

PCR in Diagnosis of Invasive Aspergillosis: a Meta-Analysis of Diagnostic Performance

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Invasive aspergillosis is a difficult-to-diagnose infection with a high mortality rate that affects high-risk groups such as patients with neutropenia and hematologic malignancies. We performed a bivariate meta-analysis of diagnostic data for an *Aspergillus* sp. PCR assay with blood specimens from high-risk hematology patients. We included all studies involving human subjects that assessed the performance of any PCR assay for invasive aspergillosis in whole blood or serum and that used the European Organization for the treatment of Cancer/Mycoses Study Group criteria as a reference standard. Three investigators independently searched the literature for eligible studies and extracted the data. Out of a total of 37 studies, 25 met strict quality criteria and were included in our evidence synthesis. Twenty-five studies with 2,595 patients were analyzed. The pooled diagnostic performance of whole-blood and serum PCR assays was moderate, with a sensitivity and specificity of 84% (95% confidence interval [CI], 75 to 91%) and 76% (95% CI, 65 to 84%), respectively, suggesting that a positive or negative result is unable, on its own, to confirm or exclude a suspected infection. The performance of a PCR assay of serum was not significantly different from that of whole blood. Notably, at least two positive PCR test results were found to have a specificity of 95% and a sensitivity of 64% for invasive infection, achieving a high positive likelihood ratio of 12.8. Importantly, the European *Aspergillus* PCR Initiative (EAP-CRI) recommendations improved the performance of the PCR even further when at least two positive specimens were used to define PCR positivity. In conclusion, two positive PCR results should be considered highly indicative of an active *Aspergillus* sp. infection. Use of the EAPCRI recommendations by clinical laboratories can further enhance PCR performance.

Despite advances in treatment and supportive care, invasive aspergillosis (IA) is associated with significant morbidity and mortality rates, especially among patients with hematologic malignancies and hematopoietic stem cell transplant (HSCT) recipients (1, 2). Systemic antifungal prophylaxis is widely used in this patient population (3, 4), and patients with recurrent or persistent fever and prolonged neutropenia frequently require empirical coverage with antifungal agents (5). Timely and accurate diagnosis of an active infection is needed in order to initiate targeted antifungal therapy and avoid unnecessary antifungal treatment, which is often accompanied by a multitude of side effects and the cumulative risk of resistance.

There is a need for the development of newer diagnostic techniques that would ideally be able to identify IA rapidly, noninvasively, and at an early stage. To aid in this endeavor, the European Organization for the treatment of Cancer/Mycoses Study Group (EORTC/MSG) developed specific criteria for the diagnosis of IA in 2002 (6), which were later revised in 2008 (7), in an attempt to provide the elusive “gold standard” to which any newer diagnostic test should be compared. Dominant among the novel diagnostic methods is the PCR assay, which allows pathogen DNA detection and identification to the species level in a variety of clinical samples. A number of PCR assays are now available to detect *Aspergillus* spp. in various clinical specimens and have a high diagnostic yield *in vitro*, but their clinical performance is debated because of contradictory reports from clinical trials. Previous systematic reviews have assessed *Aspergillus* sp. PCR assay performance with promising findings (8) but did not succeed in convincing the guideline-issuing organizations to incorporate the test in their algorithms (6, 7). The main argument has been that differences in the PCR protocols used in different studies do not allow sufficient interlaboratory comparisons to be made and thus preclude wide-

spread implementation (9). In an effort to determine the value of the *Aspergillus* sp. PCR assay in clinical diagnostics as a screening or a confirmatory tool and to evaluate different parameters that could contribute to contradictory reports regarding PCR performance from the existent literature, we performed a meta-analysis of clinical trials that evaluated the accuracy of the PCR assay for IA performed with serum and whole blood from high-risk patients.

MATERIALS AND METHODS

Literature search. A systematic literature search of the MEDLINE (1951 to December 2013) and Cochrane Central Register of Controlled Trials (2010 to June 2013) databases was conducted to identify all of the published studies involving human subjects and evaluating the performance of a PCR assay for *Aspergillus* spp. using as a reference standard the EORTC/MSG criteria. We searched only full publications and not unpublished studies or conference abstracts. We used the search term “Aspergil* AND (PCR OR PCR).” The date of last access was 20 January 2014.

Three investigators (M.A., I.M.Z., and F.N.Z.) independently identified and scrutinized studies for potential inclusion. Studies published in languages other than English were excluded. Our meta-analysis is in line with PRISMA (preferred reporting items for systematic reviews and meta-analyses) recommendations (10).

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Eligibility criteria and definitions. We included all studies involving human subjects that assessed the performance of any PCR assay for IA in whole blood or serum and that used as a reference standard the EORTC/MSG criteria (either the 2002 [6] or the revised 2008 [7] criteria). Studies of series of patients with a positive-only (probable/definite) or negative-only (no infection) reference outcome were not considered because it is impossible to extrapolate all of the pertinent diagnostic estimates from these studies. We defined as “true positive” the cases that were positive by PCR and were classified as definite or probable by the EORTC/MSG criteria and as “true negative” the cases that were negative by PCR and were classified as possible or unlikely infection by the EORTC/MSG criteria. Possible IA cases were categorized as negative in accordance with the recommendations of the Cochrane group (11). For our primary analysis, we defined as PCR positive any case that had at least one positive PCR result. In multiarm studies that used the same patients more than once, we included the strata with the highest reported performance estimates. Also, in trials that examined PCR assay performance in both a high-risk group and a control group of patients at low risk for IA, we excluded the control arm to avoid a case-control design that could spuriously increase performance estimates.

Outcomes of interest. The primary outcomes of interest were the summary sensitivity and specificity of an *Aspergillus* PCR assay of blood products compared to the reference standard. Our secondary goal was to evaluate whether the choice of sample, the type of PCR, the choice of primers, or any of the methodological aspects proposed by the European *Aspergillus* PCR Initiative (EAPCRI) (12, 13) for optimum PCR assay performance had a significant impact on the test’s accuracy. Finally, we performed a sensitivity analysis to evaluate how PCR assay performance would be affected if two positive results were used to define a positive PCR outcome.

Data extraction and quality assessment. Data from eligible studies were independently extracted by three reviewers (M.A., I.M.Z., and F.N.Z.). Discrepancies between authors were resolved by consensus. The methodological quality of each trial was evaluated independently by three of us (M.A., I.M.Z., and F.N.Z.) for potential sources of bias by using standard criteria. The QUADAS-2 score was used to test for potential sources of bias in each study (14). To have high-quality estimates, we excluded from further analysis the studies with high-bias QUADAS-2 elements. Specifically, we excluded studies that had a nonrandom population or a case-control design, and we also excluded studies that did not have a prespecified threshold of PCR positivity. We chose not to eliminate trials in which the investigators had knowledge of the EORTC/MSG classification of the patient before PCR interpretation because this knowledge cannot alter the interpretation of a positive or negative finding as long as the positivity threshold is prespecified. Finally, we excluded studies that had a >10% loss to follow-up.

Statistical analysis. We calculated the independent sensitivity and specificity of each study by using a two-by-two contingency table. Each table consisted of true positives, true negatives, false positives, and false negatives according to the results of the *Aspergillus* sp. PCR assay (the index test) compared to the classification of the patient as definite/probable versus possible/unlikely aspergillosis on the basis of the EORTC/MSG criteria (the reference standard). We estimated the combined (pooled) sensitivity, specificity, likelihood ratios (LRs), and diagnostic accuracy (area under the curve) by using a bivariate mixed-effects binomial regression model that accounts for variability both within a study and between studies (15–17). We used empirical thresholds of >10 for LR^+ and <0.1 for LR^- to rate the test as of high value in the decision to rule in (i.e., probable/definite aspergillosis) or rule out infection (18). Publication bias was assessed by using the Deeks regression test for asymmetry, with $P < 0.05$ for the slope coefficient denoting significant asymmetry (19). We did not use heterogeneity tests (16, 20), as such tests can mislead systematic reviews of diagnostic test accuracy, are not recommended by the Cochrane diagnostic test accuracy group (16), and may be misleading in diagnostic accuracy studies, since computing separate I^2

statistics for sensitivity and specificity will usually overestimate statistical heterogeneity. Effects were plotted as summary receiver operating characteristic (SROC) curves (15).

We calculated subgroup estimates to account for variation in laboratory PCR methodologies, provided that we had at least four pertinent studies in each group to draw the quadrature points and provided that we had data on the magnitude and difference of average sensitivity and specificity estimates between groups. We also evaluated whether the use of the old (2002) or the revised (2008) EORTC/MSG criteria as a reference standard affected PCR performance estimates. The significance of differences between average specificities or sensitivities was assessed by metaregression, and a P value was reported.

The bivariate meta-analysis was performed with Stata v11 (StataCorp, College Station, TX) by using the Midas set of commands (21, 22). The metaregression was performed with MetaAnalyst software (23).

RESULTS

Two thousand six nonduplicate studies were identified by the initial search, of which 37 met the inclusion criteria (Fig. 1). Twenty-three studies provided data for PCR based on the 2002 EORTC/MSG criteria (24–46), and 14 did so by using the revised (2008) criteria (47–60) (Table 1).

Quality assessment. Most of the studies were of high quality, as determined by the QUADAS-2 assessment tool (Table 2). After all of the low-quality studies were excluded, 25 studies remained and were further analyzed (25, 26, 28–37, 43–48, 50, 52–54, 56–58).

The 25 eligible studies (26 strata) included a total of 2,595 patients. The number of patients enrolled varied from 8 to 218 (median, 91). The target patient population was patients with hematologic malignancies and/or HSCT recipients in 17/25 studies, bone marrow transplant recipients in 1/25, and a mixed population of patients with hematologic malignancies and other immunocompromised individuals in 7/25. Six studies performed PCR assays of serum (23%), and the rest used whole-blood assays (77%). The setting was heterogeneous regarding quality laboratory criteria. Of note, only 3 studies (all 3 using a whole-blood PCR assay) were found to be fully compliant with all EAPCRI criteria, while 7 studies (5 using whole blood and 2 using serum) were compliant with all of the criteria but one, 10 (9 using whole blood and 1 using serum) deviated by 2 criteria, and 6 (4 using whole-blood and 2 using serum) deviated by 3 or more.

Pooled diagnostic estimates. Across the 25 eligible studies, the individual sensitivity estimates ranged from 0 to 1.0, while the individual specificity estimates ranged from 0.29 to 0.98 (Fig. 2). In bivariate meta-analysis, the pooled sensitivity was 0.84 (0.75 to 0.91) and the corresponding pooled specificity was 0.76 (0.65 to 0.84) (Table 2; Fig. 3). There was no evidence of significant funnel plot asymmetry (Deeks’ bias = -0.71 , $P = 0.9$), indicating that small study effects were not present. The combined effects show that the *Aspergillus* sp. PCR assay has a positive LR of 3.5 and a negative LR of 0.21, suggesting that its diagnostic performance is moderate (61). Specifically, in settings where the expected pretest probability of IA is low ($\leq 5\%$), PCR will yield a positive predictive value (PPV) of 0.08 (range, 0.06 to 0.11) and a negative predictive value (NPV) of 0.99 (range, 0.97 to 1.00), while in settings where IA is strongly suspected (pretest probability, $\geq 10\%$) PCR is expected to yield a PPV of 0.76 (range, 0.71 to 0.81) and an NPV of 0.71 (range, 0.66 to 0.77), respectively.

Subgroup analysis. We performed a subgroup analysis to evaluate changes in sensitivity and specificity caused by a series of different parameters in each study. We also assessed whether the

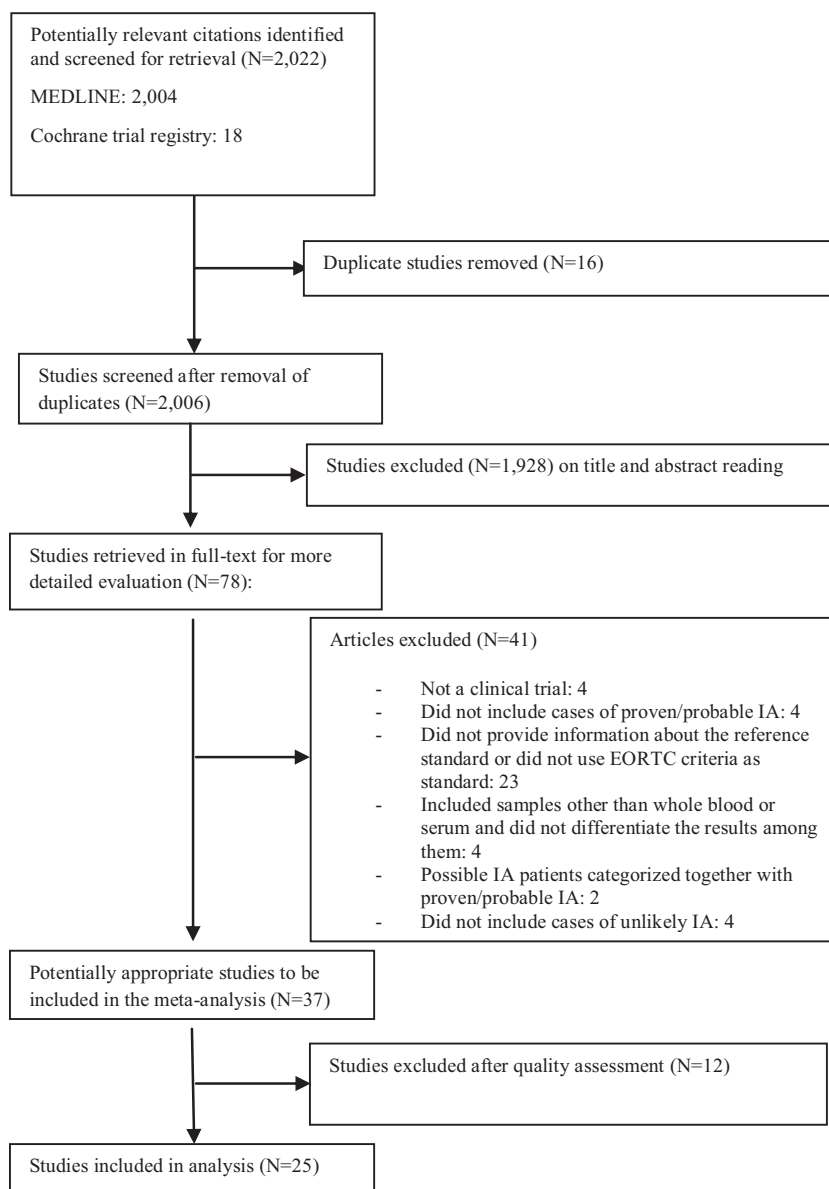


FIG 1 Flow diagram of the inclusion criteria used in this study.

effect of these covariates on the average sensitivity estimates were significant. Specifically, across studies where the updated EORTC/MSG 2008 criteria were implemented, the diagnostic estimates were 0.83 (range, 0.72 to 0.90) for sensitivity and 0.79 (range, 0.67 to 0.87) for specificity, respectively. These estimates did not differ from those of studies that used the EORTC/MSG 2002 criteria as the reference index. Interestingly, studies that used PCR assays of serum yielded higher specificity (85% versus 73%) and lower sensitivity (78% versus 86%) estimates than those that used whole-blood assays, but these differences did not reach statistical significance. It is of note that the average specificity of serum was significantly higher than that of whole blood without bead beating ($P = 0.04$).

Regarding the assay methodology used, no methodological parameter significantly affected PCR performance (Table 3). Further, when the studies that deviated by no more than one of the

EAPCRI criteria were compared to studies that deviated by two or more, no significant changes in sensitivity (87% versus 82%, respectively) or specificity (77% versus 75%, respectively) were noted. Of note, we could not assess the effect of white blood cell and red blood cell lysis steps, as these were undertaken by almost all of studies in which whole blood was used. We also were unable to evaluate the effect of the use of mitochondrial versus ribosomal primers because only two studies used the former (36, 44), while the difference in PCR performance among different ribosomal primers (18S rRNA versus others) was not significant.

Sensitivity analysis. Thirteen of our 26 studies (12 using whole blood and 1 using serum) included data on the diagnostic performance of the PCR assay when at least two PCR results were considered positive. By analyzing the performance estimates of these studies using differential criteria for PCR positivity (one versus two or more positive PCR results), we found that PCR specificity

TABLE 1 Characteristics of studies included

Reference	Study design	Population studied	PCR type	Specimen tested	Primer(s) used	No. of proven/ probable IA episodes	No. of unlikely IA episodes
25	Retrospective	218 patients with hematologic malignancies	Nested	Whole blood	18S rDNA	36	182
24	Prospective	54 patients with cancer and pulmonary infiltrates	Traditional	Whole blood	Mitochondrial DNA and alkaline protease gene	11	36
36	Prospective	17 hematology patients	Nested	Whole blood	Mitochondrial DNA	3	10
39	Retrospective	20 Patients with hematologic malignancies who had proven or probable IA	Real time	Serum	Mitochondrial DNA	20	10
33	Prospective	8 patients at risk for IFI, ^a 45 negative controls	Real time	Whole blood	18S rDNA	2	4
45	Retrospective	106 patients at risk for IFIs	Real time	Serum	5.8S rDNA	41	35
41	Prospective	96 patients at risk for IA	Real time	Whole blood	18S rRNA gene	11	125
35	Retrospective	25 patients with hematologic malignancies	PCR-ELISA	Whole blood	18S rDNA	5	20
40	Retrospective	29 patients with at least one positive galactomannan test	Real time	Serum	Mitochondrial DNA	7	11
32	Prospective	78 immunocompromised hemato-oncology patients	Real time	Whole blood	18S rDNA	1	106
34	Retrospective	77 hematologic malignancy patients or solid-organ transplant recipients at high risk for IA	Real time	Whole blood	18S rDNA	3	42
26	Prospective	29 adults and 36 children with febrile neutropenia undergoing intensive chemotherapy for hematologic malignancy or having received HSCT	Nested	Whole blood	18S rRNA gene	13	61
46	Prospective	91 febrile neutropenic pediatric patients	Traditional	Serum	18S rDNA	28	49
29	Prospective	203 patients at risk for IFI	Real time	Whole blood	28S rRNA	13	149
38	Prospective	201 patients with hematologic malignancies	PCR-ELISA	Serum	Mitochondrial DNA	33	106
27	Prospective	194 patients with hematologic malignancies	PCR-ELISA	Whole blood	rRNA	15	150
30	Prospective	62 pediatric patients at risk for IA	Real time	Whole blood	18S rRNA gene	8	54
42	Retrospective	25 patients with at least 1 GM ^b -positive serum sample	Real time	Serum	Mitochondrial DNA	11	12
43	Prospective	124 patients with hematologic malignancies undergoing chemotherapy or HSCT	Real time with fluorescent probes	Serum	28S rRNA	15	121
37	Prospective	125 hematology patients	Real time	Whole blood	18S rDNA	7	84
31	Prospective	127 patients at risk for IA	Real time	Whole blood	18S rRNA	5	105
57	Prospective	172 patients who received high-dose chemotherapy	Traditional	Serum	18S rRNA	20	173
28	Prospective	91 patients in AmBiLoad trial	Nested	Whole blood	18S rRNA	59	1

(Continued on following page)

TABLE 1 (Continued)

Reference	Study design	Population studied	PCR type	Specimen tested	Primer(s) used	No. of proven/probable IA episodes	No. of unlikely IA episodes
47	Prospective	82 bone marrow transplant recipients	Real time	Whole blood	18S rDNA	10	67
48	Prospective	46 patients receiving either allogeneic stem cell transplant or myeloablative chemotherapy	Real time	Whole blood	Multicopy ribosomal operon region from ITS1 ^d to 5.8S region	3	24
59	Retrospective	44 patients with two sequential positive serum galactomannan results and an IA risk factor	Two different real-time assays	Serum	Assay 1, mitochondrial DNA; assay 2, 18S rRNA	26	18
60	Retrospective	31 patients (10 with proven/probable IA vs 21 with no IA)	Two different real-time PCR assays	Serum	Assay 1, 28S rRNA; assay 2, 18S rRNA	10	21
49	Retrospective	38 adult patients with a high clinical suspicion of IA	Real time	Serum and whole blood	ITS1	18	4
58	Prospective	62 pediatric patients at increased risk for IA	Nested	Serum	Not reported	10	26
50	Prospective	63 patients with allogeneic stem cell transplant and myeloablative chemotherapy	Real time	Whole blood	ITS1-5.8S	3	41
51	Retrospective multicenter	47 patients with proven/probable IA and 31 controls	Various real-time PCR assays	Serum and whole blood	Various	47	31
52	Prospective (two different centers)	278 patients undergoing intensive chemotherapy or HSCT	Two different real-time PCR assays	Whole blood	28S rRNA (nested) ITS (single run)	15/30	120/83
54	Prospective	72 patients with hematologic malignancies with fever, 4 with normal temperatures, 10 healthy volunteers	Real time	Whole blood	28S-ITS2 rRNA genes	22	41
53	Prospective	51 patients with hematologic malignancies at risk for IA	Nested	Whole blood	18S	6	47
55	Prospective	103 adult hematology patients at high risk for IA	Real time	Whole blood	28S rDNA	22	59
44	Prospective	185 febrile neutropenic patients treated with chemotherapy for AML ^c	Real time	Serum	Mitochondrial DNA	11	174
56	Prospective	62 hematologic malignancy patients	Real time	Whole blood	18S rDNA	5	44

^a IFI, invasive fungal infection.

^b GM, galactomannan.

^c AML, acute myeloid leukemia.

^d ITS1, internal transcribed spacer 1.

was greatly increased (73% versus 95%, respectively) with the latter method, giving a high positive LR of 12.8, whereas sensitivity decreased (85% versus 64%, respectively). Moreover, two or more positive assays for low-risk individuals ($\leq 5\%$ pretest probability) give a PPV and an NPV of 0.23 and 0.99, respectively, while among high-risk patients ($\geq 10\%$ pretest probability), the PPV increases to 0.90 while the NPV drops to 0.62.

Interestingly, when we compared the effect of the EAPCRI recommendations on this sensitivity analysis (Table 4), we found that bead beating significantly improved the specificity of the test (96% versus 86%, metaregression $P = 0.006$), with a nonsignificant drop in sensitivity (50% versus 73%). Further, we found that compliant studies (no more than one deviation from the EAPCRI criteria) showed higher specificity (98% versus 85%, metaregression

TABLE 2 Quality assessment and final determination of inclusion versus exclusion based on QUADAS-2 tool^a

Sample type used for PCR assay and reference	Patient selection		Index test		Reference standard		Flow and timing		Applicability		No. of deviations from EAPCRI recommendations				
	Random patient selection	Case-control design	Test results without knowledge of standard threshold	Inappropriate exclusions	Reference standard interpreted without knowledge of index test	Reference standard likely to classify condition correctly	Appropriate interval between index test and reference standard	Did all patients receive a reference standard?	Was the reference standard the same for all patients?	Were all patients included in the analysis?		Are there concerns that the patients included do not match the review question?	Are there concerns that the condition defined by the reference standard does not match the review question?	Are there concerns that the index test differs from the review question?	
Whole blood															
24	No	Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	2
25	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
26	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	3
27	<i>Unclear</i>	No	<i>Unclear</i>	<i>Unclear</i>	Yes	Yes	<i>Unclear</i>	Yes	Yes	Yes	Low	Yes	Yes	Excluded	2
28	Yes	No	No	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	1
29	Yes	No	<i>Unclear</i>	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	4
30	Yes	No	No	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	1
31	Yes	No	<i>Unclear</i>	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	1
32	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	0
33	Yes	No	<i>Unclear</i>	<i>Unclear</i>	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	0
34	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	0
35	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	1
36	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	1
37	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
38	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	0
39	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
40	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
41	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
42	No	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
43	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
44	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
45	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
46	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
Serum															
38	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	2
39	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	2
40	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	3
41	Yes	No	No	No	Yes	Yes	No	Yes	Yes	Yes	Low	Yes	Yes	Excluded	3
42	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	3
43	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	3
44	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	3
45	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Excluded	3
46	Yes	No	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Low	Yes	Yes	Included	3

^a Boldface indicates characteristics that predict high risk of bias; italics indicate uncertain bias.

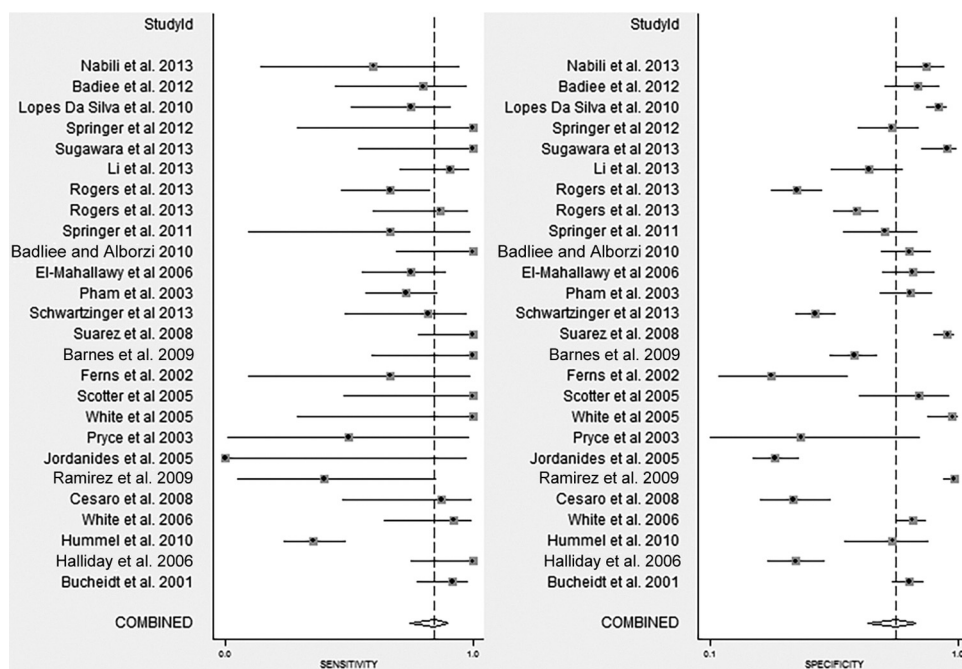


FIG 2 Forest plot of independent sensitivity and specificity estimates. The studies cited correspond to references 25, 26, 28 to 37, 43 to 48, 50, 52 to 54, and 56 to 58.

$P = 0.003$) with an additional nonsignificant increase in sensitivity (67% versus 61%).

DISCUSSION

Despite the fact that multiple *Aspergillus* sp. PCR assays of blood specimens are now available, a consensual conclusion about the role that this test can play in the diagnosis of IA has yet to be reached. Shortly after its introduction in the 1990s, it was thought

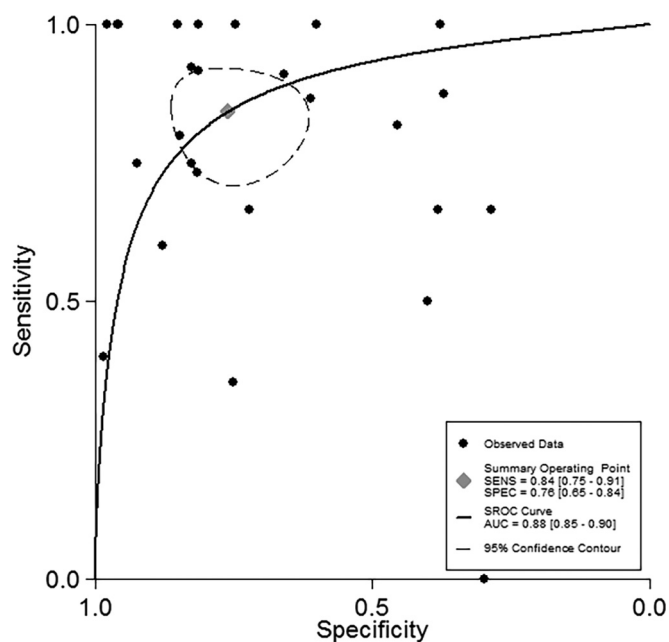


FIG 3 SROC curve of PCR performance.

that this method could, in fact, revolutionize the diagnosis and management of this severe disease (62). However, the multitude of clinical studies performed since then failed to prove beyond doubt whether the test can rule an active infection in or out, thus often leaving clinicians baffled when trying to interpret a positive or negative result. Our findings indicate that whole-blood and serum *Aspergillus* sp. PCR assays have moderate diagnostic performance, which suggests that a positive or negative result is unable, on its own, to confirm or exclude a suspected infection in high-risk patients. We should note, however, that the pooled PCR performance estimates in our study were not inferior to the performance estimates of serum galactomannan (sensitivity, 71%; specificity, 89%) (63) or beta-glucan (sensitivity, 76.8%; specificity, 85.3%) (64) tests obtained by previous meta-analyses.

Our choice to define as PCR positive all episodes that had at least one PCR-positive specimen, coupled with the fact that most of the studies tested more than one specimen per episode, may have overestimated the average sensitivity and underestimated the specificity of the method. To see whether this is true, we performed a sensitivity analysis of studies that had extractable data about at least two versus one PCR-positive specimen per patient. Indeed, we found that PCR specificity was increased dramatically by this approach to 95%, which leads to a PPV of 90% for high-risk individuals. Therefore, the presence of at least two positive whole-blood PCR specimens in a high-risk patient should be considered very indicative, if not confirmatory, of IA. On the other hand, the sensitivity of this approach was lower at 64%, which suggests that it would be more valuable as a confirmatory tool than as a screening tool. Although our results are in accordance with previous reports (8), we observed a significant variation of results reported by individual studies involved in our analysis, which led us to search for reasons behind these inconsistencies.

One of the potential explanations is the choice of the specimen,

TABLE 3 Results of subgroup analysis^c

Group or subgroup (no.) of studies with ≥ 1 positive result	Sensitivity (95% CI) ^d	Specificity (95% CI)	LR ⁺ /LR ⁻
All studies (26)	0.84 (0.75–0.91)	0.76 (0.65–0.84)	3.5/0.21
Criteria used			
2008 (10)	0.83 (0.72–0.90)	0.79 (0.67–0.87)	3.9/0.22
2002 (16)	0.85 (0.69–0.93)	0.74 (0.57–0.86)	3.2/0.21
Study design			
Prospective (20)	0.86 (0.77–0.92)	0.76 (0.63–0.85)	3.5/0.18
Retrospective (6)	0.76 (0.49–0.91)	0.77 (0.55–0.90)	3.2/0.31
Specimen type			
Serum (6)	0.78 (0.69–0.85)	0.85 (0.70–0.93)	5.1/0.25
Whole blood (10)	0.86 (0.73–0.93)	0.73 (0.59–0.83)	3.1/0.19
Elution vol (μ l) of:			
≤ 100 (13)	0.86 (0.76–0.93)	0.72 (0.59–0.82)	3.1/0.19
> 100 (8)	0.82 (0.54–0.95)	0.75 (0.46–0.91)	3.3/0.24
Sample vol			
Large ^a (18)	0.83 (0.70–0.91)	0.79 (0.68–0.87)	4.0/0.21
Small ^b (7)	0.89 (0.77–0.95)	0.64 (0.38–0.83)	2.4/0.17
Internal control			
Yes (19)	0.84 (0.73–0.91)	0.76 (0.63–0.85)	3.4/0.21
No (7)	0.79 (0.67–0.87)	0.77 (0.56–0.90)	3.4/0.27
Primer type			
18S (16)	0.84 (0.69–0.93)	0.80 (0.67–0.89)	4.3/0.19
Other ribosomal (7)	0.85 (0.70–0.93)	0.74 (0.57–0.86)	3.3/0.20
Bead beating (for whole blood only)			
Yes (8)	0.78 (0.62–0.89)	0.82 (0.64–0.92)	4.3/0.27
No (12)	0.90 (0.68–0.98)	0.64 (0.47–0.78)	2.5/0.15
Compliance with criteria			
0–1 deviation (10)	0.87 (0.74–0.94)	0.77 (0.60–0.88)	3.8/0.17
> 1 deviation (16)	0.82 (0.68–0.90)	0.75 (0.60–0.86)	3.3/0.24

^a Sample volumes used: ≥ 3 ml of whole blood, ≥ 0.5 ml of serum.

^b Sample volumes used: < 3 ml of whole blood, < 0.5 ml of serum.

^c No statistically significant differences were found in any comparison of subgroups.

^d CI, confidence interval.

namely, whole blood or serum, with which the PCR assay is performed. Two previous clinical studies comparing *Aspergillus* sp. PCR assays performed with serum and whole blood from the same high-risk patients failed to show significant differences in accuracy (49, 51), with one study suggesting a nonsignificant trend toward increased sensitivity and reduced specificity of the whole-blood PCR assay (51). In accordance with these studies, our results suggest that PCR assays performed with serum had nonsignificantly lower sensitivity and higher specificity than those performed with whole blood. Taking into consideration the facts that a serum PCR assay is faster and easier to perform and that serum can also be used for other biomarker diagnostic tests, such as galactomannan, serum has the potential to be the preferred specimen for PCR testing.

Another potential moderator of effect among different trials could be the use of different versions of the EORTC/MSG criteria

TABLE 4 Results of sensitivity analysis

Group or subgroup (no.) of studies with ≥ 2 positive results	Sensitivity (95% CI) ^f	Specificity (95% CI)	LR ⁺ /LR ⁻
All studies (13)	0.64 (0.38–0.84)	0.95 (0.88–0.98)	12.8/0.38
Criteria used			
2008 (5)	0.64 (0.28–0.89)	0.96 (0.91–0.98)	15.0/0.38
2002 (8)	0.63 (0.27–0.89)	0.94 (0.80–0.98)	10.7/0.39
Study design			
Prospective (11)	0.71 (0.45–0.88)	0.95 (0.87–0.98)	13.8/0.31
Retrospective (2)	NA ^a	NA	NA
Specimen type			
Serum (1)	NA	NA	NA
Whole blood (12)	0.62 (0.34–0.84)	0.93 (0.86–0.97)	9.5/0.41
Elution vol (μ l) of:			
≤ 100 (5)	0.74 (0.31–0.95)	0.95 (0.90–0.97)	14.1/0.28
> 100 (6)	0.68 (0.31–0.91)	0.95 (0.71–0.99)	14.2/0.34
Sample vol			
Large ^b (8)	0.45 (0.18–0.75)	0.97 (0.89–0.99)	15.2/0.57
Small ^c (5)	0.83 (0.57–0.95)	0.89 (0.76–0.95)	7.6/0.19
Internal control			
Yes (11)	0.66 (0.34–0.88)	0.95 (0.89–0.98)	14.5/0.36
No (2)	NA	NA	NA
Primer type			
18S (8)	0.64 (0.25–0.91)	0.92 (0.80–0.97)	8.0/0.39
Other ribosomal (5)	0.64 (0.34–0.86)	0.98 (0.88–1.0)	27.4/0.37
Bead beating (for whole blood only)			
Yes (6) ^d	0.50 (0.25–0.76)	0.96 (0.92–0.98)	13.7/0.51
No (6)	0.73 (0.16–0.97)	0.86 (0.71–0.94)	5.1/0.31
Compliance with criteria			
0–1 deviation (7) ^e	0.67 (0.34–0.89)	0.98 (0.93–0.99)	28.1/0.33
> 1 deviation (6)	0.61 (0.23–0.89)	0.85 (0.71–0.93)	4.2/0.46

^a NA, not applicable (too few studies to pool).

^b Sample volumes used: < 3 ml of whole blood, < 0.5 ml of serum.

^c Sample volumes used: ≥ 3 ml of whole blood, ≥ 0.5 ml of serum.

^d Average specificity significantly improves for bead beating ($P = 0.006$).

^e Average specificity significantly improves for the more compliant studies ($P = 0.003$). All other subgroup comparisons not significant.

^f CI, confidence interval.

for the diagnosis of IA. Compared to the old version, the revised criteria (7) kept the terminology but expanded the definition of probable while reducing the scope of possible IA. The adoption of the new classification may have an unclear impact on outcomes in future trials. Indeed, a recent retrospective evaluation stated that the majority of “possible” IA cases can be downgraded to “unclassifiable” but also probable cases showed 75% reductions (65). While this impact may or may not hold true in clinical practice, our data indicated that PCR performance was unaffected.

Finally, the use of different methodological parameters in DNA extraction and amplification is considered one of the most important sources of inconsistencies between different trials (66). To circumvent this problem and standardize *Aspergillus* sp. PCR assays among different laboratories, the EAPCRI has recently issued

a series of recommendations on how to perform *Aspergillus* PCR assays with whole blood and serum on the basis of the results of two multicenter studies with spiked samples (12, 13). These include the use of a larger sample volume, smaller elution volumes, an internal control, and thorough cell lysis with bead beating and enzymatic white and red blood cell lysis steps for whole blood. Notably, in the clinical setting we studied, none of these methodological characteristics significantly altered the sensitivity or specificity estimates in our primary analysis.

The interpretation of this finding is challenging, mainly because we do not know in what form *Aspergillus* spp. circulate in the blood. Of note, many studies suggest that most of the circulating *Aspergillus* DNA is present not in the form of conidia but rather as free DNA (67, 68). This provides an explanation for the fact that blood cultures have such a low sensitivity for aspergillosis (69), as well as our finding that whole-blood testing is not superior to serum testing. This would also explain why sequential cell lysis methods could not improve PCR assay performance for patients with IA, in contrast to what would be expected for samples spiked with *Aspergillus* sp. conidia (13). In addition, it is unclear whether any form of *Aspergillus* DNA can be found in the circulation at all times during an active infection. As suggested by a recent *in vitro* study, release of DNA is intermittent and happens only during certain stages of fungal growth (68). If, in fact, circulating fungal DNA is released intermittently only during mycelial breakdown, it is plausible that its levels in the blood would have such a wide fluctuation that it could make differences in sample size or elution volume relatively insignificant.

Nevertheless, it is of note that compliance with the EAPCRI recommendations significantly increased the specificity of PCR and was also accompanied by an increase in sensitivity when at least two positive results were used to define PCR positivity. An explanation for this seemingly contradictory finding would be that other factors that we were unable to assess (such as PCR testing algorithms and the use of prophylactic antifungal agents) could have served as potential confounders in the assessment of the effect of the EAPCRI recommendations on our primary analysis, and those factors should be evaluated in future trials. Compliance with these simple methodological guidelines should, in fact, be encouraged since it would allow interlaboratory comparisons and has the potential to improve PCR performance in clinical practice.

Several systematic reviews of *Aspergillus* sp. PCR assays of bronchoalveolar lavage fluid have been previously performed, with very promising findings (70). However, despite the large number of new studies on the issue, to our knowledge, there is only one meta-analysis of whole-blood and serum PCR performance to date. That study, by Mengoli et al. (8), reported that PCR assay performance is moderate but specificity can be increased when two PCR results are used to define a positive PCR finding. That study was published before the EAPCRI recommendations were issued and included a smaller number of trials, so the authors could not reach any conclusions regarding the performance of different PCR assay protocols. It is of note that our study confirms the finding that a single positive PCR assay result has moderate diagnostic accuracy, whereas a strategy that uses at least two positive PCR assay results per suspicious episode to define positivity is able to achieve superior specificity. Moreover, by investigating the effects of different protocols on PCR assay performance, our study shows that a serum PCR assay is not inferior to a whole-blood PCR

assay and that compliance with the EAPCRI recommendations has the potential to improve PCR assay performance in clinical practice.

Important limitations of our study are the facts that (i) only a subset of the studies included had extractable data regarding PCR assay performance for at least two positive specimens and (ii) the vast majority of these studies used whole blood and not serum as the test specimen. Therefore, the results of this sensitivity analysis should be interpreted with caution. Also, all of the studies included in our analysis enrolled individuals who had a high risk of invasive fungal infection, as this is the patient population for which the *Aspergillus* sp. PCR test will be the most valuable. Therefore, our performance estimates may not be accurate for the general population.

In summary, for high-risk individuals, two whole-blood or serum specimens PCR assay positive for IA should be considered very indicative, if not confirmatory, of a clinically suspected infection. Given the superior specificity of this approach, it has the potential to be used along with other circulating biomarker detection assays, such as galactomannan, as a criterion to define a probable infection. Compliance with the EAPCRI *Aspergillus* sp. PCR assay recommendations can further increase the specificity of this approach and will allow better interlaboratory comparisons. However, given the complexity of IA diagnosis, it is unlikely that a single noninvasive test will be able to be used alone in clinical decision making. Consequently, the focus of future clinical trials should include the development of decision algorithms that would take into account multiple parameters to guide the management of high-risk groups. In this context, serum and whole-blood PCR assays have a concrete potential to improve our ability to detect and diagnose IA.

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