

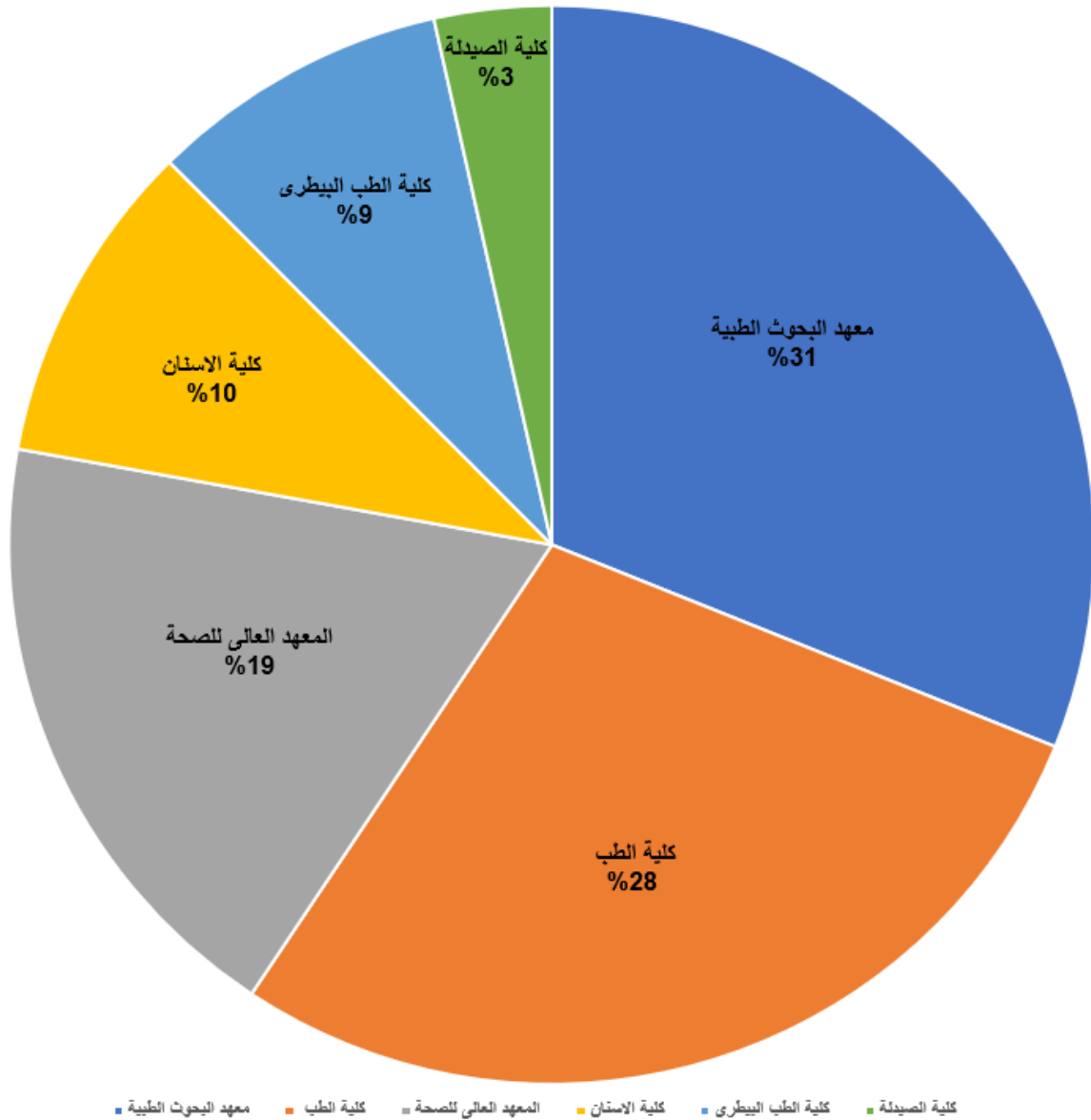
جامعة الإسكندرية
ALEXANDRIA
UNIVERSITY



GENETICS PUBLICATIONS

٢٠٥	معهد البحوث الطبية
١٨٦	كلية الطب
١٢٢	المعهد العالى للصحة
٦٤	كلية الاسنان
٥٩	كلية الطب البيطرى
٢٣	كلية الصيدلة

Registry established in August ٢٠٢١



Faculty of Medicine

Clinical Genomics Center (CGC)

- “Expanded Long Template PCR For Detection Of Full Mutation Alleles In FMR-1 Gene In Fragile X Syndrome”
Marzouk I., Kassem H., El Assi H., El Tayebani A. Expanded long template PCR for detection of full mutation alleles in FMR-1 gene in fragile X syndrome. AJM 2017; 09(4):772-84
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- “ARMS2 And HTRA1 Gene Polymorphisms In Association With Age-Related Macular Degeneration In Egypt”
https://journals.lww.com/mejmedgen/Abstract/2017/01000/ARMS2_and_HTRA1_gene_polymorphisms_in_association.3.aspx
Middle East Journal of Medical Genetics
- “Clinical Biochemical & Molecular Characterization of a Group of Egyptian PKU Patients”
<https://www.mxe.eg.net/article.asp?issn=2090-8071;year=2019;volume=8;issue=2;spage=141;epage=149;aulast=Amma>
<https://www.ingentaconnect.com/content/wk/mxe/2017/00000006/00000001/art00003>
Middle East Journal of Medical Genetics, January 2017 - Volume 6 - Issue 1 - p 22-28
- “CLINICAL GENETIC CHARACTERIZATION OF AN EGYPTIAN COHORT OF 46, XY DISORDERS OF SEXUAL DEVELOPMENT: SRD5A2 GENE ANALYSIS IN SUSPECTED STEROID 5 ALPHA REDUCTASE TYPE 2 DEFICIENCY”
http://srv1.eulc.edu.eg/eulc_v0/Libraries/Thesis/BrowseThesisPages.aspx?fn=PublicDrawThesis&BibID=12630297
- “Exploring Cancer Predisposition due to Mismatch Repair and Hereditary Diffuse Gastric Cancer-predisposing genes utilizing the 100,000 Genomes Project dataset”
- “Association Study between Catalase Gene Polymorphisms and the Susceptibility to Vitiligo in Egyptian Population.”

- “Epidemiological and Clinical Study of Genetic Skin Disorders in Kinder Garten and Primary School in Alexandria.”
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- “Detection of Human Telomerase Reverse Transcriptase Messenger RNA in Urine as a Diagnostic Tool for Urinary Bladder Cancer.”
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<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6901303/?report=classic>
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