Research paper

A comparative analysis of antibody repertoire against *Staphylococcus aureus* antigens in Patients with Deep-Seated versus Superficial staphylococcal Infections

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Immunoblot and an enzyme-linked immunosorbent assays were used to evaluate and compare IgG antibodies against *S. aureus* whole cell lysate, cell wall peptidoglycan and lipoteichoic acid to discriminate between deep-seated and superficial *S. aureus* infection. Serum samples were examined from patients with deep-seated (n = 25) and superficial (n = 25) *S. aureus* infections and 15 healthy controls. Patients with deep-seated infections exhibited a large number of immuno-reactive bands in their IgG immunoblot profile as compared to those with superficial infections and healthy controls. Anti-staphylococcal IgG antibodies that reacted with two antigens of apparent molecular weight 110 and 98 kDa were specifically present in 96% (24/25) of patients with deep-seated infections, and were absent in, superficial and control sera. Moreover other Gram-positive and Gram-negative bacteria did not share these two unique antigens. The ELISA assays revealed significantly elevated levels of IgG antibodies to peptidoglycan (PG) in 18 of 25 (72%) patients with deep infection and 15 of 20 (60%) patients with superficial staphylococcal infection. The elevated levels of IgG antibodies to teichoic acid (TA) antigen were detected in all (100%) deep-seated group patients and among 40% (10/25) patients with superficial infection. An increase in levels of antibodies to PG showed a positive correlation trend with levels of IgG antibodies to TA only in deep infection group. Thus immunoblot detection of these two unique antigens as well as detection of elevated antibodies against PG and TA may be useful for the discrimination of staphylococcal deep-seated and superficial infection in humans.

Keywords: *S. aureus*, Antibody response, Deep-seated infection, Immunoblot, ELISA

1. Introduction

*Staphylococcus aureus*, being a highly versatile organism, has the capacity to establish infections in a wide range of body sites. The infections caused by this species are often acute and pyogenic and, if untreated may spread to surrounding deeper tissues or to metastatic sites involving other organs, resulting in disseminated or deep-seated infections which are life threatening [1]. It is one of the most successful human pathogens with ability to colonize and infect both hospitalized patients with or without compromised host defenses and healthy immunologically competent people in the community [2]. Further, the recent reports of reduced susceptibility to vancomycin [3] highlight the importance of understanding molecular epidemiology of *S. aureus* infections.

*S. aureus* has a diverse arsenal of components and products that contribute to the pathogenesis of infections. A great deal is known about the contribution of these bacterial factors to the development of infection, [1] considerably less is known about the host factors. Moreover approximately 30% of healthy individuals are colonized by *S. aureus*, usually in their anterior nares which serves as the major reservoir form where the organism can spread to the other parts of the body [4]. The proposed pathogenesis for endogenous infection would be that from the nose, the skin become colonized causing subsequent infection in patients with impaired skin site [5, 6]. Recent studies have shown that mucin appears to be a critical host surface, which is colonized, involves the interactions between staphylococcal proteins and mucin carbohydrate [4, 7]. The role of other factors such as secretory IgA is not clear. Immunity to *S. aureus* infections and its role in pathogenesis remain poorly understood [8]. Although, there appears to be a little resistance to mucosal colonization by *S. aureus*, healthy individuals have a high degree of innate resistance to invasive staphylococcal infections [9]. Natural immunity in the host is attributed to epidermal and mucosal surface barriers and to intact cellular and humoral immune defense. Due to repeated natural exposure to *S. aureus* (as a constituent of normal flora) antibodies to *S. aureus* exoproteins and cell wall components peptidoglycan (PG) [10], teichoic acid (TA) [11] and capsule are prevalent in humans [9]. But titers of these anti-staphylococcal antibodies are elevated after severe infections and this view is supported by a number of studies, which have shown that significantly elevated levels of antibodies to TA or PG can be detected in the vast majority of patients with *S. aureus* invasive infections [12, 13]. Many of these studies have demonstrated the rise in titer of antibodies against TA and PG during infections, but they in general, failed to reveal a correlation between antibody titer and immunity to these infections [10]. One exception is that antibodies to the staphylococcal exoprotein TSST-1 are associated with immunity to staphylococcal toxic shock syndrome in humans [14]. The data regarding the presence of antibodies during superficial *S. aureus* infections are not evaluated systematically.

In this study we therefore aimed at analyzing antibody responses against *S. aureus* whole cell lysate and its cell wall antigens in patients with deep-seated and...
superficial staphylococcal infection. The comparison of antibody levels and responses might thus identify the patients with more risk of life threatening deep-seated infection as compare to self-limiting superficial infections. Furthermore the presence of an antibody response against these antigens may also add diagnostic information when patients with putative invasive S. aureus infection are being evaluated.

2. Material and Methods

2.1. Study population

This study was approved by Institutional Ethics Committee and Dean’s Doctoral Committee. A total of 50 patients were enrolled in the study. Group I include 25 consecutive cases admitted in Nehru Hospital, PGIMER, Chandigarh, India with clinical evidence of sepsis and bacteriological evidence of deep-seated S. aureus infections as evidenced by isolation of organism from blood or an aspirate from deep-seated normally sterile site. Cases of prior trauma or penetrating injury were excluded. Group II consisted of another 25 consecutive cases with evidence of bacteriological proven S. aureus infection limited to skin and/or subcutaneous soft tissue without any clinical or bacteriological evidence of invasion into deeper tissues. Fifteen healthy persons without any history of sepsis in the preceding three months were taken as controls. Single serum samples were obtained from all the subjects including the patients and healthy controls and preserved at -20°C till further use.

2.2. Bacterial strains

A total of 50 clinical isolate of S. aureus from the two groups of patients were obtained from clinical bacteriology laboratory, department of Medical Microbiology, PGIMER, Chandigarh, India. A standard strain of S. aureus (ATCC-12600) was obtained from Microbial Type Culture Collection (MTCC) center of Institute of Microbial Technology, Sector 39, Chandigarh. This strain was used as reference strain.

The clinical isolates were identified on the basis of colony characteristics, gram stain morphology, coagulase production and mannitol fermentation [15]. All the strains were preserved in semisolid agar butts at 4°C and in BHI broth (Hi Media) with 15% glycerol at –70°C till further use.

2.3. Preparation of whole cell lysate

All clinical isolates and the reference strain were plated on BHI agar plates and incubated at 37°C for 24h. A sweep of 4-6 colonies from these plates were inoculated into flasks containing 10ml of BHI broth and incubated at 37°C overnight in an orbital shaker. Subsequently, 1 mL of this growth culture was added to 50 mL of fresh medium and incubated at 37°C with constant shaking (150 rpm) for 8 to 10 hours, and bacterial growth was monitored by taking aliquots and measuring OD600 at various time intervals. The culture was stopped at mid-logarithmic phase (OD600 ≤ 0.5). The resulting broth cultures were centrifuged at 10,000g for 15 min at 4°C. The supernatant was discarded and the pellets were washed twice in sterile PBS (pH 7.2). The pellets were finally resuspended in equal volume of sterile PBS and disrupted by shaking with glass beads (Braun; cell disruptor) for 2-3 min under constant cooling by liquid CO2 till 90% of the cells were broken. The unbroken cells were deposited by centrifugation at 12,000g for 30 m at 4°C and the supernatant was preserved at -70°C till further use. The protein content of the samples was estimated by method of Bradford [16] and adjusted to give a final conc. of 2-3mg/ml.

2.4. Sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE)

Whole-cell protein profile of all S. aureus isolates was analysed by SDS-PAGE following the discontinuous buffer system of Laemml [17]. Samples were prepared for PAGE by mixing in proportions of one part sample to 3 parts of sample buffer and boiling for 5 min. Solubilised samples (20µl) were applied to wells in a 4% acrylamide stacking gel over a 10% acrylamide separating gel. Electrophoresis was performed using a Bio-Rad Mini-Protein-II® apparatus with vertical slab gel 7cm (L) x 8cm (W) x 0.75 mm (T). The gels were run at constant voltage of 175V until the bromophenol blue dye had reached the bottom over a period of 45 min-1hr. Broad range molecular weight markers (Bio-Rad) were also run for molecular weight estimation of bands of interest. After the completion of electrophoresis the gels were removed and stained with Coomassie Brilliant Blue R-250. The gels were photographed and protein profiles of the isolates were compared.

2.5 Immunoblotting

The whole cell lysate of the clinical isolates and reference strain were run in SDS-PAGE as mentioned earlier using a Bio-Rad Mini-PROTEAN II® apparatus following manufacturer’s instructions. Gels were equilibrated in transfer buffer and the separated proteins were transferred onto nitrocellulose (NC) paper (pore diameter, 45µm) as described by electrophoretic transfer method of Towbin et al. [18], using a Bio-Rad Mini TransBlot® cell at 100mA for 90 m at 4°C in transfer buffer.

After transfer the NC paper was cut vertically into strips, which were washed twice in PBS and soaked in blocking buffer overnight at 4°C. For characterization of serological response of patients to blotted and blocked NCP strips, patient sera diluted 1:100 in PBST containing 1% BSA were added to individual strips. After incubation at 37°C for 1 hr at 37°C, the strips were washed on a rotatory shaker with five changes of PBST for 10 m each. Anti-human IgG-HRPO conjugate (Dako) diluted 1:1000 in PBST-BSA were added to each strip and incubated for another one hour at 37°C. Following five washings for 10 m each the strips were developed by soaking for 5 m in developing solution containing 0.05%(w/v) diamobenzidine tetrahydrochloride and 1% H2O2 in PBS. The blots were dried and photographed.

2.6 Preparation of S. aureus cell wall antigens

S. aureus grown on nutrient agar plates overnight were resuspended in distilled water and heated at 70°C for 1 hr. The bacteria were centrifuged, rinsed twice with distilled water, twice with acetone and dried. Cells were disrupted in cell homogenizer (Braun, Germany) with glass beads (0.17-0.18 mm dia). Residues upon centrifugation at 3,000 rpm for 10 min were discarded and from the supernatant cell wall were sedimented by centrifugation at 13,000 rpm for 25 min and rinsed 3 times with distilled water. Cell wall pellet was resuspended in phosphate buffer (pH 7.6) containing trypsin (200 µg/ml), RNase (100 µg/ml) and DNase (50 µg/ml) and incubated at
with deep-seated infections were admitted in different

group was 21.1 years (range, 4 to 52 years). The patients

pyoderma (n = 11), folliculitis (n = 8), subcutaneous

abscesses (n= 5) and sty (n = 1) and the mean age in this

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2.6.1 Teichoic Acid extraction

Teichoic acid was extracted from cell wall fractions

with 5% TCA at room temp 3 times overnight on shaker

(1g cell wall per 250 ml, 160 ml, and 100 ml of TCA). Three

aliquots of supernatant were pooled and extracted 3 times

with double volume of ethyl ether each time and then

precipitated with 3 volumes of acetone at 4°C for 24 hrs.

Delicate precipitates of teichoic acid were centrifuged and

was washed with acetone and dried.

2.6.2 Peptidoglycan preparation

The residue remaining from cell wall extraction with

TCA was heated at 90°C for 15 min, washed with water 3
times, followed by acetone and dried. This fraction was

used as peptidoglycan in all experiments as described by

Kotani et al [19, 20].

In addition commercially available Staphylococcus

aureus peptidoglycan and teichoic acid purchased from

Sigma were also included in ELISA experiments.

2.7 ELISA

The modification of the technique originally
described by Engvall and Perlmann [21] was used to
quantitate IgG antibodies against cell walls peptidoglycan
(PG) and teichoic acid (TA) in patients and healthy control
sera. One hundred microliters of antigen solution
(10µg/ml) in carbonate buffer (pH 9.6) was added to flat
bottom microtiter plate (Maxisorp, Nunc, Denmark) wells
and left to adsorb for 1 h at 37°C and then overnight at
4°C. The plates were then washed (SLT washer, Austria)
in PBST. The unbound sites on the plastic surface were
blocked with 200µl of PBS containing 2% BSA. The
blocking agent was left for 6 h at room temperature and
then overnight at 4°C. The plates were washed 3 times
with PBST and 100µl of patient serum diluted 1:1000 in
PBST containing 1% BSA was added to each well and the
plate was kept for incubation at 37°C for 1h. After
washing (five times) 100µl of anti-human IgG-HRP
conjugate diluted in PBST containing 1% BSA was added
and incubated at 37°C for 1 h. After washing 100µl of
substrate containing 10mg of orthophenylene diamine
(OPD) in 25ml of citrate-phosphate buffer was added to
each well. After 15-20 mins at room temperature, the
reaction was stopped with 50µl of 1M H2SO4. The plates
were read in an automated ELISA reader (340 ABC, SLT,
Austria) at 492 nm using a reference wavelength of 620
nm. The results of the assay were expressed as the mean
optical density at 492 nm of the duplicate values.
Statistical analysis was performed using ANOVA and p <
0.05 were considered to be significant.

3. Results

3.1 Demographics

A total of 50 patients and 15 healthy controls were
enrolled in the study. The patients with deep-seated
infections had deep abscesses (n = 8), sepsis (n = 5),
endocarditis (n = 4), pyomyositis (n = 3), septic arthritis
and osteomyelitis (n = 2 each) and empyema (n = 1).
The mean age of patients in this group was 24.5 years (range, 4
to 65 years). Patients with superficial infections had
pyoderma (n = 11), folliculitis (n = 8), subcutaneous
abscesses (n= 5) and sty (n = 1) and the mean age in this

group was 21.1 years (range, 4 to 52 years). The patients

with deep-seated infections were admitted in different

wards of Nehru Hospital, PGIMER, Chandigarh during a
period of 2 years from Jan 1999 to March 2001, while the
majority (80%) of patients with superficial infections
enrolled in the study were taken from skin OPD over a
period of 3 months March 2001 to May 2001. The S. aureus
strains from clinical specimens of hospitalized patients
were recovered within two to three days after
hospitalization, whereas strains were isolated next day
from those appearing in OPDs (particularly skin). Serum
samples were also collected same day before the start of
antibiotic therapy.

3.2 S. aureus strains causing deep-seated and superficial
infection had identical protein profile

Recently, S. aureus strains have been shown to have
an association to the type of infection [22]. In order to
compare the whole cell protein profile of S. aureus strains
associated with deep-seated and superficial infection,
whole cell lysates of isolates of both the groups were
subjected to SDS-PAGE. Both deep and superficial isolates
showed almost identical protein profile which consisted of
25-30 bands with major bands having molecular weights
of approximately 120, 92, 80, 66, 55, 42, 36, 29, 24, 20, 18
and 16 kDa (Fig 1).

The protein profiles of isolates within the same
group were grossly similar. More than 90% of the bands
were qualitatively and quantitatively identical with minor
quantitative difference in the intensity of the others. No
significant inter strain variation was found among the
deep and superficial isolates regarding their whole cell
protein profile.

3.3 Immunoblot profile can discriminate between deep-
seated and superficial infection

The IgG antibody response to whole cell lysate of
Staphylococcus aureus during infection was examined
qualitatively by immunoblot analysis of sera from patients
of both the group. Figure 2 shows the IgG immunoblot
profile of the sera of patients with deep-seated and
superficial infections respectively. Sera from patients with
deep-seated staphylococcal infections exhibited a large
number of bands in their IgG western blot profile (Fig 2A)
as compared to those with superficial infections (Fig 2B).
The banding patterns within each group of patients were
grossly similar with minor individual variations. The most
prominent band present in both superficial and deep
patient sera had molecular weight of approximately 60
kDa. This band was also detected in healthy controls (data
not shown). In addition to this common band, the
immunoblot profile of deep-seated group of patients
showed the presence of two unique bands with molecular
weights of 110 kDa and 98 kDa which were detected in
96% of patients of this group but were totally absent in
patients with superficial infection and healthy individuals.

3.4 The 110 and 98 kDa immunodominant peptide of S.
aureus are not shared by other bacteria

In order to demonstrate the species specificity of
these two unique bands present in IgG immunoblot
profile of patient sera from deep-seated infections, pooled
sera from patients of both the groups and healthy controls
were checked against whole cell proteins of other Gram-
positive [S. epidermidis and clinical isolates of coagulase-
negative staphylococci (CONS)] and Gram-negative (E.
coli, Klebsiella spp) bacteria. The results of this experiment
showed that these two bands were present only in cases
with deep seated infections (Fig. 3A) and absent from those with superficial infections (Fig. 3B) and healthy controls (Fig. 3C). Moreover these antigens were not shared by other Gram-positive and Gram-negative bacteria tested.

3.5. Anti-staphylococcal antibodies react with PG and TA

We observed that levels of anti-staphylococcal IgG antibodies were very similar when total bacterial lysate or whole 

*S. aureus* cells were used as coating antigens (data not shown). These results suggested that the majority of antistaphylococcal IgGs reacted with the surface of the pathogen. TA and PG, the two most abundant cell wall components, have been known to be highly immunogenic in humans. Consistent with this notion, the quantitative measurement of IgG antibodies to peptidoglycan and teichoic acid in sera from patients and healthy controls was checked by ELISA.

The sera obtained from both patients and healthy controls showed great variability in levels of antibodies against *S. aureus* cell wall antigens (table 1). Sera from patients were considered positive if the OD exceeded the mean OD of healthy control sera plus one SD. Elevated levels of IgG antibodies to teichoic acid antigen were detected in 100% (25/25) patients with deep-seated infections and 40% (10/25) of patients with superficial infection. About 72% (18/25) of patients among deep group and 60% (14/25) among superficial had significantly elevated antibodies against peptidoglycan (Fig. 4). The levels of antibodies against cell wall antigens also varied within each group, amongst deep infection group 4 of 25 patients who had staphylococcal endocarditis and two patients with osteomyelitis were found to have higher antibodies to TA and PG, as compare to other patients in this group. Similarly patients with folliculitis had higher levels of antibodies to PG among superficial group. Similar results were obtained when we used the commercially available PG and TA

3.6 Correlation coefficients

We analyzed the correlation of levels of anti-peptidoglycan and teichoic acid antibodies in sera from patients with deep-seated and superficial staphylococcal infections. Sera from patients with deep-seated infections showed weak positive correlation trend between antibodies to PG and TA ($p < 0.1$), where as on the other hand no significant correlation was detected among patients with superficial infections regarding anti PG and TA antibody levels (Fig. 5).

4. Discussion

*Staphylococcus aureus* is a major human pathogen causing significant morbidity and mortality in both community and hospital acquired infections [1]. It causes a diverse array of infections ranging from relatively minor skin and wound infections to more serious and life threatening disease such as pneumonia, endocarditis, osteomyelitis, arthritis and sepsis [2]. Here in this study we investigated the prevalence of antibodies to the well-characterized staphylococcal cell wall antigens PG and TA and total cell lysate in sera from patients with deep-seated and superficial *S. aureus* infections. We demonstrated that the patients with deep-seated infection can be discriminated from those with superficial infection, which might thus identify the patients with more risk of life threatening deep-seated infection as compare to self-limiting superficial infections.

Immunoblotting of bacterial whole cell polypeptides, by the western blot technique has been used to examine the immune response of individuals to the large number of polypeptides found in whole cell extract pathogenic bacteria [23] including *Staphylococcus aureus* [24]. In a study of six patients with *S. aureus* endocarditis, antibodies to *S. aureus* and *E.coli* heat shock proteins were detected by western immunobLOTS, but a similar response was observed in sera from patients with other forms of endocarditis. While heat shock proteins may not be as specific as staphylococcal antigens studied by Colgue-Navarro and colleagues as these results emphasise that serological response to staphylococcal infection is complex partly because individuals are exposed to *S. aureus* from birth [25]. Our study has clearly demonstrated that all the individuals (patients and controls) investigated produce circulating antibodies directed against a large number of *S. aureus* polypeptides. This finding is in agreement with those of other workers who have documented those most human sera posses a variety of anti-staphylococcal antibodies [26]. Despite the complexity of immune response many polypeptides in the cell extract gave negative reaction in the western blot analysis. It is likely that the immune response against some polypeptides may be poor because they are intrinsically poor antigens, because of their structure or other factors [27]. Western blot of IgG response of patients with deep-seated infections showed antibodies against a large number of polypeptides as compared to those with superficial infections. In immunoblot profile of deep-seated patients we identified two unique polypeptides of molecular weights of approximately 110 kDa and 98 kDa, which were detected exclusively in 96% of patients with deep-seated infection, and they were absent in superficial group and healthy control group. These two unique bands are likely to be of diagnostic value. The diagnostic efficacy of antibody detection tests was not the primary objective of this study and these findings need to be further evaluated in larger group of patients and healthy individuals. In addition further characterization of these two polypeptide antigens is required to evaluate their potential role in diagnosis of these two groups of patients.

In recent years attention has been focused on the measurement of antibody to cell wall antigens in patients with proven *S. aureus* infection [28]. Teichoic acid and peptidoglycan are the major components of staphylococcal cell wall and they are known to induce inflammatory response in humans [29]. Antibodies against PG and TA antigens have been successfully titred in serological assays [28, 30, 31]. Humoral immune response to *S. aureus* infections is complex and despite the fact that α-toxin, teichoic acid and lipase are widely recognized as staphylococcal virulence determinants, the production of antibodies to one or more antigens is variable. Colgue-Navarro and colleagues acknowledge that the kinetics of the antibody response differs greatly between patients [25]. In our study, we also observed variable antibody response against *S. aureus* cell wall antigens peptidoglycan and teichoic acid between both the groups of patients and healthy controls. Studies are available which have shown the increased IgG titer against teichoic acid [32] in patients with deep-seated staphylococcal infections (especially in cases of endocarditis and osteomyelitis). We also observed that all (25/25) of the patients with deep-seated infection had elevated levels of IgG antibodies against TA, even with in
the group of patients with endocarditis and osteomyelitis showed higher levels than rest of the group. Two healthy controls also exhibited higher antibodies against this antigen. Previous staphylococcal infections or non-specific antibody responses may be a significant factor in partly explaining the high initial antibody levels [33, 34]. In contrast, failure to mount an antibody response against PG in some of the patients with deep infection might be due to impaired immunity of advancing age, poor expression of virulence determinants by the infecting strain or changes in the immune response resulting from infection itself.

Verbrugh et al. [35], in 1986 undertook one study to compare the antibody response to α-toxin, teichoic acid and peptidoglycan in patients with complicated and uncomplicated infections. Neither single test was able to detect complicated bacteraemia in all patients nor there do any test that was totally specific in this respect. Sensitivities ranged from 64% for assay of Abs against teichoic acid to 72% for assays of antibody to peptidoglycan and α-toxin. The specificity of assays for antibody to cell wall antigens (peptidoglycan and teichoic acid) was 79% whereas this specificity was 50% for α-toxin. Our data indicated a positive correlation trend between raised levels of antibodies to PG and TA among deep infection group, but surprisingly no correlation was detected among superficial infection group.

In conclusion our data indicated that patients with deep-seated and superficial staphylococcal infection can be discriminated on the basis of their immunoblot profile against total S. aureus cell lysate as well as their serum IgG response to cell wall antigens peptidoglycan and teichoic acid. Whether the presence of antibodies against these cell wall antigens of S. aureus decreases or enhances the outcome of staphylococcal infection is still not clear.

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Conflict of interest

None declared.

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Tables and Figures

Table 1. IgG antibody levels against PG and TA in sera from healthy individuals and patients with deep-seated and superficial staphylococcal infections.

| S. aureus Cell Wall Antigens | Healthy Individuals (n = 15) | Patients with Superficial Infection (n = 25) | Patients with Deep-seated Infection (n = 25) |
|-----------------------------|-------------------------------|---------------------------------------------|-----------------------------------------------|
|                            | Mean* SD* CV                  | Mean* SD* CV                                | Mean* SD* CV                                  |
| Peptidoglycan              | 0.78 0.29 0.37                | 1.05 0.57 0.54                              | 1.39 0.70 0.50                                |
| Teichoic acid              | 0.49 0.20 0.41                | 0.52 0.34 0.65                              | 1.29 0.63 0.48                                |

SD, Standard deviation; CV, coefficient of variation

* Mean and SD are given as ODs at 492 nm.

A. Deep Isolates

B. Superficial Isolates

Figure 1. Whole-cell protein profile of representative S. aureus strains. Clinical isolates of S. aureus were taken from patients with deep-seated (A) and superficial staphylococcal infection (B). The crude protein extracts were separated by SDS–PAGE and stained with Coomassie blue. Lane M; molecular weight marker; Lane A; standard S. aureus strain (ATCC 12600); Lane 1-6; whole cell lysates of representative S. aureus isolates. Figures are representative of two independent experiments.
Figure 2. Immunoblot profiles of patients with deep-seated and superficial *S. aureus* infection. Whole cell protein lysate of *S. aureus* (ATCC 12600) was separated by SDS–PAGE and transferred onto nitrocellulose membranes which were probed with serum (1:1000) from patients with deep-seated (A) and superficial (B) infection. Each lane represents serum from individual patients. Serum samples were drawn 2-3 day after detection of *S. aureus* in clinical specimens. Results are representative of two independent experiments.

Figure 3. Western blot analysis of *S. aureus* immunodominant antigens. Whole cell protein lysate of *S. epidermidis* (lane 1), *S. aureus* (lane 2), clinical isolates of coagulase negative staphylococci (lane 3-5), *E. coli* (lane 5) and *Klebsiella sp* (lane 6) were separated by SDS–PAGE and transferred onto nitrocellulose membranes and probed with pooled serum (1:1000) from patients with deep-seated (A) and superficial (B) infection and healthy controls (C). Results are representative of three independent experiments.
Figure 4. Levels of antistaphylococcal IgG antibodies. Antibodies against *S. aureus* cell wall antigen peptidoglycan (A) and teichoic acid (B) were measured ELISA in sera (1:1000) from healthy individuals and patients with superficial and deep-seated infection. Violin plots were constructed on the basis of OD 492 nm to show the distribution IgG antibody levels in each group. Data are the representative of three independent experiments.

Figure 5. Correlation plots of antibodies to PG and TA. Antibody levels against peptidoglycan and teichoic acid in patients with superficial (A) and those with deep-seated (B) as measured by ELISA were plotted against each other to elucidate correlation. Coefficient of correlation was determined by $r^2$ values as shown. Results represent the values of experiment in duplicates.