Promotion of opsonization by antibodies and phagocytosis of Gram-positive bacteria by a bifunctional polyacrylamide

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Abstract

This paper describes the application of a bifunctional polyacrylamide (pA–V–F) presenting both vancomycin and fluorescein groups, to modify the surfaces of multiple species of Gram-positive bacteria (Staphylococcus aureus, Staphylococcus epidermidis, Streptococcus pneumoniae, and Enterococcus faecalis) to control molecular recognition of these surfaces. The vancomycin groups allowed the specific recognition of a structural component of the bacterial cell wall: peptides terminated in D–Ala–D–Ala. The fluorescein groups allowed the imaging of binding of polymer to the surfaces of bacteria by fluorescence, and are representative, low molecular weight haptens; their recognition by anti-fluorescein antibodies provides proof-of-principle that bifunctional polymers can be used to introduce haptens onto the surface of the bacteria. Flow cytometry revealed that polymer-labeled S. aureus and S. pneumoniae were opsonized by anti-fluorescein antibodies ~20-fold more than were untreated bacteria; nearly all (~92%) polymer-labeled S. aureus, and a large (76%) fraction of polymer-labeled S. pneumoniae were opsonized. The bound antibodies then promoted phagocytosis of the bacteria by cultured J774 macrophage-like cells. Flow cytometry revealed that macrophages ingested S. aureus decorated with the polymer-antibody complexes ~2-fold more efficiently than S. aureus in control groups, in spite of the high background (caused by efficient antibody-independent ingestion of S. aureus by macrophages). This paper, thus, demonstrates the ability of a bifunctional polymer to carry out three distinct functions based on polyvalent molecular recognition: (i) recognition of the surface of Gram-positive bacteria, (ii) modification of this surface to generate specific binding sites recognized by an antibody, and (iii) promotion of phagocytosis of the opsonized bacteria. © 2006 Elsevier Ltd. All rights reserved.

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1. Introduction

Gram-positive bacteria are important to public health, and many can cause bacterial infections, which remain an important cause of morbidity and mortality. Immunocompromised patients are particularly susceptible to such infections [1]. The effective design of vaccines against these infections has proven challenging because bacterial surfaces are often poorly antigenic [2–5]. Further, the rise of drug-resistant strains of bacteria has highlighted the need for antibacterial agents with new mechanisms of action [6]. The approach described in this paper begins to address this need by demonstrating the effective conversion of the surfaces of several types of Gram-positive bacteria (Staphylococcus aureus, Staphylococcus epidermidis, Streptococcus pneumoniae, and Enterococcus faecalis) into surfaces presenting controllable recognition elements by treatment with a bifunctional polymer, pA–V–F (a polyacrylamide presenting both vancomycin and fluorescein groups as side chains) (Fig. 1). The polymer-labeled bacteria effectively couple with components of the immune system (antibodies and macrophages) for destruction. We believe that this approach will provide the basis for new
methods of promoting interaction between components of the immune system and surfaces (e.g., bacteria, viruses, cancer cells) not normally recognized by the immune system.

Several investigators have employed strategies, related to the one we have used here, to attach monomeric, bifunctional molecules noncovalently to the surface of pathogens or to surface-immobilized purified proteins that occur on the surfaces of pathogens, and to recruit components of the immune system to the bound species [7–10]. As an example, this bifunctional, antibody-targeting approach has been applied to the Gram-negative bacterium *Escherichia coli*, exploiting the binding of mannose by FimH, the adhesion protein on the tips of type 1 pili on the bacterial surface, using small molecule conjugates [9,10] or polymers [11]. Bertozzi and Bednarski [9,10] synthesized a mannose-biotin conjugate to which they bound avidin and an anti-avidin antibody to generate a complex that displayed both mannose and antibody molecules. They then demonstrated binding of the mannose–avidin–IgG complex to the surface of *E. coli* and subsequent recognition of the IgG portion of the complex to activate complement and enable recognition and killing of the bacteria by macrophages [10]. In a conceptually related study, Li et al. [11] used a polymer that presented both mannoside units (to bind to the bacterial surface) and \( \alpha\)-Gal units (with the potential to recruit innate anti-\( \alpha\)-Gal antibodies to the polymer adsorbed on bacteria). The investigators, however, did not demonstrate such targeting of antibodies to a bacterial surface.

We have used a related but distinct approach that, unlike the previous approaches [7–11], is not directed towards a specific protein on the surface of a particular pathogen, but rather, targets ligands that are broadly distributed in many organisms. The ligands that we have targeted are \( \delta\)-Ala–\( \delta\)-Ala residues at the carboxy termini of some of the peptides

![Fig. 1.](image)
in the peptidoglycan cell wall of many bacteria (Fig. 1(b)) [12]. These residues provide recognition sites, dispersed throughout the cell wall, for the antibiotic vancomycin, which binds to them with good affinity \((K_{d} \sim \mu \text{M})\) [13,14]. Vancomycin serves as an antibiotic by inhibiting cross-linking of the \(\text{D-Ala–D-Ala}\)-terminated peptides during formation and remodeling of the bacterial cell wall by binding to these peptides at the membrane surface, and thus inhibiting the action of transpeptidases [15]. Vancomycin-resistant Gram-positive bacteria evade the cytotoxic properties of vancomycin by expressing three essential enzymes in response to the binding of vancomycin to VanS, a transmembrane sensor protein kinase [6,16]. The net effect of the action of these enzymes is to replace \(\text{D-Ala–D-Ala}\) in the bacterial cell wall with \(\text{D-Ala–D-Lac}\); vancomycin binds to \(\text{D-Ala–D-Lac}\) with \(10^{3}\)-fold lower affinity \((K_{d} \sim \text{mM})\) than to \(\text{D-Ala–D-Ala}\), attenuating the potency of the antibiotic [16]. We included vancomycin groups in the bifunctional polymer, \(\text{pA–V–F}\), to promote specific binding of the polymer to the surfaces of Gram-positive bacteria (Fig. 1(c)).

The approach that we present here exploits polyvalency, which can generate a high-avidity binder for a given surface from low-affinity recognition units even if not all of the recognition units are involved directly in binding to their cognate surface receptors [17–25]. We previously characterized the polyvalent binding of \(\text{pA–V–F}\) to a self-assembled monolayer (SAM) that presented \(\text{D-Ala–D-Ala}\) groups using surface plasmon resonance spectroscopy (SPR) [26]. These studies demonstrated that \(\text{pA–V–F}\) formed kinetically stable \((k_{\text{off}} \sim 2 \times 10^{-6} \text{s}^{-1})\) complexes with high avidity at the surface presenting \(\text{D-Ala–D-Ala}\) groups. These studies also indicated that a thin (6.5 nm) layer of \(\text{pA–V–F}\) bound to the surface recruited anti-fluorescein antibodies to that surface almost as well as did a surface directly presenting fluorescein at a high surface density.

Our strategy in the current work was to use polyvalency and the molecular recognition of \(\text{D-Ala–D-Ala}\) by vancomycin (as opposed to the antibiotic properties of this molecule) to bind polymer with pendant vancomycin groups to the surfaces of Gram-positive bacteria; this strategy differs from that of a previous report on the use of monofunctional polymers of vancomycin as potential antibiotics [27]. The effective grafting of the polymer to the bacterial surface gives us the ability to modify the molecules displayed at the surface of the bacteria by incorporating additional groups (i.e., in addition to vancomycin) of arbitrary structure into the polymer.

As a model system, we elected to use fluorescein as a second group: this polymer \(\text{pA–V–F}\) allowed bacteria that had their surface modified by adsorption of polymer to be visualized by fluorescence, and also promoted subsequent binding of anti-fluorescein antibodies to the modified bacteria (Figs. 1(c) and (d)). These opsonized bacteria were then recognized and ingested by macrophages (Figs. 1(e) and (f)). To our knowledge, this paper is the first to report the coupling of antibodies and macrophages to the surfaces of pathogens using a bifunctional polymer.

2. Materials and methods

2.1. Materials

Unless otherwise specified, reagents were obtained from Sigma-Aldrich (St. Louis, MO) of the highest quality available and used as received. Fluorescein cadaverine, Syto-63, slow anti-flade reagent, rabbit R-phycocerythrin-conjugated anti-fluorescein IgG (polyclonal), and goat R-phycocerythrin-conjugated anti-mouse IgG (polyclonal) were obtained from Molecular Probes (Eugene, OR). Mouse anti-fluorescein IgG (clone 1.BB.75) was obtained from BD Biosciences Pharmingen (Chicago, IL). Dialysis cassettes (10,000 MWCO, 3–15 mL) were obtained from Pierce (Rockford, IL). Diffi-Quik stain set was obtained from Dade Behring (Marburg, Germany). E. coli (ATCC 53506), S. epidermidis (ATCC 35984), E. faecalis (ATCC 49332), E. faecalis VanB (ATCC 51299), and J774A.1 cells (ATCC TIB-67) were purchased from the American Type Culture Center (ATCC). S. aureus strain JL-243 was kindly provided by Dr. Jean Lee of Harvard Medical School. Confocal fluorescence microscopy was performed on a Leica confocal imaging system equipped with an Ar–Kr laser and using a 63× (N.A. = 1.4) or 100× (N.A. = 1.4) oil immersion objective. Flow cytometry was performed on a Coulter® Epics® Elite ESP flow cytometer.

2.2. Polymers

The bifunctional polycarylamide, \(\text{pA–V–F}\), and the control polyacrylamide containing only fluorescein groups, \(\text{pF–F}\), were synthesized as previously reported [26]. Briefly, fluorescein cadaverine was added to a solution of poly(s-acryloyloxysoximide) (PNAS) [18] in DMF and allowed to react. A vancomycin derivative (vancomycin conjugated to 1,4-diaminobutane [28,29]) was then added, and finally the reaction was quenched with ammonium hydroxide (30%), which reacted with remaining NHS esters to yield amides [18]. \(\text{pF–F}\) was synthesized as above for \(\text{pA–V–F}\) with the omission of the vancomycin derivative. The polymers were dialyzed against water and stored frozen at −20°C, or lyophilized and stored at −20°C. The characteristics of the polymer backbone were determined by saponifying PNAS to polyacrylic acid and analyzing this material by GPC: \(M_{s} = 96,500; M_{w} = 65,000; PDI = 1.48\); degree of polymerization ≈ 900. The side-chain % substitution of fluorescein and vancomycin on the polymer was determined by ‘H NMR and UV-Vis spectroscopies to be 1 and 5, respectively [26]. The polymers \(\text{pA–V–F}\) and \(\text{pA–F}\) were soluble in water and phosphate-buffered saline (PBS).

2.3. Culturing of bacteria

Cultures of E. coli, S. epidermidis, and S. aureus were grown as liquid cultures in 5 mL of tryptone soy broth (TSB). Cultures of E. faecalis were grown in 5 mL of brain-heart infusion broth. Cultures of E. faecalis VanB were grown in 5 mL of brain-heart infusion broth supplemented with PBS, vancomycin (5 μM), or \(\text{pA–V–F}\) (10 μM vancomycin, 2 μM fluorescein). All cultures were grown overnight at 37°C with shaking at 200 rpm in Falcon 2059 tubes. All of the bacteria were harvested by centrifugation, and washed by resuspension in PBS and centrifugation. For flow cytometry experiments, S. pneumoniae (ATCC 6303) or S. aureus (ATCC 25923) were grown on tryptic soy agar plates supplemented with 5% sheep blood for 16 h at 37°C in an atmosphere containing 5% CO₂. Following the incubation, bacterial colonies were lifted from the plates and resuspended in PBS such that the optical densities (at 600 nm) for S. pneumoniae and S. aureus were 0.3 and 1.25, respectively. Actual numbers of viable bacteria were determined by plating known serial dilutions and determining the number of colony-forming units (CFU).
2.4. Fluorescence microscopy: binding of pA–V–F to bacterial surfaces

After washing, the bacteria were re-suspended in PBS containing Syto-63 and pA–V–F. The final OD_{600} was unity for the bacteria, and the final concentration of Syto-63 was 5 μM, and pA–V–F was 10 μM in vancomycin, 2 μM in fluorescein. The bacteria were incubated for 30 min with the dye and the polymer, and then 50 μL of the suspension was transferred to an ELF spin filter (0.2 μm pore size) and centrifuged to remove the free polymer and dye. The bacteria were washed by being re-suspended in 200 μL of PBS followed by removal of the solution by centrifugation in the spin filter. Samples were washed three times. After the final washing, the bacteria were resuspended in a 1:1 mixture of PBS and SlowFade antifade reagent or in PBS; no difference was found between the two preparations. Samples of S. aureus were prepared using this protocol except that only pA–V–F was included in the binding reaction; Syto-63 was not included. Control samples in which bacteria were incubated with Syto-63 and pA–F, or Syto-63 alone were also prepared and washed following the procedure given above. Samples were prepared for microscopy by placing 2–4 μL of bacterial sample on a glass slide and covering it with a no. 1 glass cover slip. The edges of the cover slip were sealed with clear nail polish.

Treatment with pA–V–F lead to no observed increase in the tendency of the bacteria to aggregate and no special treatment was required to disperse the cells before observation. To provide accurate images, the voltage of the PMT used for detection was adjusted to prevent any pixels from being saturated. Even in the presence of the anti-fade reagent, the sample (especially fluorescein) remained susceptible to photobleaching.

2.5. Fluorescence microscopy: opsonization by antibodies of pA–V–F-labeled bacteria

S. epidermidis was grown, harvested, and washed as previously described [26]. A portion of the bacteria was exposed to Syto-63 and pA–V–F and washed. This sample was then divided. One portion was exposed to 130 μM (20 μg mL^{-1}) R-phycocerythrin-conjugated IgG (IgG(anti-fluor)) for 2 h at room temperature in PBS SuperBlock buffer. A second portion of the sample was incubated for 2 h in the same buffer but without the antibody. Another portion of the washed S. epidermidis sample was exposed to Syto-63 (but not pA–V–F) for 30 min, washed three times using an ELF spin filter, then exposed to 130 μM IgG(anti-fluor) for 2 h in PBS SuperBlock buffer. All three samples were then washed by centrifugation in ELF spin filters followed by resuspension in PBS. Samples were washed five times, resuspended in a 1:1 mixture of PBS and SlowFade, and then analyzed by fluorescence microscopy.

2.6. Flow cytometry to quantitate opsonization of polymer-labeled bacteria

For opsonization experiments, 200 μL S. pneumoniae or S. aureus solutions (prepared as for studies in microscopy) were centrifuged (10 min; 16,000 x g) and re-suspended in PBS containing no polymer, pA–F (2 μM in fluorescein), or pA–V–F (10 μM in vancomycin, 2 μM in fluorescein). Following a 30 min incubation at room temperature in the dark, bacteria were washed three times with PBS followed by centrifugation. After the third wash, bacteria were re-suspended in PBS containing no antibody, 67 nm (10 μg mL^{-1}) isotype control antibody (IgG(control)), or 67 nm (10 μg mL^{-1}) anti-fluorescein antibody (IgG(anti-fluor)). Samples were then incubated for 30 min at room temperature, washed twice, and re-suspended in PBS containing no antibody (control/no polymer group only), or 67 nm (10 μg mL^{-1}) R-phycocerythrin-conjugated goat anti-mouse IgG (IgG(anti-mouse)) for 2 h. After a final 30 min incubation, bacteria were washed twice, fixed in PBS with 4% paraformaldehyde, re-suspended in 1 mL PBS, and examined by flow cytometry.

2.7. Preparation of S. aureus and J774A.1 cells (macrophages) for in vitro phagocytosis of opsonized S. aureus

S. aureus (286 μL; OD_{600} = 1.25) treated with pA–V–F/IgG(anti-fluor) or pA–V–F/ IgG(control) were re-suspended in 1.0 mL modified Hank’s Balanced Salt Solution (HBSS+) containing 1.0% Bovine Serum Albumin (BSA). J774A.1 cells were cultured at 37 °C (+5% CO2) in RPMI medium supplemented with streptomycin (50 μg mL^{-1}), penicillin (50 U mL^{-1}), l-glutamine (2 mM), and 10% fetal bovine serum. On the day of experiments, J774 cells were centrifuged (10 min; 300 x g), washed twice with PBS and re-suspended at a concentration of 2 x 10^6 cells mL^{-1} in HBSS+ containing 1% BSA. Bacteria (250 μL) and J774A.1 cells (250 μL) were combined resulting in a bacteria CFU:cell ratio of approximately 300:1, as verified by serial dilutions and plated colony counts. J774 cells were exposed to S. aureus for one or two hours in a 37 °C incubator containing 5% CO2, washed twice with ice cold HBSS+, fixed for 30 min in PBS with 4% paraformaldehyde, and re-suspended in 0.5 mL PBS. Fixed J774A.1 cells were then examined by flow cytometry.

To visualize phagocytosis via microscopy, unfixed J774 cell monolayers were prepared by cytocentrifugation, stained using the Diff-Quik® stain set, and visualized by optical microscopy. Fluorescence microscopy was performed on unstained cells.

3. Results and discussion

3.1. Synthesis and characterization of the bifunctional polymer (pA–V–F)

The bifunctional polymer, pA–V–F, consists of a polyacrylamide backbone with pendant vancomycin and fluorescein groups (Fig. 1(a)). Vancomycin is present on the polymer at side-chain % = 5 and fluorescein is present at side-chain % = 1 (as determined by 1H NMR and UV-Vis spectroscopies). In this context, we mean the "side-chain %" of a group to represent the percentage of side chains in the polymer containing that group. We also synthesized a control polymer, pA–F, consisting of a polyacrylamide backbone with only pendant fluorescein groups (Fig. 1(a)). We have described the synthesis and characterization of these polymers elsewhere [26].

In order to bind to the exterior surface of the microorganism, these polymers did not need to diffuse through the peptidoglycan layer (a three-dimensional structure with substantial porosity and an exclusion limit of ~100 kDa) [30] to the membrane surface (where the antibiotic activity of vancomycin is manifested); they could, therefore, be large. We confirmed that pA–V–F does not serve as an antibiotic at the concentrations used in our experiments through two control studies: (i) S. aureus treated with pA–V–F formed a number of colonies similar to that formed by bacteria that were not treated with polymer and (ii) E. faecalis grew normally in brain-heart infusion with (or without) pA–V–F. The average degree of polymerization for the polyacrylamide backbone was ~900; this value corresponds to an average of 45 vancomycin groups and nine fluorescein groups per molecule of pA–V–F, a molecular weight of 136 kDa after functionalization, and an extended, end-to-end length of the polymer of approximately 100 nm.
3.2. \( p\alpha-V-F \) bound specifically to the Gram-positive bacteria S. aureus, S. epidermidis, and E. faecalis

Previously, we demonstrated that \( p\alpha-V-F \) bound to the surface of a Gram-positive bacterium, E. faecalis, using fluorescence microscopy [26]. Here we examined the generality of \( p\alpha-V-F \) labeling using two additional Gram-positive bacteria: S. epidermidis (Fig. 2) and S. aureus (data not shown). We treated a suspension of S. epidermidis with a solution of \( p\alpha-V-F \) (10 \( \mu \)M in vancomycin, 2 \( \mu \)m in fluorescein) and the cell-permeable dye for nucleic acids Syto-63 (5 \( \mu \)M), in PBS. The bacteria were then washed with PBS and examined microscopically using an Ar–Kr laser for excitation of the fluorophores. The stained nucleic acid of the S. epidermidis was clearly visible when we selectively excited Syto-63 (Fig. 2). For a large fraction of bacteria, we observed fluorescence from fluorescein (on \( p\alpha-V-F \)) that co-localized with the fluorescence from Syto-63 (Fig. 2); this result suggests a high labeling efficiency of the bacteria by \( p\alpha-V-F \).

We also investigated binding of the polymer to S. aureus (data not shown). Suspensions of this bacterium were incubated in a solution of \( p\alpha-V-F \), washed, and examined using fluorescence microscopy. Syto-63 was omitted in order to eliminate any possibility of interference from strongly fluorescent complexes of Syto-63 and nucleic acid. Similar levels of fluorescence from fluorescein were observed from the \( p\alpha-V-F \)-labeled S. aureus, S. epidermidis, and E. faecalis. A control experiment with S. aureus and \( p\alpha-F \), a polymer that does not have vancomycin side chains, showed no detectable fluorescence from fluorescein on the bacteria (data not shown).

Gram-negative bacteria (e.g., E. coli) are not susceptible to the antibiotic activity of vancomycin because their cell wall is protected by an outer membrane that is impermeable to vancomycin [15]. A control experiment using E. coli incubated with Syto-63 and \( p\alpha-V-F \), revealed fluorescence from Syto-63 but not from fluorescein (Fig. 2). The results from these two control experiments support the inference that binding of \( p\alpha-V-F \) to the surface of bacteria requires vancomycin-mediated molecular recognition.

We examined the binding of \( p\alpha-V-F \) to vancomycin-resistant E. faecalis VanB to determine whether the polymer could be effective against these clinically important bacteria. While \( p\alpha-V-F \) bound to vehicle (PBS)-treated VanB, it did not bind as effectively to VanB that had been grown overnight in 5 \( \mu \)M vancomycin (at the concentration used to display fluorescein), which displays fluorescein (at the concentration used to visualize the bacteria) and \( p\alpha-V-F \) (which displays fluorescein groups), and then rinsed with PBS to remove any material not associated with a cell. The treated bacteria were examined by confocal fluorescence microscopy using an Ar–Kr laser to excite the fluorophores. The Gram-positive bacteria, S. epidermidis and E. faecalis (vancomycin-sensitive and vancomycin-resistant VanB that had been grown with PBS or with \( p\alpha-V-F \)) displayed fluorescence from fluorescein that coincided with fluorescence from Syto-63. E. coli, a Gram-negative bacterium, and E. faecalis VanB that had been grown with vancomycin only showed fluorescence from Syto-63 (i.e., fluorescein was not observed).

VanB in the presence of \( p\alpha-V-F \) (at the concentration used in binding experiments) in order to determine whether the polymer would induce active expression of the vancomycin-resistance genes. Fig. 2 shows that \( p\alpha-V-F \) bound to the surface of these treated bacteria (after rinsing with PBS and re-treating with \( p\alpha-V-F \)); this result suggests that \( p\alpha-V-F \) does not induce expression of these genes (i.e., the
bacteria were still displaying d-Ala–d-Ala on their surfaces). The observation that (monomeric) vancomycin triggers VanB to become resistant to binding both vancomycin and pA–V–F, while pA–V–F does not induce such a resistant phenotype, suggests that pA–V–F may have advantages over (monomeric) vancomycin for treating infections caused by VanB (and potentially, by other vancomycin-resistant bacteria) using the strategy that we describe below.

3.3. Opsonization of polymer-labeled bacteria by antibodies

After demonstrating binding of the bifunctional polymer, pA–V–F, to bacterial surfaces, we examined the ability of anti-fluorescein antibodies to bind to S. epidermidis, S. pneumoniae, and S. aureus that had been labeled with polymer. For the initial studies, we used fluorescence microscopy to examine qualitatively the binding of fluorescently labeled anti-fluorescein antibodies to the pA–V–F-labeled bacteria (Fig. 3). As a control experiment, we incubated S. epidermidis with Syto-63 and pA–V–F (as for polymer labeling studies). These bacteria displayed fluorescence from Syto-63 and fluorescein (on pA–V–F) but not from R-phycoerythrin (phyco) (Fig. 3); this result demonstrates that Syto-63 and fluorescein were not fluorescent at the wavelengths used to excite and detect phycoerythrin. S. epidermidis that were incubated with pA–V–F, Syto-63, and anti-fluorescein antibody labeled with R-phycoerythrin (IgG(anti-fluor)/IgG(anti-mouse)phyco) exhibited phycoerythrin (phyco) fluorescence (from the bound IgG(anti-fluor)/IgG(anti-mouse)phyco) that co-localized with fluorescence from fluorescein (and from Syto-63) (Fig. 3). Bacteria that were exposed to IgG(anti-fluor)/IgG(anti-mouse)phyco and Syto-63 but not pA–V–F did not display any phycoerythrin fluorescence (Fig. 3). These results demonstrate that pA–V–F must be bound to the bacteria to mediate binding by the antibody, and that every bacterium labeled with pA–V–F was recognized by at least one molecule of antibody.

Flow cytometry measured the opsonization of pA–V–F-labeled bacteria (S. aureus and S. pneumoniae) by anti-fluorescein antibodies. This technique quantitatively and rapidly determines the fluorescence intensity of some thousands of bacteria. In addition, flow cytometry experiments do not require any dye for nucleic acids, because bacteria are identified by their ability to scatter light. We incubated the bacteria with polymer (pA–V–F or pA–F) and a primary antibody in a manner similar to that used in studies using microscopy. For flow cytometry, however, we used an unlabeled monoclonal IgG2a antibody (mouse anti-fluorescein antibody (IgG(anti-fluor))) so that we could use the same primary antibody in studies of both opsonization and phagocytosis (see Section 3.4). For control experiments, we used a mouse isotype-control IgG2a antibody directed towards an unrelated hapten (IgG(control)) to probe for non-specific interactions of the antibody (e.g., the Fc region) with the polymer-labeled bacteria. Finally, incubation with an anti-mouse secondary antibody labeled with R-phycoerythrin (goat IgG (anti-mouse)phyco) facilitated detection of primary antibody bound to the bacterial surface. We examined four groups of bacteria: untreated bacteria (no polymer or antibody) (1), pA–F/IgG(anti-fluor)/IgG(anti-mouse)phyco (2b), pA–V–F/IgG(control)/IgG(anti-mouse)phyco (3b), and pA–V–F/IgG(anti-fluor)/IgG(anti-mouse)phyco (4b) (Fig. 4).

We quantitated the changes in phycoerythrin fluorescence of the different groups of bacteria by determining the mean fluorescence intensity and the percentage of bacteria in each group that were more fluorescent than a threshold. The threshold was set to the highest fluorescence intensity observed for untreated bacteria (Fig. 3). Treatment of bacteria with pA–F/IgG(anti-fluor)/IgG(anti-mouse)phyco (2b) controlled for any non-specific (i.e., independent of vancomycin) binding of the bifunctional polymer to the bacterial surface. This control group for both S. aureus and S. pneumoniae demonstrated a slight increase in phycoerythrin fluorescence intensity (mean fluorescence increased by ~2-fold, Table 1), and in the fraction of bacteria above the threshold (~16%, Table 1), relative to untreated bacteria (1). The possibility of non-specific binding of the antibodies to the bacterial surfaces was eliminated with a control experiment (see below). These increases are larger than the uncertainties in the measurements, and are
consistent with a low level of non-specific binding of the polymer to the surfaces of both bacteria by interactions that are independent of vancomycin.

Bacteria treated with pA–V–F/IgG(control)/IgG(anti-mouse) Phyco (3b) displayed the same mean fluorescence intensity as untreated bacteria (within error) for both S. aureus and S. pneumoniae. The percentage of bacteria in this control group (IgG(control)) with fluorescence greater than the threshold was negligible (within error) for both bacteria. The absence of phycoerythrin fluorescence (over background) in these control experiments eliminated the possibility of non-specific binding to the bacterial surfaces of the antibodies: the Fc region of the IgG2a primary antibodies (IgG(control) and IgG(anti-fluor)) and the secondary antibody (IgG(anti-mouse) Phyco).

Both S. aureus and S. pneumoniae treated with pA–V–F/IgG(anti-fluor)/IgG(anti-mouse) Phyco (4b) had mean phycoerythrin fluorescence intensities between 10- and 20-fold greater than control groups (2b and 3b) based on statistical analysis. The percentage of bacteria that above the fluorescence threshold was high in this group for both bacteria (S. aureus, 76 ± 9%; S. pneumoniae, 92 ± 6%). These values were much higher than those values observed in any of the control groups. For S. pneumoniae, the data from flow cytometry indicate that every bacterium is labeled by pA–V–F and IgG(anti-fluor) (the percentage of bacteria with fluorescence above the threshold was ~100%). For S. aureus, the data for the group incubated with pA–V–F/IgG(anti-fluor) show a high degree of labeling of bacteria by antibody (76% of bacteria have phyco fluorescence above threshold). We cannot, however, rule out the possibility that some fraction of the population of S. aureus did not bind to either IgG(fluor) or pA–V–F, given the overlap of the histograms for control and experimental groups (Fig. 5(a)). From both the flow cytometry and fluorescence microscopy results, we conclude that the polymer must have both vancomycin and fluorescein side-chains to opsonize S. aureus and S. pneumoniae: vancomycin, to adsorb the polymer to the bacterial surface via binding to D-Ala–D-Ala in the cell wall (cf. 2b and 4b, Fig. 5 and Table 1), and fluorescein, to localize anti-fluorescein antibodies to the bacterial surface (cf. 3b and 4b, Fig. 5 and Table 1) (Figs. 1(c) and (d)).

3.4. Association with macrophages and phagocytosis of opsonized S. aureus

Having demonstrated that antibodies recognize haptens (fluorescein) on the polymers bound to bacteria, we explored the ability of the bound IgG antibodies to interact with macrophages and to promote phagocytosis of the bacteria (Figs. 1(e) and (f)). Flow cytometry allowed us to follow the association of the fluorescent polymer-labeled bacteria with macrophages (Fig. 6). We incubated S. aureus that were pre-treated with pA–V–F and primary antibody (IgG(anti-fluor): 4a, or IgG(control): 3a) with cultured J774 cells (a mouse macrophage-like cell line, which we refer to below as macrophages). After incubation, we washed the macrophages and quantitatively examined macrophages associated with pA–V–F-labeled bacteria using flow cytometry (gating on the macrophages and detecting fluorescence from fluorescein) (Fig. 6). We also examined the macrophages qualitatively by optical microscopy (Fig. 7).

In the flow cytometry experiments, we could only observe bacteria that were labeled with pA–V–F because we detected fluorescence from fluorescein (on pA–V–F). We gated on the macrophages to exclude signals from free pA–V–F and pA–V–F-labeled bacteria that were not associated with the macrophages. The control experiments for phagocytosis, therefore, used fluorescent pA–V–F-labeled bacteria treated with IgG(control) (3a) to estimate the antibody-independent phagocytosis of S. aureus [32,33]. The control experiments for opsonization had
demonstrated that IgG(control) does not bind to pA–V–F-labeled bacteria (Fig. 5, group (3b)). In the control phagocytosis experiment with IgG(control), approximately one-half (51 ± 3%) of the macrophages had fluorescence from fluorescein above the threshold, which was set as the maximum fluorescence intensity observed for macrophages interacting with untreated S. aureus (i.e., background fluorescence). These results suggest a moderate level of association between pA–V–F-labeled S. aureus and macrophages independent of specific antibodies. These data are consistent with previous reports demonstrating that macrophages have antibody-independent mechanisms for phagocytosis of S. aureus [32,33].

There was a statistically significant increase in the percentage of macrophages above the fluorescence threshold (from 51 ± 3% to 84 ± 1%) when the pA–V–F labeled S. aureus were incubated with IgG(anti-fluor) (4a) as compared to being incubated with IgG(control) (3a) (Fig. 6(b)). This increased percentage of fluorescent macrophages represents an increase in the percentage of macrophages that are associated with pA–V–F-labeled bacteria in a specific antibody (IgG(anti-fluor))-dependent manner. The high background (antibody-independent) level of association between macrophages and pA–V–F-labeled S. aureus [32,33] (Fig. 6) limited the maximum effect that we could observe (theoretical maximum of 100% fluorescent macrophages). The results, nevertheless, demonstrate that pA–V–F is able to increase the percentage of macrophages that are associated with S. aureus in a manner that requires opsonization by anti-fluorescein antibodies.

The greater than two-fold increase in mean fluorescence for macrophages in the IgG(anti-fluor) group (4a) as compared to those in the IgG(control) group (3a) (Fig. 6(c)) indicates an increase in the average number of pA–V–F-labeled bacteria associated with each macrophage when there is IgG(anti-fluor) present on the surface of the bacteria. We believe that this increase in mean fluorescence underestimates the interaction between the pA–V–F-labeled bacteria and macrophages in the IgG(anti-fluor) group due to the quenching of fluorescence from fluorescein upon binding by IgG(anti-fluor). The addition of an excess of IgG(anti-fluor) to a solution of pA–V–F quenched the fluorescence from fluorescein by >95% (data not shown). We cannot comment on the quantitative amount that this quenching reduces the observed interaction between the pA–V–F-labeled bacteria and macrophages in the IgG(anti-fluor) group because we have no estimate of the number of free (not bound by antibody) fluorescein moieties per molecule of polymer associated with bacteria (see Refs. [17–20]). This quenching does not occur in the control samples incubated with IgG(control). The results with IgG(anti-fluor), even with the likely underestimation due to quenching of fluorescence by binding of antibody, demonstrate an increased association of pA–V–F-labeled bacteria with macrophages in a manner that is dependent upon an IgG with specificity towards the hapten (fluorescein) introduced by the polymer.

Macrophage studies included both one and two hour time points, in order to evaluate whether the enhanced phagocytosis of the IgG(anti-fluor) group relative to the IgG(control) group persisted. Similar to the earlier time point, IgG with specificity for the fluorescein hapten on pA–V–F (IgG(anti-fluor)) (4a) significantly increased
Table 1
Summary of flow cytometry data for opsonization of bacteria

<table>
<thead>
<tr>
<th></th>
<th>Mean fluorescence intensity&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Percentage of bacteria above threshold&lt;sup&gt;b,c&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>S. aureus</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacteria (untreated)</td>
<td>4.5 ± 0.3</td>
<td>17 ± 16</td>
</tr>
<tr>
<td>+ pA–F/IgG(anti-fluor)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (2b)</td>
<td>9 ± 4</td>
<td>17 ± 16</td>
</tr>
<tr>
<td>+ pA–V–F/IgG(control)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (3b)</td>
<td>6 ± 2</td>
<td>6 ± 5</td>
</tr>
<tr>
<td>+ pA–V–F/IgG(anti-fluor)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (4b)</td>
<td>90 ± 40&lt;sup&gt;f&lt;/sup&gt;</td>
<td>76 ± 9&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td><strong>S. pneumoniae</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacteria (untreated)</td>
<td>4.3 ± 0.2</td>
<td>15 ± 11</td>
</tr>
<tr>
<td>+ pA–F/IgG(anti-fluor)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (2b)</td>
<td>8 ± 3</td>
<td>15 ± 11</td>
</tr>
<tr>
<td>+ pA–V–F/IgG(control)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (3b)</td>
<td>4.5 ± 0.2</td>
<td>0 ± 0</td>
</tr>
<tr>
<td>+ pA–V–F/IgG(anti-fluor)/IgG(anti-mouse)&lt;sup&gt;Phyco&lt;/sup&gt; (4b)</td>
<td>80 ± 30&lt;sup&gt;f&lt;/sup&gt;</td>
<td>92 ± 6&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a</sup>Uncertainties represent standard errors of the mean from three independent experiments.

<sup>b</sup>Threshold was set on the maximum fluorescence observed for untreated bacteria (and is shown as dotted line in Fig. 5). Percentages above threshold in group 1 are zero by definition.

<sup>c</sup><sup>p</sup><sup><0.05</sup> vs. other polymer-treated groups. Comparisons were performed using a one-way analysis of variance (ANOVA) followed by a Duncan post-hoc test. When data did not pass Levine’s test for normality and equal variance, they were logarithmically transformed (base 10) to meet ANOVA requirements.

**Fig. 6.** Flow cytometry data demonstrating greater phagocytosis of pA–V–F-labeled *S. aureus* when treated with anti-fluorescein IgG (IgG(anti-fluor)) than when treated with isotype-control IgG (IgG(control)). Bacteria were treated with pA–V–F and then with either IgG(anti-fluor) (4a) or IgG(control) (3a). Cultured J774 cells (macrophages) were incubated with these potentially opsonized bacteria for 1 or 2 h, washed, and analyzed by flow cytometry. Macrophages were identified and gated based on forward and 90° light scatter, and fluorescence intensity from fluorescein was measured for 5000 events of the macrophage population. (a) Representative histograms (with 1 h incubation of bacteria with macrophages) showing the fluorescence intensity of 5000 events of the macrophage population. The dotted line indicates the fluorescence threshold that was set on the basis of the maximum fluorescence intensity from untreated control bacteria. The fluorescein fluorescence intensity shifts to higher values for the macrophage population when incubated with IgG(anti-fluor) than when incubated with IgG(control). Summarized data from four independent experiments are expressed as (b) the percentage of macrophages that are fluorescent (based on the threshold set in (a)), and (c) mean fluorescence intensity of all events. The error bars in (b) and (c) represent standard errors of the mean from the independent measurements. The legend in (b) applies to both (b) and (c). Comparisons were made among groups within each time point and the differences between all three groups were found to be statistically significant (<sup>p</sup><sup><0.05</sup>) using one-way analysis of variance (ANOVA) followed by a Bonferroni post-hoc test.

the percentage of macrophages associated with bacteria (Fig. 6(b)) and the number of bacteria per macrophage (Fig. 6(c)) relative to non-specific IgG (IgG(control)) (3a) after 2 h of incubation. The percentage of macrophages above the fluorescence threshold and the fluorescence per macrophage changed little between one and two hours for the IgG(anti-fluor) group; this result suggests that peak or near peak bacterial uptake in the hapten-specific antibody group was achieved by 1 h. The association of pA–V–F-labeled bacteria with macrophages achieved after two hours of incubation with IgG(control) failed to reach the levels achieved after only one hour of incubation with IgG(anti-fluor). Taken together, these findings suggest that the bifunctional polymer strategy (mediated by hapten-specific IgG) increases the net amount and the rate of bacterial association with macrophages for a significant time period relative to innate phagocytosis (independent of specific IgG) by the macrophages.

Flow cytometry does not distinguish between bacteria that are actually internalized and those that are merely associated with the periphery of the cell membrane of the macrophages. To address this issue, we examined...
bacteria-associated macrophages (prepared as above) using optical microscopy (Fig. 7). Bacteria (diameter of 0.8 μm) are visible both inside of the macrophage in phagosomes (Fig. 7(a); thin black arrow) and associated with the external cell membrane of the macrophage (thick black arrow) and internalized in intracellular phagosomes of the macrophage (thin black arrow). The large, dark object in the center of the macrophage (white arrow) is its nucleus. (b) Unstained macrophages were also imaged using fluorescence microscopy. This image is a fluorescence micrograph (fluorescence from fluorescein) merged with a phase contrast micrograph of the same field. The fluorescent signals (green or white) arising from the pA–V–F-labeled S. aureus appear to originate from within the macrophage. This intensity must arise from a large cluster of cocci; single cocci are not resolved using this technique. The scale bar applies to both images.

Fig. 7. S. aureus bacteria labeled with pA–V–F and anti-fluorescein IgG (IgG(anti-fluor)) are ingested by J774 macrophages. Macrophages were treated with pA–V–F and IgG(anti-fluor) as in Fig. 6. (a) Unfixed macrophages were prepared for visualization by optical microscopy by centrifugation and by staining with the Diff-Quik® stain set (Dade Behring). In this image of a representative macrophage, the S. aureus appear as small (0.8 μm diameter), darkly stained spheres (referred to as cocci). Cocci are evident both associated with the external cell membrane of the macrophage (thick black arrow) and internalized in intracellular phagosomes of the macrophage (thin black arrow). The large, dark object in the center of the macrophage (white arrow) is its nucleus. (b) Unstained macrophages were also imaged using fluorescence microscopy. This image is a fluorescence micrograph (fluorescence from fluorescein) merged with a phase contrast micrograph of the same field. The fluorescent signals (green or white) arising from the pA–V–F-labeled S. aureus appear to originate from within the macrophage. This intensity must arise from a large cluster of cocci; single cocci are not resolved using this technique. The scale bar applies to both images.

In the specific system studied here, pA–V–F cannot target Gram-negative bacteria, viruses, or other cell types and is thus not general for all pathogens. The use of fluorescein as our functional element (i.e., hapten) required using anti-fluorescein, an antibody directed towards an unnatural hapten, in the opsonization step. These specific limitations could, theoretically, be overcome by varying the recognition and functional elements of the polymer, as discussed above. Bifunctional polymers (like all polymers) are subject to certain limitations in therapeutic applications: they are not orally bioavailable and their innate polydispersity has hampered their FDA approval [36]. We believe, however, that polymers will be particularly effective in places where their large size (and thus, low oral bioavailability) is an advantage rather than a disadvantage. Examples could include administration to appropriate compartments, such as the digestive tract, respiratory system, eye, superficial soft tissue infections, and vagina, where retaining the polyvalent ligand in that organ or structure is useful, and where release into the systemic circulation may be undesirable.

As a general approach, bifunctional polymers might be useful therapeutically to target antibodies to pathogens or cancer cells to accelerate their destruction. An appropriate recognition element for the target of interest can be incorporated into the polymer and, given the polyvalent nature of the interaction, weak monovalent interactions would be acceptable in this role. The functional element on the polymer can be either a synthetic (as demonstrated here) or a natural hapten (e.g., part of a vaccine). A synthetic hapten would require the subsequent administration of antibodies directed towards that hapten, while a natural hapten would allow antibodies of the host to target the polymer-labeled target. Bifunctional polymers may also be useful in analytical applications. Binding to cells or viruses by a molecule that has both a tunable recognition component and a tunable labeling component could be useful in approaches designed to quantitate specific populations of cells [37] (e.g., by flow-cytometry).

4. Conclusions

We used a bifunctional polyacrylamide, pA–V–F (Fig. 1(a)), to form complexes with the surfaces of several representative Gram-positive bacteria (S. aureus,
S. epidermidis, S. pneumoniae, and E. faecalis) and to “decorate” these bacteria with a synthetic molecule (fluorescein); this molecule served as a hapten that was recognized by antibodies (IgG(anti-fluor)) in a second step. These antibodies, which were bound to the bacterial surfaces, interacted with macrophages (presumably via interactions with the Fc region of the antibody) and promoted phagocytosis of the opsonized, polymer-labeled bacteria (Fig. 1).

This polyacrylamide might be useful therapeutically in the clearance from appropriate biological compartments of bacterial infections caused by Gram-positive bacteria. We are currently investigating these applications using in vivo models of bacterial infections in the respiratory system. Further, the binding of pA–V–F to E. faecalis VanB after overnight incubation with pA–V–F (but not after incubation with monomeric vancomycin itself), and preliminary results that demonstrated that pA–V–F was able to mediate opsonization of some fraction of induced VanB by anti-fluorescein antibodies, suggest that the bifunctional polymer approach using vancomycin as the recognition moiety could be successful in targeting even these vancomycin-resistant bacteria, which pose a serious public health problem, for phagocytosis in vitro and in vivo.

Most importantly, the studies reported here offer a proof-of-principle demonstration that bifunctional polymers can target microbes for antibody-mediated immunity. By appropriate selection of the recognition moiety, this polyvalent approach should allow the selective destruction of pathogens (or other cells) of interest by coating their surfaces with hapten, to which antibody-mediated host defenses can be targeted.

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References


