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Mustafa Abdalla Mohamed Salih

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19. PUBLICATIONS:

- Publications were highlighted by *Nature (Nature Index)*:
http://www.nature.com/nature/journal/v532/n7600_suppl/ni/full/532S19a.html

19.1 Theses, Supplements and Book Chapters

19.1.1 **MPCH Thesis.** Unusual Muscular Dystrophy in an Extended Sudanese Kindred. (A thesis submitted in partial fulfillment of the requirement for the Degree of MPCH, University of Khartoum, April 1980).

19.1.2 **MD Thesis.** Unusual Muscular Dystrophy in an Extended Sudanese Kindred - Doctor of Medicine Thesis (Degree granted with distinction by the University of Khartoum, December 1982).

19.1.3 **Doctor of Medical Science Thesis.** Childhood Acute Bacterial Meningitis in the Sudan: An epidemiological, clinical laboratory study. *Scandinavian Journal of Infectious Diseases* 1990; Supplement 66:1 - 103. (Uppsala University, Sweden, May 1990).

19.1.4 **Salih MAM.** Neuromuscular Disorders Among Arabs. In: Teebi AS, Farag TF, eds. *Genetic Disorders Among Arab Populations*. London : Oxford University Press, 1997.

19.1.5 **Salih MAM.** Genetic disorders in the Sudan. In : Teebi AS, Farag TF, eds. *Genetic Disorders Among Arab Populations*. London : Oxford University Press, 1997.

19.1.6 **Salih MA.** Childhood Stroke. *Saudi Medical Journal* 2006; Supplement 1(Vol.27): S1- S111.

- 19.1.7 Boerkoel C, Hirano R, **Salih M**, Takashima H (October 2007) Spinocerebellar Ataxia with Axonal Neuropathy in: GeneReviews at GeneTests: Medical Genetics Information Resource [database online]. Copyright, University of Washington, Seattle, 1997-2007. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK1105/>
- 19.1.8 Cheryl Walton, Heidrun Interthal, Ryuki Hirano, **Mustafa A.M. Salih**, Hiroshi Takashima and Cornelius Boerkel. . Spinocerebellar Ataxia with Axonal Neuropathy. In: Ahmad, Shamim, ed. Diseases of DNA Repair. New York: Springer, 2010.
- 19.1.9 **Salih MAM**. Muscular Dystrophies and Myopathies in Arab Populations. In : Teebi AS, ed. Genetic Disorders Among Arab Populations (Second edition). New York: Springer, 2010.
- 19.1.10 **Salih MAM**. Genetic Disorders in Sudan. In: Teebi AS, ed. Genetic Disorders Among Arab Populations (Second edition). New York: Springer, 2010.
- 19.1.11 **Salih MAM**. **Seven chapters** in: Elzouki AY, ed. the Textbook of Clinical Pediatrics (Second edition). New York: Springer, 2012.
- 19.1.12 Carmignac V, Suominen T, Hackman P, Udd B, **Salih MA**. Salih Myopathy (January 2012) in: GeneReviews at GeneTests: Medical Genetics Information Resource [database online]. Copyright, University of Washington, Seattle , 1997-2010. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK83297/>
- 19.1.13 Fam HK, **Salih MAM**, Takashima H, Boerkoel CF: Spinocerebellar Ataxia with Axonal Neuropathy, Autosomal Recessive (April 2012) in: GeneReviews at GeneTests: Medical Genetics Information Resource [database online]. Copyright, University of Washington, Seattle, 1997-2010. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK1105/>
- 19.1.14 **Salih MA**. Neural Tube Defects. Saudi Medical Journal 2014; Supplement (Vol.35): S1- S90.
- 19.1.15 Azzedine H, LeGuern E, **Salih MA**. Charcot-Marie-Tooth Neuropathy Type 4C. 2008 Mar 31 [Updated 2015 Oct 15]. In: Pagon RA, Adam MP, Ardinger HH, et al., editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2015. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK1340/>

19.2Papers

- 19.2.1 **Salih MAM**, Hashem N. Pendred syndrome in a Sudanese family. Sudanese Journal of Paediatrics 1978; 2: 25-34.
- 19.2.2 El Karim O, **Salih MAM**. Morbidity and mortality from measles in an urban community of the Sudan. Annals of Tropical Paediatrics, Medicine and Parasitology 1980 ; 75 : 227-30.
- 19.2.3 El Karim OA, **Salih MAM**. Extra-alveolar air collection in measles. Annals of Tropical Paediatrics 1981 ; 1 : 57-60.

- 19.2.4 **Salih MAM**, Suliman GI, Hassan HS. Complications of diphtheria seen during the 1978 outbreak in Khartoum. *Annals of Tropical Paediatrics* 1981 ; 1 : 97-101.
- 19.2.5 **Salih MAM**, Omer MIA, Karrar O, Bayoumi RA, Johnson M. Severe autosomal recessive muscular dystrophy in an extended Sudanese kindred. *Developmental Medicine and Child Neurology* 1983 ; 25 : 43-52.
- 19.2.6 **Salih MAM**, Roberts DF, Omer MIA. Reflections on muscular dystrophy in Sudanese kindred. *Clinical Genetics* 1983 ; 23 : 325- 328.
- 19.2.7 **Salih MAM**, Ekmejian A, Omer MIA. Respiratory insufficiency in a severe autosomal recessive form of muscular dystrophy. *Annals of Tropical Paediatrics* 1984 ; 4: 45-48.
- 19.2.8 **Salih MAM**. Laboratory diagnosis of meningitis : An overview in relation to the Sudan. *Sudanese Journal of Paediatrics* 1984 ; 3 : 10-19.
- 19.2.9 **Salih MAM**, Suliman GI, Hassan HS. Unusual sites of diphtheritic membrane and cervical oedema. *Sudanese Journal of Paediatrics* 1984 ; 3 : 52-62.
- 19.2.10 **Salih MAM**, Lake BD, El Hag MA, Atherton DJ. Lethal epidermolytic epidermolysis bullosa : A new autosomal recessive type of epidermolysis bullosa. *British Journal of Dermatology* 1985 ; 11 : 135-143 (Selected for abstracting by the Editors of International Synopses).
- 19.2.11 Eltom M, El Mahdi EMA, **Salih MAM**, Mukhtar E, Omer MIA. A new focus of endemic goitre in the Sudan . *Tropical and Geographical Medicine* 1985 ; 37 : 15-21.
- 19.2.12 **Salih MAM**, El Hakeem HS, Suliman GI, Khatim AS. An epidemiological study of the 1978 outbreak of diphtheria in Khartoum province. *Journal of Tropical Paediatrics* 1985; 31 : 8-12.
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- 19.2.14 **Salih MAM**. Childhood muscular dystrophy : An African review. *Annals of Tropical Paediatrics* 1985 ; 5 : 167-173.
- 19.2.15 **Salih MAM**, Bender DA, McCreanor GM. A lethal familial pellagra. - like skin lesion associated with neurological and developmental impairment and the development of cataracts. *Pediatrics* 1985 ; 76 : 787-793.
- 19.2.16 **Salih MAM**. A clinical profile of diphtheria in Sudanese children. *Sudanese Journal of Paediatrics* 1986 ; 5 : 31-36.
- 19.2.17 **Salih MAM**, Ahmed HS, Hofvander Y, Danielsson D, Olcen P. Rapid diagnosis of bacterial meningitis by an enzyme immunoassay of cerebrospinal fluid.

Epidemiology and Infection 1989 ; 103 : 301- 310. (Selected by the Editors of the Clinical Digest Series as outstanding in its field).

19.2.18 **Salih MAM**, Ahmed HS, Karrar ZA, Kamil I, Osman KA, Palmgren H, Hofvander Y, Olcen O. Features of the major epidemic of group A meningococcal meningitis in Khartoum, Sudan in 1988. *Scandinavian Journal of Infectious Diseases* 1990 ; 22 : 161- 170.

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19.2.24 **Salih MAM**, Tuvemo T. Diabetes insipidus, diabetes mellitus, optic atrophy and deafness (DIDMOAD Syndrome). A clinical study in two Sudanese families. *Acta Paediatrica Scandinavica* 1991; 8:567-572.

19.2.25 **Salih MAM**, Khaleefa OH, Bushara M, Taha ZB , Musa ZA, Kamil I, Hofvander Y, Olcen P. Long term sequelae of childhood acute bacterial meningitis in a Developing Country : A study from the Sudan. *Scandinavian Journal of Infectious Diseases* 1991; 23 : 175-182.

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19.2.39 **Salih MAM**, Ahmed AA, Ahmed HS, Olcen P. An ELISA assay for the rapid diagnosis of acute bacterial meningitis. *Annals of Tropical Paediatrics* 1995; 15 : 273-278.

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- 19.2.54 **Salih MAM**, Al Rayess M, Cutshall S, Urtizberea JA, Al Turaiki MHS, Ozo CO, Straub V, Akbar M, Abid M, Andeejani A, Campbell KP. A novel form of familial congenital muscular dystrophy in two adolescents. *Neuropediatrics* 1998;29:289-293.
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- 19.2.58 Seidahmed MZ, Rooney DE, **Salih M**, Abdel Basit OB, Shaheed MM, Abdullah MA, Abomelha A. A case of partial trisomy 2q3 with clinical manifestations of Marshall - Smith syndrome. *American Journal of Medical Genetics* 1999; 85:185-188.
- 19.2.59 **Salih MAM**, Maisonobe T, Kabiraj M, Al Rayess M, Al Turaiki MHS, Akbar M, Tahan A, Urtizberea JA, Grid D, Hamadouche T, Guilbot A, Brice A, Leguern E. Autosomal recessive hereditary neuropathy with focally folded myelin sheaths and linked to chromosome 11q23 : A distinct and homogeneous entity. *Neuromuscular Disorders* 2000;10:10-15.
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- 19.2.68 Al-Jumah M, Majumdar R, Al-Rajeh S, Chaves-Carballo E, **Salih MM**, Awada A, Al-Shahwan S, Al-Uthaim S. Deletion mutations in the dystrophin gene of Saudi patients with Duchenne and Becker muscular dystrophy. *Saudi Med J.* 2002 ; 23 : 1478-1482.
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Updated 20181211