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INTRODUCTION			PARAMETER							MEDIAN VA		
		Men							69.	6%		
<ul> <li>COVID-19 infection diagnosed by RT-PCR assays for SARS- CoV-2 on nasopharyngeal swab (NPS).</li> </ul>		Age [years]						60				
		Aspartate transaminase (AST) [u/L]						41				
<ul> <li>Sensitivity depends on illness duration and operator performance.</li> <li>10.20% false performance of PT PCP.</li> </ul>		Alanine transaminase (ALT) [u/L]							33			
		Lymphocyte count (LYMPH) [cells/mm <sup>3</sup> ]						0.84				
		Procalcitonin level (PROCAL) [ng/mL]							0.15			
<ul> <li>10-30% false negative rate of RT-PCR.</li> </ul>												
<ul> <li>We formulated and applied a clinical prediction tool for COVID-19 diagnosis [Table 1].</li> </ul>		Ferritin level (FERR) [ng/mL]						700				
		C-Reactive Protein (CRP) [mg/dL]						10.06				
		Lactate dehydrogenase level (LDH) [U/L]						365				
		SYMP (symptom duration prior to presentation) [days]							7			
METHO	DS	<u>Table</u>	<u>e <b>2:</b></u> Demogra	aphic and Me	edian Lab D	ata of VHCL	COVID-	19 patie	ents.			
Retrospective descriptive study (	13/2020-04/2020)											
conducted at a tertiary hospital in	· · · · ·	Pt	Age/Sex	AST/ALT	LYMPH	PROCAL	FERR	CRP	LDH	S١		
<ul> <li>Admitted pts. with ≥2 consecutive negative NPS COVID-19 RT-PCR tests assessed.</li> </ul>		1	69/M*	<b>63/83</b>	<b>0.57</b>	0.12	737	2.69	241			
		2	39/F* 48/M*	20/13 55/53	1.05 0.79	0.06 0.08	410 4910	2.92 1.65	NA 360			
		4	70/M	41/18	0.75	0.56	3515	16.9	249			
<ul> <li>Pts. meeting all 5 criteria for VHCL of COVID-19 infection [Table 1] included.</li> </ul>		5	55/M*	238/197	0.72	0.19	3204	9.42	413			
		6	23/F*	34/31	1.14	0.05	248	10.98				
<ul> <li>Additional data collected from electronic records.</li> </ul>		/ 8	55/M <sup>*</sup> 54/M	30/33 27/35	0.47 0.73	0.12 0.16	506 700	4.79 10.06	336 397			
		9	78/M	17/10	0.88	0.09	1306	11.22				
		10	71/M*	100/167	0.13	2.42	2143	8.30	371			
		11	61/M	148/76	0.89	0.19	3374	13.8	325			
CLINICAL CR		12	31/F	25/17	0.36	3.67	488	60.19	313			
		13	60/F	21/26	1.22	0.07	383	9.63	379			
1a. Temperature >38.3 C in first 24 hou	rs of	14	57/F*	107/87	1.16	0.08	1145	5.08	602			
hospitalization <u>OR</u>		15	84/F	31/23	1.20	0.15	795	5.40	259			
1b. Subjective fevers leading into hospi	talization	16	61/M	49/39	0.40	0.04	475	10.46				
2 New Respiratory Symptoms		<b>17</b>	<b>28/M</b> *	<b>41/32</b>	<b>0.89</b>	0.16	<b>263</b>	<b>6.81</b>	<b>333</b>			
		18 19	71/M 56/M	27/29 25/16	1.06 1.14	0.16 0.09	1110 363	25.3 11.0	355 327			
		20	89/M	42/15	1.02	0.60	409	27.9	527 417			
<b>3</b> Lymphocytopenia (lymphocyte count less than 1.25/mm <sup>3</sup> ).		20	78/M	69/34	0.27	2.22	332	17.78				
		22	61/M	75/111	0.84	0.42	1303	23	597			
4 Bilateral infiltrates without pleural effusions on chest imaging		23	25/F	121/121	0.60	0.04	343	3.27	448			
Lack of more likely diagnosis after inpat 72 hours	ient monitoring for a minimum of			and laborator d units of lab	-			D-19 pa	tients.			

**Table 1**: Clinical criteria for VHCL of COVID-19 Infection.

# **Very High Clinical Likelihood (VHCL) Of COVID-19 Infection:** Peering Beyond A Negative Nasopharyngeal Swab

**Bolded rows:** patients with 3<sup>rd</sup> NPS swab that was positive (#1,4,5,10,11,13,17). \*: Patients had negative respiratory viral panel testing.

- VHCL **[Tables 2,3]**.
- **[Table 2]**.
- 7 had 3<sup>rd</sup> positive RT-PCR **[Table 3]**.
- diagnosis.
- pandemic can:

Clinical criteria (such as VHCL COVID-19) can supplement laboratory testing for appropriate diagnosis and treatment, contact notification and infection control.

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### **RESULTS & DISCUSSION**

• 1,855 pts admitted in study period.

• 23 pts had ≥2 negative RT-PCR tests but met criteria for

• These 23 pts. had median 7 days of symptoms, lymphocytopenia; elevated PROCAL, FERR, CRP levels

• Pts with high clinical suspicion of COVID-19 with negative testing have been reported. Strict clinical criteria may help

• Incorporating clinical criteria in diagnostic algorithms in a

a) optimize infection control b) identify pts for emerging therapeutics c) aid in contact tracing to help reduce nosocomial/community transmission

### CONCLUSION

## BIBLIOGRAPHY