



## STATEMENT OF COMPLIANCE TO THE TECHNICAL SPECIFICATIO

## PROJECT: ANTIGEN FOR TESTING SERVICE FOR PARTICIPANTS AND GUEST RE: TPB STRATEGIC PLANNING AND HARMONIZATION WORKSHOP (PR No. 10.054)

## Quotation No. TPB-PR.2022.11.427

## [Bidders must state here either "Comply" or "Not Comply" against each of the individual parameters of each Specification]

ITEM	SPECIFICATION	STATEMENT OF COMPLIANCE (COMPLY/NOT COMPLY)
1	INDICATIVE DATES: 12 December 2022 (target dates of implementation) NO. OF PARTICIPANTS: 73 participants (indicative)	
SPECIF	ICATIONS – First Schedule	
2	<ul> <li>Required Service</li> <li>a. The Service Provider shall conduct on-site testing for 60 personnel who are participants to be TPB 4-day Strategic Planning and Harmonization Workshop at the TPB Office using the SARS-Cov-3 Antigen Rapid Diagnostic Test Kits.</li> <li>b. The schedule is to be determined and set by TPB with an indicative date of 12 December 2022.</li> <li>c. Site and time of the Antigen Testing will be at the designated TPB office premises</li> </ul>	
3	General Specifications a. The Service Provider must have a valid license to Operate issued by the Department of Health (DOH) following the Standards and Requirements of Administrative Order (AO) No. 2007-0027. b. The Service Provider shall designate and provide an appropriate number of duly licensed or trained medical professionals to facilitate the Antigen Testing for 60 of TPB's personnel with a names list to be provided. c. The Service Providers' deployed medical team shall submit their respective negative results for RT-PCR COVID-19 test at least one (1) day prior to the conduct of onsite testing	

	d. The Service Providers' deployed medical team shall wear	
	their company's uniform and I.D. and designate amongst the team	
	a coordinator to liaise with the assigned TPB Project Officer	
	e. During the conduct of testing and at all points of contact	
	with TPB personnel, the medical team shall wear the complete	
	prescribed Personal Protective Equipment (PPE).	
4	General Specifications	
	f. The Service Provider must have a valid license to Operate issued	
	by the Department of Health (DOH) following the Standards and	
	Requirements of Administrative Order (AO) No. 2007-0027.	
	g. The Service Provider shall designate and provide an appropriate	
	number of duly licensed or trained medical professionals to facilitate the	
	Antigen Testing for 60 of TPB's personnel with a names list to be provided.	
	h. The Service Providers' deployed medical team shall submit their	
	respective negative results for RT-PCR COVID-19 test at least one (1) day	
	prior to the conduct of onsite testing	
	i. The Service Providers' deployed medical team shall wear their	
	company's uniform and I.D. and designate amongst the team a coordinator	
	to liaise with the assigned TPB Project Officer	
	j. During the conduct of testing and at all points of contact with TPB	
	personnel, the medical team shall wear the complete prescribed Personal	
	Protective Equipment (PPE).	
5	Technical Requirements	
5	a. The Service Provider shall ensure to only use FDA	
	registered testing kits, reagents and devices for SARS-COV2 Rapid	
	Antigen Testing	
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	of at least 90% and clinical specificity of at least 97%.	
	c. The Service Provider must bring on-site adequate	
	number of testing kits, PPEs for their health care providers	
	and other consumables necessary for the testing services. The	
	Service Provider shall ensure that it has an adequate contingency	
	measure/plan to cover for any and all testing kit errors.	
	d. The SARS-COV2 Rapid Ag testing shall be collection of	
	specimens through nasopharyngeal and/or oropharyngeal	
	swabbing.	
	e. The Service Provider must ensure proper handling	
	and storage of these specimens.	
	f. The Service Provider shall take charge of the disposal of	
	sharps and other biohazardous wastes from the on-site testing	
	area.	
	g. Testing kit processing time to display result shall be	
	within thirty (30) minutes	
	h. The Service Provider must provide within two hours of	
	the conduct of antigen testing all test results, with certificates	
	issued within the same day via digital copies; hardcopies are	
	provided to clients upon request, with NO extra charge. Results	
	are to be provided via email.	

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	<ul> <li>The result must express in qualitative terms (example: positive, negative, reactive, or non-reactive).</li> <li>The Service Provider must maintain all medical results and other information in strict confidence. The Service Provider must not disclose documents and information unless authorized by TPB or the employee concerned and as</li> </ul>	
	required by a government health authority. Zero tolerance shall	
	be enforced for any violations to data privacy.	
	SPECIFICATIONS – Second Schedule	
6	Required Service a. The Service Provider shall conduct on-site testing for 15 personnel who are participants to the TPB 4-day Strategic Planning and Harmonization Workshop at the designated holding room or lounge of the designated event hotel within Clark Freeport and Special Economic Zone, Pampanga. b. The schedule is to be determined and set by TPB with an indicative date of 13 December 2022.	
	c. Site and time of the Antigen Testing is to be determined by TPB and limited to within Clark Freeport and Special Economic Zone, Pampanga	
7	General Specifications	
	<ul> <li>a. The Service Provider must have a valid license to Operate issued by the Department of Health (DOH) following the Standards and Requirements of Administrative Order (AO) No. 2007-0027.</li> <li>b. The Service Provider shall designate and provide an appropriate number of duly licensed or trained medical professionals to facilitate the Antigen Testing for 15 of TPB's VIP Guests with a names list to be provided.</li> <li>c. The Service Providers' deployed medical team shall submit their respective negative results for RT-PCR COVID-19 test at least one (1) day prior to the conduct of onsite testing</li> <li>d. The Service Providers' deployed medical team shall wearing their company's uniform and I.D. and designate amongst the team a coordinator to liaise with the assigned TPB Project Officer</li> <li>e. During the conduct of testing and at all points of contact with TPB personnel, the medical team shall wear the complete prescribed Personal Protective Equipment (PPE).</li> </ul>	
8	<ul> <li>Technical Requirements <ul> <li>a. The Service Provider shall ensure to only use FDA registered</li> <li>testing kits, reagents and devices for SARS-COV2 Rapid Antigen Testing</li> <li>b. All testing kits to be used shall be with a clinical sensitivity of at</li> <li>least 90% and clinical specificity of at least 97%.</li> <li>c. The Service Provider must bring on-site adequate number of</li> <li>testing kits, PPEs for their health care providers and other consumables</li> <li>necessary for the testing services. The Service Provider shall ensure that it</li> <li>has an adequate contingency measure/plan to cover for any and all testing</li> <li>kit errors.</li> <li>d. The SARS-COV2 Rapid Ag testing shall be collection of specimens</li> <li>through nasopharyngeal and/or oropharyngeal swabbing.</li> </ul> </li> </ul>	

	e. The Service Provider must ensure proper handling and	
	storage of these specimens.	
	f. The Service Provider shall take charge of the disposal of sharps	
	and other biohazardous wastes from the on-site testing area.	
	g. Testing kit processing time to display result shall be within	
	thirty (30) minutes	
	h. The Service Provider must provide within 30 minutes of the	
	conduct of antigen testing all test results, with certificates issued within 3	
	hours of the same day via digital copies; hardcopies are provided to clients	
	upon request, with NO extra charge. Results are to be provided via email.	
	i. The result must express in qualitative terms (example: positive,	
	negative, reactive, or non-reactive).	
	j. The Service Provider must maintain all medical results and other	
	information in strict confidence. The Service Provider must not	
	disclose documents and information unless authorized by TPB or the	
	employee concerned and as required by a government health authority.	
	Zero tolerance shall be enforced for any violations to data privacy.	
9	ELIGIBILITY REQUIREMENTS	
	1. Must comply with the legal and technical and other requirements	
	under R.A. 9184 and its Revised Implementing Rules and Regulations	
	2. Must be a Department of Health accredited medical laboratory /	
	clinic / testing facility	
	3. Must be accredited with the Philippine Government Electronic	
	Procurement System (PHILGEPS);	
	4. Must be willing to do an onsite Antigen testing at the TPB Office and	
	TPB identified testing site	
	5. Either has an existing credit line with TPB or would allow send-bill	
	arrangement.	
10	III. SPECIAL AND SPECIFIC CONDITIONS	
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	1. Should the indicative dates stated herein be not workable, then the	
	actual dates may be worked out by both parties subject to mutual	
	availability and agreement without changes to the other specifications	
	stated in this Term of Reference.	
1	2. Force Majeure:	
	If and to the extent that a Party's performance of any of its obligations	
	pursuant to this Agreement is prevented, hindered or delayed directly or	
	indirectly by fire, flood, earthquake, elements of nature or acts of God, acts	
	of war, terrorism, riots, civil disorders, rebellions or revolutions, or any	
	other similar cause beyond the reasonable control of such Party (each a	
	"Force Majeure Event"), and such non-performance, hindrance or delay	
	could not have been prevented by reasonable precautions, then the non-	
	performing, hindered or delayed Party shall be excused for such	
	nonperformance, hindrance or delay, as applicable, of those obligations	
	affected (the "Affected Services") by the Force Majeure Event for as long as	
	the Force Majeure Event continues and, except as otherwise provided in	

	this Section, such Party continues to use its commercially reasonable efforts	
	to recommence performance whenever and to whatever extent possible	
	without delay, including through the use of alternate sources, workaround	
	plans or other means. The Party whose performance is prevented, hindered	
	or delayed by a Force Majeure Event shall promptly notify the other Party of	
	the occurrence of the Force Majeure Event and describe in reasonable	
	detail the nature of the Force Majeure Event.	
	The Impacted Party shall give Notice within 14days of the Force Majeure	
	Event to the other party, stating the period of time the occurrence is	
	expected to continue. The Impacted Party shall use diligent efforts to end	
	the failure or delay and ensure the effects of such Force Majeure Event are	
	minimized. The Impacted Party shall resume the performance of its	
	obligations as soon as reasonably practicable after the removal of the cause.	
	In the event that the Impacted Party's failure or delay remains uncured for a	
	period of 90 days following Notice given by it, the other party may	
	thereafter terminate the contract agreement specific to this Terms of	
	Reference upon Notice.	
	Further, TPB and the Service Provider hereby acknowledge that while	
	current events related to the Covid-19 pandemic are known, future impacts	
	of the outbreak are unforeseeable and shall be considered a Force Majeure	
	event to the extent that they prevent the performance of a Party's	
	obligations under this Terms of Reference.	
11	TERMS OF PAYMENT:	
	1. Send bill arrangement.	
	2. Preferably has a Landbank account. Payment will be made through	
	LBP bank deposit	
	3. In case the supplier does not have a Landbank account, bank	
	charges will be shouldered by the supplier	
	For particulars, please contact:	
	WILSON R. SUBA	
	Telephone numbers: (02) 525-9318 to 27 local 209 or (02) 525-6443	
	Email address: wilson_suba@tpb.gov.ph	
	SHERYLL KARUNUNGAN	
	Telephone numbers: (02) 525-9318 to 27 local 209 or (02) 525-6443	
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I hereby certify to comply and deliver all of the above requirements.

Name of Company

Signature over Printed Name of Authorized Representative