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Published journal articles indexed by SCI, SSCI, and AHCI

- I. Long-term clinical evaluation of patients with alpha-mannosidosis – A multicenter study
KÖSE E., KASAPKARA Ç. S., İNCİ A., YILDIZ Y., Sürücü Kara İ., Kahraman A. B., TÜMER L., DURSUN A., EMİNOĞLU F. T.
European Journal of Medical Genetics, vol.68, 2024 (SCI-Expanded)
- II. Predictors of eventual requirement of phenylalanine-restricted diet in young infants with phenylalanine hydroxylase deficiency initially managed with sapropterin monotherapy
ÇIKI K., YILDIZ Y., KAHRAMAN A. B., ÖZGÜL R. K., COŞKUN T., DURSUN A., TOKATLI A., SİVRİ S.
Molecular Genetics and Metabolism, vol.140, no.3, 2023 (SCI-Expanded)
- III. COVID-19 in inherited metabolic disorders: Clinical features and risk factors for disease severity
KAHRAMAN A. B., YILDIZ Y., ÇIKI K., Erdal I., AKAR H. T., DURSUN A., TOKATLI A., SİVRİ S.
Molecular Genetics and Metabolism, vol.139, no.2, 2023 (SCI-Expanded)
- IV. Successful management of rhabdomyolysis with triheptanoin in a child with severe long-chain 3-hydroxyacyl-coenzyme A dehydrogenase (LCHAD) deficiency
KAHRAMAN A. B., YILDIZ Y., GÖKMEN ÖZEL H., KADAYIFÇILAR S., SİVRİ S.
Neuromuscular Disorders, vol.33, no.4, pp.315-318, 2023 (SCI-Expanded)
- V. Two tales of LPIN1 deficiency: from fatal rhabdomyolysis to favorable outcome of acute compartment syndrome
KAHRAMAN A. B., Karakaya B., YILDIZ Y., KAMACI S., KESİCİ S., ŞİMŞEK KİPER P. Ö., KURT ŞÜKÜR E. D., BAYRAKCİ B., Haliloglu G.
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- VI. Diagnostic distribution and postnatal evaluation of prenatally detected short femur: A single center experience
KAHRAMAN A. B., ŞİMŞEK KİPER P. Ö., ÜTİNE G. E., BODUROĞLU O. K.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.188, no.8, pp.2367-2375, 2022 (SCI-Expanded)
- VII. Novel Cranial Imaging Findings and a Splice-Site Variant in a Patient with Tyrosinemia Type III, and a Summary of Published Cases
KAHRAMAN A. B., AKAR H. T., Lafci N. G., YILDIZ Y., Tokatli A.
MOLECULAR SYNDROMOLOGY, vol.13, no.3, pp.193-199, 2022 (SCI-Expanded)

Metrics

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