Aaron S Kesselheim

List of Publications by Year
in descending order

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568 papers	15,396 citations	25423 59 h-index	43601 95 g-index
581 all docs	581 docs citations	581 times ranked	16609 citing authors

#	Article	IF	Citations
1	US Taxpayers Heavily Funded the Discovery of COVIDâ€19 Vaccines. Clinical Pharmacology and Therapeutics, 2022, 111, 542-544.	2.3	14
2	Payer-Specific Negotiated Prices for Prescription Drugs at Top-Performing US Hospitals. JAMA Internal Medicine, 2022, 182, 83.	2.6	7
3	Indication-Specific Generic Uptake of Imatinib Demonstrates the Impact of Skinny Labeling. Journal of Clinical Oncology, 2022, 40, 1102-1110.	0.8	3
4	Reporting bias in clinical trials: Progress toward transparency and next steps. PLoS Medicine, 2022, 19, e1003894.	3.9	9
5	Differences in Diabetic Prescription Drug Utilization and Costs Among Patients With Diabetes Enrolled in Colorado Marketplace and Medicaid Plans, 2014-2015. JAMA Network Open, 2022, 5, e2140371.	2.8	2
6	Medicaid Spending on Drugs Marketed Without US Food and Drug Administration Approval in 2020. JAMA Internal Medicine, 2022, 182, 342.	2.6	4
7	Extending Drug Monopolies by Patenting Safe Drug Use. JAMA Internal Medicine, 2022, , .	2.6	2
8	Competition law and pricing among biologic drugs: the case of VEGF therapy for retinal diseases. Journal of Law and the Biosciences, 2022, 9, lsac001.	0.8	5
9	The characteristics of patents impacting availability of biosimilars. Nature Biotechnology, 2022, 40, 22-25.	9.4	9
10	Performance-Linked Reimbursement and the Uncertainty of Novel Drugs. Circulation: Cardiovascular Quality and Outcomes, 2022, 15, e008642.	0.9	1
11	Patient and Caregiver Experiences With and Perceptions of Risk Evaluation and Mitigation Strategy Programs With Elements to Assure Safe Use. JAMA Network Open, 2022, 5, e2144386.	2.8	7
12	Characteristics of Clinical Trials Evaluating Biosimilars in the Treatment of Cancer. JAMA Oncology, 2022, 8, 537.	3.4	11
13	Expertsâ€™ Views on FDA Regulatory Standards for Drug and High-Risk Medical Devices: Implications for Patient Care. Journal of General Internal Medicine, 2022, , 1.	1.3	1
14	A New Way to Contain Unaffordable Medication Costs â€" Exercising the Governmentâ€™s Existing Rights. New England Journal of Medicine, 2022, 386, 1104-1106.	13.9	5
15	New Drug Postmarketing Requirements and Commitments in the US: A Systematic Review of the Evidence. Drug Safety, 2022, 45, 305-318.	1.4	7
16	Recent Orange and Purple Book legislation suggests a need to bridge drug and biologic patent regimes. Nature Biotechnology, 2022, 40, 167-169.	9.4	1
17	Aducanumab and Accelerated Approval: Where Do We Go From Here?. Clinical Pharmacology and Therapeutics, 2022, 111, 726-727.	2.3	2
18	Anticipated efficiencies, real costs: Medicaid managed care organizations and the pharmacy benefit. Journal of Managed Care & Specialty Pharmacy, 2022, 28, 354-361.	0.5	0

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19	Strategies to Manage Drugs and Devices Approved Based on Limited Evidence: Results of a Modified Delphi Panel. Clinical Pharmacology and Therapeutics, 2022, 111, 1307-1314.	2.3	2
20	Updating the Bayh-Dole Act. JAMA - Journal of the American Medical Association, 2022, 327, 923.	3.8	4
21	Switching to Over-the-Counter Availability of Rescue Inhalers for Asthma. JAMA - Journal of the American Medical Association, 2022, 327, 1021.	3.8	10
22	QALYs In Health Resource Usage Decisions: The Authors Reply. Health Affairs, 2022, 41, 610-610.	2.5	0
23	Price changes and within-class competition of cancer drugs in the USA and Europe: a comparative analysis. Lancet Oncology, The, 2022, 23, 514-520.	5.1	22
24	Making the Case for Accelerated Withdrawal of Aducanumab. Journal of Alzheimer's Disease, 2022, 87, 1003-1007.	1.2	19
25	Therapeutic Value Assessments of Novel Medicines in the US and Europe, 2018-2019. JAMA Network Open, 2022, 5, e226479.	2.8	8
26	Risks to the 340B Drug Pricing Program Related to Manufacturer Restrictions on Drug Availability. JAMA - Journal of the American Medical Association, 2022, , .	3.8	4
27	Direct-to-Consumer Generic Drugs: A Maverick Approach or Another Exposure of Market Failures?. Annals of Internal Medicine, 2022, 175, 890-891.	2.0	3
28	Use of Extrapolation in New Drug Approvals by the US Food and Drug Administration. JAMA Network Open, 2022, 5, e227958.	2.8	8
29	Medicaid Expenditures and Estimated Rebates on Line Extension Drugs, 2010â€“2018. Journal of General Internal Medicine, 2022, , .	1.3	5
30	State Laws and Generic Substitution in the Year After New Generic Competition. Value in Health, 2022, , .	0.1	1
31	Coverage of New Drugs in Medicare Part D. Milbank Quarterly, 2022, 100, 562-588.	2.1	3
32	Unwanted Advice? Frequency, Characteristics, And Outcomes Of Negative Advisory Committee Votes For FDA-Approved Drugs. Health Affairs, 2022, 41, 713-721.	2.5	7
33	Analysis of Supportive Evidence for US Food and Drug Administration Approvals of Novel Drugs in 2020. JAMA Network Open, 2022, 5, e2212454.	2.8	11
34	Institutionalizing Misinformation â€” The Dietary Supplement Listing Act of 2022. New England Journal of Medicine, 2022, 387, 3-5.	13.9	4
35	Improving the quality of US drug patents through international awareness. BMJ, The, 2022, 377, e068172.	3.0	3
36	Postâ€Marketing Requirements for Cancer Drugs Approved by the European Medicines Agency, 2004â€“2014. Clinical Pharmacology and Therapeutics, 2022, 112, 846-852.	2.3	2

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37	Trends in Prescription Drug Launch Prices, 2008-2021. JAMA - Journal of the American Medical Association, 2022, 327, 2145.	3.8	45
38	Clinical Benefit and Expedited Approval of Cancer Drugs in the United States, European Union, Switzerland, Japan, Canada, and Australia. JCO Oncology Practice, 2022, 18, e1522-e1532.	1.4	16
39	Potential Medicare Part D Savings on Generic Drugs From the Mark Cuban Cost Plus Drug Company. Annals of Internal Medicine, 2022, 175, 1053-1055.	2.0	10
40	Over-the-Counter Availability of Rescue Inhalers for Asthmaâ€"Reply. JAMA - Journal of the American Medical Association, 2022, 328, 216.	3.8	1
41	Payments for research participation: Donâ€™t tax the Guinea pig. Clinical Trials, 2022, 19, 579-583.	0.7	1
42	Federal Spending on Off-Patent Drugs That Lack Generic Competition. Journal of General Internal Medicine, 2021, 36, 821-823.	1.3	4
43	Market Exclusivity Length for Drugs with New Generic or Biosimilar Competition, 2012â€“2018. Clinical Pharmacology and Therapeutics, 2021, 109, 367-371.	2.3	26
44	Reply to Boucher et al. Clinical Infectious Diseases, 2021, 72, e422-e423.	2.9	0
45	Public funding for transformative drugs: the case of sofosbuvir. Drug Discovery Today, 2021, 26, 273-281.	3.2	10
46	An Overview Of Vaccine Development, Approval, And Regulation, With Implications For COVID-19. Health Affairs, 2021, 40, 25-32.	2.5	27
47	Changes in Erythropoiesis Stimulating Agent Use Under a Risk Evaluation and Mitigation Strategy (REMS) Program. Drug Safety, 2021, 44, 327-335.	1.4	4
48	Factors Affecting Buprenorphine Utilization and Spending in Medicaid, 2002-2018. Value in Health, 2021, 24, 182-187.	0.1	6
49	The Wrong Cure: Financial Incentives for Unimpressive New Antibiotics. Journal of Infectious Diseases, 2021, 223, 1506-1509.	1.9	3
50	Assessment of Variation in State Regulation of Generic Drug and Interchangeable Biologic Substitutions. JAMA Internal Medicine, 2021, 181, 16.	2.6	22
51	INTRODUCTION: Public Sector and Non-Profit Contributions to Drug Development â€" Historical Scope, Opportunities, and Challenges. Journal of Law, Medicine and Ethics, 2021, 49, 6-9.	0.4	4
52	A correlation analysis to assess event-free survival as a trial-level surrogate for overall survival in early breast cancer. EClinicalMedicine, 2021, 32, 100730.	3.2	9
53	Buprenorphine for opioid use disorder: The role of public funding in its development. Drug and Alcohol Dependence, 2021, 219, 108491.	1.6	6
54	ASHP Foundation Pharmacy Forecast 2021: Strategic Planning Advice for Pharmacy Departments in Hospitals and Health Systems. American Journal of Health-System Pharmacy, 2021, 78, 472-497.	0.5	12

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55	The Future of Drug-Pricing Transparency. New England Journal of Medicine, 2021, 384, 489-491.	13.9	11
56	Associations Between Copays, Coverage Limits for Opioid Use Disorder Medications, and Prescribing in Medicaid, 2018. Medical Care, 2021, 59, 266-272.	1.1	0
57	Prospects for Enforcing Prohibitions on Off-Label Drug Promotion after United States v. Caronia: An Analysis of Litigated Cases. Journal of Health Politics, Policy and Law, 2021, 46, 487-504.	0.9	3
58	Why France Spends Less Than the United States on Drugs: A Comparative Study of Drug Pricing and Pricing Regulation. Milbank Quarterly, 2021, 99, 240-272.	2.1	14
59	Trends in Medicare Part D Inhaler Spending: 2012â€“2018. Annals of the American Thoracic Society, 2021, 18, 548-550.	1.5	8
60	Frequency of First Generic Drug Approvals With â€œSkinny Labelsâ€•in the United States. JAMA Internal Medicine, 2021, 181, 995-997.	2.6	8
61	Non-warfarin oral anticoagulant copayments and adherence in atrial fibrillation: A population-based cohort study. American Heart Journal, 2021, 233, 109-121.	1.2	23
62	FDA approval standards for anticancer agents â€" lessons from two recent approvals in breast cancer. Nature Reviews Clinical Oncology, 2021, 18, 397-398.	12.5	4
63	Paying for Prescription Drugs in the New Administration. JAMA - Journal of the American Medical Association, 2021, 325, 819.	3.8	9
64	Identifying potential prescription drug product hopping. Nature Biotechnology, 2021, 39, 414-417.	9.4	2
65	Public funding and the importance of reasonable pricing for buprenorphine. Drug and Alcohol Dependence, 2021, 221, 108643.	1.6	0
66	International Reference Pricing for Prescription Drugs in the United States: Administrative Limitations and Collateral Effects. Value in Health, 2021, 24, 473-476.	0.1	7
67	A Multi-modal Approach to Evaluate the Impact of Risk Evaluation and Mitigation Strategy (REMS) Programs. Drug Safety, 2021, 44, 743-751.	1.4	7
68	Characteristics of Postmarketing Studies for Vaccines Approved by the US Food and Drug Administration, 2006-2020. JAMA Network Open, 2021, 4, e218530.	2.8	2
69	Substitution of Generic Drugs and Biosimilarsâ€"Reply. JAMA Internal Medicine, 2021, 181, 568.	2.6	0
70	Assessment of Coverage in England of Cancer Drugs Qualifying for US Food and Drug Administration Accelerated Approval. JAMA Internal Medicine, 2021, 181, 490.	2.6	32
71	Frequency Of Generic Drug Price Spikes And Impact On Medicaid Spending. Health Affairs, 2021, 40, 779-785.	2.5	4
72	Correlation Between Changes in Brand-Name Drug Prices and Patient Out-of-Pocket Costs. JAMA Network Open, 2021, 4, e218816.	2.8	15

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73	Evaluation of Aducanumab for Alzheimer Disease. JAMA - Journal of the American Medical Association, 2021, 325, 1717.	3.8	152
74	Integrating New Effectiveness Data Into US Food and Drug Administrationâ€“Approved Drug Labeling. JAMA Internal Medicine, 2021, 181, 897-898.	2.6	5
75	The timing of 30â€month stay expirations and generic entry: A cohort study of first generics, 2013â€“2020. Clinical and Translational Science, 2021, 14, 1917-1923.	1.5	5
76	Diabetes Drugs: List Price Increases Were Not Always Reflected In Net Price; Impact Of Brand Competition Unclear. Health Affairs, 2021, 40, 772-778.	2.5	5
77	Association of Californiaâ€™s Prescription Drug Coupon Ban With Generic Drug Use. JAMA - Journal of the American Medical Association, 2021, 325, 2399.	3.8	2
78	Estimating Rebates and Other Discounts Received by Medicare Part D. JAMA Health Forum, 2021, 2, e210626.	1.0	18
79	Factors Associated With Generic Drug Uptake in the United States, 2012 to 2017. Value in Health, 2021, 24, 804-811.	0.1	8
80	Continual learning in medical devices: FDA's action plan and beyond. The Lancet Digital Health, 2021, 3, e337-e338.	5.9	43
81	Fulfilling the Mandate of the US Food and Drug Administrationâ€™s Accelerated Approval Pathway. JAMA Internal Medicine, 2021, 181, 1275.	2.6	34
82	Public-sector Contributions to Novel Biologic Drugs. JAMA Internal Medicine, 2021, 181, 1522.	2.6	6
83	Assessing the Impact of US Food and Drug Administration Breakthrough Therapy Designation Timing on Trial Characteristics and Development Speed. Clinical Pharmacology and Therapeutics, 2021, 110, 1018-1024.	2.3	11
84	Cost to Medicare of Delayed Adalimumab Biosimilar Availability. Clinical Pharmacology and Therapeutics, 2021, 110, 1050-1056.	2.3	11
85	FDA Regulation and Approval of Medical Devices: 1976-2020. JAMA - Journal of the American Medical Association, 2021, 326, 420.	3.8	53
86	Barriers To US Biosimilar Market Growth: Lessons From Biosimilar Patent Litigation. Health Affairs, 2021, 40, 1198-1205.	2.5	15
87	Will Ending the Medicaid Drug Rebate Cap Lower Drug Prices?. JAMA Internal Medicine, 2021, 181, 1034.	2.6	11
88	Pursuing Valueâ€Based Prices for Drugs:ÂA Comprehensive Comparison of State Prescription Drugâ€“Pricing Boards. Milbank Quarterly, 2021, , .	2.1	3
89	Market Exclusivity and Changes in Competition and Prices Associated With the US Food and Drug Administration Unapproved Drug Initiative. JAMA Internal Medicine, 2021, 181, 1124.	2.6	4
90	Revisiting FDA Approval of Aducanumab. New England Journal of Medicine, 2021, 385, 769-771.	13.9	104

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91	Discovery and Development of Pregabalin (Lyrica). Neurology, 2021, 97, e1653-e1660.	1.5	1
92	Analysis of Launch and Postapproval Cancer Drug Pricing, Clinical Benefit, and Policy Implications in the US and Europe. JAMA Oncology, 2021, 7, e212026.	3.4	46
93	A Court Decision on â€œSkinny Labelingâ€: JAMA - Journal of the American Medical Association, 2021, 326, 1371.	3.8	6
94	Characteristics of US Patients and Prescribers Using Hydroxychloroquine During the COVID-19 Pandemic. Journal of General Internal Medicine, 2021, 36, 3918-3921.	1.3	0
95	Reimagining Pharmaceutical Market Exclusivities: Should the Duration of Guaranteed Monopoly Periods Be Value Based?. Value in Health, 2021, 24, 1328-1334.	0.1	8
96	Controversy Over Using Quality-Adjusted Life-Years In Cost-Effectiveness Analyses: A Systematic Literature Review. Health Affairs, 2021, 40, 1402-1410.	2.5	17
97	International reference pricing for prescription drugs: a landscape analysis. Journal of Managed Care & Specialty Pharmacy, 2021, 27, 1309-1313.	0.5	3
98	Regulatory and clinical consequences of negative confirmatory trials of accelerated approval cancer drugs: retrospective observational study. BMJ, The, 2021, 374, n1959.	3.0	40
99	Repurposing existing drugs for new uses: a cohort study of the frequency of FDA-granted new indication exclusivities since 1997. Journal of Pharmaceutical Policy and Practice, 2021, 14, 3.	1.1	28
100	Over-the-Counter Monograph Safety, Innovation, and Reform Act. Journal of Law, Medicine and Ethics, 2021, 49, 321-327.	0.4	1
101	COPD EXACERBATIONS AND PNEUMONIA HOSPITALIZATIONS IN NEW USERS OF COMBINATION MAINTENANCE INHALERS: A COMPARATIVE EFFECTIVENESS AND SAFETY STUDY. Chest, 2021, 160, A1795.	0.4	1
102	Medicare Spending on Drugs With Accelerated Approval, 2015-2019. JAMA Health Forum, 2021, 2, e213937.	1.0	8
103	Promoting Competition in Drug Pricing: A Review of Recent Congressional Legislation. Journal of Law, Medicine and Ethics, 2021, 49, 683-687.	0.4	0
104	Raising Medicaid Rebates For Drugs With Accelerated Approval. Health Affairs, 2021, 40, 1935-1942.	2.5	7
105	Transferrable Market Exclusivity Extensions to Promote Antibiotic Development: An Economic Analysis. Clinical Infectious Diseases, 2020, 71, 1671-1675.	2.9	11
106	US Food and Drug Administration Recommendations on the Use of Surrogate Measures as End Points in New Anti-infective Drug Approvals. JAMA Internal Medicine, 2020, 180, 131.	2.6	8
107	FDA and EMA Biosimilar Approvals. Journal of General Internal Medicine, 2020, 35, 1908-1910.	1.3	11
108	Confidentiality Orders and Public Interest in Drug and Medical Device Litigation. JAMA Internal Medicine, 2020, 180, 292.	2.6	4

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109	Novelty of Active Ingredients in High-Cost Brand-Name Drugs. Journal of General Internal Medicine, 2020, 35, 2219-2221.	1.3	1
110	Potential Medicare Savings on Inhaler Prescriptions Through the Use of Negotiated Prices and a Defined Formulary. JAMA Internal Medicine, 2020, 180, 454.	2.6	4
111	Getting the Right Evidence After Drug Approval. Frontiers in Pharmacology, 2020, 11, 569535.	1.6	10
112	Implementing U.S. Covid-19 Testing: Regulatory and Infrastructural Challenges. Journal of Law, Medicine and Ethics, 2020, 48, 608-612.	0.4	1
113	Clinical Development Times for Biosimilars in the United States. Mayo Clinic Proceedings, 2020, 95, 2152-2154.	1.4	3
114	Association between FDA and EMA expedited approval programs and therapeutic value of new medicines: retrospective cohort study. BMJ, The, 2020, 371, m3434.	3.0	56
115	Projected spending for brand-name drugs in English primary care given US prices: a cross-sectional study. Journal of the Royal Society of Medicine, 2020, 113, 350-359.	1.1	2
116	Revisiting the National Institutes of Health Fair Pricing Condition: Promoting the Affordability of Drugs Developed With Government Support. Annals of Internal Medicine, 2020, 172, 348-350.	2.0	9
117	Clinical benefit and cost of breakthrough cancer drugs approved by the US Food and Drug Administration. Cancer, 2020, 126, 4390-4399.	2.0	19
118	US Spending Associated With Transition From Daily to 3-Times-Weekly Glatiramer Acetate. JAMA Internal Medicine, 2020, 180, 1165.	2.6	7
119	Drug Prices, Rebates, and Discounts. JAMA - Journal of the American Medical Association, 2020, 324, 399.	3.8	4
120	Missed Opportunities on Emergency Remdesivir Use. JAMA - Journal of the American Medical Association, 2020, 324, 331.	3.8	15
121	Accounting for US public funding in drug development: how can we better balance access, affordability, and innovation?. BMJ, The, 2020, 371, m3841.	3.0	7
122	Understanding when real world data can be used to replicate a clinical trial: A crossâ€sectional study of medications approved in 2011. Pharmacoepidemiology and Drug Safety, 2020, 29, 1273-1278.	0.9	2
123	Regulatory Decision-making on COVID-19 Vaccines During a Public Health Emergency. JAMA - Journal of the American Medical Association, 2020, 324, 1284.	3.8	23
124	Up Is Down â€" Pharmaceutical Industry Caution vs. Federal Acceleration of Covid-19 Vaccine Approval. New England Journal of Medicine, 2020, 383, 1706-1708.	13.9	11
125	Variations in Generic Combination Opioid Use Across State Medicaid Programs. Journal of General Internal Medicine, 2020, 36, 3240-3242.	1.3	0
126	A qualitative study of biosimilar manufacturer and regulator perceptions on intellectual property and abbreviated approval pathways. Nature Biotechnology, 2020, 38, 1253-1256.	9.4	8

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127	Development of a National Public Pharmaceutical Research and Development Institute. Journal of Law, Medicine and Ethics, 2020, 48, 225-227.	0.4	0
128	Need for Transparency and Reliable Evidence in Emergency Use Authorizations for Coronavirus Disease 2019 (COVID-19) Therapies. JAMA Internal Medicine, 2020, 180, 1145.	2.6	21
129	Estimating The Cost Of Delayed Generic Drug Entry To Medicaid. Health Affairs, 2020, 39, 1011-1017.	2.5	4
130	Regulatory approval characteristics of antimicrobial versus non-antimicrobial products, 1984â€“2018: an evaluation of Food and Drug Administration flexibilities. Lancet Infectious Diseases, The, 2020, 20, e159-e164.	4.6	3
131	Specialty Drugs â€” A Distinctly American Phenomenon. New England Journal of Medicine, 2020, 382, 2179-2181.	13.9	7
132	False Negative Tests for SARS-CoV-2 Infection â€” Challenges and Implications. New England Journal of Medicine, 2020, 383, e38.	13.9	721
133	Evaluating the evidence behind the surrogate measures included in the FDA's table of surrogate endpoints as supporting approval of cancer drugs. EClinicalMedicine, 2020, 21, 100332.	3.2	80
134	Rates and Costs of Dispensing Naloxone to Patients at High Risk for Opioid Overdose in the United States, 2014â€“2018. Drug Safety, 2020, 43, 669-675.	1.4	14
135	Using realâ€world safety data in regulatory approval decisions: Sotagliflozin and the risk of diabetic ketoacidosis. Pharmacoepidemiology and Drug Safety, 2020, 29, 1322-1324.	0.9	3
136	Generating comparative evidence on new drugs and devices before approval. Lancet, The, 2020, 395, 986-997.	6.3	59
137	Decision Making Under Uncertainty: Comparing Regulatory and Health Technology Assessment Reviews of Medicines in the United States and Europe. Clinical Pharmacology and Therapeutics, 2020, 108, 350-357.	2.3	41
138	Lessons From The Impact Of Price Regulation On The Pricing Of Anticancer Drugs In Germany. Health Affairs, 2020, 39, 1185-1193.	2.5	30
139	Comparing Onset of Biosimilar Versus Generic Competition in the United States. Clinical Pharmacology and Therapeutics, 2020, 108, 1308-1314.	2.3	7
140	FDA Approval and Regulation of Pharmaceuticals, 1983-2018. JAMA - Journal of the American Medical Association, 2020, 323, 164.	3.8	197
141	Estimation of Medicare Part D Spending on Insulin for Patients With Diabetes Using Negotiated Prices and a Defined Formulary. JAMA Internal Medicine, 2020, 180, 597.	2.6	10
142	Internet Searches for Unproven COVID-19 Therapies in the United States. JAMA Internal Medicine, 2020, 180, 1116.	2.6	97
143	Prices and clinical benefit of cancer drugs in the USA and Europe: a costâ€“benefit analysis. Lancet Oncology, The, 2020, 21, 664-670.	5.1	126
144	Using Data From Routine Care to Estimate the Effectiveness and Potential Limitations of Outcomes-Based Contracts for Diabetes Medications. Value in Health, 2020, 23, 434-440.	0.1	6

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145	Insulin access and affordability in the USA: anticipating the first interchangeable insulin product. Lancet Diabetes and Endocrinology,the, 2020, 8, 360-362.	5.5	4
146	Prescribing systemic steroids for acute respiratory tract infections in United States outpatient settings: A nationwide population-based cohort study. PLoS Medicine, 2020, 17, e1003058.	3.9	13
147	Increasing Access to FDA Inspection Reports on Irregularities and Misconduct in Clinical Trials. JAMA - Journal of the American Medical Association, 2020, 323, 1903.	3.8	9
148	Incentivizing Antibiotic Development: Why Isnâ€™t the Generating Antibiotic Incentives Now (GAIN) Act Working?. Open Forum Infectious Diseases, 2020, 7, ofaa001.	0.4	19
149	The evidence landscape in precision medicine. Science Translational Medicine, 2020, 12, .	5.8	16
150	New drug approvals in oncology. Nature Reviews Clinical Oncology, 2020, 17, 140-146.	12.5	27
151	Social, cultural and economic aspects of antimicrobial resistance. Bulletin of the World Health Organization, 2020, 98, 823-823A.	1.5	16
152	Response Rates and Durations of Response for Biomarker-Based Cancer Drugs in Nonrandomized Versus Randomized Trials. Journal of the National Comprehensive Cancer Network: JNCCN, 2020, 18, 36-43.	2.3	21
153	Preferences for and Experiences With Pill Appearance Changes: National Surveys of Patients and Pharmacists. American Journal of Managed Care, 2020, 26, 340-347.	0.8	2
154	Generic Competition for Drugs Treating Rare Diseases. Journal of Law, Medicine and Ethics, 2020, 48, 789-795.	0.4	1
155	Title is missing!. , 2020, 17, e1003058.		0
156	Title is missing!. , 2020, 17, e1003058.		0
157	Title is missing!. , 2020, 17, e1003058.		0
158	Title is missing!. , 2020, 17, e1003058.		0
159	Title is missing!. , 2020, 17, e1003058.		0
160	Effect of Lawyer-Submitted Reports on Signals of Disproportional Reporting in the Food and Drug Administrationâ€™s Adverse Event Reporting System. Drug Safety, 2019, 42, 85-93.	1.4	5
161	Patent term restoration for top-selling drugs in the United States. Drug Discovery Today, 2019, 24, 20-25.	3.2	18
162	The Generic Drug Industry Embraces a Faster, Cheaper Pathway for Challenging Patents. Applied Health Economics and Health Policy, 2019, 17, 47-54.	1.0	10

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163	Why Are Biosimilars Not Living up to Their Promise in the US?. AMA Journal of Ethics, 2019, 21, E668-678.	0.4	44
164	Landscape of Cardiovascular Device Registries in the United States. Journal of the American Heart Association, 2019, 8, e012756.	1.6	3
165	Potential Medicare Savings From Generic Substitution and Therapeutic Interchange of ACE Inhibitors and Angiotensin-II-Receptor Blockers. JAMA Internal Medicine, 2019, 179, 1712.	2.6	8
166	Impact of State Laws Restricting Opioid Duration on Characteristics of New Opioid Prescriptions. Journal of General Internal Medicine, 2019, 34, 2339-2341.	1.3	27
167	Orphan Drug Designation and Exclusivity for â€œSame Drugsâ€• Journal of Law, Medicine and Ethics, 2019, 47, 347-349.	0.4	1
168	Price Increases of Protected-Class Drugs in Medicare Part D, Relative to Inflation, 2012-2017. JAMA - Journal of the American Medical Association, 2019, 322, 267.	3.8	21
169	Multimodal Analysis of FDA Drug Safety Communications: Lessons from Zolpidem. Drug Safety, 2019, 42, 1287-1295.	1.4	13
170	Reforming the Orphan Drug Act for the 21st Century. New England Journal of Medicine, 2019, 381, 106-108.	13.9	31
171	Public Approval of Exception From Informed Consent in Emergency Clinical Trials. JAMA Network Open, 2019, 2, e197591.	2.8	21
172	Competition and price among brand-name drugs in the same class: A systematic review of the evidence. PLoS Medicine, 2019, 16, e1002872.	3.9	42
173	The Impact Of Price Regulation On The Availability Of New Drugs In Germany. Health Affairs, 2019, 38, 1182-1187.	2.5	19
174	Characteristics of Recent Generic Drug Approvals by the US Food and Drug Administration. JAMA Network Open, 2019, 2, e1913029.	2.8	3
175	Transparency on Prescription Drug Research Expenditures. , 2019, , 121-131.		0
176	Physician Perceptions of Step Therapy Prescribing Requirements. Journal of Managed Care & Specialty Pharmacy, 2019, 25, 1210-1224.	0.5	4
177	Prescription Opioid Epidemic and Trends in the Clinical Development of New Pain Medications. Mayo Clinic Proceedings, 2019, 94, 2437-2443.	1.4	5
178	Randomized Study of Providing Evidence Context to Mitigate Physician Misinterpretation Arising From Off-Label Drug Promotion. Circulation: Cardiovascular Quality and Outcomes, 2019, 12, e006073.	0.9	4
179	The U.S. Insulin Crisis â€" Rationing a Lifesaving Medication Discovered in the 1920s. New England Journal of Medicine, 2019, 381, 1793-1795.	13.9	45
180	Public sector financial support for late stage discovery of new drugs in the United States: cohort study. BMJ: British Medical Journal, 2019, 367, l5766.	2.4	42

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181	Desmopressin and the risk of hyponatremia: A population-based cohort study. PLoS Medicine, 2019, 16, e1002930.	3.9	16
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