

SYMPOSIU[m]

de recherche du 175^e anniversaire

11 octobre 2018



Hôpital ophtalmique
Jules-Gonin

Service universitaire d'ophtalmologie
Fondation Asile des aveugles

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Edito

This symposium will present state-of-the-art of the translational research in ophthalmology. The first part will be dedicated to genetic of inherited retinal dystrophies and the importance to maintain a patient register. Speakers will next describe different cellular pathways involved in retinal degenerations such as AMD, dystrophies, and uveitis, as well as epigenetic alterations in retinal dystrophies and uveal melanoma. The development of new technologies to better access the retina and to visualize in vivo the retina with a cellular resolution will be presented. Finally, the program ends with a session dedicated to new therapies with an emphasis on drug, gene and cell delivery.

Comité organisateur

Prof. Yvan Arsenijevic
Dr Olivier Lorentz
Dre Irmela Mantel
Dr Alexandre Moulin
Prof. Reinier Schlingemann
Prof. Thomas J. Wolfensberger

Avec la participation de :



Programme du jeudi 11 octobre

08h15 - 08h45	<i>Welcome Coffee</i>
08h45 - 09h00	Introduction Yvan Arsenijevic, Reinier Schlingemann, Thomas J. Wolfensberger
09:00-10:05 How to deal with gene defect?	
	Moderator : Christian Grimm
09h00 - 09h25	(Patho-)Genesis of inherited cone dysfunctions Bernd Wissinger, University of Tübingen
09h25 - 09h45	Non-Mendelian inheritance in retinal degenerations Carlo Rivolta, University of Lausanne / University of Leicester
09h45 - 10h05	Molecular diagnostics in the era of clinical genomics : building national registries Pascal Escher, University of Bern
10h05 - 10h30	<i>Coffee Break</i>
10:30-12:05 Disease mechanisms I	
	Moderator : Reinier Schlingemann
10h30 - 10h40	General introduction Reinier Schlingemann, Fondation Asile des Aveugles / University of Lausanne / University of Amsterdam
10h40 - 11h05	The hypoxia challenge, a risk for AMD ? Christian Grimm, University of Zürich
11h05 - 11h20	Role of GSH in retinal diseases Raphaël Roduit, Fondation Asile des Aveugles / University of Lausanne
11h20 - 11h35	Novel target mechanisms in uveitis Yan Guex-Crosier, Fondation Asile des Aveugles / University of Lausanne

11h35 - 11h50	Other than VEGF in neovascular AMD Irmela Mantel, Fondation Asile des Aveugles / University of Lausanne
11h50 - 12h05	Biomarkers and new treatments for AMD Chiara Eandi, Fondation Asile des Aveugles / University of Lausanne
12h05 - 13h20	Lunch
13:20-14:25 Disease mechanisms II	
	Moderator : Pascal Escher
13h20 - 13h35	Epigenetic modifications during retinal dystrophies: narrowing the therapeutic targets Yvan Arsenijevic, Fondation Asile des Aveugles / University of Lausanne
13h35 - 14h00	BAP1 modulates the epigenetic landscape of uveal melanoma Nicolo Riggi, University of Lausanne et Alexandre Moulin, Fondation Asile des Aveugles / University of Lausanne
14h00 - 14h25	Use of phenotypic screening to identify candidate molecules for the treatment of glaucoma and the retinal degenerations Don Zack, Johns Hopkins University School of Medicine, Baltimore, USA
14:25-15h15 New technologies	
	Moderator : Irmela Mantel
14h25 - 14h45	Oblique illumination to visualize retinal cells Christophe Moser & Laura Kowalcuk, EPFL & Fondation Asile des aveugles / University of Lausanne
14h45 - 15h00	The artificial cornea Aleksandra Petrovic, Fondation Asile des Aveugles / University of Lausanne
15h00 - 15h15	Robotics and the eye Thomas J. Wolfensberger, Fondation Asile des Aveugles / University of Lausanne

15h15 - 15h45

Coffee Break

15:45-17:40 Drug, gene and cell therapies

	Moderator : Peter Humphries
15h45 - 16h10	On experimental molecular therapies targeting retinal degenerations and ocular hypertension Peter Humphries, University of Dublin
16h10 - 16h35	High sensitivity optogenetic to restore vision Sonja Kleinlogel, University of Bern
16h35 - 16h50	Retinal gene therapy, a long-term endeavor to success Corinne Kostic, Fondation Asile des Aveugles / University of Lausanne
16h50 - 17h15	Preclinical validation of a tissue engineered product consisting in RPE derived from human embryonic stem cells disposed on human amniotic membrane Christelle Monville, INSERM / University of Paris
17h15 - 17h40	Emerging therapies for inherited retinal degeneration Hendrik Scholl, University of Basel
17h40 - 17h45	Conclusion
18h00	<i>Aperitif</i>

Notes

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