

Focus on the renin-angiotensin, complement, coagulation and kallikrein-kinin system in inflammation: from rare diseases to specific proteins as therapeutic targets.

Rationale

The renin-angiotensin with its pro and anti-inflammatory effects, the complement system with C1 inhibitor that inhibits components of the kallikreinkinin system (KKS), such as activated FXII, FXI and plasma kallikrein, as well plasmin affecting the fibrinolytic system; the coagulation and the KKS, tightly interlinked with the contact system, which is integral to the intrinsic pathway of coagulation (FXI) and thrombus formation; each of them consist of a large number of distinct plasma and membrane bound proteins and receptors that can be activated through proteolytic cascades. Notably, all systems are excessively activated during inflammation, they have potent pro-inflammatory and prothrombotic effects and they increase vascular permeability, leading to edema.

Although the dysregulation of each protein system is involved in the pathogenesis of common disorders, only a few rare diseases such as hereditary angioedema, paroxysmal nocturnal hemoglobinuria, atypical-uremic hemolytic syndrome, C3 glomerulopathies, coagulopathies, are approved for clinical treatment targeting renin-angiotensin, complement, coagulation and KKS. The diagnosis and the treatment of these rare diseases are a challenge for health system because it is necessary to improve basic and clinical knowledge and management skills in order to offer the best response to patient's needs. In our workshop we would love to discuss the role of these complex plasma protein systems in disease pathogenesis and then share future pharmacological strategies to treat rare and common diseases with plasma contact factors and complement targeted therapies.

In order to achieve the best results within our workshop, it is of fundamental importance that everyone is an active and responsible part in our job project and in the sharing of final common choices.

ITACA BOARD

Mauro Cancian, Padova
President

Andrea Zanichelli, Milano Vice-President. Treasurer

Francesco Arcoleo, Palermo Secretary

Vincenzo Montinaro, Acquaviva delle Fonti (BA) Massimo Triggiani, Salerno Counsellors

Scientific Coordinators

Francesco Arcoleo

Director of the Unit of Clinical Pathology and Immunology Ospedale Villa Sofia - Cervello, Palermo

Pietro Accardo

Regional Reference Center Rare Diseases of the Immune System and Angioedema Haemostasis and Thrombosis Center - Ospedale Villa Sofia - Cerv<mark>ello, Pal</mark>ermo

Faculty

Pietro Accardo, Palermo
Francesco Arcoleo, Palermo
Gianluigi Ardissino, Milano
Roberta Bulla, Trieste
Mauro Cancian, Padova
Pierpaolo Coni, Cagliari
Davide Firinu, Cagliari
Claire Harris, Newcastle upon Tyne (UK)
Maurizio Margaglione, Foggia
Vincenzo Montinaro, Acquaviva delle Fonti (BA)
Paul Morgan, Cardiff (UK)
Rosario Notaro, Firenze
Santiago Rodriguez de Cordoba, Madrid (ES)
Riccardo Senter, Padova

Massimo Triggiani, Salerno

	Frida	ay, October 13 th	
	12.30	Registration and Light lunch	
	13.20	Introduction Francesco Arcoleo	
		1° WORKSHOP Chairman: Francesco Arcoleo	
	13.30	The complement system: the overture Pietro Accardo	
	14.00	Q&A	
		2° WORKSHOP Chairman: Maurizio Margaglione	e ex
	14.15	Complement genetics Santiago Rodriguez de Cordoba	To The State of th
100	14.45	The kallicrein-kinin system: genetics and biology Davide Firinu	1
	15.15	Q&A	N
4	15.30	Coffee break	
		4	



Saturday, October 14 th			
	4° WORKSHOP Chairman: <i>Santiago Rodriguez de Cordoba</i>		
09.00	Hereditary angioedema Riccardo Senter		
09.30	ITACA: a network of Italian centers involved in diagnosis and treatment of rare diseases of kallicrein-kinin system <i>Mauro Cancian</i>		
10.00	Atypical uremic syndrome Gianluigi Ardissino		
10.30	Parossismal nocturnal hemoglobinuria Rosario Notaro		
11.00	C3 glomerulopathy Vincenzo Montinaro		
11.30	Q&A		
11.45	Coffee break		
	5° WORKSHOP Chairman: Paul Morgan		
12.00	Unmet needs of rare diseases: from patient view to basic and clinical research network Massimo Triggiani		
12.30	Conformational diseases: deficit AAT model Pierpaolo Coni		
13.00	Images of rare diseases: from hereditary angioedema to acquired haemophilia Pietro Accardo		
13.30	Q&A Results of all lectures and workshops		
13.45	Round table Pietro Accardo, Francesco Arcoleo, Gianluigi Ardissino, Roberta Bulla, Mauro Cancian, Pierpaolo Coni, Davide Firinu, Claire Harris, Maurizio Margaglione, Vincenzo Montinaro, Paul Morgan, Santiago Rodriguez de Cordoba, Riccardo Senter, Massimo Triggiani		

14.00 Light lunch



General information

Official language • English

Venue • NH Hotel Palermo - Foro Italico, 22/B - 90133 Palermo

Registration • Registration is free and includes participation to the workshop, work lunches, coffee breaks, certificate of participation and the CME certificate (to those entitled to). Whomever interested must register on the website https://www.centercongressi.com/eventi/complementsystem

CME • The Workshop will be included in the Italian National program of Continuing Medical Education (CME), for the profession of **Psychologist** and **Medical Doctor** with the following specialties: Nephrologists, Molecular Biology, Genetics, Allergology, Internal Medicine, Haematology, Pathology.

To obtain the CME certificate participants must attend 90% of the workshop and correctly answer to 75% of the questions.



e congressi

info@centercongressi.com • www.centercongressi.com