



Wilson Aarhus

2026

Preliminary Program

Thursday April 30

19.00-22.00 Welcome dinner at the venue Comwell, Værkmestergade 2, DK-8000 Aarhus C

Friday May 1

Morning

9.00 Welcome *President, Thomas Sandahl (Aarhus)*

Session I: Welcome session

Chairs: Thomas Sandahl, Svetlana Lutsenko

9.10 WDGA – Wilson Disease Global Alliance *Lisbet Ottesen (Copenhagen)*

9.30 Gut-liver-brain Interactions *Valentina Medici (Sacramento)*

9.50 The Brain and Copper: Beyond Wilson Disease *Aurélia Poujois (Paris)*

10.10 ABS 1 *TBA*

10.25 Coffee break

Session II: Neurology I

Chairs: Aurélia Poujois, Petr Dusek

11.00 Modified UWDRS *Petr Dusek (Prague)*

11.20 The added value of Neurophysiology in WD with movement disorder *Mickaël Alexandre Obadia (Paris)*

11.35 Is r-TMS the future treatment for dysarthria in WD *Michaela Pernon (Paris)*

11.50 ABS 2 *TBA*

12.05 ABS 3 *TBA*

12.20 ABS 4 *TBA*

12.35 Lunch

Friday May 1
Afternoon

Session III: Epidemiology

Chairs: Tomasz Litwin, Karl Heinz Weiss

13.30	Has presenting phenotype and its relation to gender changed over the years?	<i>Karina Rewitz (Aarhus)</i>
13.45	Genotype-Phenotype Correlation and Precision Pharmacotherapy in Wilson Disease	<i>Zhi-Ying Wu (Hangzhou)</i>
14.00	Wilson disease in Brazil	<i>Marta Mitiko Deguti (Sao Paulo)</i>
14.15	ABS 5	<i>TBA</i>
14.30	ABS 6	<i>TBA</i>

Session IV: Poster session

14.45 Poster viewing and coffee break

Session V: ALF in WD

Chairs: Seema Alam, Marina Berenguer, Tudor Pop

16.00	ALF in Wilson disease	<i>Anil Dhawan (London)</i>
16.20	NCC in the diagnosis of ALF	<i>Eduardo Couchonnal (Lyon)</i>
16.40	Liver failure in Wilson disease in India	<i>Seema Alam (New Delhi)</i>
16.55	ABS 7	<i>TBA</i>
17.10	ABS 8	<i>TBA</i>
17.25	End of day	
19.00	Dinner at Martin IB, TRÆ, Kalkværksvej 5, Sydhavnen, DK-8000 Aarhus C	

Saturday May 2
Morning

Session VI: Basic science

Chairs: Hans Zischka, Roman Polishchuk

9.00	Functional Screen of Wilson Variants	<i>Svetlana Lutsenko (Baltimore)</i>
9.20	Trace Copper causes severe Iron Toxicity targeting mitochondrial Iron-Sulfur cluster enzymes	<i>Judith Sailer (Munich)</i>
9.35	ABS 9	TBA
9.50	Prion protein promotes copper toxicity in Wilson disease	<i>Roman Polishchuk (Pozzuoli)</i>
10.10	The importance of mitochondria in rescuing copper induced liver failure	<i>Hans Zischka (Munich)</i>
10.30	Coffee break	

Session VII: Screening for WD and disorders of copper metabolism

Chairs: Aftab Ala, Anil Dhawan

11.00	Current screening programs using ATP7B peptides	<i>Sihoun Hahn (Seattle)</i>
11.20	The parents' perspective	<i>Rhonda Rowland (New York)</i>
11.40	DNA-based newborn screening for disorders of copper metabolism	<i>Stephen Kaler (New York)</i>
11.55	Debate: The baby has a positive test, now what?	<i>Piotr Socha (Warsaw) /Rima Fawaz (New Haven)</i>
12.20	ABS 10	TBA
12.35	Lunch	

Saturday May 2

Afternoon

Session VIII: Bioavailable copper measurements and liver biochemistries - use for monitoring and diagnosis*Chairs: Aurélie Poujois, Anna Czlonkowska*

13.30	ANCC and CuExc	<i>Chris Harrington (Guildford)</i>
13.50	Characterization of Labile Bound Copper by a Dual Filtration Method	<i>Joshua Bornhorst (Rochester)</i>
14.05	Exchangeable copper and REC in 2026	<i>Nouzha Djebrani-Oussedik (Paris)</i>
14.25	Liver biochemistry as surrogate marker in WD	<i>Marina Berenguer (Valencia)</i>
14.40	Incorporating meaningful endpoints for treatment of WD into practice - role of NIT	<i>Michael Schilsky (New Haven)</i>
15.00	Coffee break	

Session IX: The diagnosis*Chairs: Peter Ott, Michael Schilsky*

15.30	Introduction	<i>Peter Ott (Aarhus)</i>
15.40	The value of genetic analysis	<i>Luis García Villarreal (Gran Canaria)</i>
16.00	Wilson Disease – a single gene disorder?	<i>Dorte Lildballe (Aarhus)</i>
16.15	The biopsy, total copper and metallothioneines	<i>Zoe Mariño (Barcelona)</i>
16.35	Leipzig after 2 decades of service	<i>Karl Heinz Weiss (Heidelberg)</i>
16.55	Panel discussion	
17.25	End of day	
19.00	Dinner at Havnær, Dagmar Petersens Gade 119, DK-8000 Aarhus C	

Sunday May 3

Morning

Session X: Pharmacotherapeutic advances for WD
Chairs: Zoe Mariño, Albert Stättermayer

9.00	Unmet needs in the pharmacological treatment of Wilson disease	<i>Aftab Ala (London)</i>
9.15	TDMQ20, a potential new drug for WD	<i>Bernard Meunier (Toulouse)</i>
9.30	TTM: Lost battle or a rising Phoenix?	<i>Fred Askari (Ann Arbor)</i>
9.45	ABS 11	<i>TBA</i>
10.00	⁶⁴ Cu PET update	<i>Thomas Sandahl (Aarhus)</i>
10.20	Coffee break	

Session XI: Curing WD - gene therapy and ATP7B gene repair
Chairs: Fred Askari, Tom Warner

11.00	Interim results of the ongoing Ultragenyx Phase 1-3 clinical trial of AAV9 gene therapy for Wilson Disease	<i>Andrew Grimm (Ultragenyx)</i>
11.20	Does liver fibrosis affect AAV treatment?	<i>Pasquale Piccolo (Pozzuoli)</i>
11.40	Genetic modification of ATP7B gene in human cells using RNA-based CRISPR/Cas9 technology	<i>Vanessa Sandfort (Münster)</i>
12.00	Prime editing and gene repair in WD	<i>TBA (Prime Medicine)</i>
12.15	ABS 12	<i>TBA</i>
12.30	Conference summary and take home messages	<i>Michael Schilsky (New Haven)</i>
12.45	Closing remarks	<i>Thomas Sandahl (Aarhus)</i>
12.50	Lunch	
13.50	Departure	



