Diagnosis of Pneumococcal Pneumonia: Current Pitfalls and the Way Forward

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Streptococcus pneumoniae is the most common cause of community-acquired pneumonia. However, it can also asymptotically colonize the upper respiratory tract. Because of the need to distinguish between S. pneumoniae that is simply colonizing the upper respiratory tract and S. pneumoniae that is causing pneumonia, accurate diagnosis of pneumococcal pneumonia is a challenging issue that still needs to be solved. Sputum Gram stains and culture are the first diagnostic step for identifying pneumococcal pneumonia and provide information on antibiotic susceptibility. However, these conventional methods are relatively slow and insensitive and show limited specificity. In the past decade, new diagnostic tools have been developed, particularly antigen (teichoic acid and capsular polysaccharides) and nucleic acid (ply, lytA, and Spn9802) detection assays. Use of the pneumococcal antigen detection methods along with biomarkers (C-reactive protein and procalcitonin) may enhance the specificity of diagnosis for pneumococcal pneumonia. This article provides an overview of current methods of diagnosing pneumococcal pneumonia and discusses new and future test methods that may provide the way forward for improving its diagnosis.

Key Words: Streptococcus pneumoniae, Pneumococcal pneumonia, Diagnosis, Polymerase chain reaction

Introduction

Streptococcus pneumoniae is responsible for a spectrum of diseases, including mild but common diseases such as otitis media, sinusitis, and non-bacteremic pneumonia and serious invasive pneumococcal diseases (IPD) such as bacteremia and meningitis. Among these diseases, pneumonia demands special attention because the incidence and mortality rates of community-acquired pneumonia (CAP) are high among the elderly. In the U.S., the annual incidence rate of CAP is estimated to be 5.2 to 6.1 cases per 1000 adults, and the mortality rate may reach 2.3% [1, 2]. The mortality rate for pneumococcal CAP is higher than for general CAP: < 2-5% in adults treated as outpatients, 12% of hospitalized patients, and ≥ 25% in elderly patients with bacteremia [3, 4]. Similarly, among Korean adults, the mortality rate for general CAP is estimated to be
3.2%, while the mortality rate for pneumococcal CAP is 5.9% [5]. Since *S. pneumoniae* is commonly responsible for the medically serious CAP [6], it is frequently simply referred to as “pneumococcus.”

Reflecting its status as a major human pathogen, *S. pneumoniae* was also one of the earliest pathogens to be discovered, and its microbiologic properties have been extensively investigated [7]. Pneumococci are Gram-positive bacteria with thick cell walls that contain teichoic acid (C-polysaccharide). They are capable of producing toxins (e.g., pneumolysin) as well as many surface antigens such as pneumococcal surface adhesin A (PsaA), pneumococcal choline binding protein A (PcpA), pneumococcal surface protein A (PspA), pneumococcal surface protein C (PspC), and pneumococcal autolysin A (LytA). LytA is the major autolysin responsible for lysis of pneumococci observed for mature pneumococcal cultures. However, the most prominent surface structure is the polysaccharide capsule, which is present on almost all virulent pneumococci. Pneumococci can express one of many (90+) polysaccharide capsule types that are serologically and biochemically distinct [8, 9]. Colony morphologies of two serotypes (3 and 37) are highly mucoid (Fig. 1) and distinct from other serotypes. As antibodies to the pneumococcal capsule are protective, the polysaccharide capsule is used in current vaccines. More recently, its genome sequences have been determined. Although no single gene that is unique and common to all isolates of *S. pneumoniae* has been reported, the genome sequences have been used to investigate pneumococcal evolution [10].

Despite its reputation as a pathogen, pneumococcus is a commensal that is often asymptomatically carried in the nasopharynxes of children and adults. Pneumococcal carriage occurs early in life, usually with a prevalence of about 30-60% in infants [11]; however, in some populations, > 90% of children are known to carry pneumococci [12]. The carriage rate may stay above 30-40% among children younger than 10 years of age, but it declines progressively until the rate reaches 1-10% among adults [11]. Since pneumococci are naturally present in the oro-nasopharyngeal space, the presence of pneumococcus in respiratory specimens does not necessarily indicate the presence of disease. Consequently, this commensalism must be incorporated in any diagnostic approaches to identifying pneumococcal infections.

In addition to *S. pneumoniae*, the oro- and nasopharynxes harbor Gram-negative rods and *Staphylococcus aureus* as well as many streptococcal species that resemble *S. pneumoniae*. The streptococcal species include *S. mitis*, *S. oralis*, and *S. pseudopneumoniae*, and are often referred as viridans species because they can produce α-hemolysis on blood agar plates similar to that produced by *S. pneumoniae* [13]. *S. mitis* and *S. oralis* can be responsible for subacute endocarditis and sepsis [14]. *S. pseudopneumoniae* is known to cause pneumonia or acute exacerbation in patients having a history of chronic obstructive pulmonary disease [15]. Gram-negative rods and staphylococci are mostly gentamicin-sensitive whereas viridans species and pneumococci are generally gentamicin-resistant. Consequently, the use of blood agar plates containing gentamicin improved the isolation of pneumococci and viridans species from respiratory specimens [16-19]. Although *S. pneumoniae* and viridans group are genetically related, *S. mitis* and *S. oralis* are generally resistant to optochin and bile-insoluble while *S. pneumoniae* is not (Fig. 2) [13]. *S. pseudopneumoniae* is generally bile-insoluble but is optochin-resistant only in 5% CO₂, but not in room air [15, 20].

In most clinical laboratories, *S. pneumoniae* is routinely identified by microscopic morphology (Gram-positive bacteria in the shape of slightly pointed cocci, usually in pairs), colony morphology, and characteristic phenotypes such as α-hemolysis observed on blood agar, catalase negativity, optochin susceptibility, and bile solubility [13]. These tests exclude most [13] but not all of the viridans species [21]. Conversely,
some *S. pneumoniae* strains may be bile-insoluble [22]. Also, viridans species may have *lytA* and *pneumolysin* genes, thereby limiting the usefulness of the genetic tests for *S. pneumoniae* [23]. Biochemical tests based on teichoic acid may also fail to distinguish *S. pneumoniae* because some viridans species can produce teichoic acid that is identical to pneumococcal teichoic acid. More recently, multi-locus sequence typing (MLST) has been adapted to differentiate among these streptococcus species [24-26]. This approach is called multilocus sequence analysis (MLSA) and is more reliable than previous methods in distinguishing among the streptococcal species but has been used only as a research method.

Thus, routine phenotyping methods used in clinical laboratories may be inadequate to definitely distinguish *S. pneumoniae* from viridans species. This inadequacy was illustrated by recent studies that showed that many “pneumococci” carried in the nasopharynxes of HIV patients are actually viridans species that were previously misidentified as *S. pneumoniae* [24]. Also, the presence of *S. pneumoniae* in the respiratory tract may not be pathologic in a substantial number of cases. Thus, one must be aware of these fundamental limitations when interpreting epidemiologic as well as clinical studies. This article will review conventional techniques for diagnosing pneumococcal pneumonia and recent developments in other techniques such as urinary antigen, polymerase chain reaction, and serologic tests.

**Figure 2.** *S. pneumoniae* growth is inhibited around the paper disk containing optochin (A). The test tube containing *S. pneumoniae* shows a loss of turbidity in the presence of sodium deoxycholate (bile salts) due to bacterial lysis while the test tube containing viridans species is turbid (B).

**Current methods of diagnosing pneumococcal pneumonia**

Early accurate diagnosis and treatment of pneumonia are associated with improved survival; they also reduce costs associated with unnecessary investigations and complications due to inappropriate treatment [27, 28]. In a study of 14,000 elderly patients with pneumonia treated at over 3,500 hospitals, patients who received early antibiotic therapy (within 8 hours of hospital arrival) had a lower 30-day mortality [27]. In addition, etiologic diagnosis of CAP has an important effect on our ability to provide the optimal therapy for this disease, understand its societal burden, and assess the effectiveness of pneumococcal vaccines.

The diagnosis of pneumococcal pneumonia begins with establishing the presence of pneumonia. Pneumonia is often diagnosed by clinical symptoms and radiologic evidence. The clinical symptoms include cough or difficulty breathing plus tachypnea [29]. Although there could be limitations in these approaches, this review is focused on pneumococcal pneumonia, and readers are referred to other recent reviews for the limitations of pneumonia diagnosis [30-33].

1. **Conventional microbiological methods for establishing bacterial etiology**

*S. pneumoniae* is thought to be the most common etiologic
agent of bacterial CAP. The gold standard in establishing pneumococcal CAP is to isolate bacteria from a normally sterile body fluid and then identify that bacterium as *S. pneumoniae*. To identify pneumococcal CAP from body tissue, the ideal tissue is the lung tissue obtained by biopsy or bronchoscopy; consequently, lung biopsy was used previously in research settings [34]. In addition, transtracheal aspiration has been shown to have high yields of *S. pneumoniae* [35, 36]. However, since these approaches are not practical in clinical settings, most bacteriological confirmation is performed with readily available fluid samples such as peripheral sputum, blood samples, and pleural fluids. Advantages and limitations of using these types of samples are described below.

1) Sputum Gram stains and culture

Sputum Gram stains and culture are most often the first diagnostic step for pneumonia. Gram stains of sputum could be strongly suggestive of pneumococcal pneumonia if the sputum is of high-quality (< 10 squamous epithelial cells and > 25 polymorphonuclear cells at a magnification of 100×) and shows the predominant presence of Gram-positive diplococci [37, 38]. A meta-analysis of sputum studies performed in 1966-1993 found diverse sensitivity (15-100%) and specificity (11-100%) for this diagnostic method [39]. In comparison, prospective studies with high-quality sputum samples showed relatively high sensitivity (57-82%) and specificity (93-97%) [37, 40-42]. However, direct microscopic examination of Gram-stained specimens has some limitations in clinical practice; specifically, inadequate sputum collection and antimicrobial therapy before obtaining sputum specimens lead to low diagnostic yields. In the study by Musher et al. [37], sensitivity increased in inverse proportion to the duration of prior antibiotic therapy. In primary care settings, the availability of skilled microbiologists is another limitation of sputum Gram stain [39, 43]. Additional problem with this diagnostic method is that it may not be easy to collect good quality sputum from children.

Sputum culture can further assist etiology identification. However, the diagnostic sensitivity of the culture has been reported to be quite variable, ranging between 29 and 94% [36, 44-52]. Such variable results are related to inadequate sampling of sputum, delayed processing of sputum specimens, and prior antimicrobial therapy. Sputum culture was found to be negative in about 50% of patients with concurrent pneumococcal bacteremia [35, 36]. Also, Musher et al. [37] found that the sputum culture identified pneumococci only in 44% of the persons with bacteremic pneumococcal pneumonia with the sensitivity of sputum culture increasing to 93% if the comparison is confined to adequate sputum samples obtained before antimicrobial therapy. In addition, false positives may occur due to nasopharyngeal carriage, particularly among children, so the culture results should be interpreted along with the findings from Gram stain.

2) Blood and pleural fluid culture

Since blood and pleural effusions are normally sterile, isolating pneumococci from these normally sterile tissues provides the definite diagnosis of pneumococcal pneumonia. However, the clinical utility of this diagnostic method is limited because blood cultures are estimated to be positive for *S. pneumoniae* in less than 10% of patients who actually have pneumococcal pneumonia [53-56]. The low rate of culture may be due to pneumonia without bacteremia, autolysis of *S. pneumoniae* during the stationary growth phase, use of antibiotics before the culture, or inadequate samples (e.g., insufficient blood volume) [57]. To reduce autolysis, one may have to optimize the blood culture condition and/or use an antigen-detection method or a nucleic acid amplification test (NAAT) to analyze the culture-negative broth samples from patients with a high suspicion for pneumococcal pneumonia. Nevertheless, positive blood culture unambiguously establishes pneumococcal etiology.

Empyema was once considered rare in children, but it has been increasing worldwide over the last decade [58-61]. The incidence of empyema is also increasing in adults, *S. pneumoniae* is likely to cause empyema in healthy young adults, and the *Streptococcus milleri* group is the common pathogen causing empyema in the elderly with comorbidities [62]. Conventional bacterial culture of pleural fluid is often negative among children with pneumococcal pneumonia complicated by empyema. Similar to what happens in blood cultures, the culture rate of pleural fluid may be low due to autolysis release from pneumococci during the stationary growth phase, resulting in cell death [57].

2. Pneumococcal antigen detection methods

The conventional microbiological methods of pneumococcal pneumonia detection described above have several limitations. First, the culture is often falsely negative. Second, it takes several days to culture pneumococci. To overcome these limitations, laboratory methods are designed to detect pneumococcal molecules present in diverse tissue samples such as sputum, urine, breath, pleural fluids, and peripheral blood [63-66]. These methods have the theoretical advantage of rapidly detecting pneumococci even if they are non-viable after
antibiotic treatment. In addition, using a novel molecular target such as PcpA may be highly desirable because its presence can clearly distinguish pneumonia from carriage: its expression requires low manganese levels, with those levels being high in the nasopharynx but low in blood [67, 68]. Nevertheless, below we describe clinical experiences with teichoic acid and capsular polysaccharide detection as they have been central to most previous investigations.

1) Teichoic acid (C-polysaccharide) detection
Holmberg et al. [63] compared the sensitivity and specificity of using ELISA to detect teichoic acid with the sensitivity and specificity of using latex agglutination tests to detect capsular polysaccharide antigens in sputum specimens. Both ELISA and latex agglutination tests showed favorable sensitivity (95 and 86%, respectively) and specificity (94% for both detection methods) when sputum culture was used as the standard for comparison. Recently, detection of teichoic acid has become very popular following the introduction of a rapid immunochromatographic test (Binax NOW® S. pneumoniae assay) in 2003 (US Food and Drug Administration approval). This is a point-of-care test with high analytical sensitivity for C-polysaccharide in urine specimens.

Studies of urine samples from children showed inadequate specificity for the Binax NOW assay (50-60%) due to the high rate of nasopharyngeal carriage in children, resulting in urinary excretion of teichoic acid [69-73]. When concentrated children urine was used (25-fold concentration by ultra-filtration), the specificity of the Binax NOW assay decreased even more, to about 12% [70]. Because false-positive tests are common, the Binax NOW S. pneumoniae assay is not recommended for the diagnosis of pneumococcal pneumonia in children [33].

Although the specificity of the Binax NOW assay was disappointing for urine samples from children, its sensitivity (71-96%) and specificity (71-100%) for pneumococcal empyema were high when pleural fluid samples were examined (Table 1) [74-79]. Also, Gram stain of pleural fluid is rather insensitive for pneumococcal empyema, although it has a good positive predictive value [80]. Thus, the Binax NOW S. pneumoniae assay

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**Table 1. Review of Binax NOW S. pneumoniae assay performance for the diagnosis of pneumococcal empyema**

| Study          | Country | Study design | No. | Study population | Age | Standard methods | Sensitivity (%) | Specificity (%) |
|----------------|---------|--------------|-----|------------------|-----|------------------|----------------|----------------|
| Lee et al. [78] | Korea   | Retrospective | 62  | Children         | 69 months (median) | Culture (blood or pleural fluid) or Positive PCR for lytA | 77             | 94             |
| Picazo et al. [76] | Spain  | Prospective  | 217 | Children         | <15 years old | Culture (pleural fluid or CSF) or Positive PCR for lytA | 88             | 73             |
| Strachan et al. [74] | Australia | Prospective | 137 | Children         | 4.9 years (mean) | Culture (blood or pleural fluid) or Positive PCR for lytA | 84             | 94             |
| Martinon-Torres et al. [77] | Spain | Prospective | 55  | Children         | 6.5 years (mean) | Culture (blood or pleural fluid) or Positive PCR for ply/w zg | 96             | 100            |
| Casado Flores et al. [79] | Spain | Prospective | 76  | Children         | 3.5 year (median) | Culture (blood or pleural fluid) or Positive PCR for ply | 88             | 71             |
| Porcel et al. [75] | Spain | Retrospective | 140 | Adults           | 56 years (mean) | Culture (blood or pleural fluid) | 71             | 93             |

PCR, polymerase chain reaction.
*The number includes 12 CSF (cerebrospinal fluid) samples.

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**Table 2. Sensitivity and specificity of Binax NOW urinary antigen test for the diagnosis of pneumococcal pneumonia based on meta-analysis**

| Study           | Published year | No. of included studies | Study population | Sensitivity (95% CI) | Specificity (95% CI) | Increased diagnostic yield* |
|-----------------|----------------|-------------------------|------------------|----------------------|----------------------|-----------------------------|
| Boulware et al. [83] | 2007        | 24                      | Adults           | 74% (72-77%)         | 94% (93-95%)         | 23% (range: 10-59%)         |
| Sinclair et al. [81]  | 2013        | 27                      | Adults           | 74% (67-82%)         | 97% (93-100%)        | -                           |
| Horita et al. [82]   | 2013        | 10                      | Adults           | 75% (71-79%)         | 95% (92-98%)         | -                           |
| Said et al. [6]      | 2013        | 35                      | Adults           | -                    | -                    | 11% (95% CI:10-14%)         |

CI, confidence interval.
*Increased diagnostic yield means additional diagnosis made by using Binax NOW urinary antigen test beyond conventional microbiological methods.
may be practically useful for the diagnosis of pneumococcal empyema using pleural fluid samples. The study by Le Monnier et al. [80] also found that NAAT could improve the detection of etiologic agents in 43% of patients with pneumococcal empyema.

Clinical studies with adults, however, produced better results. According to the recent meta-analyses for adults, the estimated sensitivity and specificity of the Binax NOW \textit{S. pneumoniae} assay were 74-75% and 94-97%, respectively (Table 2) [6, 81-83]. The Binax NOW \textit{S. pneumoniae} assay increased the rate of etiologic diagnosis for CAP by 11-23% beyond conventional microbiological methods (Table 2) [6, 83]. In adult studies, the sensitivity of the Binax NOW \textit{S. pneumoniae} assay was higher for bacteremic pneumococcal pneumonia (77-92%) than for non-bacteremic pneumococcal pneumonia (52-78%) [84-88]. Although the Binax NOW \textit{S. pneumoniae} assay is less sensitive in patients with non-bacteremic pneumococcal pneumonia, it can be helpful for tailoring antibiotic therapy. The 2007 IDSA/ATS guidelines for the management of CAP recommend the use of the Binax NOW \textit{S. pneumoniae} assay in the following circumstances: intensive care admission, failure of outpatient antibiotic therapy, leukopenia, active alcohol abuse, asplenia, chronic severe liver diseases, and pleural effusion [31].

One of the important advantages of the Binax NOW \textit{S. pneumoniae} assay is that prior antibiotic use has less influence on the diagnostic yield. In the meta-analysis by Said et al. [6], prior antibiotic use reduced the relative diagnostic yield for blood cultures by 67% (95% confidence interval [CI], 53-77%), for sputum cultures by 34% (95% CI, 8-53%), and for the Binax NOW \textit{S. pneumoniae} assay only by 26% (95% CI, 0-44%). However, the Binax NOW \textit{S. pneumoniae} assay has some limitations. First, false negative results may occur in relation to low levels of the C-polysaccharide antigen [83]. Second, false positive results can be induced by cross-reaction with viridans species, asymptomatic nasopharyngeal colonization of pneumococci, and previous pneumococcal infections. Detectable amounts of teichoic acid excretion persist in 40-50% of patients’ urine samples for more than 1 month following pneumococcal illness [64, 83]. If concentrated urine samples are used, the C-polysaccharide antigen can be detected in 70% of urine samples at 1 month after pneumococcal infection [89]. Also, one should be aware that a recent pneumococcal vaccination may produce a false-positive Binax NOW result [90].

2) Capsular polysaccharide detection

Since capsular polysaccharide was first detected in the urine of patients with pneumococcal pneumonia in 1917 [91], there has been considerable interest in this detection method. Using enzyme-linked immunosorbent assay (ELISA) for the capsule, Schaffner et al. [92] found the capsular polysaccharide levels in the urine of these patients to be variable (> 500-2.5 ng/mL). In addition to depending on the severity of infection, the levels were dependent on serotypes since the serotype-specific clearance rates varied more than 250 fold [92].

Although the latex agglutination test was initially developed to detect urinary capsular polysaccharide of \textit{S. pneumoniae}, the test was of limited usefulness because it was not easy to perform and could not detect all the different capsule types [13]. However, the situation recently changed with the introduction of a multiplexed immunoassay system based on the Luminex® system and monoclonal antibodies [93, 94]. Although clinical experience is limited, such multiplexed serotype-specific urinary antigen detection (UAD) assays showed excellent sensitivity (79-97%) and specificity (99-100%) for the diagnosis of bacteremic pneumococcal pneumonia based on the capsule types included in the test [94, 95].

In addition, the serotype-specific UAD assays generate additional information that allows the identification of the pneumococcal serotype causing CAP. When Bewick et al. [93] evaluated the pneumococcal serotype distribution among patients with non-invasive CAP in the U.K.; PCV13-associated serotypes were identified in 57.4% of the cases. When Sherwin et al. [96] estimated the serotype distribution of pneumococcal pneumonia using serotype-specific UAD in U.S. adults aged ≥ 50 years; PCV13-associated serotypes were detected in 80% of patients with \textit{S. pneumoniae}-positive CAP or healthcare-associated pneumonia. However, since the aforementioned UAD assays were developed by vaccine companies to study the efficacy of conjugate vaccine against pneumonia, the current UAD assays cover only a limited number of serotypes (13-14 serotypes) and are not commercially available.

Similar to the Binax NOW \textit{S. pneumoniae} assay, multiplex immunoassays for capsular polysaccharides were also used to directly detect the serotypes of pneumococcal empyema [97, 98]. Recently, Yu et al. [65] determined pneumococcal serotypes directly from pleural fluids using a multiplex serotype-specific immunoassay covering a relatively large number (36 types) of serotypes. This study found that pneumococcal empyema was associated with serotypes 1, 3, 7F/7A, and 19A, which are serotypes that were not covered by the 7-valent conjugate vaccine [65]. The clinical usefulness of these assays is still limited because the assays cover only a limited number of serotypes. However, even an assay with limited serotypes
may be useful as the diagnostic yield of conventional microbiological methods is poor with pleural fluids.

3. Nucleic acid amplification tests

As nucleic acid amplification tests (NAAT) using the polymerase chain reaction (PCR) became popular, it was hoped that NAAT would yield a sensitive molecular diagnostic test of pneumococcal pneumonia. So far, the use of NAAT for pneumococcal pneumonia focused on multiple genetic targets in blood and respiratory tract samples. The pneumolysin gene (ply), autolysin gene (lytA), pneumococcal surface adhesin A gene (psaA), wzg/cpsA, and the Spn9802 gene fragment have been used as PCR targets to detect S. pneumoniae [99]. The current experiences with these samples are described below.

1) Whole blood

Initially, the ply gene was widely used for the detection of S. pneumoniae. Compared with blood culture results, several studies detected S. pneumoniae DNA in blood samples using the ply PCR, with sensitivities ranging from 35 to 100% [88, 100-103]. The first challenging issue for the ply PCR test is the inability to differentiate S. pneumoniae from other streptococcus species. This poor specificity occurs because the ply gene is also present in viridans group streptococci [104, 105]. Considering the low specificity of the ply PCR, the use of PCR to detect the lytA gene was introduced and was found to have a higher specificity [22, 104, 106]. Regardless of carrier status, the lytA gene was not detected in the blood of healthy subjects [107]. Although the lytA gene is present in both S. pneumoniae and S. mitis, lytA gene sequences vary more among streptococcus species than among S. pneumoniae strains [22]. Thus, the use of lytA PCR with an appropriately designed primer may have a high sensitivity and specificity for the detection of S. pneumoniae. Noticeably, some (less than 2%) clinical isolates of S. pneumoniae are bile-insoluble and produce negative results for lytA PCR due to the alteration of the gene sequence [22]. Similar to the lytA PCR, the Spn9802 PCR is highly specific for S. pneumoniae, but it may also be positive for S. pseudopneumoniae [108]. In addition to the target genes themselves, the part of the gene amplified by PCR is very important because of allelic variation between closely related species [109]. A few studies quantitated the blood pneumococcal load by real-time PCR, and high bacteremic DNA load was associated with increased mortality [110, 111]. Thus, quantitative real-time PCR may be predictive of pneumonia severity.

2) Respiratory tract specimens

When PCR targeting the ply gene was applied to lower respiratory tract specimens from patients with pneumococcal pneumonia, sensitivities ranged from 68 to 100% [112-115], but specificities were quite poor. Moreover, ply PCR positive rates for throat swab specimens were almost the same in both patients with pneumonia (55%) and in control subjects (58%) [113]. In addition to the poor specificity of the ply PCR, this difference might occur because non-quantitative PCR tests could not distinguish between true pneumococcal pneumonia and nasopharyngeal colonization. To overcome this limitation, quantitative real-time PCR has been introduced with better specific gene targets. Spn9802 real-time quantitative PCR showed 71% sensitivity and 100% specificity for the diagnosis of pneumococcal pneumonia at a cutoff of >10^4 copies/mL [108]. Similarly, in a study by Albrich et al. [116], lytA quantitative real-time PCR had a sensitivity of 82.2% and a specificity of 92.0% at a density of ≥ 8,000 copies/mL for distinguishing pneumococcal CAP from asymptomatic colonization. The proportion of CAP cases attributable to pneumococcus increased from 27.1 to 52.5% using that cutoff [116]. Thus, quantitative real-time PCR of respiratory specimens may be a promising diagnostic method for pneumococcal pneumonia.

4. Antibody responses to pneumococcal infection

Epidemiologic studies investigating antibody levels in serum samples obtained at both acute and convalescent phases of pneumococcal pneumonia should reveal the presence of the disease. Such serologic studies used diverse pneumococcal antigens, which include pneumolysin, C-polysaccharide, capsular polysaccharides, and PsA, a highly immunogenic lipoprotein. Compared with blood culture, IgG antibody to PsA showed good sensitivity (85-89%) and specificity (83-98%) for pneumococcal pneumonia diagnosis in Kenyan adults [117, 118]. However, the estimated sensitivity (42%) was insufficient in children [119]. Moreover, PsA is also present in the cell walls of viridans group streptococci [120]. As the method for measuring the level of antibodies to pneumococcal capsule has been standardized by the World Health Organization (WHO) [121], Tuerlinckx et al. [122] used the assay to prospectively evaluate the etiology of CAP in children. IgG ELISA detected pneumococcal pneumonia in 80% of cases in which the diagnosis of pneumococcal pneumonia was proven with positive blood or pleural fluid culture. Although serologic tests are not affected by prior antibiotic exposure and do not require isolating bacteria, they have the potential to detect antibodies against colonized pneumococci and are not thus routinely
available in clinical diagnostic laboratories. Thus, the serologic tests are primarily useful in epidemiological surveillance studies and have limited value in clinical practice.

5. Biomarkers

The concentrations of acute-phase reactants increase in response to infection, inflammation, and tissue injury, so they may be useful as biomarkers to distinguish bacterial infection from non-infectious conditions [123] or to predict prognosis or therapeutic options. An advantage of biomarkers is that the test results can be produced rapidly. Such biomarkers include CRP, procalcitonin and triggering receptor expressed on myeloid cells (TREM-1). TREM-1 is a member of the immunoglobulin superfamily, is up-regulated by microbial products [124] and may stimulate secretion of several cytokines and chemokines [125]. The serum soluble TREM-1 was associated with bacteremic CAP [125] while soluble TREM-1 in bronchoalveolar lavage fluid was associated with bacterial pneumonia [124]. Procalcitonin is produced by the paracellular cells of the thyroid and by the neuroendocrine cells of the lungs and intestines in response to pro-inflammatory stimuli, particularly stimuli of bacterial origin. Procalcitonin may become down-regulated in the presence of viral infections [126]. The level of CRP, an acute-phase protein synthesized by the liver, can rapidly increase during acute infections or inflammations. CRP binds to phosphocholine residue on pneumococcal teichoic acid or lipoteichoic acid and can activate the complement cascade [127].

At present, there are substantial clinical data with C-reactive protein (CRP) and procalcitonin. Studies of pneumococcal vaccines have shown that a serum CRP value of >120 mg/L and a procalcitonin level of >5 ng/mL would be useful to identify cases of pneumococcal pneumonia in children with non-specific changes on chest X-rays [128, 129]. Elevated procalcitonin levels were well correlated with positive PCR, serology, and chest X-rays but not with positive urinary Binax NOW S. pneumoniae assays in children [109]. Galetto-Lacour et al. [130] also evaluated procalcitonin and CRP as predictors of a pneumococcal etiology of CAP in hospitalized children. Elevated procalcitonin and CRP values were strongly associated with pneumococcal CAP, The sensitivity was 94.4% for procalcitonin (cutoff: 1.5 ng/mL) and 91.9% for CRP (cutoff: 100 mg/L). A procalcitonin level of ≥ 1.5 ng/mL combined with the detection of positive pneumococcal urinary antigens had a diagnostic probability for pneumococcal CAP of almost 80% (positive likelihood ratio: 4.59).

Similar to the results from pediatric studies, both procalcitonin and CRP tests were also useful in distinguishing bacterial pneumonia (including pneumococcal pneumonia) from viral pneumonia in adults [131]. Moreover, procalcitonin levels were significantly higher in adult patients with pneumococcal pneumonia than in those with other bacterial pneumonias [132] and correlated with the severity of the pneumonia [133]. For these reasons, some have suggested the inclusion of quantitative assays of CRP and procalcitonin when designing clinical trials to estimate vaccine efficacy or to study the benefit of antibiotic therapy [126, 134]. However, these acute-phase reactants cannot be used as the sole determinant when distinguishing between viral and bacterial causes of CAP [33]. Yet, when combined with other pneumococcal detection methods, biomarkers may significantly enhance the specificity of diagnosis for pneumococcal pneumonia.

Summary and the way forward

The ability to rapidly and accurately diagnose pneumococcal pneumonia would improve our ability to provide appropriate therapy, assess vaccine effectiveness, and estimate the disease burden. Yet, diagnosis of pneumococcal pneumonia remains challenging. Microbiologic studies of lung tissue or bronchoalveolar lavage (BAL) fluids (yielding ≥10⁶ cfu/mL) may be the “gold standard” for CAP diagnosis, but they are too invasive to be routinely used in clinical settings. A definitive diagnosis of pneumococcal pneumonia can be established if S. pneumoniae can be isolated from blood or pleural fluid of pneumonia patients [135]. However, their culture is often negative. Positive microscopic examination and culture of high-quality sputum samples provide strong evidence for pneumococcal pneumonia [99]. However, one must be aware that the results of such tests can depend on the quality of the specimen and that positive results may be confused with normal carriage.

Recently, an assay for teichoic acid (the Binax NOW assay) has been adopted as a part of diagnostic criteria. If an adult patient is positive in both the Binax NOW assay and microbiologic studies of high-quality sputum samples, the combined results would provide the physician with solid evidence for pneumococcal pneumonia. A procalcitonin test may also be adopted as a part of diagnostic criteria. Although not yet commercially available, the serotype-specific UAD assay seems to be more sensitive and specific than the Binax NOW S. pneumoniae assay [94, 96]. Thus, serotype-specific UAD may be a promising option for the diagnosis of pneumococcal pneumo-
nia. In addition, quantitative real-time PCRs of blood and lower respiratory tract specimens appear to be sensitive and specific for pneumococcal pneumonia [108, 116]. Taken together, while there is no magic test, the increased number of sensitive and specific tests should help clinicians diagnose pneumococcal pneumonia.

In the future, additional diagnostic options may include use of metabolomics approaches such as analysis of urinary metabolites with nuclear magnetic resonance (NMR) [136] or analyzing patients’ breath or bacterial isolates with mass spectrometry. A breath test is an attractive diagnostic option for pneumococcal pneumonia; indeed, volatile metabolites are being investigated as biomarkers for specific bacterial pathogens, but, at present, data are quite insufficient to determine if such a test would be sufficiently sensitive and specific [66]. On the other hand, matrix-assisted laser desorption ionization time-of-flight mass spectrometry (MALDI-TOF MS) is used in many clinical laboratories to identify bacteria. Although MALDI-TOF-based systems have been found to be easy-to-use, cost-effective, and fast with high-throughput performances, they have shown limitations in discriminating between S. pneumoniae and the viridians group [137, 138]. However, rapid improvements are being made and Vitex MS MALDI-TOF mass spectrometry system has been reported to perform as well as the conventional identification method (e.g., optochin test) [139].

Although S. pneumoniae is the most common etiologic agent for CAP, Haemophilus influenzae, S. aureus, Klebsiella pneumoniae, and atypical pathogens (Mycoplasma pneumoniae and respiratory viruses) are also important causes of CAP. In clinical practice, therefore, it is desirable to have a diagnostic test for these pathogens as well. Some commercial multiplexed NAATs are designed to detect the DNA of these pathogens. Park et al. [140] evaluated a multiplexed NAAT designed to detect six respiratory bacterial pathogens (S. pneumoniae, H. influenzae, M. pneumoniae, Chlamydia pneumoniae, Legionella pneumophila, and Bordetella pertussis) in children. The agreement rates between multiplex PCR and cultures for S. pneumoniae and H. influenzae were 92.9% and 91.1%, respectively. However, S. pneumoniae and H. influenzae can exist as commensal organisms of the upper respiratory tract, so quantitative multiplex NAAT would be required to better detect and differentiate the etiologic agent of CAP. Indeed, a quantitative multiplex PCR to detect S. pneumoniae, H. influenzae, and Neisseria meningitidis was described in 2010 [141]. Multiplex assays for all these pathogens should potentially revolutionize both the diagnosis of pneumonia and its treatments in the future.

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