



## GUIDELINES

# SFM Interim Practice Recommendations for Zika Virus Infection in Pregnancy

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**Abstract** Zika virus (ZIKV) belongs to the family of flaviviruses and is transmitted by mosquitoes of the genus *Aedes* (*A. aegypti* and *A. albopictus*). ZIKV infection in pregnancy can have serious implications on the fetus and the neonate. Vertical transmission to the fetus can occur irrespective of symptomatic or asymptomatic infection in the pregnant woman and the risk can persist throughout pregnancy. The exact frequency of maternal-to-fetal transmission of Zika virus is difficult to determine accurately. The greatest risk of serious fetal sequelae is seen if the infection occurs within the first and second trimester. However, they can also be seen with infections in the third trimester. This document describes the presentation, fetomaternal manifestations, role of prenatal Ultrasound in diagnosis of infection, testing methods and the management in pregnancy and screening in the neonatal period. In the absence of robust evidence as to the definite effects, this document is aimed at providing practice recommendations for the management of this infection.

**Keywords** Zika virus · Pregnancy · Microcephaly

## About the Virus

Zika virus (ZIKV) is transmitted by mosquitoes of the genus *Aedes* (*A. aegypti* and *A. albopictus*). It belongs to the family of flavivirus and is closely related to other

flaviviruses including dengue virus, Japanese encephalitis, yellow fever virus and West Nile virus. Outbreaks of infection with Zika virus have occurred in Africa, South East Asia, Pacific islands, America, and the Caribbean [1, 2]. Acute onset of low-grade fever with maculopapular pruritic rash, arthralgia (particularly of small joints of hands and feet), or conjunctivitis (non-purulent) are the common clinical features of Zika virus infection. These are reported to occur in only about 20 percent of infected individuals. Zika virus may be transmitted to humans via the infected mosquito through a bite, maternal to fetal transmission, sexual transmission (including vaginal, oral and/or anal sex), transfusion of blood and/or blood products, organ transplantation and laboratory exposure [3].

## Zika Virus Outbreaks in India

Around January 2017, three confirmed cases were reported and the first outbreak was from Ahmedabad in the state of Gujarat. A second outbreak was reported in Tamil Nadu. This was followed by 94 confirmed cases in the state of Rajasthan in 2018 [4], which also included 22 pregnant women. There have been recent outbreaks in different parts of India. There were 37 confirmed cases of Zika virus infection in Kerala during July 2021 and 89 cases in Kanpur in November 2021 [5].

## Clinical Manifestations in the Mother

Pregnant women have been reported to develop symptoms in 17% to as high as 56% of cases of ZIKV infection [6]. The incubation period is 2–14 days. The common symptoms are acute onset of low grade fever (37.8 to 38.5 °C),

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pruritic rash (erythematous macules and papules on the face, trunk, extremities, palms, and soles), arthralgia (notably in the small joints of the hands and feet), and conjunctivitis (non-purulent). The less common symptoms include myalgia, headache, retro-orbital pain and vomiting. These generally resolve within 2–7 days and the infection usually has a mild course. When two or more of the symptoms are present as depicted in Table 1, then clinical illness consistent with Zika virus disease is said to be present.

There is immunity to re-infection following primary infection. Prior dengue infection may be protective against symptomatic infection with Zika virus.

## Effect of Infection with Zika Virus on the Fetus and Adverse Pregnancy Outcomes

### Risk of Vertical Transmission

Vertical transmission to the fetus can occur irrespective of symptomatic or asymptomatic infection in the pregnant woman and the risk can persist throughout pregnancy. The exact frequency of maternal-to-fetal transmission of Zika virus is difficult to determine accurately.

The greatest risk of serious fetal sequelae is seen if the infection occurs within the first and second trimester. However, they can also be seen with infections in the third trimester.

In a cohort study of 130 infants whose mothers developed symptomatic ZIKV in pregnancy which was confirmed with PCR testing, the vertical transmission rate was 65 percent [8]. Other reports have suggested lower rates. The rate was 26 percent (76/291) in a prospective cohort study from French Guiana [9].

### Consequences of Vertical Transmission

The overall risk of any abnormality or birth defect varies widely (6 to 8 percent in studies in the United States versus over 40 percent in a Brazilian study) among fetuses and infants of women with ZIKV infection during pregnancy [10–12].

Beyond birth defects, there also have been reports of fetal loss (miscarriage, stillbirth), hydrops fetalis, fetal growth restriction, neurologic and positional abnormalities, and impaired neurodevelopment.

### Spectrum of Abnormalities

#### *CNS Manifestations*

The virus shows neurotropism in-vivo and in-vitro. There are stages of affection where the infection from the mother leads to consequent infection of the placenta and the fetus. etal brain gets preferentially affected involving the neuronal progenitor cells which leads to disruption of neuronal growth, proliferation, migration, and differentiation. This leads to impaired brain development. This reflects as congenital microcephaly, Guillain Barré syndrome, myelitis, and meningoencephalitis.

The ultrasound manifestations in the fetus can appear as early as two weeks. However, these are more prominent in the second and third trimester. The three most common findings detected on a prenatal ultrasound include microcephaly, ventriculomegaly and intracranial calcifications, especially along the grey matter-white matter junction rather than punctate calcifications as seen with other congenital infections.

**Microcephaly** When the fetal head is significantly smaller than expected for gestational age it is labeled as

**Table 1** WHO Case definitions of clinical illness consistent with Zika virus disease [7]

#### Case Definitions

##### Suspected case

A person presenting with rash and/or fever and at least one of the following signs or symptoms:

Arthralgia; or

Arthritis; or

Conjunctivitis (non-purulent/hyperaemic)

##### Probable case

A suspected case with presence of IgM antibody against Zika virus and an epidemiological link

##### Confirmed case

A person with laboratory confirmation of recent Zika virus infection:

Presence of Zika virus RNA or antigen in serum or other samples (e.g. saliva, tissues, urine, whole blood); or

IgM antibody against ZIKV positive and PRNT 90 for ZIKV with titre  $\geq 20$  and ZIKV PRNT90 titre ratio  $\geq 4$  compared to other flaviviruses;

And exclusion of other flaviviruses

microcephaly. It is a sign of underlying pathology. The overall frequency of microcephaly following in utero Zika virus exposure is 5–7% and it can be primary (noted at birth) or secondary (developing in the first months of life) [13].

It generally reflects that the infection occurred early in pregnancy. Both disproportionate and proportionate microcephaly can be observed suggesting that it can affect fetal growth in addition to its destructive effect on the fetal brain. It can be detected as early as 18 weeks of gestation. Irrespective of which criteria is used to define microcephaly, it is important to note that the head circumference should be disproportionately smaller in comparison to abdominal circumference and femur length and not explained by other etiologies or congenital malformations (Table 2). Zika virus related microcephaly should be suspected if microcephaly is associated with a molecular or epidemiological link to Zika virus in the absence of other conditions known to cause microcephaly.

A molecular or epidemiological link with Zika virus is defined as:

- The pregnant woman is a confirmed case of Zika virus disease; or
- The pregnant woman had unprotected sexual contact with a confirmed case, or a history of symptoms or signs consistent with Zika virus infection and residing/traveling in an area with ongoing Zika virus transmission during her pregnancy; or
- Presence of Zika virus in amniotic fluid (identified through amniocentesis and RT-PCR assay); or
- Presence of Zika virus in fetal brain tissue (identified post-mortem through RT-PCR assay)

Regardless of the criteria used, for in utero diagnosis of Zika virus-induced microcephaly, the occipitofrontal circumference should be disproportionately small in comparison with the abdominal circumference and femur length and not explained by other etiologies or congenital disorders. If the occipitofrontal circumference is  $\geq$  3rd percentile but is notably disproportionately small compared with the abdominal circumference and fetal length or if central nervous system abnormalities are noted,

additional evaluation for ZIKV infection may be appropriate.

The various other features that can be found are [17]:

Irregular head shape including a sloping/slanted forehead, cystic lesions, intraventricular adhesions, callosal dysgenesis or agenesis, cerebellar hypoplasia or vermian dysgenesis, enlarged cisterna magna, abnormal cortical development with reduced amount of brain parenchyma, arthrogryposis, club foot and microphthalmia.

The reported sensitivity of ultrasound is 22% and specificity 98% for detection of infection [18].

In Zika virus specific associated neuroanatomic anomalies such as diffuse calcification of subcortical parenchyma and thalamus, ventriculomegaly, lissencephaly, and pachygyria have been distinctively highlighted [19], along with a constellation of particular neuro-pathological findings such as: gliosis, abnormal neuronal migration, dysmaturation of nerve cells, hypomyelination, loss of descending axons, and spinal motor neurons.

The principal clinical features of congenital Zika syndrome (CZS) in the newborn include microcephaly, facial disproportion, hypertonia/spasticity, hyperreflexia, seizures, irritability, arthrogryposis, ocular abnormalities and sensorineural hearing loss. The unique features of CZS that are rarely seen with other congenital infections are:

- Severe microcephaly with partially collapsed skull
- Thin cerebral cortices with subcortical calcifications
- Macular scarring and focal pigmentary retinal mottling
- Congenital contractures
- Marked early hypotonia

### *Fetal Growth Abnormalities*

Infection with Zika virus can result in symmetric or asymmetric fetal growth restriction [20].

In a United States cohort of women with antenatal ZIKV infection, 11.2 percent were small for gestational age, compared with 5.8 percent of controls.

**Table 2** Various standard definitions of microcephaly

Recommendation by	Definition of microcephaly
WHO [14]	Occipitofrontal circumference (head circumference) greater than 2 SD below the mean or less than the 3rd percentile based on standard growth charts for sex, age, and gestational age at birth (Postnatal)
CDC [15]	Occipitofrontal circumference below the 3rd percentile or greater than 2 SD below the mean. (Postnatal)
Society of Maternal–Fetal Medicine [16]	Head circumference equal to or greater than 3 SD below the mean for gestational age Pathologic microcephaly when head circumference is greater than 5 SD below the mean for gestational age

### Cardiac Anomalies

Zika outbreak in the Brazilian study reported that 11% (13/120) of fetuses had non-severe cardiac defects (ASD, VSD, PDA) [21]. The frequency was increased in babies whose mothers developed a rash in the second half of pregnancy or those who were noted to have abnormal postnatal central nervous system imaging and in those who were preterm.

### Trimester Specific Risks of Affection

Infection with Zika virus can result in fetal loss (miscarriage, stillbirth) and hydrops fetalis. Placental insufficiency from injury or infarction is the possible mechanism for fetal loss in the latter half of pregnancy (Table 3).

## Diagnosis of Infection

The approach to diagnosis is different in pregnant women due to longer persistence in the maternal serum.

All pregnant women should be verbally screened for any possible exposure to Zika virus at the initial antenatal visit. In case the pregnant woman resides in the endemic region or there is any history of travel related exposure, the pregnant woman should be screened verbally for any possible symptoms of the infection.

### Laboratory Diagnosis of Maternal Infection

#### Indications

Testing for Zika virus should be done for:

- Every symptomatic pregnant woman with possible ZIKV exposure
- Asymptomatic pregnant women with ongoing possible ZIKV exposure
- ZIKV-exposed pregnant women whose fetus presents with prenatal US findings consistent with congenital ZIKV infection.

- Asymptomatic pregnant women with recent possible but no ongoing exposure to ZIKV (travel history)

#### Types of Tests

- NAAT Test—RTPCR**  
Nucleic acid detection (Zika virus RNA) by reverse transcriptase-polymerase chain reaction (RT-PCR) testing of whole blood (or serum or plasma) which targets the non-structural protein is the mainstay of diagnosis during the acute phase of infection and up to even 12 weeks of symptom onset. Standard RTPCR and quantitative RT-PCR provide a rapid, specific and sensitive method for early detection of infection. If the sample was taken later in the course of infection, a negative PCR test might not exclude infection with Zika virus as the viremia levels fall rapidly. Zika virus RNA also has been detected in urine or saliva samples.
- Antibody test—IgM ELISA/Immunofluorescence test**  
Later in the infection, convalescent phase ( $\geq 5$  days) can reflect as a positive serology by testing IgM antibodies (Zika virus immunoglobulin M) in blood by ELISA. This, however, is not the mainstay of diagnosis as cross reactivity with other flaviviruses and false positive results are very high, thus making it extremely difficult to interpret serology in women who have resided in dengue endemic areas or have had a past infection with any flavivirus.  
Interpretation of serology tests:
  - If the results are positive or equivocal: Further sample may be submitted, and the pregnant woman should be referred to a fetal medicine specialist for further evaluation and follow up.
  - If the result is negative: Repeat testing may be recommended a few weeks after the last possible exposure, before Zika infection can be ruled out with confidence. It may be considered to test these individuals for other potential causes of their symptoms such as dengue, chikungunya and malaria

**Table 3** Effect on the fetus by Zika virus infection in various trimesters [22]

Effect on fetus	Percentage prevalence of defect if infection in first trimester (%)	Percentage prevalence of defect if infection in second trimester (%)	Percentage prevalence of defect if infection in third trimester (%)
Microcephaly	10	3	4
Neurologic and Ocular defects	12	3–4	5–6

- c. Plaque Reduction Neutralization Test (PRNT):
  - Confirms diagnosis in inconclusive antibody testing.
  - Discriminates between crossreacting antibodies in primary flavivirus infections
- d. Virus Isolation
  - NOT a diagnostic tool. For supplemental research studies in public health surveillance

#### *Sample Transport, Storage and Processing*

- Sample types
  - Primary* Blood / Serum (5 ml), EDTA samples (5 ml), CSF(1 ml): Suspected GBS/Myelitis.
  - Secondary samples* Urine, Semen, Saliva (To detect carriers/Late sample collection in patients).
- Transportation and storage:
  - Triple layer packaging (“Appendix 1”)
  - Sample collection form (“Appendix 2”)
  - Follow sample storage temperatures

#### **In Case of Delay**

- Keep refrigerated (2–8 °C)— To be processed / Sent to VRDL within 48 h.
- Keep frozen (– 10 to – 20 °C)— To be processed after within 48–72 h
- Keep frozen (– 70 °C)—To be processed after a 7 days. The sample can be preserved for extended periods
- Precautions
  - A BSL-2 containment level is needed for suspected samples
  - Consider all specimens as potentially infectious
  - Gloves are MANDATORY
  - Separate sample container labeled with the patient’s name and identification number, the collection site, the date of collection and the time of the collection
  - Do not contaminate the outside of the specimen container
  - Empty vials and gloves to be packed in biohazard bags, secured and loaded to autoclave for decontamination.
- Remarks
  1. The nodal laboratories for each district are decided, samples dispatched accordingly to the respective viral research and diagnostic laboratories (VRDLs)

2. NIV Pune and NCDC were identified as the Apex labs for testing the suspected Zika Viral Fever patients during the last outbreak in India.
3. The positive samples were transported to NIV, Pune for Sequencing.

#### **Ultrasound Screening for Fetal Affection**

It is recommended to perform prenatal ultrasound for screening of fetal abnormalities consistent with Zika infection for all pregnant women in whom laboratory testing (RT-PCR for Zika virus RNA or Zika virus specific IgM) is suggestive or diagnostic of possible Zika virus infection. Moreover, it is important to note that pregnant women with only history of possible exposure but without laboratory evidence of infection should only undergo routine ultrasound examination as recommended for routine antenatal evaluation. The ideal timing between exposure and first ultrasound screening and even follow up screening are not clearly known. The minimum time may be as short as two weeks in a few cases and may be seen as early as 18–20 weeks. In the majority of cases, fetal abnormalities are usually detected in the late second and early third trimesters of pregnancy despite infection in the first trimester.

Screening protocol:

- Timing of ultrasound:
  - It is reasonable to perform the first ultrasound examination four weeks from the suspected exposure, followed by serial ultrasound examinations every four weeks, ensuring that at least one ultrasound is performed between 28 and 33 weeks of gestation.
- Checklist for ultrasound:
  - Detailed fetal CNS examination to look for abnormalities
  - Fetal growth to look for any growth restriction
  - Detailed fetal echocardiography to look for any cardiac defects
  - Fetal position to look for contractures and related limb abnormalities or club foot.

It must be noted that all pregnant women residing in areas with Zika virus transmission should be advised to continue their scheduled antenatal visits in accordance with the local policy.

In case of an exposed pregnant women with no reported symptoms during travel or within 2 weeks of return or probable sexual exposure, an ultrasound for fetal growth and anatomy may be done. If this is abnormal, she should be referred to a fetal medicine specialist for review.



However, if the baseline ultrasound is normal, a repeat ultrasound in the third trimester may be considered.

## Management in Pregnancy

### Maternal Management

Currently, there is no specific management for infection with Zika virus. The woman should be advised rest and symptomatic treatment. She should drink fluids to prevent dehydration and paracetamol can be administered to relieve fever and pain. Dengue infection should be ruled out before prescribing aspirin and other (NSAIDs) and should be avoided until then to reduce the risk of hemorrhage.

### Monitoring for Fetal Affection

#### a. Ultrasound

Prenatal ultrasound should be performed to detect fetal abnormalities as enlisted above. The timing, the screening protocol as discussed above should be followed. In case of fetal growth restriction or microcephaly, serial follow up ultrasound should be based upon each clinical scenario.

#### b. Invasive testing

The exact indications, appropriate gestational age for the test and the interpretation of test results of amniocentesis are quite uncertain [23]. The sensitivity and specificity are also not exactly known. It may be performed 6–8 weeks after maternal infections and the highest sensitivity may be around 21 weeks. In addition, Zika virus RT-PCR positivity in amniotic fluid is diagnostic of fetal viral exposure but not predictive of outcome especially if the test was performed because of maternal laboratory testing and the fetus appears normal on ultrasound. The diagnostic accuracy of RT-PCR test in identifying congenital infection, and to what extent a positive test can predict subsequent fetal malformations is currently not known. Hence, the decision should be based on individual circumstances or clinical scenarios.

Amniocentesis is offered to pregnant women:

- Who have ultrasound findings suggestive of congenital Zika virus syndrome
- And/or positive or inconclusive maternal laboratory test results for Zika virus infection, when this information will impact decisions about pregnancy termination or ongoing pregnancy and delivery management.

Interpretation of amniotic fluid results:

- Positive rRT-PCR of amniotic fluid: It is considered diagnostic of fetal infection.
- Negative rRT-PCR of amniotic fluid: It does not rule out congenital infection as the virus may be detected transiently only.

### Antepartum Fetal Monitoring

As there is an increased risk of adverse pregnancy outcome due to fetal growth restriction or even hydrops fetalis, antepartum fetal monitoring may be performed based on the ultrasound affection of fetus. In case of abnormal results, the management should be based on the routine obstetric policies.

### Decision for Termination of Pregnancy

Extensive counseling should be offered to any pregnant woman who is detected positive for Zika virus infection in order to make an informed decision on termination of pregnancy, provided it is within the stipulations laid down under Medical Termination of Pregnancy Act.

Pregnant mothers, positive for Zika virus disease and beyond the period of gestation permissible by the MTP Act should be followed up for ultrasonographic evidence of microcephaly or other evidence of Zika virus affection as per the ultrasound protocol discussed above and counseled accordingly.

### Management of Labour, Delivery and Postpartum Period

- Routine obstetric standards and policies guide and determine the timing and route of delivery.
- No evidence of maternal risk from placement of neuraxial anesthesia in women with uncomplicated active infection (no Guillain Barré syndrome).
- In women with positive or inconclusive Zika virus testing results, histopathology (HPE) of the placenta and umbilical cord, Zika virus immunohistochemical (IHC) staining on fixed tissue or Zika virus RNA testing (via RTPCR) on fixed and frozen tissue can be done to document fetal infection.
- RTPCR and HPE with IHC should also be performed on fetal tissues including the umbilical cord and placenta in cases of fetal loss or stillbirths [24].
- Women should continue to breastfeed as the small potential risk of Zika virus transmission through breast milk is outweighed by the known benefits of breastfeeding.

## Post-natal Presentation

The majority of fetal or neonatal manifestations are identified postnatally. Ocular abnormalities are common including, pigmentary maculopathy, circumscribed chorioretinal atrophy, optic nerve abnormalities, microcornea, microphthalmia, cataracts, retinal dysplasia. Hearing loss has been reported in 7.1% in a large series (5/70). Neurologic and positional abnormalities can manifest in the form of hypertonia, hypotonia, spasticity and hyperreflexia [25].

## Strategies for Surveillance and Prevention

### Primary Prevention

- Adhere to mosquito protective measures
- Adhere to measures (abstinence or condoms) throughout pregnancy to protect against sexual transmission if a partner (male or female) has traveled to or lives in an area with a risk of Zika virus infection
- Adhere to standard infection precautions
- Guidance for couples planning pregnancy  
For couples residing in areas with active Zika virus transmission.
  - Inform about the risks of transmission
  - Consequences
  - Possibility of delaying pregnancy
- For couples with a history of Zika infection:
  - If the male partner is positive wait at least six months from symptom onset before attempting conception
  - If the female partner is positive wait at least eight weeks from symptom onset before attempting to conceive

### Secondary Surveillance Plan [26, 27]

1. All pregnant women in the defined area be under surveillance for febrile illness.
2. Testing strategies:

## Testing the Mother

Ask about	Travel or residence in areas with Zika virus transmission before or during the current pregnancy
	Possible sexual exposure before or during pregnancy
	Diagnosis of lab-confirmed infection before pregnancy
	Symptoms of Zika virus during current pregnancy
Counsel regarding	Personal protective measures against mosquito bites and possible sexual transmission
	Environmental protective measures for reducing proliferation of mosquitos
	Reporting signs and symptoms suggestive of Zika virus infection
Whom to test	Every symptomatic pregnant women with possible ZIKV exposure
	Asymptomatic pregnant women with ongoing possible ZIKV exposure
	ZIKV exposed pregnant women whose fetus presents with prenatal US findings consistent with congenital ZIKV infection
	Asymptomatic pregnant women with recent possible but not ongoing exposure to ZIKV (travel history)
Which test	RT-PCR in acute phase in serum
	Serology after the acute phase
Where to test	Government approved VRDLs
TAT of tests	7–10 days

3. All febrile cases in the defined geographic area should be in home isolation in a mosquito proofed room.
4. If mosquito proofed room is not available, LLIN bed nets to be used.
5. Persons who test positive for Zika virus disease to be in isolation in mosquito free surroundings for 14 days.
6. Men/ women who had tested positive for Zika virus disease to be counseled for safe sex practices to be followed for a period of six months

## Testing the New-Born and Follow-Up

Initial evaluation	<p>For infants born to mothers with laboratory evidence of Zika virus infection and/or infants with findings suggestive of congenital Zika infection in the setting of a maternal epidemiologic link:</p> <ol style="list-style-type: none"> <li>1. Thorough physical examination (with measurement of HC, length, and weight; assessment of gestational age [and examination for neurologic abnormalities and dysmorphic features</li> <li>2. Laboratory testing to detect Zika virus</li> <li>3. Head ultrasonography</li> <li>4. Standard newborn hearing assessment</li> <li>5. Additional investigations if both laboratory confirmed infection with clinical disease</li> </ol>
Whom to test	<ol style="list-style-type: none"> <li>1. Newborns born to mothers with confirmed Zika Virus infection</li> <li>2. New-born with clinical findings/ neuroimaging findings suggestive of Zika infection with or without microcephaly or with or without the mother being positive</li> </ol>
Which tests to be done	<ol style="list-style-type: none"> <li>1. Serum for Zika virus rRT-PCR</li> <li>2. Serum Zika virus IgM ELISA</li> </ol>
Follow-up plan	<ol style="list-style-type: none"> <li>1. Monthly for at least the first six months after birth: <ul style="list-style-type: none"> <li>Monitoring growth parameters, including head circumference</li> <li>Monitoring development using a standardized, validated developmental screening tool</li> <li>Providing routine immunizations</li> <li>Providing anticipatory guidance, psychosocial support, and ensuring infants receive necessary testing and consultations</li> </ul> </li> <li>2. If not performed during the birth hospitalization: <ul style="list-style-type: none"> <li>Auditory brainstem response testing within one month</li> <li>Comprehensive ophthalmologic examination should be performed within one month after birth</li> <li>Neurologist evaluation within one month</li> </ul> </li> <li>3. Monitor: <ul style="list-style-type: none"> <li>Infant feedings</li> <li>Signs of increasing intracranial pressure (eg, increasing head circumference, irritability, or vomiting)</li> </ul> </li> </ol>

## Appendix 1

### Triple Layer Packing

1. Primary Container:
  - Sterile vial -Use MICROPORE to seal the lid
  - Cover vial with cotton or tissue paper
2. Secondary container: Zip lock cover
3. Tertiary container:
  - Thermocol box lined with dry ice(ideal)
  - Unavailability of Dry Ice:Ice packs(4 to 6) to line the thermocol box in order to keep the sample at 2–8 °C

## Appendix 2

### Sample Collection Form



## DEPARTMENT OF MICROBIOLOGY

Case Information		
VRDL number: _____	Lab ID: _____	
State: _____	Country: _____	Date form completed: ____/____/____ (MM/DD/YYYY)
Medical Officer: _____		
PHC/CHC/GH/Medical College/Other: _____		
Department: _____		
Phone number: _____		
Email address: _____		
Patient Demographics and Contact Information		
Patient name: _____	IP/OP No: _____	Unit: _____ Ward: _____
Age: _____	Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female	Pregnant: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Date of Admission: _____	Occupation: _____	
Address: _____		
Phone number: _____ E-mail address: _____		
Clinical Information		
Date of symptom onset ____/____/____ OR <input type="checkbox"/> Asymptomatic		
Fever <input type="checkbox"/> Yes <input type="checkbox"/> No		
Rash <input type="checkbox"/> Yes <input type="checkbox"/> No If yes: Type: <input type="checkbox"/> Maculopapular <input type="checkbox"/> Petechial <input type="checkbox"/> Purpuric <input type="checkbox"/> Other		
Pruritic: <input type="checkbox"/> Yes <input type="checkbox"/> No Distribution: _____		
Arthralgia <input type="checkbox"/> Yes <input type="checkbox"/> No		
Conjunctivitis <input type="checkbox"/> Yes <input type="checkbox"/> No		
Other symptoms: _____		
Hospitalized <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, reason: _____ Dates/Status: _____		
Exposure Information before Symptom Onset (or specimen collection if asymptomatic)		
1. Did the patient travel to or live outside his/her city or county before symptom onset or specimen collection?		
<input type="checkbox"/> Yes <input type="checkbox"/> No		
If yes: Location(s) of travel (country, state, city): _____		
Travel start date: ____/____/____ Return date: ____/____/____		

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