THE LANCET Child & Adolescent Health

Supplementary appendix

This appendix formed part of the original submission and has been peer reviewed. We post it as supplied by the authors.

Supplement to: Oliveira EA, Colosimo EA, Simões e Silva AC, et al. Clinical characteristics and risk factors for death among hospitalised children and adolescents with COVID-19 in Brazil: an analysis of a nationwide database. *Lancet Child Adolesc Health* 2021; published online June 10. https://doi.org/10.1016/S2352-4642(21)00134-6.

Supplementary material to

Clinical characteristics and risk factors of mortality among 11,613 hospitalized children and adolescents with COVID-19: an analysis of a nationwide database Lancet Child and Adolescent Health

Supplementary material (S.1) SIVEP-Gripe (The Influenza Epidemiological Surveillance Information System)

The Influenza Epidemiological Surveillance Information System, SIVEP-Gripe (*Sistema de Informação de Vigilância Epidemiológica da Gripe*), was established by the Brazilian Ministry of Health (MS) in 2009, due to the Influenza A (H1N1) pandemic. This system has been maintained since then for the monitoring of severe acute respiratory syndrome (SARS) cases and surveillance of unusual events associated with respiratory infections in the country. Since then, this sentinel surveillance has been expanded for other respiratory viruses. In 2020, the surveillance of COVID-19 was incorporated into the system network. For the purpose of monitoring hospitalized cases of COVID-19, the Ministry of Health incorporated the testing of the SARS-CoV-2 virus to the surveillance of SARS. Case notification is mandatory, and records are stored in the computerized database SIVEP-Gripe.

In 2020, the Ministry of Health released a "Technical Note" with recommendations for the immediate notification of suspected and confirmed cases of COVID-19. It was recommended that outpatients with mild COVID-19 be notified in the e-SUS VE system and those cases hospitalized be reported in the SIVEP-Gripe system (Technical Note available at

<u>https://www.cosemsmg.org.br/site/Arquivos/PDF/nms20.pdf</u>). Consequently, SIVEP-Gripe came to be the single source for all patients admitted to both public and private hospitals with COVID-19 in Brazil. In practical terms, SIVEP-Gripe has been the primary source of information on all COVID-19 hospital admissions and deaths in the country.

To enter the SIVEP-Gripe database, the case must present flu-like syndrome and at least one of the following criteria: dyspnea or respiratory distress or O_2 saturation less than 95% in room air or cyanosis or symptoms specific for children (intercostal retractions, nasal flaring, dehydration, and inappetence). Also, they need to have been admitted to hospital or died without hospital admission.

Detailed information regarding this database, including reporting form and data dictionary, codes, and all de-identified data, such as individual participant data, are publicly available at

https://opendatasus.saude.gov.br/dataset/bd-srag-2020).

We downloaded the original database on January 10, 2021 and, for analysis purposes, the original dataset was saved in Excel, SPSS, and STATA versions. Based on the original reporting form and data dictionary, we recoded the variables using similar approach described by Ranzani et al. in a supplementary material (eTable 3) of the paper "Characterisation of the first 250 000 hospital admissions for COVID-19 in Brazil: a retrospective analysis of nationwide data". Lancet Respir Med 2021. (available at https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30560-9/fulltext#supplementaryMaterial).

In addition, we created variables for the competing-risk survival analyses. For instance, from the fields "date of admission" and "date of discharge or death" we created the variable time_event (in days). In addition, we manually revised the field "morb_desc", an open-ended one, which included some relevant clinical information, especially related with comorbidities and another risk factors (for example, prematurity). From this field, we created the close-ended field Malignancy by manually scrutinizing the field morb_desc. Of 11,613 indexed cases, 1,760 (15%) had some observation included in this field. We performed this task in the SPPS statistical package by using the diverse search mechanisms available. Thus, for instance, for the field Malignancy we searched all 1,760 cases for any oncology issue and found 583 cases of cancer in this sample. As expected, most cases were related with hematological malignancies like leukemia, lymphoma, or CNS tumors.

Supplementary material (S.2)

Sensitivity Analysis

For analysis purposes, in order to evaluate the scope of the sample in SIVEP-Gripe database, we carried on several preliminary analyses with the entire dataset of 11,603 cases < 20 years with laboratory-confirmed COVID-19.

As shown in supplementary material S.1, to enter the SIVEP-Gripe database, the case must present flulike syndrome and at least one of the following criteria: dyspnea or respiratory distress or O2 saturation less than 95% in room air or cyanosis or symptoms specific for children (intercostal retractions, nasal flaring, dehydration, and inappetence). Also, they need to have been admitted to hospital or died without hospital admission (https://opendatasus.saude.gov.br/dataset/bd-srag-2020).

Of note, our preliminary analysis has shown that only about 47%, 54%, and 41% of the pediatric patients in the sample had respectively dyspnea, respiratory distress, and oxygen saturation <95% at admission (with a significant overlapping between them). As a whole, according to the data available in SIVEP-Gripe database, 7,081 cases (61%) fulfilled the strict criteria for entry in database. In this context, to ascertain the robustness of our analysis further , we created a subset of cases including only those cases who fulfilled the strict criteria to be enrolled in SIVEP-Gripe database. In this subset, 714 (10%) had the primary outcome. Similarly. in this subset we also evaluated the primary outcome by competing risks analysis, using cumulative incidence function (CIF) and Fine and Gray proportional sub-distribution hazards model. Discharge was treated as a competing event. The complete results of sensitivity analysis are presented in supplementary tables (Appendix Tables S.6, Tables S.7, Tables S.8) and in supplementary figures (Figures S.9 and S.10A-D).

It can be observed that these results are very similar to those based on the entire data set of 11603 cases. There are very minor differences in the univariate analysis presented in Table S.5 when compared to Table S.7, for instance, regarding variable "Asian ethnicity" for both competitive outcomes. On the other hand, and more important, multivariate analysis for both datasets are in complete agreement showing the robustness of the results based on the entire data set, A final issue in this comparison should be highlighted. The analysis based on the entire data set, that is larger than the strict criterion one, is naturally more precise. The smaller length of the confidence intervals is the most notable evidence of this fact.



Figure S.3. Flow diagram of sample selection

	North (%)	Northeast (%)	Central-West (%)	Southeast (%)	South (%)
	1,841	3,388	1,356	4,065	963
Cases /million young pop.	330.4	227.0	330.0	210.5	144.5
Age (years)					
Median (IQR)	40 (1 - 130)	42 (0.7 - 130)	98 (2-170)	53 (1 - 140)	5.5 (1 - 15.0)
Mean (SD)	69 (67)	6.8 (6.2)	98(72)	7.5 (6.9)	7.8 (7.2)
Age group (years)					
0 - 1.9	692 (35.6)	1318 (38.9)	337 (24-8)	1375 (33.9)	329 (34:2)
2 - 4.9	294 (16.0)	495 (14.6)	147 (10.8)	607 (14.9)	141 (14.6)
5 - 11.9	353 (191)	661 (195)	268 (197)	830 (20:4)	177 (18·3)
12 - 199	502 (27.2)	914 (269)	604 (44:5)	1253 (30.8)	316 (32.8)
Gender (n = 11,600)					
Male	953 (51.8)	1807 (53:5)	675 (49.8)	2083 (51.3)	514 (53:4)
Female	888 (48.2)	1570 (465)	681(50:20	1980 (40.7)	449 (46.6)
Ethinicity $(n = 9,109)$					
White	157 (8.9)	343 (13·2)	286 (30.6)	1734 (560)	671(82.5)
Black / Brown	1501 (85.3)	2204 (85:4)	619 (66-2)	1326 (42.8)	134 (16:4)
Asian	11 (0.6)	23 (0.89)	15 (1.6)	27 (0.87)	3 (0.37)
Indigenous	90 (5.1)	11 (0:42)	14 (149)	7 (0.22)	5 (0.61)
Number of comorbidities					
None	1488 (80.8)	2382 (703)	1115 (82.2)	2751 (67.7)	616 (63:4)
1	309 (16.8)	898 (26.5)	211 (15.5)	1088 (267)	274 (28:4)
2	39 (2.1)	95 (2.8)	24 (1.8)	191 (4:7)	54 (5.6)
>3	5 (0.3)	13 (0.38)	6 (0:44)	35 (0.86)	19 (2.0)
ICU admission ($n = 11,074$)					
No	1495 (84.6)	2314 (73.3)	1051 (83.4)	2728 (69:4)	727 (760)
Yes	273 (15:4)	843 (267)	291 (16.6)	2015 (30.6)	229 (24:0)
Death rate $(n = 10,867)$					
No	1592 (91.5)	2743 (89.5)	1199 (963)	3657 (94.3)	882 (93.5)
Yes	147 (8.5)	320 (10.5)	46 (3.7)	220 (5.7)	61 (65)

Table	S.4.	. Demograph	ic and clir	nical chara	acteristics o	of children v	with positive	e RT-aPO	CR COVID	-19 accord	ing to the l	Brazilian ı	macroregions
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Data (n) in the first column represent the available data for those variables with missing values (gender, ethnicity, and signs and symptoms at presentation). For comorbidities, we assumed missing values as the absence of the clinical condition.

	Discharge (%)*	HR (95% CI)	P-value	Death (%)*	HR (95% CI)	P-value
	10 410 (02-2)		1 (4140	996 (7.9)		1 vulue
	10,410 (92.2)			890 (7.9)		
Age (years)						
Median (IQR)	52 (10 – 143)	098 (098, 099)	<0.001	5.5 (0.5 – 15.9)	100 (099, 101)	0.076
Mean (SD)	7.5 (68)			80(75)		
Age group (years)						
0 - 19	3,548 (34-1)	0.86 (0.83;0.90)	<0.001	354 (40:0)	1 88 (1 58; 2 24)	<0.001
2-119	3,788 (35.5)	1.0		194 (21.9)	1.0	
12-199	3,164 (30:4)	0.75 (072; 079)	<0:001	338 (38.1)	198 (165; 236)	<0.001
Gender**						
Male	5,402 (51.9)	101 (098; 105)	0.317	466 (52.7)	102 (09; 116)	0.701
Region						
Southeast	3,756 (36.1)	1.0		236 (26.6)	1.0	
South	887 (8.5)	1.03 (095;111)	0.463	63 (71)	111 (084; 146)	0.420
Central-West	1,283 (12.3)	0.75 (071; 079)	<0.001	49 (5.5)	062 (045; 084)	0.003
Northeast	2,840 (273)	0.68 (0.65; 0.71)	<0.001	377 (42.6)	2:03 (1:72; 2:38)	<0.001
North	1,644 (15.8)	0.83 (078; 088)	<0.001	161 (18·2)	1 53 (1 25; 1 87)	<0.001
Race#						
White	2,919 (35.8)	10		214 (28.9)	1.0	
Black / Brown	5,112 (62.3)	0.85 (0.81; 0.89)	<0:001	493 (66.5)	1.30 (1.11;1.53)	0.001
Asian	70 (0.9)	0.79 (0.62; 0.99)	0.045	8 (1.1)	1 50 (0 75; 3 02)	0.247
Indigenous	100 (1·2)	0.67 (0.54; 0.83)	<0:001	26 (3.5)	3 27 (2 17; 4 93)	< 0.001
Signs and symptoms						
Fever	6,984 (750)	0.98 (094, 103)	0.53	559 (73.9)	0.97 (0.82, 1.14)	0.72
Cough	6,206 (68.4)	1.11 (106, 115)	<0.001	413 (58.8)	0.68 (0.58, 0.79)	<0.001
Odynophagia	1,926 (24.8)	0.97 (092, 101)	0.22	109 (18:4)	0 69 (0 56, 0 85)	<0.001
Respiratory distress	4,385 (52.3)	0.78 (075, 081)	< 0.001	564 (75.9)	2.78 (2.35, 3.29)	<0.001
O2 saturation <95%	3,061 (381)	0.69 (0.66, 0.72)	< 0.001	504 (70.5)	3.66 (3.11, 4.30)	<0.001
Dyspnea	4,783 (45.9)	0.80 (0.72;0.83)	< 0.001	581 (65.6)	2.65 (2.23;3.14)	<0.001
Anosmia	317 (7.5)	0.97 (0.88, 1.07)	0.60	14 (5.0)	0.65 (038, 112)	0.12
Ageusia	297 (7.1)	0.99 (089, 102)	0.25	13 (4.6)	0.64 (0.36, 1.12)	0.12
Diarrhea	1,657 (21.2)	1.03 (098, 109)	0.14	117 (191)	0.88 (0.72, 1.08)	0.23
Vomit	1,821 (23.2)	1.01 (096,1.06)	0.57	147 (23:4)	100 (082, 121)	0.92
Abdominal pain	607 (14:0)	099 (091, 107)	0.87	45 (14.7)	1 04 (0 75, 1 43)	0.78
Comorbidity (yes/no)	2,715 (261)	078 (074; 081)	<0.001	448 (50.6)	2 26 (1 98; 2 58)	<0.001
Number of comorbidities						
None	7695 (73.9)	1		438 (49.4)	10	<0.001
1	2,352 (22.6)	077 (073,081)	<0.001	347 (39.2)	2.42 (2.10, 2.79)	
2	308 (3.0)	056 (050, 063)	<0:001	81 (91)	411 (325, 518)	<0.001
<u>></u> 3	55 (0.5)	044 (034, 056)	<0.001	20 (2.3)	548 (355, 846)	<0.001
Main comorbidities						<0.001
Asthma	823 (7.9)	148 (138; 159)	< 0.001	32 (3.6)	0 60 (0 42; 0 85)	<0.001
Pulmonary	204 (2.0)	074 (0.65, 0.85)	< 0.001	32 (3.6)	1.80 (1.27;2.55)	0.001
Neurology	556 (5.3)	0.61 (0.56, 0.66)	< 0.001	125 (14-1)	2.66 (2.20,3.21)	<0.001
Oncology	445 (4.3)	0.57 (0.52,0.63)	< 0.001	125 (14·1)	3.22 (2.68, 3.87)	<0.001
Cardiology	276 (2.7)	0.56 (0.50, 0.63)	< 0.001	83 (9:4)	3 36 (2 68, 4 20)	<0.001
Hematology	230 (2.2)	0.78 (0.69, 0.88)	<0.001	33 (3.7)	1.66 (1.17, 2.35)	0.004
Renal	131 (1.3)	057 (048, 0.68)	<0.001	40 (4.5)	340 (248, 466)	<0.001
Distant			-0.001	33 (3.7)	2.16(1.52, 2.06)	<0.001
Diabetes	155 (1.6)	0 /6 (0 65, 0 88)	<0.001	55 (57)	210(155.500)	<0001
Obesity	155 (1.6) 134 (1.3)	0.84(0.70, 1.00)	0.051	23 (2.7)	1.93 (1.29, 2.90)	<0001

Table S5. Univariate survival analysis competitive risk according to the demographic and clinical characteristics of children with positive RT-qPCR COVID-19

* 317 missing cases regarding primary outcome ** 13 missing; # 2,432 missing

	Overall (%)
	7,081 (100)
Age (years)	
Median (IQR)	47 (1 – 13.9)
Mean (SD)	7.3 (69)
Age group (years)	
0 - 1.9	2464 (34.8)
2 - 11.9	2526 (35.7)
12 - 199	2091 (29.5)
Gender $(n = 7,075)$	
Male	3311 (46.7)
Female	3764 (53.3)
Region	
Southeast	2553 (36.1)
South	590 (8.3)
Central-West	521 (7.4)
Northeast	2134 (30.1)
North	1283 (18.1)
Ethinicity (n = $5,821$)	
White	1912 (32.8)
Black / Brown	3776 (64.9)
Asian	39 (0.66)
Indigenous	94 (1.61)
Signs and symptoms at presentation	
Fever (n= 6,461)	4,846 (75.0)
Cough $(n = 6,460)$	4,981 (77.1)
Respiratory distress ($n = 6,371$)	5,056 (79.3)
Oxygen saturation $<95\%$ (n = 6,016)	3,685 (61.2)
Dyspneia ($n = 6,614$)	5,529 (83.6)
Odynophagia (n = 5,370)	1,367 (25.5)
Anosmia $(n = 2,890)$	213 (74)
Ageusia (n = 2,880)	188 (65)
Diarrhea $(n = 5,435)$	993 (18·3)
Vomit (n = 5,480)	1,154 (210)
Abdominal pain ($n = 3,008$)	365 (12:0)
Number of comorbidities	
None	4723 (66.7)
1	1997 (28.2)
2	298 (4.2)
>3	63 (0.9)
Main comorbidities	
Asthma	765 (10.8)
Pulmonary	217 (31)
Neurologic	530 (7·3)
Oncologic	581 (50)
Cardiopathy	285 (4.0)
Hematological	167 (2:4)
Renal	94 (13)
Diabetes	161 (23)
Obesity	130 (1.8)
Syndrome/chromosomal abn	164 (2:3)

Table S.6. Demographic and clinical characteristics of children with positive RT-qPCR COVID-19
that fulfilled SIVEP criteria

Data (n) in the first column represent the available data for those variables with missing values (gender,

ethnicity, and signs and symptoms at presentation). For comorbidities, we assumed missing values as the absence of the clinical condition.

	Discharge (%)*	HR (95% CI)	P-value	Death (%)*	HR (95% CI)	P-value
	6,166 (89.6)			714 (10.4)		
Age (years)	· · · ·					
Median (IOR)	52 (10 – 143)	0.993 (0.98, 0.996)	<0.001	5.5 (0.5 - 15.9)	101 (100, 102)	0.002
Mean (SD)	7.5 (6.8)			8.0 (7.5)		
Age group (years)						
0 - 19	2,250 (36.5)	0.80 (0.83;0.90)	<0.001	152 (21.3)	1.89 (1.55; 2.30)	<0.001
2 - 119	2.148 (34.8)	1.0		285 (39.9)	1.0	
12 - 199	1.768 (28.7)	0.75 (0.70; 0.79)	<0.001	227 (38.8)	2.20 (1.81; 2.96)	<0.001
Gender						
Male	3,292 (53.4)	1.01 (0.98; 1.05)	0.317	368 (51 ^{.5})	0.93 (080; 107)	0.33
Region						
Southeast	2,305 (37.4)	1.0		189 (26.5)	1.0	
South	528 (8.6)	0.99 (0.89;1.09)	0.91	53 (7.4)	1 20 (0 89; 1 63)	0.22
Central-West	470 (7.6)	0.87 (0.79; 0.95)	0.003	41 (5.7)	105 (075; 148)	0.73
Northeast	1,741 (28.2)	0.67 (0.63; 0.71)	<0.001	298 (41.7)	196 (164; 236)	<0.001
North	1,122 (18.2)	0.85 (0.79; 0.91)	<0.001	133 (18.6)	142 (113; 177)	0.002
Race#				· · · · ·		
White	1,703 (33.6)	1.0		170 (28.3)	10	
Black / Brown	3,259 (64.3)	0.86 (0.81; 0.91)	<0.001	401 (66.8)	121 (101, 144)	0.035
Asian	32 (0.6)	0.76 (0.52; 1.10)	0.150	7 (1.2)	208 (098; 442)	0.053
Indigenous	71 (1.4)	0.63 (0.49; 0.80)	<0.001	22 (3.7)	285 (183; 445)	< 0.001
Signs and symptoms						
Fever	4,249 (68.9)	0.96 (0.94, 1.04)	0.53	478 (67.0)	101 (084, 121)	0.86
Cough	4,485 (72.7)	0.94 (0.92, 0.98)	0.002	374 (52:4)	048 (040, 057)	<0.001
Odynophagia	1,243 (201)	1.06 (1.00, 1.13)	0.04	100 (14 0)	072 (058, 089)	0.004
Respiratory distress	4,385 (711)	0.83 (0.78, 0.89)	<0.001	564 (789)	1 67 (1 33, 2 09)	<0.001
O2 saturation <95%	3,061 (49.6)	0.69 (0.65, 0.73)	<0.001	504 (70.6)	2.81 (2.30, 3.43)	<0.001
Dyspneia	4,783 (77.6)	0.91 (0.86;0.98)	0.009	581 (81.4)	1.47 (1.16;1.86)	0.001
Anosmia	198 (3.2)	1.06 (0.92, 1.09)	0.32	12 (17)	0.66 (037, 119)	0.17
Ageusia	174 (2.8)	1.07 (0.94, 1.23)	0.28	11 (16)	069 (038, 127)	0.23
Diarrhea	880 (14.3)	0.96 (0.89, 1.02)	0.25	97 (13.6)	1 03 (0 82, 1 29)	0.76
Vomit	1,010 (16.4)	0.96 (0.90,1.02)	0.22	116 (162)	105 (086, 131)	0.28
Abdominal pain	312 (50)	0.90 (0.81, 1.00)	0.07	37 (5.2)	126 (089, 179)	0.18
	2715 (201)	0.70 (0.74, 0.01)	-0.001	110 (50 ()	226 (1.09, 2.59)	.0.001
Comorbialty (yes/no)	2,/15 (201)	0.78 (0.74; 0.81)	<0001	448 (50%)	220 (198; 258)	<0001
Number of comorbidities		1		242 (48.0)	10	
None	4,256 (69.0)	1	.0.001	343 (48.0)	10	.0.001
1	1,648 (26.7)	0.78 (0.74,0.82)	<0001	283 (39.6)	2.01 (1.72, 2.35)	<0.001
2	220 (3.6)	0.54 (0.47, 0.61)	<0001	70 (9.8)	343 (267, 441)	<0001
≥3 Main asmarkiditias	42 (0.7)	0.45 (0.34, 0.59)	<0001	18 (2.5)	431 (264, 680)	<0001
A sthma	724 (11.7)	1 56 (1 45, 1 60)	<0.001	20 (4 2)	0.24 (0.24, 0.50)	<0.001
Asuma	124 (11.7)	1.30(1.43, 1.09)	<0.001	30 (4.2)	1.40 (0.08:201)	<0001
Pullionary	1// (2.9)	0.79 (0.69, 0.91)	< 0.001	50 (4.2) 106 (14.8)	140(0.98;201)	<0.001
Neurology	404 (6.6)	0.59 (0.54, 0.65)	<0.001	106 (14.8)	2.26 (1.84,2.67)	<0001
Cardiology	443 (4.3) 100 (2.1)	0.40 (0.40,0.52)	<0.001	123(141)	322 (200, 3°87) 336 (268, 400)	<0.001
Hamatalagy	190 (5.1)	0.32 (0.43, 0.39)	<0.001	(4(104))	5 50 (2 08, 4 ² 20)	<0.004
Repel	130 (2.2)	0.74 (0.04, 0.80)	<0.001	20 (50)	100(117, 233)	0.001
Kenai Diabata	04 (1.0)	0.51 (0.40, 0.65)	<0.001	29 (4.1)	5'40 (2'48, 4'00)	<0.001
Diabetes	122 (2.0)	0.72 (0.61, 0.86)	<0.011	21 (2.9)	210 (103. 306)	<0001
Odesity	134 (1.3)	0.85 (0.70, 1.03)	0.001	23 (2·7)	195 (129, 290)	<0.001
Syncronie	120 (1.9)	0.02 (0.32, 0.74)	<0001	40 (3.0)	511 (255, 415)	<0001
						<0.001

Table S.7. Univariate survival analysis competitive risk according to the demographic and clinical characteristics of children with laboratory-confirmed COVID-19 that fulfilled SIVEP criteria

* 201 missing cases regarding primary outcome, #1,260 missing cases regarding ethnicity

<0.001

	Discharge (%)*	HR (95% CI)	P-value	Death (%)*	HR (95% CI)	P-value
	6,166 (89.6)			714 (10.4)		
Age group (years)						
0 - 1.9	2,250 (36.5)	0.76 (0.72;0.81)	<0.001	152 (21.3)	2.31 (1.85; 2.88)	<0.001
2 - 11.9	2,148 (34.8)	1.0		285 (39.9)	1.0	
12 - 199	1,768 (28.7)	0.72 (0.67; 0.76)	<0.001	227 (38.8)	2:47 (1:99 3:01)	<0.001
Region						
Southeast	2,305 (37.4)	1.0		189 (26.5)	1.0	
South	528 (8.6)	099 (088;110)	0.86	53 (7.4)	124 (089; 170)	0.177
Central-West	470 (7.6)	077 (069; 086)	<0.001	41 (5.7)	1 25 (0 85; 1 82)	0.241
Northeast	1,741 (28.2)	063 (058; 067)	<0.001	298 (41.7)	2.18 (1.74; 2.73)	<0.001
North	1,122 (18.2)	081 (075; 088)	<0.001	133 (18.6)	1 52 (1 16; 1 97)	0.002
Race#						
White	1,703 (33.6)	1.0		170 (28.3)	1.0	
Black / Brown	3,259 (64.3)	0.95 (0.89; 1.02)	0.171	401 (66.8)	1 08 (0 88;1 34)	0.438
Asian	32 (0.6)	0.82 (0.56; 1.20)	0.324	7 (1.2)	197 (090; 428)	0.086
Indigenous	71 (1.4)	063 (049; 082)	<0.000	22 (3.7)	3 24 (2 02; 5 17)	<0.001
Comorbidities						
None	4,256 (69.0)	1.0		343 (48.0)	1.0	
1	1,648 (26.7)	072 (067, 076)	<0.001	283 (39.6)	258 (2.15,309)	<0.001
2	220 (3.6)	046 (040,054)	<0.001	70 (9.8)	4.18 (3.13, 5.58)	<0.001
<u>></u> 3	42 (0.7)	036 (027,049)	<0.001	18 (2.5)	631 (385, 103)	<0.001

Table S.8. Multivariate survival analysis competing risk in children with laboratory-confirmed COVID-19 that fulfilled SIVEP criteria

* 201 missing cases regarding primary outcome, #1,260 missing cases regarding ethnicity





Time (days)

Cumulative Incidence

10



Time (days)





Time (days)



Cumulative Incidence

Time (days)

13

Supplementary figures legends

Figure S.9. Cumulative incidence functions for mortality and discharge in children and adolescents with Covid-19 that fulfilled SIVEP criteria.

Figure S.10. Cumulative incidence functions for mortality of children and adolescents with COVID-19 that fulfilled SIVEP criteria according to the (A) Age-group; (B) Region; (C) Ethnicity; and (D) Number of comorbidities