



Nurşah Yeniay Süt

MD. Specialist



 **Location:** Lokman Hekim University Ankara Hospital

 **Department:** Pediatric Neurology

 **Language:** Turkish, English

Education

Year

2013

2019

Institution

İstanbul University
SBÜ Zeynep Kamil Women's and Children's
Health Training and Research Hospital

Department

M.D
Specialization in
Pediatrics/Pediatric Neurology

Experience

Year

2025 – Still

Institution

Lokman Hekim University Ankara Hospital

Mission/ Department

Specialist in Pediatric Neurology



Publications

- Süt N.Y., Yıldırım M., Kartal A.T., Şahin S., Bektaş Ö., Teber S. Predictors associated with motor and cognitive impairment in children with motor and cognitive impairment in children with corpus callosum malformation. *Neuropediatrics*, 2025.
- Koç Yekedüz M., Köse E., İnci A., et al. Is ultrasonography a reliable approach for evaluating carpal tunnel syndrome in mucopolysaccharidosis? *Pediatric Neurology*, 2024.
- Yıldırım M., Bektaş Ö., Kartal A.T., Yeniay Süt N., Teber S. Risk of seizure relapse and long-term behavioral outcomes after antiseizure medication withdrawal in children with epilepsy. *Epilepsy & Behavior*, 2022.
- Şahin S., Botan E., Gün E., et al. Correlation between early CT findings and neurological outcome in pediatric traumatic brain injury. *Neurological Sciences*, 2024.
- Yeniay Süt N., Yıldırım M., Bektaş Ö., Kendirli T., Teber S. A case of multidrug resistant Lance-Adams syndrome successfully treated with phenobarbital. *Clinical Neuropharmacology*, 2023.
- Yılmaz Y., Türk B.G., Ser M.H., et al. Seizure treatment with olfactory training: a preliminary trial *Neurological Sciences*, 2022.
- Süt N.Y., Kartal A.T., Ertem Ş., et al. Chronic inflammatory demyelinating polyradiculoneuropathy associated with Sjögren's syndrome in a child. *Turkish Journal of Pediatrics*, 2023.
- Yeniay Süt N., Yıldırım M., Bektaş Ö., Altıntaş M., Teber S., Diagnostic challenges of congenital mirror movements in TUBB3- associated tubulinopathy. *Acta Neurologica Belgica*, 2022.
- Yılmaz Ü., Güçüyener K., Yavuz M., et al. Reexamining pediatric multiple sclerosis in the era of antibody-associated demyelination. *European Journal of Pediatric Neurology*, 2022.
- Yıldırım M., Bektaş Ö., Tunçez E., et al. Combined oxidative phosphorylation deficiency 35 with a novel TRIT1 gene variant. *Molecular Syndromology*, 2022.
- Bektaş Ö., Gülşen M., Dursun O.B., et al. Nusinersen in type 1 spinal muscular atrophy: a single-center experience. *Frontiers in Neurology*, 2025.
- Unusual Stroke Mimics: Methotrexate- Induced Reversible Stroke-Like Neurotoxicity and Leukoencephalopathy. (14th European Pediatric Neurology Society (EPNS) Congress, Glasgow, 2022.)
- Ryanodine receptor 1-related disorder presenting with malignant hyperthermia. (15th EPNS Congress, Prague, 2023.)
- Three cases of cytotoxic lesions of the corpus callosum with different etiologies. (15th EPNS Congress, Prague, 2023.)
- Episodic ataxia associated with SCN2A mutation: a case report. (15th EPNS Congress, Prague, 2023.)
- Argininosuccinate lyase deficiency causing infantile spasms: a rare metabolic disease. (17th International Child Neurology Congress (ICNC2022), Antalya, 2022)
- Rare manifestation of pediatric CIDP: hypoglossal nerve involvement. (17th ICNC, Antalya, 2022.)

