

Jan Domaradzki

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7042300/publications.pdf>

Version: 2024-02-01

25
papers

397
citations

840119

11
h-index

839053

18
g-index

25
all docs

25
docs citations

25
times ranked

283
citing authors

#	ARTICLE	IF	CITATIONS
1	Hospital chaplains facing the pandemic. A qualitative study. <i>Journal of Health Care Chaplaincy</i> , 2023, 29, 145-160.	0.7	2
2	“We are also here” Spiritual Care Practitioners’ Experiences of the COVID-19 Pandemic: A Qualitative Study from Poland. <i>Journal of Religion and Health</i> , 2022, 61, 962-992.	0.8	14
3	“Who Else If Not We” Medical Students’ Perception and Experiences with Volunteering during the COVID-19 Crisis in Poznan, Poland. <i>International Journal of Environmental Research and Public Health</i> , 2022, 19, 2314.	1.2	17
4	Treating rare diseases with the cinema: Can popular movies enhance public understanding of rare diseases?. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, 117.	1.2	4
5	Better Late Than Never: Predictors of Delayed COVID-19 Vaccine Uptake in Poland. <i>Vaccines</i> , 2022, 10, 528.	2.1	12
6	The Awareness of Rare Diseases Among Medical Students and Practicing Physicians in the Republic of Kazakhstan. An Exploratory Study. <i>Frontiers in Public Health</i> , 2022, 10, 872648.	1.3	10
7	The Werther Effect, the Papageno Effect or No Effect? A Literature Review. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 2396.	1.2	40
8	Does Religion Influence the Motivations of Future Healthcare Professionals to Volunteer During the COVID-19 Pandemic in Poland? An Exploratory Study. <i>Journal of Religion and Health</i> , 2021, 60, 1507-1520.	0.8	16
9	Medical Students' Voluntary Service During the COVID-19 Pandemic in Poland. <i>Frontiers in Public Health</i> , 2021, 9, 618608.	1.3	27
10	Knowledge and Attitudes of Future Healthcare Professionals Toward Rare Diseases. <i>Frontiers in Genetics</i> , 2021, 12, 639610.	1.1	17
11	Are rare diseases overlooked by medical education? Awareness of rare diseases among physicians in Poland: an explanatory study. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 400.	1.2	32
12	Thomas Szasz: The uncompromising rebel and critic of psychiatry. <i>Psychiatria Polska</i> , 2021, 55, 851-867.	0.2	1
13	Needs assessment study of rare diseases education for nurses and nursing students in Poland. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 167.	1.2	26
14	Public Attitudes toward Biobanking of Human Biological Material for Research Purposes: A Literature Review. <i>International Journal of Environmental Research and Public Health</i> , 2019, 16, 2209.	1.2	64
15	Medical students' knowledge and opinions about rare diseases: A case study from Poland. <i>Intractable and Rare Diseases Research</i> , 2019, 8, 252-259.	0.3	42
16	Chapter 13 Geneticization and Bioethics: Ethical Dilemmas in Genetic Counselling. <i>Philosophy and Medicine</i> , 2018, , 189-204.	0.3	0
17	Medykalizacja a genetyzacja: ci...gÅ,oÅvÅ† czy zmiana?. <i>PrzełÅ...d Socjologiczny</i> , 2017, 66, .	1.1	0
18	Definitions of health and disease among physicians and SpoÅ,em PSS employees. <i>Journal of Medical Science</i> , 2017, 86, 95-105.	0.2	0

#	ARTICLE	IF	CITATIONS
19	Family caregiversâ€™ experiences with healthcare services - a case of Huntington disease. <i>Psychiatria Polska</i> , 2016, 50, 375-391.	0.2	11
20	Behavioural genetics in Polish print news media between 2000-2014. <i>Psychiatria Polska</i> , 2016, 50, 1251-1271.	0.2	2
21	The Impact of Huntington Disease on Family Carers: a Literature Overview. <i>Psychiatria Polska</i> , 2015, 49, 931-944.	0.2	42
22	Patient rights, risk, and responsibilities in the genetic era â€“ a right to know, a right not to know, or a duty to know?. <i>Annals of Agricultural and Environmental Medicine</i> , 2015, 22, 156-162.	0.5	14
23	DNA and its Metaphors. <i>Respectus Philologicus</i> , 2015, 27, 74-84.	0.1	2
24	Caring for patients with Huntington disease â€“ A survey of caregiversâ€™ experiences and views. <i>Journal of Pre-Clinical and Clinical Research</i> , 2015, 9, 133-139.	0.2	1
25	Prawo do niewiedzy a obowiÄzek wiedzy w opiniach rodzin osÃ³b z chorobÄ... Huntingtona. <i>Etyka</i> , 0, 47, 18-33.	0.0	1