



**M Vorgerd**

**Contact**

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## Publications (1)

Vogler M, Malin J, Tegenthoff M, Lindemuth R, Pongratz D, Tettenborn B, Mortier W, Reichmann H, Burwinkel B, Kubisch C, Kilimann M. Mutation analysis in myophosphorylase deficiency (McArdle's disease). *Annals of Neurology* 1998; 43:326-331.

## Projects (0)

No results found.

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Kantonsspital St.Gallen

Rorschacher Strasse 95

CH-9007 St.Gallen

T: +41 71 494 11 11

[support.forschung@kssg.ch](mailto:support.forschung@kssg.ch)