul>
<ul> <li>decreased visual acuity, shadows, flashes, floaters, halo?</li> <li>S • Degree of deficit (e.g. ability to read, count fingers)?</li> </ul>		<ul> <li>Sensation touch and pinch in the distal leg</li> <li>Reflex</li> <li>Arm</li> <li>Patella</li> </ul>
<ul> <li>T • Constant or intermittent? Increasing?</li> <li>• Prior similar episodes?</li> <li>+ • Eye pain? Headache?</li> <li>• Fever?</li> </ul>		Coordi- nationFinger-noseNationKnee-shin• Romberg

# **19** Vision Disturbance: Clinical Diagnostic Clues

### MONOCULAR VISION DISTURBANCE

Acute monocular vision disturbance suggests a problem in the eye or the optic nerve, e.g.

- Vitreous hemorrhage
- Retinal detachment
- Temporal arteriits
- Central retinal artery occlusion
- Central retinal vein occlusion
- Optic neuritis

## **BINOCULAR VISION DISTURBANCE**

Acute binocular vision disturbance may be caused by either

- a chiasmal or post-chiasmal process
- a systemic process, e.g. temporal arteritis

Field Loss*	Terminology	Pathology
	Bitemporal (bipolar)	Midline chiasmal lesion
	hemianospia	
	Binsasal hemianospia	
	Left homonymous	Lesion affecting the right optic tract
	hemianopsia	Lesion affecting the right occipital lobe
	Left homonymous superior	Lesion affecting the lower right optic
	quadrantanopsia	radiations
	Left homonymous inferior	Lesion affecting the upper right optic
	quadrantanopsia	radiations

\* from the patient's perspective

20 Wound								
Wound. If trauma to the head or neck, see also 17 Trauma to the Head or Neck.								
BACKGROUND	PHYSICAL							
M • Current medications?	<b>Protective gear</b>	• Gloves, consider eye guard, mouth guard						
A • Allergies (e.g. to dental procedures)?	<b>Distal function</b>	• Touch (two point discrimination)?						
<b>P</b> • Past medical history?		• Motor function (specific tendon function)?						
L • Life circumstances?		• Perfusion?						
<b>E</b> • Alcohol: how often? How much?	Anesthesia	• Cleanse the skin						
<b>S</b> • Smoking: amount? Prior smoking?		• Anesthesia with lidocain +/- adrenalin						
	Inspection	• Irrigate with NaCl / tap water						
HISTORY		Hemostatic measures if needed						
When • When did the wound occur?		• Inspect for injured structures (e.g. tendons)						
What • What were you doing at the time?		Inspect for foreign material						
• Mechanism of injury?								
• Might foreign material still be present in th	e CONSIDER:							
wound?	1. Investigations to	rule-out foreign material (t ex ultrasound, X-ray)						
<b>Why</b> • Accident? Poisoning? Syncope? Assault?	2. Tetanus booster							
Self-harm?	3. Antibiotics							

# **20** Wound: Management Tips

### PRIMARY CLOSURE

Primary closure is contraindicated in the following settings:

- Wounds that are already infected
- Contamination with soil, organic matter, faeces
- Extensive tissue damage, e.g. explosion injuries, high-velocity missile injuries, complex crush injuries
- Deep or contaminated lacerations on the bottom of the foot
- Human bite wounds

Alternatives to primary closure include:

- Secondary closure (excision of the wound followed by primary closure)
- Delayed primary closure on day 4-5
- Primary healing i.e. healing by secondary intention

## **TETANUS PROPHYLAXIS**

Minimally contamined minor wound:

- Fully immunized  $\leq 10$  years since last dose: no prophylaxis
- Not fully immunized or > 10 years since last dose: tetanus toxoid

Tetanus-prone wound (contaminated or complex wound, e.g. deep puncture wound):

- Fully immunized  $\leq 5$  years since last dose: no prophylaxis
- Fully immunized 5-10 years since last dose: tetanus toxoid
- Fully immunized > 10 years since last dose OR non-fully immunized: tetanus toxoid + human tetanus immune globulin

## ANTIBIOTICS

Consider 72 hours of antibiotic treatment in the following settings:

- extremity bite wounds
- puncture-type bite wounds in any location
- intraoral lacerations that are sutured
- orocutaneous lip wounds
- wounds that cannot be cleaned or débrided satisfactorily
- highly contaminated wounds (e.g. with soil, organic matter, purulence, faeces, saliva)
- wounds involving tendons, bones, or joints
- wounds requiring extensive débridement in the operating room
- wounds in lymphedematous tissue
- distal extremity wounds when treatment is delayed for 12 to 24 hours
- patients with orthopedic prostheses
- patients at risk for the development of infective endocarditis

The choice of antibiotics depends on the cause of the wound (e.g. the species responsible for the bite) and evolving bacterial resistance.