2019 Novel Coronavirus (2019-nCoV)

Compiled by Centre for Respiratory Diseases and Meningitis and Outbreak Response, Division of Public Health Surveillance and Response, National Institute for Communicable Diseases (NICD) of the National Health Laboratory Services (NHLS)

National Department of Health, South Africa Including Communicable Diseases Cluster, Zoonotic Diseases Cluster, Port Health, Environmental Health and Emergency Medical Services

> **VERSION 4** 2020-02-06



Department: Health **REPUBLIC OF SOUTH AFRICA**



Training slides based on guidelines for case-finding, diagnosis, management and public health response in South Africa

and

Outline

- Welcome and objectives
- Microbiology, epidemiology and clinical presentation
- Laboratory diagnosis
- Infection prevention and hospital readiness
- Patient flow and actions required at each step
- Co-ordinating a public health response

Surveillance for imported cases including case definitions



BEFORE USING THIS POWERPOINT AND GUIDELINES PLEASE CHECK FOR UPDATES ON THE NICD AND NDOH WEBSITES www.nicd.ac.za and www.ndoh.gov.za

or CALL YOUR PROVINCIAL COMMUNICABLE **DISEASE CO-ORDINATOR**



THIS SITUATION IS RAPIDLY EVOLVING

Objective of training

- guidelines for
 - surveillance,
 - case detection/diagnosis
 - and management, and
 - 2019-nCoV

To familiarise attendees with RSA

 public health response to suspected and confirmed cases of infection with

Microbiology, epidemiology and clinical presentation



Introduction

- 31 December 2019, the World Health Organization (WHO) China country office reported a cluster of pneumonia cases in Wuhan, Hubei Province of China
- 7 January 2020, causative pathogen identified as a novel coronavirus (2019nCoV)
- Initially person-to-person transmission not apparent and the majority of the cases were epidemiologically linked to a seafood, poultry and live wildlife market (Huanan Seafood Wholesale Market) in Jianghan District of Hubei Province
- Number of cases continued to increase rapidly, and evidence of person-to-person transmission mounted



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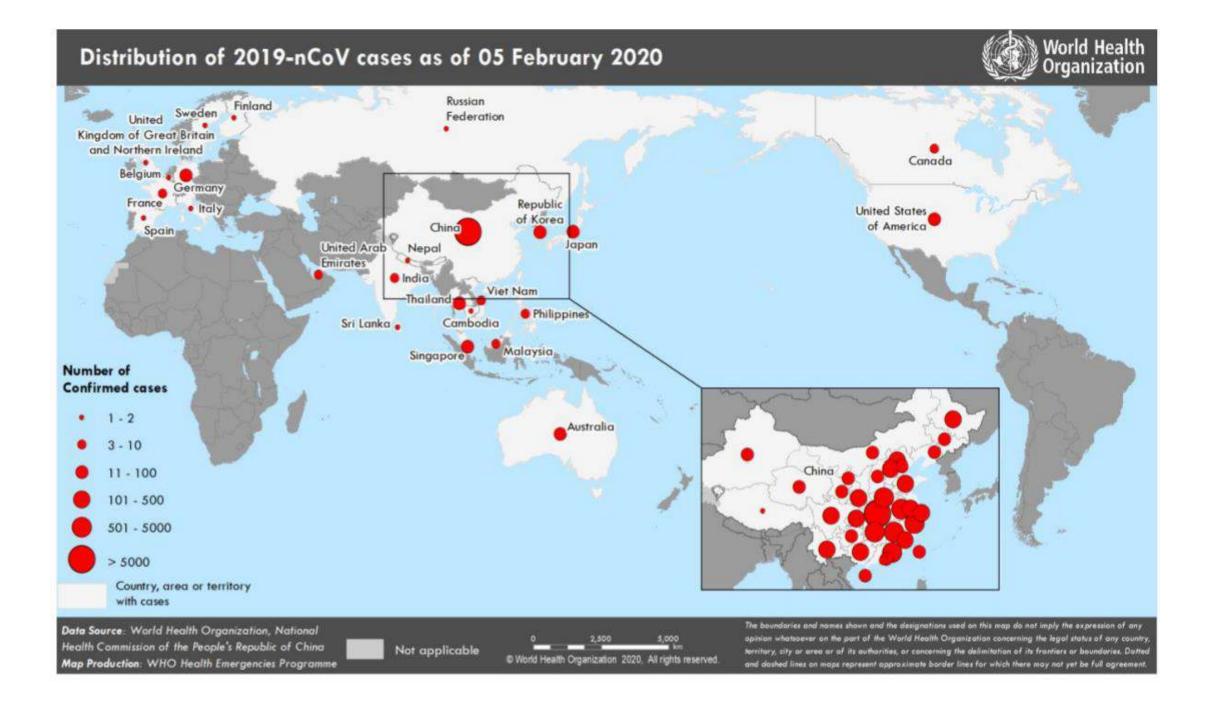
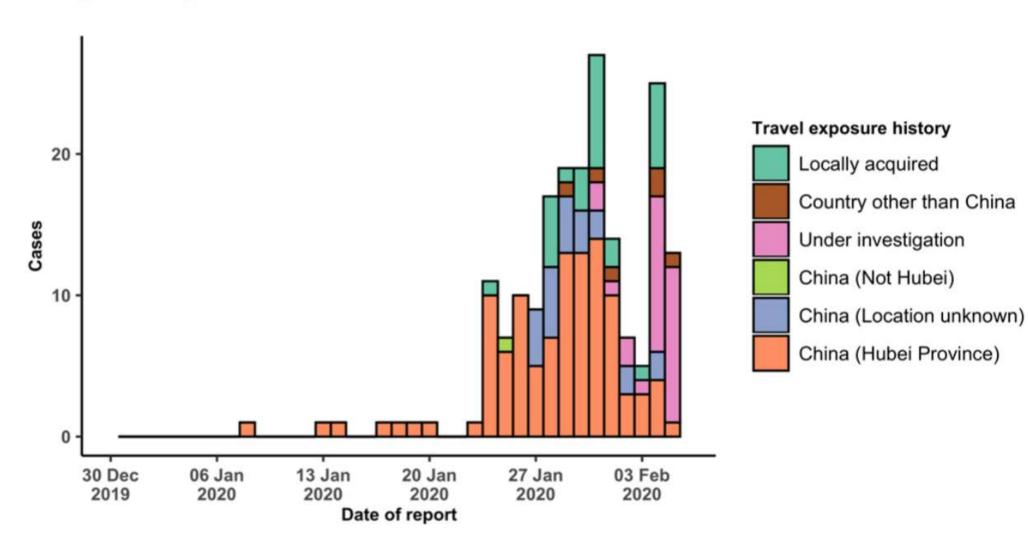


Figure 3: Epidemic curve of 2019-nCoV cases (n=191) identified outside of China, by date of reporting and travel history, 5 February 2020

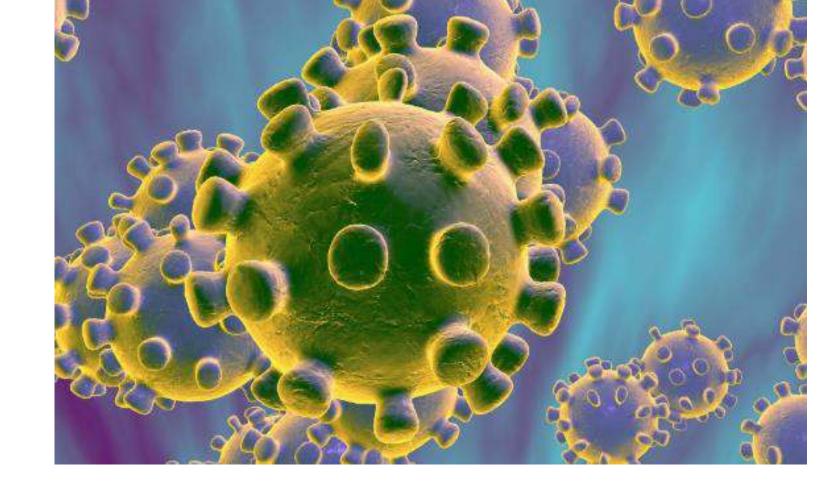


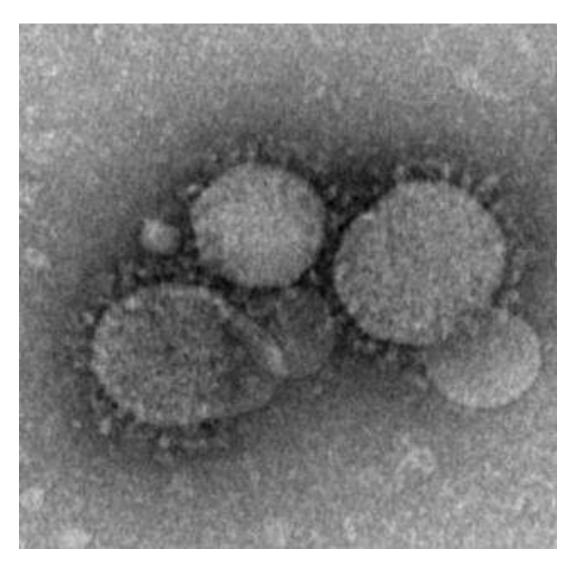




Microbiology and epidemiology

- Coronaviruses are enveloped, single-stranded positive-sense RNA viruses.
- The envelope of the coronaviruses is covered with club-shaped glycoproteins which look like 'crowns', or 'halos' – hence the name 'coronavirus.'
- Coronaviruses are responsible for the common cold, and usually cause self-limited upper respiratory tract infections.
 - Examples 229E, NL63, OC43 and HKU1





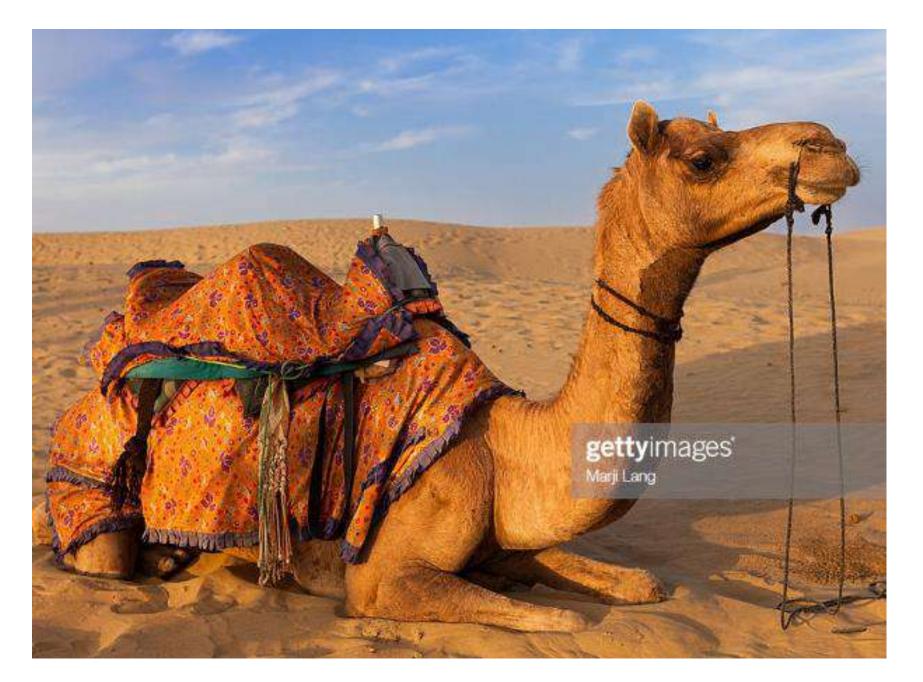




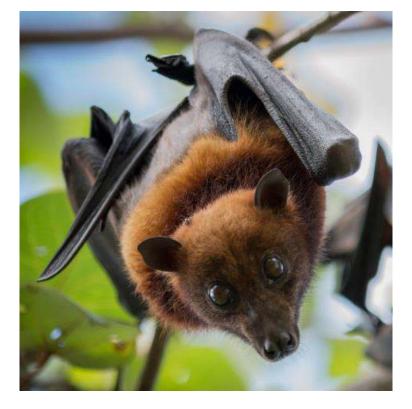
Microbiology and epidemiology

- In 2003, a new coronavirus emerged leading to the SARS (severe acute respiratory syndrome) outbreak.
- In 2012, the Middle East respiratory syndrome (MERS) was found to be caused by a coronavirus associated with transmission from camels.
- Following the identification of a cluster of pneumonia cases in Wuhan, Hubei Province of China, Chinese authorities reported on 7 January 2020 that the causative pathogen was identified as a novel coronavirus (2019-nCoV).
- These new coronaviruses have RNA sequences that are very similar to coronaviruses from animals
 - MERS-CoV = camel coronavirus
 - SARS = bat coronavirus

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Coronaviruses are a large family of viruses that cause illness ranging from the common cold to more severe diseases like pneumonia, MERS and SARS

- Sever Symptoms .
- **High Fever** .
- 38°C
- Pneumonia
- **Kidney Failure**
- Death

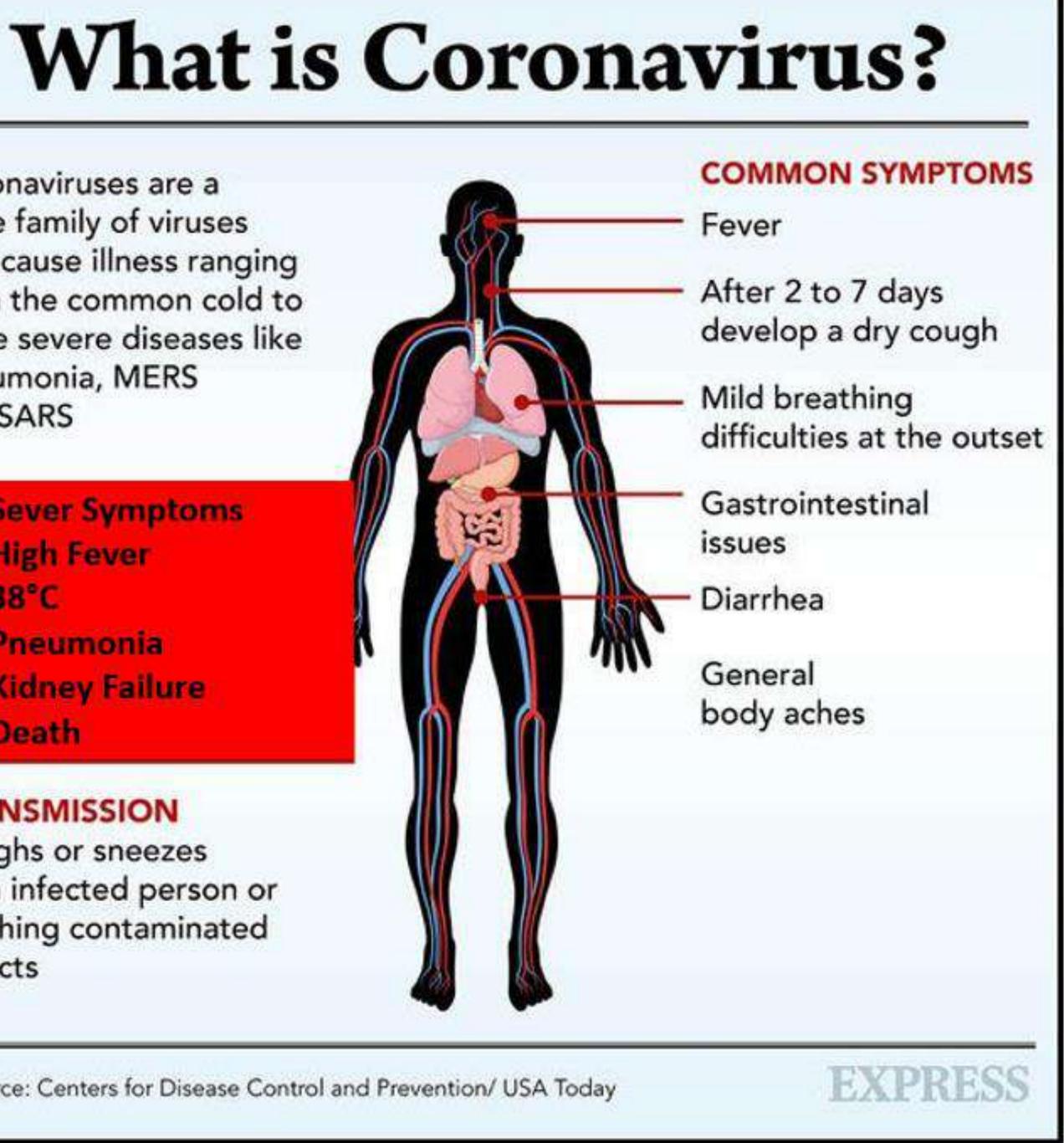
TRANSMISSION

Coughs or sneezes from infected person or touching contaminated objects

* Source: Centers for Disease Control and Prevention/ USA Today

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Transmissibility

- Main route of transmission respiratory droplets (airborne transmission has not proven)
- Excreted in stool (possibly faeco-oral)
- Mean incubation period 5.2 days (95%) confidence interval [CI], 4.1 to 7.0), 95th percentile of the distribution at 12.5 days.
- 14 days of isolation or quarantine is suggested as it allows a window of 1.5 additional days. (Li, 2020)
- In early stages, epidemic doubled in size every 7.4 days
- Basic reproductive number was estimated 2.2 (95% CI, 1.4 to 3.9) - on average each infectious case gives rise to just over 2 infectious cases.







Clinical presentation

- Who is at highest risk?
 - Largest published series to date from China 99 2019-nCoV patients with pneumonia the commonest symptoms were fever (83%), cough (82%) and shortness of breath (31%). (Chen et al Lancet 2020)
 - The majority (but not all) of severe cases are elderly or have severe underlying illness

 - Among pneumonia patients 51% had chronic diseases • 11 patients who died, 7 aged >60 years, 3 had long history of smoking and 3 had hypertension

Number of cases and deaths continue to increase

- Approximately 2% of reported confirmed cases have died
- Likely a substantial overestimation of the true case fatality ratio:
 - More severe disease tends to be reported first
 - Initial case definition in China really focused on patients with pneumonia
 - Possible backlog in testing and confirming cases in China







Surveillance and case definitions

Clinical and epidemiological criteria for person under investigation (PUI)

or not

In the 14 days prior to onset of symptoms, met at least one of the following epidemiological criteria:

- Were in close contact with a confirmed or probable case of 2019-nCoV infection;
- nCoV; i.e. China
- were being treated.



 Patients with acute respiratory infection (sudden onset of at least one of the following: cough, sore throat, shortness of breath) requiring hospitalisation

AND

OR

• Had a history of travel to areas with presumed ongoing community transmission of 2019-

OR

Worked in or attended a health care facility where patients with 2019-nCoV infections

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Who Should be tested

- Presently, the only persons who should undergo testing for 2019-nCoV are those described above under Person Under Investigation (PUI).
- All case to be discussed with NICD doctor on call before collecting samples
- The test will be free of charge for patients meeting the case definitions above



NICD Hotline 082-883-9920





If testing is indicated, what next?

- Isolate the patient using appropriate infection prevention control (see next section)
- Collect a specimen ASAP (see next section)
- Identify contacts



If testing is indicated, what next?

- Isolate the patient using appropriate infection prevention control (see next section)
- Collect a specimen ASAP (see next section)
- Identify contacts \bullet



- includes,

 - - seated.

Who is a close contact

• A person having had face-to-face contact (within 2 metres) or was in a closed environment with a 2019-nCoV case; this

• amongst others, all persons living in the same household as a 2019-nCoV case and, people working closely in the same environment as a case.

• A healthcare worker or other person providing direct care for a 2019-nCoV case.

• A contact in an aircraft sitting within two seats (in any direction) of the 2019-nCoV case, travel companions or persons providing care, and crew members serving in the section of the aircraft where the index case was

How to do contact tracing and monitoring of close contacts

- Once laboratory testing confirms 2019-nCoV infection:
- Provincial CDCC needs to identify close contacts, and make make a contact line list using Appendix in guidelines (see next slide)
- EVERY contact to complete the contact demographic section on the contact monitoring form PDF version at: <u>http://www.nicd.ac.za/diseases-a-z-</u> <u>index/novel-coronavirus-infection/ (see next slide)</u>
- Completed linelist and contact form also to be emailed to <u>ncov@nicd.ac.za</u>
- Close contacts will be asked to self-quarantine at home for 14 days since exposure to the confirmed 2019-nCoV and take their temperature daily (thermometers need to be issued)
- CDC / NICD/ delegated person will call contacts telephonically to identify if symptoms are present





Monitoring of close contacts and Health workers with occupational exposure

- monitoring dependant on the number of contacts to be followed up.
- Close contacts under monitoring should be advised to:
 - or education facilities)
 - Avoid unnecessary social contact
 - Avoid travel
 - Remain reachable for monitoring

Health Worker with occupational Exposure

- compiled by the health facility
- and tested should symptoms develop

Monitoring of close contacts may switch from telephonic monitoring to self-

• Remain at home (NICD can provide an official letter for employment

• Lists of healthcare workers with occupational exposure should be

• They should be actively monitored for symptoms and rapidly isolated

Quarantine

- \bullet persons
- lacksquarecontagious disease from healthy individuals without that contagious disease
- members of society.
- Quarantine may take place ullet
 - in the home
 - or in a designated facility.
- Depending on level of risk, and intensity of the exposure, different levels of quarantine will be \bullet employed, for example
 - If a person is expatriated from Wuhan, voluntary quarantine at a facility will be recommended. ●
 - A household member of a confirmed case will be asked to stay in their home for 14 days \bullet
 - \bullet work but would be requested to self-quarantine if symptoms develop within 14 days.

Quarantine means separating asymptomatic persons who are exposed to a disease from non-exposed

Quarantine is to be distinguished from isolation, which is the act of separating a sick individual with a

• Quarantine procedures can be effective in limiting and slowing the introduction of a novel pathogen into a population but may entail the use of considerable resources and may infringe on the rights of

if health worker wearing appropriate PEP is exposed to a confirmed case, the health worker would be allowed to

Contact line List

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2019-nCoV CONTACT LINE LIST

Complete a contact line list for every case under investigation and every confirmed case

	Details of case un	ider invest	igation	/confirmed case		Details of health of	ficial completing this form	Today's date	DD/MM/YYYY
NICD Identifier				ate Symptom nset	DD/MM/YYY	Surname		Name	
Surname			N	ame		Role		Facility name	
Contact number			A	ternative number		Email address		Telephone number(s)	
Travel (prov	vide details of all:7 day	rs before or	nset)	Travelled by	Bus Delane				
Air/bus line	15/	F	light/b	us #	Seat #				
Details of contac	cts (With close contact	¹ 7 days p	rior to	symptom onset, o	r during sympto	matic illness.)		· · · · · · · · · · · · · · · · · · ·	
Surname	First name(s)	Sex (M/F)	Age (Y)	Relation to case ²	Date of last contact with case	Place of last contact with case (Provide name and address)	Residential address (for next month)	Phone number(s), separate by semicolon	HCW? ³ (Y/N) If Yes, facility name
					DD/MM/YYYY				
					DDZMM/YYYY				
					DD/IMM/YYYY				
					DD/MM/YYYY				
					DD/MM/YYYY	S			
					DD/IMM/YYYY				×
					DDZMM/YYYY				
		0			DD/MM/YYYY				

8

2

3

5

6

¹ Close contact: A person having had face-to-face contact (<2 metres) or was in a closed environment with a 2019-nCoV case; this includes, amongst others, all persons living in the same household as a 2019-nCoV case and, people working closely in the same environment as a case. A healthcare worker or other person providing direct care for a 2019-nCoV case, while not wearing recommended personal protective equipment or PPE (e.g., gowns, gloves, NIOSH-certified disposable N95 respirator, eye protection). A contact in an aircraft sitting within two seats (in any direction) of the 2019-nCoV case, travel companions or persons providing care, and crew members serving in the section of the aircraft where the index case was seated..² Chose from: Aunt, Child, Class mate, Colleague, Cousin, Father, Friend, Grandfather, Grandmother, Healthcare worker taking care of, Mother, Nephew, Niece, Other relative, Uncle.³ Healthcare worker.

Page 1 of 2 Continues on reverse

Please refer to www.nicd.ac.za for most recent version of this document before use.

PDF version at: http://www.nicd.ac.za/diseases-a-z-index/novel-coronavirus-infection/ To be emailed to PDF version at: http://www.nicd.ac.za/diseases-a-z-index/novel-coronavirus-infection/



Version 4, 5 February 2020

Close Contact Monitoring Tool



NATIONAL INSTITUTE FOR COMMUNICABLE DISEASES 2019-nCoV DAILY SYMPTOM MONITORING TOOL

If not captured electronically at site, forward to <u>ncov@nicd.ac.za</u>, on completion of last day of monitoring.

Deta	ails of <u>contact</u> of cas	se under investi	gation/confirmed
NICD Identifier	Date last contact	DD/IMM/YYYY	Place last contact
Surname	76	Name	
Date of birth	DD/MM/YYYY	Age (Years)	Sex N
Contact #		Alternative contact #	
Relation to case		Place of contact	
Healthcare worker	Y 🗆 N 🗆	 Facility name	
Traced	Y 🗆 N 🗆	Contact type*	Close 🗆 Casual 🗆
Email		Monitoring method**	Direct Self-digital telephonic Active
Quarantine	Home 🗌 Facility 🗌	Facility where quarantined	
	Physical address	(for next month	, in South Africa
House #	Street		Suburb
Town		Municipality	
District		Province	
2	Next of kin or alt	ernative contac	t person details
Name,		Contact	
surname		number(s)	

DAY	1	2	3	4	5	6	7
Date (DD/MM)							
Fever (≥38°C)	□ Y □ N		□ Y □ N		□ Y □ N	□ Y □ N	ΟΥΟΝ
Chills	□ Y □ N		□ Y □ N	ΟΥΟΝ	□ Y □ N	□ Y □ N	ΟΥΟΝ
Cough	□ Y □ N	ΟΥΟΝ		ΟΥΟΝ	ΠΥΠΝ		Ο Υ Ο N
Sore throat	□ Y □ N	Π Y Π N			ΠΥΠΝ		Ο Υ Ο Ν
Shortness of breath		ΠΥΠΝ			ΠΥΠΝ		
Myalgia/body pains	□ Y □ N	ΠΥΠΝ	□ Y □ N		ΠΥΠΝ		
Diarrhoea	□ Y □ N			ΟΥΟΝ	ΠΥΠΝ		

DAY	8	9	10	11	12	13	14
Date (DD/MM)							
Fever (≥38°C)		ΠΥΠΝ	□ Y □ N	ΟΥΟΝ	□ Y □ N	□ Y □ N	□ Y □ N
Chills		Ο Υ Ο Ν		Ο Υ Ο Ν			
Cough	ΠΥΠΝ	□ Y □ N	Υ Ν	□ Y □ N	□ Y □ N	Υ Ν	ΟΥΟΝ
Sore throat		□ Y □ N			ΠΥΠΝ	□ Y □ N	ΠΥΠΝ
Shortness of breath		□ Y □ N		ΟΥΟΝ	ΠΥΠΝ	ΟΥΟΝ	ΠΥΠΝ
Myalgia/body pains	ΟΥΟΝ	Υ Ν	ΟΥΟΝ	ΟΥΟΝ	Υ Ν	□ Y □ N	ΟΥΟΝ
Diarrhoea		ΠΥΠΝ		ΟΥΟΝ	□ Y □ N	Ο Υ Ο Ν	ΟΥΟΝ

PDF version at: http://www.nicd.ac.za/diseases-a-z-index/novel-coronavirus-infection/



Complete for each contact of confirmed case.

Use electronic database if possible.



Details of <u>health official</u> completing form	Today's date	DD/MM/YYYY
Surname	Name	
Role	Facility name	
Email	Telephone	
address	number(s)	

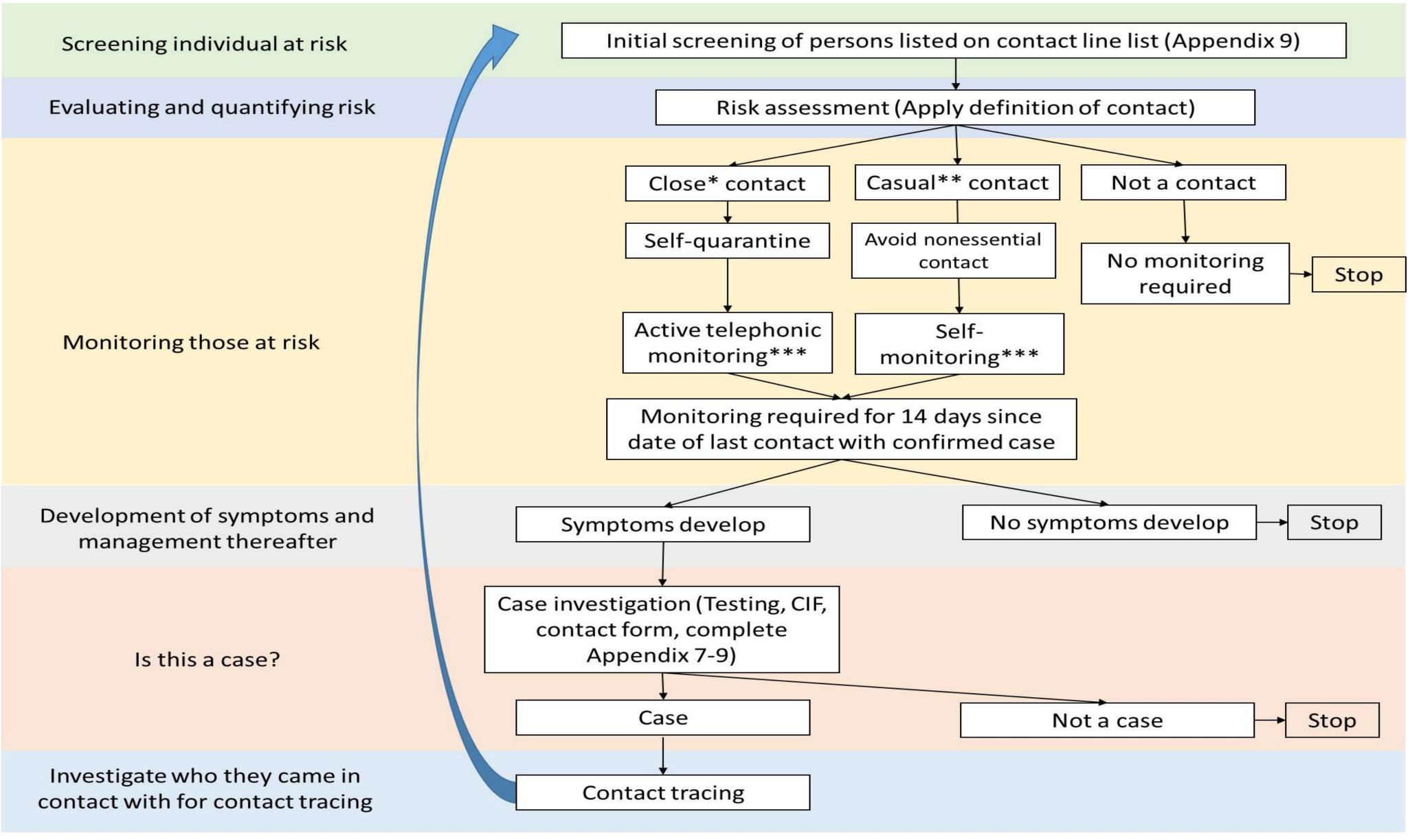
Instructions for completion: Mark "Y" if symptom present and "N" if not. If any symptoms are present collect, contact immediately and make immediate arrangements for the collection of a combined nasopharyngeal and

oropharyngeal swab. Refer to 2019-nCOV Quick Guide on the NICD website for additional details.

Management of close contacts who develop symptoms

- Should a contact develop symptoms, both the provincial CDCC and NICD call centre team should be informed
- Arrangements will be made by the provincial CDCC with assistance from NICD to visit the patient in their home on the same day to collect a specimen and to complete the required documentation.
 - Appropriate PPE should be used (e.g., gowns, gloves, NIOSH-certified disposable N95 respirator, eye protection) during home visits.
 - If a healthcare worker is not available, the patient will be requested to visit their nearest healthcare facility to have a specimen collected.
- The CDCC should inform the healthcare facility of the incoming patient in order for the healthcare facility to use appropriate infection prevention and control (IPC) measures.

Contact tracing summary



^{*} Close contact: A person having had face-to-face contact (<2 metres) or was in a closed environment with a 2019-nCoV case; this includes, amongst others, all persons living in the same household as a 2019-nCoV case and, people working closely in the same environment as a case. A healthcare worker or other person providing direct care for a 2019-nCoV case, while not wearing recommended personal protective equipment or PPE (e.g., gowns, gloves, NIOSH-certified disposable N95 respirator, eye protection). A contact in an aircraft sitting within two seats (in any direction) of the 2019-nCoV case, travel companions or persons providing care, and crew members serving in the section of the aircraft where the index case was seated. ** Casual contact: Anyone not meeting the definition for a close contact but with possible exposure. ***Monitoring methods: Active-telephonic monitoring: NICD call centre will phone person who is home-quarantined each day for a symptom report; Self-monitoring: person to consult healthcare practitioner in the event of symptom development.

Laboratory diagnostics

Who should be tested?

- Only patients under investigation (PUI) for 2019-nCoV should be tested
- Please discuss plans to collect samples with doctor on call before collecting sample: NICD hotline – 082 883 **9920**
- Rapid collection, transport and testing of appropriate specimens from PUI is a priority
- the clinical and epidemiological data strongly suggest 2019-nCoV infection





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Patients should be managed as potentially infected when



Specimen Collection

- Lower respiratory tract samples are preferred.
- Respiratory samples are the primary method if diagnosis.
- Respiratory samples include:
 - Combined nasopharyngeal and oropharyngeal swab (placed in the same tube) in ambulatory patients and
 - sputum (if produced)
 - Tracheal aspirate or Broncho alveolar lavage in patients with more severe respiratory disease.
- Serum for serological testing acute and convalescent samples may be submitted in addition to respiratory samples.
- Use universal/viral transport medium for swabs if available and if not dry swabs; sterile container for sputum and aspirates; clotted blood container for serum

preferred. method if diagnosis.



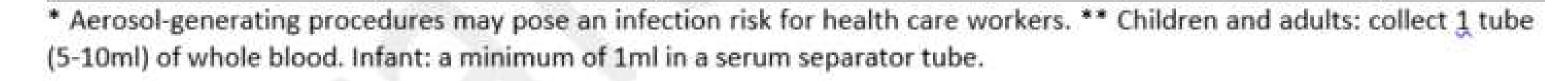


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Table 1. Type of specimens that can be collected for 2019-nCoV diagnostics and the transport requirements of these specimens

Specimen type	Collection materials	Storage and transportation	Dangerous goods shipping category	Comment
FOR SYMPTOMATIC PAT	ENTS:			
Sputum*	Deep cough sputum in sterile leak proof container	Refrigerate and ship at 2-8 °C up to 48 hrs, if >48 hrs freeze at -70°C and ship on dry ice	Biological substance, Category B	The preferred sample but need to ensure the material is from the lower respiratory tract
Bronchoalveolar lavage*	2-3 ml in sterile leak proof container	Refrigerate and ship at 2-8 °C up to 48 hrs, if >48 hrs freeze at -70°C and ship on dry ice	As above	There may be some dilution of virus but still a worthwhile specimen
(Endo)tracheal or nasopharyngeal aspirate*	2-3 ml in sterile leak proof container	Refrigerate and ship at 2-8 °C up to 48 hrs, if >48 hrs freeze at -70°C and ship on dry ice	As above	
Nasopharyngeal and oropharyngeal swab	Dacron or nylon flocked swab in Universal Transport Medium (UTM) in a sterile leak proof container	Refrigerate at 2-8 °C up to 5 days, if >5 days freeze at -70°C and ship on dry ice	As above	Nasopharyngeal and oropharyngeal swabs should be placed in the same tube to increase the viral load
Serum	Serum separator tube**	Store upright for at least 30 minutes after collection. a Refrigerate and ship at 2-8 °C within 5 days	As above	Collect paired samples: Acute – first week of illness Convalescent – 2-3 weeks later
Lung tissue from biopsy or autopsy	Sterile container with saline	Refrigerate and ship at 2-8 °C up to 24 hrs, if >24 hrs freeze at -70°C and ship on dry ice		







Collection of naso/oropharyngeal swabs for detection of respiratory viruses

COLLECTION OF NASO/OROPHARYNGEAL SWABS FOR DETECTION OF RESPIRATORY VIRUSES:

Respiratory viruses are best isolated from material that contains infected cells and secretions. Therefore, swabs should aim to brush cells and secretions off the mucous membranes of the upper respiratory tract. Good specimen quality (ie. containing sufficient cells and secretions), appropriate packaging and transport (i.e. to keep virus viable/detectable) is essential Please discuss plans to collect samples with doctor on call before collecting sample at NICD hotline - 0828839920

Step 1: Equipment and materials

- 1. Specimen submission form and case investigation form
- 2. Nasopharyngeal (NP) and oropharyngeal (OP) flocked swab
- 3. Tube containing universal transport medium (UTM)
- Tongue depressor
- 5. Gloves
- 6. N95 mask (fit tested)
- 7. Biohazard bag for disposal of non-sharp materials
- 8. Tissue for patient to wipe nose after sample collection
- 9. Cooler box and cooled ice packs
- 10. Ziploc plastic specimen bag

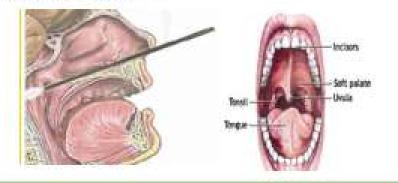
Step 2: Record keeping

- 1. Complete the specimen submission form and case investigation form (available on NICD website)
- Place the specimen submission form into a ziplock bag 3. Label the tube of universal transport media (UTM) with the patient's name and date of birth

Step 3: Collection of nasopharyngeal swab (NPS)

- 1. Don a pair of gloves, and an N95 respirator, making sure the respirator has a good fit. Open a sterile flocked swab at the plastic shaft
- 2. Ask the patient to tilt his/her head back. Estimate the distance from the patient's nose to the ear: This is how far the swab should be inserted
- Gently insert swab into the nostril and back (not upwards) to the nasopharynx until a slight resistance is met
- Rotate swab 2-3 times and hold in place for 2-3 seconds
- 5. If resistance is met remove and try another nostril
- 6. Slowly withdraw swab and without touching it, put it into a UTM
- 7. Break plastic shaft at the break point line and close the tube

Diagram: How to collect a nasopharyngeal swab (left) and oropharyngeal swab (right)



Step 4: Collection of oropharyngeal swab (OPS)

- 1. Keeping the same pair of gloves on, and holding the UTM with the nasopharyngeal swab in, take a second flocked swab and open it at the plastic shaft
- 2. Ask the patient to tilt their head back and open mouth wide
- 3. Hold the tongue down with a tongue depressor
- 4. Have the patient say "aahh" to elevate the uvula
- 5. Swab each tonsil first, then the posterior pharynx in a "figure 8" movement
- 6. Avoid swabbing the soft palate and do not touch the tongue with the swab tip as this procedure can induce the gag reflex.
- Place the swab into the same UTM tube with the NPS already in and break off the shaft at the break point line
- 8. Tightly close the tube
- 9. Place the closed tube with two swabs in the Ziploc
- 10. Remove gloves and N95 mask
- 11. Wash hands with soap and water

Step 5: Transport of specimens

- Ensure the cooler box and ice packs stay at 2-8°C
- 2. Transport to CRDM, NICD on same day as collection
- 3. Mark: Suspected Novel coronavirus, CRDM NHLS/NICD, Centre for Respiratory Disease and Meningitis (CRDM)

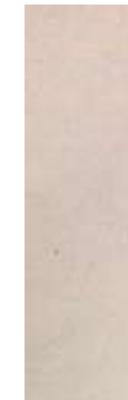
Lower North Wing, SAVP building 1 Modderfontein Rd, Sandringham, Johannesburg, 2131

- NHLS laboratories use usual overnight regional couries service
- 5. Private laboratories/clinics to organise shipment using existing systems, or contact CRDM for assistance if not available

Step 6: Contact details for additional assistance

Sample collection	Q -	
Sibongile Walaza	sibongilew@nicd.ac.za	011-386-6410/
		083-657-4741
Sample transport		
Linda de Gouveia	lindad@nicd.ac.za	011-555-0327
Amelia Buys	ameliab@nicd.ac.za	011-386-6373
Cardia Fourie	cardiaf@nicd.ac.za	011-386-6373
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http://www.nicd.ac.za/wp-content/uploads/2020/02/2019-nCov-Quick-reference-v3-03.02.2020-final.pdf



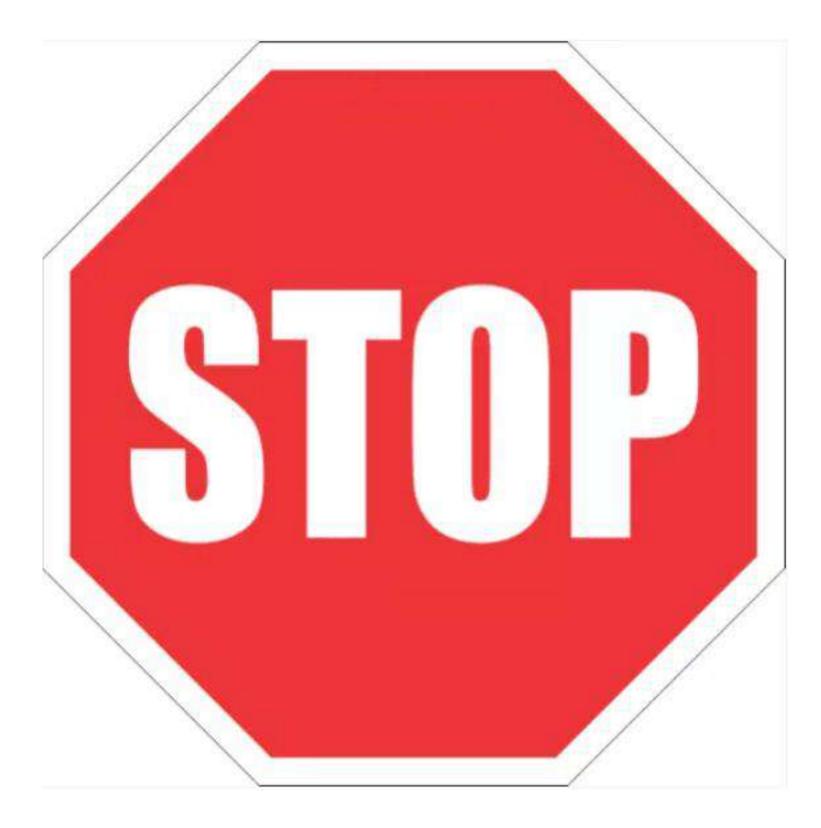
Swabs Important Information

- Clearly mark each specimen (e.g. Left Nasal Swab Tight Nasal Swab)
- If you send multiple swabs unmarked the lab has no idea where they come from
- You must identify which facility the swab comes from
- Clinicians name and contact details are important









DO NOT send any specimen to NICD without prior discussion and notification





Hand hygiene before and after any interaction with the patient





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Equipment and materials

- 1. Specimen submission form and case investigation form. 2. Nasopharyngeal (NP) and oropharyngeal (OP) flocked swab.
- 3. Tube containing universal transport medium (UTM).
- 4. Tongue depressor.
- 5. Gloves.
- 6. N95 mask (fit tested).
- 7. Biohazard bag for disposal of non-sharp materials.
- 8. Tissue for patient to wipe nose after sample collection.
- 9. Cooler box and cooled ice packs.
- 10. Ziploc plastic specimen bag.







Step 1: Report the PUI

- 1. Report the PUI to the NICD to allow a risk assessment to be carried out and guide laboratory testing
- 2. Contact the NICD Hotline +27 82 883 9920
- 3. The test will be free of charge for patients meeting the case definitions above







Step 2: Record keeping

- on NICD website). http://www.nicd.ac.za/diseases-a-z-index/novelcoronavirus-infection/
- 2. Place the specimen submission form into a ziplock bag.
- 3. Label the tube of universal transport media (UTM) with the patient's name and date of birth.



1. Complete the specimen submission form and case investigation form (available)





Complete the correct forms

- form has to be completed and submitted together with the specimens
- Always check on the NICD website that you have the current version of the forms http://www.nicd.ac.za/diseases-a-z-index/novel-coronavirus-infection/

CRDM episode no:	CRDM lab no:	Trak no:	Date received:	
		Respiratory Diseases an Specimen Submission fo		CRDM episode no:
Patient Information		Submitter Information (contact person for results)	
Identifier or Hospital no		Sumame		CRDM PCR Diagnostic
Surname		First name		
First name		Laboratory		Test name:
Age/Date of birth		City, Country		
Gender	-	Contact number (country and	*) +()	Respiratory panel
Facility/Hospital		Email address		inclusion of particular
Specimen Details		1. 	-	
Specimen collection date:	dd-mm-yygy			
	Combined NP/OP swab	Nasopharyngeal	(NP) aspirate Nasal swab	
Specimen type:	Nasopharyngeal (NP) swal	b Bronchoalveolar	lavage (BAL) Sputum	
	Oropharyngeal (OP) swab			
	Tracheal aspirate (TA)	Blood culture	☐ Serum	
	Whole blood	Other, specify:		
Laboratory Test Datails (als	ase consult with CRDM if testing other t	the second s	and in comments	
A DECK AND A DECK	p B streptococcus Hospital-ac	y-acquired pneumonia (bacter quired pneumonia (bacter Date of symptom onse	ria)*	
Clinical diagnosis: Acute Dipht Pertu	theria 🗌 Influenza-li		espiratory tract infection espiratory tract infection pecify:	
	(≥38°C) Sore Throat ness of breath Vomiting ea Other, specify:	Diarrhoea Paroxysm		Community-acquired p
and the second	aner an an anna an an an an an an an an an a	ase Diabetes [seven or well and the second of the second o	11 S.1 5.1
Underlying Risk Factors:				Hospital-acquired pnei
Underlying Risk Factors:	Heart Disease Other, specify:		Unknown None	Hospital-acquired pneu
Hospitalisation: Outpar	tient Out ent not admitted ICU ent admitted to ICU	tcome: Still hospit	alised	Atypical pneumonia
Hospitalisation: Outpa Inpatie Unpatie	tient Out ent not admitted ICU ent admitted to ICU	tcome: Still hospit	alised	Atypical pneumonia
Hospitalisation: Outpar Inpatie Inpatie Unkno Exposure History	tient Out ent— not admitted ICU ent— admitted to ICU wn	tcome: Still hospit	alised	
Hospitalisation: Outpatie Inpatie Unpatie Unkno Exposure History Did the patient travel in the	tient ent — not admitted ICU ent — admitted to ICU wwn 2 14 days prior to symptom onset?	tcome: Still hospit	alised	Atypical pneumonia
Hospitalisation: Outpatie Inpatie Unpatie Unkno Exposure History Did the patient travel in the Area/Country travelle	tient out admitted ICU ent — not admitted ICU ent — admitted to ICU own 2014 days prior to symptom onset? ed to: Date of travel	tcome: Still hospit	alised D Unknown Date of travel <u>from</u> this area	Atypical pneumonia
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Hospitalisation: Outpatie Inpatie Unpatie Unknot Exposure History Did the patient travel in the Area/Country travelle 1.	tient ent not admitted ICU ent admitted to ICU wwn 2 14 days prior to symptom onset? ed to: Date of travel	tcome: Still hospit Survived Died Unknown	alised D Unknown Date of travel <u>from</u> this area dd-mm-vyyy	Atypical pneumonia Neonatal sepsis
Hospitalisation: Outpar Inpatie Inpatie Unkno Exposure History Did the patient travel in the Area/Country travelle 1. 2. Did the patient have anima	tient out admitted ICU ent — not admitted ICU ent — admitted to ICU own 2014 days prior to symptom onset? ed to: Date of travel	tcome: Still hospit Survived Died Unknown	alised D Unknown Date of travel <u>from</u> this area	Atypical pneumonia Neonatal sepsis Bacterial meningitis
Hospitalisation: Outpatie Inpatie Unpatie Unknot Exposure History Did the patient travel in the Area/Country travelle 1. 2. Did the patient have anima Area	tient ent— not admitted ICU ent— admitted to ICU wwn 2 14 days prior to symptom onset? ed to: Date of travel dd-mm I contact in the 14 days prior to sym nimal type	tcome: Still hospit Survived Died Unknown Yes No to this area	alised Date of travel <u>from</u> this area dd-mm-vyvy No Unknown	Atypical pneumonia Neonatal sepsis
Hospitalisation: Outpar Inpatie Inpatie Unkno Exposure History Did the patient travel in the Area/Country travelle 1. 2. Did the patient have anima Area	tient ent— not admitted ICU ent— admitted to ICU wwn e 14 days prior to symptom onset? ed to: Date of travel dd-mm l contact in the 14 days prior to sym nimal type Poultry (eg. chickens, ostrich, ducks)	tcome: Still hospit Survived Died Unknown Yes No to this area	alised Date of travel <u>from</u> this area dd-mm-vyvy No Unknown	Atypical pneumonia Neonatal sepsis Bacterial meningitis

• For each person under investigation (PUI) a laboratory specimen submission form and a person under investigation (PUI)

CRDM lab no:

Trak no:

Date received:

fest Panels:

t name:	Pathogens:
piratory panel	Viruses:
	Influenza A, influenza B, influenza C, rhinovirus, human coronavirus, parainfluenza virus, human bocavirus, human metapneumovirus, enterovirus, adenovirus, parechovirus, respiratory syncytial virus (RSV)
	Bacteria:
	Mycoplasma pneumoniae, Chlamydia pneumoniae, Haemophilus influenzae, Haemophilus influenzae type B, Staphylococcus aureus, Klebsiella pneumoniae, Legionella spp., Salmonella, Bordetella pertussis, Moraxella catarrhalis Fungi:
	Pneumocystis jiroveci
nmunity-acquired pneumonia	Streptococcus pneumoniae, Staphylococcus aureus, Haemophilus influenzae, Moraxella catarrhalis
pital-acquired pneumonia	Klebsiella pneumoniae, Pseudomonas aeruginosa
pical pneumonia	Mycoplasma pneumoniae, Chlamydia pneumoniae, Legionella spp.
onatal sepsis	Group B streptococcus, Listeria monocytogenes, Staphylococcus aureus, Chlamydia trachomatics, Ureaplasma urealyticum/parvum, cytomegalovirus
terial meningitis	Streptococcus pneumoniae, Neisseria meningitidis, Haemophilus influenzae
al meningitis	Adenovirus, cytomegalovirus, epstein barr virus, herpes simplex virus 1, herpes sim- plex virus 2, varicella zoster virus, enterovirus, parechovirus, human herpesvirus 6, human herpesvirus 7, parvovirus B19, mumps virus

Person under investigation form (CIF)





MATIONAL I		A CONTRACTOR OF		health	001			Y DISEASES AND MENIN JBLIC HEALTH SURVEILL
COMMUNIC Division in the National		A CONTRACTOR OF		Department: Health REPUBLIC OF SOUTH AFR				AND RESP
		and the second		ALF OULD OF BOOTH AFR	over 11		naluse	
Patient under inv	vestigati	on (PUI) fo	rm: Re	quest for 2019-nCo	V Testine			10:
Furthe	rmore, the	e completed	case inv	e investigation forms sho estigation form must be Fax: (+27)11 8829979 H	scanned a	nd emailed to <u>ncov@ni</u>	cd.ac.za as	detailed below
Today's DD/MM	VI/YYYY	Form comp	leted by	(Name, Surname):		Contac	t number(s):
ls this a: New cl	linical que	ry□		act of a known case, e case details:	Known ca	se first name:		
Contac	t of a kno	wn case 🗆	provide	e case de talis.	Known ca	se surname:	96 	
					Known ca	se DOB:	DE)/MM/YYYY
Detected at point of	entry? Y	/ NU Unk	n	If yes, date: DD/MM/Y	YYY	Please specify the	point of en	ntry:
		DATIEN	T DETAI				DOCTOR	'S DETAILS
Patient hospital num	ber (if ava		DETAI			First name:	DOCTOR	SDETAILS
	oci fii ava	ii				10	3	
First name:			Surname			Surname: Facility name:		
DOB: DD/MIN	W/YYYY		Sex:	Male 🗆 Female 🗆			·	
Residency: SA resi	ident 🗆 N	Ion-SA reside	ent 🗆 (s	pecify)	7/	Contact number/s:	<u>108 70</u>	<u></u>
Current residential A	ddress1:	<u>24 77 -</u>	<u>88 - 05</u>	AR AL AL AL	<u> 18 - 18 -</u> 35		<u>410 80 - 70</u>	<u></u>
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lease include alternative n			_		اد <u>مین</u> منبع	Email address:	<u>ansa 10</u>	<u> </u>
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Please indicate occup tick any if apply):	pation	with		Health laborator worker	γ 🗆			
		animals Healthcare						
		worker		Facility name:				
		Other		Specify				
			NEXT O	F KIN CONTACT DETAILS	5 (alternati	ve contact details)		
irst name:				Su	imame:	17		
lelationship to the pa	atient: _			584 S.	ntact num		1 - X	40 - M 40 - M 1
Date of symptom	(000)			CLINICAL PRESENTAT				
onset:	-	MM/YYYY			urrent cons	ultation/admission:	DD/MM/YY	YY.
	Fever (a			Sore throat		Myalgia/body pains		
Symptoms (tick all	Cough	offever		Shortness of breath Nausea/vomiting		General weakness Irritability/confusior		
that apply):	Chills			Diarrhoea		Other		(specify if
	CHIIS					ouler		other)
61141-00-00	S Description			DIAGN				
Did the patient hav		n meninken s inne	revidenc	e of pheumonia				
Were chest X-rays (Y N	If yes, CXR Findings: _		
Did the patient hav distress syndrome (or radiologica	levideno	e of acute respiratory	Y N			
 Does the patient ha illness? 	ave anothe	er diagnosis∕e	tiology f	for their respiratory	Y 🗌 (speci	ý)	N	Unknown
Page 1 of 2			Please	refer to <u>www.ncd.ac.m</u> for most rec	ent version of th	is document before use.		Version 2 31 January 2

Final version 2_31JANUARY 2020

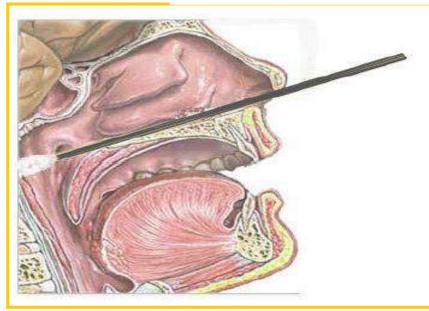
Step 3: Collection of nasopharyngeal swab (NPS)

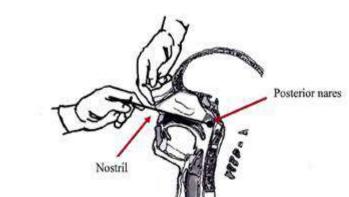
good fit. Open a sterile flocked swab at the plastic shaft

2. Ask the patient to tilt his/her head back. Estimate the distance from the patient's nose to the ear: This is how far the swab should be inserted

- 3. Gently insert swab into the nostril and back (not upwards) to the nasopharynx until a slight resistance is met
- 4. Rotate swab 2-3 times and hold in place for 2-3 seconds
- 5. If resistance is met remove and try another nostril
- 6. Slowly withdraw swab and without touching it, put it into a UTM
- 7. Break plastic shaft at the break point line and close the tube

- 1. Don a pair of gloves, and an N95 respirator, making sure the respirator has a

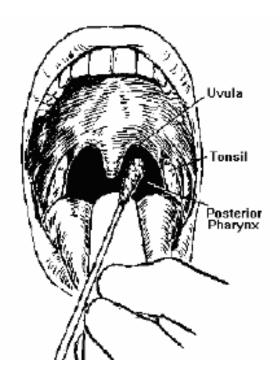




Step 4: Collection of oropharyngeal swab (OPS)

- flocked swab and open it at the plastic shaft
- Ask the patient to tilt their head back and open mouth wide 2.
- Hold the tongue down with a tongue depressor 3.
- 4. Have the patient say "aahh" to elevate the uvula
- Swab each tonsil first, then the posterior pharynx in a "figure 8" movement 5.
- Avoid swabbing the soft palate and do not touch the tongue with the swab tip as this procedure can 6. induce the gag reflex.
- line
- Tightly close the tube 8.
- Place the closed tube with two swabs in the Ziploc 9.
- 10. Remove gloves and N95 mask
- 11. Wash hands with soap and water

1. Keeping the same pair of gloves on, and holding the UTM with the nasopharyngeal swab in, take a second



7. Place the swab into the same UTM tube with the NPS already in and break off the shaft at the break point







Step 5: Transport of specimens

- 1. Ensure the cooler box and ice packs stay at 2-8 degrees Centigrade.
- 2. Transport to CRDM, NICD on same day as collection.
- 3. Mark: Suspected Novel coronavirus, CRDM NHLS/NICD, Centre for Respiratory Disease and Meningitis (CRDM) Lower North Wing, SAVP building 1 Modderfontein Rd, Sandringham, Johannesburg, 2131.
- 4. NHLS laboratories use usual overnight regional courier service.
- 5. Private laboratories/clinics to organise shipment using existing systems, or contact CRDM for assistance if not available.





Packaging of specimens for transfer to NICD

- The principle of triple layer packaging should be followed (Figure 1).
- UN/WHO approved shipping containers for hazardous specimens are commercially available, e.g. SAF-T-PAK[®] (www.saftpak.com) or PATHOPAK[®] (<u>www.intelsius.com</u>).

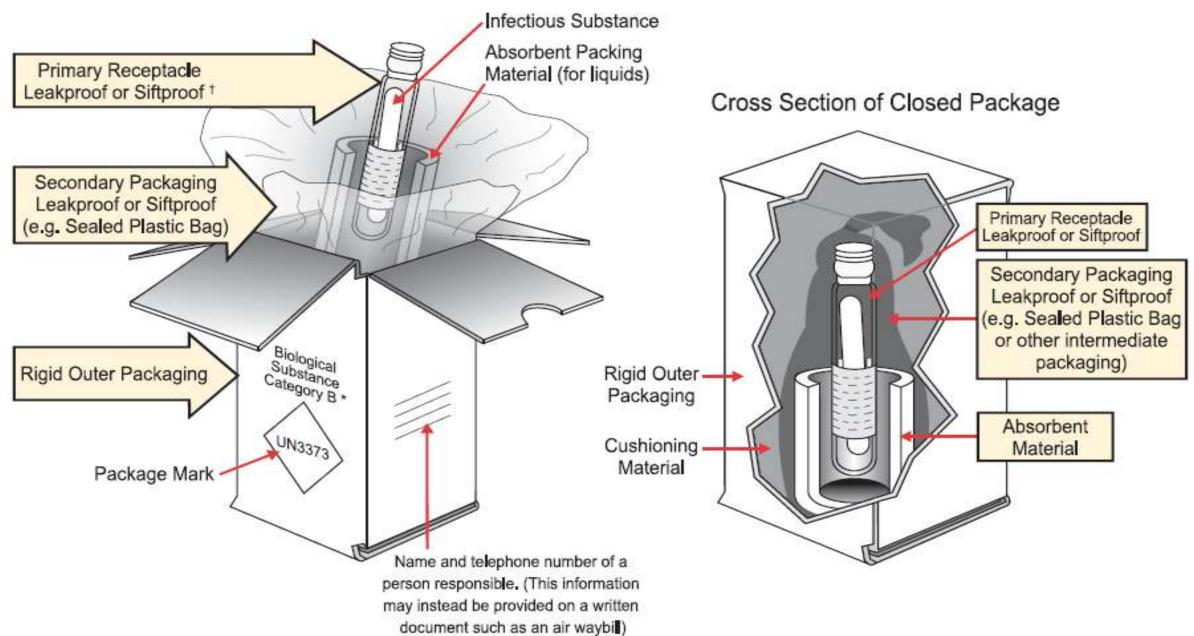


Figure 1. Example of the triple packaging system for the packing and labelling of Category B substances.



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Step 6: Contact details for additional assistance

- Sample collection
 - Sibongile Walaza sibongilew@nicd.ac.za 011-386-6410
- Sample transport
 - Linda de Gouveia lindad@nicd.ac.za 011-555-0327
 - Amelia Buys ameliab@nicd.ac.za 011-386-6373
 - Cardia Fourie cardiaf@nicd.ac.za 011-386-6373
- http://www.nicd.ac.za/wp-content/uploads/2020/02/2019-nCov-Quickreference-v3-03.02.2020-final.pdf



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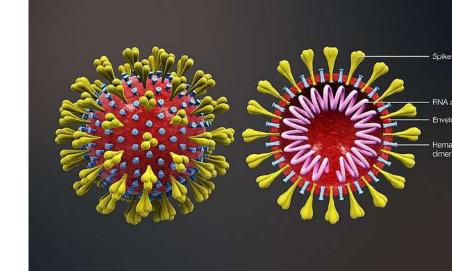


Laboratory diagnostic assays

- Real-time reverse-transcription polymerase chain reaction (rRT-PCR) - amplification and detection of unique 2019-nCoV viral nucleic acid sequences
- TAT 24 hours
- Positive specimens characterised by viral culture and whole genome sequencing

Detection of 2019 novel coronavirus (2019-nCoV real-time RT-PCR

Victor M Corman¹, Olfert Landt², Marco Kaiser², Richard Molenkamp³, Adam Meijer⁴, Daniel KW Chu⁵, Tobias Bleicker¹, Sebastian Brünink¹, Julia Schneider¹, Marie Luisa Schmidt¹, Daphne GJC Mulders³, Bart L Haagmans³, Bas van der Veer⁴, Sharon van den Brink⁴, Lisa Wijsman⁴, Gabriel Goderski⁴, Jean-Louis Romette⁶, Joanna Ellis⁷, Maria Zambon⁷, Malik Peiris⁵, Herman Goossens⁸, Chantal Reusken⁴, Marion PG Koopmans³, Christian Drosten¹



Eurosurveillance Jan 2020





Interpretation of rRT-PCR results

- Negative result does not rule out possibility of infection
- Factors that could lead to a false –negative result:
 - Poor specimen quality
 - Specimen was collected late or very early in the illness Specimen was not handled and shipped appropriately, (eg.
 - the cold chain)
 - Technical reasons inherent in the test, e.g virus mutation

suspicion for 2019-nCoV infection, especially when only upper including lower respiratory samples should be collected and tested.





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If negative results are obtained from patients with a high index of respiratory tract samples were collected, additional specimens,

Infection prevention and control

Principles of disease transmission



Direct contact

- Touching an ill persons or a contaminated surface
- E.g. agents of diarrhoea, lacksquareskin infections, common cold, ebola virus

Control

Gloves, +/- gowns, masks, visors (to prevent mucous membrane splashes, contamination of clothing)



Droplet transmission

- Inhaling droplets (up to 1/4mm in diameter)
- Persons within 2m radius are at risk. On aircraft, 2 rows behind and in front
- E.g. agents of bacterial pneumonia, Neisseria meningitidis

Control

Gloves, surgical masks, +/- gowns, masks, visors (to prevent mucous membrane splashes, contamination of clothing)

Airborne transmission

- Inhaling droplets nurclei (10-20um in diameter)
- Persons breathing the same air
- E.g. influenza, measles, chickenpox,

Control

Gloves, N95 masks, +/- gowns, masks, visors (to prevent mucous membrane splashes, contamination of clothing)



Vector transmission

- Contact with vector
- E.g. malaria, dengue, Zika,

Control

- Prevent/eliminate exposure to vector
- Chemoprophylaxis if possible



Principles of disease transmission

Coronavirus



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- E.g. agents of diarrhoea, lacksquareskin infections, common cold, ebola virus

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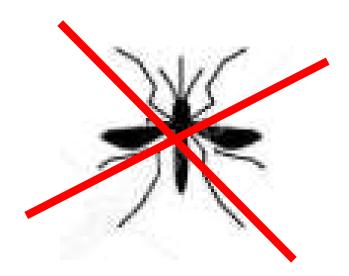


Vector transmission

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Control

- Prevent/eliminate exposure to vector
- Chemoprophylaxis if possible



IPC strategies to address suspected nCoV infection

- Ensure triage, early recognition and source control (early isolation of persons with suspected nCoV infection)
- Apply standard precautions for all patients
- Implement empiric additional precautions for suspected cases (droplet, contact and airborne where applicable)

- Implement administrative controls (IPC committee, checklist, assign responsibility for opening windows and triaging)
- Use environmental controls (open windows, UV light, ensure airflow direction protects HCW)
- Use engineering controls (ensure air-conditioning is working, Uvlight germicidal irradiation unit is functional)

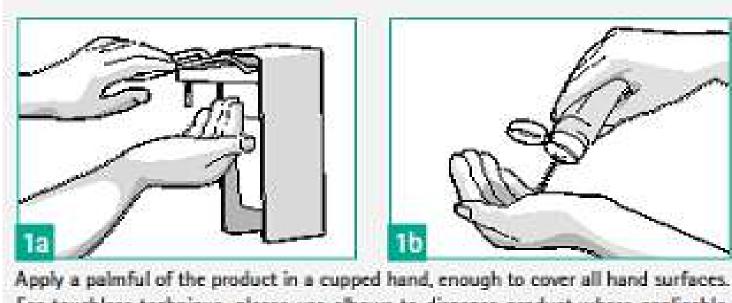
In all facilities....

- casualty / hospital
- Put a sign up asking for persons with a travel history to China in last 14 days to identify themselves to staff
- Provide surgical masks to persons who sneeze, cough etc
- See persons who have symptoms first
- Encourage hand hygiene amongst patients and HCW

 Implement screening for COUGH, respiratory symptoms and TRAVEL HISTORY at entrance to the facility / clinic /

In all facilities.....

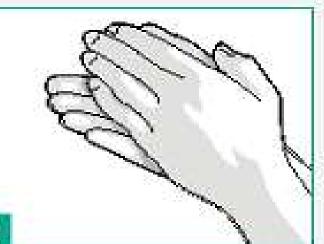
- Ensure hand hygiene for HCW and patients is possible, and done!
- Provide soap, basins
- Use posters to show 5-movements of hand hygiene
- Provide hand sanitiser
- Use health promotion staff to demonstrate hand and cough hygiene





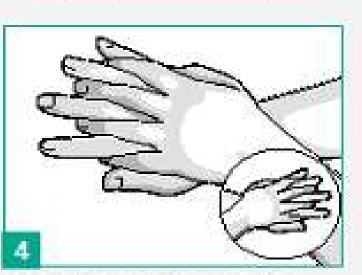


For touchless technique, please use elbows to dispense product where applicable.



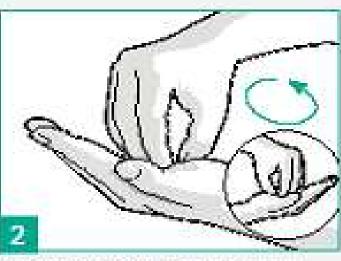
Rub hands palm to palm.

fingers interlocked.

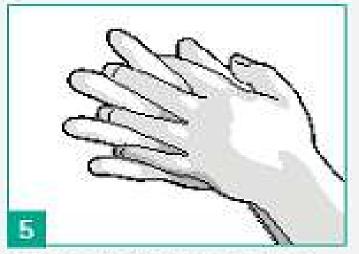


Right palm over left dorsum with interlaced fingers and vice versa.

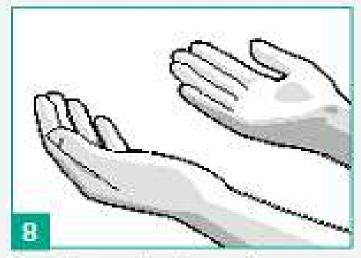
clasped in right palm and vice versa.



Rotational rubbing, backwards and forwards with clasped fingers of right hand in left palm and vice versa.



Palm to palm with fingers interlaced.



Once dry, your hands are safe.

When caring for someone with suspected nCoV-

- Implement contact and droplet precautions
 - Put in a well ventilated isolation room
 - Provide them with a mask
 - Implement contact and droplet precautions
 - Limit the number of staff who can enter the isolation room

- Implement contact and droplet precautions:
 - Surgical/medical mask
 - Disposable gown
 - Gloves
 - Eye protection
- Not required for droplet precautions
 - Boots, apron not required
 - Negative pressure respiratory isolation room not required.

When caring for someone with suspected nCoV-

- When taking a sputum specimen or nasopharyngeal swab use <u>airborne and</u> <u>contact</u> precautions are required
 - E.g. nasopharyngeal swabs, intubation, tracheal aspirate
- Use N95 respirator
- Use waterproof apron, boots
- Use a face-shield or goggles



Training in use of IPC DOFFING HAND HYGIENE PROTECTION PROTECTION EYE PROTECTION HAND HYGIENE Healthcare and Emergency Responder Organization Education through Simulation PENDED THROUGH & UNIVERSITY OF HEBRASHA PRODUCTS OF EXCELLENCE ARANT

- Ensure staff are trained and familiar with
 - Triage
 - Handwashing
 - Screening
 - Case definitions
 - Use of PPE



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Infection prevention and control during health care when novel coronavirus (nCoV) infection is suspected Interim guidance World Health Organization 25 January 2020

WHO/2019-nCoV/IPC/v2020.2

Introduction

This is the first edition of guidance on infection prevention and control (IPC) strategies for use when infection with a novel coronavirus (2019-nCoV) is suspected. It has been adapted from WHO's Infection prevention and control during health care for probable or confirmed cases of Middle East respiratory syndrome coronavirus (MERS-CoV) infection,¹ based on current knowledge of the situation in China and other countries where cases were identified and experiences with severe acute respiratory syndrome (SARS)-CoV and MERS-CoV.2

WHO will update these recommendations as new information becomes available.

This guidance is intended for healthcare workers (HCWs), healthcare managers and IPC teams at the facility level but it is also relevant for the national and district/provincial level. Full guidelines are available from WHO.²

Principles of IPC strategies associated with health care for suspected nCoV infection

To achieve the highest level of effectiveness in the response to an 2019-nCoV outbreak using the strategies and practices recommended in this document, an IPC programme with a dedicated and trained team or at least an IPC focal point should be in place and supported by the national and facility senior management.3 In countries where IPC is limited or inexistent, it is critical to start by ensuring that at least

Ensuring triage, early recognition, and source 1. control

Clinical triage includes a system for assessing all patients at admission allowing early recognition of possible 2019-nCoV infection and immediate isolation of patients with suspected nCoV infection in an area separate from other patients (source control). To facilitate the early identification of cases of suspected nCoV infection, healthcare facilities should:

- suspicion;
- staff;
- coronavirus-(2019-ncov) and
- patients to alert HCWs.

The promotion of hand hygiene and respiratory hygiene are essential preventive measures.

2. Applying standard precautions for all patients

Standard precautions include hand and respiratory hygiene, the use of appropriate personal protective equipment (PPE) according to risk assessment, injection safety practices, safe waste management, proper linens, environmental cleaning and sterilization of natient-care equipment

encourage HCWs to have a high level of clinical

establish a well-equipped triage station at the entrance of health care facility, supported by trained

institute the use of screening questionnaires according to the updated case definition (https://www.who.int/publications-detail/globalsurveillance-for-human-infection-with-novel-

post signs in public areas reminding symptomatic

- If in doubt, refer to this WHO guideline
- It is **ESSENTIAL** to distribute this guideline to your facility staff and follow up on implementati on

Management of the deceased

- Confirm the diagnosis in deceased persons who are close contacts of nCoV cases.
 - NP swabs, bronchial washings can be taken post mortem
- Use contact and droplet precautions when handling the body
 - Airborne precautions not required as the deceased do not create airborne particles
- Environmental Health Practitioners



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Health

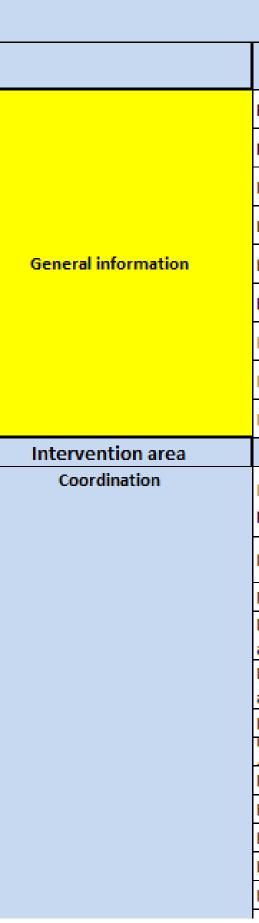
e informed following the assist with procedures

- Follow Appendix 12 of RSA guideline
 - Triple body bag, first two are transparent and sealed, and third is black and unsealed
 - A biohazard warning tab should be attached as per SOP



How can I know if my facility is ready?

- Use our facility readiness
 checklist
- Call your facility IPC committee
- Talk through the checklist
- Talk through a 'desktop simulation scenario'



Novel Coronavirus (nCoV) Readiness Checklist

	Total inFacilty	
Number of HCW employed / working at your facility		
Number of designated points of entry for ill patients		
Do you have isolation units in the Facilty if yes how many beds		
Private wards		
Private ward airborn precautions(-ve Pressure cubicles)		
Large cohort area identified Number of beds		
Isolation area in emergency department identified		
ICU Isolation cubicles number		
ICU isolation cubicles -ve pressure		
Indicators	Values	Yes/No
Is there a Facilty preparedness and response plan for events caused by respiratory pathogens?		No
Do you have a committee established in the Facilty to ensure all plans are in place		No
Do you maintain minutes of the meetings of this committee		No
Do you have and have you reviewed plans for implementation of surge capacity procedures		No No
Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities Do you have supplies of PPE for staff in front line areas		No
Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities		No No
Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities Do you have supplies of PPE for staff in front line areas		No No No
Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities Do you have supplies of PPE for staff in front line areas Do you have supplies of PPE for staff in Facility care areas in case of a positive or suspected		No No No No
Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities Do you have supplies of PPE for staff in front line areas Do you have supplies of PPE for staff in Front line areas Do you nave supplies of PPE for staff in Facility care areas in case of a positive or suspected Do you have contingency plans if the demand for PPE or other supplies exceeds supply.		No No No No No
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Facility self assessment

(SO	UTH AFRICAFacilty) Novel Coronavirus (nCoV) Checklist	Country Re	adiness
		Total inFacilty	
	Number of HCW employed / working at your facility		
	Number of designated points of entry for ill patients		
	Do you haver isolation units in the Facilty if yes how many beds		
	Private wards		
General information	Private ward airborn precautions(-ve Pressure cubicles)		
	Large cohort area identified Number of beds		
	Isolation area in emergency department identified		
	ICU Isolation cubicles number		
	ICU isolation cubicles -ve pressure		
Intervention area	Indicators	Values	Yes/No
	Is there a Facilty preparedness and response plan for events caused by respiratory pathogens?		No
			No No
	respiratory pathogens?		
	respiratory pathogens? Do you have a committee established in the Facilty to ensure all plans are in place Do you maintain minutes of the meetings of this committee Do you have and have you reviewed plans for implementation of surge capacity		No
	respiratory pathogens? Do you have a committee established in the Facilty to ensure all plans are in place Do you maintain minutes of the meetings of this committee Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with		No No
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	respiratory pathogens? Do you have a committee established in the Facility to ensure all plans are in place Do you maintain minutes of the meetings of this committee Do you have and have you reviewed plans for implementation of surge capacity procedures and crisis standards of care. Do you believe you have everything in place to identify and isolate patients with 2019-nCoV and inform key facility staff and public health authorities Do you have supplies of PPE for staff in front line areas Do you have supplies of PPE for staff in Facility care areas in case of a positive or suspected case Do you have contingency plans if the demand for PPE or other supplies exceeds supply. Have you designated an area for the isolation of patients who may be at risk for Corona virus		No No No No No No No



Hospital Name		
CEO Name		
CEO Contact details		
Cell		
Office		
email		
Fax		
Date of Report		
	Facility CEO / Manage	er Signature
Comments	Gaps	Resources



Patient and PUI* flow and actions required at each step

*PUI=person under investigation

Process Flow for detection and response to cases

Appendix 1 – process flow for detection and response to cases 1.1

DETECTION AND REPORTING OF SUSPECTED 2019-nCoV CASE

- The case definition must be strictly adhered to
- For any suspected case, isolate the patient in a suitable room/ unit for assessment, apply IPC measures, contact NICD Hotline to confirm if case definition is met and if sample collection is warranted.
- If so, collect specimen and complete accompanying documentation (Appendix 7).
- Guidelines for the collection and submission of specimens to NICD available on NICD website: <u>http://www.nicd.ac.za/diseases-a-z-index/novel-coronavirus-infection/</u> (see quick reference for healthcare workers) or appendix 5 and 6
- The facility IPC focal point, clinician or designated port health officer should complete the case investigation form and contact line list (Appendix 8, 9), forward the forms to the Provincial Communicable Disease Control and <u>ncov@nicd.ac.za</u>.
- All suspected cases who meet the case definition should be notified as Class 1 notifiable medical condition under "Respiratory Disease caused by a novel respiratory pathogen"

MEDICAL MANAGEMENT

For all cases irrespective of symptom severity, isolate the patient and apply infection precautions in accordance with site-specific standard operating procedures for this purpose. When the number of confirmed cases becomes too high, mild cases may be managed at home (selfisolation)



Contacts and details: Consultant on call for Infectious Diseases

According to site-specific protocol

> **NICD Hotline** 082-883-9920

National Health Operations Centre 012-395-9636/37

Contacts and details: see Appendix 14

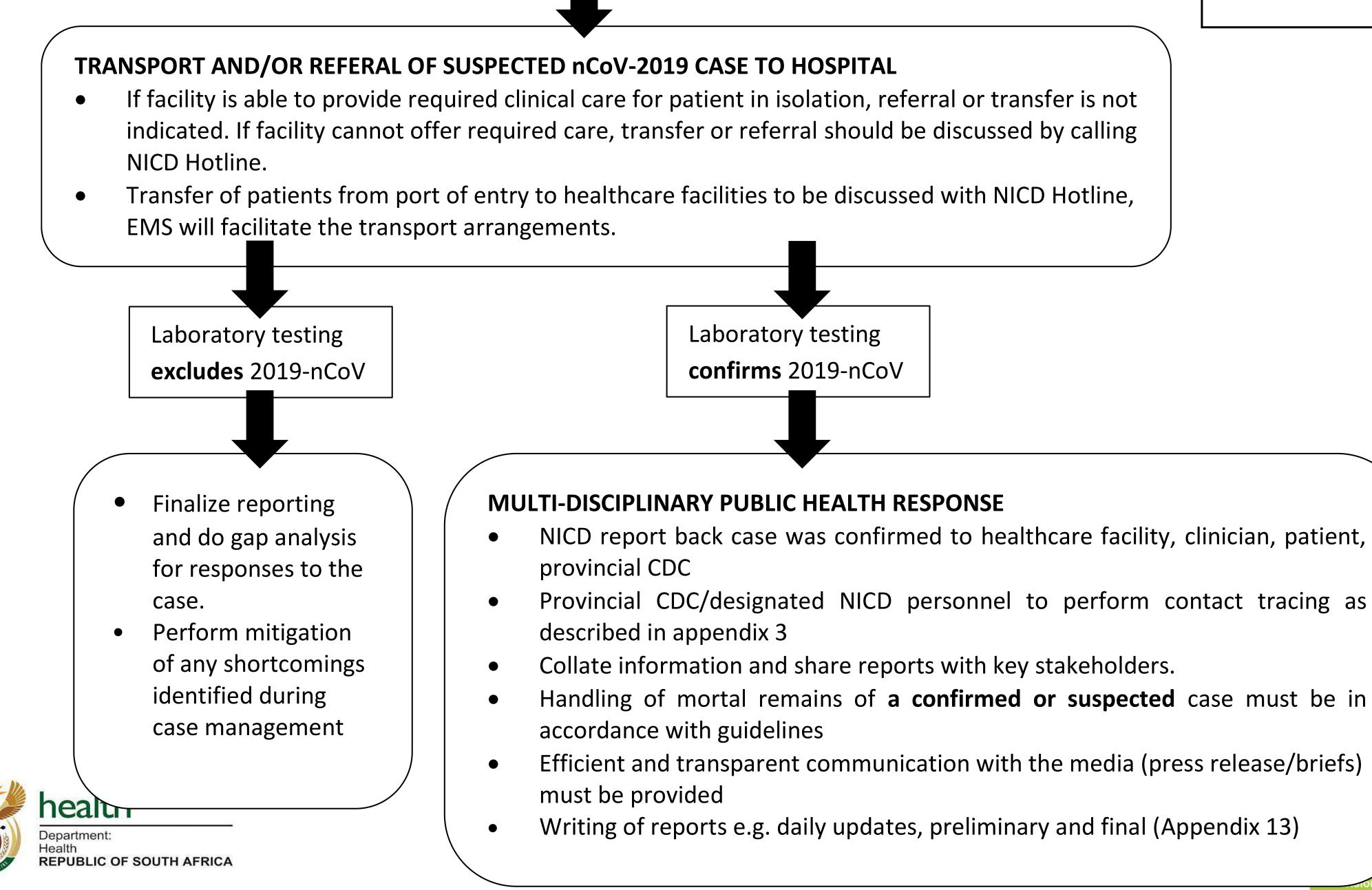
National and Provincial CDC

Provincial Port Health

EMS



Process Flow for detection and response to cases



Provincial CDC/designated NICD personnel to perform contact tracing as

Handling of mortal remains of a confirmed or suspected case must be in

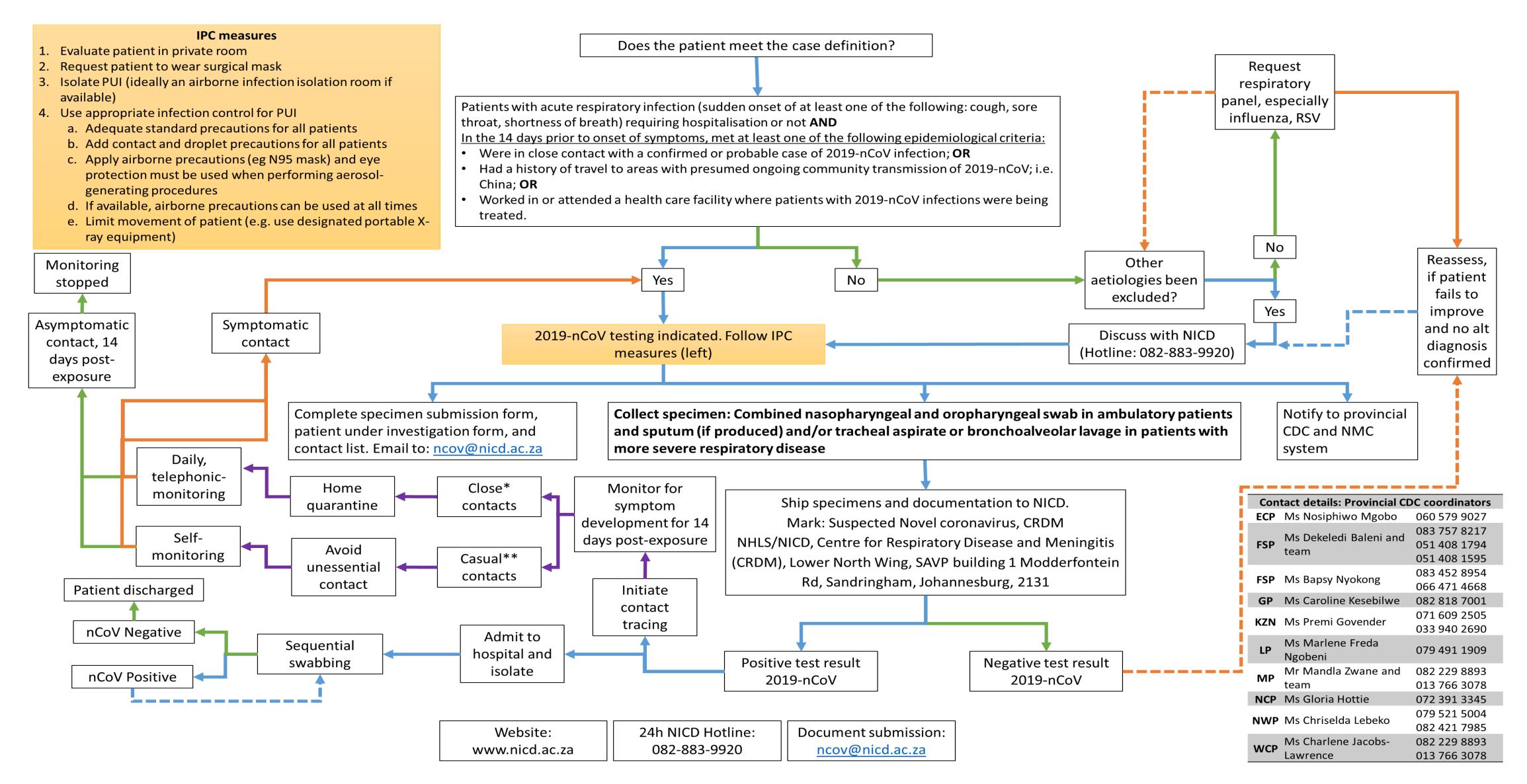
IONAL INSTITUTE FOR MMUNICABLE DISEASES

the National Health Laboratory Service



Initial diagnosis and management of suspected case (PUI), including infection control measures

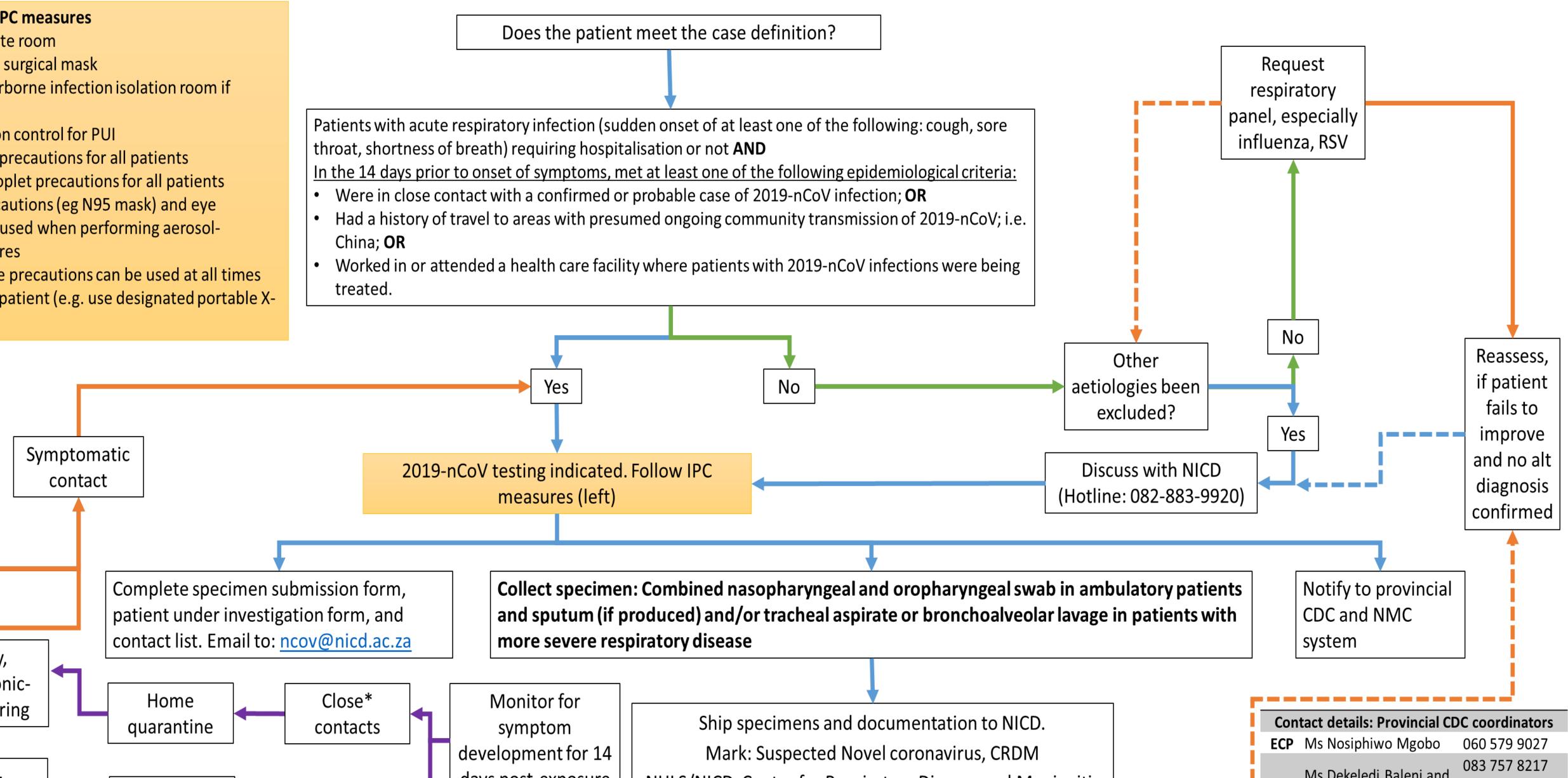
2019 novel coronavirus (2019-nCoV) process flow for use in healthcare facilities



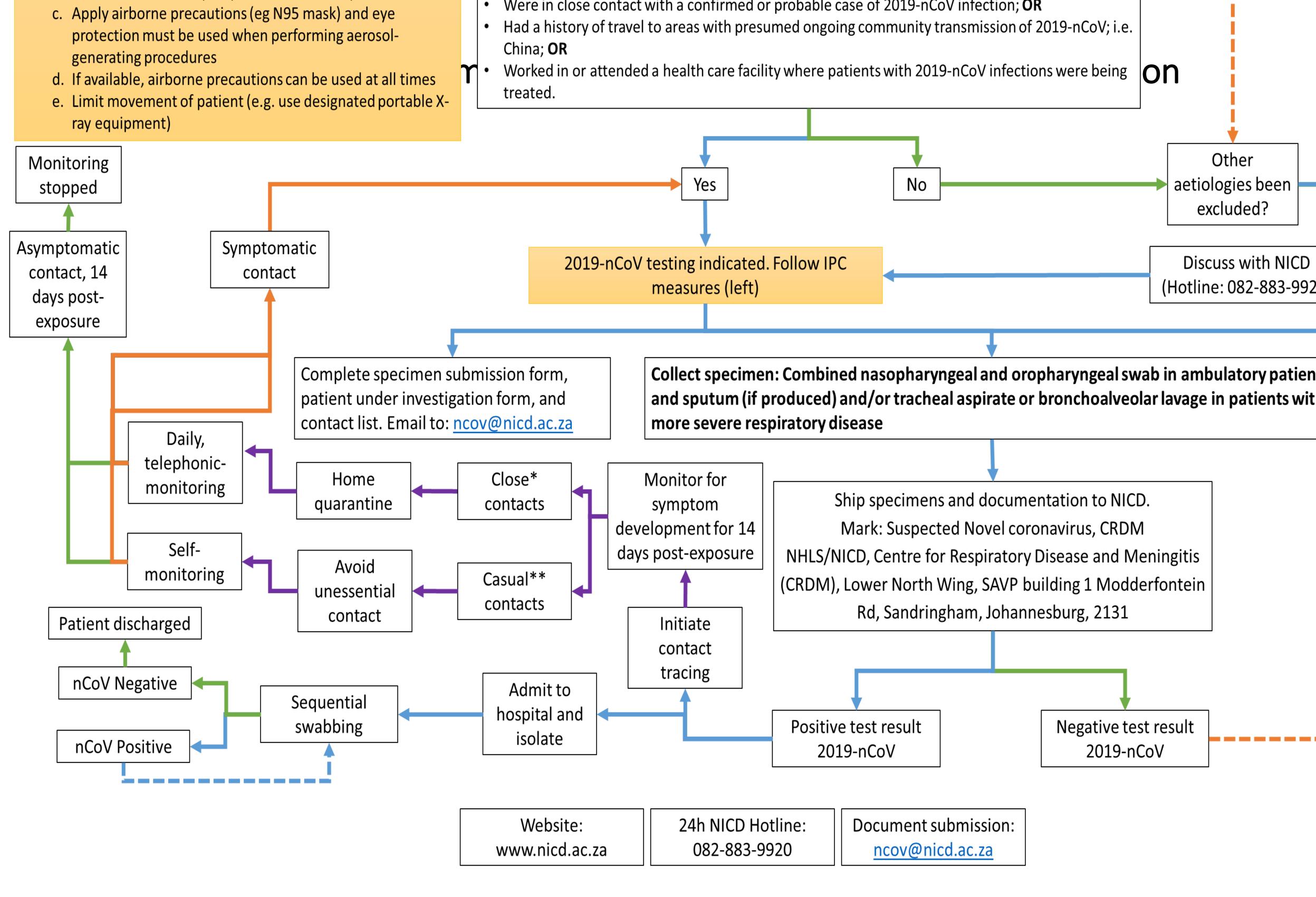
* Close contact: A person having had face-to-face contact (<2 metres) or was in a closed environment with a 2019-nCoV case; this includes, amongst others, all persons living in the same household as a 2019-nCoV case and, people working closely in the same environment as a case. A healthcare worker or other person providing direct care for a 2019-nCoV case, while **not** wearing recommended personal protective equipment or PPE (e.g., gowns, gloves, NIOSH-certified disposable N95 respirator, eye protection). A contact in an aircraft sitting within two seats (in any direction) of the 2019-nCoV case, travel companions or persons providing care, and crew members serving in the section of the aircraft where the index case was seated. ** Casual contact: Anyone not meeting the definition for a close contact but with possible exposure.

Initial diagnosis and management of suspected case (PUI), including infection control measures

2019 novel coronavirus (2019-nCoV) process flow for use in healthcare facilities







	STAGE OF ASSESSMENT OF TRAVELLERS/PERSONS UNDER INVESTIGATION FOLLOWING ARRIVAL AT PORT									
Symptom status	Arrival and disembarkat ion	Screening by Port Health	Screening by Port Health	Seen at Immigration and customs	In depth assessment at Port Health	Meets case definition, awaiting transfer by EMS	Transported by EMS to health facility	In Emergency Medicine Department (casualty)	Admission pending nCoV result	Confirmed positive test
Unknown	X			X						
No symptoms, does not meet case definition		Х		X						
Thermoscan positive			Х		X					
Meets case definition					X	X	X	X	X	Х

ACTIONS REQUIRED BY HEALTH CARE WORKERS REGARDING IPC, reporting and data collection AT THIS STAGE

Level of IPC care required by personnel	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	surgical masks, gloves, disposable gowns, eye visor/goggles if	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions [#] , incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab
Actions required	None	None	Immediately Port Health official gives patient a mask and moves traveller to private room,	None	Call NICD, collect throat swab, send to NICD Arrange transfer to medical facility	Limit staff entry to isolation room	Call ahead and request facility to prepare isolation room for clinical assessment	Take patient straight to isolation room Notify patient as suspected nCoV	Adhere to facility IPC protocols for respiratory isolation	Adhere to facility IPC protocols for respiratory isolation
References	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	#If possible, facilities should use airborne precautions

	STAGE OF ASSE	SSMENT OF TRAVELL	ERS/PERSONS UNDE	R INVESTIGATION FO	OLLOWING ARRIVAL	AT HEALTH FACILITY
Symptom status	Arrival and registration			In depth assessment by Emergency Doctor	Admission pending nCoV result	Confirmed positive test
Unknown	Х					
No symptoms, does not meet case definition		X				
Meets case definition			X	X	X	Х
	ACTIONS REQUIRE	D BY HEALTH CARE WOR	KERS REGARDING IPC, r	eporting and data collec	tion AT THIS STAGE	
Level of IPC care required by personnel	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Avoid crowds, keep 1m from people, frequent hand hygiene, MASKS not required*	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions*, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions, incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab	Droplet precautions [#] , incl surgical masks, gloves, disposable gowns, eye visor/goggles if collecting throat swab
Actions required	Screen for travel history and main complaint	Repeat screen for travel history and main complaint	Immediately provide patient with mask, and isolate patient	Collect throat swab, send to NICD	Adhere to facility IPC protocols for respiratory isolation	Adhere to facility IPC protocols for respiratory isolation; consider moving patient to designated facility
References	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	WHO guidelines 'Advice on use of masks' (*individual may choose to wear mask)	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV'	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV' (*airborne precautions if possible)	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV' (*airborne precautions if possible)	RSA Coronavirus guidelines on NICD website WHO 'IPC for NCoV' (*airborne precautions if possible)

Clinical management *prepared by Dr Jeremy Nel, Helen Joseph Hospital

Clinical management of suspected /confirmed nCoV case is essentially management of a Severe Acute Respiratory Illness (SARI)

There are two issues:



KEEP A BROAD DIFFERENTIAL DIAGNOSIS BEFORE DIAGNOSIS CONFIRMED





SUPPORTIVE CARE OF A SEVERE ACUTE RESPIRATORY ILLNESS

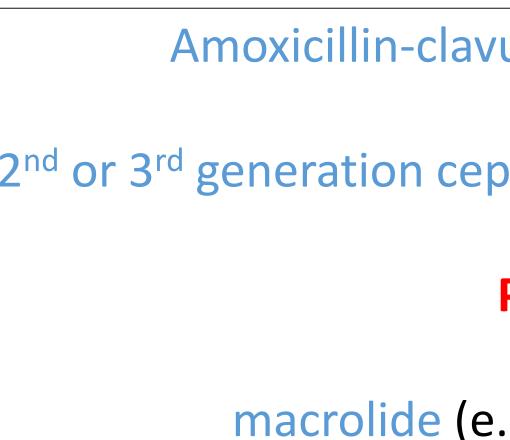
Important differential diagnosis

- Conventional bacterial pneumonia
- Atypical bacterial pneumonia
- Other viral pneumonias
- Pneumocystis pneumonia

umonia nia

Bacterial pneumonia

- Severe pneumonias generally require broad-spectrum antibiotics empirically.
- Recommended options for community-acquired pneumonia:



2017 SA Community-acquired Pneumonia Guidelines *J Thorac Dis*. 2017;9(6):1469–1502. doi:10.21037/jtd.2017.05.31

Amoxicillin-clavulanate (Augmentin)

OR 2nd or 3rd generation cephalosporin (e.g. ceftriaxone)

PLUS

macrolide (e.g. azithromycin)

Corticosteroids

- Avoid routine administration
- Although corticosteroids may be of benefit in severe prolonged viral shedding and increased mortality in influenza. (PMID: 30798570)
- Concern about possible similar effects in other viral pneumonias (including possibly 2019-nCoV)
- Should only be used if, after careful consideration, risks outweigh benefits
 - pneumonia

bacterial pneumonias, they have been associated with

• E.g. Suspected adrenal insufficiency, COPD, *Pneumocystis*

Atypical bacterial pneumonias

- Important differential diagnosis of a viral pneumonia. Like a viral pneumonia these may have:
 - Flu-like symptoms: pharyngitis, headache, myalgias, dry cough, rhinorrhoea
 - Bilateral infiltrates can appear reticulonodular / patchy don't have to have consolidation

Empiric treatment options:

- Macrolide (e.g. azithromycin) OR
- Quinolone (e.g. levofloxacin, moxifloxacin) OR
- Doxycyline

Viral pneumonia

- Influenza, parainfluenza, human metapneumovirus, respiratory syncytial virus, adenovirus, etc.
- Influenza is an important differential diagnosis to entertain, since:

 - It is potentially treatable.

• It is currently influenza season in the Northern hemisphere, where many of the 2019-nCoV suspects will have come from.

Influenza treatment

- Consider empiric oseltamivir (Tamiflu) or zanamivir
 - Are severely ill
 - asthma/COPD, etc.)
- Treatment should be started as soon as possible (best chance of benefit within 48 hours of symptom onset)

Oseltamivir 75mg po 12-hourly for 5 days

For more information, see 2019 NICD Influenza Guidelines http://www.nicd.ac.za/wp-content/uploads/2019/06/Influenza-guidelines-rev_-6-June-2019clean.pdf

treatment in patients with an influenza-like illness who:

• Are at high risk for complications (pregnant women, HIV patients, patients with

Pneumocystis pneumonia

• Consider if:

- 1. Patient significantly immunocompromised: HIV positive with CD4 < 200, chronic systemic steroid use, chemotherapy, transplant patients, etc.)
- 2. Diffuse bilateral infiltrates (often with a mid- to lower-zone predominance)
- 3. Hypoxaemia at rest (or in mild cases, with exertion)

• Consider empiric treatment if the above criteria are met:

Cotrimoxazole (Bactrim)

Prednisone if severe disease $(pO_2 < 70 \text{ mmHg}, \text{ or alveolar-arterial gradient} > 35)$

Basic work-up of patients with SARI

- Chest X-ray
- Blood cultures
- If productive of sputum: sputum MCS
- Samples for 2019-nCoV testing
- If available (private sector > public sector)

 - Urine *Legionella* antigen
- If PCP suspected:
 - Serum beta-D-glucan
 - Sputum sample / bronchoalveolar lavage (not always possible) for PCP

• Nasopharyngeal and oropharyngeal swabs for respiratory viruses and atypical pathogens

Supportive management of SARI

- Oxygen if required (titrate to $SpO_2 \ge 90\%$, or 92-95% in pregnant patients)
- Ventilatory support if required
 - protective ventilation:
 - Low tidal volumes of 6 mL/kg or less
 - Low plateau airway pressure of 30 cm H₂O or less
 - Moderate-high PEEP levels to recruit lung
- lung protective ventilation.)

• If ARDS develops, consider neuromuscular prone position, and use lung-

 Restrictive fluid management (unless shock or acute kidney injury) • ... and other standard supportive measures in critically ill patients (thromboprophylaxis, neuromuscular blockade, prone position, and

Co-ordinating a public health response

Actions to support a public health response

- Activate provincial and district outbreak response teams
 - and finance
 - Provide an overview of nCoV status globally and in RSA
 - Give an overview of RSA nCoV guidelines
 - Go through 'patient flow diagrams'
 - Emphasise importance of
 - Screening using case definitions (incl

 - \bullet for confirmation

• Ensure representation from all stakeholders especially CDC, hospitals, PHC, NHLS lab rep, NICD provincial epidemiologist and NMC nurse trainer, environmental health, EPI, EMS, port health, procurement

• Facility readiness – all facilities incl PHC can use 'Facility readiness checklist' Communication re suspected cases to NICD, and rapid transport of specimen

• Identify gaps and develop an action plan. Set date for next meeting

Resources for training

- 2-page summary document for facilities
- Specimen request form, and case investigation form (both MUST be completed when a specimen is submitted)
- Training slide set from NICD
- Training videos from NICD
- Facility readiness checklist
- NDoH / NICD nCoV guidelines
- WHO IPC for nCoV 2-page document
- NDoH communications

NDoH and NICD response structures

