

Declan Noone

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

270
citations

933447

10
h-index

940533

16
g-index

22
all docs

22
docs citations

22
times ranked

254
citing authors

#	ARTICLE	IF	CITATIONS
1	Hemophilia treatment in 2021: Choosing the "optimal" treatment using an integrative, patient-oriented approach to shared decision-making between patients and clinicians. <i>Blood Reviews</i> , 2022, 52, 100890.	5.7	22
2	Key challenges for patient registries " A report from the 1 st workshop of the EHC Think Tank Workstream on Registries. <i>The Journal of Haemophilia Practice</i> , 2022, 9, 14-19.	0.4	0
3	Key challenges for hub and spoke models of care " A report from the 1st workshop of the EHC Think Tank on Hub and Spoke Treatment Models. <i>The Journal of Haemophilia Practice</i> , 2022, 9, 20-26.	0.4	1
4	New challenges for an expanding generation of older persons with haemophilia. <i>The Journal of Haemophilia Practice</i> , 2022, 9, 1-13.	0.4	0
5	Patient agency: key questions and challenges " A report from the 1st workshop of the EHC Think Tank Workstream on Patient Agency. <i>The Journal of Haemophilia Practice</i> , 2022, 9, 27-35.	0.4	1
6	Reimbursing the value of gene therapy care in an era of uncertainty. <i>Haemophilia</i> , 2021, 27, 12-18.	2.1	7
7	Non-severe haemophilia: Is it benign? " Insights from the PROBE study. <i>Haemophilia</i> , 2021, 27, 17-24.	2.1	16
8	Vaccination against COVID-19: Rationale, modalities and precautions for patients with haemophilia and other inherited bleeding disorders. <i>Haemophilia</i> , 2021, 27, 515-518.	2.1	9
9	Clinical attributes and treatment characteristics are associated with work productivity and activity impairment in people with severe haemophilia A. <i>Haemophilia</i> , 2021, 27, 938-946.	2.1	7
10	European principles of care for women and girls with inherited bleeding disorders. <i>Haemophilia</i> , 2021, 27, 837-847.	2.1	23
11	Evaluation of the sexual health in people living with hemophilia. <i>Haemophilia</i> , 2021, 27, 993-1001.	2.1	2
12	Delivery of AAV-based gene therapy through haemophilia centres " A need for re-evaluation of infrastructure and comprehensive care: A Joint publication of EAHAD and EHC. <i>Haemophilia</i> , 2021, 27, 967-973.	2.1	29
13	Recombinant factor IX-Fc fusion protein in severe hemophilia B: Patient-reported outcomes and health-related quality of life. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12602.	2.3	4
14	Eliminating Panglossian thinking in development of AAV therapeutics. <i>Molecular Therapy</i> , 2021, 29, 3325-3327.	8.2	12
15	Evolution of Haemophilia Care in Europe: 10 years of the principles of care. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 184.	2.7	10
16	Gene therapy to cure haemophilia: Is robust scientific inquiry the missing factor?. <i>Haemophilia</i> , 2020, 26, 931-933.	2.1	24
17	Understanding minimum and ideal factor levels for participation in physical activities by people with haemophilia: An expert elicitation exercise. <i>Haemophilia</i> , 2020, 26, 711-717.	2.1	19
18	Barriers and challenges faced by women with congenital bleeding disorders in Europe: Results of a patient survey conducted by the European Haemophilia Consortium. <i>Haemophilia</i> , 2019, 25, 468-474.	2.1	38

#	ARTICLE	IF	CITATIONS
19	Exploring regional variations in the cross-cultural, international implementation of the Patient Reported Outcomes Burdens and Experience (PROBE) study. Haemophilia, 2019, 25, 365-372.	2.1	11
20	Test-retest properties of the Patient Reported Outcomes, Burdens and Experiences (PROBE) questionnaire and its constituent domains. Haemophilia, 2019, 25, 75-83.	2.1	14
21	Psychometric properties of the Patient Reported Outcomes, Burdens and Experiences (PROBE) questionnaire. BMJ Open, 2018, 8, e021900.	1.9	15
22	Issues in assessing products for the treatment of hemophilia – the intersection between efficacy, economics, and ethics. Journal of Blood Medicine, 2015, 6, 185.	1.7	6