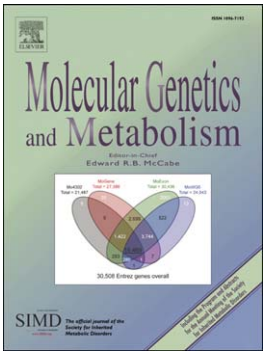

Three Rare Diseases in One Sib Pair: *RAI1, PCK1, GRIN2B* Mutations Associated with Smith-Magenis Syndrome, Cytosolic PEPCK Deficiency and NMDA Receptor Glutamate Insensitivity

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