



Republic of the Philippines Department of Health OFFICE OF THE SECRETARY

FEB 1 1 2011

DEPARTMENT MEMORANDUM NO. 2011 - 0039

TO

ALL CENTERS FOR HEALTH DEVELOPMENT (CHD), PROVINCIAL HEALTH OFFICES (PHO), CITY HEALTH OFFICES (CHO), MUNICIPAL HEALTH OFFICES (MHO)

AND CHIEFS OF HOSPITALS

SUBJECT:

Strengthening Active Surveillance for Acute Flaccid Paralysis

(AFP) in all Government and Private Hospitals

On April 23, 2010, an outbreak of poliomyelitis has occurred in the country of Tajikistan. The poliovirus was traced from Uttar Pradesh, India. The outbreak resulted to 458 paralytic cases and 27 deaths. This was the first outbreak of polio in the European Region since 2002 when the region was declared polio-free by the World Health Organization (WHO). The experience in Tajikistan could also happen in the Philippines. The wild poliovirus could be easily imported to the Philippines due to access in international travels and Overseas Filipino Workers working in polio endemic countries (Pakistan, Afghanistan, India and Nigeria).

The Philippines is one of the high risk countries in the Western Pacific Region for poliovirus importation due to pockets of low OPV immunization coverage and inadequate AFP surveillance. On November 2-3, 2010, the RCC conducted an assessment in the Philippines. The RCC found major deficiencies in the AFP surveillance.

In view of this, the following activities are recommended by the RCC to enhance AFP active surveillance:

A. Detection and Reporting of cases

- 1. All directors of government and private hospitals should designate at least one Disease Surveillance Coordinators (DSC) that performs daily case-finding of AFP and other diseases under the Philippine Integrated Disease Surveillance and Response (PIDSR) system in hospital wards.
- 2. The DSCs should submit weekly notifiable disease reports to the RESU/PESU. This report should include zero reporting of AFP.
- 3. The regional health office may designate surveillance staff to non-functioning PESU/CESUs to assist in the surveillance operation.

- 4. DSCs and surveillance officers should have access to medical records and should be allowed to obtain a copy of the pertinent records of AFP cases.
- 5. All AFP cases should have a complete history-taking, physical examination and neurological assessment indicated in the chart.
- 6. In instances where the DSC or the Disease Surveillance Officer (DSO) performed the neurological examination, the attending physician must confirm the result by affixing his/her signature on the Case Investigation Form (CIF). (See attached CIF)
- 7. If the case was detected in the community or health center, the MHO shall perform or confirm the result of the neurological exam.

B. Laboratory confirmation of cases

- 1. As soon as an AFP case is detected, **two (2) stool specimens** should be collected at 24-48 hours interval, **within 14 days from the onset of paralysis.**
- 2. Each stool specimen should be properly labeled with the patient's name, date and time of collection and stool number
- 3. After each collection, specimen should be kept in refrigerator. **DO NOT place the stool specimen in the freezer.** If a refrigerator is not available, the stool specimen can be stored in a transport box with at least 4 frozen ice packs. The ice packs should be replaced every 24 hours.
- 4. All stool specimens should be sent to the Research Institute for Tropical Medicine (RITM) within 3 days after collection with a completely filled-up PIDSR CIF.
- 5. RITM should be notified if the specimens are expected to arrive in the laboratory on a Friday, during weekends or holidays.

C. Monitoring and Supervision

- 1. The regional surveillance staff should conduct at least monthly monitoring of all retained and other tertiary hospitals that directly report to RESU. A standard monitoring tool for ESU and hospitals that was developed by the National Epidemiology Center (NEC) should be used for this activity. (See attached monitoring forms)
- 2. The DSOs should conduct weekly monitoring and supervision to hospitals that no longer submit weekly report for 3 consecutive weeks or "silent" Disease Reporting Units (DRUs). This activity should also be done by provincial and city surveillance staff.

D. Feedback

- 1. The RESU should provide monthly VPDs monitoring report to all DRUs and ESUs. This includes the following:
 - a. Completeness of reporting (number of DRUs/ESU reports over the total number of DRUs)
 - b. Timeliness of reporting by DRU (number of DRUs/ESU submits report on a weekly basis over total number of DRUs)
- 2. The RESU should give monthly AFP Surveillance Reports to the regional Expanded Program on Immunization (EPI) manager, hospitals and ESUs on the following indicators:
 - a. AFP Reporting Rate
 - b. Non-polio Rate

- c. Percentage of timely reporting
- d. Percentage of timely investigation
- e. Percentage of Adequate Stool Specimen Collection
- f. Timely 60-day follow-up
- g. NPEV (Non-polio enterovirus) Rate
- 3. The RESU should provide feedback of the laboratory results to the DRU.

By Authority of the Secretary of Health:

DAVID J. LOZADA, JR., MD, MPA, CESO I

Undersecretary of Health Health Service Delivery



Case Investigation Form



Acute Flaccid Paralysis

Name of D	RU:				Type	e: 🗆 RHU 🗀	CHO □Gov't	Hospi	ital ∏Pri	vate Hospi	tal □Clinic
Address:							oratory □F	-		•	ort/Seaport
I. PATIEN	T MATION:	Patient Num	ber:	Patient's First	Name	9	Middle Na	ame	•	La	st Name
Complete A	Address:		Į.			Sex: □Male	Date of Birth:	<u>MM</u>	<u>DD YY</u>	Age:	□Days □Months
District:		ILHZ:				□Female		/_	/	_	□Years
Patient Adn	nitted? □Ye	s 🗆 No 🖂	Jnknowr	<u> </u>		Date Admitt	ed/ Seen/Cons	ult	<u>MM</u>	<u>DD</u>	YY
Date of Rep	port:		<u>MM</u>	<u>DD</u>	<u>YY</u>	Date of Inve	estigation:		<u>MM</u>	<u>DD</u>	YY
Name of In	vestigator:						Contact Nos.:				
II. CLINIC	AL DATA (I	Put a check	[$$] in	the appropr	riate b	ox)		Salarin Salarin	e (espe	1.470	
PROD	PROME	F	ARALYS	SIS	SIT	E OF FLACCID	PARALYSIS		ensory tatus	Deep Tendon Reflexes	Motor Status
Fover C	/ ON OU	Date onset:			Right	arm: 🗆 Y 🗆 I	N DU				
				Y 🗆 N 🗆 U	Left a	arm: 🗆 Y 🗆 f	N 🗆 U				_
Diarrhea/V				Y DN DU	Right	ileg: □Y□I	V 🗆 U				
			OGRES:		Left I	eg: □Y□1	V 🗆 U		TT 1 1 1 1		
Muscle pai		Paralysis ful to 14 days fi	-		Breat	thing muscles:[JY DN DU				e grading/ atus, deep
Meningeal		DY DN I			Neck		JY DN DU	tena	lon reflexe	s and mot	or status are
OY ON	ΠU	Direction of	paralysi	s:	Facial muscles:					tnis page.	
		☐ Ascendin	_	escending	Work	king Diagnosis	5 :				
III EDIDE	MIOLOGIC	Unknown				111 (412)	. V F. S. S.		.aa		-
				<u> </u>	- 51 1 gr -	<u> </u>	e transference	- 1994			
	_				-	ify disorder:					
1						ess? 🗆 Y 🗆 I	veled: From	,	,	To /	,
				in 60 days of		Date tra i's paralysis? □		'		10/	
1	•		-	-	-	nal bite?□Y					
'	ecify type:	, ,	•								
	NIZATION H	1.00									
Total OPV/	IPV doses re	ceived:		Date last	dose c	of OPV :/		ls	s this a "Ho	ot case"?	JY 🗆 N
V. LABOI	RATORY DA	\TA			- 4		The second secon				
Stool sample #	Collected?	If YES, o		ate sent to	Date	received RITM		Re	sult		Date result
1	DYDN	/	_/				☐ NEG ☐ 1 ☐ Other, spe				
2	□Y □N	/					□ NEG □ 1 □ Other, spe				
Adequate?	□Y □N	Other Info	rmation ((Stool specime	en):						
VI. 60-DA	Y FOLLOW	-UP			in Roje		ing.				t mendet til gengen i som Late til store en er
P.E. done?	□Y□N lf	NO, reason fo days?: □ Y □	or no P.E		□Pa	w-up conducted tient died Date:	_/_/_ □ Lo	st to fol		Other, spec	

Case Investigation Form

Acute Flaccid Paralysis

VII. CLASSIFICATION (TO BE FILLED UP BY THE EXPERT PANEL ONLY)					
⊊FINAL CLASSIFICATION	IF VAPP	CLASSIFICATION CRITERIA	FINAL DIAGNOSIS		
☐ Confirmed wild polio	☐ Recipient VAPP	☐ Laboratory			
☐ Vaccine-derived paralytic polio (VDPV)	☐ Contact VAPP	☐ EPI linked			
☐ Vaccine-associated paralytic polio (VAPP)	□ Unknown	☐ Lost to follow-up			
☐ Polio compatible		☐ Death			
☐ Discarded		☐ With residual paralysis			
☐ Not AFP		☐ Without residual paralysis			
□ Sabin-like					
Date classified://					

<u> AFP Case definition:</u>

- Any child less than 15 years of age with acute flaccid paralysis, OR
- A person of any age in whom poliomyelitis is suspected by a physician.

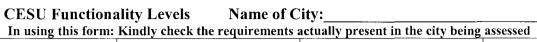
Hot Case Description:

- An AFP case that is <5 years old with < 3 doses of OPV and has fever at the onset of asymmetrical paralysis,
 OR
- An AFP case or a person of any age whose stool specimen(s) has poliovirus isolate.

Grading/Scoring of Sensory Status, Deep Tendon Reflexes and Motor Status:

- A. Sensory status is presented in percentage and categorized as follows:
 - ≤ 25% = Absent
 - ≥ 25% but <100% = Reduced
 - 100% = Normal
- B. Deep tendon reflexes (DTRs) are presented in (+) symbol and categorized as follows:
 - none or 0 = absent
 - + = reduced
 - ++ = normal
 - +++ with/without clonus = increased or exaggerated
- C. Motor status is presented in fraction and categorized as follows:
 - 0/5 = absent or no movement
 - 1/5 to 3/5 = reduced movement (with movement but not against resistance or gravity)
 - 4/5 to 5/5 = normal (movement with full resistance and against gravity)





	للشفائق		100	
Ŕ			-70	M.
e de	- 1	(Ň	
6	w.		14.1	
- 6		·	-//	
	<u> </u>		4	.
		9 1		н

0	1	2	3
Presence of Ordinance	e or Executive Order		
□ NO	□ YES	2004 2004	
Manpowe	er Present		
☐ One Designated	☐ Two Designated	☐ Three Designated	□ > Than Three Trained
Personnel Trained in	Personnel Trained in	Personnel Trained in	Designated Personnel and
PIDSR	PIDSR	PIDSR	Epidemiologist
	Presence of Office Space		
□ NO	☐ SHARED	☐ DEDICATED	
	Presence of	Computer	
□ NO	☐ SHARED	□ DEDICATED	☐ DEDICATED with
			WORKSTATION
Acc	ess to Transportation Facil	ities	
□ NO	☐ SHARED	☐ DEDICATED	And the second s
	Access to Commu	nication Facilities	
□ NO FAX NO	☐ SHARED FAX OR	☐ DEDICATED FAX	☐ DEDICATED FAX
INTERNET	INTERNET	OR INTERNET	AND INTERNET
	Presence of All	located Budget	
□ NO	☐ EXTRACTED FROM	☐ LINED UNDER	☐ DIRECT FROM LGU
	OTHER NON-HEALTH	HEALTH OFFICE/UNIT	BUDGET SEPARATE
	OFFICE/UNIT OF THE		FROM HEALTH
	LGU		UNIT/OFFICE
Presence of Work	and Financial Plan	The second of the second	
□ NO	☐ YES	The state of the s	
	Presence of Surve	eillance Activities	
□ NO CAID (collection,	☐ CAIDs data from	☐ CAIDs data from	☐ CAIDs data from
analysis, interpretation	sentinel sites and provides	sentinel sites and provides	sentinel sites and provides
and dissemination) of	feedback to stakeholders	feedback to stakeholders	feedback to stakeholders
data	irregularly	monthly	weekly
		of Laboratory Services	
□ NO	☐ Sends lab specimens	-	☐ Sends lab specimens
	to reference labs (i.e.	to labs within the region	to labs within the
	RITM) for confirmation	for confirmation	province for confirmation
	raining Activities pertainin		
□ NO	☐ YES but limited to	☐ YES and extended to	☐ YES and integrated to
	activities focused on	activities involving the	activities involving the
	surveillance and response	region	national programs' thrusts
	developed for local wide		
	use only		
		utbreak Investigations	T
□ NO	☐ YES but with the help	☐ YES but with the help	☐ YES and does
	of the national experts	of the regional experts	investigations involving
			local experts only

Scale: Add the scores of the checked boxes to get the functionality score of the CESU. The corresponding scores (0,1,2,3) are at the heads of the columns.

Example: If the CESU has an ordinance, you will check YES and the corresponding score is 1.

- 0: Your local health office doesn't have an ESU. All you have is a designated employee equivalent to a Disease Surveillance Coordinator under the PIDSR framework.
- 1 to 11: Your Local Health Office is pursuing efforts to build an ESU through shared resources and capacities with the other units in your office. Try to maximize all your efforts to achieve the category of Level 1 ESU.
- 12: Your ESU can be categorized as Level 1. Your ESU exists through shared resources and capacities. Try to maximize all your efforts to achieve the category of Level 2 ESU.
- 13 to 21: Your Level 1 ESU is already achieving resources and capacities necessary to reach the category of Level 2 ESU. Continue with your work to achieve the category of Level 2 ESU.
- 22: Your ESU can be categorized as Level 2. Your ESU exists through dedicated resources and capacities. Try to maximize all your efforts to achieve the category of Level 3.
- 23 to 29: Your Level 2 ESU is already achieving resources and capacities necessary to reach the category of Level 3 ESU. Continue with your work to achieve the category of Level 3 ESU.
- 30: Your ESU is now Level 3. Your local health office had achieved the ideal qualities of a functional ESU fit for your office. Kindly maintain the functionality for the benefit of your local government unit and the PIDSR system.

Definitions:

- 1. Presence of Ordinance or Executive Order: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of ordinance or executive order (printed document) pertaining to the creation of a functional ESU in their province, check the YES cell. Otherwise, check the No cell.
- 2. Designated Personnel: Refers to the personnel assigned FULL TIME in ESU doing FULL TIME ESU work
- 3. Shared Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is NOT SOLELY or NOT for the ESU alone
- 4. Dedicated Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is SOLELY or for the ESU alone
- 5. Presence of Work and Financial Plan: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of work and financial plan (printed document) pertaining to the functional ESU in their province, check the YES cell. Otherwise, check the No cell.



A CONTRACTOR OF THE PARTY OF TH	A
9.	
	-

DESU Functions			PUSE
1n using this form:	Kindly check the requirements a	actually present in the district be	ing assessed
	ce or Executive Order		
□ NO	☐ YES	ATT STATE OF THE S	
	er Present		
☐ One Designated	☐ Two Designated	☐ Three Designated	☐ > Than Three Trained
Personnel Trained in	Personnel Trained in	Personnel Trained in	Designated Personnel and
PIDSR	PIDSR	PIDSR	Epidemiologist
	Presence of Office Space		(A)
□ NO	☐ SHARED	☐ DEDICATED	
		f Computer	<u> </u>
□ NO	☐ SHARED	□ DEDICATED	□ DEDICATED with
			WORKSTATION
Acc	ess to Transportation Facil	ities	
□ NO	□ SHARED	☐ DEDICATED	and the second s
	. ,,,,,	nication Facilities	
□ NO FAX NO	☐ SHARED FAX OR	☐ DEDICATED FAX	□ DEDICATED FAX
INTERNET	INTERNET	OR INTERNET	AND INTERNET
II (I DIC (E)		located Budget	THIS HITEKILET
□ NO	☐ EXTRACTED FROM	☐ LINED UNDER	☐ DIRECT FROM LGU
	OTHER NON-HEALTH	HEALTH OFFICE/UNIT	BUDGET SEPARATE
	OFFICE/UNIT OF THE		FROM HEALTH
	LGU		UNIT/OFFICE
Presence of Work	and Financial Plan	At July 1	
□ NO	☐ YES	CONTRACTOR AND	All the second s
	Presence of Surv	eillance Activities	I and the second
□ NO CAID (collection,	☐ CAIDs data from	☐ CAIDs data from	☐ CAIDs data from
analysis, interpretation	sentinel sites and provides	sentinel sites and provides	sentinel sites and provides
and dissemination) of	feedback to stakeholders	feedback to stakeholders	feedback to stakeholders
data	irregularly	monthly	weekly
	Access and Availability	of Laboratory Services	-
□ NO	☐ Sends lab specimens	☐ Sends lab specimens	☐ Sends lab specimens
	to reference labs (i.e.	to labs within the region	to labs within the
	RITM) for confirmation	for confirmation	province for confirmation
Presence of T	raining Activities pertainin	g to Surveillance and Resp	onse Activities
□ NO	☐ YES but limited to	☐ YES and extended to	☐ YES and integrated to
	activities focused on	activities involving the	activities involving the
	surveillance and response	region	national programs' thrusts
	developed for local wide		
	use only		
	T	utbreak Investigations	
□ NO	☐ YES but with the help	☐ YES but with the help	☐ YES and does
	of the national experts	of the regional experts	investigations involving
			local experts only

Scale: Add the scores of the checked boxes to get the functionality score of the DESU. The corresponding scores (0,1,2,3) are at the heads of the columns.

Example: If the DESU has an ordinance, you will check YES and the corresponding score is 1.

- 0: Your local health office doesn't have an ESU. All you have is a designated employee equivalent to a Disease Surveillance Coordinator under the PIDSR framework.
- 1 to 11: Your Local Health Office is pursuing efforts to build an ESU through shared resources and capacities with the other units in your office. Try to maximize all your efforts to achieve the category of Level 1 ESU.
- 12: Your ESU can be categorized as Level 1. Your ESU exists through shared resources and capacities. Try to maximize all your efforts to achieve the category of Level 2 ESU.
- 13 to 21: Your Level 1 ESU is already achieving resources and capacities necessary to reach the category of Level 2 ESU. Continue with your work to achieve the category of Level 2 ESU.
- 22: Your ESU can be categorized as Level 2. Your ESU exists through dedicated resources and capacities. Try to maximize all your efforts to achieve the category of Level 3.
- 23 to 29: Your Level 2 ESU is already achieving resources and capacities necessary to reach the category of Level 3 ESU. Continue with your work to achieve the category of Level 3 ESU.
- 30: Your ESU is now Level 3. Your local health office had achieved the ideal qualities of a functional ESU fit for your office. Kindly maintain the functionality for the benefit of your local government unit and the PIDSR system.

Definitions:

- 1. Presence of Ordinance or Executive Order: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of ordinance or executive order (printed document) pertaining to the creation of a functional ESU in their province, check the YES cell. Otherwise, check the No cell.
- 2. Designated Personnel: Refers to the personnel assigned FULL TIME in ESU doing FULL TIME ESU work
- 3. Shared Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is NOT SOLELY or NOT for the ESU alone
- 4. Dedicated Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is SOLELY or for the ESU alone
- 5. Presence of Work and Financial Plan: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of work and financial plan (printed document) pertaining to the functional ESU in their province, check the YES cell. Otherwise, check the No cell.



Department of Health NATIONAL EPIDEMIOLOGY CENTER



ASSESSMENT FORM for the VPDISS Surveillance Operations (DRUs)

Region:			Province	Date:	
Name of DRU: Name of Respondent:				Address: Position/Designation:	
	1110		SIIDVEILI	ANCE CORE FUNCTIONS	
I.	СА	SE DETECTION	REGISTRATION & CONFIRMATION		
···			<u> </u>		
	1.		designated surveillance point persor etects VPD cases?	n/DSC?	YES NO
			spital order designating you or this p	norman.	
			inance/office order)	person	YES NO
		c. How many sta	•		1-2 3 & above
		d. Do your surve Neonatal Teta	eillance staff conduct active search anus and AEFIs?	for AFP, Measles,	YES NO
			y suspect cases did you detect from ving (previous 2 morbidity weeks during		AFP NT NEFIS
	2.	How do you dete	ect VPD cases?		
			Monitor the ER/	Review disease registers	-
			Make bedside	Others,	
	2	llaurda varileel	Review medical		
	3.	a. AFP	k for cases of the following VPDs?		— P.W
		d. AFF	<15 years old	a case of any age diagnosed as polio by a physician	Differential diagnosis
			weakness/flaccid or floppy paralysis	sudden onset (3-4 days; max. of 2 weeks)	Others,
		b. Measles	Fever/hot to touch	Cough/coryza/ conjunctivitis	Others,
			Generalized non-vesicular rash of at least 3 days	Differential Dx:	_
		c. Neonatal Tet	anus		
			1-2 days normal	Illness occurs bet. 3-28 c & generalized muscle rig	days (w/ poor suck gidity)
			others, specify		_
	4.	Where do you re	egister the cases identified?	CIF	Others,
				logbooks	
		a. Who provides	the case investigation forms?		
	_	5	Reproduce own/hospital	☐ PESU/PHO/CHO/RHU	U Others,
	5.		specimen collection for AFP & Mea		Yes No
		a. Who provide	the specimen collection kits?	☐ DRU/Hospital☐ RESU	PESU/PHO/CESO/CHO Others, specify:
		b. How do you c	collect and store specimens?	STOOL	BLOOD
			2 stool samples	Stored immediately (4-8C)	3-5 cc of blood/ 1-3 cc of serum
			thumb-size/abt. 30 ml (loose)	Properly labeled	Collected w/in 28 days
			collected 24-48 hrs apart	w/in 14 days from paralysis onset	Stored immediately in freezer/body of Ref.
			Others,		
		·			

page 2 of 4

	6.	Where do you store stool and blood samples?	Hospital Ref.	Laboratory	
			Others, specify: _	·	
	7.	Where do you send the specimens for the laboratory confirmation?	RESU	PESU	
		>	RITM		
	8.	Is there a budget for sending specimens?	☐ YES	□NO	
		a. If YES, what is the source of budget?			
III.	REP	PORTING			
	9.	Do you submit VPD data to the PESU/PHO/CHO?	☐ YES	□NO	
		a. If YES, how often?	☐ weekly	☐ monthly	
			Others:	-	
		b. What type of report do you submit?	e-copy/mdb file	Others:	
	10.	Do you practice weekly "zero" reporting to RESU?	YES	□NO	
		a. How is weekly "zero" reporting done? (specify mode of reporting if not in the choices) fax the PIDSR summary form	text the next higher I	evel	
		Send e-mail message	Others, specify		
IV.	DAT	TA MANAGEMENT & ANALYSIS			
	11.	Do you encode VPD data?	YES	□NO	
		a. What program/software are you using in encoding?			
	12.	Do you analyze VPD data?	YES	□NO	
		a. If YES, how often do you conduct data analysis?	weekly	Others,	
			monthly		
	12	How do you get a your skills in date analysis?	Fair	Good	
	13.	How do you rate your skills in data analysis?	── Very good	Excellent	
:	14	Do you keep a data bank of your cases?	YES	□ NO	
		a. If YES, what kind of databank do you have?	_		
	15	(verify by validating data from 2008 to present)			
		How do you detect VPD outbreaks? (Describe processes undertaken)			
1) /		DDACK			
IV.		DBACK			
	16.	Do you provide feedback/reports to the municipal health office/RHU?	YES	NO	
		a. If YES, how often (validate and ask for the hard copy)	weekly	monthly	
		h Danadha hairffulla an Luis (1)	Others, specify: _		
	17	b. Describe briefly the content of the report	- VC0	-	
	17.	Do you provide feedback/reports during hospital assembly? a. If YES, how often and what type of feedback?(validate & ask for hard copy)	YES	NO	
		a. If TES, now often and what type of feedback? (Validate & ask for hard copy)	weekly	monthly	
			Others, specify:	-	
	18.	Do you receive laboratory results of your cases?	YES	NO	
		a. If YES, who provides the laboratory results from RITM?	PESU/PHO/CHO	RESU	
		In the second se	Others,		
		b. What actions do you take in case of positive results for:b1. AFP/Polio	b2. Measles		

page 3 of 4

V.	RES	PONSE						
	19.	Have you established an epidemiology/surveillance unit?	YES	□ NO				
		a. If YES, how many members?	<u> </u>	3 or more				
		à: Who composes the team?	_					
	20	Do you have an epidemic response plan?	YES	□ NO				
	20.	a. If YES, ask, validate and describe						
	21.	Do you coordinate with the concern municipality/city if you notice an increase in the number of cases?	LJ YES	□NO				
		a. If YES, what type and how coordination is being done?						
	SUPPORT ACTIVITIES							
VI.	TRA	INNG						
	20.	Were you trained in PIDSR/VPD surveillance?	YES	□ NO				
		a. If YES, when was your training?	/					
		b. Who provides the training?	PESU/PHO	RESU/CHD				
			Others, specify					
	21.	Were you trained in data analysis?	YES	□ NO				
	22.	Do you conduct basic data analysis?	YES	□ NO				
	23.	Do you conduct training/orientation among hospital staff?	YES	□ NO				
		a. if YES, when was the last time you conducted training?	/					
		b. Describe briefly how the training/orientation was conducted						
VII.	RE	SOURCE/ LOGISTICS PROVISION		197				
	24.	Do you have a line item budget for surveillance/PIDSR/VPD? (validate)	☐ YES	□ NO				
	25.	Do you have a copy of the following?						
		a. PIDSR MOP	YES	NO				
		b. AO # 95 (VPD guidelines)	YES	NO				
	26.	Do you have a dedicated room for surveillance? (Note if shared with other units)	YES	□NO				
	27.							
	21.	Computer unit/s	YES, Qty	□ NO				
		Printer/s	YES, Qty	□NO				
		Internet access	YES	□NO				
		Telephone access	☐ YES	□ NO				
		Fax machine	☐ YES	□ NO				
		LCD/OH projector	□ □ YES	□ □ NO				
		Refrigerator for the specimen	☐ YES	□ NO				
i		Specimen storage boxes	☐ YES	☐ NO				
		Vehicle for surveillance	YES	□ NO				
		Other equipment available/with access:						
				- 				

page 4 of 4

X. SIGNIFICANT FINDINGS:	2 D		
`			
			The state of the s
		- N//	
			0.7001
			100
			NV 8004 J
			1.40.4
	 	40.1	

			, — 168V-17.

Legend:

DRU - Disease Reporting Unit DSC - Disease Surveillance Coordinator

RESU - Regional Epidemiology and Surveillance Unit PIDSR - Phil. Integrated Disease Surveillance and Response



MESU Functionality Levels Name of Municipality:

In using this form: Kindly check the requirements actually present in the municipality being assessed

0	1	2	3
Presence of Ordinan	ce or Executive Order		AND THE PARTY OF T
□ NO	□ YES	HALL CONTRACTOR SEA	THE RESERVE THE
Manpow	er Present		
☐ One Designated	☐ Two Designated	☐ Three Designated	□ > Than Three Trained
Personnel Trained in	Personnel Trained in	Personnel Trained in	Designated Personnel and
PIDSR	PIDSR	PIDSR	Epidemiologist
	Presence of Office Space		
□ NO	□ SHARED	□ DEDICATED	P. C.
	Presence of	f Computer	
□ NO	□ SHARED	□ DEDICATED	☐ DEDICATED with
			WORKSTATION
Acc	ess to Transportation Facil	ities	WORKSTATION
□ NO	☐ SHARED	□ DEDICATED	
	Access to Commu	nication Facilities	
□ NO FAX NO	☐ SHARED FAX OR	☐ DEDICATED FAX	☐ DEDICATED FAX
INTERNET	INTERNET	OR INTERNET	AND INTERNET
	Presence of Al	located Budget	THE THE TENT
□ NO	☐ EXTRACTED FROM	☐ LINED UNDER	☐ DIRECT FROM LGU
	OTHER NON-HEALTH	HEALTH OFFICE/UNIT	BUDGET SEPARATE
	OFFICE/UNIT OF THE		FROM HEALTH
	LGU		UNIT/OFFICE
Presence of Work	and Financial Plan		
□ NO	☐ YES	Concept of the Concep	
	Presence of Surv	eillance Activities	
□ NO CAID (collection,	☐ CAIDs data from	☐ CAIDs data from	☐ CAIDs data from
analysis, interpretation	sentinel sites and provides	sentinel sites and provides	sentinel sites and provides
and dissemination) of	feedback to stakeholders	feedback to stakeholders	feedback to stakeholders
data	irregularly	monthly	weekly
	Access and Availability	of Laboratory Services	
□ NO	☐ Sends lab specimens	☐ Sends lab specimens	☐ Sends lab specimens
	to reference labs (i.e.	to labs within the region	to labs within the
	RITM) for confirmation	for confirmation	province for confirmation
Presence of T	raining Activities pertainin	g to Surveillance and Respo	onse Activities
□ NO	☐ YES but limited to	☐ YES and extended to	☐ YES and integrated to
	activities focused on	activities involving the	activities involving the
	surveillance and response	region	national programs' thrusts
	developed for local wide		-
	use only		
	1	ıtbreak Investigations	
□ NO	☐ YES but with the help	☐ YES but with the help	☐ YES and does
	of the national experts	of the regional experts	investigations involving
			local experts only

Scale: Add the scores of the checked boxes to get the functionality score of the MESU. The corresponding scores (0,1,2,3) are at the heads of the columns.

Example: If the MESU has an ordinance, you will check YES and the corresponding score is 1.

- 0: Your local health office doesn't have an ESU. All you have is a designated employee equivalent to a Disease Surveillance Coordinator under the PIDSR framework.
- 1 to 11: Your Local Health Office is pursuing efforts to build an ESU through shared resources and capacities with the other units in your office. Try to maximize all your efforts to achieve the category of Level 1 ESU.
- 12: Your ESU can be categorized as Level 1. Your ESU exists through shared resources and capacities. Try to maximize all your efforts to achieve the category of Level 2 ESU.
- 13 to 21: Your Level 1 ESU is already achieving resources and capacities necessary to reach the category of Level 2 ESU. Continue with your work to achieve the category of Level 2 ESU.
- 22: Your ESU can be categorized as Level 2. Your ESU exists through dedicated resources and capacities. Try to maximize all your efforts to achieve the category of Level 3.
- 23 to 29: Your Level 2 ESU is already achieving resources and capacities necessary to reach the category of Level 3 ESU. Continue with your work to achieve the category of Level 3 ESU.
- 30: Your ESU is now Level 3. Your local health office had achieved the ideal qualities of a functional ESU fit for your office. Kindly maintain the functionality for the benefit of your local government unit and the PIDSR system.

Definitions:

- 1. Presence of Ordinance or Executive Order: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of ordinance or executive order (printed document) pertaining to the creation of a functional ESU in their province, check the YES cell. Otherwise, check the No cell.
- 2. Designated Personnel: Refers to the personnel assigned FULL TIME in ESU doing FULL TIME ESU work
- 3. Shared Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is NOT SOLELY or NOT for the ESU alone
- 4. Dedicated Office, Computer, Transportation or Communication Facilities: Refers to the status of the said equipment or facilities when their use is SOLELY or for the ESU alone
- 5. Presence of Work and Financial Plan: When a personnel of a province being monitored can show to the monitoring team a tangible proof of the presence of work and financial plan (printed document) pertaining to the functional ESU in their province, check the YES cell. Otherwise, check the No cell.



Department of Health

NATIONAL EPIDEMIOLOGY CENTER



ASSESSMENT FORM for the VPDISS Surveillance Operations (LGU)

LΑ	gion	•	Province) :	Date:	
Nar	ne c	of Respondent:		Position/Designati	ion:	
			SURVEILLA	NCE CORE FUNCTION	NS	
I.	CAS	SE DETECTION, RE	EGISTRATION & CONFIRMATIO	N		_
	1.	Do you have a Prov	vincial Epidemiology & Surveillan	ce Unit (PESU)?	YES NO	
		a. Date ESU estab	olished (look for the ordinance/office	ce order)	//	
		b. How many staff	do you have?		1-2 3 & above	
			ance staff conduct active search for anus in hospitals/DRUs?	or AFP, Measles	☐ YES ☐ NO	
			suspect cases did you detect from	January 2010 to	AFP NT	
			ne following?		Magalag	
		d. Do you conduct of VPDs in your	retrospective records review to ve	erify missed cases	YES NO	
		d1. If YES, hov			weekly monthly	
					Others, specify:	
	2.	What are the surve	eillance activity the unit is involved Active surveillance Passive surveillance 60-day follow-up	No. of Sites No. of Sites Case investigation	Retrospective Records Review Others, specify:	
	0	-	Specimen collection	☐ Outbreak investigation		
	3.		pitals in your catchment area	l' (D00)	Gov't Private	
			with designated disease surveillar	` '	Gov't Private	
		and Neonatal Te		Measies	Gov't Private	
	4.		nent Rural health units (RHU)?			
			h a designated DSC			
	_		ned in PIDSR Surveillance			
	5.	-	for cases of the following VPDs?			
		a. AFP	<15 years	Weakness/flaccid/ floppy paralysis	a case of any age diagnosed as polio by a physician	
			sudden onset of paralysis	Differential Dx:	others, specify	
		b. Measles	Fever/hot to touch	Cough/coryza/ conjunctiuvitis	others, specify	
			Generalized non-vesicular rash of at least 3 days	Differential Dx:	_	
		c. Neonatal Tetanus	1-2 days normal	Illness bet. 3-28 days (w/ poor suck & generalized muscle rigidity)	others, specify	
	6.	Do you conduct spe	ecimen collection for AFP and Me	• • • • • • • • • • • • • • • • • • • •	☐ YES ☐ NO	
	7.		men collection kits?		RESU RITM	
		·			Other, specify:	
	8.	How do you ensure	e adequate specimen collection? Regular coordination w/ the	DRHe		
			DSO collect specimens to al		Hosp/DRUs issue an order for the collection Others, specify:	

page 2of 4

I.	CA	SE DETECTION, REGISTRATION & CONFIRMATION (c	con't)	A			
	9.	Where do you store the specimens?		,			
		Refrigerator at the PESU		Ref. at the hospitals			
		☐ Improvised boxes		Transport boxes		Others,	
	10.	Where do you send the specimens for laboratory confirm	nation	1?		RESU	RITM
	11.	What is your source of budget for freight and handling of	spec	imens?			
		CHD/RESU provision		PESU/PHO/CHO budget		Others,	
H.	RE	PORTING				1	
	12.	Do you submit VPD data to the RESU?				YES	□ NO
1		a. If YES, how often?				weekly	monthly
					$\overline{\Box}$	Other, specify:_	
		b. What type of report do you submit?			\exists	hard copy	е-сору
	13.	Do you receive "zero reports" from your DRUs/RHUs?				YES	□ No
		A. If YES, what mode of submission is being used?			LI	. 20	□ '''
	14.	Do you practice weekly "zero" reporting to RESU?				YES	□ NO
		a. How is weekly "zero" reporting done?					
		fax the PIDSR summary form		text the next higher level			
		send email message to next higher level		Others, specify			
Ш		DATA MANAGEMENT & ANALYSIS					
	15.	Do you encode VPD & AEFI data?				YES	□ NO
		a. If YES, what program/software are you using?		PIDSR EPi-INFO		Other, specify _	11-900-04-00
	16.	Do you analyse surveillance/VPD data?				YES	☐ NO
		a. If YES, how frequent do you analyse your data?		weekly		monthly	_
				as needed		Others, specify	
		b. Ask and look for the reports generated		Graphs		Others, specify	
				Tables			
	17.	How do you rate your skills in data analysis?		Fair		Good	
				Very good	\Box	Excellent	
	18.	Do you keep a data bank of your cases?		, ,	$\overline{\Box}$	YES	□ NO
		a. If YES, what kind of databank do you have? (verify by validating data from 2008 to present)					
	19.	How do you detect VPD outbreaks/clusters based on you	ur dat	a?			
		a. AFP "HOT CASE"/Polio Compatible		. ,			
		· ·					
		b. Measles					
		c. Neonatal Tetanus					

page 3of 4

1) /	eer	DDACK		
IV.	rtt	DBACK		
	20.	Do you receive feedback (i.e. data by province) from the RESU?	YES	□ NO
	21.	Do you provide feedback/reports to the hospitals/RHU?	YES	☐ NO
		a. If YES, how often	weekly	monthly
		(validate and ask for the hard copy)	Others, specify:	
	22.	Do you provide feedback/reports to the hospitals?	YES	☐ NO
		a. If YES, how often?(validate and ask for the hard copy)	weekly	monthly
			Others, specify:	
	23.	Do you receive laboratory results from RESU / RITM?	YES	☐ NO
		a. If YES, do you provide laboratory results to the DRUs concerned?	YES	☐ NO
		b. Were you/any PESU staff invited for a PIR?	YES	☐ NO
		c. Do you conduct any form of PIR this year?	YES	☐ NO
V.	RES	SPONSE		
	24.	Have you established an epidemic response system?	YES	□ NO
	25.	Do you have a response team?	YES	☐ NO
	26.	Do you have an epidemic response plan?	YES	☐ NO
	27.	Do you conduct outbreak response for Measles, Neonatal Tetanus and AFP "hot" case?	YES	□ NO
		a. If YES, describe how do you conduct response.		
		1. AFP "HOT" case		
		2. Measles outbreak/cluster		
		3. Neonatal Tetanus outbreak/cluster		
		SUPPORT ACTIVITIES		1.00
VI.	TRA	AINNG		1 MA 1 WARMAN W
	28.	Were you trained in PIDSR/VPD surveillance?	YES	□ NO
		a. If YES, when was your training?		_
	29.	Do you conduct training/orientation on VPDS among hospital/RHU staff?	YES	☐ NO
		a. if YES, when was the last time you conducted training?		
	30.	Were you trained in data analysis?	YES	□ NO
	31.	Do you conduct basic data analysis training?	YES	□ NO
1				

Legend:
DRU - Disease Reporting Unit
DSC - Disease Surveillance Coordinator

page 4of 4

VII.	SUF	PERVISION						
	32. Do you conduct monitoring and supervisory visits to the following?							
				Hospitals	YES	□ NO		
		`		RHUs	YES	☐ NO		
		a. How frequent do	you conduct monitoring	ng?				
		<u>Hospitals</u>	weekly	monthly RHUs	weekly	monthly		
			quarterly	others,	quarterly	others,		
				SUPPORT ACTIVITIES				
VIII.		SOURCE /LOGISTIC	the state of the s		S.,			
				lance/PIDSR/VPD? (validate)	YES	NO		
	34.	Do you have a copy	y of the following?		YES	☐ NO		
				a. PIDSR MOP	YES	☐ NO		
				b. AO # 95 (VPD guidelines)	YES	NO		
	35.			ance? (Note if shared with other units)	YES	☐ NO		
	36.	Do you have the fol	lowing?					
			Computer unit/s		YES, Qty	□NO		
			Printer/s		YES, Qty	□ NO		
			Internet access		YES	□ _{NO}		
			Telephone access		YES	□NO		
			Fax machine		YES	\square_{NO}		
			LCD/OH projector		YES	□NO		
			Refrigerator for the s	specimen	YES	□ _{NO}		
			Specimen storage b	oxes	YES	□NO		
			Vehicle for surveillar	nce	YES	□no		
			Other equipment ava	ailable/with access:				
IX.	SIGI	NIFICANT FINDINGS	3:					

Legend:DRU - Disease Reporting Unit
DSC - Disease Surveillance Coordinator

RESU - Regional Epidemiology and Surveillance Unit PIDSR - Phil. Integrated Disease Surveillance and Response



Department of Health NATIONAL EPIDEMIOLOGY CENTER

ASSESSMENT FORM for the VPDISS Surveillance Operations (RESU)



Region	:	Address:		Date:				
Name o	Name of Respondent: Position/Designation							
SURVEILLANCE CORE FUNCTIONS								
I. CASE DETECTION, REGISTRATION & CONFIRMATION								
1.	Do you have a Regional E (RESU)?	pidemiology & Surveillance Unit	YES	□NO				
	a. When was the RESU e order)	stablished? (look for the ordinance/office						
	b. How many staff do you	have?	1-2	3 & above				
		aff conduct active search for AFP, nus & AEFIs in hospitals/DRUs?	YES	□ NO				
		cases did you detect from January evious 2 morbidity weeks) of the	AFP	NT				
		ective records review to verify in your catchment hospitals?	YES	□ NO				
	d.1 If YES, how often?		weekly Others, specify:	monthly				
	d.2 How many missed cases of the ff. VPDs were identifi		AFP	Measles				
	based on your last	records review	NT	AEFI				
2.	Total no. of all reporting ho	ospitals in your catchment area	Gov't	Private				
	a. No. of your sentinel hospitals/DRUs		Gov't	Private				
	b. No. of designated DSCs surveillance	s trained in PIDSR/VPD	Gov't	Private				
	c. No. that submit weekly of Tetanus and AEFIs	data on AFP, Measles, Neonatal	Gov't	Private				
3.	Total no. of provinces:	; functional PESUs:	No. of CESUs:					
	a. No. of designated DSOs							
	b. No. trained in PIDSR/VI	PD surveillance:	Provinces/PESU	City/CESU				
	c. No. that submit weekly	data on VPD & AEFI	Prov/PESU	City/CESU				
4.	Do you conduct specimen	collection for AFP and Measles cases?	YES	☐ NO				
5.	How do you define the foll	owing?						
	a. AFP	<15 years	weakness/flaccid/ floppy paralysis	a case of any age diagnosed as polio by a physician				
		sudden onset of paralysis	Differential Dx:	Others, specify				
	b. Measles	fever/hot to touch	Cough/coryza/ Conjunctivitis					
		generalized non-vesicular rash of at least 3days	Differential Dx:	Others, specify				
	c. Neonatal Tetanus	1-2 day/s normal	ill bet. 3-28 days (w/ poor suck & gen'd muscle rigidity)	Others, specify				
		Diagnosed as NT by physician						

page 2 of 5

I.	CAS	E DETECTION, REGISTRATION & CONFIRMATION (con't)							
	6. What are the surveillance activity the unit is involved in?								
		active surveillance (e.g. hospital/DRU vis	No. of sites						
		passive surveillance		No. of sites					
		60-day follow-up		case investigation	others, specify				
		specimen collection		outbreak investigation					
	7.	How do you ensure adequate specimen collection?							
		regular coordination w/ the DSCs		hospitals/DRUs issue a	n order for the collection				
		EPISO/VPD Coord. facilitates collection		others, specify					
	8.	How do you store specimens?							
		refrigerator		transport boxes					
		improvised boxes (e.g. styro)		Others,					
	9.	What is your source of budget for freight and handling of speci	mens	5?					
		CHD/RESU budget		Others,					
		NEC PIDSR sub-allotment							
II.	CAS	SE REPORTING							
	10.	Do you submit reports to NEC?		YES] NO				
		a. What is the frequency of submission?		weekly	as requested				
				monthly	others, specify				
	11.	Do you receive weekly "zero reports" from the DRUs?		YES] NO				
		a. If YES, what mode of submission is being used?							
	12.	Do you practice "zero reporting" to NEC?		YES	NO				
		a. How is weekly "zero reporting" being done?							
		fax the PIDSR summary form		text the NEC PIDSR po	oint person				
		send email message to NEC		Others, specify,					
111.	DAT	TA MANAGEMENT							
	13.	Do you encode VPD & AEFI data?		YES	NO				
	14.	What software/program do you use?		PIDSR EPI-INFO	Others, specify				
	15.	Do you analyse your surveillance data?		YES	NO				
		a. What is the frequency of your data analysis?		weekly	monthly				
				as needed	others, specify				
		b. If YES, ask and look for the reports generated?		graphs	others, specify				
				tables					
	16.	How do you detect outbreaks of the following based on your d	ata?						
		a. AFP "Hot Case"/ Polio Compatible							
		b. Measles outbreaks							
		c. NT outbreaks							

page 3 of 5

IV.	FEE	DBACK				
	17.	Do you provide feedback/reports to the municipal health of	fice/RHU?		YES	□ NO
	,	$_{\mathbf{x}}$ a. If YES, how often (validate and ask for the hard copy)	weekly		quarterly	
			monthly		others, specify:	
	18.	Do you provide feedback/reports to the hospitals?	YES		NO	
		a. If YES, how often? (validate and ask for the hard copy)	weekly		monthly	
			quarterly		others, specify: _	
	19.	Do you receive laboratory results from RESU/RITM?	YES		NO	
		a. If YES, do you provide laboratory results to the DRUs co	ncerned?		YES	☐ NO
V.	RES	PONSE			.:	
	20.	Have you established an epidemic response system?			YES	□ NO
	21.	Do you have a response team?			YES	□ NO
	22.	Do you have an epidemic response plan?			YES	□ NO
	23.	Do you conduct outbreak response for Measles, NT and Al	FP "hot" case?		YES	□ NO
		a. If YES, describe how do you conduct response.				
		a1. AFP "HOT" case				
		Inform NEC	Acquire copy important doc		dical chart & other	
		Verify investigation data (i.e. OPV status, final diagnosis, residual paralysis)	Coordinate w	/ regional	& local EPI staff	
		Other activities, specify				
		a2. Measles outbreak				
		Look for additional cases in the area	Enhance sur	veillance		
		Coordinate w/ EPI to assess the coverage of the affected and surrounding areas	Assist and corresponse	oordinate t	he EPI on the ORI/	
		Other activities, specify				
		a3. Neonatal Tetanus outbreak				
		Investigate/validate reported NT	Coordinate waffected and		ssess the coverage ing areas	of the
		Determine cases of unreported NT/NT deaths occurred in the area	Enhance sur	veillance		

page 4 of 5

VI.	TRA	JINNG		1.451				
	24.	Were you trained in PIDSR/VPD surveillance?			YES			NO
		a. If YES, when was your training?				//_	_	
	25.	Do you conduct training/orientation on PIDSR/VPDS among your	r DRUs?		YES			NO
		a. if YES, when was the last time you conducted training?				//	_	
		b. How many health facilities were trained in PIDSR?						
		Gov't hospitals			,RHUs			
		Private hospitals			clinics			
		Others:						
	26.	How many RESU surveillance staff were trained in data analysis	?					
	27.	Do you conduct basic data analysis training?			YES			NO
		a. If YES, when was the last time you conducted the training?				//_	_	
		b. Who were the participants in the training?						
	28.	List other trainings the RESU has conducted this year: (use addition	onal sheet/s if nee	ded)				
		Title of Training Category of				No. tra	ined	
			Participa	nts				
	1.)							
	1.)							
	2.)							
	2.)							
	2.)							
	2.)							
	2.)							
	2.) 3.) 4.)							
	2.) 3.) 4.)							
VII.	2.)3.)4.)5.)	PERVISION						
VII.	2.) 3.) 4.) 5.)							
VII.	2.) 3.) 4.) 5.)	PERVISION			YES			NO
VII.	2.) 3.) 4.) 5.)	PERVISION	1?					NO NO
VII.	2.) 3.) 4.) 5.)	PERVISION	? <u>Hospitals</u>		YES			
VII.	2.) 3.) 4.) 5.)	PERVISION Do you conduct monitoring and supervisory visits to the following	? <u>Hospitals</u>		YES			
VII.	2.) 3.) 4.) 5.)	PERVISION Do you conduct monitoring and supervisory visits to the following a. How frequent do you conduct monitoring?	? <u>Hospitals</u> <u>RHUs</u>		YES YES weekly			NO
VII.	2.) 3.) 4.) 5.)	PERVISION Do you conduct monitoring and supervisory visits to the following a. How frequent do you conduct monitoring? Hospitals weekly quarterly monthly	? <u>Hospitals</u> <u>RHUs</u>		YES YES weekly monthly			NO
VII.	2.) 3.) 4.) 5.)	PERVISION Do you conduct monitoring and supervisory visits to the following a. How frequent do you conduct monitoring? Hospitals weekly quarterly	? <u>Hospitals</u> <u>RHUs</u>		YES YES weekly monthly			NO

page 5 of 5

SUPPORT ACTIVITIES							
VIII.	RES	SOURCE /LOGISTICS PROVISION					
	31.	Do you have a line item budget for surveillance/PIDSR/VPD? (validate the	сору)	YES	□ NO		
	32.	Do you have a copy of the following?		YES	☐ NO		
		a. PIDSR MOP		YES	☐ NO		
		b. AO # 95 (VPD guidelines)		YES	☐ NO		
	33.	Do you have a dedicated room for surveillance?		YES	□ NO		
		(Note if shared with other units)					
	34.						
		Computer unit/s		YES, Qty	□ NO		
		Printer/s		YES, Qty	□ NO		
		Internet access		YES	NO		
		Telephone access		YES	☐ NO		
		Fax machine		YES	☐ NO		
		LCD/OH projector		YES	□ NO		
		Refrigerator for the specimen		YES	☐ NO		
		Specimen storage boxes		YES	□ NO		
		Vehicle for surveillance		YES	☐ NO		
		Other equipment available/with access:					
IX.	910	GNIFICANT FINDINGS:					
۱۸.		SNIFICANT FINDINGS:					