EAS

85th EAS Congress, Prague

Navigating Hypertriglyceridaemia: Spotlight on Familial Chylomicronaemia Syndrome

Recurrent episodes of abdominal pain, acute pancreatitis and risk of hospitalisation… These are some of the fears that patients with Familial Chylomicronaemia Syndrome (FCS) have to face every day.

Despite experiencing often severe and distressing symptoms, a cycle of misdiagnosis leaves many patients with FCS undetected, and even those who are fortunate enough that their disease has been given a name are frustrated by the restrictions it places on their everyday life. But by searching to understand the key clinical characteristics of FCS, we now have the knowledge and the tools to shine a light on this rare disease.

Join us as we explore the different diagnosis and management strategies for FCS, and the significant unmet needs in reducing the risk of complications in these patients.

Special Lecture: Recognising Familial Chylomicronaemia Syndrome

13:00–13:45, Tuesday 25 April 2017; Karel Rokytansky / Panorama Hall, 1st Floor, Prague Congress Centre

Time	Session title	Speaker
13:00–13:05	**Welcome and Introduction**	Erik Stroes University of Amsterdam, Netherlands
13:05–13:20	**Familial Chylomicronaemia Syndrome – What's in a Name?**	Erik Stroes
13:20–13:35	**Finding the One in a Million**	Željko Reiner, University Hospital Center Zagreb, Croatia
13:35–13:45	**Q&A**	Erik Stroes and Željko Reiner

Meet the Experts: Difficulties in Diagnosing and Managing the Patient with Familial Chylomicronaemia Syndrome – Case Studies

14:00–14:45, Tuesday 25 April 2017; The Hub, 2nd Floor, Prague Congress Centre

Session chairs: Erik Stroes and Željko Reiner

Time	Session title and speaker
14:00–14:05	**Welcome and Introduction**
14:05–14:25	**Case 1: Elizabeth Hughes, Sandwell Hospital, West Bromwich, UK**
14:25–14:45	**Case 2: Eric Bruckert, Pitié-Salpêtrière Hospital, Paris, France**

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