David Cassiman

List of Publications by Year
in descending order

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217 papers	7,442 citations	50244 46 h-index	69214 77 g-index
228 all docs	228 docs citations	228 times ranked	10637 citing authors

#	Article	IF	Citations
1	Expert consensus statement on acute hepatic porphyria in Belgium. Acta Clinica Belgica, 2022, 77, 735-741.	0.5	5
2	Biomarkers in Nephropathic Cystinosis: Current and Future Perspectives. Cells, 2022, 11, 1839.	1.8	2
3	Ornithine transcarbamylase deficiency: A diagnostic odyssey. Journal of Inherited Metabolic Disease, 2022, 45, 661-662.	1.7	1
4	Pyruvate and uridine rescue the metabolic profile of OXPHOS dysfunction. Molecular Metabolism, 2022, 63, 101537.	3.0	9
5	De novo loss-of-function variants in X-linked MED12 are associated with Hardikar syndrome in females. Genetics in Medicine, 2021, 23, 637-644.	1.1	16
6	An autosomal dominant neurological disorder caused by de novo variants in FAR1 resulting in uncontrolled synthesis of ether lipids. Genetics in Medicine, 2021, 23, 740-750.	1.1	25
7	Transcriptomic analysis of CFTR-impaired endothelial cells reveals a pro-inflammatory phenotype. European Respiratory Journal, 2021, 57, 2000261.	3.1	10
8	Galactokinase deficiency: lessons from the GalNet registry. Genetics in Medicine, 2021, 23, 202-210.	1.1	14
9	D-galactose supplementation in individuals with PMM2-CDG: results of a multicenter, open label, prospective pilot clinical trial. Orphanet Journal of Rare Diseases, 2021, 16, 138.	1.2	6
10	Relationship between de novo lipogenesis and serum sex hormone binding globulin in humans. Clinical Endocrinology, 2021, 95, 101-106.	1.2	11
11	Liver-Related and Cardiovascular Outcome of Patients Transplanted for Nonalcoholic Fatty Liver Disease: A European Single-Center Study. Transplantation Proceedings, 2021, 53, 1674-1681.	0.3	5
12	Estimating the broader fiscal consequences of acute hepatic porphyria (AHP) with recurrent attacks in Belgium using a public economic analytic framework. Orphanet Journal of Rare Diseases, 2021, 16, 346.	1.2	2
13	Donor Hepatectomy and Implantation Time Are Associated With Early Complications After Liver Transplantation: A Single-center Retrospective Study. Transplantation, 2021, 105, 1030-1038.	0.5	23
14	Repurposing the Antidepressant Sertraline as SHMT Inhibitor to Suppress Serine/Glycine Synthesisâ€“Addicted Breast Tumor Growth. Molecular Cancer Therapeutics, 2021, 20, 50-63.	1.9	31
15	Genotype-Phenotype Correlations in PMM2-CDG. Genes, 2021, 12, 1658.	1.0	6
16	Sorbitol Is a Severity Biomarker for <scp>PMM2â€CDG</scp> with Therapeutic Implications. Annals of Neurology, 2021, 90, 887-900.	2.8	22
17	Patents vs patients 1â€0: The case of chenodeoxycholic acid. Journal of Inherited Metabolic Disease, 2021, , .	1.7	0
18	A Patient with neonatal cholestasis. Medycyna Wieku Rozwojowego, 2021, 24, 31-33.	0.2	1

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19	Hypophosphatasia in Adults: Clinical Spectrum and Its Association With Genetics and Metabolic Substrates. Journal of Clinical Densitometry, 2020, 23, 340-348.	0.5	20
20	Obstructive sleep apnea in Hutchinson-Gilford progeria. Sleep Medicine, 2020, 66, 21-23.	0.8	0
21	Fostering practice-oriented and use-inspired science in biomedical research. Research Policy, 2020, 49, 103900.	3.3	8
22	Dietary practices in methylmalonic acidaemia: a European survey. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 147-155.	0.4	8
23	Normal liver stiffness and influencing factors in healthy children: An individual participant data metaâ€analysis. Liver International, 2020, 40, 2602-2611.	1.9	24
24	Fulminant Wilson Disease in Children. Journal of Pediatric Gastroenterology and Nutrition, 2020, 71, 720-725.	0.9	9
25	Kidney and vascular function in adult patients with hereditary fructose intolerance. Molecular Genetics and Metabolism Reports, 2020, 23, 100600.	0.4	7
26	Amino acid levels determine metabolism and CYP450 function of hepatocytes and hepatoma cell lines. Nature Communications, 2020, 11, 1393.	5.8	79
27	SGLT2 Inhibitors for Treatment of Refractory Hypomagnesemia: A Case Report of 3 Patients. Kidney Medicine, 2020, 2, 359-364.	1.0	29
28	Two cases of non-alcoholic fatty liver disease caused by biallelic ABHD5 mutations. Journal of Hepatology, 2020, 72, 1030-1032.	1.8	6
29	m.3243A > G-Induced Mitochondrial Dysfunction Impairs Human Neuronal Development and Reduces Neuronal Network Activity and Synchronicity. Cell Reports, 2020, 31, 107538.	2.9	56
30	Dietary plant stanol ester supplementation reduces peripheral symptoms in a mouse model of Niemann-Pick type C1 disease. Journal of Lipid Research, 2020, 61, 830-839.	2.0	5
31	The Role of Brown Adipose Tissue in the Development and Treatment of Nonalcoholic Steatohepatitis: An Exploratory Gene Expression Study in Mice. Hormone and Metabolic Research, 2020, 52, 869-876.	0.7	2
32	Abstract 1789: Repurposing the anti-depressant sertraline to target serine/glycine synthesis addicted cancer. , 2020, , .		0
33	The ribosomal RPL10 R98S mutation drives IRES-dependent BCL-2 translation in T-ALL. Leukemia, 2019, 33, 319-332.	3.3	50
34	LBP-36-Inhibition of glutamine synthetase in monocytes from patients with Acute-on-Chronic Liver Failure resuscitates their antibacterial and inflammatory capacity. Journal of Hepatology, 2019, 70, e159.	1.8	1
35	Oxygraphy Versus Enzymology for the Biochemical Diagnosis of Primary Mitochondrial Disease. Metabolites, 2019, 9, 220.	1.3	6
36	Patients With Aldolase B Deficiency Are Characterized by Increased Intrahepatic Triglyceride Content. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5056-5064.	1.8	30

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37	Translatome analysis reveals altered serine and glycine metabolism in T-cell acute lymphoblastic leukemia cells. Nature Communications, 2019, 10, 2542.	5.8	43
38	Case Report of Gastrointestinal Bleeding in an Adult with Chronic Visceral Acid Sphingomyelinase Deficiency. Case Reports in Gastrointestinal Medicine, 2019, 2019, 1-5.	0.2	0
39	The Metabolic Map into the Pathomechanism and Treatment of PGM1-CDG. American Journal of Human Genetics, 2019, 104, 835-846.	2.6	59
40	Clinical and biochemical footprints of inherited metabolic diseases. II. Metabolic liver diseases. Molecular Genetics and Metabolism, 2019, 127, 117-121.	0.5	32
41	Evidence for an alternative fatty acid desaturation pathway increasing cancer plasticity. Nature, 2019, 566, 403-406.	13.7	326
42	International clinical guidelines for the management of phosphomannomutase 2â€congenital disorders of glycosylation: Diagnosis, treatment and follow up. Journal of Inherited Metabolic Disease, 2019, 42, 5-28.	1.7	91
43	Age Matching of Elderly Liver Grafts With Elderly Recipients Does Not Have a Synergistic Effect on Long-term Outcomes When Both Are Carefully Selected. Transplantation Direct, 2019, 5, e342.	0.8	10
44	Emotional and behavioral problems, quality of life and metabolic control in NTBC-treated Tyrosinemia type 1 patients. Orphanet Journal of Rare Diseases, 2019, 14, 285.	1.2	19
45	Inhibition of glutamine synthetase in monocytes from patients with acute-on-chronic liver failure resuscitates their antibacterial and inflammatory capacity. Gut, 2019, 68, 1872-1883.	6.1	60
46	Measuring Rates of ATP Synthesis. Methods in Molecular Biology, 2019, 1862, 97-107.	0.4	0
47	Long-term follow-up in PMM2-CDG: are we ready to start treatment trials?. Genetics in Medicine, 2019, 21, 1181-1188.	1.1	36
48	Long-term outcome of transjugular intrahepatic portosystemic shunt for portal hypertension in autosomal recessive polycystic kidney disease. Digestive and Liver Disease, 2018, 50, 707-712.	0.4	15
49	Response by Kusters et al to Letter Regarding Article, â€œEffect of Rosuvastatin on Carotid Intima-Media Thickness in Children With Heterozygous Familial Hypercholesterolemia: The CHARON Study (Hypercholesterolemia in Children and Adolescents Taking Rosuvastatin Open Label)â€. Circulation, 2018, 137, 641-642.	1.6	1
50	Unusual yellow scaly colonic mucosal appearance: Tangier disease. Gastrointestinal Endoscopy, 2018, 88, 193-195.	0.5	1
51	Multiple Solid Organ Transplantation in Telomeropathy: Case Series and Literature Review. Transplantation, 2018, 102, 1747-1755.	0.5	17
52	Ethyl Glucuronide in Hair Is an Accurate Biomarker of Chronic Excessive Alcohol Use in Patients With Alcoholic Cirrhosis. Clinical Gastroenterology and Hepatology, 2018, 16, 454-456.	2.4	14
53	Pre-operative trans-catheter arterial chemo-embolization increases hepatic artery thrombosis after liver transplantation - a retrospective study. Transplant International, 2018, 31, 71-81.	0.8	11
54	Renal involvement in PMM2-CDG, a mini-review. Molecular Genetics and Metabolism, 2018, 123, 292-296.	0.5	19

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55	Ageing in Liver Transplantation. Transplantation, 2018, 102, S125.	0.5	0
56	Cobalamin C Deficiency Induces a Typical Histopathological Pattern of Renal Arteriolar and Glomerular Thrombotic Microangiopathy. Kidney International Reports, 2018, 3, 1153-1162.	0.4	28
57	The SLC40A1 R178Q mutation is a recurrent cause of hemochromatosis and is associated with a novel pathogenic mechanism. Haematologica, 2018, 103, 1796-1805.	1.7	19
58	Liver transplantation for very severe hepatopulmonary syndrome due to vitamin A-induced chronic liver disease in a patient with Shwachman-Diamond syndrome. Orphanet Journal of Rare Diseases, 2018, 13, 69.	1.2	3
59	The outcome of acute-on-chronic liver failure in the intensive care is similar to a propensity matched ICU population without liver disease. Journal of Hepatology, 2018, 68, S239.	1.8	0
60	The paracrine effect of visceraladipose tissue obtained at bariatric surgery on primary human hepatic stellatecells grown in human 3D healthy liver scaffolds. Journal of Hepatology, 2018, 68, S334-S335.	1.8	0
61	Pneumococcal Immunization Reduces Neurological and Hepatic Symptoms in a Mouse Model for Niemann-Pick Type C1 Disease. Frontiers in Immunology, 2018, 9, 3089.	2.2	8
62	Liver failure after long-limb gastric bypass. Clinics and Research in Hepatology and Gastroenterology, 2017, 41, e32-e37.	0.7	14
63	Dietary intervention, but not losartan, completely reverses non-alcoholic steatohepatitis in obese and insulin resistant mice. Lipids in Health and Disease, 2017, 16, 46.	1.2	19
64	On the Pathogenesis of Central Liver Nodules in Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2017, 64, e80.	0.9	1
65	Frequency and pathogenesis of central liver nodules in Alagille syndrome patients. Pediatric Radiology, 2017, 47, 1023-1024.	1.1	1
66	Proton Pump Inhibitors Decrease Phlebotomy Need in HFE Hemochromatosis: Double-Blind Randomized Placebo-Controlled Trial. Gastroenterology, 2017, 153, 678-680.e2.	0.6	29
67	Effect of Rosuvastatin on Carotid Intima-Media Thickness in Children With Heterozygous Familial Hypercholesterolemia. Circulation, 2017, 136, 359-366.	1.6	84
68	Liver disease in cystic fibrosis presents as non-cirrhotic portal hypertension. Journal of Cystic Fibrosis, 2017, 16, e11-e13.	0.3	48
69	NTCP deficiency and persistently raised bile salts: an adult case. Journal of Inherited Metabolic Disease, 2017, 40, 313-315.	1.7	27
70	Monocytes exhibit an immune and metabolic reprogramming during acute-on-chronic-liver-failure. Journal of Hepatology, 2017, 66, S100.	1.8	0
71	Galactose Supplementation in Patients With TMEM165-CDG Rescues the Glycosylation Defects. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1375-1386.	1.8	61
72	Dual loss of succinate dehydrogenase (SDH) and complex I activity is necessary to recapitulate the metabolic phenotype of SDH mutant tumors. Metabolic Engineering, 2017, 43, 187-197.	3.6	64

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73	Identification of survival-promoting OSIP108 peptide variants and their internalization in human cells. Mechanisms of Ageing and Development, 2017, 161, 247-254.	2.2	0
74	Usefulness of the single-operator cholangioscopy system SpyGlass in biliary disease: a single-center prospective cohort study and aggregated review. Surgical Endoscopy and Other Interventional Techniques, 2017, 31, 2223-2232.	1.3	73
75	Myelodysplasia and liver disease extend the spectrum of RTEL1 related telomeropathies. Haematologica, 2017, 102, e293-e296.	1.7	15
76	Nutritional Therapies in Congenital Disorders of Glycosylation (CDG). Nutrients, 2017, 9, 1222.	1.7	41
77	Fertility in adult women with classic galactosemia and primary ovarian insufficiency. Fertility and Sterility, 2017, 108, 168-174.	0.5	42
78	Pro-Inflammatory Cytokines but Not Endotoxin-Related Parameters Associate with Disease Severity in Patients with NAFLD. PLoS ONE, 2016, 11, e0166048.	1.1	52
79	<scp>ADAMTS</scp>5 deficiency protects against nonâ€alcoholic steatohepatitis in obesity. Liver International, 2016, 36, 1848-1859.	1.9	18
80	Biallelic Mutations in TMEM126B Cause Severe Complex I Deficiency with a Variable Clinical Phenotype. American Journal of Human Genetics, 2016, 99, 217-227.	2.6	57
81	Development of a Representative Mouse Model with Nonalcoholic Steatohepatitis. Current Protocols in Mouse Biology, 2016, 6, 201-210.	1.2	8
82	An Overview of Mouse Models of Nonalcoholic Steatohepatitis: From Past to Present. Current Protocols in Mouse Biology, 2016, 6, 185-200.	1.2	40
83	Off-label use of orphan medicinal products: a Belgian qualitative study. Orphanet Journal of Rare Diseases, 2016, 11, 144.	1.2	12
84	De novo Malignancy and Recurrent Alcoholic Cirrhosis Account for 70% of Deaths in Patients Transplanted for End-Stage Alcoholic Liver Disease. American Journal of Gastroenterology, 2016, 111, 436-437.	0.2	5
85	Neurocognitive outcome in tyrosinemia type 1 patients compared to healthy controls. Orphanet Journal of Rare Diseases, 2016, 11, 87.	1.2	60
86	Key-interventions derived from three evidence based guidelines for management and follow-up of patients with HFE haemochromatosis. BMC Health Services Research, 2016, 16, 573.	0.9	6
87	The orphan drug pipeline in Europe. Nature Reviews Drug Discovery, 2016, 15, 376-376.	21.5	18
88	Cause of death in patients with chronic visceral and chronic neurovisceral acid sphingomyelinase deficiency (Niemann-Pick disease type B and B variant): Literature review and report of new cases. Molecular Genetics and Metabolism, 2016, 118, 206-213.	0.5	72
89	Investigating Rare Haematological Disorders â€“ A Celebration of 10 Years of the Sherlock Holmes Symposia. European Oncology and Haematology, 2016, 12, 55.	0.0	0
90	Synergistic Activity of the Plant Defensin HsAFP1 and Caspofungin against Candida albicans Biofilms and Planktonic Cultures. PLoS ONE, 2015, 10, e0132701.	1.1	67

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91	Association of Adipose Tissue Inflammation With Histologic Severity of Nonalcoholic Fatty Liver Disease. Gastroenterology, 2015, 149, 635-648.e14.	0.6	249
92	Bone demineralisation in a large cohort of Wilson disease patients. Journal of Inherited Metabolic Disease, 2015, 38, 949-956.	1.7	22
93	Efficacy and safety of rosuvastatin therapy inÂchildren and adolescents with familial hypercholesterolemia: Results from the CHARONÂstudy. Journal of Clinical Lipidology, 2015, 9, 741-750.	0.6	42
94	Management dilemmas in pediatric nephrology: Cystinosis. Pediatric Nephrology, 2015, 30, 1349-1360.	0.9	15
95	Hepatobiliary malignancies in Wilson disease. Liver International, 2015, 35, 1615-1622.	1.9	78
96	Liver Fibrosis Associated with Iron Accumulation Due to Long-Term Heme-Arginate Treatment in Acute Intermittent Porphyria: A Case Series. JIMD Reports, 2015, 25, 77-81.	0.7	39
97	The quality of hereditary haemochromatosis guidelines: A comparative analysis. Clinics and Research in Hepatology and Gastroenterology, 2015, 39, 205-214.	0.7	11
98	Roux-en-y gastric bypass attenuates hepatic mitochondrial dysfunction in mice with non-alcoholic steatohepatitis. Gut, 2015, 64, 673-683.	6.1	64
99	The Plant Decapeptide OSIP108 Can Alleviate Mitochondrial Dysfunction Induced by Cisplatin in Human Cells. Molecules, 2014, 19, 15088-15102.	1.7	4
100	Clinical, Biochemical, and Molecular Characterization of Novel Mutations in ABCA1 in Families with Tangier Disease. JIMD Reports, 2014, 18, 51-62.	0.7	19
101	Reimbursement of orphan drugs in Belgium: what (else) matters?. Orphanet Journal of Rare Diseases, 2014, 9, 139.	1.2	20
102	The plant decapeptide OSIP108 prevents copper-induced apoptosis in yeast and human cells. Biochimica Et Biophysica Acta - Molecular Cell Research, 2014, 1843, 1207-1215.	1.9	22
103	The plant decapeptide OSIP108 prevents copper-induced toxicity in various models for Wilson disease. Toxicology and Applied Pharmacology, 2014, 280, 345-351.	1.3	14
104	Defining the phenotype and diagnostic considerations in adults with congenital disorders of N-linked glycosylation. Expert Review of Molecular Diagnostics, 2014, 14, 217-224.	1.5	17
105	Shining a light in the black box of orphan drug pricing. Orphanet Journal of Rare Diseases, 2014, 9, 62.	1.2	46
106	Hepatitis With Brown Pigment in the Liver. Gastroenterology, 2014, 147, e5-e6.	0.6	0
107	Efficacy and safety of rosuvastatin in children aged 6â€“17 years with familial hypercholesterolemia: Findings from the charon study. Atherosclerosis, 2014, 235, e34.	0.4	2
108	Effect of rosuvastatin therapy on carotid intima media thickness in children with familial hypercholesterolemia: Findings from the charon study. Atherosclerosis, 2014, 235, e18-e19.	0.4	5

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109	Angiotensin II type 1 receptor blockers increase tolerance of cells to copper and cisplatin. Microbial Cell, 2014, 1, 352-364.	1.4	2
110	Endoscopic resection of ampullary lesions: a single-center 8-year retrospective cohort study of 91 patients with long-term follow-up. Surgical Endoscopy and Other Interventional Techniques, 2013, 27, 3865-3876.	1.3	56
111	Cost-Effectiveness Assessment of Orphan Drugs. Applied Health Economics and Health Policy, 2013, 11, 1-3.	1.0	60
112	Design and baseline data of a pediatric study with rosuvastatin in familial hypercholesterolemia. Journal of Clinical Lipidology, 2013, 7, 408-413.	0.6	15
113	Clinical evidence for orphan medicinal products-a cause for concern?. Orphanet Journal of Rare Diseases, 2013, 8, 164.	1.2	26
114	Development and validation of COMPASS: clinical evidence of orphan medicinal products â€“ an assessment tool. Orphanet Journal of Rare Diseases, 2013, 8, 157.	1.2	4
115	Pivotal Studies of Orphan Medicinal Products â€“ an Analysis of Quality of Clinical Evidence. Value in Health, 2013, 16, A390.	0.1	0
116	79 THE OUTCOME OF SHUNT REDUCTION AFTER TIPS BY THE PARALLEL TECHNIQUE: A PROSPECTIVE STUDY. Journal of Hepatology, 2013, 58, S35.	1.8	0
117	Lung transplantation in cystic fibrosis normalizes essential fatty acid profiles. Journal of Cystic Fibrosis, 2013, 12, 222-228.	0.3	14
118	Misdiagnosis as asphyxiating thoracic dystrophy and CMV-associated haemophagocytic lymphohistiocytosis in Shwachman-Diamond syndrome. European Journal of Pediatrics, 2013, 172, 613-622.	1.3	16
119	Heterozygous Î±1-antitrypsin Z allele mutation in presumed healthy donor livers used for transplantation. European Journal of Gastroenterology and Hepatology, 2013, 25, 1335-1339.	0.8	16
120	Focal portal vein stenosis in an adolescent potentially related to complicated umbilical catheter insertion in the neonatal period. Acta Radiologica Short Reports, 2013, 2, 204798161349272.	0.7	0
121	Mitochondrial hepatopathy in adults. European Journal of Gastroenterology and Hepatology, 2013, 25, 892-898.	0.8	6
122	Treatment of non-alcoholic fatty liver disease: can we already face the epidemic?. Acta Gastro-Enterologica Belgica, 2013, 76, 200-9.	0.4	5
123	High-frequency vagus nerve stimulation improves portal hypertension in cirrhotic rats. Gut, 2012, 61, 604-612.	6.1	22
124	Evaluation of the interference by homogentisic acid and other organic acids on the enzymatic and JaffÃ© method creatinine assay. Clinical Chemistry and Laboratory Medicine, 2012, 50, 749-50.	1.4	4
125	Transcatheter arterial embolization for iatrogenic hemobilia is a safe and effective procedure. European Journal of Gastroenterology and Hepatology, 2012, 24, 905-909.	0.8	29
126	Acid sphingomyelinase (Asm) deficiency patients in The Netherlands and Belgium: Disease spectrum and natural course in attenuated patients. Molecular Genetics and Metabolism, 2012, 107, 526-533.	0.5	71

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127	1395 THE REDUCTION IN LIVER VOLUME IN POLYCYSTIC LIVER DISEASE WITH LANREOTIDE IS DOSE DEPENDENT AND IS MOST PRONOUNCED IN PATIENTS WITH THE HIGHEST LIVER VOLUME. Journal of Hepatology, 2012, 56, S547.	1.8	4
128	Cholestasis-induced pruritus treated with ultraviolet B phototherapy: An observational case series study. Journal of Hepatology, 2012, 57, 637-641.	1.8	50
129	Orphan Drugs for Rare Diseases. Drugs, 2012, 72, 1437-1443.	4.9	50
130	Evaluating and improving orphan drug regulations in Europe: A Delphi policy study. Health Policy, 2012, 108, 1-9.	1.4	18
131	Liver Transplantation in a Patient With an Intraabdominally Located Left Ventricular Assist Device: Surgical Aspectsâ€"Case Report. Transplantation Proceedings, 2012, 44, 2885-2887.	0.3	1
132	Septuagenarian and Octogenarian Donors Provide Excellent Liver Grafts for Transplantation. Transplantation Proceedings, 2012, 44, 2861-2867.	0.3	47
133	Outcomes of Liver Transplantations Using Donations After Circulatory Death: A Single-Center Experience. Transplantation Proceedings, 2012, 44, 2868-2873.	0.3	49
134	Risk Factors for Bleeding and Clinical Implications in Patients Undergoing Liver Transplantation. Transplantation Proceedings, 2012, 44, 2857-2860.	0.3	43
135	PHP15 What Price do we Pay for Repurposing Medicines for Rare Diseases?. Value in Health, 2012, 15, A15-A16.	0.1	3
136	Orphan Drugs for Rare Diseases: Grounds for Special Status. Drug Development Research, 2012, 73, 115-119.	1.4	14
137	HNF1Bdeficiency causes ciliary defects in human cholangiocytes. Hepatology, 2012, 56, 1178-1181.	3.6	26
138	Market uptake of orphan drugs - a European analysis. Journal of Clinical Pharmacy and Therapeutics, 2012, 37, 664-667.	0.7	29
139	Histology obtained by needle biopsy gives additional information on the prognosis of hepatocellular carcinoma. Hepatology Research, 2012, 42, 990-998.	1.8	17
140	Drugs for Rare Diseases: Influence of Orphan Designation Status on Price. Applied Health Economics and Health Policy, 2011, 9, 275-279.	1.0	40
141	Are some orphan drugs for rare diseases too expensive? A study of purchase versus compounding costs. Drugs and Therapy Perspectives, 2011, 27, 24-26.	0.3	11
142	Prospective Study Comparing Different Indirect Methods to Measure Portal Pressure. Journal of Vascular and Interventional Radiology, 2011, 22, 1553-1558.	0.2	38
143	1312 ABNORMAL PRIMARY CILIA IN CHOLANGIOCYTES CAUSE CHOLESTATIC LIVER DISEASE IN HNFÎ² DELETION IN HUMANS. Journal of Hepatology, 2011, 54, S517.	1.8	0
144	Acute-on-chronic liver failure: current concepts on definition, pathogenesis, clinical manifestations and potential therapeutic interventions. Expert Review of Gastroenterology and Hepatology, 2011, 5, 523-537.	1.4	80

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145	PHP28 Drugs for Rare Diseases: Influence of Oprhan Designation Status on Price. Value in Health, 2011, 14, A338.	0.1	1
146	PHP46 Market Uptake of Orphan Drugs â€“ A European Analysis. Value in Health, 2011, 14, A341.	0.1	0
147	1289 CHOLESTASIS-ASSOCIATED PRURITUS TREATED WITH UVB PHOTOTHERAPY: REPORT OF 13 CASES. Journal of Hepatology, 2011, 54, S508.	1.8	0
148	Acute Liver Failure Secondary to Khat (Catha edulis)â€“Induced Necrotic Hepatitis Requiring Liver Transplantation: Case Report. Transplantation Proceedings, 2011, 43, 3493-3495.	0.3	18
149	178 A PROSPECTIVE STUDY COMPARING THE DIFFERENT INDIRECT METHODS TO MEASURE PORTAL PRESSURE. Journal of Hepatology, 2011, 54, S76-S77.	1.8	0
150	PAS-positive macrophagesâ€"not always infection. Lancet, The, 2011, 377, 1890.	6.3	8
151	Giant liver hemangioma. European Journal of Gastroenterology and Hepatology, 2011, 23, 438-443.	0.8	31
152	Systematic review: the pathophysiology and management of polycystic liver disease. Alimentary Pharmacology and Therapeutics, 2011, 34, 702-713.	1.9	79
153	Left ventricular assist device as bridge to liver transplantation in a patient with propionic acidemia and cardiogenic shock. Journal of Pediatrics, 2011, 158, 866-867.	0.9	19
154	Noncirrhotic presinusoidal portal hypertension is common in cystic fibrosis-associated liver disease. Hepatology, 2011, 53, 1064-1065.	3.6	44
155	Gene Transfer for Inborn Errors of Metabolism of the Liver: The Clinical Perspective. Current Pharmaceutical Design, 2011, 17, 2550-2557.	0.9	3
156	Wilson's disease: long-term follow-up of a cohort of 24 patients treated with D-penicillamine. European Journal of Gastroenterology and Hepatology, 2010, 22, 564-571.	0.8	30
157	Alpers syndrome presenting with anatomopathological features of fulminant autoimmune hepatitis. Journal of Inherited Metabolic Disease, 2010, 33, 451-451.	1.7	3
158	Lysosomal lipid vacuoles in macrophages located in the colon. Journal of Inherited Metabolic Disease, 2010, 33, 303-304.	1.7	1
159	ADP-degrading enzymes inhibit platelet activation in bile duct-ligated rats. Journal of Thrombosis and Haemostasis, 2010, 8, 360-368.	1.9	7
160	The hepatic vagus nerve stimulates hepatic stellate cell proliferation in rat acute hepatitis via muscarinic receptor type 2. Liver International, 2010, 30, 693-702.	1.9	5
161	A Seven-Gene Set Associated with Chronic Hypoxia of Prognostic Importance in Hepatocellular Carcinoma. Clinical Cancer Research, 2010, 16, 4278-4288.	3.2	56
162	An adult male patient with multiple adenomas and a hepatocellular carcinoma: Mild Glycogen Storage Disease type Ia. Journal of Hepatology, 2010, 53, 213-217.	1.8	40

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163	Outcomes of Long-Term Administration of Intravenous Hepatitis B Immunoglobulins for the Prevention of Recurrent Hepatitis B After Liver Transplantation. Transplantation Proceedings, 2010, 42, 4399-4402.	0.3	8
164	120 A 7 GENE SET ASSOCIATED WITH CHRONIC HYPOXIA OF UNIVERSAL PROGNOSTIC IMPORTANCE IN HEPATOCELLULAR CARCINOMA. Journal of Hepatology, 2010, 52, S53.	1.8	0
165	521 THE VASOPRESSIN-2-RECEPTOR ANTAGONIST SATAVAPTAN IMPROVES PORTAL HYPERTENSION IN THE NON-ASCITIC THIOACETAMIDE CIRRHOTIC RAT WITHIN A NARROW SAFE AND EFFECTIVE DOSE-RANGE. Journal of Hepatology, 2010, 52, S209.	1.8	0
166	858 ADP-DEGRADING ENZYMES INHIBIT PLATELET ACTIVATION IN BILE-DUCT LIGATED RATS. Journal of Hepatology, 2010, 52, S334.	1.8	0
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168	Successful Conversion From Mycophenolate Mofetil to Enteric-Coated Mycophenolate Sodium (Myfortic) in Liver Transplant Patients With Gastrointestinal Side Effects. Transplantation Proceedings, 2009, 41, 610-613.	0.3	15
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