# Impact of Continuing Medical Education on Primary Care Providers' Knowledge and Confidence in Caring for Patients with Congenital Hypothyroidism

Emily Bezar, MA<sup>1</sup>, Ning Rosenthal, MD, PhD<sup>2</sup>, Lisa Feuchtbaum Dr.PH, MPH<sup>2</sup>

<sup>1</sup> Public Health Foundation Enterprises <sup>2</sup> Genetic Disease Screening Program,
California Department of Public Health

Supported by HRSA Grant # D93MC26187: "Demonstration projects for integrating newborn screening long term follow-up into primary care practice"

## The PCH Project

Long Term Follow-Up of Patients with Primary Congenital Hypothyroidism (PCH) by Primary Care Providers (PCPs)

- A three-year grant funded by the Health Resources and Services Administration (HRSA) from August 1<sup>st</sup>, 2013 through July 31<sup>st</sup>, 2016
- Participating states: California and Hawaii
- Target populations: PCPs and their patients with PCH

## **PCH Project: Overall Objectives**

- 1) Assess the willingness and capability of PCPs to provide long-term care for patients with PCH and their needs for PCH-related continuing medical education
- 2) Evaluate the current case management patterns and clinical outcomes
- 3) **Determine** PCPs' willingness to obtain informed consent and provide data to the PCH long-term follow-up (LTFU) database
- 4) Investigate the practicality of providing real time LTFU data by PCPs and identify barriers incurred
- 5) Improve PCPs' knowledge about PCH and increase their capability of providing care for patients with PCH

## Why develop a CME course for PCPs?

## **Clinical Knowledge Gaps**

Our 2014 cross-sectional survey (N=226) revealed gaps in PCPs' knowledge on treating patients with PCH:

- Only 49% knew the recommended frequency of blood tests to monitor PCH over a patient's lifespan
- Only 23% knew when to try a patient off levothyroxine treatment to determine if PCH is transient

### **Interest**

 84% of surveyed PCPs reported that they are likely or very likely to participate in CME on PCH if available

### **High incidence rate:**

PCH is the most common disorder identified in blood spot screening and affects 1 in 1,706 live births in CA

## Geographic distribution of cases:

Need to increase number of PCPs
who can provide PCH case-management
in rural areas of CA with very
few pediatric endocrinologists

## More good reasons for a PCH course....

### Relatively simple to treat:

Patients with PCH can potentially be managed well by PCPs with support from endocrinologists

## To improve access to quality care:

CME for PCPs may reduce barriers to quality follow-up care for families for whom specialty care is difficult to obtain

### Congenital Hypothyroidism

## What Every Pediatrician Needs to Know

Swati Banerjee, M.D.

Division of Pediatric Endocrinology Valley Children's Specialty Medical Group

#### Goals

By the end of this session, you will be able to:

- Initiate retesting and treatment if newborn thyroid function screening is abnormal
- Monitor treatment
   of primary congenital hypothyroidism (CH) for
   infants, children, and adolescents
- Educate families
   about the importance of adherence to treatment

Course was developed with the PCH Project's Advisory Committee of CA pediatric endocrinologists

Continuing Medical Education (CME) can be a key element in a PCP-centered follow-up model



## **Course Format and Content Summary**

- 50 minute lecture, followed by a 10 minute Q & A
- Begin by presenting 5 Case Studies in question format
- Review of California's PCH newborn screening methodology
- Review of thyroid pathophysiology and clinical presentation of PCH symptoms
- Focus on the functional priorities of PCH diagnosis,
   treatment and follow-up in the pediatric primary care context
- Recommend consultation with endocrinologist when needed
- Conclude with the same 5 Case Studies, with opportunity for audience self-assessment and discussion

## **Learning Objectives**

### At the end of this course, participants will be able to:

- Perform confirmatory testing and initiate treatment if newborn thyroid function screening is abnormal
- Assess treatment needs and monitor clinical outcomes for infants, children, and adolescents with congenital hypothyroidism
- Describe the diagnostic process for determining whether a patient has transient or permanent hypothyroidism
- Educate families about the importance of adherence to treatment for congenital hypothyroidism

#### **5 Handouts** CALIFORNIA CHILDREN'S SERVICES APPROVED ENDOCRINOLOGY CENTERS Medical Center Parents' Guide 3CDPH 5 Treating Congenital Hypothyroidism (CH) Congenital Hypothyroidism Course Evaluation 2015 Quick Post-Course 1. Please indicate how confident you are with the following tasks related to the After Receiving Positive Newborn Screening (NBS) Results After Receiving Processing a positive NBS result for CM Within 24 hours of receiving a positive NBS result for CM management of patients with confirmed congenital hypothyroidism (CH) with Document ID: 100 1="Not confident at all"; 2="Somewhat confident", 3="Confident"; 4="Very confident". After Receiving Positive Newborn Screening (1) After Receiving Positive Newborn Screening (1) After A hours of receiving a positive last seal for cool 14) After 29 AND mod seasons Test and free 14 for cool 14) For confirmation diagnosis 1a. Confirm diagnosis for patients with positive newborn Circle the Score (1-4) b. Initiate treatment for patients with confirmed CH · 210 HUM Considered normal Considered normal 1......2......3......4 Howdoon screening Tel result is: nduct follow-up for patients with confirmed CH 134 2 14 Livinit, text at for confirmatory diagnosis 10 40 HU Int. T4 for social TA) and TSH statistics from the free Taylor throughout treatment yet but do not start to order to order to be to be to order to Approximate daily dose by weight: 1......3......4 of confirmation sections. Weight (grants) | Daily Dose heck TRUE or FALSE for each of the following statements 1......3......4 Intrace treatment ample a social and refer to a social and and refer to a social and a social a 37.5 HB is collected and refer to a pediatric endocrinologist Statements 50 HE hormone (TSH) surges by 30 minutes after 2500 - 3999 10 μIU/mL within 3 days True or False Levo-thyroxine Treatment Dosing d CH cases are permanent and need Recommended to Now up schedule by 3ge. □ True □ False or book 74 5 188 dt. or serverte bue ray with a serverte bee faute for serverte bee for yet for the form of possible for the form of the f Secret are 19 12 Higher 14 C 05 mg/dl DO NOT USE LIQUID FORM ent should be initiated as soon as a Start at 10 - 15 Helve □ True flected for confirmatory testing □ False mmended initial treatment dose of levothyroxine Promotion 1544 contemporation in the Adecorporation in the Promotion from TA contemporation from the Adecorporation from the Adecorporation from the Promotion from the Adecorporation from the Adecor with confirmed CH is 15-20 µg/kg □ True Diff and returns to 210 HUlmit in □ False \* Maintain free TA (or total TA) concentration in Jupes half of the age-appoints reference have 30. The majority of confirmed CH cases are 36. For severe comments in any and the spould be initiated as 50 For severe CH, treatment should be initiated as \$00. □ True □ False 2 weeks after treatment introduce. Follow-up Frequency Lweeks after treatment intrateon. The first dinks to down of earth

### Treating Congenital Hypothyroidism (CH)

## 2015 Quick Reference

#### After Receiving Positive Newborn Screening (NBS) Results

- Within 24 hours of receiving a positive NBS result for CH (TSH ≥ 29 µIU/mL), test serum TSH and free T4 (or total T4) for confirmatory diagnosis
- If newborn screening TSH result is:

> 40 μIU/mL	29 — 40 μIU/mL
Initiate treatment as soon as a serum sample is collected and refer to a pediatric endocrinologist	May wait for the results of confirmatory serum test to initiate treatment

You are strongly encouraged to work closely with a pediatric endocrinologist to perform diagnostic evaluation, initiate treatment, and coordinate for ongoing care.

Treatment must begin within 2 weeks of age for confirmed cases.

#### If confirmatory serum TSH result is:

- > 40 μIU/mL Initiate levo-thyroxine treatment immediately
- 10 40 μIU/mL
   Repeat the free T4 (or total T4) and TSH tests
   but do not start levo-thyroxine treatment yet
- < 10 µIU/mL Considered normal, no treatment needed

#### Levo-thyroxine Treatment Dosing

Start at 10 – 15 µg/kg

(Use 15  $\mu$ g/kg if free T4 < 0.5 ng/dL or total T4 < 5  $\mu$ g/dL)

- Either brand name or generic, but stay with the same formulation, if possible. USE TABLETS, DO NOT USE LIQUID FORM
- Maintain TSH concentration in the age-specific reference range
- Maintain free T4 (or total T4) concentration in the upper half of the age-specific reference range

#### Approximate daily dose by weight:

Weight (grams)	Daily Dose
2000 – 2499	25 µg
2500 – 3999	37.5 μg
4000 or more	50 µg

#### Follow-up Frequency

2 weeks after treatment initiation:

The first clinical follow-up examination and lab tests should take place

#### Recommended follow-up schedule by age:

Age	Frequency
< 6 months	Every I-2 months

## **PCPs and PCH?? CME Evaluation Questions**



Medical Specialty of participant?

## Paired Pre- and Post- Course Questions:



**▼** CONFIDENCE?

3 questions assessing

PCP's confidence with

PCH Diagnosis, Treatment

and Follow-up



**KNOWLEDGE?** 

8 True/False PCH

clinical knowledge

questions



Intending to make changes to practice?



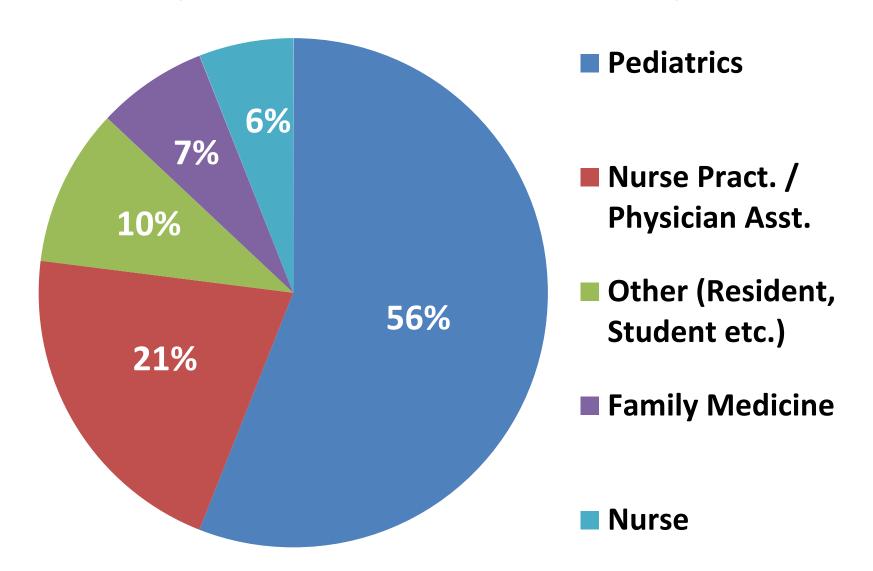
Course Ratings?



Comments and suggestions?

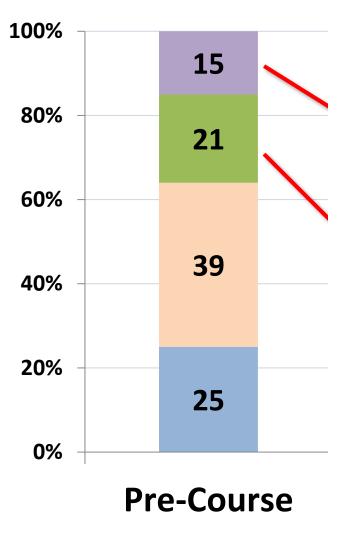


## Medical Specialties of Course Participants (N=171)



6 Events / Total attendance: ~ 300 / Evaluations Returned: 195

## **Confidence with CONFIRMING a PCH Diagnosis**

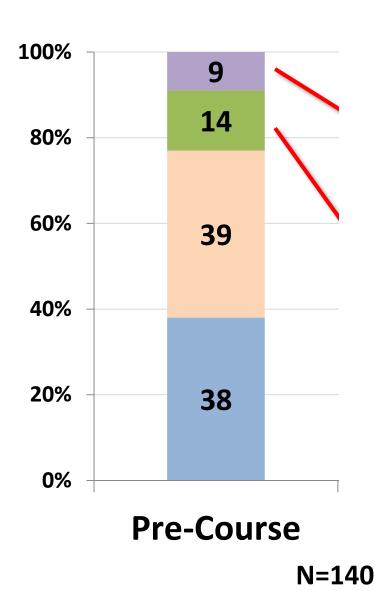


70 % improved their confidence

- Very Confident
- Confident
- Somewhat confident
- Not confident at all

N = 144

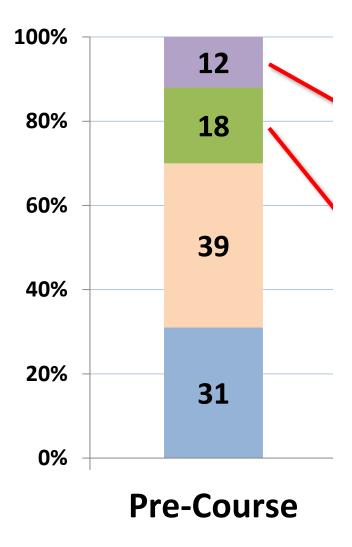
## Confidence with INITIATING TREATMENT for PCH



80 % improved their confidence

- Very Confident
- Confident
- Somewhat confident
- Not confident at all

## Confidence with PROVIDING FOLLOW-UP for PCH



69 % improved their confidence

- Very Confident
- Confident
- Somewhat confident
- Not confident at all

N = 143

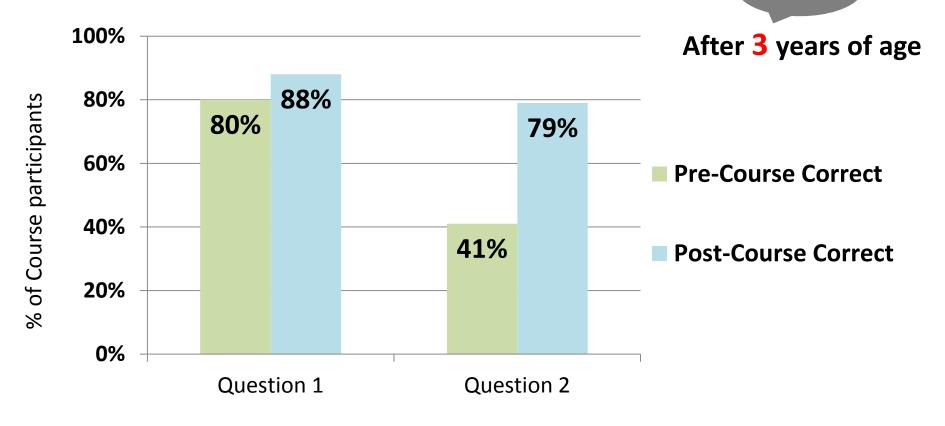
## **Improvements in Knowledge: True/False Questions**

**Question 1:** The recommended frequency of blood tests to monitor PCH patients in first 6 months of life is every 1-2 months (N=167)



**Question 2:** When transient PCH is suspected, it is safe to do a trial off levothyroxine for 4–6 weeks after 1 year of age (N=164)

False!

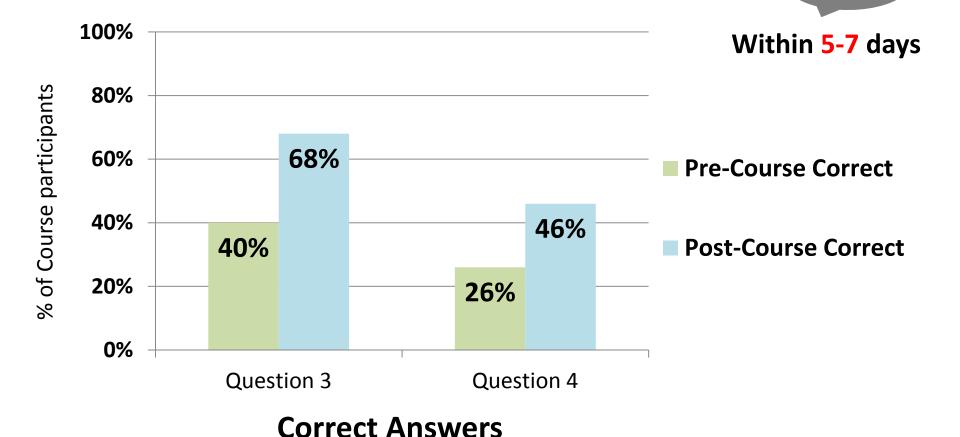


**Correct Answers** 

**Question 3:** The American Academy of Pediatrics (AAP) recommended initial treatment dose of LT4 for babies with confirmed PCH is 15–20 μg/kg (N=162) **10-15** μg/kg

Question 4: Thyroid stimulating hormone (TSH) surges by 30 minutes after birth and returns to <10  $\mu$ IU/mL within 3 days (N=159)

False!

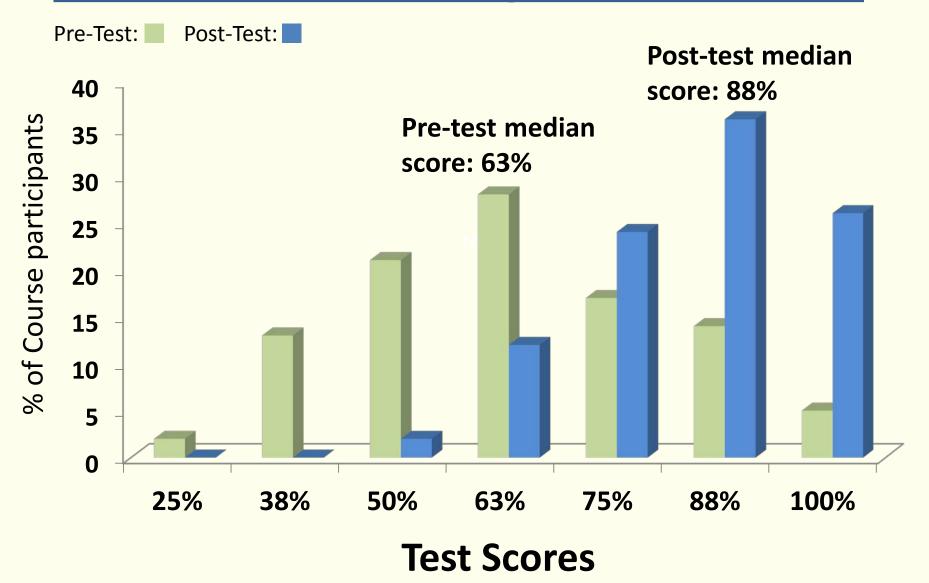


## Overall Improvement in True/False Total Score

 135 respondents answered all 8 pre-course and post-course true/false questions

Improvement on T/F Test Score (N=135)	Percentage
Improved their score	<b>72</b> %
Unchanged score	22%
Lower score	6%

## Pre- and Post- Test Score Distributions For 8 True/False Knowledge Questions (N=135)



## **Key Conclusions from True/False Results**

Results suggest **largest knowledge gaps** among PCPs about the following issues:

- Timing of post-natal TSH surge and return to normal
- Initial treatment dosage
- When and how to assess patients for transient PCH



"Speaker covered practical, useful information. Very applicable to general providers."

"I will be more conscious of following the monitoring of CH in newborns with guidelines discussed."



"I like the simple explanation of the process."

- 88% rated the course as "Outstanding"
- 75% said they plan to make changes to practice

## **Summary**

- The course was shown to be effective in improving PCPs knowledge and confidence in providing follow-up care for patients with PCH
- PCPs were especially enthusiastic about the simplicity of the course format and the practical instructions provided
- Participants indicated willingness and intention to make changes to their clinical practice in treating PCH
- Newborn screening programs may consider offering CME courses to PCPs who care for patients with PCH

## Research Team

Lisa Feuchtbaum, Dr. P.H., M.P.H.

Principal Investigator

Genetic Disease Screening Program, CDPH

#### **Co-Investigators:**

Ning Rosenthal, M.D., Ph.D.
Principal Research Scientist
Premier Inc.
Los Angeles, CA

**Sylvia Mann Au, M.S., CGC**Hawaii Department of Health, Genetic Section

Laura Bachrach, M.D.
Professor of Pediatrics
Division of Endocrinology

Stanford University Medical Center

#### **Study Manager:**

Emily Bezar, M.A.

**Public Health Foundation Enterprises** 

## **Advisory Committee**

Swati Banerjee, M.D.

Division of Pediatric Endocrinology Valley Children's Hospital, Madera, CA

Mitchell E. Geffner, M.D.

Chief, Center for Endocrinology, Diabetes and Metabolism, Children's Hospital Los Angeles, CA

Michael Gottschalk, M.D., Ph.D.

Chief, Pediatric Endocrinology
UC San Diego/Rady Children's Hospital, CA

Mary Rutherford, M.D.

Pediatrician Children's Hospital Oakland, CA

Stuart K. Shapira, M.D., Ph.D.

Chief Medical Officer and Associate Director for Science National Center on Birth Defects and Developmental Disabilities (NCBDDD) CDC, Atlanta, GA

## Thank You!

The contents, including all opinions and views expressed or implied, are entirely personal and do not necessarily represent the opinions or views of any person or organization, including the California Department of Public Health.

