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MIDDLE EAST RESPIRATORY SYNDROME CORONAVIRUS; GUIDELINES FOR HEALTHCARE PROFESSIONALS

مـــركز القيادة و الـتحــكم Command & Control Center



1. TABLE OF CONTENTS

1.	AC	KNOWLEDGMENT	2
2.	ΙΝΤ	RODUCTION	3
3.	OB	JECTIVES	3
4.	CAS	SE DEFINITION	4
4	.1	SUSPECTED CASE	4
4	.2	CONFIRMED CASE	4
5.	INF	ECTION PREVENTION AND CONTROL	5
5	.1	Administrative Interventions	5
5	.2	TRANSMISSION PRECAUTIONS	6
5	.3	PATIENT PLACEMENT	6
5	.4	PATIENT TRANSPORT	7
5	.5	Personal Protective Equipment (PPE) for HCWs	7
5	.6	Environmental Cleaning and Disinfection	8
5	.7	MEDICAL WASTE	10
5	.8	Textiles	10
5	.9	INFECTION PREVENTION AND CONTROL PRECAUTIONS FOR AEROSOL-GENERATING PROCEDURES	10
5	.10	FIT TEST AND SEAL CHECK	11
5	.11	MANAGEMENT OF EXPOSURE TO MERS-COV IN HEALTHCARE FACILITIES	12
5	.12	Outbreak Management	14
5	.13	PATIENT TRANSPORTATION AND PREHOSPITAL EMERGENCY MEDICAL SERVICES	14
5	.14	DURATION OF ISOLATION PRECAUTIONS FOR MERS-COV INFECTION	15
6.	PU	BLIC HEALTH CONSIDERATIONS	15
6	.1	Surveillance and Reporting	15
6	.2	Household and Community Contacts Management	16
6	.3	Home Isolation Guidance	17
6	.4	HUMAN ANIMAL INTERFACE	18
7.	LAE	3ORATORY DIAGNOSIS OF MERS-CoV	19
8.	ΟΤ	HER CONSIDERATIONS	20
8	.1	GENERAL OUTLINES OF MANAGEMENT	20
8	.2	Extra Corporeal Membrane Oxygenation (ECMO)	20
8	.3	MANAGING BODIES OF DECEASED MERS-COV PATIENTS	21
9.	REI	ERENCES	22
2.	AP	PENDICES	23

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2. INTRODUCTION

Middle East Respiratory Syndrome (MERS) is a viral respiratory disease caused by a novel coronavirus (Middle East Respiratory Syndrome Coronavirus, or MERS-CoV) that was first identified in Saudi Arabia in 2012.

Typical MERS-CoV symptoms include fever, cough and shortness of breath. Pneumonia is common, but not always present. Approximately 35% of reported patients with MERS-CoV have died.

Although some of human cases of MERS-CoV have been attributed to human-to-human infections in health care settings, current scientific evidence indicates that dromedary camels are a major reservoir host for MERS-CoV and an animal source of MERS-CoV infection in humans.

This is the fifth edition (updated May 2018) of the national MERS-CoV guidelines. A large group of national and international experts in epidemiology, infectious diseases, infection control, intensive care, laboratory, veterinary medicine and public health were hosted by Saudi Ministry of Health (MOH) to review current knowledge and update the guidelines.

3. OBJECTIVES

This document provides guidelines on managing MERS-CoV infection based on the best available scientific evidence and broad consensus through the following:

- Provide guidance on MERS-CoV surveillance activities in the healthcare setting and in the community.
- Provide guidance on the infection control precautions for suspected and confirmed MERS-CoV cases.
- Standardize the clinical management of MERS-CoV patients.
- Provide guidance for rational use of resources including laboratory testing.
- To act as focus for quality control, including audit.

4. CASE DEFINITION

4.1 SUSPECTED CASE¹

Age	Clinical Presentation	Epidemiologic Link
Adults	 I. Severe pneumonia (severity score ≥3 points) (Appendix A) or ARDS (based on clinical or radiological evidence) 	Not required
Adults ²	II. Unexplained deterioration ³ of a chronic condition of patients with congestive heart failure or chronic kidney disease on hemodialysis	Not required
Children and adults	 III. Acute febrile illness (T ≥38° C) with/without respiratory symptoms OR IV. Gastrointestinal symptoms (diarrhea or vomiting), AND leukopenia (WBC≤3.5×10° /L) or thrombocytopenia (platelets < 150×109/L) 	 Within 14 days before symptom onset: 1. Exposure⁴ to a confirmed case of MERS-CoV infection OR 2. Visit to a healthcare facility where MERS-CoV patients(s) has recently (within 2 weeks) been identified/treated OR 3. Contact with dromedary camels⁵ or consumption of camel products (e.g. raw meat, unpasteurized milk, urine)

4.2 CONFIRMED CASE

A Confirmed case is defined as a suspected case with laboratory confirmation of MERS-CoV infection.

¹ All suspected cases should have samples collected for MERS-CoV testing (nasopharyngeal swabs or sputum, and when intubated, lower respiratory secretions)

 $^{^2}$ Adult is defined as > 14 years old

³ Chronic renal failure and congestive heart failure patients may exhibit fever and presence of fluid overload may mask the radiological features of pneumonia

⁴ Exposure is defined as a contact within 1.5 meters with a confirmed MERS-CoV patient.

⁵ Exposure to camels include:

Direct physical contact with camels or their surroundings (milking and handling excreta are especially risky), drinking raw camel milk or other unpasteurized products derived from camel milk, and handling raw camel meat.

Indirect contact include casual contact with camel places like visiting camel market or farms without direct physical contact with camels, living with a household member who had direct contact with camels.

5. INFECTION PREVENTION AND CONTROL

5.1 ADMINISTRATIVE INTERVENTIONS

To prevent the transmission of respiratory infections in the healthcare settings, including MERS-CoV and influenza, the following infection control administrative measures should be incorporated into infection control practices and implemented:

- Triage for patients with Acute Respiratory Illness (ARI):
 - Visual triage should be used for early identification of all patients with ARI in the Emergency Room and dialysis units.
 - Visual triage station should be placed at the entry point of the healthcare facility (i.e. emergency room entrance, dialysis unit entrance) or other designated areas and attended by a trained nurse or nurse assistant.
 - All patients attending hemodialysis units and all emergency room attendees (except those with immediately life-threatening conditions) must be triaged at the entrance using predefined scoring (Appendix B).
 - Identified ARI patients should be asked to perform hand hygiene and wear a surgical mask. They should be isolated and evaluated immediately in an area separate from other patients, ideally a separate room
- Dedicate a waiting area for the ARI patients with spatial separation of at least 1.2 meter between each ARI patient and others.
- Post visual alerts (in appropriate languages) at the entrance of healthcare facilities (e.g. emergency rooms and clinics). Messages in the visual alerts include the following:
 - Cover your mouth and nose with a tissue when coughing or sneezing.
 - Dispose of the tissue in the nearest waste receptacle immediately after use.
 - Perform hand hygiene (e.g. hand washing with non-antimicrobial soap and water, alcohol-based hand sanitizer, or antiseptic hand wash) after having contact with respiratory secretions and contaminated objects or materials.
- Prevent overcrowding in clinical areas to reduce the risk of transmission between patients and to staff.

- The distance that should be maintained between patients` beds are:
 - Minimum of 1.2 meters in General words, Hemodialysis units and Emergency units.
 - Minimum of 2.4 meters in Critical care units.

5.2 TRANSMISSION PRECAUTIONS

MERS-CoV is believed to spread between humans mainly through contact and respiratory droplets. However, transmission through small particle droplet nuclei (aerosols) may occur. Environmental contamination during outbreaks in healthcare facilities can be extensive and might contributes to amplifying outbreaks, if adequate disinfection procedures are not followed.

- For patients with suspected, or confirmed MERS-CoV infection who are NOT CRITICALLY ILL, Standard, Contact, and Droplet precautions are recommended.
- For patients who are CRITICALLY ILL, Standard, Contact, and Airborne precautions are recommended due to the high likelihood of requiring aerosol-generating procedures.

5.3 PATIENT PLACEMENT

Every healthcare facility should have the capacity to care for patients with transmissible infections including airborne infections. However, the availability of single rooms and negative pressure rooms are a challenge in most facilities. The infection control teams should take the lead in managing isolation rooms.

- Patients with suspected or confirmed MERS-CoV infection who are not critically ill should be placed in single patient rooms in an area that is clearly segregated from other patient-care areas. A portable HEPA filter could be used and placed according to the manufacturer recommendations.
- Critically ill patients with suspected or confirmed MERS-CoV infection should be placed in Airborne Infection Isolation Rooms (Negative Pressure Rooms), if available.
 When negative pressure rooms are not available, the patients should be placed in

adequately ventilated private rooms with a portable HEPA filter and is placed according to the manufacturer recommendations.

When single rooms are not available, suspected or confirmed MERS-CoV patients should be placed with other patients of the same diagnosis (cohorting). If this is not possible, place patient beds at least 1.2 meters apart.

5.4 PATIENT TRANSPORT

Avoid the movement and transport of patients out of the isolation room or area unless medically necessary. The use of designated portable X-ray, ultrasound, echocardiogram and other important diagnostic machines is recommended when possible.

If transport is unavoidable, the following should be observed:

- Patients should wear a surgical mask during movement to contain secretions.
- Use routes of transport that minimize exposures of staff, other patients, and visitors.
- Notify the receiving area of the patient's diagnosis and necessary precautions as soon as possible before the patient's arrival.
- Ensure that healthcare workers (HCWs) who are transporting patients wear appropriate PPE and perform hand hygiene afterward.

5.5 PERSONAL PROTECTIVE EQUIPMENT (PPE) FOR HCWS

The following PPE should be worn by HCWs upon entry into patient rooms or care areas in the respected order:

- Gowns (clean, non-sterile, long-sleeved disposable gown).
- Surgical mask (or N95 when airborne precautions are applied)
- Eye protection (goggles or face shield).
- Gloves.

For patients on airborne precautions, any person entering the patient's room should wear a fit-tested N95 mask instead of a surgical mask. For those who failed the fit testing of N95 masks (e.g. those with beards), an alternative respirator, such as a powered air-purifying respirator (PAPR), should be used.

- Upon exit from the patient room or care area, PPEs should be removed and discarded.
- Except for N95 masks, remove PPE at the doorway or in the anteroom. Remove N95 mask after leaving the patient room and closing the door.
- Remove PPEs in the following sequence: 1. Gloves, 2. Goggles or face shield, 3.
 Gown and 4. Mask or respirator.
- The following also should be noted:
 - The outside of gloves, masks, goggles and face shield are contaminated.
 - Never wear a surgical mask under the N95 mask as this prevents proper fitting and sealing of the N95 mask thus decreasing its efficacy.
 - For female staff who wear veils, the N95 mask should always be placed directly on the face behind the veil and not over the veil. In this instance, a face-shield should also be used along with the mask to protect the veil from droplet sprays.
 - Whenever possible, use either disposable equipment or dedicated equipment (e.g. stethoscopes, blood pressure cuffs and thermometers).

5.6 ENVIRONMENTAL CLEANING AND DISINFECTION

Recent data suggested that the environment in health care facilities used for MERS-CoV patients is widely contaminated. Thorough environmental cleaning and disinfection are critical.

- Consider designating specific, well-trained housekeeping personnel for cleaning and disinfecting of MERS-CoV patient rooms/units.
- Define the scope of cleaning that will be conducted each day; identify who will be responsible for cleaning and disinfecting the surfaces of patient-care equipment (e.g. IV pumps, ventilators, monitors, etc.).

- Consider using a checklist to promote accountability for cleaning responsibilities.
- Housekeeping personnel should wear PPE as described above. Housekeeping staff should be trained by the infection control team about MERS-CoV, in proper procedures for PPE use, including removal of PPE, and the importance of hand hygiene.
- Keep cleaning supplies outside the patient room (e.g. in an anteroom or storage area).
- Keep areas around the patient free of unnecessary supplies and equipment to facilitate daily cleaning.
- Use MOH-approved disinfectants (see the <u>GCC Infection Prevention and Control</u> <u>Manual, 3rd edition</u>). Follow manufacturer's recommendations for use-dilution (i.e. concentration), contact time, and care in handling.
- Clean and disinfect MERS-CoV patients' rooms at least daily and more often when visible soiling/contamination occurs.
- Give special attention to frequently touched surfaces (e.g. bedrails, bedside and over-bed tables, TV control, call button, telephone, lavatory surfaces including safety/pull-up bars, door knobs, commodes, ventilator and monitor surfaces) in addition to floors and other horizontal surfaces.
- Wipe external surfaces of portable equipment for performing x-rays and other procedures in the patient's room with a MOH-approved disinfectant upon removal from the patient's room.
- After an aerosol-generating procedure (e.g. intubation), clean and disinfect horizontal surfaces around the patient. Clean and disinfect as soon as possible after the procedure.
- Clean and disinfect spills of blood and body fluids by current recommendations for spill management outlined in the <u>GCC Infection Prevention and Control Manual, 3rd</u> <u>edition</u>.
- Cleaning and disinfection after MERS-CoV patient discharge or transfer:
 - Follow standard procedures for terminal cleaning of an isolation room. (See the <u>GCC Infection Prevention and Control Manual, 3rd edition</u>)

- Clean and disinfect all surfaces that were in contact with the patient or may have become contaminated during patient care including items such as blood pressure cuffs, pulse oximeters, stethoscopes, etc.
- Wipe down mattresses and headboards with an MOH-approved disinfectant.
- Privacy curtains should be removed, placed in a bag in the room and then transported to be laundered.
- No special treatment is necessary for window curtains, ceilings, and walls unless there is evidence of visible soil.
- Use hydrogen peroxide vapor or UVC machines for disinfection of the room as mandatory part of the terminal cleaning process.
- If all the procedures mentioned above are followed, then the patient room can be used immediately for another patient after terminal cleaning.

5.7 MEDICAL WASTE

Housekeeping staff must wear disposable gloves and perform hand hygiene after removal of gloves when handling waste.

Collection and disposal of MERS-CoV contaminated medical waste should follow the <u>GCC Infection Prevention and Control Manual</u>, 3rd edition.

5.8 TEXTILES

General concepts when dealing with linen in MERS-CoV patient's room are outlined in the GCC Infection Prevention and Control Manual, 3rd edition.

5.9 INFECTION PREVENTION AND CONTROL PRECAUTIONS FOR AEROSOL-GENERATING PROCEDURES

An aerosol-generating procedure (AGP) is defined as any medical procedure that can induce the production of aerosols of various sizes, including small (< 5 microns) particles.

AGPs includes bronchoscopy, sputum induction, intubation and extubation, cardiopulmonary resuscitation, open suctioning of airways, Ambu bagging, nebulization therapy, high frequency oscillation ventilation and Bilevel Positive Airway Pressure ventilation- BiPAP (BiPAP is not recommended in MERS-CoV infected patients because of the high risk of generating infectious aerosols and lack of evidence for efficacy).

Additional precautions should be observed when performing aerosol- generating procedures, which may be associated with an increased risk of infection transmission:

- Perform procedures in a negative pressure room.
- Limit the number of persons present in the room to the absolute minimum required for the patient's care and support.
- Wear N95 masks: Every healthcare worker should wear a fit-tested seal check N95 mask (or an alternative respirator if fit testing failed).
- Wear eye protection (i.e. goggles or a face shield).
- Wear a clean, non-sterile, long-sleeved gown and gloves (some of procedures require sterile gloves).
- Wear an impermeable apron for some procedures with expected high fluid volumes that might penetrate the gown.
- Perform hand hygiene before and after contact with the patient and his or her surroundings and after PPE removal.

5.10 FIT TEST AND SEAL CHECK

The protection offered by a disposable particulate respirator (e.g.N95) depends on its tight fitting to the user's face. Standardized respirator fit testing helps identify the correct respirator size and shape.

- Healthcare workers are required to have a respirator fit test at least once every 2 years and if weight fluctuates or facial/dental alterations occur.
- A fit test only qualifies the specific brand/make/model of a respirator with which an acceptable fit testing result was achieved and therefore users should only wear the specific brand, model, and size he or she wore during a successful fit test.

- Each time a respirator is donned, a seal check must be performed using the procedures recommended by the manufacturer of the respirator.
- For healthcare workers who have facial hair that comes between the sealing surface of the facepiece and the face of the wearer a Powered Air Purifying Respirator (PAPR) should be used instead.

5.11 MANAGEMENT OF EXPOSURE TO MERS-COV IN HEALTHCARE FACILITIES

5.11.1 Healthcare workers exposed to a MERS-CoV case

Healthcare facilities should identify and trace all health care workers who had protected (proper use of PPE) or unprotected (without wearing PPE or PPE used improperly) exposure to patients with suspected, or confirmed MERS-CoV infection.

The decision to permit a healthcare worker to resume his/her duties after an exposure to MERS-CoV should be individualized. Infection control team will be ultimately responsible for taking that decision.

The following are general guidelines but management will depend on the infection control team risk assessment:

- a. Asymptomatic healthcare workers WITH protected exposure OR unprotected lowrisk exposure (more than 1.5 meters of the patient):
 - Testing healthcare workers for MERS-CoV is not recommended
 - o Healthcare workers can continue their duties
 - Healthcare workers shall be assessed daily for 14 days post exposure for the development of symptoms
 - Healthcare workers should delay travel until cleared by infection control team.
 - Asymptomatic healthcare workers WITH protected exposure OR unprotected low-risk exposure are considered CLEAR if they:
 - Remain asymptomatic AND
 - The observation period is over (14 days post exposure).

- b. Healthcare workers who had unprotected high-risk exposure (within 1.5 meters of the patient) or have suggestive symptoms regardless of exposure type:
 - Healthcare workers shall stop performing their duties immediately.
 - Testing (Nasopharyngeal swabs) for MERS-CoV is required (preferably 24hr or more after the exposure)
 - Healthcare workers shall not resume their duties until cleared by infection control team.
 - Healthcare workers should delay travel until cleared by infection control team.
 - Healthcare workers who test positive for MERS-CoV (regardless of the exposure type); healthcare workers who develop MERS-CoV suggestive symptoms (regardless of the exposure type) and healthcare workers who had unprotected high-risk exposure are considered CLEAR if:
 - They are asymptomatic for at least 48 hrs AND
 - The observation period is over (14 days post exposure) AND
 - Had at least one negative RT-PCR for MERS-CoV.

5.11.2 Patients exposed to a MERS-CoV case

Patients can be exposed to MERS-CoV patients prior to diagnosis or due to the failure of implementing recommended isolation precautions. The following are general guidelines but management will depend on the infection control team risk assessment:

- Patients sharing the same room (any setting e.g. ward with shared beds, open ICU, open emergency unit, etc.) with a confirmed case of MERS-CoV for at least 30 minutes:
 - Testing (Nasopharyngeal swabs or deep respiratory sample if intubated) for MERS-CoV is required (preferably 24hr or more after the exposure).
 - Patients should be followed daily for symptoms for 14 days after exposure.
 - If negative on initial testing, exposed patients should be retested with RT-PCR if they develop symptoms suggestive of MERS-CoV within the follow up period.
 - Patients discharged during the follow up period must be reported to public health department to continue monitoring for symptoms.

5.12 OUTBREAK MANAGEMENT

Healthcare facility outbreak is defined as evidence of one or more secondary transmissions of MERS-CoV within the healthcare facility. The investigation of MERS-CoV outbreak is managed by the Infection Prevention Unit of the hospital, Regional Command and Control Center (RCCC) and Central Command and Control Center and is discussed in <u>Communicable Diseases Outbreaks in Healthcare</u> <u>Facilities; Management Guidelines</u>.

Interventions such as media communication, partial or complete closure of hospitals or units, and activation of surge plan must be coordinated with the central Command and Control Center.

Contact tracing and testing shall follow approved protocols. Indiscriminate testing hamper outbreak control efforts and waste valuable resources.

5.13 PATIENT TRANSPORTATION AND PREHOSPITAL EMERGENCY MEDICAL SERVICES

Patients who may have MERS-CoV infection may be safely transported in any emergency vehicle with the proper precautions.

- Train EMS staff, including drivers, on basic infection control skills with emphasis on respiratory protection. Like other healthcare workers, respirator fit testing is also required.
- Minimize the number of people involved in the transport.
- When possible, use vehicles that have a separate driver and patient compartments and close the door/window between these compartments.
- Use a vehicle equipped with a HEPA filter incorporated into the ventilation unit especially for transporting patients on mechanical ventilation. If this unit is not available, set the regular vehicle's ventilation system to the non-circulating mode.
- Transport staff including the driver shall use PPE as described above (Personal Protective Equipment for Healthcare Workers).
- Place a surgical mask on the patient (if tolerated) and have the patient cover the mouth/nose with a tissue when coughing.

- Oxygen delivery with a non-rebreather face mask may be used to provide oxygen support during transport.
- Coordinate with the receiving facility to receive the patient at the ambulance door and limit the need for EMS personnel to enter the emergency department.
- Remove and discard PPEs in a medical waste container and follow standard operating procedures for reprocessing used linen.
- Clean and disinfect the vehicle and reusable patient-care equipment using an MOHapproved hospital disinfectant. Personnel performing the cleaning should wear a disposable gown and gloves (a respirator is generally not needed).
- Ensure appropriate follow-up and care of EMS personnel who transport MERS-CoV patients as recommended for HCWs.

5.14 DURATION OF ISOLATION PRECAUTIONS FOR MERS-COV INFECTION

The infectivity period for MERS-CoV may last as long as virus is being shed. Out of protocol testing in confirmed MERS-CoV patients is discouraged. For all patients, retesting can be done at the end of the first week of confirmation.

In order to discontinue isolation precautions, two negative lower respiratory samples 24 hours apart are required for ventilated patients and one negative respiratory sample in other patients including home isolated individuals. (*Appendix C*).

6. PUBLIC HEALTH CONSIDERATIONS

6.1 SURVEILLANCE AND REPORTING

MERS-CoV is a category I reportable infectious disease (within 24 hrs). All healthcare facilities must report suspected cases through Health Electronic Surveillance Network (HESN). Results of laboratory testing are also reported through HESN. Failure of healthcare organizations or healthcare professionals to report reportable infectious diseases will result in legal actions and may affect licensing and certification.

6.2 HOUSEHOLD AND COMMUNITY CONTACTS MANAGEMENT

The public health team at the regional health directorate is responsible for listing, tracing and follow up of household and other contacts of patients with MERS-CoV infection in the community.

A communication link with a healthcare provider should be established for the duration of the observation period.

Community and household contacts of MERS-CoV cases are defined as a person who shared the same enclosed space (e.g. room, office) for frequent or extended periods with the index case while the index case is symptomatic. Contact tracing assessment forms must be filled out for all contacts (*Appendix D*).

Contacts are categorized by the presence or absence of suggestive MERS-CoV symptoms at the first assessment:

- Contacts without suggestive MERS-CoV symptoms should be listed for follow up (<u>Appendix D</u>). Screening for MERS-CoV is not generally required. In certain situations, MERS-CoV screening may be considered:
 - If the exposed contact had intense exposure to the MERS-CoV case (e.g. direct care, sleeping in same room)
 - If exposed contact is Immunocompromised (e.g. cancer, organ failure, use of immunosuppressive medications) or has other chronic underlying conditions (e.g. diabetes, hypertension)
- Contacts with suggestive MERS-CoV symptoms should be assessed clinically and referred to a healthcare facility if admission deemed necessary (<u>Appendix D</u>). A nasopharyngeal swab should be collected by a trained personnel and sent for MERS-CoV screening.

The observation period of a MERS-CoV community and household contacts is 14 days after the last exposure. Longer observation may be required if more than one generation of transmission is identified. Contacts who develop symptoms require enhanced monitoring for disease progression. Health status must be checked by phone and if feasible, by face-to-face visits on a daily base.

6.3 HOME ISOLATION GUIDANCE

Individuals infected with MERS-CoV who are stable enough can be safely managed at their homes. The public health team at the regional health directorate should assess whether the house is suitable for home isolation.

A suitable home setting entails:

- A dedicated well ventilated bedroom for the infected individual
- An educated healthy and rapidly accessible caregiver
- A reliable communication tool (e.g. mobile phone)

Recommendations to Individuals infected and the caregivers include:

- The infected individual is instructed to limit contact with others as much as possible and to strictly adhere to respiratory etiquette and hand hygiene.
- The household members should stay in a different room or, if not possible, maintain a distance of at least one meter.
- The household members should wear a medical mask when in the same room (within one meter) with the infected individual. Masks should not be touched or handled during use. If the mask gets wet or dirty with secretions, it must be changed immediately.
- Caregiver should use disposable gloves when handling the infected individual's body secretions and perform hand hygiene after removing gloves.
- Used mask, gloves, tissues and other disposable items should be discarded in a covered waste bin, and hand hygiene performed after touching these items.
- Touched surfaces in the infected individual's room should be cleaned daily with regular household cleaners or a diluted bleach solution (1 part bleach to 99 parts water). The bathroom and toilet surfaces should be daily with regular household cleaners or a diluted bleach solution (1 part bleach to 9 parts water).

• Soiled clothes, bed sheets, and towels of the infected individual should not be shaken. They can be cleaned using regular laundry soap and water.

6.4 HUMAN ANIMAL INTERFACE

Dromedary camels (Camelus Dromedarius) are the natural reservoir for MERS-CoV. Camel to human transmission seems to occur with direct or indirect contact with camels or their surrounding environment.

All community acquired MERS-CoV infections should be investigated for direct or direct links to camel (*Appendix D*). The exposure history might not be obvious and deep inquiries are usually necessary.

Direct or indirect exposure of human MERS-CoV cases to camels are reported to the field investigation team at the ministry of environment, water and agriculture.

Interventions from the animal health side include:

- Field visit to the presumed exposure site
- If camels are identified at the presumed exposure site, they will be quarantined and tested for MERS-CoV.
- Sampling testing techniques are detailed in the MEWA manual for field investigation.
- If live virus is detected in a camel herd, the quarantine period will be extended until the live virus is no longer detected.

As a general precaution, anyone visiting farms, markets, barns, slaughterhouses or other places where dromedaries are present should practice general hygiene measures, including regular hand washing after touching animals, avoiding touching eyes, nose or mouth with hands, and avoiding contact with sick animals. People should also consider wearing protective gowns and gloves while handling animals.

Slaughterhouses and meat processing plants are required to safely dispose heads and respiratory organs (trachea and lung) of slaughtered camels.

The consumption of raw or undercooked animal products, including milk and meat carries a high risk of infection from a variety of organisms that might cause disease in humans. Animal products processed appropriately through proper cooking or pasteurization are safe for consumption but should also be handled with care, to avoid cross-contamination with uncooked foods or from contaminated environment.

Camel barns, farms and markets must be permanently relocated outside residential areas. Since 2015, Hajj and Umrah zones are declared camel free areas.

For Approved MERS-CoV Surveillance forms see (Appendix D).

7. LABORATORY DIAGNOSIS OF MERS-COV

Laboratory testing for MERS-CoV is performed to confirm a clinically suspected case and to screen contacts as per approved protocols.

- Regional MOH and selected non-MOH governmental laboratories are approved to test for MERS-CoV by using validated commercial Real-time reverse-transcription polymerase chain reaction (rRT-PCR) assays.
- Laboratory confirmation of MERS-CoV infection requires either a positive rRT-PCR result for at least two specific genomic targets; region upstream and open reading frame1a (upE and ORF1a).
- It is strongly advised that lower respiratory specimens such as sputum, endotracheal aspirate, or bronchoalveolar lavage be used when possible. If patients do not have signs or symptoms of lower respiratory tract infection or lower tract specimens are not possible or clinically indicated, nasopharyngeal specimens should be collected.
- If initial testing of a nasopharyngeal swab is negative in a patient who is strongly suspected to have a MERS-CoV infection, patients should be retested using a lower respiratory specimen or, if not possible, repeat a nasopharyngeal specimen.

For guidelines on MERS-CoV Sample collection, packaging and shipping (Appendix E).

8. OTHER CONSIDERATIONS

8.1 GENERAL OUTLINES OF MANAGEMENT

Suspected or confirmed MERS-CoV patients should be admitted to health-care facilities only if medically indicated. Clinically stable patients or asymptomatic infections can be managed at home (see Home isolation guidance below).

Confirmed MERS-CoV cases can potentially be managed at any hospital. However, in certain occasions, it might be necessary to transfer a confirmed MERS-CoV case to a higher center in coordination with central command and control center. Indication for transfer to a MERS-CoV designated hospitals (see <u>Communicable Diseases Outbreaks in</u> <u>Healthcare Facilities; Management Guidelines</u>) include:

- Inability to comply with infection control requirements as decided by the regional command and control center (e.g. staffing issues, overcrowding, lack of isolation rooms).
- Reduce the risk of outbreak during mass gathering (e.g. transfer confirmed cases outside Hajj zone during the Hajj season).
- Critically ill patients who may require sophisticated potentially lifesaving interventions (e.g. Extra-Corporeal Membrane Oxygenation).
- MERS-CoV is still a relatively uncommon cause of pneumonia. Therefore, patients admitted with suspected MERS-CoV pneumonia should be treated as per the community acquired pneumonia guidelines.

The use of non-invasive ventilation (e.g. Bi-level Positive Airway Pressure- BiPAP) should be avoided in patients with suspected or confirmed MERS-CoV pneumonia. This intervention enhances the risk of infection transmission through the aerosol generation and it lacks evidence of efficacy over endotracheal intubation and mechanical ventilation.

Meticulous supportive care is paramount to decrease mortality from MERS-CoV infection.

The use of antivirals for MERS-CoV is not recommended outside clinical trials.

8.2 EXTRA CORPOREAL MEMBRANE OXYGENATION (ECMO)

There is evidence that ECMO may offer survival benefits in some MERS-CoV patients.

ECMO may be considered in patients with following parameters:

- Age < 60 years with a potentially reversible lung pathology
- Murray score for Acute Lung Injury of 3-4 despite optimal care
- ECMO is relatively contraindicated in some situations, for example:
- Any condition that would limit the benefit of ECMO (such as severe neurologic injury or advanced malignancy).
- Any contraindication to anticoagulation.
- High FiO₂ requirements (>90) or high-pressure mechanical ventilation (P-plat >30) for 7 days or more.
- Limited vascular access
- Transfer of a MERS-CoV patient to an ECMO center shall be decided mutually between referring and accepting physicians.

Patients who meet above conditions and require transfer to an ECMO center may be considered for ECMO cannulation on-site in the correct clinical setting and then transferred.

List of centers that provide ECMO services to respiratory failure patients resulting from MERS-CoV and other etiologies can be accessed on Command and Control Center (CCC) page on the MOH website (<u>www.moh.gov.sa/ccc</u>).

8.3 MANAGING BODIES OF DECEASED MERS-COV PATIENTS

Although no postmortem transmission of MERS-CoV has ever been documented, deceased bodies theoretically may pose a risk when handled by untrained personnel.

Body washing of MERS-CoV cases should preferably be done at hospitals. However, it can be safely performed in public washing facilities attached to mosques provided that the washers have been trained on relevant infection control precautions including appropriate use of PPEs.

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2. APPENDICES

Appendix A: Pneumonia Severity Index (PSI) scoring

Appendix B: Visual triage checklist

Appendix C: Algorithm for Managing Suspected MERS-CoV Patients

Appendix D: MERS-CoV Surveillance Forms

- Form 1 : MERS CoV Hospital Based reporting Form
- Form 2 : MERS CoV Community Surveillance Form
- Form 3 : Line Listing Record for Household and Other Contacts
- Form 4 : Line Listing Record for Healthcare Workers Contacts

Appendix E: Guidelines for MERS-CoV Sample Collection, Packaging and Shipping

APPENDIX A

Severity Scores for Community-Acquired Pneumonia (CURB 65)*

Clinical Factor	Points
Confusion	1
Blood urea nitrogen > 19 mg per dL	1
Respiratory rate \geq 30 breaths per minute	1
Systolic blood pressure < 90 mm Hg OR Diastolic blood pressure ≤ 60 mm Hg	1
Age ≥ 65 years	1
Total points	

* CURB-65 = Confusion, Urea nitrogen, Respiratory rate, Blood pressure, 65 years of age and older.

APPENDIX B

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Visual Triage Checklist

Visual Triage Checklist for Acute Respiratory Illness

Date:	Time	MRN:
Name:	ID#:	Hospital:

	Points (adults)	Pints (children)	Score
A. Clinical symptom/sign			
Fever	2	1	
Cough (New or worsening)	2	1	
Shortness of breath (New or worsening)	2	1	
Nausea, vomiting, diarrhea	1	-	
Sore throat and/or runny nose	1	-	
Chronic renal failure, CAD/heart failure	1	-	
B. Risk of exposure to MERS			
Exposure to a confirmed MERS case in the last two weeks	3	3	
Exposure to camel or products (Direct or indirect*) in the last two weeks	2	2	
Visit to a healthcare facility that had MERS case in the last two weeks	1	1	
Total Score			

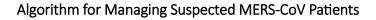
* Patient or household

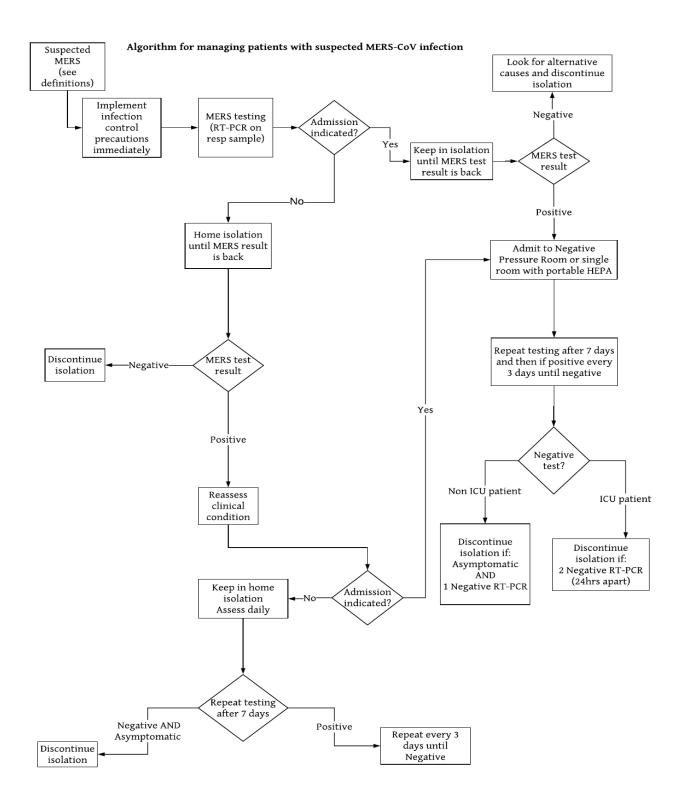
A SCORE ≥ 4, PLACE PATIENT IN AN ISOLATION ROOM AND INFORM MD FOR ASSESSMENT

MERS COV TESTING SHOULD BE DONE ONLY ACCORDING TO CASE DEFINITION

Staff name: _____ ID number: _____

APPENDIX C





APPENDIX D

MERS-CoV Surveillance Forms

MERS CoV Hospital B	ased repor	ting Form (I	Form 1)	الأوسط التنفسية		التبليغ لحالات متلاز بالمستشفيات (نموذ
Part 1. InitialNotificat (Suspect/Confirmed)	ion Form		ط التنفسية	متلازمة الشرق الأوس	بدئي لحالة	الجزء 1: التبليغ الم (مشتبة/ مؤكدة)
Date of initial notifica	ition:					تاريخ التبليغ:
1.0 Details on facility	reported s	uspect case	:	سحية: المبلغة للحالة:	لمنشأة الم	0.1 معلومات عن ا
1. Hospital Name:				1		1. اسم المستشفى:
City:		Province	2:	_ Region:		القصيم
2. Name of who comp	leted the		لاء محمد عبد العليم	الدكتور ء	عن	2. اسم من قام بالتبليغ
3. Phone Fax:			Mobile			3. رقم هاتف
4. Email:					:	4. البريد الإلكتروني
1.1 Case Information					حالة:	1.1 معلومات عن ال
اسم (Family) اسم		((First,			عبد الرحمن حسين
2. Date of Birth		dd/	mm/	уууу	عمر):	2. تاريخ الميلاد (ال
3. Gender:	Male 🗌	ذکر [Femal	انثى e		3. الجنس : ذكر
4. Nationality:						4. الجنسية :
5. Identification No.:						5. رقم الهوية :
6. Type of Identification:	نىية]_ ID	ا هوية وط	إقامة 🗌 qama	سفر [Passport]		6. نوع الهوية:
7. Hospital File numb	er (if			:	(إن توفر)	7. رقم الملف الطبي
8. Occupation: Health	Care Work	er:		:	ل الصحي	8. العمل : في المجا
lf No,				جاء تحديد نوع	ب لا ، الر	إذا كانت الإجابة
9. Phone Home:_		المنزل_	Mobile:			9. ارقام الهاتف
10. House I	lo. :	Stre	et Name :			10. عنوان الحالة:
District Name:	سم	City:		Province/R	egion: _	
11. Contact Person (f	iend,				ب	11. اسم شخص قري
12. Phone No.: Hom	e:		Mobile:			12. ارقام الهاتف:
1.2 Suspected case				ں:	كية للمريط	2.1 الخواص الإكليني
1.Date of onset of symptoms					راض:	1. تاريخ ظهور الأعر

2. F	Reason for testin عامل صحي	2. سبب طلب الفحص الـ nunity conta <u>ct</u>						
3. F	leason for testin	g					مخبر ي	3. سبب طلب الفحص ال
 1. Case criteria Fever and community- acquired Severe pneumonia (severity score ≥3 points)Appendix-A or ARDS (based on clinical or radiological evidence) 2.Case criteria Unexplained deterioration of a chronic condition of patients with congestive heart failure or chronic kidney disease on hemodialysis 		on Acute fe on ≥38° C) v	3.Case criteria Acute febrile illness (T ≥38° C) with/without respiratory symptoms		▲. Case criteria Gastrointestinal symptoms (diarrhea or vomiting), AND leukopenia (WBC≤3.5x10 ⁹ /L) or thrombocytopenia (platelets < 150x109/L)		. Infectious disease consultant recommended . Patient Asymptomatic	
1.3	3 Laboratory ME	RS CoV testing I	results			ų	لمخبري ونتائجه	3.1 معلومات الفحص ا
	e of specimen	Throat swab	Nasopharyr swab	ngeal	В	roncho-al	veolar lavage	Tracheal aspirate
col	lected:		EDTA blood	Tissue	e Biop	sy	🔲 sputum	Urine Oother
	Date sample colle	cted						تاريخ أخذ العينة:
1	Date sample sent							تاريخ إرسال العينة:
т •	Date result obtain	ed					خبري:	تاريخ ظهور نتيجة الفحص ال
	Lab Result:	Positive Negativ	ve Unclear	🗌 Rejec	ted			نتيجة الفحص المخبري:
Тур	be of specimen	☐ Throat swab	Nasopharyngeal Eswab			roncho-al	veolar lavage	Tracheal aspirate
col	lected:	Serum	EDTA blood	I Tissue Bionsy		Stool	🗌 Urine	
	Date sample colle	cted	_dd_	/_mm/_	yyyy_			تاريخ أخذ العينة:
	Date sample sent		_dd_	/_mm/_	yyyy_			تاريخ إرسال العينة:
2	Date result obtain	ed	_dd_	/_mm/_	yyyy_		خبري:	تاريخ ظهور نتيجة الفحص ال
	Lab Result:	Positive Degativ	ve Unclear	🗌 Rejec	ted			نتيجة الفحص المخبري:
			-					
	e of specimen	Throat swab	Nasopharyr swab	ngeal	В	roncho-al	veolar lavage	Tracheal aspirate
col	lected:	Serum	EDTA blood	🗌 Tissue	e Biop	sy	Stool	Urine Urine
	Date sample colle	cted	_dd_	/_mm/_	yyyy_			تاريخ أخذ العينة:
	Date sample sent		_dd_	/_mm/_	yyyy_			تاريخ إرسال العينة:
з •	Date result obtain	ed	_dd_	/_mm/_	yyyy_		خبري:	تاريخ ظهور نتيجة الفحص ال
	Lab Result:	Positive Degativ	ve Unclear	Rejec	ted			نتيجة الفحص المخبري:

Tv	pe of specimen	Throat swab	Nasopharyı swab	ngeal	Broncho-a	lveolar lavage	Tracheal aspirate	
	lected:	Serum	EDTA blood	Tissue	e Biopsy	Stool	Urine	
	Date sample collec	cted	_dd	/_mm/_	yyyy_	• .	ناريخ أخذ العينة:	
4	Date sample sent		_dd	/_mm/_	yyyy_		ناريخ إرسال العينة:	
•	Date result obtaine	ed	_dd_	/_mm/_	yyyy_	ىخبرى:	ناريخ ظهور نتيجة الفحص الم	
	Lab Result:	Positive Negativ	ve Unclear		تيجة الفحص المخبري:			
Ту	pe of specimen	Throat swab	Nasopharyi swab	ngeal	Broncho-a	lveolar lavage	Tracheal aspirate	
	be of specimen lected:		· ·		Broncho-a	lveolar lavage	Tracheal aspirate	
	•	swab	swab		e Biopsy			
col	lected:	swab	swab EDTA blood dd_	Tissue	e Biopsy		Urine [] Urine	
col	lected: Date sample collec	swab	swab EDTA blood _dd _dd		e Biopsy YYYY_ YYYY_	Stool	Urine	
	lected: Date sample collec Date sample sent Date result obtaine	swab	swab EDTA blood _dd_ _dd_ _dd_		e Biopsy YYYY_ YYYY_	Stool	Urine ناريخ أخذ العينة: ناريخ إرسال العينة:	

End of Part 1

Fill this part	when the Case	Confirmed				ص الحالة	عند تأكيد تشخي	يستكمل هذا الجزء ع
Part 2. Case	confirmation						كدة	الجزء 2: الحالة مؤ
2.1 Confirm	ation details						لمؤكدة:	1.2 تفاصيل الحالة ا
1. Is the case laboratory	e confirmed with y result.	n the positive		Yes 🗌	N نع	צ 🗌 כ	بالتحليل	 1. هل الحالة تأكدت المخبري:
the hospit have resol	to treatment guidelin al, he/she will remain ved, irrespective of the ctious disease is confin	suspected until sym e negative test resul	, ptoms	لر	اض بغضّ النو	حين إختفاء الأعر	انها مشتبهة إلى .	ملاحظة: إستناداً إلى دليل معاملة الحالة على عن النتائج السلبية مختلف.
-	cal picture makes a Mi ceed with the followin		robable,	, :	ابتها بمتلازما			إذا كانت الشواهد الإكليني الشرق الأوسط التنفسية،
2. How man househol	y people live in 1 d?	the same		(12.)		ع المريض	ذين يعيشون م	2. عدد الأشخاص ال بالمنزل:
3. Was pati was obtaine	ent hospitalized ed?	when the pos	itive r	esult	النتيجة	فی عند ظھور	منوم بالمستش	 ٤. هل كان المريض إيجابية ? نعم
	Date	e of admission:					تاريخ الدخول	
نعم 🗌 Yes	4. If hospitalize was the initial rea hospitalization? ب الرئيسي للتنويم؟	son for	ome c	onditions				المنزل غير مناسب لل
	5. Which depar م المنوم به المريض			العناية المر specify ,	:		ىدد القسم :	2
	6. If hospitalize Isolated?	d,	Yes [نعم [Vo لا	ة العزل؟	6. هل المريض بغرف
No 🗌 Y	7.lf not hospita Hospital basec إرسال نسخة من	ل form(Part 1	and P	art2)				ppy of the 7. إذا المريض لم يك كامل النموذج
	Date the case was Public Healthcare					لمنطقة/	محة العامة با	تاريخ إبلاغ إدارة ال المحافظة
	nt is not hospitalized confirmed, please sp	, ,		me G	تأكيد الحالا	المستشفى عند	,	8. في حال لم يكن ال الرجاء تحديد وضع
isolated ا زله بالمنزل		e at home بالمنزل بدو				d to another fac تنويمه بالمستشفى أو	,	Dead 🗌 متوفي
2.2 Patient	clinical informati	on on admissi	on		للمستشفى	ں عند الدخول	لينيكية للمريخ	2.2 المعلومات الإكا
1. Heig الطول_cm_	ght	2. Weight		k	الوزنg)	3. Tempe	erature:	درجة الحرارة)
4. Heart rate القلب	نبضاتع	5. Blood pres	sure:		ضغط الد	6. O ² satı رکسجین	uration, אוץ	تركيز
1. No sign	s, no movements	2 Mild		🔲 з Мо	derate	4 Seve	ere	

2.3 Other lab results pe admission or the most			نى	ذول المريد	لمخبرية الأخرى عند د. لأحدث:	3.2نتائج الفحوصات اا للمستشفى أو النتائج اا
1. White blood cells count		2. Creatinine			3. Lymphocytes %()
4. Platelet count	5. Neutroph	nils %()	6. Blood (Jrea Nitro	gen() 7.Dat	e:_dd_/_mm/_yyy
			1	·c		
2.4 History and Pre-Exi	isting Cond	itions (Comp	biete even	if patien	t is dead)	
1. Is the patient Healthcare w	vorker?	Yes		No	Unl	known
2. If yes, name of HealthCard	e facility whe	re patient is wo	rking:			
	🗌 Phy	vsician	Nurse	[X-ray technician	cleaner
3. If yes, type of the HC work	ker: 🗌 Re	spiratory rehab	ilitation the	rapist	Not patient care re	elated
	🗌 La	boratory worke	r		Other patient care	:
		U 🗌 Radio	ology	Emergen	cy room 0	utpatient
4. Department where worki	ng:	- Resp	piratory Reha	abilitation		care Dept.
5. Did the patient give care t	Dialua	is Dont				
		L	es No	Unkno	wn Non patient	care department
6. If not a healthcare worker, during the last 14 days before	did the patie		Ithcare facili	ies	Yes 🗌 No	Unknown
If yes, what healthcare facility:						
7. Did the patient have a	any pre-exi	sting				
conditions?			YES		NO	UNKNOWN
1. Diabetes Mellitus						
2. HIV/other immune deficienc	У					
a. On immune suppressive t	herapy				<u>_</u>	
b. On glucocorticoids						
c. Immune compromising d	isease					
 Hypertension Heart disease 						
5. Asthma						
6. Chronic liver disease						
7. Chronic haematological disord	der					
8. Chronic lung disease						
8. Pregnancy				s of	gestation	
9. Chronic kidney disease						
, 10. On dialysis						
11. Had kidney transplant						
12. Neoplastic disease						
14. 🔲 Other (specify):						
End of Part 2						

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Part 3	Follow up f	form - Tah	e filled unon	anv status ch	ange of the l	ocnitaliced	nation
Fails	. FOIIOW UD I	10111 - <i>100</i>	e meu ubon	univ slulus chi	unue or trie i	iosuituiiseu	bullen

3.1 Status update

S. N	Date (dd /mm/yyyy)	No change	Transferred to anotherhospital	Transfer within the hospital - Admitted to ICU	Transfer within the hospital from ICU to ward	Discharged for home isolation	Discharged to home- not isolated requires	Requires additional follow up	Died ⁶	Recovered from MERS CoV ⁷	Readmitted due to the condition worsening	Transferred from another hospital
1.												
2.	//											
3.	//											
4.												
5.												
6.	//											
7	//											
8	//											
9												
1 0	//											
1 1	//											

3.2 Complications developed during the hospitalization

	Yes	No	Unknown	Date developed
1. Pneumonia				_dd _/ _mm /_ yyyy
2. Acute renal failure				_dd _/ _mm /_ yyyy
3. ARDS (Acute Respiratory Distress				_dd_/_mm/_yyyy
4. Respiratory failure				_dd_/_mm/_yyyy
5. Cardiac failure				_dd_/_mm/_yyyy
6. Multi-organ failure				_dd_/_mm/_yyyy
Other (specify) :				dd/mm/yyy
3.3 If transferred to another hospital				
1. Name of the hospital transferred to:				2. Date of transfer:
3. Region:	4. City:			5. Sector:
3.4 When discharged from hospital				
1. Date of discharge:	2. Con	dition:	Alive	🗌 Deceased 🔲 Against Medical advice
3. Home isolation recommended.	<u> </u>	′es		No Unknown
4. Public Department of Health office	e name:		ate case trai dd_/_mm/_	nsferred to Public Department of Health
End of Part 3.		÷		

⁶ Complete the Case closure form
 ⁷ Complete the Case closure form

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Part .4 Case Close			الجزء 4: إغلاق الحالة		
4.1. Reason for case closure			1.4 أسباب إغلاق الحالة		
وفاة has bee	infectious disease en confirmed تم تأکید تشخیص	Give number of te	حدد عدد العيناتsts		
4.2 If died			2.4 إذا توفت الحالة		
1. Date of death 2. Death 2. Death ناريخ الوفاة (n certificate number: رقم وثيقة الوفاة)		
3. Place of death At home بالمنزل مكان الوفاة	ستشفى 🗌	بالم	Unknown 🗌 غیر معروف		
4. Post mortem tests performed? Yes هل تم إجراء تشريح للجثة	نعم 🗌	No 🗌 🎽			
Comment: Please attach the copy of the death certificate and send it by fax or Email to Public Health ملاحظة: الرجاء إرفاق نسخة من شهادة الوفاة وإرسالها بالفاكس أو البريد الإلكتروني إلى إدارة الصحة العامة بالمنطقة / المحافظة					
4.3 If discharged		من المستشفى	3.4 إذا خرجت الحالة		
1. Latest MERS test results Positive Negative, number of consecutive tests shown negative العينات results prior to discharge إيجابي نتيجة أخر تحليل كرونا السلبية، عدد العينات					
2. Discharge approved by:	م من اعتمد خروج المريض	لخروج اس	charge تاريخ اا (dd/_mm/yyyy)		
4.4 If another infectious disease has bee confirmed	n	ص أخر للحالة	4.4 إذا تم تأكيد تشخير		
1. Date of confirmation تاريخ تأكيد التشخيص الجديدdd/yy		causative agent التشخيص لمسبد	تم تأکید :confirmed		
End Part 4					

الإستقصاء الوبائي لحالات متلازمة الشرق الأوسط التنفسية (فيروس كرونا) (نموذج #2)

ملاحظة لمن يقوم بالاستجواب: Note for interviewer:

If you are interviewing a patient ask all questions in the first person, If you are interviewing the patient's relative or contact person ask questions in the third person.

جميع الأسئلة تخص المريض. فإذا كنت تقابل المريض فأسأله مباشرة، اما إذا كنت تقابل شخص من أقرباء المريض فتكون الأسئلة عن المريض.

Date Investigation	تاريخ إجراء
Start:	الاستقصاء:
Form completed by:	اسم من قام بإستكمال
	النموذج:
Phone number:	رقم الهاتف:
Permanent jobsite:	جهة العمل
	الأساسية:
المنطقة الصحية Health Region المنطقة الصحية:	
Part 1. Patient personal information	
1. First & Father name:	اسم
	المرء ض
	کن
2. Family name:	العائلة
	(اللقب):
3. GPS coordinates N E	إحداثيات موقع
	سكن الحالة:
4. Address in detail:	العنوان بالتفصي <u>ل:</u>
5. What type of housing? Single family Dormitory.	ما نوع المنزل الذي يسكنه
أخرى، Other, منزل home/villa	المريض ؟
specifyشقص ن/ فیلا شقةApartment	
6. Home phone: + () 8. Mobile phone:	+ ()Other
	mobile phone
9. Does the patient have another Yes Yes No کتم home?	هل للمريض منزل أخر؟

10 If yes: Address		Telephone Number				
	رقم التلفون		الإجابة نعم : ا لعنوان			
12. Is the patient the here household?	ad of Y	لا_No نعم _es	هل المريض هو رب العائلة؟			
If YES, move to part 1.2		لى الجزء 1.2 <u>-</u>	إذا كانت الإجابة بنعم، انتقل ال			
1.1 Head of Household S	ection		1.1جزء خاص برب العائلة			
 Name of head of household: 			اسم رب العائلة:			
2. Identification Number:			رقم الهوية لرب العائلة:			
هوية 🗌 National ID وطنية	اقامة 🗌 Iqama	جواز سفر 🗌 Passport	Others 🗌 اخرى			
 Relationship to patient: 	والد/والدة Parent العائل Spouse	الاطفال children أخرى، حدد Other, أخرى، حدد specify	صلة القرابة مع المريض			
4. Mobile phone numb	er: + ()	other mobile phone	رقم الجوال رب العائلة:			
1.2 Patient Social information	ation	ن	1.2 معلومات اجتماعية للمريخ			
 Education (Give highest year of school completed): 	متعلم	P دبلومDiploma متّmediate	الحالة التعليمية للمريض ماجستيرter (سجل اعلى مرحلة تعليمية وصل اليها المريض):			
2 Student طلاب Accup Employed/Government sector unemployed Unemployed Unemployed Unemployed ation: الحكومي الحكومي الحكومي Employed/Private sector Unemployed Other Other						
,	6,1	اسم المدر سة/الجامع:	إذا كان طالب، سجل			
	العملne of the employer العمل Iress العنوان:	اسم جهة:	إذا كان موظفاً، سجل التالي:			
5. Does the patient have Housemate / driver wo the home? 6. Sex الجنس 7. Age	orking in	s نعم No لا	هل المريض لديه خادمة/سائق؟			
If yes, include on contact list						
	مائهم ببيان المخالطين.	إذا كانت الإجابة نعم، اضف أس				

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Part 2 Personal Risk:	الجزء 2 عوامل الخطورة الشخصية:	
2.1 Smoking		2.1 التدخين
1. Does the patient smoke?	Yes نعم No Yes	هل المريض مدخن؟
2.lf yes, Specify: دCigarettes خان	معسل Nargghile المعسل Sh	إذا كانت الإجابة شيشة (جراك) eesha نعم، حدد
سيجارة Electronic Cigarettes [] الكترونية		
3. For how many years?		كم سنة يدخن؟
4. How many per day?		كم مرة باليوم؟
The following exposure questions before the patient developed the	•	التالي يتضمن أسئلة تغطي التعرض لعواما قبل 14 يوماً من إصابة المريض
2.2. Exposure to possible human s	ources	2.2 التعرض لمصدر عدوى إنساني محتمل
1. Did patient attend any mass gath	nerings?	هل حضر المريض إي تجمع كبير؟
Football or other large sporting	ة كرة قدم أو حدث رياضي كبير events	الجنادرية Janadria مبار
Um Rugaibah (Mazaieen-Camel	عمرة Omra	
Esterahah (extended family gatl	استراحة (تجمع عائلي كبير) (nering	☐Hajj حج □ Other Specify

2	3. Exposure to Human sources				عدوى إنساني	3.2 التعرض لمصدر
1.	Is the patient a healthcare	Y	es [] No انعم	צ	مجال	هل المريض يعمل بال
N.	B1 If patients a healthcare worker, p HOSPITAL FORM complete section		sure	*		ملاحظة 1: إذا كان المري بأن نموذج المستش
N.I	B2 If not a healthcare worker, please the following questions:	e provide a	nswers to	بال الصحي،		ملاحظة 2: إذا كان المري استكمل الجزء الت
2	Did the patient visit for any reas before onset of symptoms? بل ظهور الأعراض؟					لا_No نعم_Yes
3	Does the patient have regular v renal dialysis, diabetes manage ب غسیل کلوي، عیادة السکر ، الحمل، الخ)	ment, preg	gnancy, etc)			لا_No نعم_Yes
4	Did the patient visit a relative, r they were sick with a respirator ائه أو المدرسة اثناء مرضهم بالجهاز	y illness?				لا_No نعم

5	Where did this happe	en?	في المنزلAt home										
<u>اf yes,</u> إذا كانت الإجابة بنعم 6			في منشأة In a health care facility 🗌										
	Did the patient provi اية ذلك الشخص?person		لا_No Yes										
	mber diagnosed with ور الأعراض على المريض?k												
, ,	mber diagnosed with ونا بعد ظهور الأعراض على ا		s infection <u>After</u> patient Yes هل تم تشخير										
8. Was any other person who the patient knows personally diagnosed with Yes Yes منعم No Yes هل هنالك اشخاص يعرفهم المريض تم تشخيص حالتهم بالكورونا?MERS Corona virus													
لا_No نعم _Yes قبل ظهور الأعراض على المريض?Yes Yes													
Yes[19.2 After patient became sick? بعد ظهور الأعراض على المريض Yes[19.2 After patient became sick]													
2.4. Travel History 2.4. Travel History													
دل 14 يوم قبل ظهور الأعراض على المريض، هل سافر [1. During the 14 days before patient became محارج او داخل المملكة ؟ خارج او داخل المملكة ؟ اكانت الإجابة بنعم، أكمل بيانات الجدول ادناه: If yes, complete the following table:													
Country/City	Departure	Return date	Mode of travel										
- ,		el	خلال 3 أيام قبل ظهور الأعراض على المريض مريضاً، هل سافر خارج او داخل المملكة ؟ إذا كانت الإجابة بنعم، أكمل بيانات الجدول ادناه:										
Country/City	Departure	Return date	Mode of travel										
2.5. Exposure to CAN	IELS		5.2مخالطة الجمال										
1. Does the patient ra	ise camels? Ye	sNo	هل المريض لديه جمال ؟										
2. Is the patient's prot following:	fession one of the		هل المريض يعمل في أحد المجالات ادناه ؟ لا										
	Trader-لتاجر جمال amel	E He	erder يرعى الماشية										
SI	aughter man ينحر الماشية	🗌 Bu	جزار utcher										
Ca	يحلب الجمال amel milkier		eterinary طبيب بيطري										

	Other specify 🗌 يمتطي الجمال Camel rider 📄 يعمل ببيع اللحوم Sidewalk meat seller
3.	كا Do they have any other occupation that regularly deals Yes بعم No الجمال No No No With camels? هل المريض لديه عمل أخر يختص بالتعامل مع الجمال Specify بعدد
4.	خلال 14 يوم قبل من ظهور الأعراض على المريض، هل قام developed the illness did they :
	زار المسلخ Visit a slaughterhouse أحتك او لمس Touch a camel زار المسلخ Visit a live animal market جمل جمل
	شرب حليب ابل Drink camel milk حضر سباق Attend a camel race أمتطى Ride a camel milk جمال (الهجن) جمل
	تعامل مع لحوم جمال طازجةEat raw camel liver or partly cooked camel [] Handle raw camel meat] أكل كبدة ابل غير مطهية أو نصف مستويةliver

	List of patient's co ة من خدم وسائقين)					
S.N ت	Contact name أسماء المخالطين	Age العمر	Relationship to Patient صفة علاقته للمريض	Date swab Taken (Symptomatic) تاريخ أخذ المسحة (لمن بهم اعراض)	ldentific ation Number رقم الهوية	Type of Identification
1.						هویهٔ National I.D [قامةIqama جواز سفر Passport [
2.						هوية National I.D إقامةIqama جواز سفر Passport]
3.						هوية National I.D إقامةIqama جواز سفر Passport]
4.						هویهٔ National I.D قامةlqama جواز سفر Passport
5.						هویهٔ National I.D إقامةIqama جواز سفر Passport]
6.						هوية National I.D إقامةIqama جواز سفر Passport]
7.						هوية National I.D إقامةIqama جواز سفر Passport]
8.						هوية National I.D إقامةIqama جواز سفر Passport]
9.						هوية National I.D إقامةIqama جواز سفر Passport]
10.						هوية National I.D إقامةIqama جواز سفر Passport
Note:	Complete the table for all c includes Housemates & Dri additional page if needed.					ملاحظة: سجل جميع المخالطين ل الخادمات والسائقين. استخدم إذا استدعى الأمر.

Appendix D (count..)

					us change of the له المريض (يتم تعبئ		-						
	Data			تحديث وضع المريض الصحي Status update									
S.N ت	Date تاريخ الإتصال Method of contact بالمريض (dd (dd /mm/yyyy))		No change لا تغییر	Requires hospitalization يتطلب إحالته للمستشفى	Recovered from MERS CoV ^{8,9} تعافی من کرونا	Died ¹⁰ توفی							
1.	_/_/	Phone 🗌 ہاتف	زيارة Visit										
2.	_/_/	Phone 🗌 هاتف	زيارة []Visit										
3.	_/_/	Phone 🗌 ہاتف	زيارة []Visit										
4.	_/_/	Phone 🗌 ہاتف	زيارة []Visit										
5.	_/_/	Phone 📃 ہاتف	زيارة Visit										
6.		Phone 🗌 ہاتف	زيارة Visit										
7.	_/_/	Phone 🗌 ہاتف	زيارة []Visit										
8.	_/_/	Phone 🗌 هاتف	زيارة []Visit										
9.		Phone 🗌 هاتف	زيارة []Visit										
10.		Phone 🗌 هاتف	زيارة Visit										

⁸ No symptoms for 24 hours and latest swab MERS test is negative ⁹ Complete the Case closure form ¹⁰ Complete the Case closure form

Middle East Respiratory Syndrome Coronavirus; Guidelines for Healthcare Professionals - April 2018 - v 5.1

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Appendix

زء 4: إغلاق الحالة Part 4. Case Closure											
4.1. Reason for cas	e closure				الة	1.4 أسباب إغلاق الد					
Patient died [] وفاة المريض	Patient dis المريض	-	diseas confirm	r infectious e has been med تم تأكيد تشخيص]. G tests العينات	ive number of حدد عدد					
4.2 If died الحالة					٢	2.4 إذا توف					
_/1. Date of death (dd/ m) تاريخ الوفاة						h certificate ber: (رقم وثيقة الوفاة					
3. Place of death مكان الوفاة		At home بالمنزل	[In hospital		Unknown 📃 غیر معروف					
4. Post mortem tests p جراء تشريح للجثة			نعم 🗌 Yes		No	צ [
Comment: please a امة بالمنطقة / المحافظة	•			•							
4.3 If discharged المستشفى				لحالة من	فرجت اا	3.4 إذا					
1. Latest MERS test res بة أخر تحليل كرونا		Positive] إيجابي	negati	e, number of cons ve results prior to « قبل خروج الحالة من الم	discharge	سلبية،					
2. Discharge approved	by:	خروج المريض:	اسم من	<u>D</u> ate of discharg ريخ الخروج / / / (dd / mm / y	تار 						
4.4 If another infe للحالة	ctious diseas	e has been con	ifirmed	فيص أخر	أكيد تشك	4.4 إذا تم ت					
1. Date of confirmation / (dd / mm/yyyy)	د التشخيص الجديد _ا	تاريخ تأكي	agen	يينات 2. Specific causative agent confirmed: تم Give nur تأكيد التشخيص لمسبب أخر ،							

MERS-CoV Outbreak

Line Listing Record for Household and Other Contacts (Form 3)

Region: _____ Public Health Investigator: _____

	Record name	Daily Progress Use Legend: SF=Symptoms Free; F=Fever; C=Cough; N/V=Nausea/Vomiting; BA= Body Aches; H=Headache Died=Death HOS=Hospitalization																
	Name (To be typed in English and Arabic)	ID/ Iqama number	Age	Nationality	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1																		
2																		
3																		
4																		
5																		
6																		
7																		
8																		
9																		
10																		
11																		
12																		
13																		
14																		
15																		
16																		

MERS-CoV Outbreak

Line Listing Record for Healthcare Workers Contacts: (Form 4)

Facility: _____

Facility Contact:_____

		Perso ord name move nam	nd do n						Died=Dea	th HOS=H	ospitaliza	tion, Test	=MERS-C	oV testec	I				
	Name (To be typed in English and Arabic)	ID/ Iqama number	Age/ Sex	Nationality	Exposure risk (high or low)	Day 1 DD/ MM/ YY	Day 2 DD/ MM/ YY	Day 3 DD/ MM/ YY	Day 4 DD/ MM/ YY	Day 5 DD/ MM/ YY	Day 6 DD/ MM/ YY	Day 7 DD/ MM/ YY	Day 8 DD/ MM/ YY	Day 9 DD/ MM/ YY	Day 10 DD/ MM/ YY	Day 11 DD/ MM/ YY	Day 12 DD/ MM/ YY	Day 13 DD/ MM/ YY	Day 14 DD/ MM/ YY
1																			
2																			
3																			
4																			
5																			
6																			
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12																			
13																			
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16																			

APPENDIX E

Guidelines for MERS-CoV Sample Collection, Packaging and Shipping

APPROPRIATE COLLECTION, TRANSPORTATION AND STORAGE OF THE SAMPLE FOR MERS COV TESTING ACCORDING TO INTERNATIONAL STANDARDS PLAY A MAJOR ROLE IN THE ACCURACY OF THE RESULT

1. <u>General Considerations:</u>

- 1.1. Sample collection: Before collecting and handling specimens for Middle East Respiratory Syndrome Coronavirus (MERS-CoV), determine whether the person meets the current case definitions for Suspect, Probable or Confirmed cases.
- 1.2. Appropriate PPE should be worn by all laboratory staff handling these specimens (9.1, 9.2).
- 1.3. Proper biosafety policies and procedures should be maintained when collecting specimens (9.1, 9.2).
- 1.4. Use approved collection methods and equipment when collecting specimens.
- 1.5. Handle, store, and ship specimens following appropriate protocols.
- 1.6. It is very important to include patient national, Iqama or passport number in the request form to help trace records for patients that do doctor shopping. For illegal residents please put a note in the request which demonstrates that no Iqama is available due to illegal residency.

2. <u>Specimen type and priority:</u>

- 2.1. Best upper respiratory tract (URT) specimen is nasopharyngeal (NP) swab or combined nasopharyngeal and oropharyngeal (NP/OP) swab specimens in (9.3).
- 2.2. To increase the likelihood of detecting infection, lower respiratory Tract (LRT) specimens (Sputum, tracheal aspirate (TA), Endotracheal secretions, or Broncheo-alveolar lavage(BAL)) are preferred. Based on the current data, they are the most likely to provide positive results. However, this should not exclude another specimen from the URT to enhance viral detection in challenging samples (9.3).
- 2.3. Additional specimens such as blood and serum can be collected on presentation and in convalescence period. (Please refer to specimen collection No.3).
- 2.4. Respiratory specimens should be collected as soon as possible after symptoms start, ideally within 7 days and before antiviral medications are administered.
- 2.5. However, if more than a week has passed since onset of illness and the patient is still symptomatic, lower respiratory samples are the preferred samples.
- 2.6. Samples should not be stored in hospitals for more than 4 hours (at 4 8oC) before delivering by the courier. Delivery of MERS-CoV specimens allowed ONLY by the courier. Specimens pick up SHOULD be requested at the following number (800 6149999).
- 2.7. Label each specimen container with the unique MERS-CoV number; patient hospital ID number, specimen type, the date and the time of sample collection include patient national, Iqama or passport number.

Specimen Collection:

- 2.9. Use powder-less clean (Non-surgical) gloves when collecting specimens for MERS-CoV for PCR testing since, trace amount of powder in the sample could inhibit PCR testing producing false negative result (9.3).
- 2.10. All specimens should be regarded as potentially infectious, and HCWs, courier, laboratory personnel who collect, transport, or handle the clinical specimens should adhere rigorously to standard precautions to minimize the possibility of exposure to pathogens (9.3).
- 2.11. Ensure that HCWs who collect specimens should be properly trained on the technique and wear PPE appropriate for aerosol generating procedures.
- 2.12. Health caring facilities will assign and train personnel to perform nasopharyngeal swabbing.
- 2.13. <u>Respiratory Specimens:</u>
 - 2.13.1. Lower respiratory tract
 - 2.13.1.1. Broncheo-alveolar lavage (BAL), tracheal aspirate (TA) and/or pleural fluid should be collected whenever clinically appropriate: Collect 2-3 ml into a sterile, leak-proof, screw-cap sputum collection cup or sterile dry container. Refrigerate specimen at 2-8°C up to 48 hours; if exceeding 48 hours, freeze at-70°C and ship on dry ice.
 - 2.13.1.2. **Sputum:** (induced or spontaneous) ask the patient to rinse the mouth with water then expectorate deep cough sputum directly into a sterile, leak-proof, screw-cap sputum collection cup or sterile dry container. Refrigerate specimen at 2-8°C up to 48 hours; if exceeding 48 hours, freeze at-70°C and ship on dry ice.
 - 2.13.1.3. Mucoid specimens such as BAL, TA and sputum can be placed in VTM after collection to liquefy the specimens and preserve the trapped virus.
 - 2.13.2. Upper respiratory tract
 - 2.13.2.1. Nasopharyngeal and Oropharyngeal swabs (NP/OP swabs) MUST BE TAKEN TOGETHER. Use only synthetic fiber swabs with plastic shafts. Do not use calcium alginate swabs or swabs with wooden shafts, as they may contain substances that inactivate some viruses and inhibit PCR testing. Place swabs immediately into sterile tubes containing 2-3 ml of viral transport media. NP/OP specimens MUST BE combined, placing both swabs in the same vial. Refrigerate specimen at 2-8°C up to 48 hours; if exceeding 48 hours, freeze at- 70°C and ship on dry ice.
 - 2.13.2.2. Nasopharyngeal swabs: Insert a swab into the nostril parallel to the hard palate. Leave the swab in place for a few seconds to absorb secretions. Swab both nasopharyngeal areas.

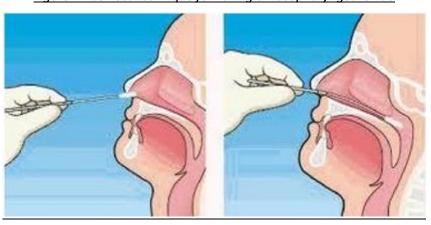


Figure 1: Correct Technique for Taking a Nasopharyngeal swab

For more information see NEJM Procedure: Collection of Nasopharyngeal Specimens with the Swab Technique: <u>http://www.youtube.com/watch?v=DVJNWefmHjE</u> <u>https://youtu.be/CcyLv67U8-Y</u>

- 2.13.2.3. Oropharyngeal swabs: Swab the posterior pharynx, avoiding the tongue.
- 2.13.2.4. Nasopharyngeal wash/aspirate or nasal aspirates: Collect 2-3 ml into a sterile, leak-proof, screw-cap sputum collection cup or sterile dry container (If highly mucoid, better collect in VTM container). Refrigerate specimen at 2-8°C up to 48 hours; if exceeding 48 hours, freeze at-70°C and ship on dry ice.

2.14. Blood Components

2.14.1. Serum (for Serological testing)

For serum antibody testing: Serum specimens should be collected during the acute stage of the disease, preferably during the first week after onset of illness, and again during convalescence \geq 3 weeks after the acute sample was collected. However, a single serum sample collected 14 or more days after symptom onset may be beneficial. Serological testing is for research/surveillance purposes and not yet for diagnostic purposes. Currently it is NOT available at the MOH regional laboratories but will be implemented soon.

2.14.2. Serum / Plasma (for rRT-PCR testing) (Not recommended for routine testing): For rRT-PCR testing (i.e., detection of the virus and not antibodies), a single serum or plasma specimen collected optimally during the first week after symptom onset, preferably within 3-4 days may be also beneficial but is not recommended for routine testing.

2.14.2.1. Serum Specimen:

2.14.2.1.1. Children and adults. Collect 1 tube (5-10 ml) of whole blood in a serum separator tube. Allow the blood to clot, centrifuge briefly, and separate sera into sterile tube container. The minimum amount of serum required for testing is 500 μl. Refrigerate the specimen at 2-8°C and ship on ice- pack; freezing and shipment on dry ice is permissible.

2.14.2.1.2. Infants. A minimum of 1 ml of whole blood is needed for testing of pediatric patients. If only 1 ml can be obtained, use a serum separator tube to achieve a minimum of 400 μl serum sample.

2.14.2.2. EDTA blood (plasma):

Collect 1 tube (10 ml) of EDTA (purple-top) blood. Avoid using heparinized (green-top) blood as this will interfere with the test and inhibit PCR. Refrigerate specimen at 2-8°C and ship on ice pack; do not freeze.

3. <u>Shipping:</u>

- 3.1. Specimens from suspected MERS-CoV cases must be packed, shipped, and transported according to the current edition of the <u>International Air Transport</u> <u>Association (IATA) Dangerous Goods Regulations</u> prepared by IATA licensed laboratory-personnel (9.4, 9.5).
- 3.2. At present MERS-CoV diagnostic specimens must be assigned to UN3373 and must be packaged as Category B infectious substances.
- 3.3. Packing responsibility is by the sample collection laboratory personnel and the shipment booking will be scheduled at the collection site in coordination with receiving laboratory.
- 3.4. Ensure that personnel who transport specimens are trained in safe handling practices and spill decontamination procedure.
- 3.5. Place specimens for transport in leak-proof specimen bags (secondary container) that have a separate sealable pocket for the specimen (i.e. a plastic biohazard specimen bag), with the patient's label on the specimen container (primary container), and a clearly written request form.
- 3.6. Ensure that health-care facility laboratories adhere to appropriate biosafety practices and transport requirements per the type of organism being handled.
- 3.7. Deliver all specimens by hand whenever possible. Do not use pneumatic tube systems to transport specimens.
- 3.8. State the name of the suspected ARI patient of potential concern clearly on the accompanying request form. Notify the laboratory as soon as possible that the specimen is being transported.
- 3.9. Shipment collection must be at Laboratory site. Time of shipment collection must be documented within AWB.

4. <u>Labeling:</u>

The outer container of all specimen packages must display the following on two opposite sides:

- Sender's name and address.
- Recipient's name and address.
- The words "Biological Substance, Category B".
- o UN 3373 label.
- Class 9 label, including UN 1845, and net weight if packaged with dry ice.

5. <u>Packaging:</u>

Specimens must be triple-packaged and compliant with IATA **Packing Instruction 650**, which is detailed in Figure 1. The maximum quantity for a primary receptacle is 500 ml or 500g and outer packaging must not contain more than 4 L or 4 kg **(9.5)**.

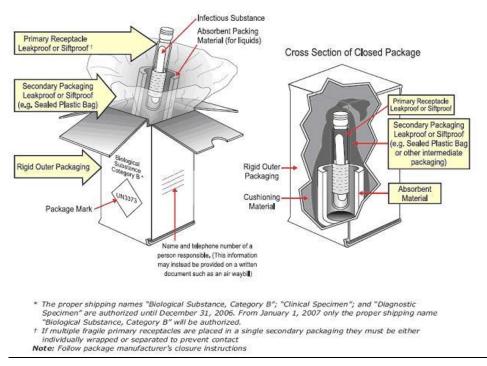


Figure 2: Packing Instruction Biological Specimens Category B

For more information on proper packaging for biological specimens "category B" see the technique on: https://youtu.be/GJK9FRT4IXM

5.1. Packing Containers

- 5.1.1. Packages must be of good quality, strong enough to withstand the rigors of transport.
- 5.1.2. Triple packaging consisting of leak proof primary receptacles (for liquid shipments), silt proof primary receptacles (for solid shipments), leak proof secondary packaging and outer packaging of sufficient strength to meet the design type test (1.2 meter drop test).
- 5.1.3. For liquid shipments, primary receptacle or secondary packaging capable of withstanding a 95Kpa internal pressure differential.
- 5.1.4. Absorbent material must be sufficient to absorb the entire contents of the shipment.
- 5.1.5. An itemized list of contents must be included between the secondary and outer packaging.
- 5.1.6. "Biological Substance, Category B" must appear on the package.
- 5.1.7. Minimum dimension is 100mm.
- 5.1.8. When large numbers of specimens are being shipped, they should be organized in a sequential manner in boxes (numerical order of patient hospital ID) with separate compartments for each specimen.
- 5.1.9. Patient Data Sheets and an Itemized List of Contents will accompany the package. The paperwork will be packaged inside the outer package NOT in the secondary container.
- 5.1.10. All specimens must be pre-packed to prevent breakage and spillage. Each specimen container should be sealed with Parafilm (after being crewed properly) and placed in a separate zip-lock bag.

- 5.1.11. Place enough absorbent material to absorb the entire contents of theSecondary Container (containing Primary Container) and separate the PrimaryContainers from each other (containing specimen) to prevent breakage.
- 5.1.12. Send specimens with cold packs or other refrigerant blocks that are selfcontained (do not use actual wet ice). This prevents the appearance of a spill due to thawed ice.
- 5.1.13. The courier will supply specimen transport container.

6. <u>Rejection of packages and samples:</u>

Apply universal rejection policy with emphasizes of the following:

- 6.1. All rejected samples will be discarded according waste management protocols in the laboratory, and immediate feedback will be given to the courier and treating physicians, the treating physician will decide if another sample is necessary).
- 6.2. Samples are not packaged according to packing instruction **P650** as **UN3373** Diagnostic Specimens.
- 6.3. An Itemized list of samples organized by Hospital Patient ID number is NOT included inside the outer package.
- 6.4. Any sample received without HESN investigations number printed clearly in request form will be rejected. In addition, results of samples received without filling MERS-CoV F117 form will be held till F117 form filled by the sender and informed to the laboratory.
- 6.5. Any mismatch or missing data between the specimen and the request form.
- 6.6. The patient data sheets are incomplete, missing or incorrectly filled out.
- 6.7. Any leakage or spillage, inside or outside the primary or secondary containers.
- 6.8. If dry ice is placed in the "Primary Container" or "Secondary Container", foam envelopes, zip-lock bags, cryo-vial boxes, or hermetically sealed containers.
- 6.9. If the Primary Containers sideways or upside down in zip-lock bags.
- 6.10. Primary containers must be packaged securely in an upright position and in the numerical order used on the Itemized List of contents.
- 6.11. If red top Secondary Containers for Category A Infectious Substances are used.
- 6.12. If any paperwork in the Secondary Containers or zip-lock bags.
- 6.13. The quality of the shipment conditions specially the temperatures of the specimens (warm).
- 6.14. Wrong swab; the swabs should not be cotton with wooden shaft as cotton will absorb the testing material (VTM) and wood could inhibit PCR testing and give false negative result.
- 6.15. Expired VTM.
- 6.16. Delay in specimen's shipment.
- 6.17. Blood samples sent in wrong tube, e.g. heparinized (green-top) tube.

7. <u>Turn Around Time (TAT) for Testing MERS CoV:</u>

- 7.1. TAT up to 24 hours.
- 7.2. A minimum of 2 runs per day.
- 7.3. For urgent samples: (Prioritizing) immediately.