

GENERAL PROGRAM

Day		Morning			Afternoon		Evening
18 th	Arri	Arriving to Havana, Transfer Havana Airport to "Havana Hotel, Cuba					Free
					ng to "Melia Peninsula Varadero Hotel"		
19 th		8:30 am- Transfer to			Registration	D	Free
	Workshop Venue				Opening Ceremony	Т	
					KNA 1		
20 th	KNA Symposium I 2-5 Basic Topics			Dia	Symposium 11 gnostic and Clinical aspects of meningitis diseases/ Poster and Rum testing	N	Free
21 th	KNA <u>Symposium III</u> 6 Meningococcal Vaccines U		U	KNA 7	Symposium IV Gonococcal vaccine	N	Free
22 th	2 th 25 years of the Proteoliposome as Human Cuban Vaccine			Stı	Symposium V reptococcus suis: disease and future challenges	E R	Free
23 th	KNA 8	Symposium VI Pneumococcal Meningitis	С Н	KNA 9	Symposium VII Correlates of Protection		20:30. GYO*
	and Vaccines				Closing Ceremony		Show
24^{th}	^h Bilateral Meetings			Che	eck out and Return to Havana		Free



Program

Day	Time	Activity
	08:30	Transfer to workshop Venue from Havana Hotels
	15:00	Arrive and Registration
19th	18:00	Opening Ceremony
	18:30	KNA 1: Carbohydrates and T cells: A sweet twosome. Kasper DL, USA
	19:30	Welcome Dinner

Time for Presentations				
Keynote address (KNA) = 35 min + 10 min for questions				
Oral Presentation = $25 \min + 5 \min$ for questions				
Short Oral Presentation (SOP) = 10 min + 5 min for				
questions				

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IV NeisseriaVaccines2013 I Workshop on Meningitis and Septicaemia

Neisseria Vaccines 2013

International Congress on Meningeril Disease Vaccines Varadero, Cuba, May 19-24, 2013

Day	Time	Activity					
		Symposium I: Basic Topics					
		Chairs: S Meri (Finland) and LM Wetzler (USA)					
	09:00-09:35	KNA 2: Could Adjuvants overcome the Thymus-independent of Polysaccharides? B Romeu, Cuba					
204h	09:45-10:20	KNA 3: Complement activating pattern recognition molecules (PRMs) of the innate immune defence and their possible role in Neisserial infections. J Chr Jensenius, Denmark					
_20th	10:30-11:05	KNA 4: Opportunities and risks of using complement regulator binding proteins as vaccines. S Meri, Finland					
	11:15-11:50	KNA 5: Mechanism of the Immune Stimulating Activity of the Neisserial Major Outer Membrane Protein PorB: Role of Antigen Presenting Cells and Systems Immunobiology Analysis. LM Wetzler , USA					
	12:00-12:25	The Meningitis Research Foundation Meningococcal Genome Library. J Lucidarme, UK					
	12	20 14:20 I unch					

12:30-14:30LunchSymposium II: Diagnostic and Clinical aspects for Meningitis DiseasesChairs: E Kaczmarski (UK) and J Diez-Dominguez (Spain)14:30-14:55Rapid diagnostics for meningitis Diseases. UK15:00-15:25Bacterial meningitis in Cuba. Clinical and epidemiological situation. J
Pérez, Cuba15:30-15:55Meningococcal disease. Beyond the figures. J Diez-Domingo, Spain16:00-16:25Lab safe working with meningococci. E Kaczmarski, UK

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	Symposium III: Meningococcal Vaccines						
	Chairs: J Holst (Norway) and G Enwere (France)						
	09:00-09:35	KNA 6: A long and winding road: towards a vaccine with broad strain- coverage against meningococcal serogroup B disease. J Holst, Norway					
	09:45-10:10	Carly development of MenBioVax vaccine with potential for universal rotection against <i>Neisseria meningitidis</i> . S Clarke, UK					
	10:15-10:40	new approach to prevention and treatment of meningococcal disease. Blackwell, Australia					
21th	10:45-11:10	ubcapsular meningococcal vaccine antigens in the common childhood ommensal, <i>Neisseria lactamica</i> . J Lucidarme, UK					
21111		OP 1: Conjugated vaccine candidate against <i>Neisseria meningitidis</i> . R Acevedo, Cuba					
		Progress and perspectives of MenAfriVac, a meningococcal A conjugate vaccine for the African meningitis belt. G Enwere, France					
	12:00-12:25	A trivalent outer membrane vesicle (OMV) vaccine against meningococcal disease for Africa. E Rosenqvist, Norway					
		SOP 2: An approach to tetravalent vaccine production from polysaccharide of <i>N. meningitidis</i> serogroups $ACYW_{135}$, to Muslim market. D González, Cuba					
	12:45-12:55	SOP 3: New vaccines strategies against <i>Neisseria meningitidis</i> serogroup X. C Zayas, Cuba					
	13	3:00-14:30 Lunch					

	Symposium IV: Gonococcal vaccine				
	Chairs: P. Rice (USA) and L. Velásquez (Chile)				
	14:30-15:15	4:30-15:15 KNA 7: Saccharide antibody protection against <i>Neisseria gonorrha</i> infections in the experimental mouse model can reversed by anti. I antibody. P Rice, USA			
21th	15:25-15:50	Role of glycocalyx and matrix extracellular proteins in the pathogenesis of <i>Neisseria gonorrhoeae</i> on fallopian tube cells. L Velásquez, Chile			
	15:55-16:20	Surfactant vesicles as a surrogate for whole cell vaccination allow for inclusion of lipooligosaccharide in vaccine preparations. D Stein, USA			
	16:25-16:50	RNA-seq Analysis of Vaginal Lavage Samples from Female Patients Identifies a Repertoire of Putative Gonococcal Vaccine Targets. C Genco, USA.			

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	Varadero,	Cuba,	May 19-24, 2013	
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		Round Table 25 years of the Proteoliposome as Human Cuban Vaccine					
	Chairs: D Cardoso (Cuba) and E Rosenqvist (Norway)						
	09:00-09:40	VA-MENGOC-BC [®] : Landmarks of a Vaccine. C Campa, G Sierra, Cuba					
	09:40-10:00	Controversies Regarding Clinical Use of VA-MENGOC-BC [®] . R Ochoa, Cuba					
22th	10:00-10:20	A-MENGOC-BC [®] : 25 years of security and efficacy preclinical ials. JF Infante, Cuba					
	10:20-10:40	Post-license surveillance of VA-MENGOC-BC [®] . I. Cuevas, Cuba					
	10:40-11:00	Technological platform for VA-MENGOC-BC [®] production. R Barberá, Cuba					
	11:00-11:20	Commercial experiences of VA-MENGOC-BC [®] . E Caro, Cuba					
	11:20-11:40	Proteoliposome as the core of VA-MENGOC-BC [®] and adjuvant platform. O Pérez, Cuba					
	11:40-	Opinions of other Finlay's Members and Colleagues					
		12:30-14:30 Lunch					

	Symposium V: Streptococcus suis: disease and future challenges				
	Chairs: V Verez (Cuba) and M Gottschalk (Canada)				
	14:30-14:55	Overview of <i>Streptococcus suis</i> epidemiology of the disease and vaccine update M Gottschalk, Canada			
	15:00-15:25	Molecular typing of <i>Streptococcus suis</i> from pigs in Cuba. I Espinosa, Cuba			
22th	15:30-15:55	The immune response against <i>Streptococcus suis</i> : sepsis and toxic shock. M Segura, Canada			
	16:00-16:25	The pathogenesis of the <i>Streptococcus suis</i> meningitis. M Gottschalk , Canada			
	16:30-16:55	New chemical methods for conjugate vaccines: is there an avenue for <i>Streptococcus suis</i> ? R Roy, Canada			
	17:00-17:25	Learning from <i>Streptococcus pneumoniae</i> conjugate vaccine for <i>S. suis</i> meningitis. Y Valdés, Cuba			



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	Symposium VI: Pneumococcal meningitis and Vaccines				
	Chairs: W Hausdorff (USA) and V Verez (Cuba)				
	09:00-09:35	KNA 8: Epidemiology of pneumococcal meningitis, serotypes responsible, and the need and prospect for protein vaccines. W Hausdorff, USA			
23th	09:45-10:10	Impact of pneumococcal conjugate vaccines on pneumococcal meningitis in Latin America. JL Di Fabio, PAHO			
	10:15-10:40	UK experience with pneumococcal meningitis and PCV. R Borrow, UK			
	10:45-11:10	Immunological considerations and lessons in comparisons of bacterial meningitis vaccines against Hib, meningococcus and pneumococcus. D Goldblatt, UK			
	11:15-11:40	<i>Streptococcus pneumoniae</i> conjugate vaccine development in Cuba. V Verez, Cuba			

12:30-14:30

Lunch

	Symposium VII: Correlates of protection for <i>Neisseria meningitidis</i> and Streptococcus pneumoniae							
		Chai	rs: R. Borrow (UK) and D. Medini (Italy)					
	14:30-15:		KNA 9: Issues with correlates of protection f meningococcal ACWY glycoconjugate vaccines. R Bo					
	15:15-15:		The preparation and use of IgG-depleted complement in the interview of the second seco					
23th	15:45-16:		Evaluating the efficacy of fHBP containing vaccines using hSBA, the surrogate of protection. L York, USA					
	16:15-16:	40 isolates conserva	Serum Bactericidal Antibody against capsular group B meningococcal solates responsible for invasive disease shows that MATS is a conservative predictor of strain coverage by the 4CMenB vaccine. D Medini, Italy.					
	16:45-17:	7:10 Carriage as an endpoint for licensing vaccines: lessons fro PneumoCarr. D Goldblatt, UK.						
	17:15-17:	30 for immu	Use of serum bactericidal antibody as an e unogenicity of meningococcal vaccines serog 5. L García, Cuba					
			Closing Ceremony					
		20:30	GYO* Show					

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24 th	09:00-11:35	Bilateral Meetings and Free interchange between Delegates
	12:00-13:00	Lunch
	14:00	Return to Havana

20th	16:25-18:25	Poster Session
20th	16:25-18:25	Poster Session

	Conferences in Havana for Pediatrician, Immunologists and Microbiologist "William Soler" Hospital		
24 th	11:00-	Pneumococcal conjugate vaccines: what have we learned and what do we have yet to learn? W Hausdorff, Belgium	
		Understanding Responses to Polysaccharide and Conjugate Vaccines in infants and adults. D Goldblatt, UK	

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Vaccines 2013

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No **Tittles y Authors** Four monoclonal antibodies against capsular polysaccharides of Neisseria 1 meningitidis serogroups A, C, Y and W_{135} : its application in identity tests. N Amín, Cuba 2 Development of sandwich ELISAs for detection and quantification of capsular polysaccharides of *Neisseria meningitidis* serogroups A, C, W₁₃₅ and Y. F Reyes, Cuba 3 Latex particles as potential reagents for diagnostic of meningococcal disease M Cuello, Cuba 4 Evaluation of a rapid test for the detection of Neisseria meningitidis serogroup B in human serum and whole blood using a novel affinity biosensor. **X Bai, UK** SDS-PAGE and densitometric analysis to determine the concentration of 5 Lipopolysaccharide from *Neisseria meningitidis* serogrupos A, W₁₃₅, Y and X. M Cuello, Cuba 6 Identification of antigenic composition of outer membrane vesicles from Neisseria meningitidis serogroup X. Y Rodríguez, Cuba 7 New methodology western blot technique in identity assay. Optimization and Validation. R Diéguez, Cuba 8 Characterization of dry polysaccharides from *Neisseria meningitidis* serogroups A, C, Y and W₁₃₅ to be used as internal controls. LD Díaz, Cuba 9 Comparative results of the individual polysaccharides A, C, Y, and W₁₃₅ and the tetravalent vaccine ACWY by using pyrogen test. JF Núñez, Cuba 10 Predicting serum bactericidal responses: sensitivity and specificity of a flowcytometric complement deposition assay. L Allen, UK Bridging of two serum bactericidal antibody assays using rabbit complement 11 performed at the Health Protection Agency and at GlaxoSmithKline Biologicals. X Bai, UK 12 A model system for assessment of opsonisation of meningococci and commensal species. CC Blackwell, Australia. 13 Establishment of Quality Assurance System for the production of polysaccharides from Neisseria meningitidis serogroups A, C, Y and W₁₃₅ under HALAL requirements. R Martínez, Cuba Development and Validation the analytical methods in Polysaccharide W_{135} of 14 Meningococcal Vaccines. I Delgado, Cuba Development and Validation of method for 15 quantification multivalent polysaccharides vaccines by Capillary Zone Electrophoresis. Y Merchán, Cuba Validation of a RI-HPLC ethanol determination method. Characterization of these 16 impurity in VA-MENGOC BC[®] vaccine. M Cuevas, Cuba



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17	Validation of the sterilization equipments for saturated steam (autoclaves) used in the meningococcal vaccines production at Finlay Institute. M Hernández, Cuba
18	Concurrent validation of the process for obtaining dry polysaccharide from <i>Neisseria meningitidis</i> serogroup C. JV Bayolo, Cuba
19	Stability studies of a <i>Neisseria Meningitidis</i> serogroups A and W_{135} vaccine based on outer membrane vesicle. A Mandiarote, Cuba
20	New process evaluation at large scale to obtain capsular polysaccharide purified from <i>Neisseria meningitidis</i> serogroup W ₁₃₅ . S Lozada, Cuba
21	Stability studies for Polysaccharides and final lots of a trivalent <i>Neisseria</i> meningitidis serogroups A, C, and W_{135} vaccine. E Pérez, Cuba
22	Clinical Trials Audit to the <i>Neisseria meningitidis</i> A, C, and W ₁₃₅ Polysaccharide vaccine from Finlay Institute. B Simón, Cuba
23	Purification of lipopolysaccharides from <i>Neisseria meningitidis</i> for use as reference material for its quantification in vaccines. RA Cabrera, Cuba
24	Production of free porcine components Neisseria meningitidis Reference Seed Lots. CA del Puerto, Cuba
25	Stability of working seed lot of <i>Neisseria meningitidis</i> serogroup X grown in culture media of non-animal origin and preserved by freezing. M Hernández, Cuba
26	<i>Neisseria meningitidis</i> serogroup X: analytical challenges and alternatives for evaluation polysaccharide content by quantitative nuclear magnetic resonance. R Garrido, Cuba
27	Conjugation of Capsular Polysaccharides from <i>Neisseria meningitidis</i> Serogroups X to Tetanus Toxoid. M González, Cuba
28	Vaccine potential of outer membrane vesicles from <i>Neisseria meningitidis</i> serogroup X. C Zayas, Cuba
29	Preparation and characterization of conjugates from <i>Streptococcus pneumoniae</i> serotypes 7F, 9V and 19A to different carrier proteins. D Mariño, Cuba
30	Adjuvation of conjugates from <i>Streptococcus pneumoniae</i> serotype 1 and 14 to tetanus toxoid in Aluminium. Y Serrano, Cuba
31	Group B meningococcal vaccine candidacy of haemoglobin receptors based on distribution and phase variable status. J Lucidarme, UK
32	Comparison of cytokine gene polymorphisms among Greek patients with invasive meningococcal disease or viral meningitis. CC Blackwell, Australia.
33	A novel factor H-Fc chimeric immunotherapeutic molecule against Neisseria gonorrhoeae. P Rice, USA
34	Antimicrobial resistance patterns in Cuban gonococcal strains. O Feliciano, Cuba
35	The utility of PCR in diagnostic of unspecified and treated bacterial meningitis in Cuba. O Feliciano, Cuba

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36	Epidemiological markers and antimicrobial susceptibility of invasive strains. Cuba, 2002-2011. E Suárez, Cuba		
37	Intrathecal synthesis of complement system proteins in patients with eosinofilic meningitis due to <i>Angiostrongylus cantonensis</i> . B Padilla, Cuba		
38	MASP2 Intrathecal synthesis in eosinophilic meningitis due to Angiostrongylus cantonensis. A Arias, Cuba		
39	MASP2: Dynamics and Intrathecal synthesis. AJ Dorta, Cuba		
40	Immunological response identified in cerebrospinal fluid in patients with neurodegenerative diseases. I Zerr, Germany		