



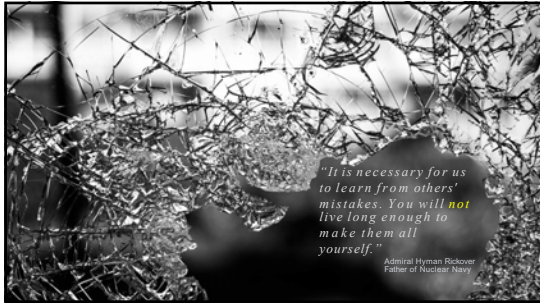
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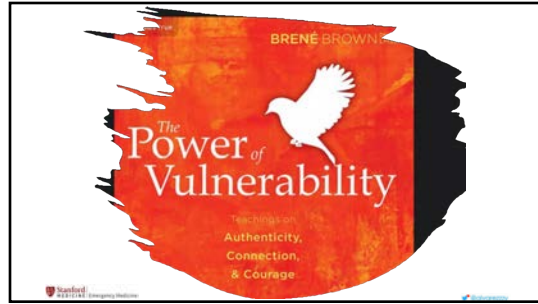
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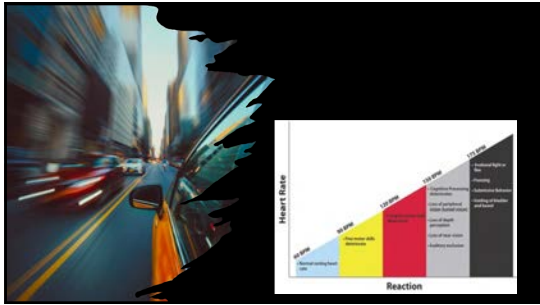
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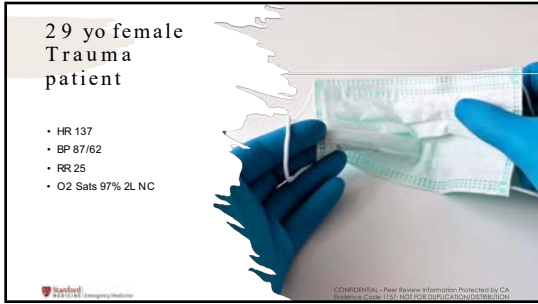
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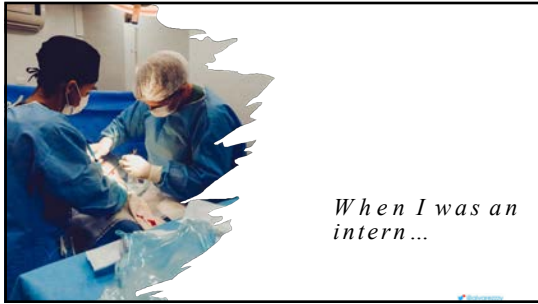
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When I was an intern...

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"After the hardest person to forgive is yourself"

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Loneliness and Isolation in Medicine

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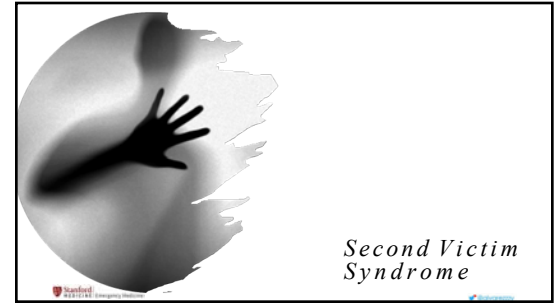
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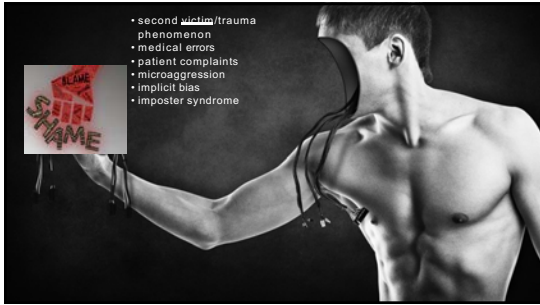
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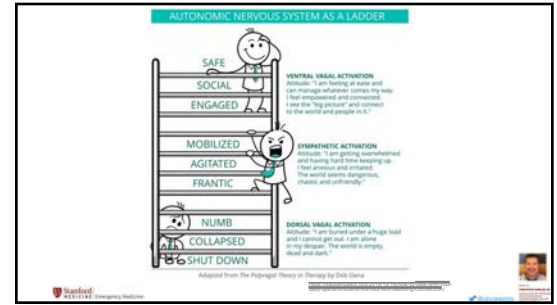
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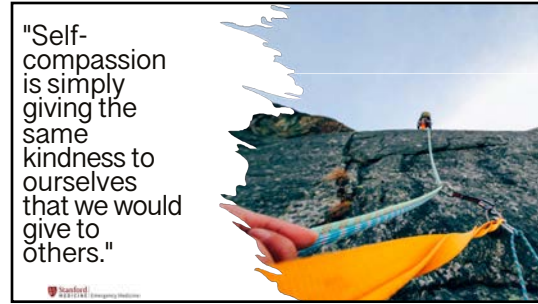
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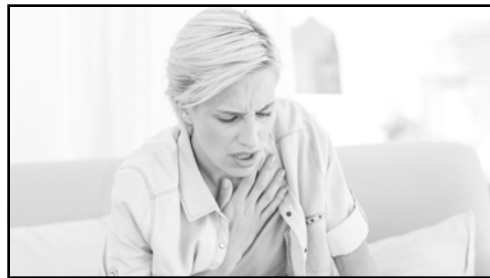
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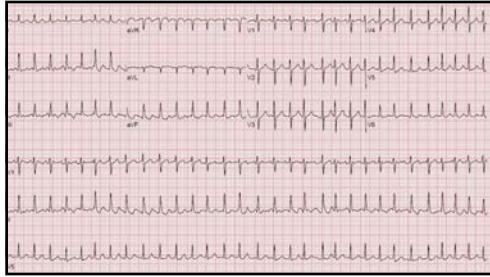
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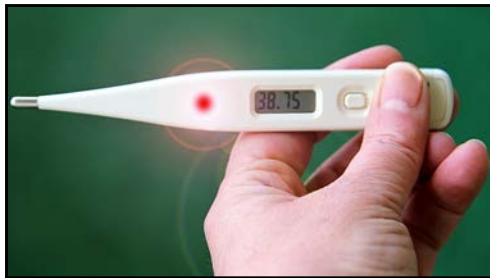
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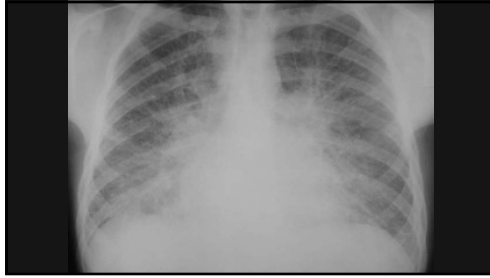
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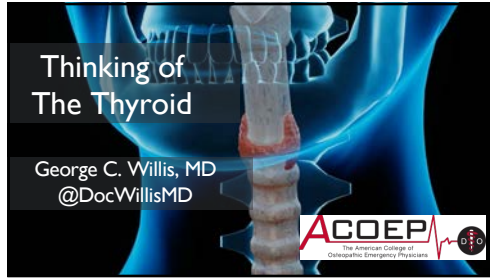
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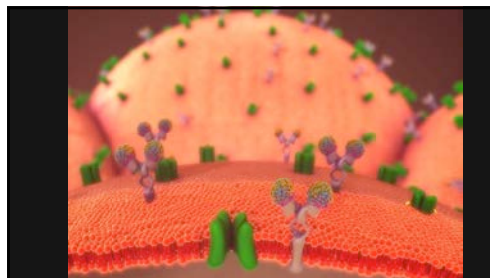
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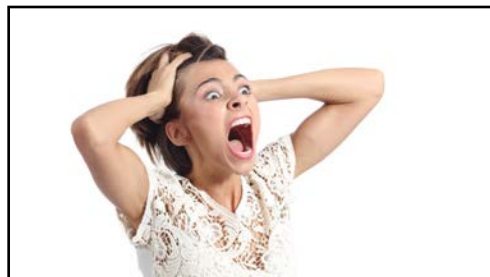
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PITFALL
Thyroid storm is often misdiagnosed!

- Sepsis
- CNS infection
- Alcohol withdrawal
- Cocaine
- Hypertensive emergency
- CHF
- Heat-related illness

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Scoring Systems?

Burch-Wartofsky scoring system

Akamizu/Japanese Thyroid Assoc. criteria

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Criteria	Points
Thermoregulatory dysfunction	
Temperature (F)	
99.0-99.9	5
100.0-100.9	10
101.0-101.9	15
102.0-102.9	20
103.0-103.9	25
≥ 104.0	30
Cardiovascular	
Tachycardia (beats per minute)	
100-109	5
110-119	10
120-129	15
130-139	20
≥ 140	25
Atrial fibrillation	
Absent	0
Present	10
Congestive heart failure	
Absent	0
Mild	5
Moderate	10
Severe	20

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Criteria	Points
Gastrointestinal-hepatic dysfunction	
Manifestation	
Absent	0
Moderate (diarrhea, abdominal pain, nausea/vomiting)	10
Severe (jaundice)	20
Central nervous system disturbance	
Manifestation	
Absent	0
Mild (agitation)	10
Moderate (delirium, psychosis, extreme lethargy)	20
Severe (seizure, coma)	30
Precedent history	
Status	
Positive	10
Negative	0

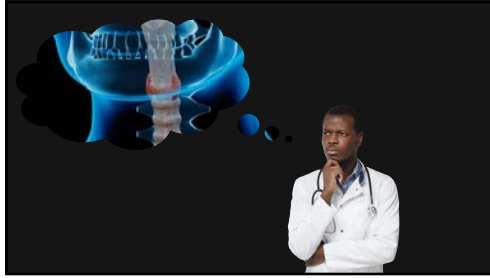
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Criterion	Burch and Wartofsky	Akamizu et al.
Thyrototoxicosis	Not included	Pre-requisite
Scoring system	Included	Not included
Fever	$\geq 37.2^\circ$	$\geq 38^\circ\text{C}$
Heart rate	≥ 90 bpm	≥ 130 bpm
Atrial fibrillation	Included	Not included
Heart failure	Pedal edema to pulmonary edema	NYHA classification class IV or Killip classification \geq III
Serum bilirubin concentrations	Not included	> 3 mg/dL
Jaundice	Included	Not included

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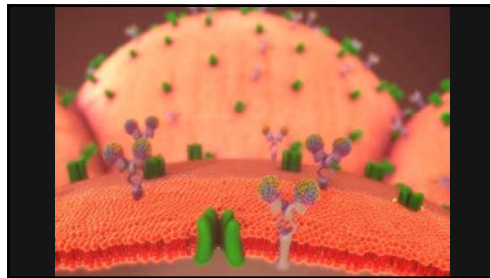
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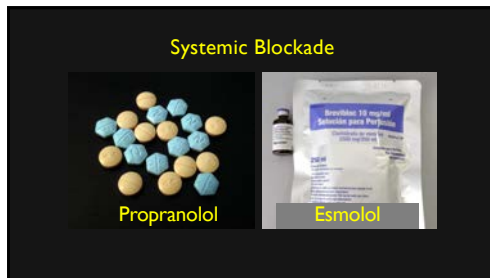
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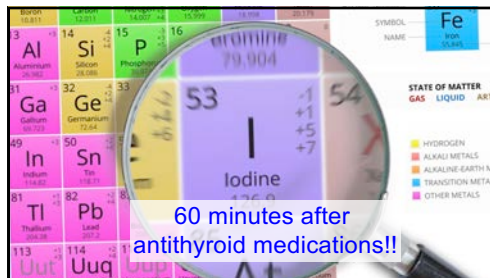
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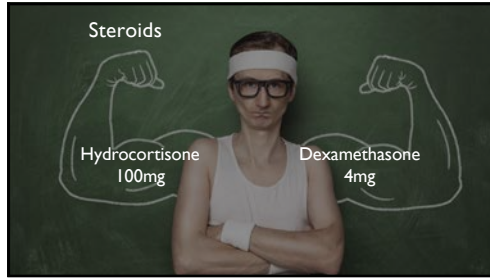
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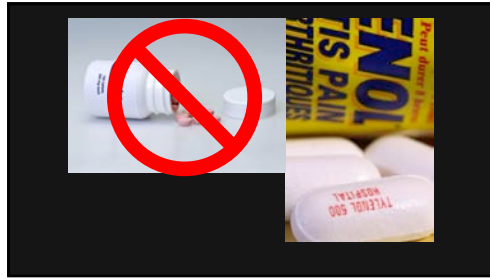
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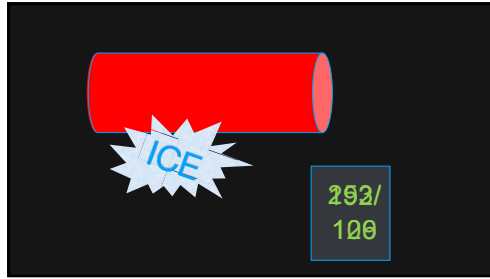
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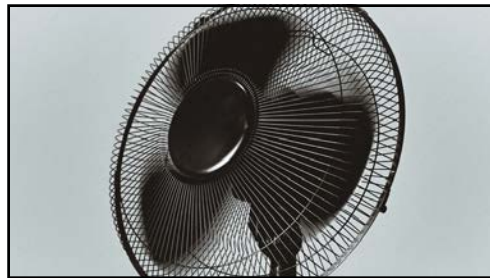
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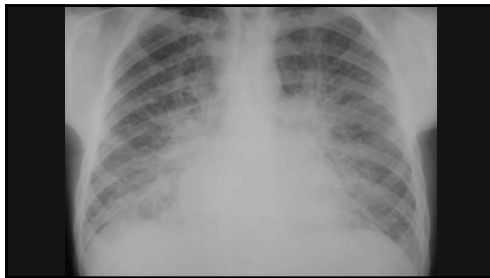
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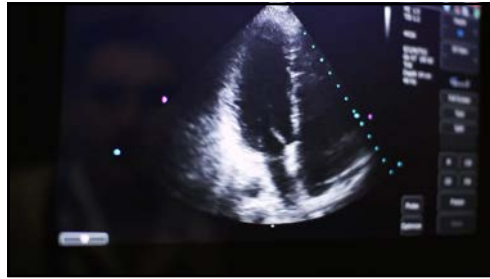
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Low Contractility	High Contractility
<ul style="list-style-type: none">• Treat like any other low output CHF• Diuretics• Afterload reduction• Preload reduction	<ul style="list-style-type: none">• Avoid preload and afterload reduction• Focus on systemic blockade• Diuretics are fine, if necessary

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Take Home Pearls

- Consider thyroid storm in patients with elevated vitals and AMS.
- Treat thyroid storm in a stepwise approach.
- Passive cooling and acetaminophen for hyperthermia. Benzos for agitation.
- Heart failure? Use your US to guide therapy.
- Don't forget to treat the stressor.

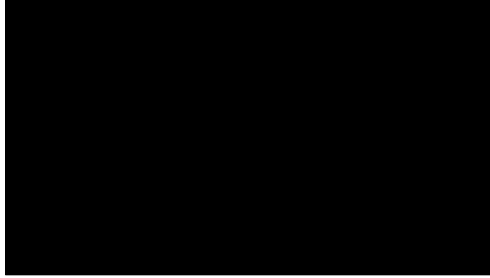
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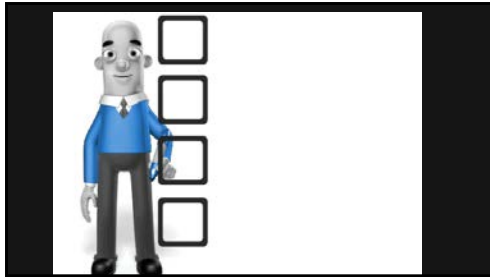
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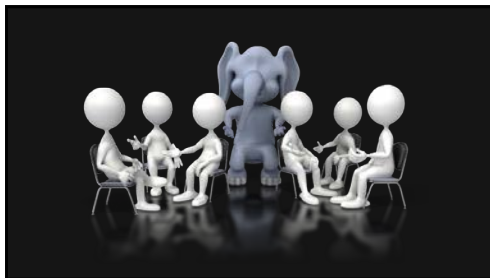
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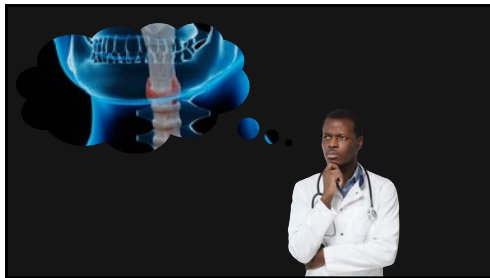
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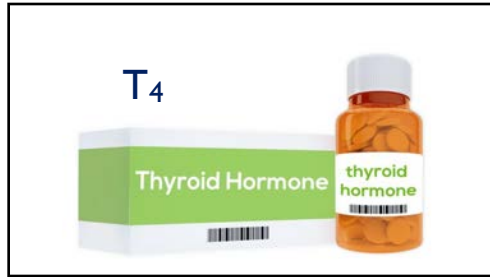
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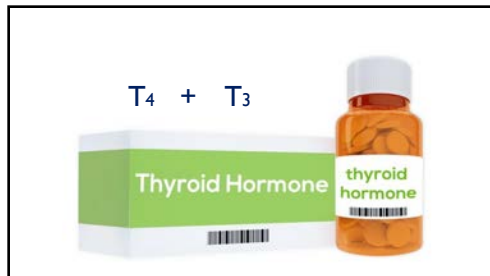
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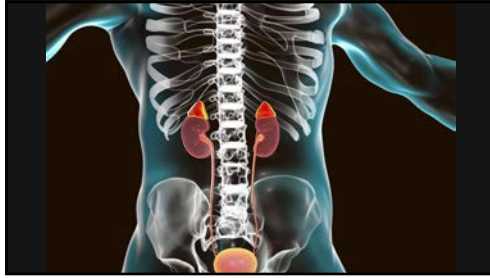
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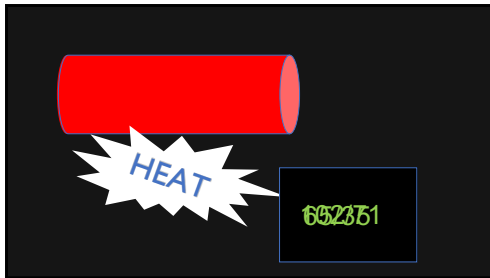
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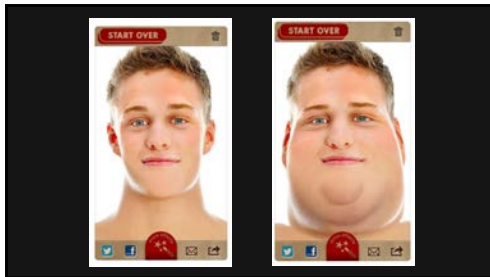
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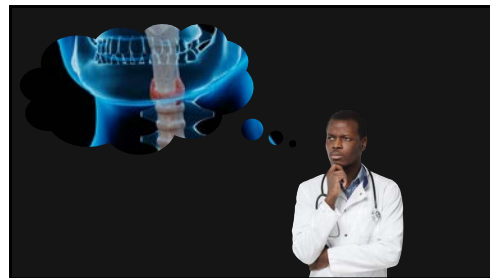


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Take Home Pearls

- Consider myxedema in patients with decreased vitals and altered sensorium.
- Consider dual thyroid replacement and administer after the stress dose steroids.
- Passive rewarming only so as to not precipitate circulatory collapse. Be careful with intubation.
- Don't forget about the stressor.

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**QRS... TUV
(MARIJUANA)**

Dan Quan, DO FAAEM FACEP FACMT
Emergency Medicine/Medical Toxicology/Addiction Medicine
Creighton University Emergency Medicine Residency
Toxicology Consultants of Arizona
Phoenix, Arizona

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DISCLOSURES

All relevant financial relationships(s) with any commercial interest to the provider's name of commercial interest(s) nature of the relationship with each

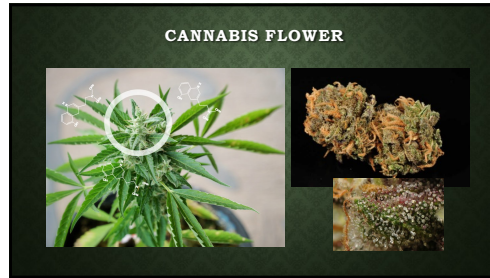
NONE

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TECHNICAL TERMS

- Cannabis
 - All products derived from the plant Cannabis sativa
- Marijuana
 - Parts of the plant that contains the most tetrahydrocannabinol (THC)
- Hemp
 - Contains little THC
- Cannabinoids
 - Chemicals found in the plant

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CANNABINOIDS

- Psychoactive
- Cannabinoid receptor 1 (CB1)
- Δ^9 -tetrahydrocannabinol or Δ^9 -THC
- Δ^8 -tetrahydrocannabinol or Δ^8 -THC
- Cannabinol
- 11-hydroxy- Δ^9 -THC
- Anandamide

- Non-psychoactive
- Cannabidiol (CBD)
- Cannabichromene
- (-) Δ^8 -THC-11-oic acid

R = H Δ^8 -Tetrahydrocannabinol (THC)
R = OH 11-Hydroxy variant (11-OH-THC)

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FDA APPROVED CANNABINOIDS

- Epidiolex
 - Purified CBD
 - Indicated for the treatment of severe seizures
- Marinol - Dronabinol (Schedule III), Syndros (Schedule II)
 - Synthetic Δ^9 THC
- Cesamet (nabilone)
 - Δ^9 THC synthetic analog
 - Used to treat nausea and vomiting caused by chemotherapy

6

CANNABINOID RECEPTORS

- Cannabinoid receptor type 1 (CB1)
 - Brain – cognition, anxiety, addiction, sleep
 - CV – negative inotropy, vasodilation, cardiac function
 - GI – mobility, secretion of gastric acid, nausea and vomiting control
- Cannabinoid receptor type 2 (CB2)
 - Immune cells, peripheral tissues, cerebellum, brain stem

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HEMP

- Fibers and oil are used to make
 - Paper, rope, textiles, plastics, animal feed, building materials
- Higher concentration of CBD vs Δ^9 THC
- Legal to grow after the 2018 Farm Bill
 - May contain up to 0.3% Δ^9 THC

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Δ^8 THC

- Derived from Cannabis sativa and mostly from hemp
- May be mixed with CBD products
- Less affinity at the CB1 receptor
- Similar "desired" effects as Δ^9 THC with less adverse effects
 - Decreases nausea and vomiting, stimulates appetite, less anxiety
 - Short-term memory problems, anxiety and paranoia, hallucinations, syncope
- Low amounts in cannabis
 - Converted from CBD using strong organic solvents
- DEA considers Δ^8 THC a Schedule I substance on 2/13/2023

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CBD

- Extracted from hemp
- Used for chronic pain and spasticity, nausea and vomiting due to chemotherapy, weight gain in HIV, sleep disorders, Tourette syndrome, seizures
- June 2018, Epidiolex was FDA approved
 - Purified CBD oral solution
- Rick Simpson's Oil (RSO)
 - www.leafly.com/news/cannabis-101/what-is-rick-simpson-oil

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NOT YOUR MAMA'S WEED

- In the 1960s and 1970s marijuana contained less than 4% of THC along with nearly the same amount of CBD
- Today, the average amount is 16% THC and lower amounts of CBD
 - Result of cross breeding, preventing pollination (plant spends time making THC rather than seeds)



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HASH OIL WITH 95% CANNABINOID, 75% Δ9-THC



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EDIBLES

- Converts Δ^9 -THC to psychoactive 11-OH-THC by the liver
 - Stronger than THC
 - Generally, 10 mg THC is for recreational use products
- Eat low and go slow
 - May take 30 minutes to 3 hours before effects show up
 - Lasts longer, peaks in 3-4 hours



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EDIBLE TOXICITY

- Drowsiness, ataxia, hypo- or hypertonia, seizures, coma, altered mental status, agitation, euphoria, mydriasis, respiratory depression
- Pediatric
 - 3 mg/kg THC - observation
 - 7 mg/kg THC - admission and moderate intervention
 - 13 mg/kg THC - ICU admission and major interventions
- Keep out of reach from children

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CANNABIS WITHDRAWAL

- Occurs in 50% of regular THC users
- May be caused by decreased CB1 receptor stimulation
- Cessation or decrease use may cause symptoms
 - Onset is 1 to 2 days
 - Peaks in 2 to 6 days and lasts up to 3 weeks
- Symptoms
 - Anxiety, irritability, aggressive, angry behavior
 - Restlessness, shakiness, insomnia, depression, anorexia
 - Sweats, headaches, diaphoresis, abdominal pain, nausea, muscle cramps

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CANNABIS WITHDRAWAL

- Withdrawal scales
 - 16-item marijuana withdrawal checklist
 - 19-item cannabis withdrawal scale
 - Not well validated
- Treatment
 - Supportive
 - Dronabinol, edible THC
 - SSRI, antipsychotics, anticonvulsants, benzodiazepines

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CANNABINOID HYPEREMESIS SYNDROME

- No risk factors other than frequent THC users
 - Any age
 - Develops with no specific timeframe
- CB1 receptors are found in enteric nerves delaying gastric emptying
- Cannabinoids accumulate in fat stores
- Genetics
 - Polymorphism in P450 may affect metabolism rate of THC
 - CYP2C9, CYP2C19, CYP3A4

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CANNABINOID HYPEREMESIS SYNDROME

- Early phase
 - Nausea, anorexia
- Hyperemetic phase
 - Persistent and painful vomiting
 - May cause electrolyte abnormalities, dehydration, acute kidney injury
- Recovery phase
 - Abstinence and symptoms resolve

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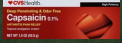
CANNABINOID HYPEREMESIS SYNDROME

- IV fluids
- Electrolyte replacement
- antiemetics
 - Ondansetron
 - Droperidol
 - 0.625 to 2.5 mg IV may help
 - Haloperidol
 - 5 mg IV, may try 5 mg PO daily prescription

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CANNABINOID HYPEREMESIS SYNDROME

- Cessation from use
- Hot showers
 - Blood flow to skin away from the enteric system
- Topical capsaicin cream 0.075% to 0.1% over the abdomen
 - Inhibits substance P and disrupts TRPV1
 - Side effects – burning sensation, skin irritation, blistering



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RAPID URINE DRUG SCREEN

- Detects 11-nor- Δ^9 -THC-9-carboxylic acid (THC-COOH)
- Positive
 - 3 to 10 days in occasional users
 - 30 to 45 days in habituated users
- Cannabidiol (CBD)
 - Does not generate THC-COOH metabolites
 - Products may contain THC (0.3% or more)
- Δ^8 -THC
 - May test positive due to cross reactivity or contamination with Δ^9 -THC
- Does NOT indicate intoxication

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METABOLISM OF CANNABIS

- Nonlinear
- Blood concentrations may drop as much as 90% in the first hour
- Cannot extrapolate backwards to determine levels
- Edibles can have a lower, gradual concentration compared to smoking

Plasma Concentration vs. Time
Smoking 3.55% THC

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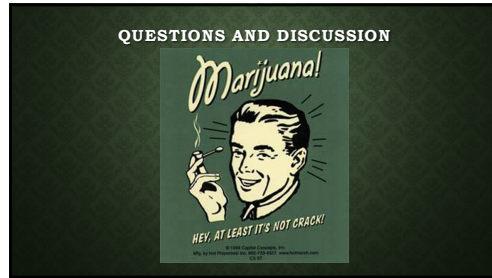
ACCURACY OF TESTING

- Affected by body habitus and type of cannabis used
- THC rapidly redistributes to fat tissue from blood
 - Smoking marijuana has a maximum plasma concentration at a mean of 8.4 minutes after the start of smoking
- Effects of THC on cognitive function do not correlate with blood concentration
 - Peak impairment occurs during the 1st hour and may last 6 hours or more
 - Chronic users may have THC levels at the limit of detection (2 ng/mL – 5 ng/mL)

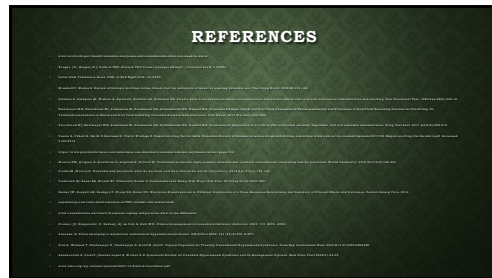
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Time Course of Standardized THC Concentration in Plasma, Performance Deficit and Subjective High after Smoking Marijuana (Adapted from Berghaus et al. 1998, Sticht and Kiferstein 1998 and Robbe 1994)

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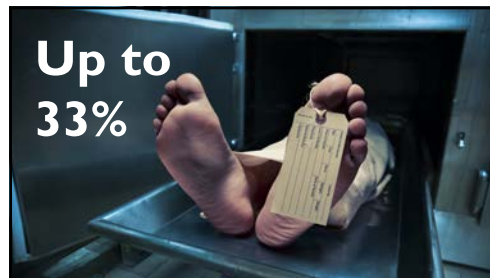
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What is syncope?

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What is syncope?

- Transient loss of consciousness
- Complete return to baseline
- No medical intervention

FAINT PASS OUT
D.F.O. BLACK
 OUT

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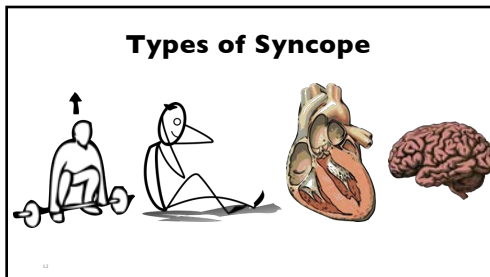
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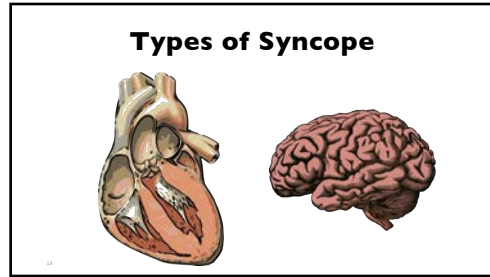
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Presyncope
=
Syncope

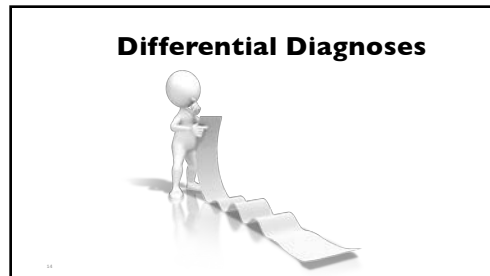
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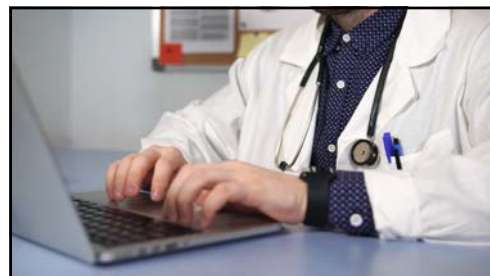
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- No or short prodrome
- Palpitations during prodrome
- Prolonged length of syncope
- Exertional
- Syncope while supine or sitting
- Worrisome symptoms
- PMH of heart disease
 - MI, CABG, Valve Dz
- Family history of SCD



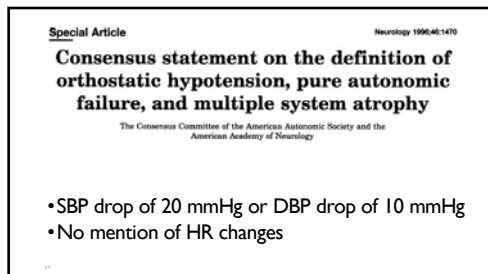
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Autonomic Neuroscience: Basic and Clinical 303 (2015) 86–90

Contents lists available at ScienceDirect

Autonomic Neuroscience: Basic and Clinical

Journal homepage: www.elsevier.com/locate/yautn

Consensus statement on the definition of orthostatic hypotension, neurally mediated syncope and the postural tachycardia syndrome

- No changes to the definition
- No mention of HR changes...again

28

been accompanied by characteristic autonomic symptoms are common, while in the elderly, typical autonomic symptoms are less frequent. Central nervous system hyperactivity, cough, and abnormal temperature control are exclusively in the elderly.

4. Postural tachycardia syndrome

4.1. Definition

The postural tachycardia syndrome (POTS) is characterized by a standing heart rate increase of ≥ 30 beats/min within 10 min of standing in the absence of orthostatic hypotension. The standing heart rate for all subjects is ≥ 120 beats/min. These criteria may not be applicable for individuals with low resting heart rates (e.g. athletes) and ≥ 25 beats/min the seated heart rate at rest. The orthostatic tachycardia may be accompanied by symptoms of central hyperadrenergic and autonomic instability that are defined by autonomic:

lack of thermoregulatory responses that include but are not limited to: chills, abnormal sweating, hyperhidrosis, anhidrosis, abnormal vasodilation, and vasoblastosis (e.g. vasodilation, flushing, and blushing).

4.2. Epidemiology and clinical features

The prevalence of POTS is unknown. The syndrome is more common in women. The syndrome's symptoms consist of light-headedness, visual blurring or tunnel vision, palpitations, lightheadedness, and weakness (especially in the legs). Other symptoms include fatigue, chronic dizziness, orthostatic intolerance, orthostatic hypotension, fainting, anxiety, chronic pain, nausea, and intolerance to heat, temperature fluctuations and heatstroke. The clinical manifestations, in addition to the heart rate increase, that present may be related and/or additive to the primary. Expert consensus may lead to revised diagnostic criteria and further research. A pathophysiological basis is present in some patients who have a mixed tachycardia, flushing, and vasodilation.

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CLINICAL STATEMENTS AND GUIDELINES

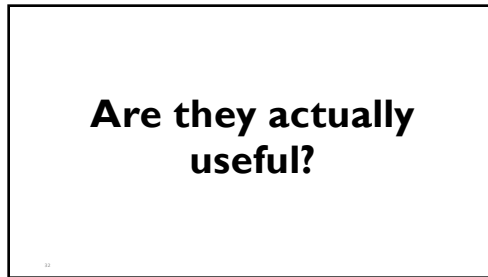
2017 ACC/AHA/HRS Guideline for the Evaluation and Management of Patients With Syncope: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Rhythm Society

- Orthostatic vital signs are recommended in every patient
- Positive orthostatic vital signs more indicative of benign etiology

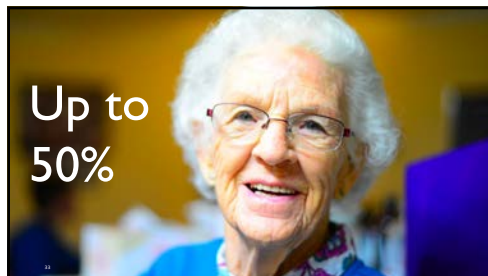
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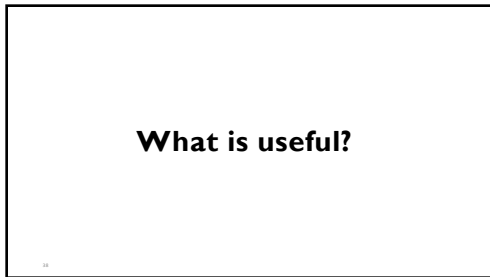
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Bottom Line

- Orthostatic vital signs do not reliably diagnose mild to moderate volume loss.
- Symptoms upon rising are more indicative of volume loss.

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“Syncope labs”

41



“if clinically indicated”

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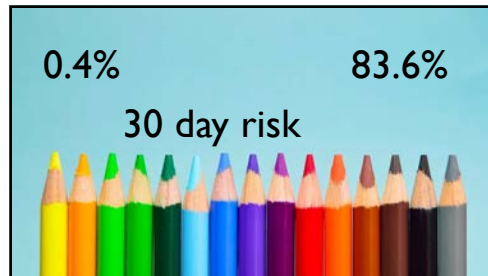


45

CSRS

Predisposition for vasovagal syncope	-1
Heart disease	1
SBP <90 or >180	2
Elevated troponin	2
Abnormal QRS axis	1
Prolonged QT interval	1
Wide QRS complexes	2
ED diagnosis of vasovagal syncope	-2
ED diagnosis of cardiac syncope	2

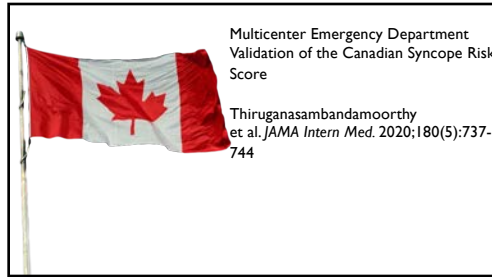
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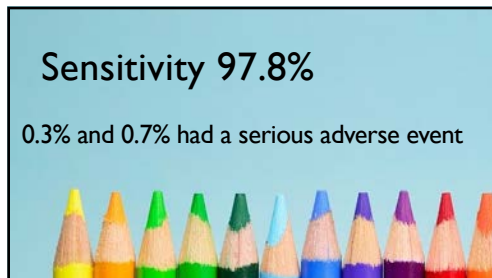
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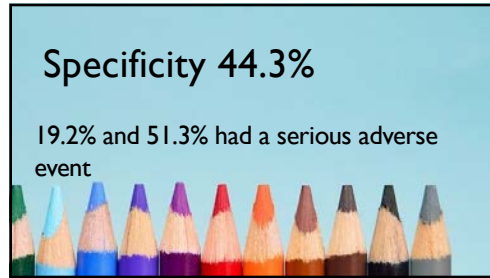
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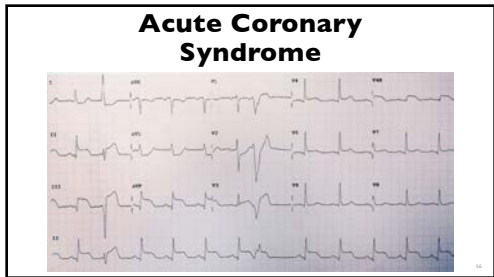
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7 things to look for on EKG!

55



56

AV Blocks

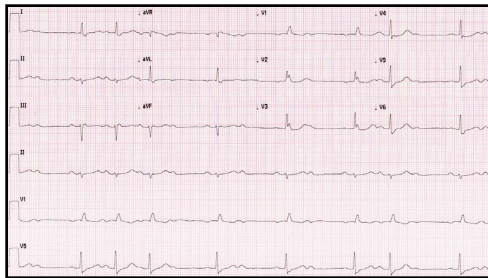
- Mostly 2nd and 3rd degree AV blocks
- Predisposes to syncope through:
 - AV dissociation
 - Sinus pause

57

2nd Degree Mobitz II

- Fixed prolonged PR interval
- Dropped QRS complex

58



2nd Degree Mobitz II


60

3rd Degree AV Block

- Complete dissociation of atria and ventricles
- P waves operate independently of QRS complexes

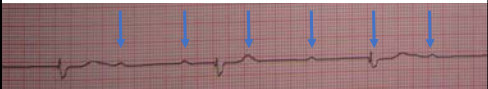
61

3rd Degree AV Block



62

3rd Degree AV Block




63

**2:1 AV
Dissociation**

- PQRS complex and then dropped QRS complex
- Think of as between 2nd and 3rd degree


64

2:1 AV Dissociation



65

2:1 AV Dissociation

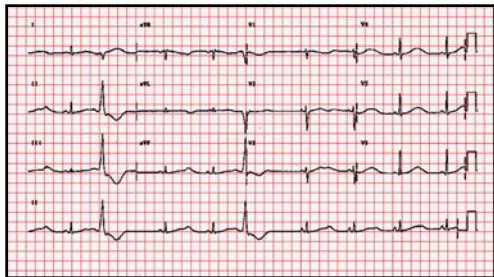


66

Prolonged QT

- QTc > 450ms
- Can be congenital or acquired
- Most commonly caused by medications
- Will progress to TdP

67



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69

Wolf-Parkinson-White

- Delta waves on ECG
- Short PR interval
- Problem=atrial fibrillation w/WPW
 - Ventricular rate of ~300
 - Wide QRS complex

70



Wolf-Parkinson-White

An ECG tracing on a grid showing multiple irregularly spaced QRS complexes. The PR interval is notably short, and the QRS complexes are wide and abnormal in morphology, characteristic of atrial fibrillation with a pre-excitation syndrome like WPW.

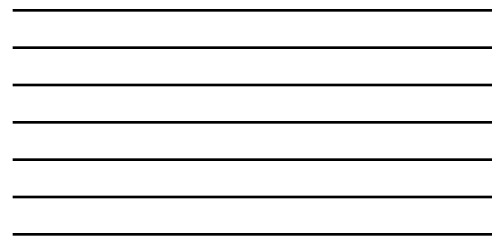
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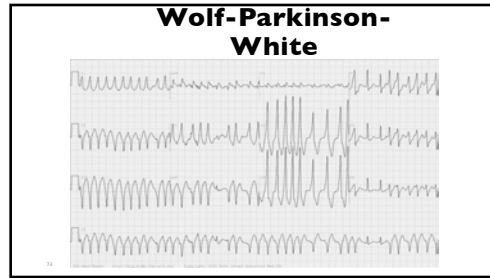


Wolf-Parkinson-White

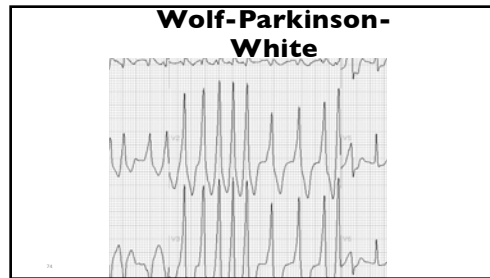
An ECG tracing on a grid showing a regular rhythm. Three blue arrows point to the initial slurred upstroke (delta wave) of the QRS complexes, which is a key diagnostic feature of WPW syndrome. The lead is labeled V4.

72





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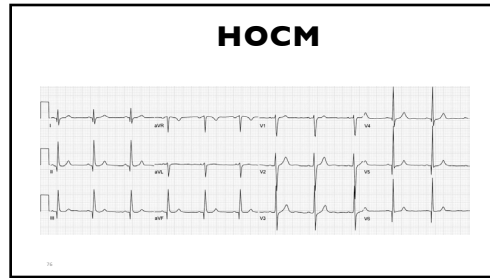
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HOCM

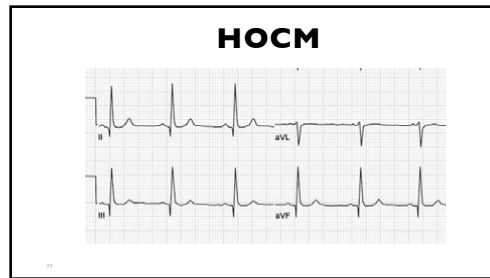
- Obstructive cardiomyopathy
- Symptoms with exertion

- LVH on EKG
- Dagger-like Q waves

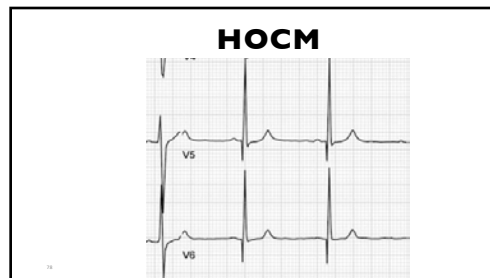
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Brugada

- Sodium channelopathy
- 3 types
- EKG findings
 - Incomplete RBBB pattern
 - ST elevation in V1 – V3
 - Coved vs. Saddle

79

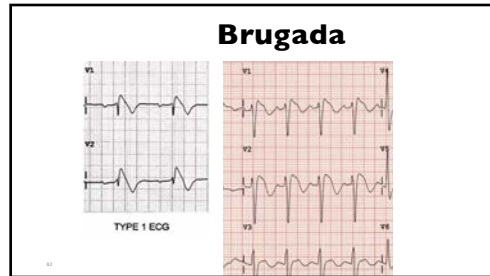
Brugada

TYPE 1 ECG TYPE 2 ECG TYPE 3 ECG

80

Brugada

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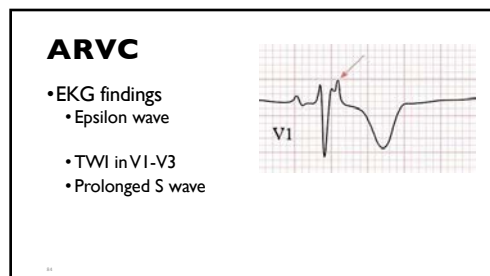


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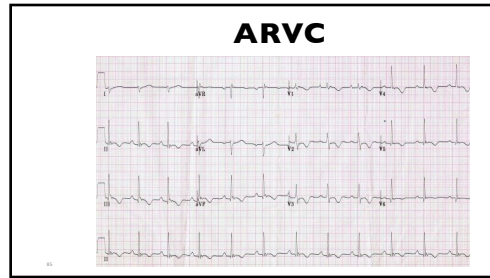
ARVC

- Arrhythmogenic right ventricular CM
- Fat deposition in the myocardium
- Causes decreased electrical conduction
- Predisposes to ventricular dysrhythmias

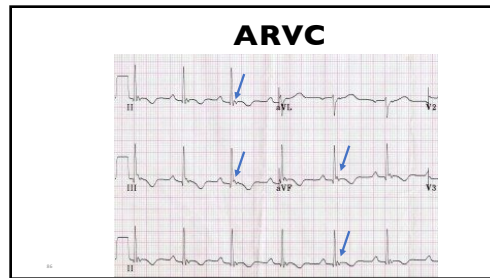
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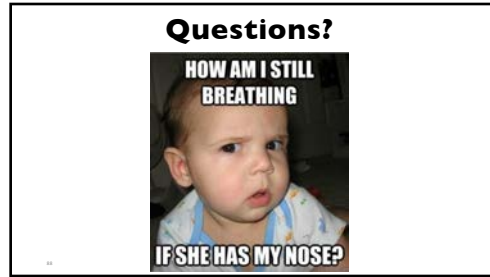
85



86

- Bottom Line**
- 7 DEADLY EKG findings in syncope
 - ACS
 - AV blocks
 - ARVC
 - Brugada
 - HOCM
 - Prolonged QT
 - WPW

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88



89

Take Home Pearls

- A good history and physical dictates the workup. There are no "syncope labs"!
- Remember the worrisome signs and symptoms for syncope. Focus on what happened surrounding the syncope.
- Orthostatics are useful for bringing on symptoms to assess for volume status. The numbers aren't helpful.
- Remember the 7 deadly EKG findings for syncope

90



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Well... For What It's Worth
*(A healthy Reassessment of What
Emergency Medicine is About)*

Presented By
Al'ai Alvarez, MD, FACEP, FAAEM
Director of Well-being
Co-Chair of the Human Potential Team
Process Improvement • Diversity/Equity and Inclusion
Design Thinking • High-Performance Team
No Disclosures



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12



Signpost Teaching

Nounou Taleghani MD, PhD
Professor, EM, Stanford

13

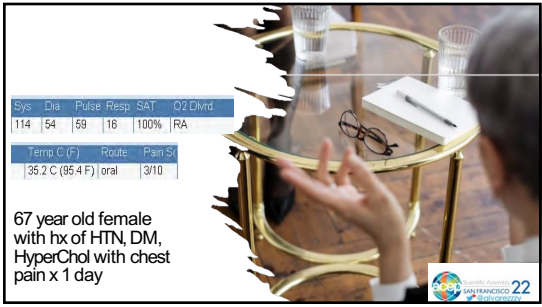


Psychological Safety

Sarah Williams, MD, MPE
Professor, EM, Stanford

PMID: 34705709

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Sys	Dia	Pulse	Resp	SAT	O2 Dlvrd
114	54	58	16	100%	RA

Temp C(F)	Route	Pain S
35.2 C (95.4 F)	oral	3/10

67 year old female with hx of HTN, DM, HyperChol with chest pain x 1 day

15



Bedside Procedures



Christine Cassidy, MD
Jacobi Medical Center

San Francisco 22

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17

Dr. Alvarez
vs.
Dr. LA



Kimberly Schertzer, MD, MS
Associate Professor
EM, Stanford



18



19

Trick: Peritonsillar abscess drainage 3.0 | All the steps with added variations


April 2018 | 100% Positive Feedback

A 22-year-old medical student comes in with a right-sided sore throat. He reports that he had a sore throat and pain on the left side of his right eye. He is currently diagnosed with peritonsillar abscess (PTA). He is currently receiving penicillin and he has not been able to eat.

Dr. Michelle Lin, founder of ALIEM, taught him how to do this procedure. He is currently and regularly using this procedure to treat PTA. He is currently using the 3.0 mm needle and he is currently using the 3.0 mm needle.


Trick of the Trade: Persistent Paracentesis Leakage 2.0


March 2018 | 100% Positive Feedback




Trick of the Trade: Angiocatheter for manual aspiration of priapism

A 25-year-old man presents with 6 hours of penile pain and swelling after recreational penile injection of Trimix (alprostadil, papaverine, and phenolamine). He denies any history of sickle cell disease or penile trauma. On exam, he is in moderate discomfort and has a tumescent penis with a soft glans. You suspect the patient is suffering from ischemic, low-flow priapism. Manual compression and ice application have been attempted with no significant improvement in the patient's clinical status.






Michelle Lin, MD
 Founder, ALIEM
 Professor, UCSF EM

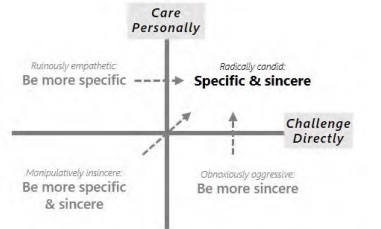


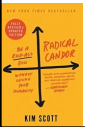
Academic Life in
Emergency Medicine

20



Barbara Fredrickson, PhD
 3:1 Positivity Ratio
 PMID: 16221001





Concept: Kim Scott 2017. www.radicalcandor.com
 Image: Taisei Yoshida, 2018

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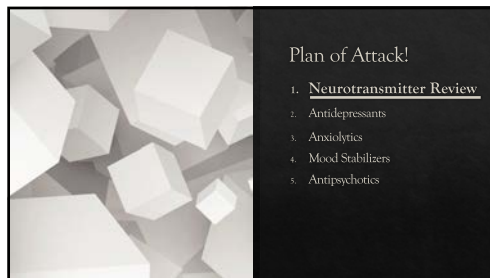
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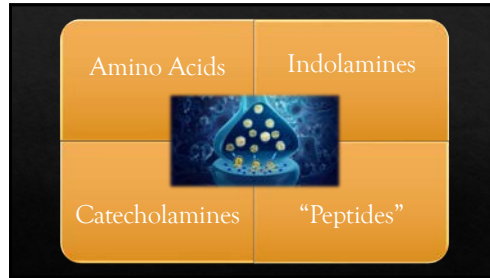
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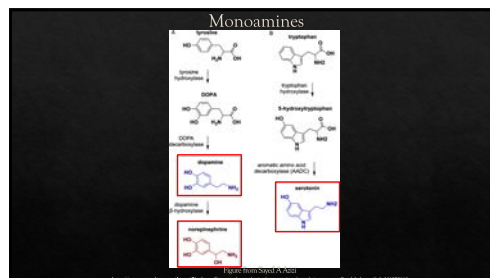
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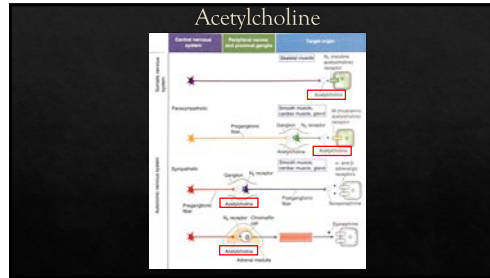
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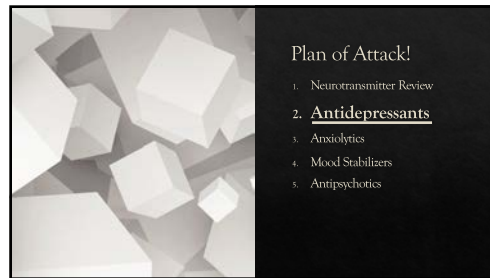
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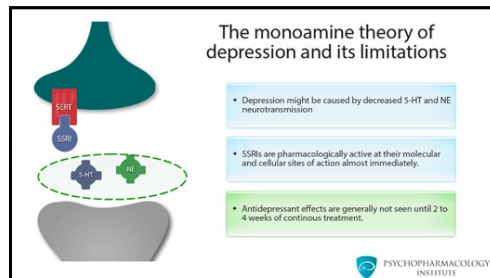
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Monoamine Oxidase Inhibitors (MAO-Is)

- ◊ Phenelzine
- ◊ Selegiline
- ◊ Isocarboxazid
- ◊ Tranylcypromine



10

Tricyclic Antidepressants

Inhibits Serotonin reuptake	Inhibits Norepinephrine reuptake
Clomipramine Imipramine	Nortriptyline Desipramine
Amitriptyline Doxepin	

- ◊ Second line: Depression, GAD, etc
- ◊ Primary uses: Chronic pain
- ◊ Side effects: anticholinergics, sodium channel blockade, "discontinuation syndrome"

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Selective Serotonin Reuptake Inhibitors

- ◊ Citalopram (Celexa)
- ◊ Escitalopram (Lexapro)
- ◊ Fluoxetine (Prozac)
- ◊ Fluvoxamine (Luvox)
- ◊ Paroxetine (Paxil)
- ◊ Sertraline (Zoloft)



First Line: Major Depression, GAD, panic disorder, OCD, PTSD, eating disorders

12

Serotonin Norepinephrine Reuptake Inhibitors (SNRIs)

- ◊ Venlafaxine (Effexor)
- ◊ Duloxetine (Cymbalta)
- ◊ Milnacipran
- ◊ Tramadol

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Mirtazapine
Serotonin and NE (?)
Often combined with SNRI

Trazodone
Orthostasis
Priapism

"Other"

Bupropion
NE + Dopamine
C.I. in seizures and
Eating disorders

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Plan of Attack!

1. Neurotransmitter Review
2. Antidepressants
3. Anxiolytics
4. Mood Stabilizers
5. Antipsychotics

15

Benzodiazepines

<p><u>Mechanism of Action</u> GABA receptor stimulation</p> <p><u>Side Effects</u> Respiratory Depression Delirium Dependency Withdrawal</p>		<p><u>Uses</u> Anxiety Acute agitation Alcohol withdrawal Seizures</p>
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Plan of Attack!

1. Neurotransmitter Review
2. Antidepressants
3. Anxiolytics
4. Mood Stabilizers
5. Antipsychotics

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Li
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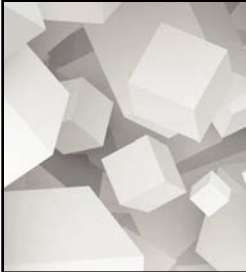
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Anticonvulsants



Hepatotoxic Pancreatitis	Danger rashes	Aplastic anemia Agranulocytosis Danger rashes
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19



Plan of Attack!

1. Neurotransmitter Review
2. Antidepressants
3. Anxiolytics
4. Mood Stabilizers
5. Antipsychotics

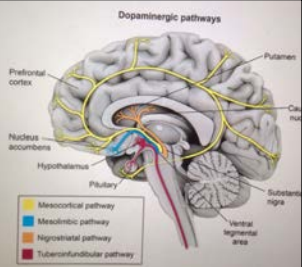
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Dopamine Theory

Meso = Midbrain
Cortical = Cortex
Limbic = Limbus system

Nigro = Substantia Nigra
Striatal = Basal Ganglia

Tubero = Hypothalamus
Infundibular = Pituitary



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Typical Antipsychotics

D2 Antagonism	Example	ADRs
Low	Thioridazine Chlorpromazine	Sedation, Postural Hypotension
High	Haloperidol Fluphenazine	Extrapyramidal symptoms

22

But why do we care???

1. First line now
2. Dirty Drugs
3. Serotonin syndrome
4. Sedation
5. Postural Hypotension

Atypical antipsychotics

Drug	D1	D2	5-HT2	α	M1	H1
Chlorazepate	++	++	+++	+++	+++	+
Risperidone	-	+++	+++	+++	-	+
Quetiapine	++	+++	+++	+++	+++	++
Olanzapine	-	++	++	+++	++	+
Ziprasidone	+/+	+++	+++	+++	++	+
Haloperidol	+	+++	++	+	-	+

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC211106/>

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ExtraPyramidal Symptoms

Signs	Description	Treat	Treatment
Acute dystonia	Prolonged periods of muscle contractions or spasms (eg. torticollis, laryngospasm, etc.)	None	<u>Anticholinergics</u> or diphenhydramine are usually the drugs of choice for acute dystonia. Anticholinergics may also be given to improve muscle relaxation during the benzodiazepines.
Dyskinesia	Paroxysmal bursts of shuffling gait, repetitive tongue.	None	Use an <u>anticholinergic</u> . Discontinuation of a dopamine agonist <u>amantadine</u> is the drug of choice for the treatment of dyskinesia.
Muscle rigidity	Subacute onset of muscle rigidity that is generalized or involving the neck.	None	Use <u>anticholinergics</u> and try <u>benzodiazepines</u> . Benzodiazepines are <u>gabapentinoids</u> that help.
Tardive dyskinesia	Intermittent, persistent, repetitive and focal movements. Usually from dopamine receptor overstimulation from chronic dopamine blockade after long-term D2R.	None	Discontinuation of the drug of choice, symptomatic treatment with more appropriate drugs, and consider changing antipsychotic drug. Anticholinergics are ineffective. <u>Drug withdrawal</u> or <u>drug withdrawal</u> may help to worsen tardive dyskinesia.
Neuroleptic malignant syndrome	Flows from rigidity, autonomic instability, elevated CPK and other abnormal laboratory tests.	Aggressive	<u>Drug withdrawal</u> provides supportive care in the ICU. Anticholinergics are <u>benzodiazepines</u> .

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Questions?

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Summary

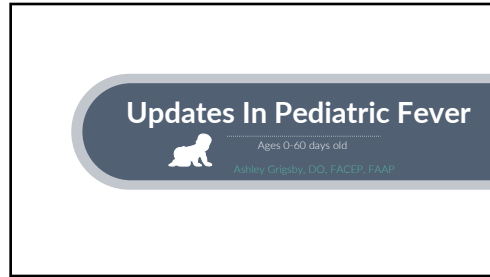
- Neurotransmitter Review
 - GABA and Glutamate
 - Monoamines
 - "Other"
- Antidepressants
 - SSRIs
 - SNRIs
 - TCAs
 - MAOIs
 - Other
- Anxiolytics
 - Benzodiazepines
- Mood Stabilizers
 - Lithium
 - Anticonvulsants
- Antipsychotics
 - First generation (typical)
 - Second generation

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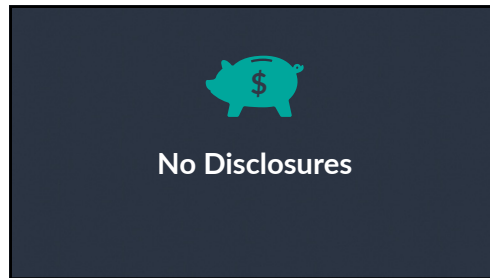
References

- ◆ Tintinalli's 8th edition
- ◆ BRS Pharmacology 6th edition
- ◆ The Human Brain An Introduction to Functional Anatomy by John N. P. Gray 6th edition
- ◆ First Aid for Step 1
- ◆ First Aid for Step 2CK
- ◆ DSM V
- ◆ Boron and Boulpaep's Medical Physiology 2nd ed

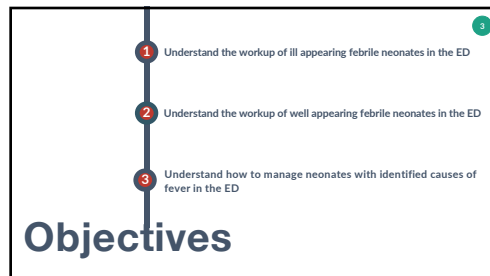
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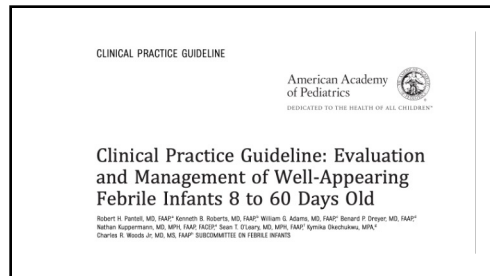
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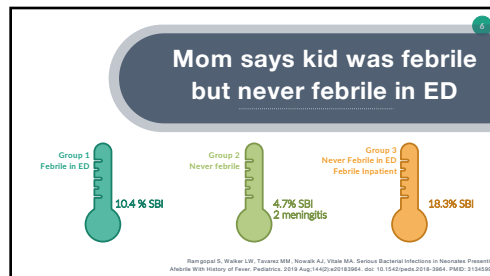
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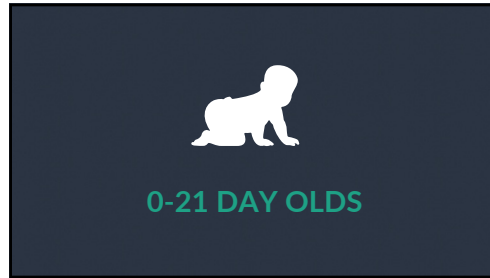
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0-21 DAYS OLD
Cookbook Medicine

BREAKDOWN	Workup	TREATMENT
Causes CSF PCR, Urinary Tract infection + TB, Bacteremia LSK Meningitis Pneumonia, sepsis, thrombocytosis	Cath urine, UA and culture Blood Cx, LFT, HSV Studies No Inflammatory Markers CSF - glucose, protein, cell count, gram stain/culture, HSV PCR, enterovirus PCR	Antibiotics Ampicillin (CSF, Urinary) Gentamicin or Carbapenem @ 10d Acyclovir @ 21 days

8

Neonatal HSV Infection
When to cover empirically?

Does History Matter?
15-30% Of neonatal HSV presented for fever alone with no other signs of HSV infection. And >75% of NHSV are acquired during delivery from newly acquired & asymptomatic mothers.

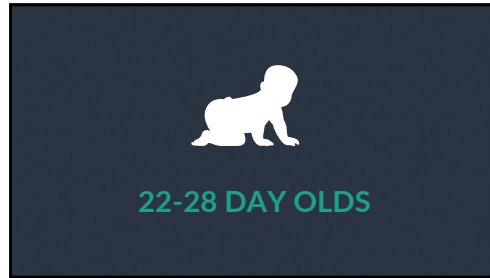
When to treat
All neonatal sepsis without any other signs of HSV
Ampicillin. Send HSV PCR on CSF!
HSV CSF PCR, HSV surface swabs, blood PCR

Highest Risk
Infants born to mothers with first-episode primary infection at time of delivery
Transmission rate 60% (no neutralizing antibodies to transmit)

>21 days
Treat any HSV if >21 days +
Maternal factors at delivery: maternal fever 48 hours before or after delivery
Infants w/ vesicles, seborrhea, hyperthermia, mucosa membrane ulcers, CSF Pleocytosis in absence of positive gram stain, leukopenia, thrombocytopenia, or elevated ALT

Allen UD, Robinson JL, Canadian Paediatric Society, Infectious Diseases and Immunization Committee. Prevention and management of neonatal herpes simplex virus infections. Paediatr Child Health. 2011;16(4):293-297. doi:10.1093/pch/nkq124

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10

What is an abnormal Inflammatory Marker (IM)?

ANC ANC significantly better than WBC ANC >4000 or <1000	CRP ≥ 20 mg/L	Procalcitonin > 0.5 ng/ml Do not use alone = 20% false infants with meningitis have normal procalcitonin	No Procalcitonin Use temp of > 38.5C (101.3F) as a positive procalcitonin
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The committee does not recommend use of abnormal inflammatory markers in LP

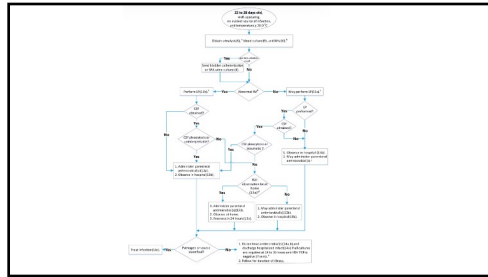
11

22-28 DAYS OLD

I thought this was cookbook?

BREAKDOWN Causes See 2019 primary care reference 10-15% bacterial Pneumonia, Sepsis, UTI, Osteomyelitis	Workup Cath urine, UA and culture Blood Cx, LFT Inflammatory Markers (IM) If LP (2-4): CRP, glucose, protein, WBC count, gram stain/culture, enterobius PCR	TREATMENT Antibiotics Use for cause of UTI Ceftriaxone 50mg/kg Meningitis = Ampicillin + Ceftriaxone Ampicillin only if no focus of infection
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LP in 22-28 day olds

Adapted to just one

Data from Step by Step study
From 2014, very similar to the new AAP guidelines

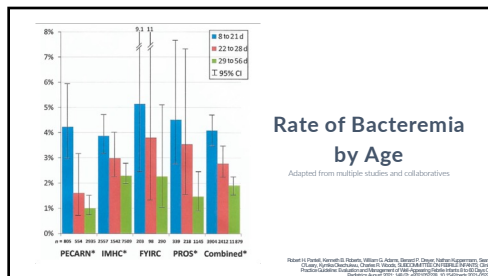
21-90 days
Study aimed to risk-stratify infants
19 prior studies included 21-28 day infants

Pathway includes
Labs: CBC, CRP, Procalcitonin
CSP and ANC

Validation Misses
7 babies were misclassified as low risk and had
IR: 4 babies non-IR (total: 1.5%)
IR: 4 or the 7 IR babies were 21-28 days

Boris Gerner, Santiago Mihalek, Shiv Bhatnagar, Linaia De Dalm, Alan Gervais, Laurence Lacroix, on behalf of the European Group for Validation of the Step-by-Step Approach. Validation of the "Step-by-Step" Approach in the Management of Young Febrile Infants. Pediatrics August 2016; 138(2): e20154381. 10.1093/peds/kpw048

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15

Can I skip the LP if the urine is dirty? 16

0-28 days

> J Pediatr. 2017 May;184:199-203. doi: 10.1016/j.peds.2017.01.022. Epub 2017 Feb 6.

Prevalence of Concomitant Acute Bacterial Meningitis in Neonates with Febrile Urinary Tract Infection: A Retrospective Cross-Sectional Study

Sowjanya S Vasbani¹, Danielle N Brown², Andrea T Cruz³

Affiliations + expand

PMID: 28186626 DOI: 10.1016/j.peds.2017.01.022

236 infants with UTI, 2 with bacterial meningitis,
1 with HSV meningoencephalitis

16

Can I skip the LP if the urine is dirty? 17

0-28 days

Comparative Study | J Pediatr Emerg Care. 1999 Dec;11(1):280-4. doi: 10.1097/00005666-199910000-00004.

Bacteremia and meningitis in neonates with urinary tract infections

NOPE!

Bachur¹, Gil-Caballero

Affiliations + expand


PMID: 8570449 DOI: 10.1097/00005666-199910000-00004

21% of \leq 28 day olds with UTI had bacteremia
4 infants <2 months had Cx positive CSF or CSF pleocytosis

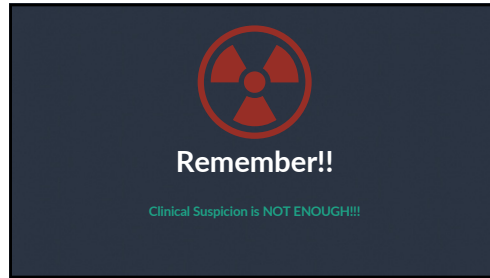
17

Age 29-60 days 18

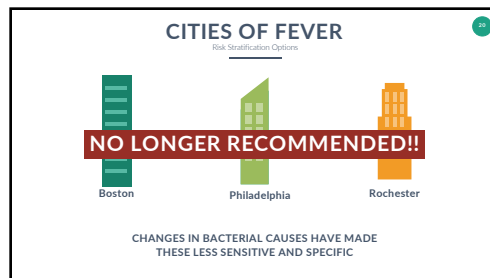
No one knows. Don't feed back.



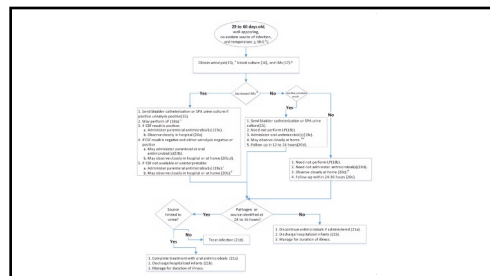
18



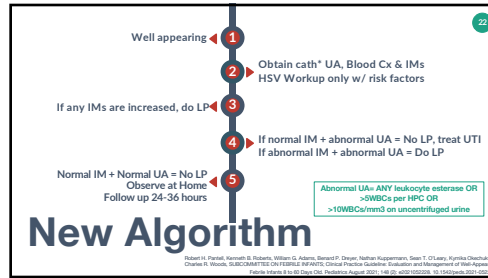
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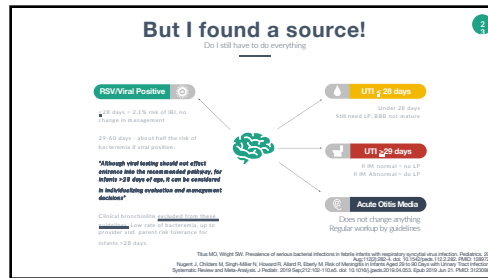
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