

Routine Laboratory Test Indices for COVID-19 Patient Management

Hematology Test Indices

			Symptom* Onset Admission	Hospitalization	Discharge**
	Test Parameter	Reference Interval	9 days (median)	12 days (median)	(Reference from COVID-19 Dessignated Hospitals, China)
W B C	White blood cells (WBC)	4.0-10.0 x10 ⁹ /L	Normal, or slightly elevated	Survivals Shifting with slight increasement within the reference range ¹ None-survivals Exceeding the upper reference range ¹	> 3.0 x10 ⁹ /L
	Lymphocyte number (Lym#)	0.8-4.0 x10 ⁹ /L	Normal, or slightly decreased	Survivals Prograssively decreasing, followed by recovering climbing back. None-survivals Persistent decrease, fluctuating at low level (below 0.8 x10 ⁹ /L) ¹	> 1.0 x10 ⁹ /L
				In Mindray COVID-19 retrospective study, Al acquired Lym# & RDW-SD parameter (unpublished, requiring further verification) > 0.794 could predict severe progression.	
	Monocyte number (Mon#)	0.12-1.2 x10 ⁹ /L	Normal, or slightly decreased	Monocyte deform to phagocyte, engulfing virus. In the deterioration process, Mon cell cluster appears some sudden change in SF CUBE (Mindray unpublished retrospective study).	0.12-1.2 x10 ⁹ /L
	Neutrophil number (Neu#)	2.0-7.0 x10 ⁹ /L	Normal, or slightly elevated	Survivals Prograssively increasing, rising slowly within the reference range ¹ None-survivals Prograssively increasing, exceeding the upper reference range ¹	> 1.5 x10 ⁹ /L
	Eosinophil number (Eos#)	0.02-0.5 x10 ⁹ /L	Normal, or slightly decreased	Progressively decreasing, some will fall out of the lower reference range ³	0.02-0.5 x10 ⁹ /L
	High fluorescent Cell number (HFC#)	0.00 x10 ⁹ /L	Normal, or slightly increased	Some results will be flagged with atypical lymphacyte.	0.00 x10 ⁹ /L
	Neutrophil-to- lymphocyte ratio (NLR)	Cutoff: 3.13 ²	Normal, or slightly increased	Elderly patients (>50 years) with NLR>3.13 are recommended to transfer to ICU ²	NA
	NLR & RDW-SD	Cutoff: 1.06⁴	Normal, or slightly increased	Patients with NLR & RDW-SD > 1.06 can be classified as the severe progression for more intervention therapy ⁴	NA
R B C	Reticulocyte number (Ret#)	0.02-0.20 x10 ¹² /L	Normal, or slightly increased,	Severe and critially ill patients will have high Ret count and IRF (Mindray unpublished retrospective study).	> 0.02 x10 ¹² /L
	Immature Reticulocyte Fraction (IRF)	0.0-25.0 %	could decrease in severe cases		0.0-25.0 %
R E	Hemoglobin (HGB)	110-160 g/L	Normal, or slightly decreased	Progressively decreasing, then rising back during recovery	> 90 g/L
T	Red blood cell distribution width – standard deviation (RDW-SD)⁵	35.0-56.0 fl	Normal, or slightly increased	Progressively increasing, can be combined with other parameters for severity indentification or prediction.	35.0-56.0 fl
P L T	Platelet count (PLT)	100-300 x10 ⁹ /L	Normal, or slightly increased, could decrease in severe cases	In the well-controlled cases, PLT rises progressively, then declining during recovery Decreasing with septic deterioration ⁶ , then rising back during recovery	> 80 x10 ⁹ /L
	Platelet Distribution Width (PDW)	6.5-12.0 fl	Normal, or slightly increased	Progressively increasing, then going down during recovery	6.5-12.0 fl
	Immature Platelet Fraction (IPF)	0.9-10.0 %	Normal, or slightly increased	Progressively increasing, then going down during recovery	0.9-10.0 %
	Platelet-large cell count (P-LCC)	30-90 x10 ⁹ /L	Normal, or slightly increased	Progressively increasing, then going down during recovery	30-90 x10 ⁹ /L
C R P	Full-range C-reactive protein (FR-CRP)	0.00-4.00 mg/L	Slightly increased	Progressively increasing, CRP > 34mg/L plus age > 60 years indicating high probability of mortability in 12 days ⁷	0.00-4.00 mg/L

^{*}Symtoms: fever, cough, breathing difficulties, headache, diarrhea

CLIA Test Indices

Test Parameter	Reference Interval	Clinical Significance	Notes	
cTnl	99th Percentile: 0.04 ng/mL cutoff of CL-series TnI assay for determining AMI is 0.5 ng/mL	 Acute cardiac injury (elevation of cTnl to >99th percentile) is associated with more severe COVID-19 disease^{1,9,10}. COVID-19 patients with elvated cTnl is associated with worse prognosis¹¹. 	Assays for troponins and BNP should be obtained only for such patients who have clinical signs of acute myocardial infarction (MI) or heart failure, respectively.	
BNP	Cutoff of CL-series BNP assay for determining heart failure is 100 pg/mL	 For COVID-19 patients with clinical evidence for heart failure, BNP could aid in the diagnosis of congestive heart failure¹². Natriuretic peptide estimates a potentially useful approach in patients with severe COVID-19 to distinguish between cardiac and pulmonary cause of dyspnea, to risk-prognosticate the patients, and to guide and monitor therapy¹³. 		
PCT	<0.05 ng/mL	 SARS-CoV-2 infection could not cause an elevated PCT concentration. Procalcitonin is typically normal on admission, but may increase among those admitted to the ICU¹⁴. Increased PCT is a risk factor associated with in-hospital death, and of the COVID-19 patients who died around half had a secondary bacterial infection leading to sepsis and death¹. 	PCT is not a substitute for good clinical judgement and cannot be used in isolation.	
FERR	Male: 27-375 ng/mL, Female: 12-135 ng/mL	 Increased ferritin level might correlate to secondary bacterial infection and associated with poor clinical prognosis¹⁵. Elevated ferritin is a marker to assess COVID-19 severity¹⁶. An elevated serum ferritin test is an indicator for systemic inflammatory response syndrome, which could cause rapid deterioration of COVID-19 patients¹⁷. 		
IL-6*	/	 In patients with COVID-19, IL-6 levels are significantly elevated and associated with adverse clinical outcomes¹⁸. Given the association of elevated IL-6 with severe COVID-19 and mortality, clinicians should use this as a potential marker to recognize severe disease¹⁹. 	Available soon on Mindray CLIA systems	

Biochemistry Test Indices

Test Parameter	Reference Interval	Clinical Significance	
LDH	Male: < 248 U/L , < 4.13 μkat/L Female: < 247 U/L, < 4.12 μkat/L	LDH level can reflect the injury degree of lung, kidney and heart. Most COVID-19 patients had elevated LDH. ¹⁸ LDH level can help evaluate the prognosis and predict the mortality of COVID-19 patients.	
CRP	<5.0mg/L	The significant increase of CRP has been reported in most COVID-19 patients. ¹⁹ CRP testing is useful in the evaluation of coronavirus infection and reflect the inflamation status.	
FER	Male: 30-400 ng/mL Female: 15-150 ng/mL	 All patients with severe COVID-19 should be screened for hyperinflammation using laboratory trends (eg, increasing ferritin, decreasing platelet counts, or increasing erythrocyte sedimentation rate). The serum ferritin has a role in predicting in-hospital mortality. Levels of serum ferritin was clearly elevated in non-survivors compared with survivors throughout the clinical course, and increased with illness deterioration.¹ 	
D-Dimer	≤1.0 μg/mL	Many studies found that D-dimer levels significantly increased with increasing severity of COVID-19. Having D-dimer level greater than 1 μg/mL is a factor that could help clinicians to identify patients with poor prognosis at an early stage.¹	

^{**}Discharge: Under the premise that the patient's nucleic acid test result is negative for two consecutive days (alveolar lavage fluid is recommended⁸)

References:

- [1]. Fei Zhou, Ting Yu, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19, a retrospective cohort study. Lancet (2020). doi: 10.1016/S0140-6736(20)30566-3 [2]. Jingyuan Liu, Yao Liu, Pan Xiang, et al. Neutrophil-to-Lymphocyte Ratio Predicts Severe Illness Patients with 2019 Novel Coronavirus in the Early Stage. Medrxiv. doi:
- 10.1101/2020.02.10.20021584
- [3]. Jin-jin Zhang, Xiang Dong, Yi-yuan Cao, et al. Clinical characteristics of 140 patients infected with SARSCoV-2, Allergy. 2020 Feb 19. doi: 10.1111/all.14238.
- [4]. Wang CZ, NLR&RDW-SD: Indices for Identifying Severe COVID-19 Patients (to be published officially). https://www.mindray.com/en/presscenter/NLR_RDW-SD__Indices_for_Identifying_Severe_COVID-19_Patients.html
- [5]. Ephrem G., Red Blood Cell Distribution Width Should Indeed Be Assessed with Other Inflammatory Markers in Daily Clinical Practice. Cardiology. 2013;124(1):61. doi: 10.1159/000345925.
- [6]. G. Lippi, M. Plebani, B. Michael Henry, Thrombocytopenia is associated with severe coronavirus disease 2019 (COVID-19) infections: A meta-analysis, Clinica Chimica Acta (2020). doi: 10.1016/j.cca.2020.03.022
- [7]. Jiatao Lu, Shufang Hu, Rong Fan, et al. ACP risk grade: a simple mortality index for patients with confirmed or suspected severe acute respiratory syndrome coronavirus 2 disease (COVID-19) during the early stage of outbreak, medRxiv. doi: 10.1101/2020.02.20.20025510
- [8]. Xiao-Hong Yao, Zhi-Cheng He, Ting-Yuan Li, Hua-Rong Zhang, et al. Pathological evidence for residual SARS-CoV-2 in pulmonary tissues of a ready-for-discharge patient. Cell Research (2020) 0:1–3. doi: 10.1038/s41422-020-0318-5
- [9]. Wang D., Hu B., Hu C., et al. Clinical characteristics of 138 hospitalized patients with 2019 novel coronavirus-infected pneumonia, JAMA, 2020.
- [10]. Lippi G., Lavie C.J., Sanchis-Gomar F. Cardiac troponin I in patients with coronavirus disease 2019 (COVID-19): Evidence from a meta-analysis. Prog Cardiovasc Dis, 2020.
- [11]. Ruan Q., Yang K., Wang W., Jiang L., Song J. Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients, Intensive Care Med, 2020.
- [12]. James L. Januzzi Jr. American College of Cardiology: Troponin and BNP use in COVID-19.
- [13]. Kunal Mahajan, Prakash Chand Negi. The role of natriuretic peptide estimation in severe COVID-19. Monaldi Archives for Chest Disease, 2020.
- [14]. Interim Clinical Guidance for Management of Patients with Confirmed Coronavirus Disease (COVID-19). US CDC.
- [15]. Bo Zhou, et al. Utility of Ferritin, Procalcitonin, and C-reactive Protein in Severe Patients with 2019 Novel Coronavirus Disease.
- [16]. Clinical guide for the management of critical care for adults with COVID-19 during the coronavirus pandemic. UK NHS.
- [17]. Colafrancesco S, et al. COVID-19 gone bad: A new character in the spectrum of the hyperferritinemic syndrome?. Autoimmun Rev. 2020.
- [18]. Coomes E, et al. Interleukin-6 in COVID-19: A Systematic Review and Meta-Analysis. doi: https://doi.org/10.1101/2020.03.30.20048058
- [19]. Aziz M, Fatima R, Assaly R. Elevated Interleukin-6 and Severe COVID-19: A Meta-Analysis. J Med Virol. 2020.

Mindray's total laboratory solutions assisted frontline health workers in fighting COVID-19 in China's Leishenshan and Huoshenshan Hospitals, two emergency specialty filed hospitals built in mere a few days in response to the coronavirus.















