

FOR IMMEDIATE PRESS RELEASE

***** Registration is now OPEN for the 1st CDG Satellite Meeting for Professionals *****

Lisbon, 28th February 2019 – To join the Rare Disease Celebrations we are opening the registrations for the **1st CDG Satellite Meeting: “Defining CDG at the international level!”**, the unique international event that will join professionals from around the world to discuss **CDG definition, nomenclature and nosology**.

The **1st CDG Satellite Meeting: “Defining CDG at the international level!”** will take place on **25th July (afternoon, pre-World Conference)**. More information [HERE](#).

Why should you come?

This Satellite Meeting is a pioneer event in which in what to promote discussion among professionals of the CDG Community.

A clear definition of the disease, the nomenclature and nosology is fundamental for a fast and accurate diagnosis and to boost therapies.

We welcome you to join us in this unique opportunity to make a difference!

Registration form - [HERE](#)

Additional information and updates will be available in social media: [Facebook](#), [Twitter](#), [LinkedIN](#) and APCDG blog [HERE](#).



**CONGENITAL
DISORDERS OF
GLYCOSYLATION
WORLD CONFERENCE**

The power of advancing patient-oriented research united
FAMILIES AND PROFESSIONALS

“We are very pleased to have Dr. Carlos Ferreira on board! Our goal is to promote discussion and to help professionals reach a consensus regarding CDG nomenclature and nosology. This is extremely important to have future breakthroughs in therapies. We hope that all CDG professionals can join us so we can push for a change together!” – said **Vanessa Ferreira, PhD, MBA (APCDG founder and volunteer scientific coordinator at CDG & Allies – PPAIN)**.

1st CDG Satellite Meeting
CDG Nomenclature

Join our main speaker
Dr. Carlos Ferreira

25th July afternoon
Lisbon

CDG PORTUGUESE ASSOCIATION

CDG & Allies - PPAIN

CONGENITAL DISORDERS OF GLYCOSYLATION WORLD CONFERENCE
The World of Glycosylation - Patient Reported Outcomes - 2019

"We are very excited to announce the **1st Satellite Meeting on CDG Nomenclature**, taking place on **25th July**. The ambitious aim of this meeting is to strive towards implementing an international and uniform definition and nomenclature for CDG." **Rita Francisco, MSc (CDG patient advocate and researcher, CDG & Allies-PPAIN)**

About the Portuguese CDG and other Metabolic Rare Disorders Association (APCDG):

Founded in 2010, APCDG (www.apcdg.com) is a patient led and centric non-profit association, whose particular goal is to stimulate new research lines that make a difference in the lives of patients and families. APCDG initiatives are developed both nationally and internationally. APCDG is committed to finding a cure for Congenital Disorders of Glycosylation (CDG) and related disorders, to improving the treatment options and to giving information and support to people with CDG, through research, education, awareness programs and advocacy. APCDG top priority is to give a complete and holistic perspective of the patient as a person. Go to our website: <http://www.apcdg.com/>

About CDG & Allies-PPAIN: With the help of a broad network of scientists, physicians, families and patient advocacy groups, we have established a patient-led national and internationally unrivaled infrastructure for research, awareness and education for CDG. The research on glycosylation disorders is primarily dedicated to Congenital Disorders of Glycosylation (CDG). The advances and innovations achieved for CDG through CDG & Allies - PPAIN will impact on a large number of patients, namely all human diseases characterized by abnormal protein glycosylation such as cancer, inflammation, Alzheimer disease and diabetes.

Contact: Sandra Brasil, PhD in Molecular Biology. Researcher at the Working Group CDG & Patient Reported Outcomes (PROMs). More information [HERE](#). Email: s.arduim@gmail.com