Research outputs

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

Views of Israeli healthcare professionals regarding communication of genetic variants of uncertain significance to patients

Views on disability and prenatal testing among families with Down syndrome and disability activists: A comparative analysis of interviews from Germany and Israel

Vergleichende Untersuchung der Debatten in Deutschland und in Israel über politische Entscheidungsprozesse am Lebensanfang: PID, NIPT und ihr Weg zur Routinisierung

Fragmented responsibility: views of Israeli HCPs regarding patient recontact following variant reclassification

A test of faith? Attitudes of ultraorthodox Jewish parents of children with down syndrome toward prenatal testing
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Cancer patients' understandings of genetic variants of uncertain significance in clinical care

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

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

Frozen: social and bioethical aspects of cryopreservation

Correction: Population screening for BRCA1/BRCA2 founder mutations in Ashkenazi Jews: proactive recruitment compared with self-referral (Genetics in Medicine, (2017), 19, 7, (754-762), 10.1038/gim.2016.182)

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

Transparency, consent and trust in the use of customers’ data by an online genetic testing company: an Exploratory survey among 23andMe users

Direct to Consumer Personal Genomic Testing and Trust: A Comparative Focus Group Study of Lay Perspectives in Germany, Israel, the Netherlands and the UK

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

Atitudes of Israeli parents of children with Down syndrome toward non-invasive prenatal screening and the scope of prenatal testing

Pandora’s pregnancy: NIPT, CMA, and genome sequencing—A new era for prenatal genetic testing

Coming to terms with the imperfectly normal child: attitudes of Israeli parents of screen-positive infants regarding subsequent prenatal diagnosis

Epigenetic metaphors: an interdisciplinary translation of encoding and decoding

Familial communication and cascade testing among relatives of BRCA population screening participants
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

Parents like me: biosociality and lay expertise in self-help groups of parents of screen-positive newborns

Representing autism: Challenges of collective representation in German and Israeli associations for and of autistic people

A symbolic interactionist user’s guide to the answering machine: 22 reflections on vocal encounters in an emerging social world

Reckless or pioneering? public health genetics services in Israel: Public health genetics services in Israel

Population screening for BRCA1/BRCA2 founder mutations in Ashkenazi Jews: Proactive recruitment compared with self-referral

Population screening for BRCA1/BRCA2 mutations: Lessons from qualitative analysis of the screening experience

Saving or Subordinating Life? Popular Views in Israel and Germany of Donor Siblings Created through PGD

Beyond cultural stereotyping: views on end-of-life decision making among religious and secular persons in the USA, Germany, and Israel

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Risks and Responsibilities: Making Plans for Life and Death

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*Cousin marriages: Between Tradition, Genetic Risk and Cultural Change*

Making responsible life plans: Cultural differences in lay attitudes toward predictive genetic testing for late-onset diseases

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