Research outputs

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


**Prediction and explainability in AI: Striking a new balance?**

Raz, A., Heinrichs, B., Avnoon, N., Eyal, G. & Inbar, Y., 1 Jan 2024, In: Big Data and Society. 11, 1

**The interplay of ethics and genetic technologies in balancing the social valuation of the human genome in UNESCO declarations**


**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**


**Engaging patients in identifying risk factors for ALS**


**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

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 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

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

Testing Fate: Tay-Sachs and the Right to be Responsible: by Shelley Z. Reuter (book review)

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

Comparative Empirical Bioethics: Dilemmas of Genetic Testing and Euthanasia in Israel and Germany
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Applying the Theoretical Tools: Being Affected, Responsibility, and Risk

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

Final Conclusion: Disentangling the Micro and the Macro in Bioethics

Introduction: Engaging in Comparative Bioethics

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Planning One’s End of Life in an Expert Biomedical Culture

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

Lay Attitudes Towards End-of-Life Decision-Making in Germany and Israel

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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 

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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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Through the looking glass: Engaging in a socio-ethical, cross-cultural dialogue

 Abortions for fetuses with mild abnormalities

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

 Transplanting management: Participative change, organizational development, and the glocalization of corporate culture

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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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 Carrier matching and collective socialization in community genetics: Dor Yeshorim and the reinforcement of stigma
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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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Alternative Forms, Same Emotions? Organizational Culture and Labor Relations in Contemporary Japan

A note on inter-viewing: Using symbolic interactionism for interview analysis

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The Gene and the Genie: Tradition, Medicalization and Genetic Counseling in a Bedouin Community in Israel

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

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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

The changing face of Japanese retail: Working in a chainstore
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The orient strikes back: A global view of cultural display.


Domesticating Disney: Onstage strategies of adaptation in Tokyo Disneyland

Riding the Black Ship: Japan and Tokyo Disneyland

Glocalization and Symbolic Interactionism

The Hybridization of Organizational Culture in Tokyo Disneyland
The Slanted Smile Factory: Emotion Management in Tokyo Disneyland

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

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

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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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Vibrational Communication in Subterranean Mole Rats (Spalax ehrenbergi)