Current Clinical Methods for Detection of Peri-Prosthetic Joint Infection

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Abstract

Background: Currently, one of the most pressing problems in the field of orthopedic surgery is peri-prosthetic joint infection [PJI]. While there are numerous ways to detect PJI, current clinical detection methods differ across institutions and have varying criteria and protocols. Some of these methods include the Modified Musculoskeletal Infection Society system, culturing, polymerase chain reaction, the determination of the presence of certain biomarkers, testing for the presence of alpha defensin peptides, and leukocyte level testing.

Methods: This review summarizes the most recent publications in the field of PJI detection to highlight current strengths as well as provide future directions to find the system for the quickest, cost-effective, and most accurate way to diagnose these types of infections.

Results: The results of this literature review suggest that, while each method of diagnosis has its advantages, each has various drawbacks as well. Current methods can be expensive, take days to weeks to complete, be prone to contamination, and can produce ambiguous results.

Conclusions: The findings in this review emphasize the need for a more comprehensive and accurate system for diagnosing PJI. In addition, the specific comparison of advantages and drawbacks can be useful for researchers and clinicians with goals of creating new diagnostic tests for PJIs, as well as in clinical scenarios to determine the correct treatment for patients.

Keywords: implant; orthopedic infection; peri-prosthetic joint infection

THE DIFFICULTY IN ESTABLISHING a diagnosis of peri-prosthetic joint infection [PJI] continues to impact adversely the field of orthopedics. The burden of misdiagnosis impedes optimal patient care and physician decision making [1]. The PJI significantly decreases patient quality of life [2] and places an economic burden on the healthcare system [3].

The current gold standard for diagnosis of PJI is the Modified Musculoskeletal Infection Society (MSIS) criteria [4–7]. The MSIS considers major and minor criteria. Infections meeting the major MSIS criteria may include two positive cultures of phenotypically identical organisms or the presence of a sinus tract, but the criteria can fail to detect borderline infections that meet fewer than three minor criteria [7,8]. Infections with low virulence organisms complicate PJI diagnosis because patients can present with reduced clinical symptoms and normal conventional laboratory values [9].

While the MSIS has led to a procedural standardization for PJI detection across hospital systems [10], the need for novel tests with high sensitivity and specificity to aid in PJI diagnosis remains pressing. An ideal detection method should be fast, accurate, specific, easy to use, inexpensive, and have a low rate of contamination. This review will summarize MSIS and the current clinical methods of microbial detection for diagnosis of PJI.

Methods

MSIS

Although other rating systems exist, MSIS is the gold standard for the definitive diagnosis of PJI and the evaluation of new tests. The two major MSIS criteria require direct visu-

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alization of infection via either [1] two or more positive cultures of phenotypically identical organisms, or [2] a visually discernible sinus tract communicates from the prosthesis to the skin and is not caused by aseptic failure. Pre-operative minor criteria include elevated serum C-reactive protein (CRP) or erythrocyte sedimentation rate (ESR), elevation of synovial white blood cell (WBC) count or change on leukocyte esterase test strip, elevation of synovial polymorphonuclear leukocytes, and as of 2018, elevated alpha defensin. If these pre-operative minor criteria are inconclusive in diagnosing PJI, intra-operative criteria of positive histologic analysis of peri-prosthetic tissue, positive purulence, or a single positive culture result can be used to confirm or deny infection.

Historically, the MSIS criteria were introduced in 2011 as the result of a collaborative review of laboratory findings designed to streamline and standardize diagnosis of PJI [11]. These criteria were re-evaluated in 2018 testing the efficacy of established criteria as well as other possible inflammatory markers, resulting in the addition of alpha-defensin to the minor criteria list [12]. These new criteria established a point scoring system for minor criteria based on beta coefficients from regression analysis to determine infection. If the pre-operative score is inconclusive, intra-operative criteria are then considered. Table 1 shows the major and minor criteria

TABLE 1. 2018 MODIFIED MUSCULOSKELETAL INFECTION SOCIETY SCORING DEFINITION FOR PERI-PROSTHETIC JOINT INFECTION

Major criteria (need only one)		Conclusion	
Two positive cultures of the same organism Sinus tract communicating to joint or visualization of the prosthesis		Infected	
Minor criteria (pre-operative)		Score	Conclusion
Serum	Elevated CRP (>1 mg/dL) or	2	≥ 6 infected
	D-dimer (>860 ng/mL)	1	
Synovial	Elevated ESR(30 mm/h)	3	2–5 possible infected
	Elevated WBC count (>3000 cells/mcg/L) or LE (++)	3	
	Positive alpha-defensin (signal-to-cutoff ratio >1)	3	≤ 1 not infected
	Elevated synovial PMN percent (>80%)	2	
	Elevated synovial CRP (>6.9 mg/L)	1	
Intra-operative scoring criteria		Score	Conclusion
Positive histology		3	≥ 6 infected
Positive purulence		3	4–5 inconclusive
Single positive culture		2	≤ 3 not infected

CRP=C-reactive protein; ESR=erythrocyte sedimentation rate; WBC=white blood cell; PMN=polymorphonuclear leukocyte.

along with their associated point values and corresponding conclusion on infection.

The 2018 criteria reported a sensitivity of 97.7% and a specificity of 99.5%, but notes that the criteria may struggle to identify infection in patients with an adverse local tissue reaction or an infection with low virulence organisms, issues that will be discussed further in this review [12].

The biggest shortcoming of the MSIS system is that its major diagnostic criteria rely on methodology that does not reflect the urgency of treatment necessitated for infection. For example, diagnosing PJI using the MSIS criteria relies on obtaining a culture of the joint to confirm the final diagnosis of infection, because this allows for visualization of the offending organism.

A culture is required even with other secondary tests, because it can determine micro-biologic resistance profiles and antibiotic susceptibility [13]. Certain slow growing bacteria, however, such as *Cutibacterium acnes* (*C. acnes*), can require up to two weeks of incubation to confirm infectious etiology [9]. This leads to delays in diagnosis, treatment, and increased complications. In addition, accurate bacterial identification can be obscured because of contamination by commensal bacteria and contamination between plates during storage and incubation [14,15].

To compensate for the time required to culture organisms, patients are treated prophylactically with broad spectrum antibiotic agents [16], a problematic practice with rising rates of antibiotic resistant strains of pathogens [17]. Specific pathogen identification and an antibiotic resistance profile are often unknown before deciding on intervention, which can lead to treatment with antibiotic agents typically reserved for more virulent strains [16].

With regard to MSIS minor criteria, current research demonstrates both promising and disappointing results. Methods using the detection of biomarkers indicative of an inflammatory response to an infection can be inconclusive at differentiating PJI from aseptic loosening [18,19]. Such biomarkers, measured in the serum and synovial fluid, include CRP detection, leukocyte counts and differential, and measurement of interleukin (IL)-6 levels. Other biomarkers such as D-dimer and fibrinogen indicate coagulation as a marker of possible infection.

While the MSIS is considered the gold standard of diagnosis for PJI, the aforementioned markers have low specificity for PJI. As such, confirming a positive result requires secondary testing, which slows the speed of results and can be subject to false negative and positive results that can be seen in culturing methods.

Last, although MSIS criteria do not differ between joints, total shoulder arthroplasty (TSA) has a higher probability of infection with low virulence pathogens such as *C. acnes*, suggesting that “joint-specific” clinical guidelines might be a consideration. [9,20,21]. Overall, problems with detecting PJI and pathogen identification based on the criteria listed above demonstrate the need for more time sensitive and accurate diagnostic methods.

Culturing

Current culturing methodology uses samples of explant, tissue, or joint aspiration fluid that are incubated on agar plates under aerobic and anaerobic conditions to identify

offending pathogens. As one of the two major MSIS criteria, culturing allows direct visualization of bacteria. In this system, two positive cultures of phenotypically identical organisms provide evidence that a specific pathogen is present. The growth of bacteria in culture, however, does not necessarily mean an infection is present. Various commensal bacteria live in and on the human body [22], meaning a single positive culture may not definitively indicate a PJI. This method can have a sensitivity as low as 9.7% [23]. Even two or more positive tissue cultures can have a sensitivity as low as 40.74% [24].

The low virulence of *Cutibacterium spp.* can cause infections with reduced clinical symptoms, normal laboratory panels, and reduced values of conventional tests for PJI [9]. Diagnosis in this setting is challenging and can delay treatment [9]. Synovial fluid cultures had only a 49% sensitivity for PJI. Renz et al. [9] noted that peri-implant tissue culture had a sensitivity of 74%, while explant sonication improved sensitivity to 81%. In addition, successfully culturing *C. acnes* requires incubation under anaerobic conditions for up to 14 days [9,25], making it a difficult pathogen to identify correctly and manage.

For TSA, culturing almost always returns monomicrobial results, most often *C. acnes* [9,22] although non-virulent microbiota are known also to be present [26]. Quantifying the interior bacterial load, differentiating between epidermal and pathogenic *C. acnes* during sampling, and the chance of sample contamination can pose potential problems for diagnosing a PJI after TSA.

Primarily, *C. acnes* colonizes pilosebaceous glands that are much more prevalent on the shoulder and back than the hip and knee. It is seen less frequently in total hip arthroplasty and total knee arthroplasty (THA and TKA). In THA and TKA, culture duration of 14 days shows no benefit compared with shorter culturing periods, except more frequent growth of contaminants [14]. Interestingly, true positive infections with *C. acnes* in this setting grew positive cultures sooner (mean of five days) rather than later (greater than 11 days). Of patients with *C. acnes* growth, 80% had only one positive culture and cultures were polymicrobial, possibly indicating contamination [14]. This reinforces previous findings [15] that extended culturing durations up to 14 days are more likely to show a false positive from non-virulent pathogens or possible contamination.

Kheir et al. [25] found that over a large cohort [$n = 711$] of hip and knee PJIs, the mean number of samples needed to yield a minimum of two positive cultures (as per MSIS requirement) was four samples. Pathogens not within the five most prevalent organisms (*Staphylococcus aureus*, *coagulase negative Staphylococcus spp.* [CoNS], *Streptococcus spp.*, *Enterococcus spp.*, and *gram negative bacilli*) required an average of seven samples, however. For *C. acnes*, the mean was 10, and for *Escherichia coli*, the mean was 25. This outlier result could be because of the propensity of *E. coli* to form a protective biofilm, which hinders its acquisition for culture.

In these cases, sonication fluid culture, an alternative to conventional methods, which removes adherent bacteria from explants, shows improved results compared with classical culturing, which may miss bacteria in biofilms [25]. Sensitivity and specificity of sonication fluid culture were 77.05% and 98.11% compared with 55.74% and 94.34% for

traditional peri-prosthetic tissue culture [27]. Yan et al. [28] examined sonication fluid cultures and found similar results, with sensitivity of 73.1% and specificity of 96.0%. These results, however, are not consistent with another study by Dudareva et al. [29] showing a reduction in sensitivity from 97% to 76% and specificity from 69% to 57% with sonication compared with tissue culture.

Inagaki et al. [30] reported similarly high statistics for conventional culturing including one positive culture, producing 88.3% sensitivity, while two positive cultures decreased the sensitivity to 80%. Decreased sensitivity with two cultures could point to the presence of contamination and the inconsistency of relying on culturing to confirm PJI. Culturing of sonication fluid should be explored further to determine its utility in the diagnosis of biofilm-forming bacteria versus conventional methods.

Other novel methods, such as tissue placement in blood culture bottles, can be financially more efficient, isolate additional micro-organisms [31], and show sensitivity and specificity similar to sonication [28]. Blood culture bottles could help visualize more non-virulent organisms that can cause less evident PJI [31]. Although time consuming, when conventional culture was combined with polymerase chain reaction (PCR) and vial culture, the specificity reached 98.57%, making it a valuable tool for ruling out infection [24].

Similar to other methods of detection, culturing can accurately detect infections with virulent pathogens such as *S. aureus* with few plates within a short period. Less virulent infections with species such as CoNS and *C. acnes* require more efficient methods to diagnose PJI accurately. Less virulent pathogens and those using protective biofilms necessitate more samples to produce positive results, although increased number of cultures can decrease sensitivity. Further, culturing is susceptible to contamination, because positive cultures in an aseptic patient are often contaminated with CoNS [23]. Culturing is time consuming, and results and protocols can vary between pathogens and sample sites, but remains the only method of PJI detection outside of the presence of a sinus tract that allows direct visualization of infection.

PCR diagnosis

The PCR is used to detect pathogen deoxyribonucleic acid (DNA) in synovial fluid by amplifying sequences of the bacterial genome. By applying this technique to multiple DNA sequences, a multiplex PCR (mPCR) can look for the presence of many pathogens and can be used as an alternative to traditional culturing to identify bacterial presence down to the specific genus and species for bacterial sequences already in a database [32]. The PCR is primarily compared with culturing because of the similar utility in isolating species and identifying antibiotic resistance. Results of PCR, however, are available much faster (within 6 h in some protocols [32]), and require a smaller amount of synovial fluid [33].

Shotgun metagenomic sequencing, a new form of PCR where every nucleic acid is sequenced and compared with a database to identify genetic traits of organisms, showed improved sensitivity compared with synovial fluid culture: 68.5% compared with 54.0%. Metagenomics found bacteria in 94.5% of culture-positive fluid samples, and most of the identified microorganisms were partially or exactly

concordant with culture results. Sequencing also detected bacteria in 40.0% of culture-negative sonicate fluid, however. More than half of these results came from patients who had received pre-operative antibiotic agents. Sequencing can detect bacteria even when a patient has already received antibiotics, which can adversely affect culturing results [34].

The mPCR also showed no adverse effects to results of patients who had received pre-operative antibiotic agents [32]. Sensitivity for mPCR was 71.1%, while culturing was 84.2%, but when these procedures are combined, sensitivity reached 92.1% [32]. Because sequencing targets specific nucleic acids, metagenomic analysis is limited by the amount of host DNA present in the sample. This can be problematic when thresholds are lowered to increase detection, ultimately increasing the specificity but lowering the sensitivity because of DNA contamination [35].

The mPCR showed 85.6% agreement on bacterial presence with culturing. It also detected more *Cutibacterium spp.* and *CoNS*, indicating its utility in detecting low virulence organisms. Low virulence infections currently missed using the standard MSIS criteria could have diagnostic improvements by including a PCR procedure as part of PJI diagnosis [32]. Fastidious and difficult to culture micro-organisms such as *C. pneumonia*, *S. maltophilia*, and *B. melitensis* were also detected by mPCR [36]. Moshirabadi et al. [36] showed vast improvement in sensitivity (from 31.6% up to 97.4%), compared with culturing methods.

Unlike mPCR, broad-range PCR as well as real-time ribonucleic acid (RNA) PCR utilize primers for the 16S rRNA sequence, which is highly conserved among bacterial species and not present in the human genome [37]. Broad-range PCR primers target the genomic sequence less specifically, giving a wider range of amplified sequences than real-time PCR, which has specific primers that amplify more limited sequences [38]. Fang et al. [37] compared the utilization of DNA qPCR with RNA qPCR to determine which proved a better diagnostic tool for detecting PJI. The RNA-based qPCR was found to be less sensitive (73.6%) than its DNA counterpart (81.5%). This is more specific than DNA and more sensitive than culture. Even though DNA qPCR provided the highest sensitivity, this ultimately creates problems with DNA contamination. These findings indicated that RNA based qPCR is a better diagnostic test, having the highest accuracy at 85.9%.

Real-time RNA PCR results agreed with culturing for joint fluid, peri-prosthetic tissue, and sonicated fluid 77.6%, 61.2%, and 83.6% of the time, respectively. The sensitivities for joint fluid (83.0%) and sonicated fluid (84.9%) were both higher than the sensitivities of their culturing counterparts. The specificities of these samples, however, followed the opposite trend. Peri-prosthetic tissue had a much lower sensitivity (34.0%), indicating little promise in its diagnostic abilities [39].

Peri-prosthetic joint infection is not only caused by bacterial colonization but can also occur from fungal presence. Real-time PCR utilizing additional primers such as 28S rRNA [40] or 18S rRNA [41] allow for the detection of fungal and eukaryotic presence. The PCR run with an 18S rRNA primer on a synovial fluid sample taken pre-operatively had a sensitivity of 55.6% and specificity of 82.0% in detection of either bacterial or fungal PJI. These values were compara-

tively less than or equal to those found for culture and CRP values of the same samples. Even with the additional fungal primer, pre-operative synovial fluid testing did not add diagnostic value [41].

Kuo et al. [40], however, found both high sensitivity (100%) and specificity (99.5%) for PCR run on synovial fluid samples collected intra-operatively. These values were significantly higher than the biomarkers used for comparison. In addition, using RNA primers over DNA primers reduced the false positive rate to 4% [40]. Reaching such high sensitivity and specificity helps to combat one of the major problems with PCR (false positive results because of contamination), thus demanding further investigation into this combination of primers.

PCR shows promise as an alternative to culturing for direct identification of bacterial species and antibiotic resistance. Different PCR methodologies have been shown to be unaffected by recent administration of antibiotic agents, unlike culturing, and allow for identification of difficult to culture pathogens. The PCR, however, can also detect small bacterial loads of "silent" colonizations of prostheses without clinical relevance, reinforcing the idea that bacteria presence does not always lead to infection [33].

The PCR can also be misleading in polymicrobial infections because sequence analysis is unable to distinguish properly the nucleic acid peaks because of intertwining from multiple species [39,40]. Missing other pathogenic bacteria in an infection can cause mistreatment or misadministration of antibiotic agents. Ultimately, PCR is a costly diagnostic test, each sample costing several hundred US dollars [35], which creates further problems if the genetic material in the sample is not read easily.

Biomarkers

Biomarkers help in the diagnosis of PJI, as they are signals of the body's response to infection. The most commonly used biomarkers, CRP and erythrocyte sedimentation rate (ESR), indicate an inflammatory response by the immune system in relation to the presence of foreign pathogens. Coagulation biomarkers, such as plasma fibrinogen and D-dimer, are being investigated currently as possible indicators of infection. These markers help determine presence of a blood clotting response in the surgical area caused by infection. Not only can markers of human immune responses be used to determine infection, but also studies indicate that identification of bacterial biomarkers confirms their presence [42]. These markers include cell wall components like lipoteichoic acid present in gram-positive cell walls and biologic byproducts of bacterial metabolism.

The CRP is an inflammatory biomarker used in distinguishing PJI from aseptic loosening. Studies have shown that CRP cannot stand alone as a diagnostic tool for distinguishing PJI, but it is helpful when paired with other inflammatory markers. Schiffner et al. [19] found that at the 1 mg/dL threshold for CRP, there was a 65% sensitivity and a 56% specificity. When CRP is measured from synovial fluid, sensitivity and specificity both increase to 88% and 82%, respectively, suggesting that synovial CRP may be a better diagnostic test for PJI than serum CRP. Alone, however, regardless of source, CRP is not specific enough to distinguish PJI from aseptic loosening [19,43]. Despite this, CRP is

a useful complementary test, because patients with higher CRP levels had significant correlation to having more positive intra-operative cultures [44].

Along with CRP, ILs are used to help determine PJI and are often paired with CRP testing. Alone, IL-6 level measurements were shown to have an 86.7% sensitivity and 89.5% specificity for PJI, but when paired with serum CRP testing, sensitivity fell to 73.7% while specificity increased to 100% [45]. This decrease in sensitivity while an increase in specificity is a common problem in pairing biomarker tests [45], because a decline in sensitivity correlates with an increase in false-negative test results, missing cases of PJI. Along with CRP, ILs are used to help determine PJI and are often paired with CRP testing. This is the reason other ILs such as IL-16, IL-18, and cysteine rich with EGF-like domains 2 (CRELD2) are being investigated as better indicators. Chen et al. [46] found that IL-16 and IL-18 and CRELD2 had statistical significance as overall better indicators in PJI detection than CRP and leukocyte count.

Other inflammatory biomarkers, including presepsin, a soluble fraction of CD14, have been shown to have positive linear correlation with both CRP and IL-6, making them comparable and possible second-line tests [47]. Sa-Ngasoonsong et al. [48] determined that procalcitonin (PCT) had a 65% sensitivity and 91.7% specificity for PJI when levels were determined from serum, but the sensitivity increased to 80% when PCT was measured from the synovial fluid.

Along with biomarkers for inflammatory response, indicators of coagulation such as D-dimer and fibrinogen levels can be used to detect PJI. Fibrinogen has been found to be a more promising marker with sensitivity and specificity similar to that of traditional inflammatory markers. Plasma fibrinogen sensitivity and specificity range from 76.3–81.0% and 25.0–86.2%, respectively [49,50]. Plasma fibrinogen has even been shown to have sensitivity as high as 90% when the threshold was lowered to 519 mg/dL, but the specificity of the test still suffered, reaching only 34% [50]. In addition, plasma fibrinogen has a negative predictive value of 98.3%, making it a reasonable marker for ruling out PJI [51].

In comparison, D-dimer had much lower values of 64.5% sensitivity and 65.0% specificity [49]. D-dimer, when combined with other biomarker tests, is seen to have an increase in specificity, because D-dimer on its own had sensitivity and specificity values of 67.44% and 44.09%. When paired with fibrinogen and CRP tests, however, the specificity increased to 90.14% and sensitivity dropped to only 56.10% [52]. Overall, plasma fibrinogen appears to be equally effective as inflammatory markers in ruling out PJI, but the clinical relevancy of D-dimer as an informative biomarker seems to be less promising.

Not only can biomarkers of host immune response be used to determine infection, but biologic markers of the bacteria also can provide an even clearer picture. Host biologic markers quantify inflammatory or coagulation responses, which are host responses to an infection, but the presence of these markers becomes clouded when a patient has inflammatory conditions like arthritis, or if the inflammation is simply because of irritation from aseptic failure [53].

On the other hand, bacterial biomarkers include the metabolic product D-lactate, which is produced almost exclusively by bacteria [54] and whose synovial fluid concen-

tration can be measured spectrophotometrically. Yermak et al. [42] found that using these D-lactate concentrations yielded 86.4% sensitivity and 81.7% specificity, values similar to those for synovial leukocyte count.

Bacteria can also be visualized using these markers, as done in fluorescence in-situ hybridization (FISH), which stains parts of the peri-prosthetic membranes that have bacterial presence. Visualizing bacteria via FISH has a 95% sensitivity and 85.42% specificity related to clinical findings, and higher values of 95.12% and 87.23% when correlated with histopathologic findings [55].

Monoclonal mouse antibodies are also used to visualize bacteria, because radio-labeled mouse antibody targets lipoteichoic acid (LTA) in gram-positive cell walls [18]. On imaging the infected mice, a weak positive correlation between the bioluminescent signal and standardized uptake values was seen with an R value of 0.767 ($R^2=0.588$), indicating specificity of the radio-labeled antibody, but little direct association between the probes [18]. These bacterial biomarkers give a true determination of the bacterial presence in a surgical site, unlike inflammatory and coagulation markers that detect the presence of a response.

Alpha defensin

Synovial fluid levels of alpha defensin have emerged as an adjunct biomarker indicative of local reaction to infection. Alpha defensin is a peptide released by neutrophils with a broad range of anti-pathogenic coverage against bacteria, fungi, and enveloped viruses. Alpha defensins induce pore formation in microbial membranes leading to depolarization and eventual inactivation [56]. As proven with its addition to the 2018 update of the MSIS criteria, clinical data suggest measuring alpha defensin levels can be effective in the diagnosis of PJI.

There are two types of tests available for synovial levels of alpha defensin: The synovasure alpha defensin lateral flow (ADLF) test, and an enzyme-linked immunosorbent assay (ELISA). The results of the ADLF test are available after 10 min while the ELISA results are available the next day after laboratory analysis.

The role of alpha defensin in the local immune response to an infection makes it a prime candidate in the diagnosis of PJI. With a higher specificity than sensitivity in almost every study examined, measurement of alpha defensin level is useful as a confirmatory test, but not a good procedure to rule out diagnosis of PJI or as the sole detection procedure [5,8]. When combined with culturing, sensitivity and specificity reach 96% and 100%, respectively [57]. Alpha defensin may miss infections with low-virulence organisms such as coagulase negative *CoNS* or *C. acnes*, with up to an 80% false negative rate [21,58].

Synovial levels of alpha defensin are unaffected by recent antibiotic therapy in patients [56,58], giving it an advantage over biomarkers such as CRP [43]. Synovasure ADLF showed statistical improvement over MSIS minor criteria including ESR/CRP level, pre-operative culture (1 positive), and synovial cell count [23,60].

Following staged treatment of PJI with insertion of an antibiotic spacer, alpha defensin testing showed low sensitivity but high specificity, furthering the conclusion that the test can be used effectively to confirm diagnosis [61,62].

In patients with adverse local tissue reaction secondary to physical prosthetic failure, Synovasure ADLF had a sensitivity and specificity of 100% and 68%, suggesting that alpha defensin may be prone to false positives in this setting [63]. Therefore, when metallosis occurs, alpha defensin testing may be unable to distinguish between septic and aseptic failure.

In the current review, six studies on the Synovasure ADLF test are grouped together because of their similarity in criteria, time period, and samples, while three on ELISA are deemed comparable. The average sensitivity and specificity of the ADLF test were 81.45% and 94.53%, respectively. The average sensitivity and specificity of the ELISA were 84.7% and 91.7% (Table 2).

In summary, alpha defensin testing remains a useful tool for the diagnosis of PJI, but its strengths and weaknesses should be noted. The averages of several studies of Synovasure ADLF resulted in a specificity of 95%, making this rapid and easy test useful to confirm the presence of infection. Alpha defensin levels have been shown to be unaffected by recent antibiotic therapy and can be used in synovial fluid contaminated with blood [8]. The lower sensitivity and possible false negatives because of low virulence pathogens, however, show that this test is not optimal for detecting infections and should not be used independently of other tests.

The current cutoff for Synovasure may prevent its use in differentiating PJI from gout, pseudogout, and other inflammatory arthropathies [4]. Further, alpha defensin testing does not provide any information about what pathogen is present, leading to broad coverage antibiotic use before return of culture results, which can narrow down the proper antibiotic therapy [15].

Leukocytes

As effector cells of the innate and adaptive immune systems, leukocytes are a critical and functional response to infection. Levels of leukocytes in the blood or synovial fluid, therefore, can be elevated both acutely and chronically to combat a pathogen. Leukocyte levels reflect the presence of inflammation as a result of infection. Several tests have been designed to detect leukocyte levels including the leukocyte esterase (LE) test, histopathologic tissue analysis, and radio-labeled scans.

The LE test rapidly detects the level of enzymes released by neutrophils in synovial fluid. Studies examining LE test

strips found sensitivities ranging from 74.22% to 80.5% and specificities above 92.7% [20,64]. The best sensitivity results were from a Synovasure lateral flow immunoassay with sensitivity of 91.95% [64]. Like other tests, in the context of TSA, LE statistics were poorer with a sensitivity of 50% and a specificity of 87%. Like alpha defensin testing, LE test strips have low sensitivity and high specificity, making LE a poor method for PJI screening, but a better confirmatory test.

Further, the test only detected pathogens in three of 15 patients specifically infected with *C. acnes*, and also had several invalid results because of contamination with blood or insufficient synovial fluid levels [20]. These problems with the LE test strip utilization make it difficult to use as a primary diagnostic test for PJI.

Intra-operatively, the LE test had a sensitivity and specificity of 100% and 93.8%, respectively, compared with 85.7% and 94.9% for frozen section histology in THA [65]. The time from sampling to the results being returned to the surgeon was on average 20.1 min for LE and 27.2 min for frozen section, a useful increase in efficiency, compared with some of the previous methods, during an operation [65].

The LE test strips can be contaminated by blood, leading to invalid readings even when high amounts of WBCs and LE are present in synovial fluid. False positive results can also occur in the setting of physical failure where metal-wear debris interferes with the test or an adverse local tissue reaction increases the level of leukocytes present [65]. Although the test is rapid, inexpensive, and commercially available, its high invalidation rate, up to 11.9%, and low sensitivity make it a poor test to rule out infection [64].

Histopathologic tissue analysis detects leukocytic infiltration as a marker of infection. One study, however, showed no statistical relationship between intra-operative histopathologic examination of neutrophils, which diagnosed 8% of infections, and post-operative hematoxylin and eosin analysis, which diagnosed 28% [66], calling into question the validity of this methodology.

Another method of measuring leukocytes is synovial fluid cell counts. Comparing effectiveness between studies is difficult because each uses a different cutoff value to determine the presence of infection. Synovial leukocyte cell counts produce promising results, but the lack of standardization limits the diagnostic relevance.

Although WBC ranges are more standardized in the settings of THA and TKA, they cannot be extrapolated to

TABLE 2. SUMMARY OF REPORTED ALPHA DEFENSIN TEST RESULTS

Author	Sample size	Sensitivity	Specificity	NPV	PPV	Type of test
Kanwar et al.	35	97.1	97.1	97.1	97.1	ELISA
Renz et al.	212	84.4	96.4	95.8	86.4	ADLF
de Saint Vincent et al.	42	88.9	90.6	96.7	72.7	ADLF
Balato et al.	51	87.5	97.1	94.4	93.3	ADLF
Riccio et al.	40	85.0	96.9			ADLF
Sigmund et al.	101	69.0	94.0	88.3	83.3	ADLF
Ding et al.	70	73.9	92.2	77.8	90.4	ADLF
Kelly et al.	39	82.0	82.0	92.0	64.0	ELISA
Ecker et al.	105	75.0	96.0	93.0	86.0	ELISA
Average (Synovasure)		81.5	94.5	90.6	85.2	
Average (ELISA)		84.7	91.7	94.0	82.4	

NPV=negative predictive value; PPV=positive predictive value; ADLF=alpha defensin lateral flow test; ELISA=enzyme-linked immunosorbant assay

TSA [67]. Further, although one might assume that low grade shoulder infections with pathogens such as *C. acnes* would produce fewer cell counts, they are, in fact, associated with high leukocyte levels [67]. Inagaki et al. [30] found that almost every case of septic knee failure had 5+ neutrophils per 400x magnification in a high power field, indicating its utility visualizing both virulent and non-virulent infections. The 5+ neutrophils under histologic analysis of TKA produced 96.7% sensitivity and 100% specificity in detection of either bacterial or fungal PJI. Neutrophil count does not provide information regarding the degree of infection or bacterial load.

Finally, radio-labeled assays can detect leukocyte accumulation. Blanc et al. used labeled leukocyte scintigraphy [LS] and found 72% sensitivity and 60% specificity [68]. The LS results were unaffected by current antibiotic treatment or type of pathogen, but the process is expensive and time consuming. Faltstie-Jensen et al. [69] found a sensitivity of 18% and a specificity of 100% in TSA using WBC/bone marrow single-photon emission computed tomography. The *C. acnes* infections, however, were not detectable by this method.

Leukocyte levels are increased in response to infection. Although secondary to detecting pathogen presence, measuring leukocyte levels are useful in the diagnosis of PJI. Similar to other secondary methods, however, leukocyte levels are not specific for PJI and can be increased because of other causes of inflammation such as gout [70].

Conclusion

The methodologies used currently in the diagnosis and detection of PJI have both benefits and drawbacks. Biomarkers can give inaccurate results when the patient has low virulence infections or complications such as metallosis or inflammatory diseases, which can raise biomarker levels without the presence of infection. Culturing becomes problematic because of the difficulty in getting a representative swab as well as the often long necessary culture time and the low sensitivity for one positive culture in diagnosis. Other tests, such as PCR and radio-labeled WBC can be expensive to perform. For more affordable tests, such as LE strips, the results are easily invalidated by blood contamination and low sensitivity.

The current clinical “gold standard,” the MSIS criteria, does not differentiate between surgical sites, and it relies mainly on culturing as a diagnostic tool. Future directions include direct visualization of bacteria at a surgical site, allowing for real quantification of bacterial presence instead of relying on antibodies and fluorescent markers. This form of rapid visualization assay could provide a solution to many of the problems other tests have with detecting PJI, including low-virulence infections, ambiguous results because of metallosis and inflammatory diseases, and getting an accurate representation of the site in question. Overall, continued research into all methods of PJI detection as well as the determination criteria is vital to most efficiently diagnose and manage PJI.

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