# **BMJ Open** Cognitive development of children with Kawasaki disease and the parenting stress of their caregivers in Taiwan: a case-control study

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#### ABSTRACT

**Objective** Kawasaki disease (KD) is an acute form of febrile vasculitis that occurs in early childhood. The multisystemic vasculitis common in patients with KD may influence blood perfusion in the brain, and thus caregivers of children with KD may feel stress with regard to caring for them. Intravenous immunoglobulin (IVIG) infusion is the standard treatment for acute KD, and the most serious complication of KD is coronary artery aneurysms (coronary artery lesion (CAL)). This study aimed to investigate the relationships between KD heterogeneity and the risk of patients' cognitive impairment or caregivers' parenting stress.

**Design** A case–control study with consecutive sampling. **Setting** A medical centre (Kaohsiung Chang Gung Memorial Hospital, Taiwan).

**Participants** This study consisted of 176 patients with KD (mean age: 5.5 years, 60.8% boys) and 85 healthy children (mean age: 6.4 years, 54.1% boys).

**Primary and secondary outcome measures** Based on the children's age, each patient with KD and control subject was administered an assessment using the Mullen Scales of Early Learning or the Wechsler Intelligence Scale, and parenting function of their caregivers was assessed using the Parenting Stress Index (PSI)-Short Form.

**Results** We observed no significant differences in any developmental index, cognitive function or parenting stress between patients with KD and controls. Among the children with KD, IVIG administration nor CAL was associated with children's cognitive scores. However, the caregivers of patients who had CAL suffered from greater PSI total scores than those of patients without CAL. Furthermore, the caregivers who had education levels of a master's degree or above showed less parenting stress than those who had an education level of college or lower. **Conclusion** Caregivers' education is associated to parenting stress, and caregivers of patients with KD who developed CAL may feel stress about the unpredictable sequela caused by CAL for their children. Such caregivers may require support to fulfil their parenting roles.

#### INTRODUCTION

Kawasaki disease (KD) manifests with acute fever, mainly affects children aged <5 years, and involves multisystemic vasculitis of unknown aetiology.<sup>1</sup><sup>2</sup> Although it occurs

## Strengths and limitations of this study

- We carried out a comprehensive study about cognitive impairment of patients with Kawasaki disease (KD) or parenting stress of caregivers.
- This is the first study indicating that caregivers of patients with KD who developed coronary artery aneurysms may feel stress about the unpredictable sequela.
- Owing to small case numbers, it is difficult to make a strong conclusion about cognitive impairment.
- We used a case-control design; therefore, no causal relationships could be identified.

around the world, KD incidence rates are especially high in East Asia, particularly Japan, Korea and Taiwan.<sup>3 4</sup> The primary clinical characteristics of KD consist of prolonged fever, diffuse mucosal inflammation, bilateral non-purulent conjunctivitis, non-suppurative cervical lymphadenopathy, indurative angioedema of the hands and feet and polymorphous skin rashes.<sup>5-7</sup> Intravenous immunoglobulin (IVIG) infusion has been established as the standard treatment for acute KD,<sup>8</sup> but 10%–20% of patients still show resistance to IVIG therapy and are at high risk for coronary artery complications,<sup>9</sup> of which, the formation of coronary artery lesions (CALs) is the most serious.<sup>10</sup> These studies suggest that 20%-24% of children with KD, even when treated with IVIG, still suffer from CAL and 4% with coronary artery aneurysm formation.<sup>1112</sup>

In addition to the harm caused to the coronary arteries, KD is also characterised by multisystemic vasculitis and may thus affect blood perfusion and cause inflammatory changes in the brain.<sup>13 14</sup> Central nervous system symptoms, such as lethargy, cranial nerve palsy and prolonged partial seizures, occur in 1%–30% of patients with KD.<sup>15–18</sup> Fortunately, these studies have consistently demonstrated that

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Dr Ho-Chang Kuo; erickuo48@yahoo.com.tw patients with KD are not associated with cognitive impairment sequelae.<sup>19–21</sup> However, different responses to IVIG treatment and the development of coronary aneurysms may be related to a discrepancy in patients' immuno-logical profiles or genetic background.<sup>22</sup> Therefore, the relationships between heterogeneity of KD and the risk of cognitive impairments warrant further investigation.

Our study revealed that children with KD suffered acute and prominent impairment in health-related quality of life, and the impairments even exceeded that of children newly diagnosed with cancer.<sup>23</sup> Furthermore, patients with KD, as well as their caregivers, may consistently worry about patients' potential risk of cardiac event-related death.<sup>24</sup> With a lack of support and adequate knowledge available at the societal level, patients' caregivers may experience stress from coping with their children's needs and fulfilling their parenting roles. Owing to the uncertainty of the long-term KD prognosis, the parents of children with coronary artery complications suffered from persistent anxiety even years after the acute phase of the illness.<sup>25</sup> The psychosocial burden of parents is associated with intensity of medical experience and family's psychosocial limitations.<sup>26</sup> However, the evidence with regarding to parenting stress among caregivers of children with KD is still scarce. Therefore, recruiting a control group is helpful to compare the cognitive profiles and parenting stress between children with KD and those without KD.<sup>27</sup>

To fill this research gap, we performed a clinical survey to explore the potential factors (eg, responses to IVIG treatment or comorbid CAL) associated with cognitive profiles in patients with KD. In addition, we investigated whether the caregivers of children with KD and healthy controls exhibited different parenting stress, and to examine the factors correlated to caregivers' parenting stress among children with KD.

## METHODS

## **Participants**

We recruited a total of 176 patients with KD from the Department of Pediatrics from June 2016 to July 2018, Kaohsiung Chang Gung Memorial Hospital, Taiwan or communities near the hospital. A senior clinician diagnosed patients with KD in accordance with the recommended universal KD criteria published by the American Heart Association.<sup>28</sup> The diagnostic criteria of KD includes fever that lasts longer than 5 days, as well as four of the following five symptoms: diffuse mucosal inflammation with strawberry tongue and fissure lips, bilateral non-purulent conjunctivitis, indurative angioedema of the hands and feet, dysmorphic skin rashes and unilateral cervical lymphadenopathy, as stated in our previous reports.<sup>29</sup> Once their symptoms were remitted, patients with KD were administered a developmental or cognitive assessment in either outpatient department or in the ward.

Because most of our patients with KD were recruited from the outpatient department, we recruited 85 healthy children from communities around Kaohsiung Chang Gung Memorial Hospital or children suffering from upper respiratory tract infection whose symptoms were currently in remission as a control group. We excluded any patients with other immunological diseases (asthma, allergic rhinitis, atopic dermatitis or allergic conjunctivitis) or major physical illnesses (such as genetic, metabolic or infectious conditions).

#### **CAL** assessment

All participants were provided with a structured questionnaire to collect demographic data, such as age, gender and age of onset. We recorded body temperature every 6 hours during the febrile stage. The CAL was defined as a luminal diameter of >3 mm in a child aged <5 years or >4 mm in those aged  $\geq$ 5 years, when the internal diameter of a segment is 1.5 times or greater than that of an adjacent segment, or when the luminal contour is clearly irregular or has a Z score>2.5 SD.<sup>30 31</sup> We estimated the Z score of the proximal right coronary artery, left main coronary artery and proximal left anterior descending artery, as well as the maximum Z score of coronary arteries both at baseline and 6-8 weeks with 2D echocardiography. The body weight and height used to calculate Z scores were obtained from the Taiwan Society of Pediatric Cardiology website (http://www.tspc.org.tw/service/z\_score.asp). The IVIG resistance was defined as persistent or recrudescent fever for at least 36 hours after the end of their IVIG infusion but not >7 days.<sup>32</sup>

## Neurocognitive assessments

Each patient with KD and control subject were administered a developmental or cognitive assessment performed by an experienced child psychologist in a room designed to reduce testing condition variables. Patients aged <4 years were assessed using the Mullen Scales of Early Learning (MSEL); patients between the ages of 4 and 7 years were examined using the Wechsler Preschool and Primary Scale of Intelligence-Fourth Edition (WPPSI-IV) and patients aged >7 years were tested using the Wechsler Intelligence Scale for Children-Fourth Edition (WISC-IV).

The MSEL, a standardised developmental assessment for newborns through children of age 68 months, offers an overall index of cognitive ability and developmental delay.<sup>33</sup> It consists of five subscales: Gross Motor (only for children aged <33 months), Visual Reception, Fine Motor, Expressive Language and Receptive Language. After scoring each item, the crude scores of each of the five scales are transformed into normalised age-specific scores, referred to as T scores. The T scores of all except the Gross Motor Scale are added together and converted into the Early Learning Composite (ELC) score. The MSEL has been applied in Taiwan's toddler samples.<sup>21 34</sup>

The WPPSI-IV is an individually administered, standardised tool for evaluating intelligence in children between the ages of 2 years 6 months and 7 years 7 months.<sup>35</sup> The test takes about 30–45 min for children between the ages of 2 years 6 months and 3 years 11 months and provides a Full-Scale Intelligence Quotient (FSIQ), Verbal Comprehension Index (VCI), Visual Spatial Index (VSI) and Working Memory Index (WMI). For children between the ages of 4 years and 7 years 7 months, it takes about 45–60 min and provides a FSIQ, VCI, VSI, Fluid Reasoning Index, WMI and Processing Speed Index (PSI).<sup>36</sup> The WPPSI-IV has been used in clinical study in Taiwan.<sup>21 37</sup>

The WISC-IV is an individually administered and normreferenced tool developed to measure the intelligence of children aged from 6 to 16 years.<sup>38</sup> The WISC-IV includes 10 core and 5 supplemental subtests. The core subtests create four factor indexes, including the VCI, the Perceptual Reasoning Index, the WMI and the PSI. Meanwhile, the 10 core subtests form the FSIQ. Each factor index, as well as the FSIQ, has a population mean of 100 and an SD of 15. The WISC-IV has been applied for Taiwanese school-aged children in several clinical studies.<sup>39 40</sup>

## **Parenting stress**

Caregivers' characteristics, including age, gender and education levels, were recorded, and their parenting function was assessed using the Parenting Stress Index-Short Form (PSI-SF). The PSI-SF, a widely adopted self-administered questionnaire for evaluating parenting stress, includes 36 items (rated on a 5-point Likert scale) and stems directly from the full-length 120-item Parenting Stress Index test.<sup>41</sup> The PSI-SF provides scores in the following subscales: (a) Parental Distress, (b) Parent-Child Dysfunctional Interaction and (c) Difficult Child. The three subscores are added together to yield a total parenting stress score.<sup>42</sup> Studies have reported that the Chinese version of the PSI-SF is a reliable assessment tool for identifying parenting stress with a need for intervention in clinical practices.<sup>43-45</sup>

#### **Statistical analysis**

All data processing and statistical analyses were performed using the SPSS software, V.14.0. Two-tailed p values <0.05 were considered statistically significant.

The sample size calculation was estimated using the G\*Power software. Based on the settings of 80% power, p=0.05, effect size=0.4 and allocation ratio N1/N2=2, the sample size should be 149 in group 1 and 75 in group 2. The common assumptions made when performing a t test include those regarding the scale of measurement, random sampling, normality of data distribution, adequacy of sample size and equality of variance in SD. The assumptions of the  $\chi^2$  include the data in the cells should be frequencies or counts of cases rather than percentages or some other transformation of the data. The levels (or categories) of the variables are mutually exclusive. Multiple linear regression analysis makes several key assumptions as following: linear relationship, multivariate normality, no or little multicollinearity, no auto-correlation and homoscedasticity.<sup>46</sup> The literature with regarding to cognitive function and parenting stress among children with KD is still scarce; therefore,

the covariates selected for the multiple linear regression model were based on the variables collected in clinical settings.

We used the  $\chi^2$  test to compare differences in categorical variables between patients with KD and those without KD. An independent t-test was adopted to compare continuous variables between the two groups. We performed a general linear model to observe the difference in children's cognition and caregivers' stress between patients with KD and controls, controlled for children's age and caregivers' education levels. The cognitive scores of patients who were assessed using the MSEL were set as the ELC scores, and the cognitive scores of those assessed with the WPPSI or WISC-IV were set as the FSIQ. The independent variables were the characteristics of the children and caregivers. Furthermore, multiple linear regression was adopted to determine the factors associated with parental stress among the patients with KD. The dependent variable was set as the total PSI score, and the independent variables were the characteristics of the children and caregivers.

#### RESULTS

The study participants consisted of 176 patients with KD (mean age: 5.5 years, 60.8% boys) and 85 healthy children (mean age: 6.4 years, 54.1% boys; table 1). The healthy children were older than patients with KD (p=0.035), and the caregivers in the control group had higher education levels than those in the KD group (p=0.018). Apart from this, no significant differences in gender or caregivers' characteristics between children with KD and controls. No significant differences in development scores measured using the MSEL or cognitive scores measured using the WPPSI or WISC-IV were observed between the children with KD and the healthy controls. When comparing the caregivers of patients with KD to those of the controls, none of the PSI subscales showed a significant difference.

Of the children with KD, the mean age of KD onset was 21.6 months; 4% of them never received IVIG treatment, and 88.6% and 7.4% received IVIG treatment once and twice, respectively; 58% of them had no CAL and 42% of them had CAL. Table 2 shows the relationship between children's and caregivers' characteristics and the cognitive score among children with KD. We found that neither IVIG administration nor CAL was associated with the cognitive score of children with KD. The characteristics of children and caregivers were also not correlated with cognitive score.

Regarding the factors related to the total scores of PSI (table 3), caregivers of patients wth KD who had CAL suffered from greater parenting stress than the caregivers of patients without CAL (p=0.019). Furthermore, the caregivers who had education levels of a master's degree or above showed less parenting stress than those who had an education level of college (p=0.010) or lower (p=0.021).

Table 1       Characteristics of children with h	CD and healthy controls			
	KD (n=176)	Controls (n=85)	Statistic†	P value
Children's characteristics				
Age (months)	66.5±39.0	77.3±38.0	-2.114	0.035*
Age of KD onset (months)	21.6±17.7	-	N/A	N/A
Sex			1.054	0.305
Male	107 (60.8)	46 (54.1)		
Female	69 (39.2)	39 (45.9)		
Intravenous immunoglobulin			N/A	N/A
None	7 (4.0)	-		
Once	156 (88.6)	-		
Twice	13 (7.4)	-		
Coronary artery aneurysms			N/A	N/A
With	74 (42.0)	-		
Without	102 (58.0)	-		
Mullen Scales of Early Learning				
Gross Motor‡	52.0±13.1	52.7±12.0	0.091	0.765
Visual Reception	54.0±13.8	53.1±17.2	0.062	0.804
Fine Motor	50.2±13.4	50.1±11.6	0.037	0.847
Receptive Language	57.0±11.5	51.2±10.7	3.838	0.054
Expressive Language	51.7±13.2	50.5±11.2	0.229	0.634
Composite scores	106.2±19.4	105.0±19.2	0.087	0.768
WPPSI				
Full Scale IQ	103.7±10.4	107.6±11.7	1.380	0.243
Verbal Comprehension Index	105.2±10.7	107.1±14.3	1.132	0.290
Visual Spatial Index	98.8±14.9	101.5±10.9	0.262	0.610
Fluid Reasoning Index	107.0±12.0	107.0±13.9	0.003	0.960
Working Memory Index	99.0±12.4	100.3±15.3	0.150	0.700
Processing Speed Index	103.3±12.5	106.9±12.7	0.683	0.411
WISC-IV				
Full Scale IQ	109.1±11.1	109.7±17.4	1.039	0.311
Verbal Comprehension Index	108.2±13.4	111.3±16.3	0.022	0.882
Perceptual Reasoning Index	106.1±14.8	106.4±17.2	0.576	0.450
Working Memory Index	110.3±13.1	111.4±15.3	0.331	0.567
Processing Speed Index	103.0±14.3	98.4±15.1	3.928	0.051
Caregivers' characteristics				
Age (years)	37.6±4.8	38.2±5.0	-0.889	0.375
Sex			1.096	0.295
Male	26 (14.9)	17 (20.0)		
Female	149 (85.1)	68 (80.0)		
Education levels			8.080	0.018*
High school or lower	81 (46.8)	29 (34.1)		
College	77 (44.5)	39 (45.9)		
Master or above	15 (8.7)	17 (20.0)		
Family expenditure (per month)§			0.515	0.773
<50 000 NT\$	94 (59.9)	42 (55.3)		
50000-100000 NT\$	46 (29.3)	24 (31.6)		

Continued

Table 1 Continued				
	KD (n=176)	Controls (n=85)	Statistic†	P value
>100000 NT\$	17 (10.8)	10 (13.2)		
PSI-SF				
Defensive Responding	18.0±5.2	18.0±5.3	0.193	0.660
Parental Distress	30.2±8.2	30.0±9.0	0.003	0.958
Parent-Child Dysfunctional Interaction	23.1±7.0	24.4±8.3	2.204	0.139
Difficult Child	27.4±8.1	29.0±8.6	3.836	0.051
Total score	80.7±20.4	83.1±21.9	1.749	0.187

Data are expressed as mean±SD or n (%).

\*p<0.05.

†Statistical values are expressed as t value or  $\chi^2$ .

‡Gross Motor is only for children aged <33 months.

§NT\$ represents new Taiwan dollars (US\$1=31.1 NT\$). There were 28 missing values (19 patients with ADHD and 9 controls).

ADHD, attention deficit hyperactivity disorder; KD, Kawasaki disease; PSI-SF, Parenting Stress Index-Short Form; WISC-IV, Wechsler

Intelligence Scale for Children-Fourth Edition; WPPSI, Wechsler Preschool and Primary Scale of Intelligence.

## DISCUSSION

This study demonstrates the potential effect of KD heterogeneity on cognitive development and parenting

stress. Our data revealed that no significant differences in developmental index, cognitive function or parenting stress between patients with KD and controls. Both IVIG

Table 2       Characteristics of children with KD	Table 2 Characteristics of children with KD and their caregivers regarding cognition development of children with KD			
	B (95% CI)	P value		
Children's characteristics				
Age (months)	0.07 (-0.01 to 0.15)	0.067		
Age of KD onset (months)	-0.01 (-0.16 to 0.13)	0.846		
Sex				
Male	1.38 (-3.43 to 6.19)	0.572		
Female	1			
Intravenous immunoglobulin				
None	3.01 (-10.86 to 16.88)	0.669		
Once	1.43 (-7.25 to 10.11)	0.746		
Twice	1			
Coronary artery aneurysms				
Without	1.32 (-3.42 to 6.07)	0.583		
With	1			
Caregivers' characteristics				
Age (years)	-0.07 (-0.66 to 0.52)	0.824		
Sex				
Male	-5.23 (-11.97 to 1.52)	0.128		
Female	1			
Education levels				
High school or lower	-3.73 (-12.25 to 4.78)	0.388		
College	-1.13 (-9.60 to 7.35)	0.794		
Master or above	1			

The dependent variable in the multiple linear regression model is a cognitive score (Early Learning Composite Score of the MSEL or the Full-Scale IQ of the WPPSI or WISC-IV). Data are expressed as B value, 95% CI and p value using multiple linear regression model. Adjusted R<sup>2</sup> of this regression model=0.054.

KD, Kawasaki disease; MSEL, Mullen Scales of Early Learning; WISC-IV, Wechsler Intelligence Scale for Children-Fourth Edition; WPPSI, Wechsler Preschool and Primary Scale of Intelligence.

	B (95% CI)	P value
Children's characteristics		
Age (months)	0.01 (-0.09 to 0.11)	0.834
Age of KD onset (months)	-0.18 (-0.37 to 0.01)	0.066
Sex		
Male	0.37 (-5.89 to 6.63)	0.907
Female	1	
ntravenous immunoglobulin		
None	1.46 (-16.60 to 19.51)	0.874
Once	-5.92 (-17.22 to 5.39)	0.303
Twice	1	
Coronary artery aneurysms		
Without	-7.41 (-13.59 to 1.23)	0.019*
With	1	
Caregivers' characteristics		
Age (years)	-0.11 (-0.88 to 0.66)	0.784
Sex		
Male	-0.57 (-9.35 to 8.21)	0.898
Female	1	
Education levels		
High school or lower	13.11 (2.03 to 24.20)	0.021*
College	14.55 (3.52 to 25.58)	0.010*
Master or above	1	

The dependent variable in the multiple linear regression model is the total score of the Parenting Stress Index. Data are expressed as B value, 95% Cl and p value using multiple linear regression model.

\*p<0.05. Adjusted  $R^2$  of this regression model=0.072.

KD, Kawasaki disease.

administration and CAL development were not associated with cognitive profiles of children with KD. In addition, the profiles of parenting stress in caregivers of patients with KD were examined in this study. We found that caregivers' education is associated to parenting stress, and the caregivers of patients who had CAL suffered from greater parenting stress than those of patients without CAL.

In our study sample, 4% never received IVIG, and 88.6% and 7.4% received IVIG treatment, once and twice, respectively. Although IVIG has been established as a standard treatment for KD,<sup>8</sup> 4% of patients still never received such intervention and 7.4% received IVIG treatment twice due to treatment resistance at the first administration. As the study sample included patients referred from local hospitals of remote areas, some of the patients may have missed the opportunity to be correctly diagnosed and receive IVIG at the critical period. Notably, 10%-20% of patients show resistance to IVIG therapy and are at risk for complications.<sup>9</sup> Our study population with KD consisted of 42% patients who exhibited CAL, a higher proportion of CAL than has been reported in this literature.<sup>10</sup> These studies suggest that approximately 20% of patients with KD suffer from CAL.<sup>47</sup> This discrepancy may be due to Kaohsiung

Chang Gung Memorial Hospital being the main medical centre in Southern Taiwan and patients recruited for this study possibly having a higher severity or greater comorbidities than those in general hospitals.

The results of this study show that KD is not related to cognitive impairment sequelae, which was generally consistent with the previous literature.<sup>19-21</sup> Because IVIG reduces vasculitis that presumably underlies any cognitive impairment, we assumed that patients with KD who had a poor response to IVIG or had developed CAL may be related to a higher severity of systemic vasculitis.<sup>13</sup><sup>14</sup> However, we found that neither IVIG administration nor CAL development was associated with cognitive performance. In our study population, most patients with KD (96%) received IVIG treatment. The case numbers of children who did not receive IVIG (only 4%) and treatment resistance (7.4%) were too small and it is difficult to make a sufficient comparison. Therefore, a future study with larger sample size is required to verify whether IVIG administration is associated with cognitive outcomes or not. Notably, a nationwide survey in Taiwan previously demonstrated that epilepsy and developmental delay were factors associated with cognitive impairments.<sup>21</sup> Whether physical comorbidities other than KD have a greater influence on cognitive function than KD itself or the disease characteristics of KD warrants further research.

We provide more scientific evidence related to parental stress or mental health of caregivers of patients with KD and keeping in mind that situation implies more health service. It is noteworthy that patients' caregivers were not assessed during patients' acute phase of KD. The clinical meaning of our findings is that parenting stress of caregivers in children with KD during follow-up was comparable those of control children. Nevertheless, our data revealed that caregivers of patients who had CAL suffered from greater parenting stress than the caregivers of patients who did not have CAL. A previous study from Canada revealed a similar finding with our study. The parents of children with coronary artery complications may have suffered from a greater anxiety level.<sup>25</sup> The psychological distress is associated with family characteristics, such as family income and maternal education.<sup>26</sup> Children with persistent CAL may develop complications,<sup>11</sup> so caregivers may worry, feel stressed and experience helplessness with regard to facing the uncertainty of their children's risk of myocardial infarction and the possibility of sudden death. This finding suggests that the parental stress or mental health of caregivers of patients with CAL require particular assistance. Furthermore, caregivers with an education level of a master's degree or above showed lower parenting stress than those who had education levels of college or lower. This finding may imply that caregivers with high education levels had greater internal or external resources to handle the patients' physical illness.<sup>26</sup> Alternatively, caregivers with lower education levels may require support or help to fulfil their parenting roles.

This study has certain limitations. First, owing to small case numbers of patients with KD without IVIG or treatment resistance, it is difficult to make a strong conclusion about cognitive impairment. Second, this is a casecontrol study, with a mean interval between KD onset and assessment for cognition and parenting stress of 40.5 months (ranged from 1 to 159 months). Therefore, the measurement for parenting stress was highly dependent on parental recall. The parenting stress reported herein did not necessarily represent the state of patients' caregivers who take care of patients during the acute onset of KD. Third, CAL was only recorded as a categorical variable (with or without), but the influence of CAL severity on cognition or parenting stress was not assessed in this study. Moreover, we did not record physical comorbidities besides KD. Whether other comorbidities (ie, developmental delay or epilepsy) actually influence or moderate children's cognitive development warrants further investigation. Moreover, an increase incidence of KD has been observed during the SARS-CoV-2 pandemic.<sup>48</sup> It suggests that SARS-CoV-2 may trigger a severe hyperinflammatory syndrome, such as paediatric inflammatory multisystem syndrome or KD-like symptoms, in childhood.<sup>49</sup> The influence of SARS-CoV-2 pandemic on the cognitive

profiles of patients with KD and parenting stress of their caregivers warrants further investigation. Fourth, the KD group and control group were not perfectly matched in age and caregivers' characteristics, and those differences may have influenced the results of this study. Finally, all participants were recruited from a single site, whether this finding can be generalised into other patient populations warrants further investigation.

#### CONCLUSION

No significant differences in developmental index, cognitive function or parenting stress between patients with KD and controls. In patients with KD, neither IVIG treatment nor CAL was associated with the cognitive profiles of patients with KD. This result is good news for caregivers and patients with KD, reassuring them that their IVIG response or CAL development will have no effect on their development milestones or cognitive function. However, caregivers of patients with KD who had CAL may feel stress about unpredictable sequela caused by CAL for their children. These caregivers may require support or help to fulfil their parenting roles.

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**Contributors** L-JW and L-SC: conceptualised and designed the study and drafted the initial manuscript. Z-YT: recruited and assessed the patients. H-CK: designed the study, recruited the patients and approved the final manuscript to be submitted.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not required.

Ethics approval Chang Gung Memorial Hospital's Internal Review Board approved this study (IRB No. 104-8261C), and we obtained the written informed consent from the parents or guardians of all participating children.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request.

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#### REFERENCES

- Newburger JW, Takahashi M, Burns JC. Kawasaki disease. J Am Coll Cardiol 2016;67:1738–49.
- 2 Hara T, Nakashima Y, Sakai Y, et al. Kawasaki disease: a matter of innate immunity. *Clin Exp Immunol* 2016;186:134–43.
- 3 Singh S, Vignesh P, Burgner D. The epidemiology of Kawasaki disease: a global update. Arch Dis Child 2015;100:1084–8.
- 4 Uehara R, Belay ED. Epidemiology of Kawasaki disease in Asia, Europe, and the United States. *J Epidemiol* 2012;22:79–85.
- 5 Zhu FH, Ang JY. The clinical diagnosis and management of Kawasaki disease: a review and update. *Curr Infect Dis Rep* 2016;18:32.

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- 6 Rowley AH. The complexities of the diagnosis and management of Kawasaki disease. *Infect Dis Clin North Am* 2015;29:525–37.
- 7 Kuo H-C, Yang KD, Chang W-C, et al. Kawasaki disease: an update on diagnosis and treatment. *Pediatr Neonatol* 2012;53:4–11.
- 8 Eleftheriou D, Levin M, Shingadia D, et al. Management of Kawasaki disease. Arch Dis Child 2014;99:74–83.
- 9 Qiu H, He Y, Rong X, et al. Delayed intravenous immunoglobulin treatment increased the risk of coronary artery lesions in children with Kawasaki disease at different status. *Postgrad Med* 2018;130:442–7.
- 10 Chen KYH, Curtis N, Dahdah N, *et al.* Kawasaki disease and cardiovascular risk: a comprehensive review of subclinical vascular changes in the longer term. *Acta Paediatr* 2016;105:752–61.
- 11 Joshi M, Tulloh R. Kawasaki disease and coronary artery aneurysms: from childhood to adulthood. *Future Cardiol* 2017;13:491–501.
- 12 Kuo H-C. Preventing coronary artery lesions in Kawasaki disease. *Biomed J* 2017;40:141–6.
- 13 Gitiaux C, Kossorotoff M, Bergounioux J, et al. Cerebral vasculitis in severe Kawasaki disease: early detection by magnetic resonance imaging and good outcome after intensive treatment. *Dev Med Child Neurol* 2012;54:1160–3.
- 14 Korematsu S, Uchiyama S-ichi, Miyahara H, et al. The characterization of cerebrospinal fluid and serum cytokines in patients with Kawasaki disease. *Pediatr Infect Dis J* 2007;26:750–3.
- 15 Hikita T, Kaminaga T, Wakita S, et al. Regional cerebral blood flow abnormalities in patients with Kawasaki disease. *Clin Nucl Med* 2011;36:643–9.
- 16 Emiroglu M, Alkan G, Kartal A, et al. Abducens nerve palsy in a girl with incomplete Kawasaki disease. *Rheumatol Int* 2016;36:1181–3.
- 17 Bailie NM, Hensey OJ, Ryan S, et al. Bilateral subdural collectionsan unusual feature of possible Kawasaki disease. Eur J Paediatr Neurol 2001;5:79–81.
- 18 Constantinescu CS, Migraine CCS. Migraine and Raynaud phenomenon: possible late complications of Kawasaki disease. *Headache* 2002;42:227–9.
- 19 King WJ, Schlieper A, Birdi N, *et al*. The effect of Kawasaki disease on cognition and behavior. *Arch Pediatr Adolesc Med* 2000;154:463–8.
- 20 Nishad P, Singh S, Sidhu M, et al. Cognitive and behaviour assessment following Kawasaki disease--a study from North India. *Rheumatol Int* 2010;30:851–4.
- 21 Wang L-J, Kuo H-C. Cognitive development after Kawasaki disease - clinical study and validation using a nationwide population-based cohort. *Circ J* 2018;82:517–23.
- 22 Dietz SM, van Stijn D, Burgner D, *et al.* Dissecting Kawasaki disease: a state-of-the-art review. *Eur J Pediatr* 2017;176:995–1009.
- 23 Kourtidou S, Slee AE, Bruce ME, et al. Kawasaki disease substantially impacts health-related quality of life. J Pediatr 2018;193:155–63.
- 24 Miura M, Kobayashi T, Kaneko T, et al. Association of severity of coronary artery aneurysms in patients with Kawasaki disease and risk of later coronary events. JAMA Pediatr 2018;172:e180030.
- 25 Chahal N, Clarizia NA, McCrindle BW, et al. Parental anxiety associated with Kawasaki disease in previously healthy children. J Pediatr Health Care 2010;24:250–7.
- 26 Chahal N, Jelen A, Rush J, et al. Kawasaki disease with coronary artery aneurysms: psychosocial impact on parents and children. J Pediatr Health Care 2017;31:459–69.
- 27 Jewell NP. Statistics for epidemiology. Boca Raton: CRC Press, 2003.
- 28 Newburger JW, Takahashi M, Gerber MA, et al. Diagnosis, treatment, and long-term management of Kawasaki disease: a statement for health professionals from the Committee on rheumatic fever, endocarditis, and Kawasaki disease, Council on cardiovascular disease in the young, American heart association. *Pediatrics* 2004;114:1708–33.

- 29 Kuo H-C, Hsieh K-S, Ming-Huey Guo M, et al. Next-Generation sequencing identifies micro-RNA-based biomarker panel for Kawasaki disease. J Allergy Clin Immunol 2016;138:1227–30.
- 30 Kuo H-C, Wang C-L, Liang C-D, et al. Association of lower eosinophil-related T helper 2 (Th2) cytokines with coronary artery lesions in Kawasaki disease. *Pediatr Allergy Immunol* 2009;20:266–72.
- 31 Kuo H-C, Yang KD, Liang C-D, et al. The relationship of eosinophilia to intravenous immunoglobulin treatment failure in Kawasaki disease. *Pediatr Allergy Immunol* 2007;18:354–9.
- 32 Kuo H-C, Chang J-C, Kuo H-C, et al. Identification of an association between genomic hypomethylation of FCGR2A and susceptibility to Kawasaki disease and intravenous immunoglobulin resistance by DNA methylation array. Arthritis Rheumatol 2015;67:828–36.
- 33 Mullen EM. *Mullen scales of early learning item administration book*. Circle Pines, MN: American Guidance Service, 1995.
- 34 Tsai J-M, Lu L, Jeng S-F, et al. Validation of the modified checklist for autism in toddlers, revised with follow-up in Taiwanese toddlers. *Res Dev Disabil* 2019;85:205–16.
- 35 Wechsler D. Wechsler preschool and primary scales of intelligence (4th ed) (WPPSI-IV). San Antonio, TX: Psychological Corporation, 2012.
- 36 Watkins MW, Beaujean AA. Bifactor structure of the Wechsler Preschool and Primary Scale of Intelligence--Fourth Edition. Sch Psychol Q 2014;29:52–63.
- 37 Lee Y-C, Lin C-H, Tsai C-H, et al. Association between executing theory of mind in a limited experimental context and executing it in daily contexts in children with autism spectrum disorder: a crosssectional study. Am J Occup Ther 2019;73:7303205150p1–11.
- 38 Baron IS. Test review: Wechsler intelligence scale for Children-Fourth edition (WISC-IV). *Child Neuropsychol* 2005;11:471–5.
- 39 Yang P, Cheng C-P, Chang C-L, et al. Wechsler intelligence scale for children 4th edition-Chinese version index scores in Taiwanese children with attention-deficit/hyperactivity disorder. *Psychiatry Clin Neurosci* 2013;67:83–91.
- 40 Wang L-J, Chan W-C, Chou M-C, et al. Polymorphisms of STS gene and SULT2A1 gene and neurosteroid levels in Han Chinese boys with attention-deficit/hyperactivity disorder: an exploratory investigation. *Sci Rep* 2017;7:45595.
- 41 Loyd BH, Abidin RR. Revision of the parenting stress index. J Pediatr Psychol 1985;10:169–77.
- 42 Haskett ME, Ahern LS, Ward CS, et al. Factor structure and validity of the parenting stress index-short form. J Clin Child Adolesc Psychol 2006;35:302–12.
- 43 Yeh CH, Chen ML, Li W, et al. The Chinese version of the parenting stress index: a psychometric study. Acta Paediatr 2001;90:1470–7.
- 44 Chen Y-C, Hwang-Gu S-L, Ni H-C, et al. Relationship between parenting stress and informant discrepancies on symptoms of ADHD/ODD and internalizing behaviors in preschool children. PLoS One 2017;12:e0183467.
- 45 Huang C-Y, Yen H-C, Tseng M-H, et al. Impacts of autistic behaviors, emotional and behavioral problems on parenting stress in caregivers of children with autism. J Autism Dev Disord 2014;44:1383–90.
- 46 Garson G. *Testing statistical assumptions*. USA: Statistical Associates Publishing, 2012.
- 47 McCrindle BW, Rowley AH, Newburger JW, et al. Diagnosis, treatment, and long-term management of Kawasaki disease: a scientific statement for health professionals from the American heart association. *Circulation* 2017;135:e927–99.
- 48 Cavallo F, Chiarelli F. An outbreak of Kawasaki-like disease in children during SARS-CoV- 2 epidemic: no surprise? *Acta Biomed* 2020;91:e2020015.
- 49 Berardicurti O, Conforti A, Ruscitti P, et al. The wide spectrum of Kawasaki-like disease associated with SARS-CoV-2 infection. Expert Rev Clin Immunol 2020;16:1205–15.