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Age-Stratified Risk of Severe COVID-19 for People With Disabilities in Korea: Nationwide Study Considering Disability Type

Boyeong Ryu , 1,2 Hoyeon Jang , 3 Jaiyong Kim , 5 Sung-il Cho , 2 and Seong-Sun Kim , 1

¹Department of Data Science, Korea Disease Control and Prevention Agency (KDCA), Cheongju, Korea ²Department of Public Health Science, Graduate School of Public Health, Seoul National University, Seoul, Korea

³Department of Big Data Research and Development, National Health Insurance Service, Wonju, Korea



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Address for Correspondence:

Sung-il Cho, MD, PhD

Department of Public Health Science, Graduate School of Public Health, Seoul National University, 1 Gwanak-ro, Gwanak-gu, Seoul 08826, Republic of Korea. Email: persontime@hotmail.com

Seong-Sun Kim, PhD

Department of Data Science, Korea Disease Control and Prevention Agency (KDCA), 187 Osongsaengmyeong 2-ro, Osong-eup, Heungdeok-gu, Cheongju 28159, Republic of Korea.

Email: sskim0719@korea.kr

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ORCID iDs

Boyeong Ryu 🕩

https://orcid.org/0000-0001-9336-8098

Hoyeon Jang 📵

https://orcid.org/0000-0002-5446-7752 Jaiyong Kim (D)

https://orcid.org/0000-0003-0985-7871

ABSTRACT

Background: Understanding disparities in severe coronavirus disease 2019 (COVID-19) outcomes between people with disabilities (PwD) and people without disabilities (PwoD) is crucial, particularly when considering the heterogeneity within PwD and age differences. This study aimed to compare severe COVID-19 outcomes including deaths between PwD and PwoD with analyses stratified by age group and further examined by disability type. Methods: This retrospective, population-based cohort study used linked data from national COVID-19 cases and health insurance for individuals aged ≥ 19 years with COVID-19 from January 2020 to October 2022 in the Republic of Korea. Severe outcomes included severe cases and deaths, with logistic regression analysis of the risk disparities between PwD and PwoD based on age group and disability types. The subgroup analysis considered epidemic periods, accounting for the severe acute respiratory syndrome coronavirus 2 variant circulation. Results: The risk of severe COVID-19 outcomes and deaths among PwD varied by age and disability type. While severe outcomes were most prevalent in the older age groups for both PwD and PwoD, younger PwD faced a markedly higher risk—up to eightfold—compared to PwoD. The risk of disability status was greater than that of comorbidities in the 19-39 age group. Among disability types, individuals with internal organs-related and intellectual disabilities showed higher risk disparities with PwoD in severe outcomes than other types of disabilities. Throughout the pandemic, the disparity in death risk remained similar, with a slight increase in disparity during the omicron period for all severe outcomes in the age groups 19-39 and 40-64 years.

Conclusion: Prioritizing younger PwD, along with older age groups and people with comorbidities, is crucial in addressing public health crises. Risk-based prioritization is important to reduce overall risk. This includes prioritizing people with nternal organs-related and intellectural disabilities, who face higher health risks among PwD during a pandemic when resources are limited and time is of the essence.

Keywords: SARS-CoV-2; People With Disabilities; Severity; Fatality; Disability Types; Age Stratification



Sung-il Cho (b)
https://orcid.org/0000-0003-4085-1494
Seong-Sun Kim (b)
https://orcid.org/0000-0001-5277-492X

Disclosure

The authors have no potential conflicts of interest to disclose.

Author Contributions

Conceptualization: Ryu B, Kim SS, Cho S.
Data curation: Ryu B, Jang H, Kim J. Formal
analysis: Ryu B. Methodology: Ryu B, Kim SS,
Cho S. Supervision: Kim SS, Cho S. Writing original draft: Ryu B. Writing - review & editing:
Jang H, Kim J, Kim SS, Cho S.

INTRODUCTION

People with disabilities (PwD) are more vulnerable to health risks than people without disabilities (PwoD). During the coronavirus disease 2019 (COVID-19) pandemic, several countries including England, the Republic of Korea, and the United States consistently reported a higher risk of death among PwD, despite variations in the definition of disability. ²⁻⁴

Regarding the diversity of health statuses and contributing factors, numerous studies have explored health disparities by disability type pre- and during pandemic. ⁵⁻⁷ A Korean study conducted before the pandemic found that the overall age-standardized mortality rate (SMR) was 2.7 times higher in PwD than in PwoD. For people with specific disabilities, including respiratory or hepatic impairment and intestinal or urinary fistulas, the SMR was 10-fold higher than that in PwoD. Socioeconomic status also varied by disability type, with 24.7% people with intestinal or urinary fistulas and 61.1% people with epilepsy disorders accounting for the lowest income group in their respective cohorts. During the pandemic, a US study found that people with intellectual and developmental disabilities (IDDs) and mobility disabilities faced a higher risk of invasive mechanical ventilation and mortality. ⁹

Disability types predominantly differ across age groups, leading to variation in health outcomes. Adjusting for age is crucial when comparing the health statuses of PwD and PwoD. Studies show that hearing impairment, with a lower SMR than other types of disability, is more prevalent in older adults, while intellectual disability, with a higher SMR, is more common in young and middle-aged adults. 8,10,11 During the COVID-19 pandemic, risk factors for severe outcomes differed by age, with older age being the most notable risk factor. 12 Given these differences, age stratified analyses are more appropriate than using age as an effect modifier, provided the sample size is large enough. 12 Some studies highlight higher risk disparities in younger age groups but often focus on specific disability types, primarily IDDs, or aggregate all types. 6,13,14 Comprehensible assessments that consider various disabilities and age concurrently are limited. Nationwide data from the Republic of Korea, including all confirmed COVID-19 cases, enable stratified analysis with a sufficient sample size. Linking these data with health insurance information, including sociodemographic and disability status, provides a detailed analysis of vulnerability to severe COVID-19 outcomes, focusing on disability types with age stratification.

Throughout the pandemic, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) continuously mutated, altering its virulence. ¹⁵ This study examined evolving risk disparities of severe COVID-19 outcomes, considering effects of the virus's virulence on PwD and PwoD according to disability status and type. All analyses were stratified by age group and epidemic period-related SARS-CoV-2 variants.

METHODS

Study design and participants

This retrospective population-based cohort study used nationwide data and included individuals aged ≥ 19 years with COVID-19 in the Republic of Korea from January 20, 2020, to October 29, 2022. According to the Infectious Disease Control and Prevention Act, the Korea Diseases Control and Prevention Agency (KDCA) received reports of all confirmed cases that



tested positive through polymerase chain reaction tests in the entire period or rapid antigen tests from March 14, 2022 onwards. ¹⁶

The reported data were combined with the COVID-19 vaccination information from the National Immunization Registry of KDCA. The combined data were linked to the National Health Information Database (NHID) of the Korean National Health Insurance Service. ¹⁷ By linking KDCA and NHID data, this study excluded unmatched cases, as well as individuals without health insurance level or with errors in disability status (**Supplementary Fig. 1**).

Variables

Study outcomes were two levels of severe COVID-19 outcomes: a wide range of severe outcomes, including severe and critical status and deaths ('all severe outcomes' hereafter), and a narrow range, including only deaths. Severe and critical cases were defined as those requiring high-flow oxygenation, non-invasive or invasive ventilation, extracorporeal membrane oxygenation, or continuous renal replacement treatment during isolation due to the COVID-19 according to the World Health Organization (WHO) and monitored by KDCA. 18,19 Death was defined as a reported COVID-19-related death based on clinician statements under the Infectious Disease Control and Prevention Act. 16

Disability is defined as official registration with the government under the Act on Welfare of Persons with Disabilities. ²⁰ The 15 main disability types were classified into four groups: physical disabilities with external bodily functions (physical disabilities, brain lesion disorders, visual and hearing impairment, language disabilities, facial disfigurement) and with internal organs (renal, cardiac, hepatic and respiratory impairment, intestinal/urinary fistula, epilepsy disorder), and mental disabilities with psychological development disorder (intellectual disabilities and autistic disorder) and with mental disease. In cases with multiple disability types, the most severe type was considered.

Potential explanatory factors that could be confounders and modifiers were included: demographic characteristics (age and sex), socioeconomic status with health insurance status, and comorbidities. To adjust for variation in the host immune status and virulence of SARS-CoV-2, we included COVID-19 vaccination, reinfection status, and epidemic periods related to SARS-CoV-2 variants in the model. Additionally, we conducted stratified analyses by epidemic periods to assess the differential impact of SARS-CoV-2 variants on severe outcomes between PwD and PwoD. Based on the time of COVID-19 confirmation, health insurance status was categorized into 11 groups, including medical aid and quintiles for self-employed and employee groups. The first quintile represented the lowest income. Comorbidities included diabetes, hypertension, kidney, pulmonary, cardiovascular, liver and neurological diseases, cancer, and others (such as mental and behavior disorders, malnutrition, and obesity). Vaccination was the last vaccination status 14 days before the COVID-19 confirmation. Reinfection was testing positive 45–90 days after the previous confirmation, with symptoms and exposure to confirmed cases or an overseas travel history or 90 days after the previous confirmation, regardless of symptoms. Epidemic periods were classified based on > 50% circulating variants of SARS-CoV-2 among tested samples by reported date: pre-delta (January 20, 2020, to July 24, 2021), delta (July 25, 2021, to January 15, 2022), and omicron (January 16, 2022, to October 29, 2022). Age-stratified models were analyzed, recognizing age as a strong effect modifier (P < 0.001).



Statistical analysis

The analyses were stratified by three age groups: 19–39 ('young-age group'), 40–64 ('middle-age group'), and ≥ 65 years ('old-age group'). For the descriptive analysis, frequency, percentage, and mean age were used. To evaluate the risk of severe COVID-19 outcomes by age group and epidemic period, age- and sex-standardized case severity rate (aCSR) and case fatality rate (aCFR) were calculated. Using the number of confirmed cases as the denominator, aCSR was calculated with the number of severe cases, critical cases, and deaths as the numerator, and aCFR with the number of deaths as the numerator. Direct standardization was conducted using the total number of confirmed cases within the 5-year age group as the standard population.

To evaluate the risk of disability for severe outcomes of COVID-19, we conducted multiple logistic regression analyses with adjustment for explanatory factors, calculating adjusted odds ratios (aORs) for disability. The analysis examined the overall risk of disability and the specific risk for each disability type. Additionally, the disability risk for severe outcomes was calculated across different epidemic periods. All models were considered with the 95% confidence interval (CI). All statistical analyses were performed using the SAS Enterprise Guide 8.2 (8.2.0.1201; SAS Institute, Cary, NC, USA).

Ethics statement

This study utilized secondary and deidentified individual data, exemption from Institutional Review Board (IRB) review and approval was received from the Korea Diseases Control and Prevention Agency IRB (IRB No. 2023-04-12). The research number of the study is NHIS-2024-1-033.

RESULTS

Baseline demographics

Of 25,504,288 confirmed cases of COVID-19, 19,035,753 individuals with COVID-19 aged \geq 19 years were included after linkage with NHID (**Supplementary Fig. 1**). Overall, PwD accounted for 5.2% (992,485); the old-age group had the highest proportion of PwD at 15.9%, and the young-age group had the lowest proportion at 1.5% (**Table 1**). Compared to PwoD, more PwD were old (\geq 65 years; 54.1% for PwD, 15.7% for PwoD), of low socioeconomic status (Medical aid; 19.0% for PwD, 1.8% for PwoD), and had comorbidities, except liver disease in the old-age group. PwD in the young-age and middle-age groups had a higher booster vaccination rate than PwoD. Simultaneously, the proportion of PwD that were not vaccinated was also higher than that of PwoD in all age groups. Despite PwD was 5.2% of overall cases, PwD among severe/critical cases and deaths were 28.3% and 32.6%, respectively, and had the highest proportion, at 44.8%, of deaths, in the 19–39-year age group.

Among PwD of all ages, physical and hearing disabilities were the most common (**Supplementary Table 1**). However, the young-age group showed different patterns, in that intellectual disabilities were the most common at 46.6%, followed by physical disabilities at 17.3%.

aCSR and aCFR

Over the entire period, the aCSR and aCFR were higher in PwD than in PwoD in all age groups (Tables 2 and 3). The aCSR was 2.1 and 6.4 per 1,000 in PwoD and PwD, respectively; 3.05 times higher than that in PwD. The aCFR was 1.2 and 3.3 per 1,000 in PwoD and PwD, respectively;



Table 1. Characteristics of coronavirus disease 2019 confirmed cases among people with and without disability based on age group (N = 19,035,753)

Characteristics	Total	Sub	-total	19-	-39 yr	40-	64 yr	> 6	65 yr
		Disabled	Non-disabled	Disabled	Non-disabled	Disabled	Non-disabled	Disabled	Non-disabled
Total	19,035,753 (100.0)	992,485 (100.0)	18,043,268 (100.0)	105,403 (100.0)	7,080,971 (100.0)	350,513 (100.0)	8,132,919 (100.0)	536,569 (100.0)	2,829,378 (100.0)
Sex									
Male	8,544,857 (44.9)	540,987 (54.5)	8,003,864 (44.4)	66,558 (63.2)	3,404,447 (48.1)	219,386 (62.6)	3,451,454 (42.4)	255,043 (47.5)	1,147,963 (40.6)
Female	10,490,903 (55.1)	451,498 (45.5)	10,039,404 (55.6)	38,845 (36.9)	3,676,524 (51.9)	131,127 (37.4)	4,681,465 (57.6)	281,526 (52.5)	1,681,415 (59.4)
Age, yr	, ,	, ,	, ,	, ,	, ,	, ,	, ,	, ,	, ,
Mean ± SD	47.0 ± 17.5	64.0 ± 17.4	46.0 ± 17.0	29.5 ± 6.0	29.2 ± 5.9	54.6 ± 7.0	51.0 ± 7.2	77.0 ± 7.8	73.7 ± 7.4
19-29	3,742,319 (19.7)	52,457 (5.3)	3,689,862 (20.5)	52,457 (49.8)	3,689,862 (52.1)	-	-	-	-
30-39	3,444,055 (18.1)	52,946 (5.3)	3,391,109 (18.8)	52,946 (50.2)	3,391,109 (47.9)	-	-	-	-
40-49	3,777,735 (19.9)	91,216 (9.2)	3,686,519 (20.4)	-	-	91,216 (26.0)	3,686,519 (45.3)	-	-
50-59	3,192,632 (16.8)	145,942 (14.7)	3,046,690 (16.9)	-	-	145,942 (41.6)	3,046,690 (37.5)	-	-
60-69	2,683,981 (14.1)	228,304 (23.0)	2,455,677 (13.6)	-	-	113,355 (32.3)	1,399,710 (17.2)	114,949 (21.4)	1,055,967 (37.3)
70-79	1,356,070 (7.1)	214,707 (21.6)	1,141,363 (6.3)	-	-	-	-	214,707 (40.0)	1,141,363 (40.3)
≥ 80	838,961 (4.4)	206,913 (20.9)	632,048 (3.5)	-	-	-	-	206,913 (38.6)	632,048 (22.3)
Insurance level	()	(20.0)	(0.0)					(00.0)	(22.0)
Medical aid	518,573 (2.7)	188,371 (19.0)	330,202 (1.8)	22,218 (21.1)	65,347 (0.9)	91,890 (26.2)	115,890 (1.4)	74,263 (13.8)	148,965 (5.3)
Self-employed 1Q		, ,	676,766 (3.8)	5,044 (4.8)	260,495 (3.7)	, ,	265,510 (3.3)	41,646 (7.8)	, ,
Self-employed 2Q	785,899 (4.1)	39,939 (4.0)	745,960 (4.1)	3,207 (3.0)	305,687 (4.3)	13,493 (3.9)	319,701 (3.9)	23,239 (4.3)	120,572 (4.3)
Self-employed 3Q	813,715 (4.3)	39,927 (4.0)	773,788 (4.3)	3,059 (2.9)	266,251 (3.8)	13,952 (4.0)	369,092 (4.5)	22,916 (4.3)	138,445 (4.9)
Self-employed 4Q	, ,	, ,	785,377 (4.4)	2,530 (2.4)	226,267 (3.2)	, ,	388,375 (4.8)	, ,	170,735 (6.0)
Self-employed 5Q		. ,	835,907 (4.6)	1,930 (1.8)	152,716 (2.2)	. ,	395,207 (4.9)	. ,	287,984 (10.2)
Employee 1Q	2,628,052 (13.8)	145,686 (14.7)	2,482,366 (13.8)	24,460 (23.2)	862,018 (12.2)	46,943 (13.4)	1,175,235 (14.5)	74,283 (13.8)	445,113 (15.7)
Employee 2Q	2,874,411 (15.1)	119,315 (12.0)	2,755,096 (15.3)	14,399 (13.7)	1,119,303 (15.8)	43,562 (12.4)	1,256,319 (15.5)	61,354 (11.4)	379,474 (13.4)
Employee 3Q	3,000,522 (15.8)	101,979 (10.3)	2,898,543 (16.1)	10,759 (10.2)	1,374,925 (19.4)	35,019 (10.0)	1,191,955 (14.7)	56,201 (10.5)	331,663 (11.7)
Employee 4Q	2,968,045 (15.6)	105,565 (10.6)	2,862,480 (15.9)	9,746 (9.3)	1,317,672 (18.6)	31,436 (9.0)	1,212,382 (14.9)	64,383 (12.0)	332,426 (11.8)
Employee 5Q	2,998,225 (15.8)	101,442 (10.2)	2,896,783 (16.1)	8,051 (7.6)	1,130,290 (16.0)	30,952 (8.8)	1,443,253 (17.8)	62,439 (11.6)	323,240 (11.4)
Comorbidities									
Diabetes	2,057,426 (10.8)	279,417 (28.2)	1,778,009 (9.9)	5,274 (5.0)	90,482 (1.3)	78,062 (22.3)	847,569 (10.4)	196,081 (36.5)	839,958 (29.7)
Hypertension	4,054,768 (21.3)	493,097 (49.7)	3,561,671 (19.7)	7,310 (6.9)	159,715 (2.3)	124,825 (35.6)	1,726,614 (21.2)	360,962 (67.3)	1,675,342 (59.2)
Kidney disease	568,311 (3.0)	112,525 (11.3)	455,786 (2.5)	4,411 (4.2)	78,339 (1.1)	34,875 (10.0)	191,427 (2.4)	73,239 (13.7)	186,020 (6.6)
Pulmonary disease	771,610 (4.1)	135,579 (13.7)	636,031 (3.5)	5,889 (5.6)	167,458 (2.4)	30,963 (8.8)	212,258 (2.6)	98,727 (18.4)	256,315 (9.1)
Cardiovascular disease	2,719,626 (14.3)	411,258 (41.4)	2,308,368 (12.8)	6,905 (6.6)	138,450 (2.0)	93,061 (26.6)	953,870 (11.7)	311,292 (58.0)	1,216,048 (43.0)
Liver disease	2,014,978 (10.6)	152,499 (15.4)	1,862,479 (10.3)	9,814 (9.3)	344,199 (4.9)	62,370 (17.8)	1,059,295 (13.0)	80,315 (15.0)	458,985 (16.2)
Neurologic disease	3,142,390 (16.5)	443,165 (44.7)	2,699,225 (15.0)	28,967 (27.5)	366,960 (5.2)	125,989 (35.9)	1,215,496 (15.0)	288,209 (53.7)	1,116,769 (39.5)
Cancer	784,316 (4.1)	80,075 (8.1)	704,241 (3.9)	1,837 (1.7)	66,375 (0.9)	20,585 (5.9)	366,736 (4.5)	57,653 (10.7)	271,130 (9.6)
Others	3,414,332 (17.9)	467,492 (47.1)	2,946,840 (16.3)	40,380 (38.3)	740,671 (10.5)	135,324 (38.6)	1,199,115 (14.7)	291,788 (54.4)	1,007,054 (35.6)
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Table 1. (Continued) Characteristics of coronavirus disease 2019 confirmed cases among people with and without disability based on age group (N = 19,035,753)

Characteristics	Total	Sub	-total	19-	39 yr	40-	64 yr	≥ 6	55 yr
		Disabled	Non-disabled	Disabled	Non-disabled	Disabled	Non-disabled	Disabled	Non-disabled
Vaccination									
Unvaccinated	1,124,760	76,132	1,048,628	8,301	497,582	24,319	406,603	43,512	144,443
	(5.9)	(7.7)	(5.8)	(7.9)	(7.0)	(6.9)	(5.0)	(8.1)	(5.1)
Incomplete	265,675	11,763	253,912	1,243	128,754	4,495	101,108	6,025	24,050
	(1.4)	(1.2)	(1.4)	(1.2)	(1.8)	(1.3)	(1.2)	(1.1)	(0.9)
Complete	5,519,212	116,823	5,402,389	22,837	3,151,398	52,471	2,046,728	41,515	204,263
	(29.0)	(11.8)	(29.9)	(21.7)	(44.5)	(15.0)	(25.2)	(7.7)	(7.2)
Booster	12,126,106	787,767	11,338,339	73,022	3,303,237	269,228	5,578,480	445,517	2,456,622
	(63.7)	(79.4)	(62.8)	(69.3)	(46.7)	(76.8)	(68.6)	(83.0)	(86.8)
Period									
Pre-delta	154,445 (0.8)	,	145,612 (0.8)	881 (0.8)	52,257 (0.7)	3,582 (1.0)	70,587 (0.9)	4,370 (0.8)	22,768 (0.8)
Delta	. ,	. ,	,	1,760 (1.7)	127,020 (1.8)	8,100 (2.3)	157,447 (1.9)	11,872 (2.2)	62,527 (2.2)
Omicron	18,512,582	961,920	17,550,662	102,762	6,901,694	338,831	7,904,885	520,327	2,744,083
	(97.3)	(96.9)	(97.3)	(97.5)	(97.5)	(96.7)	(97.2)	(97.0)	(97.0)
Reinfection									
Yes	328,150 (1.7)	21,980 (2.2)	306,170 (1.7)	2,368 (2.3)	144,037 (2.0)	7,221 (2.1)	116,898 (1.4)	12,391 (2.3)	45,235 (1.6)
Severe outcomes									
Severe and critical	26,687 (0.1)	7,553 (0.8)	19,134 (0.1)	187 (0.2)	767 (0.0)*	1,556 (0.4)	5,026 (0.1)	5,810 (1.1)	13,341 (0.5)
Death	27,269 (0.1)	8,885 (0.9)	18,384 (0.1)	82 (0.1)	101 (0.0)**	937 (0.3)	1,645 (0.0)*	7,866 (1.5)	16,638 (0.6)
Disability type (main)									
Physical: External bodily	780,540	780,540	-	39,605	-	253,426	-	487,509	-
functions	(4.1)	(78.7)		(37.6)		(72.3)		(90.9)	
Physical: Internal organs	65,438(0.3)	65,438(6.6)	-	4,255(4.0)	-	30,521(8.7)	-	30,662(5.7)	-
Mental: Psychological development disorder	97,591(0.5)	97,591(9.8)	-	57,344 (54.4)	-	33,782(9.6)	-	6,465(1.2)	-
Mental: Mental disease	48,916 (0.3)	48,916 (4.9)	-	4,199 (4.0)	-	32,784 (9.4)	-	11,933 (2.2)	-

Values are presented as number (%).

SD = standard deviation, Q = quintile (1Q: lowest)

Table 2. Age and sex adjusted case severity rate of coronavirus disease 2019 confirmed cases among people with and without disability by age group

Epidemic	Age group, yr		Non-disab	led			Disabl	ed		Risk	Risk ratio (95% CI)
period	_	Total	Severe/critical	CSR, per	aCSR, per	Total	Severe/critical	CSR, per	aCSR, per	difference	
			+ deaths	1,000	1,000		+ deaths	1,000	1,000		
Total	Total	18,043,268	32,424	1.8	2.1	992,485	14,135	14.2	6.4	4.3	3.05 (2.96-3.14)
	19-39	7,080,971	834	0.1	0.1	105,403	247	2.3	2.0	1.9	19.74 (17.04-22.87)
	40-64	8,132,919	6,041	0.7	0.8	350,513	2,198	6.3	5.2	4.4	6.82 (6.45-7.21)
	≥ 65	2,829,378	25,549	9.0	9.7	536,569	11,690	21.8	18.2	8.4	1.87 (1.83-1.91)
Pre-delta	Total	145,612	4,214	28.9	31.3	8,833	1,067	120.8	56.6	25.2	1.80 (1.63-2.00)
	19-39	52,257	131	2.5	2.5	881	13	14.8	14.6	12.1	5.84 (3.10-11.00)
	40-64	70,587	1,357	19.2	17.5	3,582	202	56.4	45.6	28.1	2.60 (2.14-3.16)
	≥ 65	22,768	2,726	119.7	127.7	4,370	852	195.0	173.8	46.1	1.36 (1.25-1.48)
Delta	Total	346,994	7,781	22.4	23.2	21,732	2,247	103.4	51.2	28.0	2.21 (2.05-2.39)
	19-39	127,020	444	3.5	3.4	1,760	31	17.6	17.0	13.7	5.05 (3.39-7.52)
	40-64	157,447	2,364	15.0	14	8,100	442	54.6	46.9	32.9	3.35 (2.94-3.81)
	≥ 65	62,527	4,973	79.5	88.5	11,872	1,774	149.4	135.0	46.4	1.52 (1.44-1.61)
Omicron	Total	17,550,662	20,429	1.2	1.4	961,920	10,821	11.2	5.0	3.6	3.55 (3.43-3.68)
	19-39	6,901,694	259	0.0*	0.0*	102,762	203	2.0	2.0	1.9	52.62 (43.59-63.54)
	40-64	7,904,885	2,320	0.3	0.3	12.68	11.80	13.64	3.8	3.5	12.68 (11.80-13.64)
	≥ 65	2,744,083	17,850	6.5	7.1	2.01	1.95	2.06	14.2	7.1	2.01 (1.95-2.06)

aCSR = age- and sex-standardized case severity rate, CI = confidence interval.

2.70 times higher than that in PwD. The aCSR and aCFR were highest in the old-age group, whereas the disparities between PwD and PwoD were highest in the young-age group. As the pandemic progressed, the aCSR and aCFR decreased, while the disparity between PwD and PwoD increased. Among 15 disability types, aCSR and aCFR were higher in people with renal, cardiac, and respiratory impairment, and brain lesion disorder (**Supplementary Table 1**).

^{*}P < 0.1, **P < 0.01.

^{*}P < 0.1.



Table 3. Age and sex adjusted case fatality rate of coronavirus disease 2019 confirmed cases among people with and without disability by age group

Epidemic	Age group,		Non-dis	sabled			Dis	abled		Risk	Risk ratio (95% CI)
period	yr	Total	Deaths	CFR, per 1,000	aCFR, per 1,000	Total	Deaths	CFR, per 1,000	aCFR, per 1,000	difference	
Total .	Total	18,043,268	18,384	1.0	1.2	992,485	8,885	9.0	3.3	2.1	2.70 (2.60-2.80)
	19-39	7,080,971	101	0.0a*	0.0*	105,403	82	0.8	0.8	0.8	55.52 (41.20-74.81)
	40-64	8,132,919	1,645	0.2	0.2	350,513	937	2.7	2.2	2.0	10.46 (9.55-11.45)
	≥ 65	2,829,378	16,638	5.9	6.4	536,569	7,866	14.7	11.6	5.2	1.81 (1.76-1.86)
Pre-delta	Total	145,612	1,461	10.0	12.3	8,833	552	62.5	24.6	12.3	2.01 (1.73-2.32)
	19-39	52,257	10	0.2	0.2	881	2	2.3	3.5	3.3	17.76 (3.88-81.26)
	40-64	70,587	138	2.0	1.7	3,582	66	18.4	15.6	13.9	9.07 (6.22-13.23)
	≥ 65	22,768	1,313	57.7	64.6	4,370	484	110.8	92.3	27.7	1.43 (1.28-1.59)
Delta	Total	346,994	3,116	9.0	10.1	21,732	1,283	59.0	23.6	13.5	2.34 (2.12-2.59)
	19-39	127,020	24	0.2	0.2	1,760	10	5.7	5.9	5.7	32.53 (14.78-71.59)
	40-64	157,447	398	2.5	2.2	8,100	157	19.4	15.3	13.1	6.81 (5.47-8.48)
	≥ 65	62,527	2,694	43.1	50.9	11,872	1,116	94.0	82.2	31.4	1.62 (1.50-1.74)
Omicron	Total	17,550,662	13,807	0.8	1.0	961,920	7,050	7.3	2.7	1.7	2.82 (2.70-2.94)
	19-39	6,901,694	67	0.0**	0.0**	102,762	70	0.7	0.7	0.7	70.52 (50.10-99.26)
	40-64	7,904,885	1,109	0.1	0.1	12.68	714	2.1	1.7	1.6	12.03 (10.82-13.38)
	≥ 65	2,744,083	12,631	4.6	5.0	2.01	6,266	12.0	9.4	4.4	1.87 (1.81-1.93)

aCFR = age- and sex-standardized case fatality rate, CI = confidence interval.

Risk disparities of severe outcomes with adjusting explanatory factors

After adjusting explanatory factors, the risk disparities of all severe outcomes and deaths between PwD and PwoD were decreased, but the risk remained higher in PwD with all severe outcomes (aOR, 1.49; CI, 1.46–1.52) and deaths (aOR, 1.40; CI, 1.36–1.44; **Tables 4** and 5). The risk of disability was highest in the young-age group (all severe outcomes, aOR, 8.02; CI, 6.64–9.68; deaths, aOR, 11.26; CI, 7.57–16.75) and lowest in the old-age group (all severe outcomes, aOR, 1.31; CI, 1.28–1.34; deaths, aOR, 1.27; CI, 1.24–1.31).

In the overall age model, older age was the most critical risk factor for all severe outcomes among the explanatory variables, followed by pulmonary (aOR, 3.44; CI, 3.37–3.52) and kidney (aOR, 1.71; CI, 1.66–1.75) diseases, and the virulence of virus (delta period, aOR, 2.02; CI, 1.94–2.11). These factors had a greater impact on all severe outcomes than disability status (aOR, 1.49; CI, 1.46–1.52). The pattern in the old age group (\geq 65 years) was consistent with that of the overall age model. However, the younger age groups showed different trends. In the 19–39 age group, disability status was the most significant risk factor, and in the 40–64 age group, pulmonary disease (aOR, 4.53; CI, 4.30–4.77) was the primary risk factor, with disability status (aOR, 2.35; CI, 2.20–2.50) being the second most significant. A similar pattern was observed in the death model, although in those aged 40–64, low-income level (Medicaid aOR, 5.84; CI, 4.71–7.24) showed a higher risk of deaths than disability status (aOR, 2.38; CI, 2.16–2.63).

The risk of severe outcomes differed by disability type (Tables 6-9, Supplementary Fig. 2). Among 15 disability types across all severe outcomes, epilepsy disorder (aOR, 3.45; CI, 2.31–5.17), renal (aOR, 2.97; CI, 2.79–3.15), and respiratory impairment (aOR, 2.88; CI, 2.47–3.36) exhibited a comparatively higher risk in all severe outcomes. Meanwhile, people with hearing impairment were at comparatively lower risk (aOR, 1.08; CI, 1.03–1.13), and those with facial disfigurement and autistic disorder had a low number of events (< 10), limiting statistical calculation. When considering deaths alone, people with hepatic impairment (aOR, 2.92; CI, 2.11–4.05), renal impairment (aOR, 2.44; CI, 2.25–2.63), and intellectual disabilities (aOR, 2.74; CI, 2.35–3.19) showed comparatively higher risk than other disability types. However,

^{*}P < 0.1, **P < 0.01.



Table 4. aOR of all severe outcomes among people with and without disability based on age group

Variables		Total	1	9-39		10-64	≥ 65		
	aORª	95% CI	aORª	95% CI	aORª	95% CI	aORª	95% CI	
Disabled									
Non-disabled	Ref.		Ref.		Ref.		Ref.		
Disabled	1.49	1.46-1.52	8.02	6.64-9.68	2.35	2.20-2.50	1.31	1.28-1.34	
Sex									
Male	Ref.		Ref.		Ref.		Ref.		
Female	0.57	0.56-0.58	0.72	0.64-0.82	0.49	0.47-0.52	0.60	0.58-0.61	
Age, yr									
19-29	Ref.		Ref.		-		-		
30-39	2.25	1.97-2.56	1.98	1.73-2.26	-		-		
40-49	4.54	4.03-5.11	-		Ref.		-		
50-59	10.93	9.76-12.24	-		1.98	1.86-2.11	-		
60-69	23.74	21.25-26.53	-		3.17	2.96-3.39	Ref.		
70-79	50.33	45.02-56.26	-		-		1.83	1.77-1.90	
≥ 80	115.80	103.61-129.43	-		-		4.23	4.08-4.39	
Income level									
Medical aid	1.48	1.43-1.54	1.49	1.10-2.01	2.07	1.87-2.29	1.24	1.18-1.29	
Self-employed 1st quintile (lowest)	1.34	1.28-1.40	1.15	0.85-1.56	1.86	1.67-2.08	1.19	1.13-1.25	
Self-employed 2nd quintile	1.32	1.26-1.39	1.20	0.90-1.60	1.70	1.52-1.91	1.18	1.12-1.25	
Self-employed 3rd quintile	1.26	1.20-1.33	1.24	0.91-1.68	1.64	1.46-1.84	1.13	1.06-1.19	
Self-employed 4th quintile	1.22	1.16-1.28	1.25	0.91-1.71	1.63	1.45-1.82	1.08	1.01-1.14	
Self-employed 5th quintile (highest)	1.05	1.00-1.10	1.32	0.92-1.88	1.39	1.23-1.56	0.94	0.89-0.99	
Employee 1st quintile (lowest)	1.15	1.11-1.20	1.06	0.82-1.36	1.37	1.24-1.51	1.07	1.02-1.12	
Employee 2nd quintile	1.08	1.04-1.13	0.93	0.72-1.20	1.13	1.02-1.26	1.06	1.02-1.11	
Employee 3rd quintile	1.01	0.97-1.06	0.90	0.70-1.16	1.05	0.95-1.18	1.00	0.95-1.05	
Employee 4th quintile	1.01	0.96-1.05	0.89	0.69-1.15	0.99	0.89-1.11	1.00	0.95-1.04	
Employee 5th quintile (highest)	Ref.	0.00 1.00	Ref.	0.00 1.10	Ref.	0.00 1.11	Ref.	0.00 1.01	
Comorbidities	noi.		itter.		no.		itoi.		
Diabetes	1.39	1.36-1.42	2.35	1.89-2.94	1.68	1.59-1.77	1.30	1.27-1.33	
Hypertension	1.07	1.04-1.09	1.65	1.33-2.04	1.19	1.13-1.25	0.99	0.96-1.01	
Kidney disease	1.71	1.66-1.75	1.72	1.32-2.24	2.52	2.35-2.70	1.60	1.55-1.64	
Pulmonary disease	3.44	3.37-3.52	5.28	4.55-6.12	4.53	4.30-4.77	3.15	3.08-3.22	
Cardiovascular disease	1.21	1.19-1.24	1.97	1.60-2.41	1.40	1.33-1.48	1.15	1.12-1.18	
Liver disease	0.94	0.92-0.97	1.49	1.25-1.79	1.06	1.00-1.12	0.88	0.85-0.91	
Neurologic disease	1.13	1.11-1.16	1.43	1.53-2.14	1.28	1.21-1.35	1.06	1.04-1.09	
Cancer	1.49	1.45-1.53	2.76	2.10-3.63	2.36	2.21-2.51	1.33	1.29-1.37	
Others	1.78	1.74-1.82	1.39	1.19-1.62	1.47	1.40-1.55	1.76	1.71-1.80	
Vaccination	1.70	1.74-1.02	1.39	1.19-1.02	1.47	1.40-1.55	1.70	1.71-1.80	
Unvaccinated	Ref.		Ref.		Ref.		Ref.		
1 or 2 doses	0.25	0.94.0.96	0.10	0.00.010	0.14	0.12.0.15		0.2F 0.27	
1 or 2 doses ≥ 3 doses	0.25	0.24-0.26 0.10-0.10	0.10	0.08-0.12 0.06-0.10	0.14	0.13-0.15 0.06-0.07	0.36 0.11	0.35-0.37 0.11-0.11	
≥ 3 doses Reinfection	0.10	0.10-0.10	0.08	0.06-0.10	0.07	0.06-0.07	0.11	0.11-0.11	
	Ref.		Dof		Dof		Dof		
None		0.00 0.70	Ref.	0.40 1.07	Ref.	0.74 1.04	Ref.	0.55.0.05	
Reinfected	0.65	0.60-0.70	0.78	0.48-1.27	0.88	0.74-1.04	0.60	0.55-0.65	
Period	D-f		D-f		D-f		D-f		
Pre-delta	Ref.	104011	Ref.	0.05.0.46	Ref.	0.00.0.55	Ref.		
Delta	2.02	1.94-2.11	2.85	2.35-3.46	2.38	2.22-2.55	1.62	1.54-1.70	
Omicron	0.26	0.25-0.27	0.19	0.15-0.23	0.22	0.20-0.23	0.30	0.28-0.31	

aOR = adjusted odds ratio, CI = confidence interval.

people with hearing impairment showed no significant disparity with PwoD (aOR, 1.02; CI, 0.97–1.08), and those with facial disfigurement, epilepsy disorder, and autistic disorder had a low number of events (< 10), limiting statistical calculation.

People aged \geq 65 years and PwD involving internal organs, including hepatic impairment (aOR, 2.11; CI, 1.34–3.32) and renal impairment (aOR, 2.04; CI, 1.87–2.23), demonstrated a higher risk disparity of death than other types of PwD. The 40–64-year age group showed a higher risk for

 $^{^{\}rm a}\textsc{Each}$ aOR is adjusted for all other variables listed in the table within each age stratum.



Table 5. aOR of deaths among people with and without disability based on age group

Variables		Total		–39 yr		0-64 yr	≥ 65 yr	
	aORª	95% CI	aORª	95% CI	aORª	95% CI	aORª	95% CI
Disabled								
Non-disabled	Ref.		Ref.		Ref.		Ref.	
Disabled	1.40	1.36-1.44	11.26	7.57-16.75	2.38	2.16-2.63	1.27	1.24-1.31
Sex								
Male	Ref.		Ref.		Ref.		Ref.	
Female	0.62	0.61-0.64	0.67	0.49-0.91	0.45	0.42-0.50	0.65	0.63-0.66
Age, yr								
19-29	Ref.		Ref.		-		-	
30-39	1.89	1.39-2.57	1.35	0.98-1.86	-		-	
40-49	5.50	4.19-7.20	-		Ref.		-	
50-59	20.11	15.56-25.98	-		2.52	2.22-2.85	-	
60-69	55.70	43.28-71.70	_		3.92	3.44-4.46	Ref.	
70-79	162.97	126.68-209.67	_		_		2.36	2.24-2.50
≥ 80	480.99	374.01-618.56	_		_		7.10	6.74-7.47
Income level		3752 020.00						3 7.17
Medical aid	1.74	1.66-1.83	4.72	2.11-10.58	5.84	4.71-7.24	1.43	1.36-1.51
Self-employed 1st quintile (lowest)	1.44	1.36-1.52	4.03	1.73-9.41	4.59	3.63-5.80	1.28	1.21-1.36
Self-employed 2nd quintile	1.41	1.32-1.50	2.89	1.17-7.12	3.76	2.94-4.81	1.27	1.19-1.36
Self-employed 3rd quintile	1.29	1.21-1.38	5.43	2.36-12.51	3.19	2.48-4.10	1.17	1.19-1.36
Self-employed 4th quintile	1.21	1.13-1.29	4.78 a	2.00-11.43 a	3.25	2.53-4.16	1.09	1.01-1.17
Self-employed 5th quintile (highest)	0.94	0.88-1.00			2.21	1.70-2.88	0.86	0.81-0.92
Employee 1st quintile (lowest)	1.26	1.20-1.33	2.20	0.97-5.01	2.63	2.10-3.29	1.19	1.13-1.25
Employee 2nd quintile	1.20	1.14-1.27	1.57	0.67-3.68	2.11	1.67-2.67	1.17	1.10-1.24
Employee 3rd quintile	1.12	1.06-1.19	1.69	0.73-3.92	1.94	1.52-2.48	1.09	1.03-1.16
Employee 4th quintile	1.07	1.01-1.13	1.15	0.46-2.86	1.70	1.32-2.18	1.04	0.98-1.10
Employee 5th quintile (highest)	Ref.		Ref.		Ref.		Ref.	
Comorbidities								
Diabetes	1.30	1.26-1.33	2.93	1.89-4.54	1.69	1.54-1.84	1.24	1.21-1.27
Hypertension	0.98	0.95-1.01	0.81	0.50-1.33	1.00	0.92-1.09	0.94	0.92-0.97
Kidney disease	1.62	1.57-1.68	2.06	1.26-3.38	2.25	2.02-2.51	1.57	1.52-1.63
Pulmonary disease	2.86	2.79-2.94	2.85	1.96-4.14	4.09	3.73-4.47	2.73	2.65-2.80
Cardiovascular disease	1.20	1.17-1.23	3.33	2.28-4.88	1.60	1.46-1.75	1.14	1.11-1.17
Liver disease	0.91	0.88-0.95	1.41	0.95-2.12	1.16	1.06-1.28	0.86	0.83-0.89
Neurologic disease	1.20	1.17-1.23	2.16	1.50-3.10	1.41	1.29-1.54	1.03	1.00-1.06
Cancer	0.91	0.88-0.95	6.59	4.30-10.10	4.34	3.96-4.75	1.39	1.34-1.45
Others	2.16	2.10-2.23	2.22	1.56-3.16	1.85	1.69-2.02	2.07	2.00-2.13
Vaccination								
Unvaccinated	Ref.		Ref.		Ref.		Ref.	
1 or 2 doses	0.35	0.34-0.36	0.14	0.09-0.21	0.24	0.22-0.27	0.59	0.54-0.63
≥ 3 doses	0.11	0.11-0.11	0.09	0.06-0.13	0.09	0.08-0.10	0.37	0.35-0.38
Reinfection								
None	Ref.		Ref.		Ref.		Ref.	
Reinfected	0.56	0.51-0.62	b	b	0.85	0.66-1.09	0.54	0.49-0.60
Period							***	21.12 21.30
Pre-delta	Ref.		Ref.		Ref.		Ref.	
Delta	1.94	1.83-2.07	2.38	1.20-4.72	2.84	2.40-3.37	1.78	1.67-1.90
Omicron	0.45	0.43-0.48	0.62	0.33-1.17	0.67	0.57-0.79	0.43	0.41-0.46
aOR = adjusted odds ratio, CI = confidence		0.70 0.70	0.02	0.00 1.17	0.07	3.37 0.73	0.43	J. 71 U. 70

aOR = adjusted odds ratio, CI = confidence interval.

PwD related to brain lesion disorder (aOR, 3.68; CI, 3.15–4.31) of external bodily function, intellectual disability (aOR, 3.22; CI, 2.53–4.11) of psychological development disorder and renal impairment (aOR, 3.11; CI, 2.56–3.77) of internal organs in COVID-19-related deaths. Among the 19–39-year age group, most types of disability had a low number of events, making statistical calculation challenging. However, a higher risk of death was observed among people with brain lesion disorder disabilities (aOR, 19.58; CI, 10.99–34.87) and

^aEach aOR is adjusted for all other variables listed in the table within each age stratum.

^baOR values were not presented due to insufficient event number (less than 10).



Table 6. Overall aORs of severe outcomes and deaths based on 15 disability types for all ages

Disability type	No.	Severe cases + deaths, %	aORª	95% CI	Deaths, %	aOR	95% CI
Without disabilities	18,043,268	32,424 (0.18)	Ref.		18,384 (0.10)	Ref.	
Physical: External bodily functions							
Physical disabilities	422,632	4,138 (0.98)	1.21	1.17-1.26	2,648 (0.63)	1.17	1.12-1.22
Brain lesion disorder	105,227	3,399 (3.23)	2.13	2.04-2.22	2,212 (2.10)	2.06	1.96-2.16
Visual impairment	90,565	1,009 (1.11)	1.26	1.18-1.35	660 (0.73)	1.22	1.12-1.33
Hearing impairment	154,253	2,481 (1.61)	1.08	1.03-1.13	1,694 (1.10)	1.02	0.97-1.08
Language disabilities	6,899	99 (1.43)	1.34	1.08-1.67	62 (0.90)	1.33	1.01-1.74
Facial disfigurement	964	6 (0.62)	b	b	3 (0.31)	b	b
Physical: Internal organs							
Renal impairment	44,302	1,781 (4.02)	2.97	2.79-3.15	943 (2.13)	2.44	2.25-2.63
Cardiac impairment	2,465	68 (2.76)	2.66	2.02-3.49	33 (1.34)	1.92	1.33-2.79
Respiratory impairment	4,141	224 (5.41)	2.88	2.47-3.36	89 (2.15)	1.70	1.35-2.13
Hepatic impairment	6,010	79 (1.31)	2.72	2.14-3.44	39 (0.65)	2.92	2.11-4.05
Intestinal/Urinary fistula	5,657	111 (1.96)	1.29	1.05-1.59	81 (1.43)	1.37	1.09-1.74
Epilepsy disorder	2,863	27 (0.94)	3.45	2.31-5.17	9 (0.31)	b	b
Mental: Psychological development disorder							
Intellectual disabilities	89,126	358 (0.40)	2.67	2.38-2.99	193 (0.22)	2.74	2.35-3.19
Autistic disorder	8,465	5 (0.06)	b	b	2 (0.02)	b	b
Mental: Mental disease							
Mental disabilities	48,916	350 (0.72)	1.43	1.27-1.60	217 (0.44)	1.70	1.48-1.96

aOR = adjusted odds ratio, CI = confidence interval.

Table 7. aORs of severe outcomes and deaths based on 15 disability types in age group of 19-39 years

Disability type	No.	Severe cases + deaths, %	aORª	95% CI	Deaths, %	aOR	95% CI
Without disabilities	7,080,971	834 (0.01)	Ref		101 (0.00)**	Ref	
Physical: External bodily functions							
Physical disabilities	18,243	40 (0.22)	8.26	5.87-11.62	10 (0.05)	11.07	5.58-21.99
Brain lesion disorder	7,295	57 (0.78)	14.74	10.55-20.59	24 (0.33)	19.58	10.99-34.87
Visual impairment	7,766	7 (0.09)	b	b	1 (0.01)	b	b
Hearing impairment	5,408	3 (0.06)	b	b	2 (0.04)	b	b
Language disabilities	723	0 (0.00)	-	-	0 (0.00)	-	-
Facial disfigurement	170	0 (0.00)	-	-	0 (0.00)	-	-
Physical: Internal organs							
Renal impairment	2,546	26 (1.02)	4.05	2.38-6.89	7 (0.27)		
Cardiac impairment	487	4 (0.82)	b	b	2 (0.41)	b	b
Respiratory impairment	84	2 (2.38)	b	b	0 (0.00)	-	-
Hepatic impairment	344	4 (1.16)	b	b	3 (0.87)	b	b
Intestinal/Urinary fistula	201	2 (1.00)	b	b	1 (0.50)	b	b
Epilepsy disorder	593	6 (1.01)	b	b	2 (0.34)	b	b
Mental: Psychological development disorder							
Intellectual disabilities	49,158	87 (0.18)	9.09	6.92-11.95	27 (0.05)	12.20	7.19-20.68
Autistic disorder	8,186	5 (0.06)	b	b	2 (0.02)	b	b
Mental: Mental disease							
Mental disabilities	4,199	4 (0.10)	b	b	1 (0.02)	b	b

aOR = adjusted odds ratio, CI = confidence interval.

intellectual disabilities (aOR, 12.20; CI, 7.19–20.68). In all age groups, people with language disabilities and facial disfigurement showed no significant disparities or limited number of events in risk of death.

Regarding the epidemic periods, the risk disparity between PwD and PwoD for all severe outcomes in the omicron period increased in all age groups, particularly in the young-age group (Fig. 1). For deaths only, risk disparities were similar across periods.

^aaOR for disability status is adjusted for all variables included in **Table 3**.

^baOR values are not presented due to insufficient events (less than 10).

^aaOR for disability status is adjusted for all variables included in **Table 3**.

^baOR values are not presented due to insufficient events (less than 10).

^{**}P < 0.01.



Table 8. aORs of severe outcomes and deaths based on 15 disability types in age group of 40-64 years

Disability type	No.	Severe cases + deaths, %	aOR	95% CI	Deaths, %	aOR	E95% CI
Without disabilities	8,132,919	6,041 (0.07)	Ref.		1,645 (0.02)	Ref.	
Physical: External bodily functions							
Physical disabilities	159,550	480 (0.30)	1.65	1.49-1.82	201 (0.13)	1.90	1.63-2.22
Brain lesion disorder	31,530	532 (1.69)	3.62	3.24-4.04	255 (0.81)	3.68	3.15-4.31
Visual impairment	33,476	95 (0.28)	1.59	1.28-1.98	37 (0.11)	1.61	1.14-2.27
Hearing impairment	25,367	71 (0.28)	1.56	1.22-2.01	29 (0.11)	1.79	1.22-2.61
Language disabilities	2,987	19 (0.64)	1.97	1.17-3.30	8 (0.27)	b	b
Facial disfigurement	516	1 (0.19)	b	b	0 (0.00)	-	-
Physical: Internal organs							
Renal impairment	21,925	579 (2.64)	4.47	3.94-5.07	213 (0.97)	3.11	2.56-3.77
Cardiac impairment	785	23 (2.93)	7.05	4.37-11.38	7 (0.89)	b	b
Respiratory impairment	1,210	39 (3.22)	4.84	3.39-6.92	9 (0.74)	b	b
Hepatic impairment	3,366	37 (1.10)	2.45	1.71-3.52	16 (0.48)	2.08	1.24-3.49
Intestinal/Urinary fistula	1,408	7 (0.50)	b	b	4 (0.28)	b	b
Epilepsy disorder	1,827	16 (0.88)	3.87	2.23-6.70	4 (0.22)	b	b
Mental: Psychological development disorder							
Intellectual disabilities	33,505	152 (0.45)	2.81	2.35-3.37	80 (0.24)	3.22	2.53-4.11
Autistic disorder	277	0 (0.00)	-	-	0 (0.00)	-	-
Mental: Mental disease							
Mental disabilities	32,784	147 (0.45)	1.44	1.19-1.73	74 (0.23)	1.69	1.31-2.18

aOR = adjusted odds ratio, CI = confidence interval.

Table 9. aORs of severe outcomes and deaths based on 15 disability types in age group ≥ 65 years

Disability type	No.	Severe cases + deaths, %	aOR ^a	95% CI	Deaths, %	aOR	95% CI
Without disabilities	2,829,378	25,549 (0.90)	Ref.		16,638 (0.59)	Ref.	
Physical: External bodily functions							
Physical disabilities	244,839	3,618 (1.48)	1.12	1.08-1.16	2,437 (1.00)	1.11	1.06-1.16
Brain lesion disorder	66,402	2,810 (4.23)	1.85	1.77-1.93	1,933 (2.91)	1.85	1.76-1.95
Visual impairment	49,323	907 (1.84)	1.19	1.11-1.28	622 (1.26)	1.18	1.08-1.28
Hearing impairment	123,478	2,407 (1.95)	1.04	0.99-1.09	1,663 (1.35)	1.00	0.95-1.05
Language disabilities	3,189	80 (2.51)	1.22	0.96-1.55	54 (1.69)	1.25	0.94-1.67
Facial disfigurement	278	5 (1.80)	b	b	3 (1.08)	b	b
Physical: Internal organs							
Renal impairment	19,831	1,176 (5.93)	2.24	2.09-2.41	723 (3.65)	2.04	1.87-2.23
Cardiac impairment	1,193	41 (3.44)	1.69	1.20-2.38	24 (2.01)	1.48	0.96-2.28
Respiratory impairment	2,847	183 (6.43)	2.48	2.10-2.93	80 (2.81)	1.61	1.27-2.04
Hepatic impairment	2,300	38 (1.65)	2.07	1.48-2.90	20 (0.87)	2.11	1.34-3.32
Intestinal/Urinary fistula	4,048	102 (2.52)	1.26	1.02-1.55	76 (1.88)	1.36	1.07-1.73
Epilepsy disorder	443	5 (1.13)	b	b	3 (0.68)	b	b
Mental: Psychological development disorder							
Intellectual disabilities	6,463	119 (1.84)	1.47	1.21-1.78	86 (1.33)	1.65	1.32-2.07
Autistic disorder	2	0 (0.00)	-	-	0 (0.00)	-	-
Mental: Mental disease							
Mental disabilities	11,933	199 (1.67)	1.37	1.18-1.59	142 (1.19)	1.60	1.34-1.90

aOR = adjusted odds ratio, CI = confidence interval.

DISCUSSION

This nationwide retrospective cohort study, encompassing all adult confirmed cases of COVID-19 until October 2022, revealed that PwD were consistently at higher risk of severe COVID-19 outcomes than PwoD. Although the risk disparities between PwD and PwoD decreased after adjusting for explanatory factors, the risk remained higher for PwD. The risk of severe outcomes was highest in the old-age group, in both PwD and PwoD, whereas that

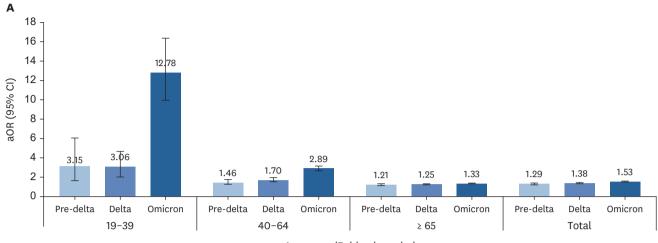
^aaOR for disability status is adjusted for all variables included in Table 3.

^baOR values are not presented due to insufficient events (less than 10).

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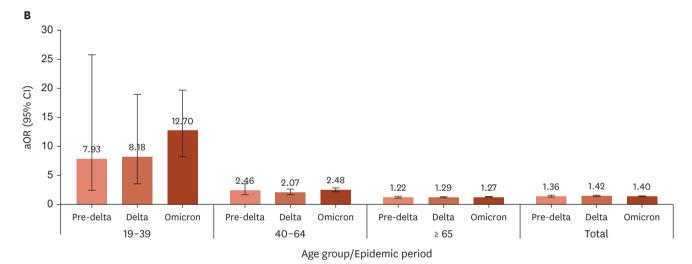


Fig. 1. Adjusted odds ratios of severe outcomes and deaths among people with and without disability based on age groups and circulating variant period, all severe outcomes (A), deaths (B).
aOR = adjusted odds ratio, CI = confidence interval.

adjusted edge ratio, or community interval.

of disparity was more marked in the young-age groups. Among 15 disability types, people with hepatic or renal impairment, intellectual disability, and brain lesion disorder showed a comparatively higher risk, while people with hearing impairment, facial disfigurement, epilepsy, and autistic disorder showed no significant disparities or a low number of events, limiting statistical calculation. As COVID-19 severity decreased in the omicron period, the risk of severe outcomes also declined in both PwD and PwoD. However, the risk disparity of all severe outcomes increased slightly in the young-age group.

This study revealed that the risk disparities between PwD and PwoD decreased after adjusting for explanatory factors associated with severe outcomes of COVID-19. This finding suggests that health vulnerability is associated with a higher prevalence of comorbidities and low socioeconomic status, which are recognized risk factors for severe COVID-19.²¹ In the real world, PwD often experiences multiple risk factors, potentially leading to higher risk and greater disparity with PwoD.¹ Moreover, barriers to accessing healthcare services, as suggested by the WHO, could contribute to an increased risk of severe COVID-19.²²⁻²⁵



During the pandemic, a Korean study reported that PwD experienced larger reductions in claims for medical services than PwoD did.²⁶ These collateral impacts worsened pre-existing health conditions and could have led to more severe outcomes following COVID-19.²⁷ A delay in COVID-19 diagnosis due to disability could exacerbate its progression in people with dementia, Down syndrome, cerebral palsy, and spinal cord injury; their disability could have obscured respiratory symptoms. People with intellectual disabilities could experience difficulty in recognizing or expressing their symptoms, and people with impaired mobility could face challenges in accessing diagnostic facilities.²⁸

Older age remains the most critical risk factor for severe COVID-19 outcomes, consistent with both our findings and previous studies, 20 with comorbidities also being well-established risk factors. However, our analysis highlights that, among adults younger than 64 years of age, disability status posed as a significant risk factor comparable to that of comorbidities, although the impact varied depending on disability type. This finding was observed in overall types of disability and IDD studies in several countries. 2 , 6 , 13 , 29 , 30 Herein, youngage and middle-age groups (19–64 years) showed greater differences in socioeconomic status and comorbidities by disability status compared to the old-age group, which could have contributed to disparities in severe outcomes. A WHO survey indicated higher unmet care needs among PwD aged 50–59 compared to those aged \geq 60 years, with significant differences in healthcare service needs between PwD and PwoD. 31 Disruptions in healthcare services due to COVID-19 could have greater impact on younger PwD than on PwoD. 32

Age-specific and disability-type analyses provided detailed insights into risk disparities, particularly in young adults. Overall, higher risks of COVID-19 deaths were observed for most internal organ and intellectual disabilities compared to other disabilities. The old-age group showed a higher risk for most internal organ disabilities. However, individuals aged 19-64 years demonstrated elevated risks for both internal organs and intellectual disabilities, and brain lesions. The latter two disabilities may account for greater differences in risks for severe outcomes observed in young-age groups by disabilities. In the US study with overall age, IDD showed the highest risk for invasive mechanical ventilation and in-hospital deaths among PwD with mobility, vision, and hearing impairment, compared with PwoD.9 Intellectual disabilities are more prevalent in young adults, who have more health risk factors and more healthcare needs than PwoD. Additionally, they show a lower average age at death, indicating that agerelated impacts on severe COVID-19 outcomes may apply earlier.^{8,30,33-35} Combined factors of COVID-19 severe outcomes, such as age, comorbidities, and healthcare disruption during the COVID-19 pandemic likely had a disproportionately greater impact on people with intellectual disabilities exacerbating health disparities. 12,21,23,26,27 For people with brain-lesion disorders, respiratory dysfunction, which contributes significantly to morbidity, mortality, high demand for medical intervention, and regular follow-up, can lead to severe COVID-19 outcomes. 36,37 Meanwhile, people with facial disfigurement, epilepsy disorder, and autistic disorder had fewer deaths, limiting statistical analysis. These findings highlight that not all disability types pose the same risk of severe COVID-19 outcomes. Therefore, targeted protective measures should be prioritized for PwD, including those with intellectual disabilities, brain lesions, physical disabilities, and renal impairment, particularly in younger populations.

This study provides valuable insights into public health. First, prioritizing PwD in the public-health crisis response is crucial, alongside well-known priorities for older adults and individuals with comorbidities. PwD of internal organs may already be prioritized due to their underlying diseases. However, PwD at high risk of severe outcomes who do not meet these



criteria, such as people with intellectual disability are suggested to be included in the priority group. Second, understanding health risks of PwD through consistent monitoring is essential for policy development. The lack of regular data collection on PwD hinders risk assessment and identification of risk factors associated with health outcomes in this vulnerable group.³⁸ During crises, when timely and labor-intensive data collection is challenging, linking previously collected data, including sociodemographic information, with new data can help identify PwD rapidly. Including disability variables in regularly monitored basic tables, alongside age and sex, can raise awareness of the need to protect the health of vulnerable groups.

The strength of this study is its nationwide scope, covering all registered PwD with COVID-19. It delineated risk disparities by age, disability types, and epidemic period, adjusting factors beyond variations in baseline health status. Nonetheless, this study had some limitations. First, it only included formally registered PwD; unregistered PwD could misclassified as PwoD. However, with 94.1% PwD registered nationwide based on 2017 statistics,³⁹ the risk of misclassification was minimized. Additional considerations also include potential discrepancies with other disability statistics due to differences in the timing of disability information, as this analysis used health insurance data. Second, severity information relied on hospital reports, possibly missing severe cases due to delayed or omitted reporting. This underestimation likely affected both PwD and PwoD groups similarly. Future studies should review medical history to better define severity and minimize underestimation.

During the COVID-19 pandemic, PwD faced higher risk of severe COVID-19 outcomes compared to PwoD. Including PwD in high-risk groups alongside older adults could minimize health impact from public health crises. Risk-based prioritization of vulnerable groups, such as young PwD, people with internal organ and intellectual disability, helps mitigate disparities in risk between PwD and PwoD, considering their health vulnerability and demand of healthcare services by disability types.

SUPPLEMENTARY MATERIALS

Supplementary Table 1

Age- and sex-adjusted case severity rates and case fatality rates of COVID-19 confirmed cases among people with and without 15 types of disabilities

Supplementary Fig. 1

Flowchart of the selection of the study population.

Supplementary Fig. 2

aORs of severe outcomes and deaths based on age groups and four type of disabilities, all severe outcomes (A), deaths (B).

REFERENCES

- World Health Organization. Global report on health equity for persons with disabilities p.312. https:// www.who.int/news-room/questions-and-answers/item/global-report-on-health-equity-for-persons-withdisabilities. Updated 2022. Accessed April 10, 2023.
- 2. Bosworth ML, Ayoubkhani D, Nafilyan V, Foubert J, Glickman M, Davey C, et al. Deaths involving COVID-19 by self-reported disability status during the first two waves of the COVID-19 pandemic in



- England: a retrospective, population-based cohort study. *Lancet Public Health* 2021;6(11):e817-25. **PUBMED | CROSSREF**
- 3. Choi JW, Han E, Lee SG, Shin J, Kim TH. Risk of COVID-19 and major adverse clinical outcomes among people with disabilities in South Korea. *Disabil Health J* 2021;14(4):101127. PUBMED | CROSSREF
- 4. Yuan Y, Thierry JM, Bull-Otterson L, Yeargin-Allsopp M, Clark KE, Rice C, et al. COVID-19 cases and hospitalizations among Medicare beneficiaries with and without disabilities United States, January 1, 2020-November 20, 2021. MMWR Morb Mortal Wkly Rep 2022;71(24):791-6. PUBMED | CROSSREF
- Landes SD, Turk MA, Wong AW. COVID-19 outcomes among people with intellectual and developmental disability in California: the importance of type of residence and skilled nursing care needs. *Disabil Health J* 2021;14(2):101051. PUBMED | CROSSREF
- Turk MA, Landes SD, Formica MK, Goss KD. Intellectual and developmental disability and COVID-19 case-fatality trends: TriNetX analysis. *Disabil Health J* 2020;13(3):100942. PUBMED | CROSSREF
- 7. Koks-Leensen MC, Schalk BW, Bakker-van Gijssel EJ, Timen A, Nägele ME, van den Bemd M, et al. Risk for severe COVID-19 outcomes among persons with intellectual disabilities, the Netherlands. *Emerg Infect Dis* 2023;29(1):118-26. **PUBMED | CROSSREF**
- 8. Bahk J, Kang HY, Khang YH. Disability type-specific mortality patterns and life expectancy among disabled people in South Korea using 10-year combined data between 2008 and 2017. *Prev Med Rep* 2022;29:101958. PUBMED | CROSSREF
- 9. Clarke KE, Hong K, Schoonveld M, Greenspan AI, Montgomery M, Thierry JM. Severity of coronavirus disease 2019 hospitalization outcomes and patient disposition differ by disability status and disability type. *Clin Infect Dis* 2023;76(5):871-80. PUBMED | CROSSREF
- Okoro CA, Hollis ND, Cyrus AC, Griffin-Blake S. Prevalence of disabilities and health care access by disability status and type among adults - United States, 2016. MMWR Morb Mortal Wkly Rep 2018;67(32):882-7. PUBMED | CROSSREF
- 11. Kirk-Wade E, Stiebahl S, Wong H. UK Disability Statistics: Prevalence and Life Experiences. London, UK: House of Commons Library; 2024.
- 12. Molani S, Hernandez PV, Roper RT, Duvvuri VR, Baumgartner AM, Goldman JD, et al. Risk factors for severe COVID-19 differ by age for hospitalized adults. *Sci Rep* 2022;12(1):6568. PUBMED | CROSSREF
- 13. Brown HK, Saha S, Chan TC, Cheung AM, Fralick M, Ghassemi M, et al. Outcomes in patients with and without disability admitted to hospital with COVID-19: a retrospective cohort study. *CMAJ* 2022;194(4):E112-21. PUBMED | CROSSREF
- 14. Henderson A, Fleming M, Cooper SA, Pell JP, Melville C, Mackay DF, et al. COVID-19 infection and outcomes in a population-based cohort of 17 203 adults with intellectual disabilities compared with the general population. *J Epidemiol Community Health* 2022;76(6):550-5. PUBMED | CROSSREF
- World Health Organization. Tracking SARS-CoV-2 variants. https://www.who.int/activities/tracking-SARS-CoV-2-variants/. Updated 2021. Accessed October 3, 2022.
- Korea Disease Control and Prevention Agency. Infectious Disease Control and Prevention Act. Cheongju, Korea: Korea Disease Control and Prevention Agency; 2022.
- Seong SC, Kim YY, Khang YH, Park JH, Kang HJ, Lee H, et al. Data resource profile: the national health information database of the National Health Insurance Service in South Korea. Int J Epidemiol 2017;46(3):799-800. PUBMED | CROSSREF
- World Health Organization. WHO R&D blueprint novel coronavirus COVID-19 therapeutic trial synopsis. https://www.who.int/docs/default-source/blue-print/covid-19-therapeutic-trial-synopsis.pdf. Updated 2020. Accessed April 10, 2023.
- 19. Ryu B, Shin E, Kim NY, Kim DH, Lee HJ, Kim A, et al. Severity of COVID-19 associated with SARS-CoV-2 variants circulating in the Republic of Korea. *Public Health Weekly Report* 2022;15(47):2873-95. CROSSREF
- 20. Ministry of Health and Welfare (KR). Act on the Welfare of Persons with Disabilities. Sejong, Korea: Ministry of Health and Welfare; 2021.
- 21. Williamson EJ, Walker AJ, Bhaskaran K, Bacon S, Bates C, Morton CE, et al. Factors associated with COVID-19-related death using OpenSAFELY. *Nature* 2020;584(7821):430-6. PUBMED | CROSSREF
- Das-Munshi J, Chang CK, Bakolis I, Broadbent M, Dregan A, Hotopf M, et al. All-cause and cause-specific
 mortality in people with mental disorders and intellectual disabilities, before and during the COVID-19
 pandemic: cohort study. Lancet Reg Health Eur 2021;11:100228. PUBMED | CROSSREF
- 23. Bigdelou B, Sepand MR, Najafikhoshnoo S, Negrete JA, Sharaf M, Ho JQ, et al. COVID-19 and preexisting comorbidities: risks, synergies, and clinical outcomes. *Front Immunol* 2022;13:890517. PUBMED | CROSSREF
- 24. World Health Organization. Disability considerations during the COVID-19 outbreak. https://www.who.int/publications/i/item/WHO-2019-nCoV-Disability-2020-1. Updated 2020. Accessed April 10, 2023.



- Centers for Disease Control and Prevention. Underlying medical conditions associated with a higher risk for severe COVID-19: information for healthcare professionals. https://www.cdc.gov/coronavirus/2019ncov/hcp/clinical-care/underlyingconditions.html. Updated 2023. Accessed May 18, 2023.
- Sohn M, Koo H, Choi H, Cho H, Han E. Collateral impact of the COVID 19 pandemic on the use of healthcare resources among people with disabilities. Front Public Health 2022;10:922043. PUBMED | CROSSREF
- 27. McBride-Henry K, Nazari Orakani S, Good G, Roguski M, Officer TN. Disabled people's experiences accessing healthcare services during the COVID-19 pandemic: a scoping review. *BMC Health Serv Res* 2023;23(1):346. PUBMED | CROSSREF
- 28. Kamalakannan S, Bhattacharjya S, Bogdanova Y, Papadimitriou C, Arango-Lasprilla JC, Bentley J, et al. Health risks and consequences of a COVID-19 infection for people with disabilities: scoping review and descriptive thematic analysis. *Int J Environ Res Public Health* 2021;18(8):4348. PUBMED | CROSSREF
- 29. Deal JA, Jiang K, Betz JF, Clemens GD, Zhu J, Reed NS, et al. COVID-19 clinical outcomes by patient disability status: a retrospective cohort study. *Disabil Health J* 2023;16(2):101441. PUBMED | CROSSREF
- 30. Cuypers M, Koks-Leensen MC, Schalk BW, Bakker-van Gijssel EJ, Leusink GL, Naaldenberg J. All-cause and cause-specific mortality among people with and without intellectual disabilities during the COVID-19 pandemic in the Netherlands: a population-based cohort study. *Lancet Public Health* 2023;8(5):e356-63.

 PUBMED | CROSSREF
- World Health Organization, World Bank. World report on disability 2011. https://www.who.int/teams/ noncommunicable-diseases/sensory-functions-disability-and-rehabilitation/world-report-on-disability. Updated 2011. Accessed April 10, 2023.
- Moynihan R, Sanders S, Michaleff ZA, Scott AM, Clark J, To EJ, et al. Impact of COVID-19 pandemic on utilisation of healthcare services: a systematic review. BMJ Open 2021;11(3):e045343. PUBMED | CROSSREF
- Australian Institute of Health and Welfare. Disability and ageing Australian population patterns and implications. https://www.aihw.gov.au/reports/disability/disability-and-ageing-australian-population/ summary. Updated 2000. Accessed April 10, 2023.
- Landes SD, Stevens JD, Turk MA. Heterogeneity in age at death for adults with developmental disability. J Intellect Disabil Res 2019;63(12):1482-7. PUBMED | CROSSREF
- National Rehabilitation Center Research Institute. 2020 Healthcare Statistics of Persons with Disabilities. Seoul, Korea: National Rehabilitation Center Research Institute; 2022.
- Brandenburg JE, Fogarty MJ, Sieck GC. Why individuals with cerebral palsy are at higher risk for respiratory complications from COVID-19. J Pediatr Rehabil Med 2020;13(3):317-27. PUBMED | CROSSREF
- 37. Kim Y, Eun SJ, Kim WH, Lee BS, Leigh JH, Kim JE, et al. A new disability-related health care needs assessment tool for persons with brain disorders. *J Prev Med Public Health* 2013;46(5):282-90. PUBMED | CROSSREF
- 38. Akobirshoev I, Vetter M, Iezzoni LI, Rao SR, Mitra M. Delayed medical care and unmet care needs due to the COVID-19 pandemic among adults with disabilities in the US. *Health Aff (Millwood)* 2022;41(10):1505-12. PUBMED | CROSSREF
- 39. Ministry of Health and Welfare, Korea Institute for Health and Social Affairs. The National survey on persons with disabilities. Sejong, Korea: Ministry of Health and Welfare; 2017.