Research outputs

High risk, mixed reward: Making genetic test results actionable in cardiology

Cultivating awareness of donation in cutting-edge allogenic cell therapies

A test of faith? Attitudes of ultraorthodox Jewish parents of children with down syndrome toward prenatal testing

Ethical and social implications of public–private partnerships in the context of genomic/big health data collection

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Non-invasive prenatal testing (NIPT): is routinization problematic?

Old and new challenges regarding comparable and viable data sharing in population-scale genomic research

Genomic uncertainty and genetic counsellors’ professional authority

AI-driven risk scores: should social scoring and polygenic scores based on ethnicity be equally prohibited?

Considering the alternatives: lessons from Israel’s meat substitutes initiatives

Editorial: Data-intensive medicine and healthcare: ethical and social implications in the era of artificial intelligence and automated decision-making
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Can not wanting to know be responsible? Conceptual analysis and meanings of not-knowing in Israeli and German prenatal genetic practices

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Challenges for precision public health communication in the era of genomic medicine

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Comparing Germany and Israel regarding debates on policy-making at the beginning of life: PGD, NIPT and their paths of routinization
Fragmented responsibility: views of Israeli HCPs regarding patient recontact following variant reclassification

"Donating with eyes shut": attitudes regarding DNA donation to a large-scale biobank in Israel

Collective representation and the founders' culture in non-profit organisations: The case of Israel's national association for autism

Who Takes Part in the Political Game? The Sex Work Governance Debate in Israel

One Size Does Not Fit All: Lessons from Israel's Covid-19 Vaccination Drive and Hesitancy

Unmet communication needs and moral work in the disposition decision concerning surplus frozen embryos: The perspectives of IVF users

Patient Advocacy in Dementia: The Culture and Ethics of Policy-Making and

The mask of autism: Social camouflaging and impression management as coping/normalization from the perspectives of autistic adults

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Divergent evolution of newborn screening: Israel and the US as gene worlds

One For All, All For One? Collective Representation in Healthcare Policy

Patient Representation and Advocacy for Alzheimer Disease in Germany and Israel

Doing gender in segregated and assimilative organizations: Ultra-Orthodox Jewish women in the Israeli high-tech labour market

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Carrier matching and collective socialization in community genetics: Dor Yeshorim and the reinforcement of stigma

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