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J. Clin. Microbiol. 2011, 49(11):3842. DOI:
10.1128/JCM.05316-11.
Published Ahead of Print 21 September 2011.

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Evaluation of *Aspergillus* PCR Protocols for Testing Serum Specimens^{∇†}

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Received 10 August 2011/Returned for modification 30 August 2011/Accepted 9 September 2011

A panel of human serum samples spiked with various amounts of *Aspergillus fumigatus* genomic DNA was distributed to 23 centers within the European *Aspergillus* PCR Initiative to determine analytical performance of PCR. Information regarding specific methodological components and PCR performance was requested. The information provided was made anonymous, and meta-regression analysis was performed to determine any procedural factors that significantly altered PCR performance. Ninety-seven percent of protocols were able to detect a threshold of 10 genomes/ml on at least one occasion, with 83% of protocols reproducibly detecting this concentration. Sensitivity and specificity were 86.1% and 93.6%, respectively. Positive associations between sensitivity and the use of larger sample volumes, an internal control PCR, and PCR targeting the internal transcribed spacer (ITS) region were shown. Negative associations between sensitivity and the use of larger elution volumes ($\geq 100 \mu\text{l}$) and PCR targeting the mitochondrial genes were demonstrated. Most *Aspergillus* PCR protocols used to test serum generate satisfactory analytical performance. Testing serum requires less standardization, and the specific recommendations shown in this article will only improve performance.

PCR testing can be used as a screening tool to exclude invasive aspergillosis (IA). High-frequency sampling is required, favoring easily obtainable specimens such as blood. In contrast, invasive specimens, such as those from bronchoalveolar lavage (BAL) or tissue, are used to confirm the presence of IA once disease is already suspected.

The European *Aspergillus* PCR Initiative (EAPCRI) reported on the process of standardizing molecular methods for the detection of *Aspergillus* in whole blood (WB) and highlighted the importance of the nucleic acid extraction protocol in achieving satisfactory analytical sensitivity (11). Compared to alternative specimens (serum and plasma), WB processing is technically demanding, requiring greater standardization. Many centers routinely test other specimens, and it is important to evaluate the processes in use and provide a standardized approach for *Aspergillus* PCR testing of these specimen types. *Aspergillus* PCR assays have been used to successfully detect DNA in serum. Although the methodology varies, the pooled sensitivity and specificity of serum PCR were 72% and 96.5%, respectively (2, 3, 10, 12, 14).

This article describes the EAPCRI process to evaluate the analytical performance of methods currently in use for the detection of *Aspergillus* DNA in serum through the blinded

distribution of a simulated serum panel to 23 centers in Europe, the United States, and Australia for molecular testing. Statistical analysis of the results returned by each participant was performed anonymously to determine factors that affected analytical performance with an emphasis on analytical sensitivity to maintain the optimal negative predictive value required for a screening assay.

MATERIALS AND METHODS

The study followed the STARD guidelines where applicable (1).

Participants. The participants comprised a core of eight laboratories and an extended group of a further 15 laboratories. To maintain impartiality throughout the analytical process, all centers were designated by a numerical code to allow blinded review of individual methodological procedures, determination of performance, and statistical analysis.

DNA source material. Conidia were harvested from a sporulating culture of *Aspergillus fumigatus* (ATCC strain 1022) using a wet microbiological loop and resuspended in sterile water containing a drop of Tween 20 to prevent clumping. Conidia were serially diluted in sterile water and quantified using a Fuchs-Rosenthal counting chamber. A volume of suspension containing the equivalent to 1.2×10^6 conidia was harvested by centrifugation at $10,000 \times g$ for 5 min. The supernatant was discarded, and conidia were exposed to mechanical disruption using the equivalent of 20 μl of ceramic beads (Roche, Burgess Hill, United Kingdom) and 30 s of bead-beating using a minibead beater (Biospec Products). After pulse centrifugation, the beads were washed with 200 μl molecular-grade water and DNA was extracted using the Roche High Pure Template DNA kit as per the manufacturer's instructions with an elution volume of 100 μl . The DNA was serially diluted using molecular-grade Tris-EDTA buffer (Sigma-Aldrich, United Kingdom) to attain a range of DNA concentrations (Table 1).

Specimens. Blood was obtained by consent from healthy volunteers, screened for the presence of infectious agents as per the protocol of the Institute of Transfusional Medicine, Wuerzburg University (Wuerzburg, Germany), and pooled. The serum was tested for the presence of contaminating *Aspergillus* DNA before processing (13). After being divided into 30-ml aliquots, two aliquots were

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[∇] Published ahead of print on 21 September 2011.

[†] The authors have paid a fee to allow immediate free access to this article.

TABLE 1. Composition and corresponding positivity of the EAPCRI serum panel^a

Initial <i>A. fumigatus</i> DNA concn (genomes/ μ l) ^b	Concn (μ l) added to 30 ml serum	Serum burden (genomes/ml)	95% variability in 1-ml sample ^c	Positivity (no. positive/tested) ^d	Mean Cq (95% CI)
12,000	25	10,000	9,803–10,197	45/45	27.8 (27.0–28.6)
1,200	25	1,000	938–1,062	45/45	31.1 (30.3–31.9)
	12.5	500	456–544	45/45	32.7 (31.9–33.5)
120	25	100	80–120	44/45	34.7 (33.9–35.7)
	12.5	50	36–64	44/47	36.0 (35.2–36.9)
12	25	10	4–16	43/47	37.5 (36.7–38.3)
	12.5	5	\leq 1–9	38/46	38.1 (37.2–39.0)
1.2	25	1	0–3	12/47	38.8 (37.1–40.5)
0	0	0	0	6/94	37.1 (35.3–38.8)

^a Each panel was composed of one aliquot of each fungal burden and two negative control serum samples.

^b DNA extracted from 1.2×10^6 *A. fumigatus* conidia and serially diluted to attain desired concentrations assuming one conidium contains a single genome and 100% DNA extraction efficiency when initially extracting DNA. Initial DNA eluted in 100 μ l.

^c Shown is the 95% confidence interval calculated on the basis of taking a 1-ml specimen from an initial 30-ml volume where the mean concentration will be as stated in the third column and assuming a Gaussian distribution.

^d Positivity defined as number of positive results per number of PCR replicates for each burden. Excluded are results from contaminated methods 10 and 21 and from unsuitable method 11 (Table 2).

retained to provide negative-control material, while eight of the aliquots were spiked with various concentrations of *A. fumigatus* DNA (Table 1). The 30-ml serum batches were further divided into 1-ml aliquots. To avoid airborne contamination, all processing of material took place in a category 2 laminar flow cabinet.

Panels. Each panel consisted of 10 1-ml serum samples (eight containing *A. fumigatus* DNA and two negative) and a PCR control consisting of *A. fumigatus* DNA equivalent to 10 genomes/ μ l to monitor PCR performance. The expediency of the panel was tested by an “in-house” *Aspergillus*-specific real-time PCR (13) and by the commercially available MycAssay Aspergillus (Myconostica, Manchester, United Kingdom). After developing the panels, the distributing center froze all panels at -80°C with no freeze-thaw intervals. To validate panels, initial blind testing was performed by the core group ($n = 8$) before distribution and blind testing by additional centers ($n = 15$). The panels were circulated on dry ice for next day delivery by courier. Participating centers were asked to confirm receipt, comment on the state of the panel (frozen or thawed), and keep specimens frozen at -80°C until testing. Centers were requested to return results within a designated time frame and provide detailed protocols for their DNA extraction and PCR amplification systems. The information required included the sample volume used, extraction method, DNA elution volume, PCR method, PCR target, PCR template input volume, PCR total reaction volume, PCR amplification platform, and internal control PCR results.

The acceptable threshold for the PCR performance panel was equivalent to 10 genomes/ml serum. It was essential that all methods evaluated were able to attain this level of detection. Methods not performing to this standard were considered suboptimal. Two concentrations below this threshold were included (5 and 1 genome/ml serum), but the lower 95% confidence interval (95% CI) for the possible concentration in each 1-ml aliquot was below 1 genome per 1-ml sample at these concentrations (Table 1).

DNA extraction. All participants were encouraged to use their own methodology. No recommendations were provided. A total of 15 different extraction systems were used (Table 2).

PCR amplification. Participating centers were asked to use their current amplification system. It was recommended that DNA extracts be tested in duplicate—preferably triplicate. The use of an internal control PCR was recommended. The use of nested-PCR assays was discouraged, as these were considered too prone to false-positive results and unlikely to attain widespread acceptance outside specialist molecular centers. Twenty-two combinations of PCR target, input volume, final reaction volume, and amplification platform were provided (Table 2).

Data analysis. A data analysis subgroup collated the results and technical information and determined covariates for blinded statistical analysis by an independent party (Table 3).

Statistical analysis. (i) **PCR positivity and genomic load.** PCR positivity was determined according to the reported real-time PCR quantification cycle (Cq

value), using a universal positivity threshold of 45 cycles, and PCR results with a Cq value of >45 cycles were considered negative. In doing so, continuous Cq data were changed into qualitative positive and negative results, and positivity at each fungal burden was determined, along with the binomial exact 95% confidence interval. The specificity was calculated using the equation $1 - \text{positivity rate for the serum samples with no fungal burden}$. In addition, a receiver operator curve (ROC) was constructed with PCR performance determined by a varying Cq threshold for positivity.

Mixed-model logistic regression analysis evaluated the probability of a positive result as a function of the genomic load using the PCR result as a binomial, positive or negative, outcome variable.

(ii) **Linear analysis.** The correlation between threshold cycle number (Cq) of real-time PCR and genomic load (\log_{10} transformed) was investigated by linear regression analysis. Samples with no fungal burden were excluded from the model, as data associated with a zero fungal burden would establish a discontinuity (genomes present/not present). To correctly evaluate the clustered nature of the data, a mixed (multilevel) random-coefficient model was used, grouping the various methods using the Stata command “xtmixed.” The basic model was Cq versus \log_{10} genomes/ml.

From the slope parameter, the mean reaction efficiency was calculated. The formula used was relative efficiency = $10^{-1/\text{slope}} - 1$, where the 100% efficiency corresponds to the doubling ($\times 2$) of the target sequence per PCR cycle. In this context, a slope of -3.32193 would correspond to a (full) relative efficiency of 100%.

The same basic model was adapted to the analysis of covariates (sample volume, elution volume, template volume, PCR volume, extraction procedure, PCR platform, DNA sequence target, use of internal control). Each covariate was added singly to the above-described model (Cq versus \log_{10} genomes/ml. A 4-level categorical variable was used for extraction evaluation, with the following levels: Roche, Qiagen, bioMérieux, and Promega. A similar coding criterion was used for the PCR platform (four levels: ABI, LightCycler, Rotorgene, and SmartCycler) and DNA sequence target (four levels: 28S, 18S, internal transcribed spacer [ITS], and mitochondria). A multivariate approach was performed using the covariates indicated above. After adjustment, the final model retained the independently significant covariates.

RESULTS

Twenty-two of the 23 centers requesting participation were able to return results, but one of these centers used a nested-PCR system, and the results were excluded from further analysis. A total of 29 different protocols (combined DNA extraction and PCR amplification) were provided by the remaining

TABLE 2. Methods used to test the EAPCRI serum panel^a

Method	Sample vol (ml)	Available target at threshold (95% CI) ^b	DNA extraction method	Elution vol (μl)	Internal control PCR used (yes/no)	PCR platform (target)	PCR vol (μl)		Positivity to threshold ^d /overall ^e (%)	False positivity (%) ^e
							Input	Final		
1	1	10 (4–16)	Roche High Pure LV	60	Yes	LC 2.0 (28S)	10	20	100/93.75	0
2	0.4	4 (1–7)	Qiagen EZ1 Virus 2.0	60	Yes	LC 2.0 (28S)	10	20	100/87.5	0
3	0.5	5 (2–8)	bioMérieux EasyMag	60	Yes	LC 2.0 (28S)	10	20	100/93.75	0
4	1	10 (4–16)	Roche MagNA Pure LV	50	Yes	LC 480 (28S)	25	50	100/93.75	0
5	1	10 (4–16)	Roche MagNA Pure LV	50	Yes	LC 480 (Mito)	5	20	100/87.5	0
6	1	10 (4–16)	Roche MagNA Pure LV	50	Yes	LC 480 (28S)	5	20	100/93.75	0
7	0.2	2 (0–5)	Qiagen Qiamp DNA	50	Yes	LC 480 (28S)	2–4	20	100/100	0
8	0.2	2 (0–5)	Roche MagNA Pure NA	50	No	LC 2.0 (18S)	10	20	92.3/70.6	0
9	0.2	2 (0–5)	Roche MagNA Pure LV	50	No	LC 2.0 (18S)	10	20	92.3/82.4	25
10	0.6	6 (3–9)	bioMérieux EasyMag	50	No	LC 2.0 (18S)	10	20	100/100	100
11	1	10 (4–16)	Cepheid GeneExpert	100	Yes	GeneExpert (18S)	25	50	50/37.5	0
12	1	10 (4–16)	Qiagen Ultrasens Virus	35	Yes	StepOnePlus (ITS2)	10	20	100/93.75	0
13	1	10 (4–16)	Qiagen Circulating NA kit	50	Yes	7500 FAST (ITS)	7	20	100/100	0
14	0.2	2 (0–5)	Qiagen EZ1 tissue	50	Yes	7000 (A. fumigatus 28S)	7.5	20	100/81.25	0
15	0.2	2 (0–5)	Qiagen EZ1 tissue	50	Yes	7000 (28S)	7.5	20	91.6/75	25
16	0.8	8 (5–11)	Qiagen Qiamp DNA kit	100	Yes	LC 1.2 (28S)	5	20	91.6/75	0
17	0.22	2 (0–5)	bioMérieux EasyMag	110	Yes	7900 FAST (28S)	10	30	100/81.25	0
18	0.5	5 (2–8)	bioMérieux EasyMag	110	Yes	7900 FAST (28S)	10	30	100/87.5	0
19	1	10 (4–16)	Promega Wizard Genomic DNA	50	Yes	Rotorgene 6000 (28S)	5	30	92.3/73.7	33.3
20	1	10 (4–16)	Qiagen QIASymphony	60	No	7500 FAST (ITS1)	5	25	100/87.5	0
21	0.4	4 (1–7)	Qiagen QIAmp DNA blood	60	Yes	Mx3000p (28S)	4	20	100/100	100
22	1	10 (4–16)	Roche MagNa Pure LV	50	Yes	LC 1.2 (18S)	10	20	100/100	0
23	0.4	4 (1–7)	Qiagen EZ1 Virus 2.0	60	Yes	SmartCycler (18S)	10	25	100/87.5	50
24	1	10 (4–16)	Promega Maxwell	30	Yes	7900 FAST (18S)	5	20	83.3/75	0
25	0.5	5 (2–8)	Roche High Pure Template DNA	100	Yes	LC 480 (28S)	20	50	100/100	0
26	1	10 (4–16)	Roche MagNA Pure Compact	50	Yes	7500 FAST (18S)	5	50	100/87.5	0
27	0.2	2 (0–5)	Qiagen QIAmp DNA kit	65	Yes	SmartCycler (18S)	10	25	100/87.5	50
28	0.4	4 (1–7)	Qiagen QIAmp DNA kit	65	Yes	SmartCycler (18S)	10	25	100/87.5	0
29	0.2	2 (0–5)	Roche MagNA Pure NA	100	No	9700 (MitoC)	5	20	100/87.5	0
Mean	0.62	6 (3–9)		62.9			9.3 ^f	26.6	92.7/86.5 ^g	13.5 ^g
Total					24/29					

^a Results for methods in boldface represent (i) procedures with less than 100% reproducibility at the designated threshold of 10 genomes/ml ($n = 5$ protocols: 8, 9, 11, 15, and 16), (ii) positivity rates below the overall positivity of 86.1% (95% CI, 82.1 to 89.5%) ($n = 8$ protocols: 8, 11, 14, 15, 16, 17, 19, and 24), or (iii) false-positivity rates ($1 - \text{specificity}$) that result in specificity being below the overall specificity of 93.6% (95% CI, 86.6 to 97.6%) ($n = 7$ protocols: 9, 10, 15, 19, 21, 23, and 27). Please note that despite reproducibly detecting the designating threshold, protocols 19 and 24 achieved 66.7% and 0% positivity, respectively, when testing the 50-genome/ml sample, resulting in <100% positivity to threshold. Please note that the overall sensitivity and specificity (stated above) were calculated excluding the data from methods 10, 11, and 21. LV, large volume; NA, nucleic acid; LC 2.0, Roche LightCycler 2.0; LC 480, Roche LightCycler 480; GeneExpert, Cepheid GeneExpert; StepOnePlus, ABI StepOnePlus; 7500 FAST, ABI 7500 FAST; 7000, ABI 7000; 7900 FAST, ABI 7900 FAST; Rotorgene 6000, Corbett Rotorgene 6000; LC 1.2, Roche LightCycler 1.2; Mx3000p, Stratagene Mx3000p; SmartCycler, Cepheid SmartCycler; 9700, ABI 9700; 28S, 28S rRNA gene; Mito, mitochondrial gene; 18S, 18S rRNA gene; ITS2, internal transcribed spacer 2; ITS, internal transcribed spacer; ITS1, Internal Transcribed Spacer 1; MitoC, mitochondrial cytochrome B.

^b The available burden and 95% CI when testing sample volumes less than 1 ml were determined assuming that the 1-ml aliquot provided contained the mid-point value of 10 genomes/ml.

^c Positivity to threshold represents the number of positive PCR results per test performed for each sample at, and above, the designated threshold of 10 genomes/ml (10,000, 1,000, 500, 100, 50, and 10 genomes/ml).

^d Overall positivity represents the number of positive PCR results per tests performed for the entire panel excluding the negative samples (10,000, 1,000, 500, 100, 50, 10, 5, and 1 genome/ml).

^e False positivity represents the number of positive PCR results per test performed for both negative samples.

^f Calculated using a mid-point value of 3 μl for method 7.

^g Includes results for all 29 protocols listed in the table.

21 centers (Table 2). Two centers contributed three protocols, four centers contributed two protocols, and 15 centers contributed a single protocol.

Aspergillus PCR performance when testing serum specimens. Of the 29 different protocols used to test the serum panel, 24 were able to reproducibly (100%) detect the designated threshold of 10 genomes/ml (Table 2). Despite detecting the designated threshold, protocols 19 and 24 achieved 66.7% and 0% positivity, respectively, when testing the 50-genome/ml sample. Protocols 8 and 9 were able to detect the threshold in two of three replicates, although protocol 9 was able to reproducibly detect the sample containing 5 genomes/ml. Protocols 15 and 16 detected the threshold in one of two replicates.

Protocol 11 was unable to detect the designated threshold; most samples were determined to be inhibitory or as extraction failures. Most protocols (21/24) attaining the threshold could also reproducibly detect the sample with 5 genomes/ml, although at this concentration protocols 14, 17, and 19 generated positivity rates of 50% and 33.3%, respectively. Eleven protocols (1, 3, 4, 6, 7, 10, 12, 13, 19, 21, and 25) were able to detect the 1-genome/ml sample. At this concentration, not every 1-ml aliquot will contain DNA (Table 1), and logistical regression analysis generated a probability of 63.21% that an individual 1-ml aliquot of the 1-genome/ml specimen would actually contain target, and this would be accentuated if volumes less than 1 ml were used for the initial DNA extraction. Protocols 10 and

TABLE 3. Covariates investigated by linear regression methods in this study

Covariate	Explanation
Sample vol.....	Vol of sample used for DNA extraction
DNA extraction system.....	Extraction platform used to extract DNA from sample—Roche, Qiagen, or combined other systems ^a
Elution vol.....	Vol used to elute DNA at the end of extraction
PCR amplification system.....	Amplification platform used to perform real-time PCR—Roche, Applied Biosystems, or combined other systems ^b
Template vol.....	Vol of DNA eluate used in each PCR
PCR vol.....	Final vol of each PCR
PCR target.....	Internal transcribed spacer target, 18S rRNA target, 28S rRNA target, or mitochondrial DNA target for <i>Aspergillus</i> PCR
Internal control.....	Internal control PCR used (yes/no)

^a bioMérieux EasyMag and Promega Wizard and Maxwell systems. Note one Qiagen system and one method using the bioMérieux EasyMag were excluded due to systemic contamination. The Cepheid GeneExpert was excluded on the basis of not being designed for use with serum.

^b Cepheid SmartCycler and Corbett Rotorgene 6000. Note one method using the Stratagene Mx3000p and one using the Roche Lightcycler 2.0 were excluded due to systemic contamination. The Cepheid GeneExpert was excluded on the basis of not being designed for use with serum.

21 generated a 100% (4/4) false-positivity rate with Cq values almost identical to those for the samples containing ≤10 genomes/ml, casting doubt on the validity of the “true”-positive results. Additional false positivity was noted for protocols 9, 15 (1/4), 19 (2/6), 23, and 27 (1/2).

In determining overall serum PCR performance, assays 10 and 21 (extensive false positivity) and 11 (commercial method designed to process WB and unsuitable for serum testing) were considered unsatisfactory, leaving 26 protocols for analysis. The overall sensitivity and specificity for *Aspergillus* PCR, including all burdens were 86.1% (95% CI, 82.1 to 89.5%) and 93.6% (95% CI, 86.6% to 97.6%), respectively. Positive and negative likelihood ratios and the diagnostic odds ratio (DOR) were 13.5, 0.15, and 90.9, respectively.

PCR performance and relationship with fungal burden. Predictably, there was a correlation between available fungal burden and PCR positivity (basic linear model: slope, -3.756; z value, -27.99; P < 0.0001). The mean PCR efficiency calculated from the slope coefficient of the regression line was 84.6% (95% CI, 77.4 to 93.3) (Fig. 1). Positivity rates and mean Cq values at the individual burdens are shown in Table 1 and

Fig. 1 and 2. One hundred percent positivity was attained when testing the sample containing the equivalent of 500 genomes/ml, although at 100, 50, and 10 genomes/ml, positivity remained above 90%, and the difference in positivity rates between the four samples was not statistically significant (Table 1). The positivity rate for detecting burdens at or greater than the predesignated detection limit of 10 genomes/ml was 97.1% (95% CI, 94.5 to 99.6%). At 5 and 1 genome/ml, the positivity rates were 82.6% and 25.5%, respectively. The latter result was significantly lower than the positivity rates for every other burden tested (difference between 5 and 1 genome/ml, 57.1%; 95% CI, 37.8 to 70.3; P = 0.0001). Logistical regression analysis determined that a sample containing a burden of 1.82 genomes/ml would be associated with a 50% positivity rate, and the positivity rate when testing samples containing 1 genome/ml would be 27.59% (Fig. 2), as predicted by the Poisson distribution for a mean of 1 genome/ml.

ROC analysis indicated that a Cq threshold of ≤43 cycles ensured a good diagnostic accuracy (DOR = 105) with a sensitivity and specificity of 85.6% and 94.7%, respectively, and using this threshold, 87.4% of samples were correctly classified

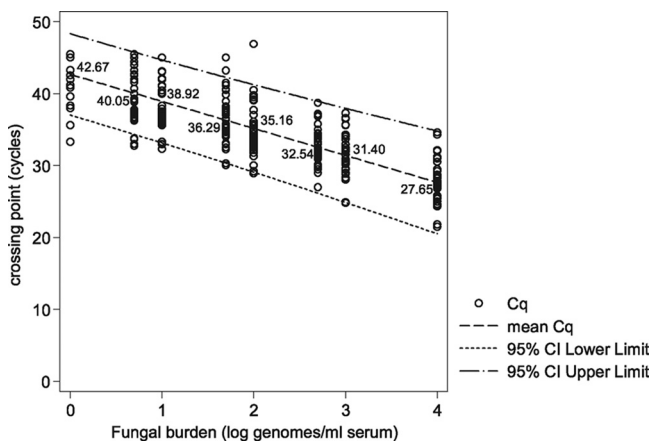


FIG. 1. Crossing point (Cq) values for *Aspergillus* PCR when testing DNA extracted from serum spiked with various fungal burdens as determined by linear regression. The linear equation for the mean Cq values is $y = -3.76 \log(x) + 42.67$, where $\log(x) = \log_{10}$ genomes/ml. The 95% CI for the slope was -3.49 to -4.02.

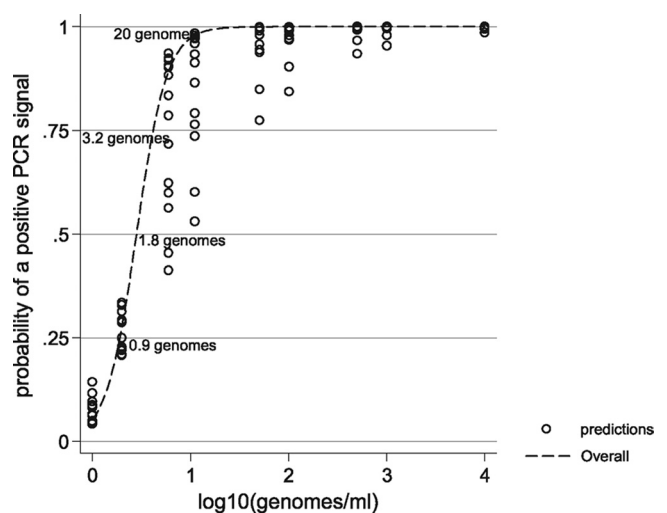


FIG. 2. Probability of PCR positivity at various fungal burdens as determined by logistic regression.

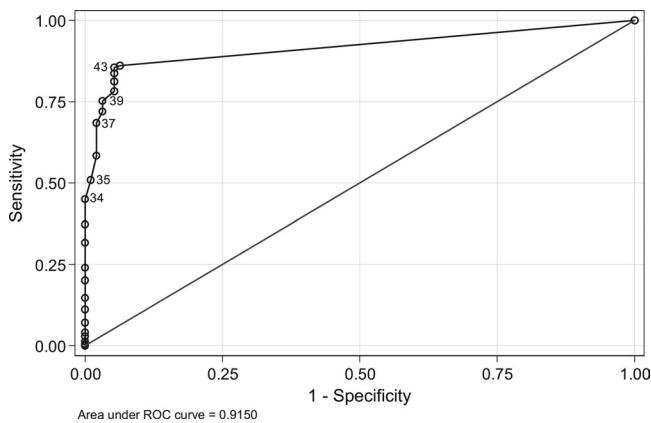


FIG. 3. Receiver operator curve for *Aspergillus* PCR methods when testing serum samples. PCR positivity was determined according to a variable Cq threshold. (Representative Cq thresholds are highlighted.) The area under the ROC was 0.9150 (95% CI, 0.89184 to 0.93806).

(Fig. 3). Using a Cq threshold of ≤ 34 cycles generated a specificity of 100%.

Linear regression analysis. Bivariate linear regression determined a statistically significant positive association between PCR sensitivity and larger sample volumes, use of an internal control, and targeting the ITS region for PCR amplification. Separating the PCR performance on the basis of sample volume used (≥ 0.5 ml or < 0.5 ml) generated sensitivities of 88.6% and 82.5% (difference, 6.1%; 95% CI, 14.9 to -1.0% ; $P = 0.1157$), respectively. When only low burdens (≤ 10 genomes/ml) were included, sensitivities of 76.9% and 54.7% for ≥ 0.5 -ml and < 0.5 -ml starting sample volumes, respectively, were generated. The difference in sensitivity at this clinically typical level was significant (difference, 21.8%; 95% CI, 5.6 to 37.1%; $P = 0.0091$).

A negative effect on the sensitivity of the PCR assay was detected for large elution volumes and for using mitochondrial targets for PCR (Table 4). When using elution volumes of ≥ 100 μ l, 40% of methods generated a positive PCR result, on at least one occasion, for burdens of ≤ 5 genomes/ml, compared to 69.0% for methods eluting DNA in < 100 - μ l volumes, the difference in positivity being 29.0% (95% CI, -3.4 to 55.1%; $P = 0.1424$). More significantly when detecting samples containing ≤ 5 genomes/ml, Cq values occurred earlier for DNA extraction methods eluting DNA in < 100 - μ l volumes. The mean Cq for < 100 - μ l elution volumes was 37.8 cycles, compared to 41.7 cycles for methods eluting DNA in volumes of ≥ 100 μ l (difference, 3.9 cycles; 95% CI, 1.9 to 5.8 cycles; $P = 0.0002$).

Multivariate analysis confirmed the negative association between sensitivity and elution volume ($P = 0.003$) and PCR amplifying the mitochondrial targets ($P = 0.011$).

DISCUSSION

The testing of serum is less technically demanding than the testing of WB (7), since lysis of red and white cells is not required. The perceived target in serum is free circulating DNA (DNAemia), released from the organism through the actions of the host's immune defenses, antifungal therapy, autolysis, and possibly exponential growth (6, 9). Any cell-associated DNA sources in serum will be limited, being trapped during blood clot formation

and/or removed during blood fractionation (12). If the target in serum is free DNA, then the extraction process is straightforward, although it is necessary to remove any serum components (proteins, electrolytes, waste products, drugs, etc.) that may interfere with the PCR amplification process.

Theoretically, targeting free circulating DNA through serum testing requires less standardization as commercial extraction methods can be employed. The commercial DNA extraction systems evaluated in the study provided comparable levels of performance. Nevertheless, each extraction system is directly associated with other variables such as sample input and elution volume, and it is essential to determine the analytical performance thresholds of any protocols utilizing commercial kits, highlighting methodologies that provide inadequate performance and should not be used in clinical practice. In this study, only five protocols (8, 9, 11, 15, and 16) failed to reproducibly detect the designated detection threshold of 10 genomes/ml serum (Table 2). Method 11 had been designed to test WB samples, providing excellent performance when testing this specimen type (11). Being fully automated, it could not be adjusted for serum testing; consequently, results were poor, and it should not be used for *Aspergillus* PCR testing of serum. Each protocol is constructed of many individual, yet linked, steps that will influence further steps downstream, and it is important to identify the critical step or steps that impact performance. For example, methods 8 and 9 use extraction platforms similar to methods 4 to 6 and 22, yet the latter methods provide better performance. It would be easy to conclude that the extraction platform provided inconsistent inter-laboratory performance. However, detailed examination of the protocols showed that methods 8 and 9 only extracted 0.2 ml of

TABLE 4. Bivariate linear regression analysis between Cq and the additional covariates, excluding \log_{10} genomes/ml^a

Covariate	z value	P value
Sample vol	-2.27	0.023
DNA extraction system		
Roche	-0.11	0.909
Qiagen	-1.15	0.250
Other	1.50	0.133
Elution vol	2.96	0.003
PCR amplification system		
Roche	-0.64	0.521
ABI	0.53	0.598
Other	0.25	0.803
Template vol	-1.07	0.286
PCR vol	0.97	0.334
PCR target		
18S rRNA gene	-0.28	0.779
ITS region	-2.48	0.013
28S rRNA gene	0.39	0.698
Mitochondrial	2.57	0.010
Internal control	-2.18	0.029

^a Significant associations are highlighted in boldface. Note that a negative z value predicts an earlier Cq, and thus a higher probability of PCR positivity, whereas a positive value has the opposite effect.

TABLE 5. Recommendations of the EAPCRI for *Aspergillus* PCR protocols when testing serum

Result	Recommendation
A positive association between sensitivity and initial sample vol was noted (z value, -2.27 ; $P = 0.023$). When detecting low fungal burdens (<10 genomes/ml), the difference in sensitivity using a starting volume of <0.5 ml serum compared to ≥ 0.5 ml is 21.8% (95% CI, 5.6 to 37.1; $P = 0.0091$).	The use of larger sample volumes (≥ 0.5 ml) will improve sensitivity. To enhance detection of low fungal burdens (<10 genomes/ml), a minimum of 0.5 ml starting material should be used.
Twenty-eight of 29 protocols evaluated were able to reach the designated threshold of detection on at least one occasion, generating a threshold positivity rate of 91.5%.	Most commercially available nucleic acid extraction systems described in Table 2 can be used to extract <i>Aspergillus</i> DNA using the protocols as described by the manufacturer. However, all kits should be screened for contamination and a limit of detection determined prior to clinical use.
Method 11 (Table 2) was specifically designed to test whole blood and was not able to efficiently process serum samples.	Protocols specifically designed for testing alternative specimen types (e.g., whole blood) should not be used.
A negative association between sensitivity and elution vol was noted (z value, 2.96 ; $P = 0.003$). When detecting low fungal burdens (<10 genomes/ml), the difference in Cq for methods eluting in <100 μ l (37.8 cycles) compared to ≥ 100 μ l (41.7 cycles) volumes was 3.9 cycles (95% CI, 1.9 to 5.8 cycles; $P = 0.0002$) and results in a 29.0% reduction in positivity.	The use of larger elution volumes (≥ 100 μ l) will reduce sensitivity. For optimal detection of low fungal burdens (<10 genomes/ml), nucleic acid should be eluted in vol of <100 μ l. Elution in vol of ≥ 100 μ l delays Cq values to the degree where real-time PCR is inconsistent and positivity will be affected.
A positive association between sensitivity and PCR targeting the ITS region was noted (z value, -2.48 ; $P = 0.013$).	The reaction kinetics for each PCR reaction will vary according to the design of specific oligonucleotides and the optimization of each particular PCR assay. Performance may not be directly associated with the target gene but exclusive to the individual assay itself. When designing a new assay, the MIQE guidelines (2a) should be followed.
A negative association between sensitivity and PCR targeting the mitochondrial regions was noted (z value, 2.57 ; $P = 0.010$).	The reaction kinetics for each PCR reaction will vary according to the design of specific oligonucleotides and the optimization of each particular PCR assay. Performance may not be directly associated with the target gene but exclusive to the individual assay itself. When designing a new assay, the MIQE guidelines (2a) should be followed.
A positive association between sensitivity and the use of an internal control PCR was noted (z value, -2.18 ; $P = 0.029$).	An internal control PCR should be used and incorporated at the start of the extraction procedure, monitoring both PCR inhibition and DNA extraction efficiency. Internal control Cq values should be representative of typical <i>Aspergillus</i> PCR-positive results (≥ 35 cycles), and human DNA targets should be avoided due to the possibility of variable target amounts within individual specimens generating varied Cq values.
Using a threshold of 43 cycles generates a DOR of 105, and 87.4% of results were correctly classified.	A PCR positivity threshold of 43 cycles provides the greatest degree of diagnostic accuracy.

the available specimen, compared to 1.0 ml for methods 4 to 6 and 22, limiting the amount of DNA target for extraction and negatively affecting final performance. In this study, a positive association between PCR positivity and the volume of starting material was shown.

False positivity was not a major issue, although two methods, 10 and 21, consistently generated false-positive results, suggesting a contamination issue. Both DNA extraction systems in these protocols had previously been associated with *Aspergillus* DNA contamination (4) (EAPCRI, group communication), but within this study, other centers had also used these systems without contamination, highlighting the transient, possibly batch-variable nature of this problem. Alternatively, the contamination issue may be associated with the individual center or other parts of these individual processes, and fungal DNA contamination of certain PCR reagents has been noted (5).

A PCR positivity threshold of 43 cycles should be used to provide the greatest degree of diagnostic accuracy, and at this threshold, 87.4% of results were correctly classified. This is in keeping with a previous report where the authors also con-

cluded that clinically the burden of DNA in serum was very low (fg/ μ l), to such a degree that positive PCR signals became inconsistent (8).

The overall sensitivities and specificities for *Aspergillus* PCR testing of serum were 86.1% and 93.6%, respectively. This compares favorably with the same performance parameters for *Aspergillus* PCR methods testing WB (11). When testing WB using protocols compliant with the EAPCRI recommendations provided to participants, the sensitivity and specificity were 88.7% and 91.6%, respectively. The probability of obtaining positive PCR results at specific fungal burdens is higher for serum than WB PCR, which is probably a consequence of the additional processing required when testing WB (results not shown). As the initial evaluation of existing serum PCR methods provides performance that is comparable to that of the agreed WB PCR protocol, it is not necessary to attempt to enhance performance through the evaluation of further serum panels with associated procedural recommendations. Indeed, our initial hypothesis that serum PCR methods would require less technical standardization appears to be correct.

Nevertheless, mixed-model linear analysis did highlight several important parameters that form the basis of the EAPCRI recommendations for *Aspergillus* PCR when testing serum samples, as highlighted in Table 5.

There were no significant associations between sensitivity and DNA template volume, PCR amplification platform, and final PCR volume, although PCR amplification of the ITS region was positively associated with sensitivity, and the opposite was true for the mitochondrial targets. This should be interpreted with caution as the reaction kinetics for each PCR will vary according to the design of specific oligonucleotides, and the optimization of each particular PCR assay and performance may not be directly associated with the target gene but exclusive to the individual assay.

To conclude, the testing of serum by *Aspergillus* PCR can be performed using commercial nucleic acid extraction methods, providing standardization and quality control. The EAPCRI previously showed PCR amplification not to be rate limiting, and commercial extraction in combination with most amplification methods will provide acceptable analytical performance(11). The use of serum is less technical than testing of WB, reducing both time to result reporting and labor, and permits fully automated nucleic acid extraction, limiting performance variability. It allows the use of a single sample for galactomannan enzyme-linked immunosorbent assay (ELISA), β -D-glucan, and PCR analysis, thereby reducing costs if high-throughput screening of high-risk patients is required. As with WB PCR testing, a multicenter clinical trial is paramount to determine the true clinical validity and utility of serum PCR testing. More so, as the clinically relevant biological target circulating in blood has not been precisely determined, a multicenter comparison of WB PCR (targeting fungal cell-associated DNA) and serum PCR (targeting free circulating DNA) is required to determine the clinically relevant biological target and consequently the optimal clinical specimen for PCR testing.

ACKNOWLEDGMENTS

We acknowledge the European Aspergillus Initiative of the ISHAM.

The EAPCRI thanks Andreas Opitz (Institute of Transfusional medicine, Wuerzburg University Hospital, Wuerzburg, Germany) for drawing and screening the blood kindly donated by our volunteers.

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We thank the following members of the participating laboratories: Vanda Chrenkova, Motol Hospital, Prague, Czech Republic; Malcolm Guiver, HPA Northwest, Manchester, United Kingdom; Catriona Halliday and Sue Sleiman, Westmead Hospital, Westmead, NSW, Australia; Chris Linton and Elizabeth Johnson, UK Mycology Reference Lab, HPA Southwest, Bristol, United Kingdom; Rebecca Gorton and Chris Kibbler, Royal Free Hospital, London, United Kingdom; Martina Lengerova, Central Molecular Biology, Gene Therapy and Haematology Clinic, Brno, Czech Republic; Eva Rosello Mayans, Vall d'Hebron University Hospital, Barcelona, Spain; Tom Rogers and Oliver Morton, Trinity College, Dublin, Ireland; Boualem Sendid, Lille University, Lille, France; Angie Caliendo, Emory University, Atlanta, GA; Tom Patterson, UTH-CSA, San Antonio, TX; Katrien Lagrou, University Hospital Gasthuisberg, Leuven, Belgium; Mark Wilks, Barts and the London Hospital, London, United Kingdom; Adrian Moody, Myconostica, Ltd., Manchester, United Kingdom; Maria J. Buitrago and Leticia Bernal-Martinez, Spanish National Centre for Microbiology, Madrid, Spain; and Jaques Billes and Phillippe Hauser, Lausanne, Switzerland.

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