Quantitative Assay of Hepatitis C Virus RNA Using an Automated Extraction System for Specific Capture with Probes and Paramagnetic Particle Separation

Hayato Miyachi,1* Atsuko Masukawa,2 Satomi Asai,1 Toshiaki Miura,3 Shigeru Tamatsukuri,3 Toru Hirose,3 and Yasuhiko Ando1

Department of Laboratory Medicine, Tokai University School of Medicine,1 and Clinical Laboratory, Tokai University Hospital,2 Bouseidai, Isehara, Kanagawa 259-1193, Japan. Phone: (81-463) 1121. Fax: (81-463) 8607. E-mail: miyachi@is.icc.u-tokai.ac.jp.

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A commercially available automated specimen preparation instrument for specific probe capture and paramagnetic separation has been developed (AmpliCap/GT-12; Roche Molecular Systems). We evaluated assay performance of the AmpliCap/GT-12 in the quantitative assay for hepatitis C virus (HCV) RNA with the AMPLICOR HCV MONITOR Test (version 2.0). Assay linearity using serial dilutions from a serum panel was observed in the range of 500 to 850,000 IU/ml, with a slightly compromised slope in the higher viral titers. The overall within-run and between-run reproducibility of the entire detection process for 3 and 5 log10 (IU/ml) of HCV RNA in samples had a standard deviation of <0.2, which was comparable to a manual method based on organic extraction and isopropanol precipitation (Roche Molecular Systems). Comparison of the test results with those obtained by the manual method showed a good correlation (R2 = 0.972, n = 86). Using heparin (3, 6.5, and 13 U/ml), dextran sulfate (0.1, 1, and 5 mM), hemoglobin (1.13, 2.25, and 4.5 g/liter), conjugated or unconjugated bilirubin (7.5, 15, and 30 mg/dl), and ATP (1.25, 2.5, and 5.0 mM) as known inhibitors, inhibition was only detected at a dextran sulfate concentration of 1 mM with the manual method but not with the AmpliCap/GT-12 extraction. In summary, the AmpliCap/GT-12 system was shown to permit a stable extraction process and accurate results for the quantitative assay of HCV RNA, successfully eliminating the inhibitory effect of dextran sulfate. This automated extraction system provides reliable and reproducible test results and saves labor; thus, it is suitable for routine diagnostic PCR.
MATERIALS AND METHODS

Clinical specimens. Serum specimens used in this study were obtained from 90 patients referred to Tokai University Hospital for chronic liver diseases. When needed, HCV RNA was quantitatively measured by the AMPLICOR HCV MONITOR Test, version 2.0 (Roche Diagnostic Systems) (7). All samples were separated from clots within 4 h of collection, divided into aliquots, and stored at −80°C until RNAs were extracted.

RNA extraction. HCV RNA was isolated from serum by an automated system consisting of the reagents (AmpliCap; Roche Molecular Systems) and a robotic processor (GT-12; Roche Molecular Systems) developed in cooperation with Precision System Science Co. Ltd. (Tokyo, Japan). Briefly, HCV RNA was isolated from 250 μl of serum by lysis of virus particles with 500 μl of guanidinium thiocyanate solution at 60°C for 20 min. The RNA was hybridized with biotinylated probes (KY78) that were specific to the 5′-untranslated region of the HCV genome (16) and identical to the downstream primer for amplification. The hybridized RNA was then captured with streptavidin-coated paramagnetic particles. The quantitation standard or an internal quantitative control was introduced into the specimen during the lysis reaction. After washing of the hybrid-particle complexes to remove nonspecifically bound materials, the particles were resuspended in 250 μl of a specimen diluent and then were ready for amplification and detection by PCR.

RNA was also isolated from 100 μl of serum by a manual method based on guanidinium thiocyanate lysis and isopropanol precipitation (Roche Molecular Systems). Briefly, 400 μl of a lysis solution containing guanidinium thiocyanate and the quantitation standard was added to the specimen. The nucleic acid extraction was precipitated by isopropanol and pelleted by centrifugation, and the pellet was washed with ethanol. The nucleic acid was resuspended in 1,000 μl of a specimen diluent, and 50 μl of extracted sample was used for amplification and detection by PCR.

The AmpliCap/GT-12 used a larger input volume of serum than the manual method; the former used 50 μl out of 250 μl of a specimen diluent derived from 250 μl of serum, and the latter used 50 μl out of 1,000 μl of a specimen diluent derived from 100 μl of serum.

COBAS AMPLICOR HCV Test. The quantitative COBAS AMPLICOR HCV Tests, and AMPLICOR HCV MONITOR Test, version 2.0, have been previously described (7, 26). All samples were extracted once, and a single amplification and detection were performed on each extract except as otherwise stated. One positive and two negative controls provided with the kit were run with each batch of patient specimens.

Sample dilutions. A serum panel from clinical specimens that was quantitated for HCV RNA levels by the AMPLICOR HCV MONITOR Test, version 2.0, was diluted serially in HCV-seronegative normal human serum. Serial dilutions of serum were prepared to provide values throughout an expected range of the HCV genome (16) and identical to the downstream primer for amplification. The quantitation standard or an internal quantitative control was introduced into the specimen during the lysis reaction. After washing of the hybrid-particle complexes to remove nonspecifically bound materials, the particles were resuspended into the specimen during the lysis reaction. After washing of the hybrid-particle complexes to remove nonspecifically bound materials, the particles were resuspended in 250 μl of serum and were then ready for amplification and detection by PCR.

DATA SHOWED THAT THE ASSAY WAS ALMOST LINEAR OVER THE RANGE OF 500 TO 850,000 IU/ml OF HCV RNA. THE LINEARITY BECAME SLIGHTLY COMPROMISED AT HIGHER TITERS OF HCV RNA, I.E., >500,000 IU/ml (Fig. 1).

RESULTS

Linearity study. A serum panel diluted serially in HCV seronegative normal human serum was isolated by the AmpliCap/GT-12 assay, and then HCV RNA was measured in duplicate by the COBAS AMPLICOR HCV MONITOR Test, version 2.0. The dilutions were evaluated in a plot of the mean of duplicate observed values versus the expected values. These data showed that the assay was almost linear over the range of 500 to 850,000 IU/ml of HCV RNA. The linearity becomes slightly compromised at higher titers of HCV RNA, i.e., >500,000 IU/ml (Fig. 1).

Within-run and between-run reproducibility. Within-run and between-run reproducibilities were evaluated by using 3 and 5 log\(_{10}\) (IU/ml) of HCV RNA in samples. For within-run reproducibility, 10-fold determinations were made on three lots of reagents of AMPLICOR HCV MONITOR Test, version 2.0. The standard deviations (SDs) of the log10 (units per milliliter) were calculated and plotted against the average log10 (units per milliliter) for each sample. Serum HCV RNA levels, tested 10 times in the same round for within-run reproducibility, showed that the SDs were 0.12 to 0.17 and 0.06 to 0.11 for 3 and 5 log10 (IU/ml) of HCV RNA in samples, respectively, and the coefficients of variation were 3.3 to 4.9% and 1.4 to 2.3%, respectively, which were comparable to those obtained in the manual method (Table 1). Because a separate sample aliquot was processed for each test performed, these variations represent the total variability of the assay.

Serum HCV RNA levels, tested on one lot of reagents over 12 days for between-run reproducibility, showed that the SD of the log10 of HCV RNA levels for 3 and 5 log10 (IU/ml) of HCV RNA in samples was 0.11 and 0.16, respectively, and the coefficients of variation were 3.1 and 3.2%, respectively, which were comparable to those obtained in the manual method (Table 1).

Comparison with the manual method. A total of 90 clinical samples were subjected to RNA extraction by either the manual method or the automated method, and measured in singlet using the version 2.0 AMPLICOR HCV MONITOR tests. Of the samples, 86 had levels of HCV RNA that quantified both assay. The overall correlation of the two assays was good (γ = 0.8877x + 0.477; \(r^2 = 0.972\)) (Fig. 2).

Interference study. Using heparin (3, 6.5, and 13 U/ml), dextran sulfate (0.1, 1, and 5 mM), hemoglobin (1.13, 2.25, and 4.5 g/liter), conjugated or unconjugated bilirubin (7.5, 15, and 30 mg/dl), and ATP (1.25, 2.5, and 5.0 mM) as known inhibi-
the log10 (units per milliliter) were calculated and plotted against the average reagents of the AMPLICOR HCV MONITOR Tests, version 2.0. The SDs of three lots of reagents of the AMPLICOR HCV MONITOR Test, version 2.0 system and tested 10 times in the same round for within-run reproducibility on log10 (units per milliliter) for each sample. CV, coefficient of variation.

In this study, we evaluated the performance of the AmpliCap/GT-12 system for the quantitative assay of HCV RNA by the COBAS AMPLICOR HCV MONITOR Test, version 2.0. Comparison of the test results with those obtained by the manual method or AmpliCap/GT-12, and measured in singlet using the conventional manual method showed that the overall correlation of the two assays was good (r² = 0.972). Assay linearity was observed in the 3-log dynamic range over HCV RNA titers of 500 to 850,000 IU/ml but with a slightly compromised slope at higher titers (i.e., >500,000 IU/ml) in comparison with that obtained by the manual method based on organic extraction and pre-
W. Kuhl. Letter, BioTechniques 9:166, 1990), who have a high prevalence of HCV infection (21). In contrast to the qualitative assay, there was a lack of inhibition in the quantitative assay of HCV RNA, when RNA was extracted by the manual method from sera in the presence of heparin. This can be explained by the impact of different preparation processes for nucleic acid amplification and detection used in the qualitative and quantitative assays of HCV RNA. In the latter, nucleic acid extracted from 100 μl of serum is diluted in 1,000 μl of a specimen diluent by 10-fold, and thus the final concentration of residual heparin could be diluted to the extent that the inhibitory effect was negligible. This is supported by our previous report that 10-fold dilution of the serum sample was enough to eliminate the inhibitory effect in the former (15). There was also a lack of inhibition when RNA was extracted by the AmpliCap/GT-12 system from sera in the presence of heparin, in spite of using a larger input volume of serum than the manual method.

One of the major advantages of automating the extraction is the ability to provide a stable extraction process among samples. Using 3 and 5 log10 (IU/ml) for within-run and between-run precision of the combination of the AmpliCap/GT-12 system and the COBAS AMPLICOR HCV MONITOR Test, the SDs were <0.2. Considering that these data are comparable to the reproducibility of the COBAS AMPLICOR HCV MONITOR Test (SD < 0.2 [7, 26]), the AmpliCap/GT-12 system has permitted a stable extraction process among samples. The system is characterized by automated separation of nucleic acid for amplification by target-specific probe capture, specimen preparation with high throughput (96 reactions in 2.5 h), and minimal hands-on reagent preparation and machine setup. The ability to automate the extraction in addition to the measurement improved the assay performance in terms of not only stability of the extraction process but also labor efficiency. The reliability of the GT-12 system has been recognized, as shown by its use in major reference laboratories and in blood screening for the Red Cross Center in Japan (9).

In summary, the AmpliCap/GT-12 system, based on the specific capture with probes and magnetic B/F separation, was shown to permit a stable extraction process and accurate results for a quantitative assay of HCV RNA, successfully eliminating the inhibitory effect of dextran sulfate from serum. This system provides reliable and reproducible test results and saves labor; thus, it is suitable for routine diagnostic PCR.

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REFERENCES