

Publications in Genetic Risk and Risk Information from the Working Group: Cultural and Ethical Aspects in Biomedicine

Mind the Risk Project Publications in Genetic Risk and Risk Information, Updated September 2018

Publications are listed from newest to oldest. Hyperlinks lead to full version of the article.

2018

S. Wöhlke, J. Perry, M. Schaper, S. Schicktanz: How uncertainty influences lay people's attitudes and risk perceptions concerning predictive genetic testing and risk communication (2018 under submission)

S. Wöhlke, Z. Alpınar Sencan, J. Perry: What triggers responsibility in lay people when dealing with genetic risk information? (2018 in preparation)

S. Wöhlke, M. Falahee, E.G. Gonzalez, A. Raz, K. Raza, M. Schaper, D. Timmermans, S. Schicktanz (2018): Public attitudes on responsibility in gene-ethics: results of a comparative focus group study in Germany, Israel, UK and Netherlands (2018 in preparation)

M. Schaper, S. Wöhlke, M. Falahee, E.G. Gonzalez, A. Raz, K. Raza, M. D. Timmermans, S. Schicktanz: Direct-to-Consumer Genetic Testing (DTC GT) in Germany, UK, Israel and the Netherlands – A Cross-national Focus Group Study of Lay Perspectives (2018 in preparation)

M. Schaper, S. Wöhlke, S. Oliveri, G. Pravettoni, S. Schicktanz: Direct access to personal genomic data – A quantitative survey of user perspectives (2018 in preparation)

M. Schaper, S. Hansen, S. Schicktanz (2018): Überreden für die gute Sache? Techniken öffentlicher Gesundheitskommunikationen und ihre ethischen Implikationen. Ethik in der Medizin (accepted)

[M. Schaper, S. Schicktanz \(2018\): Medicine, Market and Communication: Ethical Considerations in Regard to Persuasive Communication in Direct-to-Consumer Genetic Testing Services. BMC Medical Ethics](#)

[M. Schaper, S. Wöhlke, S. Schicktanz \(2018\): "I would rather have it done by a doctor"- laypeople's perceptions of direct-to-consumer genetic testing \(DTC GT\) and its ethical implications. Medicine, Healthcare and Philosophy](#)

[S. Wöhlke, J. Perry, S. Schicktanz \(2018\): Physicians' communication patterns for motivating colorectal cancer patients to biomarker research: empirical insights and ethical issues, Clinical Ethics \(accepted 4/2018\)](#)

2017

[J. Leefmann, M. Schaper, S. Schicktanz \(2017\): The concept of "Genetic Responsibility" and its meanings: A systematic review of qualitative medical sociology literature, Frontiers in Sociology ELSI in Science and Genetics.](#)

[J. Inthorn \(2017\). Genetic risk information. Journal of Risk Research, 21\(2\), 109-116.](#)

[Schicktanz, S. \(2016\). Genetic risk and responsibility: reflections on a complex relationship. Journal of Risk Research, 1-23.](#)

2016

[J. Perry, S. Wöhlke, A. Heßling & S. Schicktanz \(2016\): Why take part in personalized oncology research? Genetic misconception and genetic responsibility as critical issues from an ethical perspective. Journal of Oncology.26:e12563.](#)

[S. Schicktanz & A. Raz \(Eds.\) \(2010\): New Genetics and Society Special Issue: *Genetics and Responsibilities: Cultural perspectives, public discourses and ethical issues.*\(21:2\).](#)

[A. E. Raz & S. Schicktanz \(2016\): *Comparative Empirical Bioethics: Dilemmas of Genetic Testing and Euthanasia in Israel and Germany*, Berlin, New York, Springer](#)

Master's Thesis: Julia Perry (2016): „Dealing with the complexity of medical information in colorectal cancer treatment: Awareness, needs, and uncertainty concerning treatment procedure and participation in biomarker research – a qualitative study of ten patients in longitudinal section“ (150 pages)

2015

[S. Wöhlke, J. Perry & S. Schicktanz \(2015\): Taking it Personally: Patients' Perspectives on Personalised Medicine and its Ethical Relevance. In: Jochen Vollmann, Verena Sandow, Sebastian Wäscher, & Jan Schildmann \(Eds.\): *Personalised Medicine: Ethical, Medical, Economic and Legal Critical Perspectives*, ASHGATE, 129-149. 2](#)

2014

[A. Hessling \(2014\): Exploration medizinethischer Implikationen individualisierter Medizin beim lokal fortgeschrittenen Rektumkarzinom aus Sicht von Ärzten und Forschern. Eine empirisch-ethische Untersuchung.](#)

[J. Inthorn, P. Wehling, S. Schultz, S. Schicktanz \(2014\): *Präkonzeptionelle Anlageträgertests: Diagnostik mit Fragezeichen* \[preconceptional carrier test: a questionable diagnostic\]. Deutsches Ärzteblatt 111\(9\): A-343 / B-300 / C-285.](#)

[A. Raz, N. Rimon-Zarfaty, J. Inthorn, S. Schicktanz \(2014\): *Making Responsible Life Plans: Cultural Differences in Lay Attitudes toward Predictive Genetic Testing for Late-Onset Diseases*. In: S. Schicktanz, B. Prainsack, G. Werner-Felmayer \(ed.\): *Genetics as Social Practice. Transdisciplinary Views on Science and Culture*. Ashgate, 181-198.](#)

[S. Schicktanz \(2014\): „Individualisierte Patientenversorgung“. *Große Hoffnungen und viele offene Fragen*. In: Gerechte Gesundheit. Newsletter zur Verteilungsdebatte 26, 2-3.](#)

[S. Schicktanz mit B. Prainsack, G. Werner-Felmayer \(2014\): *Genetics as Social Practice. Transdisciplinary Views on Science and Culture*. ASHGATE.](#)

[S. Schicktanz & S. Wöhlke \(2014\): Assessing the personal perspective. In: *International Innovation 1: 18-20*.](#)

2013

[S. Wöhlke, A. Hessling, S. Schicktanz \(2013\): Wenn es persönlich wird in der „personalisierten Medizin“: Aufklärung und Kommunikation aus klinischer Forscher- und Patientenperspektive im empirisch-ethischen Vergleich Ethik in der Medizin. Ethik in der Medizin 23\(3\): 215-222.](#)

2012

A. Hessling (2012): Everything better than 50% is better than now. In: P. Dabrock, M. Braun, J. Ried (Eds.): Individualized Medicine between hype and hope. Exploring ethical and societal challenges for healthcare, Lit-Verlag: Münster, 111-135.

[S. Schicktanz, A. Hessling \(2012\): What German experts expect from individualised medicine: Problems of uncertainty and future complication in physician-patient-interaction. Clinical Ethics 7\(2\): 86-93.](#)

2009

[S. Schicktanz, A. Raz \(2009\): Diversity and Uniformity in Genetic Responsibility: Moral Attitudes of Patients, Relatives and Lay People in Germany and Israel. Medicine, Health Care and Philosophy 12\(4\): 433-442.](#)

[S. Schicktanz, A. Raz \(2009\): Lay Perceptions of Genetic Testing in Germany and Israel: The Interplay of National Culture and Individual Experience. New Genetics and Society 28\(4\): 401-414.](#)