

Kaori Muto

List of Publications by Year in descending order

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Version: 2024-02-01

39
papers

1,301
citations

516710

16
h-index

414414

32
g-index

46
all docs

46
docs citations

46
times ranked

2402
citing authors

#	ARTICLE	IF	CITATIONS
1	Overview of the BioBank Japan Project: Study design and profile. <i>Journal of Epidemiology</i> , 2017, 27, S2-S8.	2.4	451
2	Japanese citizens' behavioral changes and preparedness against COVID-19: An online survey during the early phase of the pandemic. <i>PLoS ONE</i> , 2020, 15, e0234292.	2.5	147
3	Cross-sectional analysis of BioBank Japan clinical data: A large cohort of 200,000 patients with 47 common diseases. <i>Journal of Epidemiology</i> , 2017, 27, S9-S21.	2.4	133
4	Impacts of anxiety and socioeconomic factors on mental health in the early phases of the COVID-19 pandemic in the general population in Japan: A web-based survey. <i>PLoS ONE</i> , 2021, 16, e0247705.	2.5	55
5	Overview of BioBank Japan follow-up data in 32 diseases. <i>Journal of Epidemiology</i> , 2017, 27, S22-S28.	2.4	47
6	Study design and baseline characteristics of a population-based prospective cohort study of dementia in Japan: the Japan Prospective Studies Collaboration for Aging and Dementia (JPSC-AD). <i>Environmental Health and Preventive Medicine</i> , 2020, 25, 64.	3.4	47
7	Characteristics and prognosis of Japanese colorectal cancer patients: The BioBank Japan Project. <i>Journal of Epidemiology</i> , 2017, 27, S36-S42.	2.4	38
8	Demographic and lifestyle factors and survival among patients with esophageal and gastric cancer: The BioBank Japan Project. <i>Journal of Epidemiology</i> , 2017, 27, S29-S35.	2.4	32
9	Survey on the perception of germline genome editing among the general public in Japan. <i>Journal of Human Genetics</i> , 2018, 63, 745-748.	2.3	31
10	Ethical concerns on sharing genomic data including patients' family members. <i>BMC Medical Ethics</i> , 2018, 19, 61.	2.4	30
11	Characteristics and prognosis of Japanese female breast cancer patients: The BioBank Japan project. <i>Journal of Epidemiology</i> , 2017, 27, S58-S64.	2.4	27
12	New medical big data for P4 medicine on allergic conjunctivitis. <i>Allergology International</i> , 2020, 69, 510-518.	3.3	27
13	Statin use and all-cause and cancer mortality: BioBank Japan cohort. <i>Journal of Epidemiology</i> , 2017, 27, S84-S91.	2.4	25
14	Survival of macrovascular disease, chronic kidney disease, chronic respiratory disease, cancer and smoking in patients with type 2 diabetes: BioBank Japan cohort. <i>Journal of Epidemiology</i> , 2017, 27, S98-S106.	2.4	20
15	Symptom-based stratification for hay fever: A crowdsourced study using the smartphone application AllerSearch. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2021, 76, 3820-3824.	5.7	19
16	Individual characteristics and associated factors of hay fever: A large-scale mHealth study using AllerSearch. <i>Allergology International</i> , 2022, 71, 325-334.	3.3	18
17	Characteristics of patients with liver cancer in the BioBank Japan project. <i>Journal of Epidemiology</i> , 2017, 27, S43-S48.	2.4	17
18	Characteristics and prognosis of Japanese male and female lung cancer patients: The BioBank Japan Project. <i>Journal of Epidemiology</i> , 2017, 27, S49-S57.	2.4	17

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19	Lessons for reviewing clinical trials using induced pluripotent stem cells: examining the case of a first-in-human trial for age-related macular degeneration. <i>Regenerative Medicine</i> , 2018, 13, 123-128.	1.7	15
20	Serum glucose, cholesterol and blood pressure levels in Japanese type 1 and 2 diabetic patients: BioBank Japan. <i>Journal of Epidemiology</i> , 2017, 27, S92-S97.	2.4	12
21	Risk prediction models for mortality in patients with cardiovascular disease: The BioBank Japan project. <i>Journal of Epidemiology</i> , 2017, 27, S71-S76.	2.4	11
22	Clinical and histopathological characteristics of patients with prostate cancer in the BioBank Japan project. <i>Journal of Epidemiology</i> , 2017, 27, S65-S70.	2.4	11
23	Organ Transplantation as a Family Issue: Living Liver Donors in Japan. <i>International Journal of Japanese Sociology</i> , 2010, 19, 35-48.	0.6	9
24	“Tell me what you suggest, and let me do that, doctor” Patient deliberation time during informal decision-making in clinical trials. <i>PLoS ONE</i> , 2019, 14, e0211338.	2.5	9
25	Attitudes toward genomic tumor profiling tests in Japan: patients, family members, and the public. <i>Journal of Human Genetics</i> , 2019, 64, 481-485.	2.3	6
26	Noncompliance with Human Subjects’ Protection Requirements as a Reason for Retracting Papers: Survey of Retraction Notices on Medical Papers Published from 1981 to 2011. <i>Accountability in Research</i> , 2016, 23, 123-135.	2.4	4
27	Japanese insurers’ attitudes toward adverse selection and genetic discrimination: a questionnaire survey and interviews with employees about using genetic test results. <i>Journal of Human Genetics</i> , 2021, 66, 539-542.	2.3	4
28	Risk perception of pregnancy promotes disapproval of gestational surrogacy: analysis of a nationally representative opinion survey in Japan. <i>International Journal of Fertility & Sterility</i> , 2011, 5, 78-85.	0.2	4
29	Public attitudes in the clinical application of genome editing on human embryos in Japan: a cross-sectional survey across multiple stakeholders. <i>Journal of Human Genetics</i> , 2022, 67, 541-546.	2.3	4
30	Cholesterol levels of Japanese dyslipidaemic patients with various comorbidities: BioBank Japan. <i>Journal of Epidemiology</i> , 2017, 27, S77-S83.	2.4	3
31	The Timing of Decision-Making and Informed Consent ‘1/4’ Patients’ Perspective and Experiences of Clinical Trials in Japan . <i>Japanese Journal of Clinical Pharmacology and Therapeutics</i> , 2017, 48, 31-39.	0.1	3
32	The use of human samples obtained during medicolegal autopsies in research: An introduction to current conditions and initiatives in Japan. <i>Medicine, Science and the Law</i> , 2017, 57, 75-83.	1.0	2
33	“Telling” and assent: Parents’ attitudes towards children’s participation in a birth cohort study. <i>Health Expectations</i> , 2018, 21, 358-366.	2.6	2
34	The Role of “Experienced Subjects” in the Implementation of Patient and Public Involvement Policy in Japan. <i>Iryo To Shakai</i> , 2018, 28, 129-139.	0.1	2
35	Ethical, legal and social implications of human genome studies in radiation research: a workshop report for studies on atomic bomb survivors at the Radiation Effects Research Foundation. <i>Journal of Radiation Research</i> , 2021, 62, 656-661.	1.6	1
36	A Preliminary Study Exploring Japanese Public Attitudes Toward the Creation and Utilization of Human-Animal Chimeras: a New Perspective on Animals Containing “Human Material” (ACHM). <i>Asian Bioethics Review</i> , 2017, 9, 211-228.	1.3	0

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37	Letter in reply to: "therapeutic misconception and the role of the Research Ethics Committee". Regenerative Medicine, 2019, 14, 719-720.	1.7	0
38	XVI. COVID-19 and Ethical, Legal and Social Implications: Prejudice, Discrimination and Risk Communication. The Journal of the Japanese Society of Internal Medicine, 2020, 109, 2334-2338.	0.0	0
39	Efforts to Prevent Stigmatizing and Discriminatory Behaviors Related to COVID-19 in Japan. Iryo To Shakai, 2022, 32, 83-93.	0.1	0