Performance Evaluation of the PowerChek MERS (upE & ORF1a) Real-Time PCR Kit for the Detection of Middle East Respiratory Syndrome Coronavirus RNA

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Background: Molecular detection of Middle East respiratory syndrome coronavirus (MERS-CoV) using real-time reverse transcription (rRT)-PCR assays is the method of choice for diagnosis of MERS. We evaluated the performance of the PowerChek MERS (upE & ORF1a) real-time PCR Kit (PowerChek MERS assay; Kogene Biotech, Korea) a one-step rRT-PCR assay for the qualitative detection of MERS-CoV.

Methods: We evaluated PowerChek MERS assay performance in comparison with nested RT-PCR and sequencing of the RNA-dependent RNA polymerase (RdRp) and N genes. To evaluate diagnostic sensitivity and specificity, 100 clinical specimens (50 positive and 50 negative for MERS-CoV) were simultaneously tested by using the PowerChek MERS and sequencing assays. Assay performance, including limit of detection and precision, was evaluated in vitro by using MERS-CoV RNA transcripts. Analytical specificity was evaluated with a diverse collection of 16 respiratory virus–positive clinical specimens and 14 respiratory bacterial isolates.

Results: The 95% limits of detection of the PowerChek MERS assay for the upE and the open reading frame (ORF)1a were 16.2 copies/μL and 8.2 copies/μL, respectively. No cross-reactivity was observed. The diagnostic sensitivity and specificity of the PowerChek MERS assay were both 100% (95% confidence interval, 91.1–100%).

Conclusions: The PowerChek MERS assay is a straightforward and accurate assay for detecting MERS-CoV RNA. The assay will be a useful tool for the rapid diagnosis of MERS and could prove especially important for MERS outbreak control.

Key Words: MERS-CoV, Real-time reverse-transcription PCR, Performance

INTRODUCTION

For effective control of potential outbreaks of Middle East respiratory syndrome (MERS), highly sensitive and specific laboratory diagnostic tests are needed. Nucleic acid amplification assays are the method most widely used to provide laboratory confirmation of MERS [1]. MERS cases are routinely confirmed based on the detection of unique MERS coronavirus (MERS-CoV) RNA sequences by real-time reverse-transcription (rRT)-PCR, with confirmation by nucleic acid sequencing when necessary [2]. The WHO recently published updated interim recommendations for MERS-CoV laboratory testing and recommends an rRT-PCR assay targeting the envelope (upE) gene for screening, followed by confirmation with an assay for the re-
regions encoding either the open reading frame (ORF)1a, ORF1b, or the nucleocapsid protein (N). For sequencing, two target sites on the MERS-CoV genome, the RNA-dependent RNA polymerase (RdRp) and N genes, are suggested [2-4]. Although commercial rRT-PCR assays for MERS-CoV detection have been developed by several manufacturers, including Altona Diagnostics, Fast Track Diagnostics, and Primerdesign Ltd., studies validating diagnostic performance using authentic specimens are lacking [5, 6].

We evaluated the analytical performance and clinical applicability of the PowerChek MERS (upE & ORF1a) real-time PCR Kit (PowerChek MERS assay; Kogene Biotech, Seoul, Korea). The PowerChek MERS assay is a recently developed one-step rRT-PCR assay for the qualitative detection of MERS-CoV-specific RNA that has received Conformity Europe-In Vitro Diagnostic (CE-IVD) and the Ministry of Food and Drug Safety (MFDS) of Korea approval. Probes specific for the upE, ORF1a, and internal control regions are labeled with the carboxy-fluorescein (FAM), 6-carboxy-4',5'-dichloro-2',7'-dimethoxy-fluorescein (JOE), and indodicarbocyanine (Cy5) fluorophores, respectively. This assay has been previously evaluated with only small number of clinical specimens [6]. This report presents the results of a performance evaluation of the PowerChek MERS assay using clinical respiratory specimens.

**METHODS**

1. **Clinical specimens**

The Institutional Review Board of the Samsung Medical Center, Seoul, Korea, approved this study. For analysis, 100 clinical respiratory specimens (90 sputa and 10 nasopharyngeal swabs) were collected from 100 different individuals from June to July 2015. Fifty specimens were obtained from symptomatic MERS-positive patients, and the remaining 50 were obtained from asymptomatic MERS-negative healthcare workers who were under active monitoring.

Total nucleic acid was extracted by using the QIAamp DSP Viral RNA Mini kit (Qiagen, Hilden, Germany), according to the manufacturer’s instructions. A total of 140 μL of specimen was used, and the RNA was eluted in 50 μL and stored at −70°C until testing with the PowerChek MERS and sequencing assays.

2. **PowerChek MERS (upE and ORF1a) real-time PCR, RdRp, and N gene sequencing assays**

MERS-CoV RNA was detected by using the PowerChek MERS assay for amplification of upE and ORF1a. The primers and probes were used according to the previous study [3]. All rRT-PCR reactions were performed by using the 7500 Fast Real-Time PCR System (Applied Biosystems, Foster City, CA, USA). The PCR reaction was performed in a total volume of 20 μL (15 μL PCR reaction mixture and 5 μL template RNA). Thermocycling conditions were as follows: 50°C for 30 min, followed by 95°C for 10 min, and then 40 cycles of 15 sec at 95°C and 60 sec at 60°C. Viral template–positive controls and no-template controls were included in each run. The glyceraldehyde-3-phosphate dehydrogenase (GAPDH) gene was amplified simultaneously as a heterologous endogenous internal control to monitor PCR inhibition and specimen quality. A positive test result was defined as a well-defined exponential fluorescence curve that crossed the threshold ≤37 cycles, ≤38 cycles, and ≤35 cycles for the upE, ORF1a, and GAPDH genes, respectively.

Nested RT-PCR and sequencing targeting the RdRp and N genes were performed for confirmation, according to the previous protocols [4]. Briefly, the first round of RT-PCR for the two genes was performed by using the primer sets described in the previous report. A second round was conducted if no product was visible by agarose gel electrophoresis after the first round [4]. PCR products were sequenced by using a 3730 DNA analyzer (Applied Biosystems).

3. **Analytical performance of the PowerChek MERS (upE and ORF1a) real-time PCR Kit**

MERS-CoV RNA transcripts were synthesized in vitro from the T7 promoter, as previously reported [7] and were used for analytical performance evaluation. The limit of detection (LOD), the point at which 95% of the replicates of a given viral load are detected, was determined by using in vitro RNA transcripts diluted in nuclease-free water. Serial dilutions were analyzed, with 20 replicates per dilution. Probit analysis was used to determine the 95% LOD. Reproducibility, repeatability, and lot-to-lot precision were determined by using three concentrations (50, 150, and 300 copies/μL). Reproducibility was calculated by testing duplicates of each concentration in a single run for five days at two different test sites. Repeatability was calculated by testing duplicates of each concentration in a single run for 20 days. Lot-to-lot precision was accessed by testing duplicates of each concentration in a single run for five days using three different lots. Assay specificity was evaluated with nucleic acids obtained from a diverse collection of 16 respiratory virus-positive clinical specimens and 14 other respiratory bacterial isolates (13 reference strains and one clinical isolate; Table 1).

https://doi.org/10.3343/alm.2017.37.6.494

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**Table 1. Analytical specificity of the PowerChek MERS (upE and ORF1a) real-time PCR Kit**

| Microorganism                  | Source                      | PowerChek MERS assay |
|--------------------------------|-----------------------------|----------------------|
| **Viruses**                    |                             |                      |
| Adenovirus                     | Clinical specimen*          | Negative             |
| Bovivirus                      | Clinical specimen*          | Negative             |
| Coronavirus 229E               | Clinical specimen*          | Negative             |
| Coronavirus OC43               | Clinical specimen*          | Negative             |
| Coronavirus HKU1               | Clinical specimen*          | Negative             |
| Coronavirus NL63               | Clinical specimen*          | Negative             |
| Human enterovirus              | Clinical specimen*          | Negative             |
| Human metapneumovirus          | Clinical specimen*          | Negative             |
| Human rhinovirus               | Clinical specimen*          | Negative             |
| Influenza A                    | Clinical specimen*          | Negative             |
| Influenza B                    | Clinical specimen*          | Negative             |
| Parainfluenza 1                | Clinical specimen*          | Negative             |
| Parainfluenza 2                | Clinical specimen*          | Negative             |
| Parainfluenza 3                | Clinical specimen*          | Negative             |
| Respiratory syncytial virus A  | Clinical specimen*          | Negative             |
| Respiratory syncytial virus B  | Clinical specimen*          | Negative             |
| **Bacteria**                   |                             |                      |
| Acinetobacter baumannii        | Reference strain (ATCC 19606) | Negative             |
| Escherichia coli               | Reference strain (ATCC 25922) | Negative             |
| Haemophilus influenza          | Reference strain (ATCC 9007) | Negative             |
| Klebsiella pneumonia           | Reference strain (ATCC 900603) | Negative          |
| Legionella pneumophila         | Reference strain (ATCC 33156) | Negative             |
| Mycoplasma pneumoniae          | Clinical specimen*          | Negative             |
| Mycobacterium tuberculosis     | Reference strain (ATCC 27294) | Negative             |
| Mycobacterium avium           | Reference strain (KMRC 00136-41011) | Negative |
| Mycobacterium abscessus        | Reference strain (ATCC 19977) | Negative             |
| Pseudomonas aeruginosa         | Reference strain (ATCC 27853) | Negative             |
| Staphylococcus aureus          | Reference strain (ATCC 25923) | Negative             |
| Staphylococcus epidermidis     | Reference strain (ATCC 12228) | Negative             |
| Streptococcus pneumoniae       | Reference strain (ATCC 49619) | Negative             |
| Streptococcus pyogenes         | Reference strain (ATCC 19615) | Negative             |

*Specimens that yielded positive results in two respiratory virus panel assays: the AdvanSure RV real-time PCR Kit (LG chen, Seoul, Korea) and the Allplex Respiratory Panel 1, 2, and 3 (Seegene, Seoul, Korea). *Specimens confirmed by sequencing analysis. Abbreviation: KMRC, Korea Mycobacterium Resource Center.

4. Diagnostic sensitivity and specificity of the PowerChek MERS (upE and ORF1a) real-time PCR Kit

The diagnostic performance of the PowerChek MERS assay and the RdRp and N gene sequencing assay was compared by measuring the positive percent agreement (PPA) and negative percent agreement (NPA) values and kappa coefficients. The RdRp and N gene sequencing assay was considered the reference standard method, and the diagnostic sensitivity and specificity of the PowerChek MERS assay were calculated.

5. Statistical analysis

Statistical analyses were performed by using the VassarStats website (http://vassarstats.net/) and the SPSS software, version 23.0 (SPSS Inc., Chicago, IL, USA). A P value <0.05 was considered statistically significant.

**RESULTS**

Table 2 shows the analytical sensitivity of the PowerChek MERS assay. The 95% LOD of the upE and ORF1a regions was 16.2 copies/μL (81 copies/reaction) and 8.2 copies/μL (41 copies/reaction), respectively. Table 3 shows the precision results for the PowerChek MERS assay. In the reproducibility study, the CV ranged from 0.54 to 2.10%. The assay showed acceptable repeatability, with CV results ranging from 0.54 to 2.11%. The lot-to-lot variation was acceptably low for all concentrations (CV range: 0.49–2.1%). No false-positive test results were obtained for any of these viral or bacterial isolates, but positive controls were readily detected (Table 1).

A comparison of the results of the PowerChek MERS and RdRp and N gene sequencing assays is shown in Table 4. Both methods gave positive results for all 50 clinical specimens obtained from the symptomatic MERS-positive patients. For all 50 specimens, the PowerChek MERS assay yielded positive reactions for both the upE and ORF1a targets. The range of Ct values for the upE, ORF1a, and GAPDH targets was 17.69–32.63, 19.22–34.27, and 18.51–31.17, respectively. Both methods gave negative results for all 50 specimens obtained from asymptomatic MERS-negative individuals. The two assays produced concordant results for all 100 specimens, with a kappa value of 1.00. The diagnostic sensitivity and specificity of the
PowerChek MERS assay were both 100% (95% confidence interval, 91.1–100%).

**DISCUSSION**

To date, various reports have been published on the efficacy of rRT-PCR assays for detection of MERS-CoV-specific RNA that target various regions, including the upE, ORF1a, ORF1b, and N genes [3-5, 7-9]. The interim recommendations published by the WHO suggest that the upE assay be used for screening, followed by confirmation with one of the ORF1a, ORF1b, or N gene assays [2]. Similarly, the US Centers for Disease Control and Prevention (CDC) developed and use an rRT-PCR assay composed of two independent PCR reactions; one assay (N2) combined with upE testing for screening, and a second assay (N3) for confirmation of positive results [7, 9]. Previous evaluations of the rRT-PCR assay used mock specimens spiked with cultured MERS-CoV or a small number of clinical specimens from MERS-infected patients [3-5, 7, 8, 10, 11]; this study used a total of 50 authentic clinical specimens from MERS-positive patients with various viral loads.

The PowerChek MERS assay could prevent misjudgment from sampling error, since it uses an endogenous internal control. The use of endogenous internal control would be helpful than that of artificial targets, since the endogenous internal control enables monitoring of not only the RNA extraction and reverse transcription steps but specimen quality as well [12, 13]. Monitoring specimen quality is important because assay performance for detection of MERS-CoV is dependent on specimen quality [14].

The PowerChek MERS assay is both simple and efficient, reducing the workload required for diagnosis. For rRT-PCR, the assay requires mixing of only the premix and RNA, which reduces the hands-on time and the risk of technical errors [5, 15]. The turnaround time from RNA extraction to final result is 3 hr. This rapid and simple testing technique would be especially useful in managing a large MERS outbreak.

Our study has several limitations. First, the PowerChek MERS assay was validated with only upper and lower respiratory specimens. Furthermore, the 50 clinical specimens from MERS-confirmed patients were all sputum specimens. Although lower respiratory specimens are the preferred specimen type for initial
MERS diagnosis because of their higher MERS-CoV load, rRT-PCR testing using other specimen types, including serum, urine, and stool specimens, is required in MERS-confirmed cases [2, 16, 17]. Thus, further study to evaluate performance using other specimen types is needed. Second, we did not compare assay performance across different real-time PCR platforms. The PowerChek MERS assay was developed to allow flexibility in the choice of real-time PCR platforms, and can be used with the 7500 Fast Real-Time PCR System, CFX96 real-time PCR detection system (Bio-Rad, Hercules, CA, USA), LightCycler 480 (Roche, Mannheim, Germany), Rotor-gene Q (Qiagen, Hilden, Germany), or Mx3005P thermocycler (Agilent Technologies, CA, USA); however, we evaluated the performance using only the 7500 Fast Real-Time PCR system.

In conclusion, the PowerChek MERS assay has good diagnostic performance for detecting MERS-CoV RNA. This straightforward and accurate assay is a useful tool for the rapid diagnosis of MERS and could be especially important in control of a MERS outbreak.

Authors’ Disclosures of Potential Conflicts of Interest

No potential conflicts of interest relevant to this article were reported.

Acknowledgments

This research was supported by a grant (grant number: HD15A3232) from the Korea Centers for Disease Control and Prevention.

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