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Address: Çukurova Üniversitesi Tıp Fakültesi Çocuk Metabolizma ve Beslenme Bilim Dalı 01130 Balcalı Adana

Education Information

Post Doctorate of Medicine, Cukurova University, Tıp Fakültesi, Çocuk Endokrinoloji Ve Metabolizma Bilim Dalı, Turkey
1995 - 2005

Expertise In Medicine, Cukurova University, Tıp Fakültesi, Pediatri Anabilim Dalı, Turkey 1990 - 1995

Undergraduate, Cukurova University, Tıp Fakültesi, Turkey 1982 - 1988

Foreign Languages

English, B2 Upper Intermediate

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases, Pediatric Endocrinology and Metabolism

Academic Titles / Tasks

Professor, Cukurova University, Tıp Fakültesi, Dahili Tıp Bilimleri, 1996 - Continues

Academic and Administrative Experience

Cukurova University, Hemşirelik Myo, 2001 - 2002

Courses

Metabolik karaciğer hastalıkları , Undergraduate, 2016 - 2017

Advising Theses

ÖNENLİ MÜNGAN H., Primer ailesel hiperlipidemi tanılı çocuk ve adölesanlarda demografik, klinik özellikler ve bunların mutasyon analizi sonuçlarıyla ilişkisi, Expertise In Medicine, i.kaplan(Student), 2016

ÖNENLİ MÜNGAN H., Çukurova Üniversitesi Tıp Fakültesi Pediatrik Metabolizma Hastalıkları ve Beslenme Polikliniğinde tanı alan veya takibe giren kalıtsal metabolik hastalığı olan hastaların tanılarının, klinik ve laboratuvar bulgularının analizi ile takip sonuçlarının değerlendirilmesi, Expertise In Medicine, a.kara(Student), 2012

ÖNENLİ MÜNGAN H., Çocukluk çağı obezitesinde metabolik parametrelerin diyet ve egzersizle ilişkisi, Expertise In Medicine, i.öncü(Student), 2009

Jury Memberships

Associate Professor Exam, Doçentlik sınavı, Doçentlik sınavı, June, 2016

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Real-world patient data on immunity and COVID-19 status of patients with MPS, Gaucher, and Pompe diseases from Turkey**
Kilavuz S., Kor D., BULUT F. D., Serbes M., Karagoz D., Altintas D. U., Bisgin A., Seydaoglu G., Mungan H. N. O.
ARCHIVES DE PEDIATRIE, vol.29, no.6, pp.415-423, 2022 (SCI-Expanded)
- II. **Evaluation of bone health in patients with mucopolysaccharidosis**
KOR D., BULUT F. D., Kilavuz S., Yilmaz B. S., Koseci B., KARA E., KAYA Ö., BAŞARAN S., SEYDAOĞLU G., Mungan N. O.
JOURNAL OF BONE AND MINERAL METABOLISM, vol.40, pp.498-507, 2022 (SCI-Expanded)
- III. **Diagnostic value of plasma lysosphingolipids levels in a Niemann-Pick disease type C patient with transient neonatal cholestasis**
Bulut F. D., Bozbulut N. E., Ozalp O., DALGIÇ B., Mungan N. O., Ucar H. K., BİBEROĞLU G.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.35, pp.681-685, 2022 (SCI-Expanded)
- IV. **Investigating myelin oligodendrocyte glycoprotein antibodies in hereditary citrullinemia**
Oncel I., Yousefi M., İNCİ A., ARSLAN GÜLTEN Z., Kisa P. T., Karaca M., Unal O., Gunduz M., KOR D., Mungan N. O., et al.
MEDICAL HYPOTHESES, vol.160, 2022 (SCI-Expanded)
- V. **Dysarthria, Ataxia, and Dystonia Associated with COX20 (FAM36A) Gene Mutation: A Case Report of a Turkish Child**
Ozcanyuz D. G., İNCECİK F., Herguner O. M., Mungan N. O., Bozdogan S. T.
ANNALS OF INDIAN ACADEMY OF NEUROLOGY, vol.23, no.3, pp.399-401, 2020 (SCI-Expanded)
- VI. **Defining clinical subgroups and genotype-phenotype correlations in NBAS-associated disease across 110 patients**
Staufner C., Peters B., Wagner M., Alameer S., Baric I., Broue P., Bulut D., Church J. A., Crushell E., DALGIÇ B., et al.
GENETICS IN MEDICINE, vol.22, no.3, pp.610-621, 2020 (SCI-Expanded)
- VII. **Analysis of the caregiver burden associated with Sanfilippo syndrome type B: panel recommendations based on qualitative and quantitative data**
Shapiro E., Lourenco C. M., Mungan N. O., Muschol N., O'Neill C., Vijayaraghavan S.
ORPHANET JOURNAL OF RARE DISEASES, vol.14, 2019 (SCI-Expanded)
- VIII. **A possible biomarker of neurocytolysis in infantile gangliosidoses: aspartate transaminase**
Kilic M., Kasapkara C. S., Kilavuz S., Mungan N. O., BİBEROĞLU G.
METABOLIC BRAIN DISEASE, vol.34, no.2, pp.495-503, 2019 (SCI-Expanded)
- IX. **CLINIC AND GENETIC PRESENTATION OF CHILDREN WITH CYSTINURIA**
ÖZÇELİK Ç., Anarat A., Mungan N. O., BİŞGİN A., ATMIŞ B., MELEK E., KARABAY BAYAZIT A.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1912, 2018 (SCI-Expanded)
- X. **Prospective Turkish Cohort Study to Investigate the Frequency of Niemann-Pick Disease Type C Mutations in Consanguineous Families with at Least One Homozygous Family Member**

Topcu M., Aktas D., Oztoprak M., Mungan N. O., YÜCE A., ALİKAŞİFOĞLU M.

MOLECULAR DIAGNOSIS & THERAPY, vol.21, no.6, pp.643-651, 2017 (SCI-Expanded)

- XI. **Improved metabolic control in tetrahydrobiopterin (BH4), responsive phenylketonuria with sapropterin administered in two divided doses vs. a single daily dose**
Kor D., Yilmaz B. S., BULUT F. D., Ceylaner S., Mungan N. O.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.30, no.7, pp.713-718, 2017 (SCI-Expanded)
- XII. **Propionic acidemia: a Turkish case report of a successful pregnancy, labor and lactation.**
Mungan N., KÖR D., BÜYÜKKURT S., ATMIŞ A., GÜLEÇ Ü., Satar M.
Journal of pediatric endocrinology & metabolism : JPEM, vol.29, pp.863-6, 2016 (SCI-Expanded)
- XIII. **A Case Report of a Very Rare Association of Tyrosinemia type I and Pancreatitis Mimicking Neurologic Crisis of Tyrosinemia Type I.**
UÇAR H., TUMGOR G., KOR D., KARDAŞ F., Mungan N.
Balkan medical journal, vol.33, pp.370-2, 2016 (SCI-Expanded)
- XIV. **Clinical findings and effect of sodium hydrogen carbonate in patients with glutathione synthetase deficiency.**
GÜNDÜZ M., ÜNAL Ö., KAVURT S., TÜRK E., Mungan N.
Journal of pediatric endocrinology & metabolism : JPEM, vol.29, pp.481-5, 2016 (SCI-Expanded)
- XV. **Combination of two different homozygote mutations in Pompe disease.**
ARSLAN A., POYRAZOĞLU H. H., KIRAZ A., ÖZCAN A. A., IŞIK H., ERGUL A., Mungan N., STREUBEL B., CEYLANER S., Altuner T.
Pediatrics international : official journal of the Japan Pediatric Society, vol.58, pp.241-3, 2016 (SCI-Expanded)
- XVI. **Brown-Vialetto-Van Laere syndrome: two siblings with a new mutation and dramatic therapeutic effect of high-dose riboflavin.**
HOROZ O., Mungan N., YILDIZDAS D., HERGÜNER Ö., CEYLANER S., KOR D., WATERHAM H., COSKUN T.
Journal of pediatric endocrinology & metabolism : JPEM, vol.29, pp.227-31, 2016 (SCI-Expanded)
- XVII. **Homozygous familial hypobetalipoproteinemia: A Turkish case carrying a missense mutation in apolipoprotein B.**
YILMAZ B., Mungan N., Di L., MAGNOLO L., ARTUSO L., BERNARDIS I., TUMGOR G., KOR D., TARUGI P.
Clinica chimica acta; international journal of clinical chemistry, vol.452, pp.185-90, 2016 (SCI-Expanded)
- XVIII. **A Desensitization Method to Maintain Enzyme Replacement Therapy in Mucopolysaccharidosis Type VI.**
KOR D., Şeker Y., BULUT F. D., Önenli M., Ufuk A.
Journal of investigational allergology & clinical immunology, vol.26, pp.130-2, 2016 (SCI-Expanded)
- XIX. **Genotypic and phenotypic features of the cystinosis patients from the South Eastern part of Turkey.**
Önenli-Mungan N., KOR D., KARABAY-BAYAZIT A., CENGİZ N., YAVUZ S., NOYAN A., CEYLANER G., ŞEKER-YILMAZ B., TOPALOĞLU A. K., Yuksel B., et al.
The Turkish journal of pediatrics, vol.58, pp.362-370, 2016 (SCI-Expanded)
- XX. **Primary systemic carnitine deficiency: a Turkish case with a novel homozygous SLC22A5 mutation and 14 years follow-up.**
YILMAZ B., KOR D., Mungan N., ERDEM S., CEYLANER S.
Journal of pediatric endocrinology & metabolism : JPEM, vol.28, pp.1179-81, 2015 (SCI-Expanded)
- XXI. **An asymptomatic mother diagnosed with 3-methylcrotonyl-CoA carboxylase deficiency after newborn screening**
KOR D., Mungan N. O., Yilmaz B. S., Oktem M.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.28, pp.669-671, 2015 (SCI-Expanded)
- XXII. **A 17-Year-Old Girl with Chronic Intermittent Abdominal Pain**
Mungan N. O., Yilmaz B. S., Nazoglu S., Yildizdas D., Herguner O., Turgut M., Oktem M.
PEDIATRIC ANNALS, vol.44, no.4, pp.139-141, 2015 (SCI-Expanded)
- XXIII. **TYROSINEMIA TYPE 1 AND NEUROGENIC CRISIS: A CASE REPORT**
BULUT F. D., KOR D., Onenli-Mungan N., Yükselmiş U., İNCECİK F., Yildizdas D.
JOURNAL OF INHERITED METABOLIC DISEASE, vol.35, 2012 (SCI-Expanded)

- XXIV. **Distribution of Gene Mutations Associated with Familial Normosmic Idiopathic Hypogonadotropic Hypogonadism**
GÜRBÜZ F., KOTAN L. D., Mengen E., ŞIKLAR Z., BERBEROĞLU M., Dokmetas S., Kilicli M. F., Guven A., KİREL B., Saka N., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.4, no.3, pp.121-126, 2012 (SCI-Expanded)
- XXV. **Serum IGF-1 and IGFBP-3 Levels in Healthy Children Between 0 and 6 Years of Age**
YÜKSEL B., Ozbek M. N., Mungan N. O., Darendeliler F., Budan B., BİDECİ A., Cetinkaya E., BERBEROĞLU M., Evliyaoglu O., Yesilkaya E., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.3, no.2, pp.84-88, 2011 (SCI-Expanded)
- XXVI. **Ambulatory Blood Pressure Monitoring and Serum Nitric Oxide Concentration in Type 1 Diabetic Children**
Horoz O. O., YÜKSEL B., Bayazit A., ATILGA G., SERTDEMİR Y., Mungan N. O., TOPALOĞLU A. K., OZER G.
ENDOCRINE JOURNAL, vol.56, no.3, pp.477-485, 2009 (SCI-Expanded)
- XXVII. **Serum IGF-1 and IGFBP-3 levels in healthy Turkish children between 0-6 years of age**
YÜKSEL B., Ozbek M. N., Darendeliler F., BİDECİ A., Cetinkaya E., BERBEROĞLU M., Evliyaoglu O., Bas F., Mungan N. O., Yesilkaya E., et al.
HORMONE RESEARCH, vol.72, pp.261, 2009 (SCI-Expanded)
- XXVIII. **A homozygous recurring mutation in WISP3 causing progressive pseudorheumatoid arthropathy of childhood**
Ozbek M. N., Kotan D., Lanktree M., SERİN A., Mungan N. O., CANAN H., ALPER B., YÜKSEL B., Hegele R. A., TOPALOĞLU A. K.
HORMONE RESEARCH, vol.70, pp.187, 2008 (SCI-Expanded)
- XXIX. **Bone calcium changes during diabetic ketoacidosis: A comparison with lactic acidosis due to volume depletion**
Topaloglu A. K., Yildizdas D., Yilmaz H. L., Mungan N., Yuksel B., Ozer G.
BONE, vol.37, no.1, pp.122-127, 2005 (SCI-Expanded)
- XXX. **Thyroid hormone levels and their relationship to survival in children with bacterial sepsis and septic shock**
Yildizdas D., Onenli-Mungan N., Yapicoglu H., Topaloglu A. K., Sertdemir Y., Yuksel B.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.17, no.10, pp.1435-1442, 2004 (SCI-Expanded)
- XXXI. **Growth hormone and insulin like growth factor 1 levels and their relations to survival in children with bacterial sepsis and septic shock**
ÖNENLİ Mungan H. N., YILDIZDAŞ R. D., YAPICIOĞLU YILDIZDAŞ H., TOPALOĞLU A. K., SERTDEMİR Y., YÜKSEL B.
J Paediatr Child Health, vol.40, no.4, pp.221-226, 2004 (SCI-Expanded)

Articles Published in Other Journals

- I. **Clinical features and molecular genetics of autosomal recessive ataxia in the Turkish population**
İNCECİK F., Herguner O. M., Mungan N. O.
JOURNAL OF PEDIATRIC NEUROSCIENCES, vol.15, no.2, pp.86-89, 2020 (ESCI)
- II. **Vitamin B12 levels in patients with mucopolysaccharidosis**
KOR D., Bulut D., Yilmaz B. S., Kilavuz S., Mungan N. O.
CUKUROVA MEDICAL JOURNAL, vol.45, no.2, pp.401-407, 2020 (ESCI)
- III. **A 6-Month-Old Boy with Reddish, Scaly Skin: Netherton Syndrome**
BULUT F. D., KOR D., Yilmaz B. S., YILMAZ M., ALTINTAŞ D. U., Ceylaner S., Kilavuz S., Mungan N. O.
JOURNAL OF PEDIATRIC RESEARCH, vol.5, no.1, pp.54-56, 2018 (ESCI)
- IV. **Neurological assessment of 38 late-diagnosed children with classic phenylketonuria**
HAYTOĞLU Z., Herguner O., SOYUPAK S., TOPALOĞLU A. K., YÜKSEL B., OZER G., Mungan H. N. O.
CUKUROVA MEDICAL JOURNAL, vol.41, no.1, pp.21-27, 2016 (ESCI)
- V. **The analysis of the phenylalanine hydroxylase gene mutations by sequencing and ARMS techniques**

in Turkish patients

Luleyap U., PAZARCI P., CÖMERTPAY G., Onenli H. N., PAZARBAŞI A., ALPTEKİN D., KASAP H., FROSTER U.
CUKUROVA MEDICAL JOURNAL, vol.41, no.4, pp.702-708, 2016 (ESCI)

- VI. **Brown-Vialetto-Van Laere syndrome: Two siblings with a new mutation and dramatic therapeutic effect of high dose riboflavin**
ÖNENLİ Mungan H.
J PEDIATR ENDOCRINOL METAB, vol.2015, pp.198-205, 2015 (Peer-Reviewed Journal)
- VII. **Continuous Venovenous Hemodiafiltration in Three Newborn Patients with Hyperammonemia**
YAPICIOĞLU YILDIZDAŞ H., Yildizdas D., GÜLDEREN ÖZLÜ F., Mert K., Mungan N. O.
CUKUROVA MEDICAL JOURNAL, vol.40, pp.161-166, 2015 (ESCI)
- VIII. **Fabry Disease: A Turkish Case with a Novel Mutation and Dermatological Manifestations**
Mungan N. O., Temiz F., Yilmaz B. S., Ozbek M. N., KARAKAŞ M., TOPALOĞLU A. K., YÜKSEL B.
CUKUROVA MEDICAL JOURNAL, vol.40, pp.156-160, 2015 (ESCI)
- IX. **Evaluation of Two Different Pamidronate Treatment Protocols in Children with Osteogenesis Imperfecta**
ÖNENLİ Mungan H., gürbüz F., mengen e., özgür ö., topaloğlu a. k., yüksel b.
CUKUROVA MEDICAL JOURNAL, vol.39, no.3, pp.532-539, 2014 (Peer-Reviewed Journal)
- X. **Hashimoto's Encephalopathy: Four Cases and Review of Literature.**
YÜKSEL B., İNCECİK F., HERGÜNER M., ÖZGÜR HOROZ Ö., YILDIZDAŞ R. D., ALTUNBAŞAK Ş., ÖNENLİ Mungan H., GÜL MERT G.
Int J Neurosci, vol.0, 2013 (Peer-Reviewed Journal)
- XI. **Autism symptoms related to Tyrosinemia type III: a case report**
YOLGA TAHİROĞLU A., ÖNENLİ Mungan H., FIRAT S., AVCI A.
TURKISH JOURNAL OF ENDOCRINOLOGY AND METABOLISM, vol.12, no.2, pp.55-56, 2008 (ESCI)
- XII. **Pediatric kafa travmalarında idrar antidiürik hormon seviyeleri**
YAMAN A., YÜKSEL B., AKSARAY N., ÖNENLİ Mungan H., YILDIZDAŞ R. D., Alhan A.
Ç.Ü.Sağlık Bil Derg, vol.9,10, pp.45-54, 1996 (Peer-Reviewed Journal)

Refereed Congress / Symposium Publications in Proceedings

- I. **YENİDOĞANDA BİYOKİMYASAL HİPERTROİDİSM**
YAPICIOĞLU YILDIZDAŞ H., ÖNENLİ Mungan H., AKÇALI M., GÜLDEREN ÖZLÜ F.
UNEKO 25, Antalya, Turkey, pp.182
- II. **YENİDOĞANDA BİYOKİMYASAL HİPERTROİDİSM**
YAPICIOĞLU YILDIZDAŞ H., ÖNENLİ Mungan H., AKÇALI M., GÜLDEREN ÖZLÜ F.
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- III. **YENİDOĞANDA BİYOKİMYASAL HİPERTROİDİSM**
YAPICIOĞLU YILDIZDAŞ H., ÖNENLİ Mungan H., AKÇALI M., GÜLDEREN ÖZLÜ F.
UNEKO 25, Antalya, Turkey, pp.182
- IV. **First case report of Gaucher disease and Graves' thyroiditis**
Mungan N. O., KOR D., Kilavuz S., Bulut D., Yilmaz B. S.
16th Annual Research Meeting of the WORLD Symposium(TM), Florida, United States Of America, 10 - 14 February 2020, vol.129
- V. **SON DÖNEM BÖBREK YETMEZLİĞİ VE SİSTİNOZİS: 41 YAŞINDA GÖZ MUAYENESİ İLE TANI ALABİLEN BİR OLSU SUNUMU**
DEMİR İ., ŞEKER YILMAZ B., KILAVUZ S., KÖR D., DERYA BULUT F., ERDEM E., ÖNENLİ Mungan H., PAYDAŞ S.
ULUSLARARASI KATILIMLI 6. LİZOZOMAL HASTALIKLAR KONGRESİ, Antalya, Turkey, 11 - 15 April 2018, pp.95
- VI. **Chanarin-Dorfman syndrome: A case report**
Mungan N. O., TUNCEZ E., YILMAZ B. S., LEBLEBİSATAN G., KUNT Z., BULUT D., KOR D.
12th Annual WORLD Symposium, California, United States Of America, 29 February - 04 March 2016, vol.117

- VII. **Ağır laktik asidoz gelişen MMA?lı bir olguda yüksek doz askorbik asit tedavisiyle salah**
KÖR D., ÖNENLİ MÜNGAN H., Özgür Ö., Yükselmiş U., Sarı Y., Ece Ü., Güntek S., YILDIZDAŞ R. D.
. XII. Metabolik Hastalıklar ve Beslenme Kongresi, Eskişehir, Turkey, 1 - 04 May 2013
- VIII. **Ağır laktik asidoz gelişen MMA?lı bir olguda yüksek doz askorbik asit tedavisiyle salah**
KÖR D., ÖNENLİ MÜNGAN H., Özgür Ö., Yükselmiş U., Sarı Y., Ece Ü., Güntek S., YILDIZDAŞ R. D.
. XII. Metabolik Hastalıklar ve Beslenme Kongresi, Eskişehir, Turkey, 1 - 04 May 2013
- IX. **Brown-Vialetto-Van Laere ve Fazio Londe Sendromlu Bir Olgu Sunumu.**
Gülen M., YILDIZDAŞ R. D., Sarı M., ÖZGÜR HOROZ Ö., Yükselmiş U., ÖNENLİ MÜNGAN H.
Uluslararası Katılımlı X. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Kongresi VI. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Hemşireliği Kongresi, Antalya, Turkey, 3 - 07 April 2013
- X. **Brown-Vialetto-Van Laere ve Fazio Londe Sendromlu Bir Olgu Sunumu.**
Gülen M., YILDIZDAŞ R. D., Sarı M., ÖZGÜR HOROZ Ö., Yükselmiş U., ÖNENLİ MÜNGAN H.
Uluslararası Katılımlı X. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Kongresi VI. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Hemşireliği Kongresi, Antalya, Turkey, 3 - 07 April 2013
- XI. **"A New Method For The Detection Of Ivs 10 11g-A Mutation In Phenylalanine Hydroxylase Gene With Arms Technique In Turkish Phenylketonuria Patients"**
LÜLEYAP H. Ü., ONATOGLU D., PAZARBAŞI A., GUZEL A. I., OZER G., ÖNENLİ MÜNGAN H., KASAP M., ALPTEKİN D.
XX International Congress of Genetics,, Berlin, Germany, 12 - 17 July 2008, vol.625, pp.196
- XII. **"A Modified Arms Tecnique For Detection Of P281l Mutations In Phenilalanine Hydroxylase Gene In Turkish Phenylketonuria Patients"**
LÜLEYAP H. Ü., PAZARBAŞI A., ONATOGLU D., ALPTEKİN D., GUZEL A. I., ÖNENLİ MÜNGAN H., FROSTER U.
XX International Congress of Genetics, Berlin, Germany, 12 - 17 July 2008, vol.439, pp.158
- XIII. **"A Modified Arms Tecnique For Detection Of P281l Mutations In Phenilalanine Hydroxylase Gene In Turkish Phenylketonuria Patients"**
LÜLEYAP H. Ü., PAZARBAŞI A., ONATOGLU D., ALPTEKİN D., GUZEL A. I., ÖNENLİ MÜNGAN H., FROSTER U.
XX International Congress of Genetics, Berlin, Germany, 12 - 17 July 2008, vol.439, pp.158
- XIV. **"A New Method For The Detection Of Ivs 10 11g-A Mutation In Phenylalanine Hydroxylase Gene With Arms Technique In Turkish Phenylketonuria Patients"**
LÜLEYAP H. Ü., ONATOGLU D., PAZARBAŞI A., GUZEL A. I., OZER G., ÖNENLİ MÜNGAN H., KASAP M., ALPTEKİN D.
XX International Congress of Genetics,, Berlin, Germany, 12 - 17 July 2008, vol.625, pp.196

Supported Projects

KARABAY BAYAZIT A., BİŞGİN A., ÖZÇELİK Ç., ANARAT A., ÖNENLİ MÜNGAN H., ATMIŞ B., MELEK E., Project Supported by Higher Education Institutions, ÇOCUKLUK YAŞ GRUBUNDA BAŞLAYAN SİSTİNÜRİ TANISIYLA TAKİPLİ HASTALARDA GENETİK ÇALIŞMA VE KLİNİĞE YANSIMASI, 2017 - 2018

Metrics

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