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# Identification of neurodevelopmental transition patterns from infancy to early childhood and risk factors predicting descending transition

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It is unclear whether neurodevelopmental progress from infancy to early childhood remains stable. Moreover, little is known about the risk factors, if any, affecting neurodevelopmental descending transition patterns and the relationship between these patterns and later childhood adaptive behaviours. We used data of 875 children from the Hamamatsu Birth Cohort Study in Japan. Children's neurodevelopment at 18 and 32 months and adaptive behaviours at 40 months were evaluated. Perinatal factors and infant overweight status at 18 months were investigated to identify descending-transition-associated risk factors. In the latent transition analysis, ultimately, three classes were identified for each time-point, resulting in nine transition patterns; among them, 10.4% of children showed descending class shifts (normal to delayed class). Such decelerated growth was predicted by maternal pre-pregnancy overweight status (odds ratio [OR] 2.49; 95% confidence interval [CI] 1.23, 5.02), low maternal educational history (OR 1.20; 95% CI 1.04, 1.36), and infant overweight status at 18 months (OR 5.89; 95% CI 1.26, 27.45). Children with descending transition showed poor functioning in adaptive behaviours at the age of 40 months. To prevent subsequent poor adaptive functioning, it may be necessary to consider that a certain percentage of children show decelerated growth.

An understanding of the neurodevelopmental trajectories in early life provides a clue to predict developmental pathways in later life<sup>1,2</sup>. Neurodevelopmental delays during early infancy negatively impact childhood development<sup>3</sup>. Moreover, neurodevelopmental progress from infancy to early childhood may be inconsistent, and early delays develop in very heterogeneous ways<sup>4</sup>. Some children show typical development at first time-point of measurement but deviate from typical progress at a later time-point. However, these children do not necessarily exhibit developmental regression, which is defined as loss of previously acquired skills<sup>5</sup>. Such a developmental course, which we call a descending transition, has seldom been reported. On the other hand, catching-up transitions, referring to neurological maturation retardation at certain time-points<sup>4</sup>, are well-known. However, the term 'developmental delays' has no clear definition and is often arbitrarily defined using descriptive-statistical measures such as standard deviations and percentile values<sup>6,7</sup>. Therefore, it is critically important to define the neurodevelopmental status or subtype based on the achievement level of development at each time-point to comprehend the children's neurodevelopmental transition. While many studies have comprehensively assessed early neurodevelopment and its trajectories<sup>8,9</sup>, no previous study has comprehensively evaluated a range of transition patterns which include the descending transition.

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Furthermore, little is known about risk factors that may influence such descending transitions. Hillemeier et al.<sup>6</sup> found that prenatal and obstetric factors such as low maternal education, low family income, and preterm birth had an increasing effect on descending transitions. However, the developmental pathways of children are also affected by postnatal factors, including physical growth, as well as biological and environmental factors<sup>10</sup>.

The relationship between a descending course of neurodevelopment during early childhood and untoward consequences in everyday functioning, including adaptive behaviours, at a later stage has attracted attention. While this connection has already been investigated, it only exists in a high-risk sample of siblings of children with autism spectrum disorder (ASD)<sup>11</sup>.

Hence, this study has three aims. First, in a birth cohort sample of Japanese children (18–32 months old), to apply a statistical modelling technique (latent transition analysis; LTA)<sup>12</sup> to identify classes with distinct transition patterns. Second, to explore risk factors, including pre-pregnancy maternal variables, environmental indicators, and infant physical growth variables, which could predispose children to specific transition patterns, especially focusing on a diversion from normal to delayed development. Third, to determine whether children with descending transition patterns show poorer performance in adaptive behaviours at a subsequent stage (age 40 months).

## Methods

**Ethics approval.** The study protocol was approved by the Hamamatsu University School of Medicine and University Hospital Ethics Committee (Ref Nos. 17-037 and 17-037-3) and performed in accordance with the Declaration of Helsinki. Written informed consent was obtained from each mother for her own and her infant's participation.

**Participants.** This study is part of the ongoing cohort study, the Hamamatsu Birth Cohort Study for Mothers and Children (HBC Study). Participants included mothers ( $n=1138$ ) and their infants ( $n=1258$ ) born between 24 December 2007 and 19 March 2012. The detailed recruitment procedure has been previously described<sup>9</sup>. The participants were determined to be a representative sample of the general Japanese population<sup>13</sup>.

In applying the exclusion criteria, the final sample in this study comprised 875 infants and 795 mothers (Supplementary Fig. S1); Two infants whose mothers had passed away before their first birthdays, an infant with a birth weight less than 1000 g, two infants diagnosed with Down syndrome, and twins and other multiple births ( $n=25$ ) were also excluded. Additionally, 353 infants who failed to participate in outcome assessments at 18 or 32 months of age (henceforth referred to as time 1 and time 2, respectively), or both, were excluded.

**Measures.** *Neurodevelopment.* Neurodevelopmental progress was assessed using the Mullen Scales of Early Learning (MSEL) at time 1 and time 2<sup>14</sup>. MSEL is a validated composite scale for ascertaining child neurodevelopment through direct testing and is composed of five domains: gross and fine motor, visual reception, and expressive and receptive language. A Japanese version of the associated T-score, with a mean of 50 and a standard deviation (SD) of 10, was used as the outcome. The detail of the T-score and procedures of the direct testing of MSEL and retaining the reliability of assessments among clinical evaluators have been described previously<sup>9</sup>.

*Risk factors for neurodevelopmental transition.* Risk factors associated with neurodevelopmental transition patterns in children, especially descending patterns from time 1 to time 2, were explored (Supplementary Note 1). A perinatal factor, small-for-gestational-age (birth weight < 10th percentile for gestational age)<sup>15</sup> was applied. Moreover, infant overweight status at time 1 was incorporated as another risk index<sup>16</sup>. Child weight and height at 18 months of age were obtained from records taken from periodic health check-ups. Child BMI values were transformed into a standardized body mass index (sBMI) corresponding to a BMI-for-age value with a mean of 0 and an SD of 1, based on the World Health Organization's Child Growth Standards<sup>17</sup>. Subsequently, sBMI scores were dichotomised into 'overweight' (over 1 SD) and 'others'. The mother's pre-pregnancy overweight status was also included as another biological variable; self-reported pre-pregnancy BMI ( $\text{kg}/\text{m}^2$ ) during early pregnancy was dichotomised as 'overweight' (over  $25 \text{ kg}/\text{m}^2$ ) and 'others'<sup>18</sup>. A history of maternal education (a continuous variable) obtained through an interview was also used.

**Covariates.** Infant sex, premature birth (< 37 weeks), low placenta-to-birth-weight ratio (< 10th percentile)<sup>19</sup>, parental age, paternal educational history, and annual household income were used as covariates for multinomial logistic regression. Data relating to the demographic characteristics of parents were collected during pregnancy, and perinatal variables were obtained from medical records.

*Adaptive behaviours.* The everyday functional levels of each child were quantified using the vineland adaptive behaviour scales second edition (VABS-II) at 40 months of age. The VABS is based on a semi-structured parental interview consisting of four domains: communication, daily living, socialization, and motor skills<sup>20</sup>. An overall adaptive behaviour composite standardized score, with a mean of 100 and an SD of 15, was used.

**Statistical analysis.** *Latent transition analysis.* LTA was used to compute the latent transition probabilities which suggest the likelihood of individuals changing classes or remaining in the same class across consecutive periods<sup>12</sup>. LTA may compute transition probabilities at multiple time-points, such as three time-points; however, the precision of estimation might become inferior owing to the involvement of a multiplicative increase in parameters. Therefore, we opted for two time-points (e.g., time 1 and time 2) to achieve our goal, which was to identify classes with distinct transition patterns. Prior to the main analysis, five domains of MSEL T-scores

at time 1 and time 2 were set as the outcome measures through latent class analysis (LCA). To select the appropriate class solution in latent class analysis (LCA), we used five common fit indices<sup>21</sup>: the Akaike information criterion (AIC), Bayesian information criterion (BIC), entropy, adjusted Lo-Mendell Rubin likelihood ratio test (LMR-LRT), and bootstrap likelihood ratio test (BLRT). Along with these indices, the number of classes was ultimately determined by considering theoretical justification and interpretability<sup>22</sup>. After the best solution at each time-point was identified via LCA, the transition probability by which each individual was assigned to an optimal transition pattern was computed using LTA without potential risk factors and covariates. For the LTA procedure, a slightly modified version of the three-step approach was followed (Supplementary Note 1)<sup>23,24</sup>. In this approach, potential risk factors (maternal pre-pregnancy overweight status, low maternal educational history, small-for-gestational-age, and infant overweight status at time 1) and covariates (infant sex, premature birth, placenta-to-birth-weight ratio, and household income) which differed across classes at time 1 were included in the analysis (Supplementary Note 1) and examined to determine which factors would emerge as affecting transition patterns (i.e., class shifts) using multinomial logistic regression.

*Transition pattern effect on adaptive behaviours.* The associations of transition patterns between time 1 and time 2, especially descending transitions, with adaptive behaviours at 40 months were investigated using linear regression analysis, wherein risk factors identified in LTA were included as covariates to account for their potential confounding effects. Familial clustering was controlled for using the Huber–Sandwich method. The statistical analyses in the present study were conducted using Mplus version 8 (<https://www.statmodel.com/>)<sup>25</sup> and Stata version 14.0 (<https://www.stata.com/stata14/>)<sup>26</sup>.

**Data attrition.** Infants who received MSEL evaluations at time 1 and 2 were included. The rate of missing data from the five MSEL domains was minimal (1.6% in total). The full information maximum likelihood (FIML) method, a powerful tool for missing data, was used<sup>27</sup>. Any biases arising from the assumption of missing at random for FIML were therefore considered negligible.

## Results

**Neurodevelopmental transition patterns from infancy to early childhood.** Table 1 shows the participants' demographic characteristics included in the analysis. In LCA, fit indices in latent class analysis are shown in Supplementary Table S1. We investigated up to seven class solutions. For LCA time 1 (i.e., 18 months), the adjusted LMR-LRT showed  $P < 0.05$  up to the five-class solution, whereas BIC was smallest for the six-class solution. AIC continuously reduced, while BLRT remained significant up to the seven-class solution. As for LCA time 2 (i.e., 32 months), the adjusted LMR-LRT similarly showed  $P < 0.05$  up to the five-class solution, while BIC had the smallest value for the six-class solution. AIC continued to decrease, and BLRT showed  $P < 0.001$  up to the seven-class solution. These findings indicate the optimal solution was five classes or more for both time 1 and time 2. In our previous study of the same birth cohort, we found that five classes would best delineate the distinctive trajectories of neurodevelopment over the period of infancy (1–24 months), including the current time-point (i.e., time 1)<sup>9</sup>. For comparisons and facilitation of interpretations, we opted for the five-class solution for both time 1 and time 2. Figure 1 shows the MSEL composition of the five classes identified by the LCA procedures. At time 1 (Fig. 1a), three classes within  $50 \pm 1$  SD (i.e., leftmost to the third column in the figure) were designated as 'normal', consisting of 'high normal' (21.9%), 'normal' (31.8%), and 'low normal' (40.9%), in order from left to right. The fourth group of children (2.2%) showed a downward deviation ( $-2$  SD) only in the expressive language domain (denoted as 'expressive language (EL)-delayed'). The last group (3.2%) showed low scores (below  $-1$  SD) in all five domains ('delayed'). Similarly, three normal classes, 'high normal' (7.8%), 'normal' (32.9%), and 'low normal' (46.3%), were identified at time 2 (Fig. 1b). The fourth group (9.9%) showed scores below  $-1$  SD in all five domains ('delayed'). The last group (3.1%) showed markedly low scores in all five domains ('markedly (M)-delayed').

Table 2 shows counts and proportions for each combination of class assignments from time 1 to time 2. Only a small number of children ( $n < 9$ , corresponding to less than 1%) was allocated to 12 out of the 25 observable transition classes, including the descending pattern classes. Thus, the three normal classes were amalgamated at each time-point and labelled as '3-normals' to increase statistical power during relative risk estimations of children assigned to descending transition classes in comparison with baseline control classes (i.e., normal-to-normal transition patterns). Thus, the number of transition patterns was reduced to nine. Of the 875 infants included in the analyses, 737 (84.2%) were assigned to the reference transition pattern of '3-normals to 3-normals' at both time-points, 91 (10.4%) to the descending transition patterns ("3-normals to delayed" and "3-normals to M-delayed"), while 24 (2.8%) were assigned to the catching-up transition patterns ("EL-delayed to 3-normals" and "delayed to 3-normals"), and 23 (2.6%) were assigned to the four transition patterns of remaining in delayed classes at both time-points.

**Risk factors for transition patterns.** Table 3 shows associations between the risk factors for infants with descending and catching-up transition patterns compared with infants with the reference transition pattern (i.e., "3-normals to 3-normals" at both time-points). Children whose mothers had a pre-pregnancy overweight status had a 2.49-fold increase in the risk of being assigned to the "3-normals to delayed" transition pattern than children with the reference transition pattern (odds ratio [OR] 2.49; 95% confidence interval [CI] 1.23, 5.02). Low maternal educational history was also associated with this descending transition pattern. A 1-year decrease in maternal educational history corresponded to a 20% increase in the risk of infants being assigned to the "3-normals to delayed" transition pattern (OR 1.20; 95% CI 1.04, 1.36). Overweight status at time 1 in infants was associated with being assigned to the "3-normals to M-delayed" transition pattern (OR 5.89; 95%

Infants	Mean (SD)
Birth weight (g)	2957.0 (422.2)
Gestational age at birth (weeks)	39.0 (1.4)
n (%)	
<b>Sex</b>	
Male	432 (49.4%)
Female	443 (50.6%)
<b>Prematurity</b>	
< 37 weeks	46 (5.3%)
≥ 37 weeks	829 (94.7%)
<b>Small-for-gestational-age</b>	
< 10th percentile	77 (8.8%)
10th–100th percentile	798 (91.2%)
<b>Age-standardized BMI scores at 18 months</b>	
> 1 standard deviation	50 (6.1%)
< 1 standard deviation	776 (93.9%)
<b>Parents</b>	
Maternal BMI at pre-pregnancy ( $\text{kg}/\text{m}^2$ )	n (%)
> 25.0	100 (11.4%)
< 25.0	775 (88.6%)
<b>Placenta-to birth-weight ratio</b>	
< 10th percentile	152 (17.7%)
10th–100th percentile	707 (82.3%)
Mean (SD)	
Paternal age at birth (year)	33.5 (5.7)
Maternal age at birth (year)	31.7 (5.0)
Paternal education (year)	14.1 (2.6)
Maternal education (year)	13.8 (1.8)
Household income (million JPY)	6.0 (2.7)

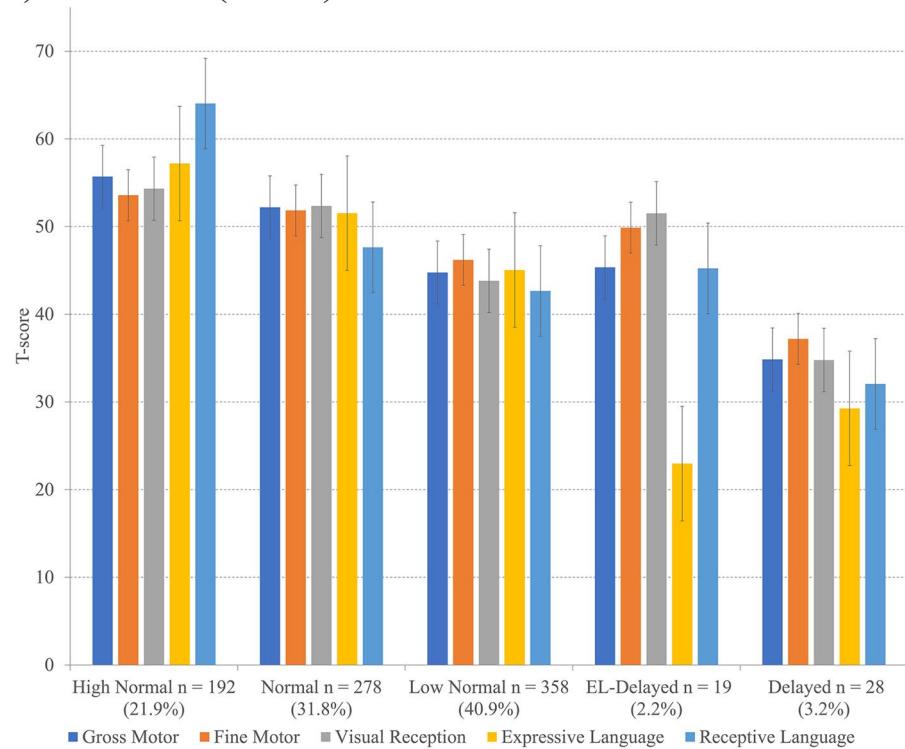
**Table 1.** Characteristics of participating infants and their parents. *SD* standard deviation, *BMI* body mass index, *JPY* Japanese Yen.

CI 1.26, 27.45). For catching-up transition patterns, low maternal educational history was also associated with assignment to EL-delayed to 3-normals” transition pattern (OR 1.21; 95% CI 1.01, 1.49), and infants with SGA were more likely to be allocated to the “delayed to 3-normals” transition pattern (OR 10.13; 95% CI 2.23, 45.85). Associations between covariates and neurodevelopmental transition patterns in the final model are shown in Supplementary Table S2 and Note 2.

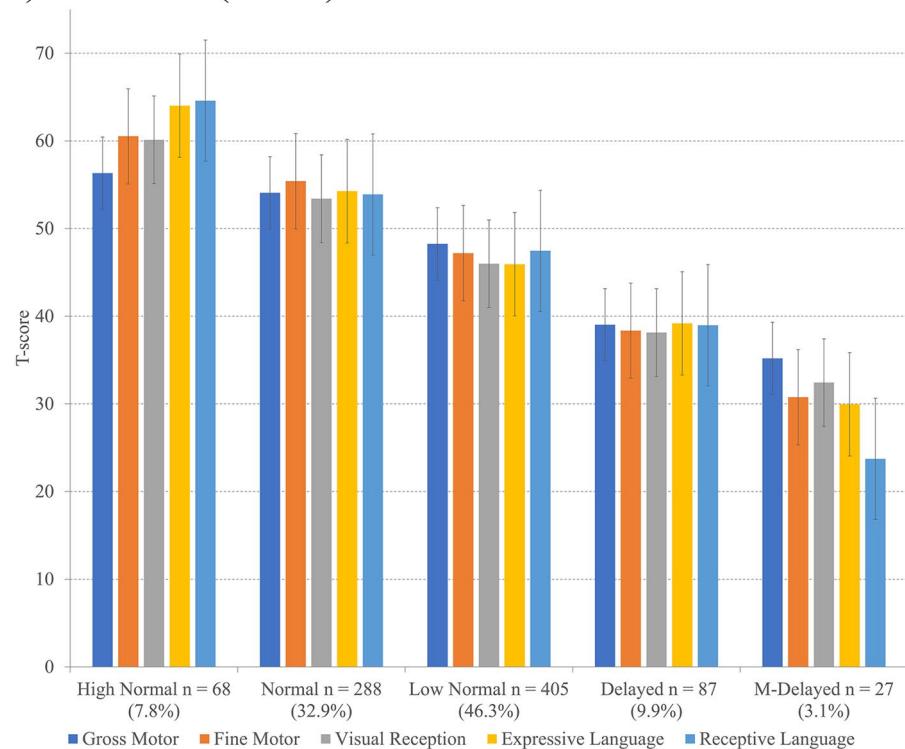
**Transition patterns and adaptive behaviour.** Table 4 presents the associations between the nine transition patterns and the adaptive behaviour composite standardized scores at 40 months of age. Compared with children with the reference transition pattern, children with the two descending transition patterns showed lower adaptive scores 8 months later (“3-normals to delayed”: coefficient = -7.88; 95% CI -9.77, -6.00 and “3-normals to m-delayed”: coefficient = -13.04; 95% CI -18.87, -7.21). Children who stayed in delayed transition classes also showed poorer levels of adaptive ability. Conversely, children who achieved the two catching-up transition patterns had no significant differences in adaptive functioning when compared with infants in the reference transition group (“EL-delayed to 3-normals”: coefficient = -2.26; 95% CI -5.04, 0.50 and “delayed to 3-normals”: coefficient = 0.50; 95% CI -3.98, 4.99).

**Data attrition.** Notably, characteristics (i.e., sex, birth weight, gestational age at birth, and prematurity) of infants and parental background characteristics (i.e., parental age and maternal educational history) differed between the infants included and those excluded from the analyses (Supplementary Table S3). These differences may have biased the findings. The inverse probability weighting method<sup>28</sup> was applied to the final model to allow for missing variations associated with these six variables. The estimates derived from the inverse probability weighting model for descending transition patterns remained statistically significant, with ORs of 2.2 (95% CI 1.32, 3.79) for maternal overweight at pre-pregnancy, 3.89 (95% CI 1.25, 12.06) for infant overweight at time 1, 4.16 (95% CI 1.40, 12.28) except for maternal educational history, which had an almost identical estimation but fell just short of statistical significance (OR 1.12; 95% CI 0.99, 1.26).

## a) 18 months (time 1)



## b) 32 months (time 2)



**Figure 1.** Latent class structure by neurodevelopmental domains at 18 months (A) and 32 months (B) ( $n=875$ ). EL-delayed expressive language delayed, M-delayed markedly delayed.

Latent class at time 1, n (%)	Combined 3-normals	Latent class at time 2, n (%)				Delayed 87 (9.9%)	M-delayed 27 (3.1%)		
		Combined 3-normals							
		High-Normal 68 (7.8%)	Normal 288 (32.9%)	Low-normal 405 (46.3%)					
		High-normal 192 (21.9%)	49 (5.6%)	87 (9.9%)	56 (6.4%)	0 (0%)	0 (0%)		
Latent class at time 1, n (%)	Combined 3-normals	Normal 278 (31.8%)	12 (1.4%)	132 (15.1%)	130 (14.5%)	0 (0%)	4 (0.5%)		
		Low-normal 358 (40.9%)	7 (0.8%)	61 (7.0%)	203 (23.2%)	74 (8.5%)	13 (1.5%)		
		EL-delayed 19 (2.2%)	0 (0%)	6 (0.7%)	9 (1.0%)	3 (0.3%)	1 (0.1%)		
		Delayed 28 (3.2%)	0 (0%)	2 (0.2%)	7 (0.8%)	10 (1.1%)	9 (1.0%)		

**Table 2.** Transition class counts and proportions derived by latent transition analysis ( $n=875$ ). *EL-delayed* expressive language delayed, *M-delayed* markedly delayed.

	Transition from 3-normals at time 1 (765)			
	To 3-normals at time 2 (681)	Delayed at time 2 (70)	Markedly delayed at time 2 (14)	
	Reference	OR [95% CI]	OR [95% CI]	
<b>Possible risk factors</b>				
Maternal overweight status at pre-pregnancy	..	2.49 [1.23, 5.02]*	1.27 [0.20, 8.02]	
Low maternal education (year)	..	1.20 [1.04, 1.36]*	1.12 [0.74, 1.69]	
Small-for-gestational-age	..	1.81 [0.72, 4.53]	0.89 [0.10, 7.93]	
Infant overweight status at 18 months	..	1.49 [0.57, 3.88]	5.89 [1.26, 27.45]*	
<b>Transition from expressive language delayed at time 1 (19)</b>				
To 3-normals at time 2 (15)		Delayed at time 2 (3)	Markedly delayed at time 2 (1)	
OR [95% CI]		OR [95% CI]	OR [95% CI]	
<b>Possible risk factors</b>				
Maternal overweight status at pre-pregnancy	0.83 [0.10, 6.69]	NA	NA	
Low maternal education (year)	1.21 [1.01, 1.49]*	0.91 [0.46, 1.81]	1.96 [1.58, 2.43]****	
Small for gestational age	0.93 [0.11, 7.53]	3.85 [0.010, 1409.03]	NA	
Infant overweight status at 18 months	NA	NA	NA	
<b>Transition from delayed at time 1 (27)</b>				
To 3-normals at time 2 (9)		Delayed at time 2 (10)	Markedly delayed at time 2 (8)	
OR [95%CI]		OR [95%CI]	OR [95%CI]	
<b>Possible risk factors</b>				
Maternal overweight status at pre-pregnancy	4.79 [0.91, 25.26]	0.90 [0.09, 8.57]	1.40 [0.17, 11.37]	
Low maternal education (year)	1.09 [0.74, 1.61]	1.26 [0.85, 1.88]	1.42 [0.86, 2.32]	
Small for gestational age	10.13 [2.23, 45.85]***	NA	NA	
Infant overweight status at 18 months	2.78 [0.24, 31.99]	2.15 [0.24, 19.18]	NA	

**Table 3.** Association between possible risk factors and neurodevelopmental transition patterns in the final model of multinomial logistic regression analysis with covariates ( $n=811$ ). Infant's sex (male), premature birth (< 37 weeks), low placenta-to-birthweight ratio (< 10th percentile), and household income. Fifty infants who had missing values for infant's sBMI at 18 months and 14 mother-infant dyads who had missing values for mother's placental weight were excluded from this analysis. *OR* odds ratio, *CI* confidence interval, *NA* not available. \* $P<0.05$ , \*\* $P<0.005$ , \*\*\* $P<0.001$ .

## Discussion

LTA was applied to multiple neurodevelopmental measures collected from infancy to early childhood in a Japanese birth cohort sample and found that 10.4% of children were diverted from normal neurodevelopment levels (3-normals) to lower levels, implying decelerated developmental patterns. Maternal pre-pregnancy overweight status, infant overweight status at time 1 (i.e., 18 months), and low maternal educational history were identified as risk factors associated with descending transition patterns. Further, children assigned to descending transition patterns or who remained in delayed neurodevelopmental classes from time 1 to time 2 (i.e., 32 months) demonstrated poorer adaptive functioning at 40 months. In contrast, children with catching-up transition patterns attained a normal range of adaptive functioning at 40 months.

Our study showed that 5.4% of infants (47/875) were assigned to the delayed classes at time 1 and 13.0% (114/875) to the delayed classes at time 2, indicating that there was an overall increase in the rate of delayed

	Adaptive behaviour composite standardized score
	Coefficient [95% CI]
<b>Transition patterns (n)</b>	
3-Normals to 3-normals (654)	Reference
3-Normals to delayed (67)	-7.88 [-9.77, -6.00]****
3-Normals to markedly delayed (13)	-13.04 [-18.87, -7.21]****
EL-delayed to 3-normals (14)	-2.26 [-5.04, 0.50]
Delayed to 3-normals (9)	0.50 [-3.98, 4.99]
EL-delayed to delayed (3)	-11.49 [-16.9, -6.01]****
EL-delayed to markedly delayed (1)	-14.13 [-15.20, -13.06]****
Delayed to delayed (10)	-9.76 [-13.39, -6.13]****
Delayed to markedly delayed (8)	-17.48 [-21.77, -13.19]****

**Table 4.** Association between neurodevelopmental transition patterns and adaptive behaviour at 40 months in linear regression analyses with covariates ( $n=779$ ). CI confidence intervals, EL-delayed expressive language delayed. \*\*\*\* $P<0.001$ . <sup>a</sup>Infant sex (male), premature birth (< 37 weeks), low placenta-to-birth-weight ratio (< 10th percentile), household income, maternal body mass index at pre-pregnancy ( $> 25 \text{ kg/m}^2$ ), small-for-gestational-age (< 10th percentile), infant's standardised body mass index at 18 months of age ( $> 1 \text{ SD}$ ), and maternal educational history.

infants over the study period. Further, 10.4% of infants were identified as having descending transition patterns from time 1 to time 2. A previous study based on a Taiwanese nationwide sample<sup>29</sup> reported that the prevalence of neurodevelopmental delay from 1 to 3 years of age increased about three to four-fold, suggesting that, akin to our findings, a substantial proportion of children show a descending developmental pattern from infancy to early childhood. Approximately half (51%) the infants ( $n=24$ ) who belonged to delayed classes (i.e., 'expressive language', EL-delayed, and 'delayed' classes) at time 1 ( $n=47$ ) moved to normal classes (i.e., '3-normals') at time 2, showing catching-up transition patterns. This finding is inconsistent with those of previous studies reporting that approximately 80–90% of children who initially belonged to a delayed class eventually caught up<sup>6,7</sup>. In these studies, however, 'delayed class' was arbitrarily defined using deviations from an average (e.g., below  $-1 \text{ SD}$ ); thus, the prevalence of developmental delays in infancy might be overestimated. Infants in the EL-delayed class in our study may include the so-called 'late talkers' (LT), or children under 3 years old who have unusually small vocabularies with unknown primary causes<sup>30</sup>. Overall, children with LT have a good prognosis, with 60–70% of children moving into the average range on language measures by preschool<sup>31</sup>. Our study supports this finding in that 79% of infants assigned to the EL-delayed class at time 1 caught up to the normal classes by time 2.

Maternal pre-pregnancy overweight status was a risk factor for a descending transition pattern. This finding supports previous studies which associated maternal pre-pregnancy overweight status with motor<sup>32</sup>, language<sup>33</sup>, and cognitive neurocognitive developmental delay in offspring<sup>34</sup>. Maternal pre-pregnancy overweight status might produce a chronic systemic inflammatory ambience in both mother and foetus, with long-lasting negative consequences for neuronal development in offspring<sup>35</sup> by affecting insulin and leptin levels and other inflammatory markers<sup>36</sup>. Girchenko et al.<sup>37</sup> indicated that higher levels of maternal inflammation of high-sensitivity C-reactive protein and glycoprotein acetyls mediated the effect of prenatal environmental adversity including maternal early pregnancy overweight on child neurodevelopmental delay.

Infant overweight status at time 1 was considered another risk factor for descending transition patterns. Overweight status in infancy is known to be related to neurodevelopmental delays<sup>16,38</sup> and neuroimaging research has accumulated evidence on the relationship between overweight status and structural changes in the brain<sup>38,39</sup>. Obesity may cause reductions in brain volume through inflammatory responses produced by adipose tissue<sup>40</sup>. Thus, overweight status during infancy, when brains are developing rapidly<sup>41</sup>, may negatively impact healthy brain development and lead to descending transition patterns in early childhood.

Low maternal education was associated with the descending transition pattern. Maternal educational history appears to be associated with variables indicating poor parenting environments, including low maternal responsiveness and lack of effective communication<sup>42</sup>, and may result in more developmental differences<sup>43</sup>. Genetic or inherent influences may become more salient during the later stage of infancy. For example, a declining trajectory of development has frequently been reported in children with neurodevelopmental disorders such as ASD<sup>44</sup>, and lower levels of maternal education may be a risk factor for neurodevelopmental disorders<sup>45</sup>.

The results of the present study showed that the risk factors are independently associated with the descending neurodevelopmental trajectory. However, it has been reported that these risk factors are associated with each other. It has been reported that maternal obesity had an adverse effect on child obesity<sup>46</sup>, and obesity in early childhood was more commonly seen among those born to mothers with low education<sup>47</sup>. Therefore, further investigation is needed to determine the association among these risk factors and child neurodevelopmental trajectories including mediation and moderation.

Small for gestational age (SGA) was found to be positively associated with the catching-up transition pattern from the Delayed class to the 3-normals class. Although SGA infants reportedly have lower levels of cognitive, language, and motor development than infants with appropriate gestational age<sup>48</sup>, a catching-up phase of neurodevelopmental progress during early childhood has been noted in these children<sup>49,50</sup>, which is compatible

with the findings of the present study. Growth hormone/insulin-like growth factors have been proposed to be possible mechanisms behind this catching-up phase in children with SGA<sup>51</sup>. Low maternal education level was also associated with catching-up transition patterns from the EL-delayed class to the 3-normals class. However, when the inverse probability weighting method was applied to the model, this finding was no longer significant (OR 1.17, 95% CI 0.91, 1.49), implying that the seeming association might reflect a bias arising from sample selection procedures.

There was no association between risk factors and persistent delayed transition patterns (i.e., from the EL-delayed class and the delayed class at time 1 to the delayed class and the markedly delayed, M-delayed, class at time 2), except that maternal education was associated with the transition pattern of EL-delayed to M-delayed. Although this exception was grounded on a small sample size ( $n=1$ ) and needs further investigations, it has been reported that there was an association between low maternal education and neurodevelopmental delay in children<sup>43</sup>.

It was found that infants with descending transition patterns and persistent delays showed lower adaptive behaviour scores at 40 months of age than infants with the reference transition pattern. Our findings regarding infants with persistent delays are consistent with a previous study, which identified an association between lower neurodevelopmental scores in infancy and poorer adaptive behaviours in siblings of children with ASD<sup>11</sup>. To the best of our knowledge, this is the first study to demonstrate the relationship between developmental transition patterns and later adaptive behaviour in a general population sample.

One of the main strengths of this study was that our population comprised a representative sample of infants; thus, our findings are generalizable. Second, this study examined five domains of neurodevelopment using established instruments and direct evaluations. Third, to the best of our knowledge, this study is the first to report risk factors for predicting descending transition patterns and subsequent poorer adaptive functioning in children at the early developmental stage.

However, this study has some limitations. First, the sample size was comparatively small. Among the 25 initially identified transition patterns, 12 were each composed of less than 1% of children; thus, the three normal classes were combined into a single “3-normals” at both time-points to avoid false-negative findings due to a lack of statistical power. Therefore, the findings, especially the risk factors, should be interpreted with caution. In fact, maternal BMI and infant sBMI were dichotomized in the present study, but when they were treated as continuous variables, the association with descending transition pattern was changed. Although the association between maternal BMI and the descending transition pattern remained significant (OR 1.08, 95% CI 1.01, 1.17), the infant sBMI did not (OR 2.10, 95% CI 0.80, 5.48). In future studies, we recommend verifying our findings through large sample sizes for possible application in clinical practice. Second, potential postnatal risk factors for predicting transition patterns were limited only to infant BMI. Further research is warranted to determine the impact of other growth-related indices, including protective factors, for infants, such as nutritional intake, interpersonal activities at early life stages, and environmental factors (i.e., living conditions and accessibility and utilization of health and social care services). Reliance on parental reports of the adaptive performance in their offspring might also have impacted the outcome measures. It is plausible that the performance of adaptive skills in children reported by parents may have been exaggerated<sup>52</sup>. Nevertheless, no evidence indicated that this bias occurred in combination with specific transition patterns of neurodevelopment; thereby, it is unlikely to have confounded the findings of the association between transition classes and adaptive functioning. However, there remains the possibility that our findings may be overrepresented by a subpopulation of infants, particularly, those with developmental impairments such as ASD. Acquired knowledge and information provided regarding the disorder may have influenced parents’ reporting attitudes in favour of the research questions in the present study. The results, however, remained virtually unchanged when children diagnosed with ASD at time 2 ( $n=26$ ) by paediatricians blinded to any research hypotheses were eliminated from the analyses (Supplementary Tables S4, S5), suggesting that our findings cannot be attributed to a specific group of developmental conditions.

In conclusion, this study underscored the dynamic nature of comprehensive neurodevelopmental progress from 18 to 32 months and identified risk factors associated with descending neurodevelopmental transition patterns during this period. In this representative sample of Japanese children, 10.4% showed descending transition patterns, which were predicted by maternal pre-pregnancy overweight status, low maternal educational history, and infant overweight status at 18 months. It was also found that children with descending transition patterns had lower adaptive behaviours at 40 months. Further studies are warranted to replicate our findings using independent samples in large studies.

## Data availability

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

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## References

1. Landa, R. J., Gross, A. L., Stuart, E. A. & Bauman, M. Latent class analysis of early developmental trajectory in baby siblings of children with autism. *J. Child Psychol. Psychiatry* **53**, 986–996 (2012).
2. Valla, L., Birkeland, M. S., Hofoss, D. & Sløning, K. Developmental pathways in infants from 4 to 24 months. *Child Care Health Dev.* **43**, 546–555 (2017).
3. Emerson, E. & Einfeld, S. Emotional and behavioural difficulties in young children with and without developmental delay: A bi-national perspective. *J. Child Psychol. Psychiatry* **51**, 583–593 (2010).

4. Taboada, E. M., Iglesias, P. M., López, S. & Rivas, R. M. Neurodevelopmental difficulties as a comprehensive category of learning disabilities in children with developmental delay: A systematic review. *Ann. Psychol.* **36**, 271–282 (2020).
5. Zhang, D. *et al.* Towards a consensus on developmental regression. *Neurosci. Biobehav. Rev.* **107**, 3–5 (2019).
6. Hillemeier, M. M., Morgan, P. L., Farkas, G. & Maczuga, S. A. Perinatal and socioeconomic risk factors for variable and persistent cognitive delay at 24 and 48 months of age in a national sample. *Matern. Child Health J.* **15**, 1001–1010 (2011).
7. McManus, B. M., Robinson, C. C. & Rosenberg, S. A. Identifying infants and toddlers at high risk for persistent delays. *Matern. Child Health J.* **20**, 639–645 (2016).
8. Toffol, E. *et al.* Infant regulatory behavior problems during first month of life and neurobehavioral outcomes in early childhood. *Eur. Child Adolesc. Psychiatry* **28**, 847–859 (2019).
9. Nishimura, T., Takei, N., Tsuchiya, K. J., Asano, R. & Mori, N. Identification of neurodevelopmental trajectories in infancy and of risk factors affecting deviant development: A longitudinal birth cohort study. *Int. J. Epidemiol.* **45**, 543–553 (2016).
10. Rutter, M., Moffitt, T. E. & Caspi, A. Gene-environment interplay and psychopathology: Multiple varieties but real effects. *J. Child Psychol. Psychiatry* **47**, 226–261 (2006).
11. Sacrey, L. R. *et al.* Developmental trajectories of adaptive behavior in autism spectrum disorder: A high-risk sibling cohort. *J. Child Psychol. Psychiatry* **60**, 697–706 (2019).
12. Muthén, B. & Muthén, L. K. Integrating person-centered and variable-centered analyses: Growth mixture modeling with latent trajectory classes. *Alcohol. Clin. Exp. Res.* **24**, 882–891 (2000).
13. Takagai, S. *et al.* Cohort profile: Hamamatsu birth cohort for mothers and children (HBC Study). *Int. J. Epidemiol.* **45**, 333–342 (2016).
14. Mullen, E. M. *Mullen Scales of Early Learning Manual* (American Guidance Service, 1995).
15. Itabashi, K. *et al.* Adoption of new reference value for birth size according to gestational age (in Japanese). *J. Jpn. Pediatr. Soc.* **114**, 1271–1293 (2010).
16. Camargos, A. C. R., Mendonça, V. A., Andrade, C. A., Oliveira, K. S. C. & Lacerda, A. C. R. Overweight and obese infants present lower cognitive and motor development scores than normal-weight peers. *Res. Dev. Disabil.* **59**, 410–416 (2016).
17. WHO Multicentre Growth Reference Study Group. *WHO Child Growth Standards: Length/Height-for-Age, Weight-for-Age, Weight-for-Length, Weight-for-Height and Body Mass Index-for-Age: Methods and Development* Vol. 312 (World Health Organization, 2006).
18. World Health Organization. *Obesity: Preventing and Managing the Global Epidemic* (World Health Organization, 2000).
19. Ogawa, Y., Iwamura, T. & Kuriya, N. Birth size standards by gestational age for Japanese neonates (in Japanese). *Acta Neonatol. Jpn.* **34**, 624–632 (1998).
20. Sparrow, S. S., Balla, D. A., Cicchetti, D. V. & Harrison, P. L. *Vineland Adaptive Behavior Scales* (American Guidance Service, 1984).
21. Ausderau, K. K. *et al.* Sensory subtypes in children with autism spectrum disorder: Latent profile transition analysis using a national survey of sensory features. *J. Child Psychol. Psychiatry* **55**, 935–944 (2014).
22. Berlin, K. S., Parra, G. R. & Williams, N. A. An introduction to latent variable mixture modeling (part 2): Longitudinal latent class growth analysis and growth mixture models. *J. Pediatr. Psychol.* **39**, 188–203 (2014).
23. Asparouhov, T. & Muthén, B. Auxiliary variables in mixture modeling: Three-step approaches using M plus. *Struct. Equ. Modeling* **21**, 329–341 (2014).
24. Nylund-Gibson, K., Grimm, R., Quirk, M. & Furlong, M. A latent transition mixture model using the three-step specification. *Struct. Equ. Modeling* **21**, 439–454 (2014).
25. Muthén, L. K. & Muthén, B. O. *Mplus User's Guide* (Muthén & Muthén, 2012).
26. StataCorp, L. *Stata Treatment-Effects Reference Manual* (A Stata Press Publication, 2015).
27. Schafer, J. L. & Graham, J. W. Missing data: Our view of the state of the art. *Psychol. Methods* **7**, 147–177 (2002).
28. Mansournia, M. A. & Altman, D. G. Inverse probability weighting. *BMJ* **352**, i189 (2016).
29. Kuo, H.-T., Muo, C.-H., Chang, Y.-T. & Lin, C.-K. Change in prevalence status for children with developmental delay in Taiwan: A nationwide population-based retrospective study. *Neuropsychiatr. Dis. Treat.* **11**, 1541 (2015).
30. Rescorla, L. Late talkers: Do good predictors of outcome exist? *Dev. Disabil. Res. Rev.* **17**, 141–150 (2011).
31. Dale, P. S., McMillan, A. J., Hayiou-Thomas, M. E. & Plomin, R. Illusory recovery: Are recovered children with early language delay at continuing elevated risk? *Am. J. Speech Lang. Pathol.* **23**, 437–447 (2014).
32. Hinkle, S. N. *et al.* Associations between maternal prepregnancy body mass index and child neurodevelopment at 2 years of age. *Int. J. Obes. (Lond.)* **36**, 1312–1319 (2012).
33. Widen, E. M. *et al.* Prepregnancy overweight and obesity are associated with impaired child neurodevelopment. *Matern. Child Nutr.* **14**, e12481 (2018).
34. Alvarez-Bueno, C., Caverio-Redondo, I., Lucas-de la Cruz, L., Notario-Pacheco, B. & Martínez-Vizcaino, V. Association between pre-pregnancy overweight and obesity and children's neurocognitive development: A systematic review and meta-analysis of observational studies. *Int. J. Epidemiol.* **46**, 1653–1666 (2017).
35. van der Burg, J. W. *et al.* The role of systemic inflammation linking maternal BMI to neurodevelopment in children. *Pediatr. Res.* **79**, 3–12 (2016).
36. Tau, G. Z. & Peterson, B. S. Normal development of brain circuits. *Neuropsychopharmacology* **35**, 147–168 (2010).
37. Girchenko, P. *et al.* Persistently high levels of maternal antenatal inflammation are associated with and mediate the effect of prenatal environmental adversities on neurodevelopmental delay in the offspring. *Biol. Psychiatry* **87**, 898–907 (2020).
38. Slining, M., Adair, L. S., Goldman, B. D., Borja, J. B. & Bentley, M. Infant overweight is associated with delayed motor development. *J. Pediatr.* **157**, 20–25 (2010).
39. Papageorgiou, I. *et al.* Abnormalities of brain neural circuits related to obesity: A diffusion tensor imaging study. *Magn. Reson. Imaging* **37**, 116–121 (2017).
40. Karlsson, H. K. *et al.* Obesity is associated with white matter atrophy: A combined diffusion tensor imaging and voxel-based morphometric study. *Obesity (Silver Spring)* **21**, 2530–2537 (2013).
41. Miller, A. L., Lee, H. J. & Lumeng, J. C. Obesity-associated biomarkers and executive function in children. *Pediatr. Res.* **77**, 143–147 (2015).
42. Blacher, J., Baker, B. L. & Kaladjian, A. Syndrome specificity and mother-child interactions: Examining positive and negative parenting across contexts and time. *J. Autism Dev. Disord.* **43**, 761–774 (2013).
43. Demirci, A. & Kartal, M. Sociocultural risk factors for developmental delay in children aged 3–60 months: A nested case-control study. *Eur. J. Pediatr.* **177**, 691–697 (2018).
44. Ozonoff, S. & Iosif, A. M. Changing conceptualizations of regression: What prospective studies reveal about the onset of autism spectrum disorder. *Neurosci. Biobehav. Rev.* **100**, 296–304 (2019).
45. DiGuiseppi, C. G. *et al.* Demographic profile of families and children in the study to explore early development (SEED): Case-control study of autism spectrum disorder. *Disabil. Health J.* **9**, 544–551 (2016).
46. Cataldo, R. *et al.* Effects of overweight and obesity on motor and mental development in infants and toddlers. *Pediatr. Obes.* **11**, 389–396 (2016).
47. Ruiz, M. *et al.* Impact of low maternal education on early childhood overweight and obesity in Europe. *Paediatr. Perinat. Epidemiol.* **30**, 274–284 (2016).
48. Savchev, S. *et al.* Neurodevelopmental outcome of full-term small-for-gestational-age infants with normal placental function. *Ultrasound Obstet. Gynecol.* **42**, 201–206 (2013).

49. Taine, M. *et al.* Early postnatal growth and neurodevelopment in children born moderately preterm or small for gestational age at term: A systematic review. *Paediatr. Perinat. Epidemiol.* **32**, 268–280 (2018).
50. Takeuchi, A. *et al.* Catch-up growth and neurobehavioral development among full-term, small-for-gestational-age children: A nationwide Japanese population-based study. *J. Pediatr.* **192**, 41–46 (2018).
51. Cho, W. K. & Suh, B. K. Catch-up growth and catch-up fat in children born small for gestational age. *Korean J. Pediatr.* **59**, 1–7 (2016).
52. Ozonoff, S. *et al.* Onset patterns in autism: Correspondence between home video and parent report. *J. Am. Acad. Child Adolesc. Psychiatry* **50**, 796–806 (2011).

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## Author contributions

T.K., T.N., and N.T. conceptualized and designed the study, carried out the analyses, drafted the initial manuscript, and revised the manuscript. N.T., Y.N., and A.S. critically reviewed the manuscript and supervised. K.J.T. coordinated and supervised data collection. Material preparation and data collection were performed by T.N., T.H., A.O., T.I., and K.J.T. All the authors critically reviewed and approved the final version of the manuscript.

## Competing interests

The authors declare no competing interests.

## Additional information

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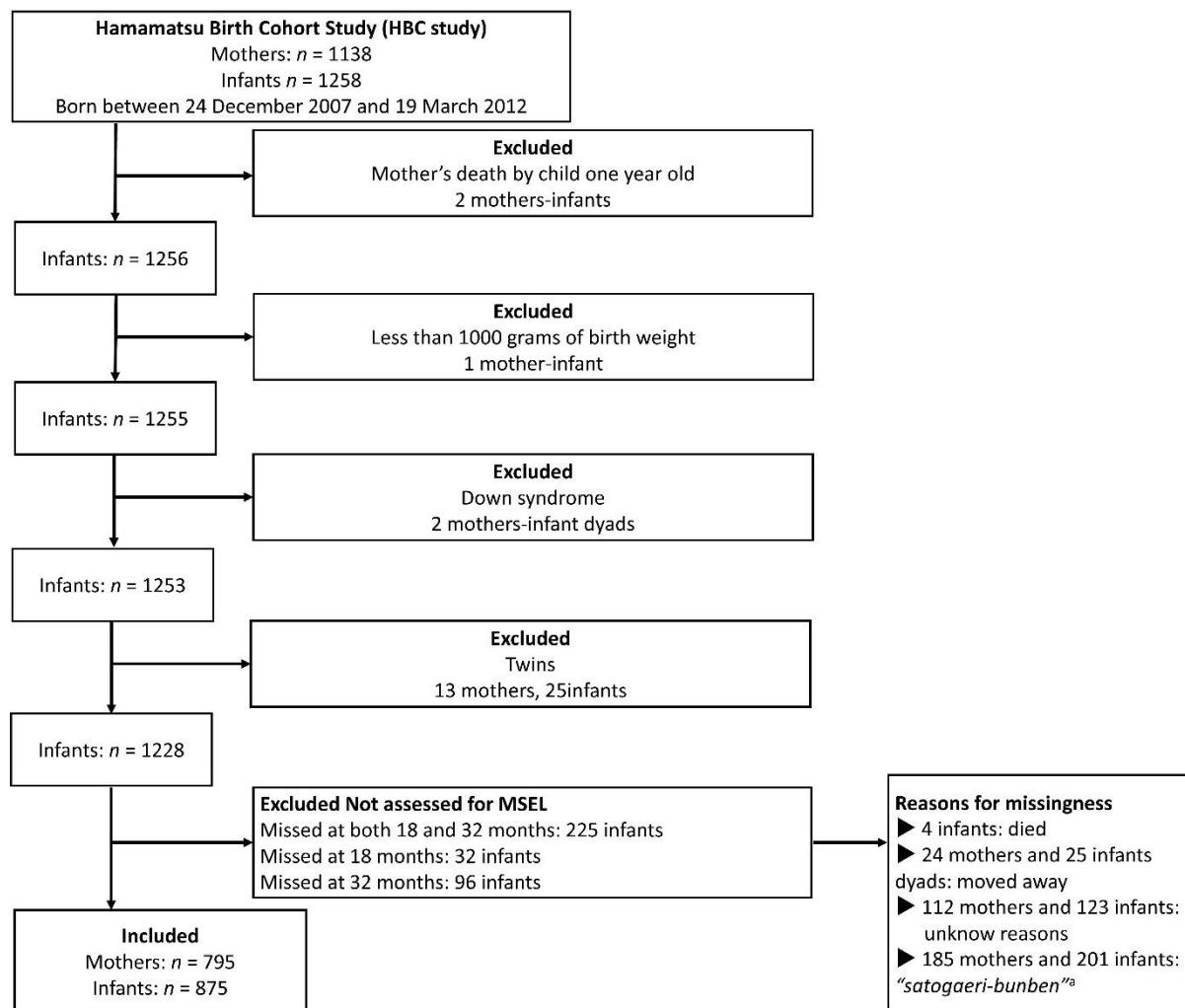
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**Online supplementary information for the “Identification of neurodevelopmental transition patterns from infancy to early childhood, and risk factors predicting descending transition**

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**Supplementary Figure S1. Participants flow chart**

<sup>a</sup>“Satogaeri-bunben” is a Japanese traditional support system for childbirth, in which some pregnant women return to their hometown for delivery while receiving family support.

MSEL, Mullen Scales of Early Learning.

**Supplementary Table S1. Fit indices in latent class analysis**

	Number of class						
	1	2	3	4	5	6	7
<b>Fit stats at time 1</b> (n = 875)							
AIC	32873.9	32233.1	31981.6	31882.7	31802.6	31737.5	31714.6
BIC	32921.9	32309.9	32087.3	32017.2	31965.9	31929.5	31935.5
Entropy	-	0.856	0.786	0.739	0.786	0.807	0.806
Adjusted LMR-LRT p value	-	< 0.001	0.067	0.132	0.029	0.167	0.351
BLRT p value	-	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001
<b>Fit stats at time 2</b> (n = 875)							
AIC	32943.5	32079.2	31680.0	31577.7	31538.6	31489.7	31478.4
BIC	32991.5	32156.0	31785.6	31712.2	31701.9	31681.8	31699.4
Entropy	-	0.705	0.799	0.758	0.753	0.706	0.722
Adjusted LMR-LRT p value	-	< 0.001	< 0.001	0.003	0.037	0.054	0.353
BLRT p value	-	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001

AIC, Akaike Information Criterion; BIC, Bayesian Information Criterion; BLRT, bootstrap likelihood ratio test; LMR-LRT, Lo-Mendell Rubin likelihood ratio test.

### Supplementary Note 1. Details of the latent transition analysis in the present study

Latent transition analysis (LTA) enables us to identify the transition of classes assigned at different time-points and investigate the association of predictors (i.e., risk factors) with class shifts. The primary aim of LTA is to compute latent transition probabilities, which represent the likelihood that individuals change classes or remain in the same class across consecutive periods<sup>1</sup>. In the present study, we followed the three-step approach recommended for LTA<sup>2,3</sup>. In step 1, the LTA model without risk factors or covariates was estimated; in step 2, individuals were assigned to optimal latent transition patterns; in step 3, a model with risk factors or covariates was fit to detect relevant risk factors. We used this approach to identify transition class assignments, as well as to investigate risk factors for predicting transition patterns using Mplus, version 8<sup>4</sup>. However, during step 3, we failed to obtain accurate and reliable estimates due to the small number of assignment memberships in many of the created cells. In effect, fewer than nine children (less than 1%) were allocated to nearly half of the total 25 cells (12/25), which was created by  $5 \times 5$  classes at time 1 and time 2 (see Table 2). Nine out of 12 belonged to descending or catching-up transition shifts, which were of interest in this study. Therefore, we modified the original step 3<sup>3</sup> and implemented the following process to enhance the precision of estimation and lead to clinically meaningful interpretations. First, we transferred the whole data on the individual latent class assignments at each time-point, transition assignments, and transition probabilities derived from Mplus to Stata software<sup>5</sup>. Then, we combined the three normal classes (i.e., High Normal, Normal, and Low Normal; see Table 2 and Fig. 1 in the main text) at each time-point into a single “3-Normals” class, resulting in reduction from five to three classes in total for both time 1 and time 2. Thus, the number of transition patterns was reduced from 25 to 9 ( $3 \times 3$ ) patterns. Second, we examined any differences in background characteristics across these three classes at time 1 to identify potential confounders (covariates) for subsequent analyses. Through the screening analytical steps, distributions of various factors such as sex, premature birth (<37 weeks), low placenta-to-birth weight ratio (<10<sup>th</sup> percentile), and household income were found to significantly differ across the three classes; thus, they were included as covariates in the analyses. Third, we used logistic regression analysis with the forward stepwise method in which the minimum number of risk factors associated with transition patterns was identified following the principle of parsimony, with covariates being compulsorily held to allow for their confounding effects. Notably, in the original three-step approach<sup>3</sup>, risk factors and covariates are entered compulsorily, thereby rendering the derived estimates unreliable due to the inflated number of parameters. In this analysis, we regarded all the variables listed in Table 1 as risk factors, except for the four covariates. Here, we used the maximum class probability from LTA (converted into odds ratio) to account for variability in the likelihood of transition pattern assignments. Since two or more children born from the same mother were enrolled, such clustering was accounted for by using robust variance estimation. The results showed that maternal pre-pregnancy body mass index ( $>25 \text{ kg/m}^2$ ), maternal education, and infant's body mass index at 18 months ( $>1$  standard deviation) were associated with descending transition patterns (see Table 3).

**Supplementary Table S2. Association between covariates and neurodevelopmental transition patterns in the final model of multinomial logistic regression analysis ( $n = 811$ ).**

<b>Transition from 3-Normals at time 1 (765)</b>			
<b>Covariates</b>	to 3-Normals at time 2 (681)	Delayed at time 2 (70)	Markedly Delayed at time 2 (14)
	Reference	OR [95% CI]	OR [95% CI]
<b>Male sex</b>			
Male sex	..	1.51 [0.88, 2.60]	6.13 [1.44, 26.01]*
<b>Premature birth before 37 weeks</b>			
Premature birth before 37 weeks	..	1.08 [0.33, 3.50]	5.12 [1.02, 25.63]*
<b>Low placenta-to-birth-weight ratio (&lt;10<sup>th</sup> percentile)</b>			
Low placenta-to-birth-weight ratio (<10 <sup>th</sup> percentile)	..	1.48 [0.76, 2.87]	NA
<b>Household income (million JPY)</b>			
Household income (million JPY)	..	0.89 [0.78, 1.01]	0.93 [0.70, 1.23]
<b>Transition from expressive language delayed at time 1 (19)</b>			
<b>Covariates</b>	to 3-Normals at time 2 (15)	Delayed at time 2 (3)	Markedly Delayed at time 2 (1)
	OR [95% CI]	OR [95% CI]	OR [95% CI]
<b>Male sex</b>			
Male sex	1.13 [0.40, 3.20]	NA	NA
<b>Premature birth before 37 weeks</b>			
Premature birth before 37 weeks	NA	0.90 [0.69, 1.18]	NA
<b>Low placenta-to-birthweight ratio (&lt;10th percentile)</b>			
Low placenta-to-birthweight ratio (<10th percentile)	NA	9.06 [0.54, 152.11]	NA
<b>Household income (million JPY)</b>			
Household income (million JPY)	0.98 [0.81, 1.19]	0.90 [0.69, 1.18]	1.11 [1.01, 1.21]*
<b>Transition from Delayed at time 1 (27)</b>			
<b>Covariates</b>	to 3-Normals at time 2 (9)	Delayed at time 2 (10)	Markedly Delayed at time 2 (8)
	OR [95%CI]	OR [95%CI]	OR [95%CI]
<b>Male sex</b>			
Male sex	2.73 [0.62, 11.96]	1.69 [0.42, 6.68]	NA
<b>Premature birth before 37 weeks</b>			
Premature birth before 37 weeks	2.40 [0.41, 13.82]	3.18 [0.27, 37.10]	NA
<b>Low placenta-to-birthweight ratio (&lt;10th percentile)<sup>a</sup></b>			
Low placenta-to-birthweight ratio (<10th percentile) <sup>a</sup>	1.38 [0.20, 9.39]	1.86 [0.39, 8.80]	2.70 [0.59, 12.31]
<b>Household income (million JPY)</b>			
Household income (million JPY)	0.61 [0.47, 0.79]****	0.73 [0.51, 1.03]	0.99 [0.77, 1.26]

\* $P < .05$ . \*\* $P < .01$ . \*\*\* $P < .005$ . \*\*\*\* $P < .001$ .

OR, odds ratio; CI, confidence interval; JPY, Japanese Yen; NA, not available.

**Supplementary Note 2. Results and inferences of covariates for neurodevelopmental transition patterns.**

The variables of male sex (OR = 6.13; 95% CI: 1.44, 26.01) and premature birth before 37 weeks (OR = 5.12; 95% CI: 1.02, 25.63) were significantly associated with the “3-Normals to M-Delayed” descending transition pattern. These variables were treated as covariates to adjust for background characteristic differences across three classes at time 1 so that they could not be considered as risk factors. Nevertheless, given that any existing confounding and risk-increasing effects in a single variable cannot be separated, there is the possibility that these two factors may in effect have contributed to the descending transition. Compatible with this inference, a previous study has reported that male sex and prematurity were associated with a diversion from normal to delayed development in the neurocognitive domain during age intervals similar to our study<sup>6</sup>.

Household income used as a covariate was negatively associated with the catching-up transition pattern from Delayed to 3-Normals; infants from *low* income were *more* likely to make the catching-up transition from the Delayed class at time 1 to the 3-Normals class at time 2, with the odds ratio (OR) being 0.61 (95% CI: 0.47, 0.79) when compared with infants who transitioned from 3-Normals at time 1 to 3-Normals at time 2. Given that higher income has been reported to be positively related to child development<sup>7</sup>, our contradictory finding is puzzling. One possible interpretation is an involvement of public health care professionals in supporting the development of infants with socioeconomically deprived backgrounds<sup>7,8</sup>. Those infants from low socioeconomic status identified as the Delayed class at time 1 in this study may have naturally raised more attention from health care providers (e.g., through nationwide infant check-ups)<sup>9</sup> and prompted intervention to foster their development.

**Supplementary Table S3. Comparison of participants' characteristics between the group included in the analysis and the group excluded from the analysis**

	<b>Included (n = 875)</b>	<b>Excluded (n = 383)</b>	<b>P value<sup>a</sup></b>
	<b>Mean (SD)</b>	<b>Mean (SD)</b>	
<b>Infants</b>			
Birth weight (g)	2957.0 (422.2)	2902.4 (472.8)	0.042
Gestational age at birth (weeks)	39.0 (1.4)	38.6 (1.8)	< 0.0001
	<b>n (%)</b>	<b>n (%)</b>	<b>P value<sup>a</sup></b>
Sex			
Male	432 (49.4%)	216 (56.4%)	0.022
Female	443 (50.6%)	167 (43.6%)	
Prematurity			
<37 weeks	46 (5.3%)	40 (10.4%)	0.0011
≥37 weeks	829 (94.7%)	343 (89.6%)	
Small for gestational age			
<10th percentile	77 (8.8%)	29 (7.6%)	0.47
10th–100th percentile	798 (91.2%)	354 (92.4%)	
Age-standardised BMI scores at 18 months			
>1 standard deviation	50 (6.1%)	3 (2.5%)	0.11
<1 standard deviation	776 (93.9%)	116 (97.5%)	
	<b>n (%)</b>	<b>n (%)</b>	<b>P value<sup>a</sup></b>
<b>Parents</b>			
Maternal BMI at pre-pregnancy (kg/m <sup>2</sup> )			
>25.0	100 (11.4%)	30 (7.8%)	0.054
<25.0	775 (88.6%)	353 (92.2%)	
Placenta-to-birth-weight ratio (twin excluded)			
<10 <sup>th</sup> percentile	152 (17.7%)	65 (19.5%)	0.46
10 <sup>th</sup> –100 <sup>th</sup> percentile	707 (82.3%)	268 (80.5%)	
	<b>Mean (SD)</b>	<b>Mean (SD)</b>	<b>P value<sup>a</sup></b>
Paternal age at birth (yr)	33.5 (5.7)	32.5 (6.0)	0.0065
Maternal age at birth (yr)	31.7 (5.0)	30.7 (5.1)	0.0005
Paternal education (yr)	14.1 (2.6)	13.8 (2.8)	0.059
Maternal education (yr)	13.8 (1.8)	13.6 (2.2)	0.028
Household income (million JPY)	6.0 (2.7)	5.9 (2.9)	0.41

SD, standard deviation; BMI, body mass index; JPY, Japanese Yen.

<sup>a</sup> *t*-test for continuous variables and the chi-square test for categorical variables were used.

**Supplementary Table S4. Association between possible risk factors and neurodevelopmental transition patterns in the final model of multinomial logistic regression after excluding infants with a clinical diagnosis of autism spectrum disorder (n = 26)**

Transition from 3-Normals at time 1 (746)			
	to 3-Normals at time 2 (673)	Delayed at time 2 (62)	Markedly Delayed at time 2 (11)
Possible risk factors	Reference	OR [95%CI]	OR [95%CI]
<b>Possible risk factors</b>			
Maternal overweight status at pre-pregnancy	..	2.61 [1.25, 5.43]**	NA
Low maternal education (yr)	..	1.23 [1.06, 1.42]	1.08 [0.78, 1.49]
Male sex <sup>a</sup>	..	1.41 [0.79, 2.51]	10.96 [2.47, 48.51]***
Small for gestational age	..	2.07 [0.82, 5.22]	0.68 [0.056, 8.50]
Premature birth before 37 weeks <sup>a</sup>	..	1.16 [0.36, 3.81]	8.69 [1.74, 43.39]**
Infant overweight status at 18 months	..	1.65 [0.62, 4.34]	8.82 [1.88, 41.36]**
Low placenta-to-birthweight ratio (<10th percentile) <sup>a</sup>	..	1.52 [0.74, 3.12]	NA
Household income (million JPY) <sup>a</sup>	..	0.90 [0.79, 1.01]	0.91 [0.66, 1.26]
Transition from Expressive Language Delayed at time 1 (17)			
	to 3-Normals at time 2 (15)	Delayed at time 2 (2)	Markedly Delayed at time 2 (0)
Possible risk factors	OR [95%CI]	OR [95%CI]	OR [95%CI]
Maternal overweight status at pre-pregnancy	0.90 [0.11, 7.31]	NA	NA
Low maternal education (yr)	1.23 [1.01, 1.49]*	3.12 [0.59, 16.66]	NA
Male sex <sup>a</sup>	1.17 [0.41, 3.33]	NA	NA
Small for gestational age	0.92 [0.11, 7.45]	33.23 [0.27, 3845.80]	NA
Premature birth before 37 weeks <sup>a</sup>	NA	140.01 [0.0094, 20652.31]	NA
Infant overweight status at 18 months	NA	NA	NA

Low placenta-to-birthweight ratio (<10th percentile) <sup>a</sup>	2.90 [0.94, 8.94]	NA	NA
Household income (million JPY) <sup>a</sup>	0.99 [0.82, 8.94]	1.16 [0.92, 1.46]	NA
<b>Transition from Delayed at time 1 (22)</b>			
	to 3-Normals at time 2 (8)	Delayed at time 2 (10)	Markedly Delayed at time 2 (4)
	OR [95%CI]	OR [95%CI]	OR [95%CI]
<b>Possible risk factors</b>			
Maternal overweight status at pre-pregnancy	5.68 [1.08, 29.77]*	0.87 [0.088, 8.52]	NA
Low maternal education (yr)	1.05 [0.68, 1.63]	1.28 [0.84, 1.92]	2.01 [1.08, 3.70]*
Male sex <sup>a</sup>	2.21 [0.48, 10.02]	1.74 [0.43, 6.93]	NA
Small for gestational age	11.86 [2.42, 58.12]***	NA	NA
Premature birth before 37 weeks <sup>a</sup>	2.65 [0.44, 15.75]	3.15 [0.26, 37.25]	NA
Infant overweight status at 18 months	3.19 [0.24, 40.98]	2.10 [0.23, 18.94]	NA
Low placenta-to-birthweight ratio (<10th percentile) <sup>a</sup>	1.58 [0.22, 11.17]	1.88 [0.40, 8.77]	2.96 [0.38, 22.89]
Household income (million JPY) <sup>a</sup>	0.66 [0.54, 0.81]****	0.74 [0.53, 1.04]	1.11 [0.86, 1.43]

OR, odds ratio; CI, confidence interval; JPY, Japanese Yen; NA, not available.

<sup>a</sup> included as covariates.

\* $P < .05$ . \*\* $P < .01$ . \*\*\* $P < .005$ . \*\*\*\* $P < .001$ .

**Supplementary Table S5. Association between neurodevelopmental transition patterns and adaptive behaviour at 40 months in linear regression analysis<sup>a</sup> after excluding infants with a clinical diagnosis of autism spectrum disorder (n=26)**

	Adaptive Behaviour Composite standardised score coefficient (95%CI)
<b>Transition patterns (n)</b>	
3-Normals to 3-Normals (646)	reference
3-Normals to Delayed (59)	-7.52 [-9.41, -5.62]****
3-Normals to Markedly Delayed (10)	-12.05 [-18.71, -5.40]****
EL-Delayed to 3-Normals (14)	-2.44 [-5.24, 0.34]
Delayed to 3-Normals (8)	0.47 [-4.44, 5.39]
EL-Delayed to Delayed (2)	-15.24 [-17.37, -13.11]****
EL-Delayed to Markedly Delayed (0)	..
Delayed to Delayed (10)	-9.85 [-13.47, -6.23]****
Delayed to Markedly Delayed (4)	-18.61 [-24.01, -13.22]****

EL-Delayed, Expressive Language Delayed.

<sup>a</sup> Covariates used in this analysis comprised infant's sex (male), premature birth (<37 weeks), low placenta-to-birthweight ratio (<10<sup>th</sup> percentile), household income, maternal body mass index at pre-pregnancy (>25 kg/m<sup>2</sup>), small for gestational age (<10<sup>th</sup> percentile), infant's standardised body mass index at 18 months of age (>1 SD), and maternal educational history.

\*\*\*\*P<.001.

## References

- 1 Collins, L. M. & Lanza, S. T. *Latent class and latent transition analysis: With applications in the social, behavioral, and health sciences*. Vol. 718 (John Wiley & Sons, 2010).
- 2 Muthén, B. & Muthén, L. K. Integrating person-centered and variable-centered analyses: growth mixture modeling with latent trajectory classes. *Alcohol. Clin. Exp. Res.* **24**, 882-891 (2000).
- 3 Nylund-Gibson, K., Grimm, R., Quirk, M. & Furlong, M. A latent transition mixture model using the three-step specification. *Struct. Equ. Modeling.* **21**, 439-454 (2014).
- 4 Muthén, L. K. & Muthén, B. O. 1998–2012. *Mplus user's guide*. (Muthén & Muthén, 2012).
- 5 StataCorp, L. *Stata treatment-effects reference manual*. (A Stata Press Publication, 2015).
- 6 Hillemeier, M. M., Morgan, P. L., Farkas, G. & Maczuga, S. A. Perinatal and socioeconomic risk factors for variable and persistent cognitive delay at 24 and 48 months of age in a national sample. *Maternal and Child Health Journal* **15**, 1001-1010 (2011).
- 7 Demirci, A. & Kartal, M. Sociocultural risk factors for developmental delay in children aged 3–60 months: a nested case-control study. *Eur. J. Pediatr.* **177**, 691-697 (2018).
- 8 Ozkan, M., Senel, S., Arslan, E. A. & Karacan, C. D. The socioeconomic and biological risk factors for developmental delay in early childhood. *Eur. J. Pediatr.* **171**, 1815-1821 (2012).
- 9 Kamio, Y. *et al.* Brief Report: Best Discriminators for Identifying Children with Autism Spectrum Disorder at an 18-Month Health Check-Up in Japan. *J. Autism Dev. Disord.* **45**, 4147-4153 (2015).