



**CONGRESO  
SAIB 2022**

Resúmenes de  
Comunicaciones Orales  
y Posters

**Abstracts of  
Oral Communications  
& Posters**



**SAIB**

Sociedad Argentina de  
Investigaciones en Bioquímica  
y Biología Molecular



LVIII Annual Meeting of the  
Argentine Society for Biochemistry  
and Molecular Biology Research



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# **DELEGATES OF SCIENTIFIC SECTIONS**

## **Cell Biology**

Pablo Aguilar

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## **Lipids**

Martín Oresti

CONICET – Universidad del Sur

## **Microbiology**

Hebe Dionisi

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## **Plants**

Elina Welchen

CONICET – Universidad Nacional del Litoral

## **Signal Transduction**

Graciela Boccaccio

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# PROGRAM AT A GLANCE

	Tuesday 8 <sup>th</sup>	Wednesday 9 <sup>th</sup>	Thursday 10 <sup>th</sup>	Friday 11 <sup>th</sup>
		<b>Oral Communications</b>	<b>Oral Communications</b>	<b>Oral Communications</b>
8:30		"Sala Magna" Microbiology	"Sala Magna" Plants	"Sala Magna" Cell Biology
10:30		"Sala Plumerillo" Plants "Sala Horcones" Lipids - Neurosciences	"Sala Plumerillo" Microbiology "Sala Horcones" Cell Biology	"Sala Plumerillo" Plants - Biotechnology "Sala Horcones" Signal Transduction – Enzymes
10:30		Coffee break	Coffee break	Coffee break
11:00		<i>"Sala Magna"</i> Plenary Lecture	<i>"Sala Magna"</i> Plenary Lecture	<i>"Sala Magna"</i> Plenary Lecture "Cono Sur"
12:30		<b>Dr. Ernesto Podesta</b>	<b>Dr. Mario Feldman</b>	<b>Dr. Rodrigo Gutierrez</b>
12:30		<i>"Sala Magna"</i> Round Table Surf your career	Free time for lunch	Free time for lunch
14:30	Registration	Free time for lunch		
		<b>Symposium</b>	<b>Symposium</b>	<b>Symposium</b>
14:30		"Sala Magna" Signal Transduction	"Sala Magna" Lipids "Sala Plumerillo" Microbiology	"Sala Magna" Cell Biology "Sala Plumerillo" Plants "Sala Horcones" Short Plenary Lectures
16:30		"Sala Plumerillo" Plants "Sala Horcones" Young Investigators	"Sala Horcones" Young Investigators	
16:30		Coffee break	Coffee break	Coffee break
17:00				
	<i>"Sala Magna"</i> <i>Opening Ceremony</i>			
17:00		POSTERS (Central Hall)	POSTERS (Central Hall)	POSTERS (Central Hall)
19:00	<i>Plenary Lecture Alberto Sols. <b>Dra. Isabel Varela Nieto</b></i>			
19:00	<i>"Sala Magna"</i> Plenary Lecture	<i>"Sala Magna"</i> Plenary Lecture Hector Torres	<i>"Sala Magna"</i> Plenary Lecture Ranwel Caputto	<i>"Sala Magna"</i> Plenary Lecture
20:30	<b>Dr. Craig Roy</b>	<b>Dra. Ana Belén Elgoyhen</b>	<b>Dra. Alejandra del Carmen Alonso</b>	<b>Dr Maximiliano Gutierrez</b>
20:30	Welcome Cocktail		SAIB Society Annual Meeting	Awards presentation and Closing Ceremony
22:00	<i>Central Hall</i>			

## LI-03

### **Fusion of micron-size vesicles: interplay between the mitochondrial Mfn2 protein and lipids**

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Membrane fusion is crucial for the coordination of mitochondrial dynamics. An imbalanced mitochondrial dynamic leads to the formation of fragmented mitochondria and a decrease in intracellular ATP levels, contributing to the development of important diseases, including neurodegenerative, cardiac or cancer conditions. The fusion process is energetically unfavorable, thereby requiring specialized proteins. In mammals, Mitofusins (Mfn) 1 and 2 are responsible for mitochondrial outer-membrane (OMM) fusion. They belong to the dynamin superfamily of multi-domain GTPases. Recent structural studies suggest that, upon GTP hydrolysis, Mfns dimerize to promote the approaching and fusion of OMM. However, the OMM fusion seems to require multiple regulatory factors that control the dynamics and kinetics of mitochondrial fusion through the formation of heterotypic Mfn1-Mfn2 dimers. In this study, we purified and functionally reconstituted the full-length mouse Mfn2 in large and giant unilamellar vesicles (LUVs and GUVs, respectively). Vesicles were prepared with POPC alone or with 30% of plasmalogen-PC or DOPE. Unlike GDP, after incubation with GTP, vesicles underwent fusion. Fast video microscopy imaged the Mfn2-dependent membrane fusion pathway which involves the formation and expansion of a membrane diaphragm and the opening of a fusion pore. The incorporation of DOPE (30% mol) in the lipid composition did not alter the fusion sequence but enhanced the fusion kinetics significantly, as revealed by a lipid-mixing assay. Our observations show that Mfn2 alone can promote the fusion of micron-sized vesicles, without the presence of other proteins in the membrane. In addition, the lipid environment is an important factor in the modulation of Mfn2-dependent membrane fusion, a process that seems to require topological lipid intermediates with negative curvature.