Role of chest radiograph in MERS-Cov pneumonia: a single tertiary referral center experience in the United Arab Emirates

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Abstract

Background: The Middle East respiratory syndrome coronavirus (MERS-Cov) continues to be a source of concern due to intermittent outbreaks. Serial chest radiographic changes in MERS-Cov patients were analyzed for various variables that could be compared to the patients’ final outcomes in a cluster of MERS-Cov patients and to identify a predictor of mortality in the United Arab Emirates.

Results: A total of 44 MERS-Cov cases were reviewed. The mean age of the patients was 43.7 ± 14.7 years. The chest radiograph was abnormal in 14/44 (31.8%). The commonest radiology features include ground-glass opacities (seven of 14, 50%), ground-glass and consolidation (seven of 14, 50%), pleural effusion (eight of 14, 57.1%), and air bronchogram (three of 14, 21.4%). The mortality rate was 13.6% (six of 44); the deceased group (6 of 44, 13.6%) was associated with significantly higher incidence of mechanical ventilation ($p < 0.001$), pleural effusion ($p < 0.001$), chest radiographic score ($8.90 ± 6.31, p < 0.001$), and type 4 radiographic progression of disease ($p < 0.001$). A chest radiographic score at presentation was seen to be an independent and strong predictor of mortality (OR [95% confidence interval] 3.20 [1.35, 7.61]). The Cohen κ coefficient for the interobserver agreement was $k = 0.89$ ($p = 0.001$).

Conclusion: The chest radiographic score, associated with a higher degree of disease progression (type 4), particularly in patients with old age or with comorbidity, may indicate a poorer prognosis in MERS-Cov infection, necessitating intensive care unit management or predicting impending death.

Keywords: MERS-Cov, Chest radiograph, ARDS

Background

The Middle East respiratory syndrome coronavirus (MERS-Cov) was first described as a zoonotic disease in Saudi Arabia in 2012 and subsequently spread to countries in the Middle East and beyond the Arabian Peninsula [1]. Since 2012, UAE has reported 88 cases (including the patient-reported above) of MERS-Cov infection and 12 associated deaths [1]. From 2012 to July 2020, the total number of laboratory-confirmed MERS-Cov-infected cases reported globally by the WHO was 2562, with 881 associated deaths [1–3]. Approximately 55% of patients with MERS-Cov require admission to the intensive care unit due to respiratory failure, including the development of adult respiratory distress syndrome (ARDS) [4, 5]. ARDS frequently leads to a clinically significant reduction in health-related quality of life and is also associated with various radiological changes [6]. A recent study described chest radiograph abnormalities associated with MERS-Cov and demonstrated that sequential chest radiographic evaluation is a valuable tool for the early diagnosis and monitoring of disease progression [4].
The purpose of this study was to determine the role of various variables such as mechanical ventilation days, chest radiographic score, disease progression on radiographic imaging, and pleural effusion in predicting mortality in clusters of MERS-Cov patients in the United Arab Emirates.

Methods
The study was conducted on 44 patients with positive MERS-Cov retrospectively from electronic medical records. The patient diagnosis was performed using RT-PCR and was based on the World Health Organization criteria [7]. All these 44 patients were recruited from the United Arab Emirates. The available chest radiographs were obtained between July 1, 2013, and September 2019.

The Institutional Review Board approved the study protocol, and the requirement for informed consent was waived due to the study’s retrospective nature. The inclusion criteria were as follows: admission to the hospital with symptoms suggestive of MERS-Cov without a history of other lung infectious diseases and has laboratory-confirmed MERS-Cov infection as determined by RT-PCR in any specimen or saliva. The exclusion criteria were the group of patients with flu-like symptoms with negative RT-PCR test for MERS-Cov and those patients without any initial chest radiographs. The study group of 44 patients was divided into two subgroups on the basis of the final outcome of recovery or death.

Study protocol and data analysis
Digital radiography equipment (Mobi1lett Plus; Siemens, Erlangen, Germany) was used to obtain portable anteroposterior (AP) projection radiographs as per the local protocol. Two chest radiographs were routinely taken per day for very sick patients in the ICU and once every other day during the recovery stage. Two radiologists with more than 20 years of experience (KMD, JK) reviewed the chest radiographs independently. Any of the discrepant reports was assessed independently by an arbitrator (KG). All patients were evaluated based on the risk factors like mechanical ventilation days, chest radiographic score, pleural effusion, and chest radiographic disease progression pattern [4, 8].

The lung parenchyma and airways were evaluated for consolidation, ground-glass opacity (GGO), pleural effusion, cavitation, and pneumothorax [4, 8]. For the purpose of calculating the lung score, each lung was divided into three zones, and each zone was evaluated for disease process involvement. The development of MERS-Cov lesions within each lung zone was assigned a score ranging from 0 (normal) to 4 (complete involvement of one zone); a score of 24 indicated complete involvement of all six zones. The scores for each of the six zones in each chest radiographic study were added together to generate a cumulative chest radiographic score ranging from 0 to 24 based on the degree of lung parenchymal involvement. At the time of initial presentation and at the peak of disease activity, scores were taken. The disease progression pattern (types 1–4) was identified from chest radiographs described by Wong et al. [9]. Radiographic deterioration followed by complete improvement was used to define type 1 disease progression. Type 2 disease progression was defined as stable radiographic changes without discernible radiographic peaks or a decrease in overall mean lung involvement of less than 25%. The progression of type 3 disease was defined as fluctuating radiographic changes with at least two radiographic peaks separated by a period of mild remission, with remission defined as a level of mean lung parenchyma involvement that was more than 25% different from the peak level. Progressive radiographic deterioration was defined as type 4 progression. Several of the above parameters, including chest radiographic disease progression pattern (types 1–4), were compared between those who recovered and those who did not.

Statistical analyses
The categorical variables were described using absolute or relative frequency distribution and quantitative variables using central tendency or location measures, such as mean (standard deviation) or median (interquartile range). The association between qualitative independent variables was assessed using the chi-square test or Fisher’s exact test as appropriate. To compare quantitative variables between the two groups (deceased and recovered), a t-test or Wilcoxon rank-sum test was used as appropriate. A logistic regression procedure was used to determine factors associated with mortality; the results are presented in the form of the odds ratio and corresponding 95% confidence interval and p value. The discrimination performance of the logistic regression was evaluated using the area under the curve (AUC). A p-value <0.05 (two tailed) was considered to be statistically significant. Statistical software, STATA/SE version 14.2 (Stata Corp LP, College Station, TX, USA), was used for the analysis. Interobserver reliability is calculated using Cohen’s kappa statistics (k) to see the degree of agreement between the two reviewers.

Results
The present cohort includes 44 MERS-Cov patients with a mean age of 43.7 ± 14.7 years, range, 11–71 years, with M: F 28 males and 16 females. The demographic and clinical characteristics of MERS-Cov cases by mortality status are shown in Table 1. Forty-four patients of MERS-Cov were divided into two groups, those who survived (n = 38) and those who died (n = 6). Patients’ age in the deceased group was significantly higher than that seen in the recovered group (59.8 ± 14.2 vs. 41.16 ± 13.36 years, p = 0.008). The most common symptoms at the time of the presentation were fever (27%), cough (27%), dyspnea (9%), rhinitis (9%), and myalgia (4.5%). Associated comorbidity was noted in
24 of 44 (54.5%) of cases with hypertension (15 of 44, 34%), diabetes (ten of 44, 22.7%), heart disease (four of 44, 9%), obesity (three of 44, 7%), end-stage renal disease (three of 44, 7%), smoker (two of 44, 4.5%), multiple myeloma (two of 44, 4.5%), and one each of asthma, COPD, cirrhosis of the liver, lung fibrosis, and SLE. The deceased patients required a significantly longer period of mechanical ventilator (Table 2) support (10.17 ± 13.66 vs. 0.76 ± 3.18 days, p ≤ 0.001). One of the six deceased patients had a secondary *Klebsiella pneumonia* infection.

The 44 patients underwent 182 chest radiographs during hospitalization; these were abnormal in 14 patients (31.8%) and normal in 30 patients (68.1%). Lung opacification was primarily located in the right mid and lower zone (nine of 14, 64.2%), followed by the left lower zone (three of 14, 21.4%) and left mid-zone (two of 14, 14.2%). The characteristics of the radiology features (Table 3) were GGO in the peripheral location (seven of 14, 50%), GGO and consolidation (seven of 14, 50%) (Figs. 1 and 2), pleural effusion (eight of 14, 57.1%), and

### Table 1 Demographic and clinical characteristics of MERS-Cov cases by final outcome (n = 44)

| Characteristics               | Disease outcome | p value* |
|-------------------------------|-----------------|----------|
|                               | Deceased (%)    | Recovered (%) |
| Frequency                     | 6 (13.6)        | 38 (86.3) |
| Age                           |                 |           |
| Mean (SD)                     | 59.8 (14.02)    | 41.16 (13.36) |
| Median (IQR)                  | 64.5 (26)       | 41 (21)   | 0.008 |
| Gender, M:F ratio             | 5:01            | 23:15     | 0.65 |
| Mechanical ventilation duration |               |           |
| Mean (SD)                     | 10.17 (13.66)   | 0.76 (3.18) |
| Median (IQR)                  | 4.5 (2)         | 0 (0)     | <0.001 |
| Hospitalization days          |                 |           |
| Mean (SD)                     | 40.33 (83.14)   | 4.76 (4.60) |
| Median (IQR)                  | 6.5 (5)         | 3 (3)     | 0.0123 |

*The Fisher exact test for categorical variables and the Mann-Whitney U test for continuous variables were used for association/comparison between the groups.

### Table 2 Distribution of predictors in 44 MERS-Cov patients by final outcome

| Characteristics | Total lesions n (%) | Disease outcome | p-value* |
|-----------------|---------------------|-----------------|----------|
| Frequency       | 44 (100)            |                 |          |
| Chest radiograph|                     |                 |          |
| Normal          | 30 (68.1)           | 0 (0.00)        | 30 (78.95) | <0.001 |
| Abnormal        | 14 (31.8)           | 6 (100.00)      | 8 (21.05) |
| Disease progression |             |                 |          |
| Normal          | 0 (0.00)            | 30 (78.95)      |          |
| T1              | 0 (0.00)            | 7 (18.42)       | <0.001 |
| T2              | 0 (0.00)            | 1 (02.63)       |          |
| T3              | 1 (16.67)           | 0 (00.00)       |          |
| T4              | 5 (83.33)           | 0 (00.00)       | <0.001 |
| Lung score      |                     |                 |          |
| At initial presentation |       |                 |          |
| Mean (SD)       | 0.47 (0.99)         | 1.82 (1.34)     | 0.25 (0.75) |
| Median (IQR)    | 0 (0.6)             | 1.48 (2.13)     | 0 (0)     | <0.001 |
| Peak of the disease |              |                 |          |
| Mean (SD)       | 1.39 (3.78)         | 8.90 (6.31)     | 0.20 (0.79) |
| Median (IQR)    | 0 (0)               | 5.09 (9.60)     | 0 (0)     | <0.001 |
| Pleural effusion|                     |                 |          |
| None            | 36 (81.82)          | 0 (00.00)       | 36 (94.74) | <0.001 |
| Unilateral      | 3 (6.82)            | 3 (50.00)       | 0 (00.00) |
| Bilateral       | 5 (11.36)           | 2 (05.26)       |          |
| Mechanical ventilation day  |                 |                 |          |
| Mean (SD)       | 10.17 (13.66)       | 0.76 (3.18)     |          |
| Median (IQR)    | 4.5 (2)             | 0 (0)           | <0.001 |
The chest radiographic score at presentation was seen to be an independent and strong predictor of mortality (OR [95% confidence interval] 3.20 [1.35, 7.61]). The AUC was 0.92 at the chest radiographic score cutoff point of 0.48 at presentation and was seen to discriminate mortality with a sensitivity and specificity of 100% and 84.21%, respectively (Fig. 3). Furthermore, the deceased group had a chest radiographic score of 8.90 ± 6.31, showing the need for intensive treatment as well as close monitoring and follow-up examination. The T4 and T3 disease progression was noted in five of the 6 (83.33%) and one of the 6 (16.67%) patients with fatal outcome (Table 2).

Patients in the deceased group (6 of 44, 13.6%) were associated with significantly higher incidence of \( p \leq 0.02 \), mechanical ventilation days \( p < 0.001 \), pleural effusion \( p < 0.001 \), chest radiographic score \( 8.90 \pm 6.31 \), \( p < 0.001 \), and type 3 and 4 radiographic progression of disease \( p < 0.001 \) (Table 2).

**Discussion**

MERS-Cov is still circulating and causing human disease in the Middle East, with isolated cases, population clusters, and nosocomial outbreaks, and there is a high risk of global spread. The current retrospective study included 44 confirmed MERS-Cov patients with a 13.6% mortality rate to establish a correlation between the mortality rate and the severity of MERS-Cov disease. The deceased group was associated with significantly higher incidence of mechanical ventilation \( p < 0.001 \), pleural effusion \( p < 0.001 \), chest radiographic score \( 8.90 \pm 6.31 \), \( p < 0.001 \), and type 4 radiographic progression of disease \( p < 0.001 \). The findings of the present cohort also indicated that for MERS-Cov patients, a chest radiographic score cutoff point of 0.48 at presentation suggests patient mortality, with a sensitivity and specificity of 100% and 84.21%, respectively. Additionally, a chest radiographic score of 8.90 ± 6.31 during the course of the disease necessitated aggressive therapy, as well as close surveillance and follow-up evaluation.

Because the chest radiograph is widely available, the ability to integrate chest radiographic results with clinical data collected routinely during ED admission is

**Table 3**: Chest radiographic findings and disease progression of MERS-Cov cases by final outcome

| Characteristics | Total lesions | Final outcome status | Disease outcome | p-value* |
|-----------------|--------------|---------------------|----------------|---------|
|                 | n (%)        | Deceased            | Recovered      |         |
| Frequency       | 44 (100)     | 6 (13.64)           | 38 (86.36)     |         |
| Type of lung opacity |       |                     |                 |         |
| Ground-glass opacity | 9 (20.45)  | 6 (100)             | 3 (7.89)       | <0.001  |
| Consolidation   | 9 (20.45)    | 6 (100)             | 3 (7.89)       | <0.001  |
| Patchy consolidation | 3 (6.82)   | 2 (33.33)           | 1 (2.63)       | 0.045   |
| Confluent consolidation | 5 (11.36) | 4 (66.67)           | 1 (2.63)       | 0.001   |
| Nodular         | 1 (2.27)     | 0                   | 1 (2.63)       | >0.999  |
| Reticular       | 1 (2.27)     | 0                   | 1 (2.63)       | >0.999  |
| Opacity central | 1 (2.27)     | 1 (16.67)           | 0              | 0.136   |
| Opacity peripheral | 10 (22.73) | 5 (83.33)           | 5 (13.16)      | 0.001   |
| Pleural effusion | 8 (18.2%)   | 6 (100)             | 2 (05.26)      | <0.001  |
beneficial for ensuring rapid stratification of pulmonary parenchymal involvement. The chest radiographic score observed in the present cohort has a close relationship with the key clinical criteria used to identify patients who require hospitalization. The chest radiographic scoring system is simple to use, repeatable, and indicative of the severity of lung parenchyma involvement. It is critical to identify the most cost-effective procedures to include in ED workflow while minimizing interaction between healthcare staff and patients as well as between patients themselves.

The majority of MERS-Cov imaging research has concentrated on CT [5, 10, 11]. Even though chest radiograph is typically the first examination for patients entering the ED with suspected MERS-Cov infection, it is also characterized by simpler logistics and use; only a small number of studies have looked into its function [4, 12, 13]. The chest radiographic score is a critical parameter associated with imaging findings that have been found to be beneficial in assessing the severity of SARS, MERS-Cov, and SARS-CoV-2 infection [13]. Increased chest radiographic scores may be used as fatal prognostic markers in patients with advanced age and comorbid lung disease infected with SARS-CoV [14]. The results in the present cohort were consistent with previous research that found a higher prevalence of chest radiographic scores, pleural effusion, and pneumothorax to be linked to a poor prognosis and short-term mortality in MERS-Cov infection [15]. Although the imaging characteristics that aid in determining the prognosis of SARS-CoV-2 infection are unknown, advanced age and progressive consolidation on imaging may indicate a poor prognosis [13]. The global mortality of MERS-Cov ranges between 32 and 67% [2, 4, 16]. The mortality rate of the current MERS-Cov cohort was 13.6%. The regional variation of death rate from previously conducted

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**Fig. 2** A 68-year-old man with Middle East respiratory syndrome coronavirus. Serial chest radiographs show a type 3 pattern of progression. **A** A frontal chest radiograph shows ground-glass opacity in the right lower zone with consolidation areas in the left mid and lower zone with GGO in the left upper zone. **B** On day 6, a follow-up frontal chest radiograph shows a marginal improvement of the left lung opacity with an increase of GGO in the right side. **C** A follow-up frontal chest radiograph on day 9 shows a considerable improvement of both lung opacities. **D** A follow-up frontal chest radiograph on day 12 shows a considerable improvement of both lung opacities. The patient died on the 13th day.
studies may be skewed due to the severity of disease and smaller sample sizes than have been investigated previously. Ahmadzadeh et al. [16] affirm that the mortality example of the MERS-Cov in Saudi Arabia is not the same as that noticed in other nations in the Middle East. The differences in the virus and the genetic background of the population affected can play a role [16]. Different reasons can include a difference in the availability or ability to implement patient isolation procedures and differences in overall medical technology among involved countries [16].

The study has some inherent limitations. First, during the disease outbreak, it was most prevalent in Dubai, Abu Dhabi, and Sharjah. The number of cases seen in the remote province of the United Arab Emirates (present cohort) was limited. Second, the relatively small cohort size limited statistical analysis, which may not reveal the actual projection. Third, the visual estimation of the lung involvement was performed in one view, and this may not reflect the actual volume of the pathology in the involved lung. Finally, autopsies were not performed on any of the deceased patients.

**Conclusion**

In conclusion, GGO and consolidation with a peripheral predominance on imaging are characteristic of MERS-Cov infection, and these lesions eventually spread to involve both lungs as the disease progresses. The extent of parenchymal abnormalities as determined by the radiographic lung score, the degree of radiographic progression, the presence of pleural effusion, and the number of days spent on mechanical ventilation all contribute to the clinical outcome or prognosis of coronavirus infection. The greater the extents of the chest radiographic score associated with a higher degree of disease progression (type 4), particularly in elderly or comorbid patients, may indicate a poorer prognosis in MERS-Cov infection, necessitating intensive care unit management or precluding death.

**Abbreviations**

MERS-Cov: Middle East respiratory syndrome coronavirus; GGO: Ground-glass opacity; SARS-CoV-2: Severe acute respiratory syndrome coronavirus 2; SARS-CoV: Severe acute respiratory syndrome coronavirus; CT: Computed tomography; ED: Emergency department; COPD: Chronic obstructive pulmonary disease; SLE: Systemic lupus erythematosus; RT-PCR: Reverse transcription polymerase chain reaction; UAE: United Arab Emirates; ARDS: Adult respiratory distress syndrome

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**Authors’ contributions**

All authors have read and approved the manuscript. Substantial contributions to the conception or design of the work, or the acquisition, analysis, or interpretation of data for the work: KMD, JAA, SAA, MTA, YS, ESS, KVG. Drafting the work or revising it critically for important intellectual content: KMD, RS, VS. Final approval of the version to be published: KMD, JAA, SAA, MTA, YS, ESS, KVG. Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved: KMD. All the authors read and approved the final manuscript.

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**Availability of data and materials**

Yes
Declarations

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Consent for publication
Waiver was obtained from the ethical committee.

Competing interests
The authors declare that they have no competing interests.

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References

1. Yusof MF, Eltahir YM, Serhan WS, Hashem FM, Elsayed EA, Marzoug BA, Abdelazim AS, Benalshah OA, al Muhairi SS (2015) Prevalence of Middle East respiratory syndrome coronavirus (MERS-CoV) in dromedary camels in Abu Dhabi emirate, United Arab Emirates. Virus Genes 50(3):509–513. https://doi.org/10.1007/s11262-015-1174-0
2. Hunter JC, Nguyen D, Aden B, al Bandar Z, al Dhaheri W, Abu Elkheir K, Khudairi A, al Mulla M, al Saleh F, Imambaccus H, al Kaabi N, Sheik FA, Sasse J, Turner A, Abdel Wareth L, Weber S, al Ameri A, Abu Amer W, Alami NN, Bunga S, Haynes LM, Hall AJ, Kallen AJ, Kuhar D, Pham H, Pringle K, Tong S, Whitaker BL, Gerber SI, al Hosani FI (2016) Transmission of Middle East respiratory syndrome coronavirus infections in healthcare settings, Abu Dhabi emirate, United Arab Emirates. Virus Genes 50(3):509–513. https://doi.org/10.1007/s11262-015-1174-0
3. WHO (2020) Middle East respiratory syndrome coronavirus (MERS-CoV) – Saudi Arabia. https://www.who.int/csr/don/02-jul-2020-mers-saudi-arabia/en/
4. Das KM, Lee EY, Al Jawder SE et al (2015) Acute Middle East respiratory syndrome coronavirus: temporal lung changes observed on the chest radiographs of 55 patients. AJR Am J Roentgenol 205(3):W267–W274. https://doi.org/10.2214/AJR.15.14445
5. Das KM, Lee EY, Enani MA, AlJawder SE, Singh R, Bashir S, al-Nakshbandi N, AlDosani K, Larsson SG (2015) CT correlation with outcomes in 15 patients with acute Middle East respiratory syndrome coronavirus. Am J Roentgenol 204(4):736–742. https://doi.org/10.2214/AJR.14.13671
6. Davidson TA, Caldwell ES, Curtis JR, Hudson LD, Steinberg KP (1999) Reduced quality of life in survivors of acute respiratory distress syndrome compared with critically ill control patients. Jama 281(4):354–360. https://doi.org/10.1001/jama.281.4.354
7. Group WM-CR (2013) State of knowledge and data gaps of Middle East respiratory syndrome coronavirus (MERS-CoV) in humans. PLoS Curr. PLoS Curr 12:5. https://doi.org/10.1371/currents.outbreaks.0bf719e352e747f8ba488fa13ca0127dbb8.
8. Hansell DM, Bankier AA, MacMahon H, McCloud TC, Müller NL, Remy J (2008) Fleischner society: glossary of terms for thoracic imaging. Radiology 246(3):697–722. https://doi.org/10.1148/radiol.2462070712
9. Wong KT, Antonio GE, Hui DSC, Lee N, Yuen EHY, Wu A, Leung CB, Rainer TH, Cameron P, Chung SSC, Sung SW, Ahuja AT (2003) Severe acute respiratory syndrome: radiographic appearances and pattern of progression in 138 patients. Radiology 228(2):401–406. https://doi.org/10.1148/radiol.2282030593
10. Hamimi A (2016) MERS-CoV: Middle East respiratory syndrome corona virus can radiology be of help? Initial single center experience. Egypt J Radiol Nucl Med 47(1):95–106. https://doi.org/10.1016/j.ejrnm.2015.11.004
11. AlJawder SE, Jamoom LG, Alharthy A, Madani TA (2014) Middle East respiratory syndrome coronavirus (MERS-CoV) infection: chest CT findings. Am J Roentgenol 203(4):782–787. https://doi.org/10.2214/AJR.14.13021
12. AlGhamdi M, Muhitfaq F, Awn N, Shahhoub S et al (2015) MERS CoV infection in two renal transplant recipients: case report. Am J Transplant 15(8):1101–1104. https://doi.org/10.1111/ajt.13085
13. Franquet T, Jeong YJ, Lam HYS, Wong HYF, Chang YC, Chung MJ, Lee KS (2020) Imaging findings in coronavirus infections: SARS-CoV–MERS-CoV, and SARS-CoV-2. Br J Radiol 93(112):20200515. https://doi.org/10.1259/bjr.2020515
14. Ko S-F, Lee T-Y, Huang C-C, Cheng Y-F, Ng SH, Kuo SY, Lin MC, Liu JW, Yang KD, Chen MC, Chen CL (2004) Severe acute respiratory syndrome: prognostic implications of chest radiographic findings in 52 patients. Radiology 233(1):173–181. https://doi.org/10.1148/radiol.2323031547
15. Cha MJ, Chung MJ, Kim K, Lee KS, Kim TJ, Kim TS (2018) Clinical implication of radiographic scores in acute Middle East respiratory syndrome coronavirus pneumonia: report from a single tertiary-referral center of South Korea. Eur J Radiol 107:196–202. https://doi.org/10.1016/j.ejrad.2018.09.008
16. Ahmadzadeh J, Mobarski K, Mousavi SJ, Aghazadeh-Attari J, Mirza-Aghazadeh-Attari M, Mohebbl I (2020) The risk factors associated with MERS-CoV patient fatality: a global survey. Diagn Microbiol Infect Dis 96(3):114876. https://doi.org/10.1016/j.diagmicrobio.2019.114876