## **ERRATUM /** *ERRATA*

In the manuscript "Evaluation of the initial response in clinical trial efforts for COVID-19 in Brazil", DOI: 10.1590/1980-549720200104, published in the Rev bras epidemiol. 2020; 23: e200104. On page 7, Table 1.

## Where it reads:

Table 1. Characteristics of COVID-19 research protocols in development in Brazil available at clinical trials registry databases until May 2020.

Interventions for the tr	eatment or prophylaxis of COVID-19 (n=22)	n	(%)
Study design	Randomized clinical trial	18	(82)
	Non-randomized clinical trial	4	(18)
Trial phase	Phase 1	3	(14)
	Phase 2	5	(23)
	Phase 3	8	(36)
	Phase 4	2	(9)
	NM	4	(18)
Masking	Open-label	13	(59)
	Single-blind	1	(5)
	Double-blind	2	(9)
	Quadruple-blind	6	(27)
Control group	Placebo/standard treatment comparator	13	(59)
Multicenter <sup>a</sup>	Yes	7	(32)
	Hospital	16	(73)
Recruitment site	Outpatient clinic	2	(9)
	NM	4	(18)
Power calculation <sup>b</sup> Scenario of large effect size RR = 0.4; incidence: 10 vs. 25%)	High ≥ 90% <sup>c</sup>	1	(5%)
	Good (< 90 and ≥ 80%) <sup>d</sup>	1	(5%)
	Low (< 80 and > 50%) <sup>e</sup>	2	(9%)
	Very low (< 50%) <sup>f</sup>	17	(81%
Power calculation <sup>b</sup> Scenario of moderate effect size RR = 0.6; incidence: 7.2 vs. 12%)	High ≥ 90% <sup>g</sup>	11	(52%
	Good (< 90 and ≥ 80%) <sup>h</sup>	1	(5%)
	Low (< 80 and > 50%) <sup>i</sup>	3	(14%
	Very low (< 50%) <sup>j</sup>	6	(29%
Interven	tions for the treatment of COVID-19 (n=21)		
Inclusion criterion - COVID-19	Confirmed diagnosis (test+)	5	(24)
	Suspected diagnosis - Clinical suspicion or test+	12	(57)
	Suspected diagnosis - Clinical suspicion only	4	(19)
Severity of the patients included	Hospitalized - moderate or severe	4	(19)
	Hospitalized - severe	5	(24)
	Hospitalized - moderate	1	(5)
	Hospitalized - nonspecific	6	(28)
	Not hospitalized	1	(5)
	NM	4	(19)
Use of important outcomes for COVID-19k	As primary outcome	8	(38)
	Among secondary outcomes	18	(86)

°More that one recruitment center; 'power calculation did not include the study of the World Health Organization due to access limitations of the sample size in each arm; 'sample size (n) = 1,986; 'n = 130; 'n between 400 and 1,000; 'n between 20 and 600; 'n between 630 and 1,968; 'n between 200 and 290; 'n between 66 and 210; 'n between 22 and 50; 'simportant outcomes according to the consensus document published on the initiative Core Outcome Measures in Effectiveness Trials for hospitalized patients; NM: not mentioned; RR: relative risk.

## It should read:

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	Low (< 80 and > 50%) <sup>e</sup>	2	(9%)
	Very low (< 50%) <sup>f</sup>	17	(81%)
Power calculation <sup>b</sup> Scenario of large effect size RR = 0.4; incidence: 10 vs. 25%)	High ≥ 90% <sup>g</sup>	10	(48%)
	Good (< 90 and ≥ 80%) <sup>h</sup>	2	(9.5%)
	Low (< 80 and > 50%) <sup>i</sup>	2	(9.5%)
	Very low (< 50%) <sup>j</sup>	7	(33%)
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Inclusion criterion - COVID-19	Confirmed diagnosis (test+)	5	(24)
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 $^{\rm o}$ More that one recruitment center;  $^{\rm b}$ power calculation did not include the study of the World Health Organization due to access limitations of the sample size in each arm;  $^{\rm c}$ sample size (n) = 1,986;  $^{\rm d}$ n = 130;  $^{\rm c}$ n between 400 and 1,000;  $^{\rm f}$ n between 20 and 600;  $^{\rm o}$ n between 630 and 1,968;  $^{\rm h}$ n between 200 and 290;  $^{\rm h}$ n between 66 and 210;  $^{\rm h}$ n between 22 and 50;  $^{\rm k}$ important outcomes according to the consensus document published on the initiative Core Outcome Measures in Effectiveness Trials for hospitalized patients; NM: not mentioned; RR: relative risk.