

# Full list of Publications Karin Jongasma

## SCIENTIFIC ARTICLES:

1. Gerhards H, Jongasma K, Schicktanz S. (2017). The relevance of different trust models for representation in patient organizations: conceptual considerations. BMC Health Services Research, 17(1):474. DOI : 10.1186/s12913-017-2368-z
2. Jongasma KR and Sand M. (2017). The usual suspects - why techno-fixing Dementia is flawed. Medicine Health Care and Philosophy, 20:119–130.
3. Jongasma KR. (2016). Patientenverfügungen für die Demenzforschung aus ethischer Perspektive betrachtet. Ärzteblatt BW, 12, 612-615.
4. Sand M and Jongasma KR. (2016). Why neural determinism is not real determinism and why mental states cannot act. AJOB Neuroscience, 7(4), 205-207.
5. Jongasma KR. (2016). Advance Directives in Dementia Research - A medical ethical inquiry. ISBN: 978-94-6050-022-0
6. Jongasma KR, Sprangers M and van de Vathorst S. (2016). The implausibility of response shifts in dementia patients. Journal of Medical Ethics, 42:597–600
7. Sand M and Jongasma KR. (2016). Towards an Ageless Society. In: Dominguez-Rue and Nierling: Ageing and Technology: Perspectives of the Social Sciences. Transcript (Aging Studies, 9), pp. 291-310.
8. Andorno R, Gennet E, Jongasma KR and Elger B. (2016) Integrating Advance Research Directives into the European Legal Framework. 23, 49-64.
9. Jongasma KR, van Bruchem-Visser RL, van de Vathorst S and Mattace-Raso FUS. (2016). Has dementia research lost sense of reality? - A descriptive analysis of eligibility criteria of Dutch dementia research protocols. Netherlands Journal of Medicine, 74(5), 201-209
10. Jongasma KR and van de Vathorst S. (2015). Beyond Competence: Advance Directives in Dementia Research. Monash Bioethics, 33(2), 167-18.
11. Jongasma KR and van de Vathorst S. (2015). Dementia research and advance consent: it is not about critical interest. Journal of Medical Ethics; 41(8):708-709.
12. Jongasma KR, Bos W and van de Vathorst S. (2015). Morally relevant similarities and differences between children and dementia patients as research subjects - Representation in legal documents and ethical guidelines. Bioethics, 29, 9, 662-670.

13. Jongsma KR and van de Vathorst S. (2015). Advance directives in dementia research: The opinions and arguments of clinical researchers – an empirical study. *Research Ethics*; 11(1):4- 14

#### PUBLIC OUTREACH & POLICY PAPERS

14. Jongsma KR and Klar V. (2016). Empathy is an overrated skill when dispensing medical care. *AEON*. <https://aeon.co/ideas/empathy-is-an-overrated-skill-when-dispensing-medical-care>
15. Jongsma KR (2016). How people with sports addiction are like drug addicts. *AEON*. <https://aeon.co/ideas/how-people-with-sports-addiction-are-like-drug-addicts>
16. Aartsen JGM and Jongsma KR. (2015). Ouderenmishandeling en de rol van de mentor. *Podium voor Bio-ethiek*. 22(4), 12-14.
17. Struijs AJ en Jongsma KR. (2013). Gezamenlijke besluitvorming door zorgverlener en patiënt - normatieve achtergrond. *Centrum voor Ethiek en Gezondheid (CEG)*.
18. Jongsma KR. (2012). Lost your mind? Wilsonbekwame demente ouderen in de onderzoekscontext. *Podium voor Bio-Ethiek*. 19(4), 27-28.
19. Struijs AJ en Jongsma KR (ed). (2012). De mens centraal? - Ethische dilemma's bij gezondheidsbeleid met goede zorg voor dier en natuur. *Centrum voor Ethiek en Gezondheid (CEG); Raad voor de Leefomgeving en de Infrastructuur (RLI), de Raad voor Dierenaangelegenheden (RDA)*.

#### PRESENTATIONS ON CONFERENCES

1. Jongsma KR. ARDs in dementia research-what do they solve? Colloquium Münster, Department of Medical Ethics and History of Medicine, Bochum, 10.07.2017
2. Jongsma KR. ARDs in dementia research-what do they solve? Colloquium Bochum, Department of Medical Ethics and History of Medicine, Bochum, 30.01.2017
3. Jongsma KR, Späth E, Schicktanz S. Epistemic injustice in Autism Patient organizations. Workshop Collective Representation of Autism, 27.04.2017
4. Jongsma KR. Advance Research Directives- What do they solve? Minerva Workshop, Judean Hills, Israel, 04.06.2017

5. Jongsma KR. Speaker at panel: Decisional Competence. : Incompetent research participants: morally relevant differences and similarities between children and dementia patients  
OZSW Conference 2016 (Dutch Graduate school for Philosophy),  
Groningen 9-10.12.2016
6. Jongsma KR. The more the merrier? Concepts of representation and shared decision-making in health care policy.  
Conference: Shared decision-making in personalized medicine, Göttingen,  
5-6.12.2016
7. Jongsma KR. The relevance of different trust models for representation in patient organisations.  
Conference: Trust in Organisations, Copenhagen 24-25.11.2016
8. Jongsma KR. The implausibility of response shifts in dementia patients.  
IAB World Congres 2016 Edinburgh 14-17.06.2016
9. Jongsma KR & Schicktanz S. Empirical-ethical reflection on collective representation -Autism/Dementia and their representatives.  
IAB World Congres Edinburgh 14-17.06.2016, &  
International Workshop One for All, All for one 20.10.2016, Göttingen
10. Jongsma KR & Sand M. Techno fixing Dementia, what we can learn from STS and TA. Colloquium, Department of Medical Ethics and History of medicine, Mainz  
29.01.2016
11. Jongsma KR & Sand M. Towards an Ageless Society.  
Workshop Homo Senescens Hannover 1-3.10.2015
12. Jongsma KR. ARDs in dementia research-what do they solve?  
IAB World Congres Mexico City 25-28.06.2014
13. Jongsma KR, Bos W, van de Vathorst S. Morally relevant differences and similarities of vulnerable research populations: Representation in legal documents and ethical guidelines  
IAB World Congres Mexico City 25.28.06.2014
14. Jongsma KR: Advance Directives in dementia research  
University of Basel 29.01.2015  
University of Tübingen 13.05.2013
15. Jongsma KR. Have you lost your mind? - Dementia, identity and advance directives.

IAB World Congress Rotterdam, 26-29.06.2012

16. Jongsma KR. Advance Research Directives in Dementia Research: Ethical and Legal Considerations.

PGBC conference, Kings College, London, 22-24.05.2013

17. Jongsma KR & Bos W: Regulation of research with vulnerable groups.

Dutch Central Committee on Research with Human Subjects (CCMO). 24-09.2013