

Mitochondrial Medicine 2018

Wellcome Genome Campus Hinxton, Cambridge, UK

9-11 May 2018

Draft Conference Programme

Wednesday 9th May 2018

11:30-12:50 Registration with Lunch

Wellcome Genome Campus Conference Centre

12:50-13:00 Welcome and Introductions

Francis Crick Auditorium

Patrick Chinnery

University of Cambridge, UK

13:00-14:30 Session 1: Precision diagnostics and new clinical biomarkers

Chair: Patrick Chinnery, University of Cambridge, UK

13:00 Defining mitochondrial protein function through systems

biochemistry

David Pagliarini

University of Wisconsin-Madison, USA

13:30 Integrated 'omics

Holger Prokisch

Institute of Human Genetics, Germany

14:00 Selected talk

14:15 Selected talk

14:30-14:45 Session Discussion

14:45 15:00 **Afternoon Tea**

15:00-16:30 Session 2: What determines the phenotype of mitochondrial

diseases?

Chair. Jan Smeitink, Radboud UMC, The Netherlands

15:00 Mitochondrial diseases - pathophysiology beyond the

organelle

Anu Suomalainen

University of Helsinki, Finland

15:30 Post-transcriptional regulation of mitochondrial metabolism

during stress Elena Rugarli

University of Cologne, Germany

16:00 Selected talk

16:15 Selected talk

16:30-16:45 Session Discussion

16:45 – 17:30 Lightening Talks (1 minute per poster – odd number only)

17:30-18:45 Poster Session I (odd numbers) with drinks reception

19:00 prompt **Dinner**

Hinxton Hall Restaurant

19:00-23:00 **Cash Bar**

Wellcome Genome Campus Conference Centre

Thursday 10th May 2018

07:30-09:00 Breakfast

Hinxton Hall Restaurant

09:00-10:30 Session 3: Autophagy-mitophagy: is it important for

mitochondrial disease?

Chair: Massimo Zeviani, MRC Mitochondrial Biology Unit, UK

09:00 Using small molecule modulators for protein import to

regulate mitophagy Carla Koehler UCLA, USA

09.30 Resolving the pervasive nature of mammalian

mitophagy lan Ganley

University of Dundee, UK

10:00 Selected talk

10:30 – 10:45	Session Discussion
10:45-11.15	Morning Coffee
11:15-12:45	Session 4: Fission/fusion defects in mitochondrial disease: cause or effect? Chair: Marni Falk, The Children's Hospital of Philadelphia, USA
	11:15 Linking mitochondrial cell biology to mitochondrial diseases Julian Prudent MRC Mitochondrial Biology Unit, UK
	11:45 Mitochondrial dynamics Orian Shirihai Boston University, USA
	12:15 Selected talk
	12:30 Selected talk
12:45-13:00	Session Discussion
13:00-14:30	Lunch Hinxton Hall Restaurant
4.4.00.40.00	
14.30-16:00	Session 5: Mitochondrial biogenesis: the key to therapy? Chair: Anu Suomalainen, University of Helsinki, Finland
14.30-16:00	
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19:00 prompt Conference Dinner

Hinxton Hall Restaurant

19:00-23:00 Cash Bar

Hinxton Hall Bar and Wellcome Genome Campus Conference

Centre

Friday 11 May 2018

07:30-09:00 Breakfast

Hinxton Hall Restaurant

09:00-10:30 Session 6: Gene editing to treat mitochondrial diseases

Chair: Vasmi Mootha, Harvard Medical School, USA

09:00 Title tbc *Michio Hirano*

Columbia University, USA

09:30 Protein replacement therapy for mitochondrial genetic

diseases

Haya Louberboum-Galski Hebrew University, Israel

10:00 Selected talk

10:15 Selected talk

10:30-10:45 **Session Discussion**

10:45-11:15 Morning Coffee

11:15-12:15 **Session 7: Trial design in rare disease:**

Chair: Patrick Chinnery, University of Cambridge, UK

11:15 Trial design in rare disease

Shamima Rahman

University College London, UK

11:45 Trial design in rare disease

Patrick Yu Wai Man

University of Cambridge, UK

12:15-12:45 Session Discussion

12:45-13:30 Session 8: Clinical intervention studies - latest updates

Chair: Vasmi Mootha, Harvard Medical School, USA

Oral presentations selected from submitted abstracts

13:	:30	Conference Summary
13:	:45	Conference Closure with Grab Bag lunch
14:	:00	Complimentary shuttle to Cambridge train station, Cambridge City Centre, London Stansted Airport and London Heathrow Airport