## Martine C De Vries

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Medical decision-making in children and adolescents: developmental and neuroscientific aspects. BMC Pediatrics, 2017, 17, 120.	1.7	216
2	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.	2.4	106
3	Early Medical Treatment of Children and Adolescents With Gender Dysphoria: An Empirical Ethical Study. Journal of Adolescent Health, 2015, 57, 367-373.	2.5	78
4	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.	2.4	71
5	Standards of practice in empirical bioethics research: towards a consensus. BMC Medical Ethics, 2018, 19, 68.	2.4	62
6	Use of Fertility Preservation Among a Cohort of Transgirls in the Netherlands. Journal of Adolescent Health, 2019, 64, 589-593.	2.5	56
7	Trajectories of Adolescents Treated with Gonadotropin-Releasing Hormone Analogues for Gender Dysphoria. Archives of Sexual Behavior, 2020, 49, 2611-2618.	1.9	50
8	Physical Changes, Laboratory Parameters, and Bone Mineral Density During Testosterone Treatment in Adolescents with Gender Dysphoria. Journal of Sexual Medicine, 2019, 16, 1459-1468.	0.6	45
9	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.	2.0	45
10	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.	1.8	44
11	Perceptions of Sex, Gender, and Puberty Suppression: A Qualitative Analysis of Transgender Youth. Archives of Sexual Behavior, 2016, 45, 1697-1703.	1.9	43
12	REFLECTIVE EQUILIBRIUM AND EMPIRICAL DATA: THIRD PERSON MORAL EXPERIENCES IN EMPIRICAL MEDICAL ETHICS. Bioethics, 2009, 24, 490-498.	1.4	34
13	Clinical Trial Decisions in Difficult Circumstances: Parental Consent Under Time Pressure. Pediatrics, 2015, 136, e983-e992.	2.1	32
14	Comic strips help children understand medical research. Patient Education and Counseling, 2015, 98, 518-524.	2.2	31
15	Parents' Perspectives and Societal Acceptance of Implementation of Newborn Screening for SCID in the Netherlands. Journal of Clinical Immunology, 2021, 41, 99-108.	3.8	25
16	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.	5.0	23
17	A systematic breakdown of the levels of evidence supporting the European Society of Cardiology guidelines. European Journal of Preventive Cardiology, 2019, 26, 1944-1952.	1.8	22
18	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.	1.0	22

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19	Assessing Medical Decision-Making Competence in Transgender Youth. Pediatrics, 2021, 148, .	2.1	21
20	The ethics of deferred consent in times of pandemics. Nature Medicine, 2020, 26, 1328-1330.	30.7	20
21	Dilemma of Reporting Incidental Findings in Newborn Screening Programs for SCID: Parents' Perspective on Ataxia Telangiectasia. Frontiers in Immunology, 2019, 10, 2438.	4.8	19
22	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.	1.5	18
23	Why do children decide not to participate in clinical research: a quantitative and qualitative study. Pediatric Research, 2015, 78, 103-108.	2.3	17
24	Research information for minors: Suitable formats and readability. A systematic review. Journal of Paediatrics and Child Health, 2015, 51, 505-511.	0.8	14
25	Fetoscopic myelomeningocoele closure: Is the scientific evidence enough to challenge the gold standard for prenatal surgery?. Prenatal Diagnosis, 2021, 41, 949-956.	2.3	14
26	Readability and Visuals in Medical Research Information Forms for Children and Adolescents. Science Communication, 2015, 37, 89-117.	3.3	13
27	The Use of Dogmas in Pediatric Research Ethics. American Journal of Bioethics, 2015, 15, 18-19.	0.9	11
28	Perspectives of adolescents on decision making about participation in a biobank study: a pilot study. BMJ Paediatrics Open, 2017, 1, e000111.	1.4	11
29	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.	1.9	11
30	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
31	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.	1.9	10
32	Parental experiences in endâ€ofâ€life decisionâ€making in allogeneic pediatric stem cell transplantation. Pediatric Blood and Cancer, 2020, 67, e28229.	1.5	10
33	When is it impractical to ask informed consent? A systematic review. Clinical Trials, 2022, 19, 545-560.	1.6	10
34	Reviewing recordings of neonatal resuscitation with parents. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2021, 106, 346-351.	2.8	9
35	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.	0.9	7
36	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.	2.8	7

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37	Is a New Protocol for Acute Lymphoblastic Leukemia Research or Standard Therapy?. Pediatrics, 2015, 136, 566-570.	2.1	6
38	Iron status and its association with HbA1c levels in Dutch children with diabetes mellitus type 1. European Journal of Pediatrics, 2018, 177, 603-610.	2.7	6
39	Pediatric oncology as a <scp>Learning Health System: Ethical</scp> implications for best available treatment protocols. Learning Health Systems, 2018, 2, e10052.	2.0	5
40	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.	1.5	5
41	How Traditional Informed Consent Impairs Inclusivity in a Learning Healthcare System: Lessons Learned from the Utrecht Cardiovascular Cohort. Journal of Clinical Epidemiology, 2022, , .	5.0	4
42	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.	2.4	3
43	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.	1.9	3
44	On classifying the field of medical ethics. BMC Medical Ethics, 2017, 18, 30.	2.4	1
45	The Duty to Support Learning Health Systems: A Broad Rather than a Narrow Interpretation. American Journal of Bioethics, 2021, 21, 14-16.	0.9	1
46	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.	2.8	1
47	Serum hepcidin concentrations in relation to iron status in children with type 1 diabetes. Pediatric Hematology and Oncology, 2021, 38, 108-123.	0.8	0
48	Antimicrobial guidelines in clinical practice: incorporating the ethical perspective. JAC-Antimicrobial Resistance, 2021, 3, dlab074.	2.1	0
49	Consent for Delivery Room Studies: What Can Be Learned from Perceptions of Parents. Neonatology, 2022, 119, 214-221.	2.0	0