

Kişisel Bilgiler

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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Chromosomal findings and sequence analysis of target exons of calcium-sensing receptor (CaSR) gene in patients with Sagliker syndrome**
Tunc E., DEMİRHAN O., Sağlıkler Y., Yıldız I., Paylar N., Guzel A. I.
TURKISH JOURNAL OF MEDICAL SCIENCES, cilt.47, sa.1, ss.13-21, 2017 (SCI-Expanded)
- II. **Chronic Renal Failure: Unexpected Late Sequela of Pulmonary Tuberculosis After 30 Years**
Zumrutdal A., Yildiz I., Ozelsancak R., Canpolat T.
MIKROBIOLOJİ BULTENİ, cilt.45, sa.2, ss.366-370, 2011 (SCI-Expanded)
- III. **Audiological Findings in Chronic Kidney Disease Patients With Sagliker Syndrome**
Erkan A. N., Sağlıkler Y., YILDIZ İ., Ozluoglu L.
JOURNAL OF RENAL NUTRITION, cilt.20, sa.5, 2010 (SCI-Expanded)
- IV. **A very rare case of encephalopathy in a patient with end-stage renal disease: contrast agent, ioversol**
Ozelsancak R., Erken E., Yildiz I., Giray S., Yildirim T., Micozkadioglu H.
RENAL FAILURE, cilt.32, sa.9, ss.1128-1130, 2010 (SCI-Expanded)
- V. **Left kidney starts nephrosclerosis first in primary hypertension. This sign may be called ergun sign**
Ergun Y., Sağlıkler Y., Paylar N., Aikimbaer K., YILDIZ İ.
NEPHROLOGY, cilt.13, 2008 (SCI-Expanded)
- VI. **International study on Sagliker syndrome and uglifying human face appearance in severe and late secondary hyperparathyroidism in chronic kidney disease patients**
SAGLIKER Y., ACHARYA V., LING Z., GOLEA O., SABRY A., EYUPOGLU K., OOKALKAR D. S., TAPIAWALA S., DURUGKAR S., KHETAN P., et al.
JOURNAL OF RENAL NUTRITION, cilt.18, sa.1, ss.114-117, 2008 (SCI-Expanded)
- VII. **Is survival enough for quality of life in Sagliker Syndrome-uglifying human face appearances in chronic kidney disease?**
Sağliker Y., Acharya V., Golea O., Sabry A., Bali M., Eyupoglu K., Ookalkar D., Tapiawala S., Durugkar S., Khetan P., et al.
JOURNAL OF NEPHROLOGY, cilt.21, 2008 (SCI-Expanded)
- VIII. **Neurologic manifestations in Sagliker syndrome: Uglifying human face appearance in severe and late secondary hyperparathyroidism in chronic renal failure patients**
GIRAY S., SAGLIKER Y., Yildiz I., Halvacı I., PAYLAR N., OCAL F., Balal M., OZKAYNAK P. S., Paydas S., SAGLIKER C., et al.
JOURNAL OF RENAL NUTRITION, cilt.16, sa.3, ss.233-236, 2006 (SCI-Expanded)
- IX. **Cephalometric evaluation of patients with Sagliker syndrome: Uglifying human face appearance in severe and late secondary hyperparathyroidism in chronic renal failure patients**
Uzel A., UZEL I., SAGLIKER Y., Yildiz I., Halvacı I., PAYLAR N., OCAL F., Balal M., OZKAYNAK P. S., Paydas S., et al.
JOURNAL OF RENAL NUTRITION, cilt.16, sa.3, ss.229-232, 2006 (SCI-Expanded)

Diğer Dergilerde Yayınlanan Makaleler

- I. **Case Image: Left ventricular pseudoaneurysm as a silent complication of non-ST segment elevation myocardial infarction.**
YILDIZ I., Özmen Y., GÜRBAK İ., Kaya B.
Turk Kardiyoloji Dernegi arsivi : Turk Kardiyoloji Derneginin yayin organidir, cilt.46, ss.164, 2018 (ESCI)
- II. **Diffuse Alveolar Hemorrhage Associated with Warfarin Therapy.**
Kaya B., YILDIZ I., BAHA R., ZEYTUN N., YETISGEN A.
Case reports in medicine, cilt.2015, ss.350532, 2015 (ESCI)
- III. **The Effect of Serum Mannose-Binding Lectin Levels on Dialysis-Related Peritonitis and Catheter-Related Bacteremia**
Erken E., Torun D., Sezgin N., Micozkadioglu H., Zumrutdal A., Ozelsancak R., Yildiz I.
TURKISH NEPHROLOGY DIALYSIS AND TRANSPLANTATION JOURNAL, cilt.24, sa.2, ss.189-194, 2015 (ESCI)

Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

- I. **WHOLE 13 EXONS OF GNAS1 GENE IN SAGLIKER SYNDROME(SS). COMBINATION-COMPULSION OF BONE DYSPLASIAS-HEREDITARY OSTEODISTROPHIES(BD), CHRONIC KIDNEY DISEASES (CKD) AND SECONDARY HYPERPARATHYROIDISM(SH)**
DEMİRHAN O., SAGLIKER Y., AKBAL E., PAYLAR N., Sagliker H. S., Ozkaynak P. S., YILDIZ İ., INANDIKLIOGLU N., TUNC E., BALAL M., et al.
52nd Congress of the European-Renal-Association-European-Dialysis-and-Transplant-Association, London, Kanada, 28 - 31 Mayıs 2015, cilt.30
- II. **THE WHOLE 13 EXONS OF GNAS1 GENE AND THE GENES FOR HEREDITARY OSTEODISTROPHIAS(HO) IN SAGLIKER SYNDROME(SS). COMBINATION-COMPULSION OF HEREDITARY OSTEODISTROPHIAS AND CHRONIC KIDNEY DISEASES (CKD) ?**
SAGLIKER Y., DEMİRHAN O., YILDIZ İ., PAYLAR N., INANDIKLIOGLU N., AKBAL E., TUNC E.
51st Congress of the European-Renal-Association(ERA)/European-Dialysis-and-Transplant-Association (EDTA), Amsterdam, Hollanda, 31 Mayıs - 03 Haziran 2014, cilt.29, ss.381
- III. **Cephalometric evaluation of the patients with Sagliker Syndrome: A preliminary report**
Sagliker Y., Uzel A., Uzel I., Yildiz I., Paylar N., Halvacı I.
44th ERA-EDTA Congress, Barcelona, İspanya, 22 - 24 Haziran 2007, cilt.22, ss.361
- IV. **Nephrosclerosis of primary hypertension starts first in the left kidney and this sign could be called as Ergun sign.**
Ergun Y., Sagliker Y., Aikimbaer K., Yildiz I., Firincioglu H.
21st Scientific Meeting of the International-Society-of-Hypertension/5th Asian-Pacific Congress of Hypertension/29th Annual Scientific Meeting of the Japanese-Society-of-Hypertension, Fukuoka, Japonya, 15 - 19 Ekim 2006, cilt.24, ss.340-341

Metrikler

Yayın: 16

Atıf (WoS): 81

Atıf (Scopus): 70

H-İndeks (WoS): 5

H-İndeks (Scopus): 5