

CIRRUCULUM VITAE: Professor Ilhem Ben Youssef Turki

Research Laboratory of Neurological Diseases of the Child (LR18SP04), National Institute Mongi Ben Hmida of Neurology of Tunis, Faculty of Medicine, University of Tunis El Manar (UTM), Tunisia



	DOGUDION & DUDI D		
NAME: Ilhem BEN YOUSSEF-TURKI, Female, Married, Mother of 3, Date of birth: 01/05/1958, Beja, Tunisia	 POSITION & TITLE : Head of the department of pediatric neurology- National Institute Mongi Ben Hmida of Neurology- Tunis- Tunisia (NINT) Professor in Neurology: University Tunis El Manar (UTM)- Faculty of medicine- Tunis- Tunisia 		
RESEARCH GATE <u>researchgate.net/profile/Ilhem_Turki2</u> ORCID iD: <u>https://orcid.org/0000-0002-4994-</u> <u>9925</u>	 H-index: 10 (Scopus) Publications: 60 (Scopu Citations: 378 (RG) ORCID: 0000-0002-499 	1s) 94-9925	204) of Child Neurology Diseases
INSTITUTION AND STUDY LOCATION	Department of Pediatric Neurolo Neurology-Faculty of Medecine	estreet-Rabta-	
BUSINESS PHONE NUMBER	+216 71570052/ +216 98557716 Ilhem.byt@fmy.utm.tn/secretari	atneuroped@gn	
PROFESSIONAL LICENCE	National Council of Health Profe Unique identifier: 0090720864	essionals numb	er: 4845
ENGLISH LEVEL	Intermediate (Read, Spoken, wri	itten)	
ACADEMIC QUALIFICATIONS	DEGREE:	YEAR	FIELD OF STUDY/ SPECIFIC SKILLS
Faculty of medicine of Tunis University Tunis El Manar- Tunis- Tunisia	MD Professor of neurology	06/1986 12/2014	Neurologist. Child Neurology option: Neurometabolic disorders, Hereditary ataxia, Neuromuscular disorders, Epilepsy, child neuroinflamatory and Autoimmune diseases, Infectious diseases Competence and long experience in Child neurophysiology: Video- EEG-ENMG, PE
Key achievements	 Education and Professional experience: 1976: Bachelor's degree 1976-1982: Medical Study at the Faculty of Medicine of Tunis-Tunis El Manar University 1982-1986: Specialty curriculumin neurology 1986: doctorate in medicine, faculty of medicine of Tunis, UTM. 1989: specialty degree in neurology and recruitment at the National Institute of Neurology of Tunis - Tunisia 1998: teacher in neurology in faculty of medicine of Tunis, UTM. 2008: associate professor in neurology. 2015: professor in neurology. 2012 to date: head of the department of Pediatric Neurology, NINT. 2018: Chief of Research Unit 12SP24, NINT. 2018 to date: Chief of the Research Laboratory LR18SP04, NINT. Awards: 2016: award of the Société Française d'ENMG, Strasbourg. 2016: award in 11th Maghreb Congress of Neurology, Algeria. 		

 2017: award in the 15th Congress of the Pan Arab Union of Neurology: Turis ia. 2018: award in 16th Pan Arab Union of Neurology. Turis ia. 2018: award in 16th Pan Arab Union of Neurology. Fest, Morocco-November 4-6 2010. (bh Maghrebian Congress of Neurology - November 13-15-2014: Agadir. Morocco 2018: (bh Maghrebian Congress of Neurology and the 21st National Congress of Neurology November 13-15-2014: Agadir. Morocco 2018: (bh Maghrebian Congress of Neurology and the 21st National Congress of Neurology-December, 3-5, Turis 2016: (bh Maghreb Congress of Neurology November 13-15-2014: Agadir. Morocco 2017: the INMG Francophone Days-Lune01-03, 2016 - Strashourg. France 2017: the INMG Francophone Days-Lune01-03, 2016 - Strashourg. France 2017: the INMG Francophone Days-Lune01-03, 2016 - Strashourg. France 2018: 101 Maghreb Long of Neurology: A Matchead Morocco 2017: the Ist meeting of the Arisan Academy of Neurology, the 15th Congress of the Pune Arab Union of Neurology: Martha 15th Congress of Neurology (10-19). Strashourg. France 2018: 101 Maghreb Line on Okuronology: Marthaken-Morocco December 13-15, 2018. A Manama-Jordan. 2018: 110 Maghrebian Congress of Neurology: Marthaken-Morocco December 13-15, 2018. A Manama-Jordan. 2018: 121 Maghrebian Congress of Peduatric Neurology (NCA) with ICNA endorsement in 2020 and 2021. Peduatric Neurology (NCA) with ICNA endorsement in 2020 and 2021. Peduatric Neurology (NCA) with ICNA endorsement in 2020 and 2021. Peduatric Neurology-neurosacciation for the Study of Herediary Metadoir Diseases (A TTMMHI)). 2021: Sth Moroccun Congress of Peduatric Neurology (ATI). 2010: 2021: Metador of the autural connitice of NINT. 2010: 2020: Metador of the autural connitice of Chief and Adolescent with pedlesy PTT AVIA. 2013: to date: Isonad of administration	
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• 2019 to date: president of the National expert committee for the eradication of	

	• 2019 to date: Member of the creation committee of the website of NINT.
	• 2020-2023: President elected of the Tunisian Child Neurology Association
	(TCNA): tcna.org
	International activities:
	• 1992- 2010: : Member of th French-speaking ENMG group member
	• 2010 to date: Member of French-speaking ENMGass ociation member
	• 2010 to date: Member of the European Society of Pediatric Neurology (SENP)
	• 2012 to date: Member of the French League against Epilepsy (LFCE)
	• 2012 to date: Member of ICNA-ACNA
	• 2013 to date: member of the Pediatric Multiple Sclerosis Study Group (PMSSG)
	2020 to date: Member of Europeen Pediatric Neurology Society (EPNS)
	• 2005-2010: National study: Clinical Study: Federated Research Project (PRF) on National Scholar Learning - My Position: Clinical Investigator.
	• 2018: collaborative networks with Pasteur Institute of Tunis (Sonia ABDELHAK, Rym KEFI) and Mitochondrial Medicine Research Center (LENAERBS, Guy) to
	characterize the clinical and genetic aspects of Leigh syndrome in Tunisia.
Clinical Research experience	• PTR_Rejuvenage 2018_2019: study of mitochondrial dysfunction in Cockayne
	syndrome (model of early cell aging which involve oxidative stress and
	mitochondria), with the Pasteur Institute of Tunis Network (Houda YACOUB,
	Tunis and Paris via PHP Strasbourg).
	• 2020-2023: Project Tunisia- Maroco 20/ PRD-18: Clinical, Biochemical and
	Molecular Investigations of Mitochondrial Cytopathies in Children.
	• 2006: International Therapeutic Trial: Gaucher Disease N°: GZGD02507: 2006-
	My Position: Clinical investigator
	• 2014: One of the research project, led by Said Galai (LR18SP04 Research coordinator) is the biochemical approach for the diagnosis of mitochondrial
	metabolic abnormalities (redoxpoint assay: lactate / pyruvate and acetoacetate / 3
	hydroxybutyrate and enzymatic assay of mitochondrial respiratory complexes) in
	mitochondrial cytopathies. This project is the subject of research master's thesis
	and science thesis. The goal of this project is to set up biochemical tests for
	mitochondrial diseases in our laboratory. My position: support as chief of the
	Laboratory.
	 2013- 2021: Clinical and molecular study of isolated and syndromic
	leukodystrophies in Tunisia- Es Sciences Thesis- My Position: Codirector
	• 2015 (OG) to date: International Therapeutic Trail on ataxia Telangiectasia:
	IEDAT- Study IEDAT-02-2015_TU2_SIV 16-17 Jul 2019 : My Position: Co-
Research Project:	investigator
, , , , , , , , , , , , , , , , , , ,	• 2019- 2021: Retrotope Phase 2/3 Clinical Trial of RT001 and Concurrent Natural
	History Study in Patients with Infantile Neuroaxonal Dystrophy (INAD)- My
	position: Principal investigator (PI).
	• 2019- 2023: Project entitled: Cockayne syndrome and aging and the risks to them-
	Clinical and genetic study and Investigation of enzymatic defects in CS patients
	and implementation of a non-invasive diagnostic test-My Position: Co-
	investigator
	• 2020-2022: Project entitled: "Implication du stress oxydatif dans l'évolution des
	pathologies neurologiques de l'enfant : Focus sur les cytopathies mitochdriales"-
	My position: Co-investigator
	2020-2023: Tunisian-Maroccan cooperation project IN THE FIELD OF
	RESEARCH AND INNOVATION, entitled clinical, Biochemical and Molecular
	 2020-2023: Tunisian-Maroccan cooperation project IN THE FIELD OF RESEARCH AND INNOVATION, entitled clinical, Biochemical and Molecular Investigations of Mitochondrial Cytopathies in children. A ccepted on Mars 2020. My position: Principal investigator (PI).

Selected peer-reviewed publications	s:
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<u>1)</u>	Epilepsy aspects and EEG patterns in Neuro-metabolic Diseases. Turki I , Kraoua I, Kechaou M, Smirani S, BenRhouma H, Rouissi A, Gouider-
	Khouja N. Journal of Behavioral and Brain Science. 2011; 1: 69-74.
<u>2)</u>	3-Phosphoglycerate dehydrogenase deficiency: description of two new cases
	in Tunisia and review of the literature. Kraoua I, Wiame E, Kraoua L, Nasrallah F, Benrhouma H, Rouissi A, Turki I , Chaabouni H, Briand G,
	Kaabachi N, Van Schaftingen E, Gouider-Khouja N. Neuropediatrics. 2013;
	44(5) :281-5
<u>3)</u>	Infantile and childhood onset PLA2G6-associated neurodegeneration in a
	large North African cohort. Romani M, Kraoua I, Micalizzi A, Klaa H,
	Benrhouma H, Drissi C, Turki I , Castellana S, Mazza T, Valente EM, Gouider-Khouja N. Eur J Neurol. 2015 Jan ;22(1) :178-86
<u>4)</u>	Elevated as partate aminotransferase and lactate dehydrogenase levels are a
	constant finding in PLA2G6-associated neurodegeneration. Kraoua I, Romani
	M, Tonduti D, BenRhouma H, Zorzi G, Zibordi F, Ardissone A, Gouider-
	Khouja N, Ben Youssef-Turki I , Nardocci N, Valente EM. Eur J Neurol.
<u>5)</u>	2016 Apr; 23(4) :e24-5 Pyridoxine-dependent epilepsy: a novel mutation in a Tunisian child
<u>6)</u>	T. Ben Younes, I. Kraoua, H. Benrhouma, F. Nasrallah, N. Ben Achour, H.
_	Klaa, A. Hassen-Rouissi, C. Drissi, JF. Benoist, I. Ben Youssef-Turki. Arch
	Pediatr. 2017 Mar; 24(3):241-243
<u>7)</u>	Homozygous 2p11.2 deletion supports the implication of ELMOD3 in hearing loss and reveals the potential association of CAPG with ASD/ID etiology.
	Lahbib S, Leblond CS, Hamza M, Regnault B, Lemée L, Mathieu A, Jaouadi
	H, Mkaouar R, Youssef-Turki IB, Belhadj A, Kraoua I, Bourgeron T,
	Abdelhak S. J Appl Genet. 2019 Feb; 60(1):49-56. doi: 10.1007/s13353-018-
8)	0472-3. Epub 2018 Oct 4. PMID: 30284680 ABCB1 Polymorphisms and Drug-Resistant Epilepsy in a Tunisian
<u>8)</u>	Population. Malek Chouchi, Hedia Klaa, Ihem Ben-Youssef Turki, Lamia
	Hila. Hindawi Disease Markers, Volume 2019 Article ID 1343650 16 pages
	https://doi.org/10.1155/2019/1343650
<u>9)</u>	Novel POLR1C mutation in RNA polymerase III-related leukodystrophy with severe myoclonus and dystonia. Kraoua I, Karkar A, Drissi C, Benrhouma H,
	Klaa H, Samaan S, Renaldo F, Elmaleh M, Ben Hamouda M, Abdelhak S,
	Boespflug-Tanguy O, Ben Youssef-Turki I, Dorboz I. Mol Genet Genomic
10)	Med. 2019 Sep;7(9):e914.
10)	SQSTM1 mutation: Description of the first Tunisian case and literature review. M. Akkari, I. Kraoua, H. Klaa, H. Benrhouma, T. Ben Younes, A.
	Rouissi, M. Chaabouni, I. Ben Youssef Turki. Molecular Genetics &
	Genomic Medicine 02 November 2020
11\	https://doi.org/10.1002/mgg3.1543
<u>11)</u>	Hypomyelination and Congenital Cataract:Clinical, Imaging, and Genetic Findings in ThreeTunisian Families and Literature Review
	lchraf Kraoua* Yosra Bouyacoub,* Cyrine Drissi* Mariem Chargui, lbtihel
	Rebai, Ahmed Chebil, Hédia Klaa, Hanene Benrhouma, Aida Hassen, Neziha
	Gouider-Khouja, Sonia Abdelhak, odile Boespflug-Tanguy, Ilhem Ben
12)	Youssef-Turki , lmen Dorboz- Neuropediatrics 2021 ;00:1-7 book: Epilepsies dans les maladies neuro-métaboliques : Hedia Klaa- Ichraf
<u>12)</u>	Kraoua- Ilhem Turki
	ISBN-13: 978-3-639-54331-5- ISBN-10: 3639543319- EAN:
	9783639543315- Book language: French-Publishing house: Editions
	universitaires europeennes (2016-11-23) - Website: http://www.editions- ue.com/
13)	Book: Les leucodystrophies sans marqueurs biochimiques : Ichraf Kraoua-
	Thyouraya Ben Younes- Ihem Ben Youssef Turki
	ISBN-13: 978-620-3-41886-6-ISBN-10:6203418862-EAN:9786203418866-
	4/5

Book language : French-Editions universitaires europeennes (02-07-2021)-Website: http://www.editions-ue.com/

Date: 11/19/2021

Signature:

