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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., Sağliker 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., Yıldız I., Ozelsancak R., Canpolat T.  
MIKROBİYOLOJİ BULTENİ, 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., Yıldız I., Giray S., Yıldırım 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., Yıldız I., Halvaci 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., Yıldız I., Halvaci 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 Derneği arsivi : Turk Kardiyoloji Derneğinin yayın organıdır, vol.46, pp.164, 2018 (ESCI)
- II. **Diffuse Alveolar Hemorrhage Associated with Warfarin Therapy.**  
Kaya B., YILDIZ I., BAHÀ R., ZEYTUN N., YETİSGEN 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., INANDIKLOGLU N., TUNC E., BALAL M., et al.  
52nd Congress of the European-Renal-Association-European-Dialysis-and-Transplant-Assocation, 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., INANDIKLOGLU 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

Publication: 16  
Citation (WoS): 81  
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H-Index (Scopus): 5