

Non-reassuring foetal status and sleep problems in 1-year-old infants in the Japan Environment and Children's Study: a cohort study

Nakahara, Kazushige

Department of Obstetrics and Gynecology, School of Medical Sciences, Kyushu University

Michikawa, Takehiro

Department of Environmental and Occupational Health, School of Medicine, Toho University

Morokuma, Seiichi

Department of Health Sciences, Graduate School of Medical Sciences, Kyushu University

Ogawa, Masanobu

Research Center for Environmental and Developmental Medical Sciences, Kyushu University

他

<https://hdl.handle.net/2324/4476150>

出版情報 : Scientific reports. 10 (11432), 2020-07-10. Nature Publishing Group

バージョン :

権利関係 : Creative Commons Attribution 4.0 International





OPEN

Non-reassuring foetal status and sleep problems in 1-year-old infants in the Japan Environment and Children's Study: a cohort study

Kazushige Nakahara^{1,26}, Takehiro Michikawa^{2,26}, Seiichi Morokuma^{3,4}✉, Masanobu Ogawa⁴, Kiyoko Kato^{1,4}, Masafumi Sanefuji^{4,5}, Eiji Shibata^{6,7}, Mayumi Tsuji^{6,8}, Masayuki Shimono^{6,9}, Toshihiro Kawamoto⁶, Shouichi Ohga⁵, Koichi Kusuhara^{6,9} & The Japan Environment and Children's Study Group*

Abnormal autonomic function may cause false-positive non-reassuring foetal status (*fpNRFS*) and may also cause sleeping problems after birth. However, an association between *fpNRFS* and sleeping problems in infants has not been reported. We previously showed an association of NRFS with temperament, including bad mood and frequent crying for long durations in 1-month-old infants. In the present study, we aimed to assess this association in 1-year-old infants. A total of 62,612 single pregnant women were included in the analysis. *fpNRFS* was identified from medical records. Sleep problems, such as short sleep duration or crying at night, were investigated in 1-year-old infants using a questionnaire for mothers. We used a log-binominal regression model to explore the association of *fpNRFS* with each sleep problem and to estimate risk ratios (RRs). The number of *fpNRFS* cases was 2,071, with a frequency of 3.3%. We observed an association of *fpNRFS* with shorter sleep duration of less than 8 h a night (RR 1.30, 95% confidence intervals [CI] 1.10–1.54), crying at night (RR 1.19, 95% CI 1.03–1.39), and bedtime after 22:00 (RR 1.09, 95% CI 1.00–1.18). *fpNRFS* may be associated with sleep problems in 1-year-old infants.

Children with developmental disorders tend to have sleep problems and abnormal temperaments^{1,2,3,4}. Foetal distress is one of the prenatal risk factors for developmental disorders, such as autism spectrum disorder (ASD)⁵.

No non-invasive methods have been developed to determine actual foetal distress with foetal acidosis. Obstetricians clinically use non-reassuring foetal status (NRFS) by monitoring foetal heart rate using cardiotocography (CTG) instead of foetal distress. Unfortunately, the false-positive rate of NRFS assessment by CTG is high^{6–8}.

¹Department of Obstetrics and Gynecology, Graduate School of Medical Sciences, Kyushu University, Fukuoka, Japan. ²Department of Environmental and Occupational Health, School of Medicine, Toho University, Tokyo, Japan. ³Department of Health Sciences, Graduate School of Medical Sciences, Kyushu University, Fukuoka 812-8582, Japan. ⁴Research Center for Environmental and Developmental Medical Sciences, Kyushu University, Fukuoka, Japan. ⁵Department of Pediatrics, Graduate School of Medical Sciences, Kyushu University, Fukuoka, Japan. ⁶Japan Environment and Children's Study, UOEH Subunit Center, University of Occupational and Environmental Health, Kitakyushu, Fukuoka, Japan. ⁷Department of Obstetrics and Gynecology, School of Medicine, University of Occupational and Environmental Health, Kitakyushu, Fukuoka, Japan. ⁸Department of Environmental Health, School of Medicine, University of Occupational and Environmental Health, Kitakyushu, Fukuoka, Japan. ⁹Department of Pediatrics, School of Medicine, University of Occupational and Environmental Health, Kitakyushu, Japan. ²⁶These authors contributed equally: Kazushige Nakahara and Takehiro Michikawa. *A comprehensive list of consortium members appears at the end of the paper. ✉email: morokuma@med.kyushu-u.ac.jp

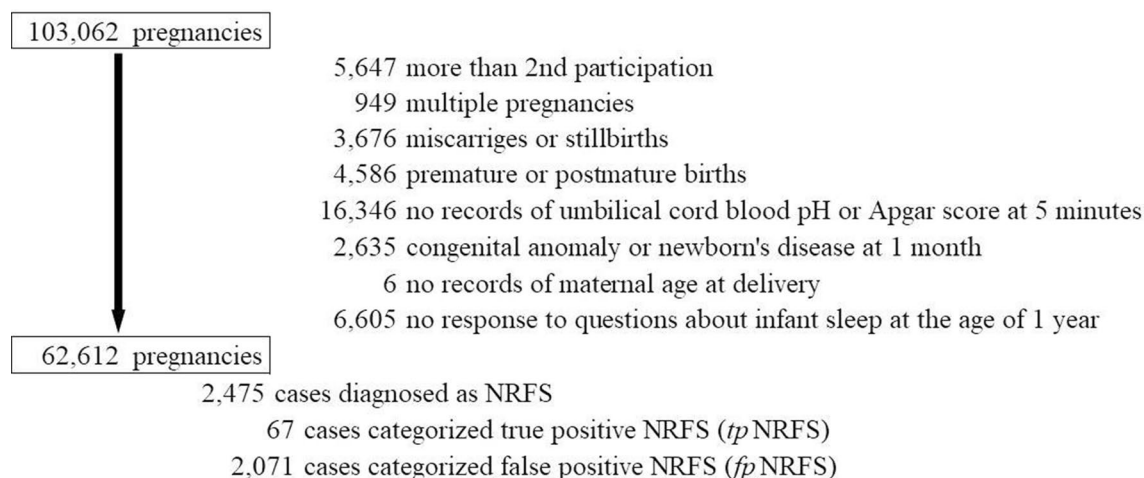


Figure 1. Population flow chart. NRFS: non-reassuring foetal status; true-positive NRFS (*tpNRFS*): cases with umbilical cord pH < 7.2 and Apgar score of < 7 at 5 min after birth of all NRFS cases; false-positive NRFS (*fpNRFS*): cases with umbilical cord pH ≥ 7.2 and Apgar score of ≥ 7 at 5 min after birth of all NRFS cases.

Many diagnosed NRFS cases do not show actual foetal distress. They demonstrate normal umbilical cord blood pH and normal Apgar scores.

Babies with normal umbilical cord blood and normal Apgar scores may be categorised into two groups. Some babies show abnormal heart rate patterns and are diagnosed with false-positive NRFS (*fpNRFS*), whereas others do not. Babies showing *fpNRFS* may inherently have abnormal regulation of heart rate. Autonomic nerves play an important role in heart rate regulation⁹ and are also closely related to sleep¹⁰. Children with ASD tend to have different heart rate variability from that of normal children¹¹, and they frequently have sleep disorders, such as short sleep during the night and frequent night crying^{12,13}. Therefore, children with *fpNRFS* may have abnormal autonomic functions and could show abnormal sleep patterns after birth.

We previously showed an association of NRFS with temperaments, including bad mood and frequent crying for a long duration in 1-month infants¹⁴. These findings suggest that NRFS may be associated with sleep disorders; nevertheless, an association between the two has not been reported. Therefore, this study aimed to investigate the presence of an association between NRFS, especially *fpNRFS*, and sleep problems in 1-year-old infants.

Methods

Data used in this study were obtained from the Japan Environment and Children's Study (JECS), an ongoing large-scale cohort study. The JECS was designed to follow-up children from the prenatal period to the age of 13 years. The baseline profile of participants in the JECS was reported previously¹⁵.

Ethics of research. The JECS study protocol was approved by the Ministry of the Environment's Institutional Review Board on Epidemiological Studies (No. 100406001) and the Ethics Committee of all participating institutions: the National Institute for Environmental Studies that leads the JECS, the National Center for Child Health and Development, Hokkaido University, Sapporo Medical University, Asahikawa Medical College, Japanese Red Cross Hokkaido College of Nursing, Tohoku University, Fukushima Medical University, Chiba University, Yokohama City University, University of Yamanashi, Shinshu University, University of Toyama, Nagoya City University, Kyoto University, Doshisha University, Osaka University, Osaka Medical Center and Research Institute for Maternal and Child Health, Hyogo College of Medicine, Tottori University, Kochi University, University of Occupational and Environmental Health, Kyushu University, Kumamoto University, University of Miyazaki, and University of Ryukyus. Written informed consent was obtained from all participants. All methods were performed in accordance with approved guidelines. The detailed protocol has been reported elsewhere¹⁶.

Study participants. Between 2011 and 2014, 103,062 pregnancies were registered from 15 regions throughout Japan (Fig. 1). Of those, we excluded 40,450 pregnancies due to the following reasons: prior participation in the study ($n = 5,647$), multiple pregnancies ($n = 949$), miscarriage or stillbirth ($n = 3,676$), pre-term or post-term birth ($n = 4,586$), no records of umbilical cord blood pH or Apgar score at 5 min ($n = 16,346$), congenital anomaly or newborn disease at 1 month old ($n = 2,635$), missing information on maternal age at delivery ($n = 6$), and no response to the questions about infant sleep at the age of 1 year ($n = 6,605$). Finally, 62,612 single pregnant women were included in our analysis.

Exposure (NRFS). In Japan, abnormal patterns in foetal heart rate are categorised into five levels based on baseline heart rate, variability, kinds, severity of deceleration, and so on¹⁷. Level 1 is considered normal, while levels 3 or higher are usually diagnosed as NRFS, and require preparation and execution of forced delivery.

In the present study, the NRFS cases were selected in two ways, based on the medical record transcription at birth. First, we selected cases diagnosed as NRFS by obstetricians. In the cohort data, only information on

	Without non-reassuring foetal status (n = 60,137)		With non-reassuring foetal status (n = 2,475)	
	n ^a	%	n ^a	%
Maternal characteristics				
Age at delivery (years)				
< 25	5,437	9.0	202	8.2
25–29	16,536	27.5	654	26.4
30–34	21,513	35.8	887	35.8
≥ 35	16,651	27.7	732	29.6
Smoking habits				
Never smoked	35,745	59.6	1,512	61.3
Ex-smokers who quit before pregnancy	14,175	23.7	505	20.5
Smokers during early pregnancy	10,028	16.7	451	18.3
Alcohol consumption				
Never drank	20,847	34.7	804	32.6
Ex-drinkers who quit before pregnancy	11,001	18.3	401	16.2
Drinkers during early pregnancy	28,161	46.9	1,265	51.2
Pre-pregnancy body mass index, kg/m ²				
< 18.5	9,613	16.0	401	16.2
18.5–24.9	44,463	74.0	1,803	72.9
≥ 25.0	6,041	10.1	270	10.9
Parity				
0	25,965	43.3	1,874	75.9
≥ 1	33,997	56.7	594	24.1
Infertility treatment				
No	56,104	93.3	2,173	87.9
Ovulation stimulation/artificial insemination by sperm from husband	2,203	3.7	139	5.6
Assisted reproductive technology	1,798	3.0	161	6.5
Type of delivery				
Vaginal	50,555	84.1	1,201	48.5
Caesarean	9,532	15.9	1,274	51.5
Gestational age (weeks)				
Early term (37–38)	19,779	32.9	558	22.6
Full term (39–41)	40,358	67.1	1,917	77.5
Educational background (years)				
< 10	2,428	4.1	92	3.8
10–12	18,404	30.9	683	27.8
13–16	37,813	63.5	1,641	66.9
≥ 17	876	1.5	37	1.5
Household income (million Japanese-yen/year)				
< 2	2,908	5.2	119	5.2
2 to < 4	18,989	34.1	747	32.4
4 to < 6	18,531	33.3	748	32.5
6 to < 8	9,060	16.3	412	17.9
8 to < 10	3,769	6.8	181	7.9
≥ 10	2,399	4.3	97	4.2
Kessler six-item psychological distress scale at 1 year				
0–4	46,845	78.0	1,983	80.3
> = 5 (psychological distress)	13,183	22.0	487	19.7
Birth weight				
Mean (SD) (g)	3,065 (352)		2,987 (407)	
Small for gestation age	4,121	6.9	395	16.0
Infant sex				
Male	30,418	50.6	1,429	57.7
Female	29,719	49.4	1,046	42.3
Continued				

	Without non-reassuring foetal status (n = 60,137)		With non-reassuring foetal status (n = 2,475)	
	n ^a	%	n ^a	%
Infant characteristics				
Doctor diagnosis at 1 year old				
Asthma	1,528	2.5	38	1.5
Atopic dermatitis	2,588	4.3	105	4.2
Feeding status				
Formula feeding	1,298	2.2	47	1.9
Partial breastfeeding	38,258	63.6	1,758	71.0
Exclusive breastfeeding	20,581	34.2	670	27.1

Table 1. Baseline characteristics of the study population. ^aNumbers in subgroups do not equal the overall number because of missing data.

the presence or absence of NRFS was available. Second, we selected cases that were not diagnosed as NRFS but showed umbilical cord blood pH < 7.2 and Apgar scores < 7 at 5 min after birth.

Of all the NRFS cases, those showing umbilical cord blood pH < 7.2 and Apgar scores of < 7 at 5 min were categorised as true-positive NRFS (*tp*NRFS) cases, and those showing umbilical cord blood pH ≥ 7.2 and Apgar scores ≥ 7 at 5 min were categorised as false-positive NRFS (*fp*NRFS) cases. Only cases that met either umbilical cord blood pH or Apgar scores at 5 min were excluded from the stratified analysis with *tp*NRFS and *fp*NRFS.

Outcome (infant sleep and crying at night). At 1 year after delivery, information on infant sleep and crying at night was collected via a parent-reported questionnaire. In this analysis, we focused on five points. First, from the responses regarding infant’s sleeping period the previous day, we determined the number of nocturnal awakenings. We defined ≥ 3 awakenings as too many because a previous study reported that the upper limit of number of awakenings during the night is 2.5 for 1-year-old infants.¹⁸ Second, we analysed whether the infants awoke more than once and kept awake for more than 1 h during the night. Third, we analysed the duration of sleep during the night (20:00–7:59). We regarded less than 8 h as unusual. Fourth, we determined bedtime of infants. We defined bedtime after 22:00 as too late. Fifth, we analysed crying at night in the past month. If the mother answered that her infant awoke and cried during the night and that the frequency of crying at night was more than 5 times in a week, we defined the case as crying at night.

Covariates. Information about maternal age at delivery, smoking habits, alcohol consumption, pre-pregnancy body mass index (BMI), parity, gestational age at birth, infertility treatment, type of delivery, small for gestational age, infant sex, maternal psychological distress at 1 year after delivery, physician diagnosis of asthma and atopic dermatitis at 1 year old, and feeding status were collected via self-administered questionnaires and/or medical records. Maternal psychological distress was assessed using the Kessler 6^{19,20}, including the questionnaire at 1 year after delivery. According to previous studies, participants with scores of 5 or more were categorised as having distress²¹.

Statistical analyses. We used a log-binominal regression model to explore the association of NRFS with each outcome and to estimate the risk ratio (RRs) of each outcome and 95% confidence intervals (CIs). We initially adjusted for maternal age at delivery and then further adjusted for smoking habits (never smokers, ex-smokers who quit before pregnancy, smokers during early pregnancy), alcohol consumption (never drinkers, ex-drinkers who quit before pregnancy, drinkers during early pregnancy), pre-pregnancy BMI (< 18.5, 18.5–24.9, ≥ 25.0 kg/m²), parity (0, ≥ 1), infertility treatment (no, ovulation stimulation/artificial insemination by sperm from husband, assisted reproductive technology), type of delivery (vaginal, caesarean section), gestational age at birth (37–38, 39–41 weeks), small for gestational age (yes, no), psychological distress at 1 year after delivery (yes, no), doctor diagnosis of asthma and atopic dermatitis at 1 year old, and feeding (breast milk, synthetic milk, both).

We used a fixed dataset “jecs-an-20180131,” which was released in March 2018. Stata version 15 (StataCorp LP, College Station, Texas, USA) was used for all the analyses.

Results

The baseline characteristics of the present study population with or without NRFS are shown in Table 1. The number of all NRFS cases was 2,475, with a frequency of 4.0%. Among the NRFS cases, 67 cases (2.7%) were *tp*NRFS, 2,071 cases (83.7%) were *fp*NRFS. The other 337 cases (13.7%) did not belong to any groups because these cases met only either umbilical cord pH or Apgar scores at 5 min.

Table 2 shows the RRs for NRFS and infant sleep and crying at night. In the multivariable model, we observed the association of all NRFS with shorter sleep time less than 8 h during the night (RR 1.28, 95% CI 1.10–1.49), crying at night (RR 1.17, 95% CI 1.02–1.34) and bedtime after 22:00 (RR 1.10, 95% CI 1.01–1.18). The same associations were also observed only in the *fp*NRFS cases (RR for short sleep 1.30, 95% CI 1.10–1.54, RR for

	n	Number of outcomes	Frequency of outcome (%)	Maternal age adjusted model			Multivariable model ^a		
				RR	95% CI		RR	95% CI	
All NRFS									
Waking up 3 or more times in a night									
No NRFS	59,697	1,426	2.4	Ref			Ref		
NRFS	2,456	67	2.7	1.14	0.89	1.45	1.20	0.93	1.55
Waking up 1 or more times and remaining awake for more than 1 h									
No NRFS	59,697	3,435	5.8	Ref			Ref		
NRFS	2,456	153	6.2	1.08	0.93	1.27	1.01	0.86	1.19
Sleep for less than 8 h during the night (20:00–7:59)									
No NRFS	59,697	3,067	5.1	Ref			Ref		
NRFS	2,456	174	7.1	1.37	1.19	1.59	1.28	1.10	1.49
Sleep at 22:00 or later									
No NRFS	59,697	11,929	20.0	Ref			Ref		
NRFS	2,456	573	23.3	1.17	1.09	1.26	1.10	1.01	1.18
Crying for 5 days or over in a week									
No NRFS	60,054	4,452	7.4	Ref			Ref		
NRFS	2,474	207	8.4	1.13	0.99	1.29	1.17	1.02	1.34
True-positive NRFS (tpNRFS)									
Waking up 3 or more times in a night									
No NRFS	59,697	1,426	2.4	Ref			Ref		
NRFS	66	3	4.6	1.95	0.64	5.88	2.08	0.69	6.26
Waking up 1 or more times and remaining awake for more than 1 h									
No NRFS	59,697	3,435	5.8	Ref			Ref		
NRFS	66	6	9.1	1.56	0.73	3.36	1.36	0.63	2.91
Sleep for less than 8 h during the night (20:00–7:59)									
No NRFS	59,697	3,067	5.1	Ref			Ref		
NRFS	66	2	3.0	0.59	0.15	2.29	0.50	0.13	1.95
Sleep at 22:00 or later									
No NRFS	59,697	2,409	4.0	Ref			Ref		
NRFS	66	12	18.2	0.92	0.55	1.53	0.83	0.50	1.38
Crying for 5 days or over in a week									
No NRFS	60,054	4,452	7.4	Ref			Ref		
NRFS	67	8	11.9	1.63	0.85	3.12	1.64	0.86	3.12
False-positive NRFS (fpNRFS)									
Waking up 3 or more times in a night									
No NRFS	59,697	1,426	2.4	Ref			Ref		
NRFS	2,055	60	2.9	1.21	0.94	1.57	1.28	0.98	1.68
Waking up 1 or more times and remaining awake for more than 1 h									
No NRFS	59,697	3,435	5.8	Ref			Ref		
NRFS	2,055	131	6.4	1.11	0.94	1.31	1.04	0.87	1.23
Sleep for less than 8 h during the night (20:00–7:59)									
No NRFS	59,697	3,067	5.1	Ref			Ref		
NRFS	2,055	148	7.2	1.40	1.19	1.64	1.30	1.10	1.54
Sleep at 22:00 or later									
No NRFS	59,697	11,929	20.0	Ref			Ref		
NRFS	2,055	474	23.2	1.16	1.07	1.26	1.09	1.00	1.18
Crying at night for 5 days or over in a week									
No NRFS	60,054	4,452	7.4	Ref			Ref		
NRFS	2,071	176	8.5	1.14	0.99	1.32	1.19	1.03	1.39

Table 2. Association between non-reassuring foetal status and infantile sleep at 1 year of age. *CI* confidence interval, *RR* risk ratio. ^aAdjusted for maternal age at delivery, smoking habits, alcohol consumption, pre-pregnancy body mass index, parity, infertility treatment, type of delivery, gestational age at birth, small for gestational age, infant sex, psychological distress at 1 year after delivery, doctor diagnosis of asthma and atopic dermatitis at 1 year old, and feeding status.

crying at night = 1.19, 95% CI 1.03–1.39, RR for bedtime after 22:00 = 1.09, 95% CI 1.00–1.18). *tpNRFS* was not associated with any outcomes.

Discussion

This study showed that children with NRFS before birth tended to have sleep problems at 1 year old. The association of *fpNRFS* with sleep problems was similar to that of all NRFS with sleep problems. This finding was attributed to children with *fpNRFS* comprising most of the group with all NRFS. *tpNRFS* was not associated with any outcomes; however, the number of *tpNRFS* cases was few, limiting the ability to make conclusions.

The present study had two major limitations. First, there could be unmeasured confounding factors, such as parental life rhythm and other siblings. Second, child sleep was evaluated using a questionnaire for mothers, which might introduce some bias. On the other hand, a strong point of the present study was the large sample size collected nationwide.

We previously reported an association between NRFS and temperament at 1 month after birth¹⁴. The present study showed that NRFS before birth appeared to influence sleep at 1 year. The possible reason for this is that temperament and sleep are formed prenatally^{22,23}, and foetal temperament affects heart rate reaction to stress.

It is conceivable that children with sleep problems in early infancy are more likely to have developmental disorders than those without sleep problems. Thus, careful follow up of neonates showing NRFS may lead to early detection of developmental disorders.

On the other hand, the RRs for sleeping problems in the NRFS group for sleep problems were not very large. Further investigations are needed to confirm an association between NRFS and sleeping problems in infants.

In conclusion, *fpNRFS* may be associated with sleep problems in 1-year-old infants.

Ethical approval. The study protocol was approved by the Ministry of the Environment's Institutional Review Board on Epidemiological Studies (No. 100406001) and the Ethics Committee of all participating institutions. Written informed consent was obtained from all participants.

Received: 20 January 2020; Accepted: 15 June 2020

Published online: 10 July 2020

References

- Paterson, S. J. *et al.* The importance of temperament for understanding early manifestations of autism spectrum disorder in high-risk infants. *J. Autism Dev. Disord.* <https://doi.org/10.1007/s10803-019-04003-2> (2019).
- Sacrey, L.-A.R. *et al.* Temperament and its association with autism symptoms in a high-risk population. *J. Abnorm. Child Psychol.* **44**, 757–769 (2015).
- Humphreys, J. S. *et al.* Sleep patterns in children with autistic spectrum disorders: A prospective cohort study. *Arch. Dis. Child.* **99**, 114–118 (2014).
- Nguyen, A. K. D., Murphy, L. E., Kocak, M., Tylavsky, F. A. & Pagani, L. S. Prospective associations between infant sleep at 12 months and autism spectrum disorder screening scores at 24 months in a community-based birth cohort. *J. Clin. Psychiatry* **79**, 16m11127 (2018).
- Sun, Y., Hons, B., Cistulli, P. A. & Hons, M. Childhood Health and Educational Outcomes Associated With Maternal Sleep Apnea: A Population Record-Linkage Study (2014).
- Tasnim, N., Mahmud, G. & Akram, S. Predictive accuracy of intrapartum cardiotocography in terms of fetal acid base status at birth. *J. Coll. Physicians Surg. Pak.* **19**, 632–635 (2009).
- Curzen, P., Bekir, J. S., McLintock, D. G. & Patel, M. Reliability of cardiotocography in predicting baby's condition at birth. *Obstet. Gynecol. Surv.* **40**, 344–345 (1985).
- Beard, R. W., Filshie, G. M., Knight, C. A. & Roberts, G. M. The significance of the changes in the continuous fetal heart rate in the first stage of labour. *BJOG An Int. J. Obstet. Gynaecol.* **78**, 865–881 (1971).
- Ernst, G. Heart-rate variability—more than heart beats?. *Front. Public Heal.* **5**, 1–12 (2017).
- Fink, A. M., Bronas, U. G. & Calik, M. W. Autonomic regulation during sleep and wakefulness: a review with implications for defining the pathophysiology of neurological disorders. *Clin. Auton. Res.* **28**, 509–518 (2018).
- Harder, R. *et al.* Heart rate variability during sleep in children with autism spectrum disorder. *Clin. Auton. Res.* **26**, 423–432 (2016).
- Precenzano, F., Ruberto, M. & Parisi, L. Sleep habits in children affected by autism spectrum disorders: a preliminary case-control study. *Acta Medica Mediterr.* **33**, 405–409 (2017).
- Richdale, A. L. & Prior, M. R. The sleep/wake rhythm in children with autism. *Eur. Child Adolesc. Psychiatry* **4**, 175–186 (1995).
- Morokuma, S. *et al.* Non-reassuring foetal status and neonatal irritability in the Japan Environment and Children's Study: a cohort study. *Sci. Rep.* **8**, 1–7 (2018).
- Michikawa, T. *et al.* Baseline profile of participants in the Japan Environment and Children's Study (JECS). *J. Epidemiol.* **28**, 99–104 (2018).
- Kawamoto, T. *et al.* Rationale and study design of the Japan environment and children's study (JECS). *BMC Public Health* **14**, 25 (2014).
- Okai, T. *et al.* Intrapartum management guidelines based on fetal heart rate pattern classification. *J. Obstet. Gynaecol. Res.* **36**, 925–928 (2010).
- Galland, B. C., Taylor, B. J., Elder, D. E. & Herbison, P. Normal sleep patterns in infants and children: a systematic review of observational studies. *Sleep Med. Rev.* **16**, 213–222 (2012).
- Kessler, R. C. *et al.* Screening for serious mental illness in the general population. *Arch. Gen. Psychiatry* **60**, 184–189 (2003).
- Toshia, F., Norito, K., Mari, S. & Yutaka, O. The performance of the Japanese version of the K6 and K10 in the World Mental Health Survey Japan. *Int. J. Methods Psychiatr. Res.* **17**, 152–158 (2008).
- Sakurai, K., Nishi, A., Kondo, K., Yanagida, K. & Kawakami, N. Screening performance of K6/K10 and other screening instruments for mood and anxiety disorders in Japan. *Psychiatry Clin. Neurosci.* **65**, 434–441 (2011).
- DiPietro, J. A., Ghera, M. M. & Costigan, K. A. Prenatal origins of temperamental reactivity in early infancy. *Early Hum. Dev.* **84**, 569–575 (2008).
- Mirmiran, M., Maas, Y. G. H. & Ariagno, R. L. Development of fetal and neonatal sleep and circadian rhythms. *Sleep Med. Rev.* **7**, 321–334 (2003).

Acknowledgements

We would like to express our gratitude to all the participants in this study and all the individuals involved in data collection. The idea of this work was obtained from other works supported by RIKEN Healthcare and Medical Data Platform Project and JSPS KAKENHI (Grant numbers: JP 16H01880, JP 16K13072, JP 18H00994, JP18H03388). The Japan Environment and Children's Study was funded by the Ministry of the Environment, Japan. The findings and conclusions of this article are solely the responsibility of the authors and do not represent the official views of the Ministry of the Environment.

Author contributions

Study conception and design: S.M. Statistical analyses: T.M. Drafting of the manuscript and approval of the final content: K.N., S.M., and T.M. Critical revision of the manuscript for important intellectual content and manuscript review: K.N., T.M., S.M., M.O., K.K.(Kiyoko Kato), M.S.(Masafumi Sanefuji), E.S., M.T., M.S.(Masayuki Shimono), T.K., S.O., K.K.(Koichi Kusuhara), and the JECS Group members.

Competing interests

The authors declare no competing interests.

Additional information

Correspondence and requests for materials should be addressed to S.M.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2020

The Japan Environment and Children's Study Group

Michihiro Kamijima¹⁰, Shin Yamazaki¹¹, Yukihiro Ohya¹², Reiko Kishi¹³, Nobuo Yaegashi¹⁴, Koichi Hashimoto¹⁵, Chisato Mori¹⁶, Shuichi Ito¹⁷, Zentaro Yamagata¹⁸, Hidekuni Inadera¹⁹, Takeo Nakayama²⁰, Hiroyasu Iso²¹, Masayuki Shima²², Youichi Kurozawa²³, Narufumi Suganuma²⁴ & Takahiko Katoh²⁵

¹⁰Nagoya City University, Nagoya, Japan. ¹¹National Institute for Environmental Studies, Tsukuba, Japan.

¹²National Center for Child Health and Development, Tokyo, Japan. ¹³Hokkaido University, Sapporo, Japan.

¹⁴Tohoku University, Sendai, Japan. ¹⁵Fukushima Medical University, Fukushima, Japan. ¹⁶Chiba University, Chiba, Japan.

¹⁷Yokohama City University, Yokohama, Japan. ¹⁸University of Yamanashi, Chuo, Japan. ¹⁹University of Toyama, Toyama, Japan.

²⁰Kyoto University, Kyoto, Japan. ²¹Osaka University, Suita, Japan. ²²Hyogo College of Medicine, Nishinomiya, Japan.

²³Tottori University, Yonago, Japan. ²⁴Kochi University, Nankoku, Japan.

²⁵Kumamoto University, Kumamoto, Japan.