Teaching scripts via smartphone app facilitate resident-led teaching of medical students

Nicholas R. Zessis*¹, Amanda R. Dube², Arhanti Sadanand³, Jordan J. Cole⁴, Christine M. Hrach², and Yasmeen N. Daud²

Nicholas R. Zessis Department of Pediatrics, Northwestern University Feinberg School of Medicine 225 East Chicago Avenue, Box 152 Chicago, Illinois, USA 60611

Telephone: 312-227-7410

Fax: 312-227-9525

Email: nzessis@northwestern.edu

¹Department of Pediatrics, Northwestern University Feinberg School of Medicine, Chicago, Illinois, USA

²Department of Pediatrics, Washington University School of Medicine, Saint Louis, Missouri, USA

³Department of Pediatrics, Emory University School of Medicine, Atlanta, Georgia, USA

⁴Department of Neurology, Washington University School of Medicine, Saint Louis, Missouri, USA

^{*}Corresponding Author:

Additional file 4: Teaching Script Example: Kawasaki Disease PowerPoint © (Lecture) Format

This format was utilized by residents who desired to teach with a visual aid, which could be presented either via a smartphone or a computer monitor. The content was a hybrid of the outline format (described above) and question-and-answer cards (below), but generally was the most detailed of the three formats. The main talking point appears first at the top of the slide (i.e. "Pathophysiology"), the resident would then be able to discuss this point independently without additional information, then supplement it (if deemed necessary by the resident) with our provided teaching points by progressing the presentation to reveal additional details that would then appear at the bottom of the slide.

Kawasaki Disease

Learning Objectives

By the end of this presentation, third-year medical students will be able to:

Why do we care?

· Symptoms overlap with many other

diagnosed in a timely matter

· Significant morbidity and mortality if not

conditions

- Obtain a focused history for a child with Kawasaki Disease
- Recognize the clinical presentation
- Understand basic management

1

2

Epidemiology

- One of the most common vasculitides in childhood
- Most common in children living in east Asia or of Asian descent
- More common in boys than girls
- >90% of cases in children <5 years
- · Rare in children <6 months

-

6

3

Pathophysiology

- Systemic, inflammatory illness that affects medium-sized arteries, especially the coronary arteries
- Inflammation of multiple tissues, but longterm sequelae in the arteries
- Destruction of elastin and collagen fibers, loss of structural integrity of the arterial wall -> dilatation and aneurysm formation

Pathology

- Inflammatory cell infiltration into vascular tissues
- Destruction of luminal endothelial cells, elastic lamina, and medial smooth muscle cells in severe cases

Case

A 3 yo boy presents with 5 days of fever that does not improve with antipyretics

What other signs and symptoms would you look for?

History and Physical

with at least bur of the five following physical findings, without an

+Bilateral bulbar conjunctival injection

+Oral muscus membrane changes, including injected or fasured lips, injected pharyts, or strawberry tongue

 Peripheral extremity changes, including erythems of paints or soles, edema of hands or feet (acute phase), and perlungual desquemetion (convelescent phase)

◆Polymorphous resh

 Carnical lymphedenopathy (at least one lymph node >1.5 cm in diameter)



7

8

Initial Labs

- · Complete blood counts with differential white blood cell (WBC) counts
- · Aspartate transaminase (AST) and alanine transaminase (ALT)
- · C-reactive protein (CRP) and erythrocyte sedimentation rate (ESR)
- Urinalysis

Lab Findings

- · Elevation of acute-phase reactants (ESR, CRP)
- · Thrombocytosis that generally develops after the seventh day of liness
 - Leukocytosis, and a left-shift
- Sterile pyurfa (missed on UA because leuk esterase will be negative)
- Mild to moderate increase in transaminases

9

10

Other Possible Lab Findings

CSF may display a mononuclear pleocytosis without hypoglycorrhachia (low CSF glucose)

- · Arthrocentesis of involved joints typically demonstrates a pleocytosis
- · Hyponatremia (Increased risk of coronary artery ańeurysms)

Other than Kawasaki...

· What are some possible etiologies for his symptoms?

Differential Diagnosis

- Viral illness
- Toxic Shock Syndrome
- Rocky Mountain Spotted Fever
- Stevens-Johnson Syndrome
- Systemic Juvenile Idiopathic Arthritis

Complications

- Shock
- Cardiovascular complications
 - -Coronary artery aneurysms (1 in 5 patients)
 - -Myocardial abnormalities
 - -Infants <1 year at greatest risk</p>
- Macrophage activation syndrome, also called secondary
- hemophagocytic lymphohisticcytosis [HLH]

 –May lead to severe complications including disseminated intravascular coagulopathy, cytopenias, and thrombosis

13 14

Initial Treatment

- IVIG within the first 10 days decreases risk of coronary aneurysms (CA) 5 fold
- Decreased CA aneurysms if given IVIG with aspirin compared to aspirin alone
- · IVIG given once and patient observed to confirm resolution of fever

Referrals/Consults

- Pediatric rheumatologists
- Infectious disease specialists
- Cardiologists, and/or
- Hospitalists at a pediatric institution with experience in treating Kawasaki disease can aid in diagnosis and management

15 16

Prognosis

- · Mortality due to KD is rare among children treated with intravenous IVIG
- · Long-term morbidity is primarily related to the degree of coronary artery (CA) involvement
- Recurrence of KD is uncommon.

Follow-up

- · Monitoring for recurrence of fever
- · Repeat echocardiograms to assess for cardiac involvement

17 18

References

- Lin M.T., Wu M.H. The global epidemiology of Kawasaki disease: Review and future perspectives. Glob Cardiol Sci Pract 3 (2017). Doi: 10.21542/gcsp.2017.20
- Dietz, S.M., van Stjin, D., Burgner, D. et al. Dissecting Kawasaki disease: a state-of-the-art review. Eur J Pediatr 178, 995–1009 (2017). Doi: 10.1007/s00431-017-2937-5
- Newburger J.W., Takahasi M., Burns J.C. Kawasaki Disease. J Am Coll Cardiol 67 1738-1749 (2016).