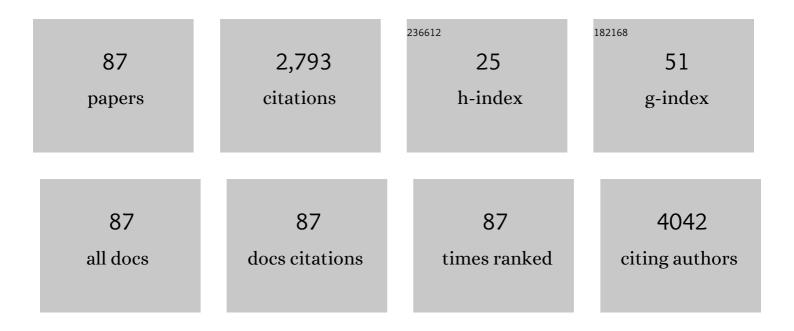
List of Publications by Year in descending order

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Μιιανι Ιισςα

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | A mutation in the SAA1 promoter causes hereditary amyloid A amyloidosis. Kidney International, 2022, 101, 349-359. | 2.6 | 10 |
| 2 | The interactions between DNA nanostructures and cells: A critical overview from a cell biology perspective. Acta Biomaterialia, 2022, 146, 10-22. | 4.1 | 10 |
| 3 | Liver stiffness measured by two-dimensional shear-wave elastography predicts hepatic vein pressure gradient at high values in liver transplant candidates with advanced liver cirrhosis. PLoS ONE, 2021, 16, e0244934. | 1.1 | 6 |
| 4 | Expression of Interferons Lambda 3 and 4 Induces Identical Response in Human Liver Cell Lines Depending Exclusively on Canonical Signaling. International Journal of Molecular Sciences, 2021, 22, 2560. | 1.8 | 5 |
| 5 | Alpha-1 Antitrypsin and Hepatocellular Carcinoma in Liver Cirrhosis: SERPINA1 MZ or MS Genotype Carriage Decreases the Risk. International Journal of Molecular Sciences, 2021, 22, 10560. | 1.8 | 8 |
| 6 | Light-induced modulation of the mitochondrial respiratory chain activity: possibilities and limitations. Cellular and Molecular Life Sciences, 2020, 77, 2815-2838. | 2.4 | 29 |
| 7 | ABCB4 disease: Many faces of one gene deficiency. Annals of Hepatology, 2020, 19, 126-133. | 0.6 | 35 |
| 8 | Donor PNPLA3 and TM6SF2 Variant Alleles Confer Additive Risks for Graft Steatosis After Liver Transplantation, 2020, 104, 526-534. | 0.5 | 16 |
| 9 | Analyzing the mechanisms of iron oxide nanoparticles interactions with cells: A road from failure to success in clinical applications. Journal of Controlled Release, 2020, 328, 59-77. | 4.8 | 72 |
| 10 | Hepatic Tumor Cell Morphology Plasticity under Physical Constraints in 3D Cultures Driven by YAP–mTOR Axis. Pharmaceuticals, 2020, 13, 430. | 1.7 | 5 |
| 11 | Iron Oxide Nanoparticle-Induced Autophagic Flux Is Regulated by Interplay between p53-mTOR Axis and Bcl-2 Signaling in Hepatic Cells. Cells, 2020, 9, 1015. | 1.8 | 25 |
| 12 | Progressive lysosomal membrane permeabilization induced by iron oxide nanoparticles drives hepatic cell autophagy and apoptosis. Nano Convergence, 2020, 7, 17. | 6.3 | 19 |
| 13 | Olgu sunumu: moleküler düzeyde tanısı konulmuş olan ilk Türk Rotor sendromlu aile. Turk Pediatri Arsivi, 2020, 55, 430-433. | 0.9 | 0 |
| 14 | ABCB4 disease mimicking morbus Wilson: A potential diagnostic pitfall. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2020, 164, 121-125. | 0.2 | 2 |
| 15 | Dubin-Johnson syndrome in Tunisia: Spectrum of a rare disease. Presse Medicale, 2019, 48, 81-82. | 0.8 | 1 |
| 16 | PNPLA3 rs738409 G allele carriers with genotype 1b HCV cirrhosis have lower viral load but develop liver failure at younger age. PLoS ONE, 2019, 14, e0222609. | 1.1 | 3 |
| 17 | IL28B rs12979860 T allele protects against CMV disease in liver transplant recipients in the postâ€prophylaxis and late period. Transplant Infectious Disease, 2019, 21, e13124. | 0.7 | 5 |
| 18 | Beyond an Obvious Cause of Cholestasis in a Toddler: Compound Heterozygosity for <i>ABCB11</i> Mutations. Pediatrics, 2019, 143, . | 1.0 | 8 |

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|----|--|------------------|--------------|
| 19 | Remote Actuation of Apoptosis in Liver Cancer Cells via Magneto-Mechanical Modulation of Iron Oxide Nanoparticles. Cancers, 2019, 11, 1873. | 1.7 | 40 |
| 20 | Functional evaluation of variants of unknown significance in the <i>BRCA2</i> gene identified in genetic testing. Cancer Biology and Therapy, 2019, 20, 633-641. | 1.5 | 1 |
| 21 | Targeting the mTOR Signaling Pathway Utilizing Nanoparticles: A Critical Overview. Cancers, 2019, 11, 82. | 1.7 | 34 |
| 22 | Non-Thermal Plasma, as a New Physicochemical Source, to Induce Redox Imbalance and Subsequent Cell Death in Liver Cancer Cell Lines. Cellular Physiology and Biochemistry, 2019, 52, 119-140. | 1.1 | 33 |
| 23 | Donor PNPLA3 rs738409 genotype is a risk factor for graft steatosis. A post-transplant biopsy-based study. Digestive and Liver Disease, 2018, 50, 490-495. | 0.4 | 25 |
| 24 | Variable X-chromosome inactivation and enlargement of pericentral glutamine synthetase zones in the liver of heterozygous females with OTC deficiency. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 1029-1039. | 1.4 | 12 |
| 25 | Manipulating the mitochondria activity in human hepatic cell line Huh7 by low-power laser irradiation. Biomedical Optics Express, 2018, 9, 1283. | 1.5 | 21 |
| 26 | New Insights in Genetic Cholestasis: From Molecular Mechanisms to Clinical Implications. Canadian Journal of Gastroenterology and Hepatology, 2018, 2018, 1-12. | 0.8 | 59 |
| 27 | Chemically different non-thermal plasmas target distinct cell death pathways. Scientific Reports, 2017, 7, 600. | 1.6 | 36 |
| 28 | Cytotoxic bilirubin metabolites overlooked so far. Journal of Hepatology, 2017, 67, 214-215. | 1.8 | 2 |
| 29 | Nanoparticle core stability and surface functionalization drive the mTOR signaling pathway in hepatocellular cell lines. Scientific Reports, 2017, 7, 16049. | 1.6 | 38 |
| 30 | Hepcidin knockout mice spontaneously develop chronic pancreatitis owing to cytoplasmic iron overload in acinar cells. Journal of Pathology, 2017, 241, 104-114. | 2.1 | 36 |
| 31 | The use of pulsed magnetic fields to increase the uptake of iron oxide nanoparticles by living cells. Applied Physics Letters, 2017, 111, . | 1.5 | 19 |
| 32 | Prevalence and risk factors of steatosis after liver transplantation and patient outcomes. Liver Transplantation, 2016, 22, 644-655. | 1.3 | 56 |
| 33 | Modulation of collective cell behaviour by geometrical constraints. Integrative Biology (United) Tj ETQq1 1 0.784 | 1314 rgBT 0.6 | /Overlock 10 |
| 34 | HGSNAT has a TATA-less promoter with multiple starts of transcription. Gene, 2016, 592, 36-42. | 1.0 | 2 |
| 35 | Successful Treatment of Iron-Overload Cardiomyopathy in Hereditary Hemochromatosis With Deferoxamine and Deferiprone. Canadian Journal of Cardiology, 2016, 32, 1574.e1-1574.e3. | 0.8 | 17 |
| 36 | Rare variants in known and novel candidate genes predisposing to statin-associated myopathy. Pharmacogenomics, 2016, 17, 1405-1414. | 0.6 | 17 |

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|----|--|-----|-----------|
| 37 | Radionuclide cholescintigraphy in genetically confirmed Rotor syndrome. Pediatrics International, 2015, 57, 981-985. | 0.2 | 1 |
| 38 | USP18 downregulation in peripheral blood mononuclear cells predicts nonresponse to interferon-based triple therapy in patients with chronic hepatitis C, genotype 1: a pilot study. Therapeutics and Clinical Risk Management, 2015, 11, 1853. | 0.9 | 2 |
| 39 | Hereditary haemochromatosis caused by homozygous <i><scp>HJV</scp></i> mutation evolved through paternal disomy. Clinical Genetics, 2015, 87, 96-98. | 1.0 | 1 |
| 40 | NKD1 marks intestinal and liver tumors linked to aberrant Wnt signaling. Cellular Signalling, 2015, 27, 245-256. | 1.7 | 19 |
| 41 | Alagille Syndrome Mimicking Biliary Atresia in Early Infancy. PLoS ONE, 2015, 10, e0143939. | 1.1 | 21 |
| 42 | Relevance of low viral load in haemodialysed patients with chronic hepatitis C virus infection. World Journal of Gastroenterology, 2015, 21, 5496. | 1.4 | 8 |
| 43 | Progressive familial intrahepatic cholestasis type 2 – paediatric patients followed at the Paediatric Clinic of the 2nd Medical Faculty, University Hospital Motol, Prague. Gastroenterologie A Hepatologie, 2015, 69, 547-553. | 0.0 | 0 |
| 44 | Down-regulation of OATP1B proteins correlates with hyperbilirubinemia in advanced cholestasis. International Journal of Clinical and Experimental Pathology, 2015, 8, 5252-62. | 0.5 | 6 |
| 45 | Treatment of pruritus with <scp>P</scp> rometheus dialysis and absorption system in a patient with benign recurrent intrahepatic cholestasis. Hepatology Research, 2014, 44, E304-E308. | 1.8 | 10 |
| 46 | Genetic variation in TNFA predicts protection from severe bacterial infections in patients with end-stage liver disease awaiting liver transplantation. Journal of Hepatology, 2014, 60, 773-781. | 1.8 | 7 |
| 47 | <i>ABCB4</i> mutations underlie hormonal cholestasis but not pediatric idiopathic gallstones. World Journal of Gastroenterology, 2014, 20, 5867. | 1.4 | 21 |
| 48 | Troy, a Tumor Necrosis Factor Receptor Family Member, Interacts With Lgr5 to Inhibit Wnt Signaling in Intestinal Stem Cells. Gastroenterology, 2013, 144, 381-391. | 0.6 | 94 |
| 49 | Ornithine carbamoyltransferase deficiency: molecular characterization of 29 families. Clinical Genetics, 2013, 84, 552-559. | 1.0 | 6 |
| 50 | New insights in bilirubin metabolism and their clinical implications. World Journal of Gastroenterology, 2013, 19, 6398. | 1.4 | 151 |
| 51 | Dubin-Johnson syndrome coinciding with colon cancer and atherosclerosis. World Journal of Gastroenterology, 2013, 19, 946. | 1.4 | 12 |
| 52 | Nasobiliary Drainage in an Episode of Intrahepatic Cholestasis in a Child With Mild <i>ABCB11</i> Disease. Journal of Pediatric Gastroenterology and Nutrition, 2012, 55, 88-90. | 0.9 | 9 |
| 53 | Complete OATP1B1 and OATP1B3 deficiency causes human Rotor syndrome by interrupting conjugated bilirubin reuptake into the liver. Journal of Clinical Investigation, 2012, 122, 519-528. | 3.9 | 321 |
| 54 | FAT MASS AND OBESITYâ€ASSOCIATED (FTO) GENE AND ALCOHOL INTAKE. Addiction, 2012, 107, 1185-1186. | 1.7 | 7 |

| ATP8B1 Gene Expression Is Driven by a Housekeeping-Like Promoter Independent of Bile Acids a Farnesoid X Receptor. PLoS ONE, 2012, 7, e51650. Glucocerebrosidase gene has an alternative upstream promoter, which has features and express characteristic of housekeeping genes. Blood Cells, Molecules, and Diseases, 2011, 46, 239-245. Disruption of OTC promoter-enhancer interaction in a patient with symptoms of ornithine carbamoyltransferase deficiency. Human Mutation, 2010, 31, E1294-E1303. Serum bilirubin levels and UGT1A1 promoter variations in patients with schizophrenia. Psychiatr Research, 2010, 178, 449-450. Do common genetic variants in endotoxin signaling pathway contribute to predisposition to alcoholic liver cirrhosis?. Clinical Chemistry and Laboratory Medicine, 2009, 47, 398-404. | sion | 4 6 |
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| ⁵⁶ characteristic of housekeeping genes. Blood Čells, Molecules, and Diseases, 2011, 46, 239-245. ⁵⁷ Disruption of OTC promoter-enhancer interaction in a patient with symptoms of ornithine carbamoyltransferase deficiency. Human Mutation, 2010, 31, E1294-E1303. ⁵⁸ Serum bilirubin levels and UGT1A1 promoter variations in patients with schizophrenia. Psychiatr Research, 2010, 178, 449-450. ⁵⁰ Do common genetic variants in endotoxin signaling pathway contribute to predisposition to | | 6 |
| ⁵⁷ carbamoyltransferase deficiency. Human Mutation, 2010, 31, E1294-E1303. Serum bilirubin levels and UGT1A1 promoter variations in patients with schizophrenia. Psychiate Research, 2010, 178, 449-450. Do common genetic variants in endotoxin signaling pathway contribute to predisposition to | 1.1 | |
| ⁵⁸ Research, 2010, 178, 449-450. Do common genetic variants in endotoxin signaling pathway contribute to predisposition to | 1.1 | 14 |
| ⁵⁹ Do common genetic variants in endotoxin signaling pathway contribute to predisposition to alcoholic liver cirrhosis?. Clinical Chemistry and Laboratory Medicine, 2009, 47, 398-404. | у 1.7 | 31 |
| | 1.4 | 10 |
| 60 N -acetyl cysteine averted liver transplantation in a patient with liver failure caused by erythropoprotoporphyria. Liver Transplantation, 2009, 15, 352-354. | pietic 1.3 | 7 |
| 61 Wilson disease as a cause of liver injury in cystic fibrosis. Journal of Cystic Fibrosis, 2009, 8, 63-6 | 55. 0.3 | 3 |
| Novel ABCB11 mutations in a Thai infant with progressive familial intrahepatic cholestasis. Wor Journal of Gastroenterology, 2009, 15, 4339. | ld 1.4 | 13 |
| ⁶³ 1H MR spectroscopy as a diagnostic tool for cerebral creatine deficiency. Magnetic Resonance Materials in Physics, Biology, and Medicine, 2008, 21, 327-332. | 1.1 | 20 |
| Highly sensitive method for quantitative determination of bilirubin in biological fluids and tissue Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2008 37-42. | | 67 |
| 65 Severe Bile Salt Export Pump Deficiency: 82 Different ABCB11 Mutations in 109 Families. Gastroenterology, 2008, 134, 1203-1214.e8. | 0.6 | 331 |
| 66 CYP7A1 promoter polymorphism â^203A>C affects bile salt synthesis rate in patients after il resection. Journal of Lipid Research, 2008, 49, 2664-2667. | eal 2.0 | 15 |
| Revised King's College score for liver transplantation in adult patients with Wilson's disease. Liv Transplantation, 2007, 13, 55-61. | er 1.3 | 51 |
| Rotor-type hyperbilirubinaemia has no defect in the canalicular bilirubin export pump. Liver International, 2007, 27, 485-491. | 1.9 | 22 |
| Linkage between A(TA)7TAA and ?3279T>G mutations in UGT1A1 is not essential for pathogene Gilbert syndrome. Liver International, 2006, 26, 1302-1303. | esis of 1.9 | 8 |
| 70 Dual Hereditary Jaundice: Simultaneous Occurrence of Mutations Causing Gilbert's and Dubin-Johnson Syndrome. Gastroenterology, 2005, 129, 315-320. | 0.6 | 33 |
| ⁷¹ Indel in the FIC1/ATP8B1 gene?a novel rare type of mutation associated with benign recurrent intrahepatic cholestasis. Hepatology Research, 2004, 30, 1-3. | 1.8 | 1 |
| 72 Effect of acute hyperglycaemia on selected plasma and urinary cytokine antagonists in Type 1 d mellitus. Diabetologia, 2003, 46, 470-474. | iabetes 2.9 | 3 |

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|----|--|-----|-----------|
| 73 | Relation between hepatic expression of ATP-binding cassette transporters G5 and G8 and biliary cholesterol secretion in mice. Journal of Hepatology, 2003, 38, 710-716. | 1.8 | 78 |
| 74 | Genetic background of cholesterol gallstone disease. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2003, 1637, 1-19. | 1.8 | 47 |
| 75 | Gilbert syndrome and ischemic heart disease: a protective effect of elevated bilirubin levels. Atherosclerosis, 2002, 160, 449-456. | 0.4 | 383 |
| 76 | Role of biliary proteins and non-protein factors in kinetics of cholesterol crystallisation and gallstone growth. Frontiers in Bioscience - Landmark, 2001, 6, e154. | 3.0 | 18 |
| 77 | Carcinoembryonic antigen-related cell adhesion molecule 1 is the 85-kilodalton pronase-resistant biliary glycoprotein in the cholesterol crystallization promoting low density protein-lipid complex. Hepatology, 2001, 34, 1075-1082. | 3.6 | 9 |
| 78 | Soluble Adhesion Molecules and Cytokines in Patients with Myasthenia Gravis Treated by Plasma Exchange. Blood Purification, 2000, 18, 115-120. | 0.9 | 22 |
| 79 | LONG TERM TREATMENT OF IgA NEPHROPATHY WITH CYCLOSPORINE A. Renal Failure, 2000, 22, 55-62. | 0.8 | 7 |
| 80 | Intestinal Colonization Leading to Fecal Urobilinoid Excretion May Play a Role in the Pathogenesis of Neonatal Jaundice. Journal of Pediatric Gastroenterology and Nutrition, 2000, 30, 294-298. | 0.9 | 45 |
| 81 | Influence of Plasma Exchange on Serum Levels of Cytokines and Adhesion Molecules in ANCA-Positive Renal Vasculitis. Blood Purification, 1998, 16, 72-80. | 0.9 | 56 |
| 82 | Long-term treatment of IgA nephropathy with cyclosporin A-a preliminary report. Nephrology Dialysis Transplantation, 1997, 12, 2206-2207. | 0.4 | 10 |
| 83 | Bacterial oxidation of cholic acid byArthrobacter sp. Folia Microbiologica, 1995, 40, 621-626. | 1.1 | 1 |
| 84 | Hyperbaric oxygen and photodynamic therapy in tumour-bearing nude mice. European Journal of Cancer & Clinical Oncology, 1991, 27, 109. | 0.9 | 41 |
| 85 | Influence of hyperbaric oxygenation on bilirubin and ditaurobilirubin auto-oxidation and porphyrin-sensitized photo-oxidation. Journal of Photochemistry and Photobiology B: Biology, 1990, 5, 295-302. | 1.7 | 5 |
| 86 | The process of continuous production and isolation of coproporphyrin. Biotechnology Letters, 1988, 10, 273-276. | 1.1 | 0 |
| 87 | Solid phase extraction and isocratic separation of urinary porphyrins by HPLC. Biomedical Chromatography, 1986, 1, 159-162. | 0.8 | 12 |