Maternal age and birth defects after the use of assisted reproductive technology in Japan, 2004–2010

Syuichi Ooki

Department of Health Science,
Ishikawa Prefectural Nursing
University, Ishikawa, Japan

Background: Older mothers are becoming more common in Japan. One reason for this is the widespread use of assisted reproductive technology (ART). This study assesses the relationship between maternal age and the risk of birth defects after ART.

Methods: Nationwide data on ART between 2004 and 2010 in Japan were analyzed. Diseases that were classified as code Q00-Q99 (ie, congenital malformations, deformations, and chromosomal abnormalities) in the International Classification of Diseases, tenth edition, were selected. There were 219,185 pregnancies and 153,791 live births in total ART. Of these, 1943 abortions, stillbirths, or live births with birth defects were recorded. Percentage of multiple birth defects in total birth defects, the prevalence, crude relative risk and 95% confidence interval per 10,000 pregnancies and per 10,000 live births were analyzed according to the maternal age class (ie, 25–29, 30–34 (reference), 35–39, and 40+ years).

Results: Multiple birth defects were observed among 14% of the 25–29 year old class, and 8% among other classes when chromosomal abnormalities were excluded. The prevalence of chromosomal abnormalities per pregnancy and per live birth became significantly and rapidly higher in mothers in the age classes of 30–35 and 40+ years. Nonchromosomal birth defects per pregnancy decreased linearly with advanced maternal age, while the number of nonchromosomal birth defects per live birth formed a gradual U-shaped distribution. The prevalence per pregnancy of congenital malformations of the nervous system was significantly lower with advanced maternal age. The relative risk per live birth was significant regarding congenital malformations of the circulatory system for a maternal age of 40+ years. Some other significant associations between maternal age and birth defects were observed.

Conclusion: Maternal age is associated with several birth defects; however, older maternal age in itself does not produce noticeable extra risk for nonchromosomal birth defects overall.

Keywords: birth defects, assisted reproductive technology, maternal age, pregnancy, live births, epidemiologic prevalence study

Introduction

The percentage of mothers over 40 years old has been increasing rapidly, peaking in 2011 at 3.6%. The recent tendency in Japan to marry later is a likely factor in this increase, but so is the widespread use of fertility treatment, including assisted reproductive technology (ART). According to the Japanese ART statistics presented by the Japan Society of Obstetrics and Gynecology (JSOG), the percentage of total ART live births itself increased from 0.2% in 1992 to 2.7% in 2010. The percentages

Correspondence: Syuichi Ooki
Department of Health Science, Ishikawa
Prefectural Nursing University,
1-1, Gakuendai, Kahoku,
Ishikawa, 929-1210, Japan
Tel/Fax +81 762 818 377
Email sooki@ishikawa-nu.ac.jp

of mothers of live birth neonates who were over 40 years of age in the general population and in the ART population were 2.6% and 10.1% in 2008 and 3.6% and 12.2% in 2010, respectively, a finding that supports the supposition of a recent tendency to marry later due to the availability of fertility treatment.

It is well established that the increasing maternal age has contributed to some adverse obstetric outcomes,^{1–8} including birth defects.^{9,10} The effect of older maternal age on obstetric outcomes may be pronounced in the ART population. Both lower and higher maternal age may pose increased risks for birth defects.^{11–18} For example, older mothers have a higher risk of chromosomal abnormalities,^{19–22} such as Down syndrome,^{23–25} but whether they are at excess risk of nonchromosomal birth defects is less clear.¹⁷ It is important to have accurate information on maternal age-specific risks of birth defects after ART to gauge the implications of this rise in maternal age for public health, to gauge clinical care needs, and to provide information to couples of childbearing age.¹⁶

The purpose of the present epidemiologic prevalence study was to analyze birth defects after ART according to

the maternal age class and to examine the excess risk of nonchromosomal birth defects among older mothers.

Materials and methods

Outline of Japanese birth defects data after ART

The method for collecting data is described in detail elsewhere.^{26–29} Almost all medical institutions performing ART are registered with the JSOG. The JSOG administers questionnaire surveys for these medical institutions. Some of the survey data are presented in simple annual reports of aggregate, not individual, data.

From 2004 to 2010 (the latest), the individual list of all ART pregnancies resulting in birth defects was presented in the annual reports. The presented items are shown in Table 1. The mean response rate throughout the 7 years was 99.0% (4233/4274), meaning that a nearly complete database reflecting the current situation of ART and birth defects in Japan could be constructed. The methods of fertility treatment were divided into in vitro fertilization, microinsemination, frozen embryo transfer and others (duplicative methods), and do not include simple ovulation stimulation/enhancement. All methods of

Table 1 Demographic and perinatal outcome data of ART pregnancies with birth defect

		Maternal age class		25–29 (n = 171)		30–34 (n = 658)		35–39 (n = 796)		40+ (n = 292)		P ^a
		n	%	n	%	n	%	n	%			
Method of treatment	IVF	34	19.9	166	25.2	176	22.1	71	24.3	0.508		
	ICSI	47	27.5	136	20.7	179	22.5	63	21.6			
	IVF and ICSI	7	4.1	25	3.8	32	4.0	6	2.1			
	Frozen embryo transfer	83	48.5	330	50.2	408	51.3	152	52.1			
	Unknown/missing values	0	0.0	1	0.2	1	0.1	0	0.0			
Blastocyst transfer	Yes	94	55.0	365	55.5	420	52.8	156	53.4	0.731		
	No	76	44.4	289	43.9	373	46.9	135	46.2			
	Unknown/missing values	1	0.6	4	0.6	3	0.4	1	0.3			
Gestational weeks	Range	11–42		10–42		13–42		13–42				
	Mean ± SD	36.5 ± 5.6		36.1 ± 6.0		36.4 ± 5.1		35.8 ± 5.6				
	Median	38		38		38		37				
	Unknown/missing values	15	8.8	57	8.7	90	11.3	62	21.2			
Sex (fetuses/neonates)	Male	93	54.1	329	49.3	397	49.4	109	37.1	0.046		
	Female	58	33.7	260	38.9	307	38.2	119	40.5			
	Unknown/missing values	21	12.2	79	11.8	100	12.4	66	22.4			
Perinatal outcome (fetuses/neonates)	Abortion (<22 weeks)	8	4.7	42	6.3	36	4.5	15	5.1	0.394		
	Stillbirths (≥22 weeks)	2	1.2	13	1.9	15	1.9	7	2.4			
	Live births	153	89.0	566	84.7	684	85.1	217	73.8			
	Unknown/missing values	9	5.2	47	7.0	69	8.6	55	18.7			
Abortion	Spontaneous	0	0	2	4.8	1	2.8	1	6.7			
	Artificial	7	87.5	28	66.7	25	69.4	11	73.3			
	Unknown/missing values	1	12.5	12	28.6	10	27.8	3	20.0			
Early neonatal death (neonatal death up to day 6 after birth)	Yes	5	3.3	18	3.2	30	4.4	12	5.5	0.264		
	No	113	73.9	413	73.0	497	72.7	134	61.8			
	Unknown/missing values	35	22.9	135	23.9	157	23.0	71	32.7			

Notes: ^aComparison among all four groups by the χ^2 test. Unknown/missing values were excluded from the statistical tests.

Abbreviations: ART, assisted reproductive technology; n, number; IVF, in vitro fertilization; ICSI, intracytoplasmic sperm injection; SD, standard deviation.

fertility treatment were treated as ART in the present secondary data analyses. There were a total of 219,185 pregnancies and 153,791 live births between 2004 and 2010.

Diseases that were classified under the International Classification of Diseases, tenth edition (ie, ICD-10, 2003 version) codes Q00-Q99 (ie, congenital malformations, deformations, and chromosomal abnormalities) were regarded as birth defects in the present study. Other congenital diseases that were not classified under Q00-Q99, such as congenital hypothyroidism, were excluded. In total, 1943 abortions, stillbirths, or live births with birth defects (number of fetuses or neonates) consisting of 1637 (84.3%) singletons, 292 (15.0%) twins, nine (0.5%) triplets, and five (0.3%) unknown plurality were included.

Maternal age was distributed normally from 23–46 years, with a mean of 35.2 years, a standard deviation of 4.1 years, a median of 35 years, and a mode of 36 years. The maternal age classes used were as follows: 25–29, 30–34, 35–39, and 40+ years. Patients with a maternal age less than or equal to 24 ($n = 5$) were excluded. Thus, 1938 fetuses/neonates from 1917 mothers were analyzed. The numbers of pregnancies and live births according to maternal age class were estimated using the age distributions (percentage of each maternal age class) from the 2008 and 2010 data (the only published data). Regression lines for the percentage of each maternal age class were calculated using data from both years, and applied to other years.

Statistical analyses

First, demographic and perinatal outcome data were analyzed according to the maternal age class. Then, the percentages of multiple birth defects were analyzed. Multiple birth defects are defined as two or more unrelated major structural malformations that cannot be explained by an underlying syndrome or sequence.³⁰ Multiple birth defects were analyzed using European Surveillance of Congenital Anomalies (EUROCAT) algorithms,³⁰ that was based on the ICD-10 codes. This algorithm gives explicit coding rules for classification and picks out a small subset of cases of potential multiple birth defects. The etiologic classification was translated into an algorithm that imposes a hierarchical classification according to the ICD codes given for each case.³⁰ All cases are classified into the following four main categories: chromosomal case, syndromes, isolated single anomalies, and potential multiple birth defects.

For the comparison between maternal age class and qualitative variables, such as method of treatment, sex of fetuses/neonates, perinatal outcome, and multiple birth

defects, the χ^2 test was performed. The significance level was set at 0.05.

Next, the crude prevalence of birth defects per 10,000 pregnancies (number of mothers) and 10,000 live births by disease classification were calculated according to maternal age class. Abortion/stillbirths and plurality were not directly considered in the present study, since these data for the total ART population were not available according to maternal age. In the analysis of birth defects per pregnancy, concordant multiple pairs (cases in which both twins have birth defects or at least two triplets have birth defects, $n = 21$ pairs) were treated as a whole (namely, 21 pregnancies with birth defects).

The relative risk (RR) and 95% confidence interval (CI) for the birth defect categories according to the ICD-10 were calculated with 30–34 years of maternal age as the reference group. The RR and 95% CI of any birth defects were calculated by including or excluding chromosomal abnormalities. All eleven major categories were analyzed regardless of the number of cases in each. Analyses were performed for birth defects subcategories containing five or more cases in the 40+ age group.

Statistical analysis was conducted using Microsoft Excel 2010 (Microsoft Corporation, Redmond, WA, USA) and SAS for Windows version 9.2 (SAS Institute Inc, Cary, NC, USA).

Results

Demographic and perinatal outcome data of ART pregnancies with birth defects are summarized according to maternal age class in Table 1. Sex of the fetuses/neonates differed marginally significantly among maternal age class ($P = 0.046$). Males were more frequent in the 25–29, 30–34, and 35–39 year age classes, while females were more frequent in the 40+ year age class.

The percentage of multiple birth defects is shown in Table 2. The percentage of multiple birth defects gradually and linearly decreased with maternal age class. Isolated single anomalies decreased at ages greater than 35–39 years. The percentage of multiple birth defects was 14% (22/159) in the 25–29 year age class, and all 8% among the 30–34 (47/613), 35–39 (48/625), and 40+ (12/157) year age classes when chromosomal abnormalities were excluded. The difference between the numbers of multiple birth defects and isolated single anomalies according to maternal age class was not statistically significant.

The RRs of any birth defects per pregnancy and per live birth according to maternal age class are shown in Figure 1A and B, respectively. Regarding the presence

Table 2 Distribution of classification of cases using the EUROCAT computer algorithm according to maternal age class

Final classification	25–29		30–34		35–39		40+		P
	n	%	n	%	n	%	n	%	
Chromosomal cases	13	7.6	55	8.2	179	22.3	137	46.6	0.092
Syndromes	2	1.2	25	3.7	23	2.9	10	3.4	
Isolated single anomalies	135	78.5	541	81.0	554	68.9	135	45.9	
Multiple birth defects	22	12.8	47	7.0	48	6.0	12	4.1	

Note: The χ^2 test was performed between multiple birth defects and isolated birth defects according to maternal age class (the 2×4 contingency table).

Abbreviations: EUROCAT, European Surveillance of Congenital Anomalies; n, number.

of any birth defects per pregnancy, the prevalence of chromosomal abnormalities was constant in the 25–29 and 30–34 year age classes and increased rapidly thereafter, while nonchromosomal birth defects decreased linearly with advanced maternal age; thus the total prevalence across the maternal age distribution had a gradual U-shape. Regarding the prevalence of any birth defect per live birth, the prevalence of chromosomal abnormalities was constant in the 25–29 and 30–34 year maternal age classes and increased rapidly thereafter, while the prevalence of nonchromosomal birth defects was gradual and U-shaped; thus the total prevalence across the maternal age distribution was J-shaped.

The number, prevalence (per 10,000 pregnancies), RR, and 95% CI of birth defects in different organ systems are shown in Table 3. The prevalence of chromosomal abnormalities became significantly higher with advanced maternal age (RR = 2.71, 95% CI 2.01–3.65 for 35–39

years and RR = 6.93, 95% CI 5.08–9.45 for 40+ years). In contrast, the prevalence of nonchromosomal birth defects became significantly lower with advanced maternal age (RR = 0.88, 95% CI 0.79–0.99 for 35–39 years and RR = 0.80, 95% CI 0.68–0.95 for 40+ years). The prevalence of congenital malformations of the nervous system became significantly lower with advanced maternal age (RR = 0.71, 95% CI 0.52–0.98 for 35–39 years and RR = 0.42, 95% CI 0.23–0.78 for 40+ years). The RR was also significant with regard to other congenital malformations of the digestive system in the 25–29 year age class (RR = 1.64, 95% CI 1.01–2.64).

The number, prevalence (per 10,000 live births), RR, and 95% CI of birth defects in different organ systems are shown in Table 4. The prevalence of chromosomal abnormality became significantly higher with advanced maternal age (RR = 3.28, 95% CI 2.22–4.85 for 35–39 years and RR = 9.48, 95% CI 6.27–14.35 for 40+ years). The RR was

Figure 1 Crude prevalence of birth defects according to maternal age, 2004–2010. (A) Prevalence per 10,000 pregnancies. (B) Prevalence per 10,000 live births.

also significant with regard to nonchromosomal birth defects in the 40+ year age group (RR = 1.20, 95% CI 1.01–1.43). The RR was significant with regard to congenital malformations of the circulatory system in the 40+ year age group (RR = 1.40, 95% CI 1.10–1.79). The RR of congenital malformations of the aortic and mitral valves was significant for both the 25–29 year and 40+ year age groups, giving a clear U-shaped prevalence curve. The RR of congenital absence, atresia, and stenosis of the small intestine was also significant for the 40+ year age class (RR = 4.10, 95% CI 1.44–11.69).

Discussion

Given the very limited information provided by the present data, the main purpose of this study was to describe the epidemiologic features of birth defects after ART and maternal age (ie, a descriptive epidemiologic study), not to analyze the possible causes or mechanisms of birth defects controlling for the confounding factors (ie, an analytical epidemiologic study).

Demographic and perinatal outcomes

No special feature of birth defects cases was observed regarding demographic and perinatal outcomes according to the maternal age class, except that female cases were more common in the 40+ year age class. This result does not necessarily mean that the prevalence of birth defects in females in the 40+ year age class was higher compared to other maternal age classes, since the total ART population according to maternal age class and sex was unclear.

Multiple birth defects

The percentage of multiple birth defects was slightly higher in 25–29 year old mothers, but the difference was not statistically significant. The reason for this result was unclear. The classification method of multiple birth defects was not necessarily constant across studies.³⁰ Moreover, the relationship between maternal age and multiple birth defects was not well established.

Maternal age-specific risk of nonchromosomal birth defects

The present percentage of total birth defects after ART may be lower overall compared with that found in other studies, as seen in many reviews.^{31–33} One possible reason for this was that the cases in the present study were strictly restricted to only ICD-10 Q codes. Nevertheless, the main objective of this study was to evaluate the prevalence of birth defects in different maternal age groups and not to compare the birth

defect prevalence across different populations. Therefore, the comparison of birth defects across different maternal age classes may be biased only if there is differential reporting according to maternal age class, which is not likely to occur.

There have been several population-based studies of the relationship between maternal age and total birth defects.^{11–18} Recently Loane et al¹⁶ performed a population-based prevalence study using data from EUROCAT registers in 23 regions of Europe in 15 countries, covering a total of 1.75 million births from 2000 to 2004. According to their results, teenage mothers had the highest prevalence of all nonchromosomal congenital anomalies. The prevalence decreased as maternal age increased, until a slight increase was observed in mothers 40–44 years of age and a further increase was observed in mothers 45+ years of age. The prevalence of chromosomal abnormalities increased extremely with maternal age, especially in the 35–39 year and 40+ year groups. They concluded that higher maternal age is a negligible risk factor for nonchromosomal congenital anomalies, especially when compared to chromosomal abnormalities. Reefhuis and Honein¹⁵ analyzed a total of 1,050,616 singleton infants born from 1968 through 2000. They found that all birth defects combined and all nonchromosomal birth defects showed increased odds ratios for young and advanced maternal age, with a more pronounced increase for the group that included chromosomal defects. Overall, the relationship between maternal age and the prevalence of birth defects per birth or per live birth was similar in many previous studies in that these studies showed U-curves or gradual J-curves for nonchromosomal birth defects and extreme J-curves for chromosomal abnormalities.^{11–17} The present results also support previous findings that the effect of advanced maternal age, especially over 40, on nonchromosomal birth defects was not large compared to the effect on chromosomal abnormalities.

However, the maternal age with the lowest risks varied across studies, ie, 20–24 years,¹³ 25–29 years^{12,15} and 35–39 years.¹⁶ In the present study, the lowest RR for nonchromosomal birth defects per live birth was obtained in the 30–34 year group. One possible explanation for the present result is the essential difference in age-specific risks of nonchromosomal birth defects in infertile couples compared to the general population. However, according to Loane et al,¹⁶ the pattern of maternal age-specific risk widely differs between countries with some exceptional patterns. For example, the RR of any nonchromosomal birth defects compared with mothers aged 25–29 years increased consistently with

Table 3 Crude birth defect prevalence, RR, and 95% CI in ART according to maternal age class (prevalence per 10,000 pregnancies)

	25–29 n	Prevalence	RR	95% CI	30–34 n	Prevalence	RR
Q00-Q07							
Congenital malformations of the nervous system	27	14.9	1.47	0.95–2.27	80	10.2	1 (reference)
Q10-Q18							
Congenital malformations of eye, ear, face and neck	17	9.4	1.61	0.92–2.80	46	5.9	1 (reference)
Q17 Other congenital malformations of ear	11	6.1	1.41	0.71–2.77	34	4.3	1 (reference)
Q20-Q28							
Congenital malformations of the circulatory system	69	38.1	1.14	0.87–1.48	264	33.6	1 (reference)
Q20 Congenital malformations of cardiac chambers and connections	4	2.2	Insufficient		18	2.3	1 (reference)
Q21 Congenital malformations of cardiac septa	32	17.7	0.96	0.65–1.41	145	18.5	1 (reference)
Q210 Ventricular septal defect	23	12.7	1.00	0.64–1.57	100	12.7	1 (reference)
Q211 Atrial septal defect	6	3.3	0.84	0.35–2.01	31	3.9	1 (reference)
Q23 Congenital malformations of aortic and mitral valves	5	2.8	3.10	0.99–9.77	7	0.9	1 (reference)
Q24 Other congenital malformations of heart	1	0.6	Insufficient		17	2.2	1 (reference)
Q25 Congenital malformations of great arteries	19	10.5	1.68	0.99–2.86	49	6.2	1 (reference)
Q250 Patent ductus arteriosus	10	5.5	1.50	0.73–3.07	29	3.7	1 (reference)
Q30-Q34							
Congenital malformations of the respiratory system	4	2.2	Insufficient		12	1.5	1 (reference)
Q35-Q37							
Cleft lip and cleft palate	14	7.7	1.11	0.62–1.99	55	7.0	1 (reference)
Q37 Cleft palate with cleft lip	3	1.7	Insufficient		31	3.9	1 (reference)
Q38-Q45							
Other congenital malformations of the digestive system	23	12.7	1.64	1.01–2.64	61	7.8	1 (reference)
Q41 Congenital absence, atresia and stenosis of small intestine	4	2.2	Insufficient		7	0.9	1 (reference)
Q50-Q56							
Congenital malformations of genital organs	11	6.1	1.49	0.75–2.96	32	4.1	1 (reference)
Q60-Q64							
Congenital malformations of the urinary system	7	3.9	0.95	0.42–2.15	32	4.1	1 (reference)
Q62 Congenital obstructive defects of renal pelvis and congenital malformations of ureter	2	1.1	Insufficient		14	1.8	1 (reference)
Q620 Congenital hydronephrosis	0	0.0	Insufficient		11	1.4	1 (reference)
Q65-Q79							
Congenital malformations and deformations of the musculoskeletal system	34	18.8	1.13	0.77–1.64	131	16.7	1 (reference)
Q69 Polydactyly	9	5.0	1.15	0.55–2.40	34	4.3	1 (reference)
Q70 Syndactyly	2	1.1	Insufficient		16	2.0	1 (reference)
Q79 Congenital malformations of the musculoskeletal system, not elsewhere classified	3	1.7	Insufficient		28	3.6	1 (reference)
Q790 Congenital diaphragmatic hernia	1	0.6	Insufficient		12	1.5	1 (reference)
Q80-Q89							
Other congenital malformations	4	2.2	Insufficient		38	4.8	1 (reference)
Q87 Other specified congenital malformation syndromes affecting multiple systems	2	1.1	Insufficient		8	1.0	1 (reference)
Q90-Q99							
Chromosomal abnormalities, not elsewhere classified ^a	13	7.2	1.01	0.55–1.84	56	7.1	1 (reference)

35–39 n	Prevalence	RR	95% CI	40+ n	Prevalence	RR	95% CI
68	7.3	0.71	0.52–0.98	12	4.3	0.42	0.23–0.78
51	5.4	0.93	0.62–1.38	10	3.6	0.62	0.31–1.22
36	3.8	0.89	0.56–1.42	7	2.5	0.58	0.26–1.32
306	32.6	0.97	0.82–1.15	90	32.4	0.97	0.76–1.23
15	1.6	0.70	0.35–1.39	8	2.9	1.26	0.55–2.89
183	19.5	1.06	0.85–1.32	45	16.2	0.88	0.63–1.23
119	12.7	1.00	0.76–1.30	29	10.5	0.82	0.54–1.24
43	4.6	1.16	0.73–1.84	10	3.6	0.91	0.45–1.86
6	0.6	0.72	0.24–2.14	7	2.5	2.83	0.99–8.07
25	2.7	1.23	0.67–2.28	6	2.2	1.00	0.39–2.53
47	5.0	0.80	0.54–1.20	17	6.1	0.98	0.57–1.71
30	3.2	0.87	0.52–1.44	12	4.3	1.17	0.60–2.30
16	1.7	1.12	0.53–2.36	1	0.4	Insufficient	
57	6.1	0.87	0.60–1.26	15	5.4	0.77	0.44–1.37
26	2.8	0.70	0.42–1.18	9	3.2	0.82	0.39–1.73
62	6.6	0.85	0.60–1.21	21	7.6	0.97	0.59–1.60
7	0.7	0.84	0.29–2.39	7	2.5	2.83	0.99–8.07
30	3.2	0.79	0.48–1.29	3	1.1	Insufficient	
39	4.2	1.02	0.64–1.63	12	4.3	1.06	0.55–2.06
17	1.8	1.02	0.50–2.06	7	2.5	1.42	0.57–3.51
15	1.6	1.14	0.53–2.49	6	2.2	1.54	0.57–4.18
142	15.1	0.91	0.72–1.15	34	12.3	0.73	0.50–1.07
34	3.6	0.84	0.52–1.35	7	2.5	0.58	0.26–1.32
15	1.6	0.79	0.39–1.59	5	1.8	0.88	0.32–2.42
30	3.2	0.90	0.54–1.50	7	2.5	0.71	0.31–1.62
10	1.1	0.70	0.30–1.62	5	1.8	1.18	0.42–3.35
33	3.5	0.73	0.46–1.16	13	4.7	0.97	0.52–1.82
7	0.7	0.73	0.27–2.02	6	2.2	2.12	0.74–6.12
181	19.3	2.71	2.01–3.65	137	49.4	6.93	5.08–9.45

(Continued)

Table 3 (Continued)

	25–29 n	Prevalence	RR	95% CI	30–34 n	Prevalence	RR
Q90 Down syndrome	5	2.8	0.72	0.28–1.87	30	3.8	1 (reference)
Q91 Edwards syndrome and Patau syndrome	7	3.9	2.34	0.93–5.86	13	1.7	1 (reference)
Q913 Edwards syndrome, unspecified	5	2.8	2.17	0.74–6.35	10	1.3	1 (reference)
Q917 Patau syndrome, unspecified	2	1.1	Insufficient		3	0.4	1 (reference)
Q99 Other chromosome abnormalities, not elsewhere classified	0	0.0	Insufficient		8	1.0	1 (reference)
Any birth defects (number of mothers)	171	94.5	1.13	0.96–1.33	658	83.7	1 (reference)
Nonchromosomal birth defects (number of mothers)	158	87.3	1.13	0.95–1.34	609	77.5	1 (reference)

Notes: The estimated numbers of pregnancies according to maternal age class were as follows: 18,095 for 25–29 years; 78,576 for 30–35 years; 93,787 for 35–39 years; and 27,751 for 40+ years. The numbers in boldface represent a statistically significant association. ^aThe values were different from those of chromosomal cases given in Table 2, since some fetuses/neonates had two different chromosomal abnormalities.

Abbreviations: RR, relative risk; CI, confidence interval; ART, assisted reproductive technology; n, number.

advanced maternal age in Germany, while it decreased consistently in France. These results suggest that it is not biological age that is associated with the risk of nonchromosomal birth defects, but reproductive age, including parity and use of ART, social, ethnic exposure, and lifestyle factors that have different relationships with maternal age in different countries.¹⁶ Differences in study design, case determination, and potential confounders may play a role in the discrepancies observed.¹⁷

In the present study, the prevalence of nonchromosomal birth defects per pregnancy decreased linearly with maternal age class, while the prevalence of nonchromosomal birth defects per live birth showed a gradual U-shape. This difference could be explained by the fact that the percentage of abortions after ART in Japan rapidly increases in mothers aged over 35 years (data not shown). Most previous studies calculated the prevalence of birth defects per birth or live birth, since the total number of pregnancies in the general population is not always obtained. It is possible that different maternal age effects would be found if all incident cases during pregnancy could be considered, because of differential *in utero* survival effects.¹⁶ On this point, ART statistics offer a unique opportunity to analyze the epidemiological indicators per pregnancy, since the total number of pregnancies is recorded.

Maternal age-specific risk of several birth defects

Several birth defects showed a significant effect of maternal age. Given the limited information on other potential risk factors in the present study, the precise reason for the maternal age effect was unclear. Further epidemiological study with

a larger sample size and information on environmental and genetic factors is needed to clarify the direct and indirect maternal age effects on specific birth defects.

The effect of advanced maternal age was, as expected, largest regarding chromosomal abnormalities in total or specific birth defects, such as Down syndrome. This is a well-established finding in the general population.^{22,24,25} Moreover, according to Chen et al,³⁴ this finding was also observed in the ART population.

Several studies found an effect of maternal age on the nervous system,^{12,16} specifically on neural tube defects^{8,35–37} such as spina bifida,^{14,17} anencephalus,¹⁶ and encephalocele.¹⁶ The risk of congenital malformations of the nervous system per pregnancy was significantly lower with advanced maternal age. This tendency was also observed per live birth, although it was not statistically significant. Petrova and Vaktskjold³⁶ reported an inverse association between maternal age and the prevalence of neural tube defects. This tendency would be partly explained by the lower folic acid status resulting from poorer nutrition or lower periconceptional folic acid supplementation rates.^{16,36} It was also pointed out that in Japan younger women did not recognize the importance of folic acid intake to prevent neural tube defects as strongly as older women, and they were apt to take less folic acid, regardless of the precautions taken by the Ministry of Health, Labor and Welfare of Japan.³⁸ The results of the National Health and Nutrition Examination Survey performed in 2010³⁹ also supported this tendency, namely by finding that folic acid intake is lowest in women aged 20–29 compared to other age classes.

35–39 n	Prevalence	RR	95% CI	40+ n	Prevalence	RR	95% CI
110	11.7	3.07	2.05–4.60	83	29.9	7.83	5.16–11.89
46	4.9	2.96	1.60–5.49	40	14.4	8.71	4.66–16.29
37	3.9	3.10	1.54–6.23	34	12.3	9.63	4.76–19.48
9	1.0	Insufficient		6	2.2	Insufficient	
14	1.5	1.47	0.62–3.50	6	2.2	2.12	0.74–6.12
796	84.9	1.01	0.91–1.12	292	105.2	1.26	1.10–1.44
643	68.6	0.88	0.79–0.99	172	62.0	0.80	0.68–0.95

The risk of congenital malformations of the circulatory system per live birth significantly increased for mothers in the 40+ year age class. This result was in accordance with many other studies.^{13–15,37,40–42} According to Gill et al,¹⁷ increased risks for several cardiac defects, including ventricular septal defects, atrial septal defects, and Tetralogy of Fallot, were observed for mothers who were 40+ years of age. In the present study, a U-shaped maternal age effect on congenital malformations of the aortic and mitral valves was observed.

The RR of congenital absence, atresia, and stenosis of the small intestine was also significant for mothers in the 40+ year age class. The reason for this result was unclear. Several studies reported an association between older maternal age and esophageal atresia.^{16,17} However, no relationship between maternal age and intestinal atresia/stenosis was observed.^{17,37} It is possible that certain risk factors may differ among countries and may contribute greatly to the susceptibility to certain birth defects.

Over-ascertainment bias for older mothers should be considered. The frequency of medical checks increases in older mothers, since they are a potential high-risk group.^{13,15,17} Moreover, the effect of parity might be more important than maternal age for some birth defects,¹³ since maternal age and parity are strongly correlated. The obstetric outcomes, such as prematurity, low birth weight, and Caesarean section of women aged 40 years or older have been found to be influenced by parity.^{5,43} Few studies have focused on the possibility that parity is an independent risk factor for birth defects.⁴⁴

Limitations

This study has the following limitations, most of which could be attributed to the dataset, namely to the fact that individual information was obtained only from subjects with birth defects after ART (numerator of prevalence) and not for all of the ART population (denominator of prevalence). The estimation of the total ART population according to maternal age class may be biased if the percentage of mothers belonging to each maternal age class did not increase or decrease in a linear fashion. However, given the relatively small change in the percentage of all mothers contained in each maternal age class between 2008 and 2010, as well as given the low value of the numerator (number of fetuses/neonates with birth defects) and the large value of the denominator (total ART population), the influence seemed to be small.

The first and greatest limitation is that the author could not check the reliability of the data directly. This is an essential limitation of secondary data analyses. Second, although the present dataset was from a multiyear nationwide survey, it still did not have sufficiently high statistical power to detect the statistical significance of several birth defects with high RR, especially for the group with a maternal age over 40 years. On the other hand, some significant relationships between maternal age and RR could be produced by chance alone. Third, the author could not control for confounding factors that can affect ART and/or birth defects,^{32,45} such as multiple births, parity, smoking, socioeconomic status, medical history, and prenatal care. Fourth, follow-up after birth was limited to the neonatal period, and was incomplete. Therefore, some birth defects might not be detected, since they are not apparent within a few days after birth.

Table 4 Crude birth defect prevalence, RR, and 95% CI in ART according to maternal age class (prevalence per 10,000 live births)

	25–29 n	Prevalence	RR	95% CI	30–34 n	Prevalence	RR
Q00-Q07							
Congenital malformations of the nervous system	16	11.2	1.66	0.93–2.97	40	6.7	1 (reference)
Q10-Q18							
Congenital malformations of eye, ear, face and neck	16	11.2	1.58	0.89–2.82	42	7.1	1 (reference)
Q17 Other congenital malformations of ear	10	7.0	1.34	0.66–2.73	31	5.2	1 (reference)
Q20-Q28							
Congenital malformations of the circulatory system	69	48.2	1.12	0.86–1.46	257	43.2	1 (reference)
Q20 Congenital malformations of cardiac chambers and connections	4	2.8	Insufficient		18	3.0	1 (reference)
Q21 Congenital malformations of cardiac septa	32	22.3	0.93	0.63–1.36	143	24.0	1 (reference)
Q210 Ventricular septal defect	23	16.1	0.97	0.61–1.52	99	16.6	1 (reference)
Q211 Atrial septal defect	6	4.2	0.80	0.34–1.93	31	5.2	1 (reference)
Q23 Congenital malformations of aortic and mitral valves	5	3.5	3.46	1.06–11.35	6	1.0	1 (reference)
Q24 Other congenital malformations of heart	1	0.7	Insufficient		15	2.5	1 (reference)
Q25 Congenital malformations of great arteries	19	13.3	1.61	0.95–2.74	49	8.2	1 (reference)
Q250 Patent ductus arteriosus	10	7.0	1.43	0.70–2.94	29	4.9	1 (reference)
Q30-Q34							
Congenital malformations of the respiratory system	4	2.8	Insufficient		9	1.5	1 (reference)
Q35-Q37							
Cleft lip and cleft palate	14	9.8	1.10	0.61–1.98	53	8.9	1 (reference)
Q37 Cleft palate with cleft lip	3	2.1	Insufficient		29	4.9	1 (reference)
Q38-Q45							
Other congenital malformations of the digestive system	23	16.1	1.59	0.99–2.58	60	10.1	1 (reference)
Q41 Congenital absence, atresia and stenosis of small intestine	4	2.8	Insufficient		7	1.2	1 (reference)
Q50-Q56							
Congenital malformations of genital organs	11	7.7	1.43	0.72–2.83	32	5.4	1 (reference)
Q60-Q64							
Congenital malformations of the urinary system	5	3.5	0.83	0.32–2.17	25	4.2	1 (reference)
Q62 Congenital obstructive defects of renal pelvis and congenital malformations of ureter	2	1.4	Insufficient		14	2.4	1 (reference)
Q620 Congenital hydronephrosis	0	0.0	Insufficient		11	1.8	1 (reference)
Q65-Q79							
Congenital malformations and deformations of the musculoskeletal system	31	21.6	1.17	0.79–1.74	110	18.5	1 (reference)
Q69 Polydactyly	9	6.3	1.13	0.54–2.37	33	5.5	1 (reference)
Q70 Syndactyly	2	1.4	Insufficient		16	2.7	1 (reference)
Q80-Q89							
Other congenital malformations	4	2.8	Insufficient		31	5.2	1 (reference)
Q87 Other specified congenital malformation syndromes affecting multiple systems	2	1.4	Insufficient		8	1.3	1 (reference)
Q90-Q99							
Chromosomal abnormalities, not elsewhere classified	9	6.3	1.17	0.56–2.45	32	5.4	1 (reference)
Q90 Down syndrome	4	2.8	Insufficient		23	3.9	1 (reference)
Q91 Edwards syndrome and Patau syndrome	4	2.8	Insufficient		8	1.3	1 (reference)
Q913 Edwards syndrome, unspecified	2	1.4	Insufficient		6	1.0	1 (reference)
Any birth defects	153	106.8	1.12	0.94–1.34	566	95.1	1 (reference)
Nonchromosomal birth defects	144	100.5	1.11	0.92–1.33	540	90.7	1 (reference)

Notes: The estimated numbers of live births according to the maternal age class were as follows: 14,322 for 25–29 years; 59,532 for 30–35 years; 64,676 for 35–39 years; and 14,521 for 40+ years. The numbers in boldface represent a statistically significant association.

Abbreviations: RR, relative risk; CI, confidence interval; ART, assisted reproductive technology; n, number.

Conclusion

The present results elucidate the current situation regarding birth defects after ART according to maternal age class. The risk of chromosomal abnormalities increased notably at a maternal age of 30 or more. Early detection of chromosomal abnormalities is essential when performing ART for

advanced-age couples. On the other hand, the risk of non-chromosomal birth defects in ART live births is relatively small between maternal age groups. This information would be important to include in the counseling provided before ART is performed. Reassurance can be given to infertile couples of advanced age in that maternal age in itself does

35–39 n	Prevalence	RR	95% CI	40+ n	Prevalence	RR	95% CI
37	5.7	0.85	0.54–1.33	7	4.8	0.72	0.32–1.60
50	7.7	1.10	0.73–1.65	10	6.9	0.98	0.49–1.95
35	5.4	1.04	0.64–1.69	7	4.8	0.93	0.41–2.10
301	46.5	1.08	0.91–1.27	88	60.6	1.40	1.10–1.79
15	2.3	0.77	0.39–1.52	8	5.5	1.82	0.79–4.19
183	28.3	1.18	0.95–1.47	44	30.3	1.26	0.90–1.77
119	18.4	1.11	0.85–1.44	28	19.3	1.16	0.76–1.76
43	6.6	1.28	0.81–2.03	10	6.9	1.32	0.65–2.70
5	0.8	0.77	0.23–2.51	7	4.8	4.78	1.61–14.23
22	3.4	1.35	0.70–2.60	5	3.4	1.37	0.50–3.76
47	7.3	0.88	0.59–1.32	17	11.7	1.42	0.82–2.47
30	4.6	0.95	0.57–1.59	12	8.3	1.70	0.87–3.32
16	2.5	1.64	0.72–3.70	1	0.7	Insufficient	
57	8.8	0.99	0.68–1.44	13	9.0	1.01	0.55–1.84
26	4.0	0.83	0.49–1.40	7	4.8	0.99	0.43–2.26
61	9.4	0.94	0.66–1.34	21	14.5	1.43	0.87–2.36
7	1.1	0.92	0.32–2.62	7	4.8	4.10	1.44–11.69
30	4.6	0.86	0.52–1.42	3	21	Insufficient	
36	5.6	1.33	0.80–2.21	11	7.6	1.80	0.89–3.67
17	2.6	1.12	0.55–2.27	7	4.8	2.05	0.83–5.08
15	2.3	1.26	0.58–2.73	6	4.1	2.24	0.83–6.05
120	18.6	1.00	0.78–1.30	28	19.3	1.04	0.69–1.58
33	5.1	0.92	0.57–1.49	7	4.8	0.87	0.39–1.97
15	2.3	0.86	0.43–1.75	5	3.4	1.28	0.47–3.50
31	4.8	0.92	0.56–1.51	11	7.6	1.45	0.73–2.89
7	1.1	0.81	0.29–2.22	5	3.4	2.56	0.84–7.83
114	17.6	3.28	2.22–4.85	74	51.0	9.48	6.27–14.35
83	12.8	3.32	2.09–5.27	50	34.4	8.91	5.44–14.60
16	2.5	1.84	0.79–4.30	17	11.7	8.71	3.76–20.18
14	2.2	2.15	0.83–5.59	13	9.0	8.88	3.38–23.37
684	105.8	1.11	0.99–1.24	217	149.4	1.57	1.35–1.84
595	92.0	1.01	0.90–1.14	158	108.8	1.20	1.01–1.43

not produce noticeable extra risk for nonchromosomal birth defects.

Acknowledgments

I would like to thank Toshimi Ooma for assistance with the data analysis. This work was supported in part by a

Grant-in-Aid for Scientific Research (B) from the Japan Society for the Promotion of Science (grant number 24390167).

Disclosure

The author reports no conflicts of interest in this work.

References

- Johnson JA, Tough S; for Society of Obstetricians and Gynaecologists of Canada. Delayed child-bearing. *J Obstet Gynaecol Can.* 2012;34(1):80–93.
- Alshami HA, Kadasne AR, Khalfan M, Iqbal SZ, Mirghani HM. Pregnancy outcome in late maternal age in a high-income developing country. *Arch Gynecol Obstet.* 2011;284(5):1113–1116.
- Carolan M, Frankowska D. Advanced maternal age and adverse perinatal outcome: a review of the evidence. *Midwifery.* 2011;27(6):793–801.
- Jahromi BN, Hussein Z. Pregnancy outcome at maternal age 40 and older. *Taiwan J Obstet Gynecol.* 2008;47(3):318–321.
- Chan BC, Lao TT. Effect of parity and advanced maternal age on obstetric outcome. *Int J Gynaecol Obstet.* 2008;102(3):237–241.
- Cleary-Goldman J, Malone FD, Vidaver J, et al; for FASTER Consortium. Impact of maternal age on obstetric outcome. *Obstet Gynecol.* 2005;105(5 Pt 1):983–990.
- Schoen C, Rosen T. Maternal and perinatal risks for women over 44 – a review. *Maturitas.* 2009;64(2):109–113.
- Czeizel A. Maternal mortality, fetal death, congenital anomalies and infant mortality at an advanced maternal age. *Maturitas.* 1988;Suppl 1:73–81.
- Salem Yaniv S, Levy A, Wiznitzer A, Holcberg G, Mazor M, Sheiner E. A significant linear association exists between advanced maternal age and adverse perinatal outcome. *Arch Gynecol Obstet.* 2011;283(4):755–759.
- Yogev Y, Melamed N, Bardin R, Tenenbaum-Gavish K, Ben-Shitrit G, Ben-Haroush A. Pregnancy outcome at extremely advanced maternal age. *Am J Obstet Gynecol.* 2010;203(6):558. e1–e7.
- Goldberg MF, Edmonds LD, Oakley GP. Reducing birth defect risk in advanced maternal age. *JAMA.* 1979;242(21):2292–2294.
- Croen LA, Shaw GM. Young maternal age and congenital malformations: a population-based study. *Am J Public Health.* 1995;85(5):710–713.
- Hollier LM, Leveno KJ, Kelly MA, McIntire DD, Cunningham FG. Maternal age and malformations in singleton births. *Obstet Gynecol.* 2000;96(5 Pt 1):701–706.
- Hay S, Barbano H. Independent effects of maternal age and birth order on the incidence of selected congenital malformations. *Teratology.* 1972;6(3):271–279.
- Reefhuis J, Honein MA. Maternal age and non-chromosomal birth defects, Atlanta, 1968–2000: teenager or thirty-something, who is at risk? *Birth Defects Res A Clin Mol Teratol.* 2004;70(9):572–579.
- Loane M, Dolk H, Morris JK; for EUROCAT Working Group. Maternal age-specific risk of non-chromosomal anomalies. *BJOG.* 2009;116(8):1111–1119.
- Gill SK, Broussard C, Devine O, Green RF, Rasmussen SA, Reefhuis J; and the National Birth Defects Prevention Study. Association between maternal age and birth defects of unknown etiology – United States, 1997–2007. *Birth Defects Res A Clin Mol Teratol.* 2012;94(12):1010–1018.
- Baird PA, Sadovnick AD, Yee IM. Maternal age and birth defects: a population study. *Lancet.* 1991;337(8740):527–530.
- Savva GM, Walker K, Morris JK. The maternal age-specific live birth prevalence of trisomies 13 and 18 compared to trisomy 21 (Down syndrome). *Prenat Diagn.* 2010;30(1):57–64.
- Hassold T, Chiu D. Maternal age-specific rates of numerical chromosome abnormalities with special reference to trisomy. *Hum Genet.* 1985;70(1):11–17.
- Sherman SL, Allen EG, Bean LH, Freeman SB. Epidemiology of Down syndrome. *Ment Retard Dev Disabil Res Rev.* 2007;13(3):221–227.
- Hagen A, Entezami M, Gasiorek-Wiens A, et al. The impact of first trimester screening and early fetal anomaly scan on invasive testing rates in women with advanced maternal age. *Ultraschall Med.* 2011;32(3):302–306.
- Sipek A, Gregor V, Horáček J, Chudobová M. Down syndrome in the Czech Republic during 1961–1997: prevalence, prenatal diagnosis and maternal-age-specific rates. *J Obstet Gynaecol.* 2001;21(3):266–269.
- Dolk H, Loane M, Garne E, et al. Trends and geographic inequalities in the prevalence of Down syndrome in Europe, 1980–1999. *Rev Epidemiol Sante Publique.* 2005;53 Spec No 2:S87–S95.
- Morris JK, Wald NJ, Mutton DE, Alberman E. Comparison of models of maternal age-specific risk for Down syndrome live births. *Prenat Diagn.* 2003;23(3):252–258.
- Ooki S. Birth defects in singleton versus multiple ART births in Japan (2004–2008). *J Pregnancy.* 2011;2011:285706.
- Ooki S. Birth defects after assisted reproductive technology in Japan: comparison between multiples and singletons, 2004–2009. In: Shi Z, editor. *In vitro Fertilization (Special Issue). Reproductive System and Sexual Disorders.* 2012;S:5.
- Ooki S. Concordance rates of birth defects after assisted reproductive technology among 17,258 Japanese twin pregnancies: a nationwide survey, 2004–2009. *J Epidemiol.* 2013;23(1). doi:10.2188/jea.JE20120103.
- Ooki S. Multiple congenital anomalies after assisted reproductive technology in Japan (between 2004 and 2009). *ISRN Epidemiology.* Epub 2013. doi:10.5402/2013/452085.
- Garne E, Dolk H, Loane M, et al; for EUROCAT Working Group. Paper 5: Surveillance of multiple congenital anomalies: implementation of a computer algorithm in European registers for classification of cases. *Birth Defects Res A Clin Mol Teratol.* 2011;Suppl 1:S44–S50.
- Rimm AA, Katayama AC, Diaz M, Katayama KP. A meta-analysis of controlled studies comparing major malformation rates in IVF and ICSI infants with naturally conceived children. *J Assist Reprod Genet.* 2004;21(12):437–443.
- Källén B, Finnström O, Lindam A, Nilsson E, Nygren KG, Otterblad PO. Congenital malformations in infants born after in vitro fertilization in Sweden. *Birth Defects Res A Clin Mol Teratol.* 2010;88(3):137–143.
- Shebl O, Ebner T, Sir A, Sommergruber M, Tews G. The role of mode of conception in the outcome of twin pregnancies. *Minerva Ginecol.* 2009;61(2):141–152.
- Chen BY, Hwang BF, Guo YL. Epidemiology of congenital anomalies in a population-based birth registry in Taiwan, 2002. *J Formos Med Assoc.* 2009;108(6):460–468.
- Sipek A, Horáček J, Gregor V, Rychtářková J, Dzurová D, Masátová D. Neural tube defects in the Czech Republic during 1961–1999: incidences, prenatal diagnosis and prevalences according to maternal age. *J Obstet Gynaecol.* 2002;22(5):501–507.
- Petrova JG, Vaktskjold A. The incidence of neural tube defects in Norway and the Arkhangelskaja Oblast in Russia and the association with maternal age. *Acta Obstet Gynecol Scand.* 2009;88(6):667–672.
- Materna-Kiryluk A, Wiśniewska K, Badura-Stronka M, et al. Parental age as a risk factor for isolated congenital malformations in a Polish population. *Paediatr Perinat Epidemiol.* 2009;23(1):29–40.
- The Ministry of Health, Labor and Welfare of Japan. Available from: http://www1.mhlw.go.jp/houdou/1212/h1228-1_18.html. (In Japanese) Accessed December 27, 2012.
- The National Health and Nutrition Examination Survey. Available from: <http://www.mhlw.go.jp/bunya/kenkou/eiyou/h22-houkoku.html>. (In Japanese) Accessed December 27, 2012.
- Kim MA, Yee NH, Choi JS, Choi JY, Seo K. Prevalence of birth defects in Korean livebirths, 2005–2006. *J Korean Med Sci.* 2012;27(10):1233–1240.
- Reller MD, Strickland MJ, Riehle-Colarusso T, Mahle WT, Correa A. Prevalence of congenital heart defects in metropolitan Atlanta, 1998–2005. *J Pediatr.* 2008;153(6):807–813.
- Miller A, Riehle-Colarusso T, Siffel C, Frías JL, Correa A. Maternal age and prevalence of isolated congenital heart defects in an urban area of the United States. *Am J Med Genet A.* 2011;155A(9):2137–2145.
- Ziadeh S, Yahaya A. Pregnancy outcome at age 40 and older. *Arch Gynecol Obstet.* 2001;265(1):30–33.
- Duong HT, Hoyt AT, Carmichael SL, et al; for National Birth Defects Prevention Study. Is maternal parity an independent risk factor for birth defects? *Birth Defects Res A Clin Mol Teratol.* 2012;94(4):230–236.
- Källén B, Finnström O, Nygren KG, Olausson PO. In vitro fertilization (IVF) in Sweden: risk for congenital malformations after different IVF methods. *Birth Defects Res A Clin Mol Teratol.* 2005;73(3):162–169.

International Journal of Women's Health**Dovepress****Publish your work in this journal**

The International Journal of Women's Health is an international, peer-reviewed open-access journal publishing original research, reports, editorials, reviews and commentaries on all aspects of women's healthcare including gynecology, obstetrics, and breast cancer. The manuscript management system is completely online and includes

a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Submit your manuscript here: <http://www.dovepress.com/international-journal-of-womens-health-journal>