Research outputs

Can not wanting to know be responsible? Conceptual analysis and meanings of not-knowing in Israeli and German prenatal genetic practices

Commentary - "Yes, but..." vs. "no, but...": Ambivalences towards prenatal diagnosis in Israel and Germany

Comparison through conversation: Thinking with different differences

PND in Israel: Public health services and uptake in cultural context

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

What does prenatal testing mean for women who have tested?

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

Cancer patients’ understandings of genetic variants of uncertain significance in clinical care

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

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

Genomic uncertainty and genetic counsellors' professional authority

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 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

Attitudes 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

What a difference a role makes: Occupational and organizational characteristics related to the HR strategic role among human resource managers

Donation of surplus frozen pre-embryos to research in Israel: Underlying motivations

Applying the Theoretical Tools: Being Affected, Responsibility, and Risk

Contextualizing the Cultural and Medico-Legal Debate on Adult Genetic Testing

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Making Responsible Life Plans: Cultural Differences in Lay Attitudes in Germany and Israel Towards Predictive Genetic Testing for Late-Onset Diseases

Planning One’s End of Life in an Expert Biomedical Culture
Risks and Responsibilities: Making Plans for Life and Death

Setting the Methodological Scene: The Value of Explication and Pluralization of Moral Grammars

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Premarital carrier testing and matching in Jewish communities

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

Successful implementation of large-scale drip irrigation projects: An exploratory study of the socio-economic impact of the apmip among smallholders in India

Exploring the positions of German and Israeli patient organizations in the bioethical context of end-of-life policies
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"The Most Important Test You'll Ever Take"? Attitudes toward confidential carrier matching and open individual testing among modern-religious Jews in Israel

When does a fetus become a person? An Israeli viewpoint

The cultural context of patient's autonomy and doctor's duty: Passive euthanasia and advance directives in Germany and Israel

The cultural context of end-of-life ethics: A comparison of Germany and Israel

Between social hypocrisy and social responsibility: Professional views of eugenics, disability and repro-genetics in Germany and Israel

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Abortion committees as agents of eugenics: Medical and public views on selective abortion following mild or likely fetal pathology

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Can population-based carrier screening be left to the community?

Diversity and uniformity in genetic responsibility: moral attitudes of patients, relatives and lay people in Germany and Israel.

A Life (un)Worthy of Living:

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Poetry of unadulterated imagination: The late style of akira kurosawa

The Right Blend: Integrating Female Cadets in the Israeli Air Force—An Organizational Culture Perspective

Communities of practice or communities of coping? Employee compliance among CSRs in Israeli call centres


Ambiguous professionalism: Managing efficiency and service quality in an Israeli call centre

"We came to talk with the people behind the disease." Communication and control in medical education

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The gene and the genie: tradition, medicalization, and genetic counseling in a Bedouin community in Israel

Upright generations of the future: Tradition and medicalization in community genetics

Cousin marriage and premarital carrier matching in a Bedouin community in Israel: Attitudes, service development and educational intervention

"Important to test, important to support": Attitudes toward disability rights and prenatal diagnosis among leaders of support groups for genetic disorders in Israel

Between acculturation and ambivalence: Knowledge of genetics and attitudes towards genetic testing in a consanguineous Bedouin community

'Āysha': Genetics education and community engagement in a consanguineous Arab-Bedouin population in Israel

Nondirectiveness and its lay interpretations: The effect of counseling style, ethnicity and culture on attitudes towards genetic counseling among Jewish and bedouin respondents in Israel

Status disclosure: Genetic counseling as an arena for negotiation

The Slanted Smile Factory: Emotion Management in Tokyo Disneyland

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The Hybridization of Organizational Culture in Tokyo Disneyland

The Slanted Smile Factory: Emotion Management in Tokyo Disneyland

The authorized self: How middle age defines old age in the postmodern

'America' Meets 'Japan': A Journey for Real between Two Imaginaries

The mask of dementia: Images of 'demented residents' in a nursing ward

Life Stories, Status and Symbolic Typing: Exploring Beggars as an Interpretive Community

Rituals of Exchange In the Social World of Israeli Beggars: An Exploratory Study

The Discourse of Aging and Other Age-Related Languages: Social Identity and Life-Span Communication


Rewriting the Holocaust: an Israeli Case Study in the Sociology of the Novel: an Israeli case study in the sociology of the novel
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