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Risk of nosocomial transmission of coronavirus disease 2019: an experience in a general ward setting in Hong Kong

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\textbf{SUMMARY}

\textbf{Background:} Coronavirus disease 2019 (COVID-19) was first reported in Wuhan in December 2019 and has rapidly spread across different cities within and outside China. Hong Kong started to prepare for COVID-19 on 31\textsuperscript{st} December 2019 and infection control measures in public hospitals were tightened to limit nosocomial transmission within healthcare facilities. However, the recommendations on the transmission-based precautions required for COVID-19 in hospital settings vary from droplet and contact precautions, to contact and airborne precautions with placement of patients in airborne infection isolation rooms.

\textbf{Aim:} To describe an outbreak investigation of a patient with COVID-19 who was nursed in an open cubicle of a general ward before the diagnosis was made.

\textbf{Method:} Contacts were identified and risk categorized as 'close' or 'casual' for decisions on quarantine and/or medical surveillance. Respiratory specimens were collected from contacts who developed fever, and/or respiratory symptoms during the surveillance period and were tested for SARS-CoV-2.

\textbf{Findings:} A total of 71 staff and 49 patients were identified from contact tracing, seven staff and 10 patients fulfilled the criteria of 'close contact'. At the end of 28-day surveillance, 76 tests were performed on 52 contacts and all were negative, including all patient close contacts and six of the seven staff close contacts. The remaining contacts were asymptomatic throughout the surveillance period.

\textbf{Conclusion:} Our findings suggest that SARS-CoV-2 is not spread by an airborne route, and nosocomial transmissions can be prevented through vigilant basic infection control measures, including wearing of surgical masks, hand and environmental hygiene.

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started in Wuhan since December 2019 but has now spread throughout different provinces in China, and has since been declared a Public Health Emergency of International Concern by the World Health Organization (WHO) on 30th January 2020 [1–3]. Hong Kong, in close vicinity with China and ruthlessly affected by the Severe Acute Respiratory Syndrome (SARS) 17 years ago, started preparation for COVID-19 on 31st December 2019, when an alert of clustered pneumonia of unknown origin was announced by the Wuhan Municipal Health Commission [4]. The first imported case of COVID-19 in Hong Kong was reported on 23rd January 2020, and the first local case (with no known travel history or travel-related contact history) was reported on 31st January 2020. The WHO currently recommended droplet and contact precautions for patients with suspected or known COVID-19, while applying airborne precautions when performing aerosol-generating procedures (AGPs) [5]. While the Australian Government has adopted the WHO recommendations [6], the Centers for Disease Control and Prevention, Public Health England and the Hong Kong Hospital Authority (HA), the statutory body responsible for managing Hong Kong’s public hospital service, recommended contact and airborne precautions for patients with suspected or known COVID-19, including placement in an airborne infection isolation room (AIIR) [7–9]. The potential risk of nosocomial transmission of SARS-CoV-2 has posed great stress and anxiety to healthcare workers, a shadow cast by the SARS epidemic in 2003 [10,11]. For COVID-19, current reports from China reveal an attack rate of 2.09–29% among healthcare workers [3,12,13] Although the major routes of transmission of SARS-CoV-2 is believed to be droplet and contact, some healthcare workers remained extremely mindful of the potential ‘super-spreading events’ due to opportunistic airborne transmission through various healthcare-related activities [11,14]. While opening suction of respiratory tract, intubation, bronchoscopy, and cardiopulmonary resuscitation were well accepted as AGPs [5], the extent of infectious aerosols generated by the use of nebulizers, high-flow oxygen (especially through venturi-type masks) and non-invasive positive pressure ventilations were more controversial [15,16]. These activities were avoided during the SARS epidemic in view of the potential risks [15], but a subsequent systematic review found no statistically significant increase in SARS transmission risk with therapeutic activities such as suction before or after intubation, nebulizer treatment, oxygen mask manipulation or chest compression [16]. Here, we report the outbreak investigation and outcome of 49 patients and 71 healthcare workers exposed to a patient with severe pneumonia due to SARS-CoV-2 in a general ward setting before the diagnosis was made.

Material and methods

Setting

Queen Elizabeth Hospital is a major acute hospital with over 2000 beds with a 24-h Accident and Emergency Service. The Accident and Emergency Service has an attendance of approximately 14,700 patients per month. Ward A is a 53-bedded female medical ward for renal patients, with five open cubicles and two three-bedded rooms (Figure 1). Ward A has an average of 260 admissions per month.

Epidemiological investigation

On 3rd February 2020, an epidemiological investigation was convened when a patient was diagnosed with COVID-19. The patient had stayed in an open cubicle (bed 2) of ward A with 10 other patients for 35 h (including 18.5 h of oxygen therapy at 8 L/min) before transferring to AIIR (12 air changes per hour (ACH)) for intubation with implementation of contact and airborne precautions. Ethical approval was obtained from the Research Ethics Committee of the Kowloon Central/Kowloon East Cluster, HA. Written consent for publication was obtained from the index patient.

Contact tracing and outbreak management

Immediate infection control measures were implemented, and contact tracing was conducted to search for all staff and patients who had been exposed to the index patient before the diagnosis of COVID-19 was made. A contact case was defined as a patient or staff who stayed or worked in the same ward as the index patient. Patients would be identified through the Patient
Administration Contact Tracing System from the index patient admission until the diagnosis of COVID-19 was made, while staff contacts were identified through ward managers. These contacts were interviewed and risk categorized according to the nature of activities, duration of exposure, personal protective equipment (PPE) worn at the time of exposure. 'Staff close contact' was defined as staff who had contact within 2 m of the index case for a cumulative time of >15 min, or had performed AGPs, without 'appropriate' PPE. ‘Appropriate’ PPE in the above contact episodes referred to the use of N95 respirator, face shield/goggles, gown and gloves. Patients who shared the same cubicle with the index case were considered as 'patient close contact'. Staff close contacts were subjected to a 14-day work exclusion and quarantined at a designated camp site, followed by medical surveillance for another 14 days. Patient close contacts were quarantined into an AIIR (or quarantine camp if the patient was deemed clinically stable to be discharged from hospital) for 14 days, followed by medical surveillance for 14 days. Other staff and patient contacts ('casual contacts') were subjected to medical surveillance for 28 days with no restriction to work or discharge from hospital. Body temperature and respiratory symptoms were monitored daily throughout the 28-day period. Any abnormalities were reported to the medical personnel at the quarantine camp, or to the hospital infection control team, with hospitalization into an AIIR and testing of SARS-CoV-2 where indicated.

Real-time reverse transcription polymerase chain reaction assay for SARS-CoV-2

Upper respiratory tract specimens, e.g., nasopharyngeal aspirates (NPA), nasopharyngeal swabs (NPS) with or without concurrent throat swabs, or lower respiratory tract specimens, e.g., sputum, tracheal aspirate or bronchoalveolar lavage, were all acceptable specimen types for the real-time reverse transcription-polymerase chain reaction (RT-PCR) assay. All specimens were preserved in viral transport medium before further processing. Total nucleic acid extraction was performed using MagNA Pure LC 2.0 (Roche, Switzerland) (from 3rd to 5th February 2020) or MagNA Pure 96 (Roche, Switzerland) (from 6th February 2020), and the RT-PCR was performed using LightMix® Modular SARS and Wuhan CoV E-gene, EAV RNA extract control (TIB-MOLBIOL, Berlin, Germany), and the LightCycle® Multiplex RNA Virus Master (Roche Diagnostics, Mannheim, Germany) on cobas z 480 real-time PCR analyser (Roche Diagnostics, Mannheim, Germany) according to the manufacturer’s instructions. Briefly, each 20-μL reaction mixture contained 4.0 μL of Roche Master, 0.1 μL of RT enzyme, 0.5 μL of reagent mix, 5.4 μL of water and 10 μL of extracted RNA template. RT-PCR was performed under the following conditions: RT step at 55°C for 5 min and 95°C for 5 min, then 45 thermal cycling at 95°C for 3 s, 60°C for 12 s and 72°C for 3 s, followed by cooling at 40°C for 30 s. Specimens with a Cₚ-value of lower than 39 were be sent to the Public Health Laboratory Service Branch, Centre for Health Protection, Department of Health, Hong Kong Special Administrative Region, for confirmation.

Results

Outbreak investigation

The index patient, a 64-year-old woman, attended the Department of Accident and Emergency at 23:42 h on 1st
| Type of contact | >15 min contact time (duration of contact if known, hours) | Activity performed (staff)/bed number at time of exposure (patient) | Patient on 8 L/min oxygen | PPE used during contact | SARS-CoV-2 test during surveillance period (day of testing from last exposure date*) |
|----------------|----------------------------------------------------------|---------------------------------------------------------------------|---------------------------|-------------------------|--------------------------------------------------------------------------------|
| Staff (Doctor) | Yes                                                      | History taking, chest auscultation                                  | No                        | N95                     | None                                                                             |
| Staff (Doctor) | Yes                                                      | Blood taking                                                       | No                        | Surgical mask           | Yes (1)                                                                          |
| Staff (Nurse)  | Yes                                                      | Administration of medications                                      | Yes                       | N95, goggles            | Yes (1)                                                                          |
| Staff (Nurse)  | Yes                                                      | Obtained nasopharyngeal swab                                       | Yes                       | N95, gloves, cap         | Yes (1)                                                                          |
| Staff (Nurse)  | Yes                                                      | Escorted patient from general ward to airborne isolation ward (AIIR) | Yes                       | N95                     | Yes (1)                                                                          |
| Staff (Nurse)  | Yes                                                      | Nurse-in-charge for the affected cubicle                           | Yes                       | N95                     | Yes (1)                                                                          |
| Staff (Nurse)  | Yes                                                      | Escorted patient from general ward to AIIR                        | Yes                       | N95                     | Yes (1)                                                                          |
| Patient        | Yes (35)                                                 | 01                                                                  | Yes                       | Consistent use of surgical masks       | Yes (1, 4, 8, 11, 13)                                                             |
| Patient        | Yes (33)                                                 | 51                                                                  | Yes                       | —                        | Yes, (1, 13)                                                                     |
| Patient        | Yes (35)                                                 | 03                                                                  | Yes                       | —                        | Yes, (1, 3, 13)                                                                  |
| Patient        | Yes (35)                                                 | 04                                                                  | Yes                       | Inconsistent use of surgical masks    | Yes (1, 2, 4, 14)                                                                |
| Patient        | Yes (35)                                                 | 05                                                                  | Yes                       | Consistent use of surgical masks      | Yes (1, 14)                                                                     |
| Patient        | Yes (35)                                                 | 06                                                                  | Yes                       | —                        | Yes (1, 6, 9, 13)                                                               |
| Patient        | Yes (35)                                                 | 07                                                                  | Yes                       | —                        | Yes (2, 4, 9, 11, 14)                                                           |
| Patient        | Yes (35)                                                 | 08                                                                  | Yes                       | Inconsistent use of surgical masks    | Yes (1, 2, 5, 7, 11, 13)                                                       |
| Patient        | Yes (35)                                                 | 41                                                                  | Yes                       | Consistent use of surgical masks      | Yes (13)                                                                        |
| Patient        | Yes (35)                                                 | 42                                                                  | Yes                       | Inconsistent use of surgical masks    | Yes (14)                                                                        |

—, not available due to inability of patient to provide information on compliance of surgical mask wearing; AIIR, airborne infection isolation room; PPE, personal protective equipment.

* Last exposure date, i.e. Feb 3, 2020, was counted as day 0.
| Close/high- to medium-risk contact | World Health Organization | Centers for Disease Control and Prevention | Public Health England [28] | European Centres for Disease Prevention and Control [29] | Communicable Disease Network Australia [6] |
|----------------------------------|--------------------------|-------------------------------------------|-----------------|------------------------------------------------|--------------------------|
| (No explicit differentiation between close and casual contact) Contacts include provision of direct care for patients with COVID-19, or visiting patients or staying in the same environment as a patient with COVID-19, or working with HCWs with COVID-19 disease | Within 2 m of a person with COVID-19 for a prolonged period of time (>1–2 min). Taking into account the clinical syndrome patients (e.g., coughing) and whether patient wore facemask High risks: perform or present in the same room for AGPs when the HCW eyes, nose or mouth were not protected. Medium risks: perform or present in the same room for AGPs without using gown and gloves (but eyes, nose and mouth were protected), or HCWs with unprotected eyes, nose or mouth with prolonged close contact with patients with COVID-19 who was or was not wearing a facemask, or HCW with direct contact with secretion/excretion of COVID-19 patient without wearing gloves and failed to perform immediate hand hygiene | Face-to-face contact, or spending >15 min within 2 m of an infected person | Direct physical contact, or unprotected direct contact, or face-to-face contact within 2 m for >15 min, or closed environment for 15 min at a distance of <2 m as a COVID-19 case, or laboratory workers handling specimens from a COVID-19 case; AND without recommended PPE or with a possible breach of PPE | Face-to-face contact for >15 min in any setting with a confirmed case in the period extending from 24 h before onset of symptoms in the confirmed case, or sharing of a closed space with a confirmed case for >2 h in the period extending from 24 h before onset of symptoms in the confirmed case |

| Casual-/low-risk contacts | HCW wearing a facemask or respirator only and have prolonged close contact with a patient who was wearing a facemask, or HCW using all recommended PPE (i.e., a respirator, eye protection, gloves and a gown) while caring for or having contact with the secretions/excretions of a patient, or HCW (not using all recommended PPE) who have brief interactions with a patient regardless of whether patient was wearing a facemask (e.g., brief conversation at a triage desk; briefly entering a patient room but not having direct contact with the patient or their secretions/excretions; entering the patient room immediately after they have been discharged) | Not defined | Person in a closed environment with a COVID-19 case for <15 min, or at a distance of more than 2 m, or a person having had face-to-face contact with a COVID-19 case for <15 min and at within 2 m | <15 min face-to-face contact with a symptomatic confirmed case in any setting, or sharing a closed space with a symptomatic confirmed case for less than 2 h |

(continued on next page)
February 2020 with fever, productive cough and breathlessness for 2 days. She developed flu-like symptoms on 24th January 2020 with transient improvement after taking antibiotics and symptomatic treatment prescribed by a general practitioner. However, her fever relapsed on 30th January 2020 with productive cough and dyspnea. She had no history of travel in the preceding month, but she owned a fashion boutique with many mainland Chinese customers owing to its proximity to the West Kowloon high-speed rail station. On admission, she had a fever of 39.6°C with sinus tachycardia of 126 bpm and blood pressure of 198/92 mmHg. She was tachypnoeic with oxygen saturation of 80% in room air. Bilateral crepitations were heard on auscultation and chest X-ray revealed multi-lobar pulmonary infiltrates (Figure 2(a)). In the absence of history of travel to China or contact with a confirmed COVID-19 patient, she was admitted to ward A in an open cubicle under standard precautions for community-acquired pneumonia (6 ACH, normal pressure setting) at 01:22 h on 2nd February 2020. She could not wear a surgical mask as she was on oxygen therapy through a simple facemask (Soundway®, Ningbo Shengyurui Medical Appliances Co. Ltd, Ningbo, China; Figure 2(b)). She also had frequent coughs while residing in the ward. She became more hypoxic around 18:00 h and required an increase in oxygen therapy to 8 L/min, delivered through the same facemask. On 3rd February 2020, she was transferred to an AIIR at 12:35 h and was electively intubated at 13:00 h for progressive respiratory failure. Nasopharyngeal swab for multiplex PCR FilmArray® RP2 panel (Biofire Diagnostics, bioMérieux, Marcy-l’Étoile, France), urine for BinaxNOW Legionella and pneumococcal antigen (Abbott, IL, USA) were negative. The patient was tested for COVID-19 as enhanced laboratory surveillance on 3rd February 2020, where both combined NPA with throat swab, and tracheal aspirate showed detectable SARS-CoV-2 RNA, with a Cp value of 22.8 and 26.1, respectively. The patient was subsequently transferred to the Infectious Diseases Centre, Princess Margaret Hospital, for further management.

Contact tracing and outbreak management

Terminal disinfection and changing of all curtains were performed in ward A immediately after the diagnosis of COVID-19 in the index patient, followed by regular environmental disinfection twice daily with 1000 ppm sodium hypochlorite. Also, ward A was closed to new admissions for 14 days. Enhanced infection control measures in accordance with the Emergency Response level in the HA since 26th January 2020 were reinforced, including the wearing of surgical masks (ASTM F2100 level 1) by all staff, patients and visitors in all hospital areas, as well as the minimization of traffic by suspension of visiting hours, volunteer service and clinical attachments.

A total of 71 staff and 49 patients were identified from contact tracing. Seven staff and 10 patients fulfilled the criteria of ‘close contact’ (Table I). All staff and patients who did not fulfill the pre-defined criteria for close contacts were managed as ‘casual contacts’ (Supplementary Tables S1 and S2). Thirty staff and 22 patients developed fever and/or reported respiratory symptoms during the surveillance period, with 76 respiratory specimens sent for RT-PCR for SARS-CoV-2. All specimens from the 52 contacts were negative.
Discrimination

More than 100,000 cases of COVID-19 have been reported worldwide at the time of writing, after just over 2 months after the identification of this novel coronavirus [17,18]. While it is believed that person-to-person spread of SARS-CoV-2 occurs predominantly through droplet and contact transmissions, the actual dynamics remained uncertain. The reproduction number of SARS-CoV-2 appears to be higher than SARS, with an initial estimation of 1.4–6.49, and median of 2.79 [19–21]. The high attack rates among household contacts [22–24], and passengers on the Diamond Princess Cruise are most concerning [25]. In the described scenario, our patient coughed frequently and a high viral load (as implicated by the low Cp-value) detected from her respiratory specimens; moreover, she could not put on a surgical mask during her stay in the general medical ward. All these factors could have contributed to a great deal of droplet generation from our index patient, with secondary contamination of her surroundings. Moreover, she received oxygen therapy at 8 L/min through a simple oxygen facemask for a significant amount of time. Oxygen therapy with flow of ≥6 L/min was considered an AGP in the HA hospitals. Despite the aforementioned, none of the neighbouring patients or staff contacts were infected. While there is no consensus on what constitutes ‘close’ and ‘casual’ contacts, our risk assessment approach was similar to the subsequent published guidance for contact tracing [6,26–29] (Table II). The authors believe that the universal mask wearing of staff, patients and visitors as a component of the Emergency Response in HA hospitals, as well as the heightened alertness to hand hygiene and environmental cleanliness have played a major part in halting further transmissions in this incident. In fact, frequent hand hygiene and appropriate use of surgical masks have previously been shown to be a protective factor against SARS, and influenza [30–33]. In the Shenzhen family cluster of COVID-19, the uninfected family member was the only person reported to have consistent use of surgical masks [22]. Also, the lack of secondary transmission in our experience may imply that oxygen therapy ≥6 L/min with a simple facemask poses a low risk of aerosol generation, consistent with the finding from a previous study [34]. Another local study on aerosol dispersion in various respiratory therapies demonstrated that the air leak through the side vents of the simple oxygen facemask during delivery of oxygen for 6, 8, 10 L/min were limited to 0.22, 0.3 and 0.4 m [35], respectively, hence the neighbouring patients who should be at least 1 m from the index patient were not at increased risk of droplet transmission despite the use of high-flow oxygen.

There are a few limitations to our study. Firstly, our sample size was small with only one index patient. However, the detailed description of contact tracing including nature of exposure and PPE worn involving 120 contacts may still be a useful observation of transmission dynamics in a normal ward environment, because the majority of the subsequent patients were admitted directly into AIIR. Secondly, we only tested contacts with fever or respiratory symptoms with RT-PCR, hence the possibility of asymptomatic infection among the staff close contacts can not be entirely excluded. None the less, all 10 close patient contacts, who had stayed in the same cubicle as the index patient for > 20 h had multiple negative SARS-CoV-2 RT-PCR results, up to days 13 and 14 from the last exposure. Thirdly, quantitative RT-PCR and viral culture were not available at our centre, thus the exact viral load and the viability of the virus can not be ascertained.

In conclusion, the route of transmission of SARS-CoV-2 remains to be confirmed, however there is no reason to suspect its physical property would differ greatly from SARS, MERS or any other coronavirus. Our index patient with COVID-19, despite a 35-h stay in an open cubicle in a general ward, did not result in any secondary nosocomial infection in any contacts at the time of writing (37 days after last exposure to the index case). Vigilance with basic infection control measures, including wearing of surgical masks, hand hygiene and environmental hygiene continues to remain fundamental and essential in the prevention of human-to-human transmissions of SARS-CoV-2.

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Conflict of interest statement

None declared.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.jhin.2020.03.036.

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