Radiology Education: Past, Present & Future

1950's-1960's



1 talk/1 audience/5 lbs

EBD Neuhauser



1970's-1990's











2 TB flash drive



20 talks/ 1 audience/ 20

gms + 2.4 lbs (MAC)

GA Taylor

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2016+ Live streaming

Google Hangouts Messaging, Voice and Video Calls



ADOBE CONNECT

WFPI





Trends

- Local Audience → Widespread, Distant Audience
- One-time \rightarrow Enduring
- Hard-copy → Electronic
- One-size fits all \rightarrow Customizable
- Long-interval updates → Short interval (continuous) updates

2016+: Where Should we go??

- Shorter, focused content
- Based on adult learning
- Relevant to broad range of environments
- Curated
- Current
- Several languages
- Easily accessible (web-based)

Initial Strategy/ What are we looking for?

- **1.** Develop high-quality, relevant and accessible content
- 2. All areas of Pediatric Radiology/Imaging
- 3. Keynote / Powerpoint presentations, 10 to 15 minutes in duration
- 4. Basic to Advanced
- **5.** Resource rich and poor environments
- **6.** English y Español (inicialmente, otros a seguir)

2016+: How do we get there??

- Consider full engagement and support for web-based educational content
- WFPI
- E-book self-publication
- Sanjay











Disclosures

- No financial disclosures
- Self-confessed technology geek/nerd
- On various social media platforms for education
- •WFPI webmaster

Outline

- What constitutes "Digital education"
- •Why now?
- •Why WFPI?
- Challenges and Solutions
- •How do we do this?
- What do we need to accomplish this?



Why now?

- Low cost web/app platforms for content creation and consumption
- Access to web universal (...almost)
- Feedback and metrics readily available
- Wide global reach without large overhead
- Low bar to entry
- Interactive
- Readily updatable
- Generation X and Y audience demands it!

Why WFPI?

- Global reach
- Ability to break barriers
- Social media following
- Potential to be non US-centric
- Bring up issues important to non-US audiences
- Willingness to innovate
- Altruism at the core of the WFPI

Challenges and solutions

- Creating content- who, how and why?
- Reliance on a platform e.g. Facebook
- Peer review
- Curation of comments
- Content management
- Broad target audience- positives and negatives

for? What kind of content are we looking

- Anything you teach residents and fellows
- Anything new you saw or learnt on a day or during the week (e.g. my "TIL" book)
- Anything you wish someone told you when you were young!
- Summary of literature

Social media and education- mixing in the education with the "entertainment"



Opportunity is knocking...so



Creating a Community of Practice for the WFPI

Through

Leveraging + Preserving Our Present

University of Iowa College of Medicine / University of Iowa Children's Hospital Michael P. D'Alessandro, M.D.

Radiopaedia

Successful media strategy

- Radiology textbook Web site is content hub
- Social media used to promote site cases / topics + drive users to Web



WFPI Textbook of Pediatric Imaging

Successful media strategy

- Pediatric radiology textbook Web site is content hub
- Social media used to promote site cases / topics + drive users to Web



n most cases the outcome of a fetus with CPAM is very good. In rare cases in every 30,000 pregnancies.[1] sue. The underlying cause for CPAM is unknown. It occurs in approximately ually an entire lobe of lung is replaced by a non-working cystic piece of normal lung tissue. This abnormal tissue will never function as normal lung der of the lung similar to bronchopulmonary sequestration. In CPAM

the cystic mass grows so large as to limit the growth of the surrounding lung and cause pressure against the hand. In these estimations, the CPAM can be submitted and the fault of the faults. CPAM type 1 faits the most common, with clinical and pathologic features.^{[21} CPAM type 1 is the most common, with large cysts and a good prognost. CPAM type 2 (with medium-sized cysts) often has a poor prognosis, owing to its frequent association with other significant anomalies. Other types are rare $^{(3)}$

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Diagnosis and treatment criteria [odt]

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Deutsch

Polski

CPAMs are often identified during routine prenatal ultrasonography. Identifying everted (pushed downward) diaphragm, or the absence of visible lung tissue he chest of the fetus, displace stics on the sonogram include: an echogenic (bright) mass appearing in ment of the heart from its normal position, a flat or

CPAMs are classified into three different types based largely on their gross appearance. Type I has a large (>2 cm) multiloculated cysts. Type II has smaller uniform cysts. Type III is not grossily cystic, referred to as the "adenomator" type Microscopically, the lesions are not rule cysts, but communicate with the Microscopically. surrounding parenchyma. Some lesions have an abnormal connection to a blood vessel from an aorta and are referred to as "hybrid lesions."

Imaging [odit]

The earliest point at which a CPAM can be detected is by prenatal ultrasound. The issic description is of an ech isic description is of an echogenic lung mass that gradually disappears over sequent ultrasounds. The disappearance is due to the malformation becoming filled with fluid over the course of the CPAM on chest radiograph in a our newborn. Large cystic changes in the left lung, leading to a mediastinal shift to the right due to their mass effect.

gestation, allowing the utrasound waves to penetrate it more easily and rendering it invisible on norographic importance. When a CPAM is targetly growing, either solid or with a dominant royst. They have a higher incidence of developing venous outflow obstruction, cardiac failure and utimately *hydrops fetalis*. If *hydrops* is not present, the fetus has a 95% chance of outflow obstruction, cardiac failure and utimately *hydrops fetalis*. If *hydrops* is not present, the fetus has a 95% chance of outflow obstruction. survival. If it is seen, the fetus will die without *in utero* surgery, or delivery if it development after 32 weeks. The greatest period of growth is during the end of the second trimester, between 20-26 weeks.

has been developed to predict the risk of lydrops. The lung mass volume is determined using the formula (length : width × anteroposterior diameter + 2), divided by head circumference. With a CVR greater than 1.6 being considered high risk. Fetuses with a CVR least than 1.6 and without a dominant cyst have less than a 3% risk of lydrops. After delarey, if the I measure of mass volume divided by head circumference, termed cystic adenomatoid malformation volume ratio (CVR

Where Do You Get the Content?

- Educational posters of meetings from 13 member societies
- Submitted in MediaWiki format (rather than PowerPoint)
- Peer reviewed by educational poster session reviewers
- **Edited by authors**
- arter Published online for meeting + then forever
- Readers send corrections + comments to authors
- Editing restricted to authors + site editors
- Content serves as basis of next year's social media campaign



Diagnosis and treatment criteria [odit]

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Advantages

- Authors get educational poster + textbook chapter in CV
- Receive impact of chapter in social media measured by ImpactStory **Receive altmetrics yearly for chapter (pages read, users)**
- Great project for
- Senior radiologists to select poster / chapter topics + oversee
- Junior radiologists to write chapters, do tech + social media
- Users free textbook of pediatric imaging
- Start small partner at first with one member society
- Could be multilingual as you partner with more societies
- Cost almost nothing
- Takes advantage of pre-existing content workflow in form of educational posters + preserves it
- **Result is tangible permanent pediatric imaging reference**
- Build community of practice of members (authors/editors) around it