

CURRICULUM VITAE

PERSONAL DATA

Personal Details

Name : Nahla Abdel Rahman Nazmy Hafez
Birth Date & Place : February 7th 1961, Alexandria, Egypt

Contact

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JOB POSTION:

- ❖ Professor in Human Genetics Department .
Medical Research institute, Alexandria University, Egypt
General speciality: Human Genetics
Fine speciality: Clinical Cytogenetics .
- ❖ PhD program coordinator, Human Genetics Department
- ❖ Academic advisor for PhD students, Human Genetics Department
- ❖ Cytogenetic lab director, Human Genetics Department

QUALIFICATIONS:

- MBBCh, November 1984, from faculty of Medicine. Alexandria University, Egypt with estimation, very good,with price of honour.
- Master degree in –Genetics- from Medical Research institute Alexandria University Egypt, in 1992 with estimation – Excellent

- The Master degree thesis was – Cytogenetic studies in primary amenorrhoea.
- Doctor degree in –Genetics - from Medical Research institute Alexandria University Egypt in 1998.
- The doctor degree thesis was – Study of NOR heteromorphisms in cases with Down syndrome and their parents.

POSITIONS HELD:

- House officer in Alexandria University hospitals, from 1/3/1985 to 28/2/1986.
- Demonstrator in Human Genetics Department, Medical Research institute, Alexandria University from 28/8/1986 to 21/12/1992.
- Assistant Lecturer of Human Genetics in Human Genetics Department, Medical Research institute, Alexandria University, from 22/12/1992 to 29/11/1998.
- Lecturer of Human Genetics, Human Genetics Department Medical Research institute, Alexandria University, from 30/11/1998 .
- Assistant professor of Human Genetics in Human Genetics Department, Medical Research Institute, Alexandria University, from 30/12/2003.
- Professor of Human Genetics, Human Genetics Department Medical Research institute, Alexandria University, from 28/7/2009 till now .
- Cytogenetics Lab director since 2012 till now.
- Head of Human Genetics Department since 2019 till 2021.
- Emirate professor of Human Genetics, Human Genetics Department Medical Research institute, Alexandria University, from 2021 till now .

Previous Projects:

- Participated in the National Project for Prevention of Genetic Diseases and Congenital Anomalies in Newborns, 1994-1995.
- Participated in Screening for Congenital Malformations in the Newborn. January 1996-July 1996 (WHO).
- Participated in the National Project for Early Detection and Primary Intervention of Inherited Childhood Neurometabolic Diseases. September 2006-December 2008(National council for childhood and motherhood).
- Executive manager of HLAP Grant LP7-009 Alex 2013-2016 project (Development of the Cytogenetic Lab, Human Genetics Department , MRI and its qualification for accreditation) .

CONSTRUCTIVE ACTIVITY

- Participated in establishing the cytogenetic Lab G-banding, C-banding, NOR stain and FISH techniques.
- Participated in establishing the Human Genetics Clinic.
- Participated in accreditation of Cytogenetic Lab by EGAC July 2017 (iso 15189 – 2012).

SERVICES

- Participated in Lab and clinical diagnosis of patients with genetic disorders
- Supervising junior staff training in the cytogenetics lab
- Participating in training courses provided by the department
- Participates in conferences in Human Genetics

TEACHING AND ACADEMIC RESPONSIBILITIES

- *Teaching activities:*
 - MSc Human Genetics
 - PhD Human Genetics
- *Training Courses :*

Participated in training courses provided by the Human Genetics Department in:

- Cytogenetics
- Clinical cytogenetics
 - Academic Advisor for PhD students in Human Genetics
 - PhD Program coordinator in Human Genetics

NGO MEMBERSHIP

- Genetic Counseling Society.
- The Egyptian Society of Human Genetics.

PUBLICATIONS

1. A genetic study of chromosomal translocations with phenotype-karyotype correlations. The Second International Conference on Population and Molecular Genetics Update. Cairo Egypt, 30/11 to 2/12/1999.
2. Genetic aspects of delayed puberty. Bulletin Of High Institute Of Public Health, Vol 32(2), April 2002.
3. Genetic study of abnormal sexual differentiation in Alexandria ,Egypt. The Egyptian Journal Of Medical Human Genetics, Vol3(1), May 2002.
4. Cytogenetic profile of Down syndrome in Alexandria, Egypt. Eastern Mediterranean Health Journal, Vol 9(1,2), 2003.
5. Cytogenetic study in infertile patients with idiopathic oligozoospermia. Egyptian Journal of Andrology & Reproduction, Vol 16(2), July 2002.
6. C-band heteromorphism in breast cancer patients. Bulletin Of High Institute of Public Health, Vol 33(2), April 2003.
7. Cytogenetic abnormalities among children with genetic disorders attending the genetic clinic in Alexandria. Alexandria Journal of Pediatrics, Vol 17(2), July 2003.
8. Detection of microdeletions involving the DAZ locus in idiopathic male infertility. Journal of The Medical Research Institute, Vol 24(3), 2003.
9. Clinical , cytogenetic and molecular studies_ of children with autistic features. Journal of The Medical Research Institute, Vol 28(2), 2007.
10. Chromosome anomalies in a population referred for

cytogenetic analysis. A five year experience in Alexandria, Egypt. Journal of The Medical Research Institute, Vol 28(3), 2007.

11. Phenotypic variability in patients with isodicentric Y(p11.32). A clinical , molecular and cytogenetic study. The Egyptian Journal of Medical Human Genetics, Vol 9(2), Nov 2008.
12. Cytogenetic study of couples with reproductive failure in Alexandria, Egypt. The Journal of the Egyptian Public Health Association, Vol 83(3&4), 2008.
13. Disease associated balanced chromosome rearrangement: Report on 12 cases. Alexandria Journal of Pediatric, Vol 22, No22, July 2008
14. Risk factors for hypospadias - A case control study. Bulletin Of High Institute of Public Health, Vol.38(2), April 2008
15. Pericentric inversion chromosome 9 : problem and significance in clinical genetics. Alexandria Journal of Pediatrics, Vol 23(1), January 2009.
16. Detection of low level sex chromosome mosaicism in Turner syndrome by FISH. Bulletin of Alexandria Faculty of Medicine, Vol 45(1), 2009
17. Neonatal screening and molecular genetic characterization of glucose-6-phosphate dehydrogenase deficiency in Alexandria. HUGO Journal 5:229, March 2011.
18. Genetic diagnosis of Prader Willi syndrome. Middle East Journal of Medical Genetics, 5(2):45-53, July 2016.
19. Familial Reciprocal non Robertsonian translocation t(14;22) resulting in 22q11.2 deletion syndrome. Turkish Journal of Pediatrics, 61(5):780, January 2019.
20. Detection Of SHOX gene deletions in Egyptian Children with idiopathic short stature using FISH. Meta gene, 24:100697, June 2020.

- 21. Ghada Elhady, Soha Kholeif, Nahla Nazmy. Chromosomal Aberrations in 224 Couples with Recurrent Pregnancy Loss. Journal of Human Reproductive Science. 13(4):340 .October 2020.**
- 22. Descriptive study of the different phenotypes of congenital heart disease in a cohort of Egyptian patients diagnosed with Down syndrome. Medical Journal of Cairo University, Vol 91, No. 3, September: 1061-1065, 2023.**
- 23. Influence of oxytocin level and oxytocin receptor gene expression on the status and severity of autism spectrum disorder in a group of Egyptian children. Egyptian Pharmaceutical Journal. Vol 24, No. 1. January-March: 122-130, 2025.**