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Biography:

Teaching courses:

- **Human Genetics**
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- **General Biology**
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Research interests:

Genetic diseases in Palestine

Publications:

- **Ghanim M**, Rossignol S, Delobel B, Irving M, Miller O, Devisme L, Plennevaux JL, Lucidarme-Rossi S, Manouvrier S, Salah A, Chivu O, Netchine I, Vincent-Delorme C. Possible association between complex congenital heart defects and 11p15 hypomethylation in three patients with severe Silver-Russell syndrome. **Am J Med Genet A. 2013 Mar;161(3):572-7. doi: 10.1002/ajmg.a.35691.**

- **Ghanim M**, Netchine I, Rossignol S, Devisme L, Delobel B, Irving M, Miller O, Plennevaux JL, Lucidarme-Rossi S, Manouvrier S, Chivu O, Vincent-Delorme C. Fetal presentation of Silver-Russell Syndrome. **European Journal of Human Genetics volume 19 supplement 2, May 2011.**

- **Ghanim M**, Guillot-Noel L, Pasquier F, Jornea L, Deramecourt V, Dubois B, Le Ber I, Brice A; French Research Network on FTD and FTD/MND. CHMP2B mutations are rare in French families with frontotemporal lobar degeneration. **J Neurol.** 2010 Dec; **257(12):2032-6.**

- Le Ber I, Camuzat A, Hannequin D, Pasquier F, Guedj E, Rovelet-Lecrux A, Hahn-Barma V, van der Zee J, Clot F, Bakchine S, Puel M, **Ghanim M**, Lacomblez L, Mikol J, Deramecourt V, Lejeune P, de la Sayette V, Belliard S, Vercelletto M, Meyrignac C, Van Broeckhoven C, Lambert JC, Verpillat P, Champion D, Habert MO, Dubois B, Brice A; French research network on FTD/FTD-MND. Phenotype variability in progranulin mutation carriers: a clinical, neuropsychological, imaging and genetic study. **Brain.** 2008 Mar; **131(Pt 3):732-46.**