Programme EPOS 2006 Villamoura

No.	Time Thursday	Author 05.10.2006	Title
	18.00 – 20.00 h		Registration
	19.00 – 22.00 h		Opening Reception
	Friday	06.10.2006	
	8.00	B. Lorenz	Presidential Welcome
	8.10	E. Silva	Welcome by the local host
			Keynote Lecture:
L1	8.20	Sieving, P.	Ciliary Neurotrophic Factor (CNTF): Background
			and Outcome of a Phase 1-2 Clinical Trial for
			Retinal Degeneration
	9.00 – 9.40 h	I. Scientific Session	Neuroophthalmology – Function and
	3.00 – 3.40 H	i. Ocientino Ocasion	Functional Testing
	Chairman		
T1	9.00	Berk, A.	Horizontal Gaze Palsy And Scoliosis
			PHACE syndrome presentating with cerebral
T2	9.10	Hildebrand, G.D.	vasculopathy, congenital oculomotor nerve
. –	0.1.0		paresis and segmental, plaque-like facial
			hemangiomata in an infant
Т3	9.20	Yaman, A.	Cranial nerve palsies in children
			Impaired horizontal saccades and jerky head
T4	9.30	Carrasquinho, S.	movements – a clinical case of Congenital
			Oculomotor Apraxia?
			Discussion
	9.40 – 10.00 h		Coffee Break
	10.00 – 11.30 h	II. Scientific Session	Neuroophthalmology – Causes and Development
	Chairman		
L2	10.00	Brodsky, M.	Neurodiagnostic Malformations of the Optic Disc
T5	10.40	Gupta, A.	Autosomal dominant microcephaly-

			lymphoedema-chorioretinal dysplasia syndrome
Т6	10.50	Fonseca, A.	Parry-Romberg syndrome: report of two cases
T7		Hodgkins, P.	Swollen optic discs with headaches in children: Is it papilloedema Doc?
Т8	11.10	Sandfeld Nielsen, L.	Risk factors of ocular disorders in children with developmental delay Discussion
	11.30 – 12.30 h		1. Poster Session
			Neuroophthalmology – Function and
			Functional Testing Poster
5.4			Visual function, brain MRI findings and motility
P1		Kozeis, N.	status in children with spastic quadriplegia
			The role of visual evoked potentials (VEPs) as a
P2		Kozeis, N.	prognostic factor in children with bilateral spastic
			cerebral palsy
			Light flash evoked VEPs in children with
P4		Sjöström, A.	Dystrophia Myotonica
P5		Tekavcic Pompe, M.	Diagnosing optic neuritis in children
P6		Wittebol-Post, D.	Plasticity of the visual system in cerebral palsy
P7		Wittebol-Post, D.	The importance of visual symptoms and
1 7		Wittebol-1 Ost, D.	ophthalmic signs in hydrocephalus
			Neuroophthalmology – Causes and
			Development Posters
P8		Hodgkins, P.	Cerebral sinus venous thrombosis in children
P9		Gore, D.	Presumed infective meningo-encephalitis
10		0010, D.	complicated by bilateral optic neuritis
P10		Bangsgaard, R.	Isolated bilateral optic neuritis in demyelinating
			diseases in children
P11		Hodgkins, P.	Idiopathic Intracranial Hypertension or Optic
		-	Neuritis: A diagnostic dilemma in children
P12		Knezy, K.	Visual loss associated with scaphocephaly- case
			report Pariyantricular loucomalacia: an important causa
P13		Maduro, V.	Periventricular leucomalacia: an important cause of visual dysfunction in preterm infants
			or visual dysturiction in preterm infants

P14		Yaman, A.	Optic neuropathy associated with voluntary globe luxation in a 14-year-old child
	12.30 – 13.30 h		Lunch Break
	13.30 – 14.30 h		2. Poster Session
			ROP Poster
			Retinopathy of Prematurity Screening – A 9 year
P15		Patrício, M	retrospective study in Hospital Fernando
			Fonseca, Portugal
			Tumor Posters
P16		Casaer, P.	Retinal hamartoma in a healthy 5-year-old boy
P17		Teixeira, S.	Combined chemotherapy and adjuvant
1 17		Teixella, G.	treatment for intraocular retinoblastoma
			Cataract Posters
			Regression of cataract in a four-year-old girl with
P18		Gade, E.	Alpha-mannosidosis after bone marrow
			transplantation
			Pediatric visual rehabilitation by scleral-fixated
P19		Ganesh, A.	posterior chamber intraocular lenses in a
			developing country
P21		Nair, R.	Long-term review of 125 Paediatric cataract
1 2 1		Naii, IX.	patients
P22		Nair, R.	Contact lens management of Paediatric
1 22		rvan, rv.	aphakia-different stages
			Glaucoma Posters
			Clinical evaluation of drainage implants in the
P23		Autrata, R.	treatment of refractory pediatric glaucoma: 15-
			year experiences
P24		Maka, E.	Results of Trabeculectomy for Congenital
1 44		mana, L.	Glaucoma
P25		Serra, A.	Congenital Glaucoma Associated With
. 20		Jona, 7	Congenital Ipsilateral Lid Neurofibroma In NF-1:

Case Report

			Amblyopia Posters
P27 P28		Aznauryan, I. Balasanyan, V.	Objective methods of refractive amblyopia meridional form diagnostics with children Transdermal stimulation of visual nerve in amblyopia treatment
P30		lqbal, Z.	Effect of occlusion therapy on visual functions in children with amblyopia.
	14.30 – 15.10 h Chairman:	III. Scientific Session	ROP
Т9	14.30	Pacheco, T.	Retinopathy Of Prematurity: The Experience Of The Last Decade
T10	14.40	Ciomartan, T.	The use of RetCam II as a diagnostic instrument in telemedicine – the first experience in Romania
T11	14.50	Hellgren, K.	VLBW is associated with optic nerve and visual field abnormalities
T12	15.00	Teixeira, S.	Retinopathy of Prematurity - Consequences and Controversies of ETROP
			Discussion
	15.20 – 15.50 h		Coffee Break
	15.50 - 17.00 h	IV. Scientific Session	Tumours
	Chairman:		
L3	15.50	Désjardins, L.	Review of ophthalmic tumors relevant in neuroophtalmology
T13	16.20	lqbal, Z.	The Management and Dilemma of Retinoblastoma in the Developing Countries
T14	16.30	Schalij-Delfos, N.	Diagnostic pitfalls in intracranial germinomas

Eye involvement in patients with neuroblastoma

Discussion

17.00 – 18.30 h V. Scientific Session Cataract

16.40 Wenniger-Prick L

T15

	Chairman:				
T16	17.00	Patrício, M.	Combined Aqualase Liquefaction and Acrysof ReSTOR in the management of cataract in young patients		
T17	17.10	Edelson, C.	Refractive outcomes in pediatric cataract surgery		
T18	17.20	Pinello, L.	Visual outcome in bilateral congenital cataract associated with ocular and systemic diseases		
T19	17.30	Ashworth, J.	Outcome of intraocular lens implantation in infants		
T20	17.40	Dureau, P.	Echographic examination of corneal opacities in children		
T21	17.50	Morales Ballús, M.	Refractive change in pediatric pseudophakia		
T22	18.00	Serra, A.	Primary Juvenile Glaucoma In Pediatric Patients Discussion		
	19.30 h		Shuttle Bus to Pine Cliffs Resort		
	20.00 – 22.30 h		Dinner Party at Pine Cliffs Resort		
	Saturday	07.10.06			
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	8.00 – 9.20 h	VI. Scientific Session	Genetics
	Chairman:		
T23	8.00	Casteels, I.	Ocular findings in children with a microdeletion
			in chromosome 22q11.2
T24	8.10	de Ravel, T.	The Nance-Horan Syndrome: Underdiagnosed?
			A unique BBS family allows the identification of
T25	8.20	Dollfus, H.	a novel major BBS gene encoding for vertebrate
			specific chaperonine-like protein
			Variation of phenotype in patients with
T26	8.30	Friedburg, C.	compound heterozygous mutations of RetGC1
			depending on the affected domains
			Pseudoxanthoma Elasticum: Variability in
T27	8.40	Plomp, A.	expression both in patients homozygous and in
121			those heterozygous for the ABCC6 3775delT
			mutation
T28	9.50	Preising, M.	Mutational Screening in Choroideremia –
120	8.50	i reising, ivi.	Identification of an Innovative Spectrum of

T29	9.00	Votruba, M.	A mouse model of opa1 optic atrophy
			Discussion
	9.20 – 9.50 h		Coffee Break
	9.50 – 11.00 h Chairman:	VII. Scientific Session	Neuroophthalmology – Metabolics
L4	9.50	Diogo	Mitochondrial respiratory chain diseases in children – a most challenging issue
T30	10.20	Monteiro Magalhães, A.	Ocular manifestations of rare metabolic diseases
T31	10.30	Benevides de Melo, A.	The ocular manifestations of the Mucopolysaccharidoses
T32	10.40	Monteiro Magalhães, A.	Maroteaux-Lamy Syndrome: clinical presentation and evolution in 11 patients. Discussion
			Discussion
	11.00 – 12.00 h		EPOS General Assembly
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	12.00 – 13.00 h		Lunch Break
	12.00 – 13.00 h 13.30 – 14.30 h		Lunch Break 3. Poster Session Genetics Poster
P32		Andersson Grönlund, M.	3. Poster Session Genetics Poster Mitochondrial myopathy with retinal dystrophy and exercise intolerance in a sporadic patient with a G583A mutation in the mt tRNAphe gene
P32		•	3. Poster Session Genetics Poster Mitochondrial myopathy with retinal dystrophy and exercise intolerance in a sporadic patient
		M.	3. Poster Session Genetics Poster Mitochondrial myopathy with retinal dystrophy and exercise intolerance in a sporadic patient with a G583A mutation in the mt tRNAphe gene Familial exudative vitreoretinopathy (FEVR) in a child with pyruvate dehydrogenase (PDH)
P33		M. Bangsgaard, R.	3. Poster Session Genetics Poster Mitochondrial myopathy with retinal dystrophy and exercise intolerance in a sporadic patient with a G583A mutation in the mt tRNAphe gene Familial exudative vitreoretinopathy (FEVR) in a child with pyruvate dehydrogenase (PDH) deficiency Autozygosity mapping of Bardet-Biedl syndrome to 12q21.2 and confirmation of FLJ23560 as

Truncating Mutations

		Neuroophthalmology – Metabolics
		Optic neuritis in pediatric patients- clinical
P37	Autrata, R.	manifestations, etiology, therapy and visual
		outcomes in the long-term follow-up.
		Asymptomatic shunt malfunction detected
P38	Urban, B.	fortuitously by observation of papilloedema in 9-
		year-old boy
P39	Dutton, G.	Identifying Cerebral Visual Impairment
F 39	Dutton, G.	Symptoms in Orthoptic Clinics
		Problems experienced by Children with
		Cognitive Visual Dysfunction due to Cerebral
P40	Dutton, G.	Visual Impairment – and the Approaches which
		Parents Have Adopted to Deal with these
		Problem
		Case Reports - Posters
		Visual effects of ocular fundus abnormalities in
P41	Alves do Vale, P.	children with fetal alcohol syndrome
7.10		Ocular toxoplasmosis associated with Fuchs
P42	Alves do Vale, P.	heterochromic iridocyclitis
P43	Biswas, S.	Auto-immune Corneal Endotheliopathy
		Acute retinal necrosis and ipsilateral cerebral
D.1.1	0.44.5	infarction associated with vasculitis due to
P44	Sekhri, R.	varicella zoster virus in a patient with human
		immunodeficiency virus
D .1-		Blau syndrome in a 2 year old with devasting
P45	Houtman, A.	visual loss
P46	Fonseca, A.	Myastenia Gravis: Early onset in a Chinese Girl
P47	Maduro, V.	Joubert Syndrome – A case report
D40	Madura V	Ocular manifestations of congenital
P48	Maduro, V.	toxoplasmosis – 3 cases report
D40	Mantaina Manalla a a A	Topical Apraclonidine in the Diagnosis of Horner
P49	Monteiro Magalhães, A.	Syndrome
P50	Patrício, M.	Marcus-Gunn syndrome – our experience
D. 1		·
D51	Pathai S	Introduction of the Plusoptix Photoscreener into
P51	Pathai, S.	Introduction of the Plusoptix Photoscreener into a District General Hospital Ophthalmology

P52		Bakunowicz – Lazarczyk, A.	Service Transscleral intraocular lens fixation for the management of dislocated lenses in children with Marfan's syndrome Curettage of a periocular giant congenital
P53		Deconinck, H.	melanocytic naevus in the neonate : a case report
P54		Carrasquinho, S.	Treatment of Coats disease with indirect laser photocoagulation and cryotherapy – our experience
P55		Kozeis, N.	Developmental delays and severe eye problems: Is there any way out to our world? - Thessaloniki program
	14.30 – 15.50 h	VIII. Scientific Session	Neuroophthalmology – Assessment 1
	Chairman:		
L5	14.30	Hyvärinen, L.	Children with a different visual world
T33	15.10	Aring, E.	Abnormal development of ocular fixation in children with hydrocephalus
T34	15.20	Taylor, R.	Visual Field Assessment In Small Children
T35	15.30	Moser, E.	Optic Nerve Hypoplasia and Aplasia – Incidence and Clinical Aspects Discussion
	15.50 – 16.20 h		Coffee Break
	16.20 – 17.30 h Chairman:	IX. Scientific Session	Neuroophthalmology – Assessment 2
T36	16.20	Alves do Vale, P.	Optic nerve morphometry in children with fetal alcohol syndrome
T37	16.30	Audren, F.	Nonorganic Visual Disorders in Children Problems experienced by Children with Cognitive Visual Dysfunction due to Cerebral
T38	16.40	Mckillop, E.	Visual Impairment – and the Approaches which Parents Have Adopted to Deal with these

Problem

				Cognitive visual impairment (dorsal stream
		40.50	Saidkasimova, S.	dysfunction) but good visual acuities in
T39		16.50		premature and term children with mild occipito-
				parietal periventricular white matter pathology
T40		17.00	Harris Director E	Motor Rehabilitation And Visual Rehabilitation,
140		17.00	Hoogslag-Bienfait, F.	Where Two Experts Meet
				The role of index K (a global index of visual
T41		17.10	Kozeis, N.	function) as a prognostic factor in children with
				spastic cerebral palsy (CP)
T42		17.20	Stellingwerf, C.	Charles Bonnet syndrome in children
				Discussion
	17.40 h		B. Lorenz	Scientific Awards
	17.50 h		B. Lorenz	Closing Remarks
	18.00 h			End of Meeting