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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

I. **Effect of high-dose oral cholecalciferol on cardiac mechanics in children with chronic kidney disease**
Deveci M., AYTAÇ M. B. , Altun G., Kayabey O., Babaoglu K.
CARDIOLOGY IN THE YOUNG, cilt.27, sa.9, ss.1807-1814, 2017 (SCI İndekslerine Giren Dergi)

II. **Effect of cholecalciferol on local arterial stiffness and endothelial dysfunction in children with chronic kidney disease**
AYTAÇ M. B. , Deveci M., BEK K., Kayabey O., Ekinci Z.
PEDIATRIC NEPHROLOGY, cilt.31, sa.2, ss.267-277, 2016 (SCI İndekslerine Giren Dergi)

III. **Spironolacton Is Not A Safe And Sensitive Drug For Distinction Between Liddle Syndrome And Apperent Mineralocorticoid Excess**
Ekinci Z., AYTAÇ M. B. , Wudy S. A. , Cheong H. I.
PEDIATRIC NEPHROLOGY, cilt.29, sa.9, ss.1821, 2014 (SCI İndekslerine Giren Dergi)

IV. **Nutcracker Syndrome: Different Presentations Of The Same Disease**
AYTAÇ M. B. , Ekinci Z.
PEDIATRIC NEPHROLOGY, cilt.29, sa.9, ss.1799, 2014 (SCI İndekslerine Giren Dergi)

V. **Unusual InheritanceOf Atp6v1b1 Mutations In Four Children From Two Unrelated Families**
Ekinci Z., AYTAÇ M. B. , Cheong H. I.
PEDIATRIC NEPHROLOGY, cilt.29, sa.9, ss.1673, 2014 (SCI İndekslerine Giren Dergi)

VI. **A case of SCNN1A splicing mutation presenting as mild systemic pseudohypoaldosteronism type 1.**
Ekinci Z., Aytac M. B. , Cheong H.
Journal of pediatric endocrinology & metabolism : JPEM, cilt.26, ss.1197-200, 2013 (SCI İndekslerine Giren Dergi)

VII. **Colchicine can Induce Remission in Atypical Hemolytic Uremic Syndrome**
Ekinci Z., AYTAÇ M. B.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1679, 2012 (SCI İndekslerine Giren Dergi)

VIII. **Heterozygous factor H mutation presented as diarrhea-associated hemolytic uremic syndrome**
Ekinci Z., AYTAÇ M. B. , Hancer V. S. , Kucukkaya R. D.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1678, 2012 (SCI İndekslerine Giren Dergi)

IX. **A New Splicing Mutation of SCNN1A: Severe Neonatal Pseudohypoaldosteronism Type 1 with Normal Growth**
Ekinci Z., AYTAÇ M. B. , Cheong H. I.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1672, 2012 (SCI İndekslerine Giren Dergi)

X. **A severe case of diarrhea-associated hemolytic uremic syndrome: therapy with eculizumab**
Ekinci Z., AYTAÇ M. B.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1678, 2012 (SCI İndekslerine Giren Dergi)

XI. **Eculizumab therapy for aHUS associated with heterozygous factor I mutation**
Ekinci Z., AYTAÇ M. B. , Hancer V. S. , Kucukkaya R. D.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1680, 2012 (SCI İndekslerine Giren Dergi)

XII. **An Unusual Case of Henoch Schonlein Purpura**

AYTAÇ M. B. , Aydogan A., Yildiz K., Ekinci Z.
PEDIATRIC NEPHROLOGY, cilt.27, sa.9, ss.1693-1694, 2012 (SCI İndekslerine Giren Dergi)

XIII. **Hepatitis B vaccination in juvenile systemic lupus erythematosus.**
Aytac M. B. , Kasapcopur O., Aslan M., Erener-Ercan T., Cullu-Cokugras F., Arisoy N.
Clinical and experimental rheumatology, cilt.29, ss.882-6, 2011 (SCI İndekslerine Giren Dergi)

XIV. **Mutations in the human laminin beta2 (LAMB2) gene and the associated phenotypic spectrum.**
Matejas V., Hinkes B., Alkandari F., Al-Gazali L., Annexstad E., Aytac M. B. , Barrow M., Bláhová K., Bockenhauer D., Cheong H., et al.
Human mutation, cilt.31, ss.992-1002, 2010 (SCI İndekslerine Giren Dergi)

Diğer Dergilerde Yayınlanan Makaleler

I. **Renal outcome with eculizumab in two diarrhea-associated hemolytic–uremic syndrome cases with severe neurologic involvement**
Ekinci Z., BEK K., AYTAÇ M. B. , KARADENİZLİ A., Hancer V. S.
Hong Kong Journal of Nephrology, cilt.16, sa.2, ss.46-49, 2014 (Diğer Kurumların Hakemli Dergileri)

Atıflar

Toplam Atıf Sayısı (WOS):132
h-indeksi (WOS):3