



Scientific Letter

Clinical Characteristics of Omicron (B.1.1.529) Variant in Children: A Multicenter Study in Spain



To the Director,

The severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) disease (COVID-19) was first reported in Wuhan (China) on December 31, 2019.¹ When the first patients infected with SARS-CoV-2 were analyzed, the genomic sequences of the single-stranded RNA virus were almost identical.² However, as the pandemic has evolved, so has the virus genome.³

The SARS-CoV-2 Omicron variant (B.1.1.529) was first identified in November 2021. On November 26, 2021, the WHO defined the strain as a variant of concern and named it Omicron. This new variant is associated with high transmissibility leading to high infectivity and probably increased reinfection rates.⁴ It is currently being proven that cases of severe COVID-19 are less frequent than with previous variants. For this reason, both governmental and scientific bodies are beginning to consider the possibility that in the more or less near future, SARS-CoV-2 infection should begin to be managed as a common endemic respiratory viral disease.^{5,6}

On November 29, the first case of infection by the Omicron variant was confirmed in Spain. According to data provided by the Center for Coordination of Alerts and Health Emergencies under the Ministry of Health of Spain, the Delta variant of the SARS-CoV-2 virus was the predominant one in our country until December 2021. After that time, it was displaced by the Omicron variant. In random sampling by specific PCR from 13 Spanish Autonomous Communities, carried out in the epidemiological week 52 of the year 2021, the proportion of infections by Omicron ranged from 48.7% (Murcia) to 93.8% (Madrid) of the total number of samples analyzed.⁷

A multicenter observational descriptive study was conducted in 5 secondary and tertiary hospitals of Madrid (Spain) between December 27, 2021, and January 2, 2022. Consecutive patients aged 0–16 attending the corresponding Pediatric Emergency Departments, with a positive result in the real-time polymerase chain reaction test (RT-PCR) or antigenic test to detect SARS-CoV-2 in the nasopharyngeal sample were included.

Quantitative variables were expressed by central tendency and dispersion [mean and standard deviation (\pm SD)]; absolute frequencies and percentages measured qualitative variables. Comparison between categorical variables was performed using the χ^2 test or Fisher's exact test. A p value of <0.05 was considered statistically significant.

Seven hundred seventy-four patients were diagnosed with SARS-CoV-2 infection during the study period. Overall, 68.9% reported having been in contact with a confirmed case of COVID-19. The mean age was 6.3 ± 4.7 years. Males accounted for 52.2% and females for 47.8%.

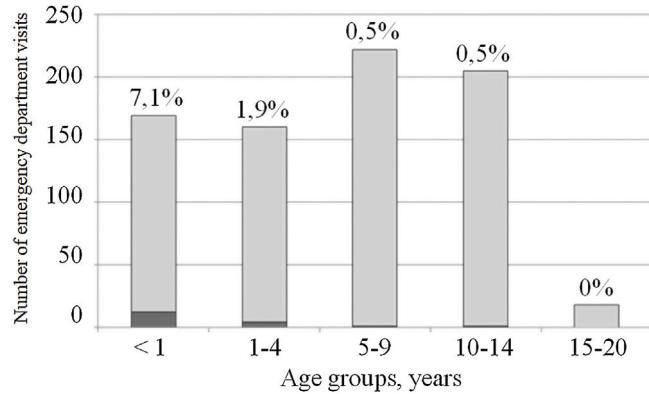


Fig. 1. Rates of hospitalization for COVID-19.

The most frequent clinical manifestation was fever (62.8%), with a mean duration time of 25.2 ± 39.9 h. Temperatures $\geq 39^\circ\text{C}$ were reported in 14.3% of cases, with children aged 1–4 years presenting the highest temperatures in a higher proportion (21.8%; $p = 0.016$) (Table 1).

Respiratory symptoms were predominant in all cases, regardless of age. The presence of cough and nasal discharge was observed in 59.3% and 53.5% of cases, respectively, and these percentages were very similar in all age ranges. Less than 3% of the cases presented with respiratory distress, mainly low respiratory distress, especially among children under 1. During their stay, chest X-rays were requested in 9 cases, 7 of which in patients who were hospitalized. In 6/9 cases (66.6%), the X-ray was considered abnormal (3 cases of interstitial infiltrates and 3 of consolidation).

Among gastrointestinal symptoms, the most frequent was vomiting (15.4%). Diarrhoea and abdominal pain were more frequent among children under 1-year-old (11.8%; $p = 0.019$) and 5–9 years old (13%; $p < 0.001$), respectively. Slightly more than half of the patients over 15 years of age (55.5%; $p < 0.001$) reported sore throat. Headache (33.3%; $p < 0.001$) and myalgia (11.1%; $p = 0.001$) were also more frequent in this age group. Only three patients had seizures at the time of diagnosis.

A total of 17/774 patients were hospitalized (2.3%). The rate of hospitalization was inversely proportional to the age of the cases, with the maximum rate observed being 7.1% among those under 1 year of age (Fig. 1). The main reason for admission was respiratory distress (8/17; 47%): 7 patients were admitted with a bronchiolitis diagnosis and one with severe laryngitis. In 3/17 (17.6%), the cause of hospitalization was the patients' underlying medical conditions: one case of short bowel and two patients with hemato-oncologic pathology. In all of them, the initial reason for consultation was fever. The other reasons for admission were: 2 cases in newborns less than 1-month-old, a 2-month-old infant

Table 1

Epidemiological and clinical characteristics, n(%).

Characteristics	All cases (n = 774)	Age groups					p value
		<1 year (n = 169)	1-4 years (n = 160)	5-9 years (n = 222)	10-14 years (n = 205)	15-20 years (n = 18)	
Sex							
Male	404 (52.2)	91 (53.8)	81 (50.6)	128 (57.6)	102 (49.7)	2 (11.1)	0.004
Female	370 (47.8)	78 (46.1)	79 (49.3)	94 (42.3)	103 (50.2)	16 (88.8)	
Presenting features							
Fever	486 (62.8)	120 (71)	113 (70.6)	126 (56.7)	119 (58)	8 (44.4)	0.001
≥39 °C	111 (14.3)	26 (15.3)	35 (21.8)	22 (9.9)	25 (12.1)	3 (16.6)	0.016
Respiratory symptoms							
Cough	459 (59.3)	103 (60.9)	106 (66.2)	118 (53.1)	121 (59)	11 (61.1)	0.141
Runny nose	414 (53.5)	102 (60.3)	98 (61.2)	96 (43.2)	109 (53.1)	9 (50)	0.002
Difficulty breathing							0.297
No difficulty breathing	756 (97.7)	161 (95.2)	156 (97.5)	219 (98.6)	202 (98.5)	18 (100)	
Bronchiolitis/Bronchospasm	11 (1.4)	6 (3.5)	3 (1.8)	1 (0.5)	1 (0.4)	0 (0)	
Laryngitis	4 (0.5)	1 (0.5)	1 (0.6)	0 (0)	2 (0.9)	0 (0)	
Gastrointestinal symptoms							
Vomiting	119 (15.4)	28 (16.5)	24 (15)	40 (18)	27 (13.1)	0 (0)	0.246
Diarrhea	51 (6.6)	20 (11.8)	12 (7.5)	9 (4)	9 (4.3)	1 (5.5)	0.019
Abdominal pain	54 (7)	1 (0.5)	11 (6.8)	29 (13)	12 (5.8)	1 (5.5)	0.000
Otorhinolaryngological symptoms							
Sore throat	127 (16.4)	5 (2.9)	17 (10.6)	32 (14.4)	63 (30.7)	10 (55.5)	0.000
Otalgia	16 (2.1)	0 (0)	7 (4.3)	5 (2.2)	3 (1.4)	1 (5.5)	0.055
Neurological manifestations							
Headache	133 (17.1)	0 (0)	14 (8.7)	54 (24.3)	59 (28.7)	6 (33.3)	0.000
Seizures	3 (0.4)	0 (0)	2 (1.8)	1 (0.5)	0 (0)	0 (0)	0.327
Anosmia	2 (0.3)	0 (0)	0 (0)	0 (0)	2 (0.9)	0 (0)	0.234
Mucocutaneous Manifestations							
Skin rash	9 (1.2)	2 (1.1)	5 (3.1)	0 (0)	2 (0.9)	0 (0)	0.083
Conjunctivitis	6 (0.8)	2 (1.1)	1 (0.6)	1 (0.4)	2 (0.9)	0 (0)	0.915
Others							
Myalgia	19 (2.4)	0 (0)	5 (3.1)	1 (0.4)	11 (5.3)	2 (11.1)	0.001
Thoracic pain	10 (1.2)	0 (0)	1 (0.6)	4 (1.8)	4 (1.9)	1 (5.5)	0.356
Arthralgia	2 (0.3)	0 (0)	1 (0.6)	0 (0)	1 (0.4)	0 (0)	0.678

admitted for a 4-day persistent fever associated with rash, and a 7-year-old child admitted progressive hypoxia. Finally, in one case, the diagnosis of SARS-CoV-2 infection was done by screening for admission for diabetes debut.

Since its identification, the Omicron variant of the SARS-CoV-2 virus has shown a more rapid spread than previous variants.⁸ Omicron is characterized by the high number of mutations identified in its genome,⁹ that appear to be responsible for its high capacity to cause reinfection and its partial resistance to existing vaccines.^{10,11} These characteristics have enabled it to spread rapidly throughout the world,¹² leading to higher levels of COVID-19 incidence than in previous pandemic periods. However, as we have also seen in our series, indicators of disease severity are lower compared to previous pandemic peaks.^{13,14}

As with previous variants,¹⁵ fever and respiratory symptoms are the predominant symptoms of Omicron infection. However, some authors have begun to report seizures associated with infection by this variant.¹⁶ In the study by Cloete et al.¹⁷ conducted in the pediatric population of the Gauteng province in South Africa, the origin of the Omicron wave, 20% of hospitalizations were for febrile seizures.

Our hospitalization rate was 2.3%, mainly concentrated in children under 4 years of age. A multicenter study in 14 North American states¹⁸ analyzed the evolution of hospitalizations due to COVID-19, specifically in children in this age range, during the period of the predominance of Omicron; the weekly hospitalization rates were approximately five times lower than those during the period of the predominance of Delta variant.

The main reason for the hospitalization of our patients was the presence of respiratory distress. Most were diagnosed with bron-

chiolitis and, in one case, with laryngitis; both conditions have been previously described in the context of SARS-CoV-2 infection and are associated explicitly with the Omicron variant.¹⁹ In our series, only in one case was the diagnosis of COVID-19 made by screening at admission for other condition different than COVID-19. Other studies have found up to 63% of the incidental diagnosis of COVID-19 in diagnostic tests prior to hospitalization.²⁰

Our study has some limitations: the diagnostic cases of COVID-19 analyzed were assumed to be caused by the Omicron variant based on the epidemiological analyses carried out by the Community of Madrid, which reported that Omicron was the predominant circulating variant in the study period. The variables analyzed did not include the vaccination status of the children, a factor that can undoubtedly condition the severity of the disease. However, according to data provided by the Spanish Ministry of Health, by the end of January 2022, 90.8% of the population over 12 years of age had a complete vaccination schedule. As of December 15, 2021, vaccination of children between 5 and 11 years of age was initiated.

In conclusion, Omicron variant infection in pediatric patients causes mild symptoms and is associated with a low hospitalization rate, especially in children who have not yet received SARS-CoV-2 vaccination.

Authors' contributions

Conception and design of the study: MAM. Acquisition of data: BGC, CMDR, JAR, JTRA, ICC, LST, MAGH, MAM, MDC, and MJP. Analysis and interpretation of data: JRD. All authors contributed in drafting the article.

Funding

The present investigation has not received specific support from public sector agencies, the commercial sector, or non-profit organizations.

Conflict of interest

The authors declare no conflicts of interest.

References

1. Wang C, Horby PW, Hayden FG, Gao GF. A novel coronavirus outbreak of global health concern. *Lancet*. 2020;395:470–3.
2. Lu R, Zhao X, Li J, Niu P, Yang B, Wu H, et al. Genomic characterisation and epidemiology of 2019 novel coronavirus: implications for virus origins and receptor binding. *Lancet*. 2020;395:565–74.
3. Tracking SARS-CoV-2 variants. Organización Mundial de la Salud (OMS). <https://www.who.int/en/activities/tracking-SARS-CoV-2-variants/> [accessed 12.01.22].
4. Papanikolaou V, Chrysovergis A, Ragos V, Tsiambas E, Katsinis S, Manoli A, et al. From delta to Omicron: S1-RBD/S2 mutation/deletion equilibrium in SARS-CoV-2 defined variants. *Gene*. 2022;814:146134.
5. Hacia el fin de la excepcionalidad (Editorial). AMF 2022;4. <https://amf-semfyc.com/web/article/3063> [accessed 12.01.22].
6. Romero, JM. Sánchez plantea evaluar el covid como una gripe y controlará el precio de los antígenos. 10 de enero de 2022. El Periódico. <https://www.elperiodico.com/es/sociedad/20220110/sanchez-estrategia-seguimiento-covid-gripe-precio-antigenos-13076306> [accessed 12.01.22].
7. Centro de Coordinación de Alertas y Emergencias Sanitarias. Actualización de la situación epidemiológica de las variantes de SARS-CoV-2 en España (10 de enero de 2022). Ministerio de Sanidad. https://www.sanidad.gob.es/profesionales/saludPublica/ccayes/alertasActual/nCov/documentos/COVID19_Actualizacion_variantes_20220110.pdf.
8. Enhancing response to Omicron SARS-CoV-2 variant. Technical document. Organización Mundial de la Salud (OMS). [https://www.who.int/publications/m/item/enhancing-readiness-for-omicron-\(b.1.1.529\)-technical-brief-and-priority-actions-for-member-states](https://www.who.int/publications/m/item/enhancing-readiness-for-omicron-(b.1.1.529)-technical-brief-and-priority-actions-for-member-states) [accessed 03.04.22].
9. Bansal K, Kumar S. Mutational cascade of SARS-CoV-2 leading to evolution and emergence of omicron variant. *Genomics*. 2021; <http://dx.doi.org/10.1101/2021.12.06.471389> [cited 2.4.22].
10. Araf Y, Akter F, Tang Y, Fatemi R, Parvez MdSA, Zheng C, et al. Omicron variant of SARS-CoV-2: genomics, transmissibility, and responses to current COVID-19 vaccines. *J Med Virol*. 2022;94:1825–32.
11. Chen LL, Chua GT, Lu L, Chan BPC, Wong JSC, Chow CCK, et al. Omicron variant susceptibility to neutralizing antibodies induced in children by natural SARS-CoV-2 infection or COVID-19 vaccine. *Emerg Microb Infect*. 2022;11:543–7.
12. Elliott P, Bodinier B, Eales O, Wang H, Haw D, Elliott J, et al. Rapid increase in Omicron infections in England during December 2021: REACT-1 study. *Science*. 2022;375:1406–11.
13. Wang L, Berger NA, Kaelber DC, Davis PB, Volkow ND, Xu R. Comparison of outcomes from COVID infection in pediatric and adult patients before and after the emergence of Omicron. *Infect Dis (except HIV/AIDS)*. 2022; <http://dx.doi.org/10.1101/2021.12.30.21268495> [scheduled for April 2, 2022].
14. Juliano AD, Brunkard JM, Boehmer TK, Peterson E, Adjei S, Binder AM, et al. Trends in disease severity and health care utilization during the early omicron variant period compared with previous SARS-CoV-2 high transmission periods—United States, December 2020–January 2022. *Morb Mortal Wkly Rep*. 2022;71:146–52.
15. A multicenter study of confirmed COVID-19 cases: preliminary data on 2690 pediatric patients in Argentina during the first year of the pandemic. *Arch Argent Pediatr*. 1 de abril de 2022 [scheduled for April 3, 2022];120. Available from: <https://www.sap.org.ar/docs/publicaciones/archivosarg/2022/v120n2a04e.pdf>.
16. Ludvigsson JF. Convulsions in children with COVID-19 during the Omicron wave. *Acta Paediatr*. 2022; apa.16276.
17. Cloete J, Kruger A, Mashu M, du Plessis NM, Mawela D, Tshukudu M, et al. Pediatric hospitalisations due to COVID-19 during the first SARS-CoV-2 omicron (B.1.1.529) variant wave in South Africa: a multicentre observational study. *Lancet Child Adolesc Health*. 2022;6:294–302.
18. Marks KJ, Whitaker M, Agathis NT, Anglin O, Milucky J, Patel K, et al. Hospitalization of infants and children aged 0–4 years with laboratory-confirmed COVID-19—COVID-NET, 14 states, March 2020–February 2022. *Morb Mortal Wkly Rep*. 2022;71:429–36.
19. Milani GP, Bollati V, Ruggiero L, Bosisi S, Pinzani RM, Lunghi G, et al. Bronchiolitis and SARS-CoV-2. *Arch Dis Child*. 2021;106:999–1001.
20. Abdullah F, Myers J, Basu D, Tintinger G, Ueckermann V, Mathebula M, et al. Decreased severity of disease during the first global omicron variant covid-19 outbreak in a large hospital in Tshwane, South Africa. *Int J Infect Dis*. 2022;116:38–42.

Miguel Ángel Molina Gutiérrez ^{a,*}, Lara Sánchez Trujillo ^b, José Antonio Ruiz Domínguez ^a, Ignacio Callejas Caballero ^c, Beatriz García Cuartero ^d, María Ángeles García-Herrero ^b, María Jesús Pascual Marcos ^e, José Tomás Ramos Amador ^f, Carmen Martínez del Río ^f, María de Ceano-Vivas La Calle ^a

^a Pediatric Emergency Department, Hospital Universitario La Paz, Madrid, Spain

^b Pediatric Emergency Department, Hospital Universitario Príncipe de Asturias, Alcalá de Henares, Spain

^c Department of Paediatrics, Hospital Universitario de Getafe, Madrid, Spain

^d Unidad de Endocrinología y Diabetes Pediátrica, Servicio de Pediatría, Hospital Universitario Ramón y Cajal, Madrid, Spain

^e Department of Paediatrics, Hospital Vithas Madrid Aravaca, Madrid, Spain

^f Department of Paediatrics, Hospital Universitario Clínico San Carlos, Madrid, Spain

Corresponding author.

E-mail address: malacatin@hotmail.com (M.Á.M. Gutiérrez).