

“Market withdrawals” of medicines in Germany and clinical guideline recommendations

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#	ARTICLE	IF	CITATIONS
3	Varying results of early benefit assessment of newly approved pharmaceutical drugs in Germany from 2011 to 2017: A study based on federal joint committee data. <i>Journal of Evidence-Based Medicine</i> , 2019, 12, 9-15.	2.4	3
4	Different interpretation of additional evidence for HTA by the commissioned HTA body and the commissioning decision maker in Germany: whenever IQWiG and Federal Joint Committee disagree. <i>Health Economics Review</i> , 2019, 9, 35.	0.8	6
5	A decade of early benefit assessment of ophthalmic drugs in Germany: success story or not?. <i>Expert Review of Pharmacoeconomics and Outcomes Research</i> , 2021, , 1-15.	0.7	0
6	Zehn Jahre AMNOG – Rückblick und Ausblick aus Sicht der Arzneimittelkommission der deutschen Ärzteschaft. , 2020, , 185-200.		6
8	Multicenter, cross-sectional study of the costs of illness and cost-driving factors in adult patients with epilepsy. <i>Epilepsia</i> , 2022, 63, 904-918.	2.6	20
9	How Far is Germany From Value-Based Pricing 10 Years After the Introduction of AMNOG?. <i>Applied Health Economics and Health Policy</i> , 2022, 20, 287-290.	1.0	3
10	Does health technology assessment compromise access to pharmaceuticals?. <i>European Journal of Health Economics</i> , 2023, 24, 437-451.	1.4	2
12	Getting the Price Right: Lessons for Medicare Price Negotiation from Peer Countries. <i>Pharmacoeconomics</i> , 2022, 40, 1131-1142.	1.7	2
13	Health App by Prescription: The German Nation-Wide Model. , 2022, , 63-79.		0
14	Qualität der Arzneimittelversorgung – Theoretischer und konzeptueller Rahmen. , 2022, , 13-29.		0