Prof. PINAR ATA

Personal Information

Office Phone: +90 6254545 Extension: 4714
Email: pinar.ata@marmara.edu.tr
Other Email: pinaren@gmail.com
Web: https://avesis.marmara.edu.tr/pinar.ata

Education Information

Doctorate, Marmara University, School of Medicine, Internal Medical Sciences, Turkey 1999 - 2006
Expertise In Medicine, Sağlık Bakanlığı Haydarpaşa Numune Eğitim Ve Araştırma Hast, Turkey 1997 - 2001
Under Graduate, Marmara University, School of Medicine, Turkey 1988 - 1994

Foreign Languages

English, C1 Advanced

Certificates, Courses and Trainings

Health&Medicine, Herediter Kanser Hastalıklarında Aileye Yaklaşım, Marmara Üniversitesi Tıp Fakültesi Pendik EĞİTİM VE ARAŞTIRMA HASTANESİ, 2019

Dissertations

Doctorate, Düşük dereceli gliomalarda egfr, pten, pdgfr-alfa, MGMT ifadelerindeki, TP53 ve P53 yolağı (MDM-2 ve P14arf) genlerindeki düzensizlikler ve klinik etkileri, Marmara Üniversitesi, Tıp Fakültesi, 2006
Expertise In Medicine, Trisomi 21'in ve Nöral Tüp Defektinin Prenatal Tanısında İkinci Trimester AFP, hCG, uE3 Testlerinin Klinik Sonuçları, University Of Health Sciences, Ankara Keçiören Educatıon And Research Hospıtal, Department Of Internal Medıcıne, 2001

Research Areas

Health Sciences, Natural Sciences

Academic Titles / Tasks

Professor, Marmara University, School of Medicine, Internal Medical Sciences, 2018 - Continues
Associate Professor, Marmara University, School of Medicine, Internal Medical Sciences, 2012 - Continues

Advising Theses

ATA P., Ankilozan spondilit'te mikrorna ekspresyonu ve etklili epigenetik mekanizmaların incelenmesi, Expertise In Medicine, A.TÜRKYILMAZ(Student), 2018
ATA P., Kronik böbrek yetersizliği hastalarında idrar bulundurmayan mesane dokusunda IL-8, CXR1 ve CXR2 nin gen ifadelerinin araştırılması, Post Graduate, K.GÜNDOĞDU(Student), 2013
ATA P., Kronik böbrek yetersizliği hastalarında idrar bulundurmayan mesane dokusunda apoptotik bulguların incelenmesi, Post Graduate, H.FINDIK(Student), 2013

Jury Memberships

Associate Professor Exam, Associate Professor Exam, Marmara Üniversitesi, August, 2020
Post Graduate, Post Graduate, Marmara Üniversitesi, July, 2020
Appointment Academic Staff, Appointment Academic Staff, Sağlık Bilimleri Üniversitesi, July, 2020
Associate Professor Exam, Associate Professor Exam, Çukurova Üniversitesi, June, 2020

Taught Courses And Trainings

Türköz H. K. , Atagündüz I., Bozkurt S., Ata P., Toptaş T., Myelodisplastik sendrom: patogenezden tedaviye yenilikler sempozyumu, 2016 - 2016

Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

I. **A large Turkish pedigree with multiple endocrine neoplasia type 1 syndrome carrying a rare mutation: c.1680_1683 del TGAG**
DEMİRTAŞ C. Ö. , ATA P., ÇETİN A., Turkyilmaz A., DUMAN D.
TURKISH JOURNAL OF GASTROENTEROLOGY, vol.31, pp.508-514, 2020 (Journal Indexed in SCI)

II. **A Rare Cause of Hypophosphatemia: Raine Syndrome Changing Clinical Features with Age**
ELTAN M., ALAVANDA C., Yavas Abali Z., Ergenekon P., Yalindag Ozturk N., SAKAR M., DAĞÇINAR A., KIRKGÖZ T., KAYGUSUZ S. B. , GÖKDEMİR Y., et al.
CALCIFIED TISSUE INTERNATIONAL, vol.107, pp.96-103, 2020 (Journal Indexed in SCI)

III. **THE EXPRESSION LEVELS OF microRNAs ASSOCIATED WITH T AND B CELL DIFFERENTIATION/STIMULATION IN ANKYLOSING SPONDYLITIS**
Turkyilmaz A., Ata P., Akbas F., Yagci I.
BALKAN JOURNAL OF MEDICAL GENETICS, vol.23, pp.25-31, 2020 (Journal Indexed in SCI)

IV. **Cinacalcet as a First-Line Treatment in Neonatal Severe Hyperparathyroidism Secondary to Calcium Sensing Receptor (CaSR) Mutation**
Gulcan-Kersin S., Kirkgoz T., ELTAN M., Rzayev T., ATA P., Bilgen H., ÖZEK E., BEREKET A., Turan S.
Hormone Research in Paediatrics, 2020 (Journal Indexed in SCI Expanded)

V. **Fibrodysplasia ossificans progressiva: lessons learned from a rare disease.**
Akyuz G. D. , Gencer-Atalay K., Ata P.
Current opinion in pediatrics, vol.31, pp.716-722, 2019 (Journal Indexed in SCI)

VI. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**
KAYGUSUZ S. B. , ARMAN A., ABALI S., ATA P., KIRKGÖZ T., ELTAN M., Abali Z. Y. , Helvacioglu D., Tosun B. G. , Menevse T. S. , et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.214, 2019 (Journal Indexed in SCI)

VII. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**
KAYGUSUZ S. B. , ATA P., KIRKGÖZ T., Abali Z. Y. , ELTAN M., Tosun B. G. , Menevse T. S. , Helvacioglu D., GÜRAN T., ARMAN A., et al.

HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.215, 2019 (Journal Indexed in SCI)

VIII. **A rare cause of hypophosphatemia: Raine Syndrome**
ELTAN M., ATA P., KIRKGÖZ T., ALAVANDA C., KAYGUSUZ S. B. , Menevse T. S. , Tosun B. G. , Abali Z. Y. , Helvacioglu D., GÜRAN T., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.384, 2019 (Journal Indexed in SCI)

IX. **Novel mutations and deletions in cystic fibrosis in a tertiary cystic fibrosis center in Istanbul.**
Atag E., Bas I., Ergenekon A., Gokdemir Y., Eralp E., Ata P., Ersu R., Karakoc F., Karadag B. T.
Pediatric pulmonology, vol.54, pp.743-750, 2019 (Journal Indexed in SCI)

X. **Association between occurrence of ossicular chain defect and osteoprotegerin gene expression in patients with chronic otitis media**
Keskin S., Tatlipinar A., ATA P., Uzun S., Kinal M. E. , Erkal B.
EUROPEAN ARCHIVES OF OTO-RHINO-LARYNGOLOGY, vol.276, pp.1321-1325, 2019 (Journal Indexed in SCI)

XI. **Challenges in the treatment of fibrodysplasia ossificans progressiva**
Gencer-Atalay K., Ozturk E. C. , YAĞCI İ., ATA P., DELİL K., Ozgen Z., Akyuz G.
RHEUMATOLOGY INTERNATIONAL, vol.39, pp.569-576, 2019 (Journal Indexed in SCI)

XII. **THE CO-EXISTENCE OF TWO RARE DISEASES: A CASE REPORT**
Saki M., KIRKGÖZ T., GÖKCE İ., Cicek N., ATA P., Turan S., YILDIZ N., ALPAY H.
PEDIATRIC NEPHROLOGY, vol.33, pp.1881, 2018 (Journal Indexed in SCI)

XIII. **NOVEL MUTATIONS AND VARIATIONS IN TURKISH CHILDREN WITH ALPORT SYNDROME: ARE THE BENIGN VARIATIONS ALWAYS HARMLESS?**
Cicek N., YILDIZ N., ATA P., GÖKCE İ., SAK M., KAYA H., ALPAY H.
PEDIATRIC NEPHROLOGY, vol.33, pp.1918-1919, 2018 (Journal Indexed in SCI)

XIV. **Intraoperative Tissue-Immunosuppressive Therapy Reduces Rejection Episodes in Heart Transplant Recipients.**
Rabus M., Cekmecelioglu D., Ata P., Salihi S., Selcuk E., Balkanay M.
Experimental and clinical transplantation : official journal of the Middle East Society for Organ Transplantation, 2018 (Journal Indexed in SCI Expanded)

XV. **THE ROLE OF SOLUBLE CTLA-4 AS A NON-INVASIVE BIOMARKER FOR DIAGNOSIS OF KIDNEY ALLOGRAFT REJECTION: A PRELIMINARY STUDY**
Ruhi C. B. , ATA P., Titiz I. M.
TRANSPLANT INTERNATIONAL, vol.30, pp.303, 2017 (Journal Indexed in SCI)

XVI. **THE INFLUENCE OF PREFORMED HLA CLASS I AND II PANEL REACTIVE ANTIBODIES ON CLINICAL AND PATHOLOGICAL OUTCOMES OF KIDNEY ALLOGRAFT**
Ruhi C., Tugcu M., Kasapoglu U., Boynuegri B., Gumrukcu G., ATA P., Apaydin S., Titiz I. M.
TRANSPLANT INTERNATIONAL, vol.30, pp.318, 2017 (Journal Indexed in SCI)

XVII. **COMPARISON OF THE TREATMENT EFFICACY OF RITUXIMAB AND PLASMAPHERESIS/INTRAVENOUS IMMUNOGLOBULIN COMBINATION WITH HISTORICAL CONTROL IN CHRONIC ANTIBODY MEDIATED REJECTION**
Ruhi C., TugCu M., KasapogLu U., Gokce A. M. , ATA P., Titiz I. M.
TRANSPLANT INTERNATIONAL, vol.30, pp.300-301, 2017 (Journal Indexed in SCI)

XVIII. **DOES C.3979 G > A/P.VAL1327MET VARIANT OF COL4A4 HAS ANY PATHOGENIC EFFECT IN TURKISH PATIENTS WITH ALPORT SYNDROME?**
YILDIZ N., ATA P., ALPAY H., GÖKCE İ., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.32, pp.1773, 2017 (Journal Indexed in SCI)

XIX. **Cytotoxic Antibody Detection by Means of Flow-Cytometric Cross-Match**
Bilgen T., Ata P., Tozkir J., Tozkir H., Titiz M. I.
TRANSPLANTATION PROCEEDINGS, vol.49, pp.440-444, 2017 (Journal Indexed in SCI)

XX. **Evaluation of Pre-Transplant Panel Reactive Antibody Levels and Sensitization: A Single-Center Study**
Can O., Gokce A. M. , Canbakan M., ATA P., Sahin G. M. , TİTİZ M. İ. , Apaydin S.
ANNALS OF TRANSPLANTATION, vol.21, 2016 (Journal Indexed in SCI)

XXI. **Determination of CYP2C19 Polymorphism, Side Effects, and Medication Adherence in Patients Who have Utilized Selective Serotonin Reuptake Inhibitors**
Deniz S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., Talas A., Gunes T., Caliskan M., Izzettin F. V.
KLINIK PSIKOFARMAKOLOJI BULTENI-BULLETIN OF CLINICAL PSYCHOPHARMACOLOGY, vol.26, pp.152-160, 2016 (Journal Indexed in SCI)

XXII. **A Rare Reason of Ileus in Renal Transplant Patients With Peritoneal Dialysis History: Encapsulated Peritoneal Sclerosis.**
Gökçe A., Özel L., İbişoğlu S., Ata P., Şahin G., Gücün M., Kara V., Özdemir E., Titiz M.
Experimental and clinical transplantation : official journal of the Middle East Society for Organ Transplantation, vol.13, pp.588-92, 2015 (Journal Indexed in SCI Expanded)

XXIII. **Cancer Screening of Renal Transplant Patients Undergoing Long-Term Immunosuppressive Therapy**
Demir T., Ozel L., Gokce A. M. , Ata P., Kara M., Eris C., Ozdemir E., Titiz M. I.
TRANSPLANTATION PROCEEDINGS, vol.47, pp.1413-1417, 2015 (Journal Indexed in SCI)

XXIV. **Relationship of Urothelial Gene Expressions in Urine-Deprived Bladders of Renal Recipients With Posttransplant Urinary Infections**
Gokce A. M. , Findik H., Ata P., Gumrukcu G., Ozel L., Gundogdu K., Yazicioglu B., Titiz M. I.
TRANSPLANTATION PROCEEDINGS, vol.47, pp.1331-1335, 2015 (Journal Indexed in SCI)

XXV. **Factors Affecting the Selection of Patients on Waiting List: A Single Center Study.**
Can Ö., Kasapoğlu U., Boynueğri B., Tuğcu M., Çağlar R., Canbakan M., Murat G., Ata P., İzzet T., Apaydın S.
Transplantation proceedings, vol.47, pp.1265-8, 2015 (Journal Indexed in SCI Expanded)

XXVI. **Determination of CYP2C19 polymorphisms, adverse drug reaction, and medication adherence in patients utilized selective serotonin reuptake inhibitors**
Deniz S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., Talas A., Gunes T., Caliskan M., Izzettin F. V.
INTERNATIONAL JOURNAL OF CLINICAL PHARMACY, vol.37, pp.231-232, 2015 (Journal Indexed in SCI)

XXVII. **Proteinuria associated with mTOR inhibitors after kidney transplant.**
Guney M., Sahin G., Yilmaz B., Canbakan M., Gucun M., Kayatas K., Eren P., Titiz I.
Experimental and clinical transplantation : official journal of the Middle East Society for Organ Transplantation, vol.12, pp.539-42, 2014 (Journal Indexed in SCI Expanded)

XXVIII. **The GABA(A) Receptor gamma 2 Subunit (R43Q) Mutation in Febrile Seizures**
Hancili S., Onal Z. E. , Ata P., Karatoprak E. Y. , Gurbuz T., Bostanci M., Pacal Y., Nuhoglu C., CERAN Ö.
PEDIATRIC NEUROLOGY, vol.50, pp.353-356, 2014 (Journal Indexed in SCI)

XXIX. **TLR4 gene polymorphism in patients with nonalcoholic fatty liver disease in comparison to healthy controls.**
Kiziltas S., Ata P., Colak Y., Mesçi B., Senates E., Enc F., Ulasoglu C., Tuncer I., Oguz A.
Metabolic syndrome and related disorders, vol.12, pp.165-70, 2014 (Journal Indexed in SCI Expanded)

XXX. **EFFECTIVENESS OF PLASMAPHERESIS AS A TREATMENT OF CHOICE IN SENSITIZED RENAL RECIPIENTS**
Ata P., Ibisoglu S., Canbakan M., Gucun M., Gokce A. M. , Kara M., Gumrukcu G., Titiz I.
TRANSPLANT INTERNATIONAL, vol.26, pp.79, 2013 (Journal Indexed in SCI)

XXXI. **Association of COL1A1 polymorphism in Turkish patients with otosclerosis.**
Ertugay O., Ata P., Kalaycik E., Kaya K., Tatlipinar A., Kulekci S.
American journal of otolaryngology, vol.34, pp.403-6, 2013 (Journal Indexed in SCI Expanded)

XXXII. **Monitoring of CD3(+) T-cell count in patients receiving antithymocyte globulin induction after cadaveric renal transplantation.**
Ata P., Kara M., Özdemir E., Canbakan M., Gökçe A., Bayraktar F., Şahin G., Özel L., Titiz M.
Transplantation proceedings, vol.45, pp.929-31, 2013 (Journal Indexed in SCI Expanded)

XXXIII. **The impact of preoperative immunonutrition and other nutrition models on tumor infiltrative lymphocytes in colorectal cancer patients.**
Caglayan K., Oner I., Gunerhan Y., Ata P., Koksal N., Ozkara S.
American journal of surgery, vol.204, pp.416-21, 2012 (Journal Indexed in SCI Expanded)

XXXIV. **Serum flow cytometric C1q binding antibody analysis of renal recipients with low levels of**

sensitization.
Ata P., Canbakan M., Kara M., Özel L., Ünal E., Titiz M.
Transplantation proceedings, vol.44, pp.1652-5, 2012 (Journal Indexed in SCI Expanded)

XXXV. **The impact of C4d staining as a humoral injury marker.**
Kara M., Demir F., Ata P., Ozel L., Gumrukcu G., Unal E., Canbakan M., Gucun M., Esadoglu V., Ozdemir E., et al.
Transplantation proceedings, vol.44, pp.1694-6, 2012 (Journal Indexed in SCI Expanded)

XXXVI. **Flow Cytometric Detection of Anti-AB Antibody Titers in Blood Group O Recipients of Blood Group A2 Donor Kidneys**
Ata P., Cetinkaya F., Ozgezer T., Ozel L., Tulunay A., Eksioglu E., Titiz M. I.
TRANSPLANTATION PROCEEDINGS, vol.44, pp.1706-1709, 2012 (Journal Indexed in SCI)

XXXVII. **Ramsay Hunt syndrome with atypical progress in a renal transplant recipient: a case report.**
Ozel L., Toros S., Unal E., Kara M., Eren P., Canbakan M., Kucuk M., Titiz I.
Experimental and clinical transplantation : official journal of the Middle East Society for Organ Transplantation, vol.9, pp.413-6, 2011 (Journal Indexed in SCI Expanded)

XXXVIII. **Pyoderma gangrenosum in a renal transplantation patient having immunosuppressive treatment for 5 years.**
Serdar Z., Ata P., Titiz M.
Transplant international : official journal of the European Society for Organ Transplantation, vol.24, 2011 (Journal Indexed in SCI)

XXXIX. **Risk factors for osteoporosis after renal transplantation and effect of vitamin D receptor Bsm I polymorphism.**
Ozel L., Ata P., Ozel M., Toros A., Kara M., Unal E., Canbakan M., Erdogrul G., Aktas G., Titiz M.
Transplantation proceedings, vol.43, pp.858-62, 2011 (Journal Indexed in SCI)

XL. **Elective and emergency surgery in chronic hemodialysis patients.**
Ozel L., Krand O., Ozel M., Toros A., Sağıroğlu J., Kara M., Erdoğdu E., Yiğit B., Ata P., Çavdar F., et al.
Renal failure, vol.33, pp.672-6, 2011 (Journal Indexed in SCI)

XLI. **Effects of intra- and extracellular factors on anti-aging klotho gene expression**
TURAN K., ATA P.
GENETICS AND MOLECULAR RESEARCH, vol.10, pp.2009-2023, 2011 (Journal Indexed in SCI)

XLII. **Dermatologic Findings in Renal Transplant Recipients: Possible Effects of Immunosuppression Regimen and p53 Mutations**
Serdar Z. A. , Eren P., Canbakan M., Turan K., Tellioglu G., Gulle S., Ozgezer T., Kara M., Berber I., Titiz M. I.
TRANSPLANTATION PROCEEDINGS, vol.42, pp.2538-2541, 2010 (Journal Indexed in SCI)

XLIII. **Evaluation of intraoperative parathormone measurement for predicting successful surgery in patients undergoing subtotal/total parathyroidectomy due to secondary hyperparathyroidism.**
Kara M., Tellioglu G., Bugan U., Krand O., Berber I., Seymen P., Eren P., Ozel L., Titiz I.
The Laryngoscope, vol.120, pp.1538-44, 2010 (Journal Indexed in SCI Expanded)

XLIV. **Determination of the Risk Group in Patients with Venous Thrombosis**
Eren P., Denizli N., Sokmen H. M. , Erdem S., Solak M.
TURKIYE KLINIKLERI TIP BILIMLERI DERGISI, vol.29, pp.1430-1434, 2009 (Journal Indexed in SCI)

XLV. **The clinical significance of parathyroid tissue calcium sensing receptor gene polymorphisms and expression levels in end-stage renal disease patients.**
ATA EREN P., TURAN K., BERBER İ., CANBAKAN M.
Clinical nephrology, vol.72, pp.114-21, 2009 (Journal Indexed in SCI)

XLVI. **The clinical significance of parathyroid tissue calcium sensing receptor gene polymorphisms and expression levels in end-stage renal disease patients.**
Eren P., Turan K., Berber I., Canbakan M., Kara M., Tellioglu G., Bugan U., Sevinç C., Turkmen F., Titiz M.
Clinical nephrology, vol.72, pp.114-21, 2009 (Journal Indexed in SCI Expanded)

XLVII. **Calcium-sensing receptor gene polymorphisms and cardiac valvular calcification in patients with chronic renal failure: a pilot study.**
Turkmen F., Ozdemir A., Sevinc C., Eren P., Demiral S.

Hemodialysis international. International Symposium on Home Hemodialysis, vol.13, pp.176-80, 2009 (Journal Indexed in SCI Expanded)

XLVIII. **Evaluation of Gadolinium Pre-Treatment with or without Splenectomy in the Setting of Renal Ischemia Reperfusion Injury in Rats**
Kara M., Tellioglu G., Sehirli O., Yildar M., Krand O., Berber I., ÇETİNEL Ş., Eren P., Sener G., Titiz I.
RENAL FAILURE, vol.31, pp.956-963, 2009 (Journal Indexed in SCI)

XLIX. **Cerebellar liponeurocytoma/lipidized medulloblastoma - Case report and review of the literature**
Aker F., Ozkara S., Eren P., Peker O., Armagan S., Hakan T.
JOURNAL OF NEURO-ONCOLOGY, vol.71, pp.53-59, 2005 (Journal Indexed in SCI)

Articles Published in Other Journals

I. **Association of TBX21 gene polymorphism with nasal polyposis**
KAYA K. S. , ATA P., ERTUGAY Ö. Ç. , KÜLEKÇİ S., ZER TOROS S.
Praxis of ORL, vol.7, pp.145-150, 2019 (Refereed Journals of Other Institutions)

II. **Comparison of the Treatment Efficacy of Rituximab and Plasmapheresis/Intravenous Immunoglobulin Combination with Historical Control in Chronic Antibody Mediated Rejection**
Ruhi C., Tugcu M., Kasapoglu U., Gokce A. M. , ATA P., Titiz M. I.
TURKISH NEPHROLOGY DIALYSIS AND TRANSPLANTATION JOURNAL, vol.26, pp.48-54, 2017 (Journal Indexed in ESCI)

Books & Book Chapters

I. **Bölüm 1. İmmunolojik Terimler Sözlüğü**
ATA P.
in: Laboratuvardan Kliniğe Transplantasyon Pratiği, ATA PINAR, Editor, NAMIK KEMAL ÜNİVERSİTESİ, pp.17-20, 2017

II. **Bölüm 2. Bağışıklık Sistemi ve Antikorlar**
ATA P.
in: Laboratuvardan Kliniğe Transplantasyon Pratiği, ATA PINAR, TİTİZ İZZET TİTİZ, Editor, NAMIK KEMAL ÜNİVERSİTESİ, pp.20-48, 2017

III. **Transplantasyon Antijenleri ve Genetiği**
ATA P.
in: Renal Transplantasyona Pratik Yaklaşım, MESUT İZZET TİTİZ, Editor, NOON TANITIM, İstanbul, pp.47-58, 2010

IV. **Transplantasyon İmmünolojisine Giriş,**
ATA P.
in: Renal Transplantasyona PratikYaklaşım, Mesut İzzet Titiz, Editor, NOON TANITIM, İstanbul, pp.29-42, 2010

Refereed Congress / Symposium Publications in Proceedings

I. **Expression Profile Of Complement Activation At Decidual Tissue In Patients With Recurrent Pregnancy Loss**
YILMAZ İ., ATA P., gürkan karakaş n., ESİM BÜYÜKBAYRAK E.
European Human Genetics Virtual Conference 2020, 6 - 09 June 2020

II. **GENETIC ANALYSIS IN TURKISH CHILDRENWITH ALPORT SYNDROME: NOVEL MUTATIONS AND VARIATIONS**
çiçek n., YILDIZ N., ATA P., GÖKCE İ., KAYA H., ALPAY H.
18 th Congress of the International Pediatric Nephrology Association-IPNA, Italy, 17 - 21 October 2019, vol.34,

pp.1821-2260

III. **A NOVEL MUTATION IN ATP6V0A4 GENE IN APATIENT WITH DISTAL RENAL TUBULAR ACIDOSIS**
SAK M., YILDIZ N., ÇİÇEK N., GÜVEN S., ATA P., GOKCE İ., ALPAY H.
18 th Congress of the International Pediatric Nephrology Association-IPNA, 17 - 21 October 2019

IV. **A NOVEL MUTATION IN ATP6V0A4 GENE IN APATIENT WITH DISTAL RENAL TUBULAR ACIDOSIS**
SAK M., YILDIZ N., Çiçek N., GÜVEN S., ATA P., GÖKCE İ., ALPAY H.
18 th Congress of the International Pediatric Nephrology Association-IPNA, 17 - 21 October 2019

V. **Schaaf Yang sendromu**
ALAVANDA C., ARSLAN ATEŞ E., POLAT H., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P., ARMAN A.
4. Ulusal Çocuk Genetik kongresi, Turkey, 25 - 27 September 2019

VI. **Nadir bir iskelet displazisi: Piknodizostoz tanısı alan iki kız kardeş**
ALAVANDA C., GEÇKİNLİ B. B. , ARSLAN ATEŞ E., POLAT H., SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P., ARMAN A.
4. Ulusal Çocuk genetik kongresi, Turkey, 25 - 27 September 2019

VII. **Rare Causes of Osteogenesis Imperfecta are Commonin Consanguineous Pedigrees**
KAYGUSUZ S. B. , ARMAN A., ABALI S., ATA P., KIRKGÖZ T., YAVAŞ ABALI Z., HELVACIOĞLU D., GÜRPINAR T. B. , SEVEN M. T. , GÜRAN T., et al.
European Society for Paediatric Endocrinology (ESPE)58th Annual Meeting, Vienna, VİYANA, Austria, 19 - 21 September 2019

VIII. **A rare cause of hypophosphatemia: Raine Syndrome**
ELTAN M., ATA P., KIRKGÖZ T., ALAVANDA C., KAYGUSUZ S. B. , SEVEN M. T. , GURPINAR TOSUN B., YAVAŞ ABALI Z., GÜRAN T., ELÇİOĞLU H. N. , et al.
European Society for Paediatric Endocrinology (ESPE)58th Annual Meeting, Vienna, VİYANA, Austria, 19 - 21 September 2019, vol.91

IX. **Genotype and Phenotype Characterization ofTurkish Patients with Vitamin D Dependent RicketsType IA**
KAYGUSUZ S. B. , ATA P., KIRKGÖZ T., YAVAŞ ABALI Z., ELTAN M., GÜRPINAR T. B. , SEVEN M. T. , HELVACIOĞLU D., ARMAN A., GÜRAN T., et al.
European Society for Paediatric Endocrinology (ESPE)58th Annual Meeting, Vienna, VİYANA, Austria, 19 - 21 September 2019, vol.91

X. **A novel intronic ATM gene mutation affecting splicing in a patient with Ataxia-Telangiectasia**
Ates E. A. , Turkyilmaz A., SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , ATA P., ARMAN A., GÜNEY A. İ.
51st Conference of the European-Society-of-Human-Genetics (ESHG) in conjunction with the European Meeting on Psychosocial Aspects of Genetics (EMPAG), Milan, Italy, 16 - 19 June 2018, vol.27, pp.255-256

XI. **FGF3 gene mutations related to two syndromic Congenital deafness cases: Congenital deafness with inner ear agenesis (Michel aplasia), microtia, and microdontia and Otodental dysplasia**
Turkyilmaz A., GEÇKİNLİ B. B. , Ates E. A. , SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P., ARMAN A.
51st Conference of the European-Society-of-Human-Genetics (ESHG) in conjunction with the European Meeting on Psychosocial Aspects of Genetics (EMPAG), Milan, Italy, 16 - 19 June 2018, vol.27, pp.893

XII. **The effects of calcium-sensing receptor CASR genotypes, treatment duration, gender bone health and mineral metabolism in chronic renal failure patients**
ATA P., Erkal B., Gultekin D., Altas B., ÇELİK B., Kayir D., Low A., Eren A., Tuglular S.
51st Conference of the European-Society-of-Human-Genetics (ESHG) in conjunction with the European Meeting on Psychosocial Aspects of Genetics (EMPAG), Milan, Italy, 16 - 19 June 2018, vol.27, pp.97

XIII. **The prevalence in coeliac related hla haplotypes in paediatric IBD patients**
Ertem Şahinoğlu D., Akkelle B., Şengül Ö., Volkan B., Tutar E., Ata P.
52. ESPGHAN, Glasgow, United Kingdom, 5 - 08 June 2019, vol.68, no.1, pp.592

XIV. **Fokal Segmental Glomeruloskleroz, son tanı mı?**
Çiçek N., Yıldız N., Ata P., Gökce İ., Güven S., Sak M., Alpay H.
10.Uluslararası katılımlı Çocuk Nefroloji Kongresi, Muğla, Turkey, 1 - 04 May 2019, pp.194

XV. **Alport Sendromu Tanılı Hastalarda Genetik Analiz ve Yeni Mutasyonlar: Varyasyonlar Her Zaman Masum Mu?**

çiçek n., YILDIZ N., ATA P., GÖKCE İ., SAK M., KAYA H., ALPAY H.
Uluslararası katılımlı 10. Çocuk Nefroloji Kongresi, Turkey, 1 - 04 May 2019

XVI. **Acute Pancreatitis Severity Related With Macrophage Migration Inhibitory FactorGene - 173 G/C Polymorphism and Serum MIF Level**
UZUN M. A. , erkal b., ATA P., DOĞAN T., BERK OCAK S., ALKAN KAYAOĞLU S., TİLKİ M., YILMAZ İ., Arslan E.
13. balkan genetik kongresi, Edirne, Turkey, 16 - 20 April 2019

XVII. **TWO NOVEL MUTATIONS IN THREE DIFFERENT GENES ASSOCIATED WITHRETINITIS PIGMENTOSA/LEBER CONGENITAL AMAROSIS IN ONE PATIENT**
ALAVANDA C., ATA P., YILMAZ Ö., POLAT H., GEÇKİNLİ B. B. , ARMAN A.
13. BALKAN GENETİK KONGRESİ, Edirne, Turkey, 16 - 20 April 2019

XVIII. **Von Hippel Lindau Patients**
ALAVANDA C., TÜRKYILMAZ A., POLAT H., GEÇKİNLİ B. B. , GÜNEY A. İ. , ATA P., SÖYLEMEZ M. A. , ARMAN A.
13. BALKAN GENETİK KONGRESİ, Edirne, Turkey, 16 - 20 April 2019

XIX. **Yenidoğanda Nadir Bir Hiperkalsemi nedeni : Kalsiyum Duyarlı Reseptör Mutasyonuna bağlı Ağır neonatal Hiperparatiroidizm**
Rzayev T., KIRKGÖZ T., Özdemir H., ATA P., BİLGEN H. S. , MEMİŞOĞLU A., ELTAN M., ÖZEK E., Turan S.
27. Ulusal Neonatoloji Kongresi, Antalya, Turkey, 3 - 07 April 2019

XX. **FARKLI GENLER,FARKLI MUTASYONLAR,FENOTIPTE FARKLILIK YARATIYOR MU ?: ALPORT SENDROMU**
ATA P., ALAVANDA C., TÜRKYILMAZ A., YILDIZ N., ALPAY H.
Uluslararası Katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Turkey, 7 - 11 November 2018

XXI. **IKI LAMINOPATI OLGUSU: LMNA GENI MUTASYONLARI ILE ILIŞKILI FENOTIPLER**
GEÇKİNLİ B. B. , TÜRKYILMAZ A., SÖYLEMEZ M. A. , ATES E., YILDIRIM Ö., ATA P., ARMAN A.
Uluslararası Katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Antalya, Turkey, 07 November 2018 - 11 November 1918

XXII. **GENETIK KLINIĞINE PSO'DOBARTTER SENDROMU BULGULARIYLA BAŞVURAN KISTIK FIBROZLU IKI KIZ KARDEŞ**
ALAVANDA C., ATA P., TÜRKYILMAZ A., Arslan E., YILDIZ N., ALPAY H.
Uluslararası Katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Turkey, 7 - 11 November 2018

XXIII. **ANKILOZAN SPONDILIT'xxTE MIKRO RNA EKSPRESYONU**
TÜRKYILMAZ A., AKBAŞ F., YAĞCI İ., ATA P.
Uluslararası Katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Turkey, 7 - 11 November 2018

XXIV. **PEDIATRIK AKUT LENFOBLASTIK LO'SEMIDE TEŞHISSEL TESTLERIN GEÇERLILIĞI**
Yılmaz İ., ATA P., ALAVANDA C., Arslan E., Eren R., Yılmaz B., timur ç., pala S., Acıyiyen Y., sökmen b., et al.
13. Uluslararası Katılımlı Ulusal Tıbbi Genetik Kongresi, Antalya, Turkey, 7 - 11 November 2018

XXV. **Chromosomal array-CGH analysisin patients having neurodevelopmental delay and dysmorphic features**
ALAVANDA C., ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , ATA P., GÜNEY A. İ. , SÖYLEMEZ M. A. , ÖZYAVUZ ÇABUK P., ARMAN A.
13 ULUSAL TIBBİ GENETİK KONGRESİ, Turkey, 7 - 11 November 2018

XXVI. **REVERSE GENETIK ILE TANI KOYULAN NADIR BIR SPASTIK PARAPLEJI AILESI**
ALAVANDA C., ates e., TÜRKYILMAZ A., GÜNEY A. İ. , GEÇKİNLİ B. B. , ATA P., ARMAN A.
13.ULUSLARARASI KATILIMLI ULUSAL TIBBİ GENETİK ANABİLİM DALI, Turkey, 7 - 11 November 2018

XXVII. **NO'ROMOTOR GELIŞME GERILIĞI VE DISMORFIK BULGULARI OLAN HASTALARDA ARRAY-CGH ANALIZI**
ATA P., ALAVANDA C., ATEŞ E., GEÇKİNLİ B. B. , GÜNEY A. İ. , SÖYLEMEZ M. A. , ÇABUK P., ARMAN A.
Uluslararası Katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Antalya, Turkey, 7 - 11 November 2018

XXVIII. **Genotype and phenotype correlation in Turkish Alport syndrome patients with different COL4A3, COL4A4 and COL4A5 mutations.**
ATA P., CELİK T., PAK T., ISIK A., YILDIRIM S., TÜRKYILMAZ A.
ASHG 2018, 16 October 2018 - 20 January 2019

XXIX. **Novel mutations and variations in turkish children with Alport syndrome: are the benign variations**

always harmless?
Çiçek N., YILDIZ N., ATA P., Gökçe İ., SAK M., KAYA H., ALPAY H.
51th Annual Scientific Meeting of the European Society for Paediatric Nephrology, Antalya, Turkey, 3 - 06 October 2018

XXX. **Novel Large CFTR Gene Deletions in Turkish Patients with Increased Morbidity**
ATA P., Atag E., GÖKDEMİR Y., Ikizoglu N. B. , DELİL K., Eralp E., Ergenekon P., KARADAĞ B. T.
50th European-Society-of-Human-Genetics (ESHG) Conference, Copenhagen, Denmark, 27 - 30 May 2017, vol.26, pp.980

XXXI. **A novel intronic ATM genemutation a×ecting splicing in a patient withAtaxia-Telangiectasia**
ATES E., TÜRKYILMAZ A., ATA P., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , GÜNEY A. İ. , ARMAN A.
ESHG2018, 16 - 19 June 2018

XXXII. **HLA and non-HLA genetic predisposing factors and environmental effects at Celiac Disease pathogenesis**
ATA P., erkal b., ŞAHİN AKKELLE B., SİSKO S., ERTEM ŞAHİNOĞLU D.
EUROPEAN SOCIETY OF HUMAN GENETICS, 16 - 19 June 2018

XXXIII. **The relationship betweendefects at the middle ear bone chain andNLRP3 and OPG gene polymorphisms atchronic otitis media patients**
KESKIN S., ATA P., erkal b., DOGAN T., EREN A., TATLIPINAR A.
ESHG 2018, 16 June 1918 - 19 June 2018

XXXIV. **Herediter parapleji hastalık ailesine olgularla yaklaşım**
SAĞER S. G. , ÖZTÜRK THOMAS G., TÜRKYILMAZ A., ATA P., ÜNVER O., TÜRKDOĞAN D.
20. Çocuk Ulusal Nörolojisi Kongresi, Cyprus (Kktc), 2 - 06 May 2018

XXXV. **Hepatosellüler karsinom gelişiminde moleküleryolaklar**
ATA P.
M.Ü. Tıp Fakültesi III. Uluslararası Onkolojive Cerrahi Günleri Sempozyumu, 3 - 04 May 2018

XXXVI. **Hepatosellüler karsinom tanısında genetikbiyobelirteçler**
ATA P.
MARMARA ÜNİVERSİTESİ TIP FAKÜLTESİIII. ULUSLARARASI ONKOLOJİ VE CERRAHİ GÜNLERİHEPATO-PANKREATO-BİLİYER ve GASTROİNTESTİNAL KANSERLERDESİSTEMİK VE CERRAHİ TEDAVİLERDE GELİŞMELER, 3 - 05 May 2018

XXXVII. **Makrofaj migrasyon inhibitör faktör -173 G/C gen polimorfizmi ile akut pankreatit gelişimi ve şiddeti arasındaki ilişki**
UZUN M. A. , ALKAN KAYAOĞLU S., ATA P., TİLKİ M., BERK OCAK S., DOĞAN T., ERKAL B.
13. Türk Hepatopankreatobilier Cerrahi Kongresi, Antalya, Turkey, 1 - 04 November 2017, vol.33, pp.1-92

XXXVIII. **THE ROLE OF SOLUBLE CTLA-4 AS A NON-INVASIVEBIOMARKER FOR DIAGNOSIS OF KIDNEY ALLOGRAFTREJECTION: A PRELIMINARY STUDY**
Ruhi Borçak ç., ATA P., TİTİZ M. İ.
18th Congressof the European Society for Organ Transplantation, 24 - 27 September 2017

XXXIX. **COMPARISON OF THE TREATMENT EFFICACY OFRITUXIMAB AND PLASMAPHERESIS/INTRAVENOUSIMMUNOGLOBULIN COMBINATION WITH HISTORICALCONTROL IN CHRONIC ANTIBODY MEDIATEDREJECTION**
Ruhi Borçak ç., ATA P., TİTİZ M. İ.
18th Congressof the European Society for Organ Transplantation, 24 - 27 September 2017

XL. **THE INFLUENCE OF PREFORMED HLA CLASS I AND II PANEL REACTIVE ANTIBODIES ON CLINICAL AND PATHOLOGICAL OUTCOMES OF KIDNEY ALLOGRaft**
RUHİ Ç., TUĞCU M., KASAPOĞLU U., BOYNUEĞRİ B., GÜMRÜKÇÜ G., ATA P., APAYDIN S., TİTİZ M. İ.
ESOT, 15 - 18 September 2017, vol.30, pp.318

XLI. **Does C.3979 G A/P.VAL1327MET variant of COL4A4 has any pathogenic effect in Turkish patients with Alport Syndrome ?**
YILDIZ N., ATA P., ALPAY H., GÖKCE İ., ÖZALTIN F.
48th Anniversary Meeting of the European Society for Paediatric Nephrology, 6 - 09 September 2017

XLII. **A Novel HNF1B mutation in a family with two MODY patients**
ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , DELİL K., ATA P., GÜNEY A. İ.
The European Society of Human Genetics, European Human Genetics Conference, 27 - 30 May 2017

XLIII. **BBS10 frameshift mutation in a Turkish girl with Bardet Biedl Syndrome.**
SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , DELİL K., ARSLAN ATEŞ E., TÜRKYILMAZ A., AVŞAR M., GİRGİN G., ATA P., ARMAN A., GÜNEY A. İ.
The European Society of Human Genetics, European Human Genetics Conference, 27 - 30 May 2017

XLIV. **Evaluation of BRCA1/BRCA2 test results for Turkish breast cancer families.**
ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , GİRGİN ÖZGÜMÜŞ G., SÖYLEMEZ M. A. , DELİL K., ATA P., ARMAN A., GÜNEY A. İ.
The European Society of Human Genetics, European Human Genetics Conference, 27 - 30 May 2017

XLV. **BBS10 frameshift mutation in a turkish girl with bardet biedl syndrome**
SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , DELİL K., TÜRKYILMAZ A., AVŞAR M., GİRGİN G., ATA P., ARMAN A., GÜNEY A. İ.
The European Society of Human Genetics 2017, KOPENHAGEN, Denmark, 25 - 30 May 2017

XLVI. **Inversion Y Having Different Phenotypic Expressions at Three Brothers**
TÜRKYILMAZ A., ATA P., DELİL K., SOYSAL S., ARSLAN ATEŞ E., GÜNEY A. İ.
The European Society of Human Genetics, KOPENHAGEN, Denmark, 25 - 30 May 2017

XLVII. **Evaluation of BRCA1/2 test results for Turkish breast cancer families.**
ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , ÖZGÜMÜŞ GİRGİN G., DELİL K., ATA P., ARMAN A., GÜNEY A. İ.
The European Society of Human Genetics 2017, KOPENHAGEN, Denmark, 25 - 30 May 2017

XLVIII. **A novel HNF1B mutation in a family with two MODY patients**
ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , DELİL K., SÖYLEMEZ M. A. , ATA P., GÜNEY A. İ.
The European Society of Human Genetics 2017., KOPENHAGEN, Denmark, 25 - 28 May 2017

XLIX. **Novel Large CFTR Gene Deletions in Turkish Patients with Increased Morbidity,**
ATA P., ATAG E., GOKDEMİR Y., NE B. I. , DELİL K., ERALP E., ERGENEKON A. P. , KARADAĞ B. T.
The European Society of Human Genetics, KOPENHAGEN, Denmark, 26 - 30 May 2017

L. **BBS-10 frameshift mutation in a Turkish girl with bardet biedl syndrome**
GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , TÜRKYILMAZ A., ARSLAN ATEŞ E., DELİL K., ATA P., GÜNEY A. İ.
ERCİYES TIP GÜNLERİ 2017, Kayseri, Turkey, 11 - 13 May 2017

LI. **EVALUATION OF THE BRCA1/BRCA2 MUTATIONS AT BREAST CANCER PATIENTS**
ARSLAN ATEŞ E., TÜRKYILMAZ A., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , GİRGİN ÖZGÜMÜŞ G., DELİL K., ATA P., GÜNEY A. İ.
ERCİYES TIP GÜNLERİ, Kayseri, Turkey, 11 - 13 May 2017

LII. **Largest family in Turkey with multiple endocrine neoplasia-type 1 and a newly discovered mutation.**
DUMAN D., ÇETİN A., Tütüncü Y., Şimşek H., ATA P.
16. Eurasian Congress of Gastrtoenterology and Surgery, Mostar, Bosnia And Herzegovina, 27 - 30 April 2017, vol.7, pp.117-118

LIII. **May Intraoperative Immunosuppressive Therapy Reduce the Rejection Episodes at Cardiac Transplant Recipients?**
Rabus M. B. , Cekmecelioglu D., Ata P., Salihi S., Selcuk E., Balkanay M.
37th Annual Meeting and Scientific Sessions of the International-Society-for-Heart-and-Lung-Transplantation (ISHLT), California, United States Of America, 5 - 08 April 2017, vol.36

LIV. **A New Flowcytometric crossmatch Technique with 7AAD For Cytotoxic Antibody Detection**
BİLGEN T., ATA P., TOZKIR J., TOZKIR H., TİTİZ M. İ.
TÜRKİYE ORGAN NAKLİ KOORDİNATÖRLERİ KONGRESİ, Konya, Turkey, 13 - 15 October 2016

LV. **BREAST CANCER INHERITANCE FAMILY PATIENTS POINT OF VIEW**
DASTAN B., ERGİN Z., GÜLTEKİN H. D. , AKIN S., HELVACI M. S. , NAZLI Y., GULLUOGLU M. B. , ATA P.
International Istanbul Breast Cancer Conference - BREASTANBUL 2016, İstanbul, Turkey, 10 - 12 October 2016

LVI. **Von Hippel -Lindau sendromu ailesindeki asemptomatik 3 çocoğun genetik test sonuçlarının değerlendirilmesi**

DELİL K., GEÇKİNLİ B. B. , ŞİMŞEK H., TÜRKYILMAZ A., ARSLAN ATEŞ E., SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P., ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LVII. **Silver Russel Sendromlu bir olgu**

GEÇKİNLİ B. B. , DELİL K., ŞİMŞEK H., SÖYLEMEZ M. A. , TÜRKYILMAZ A., ARSLAN ATEŞ E., KARAKAYA T., GÜNEY A. İ. , ATA P., ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LVIII. **Beckwith Wiedemann sendromlu olgunun klinik bulguları ve moleküler tanısı**

KARAKAYA T., SÖYLEMEZ M. A. , ARSLAN ATEŞ E., TÜRKYILMAZ A., ŞİMŞEK H., DELİL K., GEÇKİNLİ B. B. , ATA P., GÜNEY A. İ. , ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LIX. **RAB3GAP1 geninde bilinen bir splice-site mutasyonunun fonksiyonel değerlendirilmesi**

TÜRKYILMAZ A., SÖYLEMEZ M. A. , ARSLAN ATEŞ E., ERGÜNER B., ŞİMŞEK H., KARAKAYA T., DELİL K., GEÇKİNLİ B. B. , ATA P., GÜNEY A. İ. , et al.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LX. **Chromosome 17p11.2 deletion in a Turkish girl with Smith -Magenis Syndrome**

GEÇKİNLİ B. B. , DELİL K., TÜRKYILMAZ A., SÖYLEMEZ M. A. , ŞİMŞEK H., KARAKAYA T., ARSLAN ATEŞ E., GÜNEY A. İ. , ATA P., ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LXI. **Nadir görülen bir genetik sendrom: Meacham sendromu**

DELİL K., TÜRKYILMAZ A., ŞİMŞEK H., KARAKAYA T., ARSLAN ATEŞ E., SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , GÜNEY A. İ. , ATA P., ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LXII. **Geç tanı almış Williams sendromlu üç olgunun klinik değerlendirmesi**

ARSLAN ATEŞ E., GEÇKİNLİ B. B. , KARAKAYA T., TÜRKYILMAZ A., ŞİMŞEK H., SÖYLEMEZ M. A. , DELİL K., ATA P., GÜNEY A. İ. , ARMAN A.

12. Ulusal Tıbbi Genetik Kongresi, Turkey, 5 - 09 October 2016

LXIII. **Clinical Evaluation of a MDC1A Case Carrying LAMA2 Mutation**

SÖYLEMEZ M. A. , TÜRKYILMAZ A., ŞİMŞEK H., DELİL K., GEÇKİNLİ B. B. , ARMAN A., GÜNEY A. İ. , ATA P., OKUYAN B.

ULUSAL TIBBİ GENETİK KONGRESİ, İZMİR ÇEŞME, Turkey, 5 - 09 October 2016

LXIV. **Yaşlanma ile İlişkili Klotho Geni ve Bu Genin Ekspresyonunda Epigenetik Faktörlerin Rolü**

TURAN K., Çağlayan E., ATA P.

Ulusal Moleküler Tıp Sempozyumu: Moleküler Bakıştan Kliniğe, İstanbul, Turkey, 1 - 03 June 2016

LXV. **A case of Weaver Syndrome caused by a novel frameshift EZH2 mutation**

TÜRKYILMAZ A., KARAKAYA T., ŞİMŞEK H., SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , ARMAN A., DELİL K., ATA P.

European Society of Human Genetics, 21 - 24 May 2016

LXVI. **Analysis of The Molecular Markers in 49 AML Patients**

TÜRKYILMAZ A., SÖYLEMEZ M. A. , DELİL K., ŞİMŞEK H., GÜNEY A. İ. , ATA P.

European Society of Human Genetics Congress, Barcelona, Spain, 21 - 24 May 2016

LXVII. **A novel splice site JAG1 mutation in a Turkish girl with Alagille Syndrome**

TÜRKYILMAZ A., GEÇKİNLİ B. B. , DELİL K., SÖYLEMEZ M. A. , GÜNEY A. İ. , ARMAN A., ATA P.

European Society Of Human Genetics Congress, Barcelona, Spain, 21 - 24 May 2016

LXVIII. **Recurrent miscarriage and implantation failure Could the etiology be maternal intolerance itself**

ATA P., SOYSAL S., TÜRKYILMAZ A., DELİL K., ANIK İLHAN G., GEÇKİNLİ B. B.

European Society of Human Genetics Congress, BARSELONA, Spain, 21 - 24 May 2016

LXIX. **A case of Weaver Syndrome caused by a novel frameshift EZH2mutation**

ŞİMŞEK H., GEÇKİNLİ B. B. , DELİL K., SÖYLEMEZ M. A. , TÜRKYILMAZ A., Arslan Ateş E., AVŞAR M., ATA P., GÜNEY A. İ. , ARMAN A.

ESHG 2016 | BARCELONA, SPAIN, 21 - 24 May 2016

LXX. **Determination of Potential Drug Drug Interactions by Using Various Drug Interaction Software**

Programs at Hospital Pharmacy Setting
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
44th European Symposium on Clinical Pharmacy, LİZBON, Portugal, 28 - 30 October 2015

LXXI. **Assessment of Attitude and Knowledge in Patients Utilized Disposable Insulin Pens at Community Pharmacy Setting**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
44th European Symposium on Clinical Pharmacy, LİZBON, Portugal, 28 - 30 October 2015

LXXII. **Implementation of Medication Reconciliation and Medication Review Services Conducted by Pharmacist in Hospitalized COPD Patients**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
44th European Symposium on Clinical Pharmacy, LİZBON, Portugal, 28 - 30 October 2015

LXXIII. **Determination of Potential Drug Drug Interactions by Using Various Software Programs at Community Pharmacy Setting**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
44th European Symposium on Clinical Pharmacy, LİZBON, Portugal, 28 - 30 October 2015

LXXIV. **Evaluation of Potentially Inappropriate Medication Use and Drug Burden Index in Elderly Patients with Cancer**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
44th European Symposium on Clinical Pharmacy, LİZBON, Portugal, 28 - 30 October 2015

LXXV. **Recurrent Fetal Loss Family with Translocation t 18 19 p11 2 p13 1 and its Clinical İmplications**
TURKYILMAZ A., ŞİMSEK H., MAMADOV e., DELİL K., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P.
European Society of Human Genetics, 6 - 09 June 2015

LXXVI. **Hereditary Heterotopic Ossification Syndromes Effect of GNAS inactivation in progressive osseousheteroplasia A case report**
ATA P., TURKYILMAZ A., DELIL K., YAĞCI İ., GENCER K., OZTURK E., AKYÜZ G. D.
European Society of Human Genetics, Milan, Italy, 6 - 09 June 2015

LXXVII. **A Case of Sotos Syndrome with a Novel Mutation of NSD1 Gene**
ŞİMŞEK H., GEÇKİNLİ B. B. , DELİL K., SÖYLEMEZ M. A. , TURKYILMAZ A., ATA P., GÜNEY A. İ. , ARMAN A.
European Society ofHuman Genetics, 6 - 09 June 2015

LXXVIII. **Translocation t 18 19 p11 2 p13 1 and its Clinical Implications Case report and Mechanism of Pathogenesis**
TÜRKYILMAZ A., ŞİMŞEK H., Mamadov E., DELİL K., GEÇKİNLİ B. B. , SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P.
ESHG Congress 2015, 6 - 09 June 2015

LXXIX. **PROGRESİVA ANKİLOZAN SPONDİLİT İLE KARIŞAN NADİR BİR HASTALIK**
AKYÜZ G. D. , GENCER K., YAĞCI İ., ÖZTÜRK E., DELİL K., TÜRKYILMAZ A., ATA P.
25.UlusalFiziksel TipveRehabilitasyonKongresi, Turkey, 22 - 26 April 2015

LXXX. **KALITIMSAL HETEROTOPI K OSSI FI KASYON SENDROMLARI FI BRODI SPLAZI A OSSI FI KANS PROGRESSI VAMI PROGRESI F OSSÖZ HETEROPLAZI MI OLGU SUNUMU**
YAĞCI İ., GENCER K., AKYÜZ G. D. , DELİL K., TÜRKYILMAZ A., ATA P.
25.UlusalFizikselTipveRehabilitasyonKongresi, Turkey, 22 - 26 April 2015

LXXXI. **KALITIMSAL HETEROTOPİK OSSİFİKASYON SENDROMLARI FİBRODİSPLAZİA OSSİFİKANS PROGRESSİVA MI PROGRESİF OSSÖZ HETEROPLAZİ Mİ OLGU SUNUMU**
YAĞCI İ., GENÇER Z. K. , AKYÜZ G. D. , DELİL K., TÜRKYILMAZ A., ATA P.
24. Ulusal Fiziksel Tıp ve Rehabilitasyon Kongresi, Turkey, 23 - 26 April 2015

LXXXII. **Determination of drug related problem and drug burden index in elderly patients**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.208-209

LXXXIII. **Assessment of knowledge and attitude towards osteoporosis among women attending community pharmacy settings**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.

43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.213

LXXXIV. **The role of pharmacist in detecting angiotensin converting enzyme inhibitors induced dry cough at community pharmacy setting**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.207

LXXXV. **Evaluation of drug burden index in elderly patients utilized disposable insulin pen**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.203-204

LXXXVI. **Assessment of patients knowledge and attitude towards methotrexate utilization**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.256

LXXXVII. **Evaluation of drug burden index in elderly patients utilized inhaler**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.205

LXXXVIII. **Evaluation of drug burden index and medication utilization in geriatric patients at community pharmacy**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GUNES T., CALISKAN M., İZZETTİN F. V.
43rd European Symposium on Clinical Pharmacy, KOPENHANG, Denmark, 22 - 24 October 2014, vol.37, pp.206-207

LXXXIX. **Recurrent fetal loss family with translocation t 18 19 p11 2 p13 1 and its clinical implications**
TÜRKYILMAZ A., ŞİMŞEK H., MAMADOV E., DELİL K., GEÇKİNLİ B. B. , GÜNEY A. İ. , SÖYLEMEZ M. A. , ATA P.
11. Ulusal Tıbbi Genetik Kongresi, 24-27 Eylül 2014, Turkey, 24 - 27 September 2014

XC. **FBN1 mutasyonu ve tipik klinik bulguları olan Marfan sendromlu üç olgu**
DELİL K., SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , AVŞAR M., ŞİMŞEK H., TÜRKYILMAZ A., GÜNEY A. İ. , ATA P., ARMAN A.
11. Ulusal Tıbbi Genetik Kongresi, Turkey, 24 - 27 September 2014

XCI. **Alt extremite hemihipertrofisi olan bir Prader Willi Sendromu Olgusu**
ŞİMŞEK H., DELİL K., GEÇKİNLİ B. B. , TÜRKYILMAZ A., AVŞAR M., SÖYLEMEZ M. A. , GÜNEY A. İ. , ATA P., ARMAN A.
11. Ulusal Tıbbi Genetik Kongresi, Turkey, 24 - 27 September 2014

XCII. **Duchenne Musküler Distrofi DMD tipik klinik bulguları olan ve DMD geninde nonsense mutasyon saptanan olgu**
TÜRKYILMAZ A., DELİL K., GEÇKİNLİ B. B. , ŞİMŞEK H., SÖYLEMEZ M. A. , AVŞAR M., GÜNEY A. İ. , ATA P., ARMAN A.
11. Ulusal Tıbbi Genetik Kongresi, Turkey, 24 - 27 September 2014

XCIII. **Determination of CYP2C19 polymorphisms adverse drug reaction and medication adherence in patients utilized selective serotonin reuptake inhibitors**
DENİZ S., SANCAR M., OKUYAN B., ATA P., BİNGÖL ÖZAKPINAR Ö., TALAS A., GÜNEŞ T., ÇALIŞKAN M., İZZETTİN F. V.
Four International Meeting on Pharmacy and Pharmaceutical Sciences (IMPPS-4), 18 - 21 September 2014

XCIV. **Is the TLR-4 Gene Polymorphism Play a Protective Role in the Development of NAFLD in Humans ?**
Kiziltas S., Ata P., Colak Y., Mesci B., Senates E., Enc F., Ulasoglu C., Tuncer I., Oguz A.
Digestive Disease Week / 28th Annual Residents and Fellows Research Conference of the Society-for-Surgery-of-the-Alimentary-Tract (SSAT), Florida, United States Of America, 18 - 21 May 2013, vol.144

XCV. **Anti aging Klotho geni ekspresyonuna hücre içi ve hücreler arası faktörlerin etkisi**
ATA P., TURAN K.
10.Ulusan Tıbbi Genetik Kongresi, Bursa, Turkey, 19 - 23 December 2012

XCVI. **Early posttransplant clinical status of desensitized live-donor kidney recipients sensitized with pregnancy and blood transfusions**
Ata P., Canbakan M., Ozel L., Kara M., Krand O., Unal E., Demiralp E., Titiz M. I.
25th Conference on European Immunogenetics and Histocompatibility, Prague, Czech Republic, 4 - 07 May 2011, vol.77, pp.453

XCVII. **Impact of sensitization to HLA antigens on receiving a second kidney transplant after a failed graft**
Eren P., Goral S., Bloom R., Doyle A., Grossman R., Israni A., Naji A., Sellers M., Markmann J., Kearns J., et al.

42nd Annual Meeting of the European-Renal-Association/European-Dialysis-and-Transplant-Association (ERA-EDTA), İstanbul, Turkey, 4 - 07 June 2005, vol.20

XCVIII. **Impact of sensitization to HLA class I and class II antigens (AG) on receiving a 2ND kidney transplant (tx) after a failed graft: Report from a transplant center using a stringent HLA crossmatching (XM) strategy**
Goral S., Eren P., Bloom R., Doyle A., Grossman R., Israni A., Naji A., Sellers M., Markmann J., Kearns J., et al.
6th American Transplant Congress, Washington, United States Of America, 21 - 25 May 2005, vol.5, pp.351

Supported Projects

Ata P., Bilgen T., Titiz M. İ. , TUBITAK Project, Transplantasyon Başarısında Etkili Olan Donöre ÖzgünAlloreaktif Hafıza B Hücre Rezervinin Değerlendirilmesi, 2018 - 2020
Ata P., Alpay H., Ürer A., Project Supported by Higher Education Institutions, CAKUT Hastalarının Demografik Özellikleri ve Organogenezde Etkili Faktörlerinin Değerlendirilmesi, 2018 - 2019

Scientific Refereeing

TUBITAK Project, 1001 - Program for Supporting Scientific and Technological Research Projects, Marmara University, Turkey, September 2019
TUBITAK Project, 1001 - Program for Supporting Scientific and Technological Research Projects, Marmara University, Turkey, June 2019

Citations

Total Citations (WOS):176
h-index (WOS):8