ISA 2022

XVIII. INTERNATIONAL SYMPOSIUM ON AMYLOIDOSIS

4TH – 8TH SEPTEMBER 2022 HEIDELBERG

QUO VADIS AMYLO\DOSIS?

CONFERENCE GUIDE







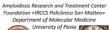


for rare or low prevalence complex diseases

Network
 Hematological
 Diseases (ERN EuroBloodNet)



















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ORGANIZATION

ISA 2022 Committees

Hosts

Stefan Schönland and Ute Hegenbart, Amyloidosis Centre, Medical Clinic V, Medical Faculty of Heidelberg University Im Neuenheimer Feld 410 69120 Heidelberg

Co-Organizers

Carlos Fernández de Larrea and Maria Teresa Cibeira, *Barcelona, Spain* Giovanni Palladini, *Pavia, Italy*

Local Organizing Committee

Markus Weiler and Fabian Siepen on behalf of the members of the Amyloidosis Center, Heidelberg, Germany

Scientific Advisory Committee Board of International Society of Amyloidosis, ISA

Giovanni Palladini, President
Stefan Schönland, President Elect,
Representative of the German Society
Yukio Ando, Past President
Shaji Kumar, Treasurer
Vaishali Sanchorawala, Secretary
Per Westermark, Editor-in-Chief of AMYLOID
Giampaolo Merlini, Representative of
the Italian Society

Ute Hegenbart, Member at Large
Paolo Milani, Member at Large
Ashutosh Wechalekar, Member at Large
Hironobu Naiki, Representative of the
Japanese Society

Board of the German Society of Amyloid Diseases, DGAK

Stefan Schönland, President Ute Hegenbart, Treasurer Alexander Carpinteiro, Secretary

Organizational Unit

Congress and Conference Management (UniKT) Heidelberg University Seminarstraße 2 69117 Heidelberg

Acknowledgements

We thank Marie Brumma (Amyloidosis Centre, Medical Clinic V) for her assistance in all organizational matters and especially for the intensive sponsor support in cooperation with UniKT.

We would like to thank our in-house agency UniKT, which brought the congress from conception to implementation with the tireless efforts of the entire team.

We are also grateful to our colleagues who will be available throughout the conference to help us manage the many tasks of a hybrid congress.

WELCOME ADDRESS

Dear all,

On behalf of the International Society of Amyloidosis we want to welcome you to the "XVIII. International Symposium on Amyloidosis" which will be held in the beautiful city center of Heidelberg. Many scientists, philosophers and poets from all over the world have a strong relationship to this unique place. We are proud that the meeting is hosted by the Ruperto Carola University, the one with the longest tradition in Germany, as it was already founded in 1386.



Stefan Schönland President elect of ISA



Ute Hegenbart
Member of ISA Board

The motto of the XVIII. meeting is "Quo vadis, amyloidosis?" The scientific and clinical developments have been overwhelming in the last years. The general interest is growing as a number of very effective drugs became available in systemic amyloidoses. This is reflected by the scientific program which will feature distinguished keynote speakers, round tables with debates, and compelling presentations by junior scientists as well as established scientists from various fields of amyloidoses. Furthermore, we have created new items like a "best abstracts" session on Thursday. The conference will be held as a hybrid meeting with more than 1100 attendees after the fully virtual conference in 2020.

We heartily thank all the members of the ISA 2022 Committee, the ISA Board and the abstract reviewers for their support in addressing the many challenges of the Corona Pandemic era and the complex circumstances in which we live. We also would like to thank the sponsors for their generous financial support and the organization of industry symposia, as well as the Amyloidosis Foundation, which again provided a large number of grants for young scientists. The winners of the awards will be honored at the Gala Dinner on Wednesday evening in the "halle02", a former building of the freight depot located in the newest district of Heidelberg named "Bahnstadt"

We wish you a pleasant and stimulating time with us in Heidelberg, which encompasses academic excellence, cutting-edge medicine, modern urban development and, last but not least, tourist highlights and German Romanticism.

Stefan Schönland and Ute Hegenbart

Abstract Book

The Book of Abstracts and also the Conference Guide can be viewed and downloaded as a flipping book on the conference website.

Badges

Participants receive their conference name badges with their registration material at the conference office or at the booth of the inviting companies and are kindly asked to wear their badges throughout the entire conference. Please use your badge to identify yourself at the external events in the old assembly hall (*Monday*), the inner courtyard of the Triplex Canteen (*Tuesday*) or the Event at halle02 (*Wednesday*).

Certificate of Attendance

You will receive a certificate of attendance with your registration documents.

CME Credits for german attendees

The ISA conference is accredited with CME Credits by the State Chamber of Physicians (Germany). 27 credits are awarded for complete attendance. In order to receive CME Credits for the conference, please contact the registration counter. Corresponding certificates of participation can be picked up at the registration counter after the event.

Coffee Breaks and Lunch

Free coffee and light refreshments will be served during the designated breaks for all participants in the foyers ground floor, second floor and in the courtyard of Neue Universität (New University). Lunch will be available at Triplex Canteen located right next

to Heidelberg University at University Square in the foyer ground floor, in the courtyard of Neue Universität (New University) and Zeughaus Canteen.

Conference office/Registration

The conference office can be found in the foyer of the New University building (Neue Universität; see Floor Plan). Opening hours will be from Sunday 4th September, 14:00 PM to Thursday, 8th September 2:00 PM during the conference hours.

Conference venue

The XVIII. International Symposium on Amyloidosis will take place in the New University building (Neue Universität), one of the central buildings of Heidelberg University. It is located at the University Square (Universitätsplatz), right in the historic center of Heidelberg city. (Address: Grabengasse 1, 69117 Heidelberg)

Copy & Print Service

If you need to copy or print a manageable amount of documents, feel free to contact our staff at the registration desk. For greater print or copy jobs we would also be pleased to refer you to one of the local copy shops in the area.

Getting around

The conference will take place right next to the University Square (*Universitätsplatz*) which is located in the very heart of the historic center of Heidelberg. Starting from there, you can reach many interesting spots within a few minutes' walk (*see Heidelberg Beyond the Conference*).

There are restaurants, hotels and souvenir shops close to the conference venue.

Therefore, you not necessarily have to rely on public transportation in order to get around.

HeidelbergCARD

The HeidelbergCARD offers you free travel on public transport and a one-time entry to the Heidelberg castle courtyard with the barrel cellar and the possibility to visit the German Pharmacy Museum. The HeidelbergCARD also enables several discounts on guided tours, in museums and exhibitions, restaurants as well as on shopping and cultural events. The HeidelbergCARD is available at the conference office.

Internet Access

During the conference all ISA2022 participants have free access to the wireless network of Heidelberg University. Please choose:

Network SSID: Uni-Webaccess

User name:

isa00000@webaccess@uni-heidelberg.de Password: Z*Kg3oRQdpbd

If your home institution participates in the eduroam project, you can also use your eduroam network login.

Lunch

The lunch will be offered in three locations. You will receive three vouchers with your registration documents, stating on which day, in which location you will receive your lunch.

- a) Triplex Canteen: This is right next to the congress building on the other side of the street. The poster presentations and viewings will also take place here.
- b) New University: In the conference building itself.

- c) Zeughaus Canteen: In the Marstallhof
- d) For all participants dessert will be served in the conference building on the ground floor and on the 2nd floor in front of the main lecture hall.

Pharmacy/Emergency phone numbers

The closest pharmacy, the "Universitäts-Apotheke" is located: Hauptstraße no. 114 opening hours: Mo – Fr, 9:00 AM – 7:00 PM,

Saturday 10:00 AM - 5:00 PM phone: +49 6221 22 514

Police phone: 110

Fire brigade and medical emergency phone: 112

Ambulance phone: +49 6221 19 222

Public Transportation

In order to get from the conference venue to Heidelberg Central Station or Bismarck Square, please take any bus from "Heidelberg, Universitätsplatz" to "Heidelberg, Bismarckplatz" or "Heidelberg, Hauptbahnhof".

For detailed timetable and fare information please visit: http://www.vrn.de

Tourist information

The tourist information can be found at the ground floor of the Town Hall, within 5 minutes walking distance from the conference venue. (Address: Marktplatz 10)

Wheelchair users

The New University building is designed to be wheelchair accessible. Access is via the inner courtyard to the ground floor or via the lift to the upper floors (*see Floor Plan*). If you require assistance, please approach our staff at the Registration Desk.

Heidelberg Beyond the Conference

- 1. Heiliggeistkirche: The Church of the Holy Spirit was constructed in 1398. Facing the Town Hall, the late Gothic church is located at the Market place (Marktplatz) of Heidelberg and has been both Catholic and Protestant during its long history.
- 2. Alte Brücke: The famous Old Bridge was erected in 1786 1788 by Elector Karl Theodor and consists of barrel vaults of red sandstone.
- 3. Studentenkarzer & Universitätsmuseum: In the so-called Student Prison students were imprisoned between 1778 and 1914. The university could place students under arrest for up to four weeks.
- 4. Universitätsbibliothek: The University Library was constructed in 1905 combining elements of German Mannerism of the late 16th century with Art Nouveau motifs. The University Library has over 3 million books. 1.5 million loans each

- year and is one of the finest buildings of the university- definitely worth a look.
- 5. Heidelberger Schloss: Presumably built in between the 13th and 15th century, Heidelberg Castle served as one of the main seats of the Prince Electors of the Palatinate. It was continually enlarged by Ruprecht III and his successors, but destroyed by French troops during the War of Palatine Succession. Even today, the castle ruins still rise majestically over the roofs of the Old Town one of the absolute must-see sights during a stay in Heidelberg.
- **6. Philosophenweg:** The Philosopher's Path is one of Germany's most famous walks. Especially during the Romantic era many poets, famous thinkers and other "children of Heidelberg" like Hölderlin, Eichendorff and Scheffel, to name a few examples came here to relax, stroll around, as well as to be inspired by the nature and by the stunning view over the Neckar valley.



Special Events

Conference Welcome Reception

Sunday, September 4, 7:00 PM after the opening lectures Conference Building Neue Universität

Merlini Award Ceremony

Monday, September 5, 7:05 PM Building Alte Universität

Get Together with Poster Presentations and Viewing of late breaking abstracts, Challenging Cases and Junior Meets Senior

Tuesday, September 6, 7:00 PM
Triplex Canteen, foyer, 1st floor and courtyard

Award Ceremony, Conference Dinner

Wednesday, September 7, 7:00/7:30 PM halle02

Zollhofgarten 2, 69115 Heidelberg How to get there? Please see the information distributed at the congress.

Poster Presentations and Poster Viewings

There are four Poster Viewings between Monday and Wednesday (*see program*).

The poster presentations and viewings will take place in the neighbouring building Triplex Canteen on the 1st floor.

Poster Viewing hours:

Monday – Wednesday 11:30 AM – 4:30 PM Tuesday 6:00 PM – 10:00 PM

The posters will be mounted by the presenters themselves. An assistant will be on site at the following times:

Monday-Wednesday: 7:00~AM-8:00~AM

and 11:30 AM – 12:00 PM Tuesdav: 6:00 PM – 7:00 PM

Prices and tips

In restaurants, cafes, bars, etc. prices include taxes and service. A tip of approximately 5-10% is appreciated but not obligatory.

Smoking

Due to the strict non-smoking policy with regard to public buildings, smoking is prohibited in all conference venues as well as in most of the restaurants, cafes, etc. Outside the New University building, there are smoking areas and smoking is generally permitted in the street.

Virtual Venue - Venueless

You will find a link to the virtual venue via your Conftool registration access data. Please go to https://www.conftool.com/isaheidelberg2022/ and login. You will find the link on your Conftool Welcome page.

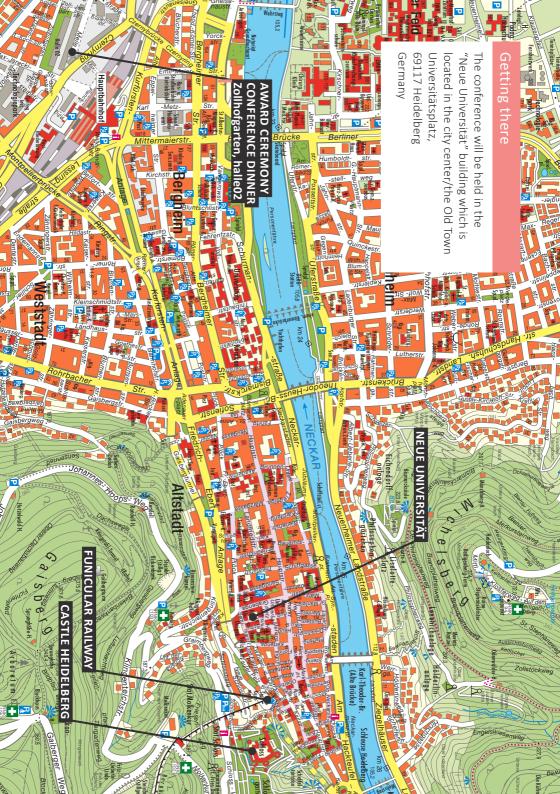
Wardrobe

There is an unguarded wardrobe in H02 on the ground floor (see Floor Plan).

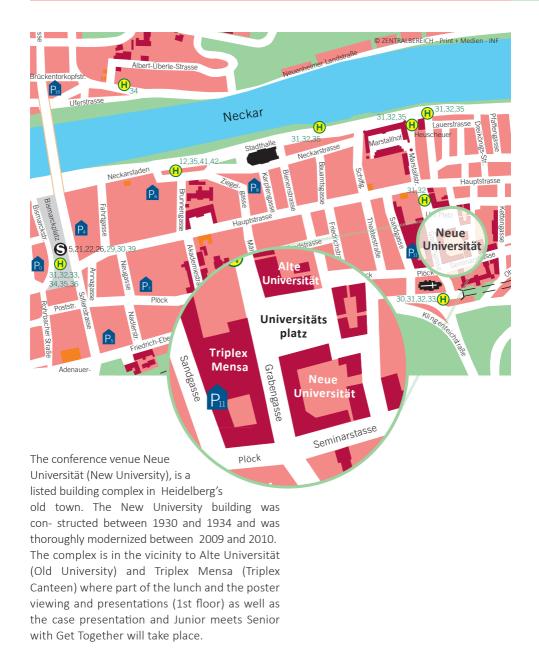
Lockers – if necessary – are to be found in the University Library (*Universitätsbibliothek*), just across the road (*about 50 m*).

Additional lecture room H14

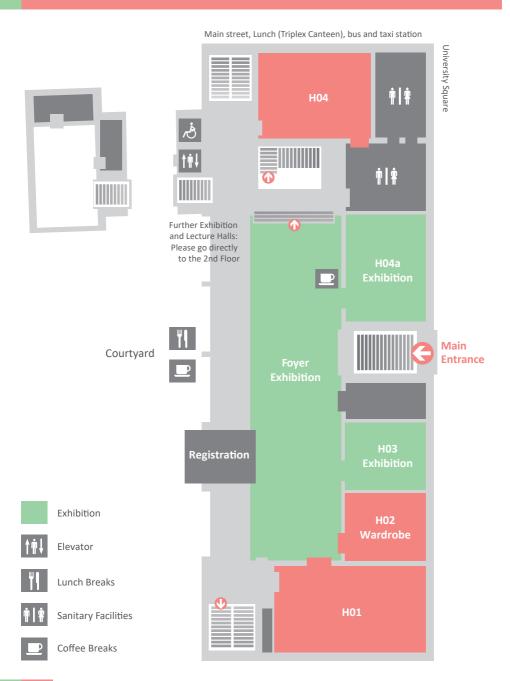
The entire program from Main Lecture Hall will be streamed to H14. Exceptions are the few cases when H14 is used for a parallel event.



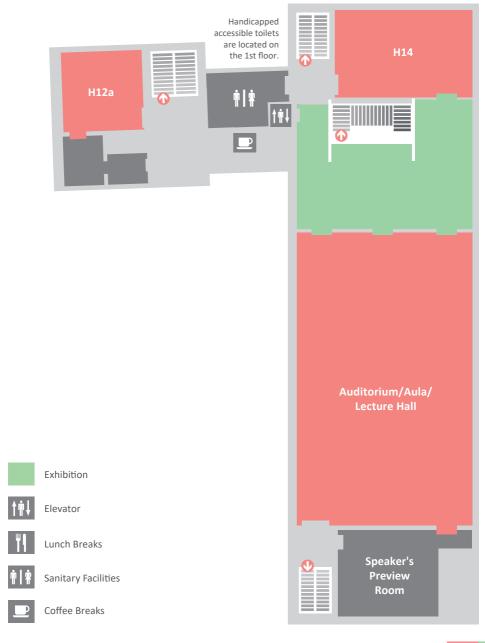
VENUE



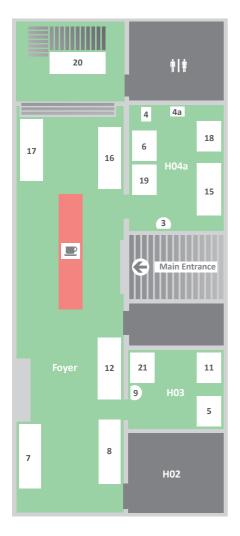
FLOORPLAN - GROUND FLOOR



FLOORPLAN – 2ND FLOOR



EXHIBITION FLOORPLAN & EXHIBITOR LIST



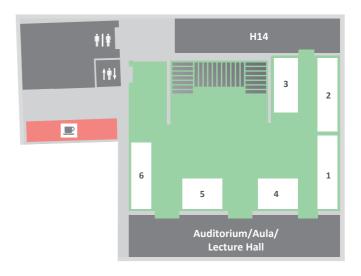
GROUND FLOOR

- 3 Life Molecular Imaging
- 4 Amyloidose Alliance
- 4a Myeloma UK
- 5 AbbVie
- 6 Sanofi Aventis
- 7 Alnylam
- 8 Janssen-Cilag
- 9 Attralus
- 11 Binding Site
- 12 Alexion/AstraZeneca
- **15** Sobi™
- 16 Ionis/AstraZeneca UK
- 17 Pfizer
- 18 Bristol Myers Squibb
- 19 GlaxoSmithKline
- 20 International Society of Amyloidosis/ Mayo Clinic
- 21 Protego

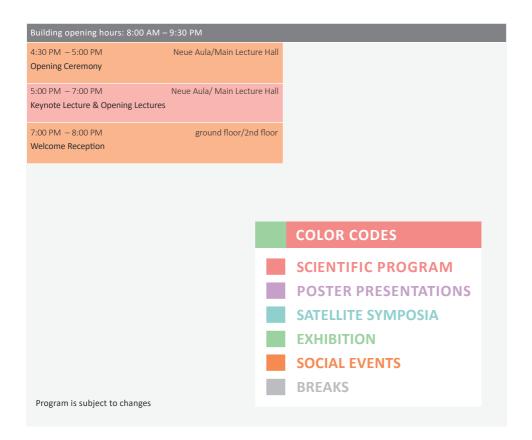
EXHIBITION FLOORPLAN & EXHIBITOR LIST

2nd FLOOR

- 1 Ionis/AstraZeneca UK
- 2 Prothena
- 3 BridgeBio
- 4 Alnylam
- 5 Alexion/AstraZeneca
- 6 Pfizer



PROGRAM OVERVIEW – SUNDAY, 4TH



PROGRAM OVERVIEW – MONDAY, 5TH

Building opening hours: 7:00 AM – 9:30 PM		
	7:00 AM – 7:45 AM Sponsored breakfast	2nd floor
7:15 AM – 8:30 AM Neue Aula/Main Lecture Hall IKMG/ISA – Challenges in Monoclonal Gammopathy of Renal Significance	7:30 AM – 8:30 AM Global Bridges: Building Diagno:	H14 stic Capacity Worldwide
8:30 AM — 10:05 AM Neue Aula/Main Lecture Hall Basic research — Light chains including cardiotoxicity OP001 — 008		9:30 AM – 5:00 PM ground floor/2nd floor
10:05 AM – 10:30 AM Virtual Venue Virtual Speaker Corner: Talk with industry partners	10:05 AM – 10:30 AM New University, ground floor/2nd floor/courtyard Coffee Break	Exhibition on-site and virtual
10:30 AM – 12:05 PM Neue Aula/Main Lecture Hall ATTRwt – Clinical aspects OP009 – 016		
12:05 PM — 1:20 PM Triplex Canteen/1st floor Poster Presentation: AL and ATTR amyloidosis	12:05 PM – 1:20 PM New University, ground floor/ courtyard and Triplex Canteen Lunch Break	
1:20 PM – 3:00 PM Neue Aula/Main Lecture Hall Imaging in amyloidosis OP017 – 024		
3:05 PM — 4:10 PM Neue Aula/Main Lecture Hall Preclinical models of systemic amyloidosis		
	4:10 PM – 4:30 PM New University, ground floor/2nd floor/courtyard Coffee Break	
4:30 PM – 6:00 PM Neue Aula/Main Lecture Hall Janssen: Navigating the patient journey in AL amyloidosis: a multidisciplinary approach		
6:00 PM $-$ 7:00 PM Neue Aula/Main Lecture Hall Pfizer: How can one solution solve the multifaceted challenges of ATTR amyloidosis?		
7:05 PM — 8:05 PM Alte Aula/Building "Alte Universität" Merlini Award Ceremony		

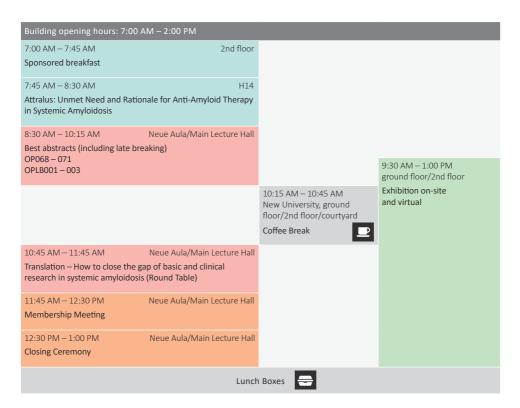
PROGRAM OVERVIEW – TUESDAY, 6TH

Building opening hours: 7:00	AM – 9:30 PM		
7:00 AM – 7:30 AM	2nd floor		
Sponsored breakfast			
7:30 AM – 8:30 AM	Neue Aula/Main Lecture Hall		
Prothena: Addressing the Unm AL Amyloidosis: Key Insights fro			
8:30 AM – 10:05 AM	Neue Aula/Main Lecture Hall		
AL – Clinical aspects OP025 – 0	32		
		10:05 AM – 10:30 AM New University, ground floor/2nd floor/courtyard Coffee Break	9:30 AM – 5:00 PM ground floor/2nd floor Exhibition on-site and virtual
10:30 AM – 12:05 PM	Neue Aula/Main Lecture Hall		
Basic Research – New treatmen OP033 – 040	nt targets and biomarkers		
12:05 PM – 1:15 PM Triplex Canteen/1st floor	12:05 PM – 1:15 PM Virtual Venue	12:05 PM – 1:15 PM New University, ground floor/	
Poster Presentation: Basic science and imaging	Virtual Speaker Corner: Talk with industry partners	courtyard and Triplex Canteen Lunch Break	
1:15 PM – 2:30 PM AA – Clinical aspects OP041 – C	Neue Aula/Main Lecture Hall 043		
2:30 PM — 3:30 PM BridgeBio: Improving patient o	Neue Aula/Main Lecture Hall utcomes in ATTR-CM		
3:30 PM - 5:00 PM	Neue Aula/Main Lecture Hall		
Ionis/AstraZeneca: Optimizing Patients With ATTR	Multidisciplinary Care in		
5:00 PM – 7:00 PM Free Time/Explore Heidelberg			Heidelberg
7:00 PM – 7:30 PM Poster Presentation: Late break	Triplex Canteen/1st floor king abstracts	7:00 PM – 10:00 PM Get Together	Triplex Canteen/Foye
7:30 PM – 8:15 PM Challenging Cases	Triplex Canteen/Foyer		
8:15 PM – 9:15 PM Junior Meets Senior (Round Tal	Triplex Canteen/Foyer		

PROGRAM OVERVIEW – WEDNESDAY, 7TH

Building opening hours: 7:00 AM – 9:30 PM		
7:00 AM – 7:45 AM 2nd floor		
Sponsored coffee		
7:45 AM – 8:30 AM Neue Aula/Main Lecture Hall		
Sobi™: Expert Discussion: Is Disease Remission in hATTR a Realistic Goal?		
8:30 AM – 10:05 AM Neue Aula/Main Lecture Hall		
ATTRv – Clinical aspects OP044 – 051		9:30 AM – 5:00 PM ground floor/2nd floor
9:00 AM – 10:00 AM H14		Exhibition on-site
EHA/ISA – Better understanding and targeting the clone in AL amyloidosis		and virtual
	10:05 AM – 10:30 AM New University, ground floor/2nd floor/courtyard	
	Coffee Break	
10:30 AM – 12:05 PM Neue Aula/Main Lecture Hall		
Pathways to diagnosis OP052 – 059		
12:05 PM – 1:20 PM Triplex Canteen/1st floor	12:05 PM – 1:20 PM	
Poster Presentation: Pathway to diagnosis, Innovative drugs and non-AL / non-ATTR amyloidosis	New University, ground floor/ courtyard and Triplex Canteen	
and non-AL / non-ALTR arryioloosis	Lunch Break	
1:20 PM – 3:00 PM Neue Aula/Main Lecture Hall		
Basic research – Amyloid fibril formation including proteolysis and tissue interactions OP060 – 067		
3:00 PM – 4:30 PM Neue Aula/Main Lecture Hall		
Alexion: Beyond Survival: Unmet Medical Needs in AL Amyloidosis		
4:30 PM – 5:00 PM Virtual Venue	4:30 PM - 5:00 PM	
Virtual Speaker Corner: Talk with industry partners	New University, ground floor/2nd floor/courtyard	
	Coffee Break	
5:00 PM – 6:30 PM Neue Aula/Main Lecture Hall		
Alnylam: Meeting the needs of patients with hATTR amyloidosis: innovation in practice		
7:00/7:30 PM – open end "halle02" Heidelberg Conference Dinner/ Award Ceremony		

PROGRAM OVERVIEW - THURSDAY, 8TH





"One thinks Heidelberg by day

- with its surroundings —
is the last possibility of the beautiful;
but when he sees Heidelberg by night,
a fallen Milky Way, with that glittering
railway constellation pinned to the
border, he requires time to consider
upon the verdict."

Mark Twain

DETAILED PROGRAM

SUNDAY, 4TH – THURSDAY, 8TH

DETAILED PROGRAM – SUNDAY, 4TH

4:30 PM - 5:00 PM

SOCIAL EVENT

Neue Aula/Main Lecture Hall

Opening Ceremony

Musical accompaniment

Kathrin Christians, Flute, Sonia Akchar, Piano

Ute Hegenbart, host ISA 2022, Amyloidosis Center of the University

Hospital, Heidelberg, Germany

Giovanni Palladini, ISA president, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo,

Pavia, Italy

Christian Schaaf, Representative of the Center of Rare Diseases,

University Hospital, Heidelberg, Germany

5:00 PM - 7:00 PM

SCIENTIFIC SESSION

Neue Aula/Main Lecture Hall

Keynote Lecture & Opening Lectures

Chairs

Stefan Schönland, host ISA 2022, Amyloidosis Center of the University

Hospital, Heidelberg, Germany

Giovanni Palladini, ISA president, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San

Matteo, Pavia, Italy

Introduction

Stefan Schönland

Keynote Lecture

Aging (of the Immune System)

Cornelia Weyand, Stanford University, Stanford, and Mayo Clinic,

Rochester, USA

Opening Lecture

AL Amyloidosis model – leading the way to cure

Giampaolo Merlini, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

Introduction

Giovanni Palladini

Opening Lectures

Turning swords into plowshares: the case of functional amyloid

Daniel Otzen, Aarhus University, Aarhus, Denmark

New horizons of gene targeting therapy in ATTRv amyloidosis

Julian Gillmore, National Amyloidosis Centre, London, United Kingdom

7:00 PM - 8:00 PM

SOCIAL EVENT

ground floor/2nd floor

Welcome Reception

Get Together with served snacks and beverages

9:30 AM – 5:00 PM EXHIBITION ground floor/2nd floor

Exhibition on-site and virtual

7:30 AM – 8:30 AM SATELLITE SYMPOSIUM H14
Sponsored breakfast starts at 7:00 AM

Global Bridges: Building Diagnostic Capacity Worldwide

Global Bridges: Building Diagnostic Capacity Worlds

Amyloidosis Grantee Presentations

Chairs Morie Gertz, Mayo Clinic, Rochester, USA

Martha Grogan, Mayo Clinic, Rochester, USA

Introduction Morie Gertz

Presentations Building Awareness and Improving the Quality of Diagnosis

Sharmila Dorbala, Brigham and Women's Hospital, USA

Firas Al Badarin, Cleveland Clinic Abu Dhabi, UAE

Bruno Bueno, Heart Institute, Hospital Das Clinicas, U Sao Paolo, Brazil

Isabel Carvajal Juárez, ASNC Latin America, Mexico Special patient populations/leveraging technology

Saurabh Malhotra, Cook County Health, Illinois, USA

Paolo Milani, Amvloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

Establishing Regional Centers of Excellence Ramzi Tabbalat, Abdali Medical Center, Jordan

Silvia Lupu, Emergency Institute for Cardiovascular Disease and

Transplant, Romania

Kwaku Appiah-Kubi, C.K. Tedam University of Technology and Applied

Science, Ghana

Summary/Close Martha Grogan

7:15 AM – 8:30 AM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

IKMG/ISA - Challenges in Monoclonal Gammopathy of Renal

Significance Joint Session

Chairs Ute Hegenbart, host ISA 2022, Amyloidosis Center of the University

Hospital, Heidelberg, Germany

Nelson Leung, Mayo Clinic, Rochester, USA

Lectures Differential diagnosis of MGRS

Guillermo Herrera, University of South Alabama, USA

Treatment endpoints in MGRS

Angela Dispenzieri, Mayo Clinic, Rochester, USA

MGRS: directions for future research

Frank Bridoux, Hôpital Jean Bernard, Poitiers, France

8:30 AM - 10:05 AM SCIENTIFIC SESSION

Neue Aula/Main Lecture Hall

Basic research - light chains including cardiotoxicity

Chairs Stefano Ricagno, Università di Milano, Milano, Italy

Marina Ramirez-Alvarado, Mayo Clinic, Rochester, USA

State-of-the-art lecture Stefano Ricagno

OP001 Next Generation Sequencing Identifies AL-related IGLV Genes in

Patients with λ -isotype MGUS or Smoldering Multiple Myeloma

Ray Comenzo, Tufts Medical Center, Boston, USA

OP002 Elucidation of the cardiotoxicity of full-length light chains derived

from patients with cardiac light chain amyloidosis in comparison to

other plasma cell dyscrasias

Panagiota-Efstathia Nikolaou, National and Kapodistrian University,

Athens. Greece

OP003 Investigation of IGLV2-14 AL amyloidosis and multiple myeloma light

chain sequences

Natalie Berghaus, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

OP004 Sequence diversity of the kappa light chains from patients with AL

amyloidosis and multiple myeloma

Sarah Schreiner, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

OP005 Single Molecule Real-Time Sequencing of the M protein (SMaRT

M-Seg): toward personalized medicine approaches in monoclonal

gammopathies

Mario Nuvolone, Amyloidosis Research and Treatment Center, Univer-

sity of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

OP006 An N-glycosylation hotspot in immunoglobulin κ light chains is

associated with AL amyloidosis

Alice Nevone, Amyloidosis Research and Treatment Center, University of

Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

OP007 Amyloidogenesis by mesangial cells involves active participation of

Ivsosomes

Guillermo A. Herrera, University of South Alabama, Alabama, USA

OP008 Mapping and modelling the molecular mechanisms that drive

amyloidogenic light chain induced cardiotoxicity

Camille Vanessa Edwards, Boston University, Boston, USA

10:05 AM - 10:30 AM



New University, ground floor/2nd floor/courtyard

10:05 AM - 10:30 AM EXHIBITION

Virtual Venue

Virtual Speaker Corner Talk with industry partners

10:30 AM -12:05 PM

Neue Aula/Main Lecture Hall

ATTRwt - Clinical aspects

Chairs: Thibaud Damy, Referral Center for Cardiac Amyloidois, Henri Mondor

Hospital, Creteil, France

Matthew Maurer, Columbia University, New York, USA

State-of-the-art lecture Thibaud Damy

OP009 Long-term tafamidis treatment reduces the decline in quality of life

among patients with transthyretin amyloid cardiomyopathy

Martha Grogan, Mayo Clinic, Rochester, USA

Atrial fibrillation as a prognostic factor for all-cause mortality in **OP010**

> patients with transthyretin amyloid cardiomyopathy Ronald Witteles, Stanford University, Stanford, USA

OP011 Long-term safety and tolerability of acoramidis (AG10) in

symptomatic transthyretin amyloid cardiomyopathy: Updated analysis

from an ongoing phase 2 open-label extension study

Ahmad Masri, OHSU, Portland, USA

OP012 Chronic Intravenous Inotropic Therapy in Cardiac Amyloidosis

Johana Rocio Fajardo, Medstar Heart & Vascular Institute, Washington,

USA

OP013 Tolerability and side-effects of therapy in an open-label trial of

inotersen for transthyretin amyloid cardiomyopathy

Leo C. Samuels, Brigham and Women's Hospital, Boston, USA

OP014 Looking over your shoulder to catch amyloidosis earlier:

shoulder pathologies are significantly more prevalent in patients with

transthyretin cardiac amyloidosis

Alyssa Basdavanos, Cleveland Clinic Foundation, Cleveland, USA SARS-CoV-2 infection in systemic amyloidosis: the International

Society of Amyloidosis' survey

Paolo Milani, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy Cardiac Transplantation in Transthyretin Amyloid Cardiomyopathy:

Outcomes from Three Decades of Tertiary Centre Experience Yousuf Razvi, National Amyloidosis Centre, London, United Kingdom

12:05 PM - 1:20 PM

OP015

OP016

LUNCH New University, ground floor/courtyard & Triplex Canteen

12:05 PM - 1:20 PM

POSTER PRESENTATIONS

Triplex Canteen, 1st floor

Poster Numbers: P001 - P121, P208: AL and ATTR amyloidosis For the virtual audience the poster and an audio summary by the presenting authors will be provided in the Virtual Venue.

AL amyloidosis

Chairs Vaishali Sanchorawala, Amyloidosis Center at Boston University School

of Medicine, Boston, USA

Paolo Milani, Amyloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

Maite Cibeira, Amyloidosis and Myeloma Unit, Barcelona, Spain Timon Hansen, Haematology-Oncology Clinic, Altona, Hamburg,

Germany

Hermine Agis, Department of Medicine, Medical University of Vienna,

Vienna, Austria

Angela Dispenzieri, Mayo Clinic, Rochester, USA Jeffrey Zonder, Karmanos Cancer Institute, Detroit, USA

ATTRwt amvloidosis

Chairs Thibaud Damy, Referral Center for Cardiac Amyloidois, Henri Mondor

Hospital, Creteil, France

Matthew Maurer, Columbia University, New York, USA Pablo Garcia-Pavia, Hospital Universitario Puerta de Hierro

Majadahonda, Madrid, Spain

Maria Papathanasiou, University Hospital Essen, Germany

ATTRv amyloidosis

Chairs Ernst Hund, Amyloidosis Center of the University Hospital, Heidelberg,

Germany

Katrin Hahn, Amyloidosis Center, Charité, Berlin, Germany Isabel Maria Conceicao, Centro Hospitalar Universitário Lisboa

Norte – Hospital de Santa Maria, Lisboa, Portugal

Markus Weiler, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

Imaging in amyloidosis

Chairs Ashutosh Wechalekar, National Amyloidosis Centre, London, United

Kingdom

Sharmila Dorbala, Brigham and Women's Hospital, Boston, USA

State-of-the-art lecture Amyloid Imaging

Ashutosh Wechalekar

State-of-the-art lecture Cardiac Imaging

Sharmila Dorbala

OP017 Results of the first-in-human PET/CT imaging study of the amyloid-

reactive peptide 124I-AT-01 (124I-p5+14) for the detection of

systemic amyloidosis

Jonathan Wall, University of Tennessee, Knoxville, USA

OP018 Pan-amyloid reactivity of radioiodinated peptide 124I-AT-01 in

patients with systemic amyloidosis demonstrated by PET/CT imaging

Emily Martin, University of Tennessee, Knoxville, USA

OP019 Tracking multi-organ treatment response in systemic AL-amyloidosis

with cardiac magnetic resonance derived extracellular volume mapping

Adam Ioannou, Royal Free Hospital, London, United Kingdom

OP020 Regression of Cardiac Bone-Tracer Uptake in Patients with Hereditary

Transthyretin Amyloidosis after One Year Treatment with Patisiran.

An early Marker of Treatment Response?

Hendrea S.A. Tingen, University Medical Center, Groningen, Netherlands

OP021 Real World Experience of Tc-DPD scintigraphy as a diagnostic imaging

tool in amyloidosis

Muhammad Umaid Rauf, National Amyloidosis Centre, London, United

Kingdom

OP022 Assessing left ventricular strain in cardiac amyloidosis:

the importance of accurate measurement technique

Rodney H. Falk, Brigham and Women's Hospital, Boston, USA

OP023 Quantitative magnetic resonance neurography biomarkers: Cross-

> sectional results from a single center study in 80 subjects with symptomatic or asymptomatic hereditary transthyretin amyloidosis

Jennifer Hayes, Amyloidosis Center of the University Hospital, Heidelberg

Assessment of the clinical value of whole-body MRI in untreated OP024

patients with systemic light chain amyloidosis

Simone Christine Brandelik, Amyloidosis Center of the University

Hospital, Heidelberg, Germany

3:05 PM - 4:10 PM

Neue Aula/Main Lecture Hall

Preclinical models of systemic amyloidosis

Chairs Alexander Carpinteiro, Amyloidosis Center of the University Hospital,

Essen, Germany

Gunilla Westermark, Uppsala University, Uppsala, Sweden

Introduction Alexander Carpinteiro

A mouse model of AL amyloidosis Lectures

Christophe Sirac, University of Limoges, Limoges, France

Insights into ATTR amyloidosis from a new transgenic mouse model Paul Simons, National Amyloidosis Centre, London, United Kingdom

AL amyloidosis modelled in Caenorhabditis elegans Luisa Diomede, Mario Negri IRCCS, Milano, Italy

Debate Christophe Sirac, Paul Simons, Luisa Diomede

4:10 PM - 4:30 PM



New University, ground floor/2nd floor/courtyard

4:30 PM - 6:00 PM

Neue Aula/Main Lecture Hall

Janssen: Navigating the patient journey in AL amyloidosis:

a multidisciplinary approach

Chairs Giovanni Palladini, Amyloidosis Research and Treatment Center, Univer-

> sity of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy Marianna Fontana, National Amyloidosis Centre, London, United

Kingdom

Introduction Giovanni Palladini

Lecture Diagnostic pit-falls and risk stratification in AL amyloidosis

Marianna Fontana

Panel Discussion Frank Bridoux, Hôpital Jean Bernard, Poitiers, France

Darren Foard, National Amyloidosis Centre, London, United Kingdom

Christoph Röcken, University Hospital, Kiel, German

Lecture Key considerations when selecting treatment for patients with

AL amyloidosis
Giovanni Palladini

Panel Discussion Frank Bridoux, Darren Foard, Christoph Röcken

Summary/Close Giovanni Palladini

6:00 PM - 7:00 PM SATELLITE SYMPOSIUM Neue Aula/Main Lecture Hall

Pfizer: How can one solution solve the multifaceted challenges of

ATTR amyloidosis?

Chairs Arnt Kristen, Amyloidosis Center of the University Hospital, Heidelberg,

Germanv

Introduction Arnt Kristen

Lectures Transthyretin in health and disease

Laura Obici, Amyloidosis Research and Treatment Center, University of

Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

Improving patient outcomes in ATTR-PN

Laura Obici

Addressing cardiac involvement in patients with ATTR amyloidosis

Arnt Kristen

Clinical clues to cardiac involvement in ATTR amyloidosis –

how to identify patients early

Arnt Kristen, Rachele Bonfiglioli, Azienda Ospedaliero-Universitaria,

Bologna, Italy

Panel Discussion/Close Arnt Kristen, Laura Obici, Rachele Bonfiglioli

7:05 PM — 8:05 PM SOCIAL EVENT

Alte Aula/Building "Alte Universität"

Merlini Award Ceremony Musical accompaniment

Kathrin Christians, Flute, Sonia Akchar, Piano

Laudatio Giovanni Palladini, Amyloidosis Research and Treatment Center,

University of Pavia and IRCCS Fondazione Policlinico San Matteo,

Pavia, Italy

Lecture Amyloidosis: Reflections on passed and coming times

Per Westermark, University Hospital, Uppsala, Sweden

Merlini Award Winner 2022



DETAILED PROGRAM - TUESDAY, 6TH

9:30 AM – 5:00 PM EXHIBITION ground floor/2nd floor

Exhibition on-site and virtual

7:30 AM – 8:30 AM SATELLITE SYMPOSIUM Neue Aula/ Main Lecture Hall

Sponsored breakfast starts at 7:00 AM

Prothena: Addressing the Unmet Need in Advanced AL Amyloidosis:

Key Insights from a Panel of Experts

Chair Morie Gertz, Mayo Clinic, Rochester, USA

Introduction Morie Gertz

Lectures Epidemiology and Unmet Need in Advanced AL Amyloidosis

Maria Papathanasiou, *University Hospital Essen, Germany*

Diagnosis and Current Treatment Approaches in AL Amyloidosis Giovanni Palladini, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

Emerging Therapies in Advanced AL Amyloidosis

Morie Gertz

Panel Discussion Morie Gertz, Maria Papathanasiou, Giovanni Palladini

8:30 AM – 10:05 AM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

AL - Clinical aspects

Chairs Shaji Kumar, Mayo Clinic, Rochester, Minnesota, USA

Vaishali Sanchorawala, Amyloidosis Center at Boston University School

of Medicine, Boston, USA

State-of-the-art lecture Shaji Kumar

OP025 Predictors of hematologic response and survival with stem cell

transplantation in AL amyloidosis: a 25-year longitudinal study Joshua Gustine, *Amyloidosis Center at Boston University School of*

Medicine, Boston, USA

OP026 The prognostic importance of flow cytometry-based measurable

residual disease (MRD) in patients with systemic light chain

amyloidosis

Andrew Staron, Amyloidosis Center at Boston University School of

Medicine, Boston, USA

DETAILED PROGRAM – TUESDAY, 6TH

OP027	Prognostic Impact of	Translocation t(11:14) and of other cytogenetic

abnormalities in patients with AL amyloidosis in the era of

contemporary therapies

Despina Fotiou, National and Kapodistrian University, Athens, Greece

OP028 Incidence and risk factors of sudden death in patients with

cardiacamyloidosis

Fernando de Frutos, Hospital Universitario Puerta de Hierro,

Majadahonda, Spain

OP029 Clonal features affect survival of patients with non-cardiac light chain

(AL) amyloidosis: a European study of 386 patients

Paolo Milani, Amyloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

Light chain deposition disease: an international study in 523 patients OP030

Paolo Milani, Amyloidosis Research and Treatment Center,

University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia,

Italy

OP031 Elotuzumab in combination with IMIDS for AL amyloidosis patients

with relapsed/refractory plasma cell dyscrasia and advanced organ

involvement

Tobias Dittrich, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

OP032 First Report on Outcome of Patients with Newly Diagnosed Stage IIIb

Cardiac Light Chain Amyloidosis Treated with Daratumumab-based

Frontline Therapy

Rajshekhar Chakraborty, Irving Medical Center, Columbia, USA



10:05 AM – 10:30 AM COFFEE BREAK New University, ground floor/2nd floor/courtyard

10:30 AM - 12:05 PM

SCIENTIFIC SESSION

Neue Aula/Main Lecture Hall

Basic Research - New treatment targets and biomarkers

Chairs Gareth Morgan, Amyloidosis Center at Boston University School of

Medicine, Boston, USA

Carlos Fernández de Larrea, Amyloidosis and Myeloma Unit,

Barcelona, Spain

DETAILED PROGRAM – TUESDAY, 6TH

State-of-the-art lecture Gareth Morgan

OP033 Collagen associated with AL amyloid inhibits fibril phagocytosis -

Collagen degradation renders amyloid sensitive to uptake by the

innate immune system

Joseph W. Jackson, University of Tennessee, Knoxville, USA

OP034 In-vitro ultrasonic assay indicates importance of extracellular

chaperon-like effect of serum albumin to protect dialysis patients

from dialysis-related amyloidosis

Kichitaro Nakajima, Osaka University, Osaka, Japan

OP035 Dissecting FAM46C-dependent tuning of antibody secretion in

systemic AL amyloidosis

Enrico Milan, San Raffaele Scientific Institute, Milano, Italy

Development of novel human chimeric antigen receptor-**OP036**

macrophages (CAR-M) as a potential therapeutic for amyloid

clearance

Manasi Balachandran, University of Tennessee, Knoxville, USA

OP037 Preclinical characterization of AT-02, a pan-amyloid-binding

immunoglobulin-peptide fusion protein capable of inducing

enhanced phagocytosis of amyloid

Jonathan Wall, University of Tennessee, Knoxville, USA

OP038 Regulation of BCL2 family members by microRNA-9 and microRNA-

181a in AL amyloidosis

Oshrat Hershkovitz-Rokah, Assuta Medical Center, Tel Aviv, Israel

OP065 AA amyloid-containing diet potentiates Aβ42 induced effects in

transgenic Drosophila melanogaster

Gunilla Westermark, Uppsala University, Department of Medical Cell

Biology, Uppsala University, Uppsala, Sweden

OP040 Targeting protein secretion as a therapeutic strategy in AL amyloidosis

Maria Moscvin, Brigham and Women's Hospital, Boston, USA

12:05 PM - 1:15 PM



LUNCH New University, ground floor/courtyard & Triplex Canteen

DETAILED PROGRAM – TUESDAY, 6TH

12:05 PM – 1:15 PM POSTER PRESENTATIONS

Triplex Canteen, 1st floor

Poster Numbers: P122 – P190: Basic science and Imaging
For the virtual audience the poster and an audio summary by the

presenting authors will be provided in the Virtual Venue.

Fibril and Amyloid Formation

Chairs Mario Nuvolone, Amyloidosis Research and Treatment Center,

University of Pavia and IRCCS Fondazione Policlinico San Matteo,

Pavia, Italy

Christoph Röcken, University Hospital, Kiel, Germany

Joost Schymkowitz, Leuven Center for Brain & Disease Research,

Leuven, Belgium

Mechanism of organ dysfunction and damage

Chairs Marina Ramirez-Alvarado, Mayo Clinic, Rochester, USA

Stefano Ricagno, Università di Milano, Milano, Italy

Pre-clinical disease models

Chairs Gunilla Westermark, Uppsala University, Uppsala, Sweden

Alexander Carpinteiro, University Hospital Essen, Germany

Imaging

Chairs Simon Gibbs, The Victorian and Tasmanian Amyloidosis Service,

Australian Amyloidosis Network, Australia

Jennifer Hayes, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

Jonathan Wall, University of Tennessee, Knoxville, USA

Martha Grogan, Mayo Clinic, Rochester, USA

12:05 PM - 1:15 PM

EXHIBITION

Virtual Venue

Virtual Speaker Corner
Talk with industry partners

DETAILED PROGRAM – TUESDAY, 6TH

1:15 PM – 2:30 PM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

AA - Clinical aspects

Chairs Norbert Blank, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

Sophie Georgin-Lavialle, Sorbonne University, Paris, France

State-of-the-art lecture Norbert Blank

Lectures AA Amyloidosis in Europe, Perspectives and Challenges

Sophie Georgin-Lavialle

AA Amyloidosis in Africa, Perspectives and Challenges Ghalia Khellaf, *Centre Hospitalo-Universitaire*, *Algeria*

OP041 Tocilizumab can prevent the progression of renal AA amyloidosis to

end stage renal disease

Peter Kvacskay, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

OP042 Natural history and risk stratification of AA amyloidosis based on a

40-year experience in the United States

Tracy Joshi, Amyloidosis Center at Boston University School of Medicine,

Boston, USA

OP043 Long-term transplant outcomes in recipients with renal amyloidosis

Tale Norbye Wien, Bærum Hospital Vestre Viken, Drammen, Norway

2:30 PM — 3:30 PM SATELLITE SYMPOSIUM Neue Aula/Main Lecture Hall

BridgeBio: Improving patient outcomes in ATTR-CM

Chairs Julian Gillmore, National Amyloidosis Centre, London, United Kingdom

Matthew S. Maurer, Columbia University, New York, USA

Introduction Julian Gillmore, Mathew S. Maurer Lectures ATTR-CM consensus – what's new?

Claudio Rapezzi, University of Ferrara, Ferrara, Italy

Present and future in ATTR-CM

Pablo Garcia-Pavia, Hospital Universitario Puerta de Hierro

Majadahonda, Madrid, Spain

Q&A/Close Julian Gillmore, Mathew S. Maurer

DETAILED PROGRAM – TUESDAY, 6TH

3:30 PM – 5:00 PM SATELLITE SYMPOSIUM Neue Aula/Main Lecture Hall

Ionis/AstraZeneca: Optimizing Multidisciplinary Care in Patients

With ATTR

Chair Arnt Kristen, *Amyloidosis Center of the University Hospital, Heidelberg,*

Germany

IntroductionArnt KristenPatient Journey VideoArnt Kristen

ATTR: Disease Overview

Arnt Kristen

Lecture Optimizing Patient Care: A Neurologist's Perspective

Teresa Coelho, Centro Hospitalar Universitário, Porto, Portugal

Interactive Case Studies: Neurology

Panel Discussion Arnt Kristen, Teresa Coelho,

Pablo Garcia-Pavia, Hospital Universitario Puerta de Hierro

Majadahonda, Madrid, Spain

Lecture Optimizing Patient Care: A Cardiologist's Perspective

Pablo Garcia-Pavia

Interactive Case Studies: Cardiology

Panel Discussion/Q&A Arnt Kristen, Teresa Coelho, Pablo Garcia-Pavia

Close Arnt Kristen

5:00 PM - 7:00 PM Heidelberg

Free Time/Explore Heidelberg

We have posted some nice options for sightseeing and leisure tours on

our ISA 2022 homepage.

7:00 PM – 10:00 PM SOCIAL EVENT Triplex Canteen Foyer/1st floor/Courtyard

Get Together

Get Together with served snacks and beverages

07:00 PM – 07:30 PM POSTER PRESENTATIONS Triplex Canteen, 1st floor

Poster Numbers: PLB001 - 017: Late breaking abstracts

For the virtual audience the poster and an audio summary by the

presenting authors will be provided in the Virtual Venue.

Chairs Suzanne Lentzsch, Columbia University, New York, USA

Per Westermark, Uppsala University, Uppsala, Sweden

7:30 PM – 8:15 PM SCIENTIFIC SESSION Triplex Canteen, Foyer

Challenging Cases

Three cases of diagnostic and therapeutic challenges will be presented during this session. Panelists and audience will participate to solve these

difficult cases.

Chairs Vaishali Sanchorawala, Amyloidosis Center at Boston University School

of Medicine, Boston, USA

Maite Cibeira, Amyloidosis and Myeloma Unit, Barcelona, Spain Paolo Milani, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo,

Pavia, Italy

Panel of Experts Angela Dispenzieri, Mayo Clinic, Rochester, USA

Julian Gillmore, National Amyloidosis Centre, London, United Kingdom

Pablo Garcia-Pavia, Hospital Universitario Puerta de Hierro

Majadahonda, Madrid, Spain

DETAILED PROGRAM – TUESDAY, 6[™]

8:15 PM - 9:15 PM

SOCIAL EVENT

Triplex Canteen, Foyer

Junior Meets Senior
Round Table

The path of a scientific career is often long and sometimes difficult. The goal of this session is to provide junior physicians and scientists in the audience a chance to interact with experts in the field of Amyloidosis and beyond. Both, young and established colleagues might profit from sharing their experience in diverse areas, including their academic careers, approach to research and publishing, and also more personal aspects like work-life balance.

Chairs

Fabian aus dem Siepen, Amyloidosis Center of the University Hospital, Heidelberg, Germany Eloisa Riva, Hospital de Clínicas "Dr. Manuel Quintela", Montevideo, Uruguay

Senior Experts

Per Westermark, Uppsala University, Uppsala, Sweden Suzanne Lentzsch, Columbia University, New York, USA Cornelia Weyand, Stanford University, Stanford, and Mayo Clinic, Rochester, USA

Jörg Goronzy, Stanford University, Stanford and Mayo Clinic, Rochester, USA Martha Grogan, Mayo Clinic, Rochester, USA

Carsten Müller-Tidow, University Hospital, Heidelberg, Germany Norbert Frey, University Hospital, Heidelberg, Germany



DETAILED PROGRAM - WEDNESDAY, 7TH

9:30 AM – 5:00 PM EXHIBITION ground floor/2nd floor
Exhibition on-site and virtual

7:45 AM – 8:30 AM SATELLITE SYMPOSIUM Neue Aula/Main Lecture Hall

Sponsored coffee starts at 7:00 AM

Sobi™: Expert Discussion: Is Disease Remission in hATTR a Realistic

Goal?

Chair Teresa Coelho, Centro Hospitalar Universitário, Porto, Portugal

Introduction Teresa Coelho

Lectures Evolution of diagnosis and management of hATTR

Katrin Hahn, Amyloidosis Center, Charité, Berlin, Germany

"Flipping the pyramid": lessons learned from MS

Gavin Giovannoni, Queen Mary University, London, United Kingdom

Are we primed to make remission a reality?

Marco Luigetti, *Università Cattolica del Sacro Cuore, Rome, Italy* Teresa Coelho, Katrin Hahn, Gavin Giovannoni, Marco Luigetti

Conclusion Teresa Coelho

Panel Discussion

8:30 AM – 10:05 AM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

ATTRv - Clinical aspects

Chairs David Adams, Université Paris Saclay, Paris, France

Marcia Waddington-Cruz, Federal University of Rio de Janeiro, Rio de

Janeiro, Brazil

State-of-the-art lecture David Adams

OP044 Patisiran Global Open-label Extension Study at 36 Months: Effect

of Long-term Treatment on Mortality and Ambulatory Function in

Patients with hATTR Amyloidosis with Polyneuropathy

Jonas Wixner, Umeå University, Umeå, Sweden

OP045 Progression and distribution pattern of cerebral amyloid angiopathy

in hereditary ATTR amyloidosis patients visualized by 11C-PiB-PET

imaging

Yusuke Takahashi, Shinshu University, Matsumoto, Japan

OP046 An international Delphi survey for the definition of a multidisciplinary

holistic approach to the care of hereditary ATTR amyloidosis

Laura Obici, Amvloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

OP047 Eplontersen in ATTR-polyneuropathy: results from the 35-week

interim analysis of NEURO-TTRansform

Teresa Coelho, Centro Hospitalar Universitário, Porto, Portugal Diflunisal treatment for hereditary transthyretin amyloidosis –

the Swedish DENS-02 trial

Jonas Wixner, Umeå University, Umeå, Sweden

OP049 Kidney phenotype and immune activation in hereditary ATTR

amyloidosis during inotersen therapy

Joana Tavares, Centro Hospitalar Universitário do Porto, Porto, Portugal

Association of hereditary V122I amyloidogenic transthyretin variant OP050

with heart failure: A systematic review and meta-analysis Kwaku Appiah-Kubi, C. K. Tedam University, Navrongo, Ghana

OP051 Tafamidis reduces skin denervation and amyloid in skin biopsies of

very early symptomatic hATTRv patients after one year of treatment

Aleiandra Gonzalez-Duarte, INCMNSZ, Ciudad de México, Mexico

9:00 AM - 10:00 AM SCIENTIFIC SESSION

EHA/ISA - Better understanding and targeting the clone in

AL amyloidosis

Chairs Efstathios Kastritis, National and Kapodistrian University, Athens,

Monique Minnema, University Medical Center, Utrecht, Netherlands

Lectures Future Myeloma therapies suitable for AL amyloidosis

Marc Raab, Myeloma Center, University Heidelberg, Germany

Marginal zone lymphoma and M. Waldenström as underlying disease

in AL amyloidosis

Davide Rossi, Oncology Institute of Southern Switzerland, Bellinzona,

Switzerland



10:05 AM – 10:30 AM COFFEE BREAK New University, ground floor/2nd floor/courtyard

H14

OP048

DETAILED PROGRAM – WEDNESDAY, 7TH

10:30 AM - 12:05 PM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

Pathways to diagnosis

Chairs Hironobu Naiki, University of Fukui, Fukui, Japan

Kerstin Amann, University Hospital, Erlangen, Germany

State-of-the-art lecture Hironobu Naiki

OP052 Automated cardiac amyloidosis risk detection on whole body bone

scintigraphy using deep-learning approach

Marc-Antoine Delbarre, University Hospital, Amiens, France

OP053 Establishment of the nation-wide pathology consultation system for

the typing diagnosis of amyloidosis in Japan: Steep increase in the

number of transthyretin-positive cardiac biopsy cases

Hironobu Naiki

OP054 The amyloid proteome: a two-way approach for a protein

classification system

Juliane Gottwald, University Hospital, Kiel, Germany

OP055 Fat aspiration as a screening tool for symptomatic systemic

amyloidosis - sensitivity and specificity analysis with better sensitivity

results for females with all major subtypes

Christoph Richard Kimmich, University Hospital, Oldenburg, Germany

OP056 Sequence of diagnostic testing in cardiac amyloidosis patients: early

monoclonal protein study is associated with better outcomes in AL

amvloidosis

Francesca Fabris, Amyloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

OP057 Artificial intelligence modeling for earlier identification of cardiac

amyloidosis

Surendra Dasari, Mayo Clinic, Rochester, USA

A novel mass spectrometry-based method for the identification of OP058

> subtype specific amyloidogenic proteins from fat aspirates Hans Christian Beck, University Hospital, Odense, Denmark

OP059 Pre-symptomatic diagnosis of systemic AL amyloidosis by biomarker-

based screening in patients with MGUS

Silvia Mangiacavalli, Amyloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

12:05 PM - 1:20 PM



LUNCH New University, ground floor/courtyard & Triplex Canteen

12:05 PM - 1:20 PM

POSTER PRESENTATIONS

Triplex Canteen, 1st floor

Poster Numbers: P191 - P295 (except P208, moved to Monday):

Pathway to diagnosis, Innovative drugs and non-AL/

non-ATTR amyloidosis

For the virtual audience the poster and an audio summary by the presenting authors will be provided in the Virtual Venue.

Pathway to Diagnosis

Chairs Eli Muchtar, Mayo Clinic, Rochester, USA

Rahel Schwotzer, University Hospital, Zurich, Switzerland

Fabian aus dem Siepen, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

Eloisa Riva, Hospital de Clínicas "Dr. Manuel Quintela", Montevideo,

Uruguay

Sandra Ihne, Amyloidosis Center, University Hospital Würzburg, Germany

Hans Nienhuis, Amyloidosis Centre of Expertise, University Medical

Center, Groningen, Netherlands

Kerstin Amann, University Hospital, Erlangen, Germany

Hironobu Naiki, University of Fukui, Fukui, Japan

Arnaud Jaccard, CHU, Limoges, France

AA Amyloidosis

Chair Norbert Blank, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

Biology, Clinics and Therapeutics in other types of amyloidosis

Laura Obici, Amyloidosis Research and Treatment Center, University of

Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

Innovative Therapies

Chairs Efstathios Kastritis, National and Kapodistrian University, Athens, Greece

Gareth Morgan, Amyloidosis Center at Boston University School of

Medicine, Boston, USA

Peter Mollee, Princess Alexandra Hospital, Brisbane, Queensland,

Australia

Chairs

DETAILED PROGRAM – WEDNESDAY, 7TH

1:20 PM – 3:00 PM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

Basic research – Amyloid fibril formation including proteolysis and

tissue interactions

Chairs Matthias Schmidt, *University Ulm, Ulm, Germany*

Christoph Röcken, University Hospital, Kiel, Germany

Stefan Schönland, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

State-of-the-art lecture Matthias Schmidt, Christoph Röcken, Stefan Schönland

OP060 Kinetics of the aggregation process of the human λ -III

immunoglobulin light chain FOR005 involved in AL amyloidosis at

atomic resolution

Tejaswini Pradhan, Technical University, Munich, Germany

OP061 The ex vivo cryo-EM structure of AA amyloid from a domestic short

hair cat

Tim Schulte, Gruppo San Donato, San Donato Milanese, Italy

OP062 The interplay between protein dynamics and proteolysis in

LC amyloid aggregation

Stefan Ricagno, Università di Milano, Milano, Italy

OP063 Time-resolved nano-spectroscopy with single-molecule sensitivity for

blood-based non-immune diagnosis of amyloid diseases

Ann Tiiman, Center for Molecular Medicine (CMM), Department of Clinical Neuroscience, Karolinska Institute, Stockholm, Sweden

OP064 In vitro and in vivo effects of SerpinA1 on the modulation of

transthyretin proteolysis

Maria Rosario Almeida, Universidade do Porto, Molecular Biology

Department, Porto, Portugal

OP039 Elevated fibrosis associated biomarkers in ATTR amyloidosis patients

are associated with impaired cardiovascular outcome

Selina Julia Hein, Amyloidosis Center of the University Hospital,

Heidelberg, Germany

OP066 Dichotomy of Circulating Non-native TTR (NNTTR) Levels in

Polyneuropathy and Cardiomyopathy Patients Provides a Glimpse to

ATTR Tissue Specificity

Xin Jiang, Protego Biopharma, San Diego, USA

OP067 Heterotypic amyloid interactions and their effect on amyloid assembly

Frederic Rousseau, VIB-KU Leuven Center for Brain and Disease

Research, Leuven, Belgium

3:00 PM – 4:30 PM SATELLITE SYMPOSIUM Neue Aula/Main Lecture Hall

Alexion: Beyond Survival: Unmet Medical Needs in AL Amyloidosis

Chair Vaishali Sanchorawala, Amyloidosis Center at Boston University School

of Medicine, Boston, USA

Introduction Vaishali Sanchorawala

Lecture Looking Beyond Plasma Cell Dyscrasia in AL Amyloidosis

Rodney Falk, Brigham and Women's Hospital, Boston, USA

Panelists Julian Gillmore, National Amyloidosis Centre, London, United Kingdom

Giovanni Palladini, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo.

Pavia, Italy

Vaishali Sanchorawala

Ashutosh Wechalekar, National Amyloidosis Centre, London, United

Kingdom

Lecture Unmet Medical Needs in Managing Tissue Toxicity and Organ

Dysfunction

Giovanni Palladini

Panelists Rodney Falk, Julian Gillmore, Vaishali Sanchorawala,

Ashutosh Wechalekar

Lecture Beyond Survival - Assessing and Monotoring Organ and Tissues

Ashutosh Wechalekar

Panelists Rodney Falk, Julian Gillmore, Giovanni Palladini, Vaishali Sanchorawala

Lecture Continuing Conversation to Learn from Patient Insights

Giovanni Palladini

Panelists Deborah Boedicker, Amyloidosis Speakers Bureau, Washington, DC,

USA

Kristen Hsu, Amyloidosis Research Consortium, Newton, USA

Q&A Vaishali Sanchorawala

Rodney Falk, Giovanni Palladini, Ashutosh Wechalekar

Close Vaishali Sanchorawala

COFFEE BREAK New University, ground floor/2nd floor/courtyard

4:30 PM - 5:00 PM

4:30 PM - 5:00 PM Virtual Venue Virtual Speaker Corner Talk with industry partners 5:00 PM - 6:30 PM Neue Aula/Main Lecture Hall Alnylam: Meeting the needs of patients with hATTR amyloidosis: innovation in practice Chair Thomas Skripuletz, Department of Neurology, Hannover Medical School, Hannover, Germany Introduction Thomas Skripuletz The origin and evolution of treatment Thomas Skripuletz Evolving experience with RNAi therapies in a multisystem disease – Experience of managing hATTR amyloidosis at a reference centre Marcus Anthony Urey, Division of Cardiology, Department of Medicine, University of California, San Diego, USA Evolving experience with RNAi therapies in a multisystem disease – A cardiologist's perspective on hATTR amyloidosis

Neurofilament Light Chain (NfL) as a biomarker in hATTR amyloidosis
Hans Nienhuis, *Amyloidosis Centre of Expertise, University Medical*Contag Croningen Natherlands

Andoni Echaniz-Laguna, APHP, CHU de Bicêtre, Le Kremlin-Bicêtre,

Center, Groningen, Netherlands

Redefining standards of care

France

Laura Obici, Amyloidosis Research and Treatment Center, University of Pavia and IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

7:00/7:30 PM – open end

SOCIAL EVENT

"halle02" Heidelberg

Conference Dinner/Award Ceremony

The Conference Dinner is held in the largest event venue in Heidelberg located in the building of the former freight station. It offers conference participants an excellent chance to enjoy delicious food in a unique industry ambience while getting to know each other and making new connections outside the main conference environment. Finally, the winner of the ISA 2022 presidential awards and poster awards will be presented.

The halle02 offers more than 1,500 sqm of inspiring rooms and 1,200 sqm of outdoor space for events of all kinds and is exclusively available to conference participants for this event.

How to get there? *Please see the information distributed at the congress*



DETAILED PROGRAM – THURSDAY, 8TH

ground floor/2ndfloor

Exhibition on-site and virtual 7:45 AM - 8:30 AM H14 Sponsored coffee starts at 7:00 AM Attralus: Unmet Need and Rationale for Anti-Amyloid Therapy in **Systemic Amyloidosis** Chair Ashutosh Wechalekar, National Amyloidosis Centre, London, United Kingdom Ashutosh Wechalekar Introduction Lectures Unmet need in Systemic Amyloidosis Morie Gertz, Mayo Clinic, Rochester, USA Preclinical Rationale for Anti-Amyloid therapy Marina Ramirez-Alvarado, Mayo Clinic, Rochester, USA Review of Efficacy with Anti-Amyloid therapy Ashutosh Wechalekar Safety of Anti-Amyloid Approach Arnt Kristen, Amyloidosis Center of the University Hospital, Heidelberg,

Discussion/Close Ashutosh Wechalekar, Morie Gertz, Marina Ramirez-Alvarado,

Arnt Kristen

Germany

9:30 AM - 1:00 PM

8:30 AM – 10:15 AM SCIENTIFIC SESSION Neue Aula/Main Lecture Hall

Best abstracts (including late breaking)

Chairs Markus Weiler, Amyloidosis Center Heidelberg, Germany

Peter Mollee, University of Queensland, Australi

OP068 Feasibility of a novel academic BCMA-CART (HBI0101) for the

treatment of relapsed and refractory AL amyloidosis Moshe Gatt, *Hadassah Medical Center, Jerusalem, Israel*

DETAILED PROGRAM – THURSDAY, 8TH

OP069 A Proteomic Atlas of Renal Amyloid Plagues Provides Insights Into

Disease Pathogenesis

Charalampos Charalampous, Mayo Clinic, Rochester, USA

Cryo-EM study of cardiac ATTR fibrils and structure-based **OP070**

development of detection probes and anti-seeding inhibitors

Lorena Saelices, UT Southwestern, Dallas, USA

OP071 An European collaborative study on 476 patients with AA amyloidosis:

identification and validation of survival and renal staging systems

Marco Basset, Amyloid Research and Treatment Center, IRCCS

Fondazione Policlinico San Matteo, Pavia, Italy

OPI B001 Primary Results From APOLLO-B, A Phase 3 Study Of Patisiran In

Patients With Transthyretin-Mediated Amyloidosis With

Cardiomyopathy

Matthew Maurer, Columbia University, New York, USA

OPLB002 Venetoclax targeted therapy in t(11;14) AL amyloidosis patients:

a retrospective analysis from the French Amyloidosis Network

Murielle Roussel, CHU Duputyren, Limoges, France

OPLB003 Amyloidosis-related orthopedic events, low plasma transthyretin,

and risk of cardiac events

Anders Moller Greve, Rigshospitalet, Copenhagen, Denmark

10:15 AM - 10:45 AM COFFEE BREAK



New University, ground floor/2nd floor/courtyard

10:45 AM - 11:45 AM SCIENTIFIC SESSION

Neue Aula/ Main Lecture Hall

Translation – How to close the gap of basic and clinical research in systemic amyloidosis

Round Table

Chairs Stefan Schönland, host ISA 2022, Amyloidosis Center of the University

Hospital, Heidelberg, Germany

Vittorio Bellotti, IRCCS Fondazione Policlinico San Matteo, Pavia, Italy

Panel of Experts Marina Ramirez-Alvarado, Mayo Clinic, Rochester, USA

Joost Schymkowitz, Leuven Center for Brain & Disease Research,

Leuven, Belgium

Jonathan Wall, University of Tennessee, Knoxville, USA

Arnaud Jaccard, CHU, Limoges, France

Gunilla Westermark, Uppsala University, Uppsala, Sweden

DETAILED PROGRAM – THURSDAY, 8TH

11:45 AM - 12:30 PM SOCIAL EVENT

Neue Aula/Main Lecture Hall

ISA Membership Meeting

The outgoing President, Giovanni Palladini, will report on the scientific and organisational activities of ISA during the last 2 years.

Shaji Kumar, treasurer will give his financial report.

Per Westermark, editor in chief of Amyloid, will give a summary on the recent developments of our journal.

Vaishali Sanchorawala, secretary, will present the election results (board and bylaws).

Stefan Schönland, President elect, will acknowledge the outgoing Board members, welcome and introduce the new Board members and give his prospects for the next 2 years.

12:30 PM - 1:00 PM

SOCIAL EVENT

Neue Aula/Main Lecture Hall

Closing Ceremony

Chairs

ISA 2022 Organizing Committee

Summary ISA 2022 Heidelberg

Bouke Hazenberg, University Medical Center, Groningen, Netherlands

Outlook ISA 2024

Morie Gertz, Martha Grogan, Angela Dispenzieri, Mayo Clinic,

Rochester, USA

LUNCH BOXES



Thank you for your participation and have a safe trip back home!

Detailed information will be provided in the abstract book which will be published on the ISA website.

Poster Presentations – Monday, 5th – 12:05 PM – 1:20 PM

P001	Morbidity and mortality measured through "Days Alive and Out of Hospital" (DAOH) in patients with AL amyloidosis according to cardiac involvement and specific treatment. María LOURDES POSADAS MARTINEZ
P002	Selection of appropriate quality of life instrument to measure patient- reported outcomes in systemic light chain (AL) amyloidosis Anita D'SOUZA
P003	Symptom burden and quality of life in AL amyloidosis patients among recently diagnosed and long-term survivors Anita D'SOUZA
P004	Performance of a survival staging system incorporating sST2 in patients with light chain amyloidosis Darae KIM
P005	Evaluation of the Patterns Leading to Diagnosis in Patients With Amyloid Light-Chain Amyloidosis Using the Komodo Database Julia CATINI
P006	Healthcare resource utilization in patients with light chain amyloidosis in Europe Efstathios KASTRITIS
P007	Reduction of cardiac AL amyloid deposition after complete response visualized by PiB-PET imaging Nagaaki KATOH
P008	The impact of longitudinal strain (LS) response in patients with advanced cardiac AL amyloidosis Joshua BOMSZTYK
P009	CMR with T1 mapping in systemic light-chain (AL) amyloidosis: from cardiac amyloid regression to refining treatment response Ana MARTINEZ-NAHARRO
P010	Outcomes for patients with systemic light chain (AL) amyloidosis and Mayo stage 3B disease Foteini THEODORAKAKOU
P011	Finding a needle in a haystack: oligomer detection via urinary extracellular vesicles in AL light chain amyloidosis Shawna Ann COOPER

P012	Longitudinal testing for measurable residual disease (MRD) using multiparametric flow cytometry in patients with systemic light chain amyloidosis Andrew STARON
P013	Racial differences in the cytogenetic underpinnings of light chain amyloidosis Andrew STARON
P014	Evaluation of NT-proBNP as surrogate endpoint in AL amyloidosis: development of a platform for federated, multi-institution meta-analysis of randomized trials Angela DISPENZIERI
P015	Patterns of target organ amyloid deposition in patients with AL amyloidosis; role for diagnosis and prognosis Despina FOTIOU
P016	Diagnostic Hospitalization and Associated Costs in Patients with Amyloid Light-Chain Amyloidosis Tiffany P. QUOCK
P017	Atypical neurological presentation of immunoglobulin light-chain amyloidosis Michelle PIROTTE
P018	Characterization and outcome of patients with systemic AL amyloidosis requiring dialysis prior to initial therapy Lilli Sophie SESTER
P019	Beta-2-microglobulin and lactate dehydrogenase as prognostic parameters in light-chain amyloidosis Sara OUBARI
P020	Diagnostic value of liver stiffness as marker of hepatic amyloid deposition in systemic AL amyloidosis Anne Floor BRUNGER
P021	The Pattern of Organ Responses Varies in Patients with Systemic Light- chain Amyloidosis (AL) and Heart or Kidney or Heart and Kidney Involvement Who Achieve Deep Hematologic Responses Diana ZHANG
P022	Outcomes of patients with AL amyloidosis and end- stage renal disease requiring dialysis Foteini THEODORAKAKOU
P023	Identifying symptoms of AL amyloidosis in electronic health records using natural language processing, ICD codes, and manually abstracted registry data Eli SILVERT
P024	A revised renal staging system for long-term renal survival in patients with AL amyloidosis with renal involvement Marco ALLINOVI

 Functional Status and Heart Failure Quality of Life Provide Incremental Prognostic Value in Light Chain Amyloidosis Jocelyn CANSECO NERI Heart and autologous stem cells transplantation in AL amyloidosis. María Adela AGUIRRE P028 Droxidopa for Treatment of Refractory Orthostatic Hypotension in Patients with AL Amyloidosis: A Case Series Jorge Nicolas RUIZ LOPEZ P029 AL Amyloidosis – a reason to transplant MGUS phenotype? Maximilian Johannes STEINHARDT P030 Daratumumab for the treatment of Relapsed/Refractory AL amyloidosis: Experience from the Amyloidosis Program of Calgary (APC) Daniel LEVIN P031 Efficacy and safety of daratumumab monotherapy in newly diagnosed patients with stage 3b light chain amyloidosis: a phase 2 study by the European Myeloma Network Efstathios KASTRITIS P032 Role of Doxycycline in the treatment of patients with AL amyloidosis receiving Bortezomib-containing regimens in the frontline setting: Experience from the Amyloidosis Program of Calgary Ellen LEWIS P033 Effect of the presence of t(11;14) for patients with AL amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary Ellen LEWIS P034 Cyclophosphamide, Bortezomib and Methylprednisolone for the treatment AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis 	P025	Role of subcutaneous abdominal fat tissue aspiration in the diagnosis of systemic immunoglobulin light-chain amyloidosis David F MORENO
 P027 Heart and autologous stem cells transplantation in AL amyloidosis. María Adela AGUIRRE P028 Droxidopa for Treatment of Refractory Orthostatic Hypotension in Patients with AL Amyloidosis: A Case Series Jorge Nicolas RUIZ LOPEZ P029 AL Amyloidosis – a reason to transplant MGUS phenotype? Maximilian Johannes STEINHARDT P030 Daratumumab for the treatment of Relapsed/Refractory AL amyloidosis: Experience from the Amyloidosis Program of Calgary (APC) Daniel LEVIN P031 Efficacy and safety of daratumumab monotherapy in newly diagnosed patients with stage 3b light chain amyloidosis: a phase 2 study by the European Myeloma Network Efstathios KASTRITIS P032 Role of Doxycycline in the treatment of patients with AL amyloidosis receiving Bortezomib-containing regimens in the frontline setting: Experience from the Amyloidosis Program of Calgary Ellen LEWIS P033 Effect of the presence of t(11;14) for patients with AL amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary Ellen LEWIS P034 Cyclophosphamide, Bortezomib and Methylprednisolone for the treatment AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis 	P026	Functional Status and Heart Failure Quality of Life Provide Incremental Prognostic Value in Light Chain Amyloidosis
in Patients with AL Amyloidosis: A Case Series Jorge Nicolas RUIZ LOPEZ P029 AL Amyloidosis — a reason to transplant MGUS phenotype? Maximilian Johannes STEINHARDT P030 Daratumumab for the treatment of Relapsed/Refractory AL amyloidosis: Experience from the Amyloidosis Program of Calgary (APC) Daniel LEVIN P031 Efficacy and safety of daratumumab monotherapy in newly diagnosed patients with stage 3b light chain amyloidosis: a phase 2 study by the European Myeloma Network Efstathios KASTRITIS P032 Role of Doxycycline in the treatment of patients with AL amyloidosis receiving Bortezomib-containing regimens in the frontline setting: Experience from the Amyloidosis Program of Calgary Ellen LEWIS P033 Effect of the presence of t(11;14) for patients with AL amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary Ellen LEWIS P034 Cyclophosphamide, Bortezomib and Methylprednisolone for the treatment AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis	P027	Heart and autologous stem cells transplantation in AL amyloidosis.
P030 Daratumumab for the treatment of Relapsed/Refractory AL amyloidosis: Experience from the Amyloidosis Program of Calgary (APC) Daniel LEVIN P031 Efficacy and safety of daratumumab monotherapy in newly diagnosed patients with stage 3b light chain amyloidosis: a phase 2 study by the European Myeloma Network Efstathios KASTRITIS P032 Role of Doxycycline in the treatment of patients with AL amyloidosis receiving Bortezomib-containing regimens in the frontline setting: Experience from the Amyloidosis Program of Calgary Ellen LEWIS P033 Effect of the presence of t(11;14) for patients with AL amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary Ellen LEWIS P034 Cyclophosphamide, Bortezomib and Methylprednisolone for the treatment AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis	P028	in Patients with AL Amyloidosis: A Case Series
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receiving Bortezomib-containing regimens in the frontline setting: Experience from the Amyloidosis Program of Calgary Ellen LEWIS P033 Effect of the presence of t(11;14) for patients with AL amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary Ellen LEWIS P034 Cyclophosphamide, Bortezomib and Methylprednisolone for the treatment AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis	P031	diagnosed patients with stage 3b light chain amyloidosis: a phase 2 study by the European Myeloma Network
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AL amyloidosis: Updated report from the Amyloid Program of Calgary (APC) Ellen LEWIS P035 Safety and Efficacy of Propylene Glycol-Free Melphalan in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis	P033	amyloidosis treated with Bortezomib-containing regimens: Experiences from the Amyloidosis Program of Calgary
Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study Michelle H. LEE P036 Treatment outcomes according to salvage chemotherapy modalities for relapsed/refractory AL amyloidosis	P034	
modalities for relapsed/refractory AL amyloidosis	P035	Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Results of a phase II study
YOUN SAING EUIN	P036	

P037	Autologous stem cell transplantation in primary amyloidosis: a single centre experience
	Tugrul ELVERDI
P038	Bortezomib-based induction therapy is associated with superior hematologic responses and survival after stem cell transplantation in patients with AL amyloidosis
P039	Joshua N GUSTINE Venetoclax in Relapsed or Refractory AL Amyloidosis with t(11;14) and BCL2 overexpression. Pamella PAUI
P040	Birtamimab in Patients with Mayo Stage IV AL Amyloidosis: Rationale for Confirmatory AFFIRM-AL Phase 3 Study Morie A. GERTZ
P041	Assessing clinical outcomes in patients with AL amyloidosis across different criteria for hematologic complete response: Results from ANDROMEDA Giovanni PALLADINI
P042	Efficacy of bortezomib based regimens in elderly patients with newly diagnosed AL amyloidosis and heart failure. David GARRIDO
P043	Time to next treatment (TtNT) is an independent prognostic marker for outcome in newly diagnosed patients with AL amyloidosis Despina FOTIOU
P044	Second autologous stem cell transplantation as salvage therapy in selected patients with relapsed/progressed light-chain (AL) amyloidosis M. Teresa CIBEIRA
P045	IsAMYP: a phase 2 single-stage study to evaluate the efficacy of isatuximab, pomalidomide and dexamethasone, in patients with AL amyloidosis not in VGPR or better after any previous therapy. Murielle ROUSSEL
P046	Epidemiology of light-chain amyloidosis in Latin America: a retrospective analysis of 212 patients. Grupo Latinoamericano de Estudio del Mieloma Múltiple (GELAMM) Erika BRULC
P047	Timeline change of AL amyloidosis treatment response and clinical outcomes; a single-center experience from Turkey Metban GUZEL MASTANZADE
P048	Autologous Stem Cell Transplantation in AL amyloidosis in two centers from Latin America Eloísa RIVA

P049	based therapy in patients with light chain (AL) amyloidosis and high plasma cell burden evaluated at 3 months Claudia BELLOFIORE
P050	Comparison of bortesomib-based induction regimens with other treatment modalities in patients with newly diagnosed systemic light chain amyloidosis Olga V. KUDYASHEVA
P051	Autologous stem cell transplantation (ASCT) remains effective therapy for systemic immunoglobulin light chain amyloidosis (AL): Experience from a single Australian Amyloidosis Service Tran Bing Andrew GIANG
P052	Daratumumab, pomalidomide and dexamethasone (DPd) in relapsed/refractory light chain amyloidosis previously exposed to daratumumab (NCT04270175): Interim results Cara A. ROSENBAUM
P053	Functional Status and Heart Failure Quality of Life Improve Following Therapy in Light Chain Amyloid Cardiomyopathy Jocelyn CANSECO NERI
P054	Atrial Fibrillation Does Not Influence the Occurrence of Cerebrovascular Accidents Among Patients with Amyloidosis Ramtej ATLURI
P055	Morbidity and mortality measured through "Days Alive and Out of Hospital" (DAOH) in patients with amyloidosis. María Lourdes POSADAS MARTÍNEZ
P056	Factors associated with morbidity and mortality, measured through "Days Alive and Out of Hospital" (DAOH) in patients with AL and ATTR amyloidosis. María Lourdes POSADAS MARTÍNEZ
P057	Significant tricuspid regurgitation is associated with adverse outcomes in patients with transthyretin amyloidosis. Eugenia VILLANUEVA
P058	Prevalence of hereditary transthyretin amyloidosis among elderly patients with transthyretin cardiomyopathy Eva CABRERA ROMERO
P059	1 Clinical findings and comorbidities in wtATTR patients with suspected amyloid neuropathy Sasha ZIVKOVIC
P060	Identification of wild-type transthyretin cardiac amyloidosis in patients with recent CTS surgery Bertil LADEFOGED

P061	Evolution of demographics of patients with transthyretin amyloid cardiomyopathy over time: implications for disease awareness strategies and future trial design Claudio RAPEZZI
P062	Descriptive analysis of women with transthyretin amyloid cardiomyopathy: examining the patient demographics of a growing patient population Martha GROGAN
P063	Assessment and Management of Older Patients with Transthyretin Amyloidosis Cardiomyopathy: Geriatric Cardiology, Frailty Assessment and Beyond Biobelemoye IRABOR
P064	Preservation of Left Ventricle Stroke Volume in Patients with ATTRwt Cardiac Amyloidosis Treated with Selective TTR Stabiliser Tafamidis Dariusz KORCZYK
P065	Implementation of a machine learning model to assess transthyretin amyloid cardiomyopathy risk in an external platform George KOUTITAS
P066	Clinical and Socioeconomic Differences Among Patients with Transthyretin Cardiac Amyloidosis Belonging to North and South of Chicago. Mukunthan MURTHI
P067	Relationship of tafamidis binding site occupancy, transthyretin stabilization, and disease modification in tafamidis treated transthyretin amyloid cardiomyopathy patients Amy T. MOODY
P068	Diagnostic Path, Clinical Characteristics and Outcomes of Patients With ATTR Cardiomyopathy in Greece Foteini THEODORAKAKOU
P069	Screening for transthyretin-related amyloidosis in patients with aortic stenosis undergoing aortic valve replacement Isabel MATTIG
P070	Case series of the treatment journeys of patients who underwent heart transplantation for transthyretin (ATTR) cardiac amyloidosis, with subsequent confirmed orthopedic disease Jill WALDRON
P071	Factors Associated with Financial Toxicity in Patients with ATTR: Results From Amyloidosis Research Consortium's ATTR Treatment Affordability Patient and Caregiver Survey Sabrina REBELLO
P072	Characterisation of Austrian Transthyretin Amyloid Cardiomyopathy (ATTR-CM) patients enrolled in a Tafamidis (61mg) early access program Christoph GROJER

P073	Incidence and Risk Factors for Pacemaker Implantation in Light Chain and Transthyretin Cardiac Amyloidosis Aldostefano PORCARI
P074	The relationship between NT-proBNP and perception of the severity of cardiac symptoms in TTR-CA: the moderating role of anxious and depressive symptoms Francesco CAPPELLI
P075	Neurological symptoms of patients with transthyretin amyloidosis at first neurological presentation at the Amyloidosis Center of Lower Saxony Janna Margaretha SIEMER
P076	Incidence and factors associated with de novo atrial fibrillation in patients with wild-type transthyretin cardiac amyloidosis Carlo FUMAGALLI
P077	Quality of Life in Patients with Transthyretin Amyloid Cardiomyopathy Treated with Inotersen Sloan M COUGHLIN
P078	Prospective evaluation of an applied wt-ATTR-CM machine learning model to a United States (U.S.) health system electronic health record Ahmad FARAZ
P079	Prognostic Value of Cardiopulmonary Exercise Testing in Patients with Transthyretin Cardiac Amyloidosis. Francesco CAPPELLI
P080	Tafamidis 61 mg for treatment of ATTR cardiomyopathy in daily clinical practice: an observational study Fabian AUS DEM SIEPEN
P081	Transthyretin Cardiac Amyloidosis (ATTR-CA) and its rising awareness: Patient characteristics and survival in the Australian context Boyoun CHOI
P082	Significant survival benefits with Diflunisal in patients with Transthyretin (TTR) Amyloidosis Cardiomyopathy (ATTR-CM); A retrospective analysis Boyoun CHOI
P083	Effect of inotersen on global longitudinal strain in transthyretin cardiac amyloidosis Gerard T GIBLIN
P084	Inotersen treatment in transthyretin amyloid cardiomyopathy results in early and sustained serum transthyretin knockdown Gerard T GIBLIN
P085	Baseline ECG characteristics in ATTR-CM Dominic Richard PIMENTA

P086	Real-world experience with tafamis at ceparm University hospital. Federal University of Rio de Janeiro, Brazil.
P087	Márcia WADDINGTON-CRUZ Real-world experience with tafamis at ceparm University hospital. Federal University of Rio de Janeiro, Brazil. Márcia WADDINGTON-CRUZ
P088	Neurofilament Light Chain as a Biomarker in Hereditary Transthyretin-Mediated Amyloidosis: 36-Month Data from the Patisiran Global Open-Label Extension Emre ALDINC
P089	Characteristics of patients with ATTR amyloidosis and the Ile107Val mutation: insights from the Transthyretin Amyloidosis Outcomes Survey (THAOS) Márcia WADDINGTON-CRUZ
P090	Analysis of cardiac involvement in hereditary transthyretin amyloidosis after liver transplantation Yohei MISUMI
P091	4 Effect of Patisiran on Polyneuropathy and Cardiomyopathy in Patients with hATTR Amyloidosis with V122I/T60A Variants: A Phase 4 Observational Study Francy SHU
P092	Transthyretin amyloid polyneuropathy in mainland China: a unicentric study Lingchao MENG
P093	Phenotype and Clinical Outcomes of Hereditary Transthyretin Amyloidosis caused by p.Glu109Lys TTR Variant. A new endemic variant in Spain Fernando DE FRUTOS
P094	Comparison of disability: intradermal vasomotor nerves, intradermal sudomotor nerves, and intraepidermal sensory nerves in hereditary transthyretin amyloidosis Konen OBAYASHI
P095	A descriptive analysis of patients with ATTR amyloidosis and a mixed phenotype from the Transthyretin Amyloidosis Outcomes Survey (THAOS) Juan GONZÁLEZ-MORENO
P096	Hereditary transthyretin amyloidosis in middle-aged and elderly patients with idiopathic polyneuropathy: a nationwide prospective study. Guillaume FARGEOT
P097	Characterization of patients with hereditary Transthyretin Amyloidosis in a Register Study in Germany Helena Franziska PERNICE
P098	Hereditary Transthyretin Amyloidosis (ATTRv) in the Middle East: a short report of two confirmed cases Ramzi Ali TABBALAT

P099	How occupational needs can be obtained? A Semi-Structured Interview with Hereditary Transthyretin Amyloidosis patients. Aina Isabel GAYÁ BARROSO
P100	A rare case of late-onset hereditary ATTR Amyloidosis
P100	with cardiac and neurologic manifestation
	Vladimir CEJKA
P101	5 Characteristics of patients with ATTR amyloidosis and the Ser77Tyr mutation: insights from the Transthyretin Amyloidosis Outcomes Survey (THAOS) Violaine PLANTE-BORDENEUVE
P102	Effectiveness of patisiran following switch from tafamidis for the treatment of hereditary transthyretin-mediated (hattr) amyloidosis with polyneuropathy Céline LABEYRIE
P103	Characteristics of patients with ATTR amyloidosis and the Ile68Leu mutation: insights from the Transthyretin Amyloidosis Outcomes Survey (THAOS) Igor DIEMBERGER
P104	8 Characteristics of Patients With ATTR Amyloidosis and the Phe64Leu Mutation: Insights From the Transthyretin Amyloidosis Outcomes Survey (THAOS) Marco LUIGETTI
P105	Effect of tafamidis on disease progression in patients with non- Val30Met transthyretin amyloid polyneuropathy: a sub-study of the Transthyretin Amyloidosis Outcomes Survey (THAOS) Alejandra GONZÁLEZ-DUARTE
P106	Cardiological evolution of hereditary transthyretin amyloidosis (AhTTR) in patients with liver transplant Alvaro GRAGERA MARTINEZ
P107	Hereditary Transthyretin Amyloidosis caused by p.Ser43Asn TTR Variant. Insights and possible founder effect in Ecuador. Fernando DE FRUTOS
P108	A-V block as presentation of cardiac amyloid caused by conduction tissue infiltration Cristina CHIMENTI
P109	Comparing patient and clinician perspectives of ATTR amyloidosis: Insights from the development of the Transthyretin Amyloidosis Quality of Life (ATTR-QOL) Questionnaire Meaghan O'CONNOR
P110	Peritoneal dialysis is a valid treatment option in hereditary transthyretin amyloidosis João FERNANDES

P111	Transthyretin Familial Amyloid Polyneuropathy (TTR-FAP):
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P112	Case series: p.Leu131Met transthyretin amyloidosis in
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P114	Cardiac Screening of Amyloid TTR Pathogenic Variant
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P115	Comparison of Amyloid Detection in the Skin and
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P116	A rare TTR mutation determining severe cardiac and neurological amyloidosis
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P117	Lung volume restriction and abnormal cardiopulmonary response to
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P118	Evaluation and follow-up of the sudomotor function in a
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P119	The Relation between African American Race, Genotype,
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P120	Pre-symptomatic genetic testing for hereditary transthyretin
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D121	Peripheral nerve and cardiac features in hATTR patients presenting
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P208	Prospective MALDI-TOF analysis of blood serum peptidome to predict
P200	the onset and progression of hereditary transthyretin amyloidosis.
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P123	An additive destabilizing effect of two substitutions, T60I/V122I, in heterozygous compound TTRv amyloidosis
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P124	Selective recognition of human small transthyretin
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P125	Domain-domain interactions and dimerization of the human λ -III
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P126	ALBase: an updated platform to study immunoglobulin light chain sequences Tatiana PROKAEVA
P127	Structure-based peptides as novel therapeutic and
1127	detection tools in cardiac amyloidosis
	Rose M. PEDRETTI
P128	Alternative pathogenic mechanisms and novel
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	Luisa DIOMEDE
P129	Coagulative and fibrinolytic blood proteases efficiently cleave human transthyretir (hTTR) oligomers in vitro to generate the amyloidogenic fragment hTTR(49-127) Barbara SPOLAORE
P130	Mechanism of Misfolding and Amyloid Aggregation of the λ6 Light Chains Luis DEL POZO-YAUNER
P131	Filling the gap for transthyretin amyloidosis: biochemical and structural
	studies of in vitro and in vivo assembled amyloid fibrils
	Valentina MONDANI
P132	Modulation of transthyretin aggregation: role of preformed fibrils and heparin Guglielmo VERONA
P133	Circulating forms of plasma transthyretin in patients with wild-
	type transthyretin amyloidosis and effects of tafamidis
	Chiara SANGUINETTI
P134	Early Events of Immunoglobulin Light Chain Aggregation:
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	Valeria PIFRI UISSI-RUI7

P135	Recommendations for Addressing the Translational Gap between Experimental and Clinical Research on Amyloid Diseases Miriam SOLOMON
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P137	Clusterin in Alzheimer's disease: friend or foe? Georgia NASI
P138	Study of berry distinct polymorphism in ATTR amyloidosis fibrils by cryo-EM Binh NGUYEN
P139	Cardiac proteotoxicity is resulting from a complex interplay of several molecular properties of amyloidogenic Light Chains Rosaria RUSSO
P140	Assessing Immunoglobulin Light Chain Protein Stability and Stabilization by Pharmacological Chaperones Using Differential Scanning Fluorimetry Jianying WANG
P141	Limited cardiomyocytes' growth response to stimulation with human plasma and phenylephrine predicts poor outcome in ATTR cardiomyopathy in an in-vitro model. Selina Julia HEIN
P142	When Amyloid Occupies the Bone Marrow: Are the Responses of CD138-depleted Cells from Marrows with Interstitial Amyloid Archetypal? Ping ZHOU
P143	Exploring the in vitro effects of light chain-induced proteotoxicity on primary human cardiovascular system cells Paola ROGNONI
P144	Exploring Alzheimer's disease (AD) related human brain proteome with MALDI Imaging Mass Spectrometry in combination with shotgun proteomics Yumiko TOYAMA
P145	Transthyretin (hTTR) Amyloid Fibrils Trigger Plasma Clotting by Activating the Intrinsic Pathway of Blood Coagulation: Implications in Cardiac Senile Systemic Amyloidosis Laura ACQUASALIENTE
P146	The abstract will not be published
P147	V30M TTR animal model displayed a downregulated expression of several chemokines in different immune cell populations João MOREIRA
P148	In vitro treatment of light-chain amyloidosis plasma cells to characterize response to venetoclax Niels VAN NIEUWENHUIJZEN

P149	Frequent Occurrence of Fibrinogen Amyloidosis in Japanese Squirrels (Sciurus lis) Susumu IWAIDE
P150	Exercise suppresses mouse systemic AApoAII amyloidosis through
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P151	Characterization of heterozygous ATTR Y114C amyloidosis-specific iPS cells
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P152	BiP deletion leads to decreased cell viability and antibody
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P153	Age-related amyloid deposition in C57BL/6 mice:
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P154	Hemostasis dysfunction induces senile apoa2 amyloidosis in a mouse model G. Roussine CODO
P155	Endocytic inhibitory drugs protect C. elegans from
	the toxicity of amyloidogenic light chains
	Margherita ROMEO
P156	Is C. elegans a good model of β2-microglobulin related amyloidosis? Giulia FARAVELLI
P157	Poster retracted
P158	Efficient transient expression of exogenous immunoglobulin light
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P159	Strategies to induce amyloid light chain deposition in a transgenic mouse model
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P160	Prognostic Implication of Longitudinal Changes of Left Ventricular Global
	Strain after Chemotherapy in Cardiac Light Chain Amyloidosis
	Minjung BAHK
P161	Amyloid Valvular Heart Disease: A Look Beyond the Ventricular Walls
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P162	Scintigraphy Scan with planar and SPECT imaging of the chest
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	Amyloidosis: Comparison with an ATTR cohort of patients
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P163	A kit method for direct radiolabeling the amyloid reactive
	peptide p5+14 with technetium-99m (99mTc) for the detection
	of cardiac amyloidosis by SPECT/CT imaging
	Stephen KENNEL

P164	Quantitative changes in organ-specific amyloid load in a patient with AL amyloidosis, measured by 124I-AT-01 PET/ CT imaging, correlate with serum biomarkers Alan STUCKEY
P165	The role of speckle tracking echocardiography in the diagnostic assessment of cardiac amyloidosis and Fabry disease Isabel MATTIG
P166	Assessment of Incidental Cardiac Uptake in Bone Scintigraphy across Spain. ECCINGO Study Patricia TARILONTE
P167	Valve disease in cardiac amyloidosis: an echocardiographic score Alberto AIMO
P168	Coronary flow reserve by PET 13N-Ammonia in patients with Hereditary Transthyretin Amyloidosis with and without cardiac involvement. Aristóteles Comte DE ALENCAR NETO
P169	Differentiation of ATTR amyloidosis based on abdominothoracic organ- specific uptake of 124I-AT-01 (124I-p5+14) assessed by PET/CT imaging R. Eric HEIDEL
P170	Detection of extracardiac amyloid in patients with ATTR amyloidosis by PET/CT imaging using the amyloidophilic radiotracer 124I-AT-01 (124I-p5+14) Jonathan WALL
P171	Myocardial stiffness evaluation using atrial kick in healthy controls and patients with cardiac amyloidosis: a pilot study Ali SADEGHI
P172	[99mTc]Tc-DPD Scintigraphy Associating Semi-Quantitative Methods for the Diagnosis of Cardiac Amyloidosis: Experience in an Endemic Area Núria ORTA
P173	Myocardial Contraction Fraction (MCF): A Simple Measure of Myocardial Shortening That is Associated with Longitudinal and Circumferential Strain in Transthyretin Cardiac Amyloidosis. Dia SMILEY
P174	Histological validation of cardiac 99mTc-DPD uptake in patients with cardiac transthyretin amyloidosis Maria UNGERICHT
P175	Is a change of the current echocardiographic red flag for left ventricular wall thickness useful in cardiac amyloidosis screening? Maria UNGERICHT

P176	Prognosis of light chain amyloidosis with biopsy-proven cardiac involvement Matthias AURICH
P177	Multi-imaging characterisation of cardiac phenotype in different types of amyloidosis Adam IOANNOU
P178	Prognostic Implications of Clinical Phenotype and Severity of Cardiac Involvement in Patients Presenting with Immunoglobulin Light Chain Amyloidosis Aldostefano PORCARI
P179	Diagnostic performance characteristics of quantitative and semi- quantitative parameters of Tc99m pyrophosphate imaging for diagnosis of transthyretin (ATTR) cardiac amyloidosis: The SCAN-MP Study Shivda PANDEY
P180	DPD scintigraphy – a biomarker of microcalcifications rather than amyloid Ulrika THELANDER
P181	Prognostic Role of Echocardiographic Right Ventricular Parameters in Patients with Wild-Type ATTR Cardiac Amyloidosis Shravya VINNAKOTA
P182	False positive bone-scintigraphy in elderly hypertrophic cardiomyopathy Cristina CHIMENTI
P183	Dual-echo turbo-spin-echo and 12-echo multi-spin-echo sequences are equivalent techniques for obtaining T2-Relaxometry data in hereditary transthyretin amyloidosis Anysia PONCELET
P184	Cardiac imaging for assessing involvement in al amyloidosis patients: Experience in a single tertiary hospital Ana LLAMAZARES DE LA MORAL
P185	Phenotyping of hypertrophic cardiomyopathies using echocardiography: amyloid, Anderson-Fabry and hypertensive heart disease Aaisha FERKH
P186	Multimodality cardiac imaging in differential diagnosis of infiltrative cardiomyopathy Per ELDHAGEN
P187	Quantification of left ventricular amyloid using 124I-p5+14 (AT-01) and 18F-florbetapir positron emission tomography in AL and ATTR amyloidosis Olivier F. CLERC
P188	Changes in Left Ventricular Myocardial Composition Following Targeted Plasma Cell Therapy in Light Chain Amyloidosis: A Cardiac Magnetic Resonance Study Olivier F. CLERC

P189 Evaluation of Echocardiographic Parameters for Prognostication

in Patients with Systemic Light Chain Amyloidosis

Ahmet Mursel ULUSAN

P190 Central Nervous System damage in hereditary Transthyretin

Amyloidosis: A multimodal MRI study Renan Flávio DE FRANÇA NUNES



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	Natasha GORRIE
PLB002	Burden of transthyretin amyloid cardiomyopathy in patients and caregivers: interim analysis of a large, ongoing, non-interventional study Lucia PONTI
PLB003	Epidemiology of cardiac amyloidosis in Germany: a retrospective analysis from 2009 to 2018 Roman PFISTER
PLB004	Ixazomib maintenance following initial therapy in patients with high-risk immunoglobulin light chain (AL) amyloidosis. Roman PFISTER
PLB005	Poster retracted
PLB006	Mid-term analysis of the Clinical Amyloidosis Registry in Germany Niklas FUHR
PLB007	On bead de-glycosylation coupled with MALDI-TOF mass spectrometry provides a simple method for confirming light chain glycosylation and provides a sensitive method for residual disease detection Hannah V. GILES
PLB008	Real World Patient, Advocate, and Caregiver Perspectives on Amyloidosis: Awareness, Knowledge Gaps, and Psychosocial Impact Mukund NORI
PLB009	Renal histopathological scoring of amyloid deposits is crucial to assess disease progression in light-chain (AL) amyloidosis: a multicentre retrospective study Marco ALIINOVI
PLB010	Response rates to second line treatment with Daratumumab Bortezomib (Velcade) Dexamethasone (DVD) in relapsed/refractory light chain (AL) amyloidosis after initial bortezomib based regime Joshua BOMSZTYK
PLB011	Retrospective Cohort Study of treatment with BCL-2 inhibitor Venetoclax in relapsed or refractory AL amyloidosis Kaya VEELKEN
PLB012	Tafamidis medication adherence in patients with transthyretin cardiac amyloidosis (ATTR-CM) in a Japanese medical claims database

PLB013	The impact of renal histopathology on the renal outcome
	for newly diagnosed patients with AL amyloidosis
	Tarek ASHOUR
PLB014	The patient voice: development and results of a pilot
	patient experience data (PED) survey
	Mukund NORI
PLB015	The role of local complement expression in renal
	amyloidogenic light chain amyloidosis
	Nathalie KRIEGLSTEIN
PLB016	Transthyretin tetramer destabilization, marked by lower plasma
	transthyretin, is causally associated with increased risk of all-
	cause and cardiovascular mortality in the general population
	Mette CHRISTOFFERSON
PLB017	Two-year follow-up of the first case of systemic light chain amyloidosis
	treated with anti-B cell maturation antigen -CAR T cells
	Aina OLIVER-CALDÉS

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P192	The Journey to Diagnosis of ATTR Amyloidosis: Burden of Early Disease Chafic KARAM
P193	Manifestations of chronic heart failure in patients with AL-amyloidosis and ATTR-amyloidosis. Prospective observation data. Maria KUDRYAVTSEVA
P194	Artificial Intelligence Enhanced Electrocardiogram: A Simple Tool to Monitor for Clinical Improvement in Cardiac Amyloidosis? Awais MALIK
P195	Diagnostic and therapeutic center for amyloidosis at Kumamoto University Mitsuharu UEDA
P196	Experiences and decision-making in confirmed and potential carriers of attr- related genetic variants Jocelyn ASHFORD
P197	Potential sources of error in the identification and referral of amyloidosis to a tertiary center Lisa MENDELSON
P198	Higher Length of Stay and Readmission Burden in Heart Failure Patients with Cardiac Amyloidosis Than Those without Abdallah MASRI
P199	Management and patient estimation of amyloid light-chain (al) amyloidosis in Portugal: Results from a physicians' survey Susana SANTOS
P200	Prevalence of pauci-symptomatic amyloid transthyretin cardiac amyloidosis in the general population Alberto AIMO
P201	Redefining the epidemiology of cardiac amyloidosis. A systematic review and meta-analysis of screening studies Alberto AIMO
P202	Finding a balance between specialist and local care: amyloidosis patient perspectives on a single-centre approach Faye Amelia SHARPLEY

P203	Multidisciplinary approach for the early detection of amyloid in patients who undergo carpal tunnel syndrome or lumbar stenosis surgery. Preliminary results of an ongoing study. Núria ORTA
P204	Artificial Intelligence-Enhanced Models To Predict Light Chain Amyloidosis From Patients With Monoclonal Gammopathy Of Undetermined Significance And Smoldering Multiple Myeloma Eli MUCHTAR
P205	Lumbar spinal stenosis syndrome as surrogate for transthyretin cardiac amyloidosis. Laura DE MICHIELI
P206	Expert recommendations for improving the implementation of nuclear scintigraphy to support accurate diagnosis of cardiac amyloidosis in a non-specialist setting Julian GILLMORE
P207	Amyloidosis and its multifaces: A case report Diane Xavier ÁVILA
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P209	Real-life evaluation of an algorithm for the diagnosis of cardiac amyloidosis Mélanie BÉZARD
P210	Optimal Patient Selection for Referral to Tc-99m-PYP Scanning for Transthyretin Cardiac Amyloidosis Sarah CUDDY
P211	Cardiac amyloidosis screening in a cohort of patients with spinal stenosis: a case series. Pierpaolo Mattia MENNILLI
P212	Changes in the Journey to Diagnosis of Cardiac Amyloidosis in the Past 10 Years: Results from the Amyloidosis Research Consortium's 2017 Cardiac Survey and 2022 Amyloidosis Community Survey Sabrina REBELLO
P213	Prognostic Value of an Artificial Intelligence Enhanced ECG Model in Cardiac Amyloidosis Surendra DASARI
P214	Retrospective analysis of a patient cohort with suspicion of systemic amyloidosis, finally not confirmed Philine RITTER
P215	Validation of Amylo-AFFECT, a self-reported questionnaire to assess health-related quality of life and to determine the prognosis in cardiac amyloidosis

P216	The landscape of amyloidosis in Switzerland: Report of the Amyloidosis Registric Sofie BROUWERS
P217	Low QRS voltages in cardiac amyloidosis: echocardiographic correlates and prognostic value Alberto CIPRIANI
P218	Clinical impact of musculoskeletal pathology in patients with transthyretin-associated amyloidosis (ATTR): retrospective analysis of the case series from our center Andreu TRIGUERO
P219	Early detection of hereditary amyloidosis Ligia Rocha ANDRADE
P220	Role of combining ai-ecg to clinical risk scores for the prediction of transthyreti amyloid cardiomyopathy in heart failure with preserved ejection fraction Daniel DAVIES
P221	Prevalence and Implications of Classic ECG Findings in a Contemporary ATTR Cohort Genise GREEN
P222	Prevalence of Daytime and Nighttime Central Apneas in Patients with Cardiac Amyloidosis Francesco GENTILE
P223	The distribution of amyloidosis diseases in Germany: National Clinical Amyloidosis Registry Ute HEGENBART
P224	Amyloidosis diagnoses and shifting distribution of ATTR and AL from 2019 to 2021: a German single center experience. Timon HANSEN
P225	Initial Experience of a Private Amyloidosis Center Diane Xavier ÁVILA
P226	From symptoms and signs to diagnosis – Development of a simple screening tool for hereditary transthyretin amyloidosis (AmyloScan©) Juliane SACHAU
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P228	Cardiac amyloidosis in Latin America: Opportunities to increase disease awareness among clinicians. Findings from the AMILO-LATAM research group Estela Isabel CARVAJAL-JUAREZ

P229	Quality assessment of teaching in transthyretin amyloidosis Bruno Vaz Kerges BUENO
P230	Glomerular and tubular renal function in patients with Hereditary Transthyretin Amyloidosis (ATTR) Moisés Dias SILVA
P231	Targeted sequencing of functionally selected genes in patients with wild-type transthyretin amyloidosis Inmaculada MORENO GÁZQUEZ
P232	Detection of amyloidogenic FLC in serum as a non-invasive tool to facilitate the diagnosis of AL amyloidosis Rivka GOLDIS
P233	Immunoglobulin high-throughput sequencing in Monoclonal Gammopathy of Clinical Significance (MGCS): experience of the French Amyloidosis center Sébastien BENDER
P234	Neurofilament light chain, an early biomarker for polyneuropathy in hereditary transthyretin-related (ATTRv) amyloidosis Milou BERENDS
P235	The Clusterin/von Willebrand Factor Ratio Is Significantly Lower in Marrow Plasma from AL λ -type Than from λ -isotype Monoclonal Gammopathy Patients Stephanie SCALIA
P236	A spectrum of clinical phenotypes associated with p.(Arg54Gly) TTR variant. Dorota ROWCZENIO
P237	Neurofilament light chain as early biomarker for hereditary transthyretin amyloidosis – the Swedish experience Malin OLSSON
P238	Neurofilament light chain measurement in hereditary transthyretin- related amyloidosis patients with myocardial sympathetic neuronal damage: substitution for 123I-meta-iodobenzylguanidine scintigraphy? Milou BERENDS
P239	Prevalence of variant genotype in patients with suspected cardiac ATTR amyloidosis Roberta MUSSINELLI
P240	Engraftment Syndrome After ASCT is Associated elevated IL-10 and IP-10 (CXCL10) levels Amer ASSAL
P241	Genetic modifiers in hereditary and acquired TTR amyloidosis: a genome-wide association study

P242	Skin Biopsy Has A High Diagnostic Yield In Patients With Systemic Amyloidosis And Neuropathy.
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P243	Assessing amyloid prevalence, type, and extent of burden in the ligamentum flavum of patients with spinal stenosis undergoing routine laminectomy Emily MARTIN
P244	Prevalence of Localized Amyloid in Ligamentum Flavum of Patients with Lumbar Spinal Stenosis Francesco MARCHI
P245	Definition of Bone marrow biopsy involvement in amyloidosis – proposal for reporting Maria M. PICKEN
P246	Detection of amyloid deposits in skin biopsies of patients with clinically suspected variant transthyretin amyloidosis Rivka GOLDIS
P247	Amyloid deposits are common in different spinal structures among patients with spinal stenosis referred for decompression surgery Navya KOTTURU
P248	Diagnosis and typing of pulmonary amyloidosis: A series of cases Paola CASTILLO
P249	Concurrent light chain amyloidosis and proximal tubulopathy: insights into different aggregation behavior. Simone FEURSTEIN
P250	Clinicopathological analyses of an autopsied case with hereditary ATTR amyloidosis 22 years after liver transplantation Shiori YAMAKAWA
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P254	Glomerular proteomics – unmasking sub-clinical amyloid or false positive? Janet GILBERTSON
P255	Complement in amyloidosis – immunohistochemical verification of activation of the complement system in renal light-chain amyloidosis Helene MÜHLE

P256	the Norwegian Kidney Biopsy Registry 1988 – 2017 Hilde Jæger VASSTRAND
P257	Clinical and Morphological Phenotypes of AL and ATTR Amyloidosis Detected via Gastrointestinal Biopsies Catherine E. HAGEN
P258	Amyloid deposition in granuloma of tuberculosis patients: A single-center pilot study Shreya GHOSH
P259	The role of minor salivary gland biopsy in the diagnosis of systemic amyloidosis: results of a prospective study in 332 patients Martina NANCI
P260	Managing patients with carpal tunnel syndrome and positive congo red stain: a work in progress Ankita TANDON
P261	RNA-sequencing reveals similarity of AL amyloidosis and MGUS aberrant cells along with several potential target genes Zuzana CHYRA
P262	Presentation and outcome of patients with coexisting cardiac AL and ATTR amyloidosis Marco BASSET
P263	Validation of a mouse anti-human Serum Amyloid A antibody immunohistological test for diagnostic clinical practice. Nicola Amy BOTCHER
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P265	The use of abdominal fat aspiration biopsy to quantify the amyloid burden during long-term follow-up of patients with AA amyloidosis Norbert BLANK
P266	Renal amyloidosis in people who inject drugs in Oslo, Norway. Helga GUDMUNDSDOTTIR
P267	Properties of generic monoclonal AA antibodies for classifying most vertebrate AA amyloidoses, and as probes for innate functions of SAA Reinhold Paul LINKE
P268	SARS-Cov2 infection in AA amyloidosis: a high rate of mortality Rim BOURGUIBA
P269	Clinicopathological characteristics of a Japanese case with fibrinogen $A\alpha$ -chain amyloidosis Toshiya NOMURA

P270	Hereditary Gelsolin amyloidosis: Clinical features from a large cohort in the US Lisa MENDELSON
P271	Siblings with a novel type of amyloid neuropathy, hereditary Aβ2M amyloid neuropathy: Report of the second family in the world Nagaaki KATOH
P272	Apolipoprotein A-IV amyloidosis in a Cotton-top tamarin (Saguinus oedipus) Tomoaki MURAKAMI
P273	Histological and proteomic analyses of semenogelin 1 amyloid deposits Masayoshi TASAKI
P274	Mass spectrometry-based proteomic analysis of adsorbed molecules related with dialysis-related amyloidosis in hexadecyl-immobilized cellulose beads. Suguru YAMAMOTO
P275	Echocardiographic findings in subjects with an amyloidogenic apolipoprotein A1 mutation Daniela TOMASONI
P276	latrogenic cerebral amyloid angiopathy Jan Christoph PURRUCKER
P277	Fibrinogen A alpha-chain amyloidosis journey: dialysis and kidney transplantation interface José Pedro ESCALEIRA
P278	NNC6019-0001, a humanized monoclonal antibody, in patients with transthyretin amyloid cardiomyopathy (ATTR-CM): rationale and study design of a phase 2, randomized, placebo-controlled trial Matthew MAURER
P279	Efficacy and safety of belantamab mafodotin monotherapy in patients with relapsed or refractory light chain amyloidosis: a phase 2 study by the European Myeloma Network Efstathios KASTRITIS
P280	Glavonoid, a possible prophylactic supplement for ATTR amyloidosis Hiroaki MATSUSHITA
P281	Safety and tolerability of CAEL-101, an anti-amyloid monoclonal antibody, combined with anti-plasma cell dyscrasia therapy in patients with light-chain (AL) amyloidosis: results from a phase 2 study Cristina QUARTA
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BindingSite	15 000 €	Support congress, exhibition, logo		
BMS	15 000 €	Support congress, exhibition, logo		
GSK	15 000 €	Support congress, exhibition, logo		
Intellia	15 000 €	Support congress, logo		
Protego	15 000 €	Support congress, exhibition, logo		
Sanofi	15 000 €	Support congress, exhibition, logo		
Zentalis	15 000 €	Support congress, exhibition, logo		
Biotec-Level Signature 1				
Neurimmune	5 000 €	Support congress, exhibition, logo		
Life Molecular Imaging	5 000 €	Support congress, exhibition, logo		

SAVE THE DATE

The 2024 International Symposium on Amyloidosis will be held at

Mayo Clinic in Rochester Minnesota, USA on June 9 – 13, 2024