

D-galactose

- Correction for: D-galactose induces senescence of glioblastoma cells through YAP-CDK6 pathway
 - Protective effect of oleuropein on the brain tissue in D-Galactose-induced aging in rat model
 - GOS enhances BDNF-mediated mammary gland development in pubertal mice via the gut-brain axis
 - Targeting the Leloir Pathway with Galactose-Based Antimetabolites in Glioblastoma
 - Utility of an Untargeted Metabolomics Approach Using a 2D GC-GC-MS Platform to Distinguish Relapsing and Progressive Multiple Sclerosis
 - Schizophyllum commune fruiting body polysaccharides inhibit glioma by mediating ARHI regulation of PI3K/AKT signalling pathway
 - Neurosurgical Management of Patients with Alpha-Gal Syndrome
 - Splenic nerve denervation attenuates depression-like behaviors in Chrna7 knock-out mice via the spleen-gut-brain axis
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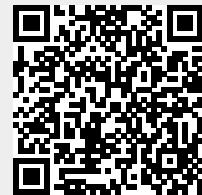
MOGHE is defined as mild **malformation** of **cortical development** with **oligodendroglial hyperplasia** in epilepsy. Approximately half of the patients with histopathologically confirmed MOGHE carry a brain somatic variant in the **SLC35A2** gene encoding a **UDP-galactose transporter**. Previous research showed that D-galactose supplementation results in clinical improvement in patients with a congenital disorder of **glycosylation** due to germline variants in SLC35A2. Aledo-Serrano et al. aimed to evaluate the effects of D-galactose **supplementation** in patients with histopathologically confirmed MOGHE, with uncontrolled **seizures** or **cognitive impairment** and epileptiform activity at the **EEG** after **epilepsy surgery** (NCT04833322). Patients were orally supplemented with D-galactose for 6 months in doses up to 1.5 g/kg/day and monitored for seizure frequency including 24-h video-EEG recording, cognition and behavioral scores, i.e., WISC, BRIEF-2, SNAP-IV, and SCQ, and quality of life measures, before and 6 months after treatment. Global response was defined by > 50% improvement of seizure frequency and/or cognition and behavior (clinical global impression of "much improved" or better). Twelve patients (aged 5-28 years) were included from three different centers. Neurosurgical tissue samples were available in all patients and revealed a brain somatic variant in SLC35A2 in six patients (non-present in the blood). After 6 months of supplementation, D-galactose was well tolerated with just two patients presenting abdominal discomfort, solved after dose spacing or reduction. There was a 50% reduction or higher of seizure frequency in 3/6 patients, with an improvement at EEG in 2/5 patients. One patient became seizure-free. An improvement of cognitive/behavioral features encompassing impulsivity (mean SNAP-IV - 3.19 [- 0.84; - 5.6]), social communication (mean SCQ - 2.08 [- 0.63; - 4.90]), and executive function (BRIEF-2 inhibit - 5.2 [- 1.23; - 9.2]) was observed. Global responder rate was 9/12 (6/6 in SLC35A2-positive). The results suggest that supplementation with **D-galactose** in patients with MOGHE is safe and well tolerated and, although the **efficacy** data warrant larger studies, it might build a rationale for **precision medicine** after **epilepsy surgery**¹⁾.

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Aledo-Serrano Á, Valls-Carbó A, Fenger CD, Groepel G, Hartlieb T, Pascual I, Herraez E, Cabal B, García-Morales I, Toledano R, Budke M, Beltran-Corbellini Á, Baldassari S, Coras R, Kobow K, Herrera DM, Del Barrio A, Dahl HA, Del Pino I, Baulac S, Blumcke I, Møller RS, Gil-Nagel A. D-galactose Supplementation for the Treatment of Mild Malformation of Cortical Development with Oligodendroglial Hyperplasia in Epilepsy (MOGHE): A Pilot Trial of Precision Medicine After Epilepsy Surgery. Neurotherapeutics. 2023 Jun 6. doi: 10.1007/s13311-023-01395-z. Epub ahead of print.

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