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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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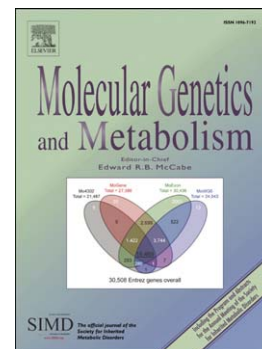
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Smith-Magenis Syndrome, Cytosolic PEPCK Deficiency and NMDA Receptor Glutamate
Insensitivity**

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