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Yoksis Araştırmacı ID: 34569

Eğitim Bilgileri

Doktora, Atatürk Üniversitesi, Fen Bilimleri Enstitüsü, Kimya (Dr), Türkiye 1998 - 2002

Yüksek Lisans, Atatürk Üniversitesi, Fen Bilimleri Enstitüsü, Kimya (YL) (Tezli), Türkiye 1996 - 1998

Lisans, Van Yüzüncü Yıl Üniversitesi, Fen-Edebiyat Fakültesi, Kimya Bölümü, Türkiye 1991 - 1995

Yaptığı Tezler

Doktora, Koyun eritrositleri ve göz lensinden glukoz-6-fosfat dehidrogenaz enziminin saflaştırılması, karakterizasyonu, bazı ilaç ve kimyasal maddelerin inhibisyon veya aktivasyon kinetiklerinin incelenmesi, Atatürk Üniversitesi, Fen Bilimleri Enstitüsü, Kimya (Dr), 2002

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Araştırma Alanları

Biyokimya, Kimya, Biyokimya

Akademik Unvanlar / Görevler

Prof. Dr., Anadolu Üniversitesi, ECZACILIK FAKÜLTESİ, TEMEL ECZACILIK BİLİMLERİ BÖLÜMÜ, 2016 - Devam Ediyor

Prof. Dr., Atatürk Üniversitesi, Fen Fakültesi, Kimya Bölümü, 2012 - 2016

Doç. Dr., Atatürk Üniversitesi, Fen Fakültesi, Kimya Bölümü, 2006 - 2012

Yrd. Doç. Dr., Atatürk Üniversitesi, Fen Fakültesi, Kimya Bölümü, 2002 - 2006

Araştırma Görevlisi, Atatürk Üniversitesi, Fen Fakültesi, Kimya Bölümü, 1996 - 2002

Akademik İdari Deneyim

Bilecik Şeyh Edebali Üniversitesi, 2020 - Devam Ediyor

Bursa Teknik Üniversitesi, 2016 - 2016

İğdır Üniversitesi, 2015 - 2016

Atatürk Üniversitesi, 2010 - 2012

Atatürk Üniversitesi, 2010 - 2011

Verdiği Dersler

GENEL KİMYA, Lisans, 2015 - 2016, 2014 - 2015, 2013 - 2014, 2012 - 2013, 2011 - 2012, 2010 - 2011, 2009 - 2010

BİYOKİMYA, Lisans, 2015 - 2016, 2011 - 2012, 2010 - 2011

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Bitki Biyokimyası, Lisans, 2010 - 2011

Laboratuar Deneyimi, Yüksek Lisans, 2010 - 2011

Biyokimya Laboratuvarı, Lisans, 2009 - 2010

Yönetilen Tezler

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ŞÜKRÜ B., İnsan trombositlerinden nitrik oksit sentaz enziminin ve insan serumundan Paraoksonaz-1 enziminin saflaştırılması, bazı ilaçların bu enzimlerin aktiviteleri üzerine etkilerinin incelenmesi, Doktora, Z.ALIM(Öğrenci), 2015

ŞÜKRÜ B., Koyun karaciğerinden aldoz reduktaz ve sorbitol dehidrogenaz enzimlerinin saflaştırılması ve bazı fenolik asitlerin enzimlerin aktivitesi üzerine etkilerinin incelenmesi, Yüksek Lisans, H.ESRA(Öğrenci), 2015

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ŞÜKRÜ B., Hamsi (*Engraulis encrasiculus*) ve mezgit (*Merlangius merlangus euxinus*) balıklarının solungaç dokularından karbonik anhidraz enziminin saflaştırılması, karakterizasyonu ve bazı metal iyonlarının enzim aktivitesi üzerine

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ŞÜKRÜ B., Nitrik oksit sentaz enziminin siğır böbrek ve trombositlerinden saflaştırılması, karakterizasyonu ve bazı ilaçların enzim aktivitesi üzerine etkilerinin incelenmesi, Doktora, M.MUSTAFA(Öğrenci), 2013

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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

I. New naphthoquinone thiazole hybrids as carbonic anhydrase and cholinesterase inhibitors: Synthesis, crystal structure, molecular docking, and acid dissociation constant

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JOURNAL OF MOLECULAR STRUCTURE, cilt.1301, 2024 (SCI-Expanded)

II. Protective effects of esculetin against doxorubicin-induced toxicity correlated with oxidative stress in rat liver: In vivo and in silico studies

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III. Novel spiroindoline derivatives targeting aldose reductase against diabetic complications: Bioactivity, cytotoxicity, and molecular modeling studies

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IV. The impact of some metals, molecular docking and molecular dynamic calculations on glucose 6-phosphate dehydrogenase activity in *< i> Capoeta</i>< i> trutta</i>* (Heckel, 1843) tissue

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V. New *< i> N</i>- (1,3,4-thiadiazole-2-yl)acetamide derivatives as human carbonic anhydrase I and II and acetylcholinesterase inhibitors*

- Dawbaa S., Tuerkes C., Nuha D., DEMİR Y., Evren A. E., YURTTAŞ L., BEYDEMİR \$.
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- VI. Isolation of phenolic compounds from eco-friendly white bee propolis: Antioxidant, wound-healing, and anti-Alzheimer effects
NECİP A., Demirtas I., Tayhan S. E., İŞIK M., BİLGİN S., Turan I. F., Ipek Y., Beydemir \$.
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- IX. Design and Synthesis of Pyrazole Carboxamide Derivatives as Selective Cholinesterase and Carbonic Anhydrase Inhibitors: Molecular Docking and Biological Evaluation
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- XI. Synthesis, Theoretical, in Silico and in Vitro Biological Evaluation Studies of New Thiosemicarbazones as Enzyme Inhibitors
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CHEMISTRY & BIODIVERSITY, cilt.20, sa.11, 2023 (SCI-Expanded)
- XII. Novel bis-ureido-substituted sulfaguanidines and sulfisoxazoles as carbonic anhydrase and acetylcholinesterase inhibitors
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- XIII. Synthesis and characterization of novel acyl hydrazones derived from vanillin as potential aldose reductase inhibitors
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- XV. Novel beta-lactam substituted benzenesulfonamides: in vitro enzyme inhibition, cytotoxic activity and in silico interactions
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- XVI. Discovery of novel benzenesulfonamides incorporating 1,2,3-triazole scaffold as carbonic anhydrase I, II, IX, and XII inhibitors
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- XVII. A novel series of thiosemicarbazone hybrid scaffolds: Design, synthesis, DFT studies, metabolic enzyme inhibition properties, and molecular docking calculations
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- XVIII. **A new series of hydrazones as small-molecule aldose reductase inhibitors**
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- XIX. **Novel acetic acid derivatives containing quinazolin-4(3H)-one ring: Synthesis, in vitro, and in silico evaluation of potent aldose reductase inhibitors**
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- XX. **Enzyme inhibition, molecular docking, and density functional theory studies of new thiosemicarbazones incorporating the 4-hydroxy-3,5-dimethoxy benzaldehyde motif**
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- XXI. **The synthesis, biological evaluation and in silico studies of asymmetric 3,5-diaryl-rhodanines as novel inhibitors of human carbonic anhydrase isoenzymes**
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- XXII. **Exploration of Some Bis-Sulfide and Bis-Sulfone Derivatives as Non-Classical Aldose Reductase Inhibitors**
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Duran H. E., BEYDEMİR Ş.
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- XXVI. **Molecular Docking Studies and the Effect of Fluorophenylthiourea Derivatives on Glutathione-Dependent Enzymes**
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- XXVII. **In Vitro Inhibitory Activity and Molecular Docking Study of Selected Natural Phenolic Compounds as AR and SDH Inhibitors**
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- XXVIII. **N-substituted phthalazine sulfonamide derivatives as non-classical aldose reductase inhibitors**
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- XXIX. **Biological evaluation and in silico study of benzohydrazide derivatives as paraoxonase 1 inhibitors**
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- XXX. **Methyl benzoate derivatives: in vitro Paraoxonase 1 inhibition and in silico studies**
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- XXXI. **Cytotoxic effect, enzyme inhibition, and in silico studies of some novel N-substituted sulfonyl amides incorporating 1,3,4-oxadiazol structural motif**

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- XXXII. **Design, synthesis, and biological activity of novel dithiocarbamate-methylsulfonyl hybrids as carbonic anhydrase inhibitors**
OSMANİYE D., TÜRKEŞ C., Demir Y., ÖZKAY Y., BEYDEMİR \$., KAPLANCIKLİ Z. A.
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Akdag M., ÖZCELİK A. B., Demir Y., BEYDEMİR \$.
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- XXXIV. **Synthesis, biological evaluation, and in silico study of novel library sulfonates containing quinazolin-4(3H)-one derivatives as potential aldose reductase inhibitors**
TOKALI F. S., Demir Y., Demircioglu I. H., TÜRKEŞ C., Kalay E., Sendil K., BEYDEMİR \$.
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- XXXV. **In vitro and in silico interactions of antiulcer, glucocorticoids and urological drugs on human carbonic anhydrase I and II isozymes**
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- XXXVI. **AChE mRNA expression as a possible novel biomarker for the diagnosis of coronary artery disease and Alzheimer's disease, and its association with oxidative stress**
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- XXXVII. **Oxidative Stress and Changes of Important Metabolic Gene Expressions as a Potential Biomarker in the Diagnosis of Atherosclerosis in Leukocytes**
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- 2nd European Chemistry Conference (ECC-2019), Amsterdam, Hollanda, 18 - 19 Mayıs 2019
- XXXII. **In vitro Effects and In silico Studies on Paraoxonase 1 (PON1) of Some Plant Extracts from Mavrovo (Macedonia) Region**
BEYDEMİR Ş., BÜYÜKEMİR O., NEZİR D., TÜRKEŞ C.
2nd European Chemistry Conference, Amsterdam, Hollanda, 15 - 16 Mayıs 2019, ss.33
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- XXXIV. **Effects of Some Food Additives on Carbonic Anhydrase Activity**
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1st International Technology Sciences and Design Symposium, 27 - 29 Haziran 2018
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- XXXVIII. **Investigation Inhibition Effects of N-Benzyl-N-(4-Methoxyphenyl) Acetamide, N-Benzyl-N-(4-Methoxyphenyl)-4 Methylbenzenesulfonamide, N-Benzyl-4-Methoxyaniline and (E)-N-(4-Methoxyphenyl)-1-Phenylmethanimine Derivatives on the Aldose Reductase Enzyme and Determination of Enzyme-inhibitor Interaction with Docking Method**
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- XL. **Oxidative Stress Biomarkers in Coronary Artery and Cerebrovascular Disease**
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- XLI. **Determination of mRNA expression in the leukocytes of coronary artery and cerebrovascular patients**
İŞIK M., TUNÇ A., BEYDEMİR Ş.
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- XLII. **İnsan Serum Paraoksonaz-I (hPON1): Bazı Antibakteriyel İlaçların In Vitro Etkileri**
TÜRKEŞ C., BEYDEMİR Ş.
3. Ulusal Uygulamalı Biyolojik Bilimler Kongresi - UBBK 2018, Eskişehir, Türkiye, 3 - 05 Mayıs 2018, ss.33
- XLIII. **5-Hydroxy-1,4-naphthoquinone, Morin hydrate and Valeric acid Inhibit Human Serum Paraoxonase-I**
TÜRKEŞ C., BEYDEMİR Ş.
4th International Congress on Applied Biological Sciences - ICABS 2018, Eskişehir, Türkiye, 3 - 05 Mayıs 2018, ss.165
- XLIV. **The Effects of Antimycotic Drugs on Human Serum Paraoxonase-I**
TÜRKEŞ C., BEYDEMİR Ş.
4th International Congress on Applied Biological Sciences - ICABS 2018, Eskişehir, Türkiye, 3 - 05 Mayıs 2018, ss.57
- XLV. **Inhibition Impacts of Some Anesthetics on Human Carbonic Anhydrase Isoenzymes (hCA I - II)**

Activities

TÜRKES C., DEMİR Y., BEYDEMİR Ş.

International Eurasian Conference on Biological and Chemical Sciences, 26 - 27 Nisan 2018

XLVI. Inhibition of Glutathione S-Transferase by Usnic and Carnosic Acid

CEYLAN H., DEMİR Y., BEYDEMİR Ş.

International Biochemistry Congress/28th National Biochemistry Congress, Erzurum, Türkiye, 19 - 23 Eylül 2017,
cilt.42

XLVII. The Effect of 8-Methoxypsoralen on Polyol Pathway Enzymes

DEMİR Y., CEYLAN H., BEYDEMİR Ş.

International Biochemistry Congress/28th National Biochemistry Congress, Erzurum, Türkiye, 19 - 23 Eylül 2017,
cilt.42

XLVIII. Inhibition Effects and Molecular Docking Studies of Novel Aldose Reductase Inhibitory Class N-Benzyl-4-Methoxyaniline Derivatives

KILINÇ N., YILDIZHAN G., BAYRAK Ç., MENZEK A., DURDAĞI S., BEYDEMİR Ş.

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XLIX. Investigation of in Vivo Effects of Arachidonoyl Dopamine on Lipid Peroxidation and Some Enzyme Activities in Rat Tissue and Erythrocytes

HUYUT Z., BEYDEMİR Ş., GÜLÇİN İ., HUYUT M. T.

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L. Synthesis and carbonic anhydrase inhibitory activity of some novel sulfonamides

OSMANİYE D., LEVENT S., ACAR ÇEVİK U., SAĞLIK B. N., DEMİR Y., ÖZKAY Y., ILGIN S., KORKUT B., BEYDEMİR Ş., KAPLANCIKLİ Z. A.

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LII. Influence of Calcium Channel Blockers on Sheep Kidney Aldose Reductase Enzyme

TÜRKES C., DEMİR Y., BEYDEMİR Ş.

3rd International Multidisciplinary Symposium on Drug Research and Development, 5 - 07 Ekim 2017

LIII. In vitro effects of some metals on glucose 6-phosphate dehydrogenase purified from freshwater fish Capoeta umbla gill.

KIRICI M., ATAMANALP M., KIRICI M., BEYDEMİR Ş.

The 4th International Scientific Conference Current Problems of Biochemistry and Cell Biology, Dnipro, Ukrayna, 5 - 06 Ekim 2017, ss.217-218

LIV. Why There is No Difference in Antioxidant Levels Between Acute Perforated and Suppurative Appendicitis?

Temiz A., İŞIK M., ALBAYRAK Y., Aslan H. E., Aslan A., Demir Y., Atasoy G., ALBAYRAK A., Özdemir Y., BEYDEMİR Ş.

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LV. Evaluation of chalcone derivatives as inhibitors of glutathione S-transferase

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LVII. Synthesis of The 4-(2-Aminoethyl)-Benzenesulfonamide Derivatives and Inhibitory Activity Against Human PON1

DEMİR Y., İŞIK M., DURGUN M., NECİP A., aslan h. e., BEYDEMİR Ş.

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LVIII. Inhibition Effects of Chemotherapy Drugs on Human PON1

DEMİR Y., TÜRKES C., BEYDEMİR Ş.

- International Congress on Fundamental and Applied Sciences, 21 - 25 Ağustos 2017
- LIX. **Investigation Effects of Some Sulfonamide Derivatives on ACHE Enzyme**
İŞİK M., BEYDEMİR Ş., DEMİR Y., DURGUN M., NECİP A.
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- LX. **INHIBITION EFFECTS OF CHEMOTHERAPY DRUGS ON HUMAN PON1**
DEMİR Y., TÜRKEŞ C., BEYDEMİR Ş.
International Congress on Fundamental and Applied Sciences 2017 (ICFAS2017), 21 - 25 Ağustos 2017
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- LXII. **Biosensor Applications of Enzyme Immobilized to Cationic Octadecylamine by Langmuir-Blodgett Techniques**
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