Date of Preparation: 06/12/2023

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Signature

DORIS TAHA, MD

Office Address: 3950 Beaubien St
Pediatric Endocrinology
Children's Hospital of Michigan
Detroit, MI 48201

EDUCATION **DATES**

- Doctor of Medicine (M.D.) July 1990-June 1994
  American University of Beirut, Beirut, Lebanon

- Bachelor of Science (with Distinction) Mar. 1987-Apr. 1990
  American University of Beirut, Beirut, Lebanon

POSTGRADUATE TRAINING

- Fellowship in Pediatric Endocrinology (PGY IV-VI) July 1998-June 2001
  State University of New York Health Science Center at Brooklyn, Brooklyn, NY

- Residency in Pediatrics (PGY I-III) Apr. 1995-Apr. 1998
  State University of New York Health Science Center at Brooklyn, Brooklyn, NY

HOSPITAL OR OTHER PROFESSIONAL APPOINTMENTS

- **Division Chief, Pediatric Endocrinology** **Aug 2017- present**
  Children's Hospital of Michigan (CHM)
  University Pediatricians, Detroit, MI

- **Interim Chief, Division of Pediatric Endocrinology** **June 2016- July 2017**
  Children's Hospital of Michigan (CHM)
  University Pediatricians, Detroit, MI

- **Associate Professor of Pediatrics** — **July 2014- 2020**
  Pediatric Endocrinologist
  Wayne State University, Detroit, MI

- **Associate Professor of Pediatrics** — **2020- Present**
  Pediatric Endocrinologist
  Central Michigan University, Detroit, MI

- Attending, Pediatric Endocrinology — Aug. 2013- present
  Children's Hospital of Michigan
  University Pediatricians, Detroit, MI

- Consultant Pediatrician and Pediatric Endocrinologist — Oct. 2009-June 2012
  Cape Breton Regional Hospital, Sydney, Nova Scotia, Canada

- Consultant Pediatric Endocrinologist — Mar 2003-Sep. 2009
  King Faisal Specialist Hospital and Research centre- Jeddah, Saudi Arabia

- Associate Consultant Pediatric Endocrinologist — Feb. 2002-Mar. 2003
  King Faisal Specialist Hospital and Research centre- Jeddah, Saudi Arabia

- Clinical Instructor — 1995- 2001
  State University of New York, Downstate Medical Center, Brooklyn, NY

MAJOR PROFESSIONAL SOCIETIES

- **Member of the U.S. News & World Report's Best Children's Hospitals working group** — July 2016- present
- Member of the Endocrine Society
- Member of Pediatric Endocrine Society
- Fellow of the American Academy of Pediatrics
- Member of the Canadian Pediatric Endocrine Group
- Member of Doctors Nova Scotia (2009-2012)

MEDICAL LICENSURE

- Indiana State Physician License (Inactive) — Nov.2000-Jun. 2007
- New York State Physician License (Inactive) — Sep.2001-Aug. 2004

- Michigan Physician License (Active) Dec.2012-Dec. 2023
- Michigan Pharmacy License (Active) Dec. 2012-Dec. 2023
- Controlled Substance License (Active) 2018-2023

BOARD CERTIFICATION

- **American Board of Pediatric Endocrinology- Certificate # 921 Meeting MOC requirements until 12/2027** **Aug. 2001-Present**
- American Board of Pediatrics- Certificate # 65939 Oct. 1998- Dec 2015

HONORS/AWARDS **DATES**

- **Detroit Hour Top Docs** **2018-present**
- Lilly Travel grant to attend and present at the 83rd Annual Endocrine Society meeting- Denver- CO June 2001
- Division travel grant to attend and present at the Pediatric Academic Societies meeting- Baltimore- MD Apr. 2001
- Genentech Travel grant to attend and present at the 14th Annual NCGS Investigators Meeting- San Francisco- CA Sep. 2000
- Lilly Travel grant to attend and present at the Society of Pediatric Research meeting- Boston- MA May 2000
- Dean's Honor List- Medical School- American University of Beirut 1991 & 1993
- Dean's Honor List- School of Arts and Sciences- American University of Beirut 1987- 1990
- Graduated BS with Distinction Apr 1990

SERVICE

- **Division Chief- Pediatric Endocrinology- CHM** **Aug 2017- present**
  Contributing to growing the Division and introducing new programs, protocols, and clinics and physician and staff recruitment
- Member of the **U.S. News & World Report's Best Children's** **July 2016- present**

Hospitals working group: Reviews survey questions to rank Best Children's Hospitals

- **Medical Staff Operations Committee (MSOC)- CHM** **Jan 2021- present**

- Research Day Planning Committee 2017 and 2018

- Endocrine Stimulation tests power plans Committee June 2016- present

- Member of EMR documentation committee- Pediatric Endocrinology Division- Children's Hospital of Michigan Jan 2016- present

- Development and revision of protocols for Endocrine Division at the Children's Hospital of Michigan 2016- present
  - DKA protocol (Revision with 2 bag system implementation) 2018; revision 2020
  - High risk diabetes protocol 2019
  - Hypoglycemia protocol (with ED and Hospitalists) 2022
  - Bisphosphonate protocol (with infusion center) 2022
  - T-Zield protocol (with infusion center) 2023
  - Traumatic brain injury (TBI) protocol 2023

- Organizing and participation in Diabetes School Nurse Education Program in Metro Detroit area 2016- present

- Overseeing and help organizing Endocrine Division Teaching Activities 2015- present

- Member of the Institutional Research Review Board (IRB)- King Faisal Specialist Hospital and Research Centre- Jeddah- Saudi Arabia 2004-2009

- Chairperson of the Departmental Research Committee- Department of Pediatrics- King Faisal Specialist Hospital and Research Centre- Jeddah- Saudi Arabia 2005-2009

- Member of the Departmental Education Committee- Department of Pediatrics- King Faisal Specialist Hospital and Research Centre- Jeddah- Saudi Arabia 2002-2005

- Chairperson (3 times) and member (3 times) of Scientific organizing committees of 6 different scientific symposia/ conferences- King Faisal Specialist Hospital and Research Centre- Jeddah- Saudi Arabia 2003-2009

PHYSICIAN PRACTICE PLAN

University Pediatricians **Aug 2013-Present**

AFFILIATE MEDICAL ORGANIZATIONS

DMC- Children's Hospital of Michigan	**Aug 2013-Present**
Cape Breton Regional Hospital- Cape Berton- NS- Canada	2009-2012
King Faisal Specialist Hospital and Research Centre- Jeddah- Saudi Arabia	2002-2009
Downstate Medical Center	1995-2001

COMMUNITY SERVICES/ EDUCATIONAL PROGRAMS

Manage diabetes in your child and help them stay healthy. Expertbeacon.com	2013
School Nurse Diabetes Training Program at CHM- yearly	**2017- Present**
Educational materials for use at CHM clinics- Review and help with translation	**2013- Present**

SERVICE FOR PEER-REVIEWED JOURNALS AND NATIONAL/INTERNATIONAL MEETINGS

Review of Manuscripts

Reviewed articles submitted to: 2004- Present

- Metabolic Syndrome and Related Disorders
- The Annals of Saudi Medicine
- Saudi Medical Journal
- Journal of Medical genetics
- European Journal of Pediatrics

Review of Abstracts:

Pediatric Endocrine Society Meeting	2020- Present
Great Lakes Pediatric Research Day abstract review	**2022-** Present
International Meeting of Pediatric Endocrinology- September 2017	2017

Moderator for Scientific Sessions:

The Fifth International Pediatric Update Symposium-2009. Endocrinology Session. Jeddah, KSA.	April 2009
The Second International Pediatric Update Symposium-Pediatric Update 2005. Endocrinology Session. Jeddah, KSA.	April 2005
The First International Pediatric Update Symposium-Pediatric Update 2003. Endocrinology Session. Jeddah, KSA.	April 2003

Mentorship

Mentoring junior faculty in the Endocrine Division as CHM Division Chief	**2016- present**
Jawaher Bardise, MD. Fellow. Mentorship and training in Pediatric Endocrinology at King Faisal Specialist and Research Center- Jeddah, Saudi Arabia.	2008-2009
Lina Saadeh, MD. Resident. Mentorship on developing a research project on patients with Septo-optic dysplasia, writing an IRB protocol, presentation and abstract publication- CHM.	2015- 2016
Noelle Fabie, MD. Resident. Mentorship on developing a research project on patients with Septo-optic dysplasia, writing an IRB protocol, presentation and abstract publication- CHM.	2016-2018
Shivali Patel, WSU medical student. Mentorship during Pediatric rotation	2017
Yafa Davydova and Andrea Kendall; WSU medical students. Mentorship on retrospective study on patients with 45X/46XY mosaicism and writing IRB protocol- CHM.	2017
Pooja Dave- Resident at CHM. Mentorship on writing a case report and preparing abstract for presentation at the Pediatric Endocrine Society Meeting.	2019-2020
Jenny Huang- Immunology fellow. Project on Adrenal insufficiency and Asthma.	2019-2020

COURSES/ EDUCATION PROGRAMS:

- **Diabetes Educator Course**

King Abdulaziz Medical City- National Guard Hospital- Jeddah, Saudi Arabia. Insulin types and their actions. Diabetes Educator course. 2006-2008

- **CHM Diabetes School Nurse Education Program- Curriculum Development** 2016- present

PATENTS

USE OF GLIS3 FOR PREPARING FUNCTIONAL PANCREATIC BETA-CELLS- Application number: 20110112015

Filed: April 16, 2007
Issued: May 12, 2011

Abstract: The present invention relates to a method of diagnosis of neonatal diabetes, congenital hypothyroidism and congenital glaucoma,

comprising detecting the presence of a mutation in the GLIS3 gene and to a method for preparing functional pancreatic beta-cells by culturing human multipotent/ pluripotent cells, such as embryonic stem cells (ES-cells) or human somatic stem cells in a culture medium comprising a effective amount of GLIS3 to induce cells differentiation into functional pancreatic beta-cells producing insulin. It relates to a pharmaceutical composition comprising GLIS3 as active ingredient. **Type:** Application **Inventors:** Cecile Julier, Doris Taha, Marc Nicolino, Douglas Cavene *(co-inventor).*

PEER-REVIEWED PUBLICATIONS:

Reports of Original Work:

1. An emerging, recognizable facial phenotype in association with mutations in GLI-similar 3 (GLIS3). Dimitri P, De Franco E, Habeb AM, Gurbuz F, Moussa K, **Taha D**, Wales JK, Hogue J, Slavotinek A, Shetty A, Balasubramanian M. **Am J Med Genet A**. **2016** Jul;170(7):1918-23. PMID 27148679.

2. Expanding the clinical spectrum associated with GLIS3 mutations. Dimitri P, Habeb AH, Garbuz F, Millward A, Wallis S, Moussa K, Akcay T, **Taha D**, Hogue J, Slavotinek A, Shetty A, Hawkes D, Hattersley AT, Ellard S, De Franco E. 2015. **J Clin Endocrinol Metab**. **2015**;100(10):E1362-9. PMID 26259131.

3. Vitamin D receptor mutations in patients with hereditary 1,25-dihydroxyvitamin D-resistant rickets. Malloy PJ, Tasic V, **Taha D**, Tütüncüler F, Ying GS, Yin LK, Wang J, Feldman D. **Mol Genet Metab**. **2014** ; 111 (1): 33-40. PMID 24246681.

4. Reduced elastogenesis: a clue to the arteriosclerosis and emphysematous changes in Schimke immuno-osseous dysplasia? Morimoto M, Yu Z, Stenzel P, Clewing JM, Najafian B, Mayfield C, Hendson G, Weinkauf JG, Gormley AK, Parham DM, Ponniah U, André JL, Asakura Y, Basiratnia M, Bogdanović R, Bokenkamp A, Bonneau D, Buck A, Charrow J, Cochat P, Cordeiro I, Deschenes G, Fenkçi MS, Frange P, Fründ S, Fryssira H, Guillen-Navarro E, Keller K, Kirmani S, Kobelka C, Lamfers P, Levtchenko E, Lewis DB, Massella L, McLeod DR, Milford DV, Nobili F, Saraiva JM, Semerci CN, Shoemaker L, Stajić N, Stein A, **Taha D**, Wand D, Zonana J, Lücke T, Boerkoel CF. **Orphanet J Rare Dis**. **2012** Sep 22;7:70. doi: 10.1186/1750-1172-7-70. PMID 22998683.

5. Schimke immunoosseous dysplasia: defining skeletal features. Hunter KB, Lücke T, Spranger J, Smithson SF, Alpay H, André JL, Asakura Y, Bogdanovic R, Bonneau D, Cairns R, Cransberg K, Fründ S, Fryssira H, Goodman D, Helmke K, Hinkelmann B, Lama G, Lamfers P, Loirat C, Majore S, Mayfield C, Pontz BF, Rusu C, Saraiva JM, Schmidt B, Shoemaker L, Sigaudy S, Stajic N, **Taha D**, Boerkoel CF.**Eur J Pediatr**. **2010** Jul;169(7):801-11. Epub 2009 Dec 15. PMID 20013129.

6. The Prevalence of the Metabolic Syndrome and Cardiovascular Risk Factors in a Group of Obese Saudi Children and Adolescents: A Hospital-based study. **Taha D**, Ahmed O. **Ann Saudi Med**. **2009** Sep-Oct;29(5):357-60. PMID 19700892.

7. Hyperglycemia and Hypoinsulinemia in Patients with Fanconi-Bickel Syndrome. **Taha D**, AL-Harbi N, AL-Sabban E. **J Pediatr Endocrinol Metab**. **2008**; 21(6):581-6. PMID 18717244.

8. Schimke immunoosseous dysplasia: suggestions of genetic diversity. Clewing JM, Fryssira H, Goodman D, Smithson SF, Sloan EA, Lou S, Huang Y, Choi K, Lücke T, Alpay H, André JL, Asakura Y, Biebuyck-Gouge N, Bogdanovic R, Bonneau D, Cancrini C, Cochat P, Cockfield S, Collard L, Cordeiro I, Cormier-Daire V, Cransberg K, Cutka K, Deschenes G, Ehrich JH, Fründ S, Georgaki H, Guillen-Navarro E, Hinkelmann B, Kanariou M, Kasap B, Kilic SS, Lama G, Lamfers P, Loirat C, Majore S, Milford D, Morin D, Ozdemir N, Pontz BF, Proesmans W, Psoni S, Reichenbach H, Reif S, Rusu C, Saraiva JM, Sakallioglu O, Schmidt B, Shoemaker L, Sigaudy S, Smith G, Sotsiou F, Stajic N, Stein A, Stray-Pedersen A,**Taha D**, Taque S, Tizard J, Tsimaratos M, Wong NA, Boerkoel CF. **Hum Mutat**. **2007** Mar;28(3):273-83. PMID 17089404.

9. Mutations in *GLIS3* are responsible for a rare syndrome with neonatal diabetes mellitus and congenital hypothyroidism. Senée V, Chelala C, Duchatelet S, Feng D, Blanc H, Cossec JC, Charon C, Nicolino M, Boileau P, Cavener DR, Bougnères P, **Taha D** & Julier C. **Nature Genetics 2006**; 38(6):682-7. Epub 2006 May 21. PMID 16715098.

10. Type 2 Diabetes in African – American Adolescents: Impaired cell function in the face of severe insulin resistance. **Taha D**, Umpaichitra V, Banerji MA, Castells S. **Journal of Pediatric Endocrinology & Metabolism 2006**;19(2):135-42. PMID 16562586 Clinical Trial.

11. Absent or delayed adrenarche in patients with Pit-1/POU1F1 deficiency. **Taha D.**, Mullis PE, Ibanez L, de Zegher F. **Horm Res 2005**;64(4):175-179. PMID 16210857.

12. Association of migraine-like headaches with Schimke immuno-osseous dysplasia.
Kilic SS, Donmez O, Sloan EA, Elizondo LI, Huang C, André JL,Bogdanovic R, Cockfield S, Cordeiro I, Deschenes G, Fründ S, Kaitila I, Lama G, Lamfers P, Lücke T, Milford DV, Najera L, Rodrigo F, Saraiva JM, Schmidt B,Smith GC, Stajic N, Stein A, **Taha D**, Wand D, Armstrong D, Boerkoel CF. **Am J Med Genet A**. **2005** Jun 1;135(2):206-10. PMID 15884045.

13. EIF2AK3 mutations and clinical variability in Wolcott-Rallison syndrome, and suggestion of genetic heterogeneity. Senee V, Vattem KM, Delepine M, Rainbow LA, Haton C, Lecoq A, Shaw NJ, Robert JJ, Rooman R, Diatloff-Zito C, Michaud JL, Bin-Abbas B, **Taha D**, Zabel B,Franceschini P,Topaloglu AK,Lathrop GM, Barrett TG, Nicolino M,Wek RC, Julier C. **Diabetes 2004**;53(7):1876-83. PMID 15220213.

14. Hyperlipidemia in children with type 2 diabetes mellitus. **Taha D**. **Journal of Pediatric Endocrinology and Metabolism 2002** Apr;15 Suppl 1:505-7. PMID 12017224.

15. Beta Cell Response to Intravenous Glucagon in African- American and Hispanic Children with Type 2 Diabetes Mellitus. **Taha D**, Castells S, Umpaichitra V, Bastian W, Banerji MA. **Journal of pediatric Endocrinology and Metabolism 2002** Jan; 15(1): 59-67. PMID 11822582.

16. C-peptide and Glucagon Profiles in Minority Children with Type 2 Diabetes Mellitus. Umpaichitra V., Bastian W, **Taha D** , Banerji M, AvRuskin TW, Castells S. **Journal of Clinical Endocrinology and metabolism 2001** Apr; 86(4):1605-1609. PMID 11297591.

Case Reports:

17. Familial Neonatal Non-autoimmune Hyperthyroidism Due to A (D619G) Thyrotropin Receptor Mutation. **Taha D**, Adhikari A, Flore LA. **J Pediatr Endocrinol and Metab. 2020; 34 (2): 267-271.** PMID 33180037.

18. A Case of Congenital Hypopituitarism Associated With a 1p31 Microdeletion: A Possible Role for LEPR and JAK1. Thakur M. **Taha D**, Misra V. **Journal of the Endocrine Society**, Volume 1, Issue 4, 1 **April 2017**, Pages 278–282. PMID 29264484.

19. Neonatal diabetes mellitus because of pancreatic agenesis with dysmorphic features and recurrent bacterial infections. **Taha D**, Bardise J*, Hegab A, Bonnefond A, Marchand M, Drunat S, Vaxillaire M, Polak M. **Pediatr Diabetes**. **2008**; 9(3Pt1): 240-4. PMID 18547237.

20. A Novel VPS33B Mutation in an ARC Syndrome Patient Presenting with Osteopenia and Fractures at Birth. **Taha D**, Khider A*, Cullinane A, Gissen P. **Am J Med Genet A**. **2007**;143A(23):2835-2837. PMID 17994566.

21. Arthrogryposis, Renal tubular acidosis & Cholestasis (ARC) Syndrome: Two new cases & review. Abu-Sa'da O*, Barbar M, Al-Harbi N, **Taha D**. **Clin Dysmorphology**. **2005**; 14(4):191-196. PMID 17159523.

22. Nephrocalcinosis in glucose-galactose malabsorption, association with renal tubular acidosis. El-Naggar W*, Balfe JW, Barbar M, **Taha D**. **Pediatr Nephrol**. **2005**; 20(9):1336-9. Epub 2005 Jul 12. PMID 16010597.

23. Fatal Lymphoproliferative Disorder in a child with Schimke Immuno-osseous Dysplasia. **Taha D,** Boerkoel CF, Balfe JW, Khalifah M, Sloan EA, Barbar M, Haider A, Kanaan H. **American Journal of Medical Genetics 2004**; 131A(2):194-9. PMID 15523612.

24. Neonatal diabetes mellitus, congenital hypothyroidism, hepatic fibrosis, polycystic kidneys, and congenital glaucoma: A new autosomal recessive syndrome? **Taha D**, Barbar M, Kanaan H, and Balfe J.W. **American Journal of Medical Genetics 2003**, Oct 15;122A(3):269-73. PMID 12966531.

25. Monozygotic twins with congenital adrenal hyperplasia: long-term endocrine evaluation and gene analysis. AvRuskin TW, Witchel SF, **Taha DR**, Juan CS. **Journal of Pediatric Endocrinology and Metabolism 2003**; 16(4):565-70. PMID 12793610.

26. Growth Hormone Replacement Therapy in Children With Leukemia In Remission. **Taha D**, Bastian W, Castells S. **Clinical Pediatrics (Phila) 2001** Aug.; 40(8): 441-445. PMID 11516051.

Letters to the Editor:

27. EIF2AK3 mutations in Patients with Wolcott-Rallison Syndrome. **Taha D.** Annals of Saudi Medicine **2005**; 25(4):350. PMID16212134.

Published Abstracts:

1- Maturity onset diabetes of the young (MODY) due to a new *BLK* gene mutation. **Taha D**, Thirunagari R, Adhikari A. Accepted for publication in the **Journal of the Endocrine Society**. **2023**.

2- Comsumptive hypothyroidism secondary to hemangioma. Thirunagari R, **Taha D**. Accepted for publication in the **Journal of the Endocrine Society, 2023**.

3- Isolated Growth Hormone Deficiency in A Patient With Gitelman Syndrome. Dave P, **Taha D**. **Hormone Research in Pediatrics 2020**; vol 93 suppl 1: 125-126.

4- Familial Neonatal Nonautoimmune Hyperthyroidism Due To A Thyrotropin-Receptor Gene Mutation (A619G). **Taha D**, Adhikari A, Thirunagari R, Senguttuvan R, Flore LA. **Journal of the Endocrine Society 2019**; vol 3 suppl 1: MON-264.

5- A Case of a Forgotten Histrelin Acetate Implant: Can Histrelin Acetate Implants Be Used for More Than a Year? Krishnaraj BK, Adhikari A, **Taha D**, Thirunagari R, Senguttuvan R. **Journal of the Endocrine Society 2019**; vol 3 suppl 1: SAT-298.

6- Mutation in SRY Gene Presenting as Syndromic 46XY Disorder of Sexual differentiation (DSD). Adhikari A, **Taha D**, Rajan S, Yegappan L. **Journal of the Endocrine 2019**; vol 3 suppl 1: SAT-297.

7- Nutritional Rickets: Still A Problem in The 21st Century. Thirunagari R, **Taha D**. **Journal of the Endocrine Society2019**; vol 3 suppl 1: SAT-275.

8- Clinical and Endocrinological Characteristics of Patients with Septo-optic Dysplasia at the Children's Hospital of Michigan. Fabie N, Saadeh L, **Taha D**. **Hormone Research in Paediatrics 2017;** vol 88 suppl 1; P1-1002.

9- Prevalence of the Metabolic Syndrome Risk Factors in Obese Saudi Children and Adolescents. **Taha D**, Ahmed O. **Journal of Diabetes 2009**;1 (s1):A177.

10- Treatment of hereditary vitamin D resistance rickets (HVDRR) with intermittent intravenous calcium. **Taha D**, Al-zaydi A, Basha T, Al-Ghamdi M, Bin-Abbas B, Al-Ashwal A. **Pediatric Research 2003**; 53(4): 150A (856).

11- Management of Minority Children with Type 2 Diabetes. Castells S, **Taha D**, Umpaichitra V, Bastian W, and Banerji M. **Pediatric Research 2001**; 49(4): 100A(561).

12- Autoantibodies in Children with Type 2 Diabetes Mellitus. Umpaichitra V, Jackson R, **Taha D**, Bastian W, and Castells S. **Pediatric Research 2001**; 49(4): 100A(560).

13- Single-Stranded Conformation Polymorphisms of the Glucokinase Gene in Type 2 Diabetes of Adolescence. Bastian W, Umpaichitra V, **Taha D**, Castells S. **Journal of Pediatric Endocrinology & Metabolism 2000**; 13 suppl4: Abstract 4(p1209).

14- C-peptide and Glucagon profiles in minority Children with type 2 Diabetes Mellitus. Umpaichitra V, Bastian W, **Taha D**, Banerji MA, AvRuskin TW, Castells S. **Pediatric Research 2000**; 47(4): 141A (829).

15- The serum C-peptide Response to Stimulation with Intravenous Glucagon in Type 2 Diabetes in Children. **Taha D**, Umpaichitra V, Bastian W, Banerji MA, Castells S. **Pediatric Research 2000**; 47(4): 140A (824).

16- Diabetic Adolescent: type 1 or type 2? **Taha D**, Sadagopan R, Bastian W, Castells S. **Pediatric Research 1999**; 45: 98A (570).

PRESENTATIONS TO NATIONAL AND INTERNATIONAL MEETINGS:

1-Maturity onset diabetes of the young (MODY) due to a new *BLK* gene mutation. **Doris Taha**, Rajeev Thirunagari, Amita Adhikari. **Presented at the Endocrine Society Meeting, Chicago, June 2023.**

2-Comsumptive hypothyroidism secondary to hemangioma. Rajeev Thirunagari, **Doris Taha**. **Presented at the Endocrine Society Meeting, Chicago, June 2023**.

3- Isolated Growth Hormone Deficiency in A Patient With Gitelman Syndrome. Pooja Dave, **Doris Taha**. Accepted for presentation at the **Pediatric Endocrine Society Meeting**, Texas, **April 2020,** conference canceled.

4- Familial Neonatal Nonautoimmune Hyperthyroidism Due To A Thyrotropin-Receptor Gene Mutation (A619G). **Doris Taha**, Amita Adhikari, Rajeev Thirunagari, Rajan Senguttuvan, Leigh A Flore. Presented at the **Endocrine Society Meeting**, New Orleans, **March 2019**.

5- A Case of a Forgotten Histrelin Acetate Implant: Can Histrelin Acetate Implants Be Used for More Than a Year? Krishnaraj BK, Adhikari A, **Doris Taha**, Thirunagari R, Senguttuvan R. Presented at the **Endocrine Society Meeting**, New Orleans, **March 2019**.

6- Mutation in SRY Gene Presenting as Syndromic 46XY Disorder of Sexual differentiation (DSD). Adhikari A, **Doris Taha**, Rajan S, Yegappan L. Presented at the **Endocrine Society Meeting**, New Orleans, **March 2019**.

7- Nutritional Rickets: Still A Problem in The 21st Century. Thirunagari R, **Doris Taha**. Presented at the **Endocrine Society Meeting**, New Orleans, **March 2019**.

8- Clinical and Endocrinological Characteristics of Patients with Septo-optic Dysplasia at the Children's Hospital of Michigan. Noelle Fabie, Lina Saadeh, Amita Adhikari, Colleen Buggs-Saxton, **Doris Taha**. Presented at the **10th International meeting of Pediatric Endocrinology**, Washington, DC. **September 2017**.

9- Prevalence of the Metabolic Syndrome Risk Factors in Obese Saudi Children and Adolescents. **Doris Taha**, Omaima Ahmed. Presented at the **3rd International Congress on Prediabetes and the Metabolic Syndrome**, Nice, France. **April 2009**.

10- Clinical Variability in Wolcott-Rallison Syndrome (Permanent Neonatal Diabetes and Spondyloepiphyseal Dyspalsia). MP Nicolino, V Senee, M Delepine, L Rainbow, C Haton, **Doris Taha,** et al. Presented at The **Endocrine Society's 86th Annual Meeting**, New Orleans, LO**. June 2004**.

11- Abnormal Carbohydrate metabolism in patients with Fanconi-Bickel Syndrome. **Doris Taha**, N. Al-Harbi, E. Al-Sabban. Presented at The **Endocrine Society's 86th Annual Meeting**, New Orleans, LO. **June 2004**.

12- Treatment of hereditary vitamin D resistance rickets (HVDRR) with intermittent intravenous calcium. **Doris Taha**, A. Al-zaydi, T. Basha, M. Al-Ghamdi, B. Bin-Abbas, A. Al-Ashwal. Presentated at the **2003 Pediatric Academic Societies Meeting**, Seattle, WA. **May 2003**.
Also, presented (oral presentation) at the **2nd international Pediatric Symposium, King Faisal Specialist Hospital & Research Centre-Jeddah, April 2003**, and presented (oral presentation) at the **5th Pan-Arab Congress on Endocrinology and Diabetes**, Riyadh, KSA, **Feb 2004**.

13- Familial glucocorticoid deficiency type 2: A case report. W. Al-Najjar, **Doris Taha**. Presented at the **2nd international Pediatric Symposium**, King Faisal Specialist Hospital & Research Centre-Jeddah, **April 2003**.

14- Fasting lipid profiles in Children With Type 2 Diabetes. V. Umpaichitra, **Doris Taha**, Y. Ahmed, R. Walia, M. Banerji, W. Bastian, and S. Castells. Presented at The **Endocrine Society's 83rd Annual Meeting**, Denver, CO. **June 22, 2001**.

15- C-peptide and Insulin Responses to Oral Glucose Load in Type 2 Diabetes in Children. **Doris Taha**, V. Umpaichitra, Y. Ahmed, R. Walia, M. Banerji, W. Bastian, and S. Castells. Presented at the The **Endocrine Society's 83rd Annual Meeting**, Denver, CO. **June 21, 2001**.

16- Autoantibodies in Children with Type 2 Diabetes. V. Umpaichitra, R. Jackson, **Doris Taha**, W. Bastian, and S. Castells. Presented at the **2001 Pediatric Academic Societies Meeting**, Baltimore, MD. **April 28, 2001**.

17- Management of Minority Children with Type 2 Diabetes. S. Castells, **Doris Taha**, V. Umpaichitra, W. Bastian, and M. Banerji. Presented at the **2001 Pediatric Academic Societies Meeting**, Baltimore, MD. **April 28, 2001**.

18- Single stranded conformation polymorphisms of the glucokinase gene in type 2 diabetes of adolescence. William Bastian, Vatcharapan Umpaichitra, **Doris Taha**, Salvador Castells. Presented at **the 26th Annual Meeting of the International Society for Pediatric and Adolescent Diabetes (ISPAD).** Hollywood, CA. **Oct31-Nov3, 2000**.

19- Growth Hormone therapy during remission of leukemia. **Doris Taha**, William Bastian, Salvador Castells. Presented at the **Fourteenth Annual NCGS Investigators Meeting**, San Francisco, CA. **Sept. 2000**.

20- C-peptide and Glucagon Profiles in Minority Children with Type 2 Diabetes Mellitus. V. Umpaichitra, W. Bastian, **Doris Taha**, M. Banerji, T.W. AvRuskin, S. Castells. Presented at the meeting of the Society of Pediatric Research (SPR), Boston, MA. **May 2000**.

21- The Serum C-peptide Response to Intravenous Glucagon stimulation in Children with Type 2 Diabetes. **Doris Taha**, Vatcharapan Umpaichitra, William Bastian, M. Banerji, Salvador Castells. Presented at the **meeting of the Society of Pediatric Research (SPR),** Boston, MA. **May 2000**.

INVITED LECTURES//PRESENTATIONS/ MODERATOR:

- Moderates Lectures/ Clinical presentations/Journal clubs- Pediatric Endocrine Division- CHM- **2016 to present**
- Yearly presentation for School Nurse Diabetes Education Program-CHM- **2018 to present**
- Lecture for Pediatric Residents, Children's Hospital of Michigan. Management of Girls with Turner Syndrome. Detroit, MI. **2016, 2017, 2018.**
- Pediatric Endocrine Division Lecture, Children's Hospital of Michigan. Guidelines for management of differentiated thyroid cancer in children. Detroit, MI. **2016, 2017, 2018, 2019.**
- Pediatric Endocrine Division Lecture, Children's Hospital of Michigan. Guidelines for management of thyroid nodules in children. Detroit, MI. **2015**
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- The First International Symposium on the Metabolic Syndrome. The Metabolic Syndrome in Children. Jeddah, Saudi Arabia. **26 April 2006**.
- The Fourth International Conference- Women's Health Update 2006. Growth hormone in Short Stature. Jeddah, Saudi Arabia. **22 March 2006**.
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- Neonatology Symposium, KFSH&RC-Jeddah. Neonatal Hypoglycemia. Jeddah, Saudi Arabia. **October 2004**.
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