Martine C De Vries

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Consent for Delivery Room Studies: What Can Be Learned from Perceptions of Parents. Neonatology, 2022, 119, 214-221.	0.9	0
2	Towards Culturally Sensitive Shared Decision-Making in Oncology A Study Protocol Integrating Bioethical Qualitative Research on Shared Decision-Making Among Ethnic Minorities With Ethical Reflection. International Journal of Qualitative Methods, The, 2022, 21, 160940692210867.	1.3	1
3	How Traditional Informed Consent Impairs Inclusivity in a Learning Healthcare System: Lessons Learned from the Utrecht Cardiovascular Cohort. Journal of Clinical Epidemiology, 2022, , .	2.4	4
4	When is it impractical to ask informed consent? A systematic review. Clinical Trials, 2022, 19, 545-560.	0.7	10
5	Text-mining in electronic healthcare records can be used as efficient tool for screening and data collection in cardiovascular trials: a multicenter validation study. Journal of Clinical Epidemiology, 2021, 132, 97-105.	2.4	23
6	Serum hepcidin concentrations in relation to iron status in children with type 1 diabetes. Pediatric Hematology and Oncology, 2021, 38, 108-123.	0.3	0
7	Parents' Perspectives and Societal Acceptance of Implementation of Newborn Screening for SCID in the Netherlands. Journal of Clinical Immunology, 2021, 41, 99-108.	2.0	25
8	Antimicrobial guidelines in clinical practice: incorporating the ethical perspective. JAC-Antimicrobial Resistance, 2021, 3, dlab074.	0.9	0
9	Fetoscopic myelomeningocoele closure: Is the scientific evidence enough to challenge the gold standard for prenatal surgery?. Prenatal Diagnosis, 2021, 41, 949-956.	1.1	14
10	Decision making for hematopoietic stem cell transplantation in pediatric, adolescent, and young adult patients with a hemoglobinopathy—Shared or not?. Pediatric Blood and Cancer, 2021, 68, e29099.	0.8	5
11	How to Facilitate Decision-Making for Hematopoietic Stem Cell Transplantation in Patients With Hemoglobinopathies. The Perspectives of Healthcare Professionals. Frontiers in Pediatrics, 2021, 9, 690309.	0.9	3
12	Reviewing recordings of neonatal resuscitation with parents. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2021, 106, 346-351.	1.4	9
13	Assessing Medical Decision-Making Competence in Transgender Youth. Pediatrics, 2021, 148, .	1.0	21
14	The Duty to Support Learning Health Systems: A Broad Rather than a Narrow Interpretation. American Journal of Bioethics, 2021, 21, 14-16.	0.5	1
15	SERIES: eHealth in primary care. Part 2: Exploring the ethical implications of its application in primary care practice. European Journal of General Practice, 2020, 26, 26-32.	0.9	45
16	Ethical Advice for an Intensive Care Triage Protocol in the COVID-19 Pandemic: Lessons Learned from The Netherlands. Public Health Ethics, 2020, 13, 157-165.	0.4	22
17	Money-oriented risk-takers or deliberate decision-makers: a cross-sectional survey study of participants in controlled human infection trials. BMJ Open, 2020, 10, e033796.	0.8	10
18	The ethics of deferred consent in times of pandemics. Nature Medicine, 2020, 26, 1328-1330.	15.2	20

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19	Trajectories of Adolescents Treated with Gonadotropin-Releasing Hormone Analogues for Gender Dysphoria. Archives of Sexual Behavior, 2020, 49, 2611-2618.	1.2	50
20	Parental experiences in endâ€ofâ€life decisionâ€making in allogeneic pediatric stem cell transplantation. Pediatric Blood and Cancer, 2020, 67, e28229.	0.8	10
21	Dealing with Moral Challenges in Treatment of Transgender Children and Adolescents: Evaluating the Role of Moral Case Deliberation. Archives of Sexual Behavior, 2020, 49, 2619-2634.	1.2	11
22	A systematic breakdown of the levels of evidence supporting the European Society of Cardiology guidelines. European Journal of Preventive Cardiology, 2019, 26, 1944-1952.	0.8	22
23	Physical Changes, Laboratory Parameters, and Bone Mineral Density During Testosterone Treatment in Adolescents with Gender Dysphoria. Journal of Sexual Medicine, 2019, 16, 1459-1468.	0.3	45
24	Use of Fertility Preservation Among a Cohort of Transgirls in the Netherlands. Journal of Adolescent Health, 2019, 64, 589-593.	1.2	56
25	Deferred consent for the enrolment of neonates in delivery room studies: strengthening the approach. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2019, 104, fetalneonatal-2018-316461.	1.4	7
26	Dilemma of Reporting Incidental Findings in Newborn Screening Programs for SCID: Parents' Perspective on Ataxia Telangiectasia. Frontiers in Immunology, 2019, 10, 2438.	2.2	19
27	Iron status and its association with HbA1c levels in Dutch children with diabetes mellitus type 1. European Journal of Pediatrics, 2018, 177, 603-610.	1.3	6
28	Pediatric oncology as a <scp>Learning Health System: Ethical</scp> implications for best available treatment protocols. Learning Health Systems, 2018, 2, e10052.	1.1	5
29	Participation in a single-blinded pediatric therapeutic strategy study for juvenile idiopathic arthritis: are parents and patient-participants in equipoise?. BMC Medical Ethics, 2018, 19, 96.	1.0	3
30	ESPE and PES International Survey of Centers and Clinicians Delivering Specialist Care for Children and Adolescents with Gender Dysphoria. Hormone Research in Paediatrics, 2018, 90, 326-331.	0.8	44
31	Standards of practice in empirical bioethics research: towards a consensus. BMC Medical Ethics, 2018, 19, 68.	1.0	62
32	Medical decision-making in children and adolescents: developmental and neuroscientific aspects. BMC Pediatrics, 2017, 17, 120.	0.7	216
33	On classifying the field of medical ethics. BMC Medical Ethics, 2017, 18, 30.	1.0	1
34	Perspectives of adolescents on decision making about participation in a biobank study: a pilot study. BMJ Paediatrics Open, 2017, 1, e000111.	0.6	11
35	Perceptions of Sex, Gender, and Puberty Suppression: A Qualitative Analysis of Transgender Youth. Archives of Sexual Behavior, 2016, 45, 1697-1703.	1.2	43
36	Balancing research interests and patient interests: A qualitative study into the intertwinement of care and research in paediatric oncology. Pediatric Blood and Cancer, 2015, 62, 816-822.	0.8	18

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37	Informed consent instead of assent is appropriate in children from the age of twelve: Policy implications of new findings on children's competence to consent to clinical research. BMC Medical Ethics, 2015, 16, 76.	1.0	106
38	Readability and Visuals in Medical Research Information Forms for Children and Adolescents. Science Communication, 2015, 37, 89-117.	1.8	13
39	Research information for minors: Suitable formats and readability. A systematic review. Journal of Paediatrics and Child Health, 2015, 51, 505-511.	0.4	14
40	Comic strips help children understand medical research. Patient Education and Counseling, 2015, 98, 518-524.	1.0	31
41	Early Medical Treatment of Children and Adolescents With Gender Dysphoria: An Empirical Ethical Study. Journal of Adolescent Health, 2015, 57, 367-373.	1.2	78
42	Why do children decide not to participate in clinical research: a quantitative and qualitative study. Pediatric Research, 2015, 78, 103-108.	1.1	17
43	Clinical Trial Decisions in Difficult Circumstances: Parental Consent Under Time Pressure. Pediatrics, 2015, 136, e983-e992.	1.0	32
44	ls a New Protocol for Acute Lymphoblastic Leukemia Research or Standard Therapy?. Pediatrics, 2015, 136, 566-570.	1.0	6
45	The Use of Dogmas in Pediatric Research Ethics. American Journal of Bioethics, 2015, 15, 18-19.	0.5	11
46	What Constitutes the Best Interest of a Child? Views of Parents, Children, and Physicians in a Pediatric Oncology Setting. American Journal of Bioethics Primary Research, 2013, 4, 1-10.	1.5	10
47	Ethical issues at the interface of clinical care and research practice in pediatric oncology: a narrative review of parents' and physicians' experiences. BMC Medical Ethics, 2011, 12, 18.	1.0	71
48	REFLECTIVE EQUILIBRIUM AND EMPIRICAL DATA: THIRD PERSON MORAL EXPERIENCES IN EMPIRICAL MEDICAL ETHICS. Bioethics, 2009, 24, 490-498.	0.7	34
49	A Case Against Something That Is Not the Case: The Groningen Protocol and the Moral Principle of Non-Maleficence. American Journal of Bioethics, 2008, 8, 29-31.	0.5	7