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## Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Chromosomal findings and sequence analysis of target exons of calcium-sensing receptor (CaSR) gene in patients with Sagliker syndrome**  
Tunc E., DEMİRHAN O., Sağlık Y., Yıldız I., Paylar N., Guzel A. I.  
TURKISH JOURNAL OF MEDICAL SCIENCES, vol.47, no.1, pp.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.  
MIKROBIYOLOJİ BÜLTENİ, vol.45, no.2, pp.366-370, 2011 (SCI-Expanded)
- III. **Audiological Findings in Chronic Kidney Disease Patients With Sagliker Syndrome**  
Erkan A. N., Sagliker Y., YILDIZ İ., Ozluoglu L.  
JOURNAL OF RENAL NUTRITION, vol.20, no.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, vol.32, no.9, pp.1128-1130, 2010 (SCI-Expanded)
- V. **Left kidney starts nephrosclerosis first in primary hypertension. This sign may be called ergun sign**  
Ergun Y., Sagliker Y., Paylar N., Aikimbaer K., YILDIZ İ.  
NEPHROLOGY, vol.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, vol.18, no.1, pp.114-117, 2008 (SCI-Expanded)
- VII. **Is survival enough for quality of life in Sagliker Syndrome-uglifying human face appearances in chronic kidney disease?**  
Sagliker Y., Acharya V., Golea O., Sabry A., Bali M., Eyupoglu K., Ookalkar D., Tapiawala S., Durugkar S., Khetan P., et al.  
JOURNAL OF NEPHROLOGY, vol.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, vol.16, no.3, pp.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, vol.16, no.3, pp.229-232, 2006 (SCI-Expanded)

## Articles Published in Other Journals

- 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, vol.46, pp.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, vol.2015, pp.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, vol.24, no.2, pp.189-194, 2015 (ESCI)

## Refereed Congress / Symposium Publications in Proceedings

- 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, Canada, 28 - 31 May 2015, vol.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, Netherlands, 31 May - 03 June 2014, vol.29, pp.381
- III. **Cephalometric evaluation of the patients with Sagliker Syndrome: A preliminary report**  
Sagliker Y., Uzel A., Uzel I., Yildiz I., Paylar N., Halvaci I.  
44th ERA-EDTA Congress, Barcelona, Spain, 22 - 24 June 2007, vol.22, pp.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, Japan, 15 - 19 October 2006, vol.24, pp.340-341

## Metrics

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