Antibody-recruiting protein-catalyzed capture agents to combat antibiotic-resistant bacteria†

Matthew N. Idso, †a Ajay Suresh Akhade, a Mario L. Arrieta-Ortiz, a Bert T. Lai, b Vivek Srinivas, a James P. Hopkins Jr., a Ana Oliveira Gomes, a Naeha Subramanian, a Nitin Baliga a and James R. Heath† a

Antibiotic-resistant infections are projected to cause over 10 million deaths by 2050, yet the development of new antibiotics has slowed. This points to an urgent need for methodologies for the rapid development of antibiotics against emerging drug resistant pathogens. We report on a generalizable combined computational and synthetic approach, called antibody-recruiting protein-catalyzed capture agents (AR-PCCs), to address this challenge. We applied the combinatorial protein catalyzed capture agent (PCC) technology to identify macrocyclic peptide ligands against highly conserved surface protein epitopes of carbapenem-resistant Klebsiella pneumoniae, an opportunistic Gram-negative pathogen with drug resistant strains. Multi-omic data combined with bioinformatic analyses identified epitopes of the highly expressed MrkA surface protein of K. pneumoniae for targeting in PCC screens. The top-performing ligand exhibited high-affinity (EC50 ~50 nM) to full-length MrkA, and selectively bound to MrkA-expressing K. pneumoniae, but not to other pathogenic bacterial species. AR-PCCs that bear a hapten moiety promoted antibody recruitment to K. pneumoniae, leading to enhanced phagocytosis and phagocytic killing by macrophages. The rapid development of this highly targeted antibiotic implies that the integrated computational and synthetic toolkit described here can be used for the accelerated production of antibiotics against drug resistant bacteria.

Introduction

The emergence of antibiotic-resistant bacteria represents a major threat to human health,1−4 with both increasing mortality rates and costs of care.5−8 One example is carbapenem-resistant Klebsiella pneumoniae (K. pneumoniae) strains that harbor resistance against many or all antibiotics1,6,9,10 and frequently cause hospital-acquired infections11 with high mortality rates.12 Compounding the general problem of antibiotic resistance are challenges in developing and approving new antibiotics.13 This has spurred research into understanding resistance mechanisms,14,15 host defenses,16−18 diagnostics19 and antibiotic generation.20,21 Ultimately, without a rapid and perhaps general method to develop new targeted antibiotics, therapies might relapse towards those of the pre-antibiotic era.4,13

Immunomodulation by synthetic molecules is an emerging therapeutic strategy to combat bacterial and fungal infections,22−25 as well as other human diseases including cancer.26,27 The guiding principle is to employ an immunogenic agent to elicit a targeted immune response against a particular pathogen. The archetypical immune agent is an antibody and, while many therapeutic antibodies neutralize pathogens by direct interaction, a few operate by enhancing immune responses against the antibody target.22,28 Examples include antibodies against the Ebola virus,29 Staphylococcus aureus,30 Pseudomonas aeruginosa,31,32 and K. pneumoniae.33,34 While effective, antibodies can be challenging to produce and globally distribute at scale.35−37 Alternative compelling strategies include employing synthetic molecules that bind bacterial surface proteins36,37 or peptidoglycans,36,38 and present haptens so as to recruit the native immune system to promote bacterial clearance. Other synthetic approaches include metabolically incorporating non-native haptens into bacterial surface components.39,40 While inexpensive and scalable, these technologies can be challenging to adapt to different pathogens, or they can be non-selective, raising concerns about deleterious off-target effects.

Guided by these approaches, we envisioned a new class of highly targeted antibiotics called antibody-recruiting protein-catalyzed capture agents (AR-PCCs) that could be rapidly developed against a specified drug-resistant bacterium (Fig. 1). AR-PCCs consist of two molecular motifs. The first is a macrocyclic polypeptide ligand (the protein-catalyzed capture agent, or PCC) developed against a designated epitope of a specific...
bacterial surface protein. The second is an antibody-recruiting (AR) label on the PCC that promotes phagocytosis of the pathogen by innate immune cells (Fig. 1). To generate the AR-PCC ligand, we employed the recently reviewed, all synthetic epitope-targeted PCC method coupled with a bioinformatics approach to identify epitopes for targeting. By targeting highly exposed epitopes of the type 3 Fimbrial Shaft (MrkA) surface protein of K. pneumoniae, we developed a small macrocyclic peptide binder in a single generation screen. That binder exhibits high affinity for the MrkA protein, high selectivity for carbapenem-resistant K. pneumoniae, and, when labeled with the AR tag, promotes macrophage-mediated phagocytosis of the pathogen. This work demonstrates that AR-PCCs can be used to target multi-drug resistant K. pneumoniae, and that the basic technology might provide a route towards drugging “undruggable” pathogenic bacteria.

Results and discussion
Multi-omic analyses to select a target protein on K. pneumoniae
Here we describe the algorithm used to identify protein targets, and epitopes on those targets, for drugging K. pneumoniae using AR-PCCs. Traditional drugging strategies tend to rely on disrupting the function of, for example, an enzyme by competing for occupancy within a strategic hydrophobic binding pocket. Our requirements are very different. Instead, favorable aspects of target proteins are high expression levels on only the pathogen of interest, plus localization of that protein to the outer membrane or extracellular space of the pathogen. Further, once such a target protein is identified, there are additional considerations regarding which epitopes of that protein present the greatest opportunities for exploiting AR-PCCs.

The flow diagram in Fig. 2A delineates the strategy for identifying the major fimbrial subunit, MrkA, as an ideal target protein for K. pneumoniae. Protein expression levels can vary across environments and growth phases, so we analyzed the transcriptional data reported by Guilhen et al. to identify proteins with consistently high transcript levels across three major life phases of K. pneumoniae: exponential phase, stationary phase, and biofilms (including detached cells) (see Materials and Methods section). Briefly, we focused on the top 10% of highly-expressed genes across these life phases (515 genes out of 5146 genes in the transcriptomics dataset). Subsequent cross-referencing of these 515 genes with proteomics-derived information about the localization of K. pneumoniae proteins elucidated 13 highly expressed genes that encode proteins localized to the outer membrane or extracellular space. A literature search (see Materials and Methods section for specific references) then narrowed the selection by prioritizing essential virulence- and pathogenicity-related genes as well as protein orientation in the outer membrane, to five proteins: (i) FhuA, a siderophore, (ii) Lpp, a lipoprotein, (iii) Pal, a peptidoglycan-associated lipoprotein, (iv) NlpD, a lipoprotein, (v) MrkA, a subunit of the type 3 fimbriae. Ultimately, MrkA was chosen as a target due to its key roles in infection and persistence, its presence in the majority of sequenced K. pneumoniae strains, and, critically, its location in fimbrial rods. These rods are large extracellular structures (0.5–2 μm long, 4-to-5 nm in diameter) that are each comprised of up to 1000s of MrkA copies.

The selection of target epitopes on MrkA is illustrated in Fig. 2B. Selection of surface exposed epitopes would be aided by protein structure, but no published structures exist for MrkA. As a surrogate, we employed bioinformatics tools to survey MrkA for epitopes with high surface exposure and low homology with the human proteome. B-cell antigenicity was also mapped, but wasn’t a critical selection consideration.
The surface exposure, homology, and antigenicity of all 14-residue epitopes on the MrkA sequence were predicted and superimposed in Fig. 2B. Surface-exposure and antigenicity were calculated by averaging the predicted values assigned to each residue in the full-length MrkA sequence (by NetSurfP-2.0 or BepiPred-2.0) over the entire 14-residue epitope, while homology was predicted by comparing each 14-residue epitope with the human proteome using BlastP 2.0 with parameters defined in the heuristic string method by Berglund et al. This homology search yielded many partial matches per MrkA epitope, and the “Maximum homology” value plotted in Fig. 2B represents the percentage overlap with the best match. Predicted and averaged values from these analyses are tabulated in Table S1.

The plot in Fig. 2B reveals several regions containing epitopes of high (>60%) predicted surface exposure (Fig. 2B, green line) and relatively low (~50–55%) maximum homology to the human proteome (Fig. 2B, blue bars). Note that epitopes in surface-exposed regions range in predicted antigenicity from 0 to 100%.

Four epitopes, indicated by arrows in Fig. 2B, were selected based on their predicted high surface exposures (>60%), limited maximum homology (57–71%), and broad range of antigenicity values. The sequences of these candidate epitopes are shown in Fig. 2B. Importantly, the MrkA protein and candidate epitopes are highly conserved among K. pneumoniae strains (Fig. 2C). In fact, the MrkA sequence (with 202 amino acids) is 95% conserved across 380 analyzed K. pneumoniae strains.

**Epitope-targeted PCC ligands against the MrkA protein**

PCC ligands against the four selected epitopes were identified from a combinatorial library of macrocyclic peptides by using the epitope-targeted PCC method. This method exploits non-catalyzed click chemistry via an in situ click screen. For the screen, an alkyne-presenting, one-bead-one-compound (OBOC) combinatorial library of approximately 1 M peptide macrocycles with a 5-residue variable region is screened against synthetic variants of the epitopes (SynEps). Each SynEp is designed with a biotin assay handle and a strategically incorporated azide click handle (ESI Fig. S1 and S2† contain the structures and characterization data for all MrkA SynEps). The concept behind the screen is that select OBOC library elements will bind to a SynEp in just the right orientation so as to promote the azide-acetylene click reaction, thus covalently linking the SynEp to the bead. This product can be detected using the biotin assay handle on the SynEp, coupled with enzymatic amplification, to add color to the hit bead. Prior to screening SynEps, the library is cleared of beads that bound the detection antibody, streptavidin-alkaline phosphatase (SAv-AP). Hit beads are...
cy(LLFFF), conjugated to a 2,4-dinitrophenyl (DNP) moiety, was biotinylated. SynEps were immobilized. The lead ligand cy(LLFFF) binds. We performed a sandwich ELISA in which the second ligand binds the same epitope as cy(LLFFF), but cy(YRHLG) and cy(TTFFF) were also discovered, with sequences cy(TTFFF), cy(YRHLG) and cy(GVHRl). Based upon chemical homology, cy(TTFFF) likely binds the same epitope as cy(LLFFF), but cy(YRHLG) and cy(GVHRl) presumably bind a different one.

We next identified the particular epitope target to which cy(LLFFF) binds. We performed a sandwich ELISA in which the biotinylated SynEps were immobilized. The lead ligand cy(LLFFF), conjugated to a 2,4-dinitrophenyl (DNP) moiety, was titrated at a 500 nM concentration, and anti-DNP was used as a detection antibody. The results in Fig. 3B show the highest signals from SynEp 2 (TEVKAAADTYLKP). A background level for this assay was established using a biotinylated 6-mer polyethylene glycol (biotin-PEG6). Negative signals observed for the case of SynEp 1 (FFGKVTDSCTVSV) indicate that this SynEp blocks non-specific binding of cy(LLFFF) or anti-DNP more than the biotin-peg6 control. As the native epitope for SynEp1 (FFGKVTDSCTVSV) has a GRAVY score of 1.01, while the native versions of the other SynEps range between –0.9 and –0.28, the negative ELISA signals may arise from the relatively high hydrophobicity of SynEp 1. The blocking may also result from the reactivity of the cysteine residue in SynEp 1. Overall, these findings suggest that cy(LLFFF) binds the epitope TEVKAAADTYLKP in native MrkA. This epitope is 100% conserved across the 380 K. pneumoniae isolates analyzed (Fig. 2C), suggesting that cy(LLFFF) would bind the entire cohort of these K. pneumoniae strains.

An alanine scan was performed to establish which residues of cy(LLFFF) contribute most to MrkA binding. For this experiment, a sandwich ELISA is used to quantify the affinity of several cy(LLFFF) analogues, in which one residue is substituted with an alanine, towards full-length MrkA protein. The ELISA results in Fig. 4 show lower signal for every alanine-substituted cy(LLFFF) analogue compared to unmodified cy(LLFFF), establishing that the native compound has the highest affinity. Modestly lower binding is observed for cy(ALFFF), cy(LAFFF), and cy(LLAFF), but binding is substantially reduced for analogues with alanine substitutions at the C-terminal diphenylalanine motif. Thus, these two C-terminal phenylalanine residues (Fig. 3A, red arrows in inset) play critical roles in MrkA binding, and are targets for modifications to improve affinity. Another important detail of these alanine scan results is a strong dependence of MrkA binding on the residue position, independent of residue type. For example, cy(LLAFF) and cy(LLFAA) have very different binding affinities, despite having the same phenylalanine-to-alanine substitution. This indicates that MrkA-binding is sequence dependent, suggesting selectivity of cy(LLFFF) for MrkA at the TEVKAAADTYLKP epitope.

**Synthetic modifications to optimize AR-PCC pharmacokinetics and avidity**

The efficacy of AR-PCCs *in vivo* will not only depend on target binding, but also pharmacokinetic (PK) properties, of which clearance pathway is a critical parameter. Separate mouse studies in our labs indicate that PCCs with isocratic points (IPs) above 35% predominantly clear via the liver (*i.e.*, hepatic clearance), while more hydrophilic compounds with lower isocratic points clear by the kidneys (*i.e.*, renal clearance). Thus, the highly hydrophobic cy(LLFFF) ligand would likely exhibit hepatic clearance. To afford greater control over PK properties, we explored synthetic modifications to improve the hydrophilicity of cy(LLFFF), to favor renal clearance, while retaining the desired avidity characteristics. Modifications include single- and double-residue substitutions, residue removal, and the addition of non-ionic PEG and charged poly-arginine tags (R$i$, where $i$ represents the number of arginine). Results of this
optimization are shown in Fig. 4B, which plots IP versus the affinity of the compound to MrkA, relative to cy(LLFFF). The same results are tabulated in Table S2.†

Residue substitutions and removals categorically reduced affinity to MrkA, indicating that the structure of cy(LLFFF) compound is somewhat optimized to bind the TEVKAADTYLKP epitope. These residue substitutions and removals provided up to 30% reductions in IP versus cy(LLFFF). By comparison, the addition of polyarginine tags reduced the IP to 40% without a substantial loss in MrkA avidity (three leftmost data points in Fig. 4B). In fact, cy(LLFFF) with a hexa-arginine tag, "cy(LLFFF)+R6", shows a 22% stronger affinity to MrkA over cy(LLFFF) and had an isocratic point near 40%, as indicated by the red arrow in Fig. 4B. This represents an improvement of ~35% versus the IP of unmodified cy(LLFFF). An additional sandwich ELISA conducted with MrkA protein at 250 nM in solutions of 10 and 100% mouse sera demonstrate that cy(LLFFF)+R6 binds MrkA protein in the context of sera (Fig. 4B inset), making this molecule promising for in vivo translation.

AR-PCC binding to multidrug resistant *Klebsiella pneumoniae* surfaces

While cy(LLFFF) exhibits high affinity towards recombinant full-length MrkA protein in solution, the structure of MrkA is likely much different in native fimbriae, in which MrkA proteins oligomerize into a helix-like assembly.25,26 The affinity of cy(LLFFF) towards MrkA in the fimbriae of *K. pneumoniae* was interrogated using whole-cell ELISAs (Fig. 5A). In this assay, bacteria are exposed to biotinylated PCC ligands, then to SAv-horseradish peroxidase (SAv-HRP), and then developed to produce a colorimetric signal that correlates with SAv-HRP binding.

Whole-cell ELISA assays were conducted on the wild type version of the clinically-relevant strain of *K. pneumoniae*, KPPR1 (ATCC 43816). Wild type KPPR1 cells were cultured in G-CAA media to promote fimbrial expression,24 and MrkA expression in KPPR1 was confirmed by Western blot (Fig. 5B inset, full blot in Fig. S5†). The ELISA results in Fig. 5B (solid orange bars) show strong signals from wild type KPPR1 cells exposed to cy(LLFFF) at concentrations ranging from 5 μM down to 5 nM. By comparison, KPPR1 cells treated with a control compound cy(LLFFA), which differs from cy(LLFFF) by a single phenyl moiety, produce much lower intensities (Fig. 5B, solid blue bars) that are comparable to those of untreated KPPR1 cells (*"No PCC", Fig. 5B). These data not only demonstrate that cy(LLFFF) binds MrkA-expressing cells, but also reveal a high sensitivity of cy(LLFFF) structure on cell binding, establishing a strong structure–activity relationship for this PCC.

The specificity of cy(LLFFF) towards MrkA in particular was further interrogated by affinity tests to KPPR1 cells in which the gene encoding MrkA was knocked out (along with *mrkB* and *mrkc*).24 Western blot analysis confirmed that this modified KPPR1 strain does not produce MrkA (Fig. 5B inset, full blot in Fig. S5†). Knockout KPPR1 cells treated with cy(LLFFF) and control cy(LLFFA) yielded ELISA intensities comparable to baseline (Fig. 5B, dashed bars). This clearly establishes that cy(LLFFF) binds MrkA on *K. pneumoniae* cells. The specificity of cy(LLFFF) towards MrkA is further borne out by separate whole-cell ELISAs performed with *K. pneumoniae* cells (strain BAA 1705) cultured in minimal media containing 1% glycerol and 0.3% casamino acids (G-CAA media, which elicits high MrkA expression, or Lysogeny Broth (LB), which yields almost undetectable quantities of MrkA (Fig. S6 and S7†). ELISA signals were only observed for *K. pneumoniae* cells (strain BAA 1705) cultured in G-CAA media (Fig. S7†), when treated with cy(LLFFF) at concentrations of μM or lower. However, when higher 50 μM PCC concentrations were used, both cy(LLFFF) and control cy(LLFFA) bound *K. pneumoniae* cells regardless of MrkA-expression level (Fig. S7B†), suggesting non-selective hydrophobic binding at this high concentration.

The MrkA-specific binding of cy(LLFFF) anticipates selectivity towards MrkA-expressing *K. pneumoniae* versus bacteria devoid of MrkA. To test this, whole-cell ELISAs were performed on MrkA-producing multidrug-resistant *K. pneumoniae* (strains BAA 1705 & 2146) and the bacteria *Escherichia coli* (*E. coli*) and *Salmonella enterica* serovar Typhimurium (*S. Typhimurium*) that do not express MrkA, as confirmed by Western blot (Fig. S6†). Whole-cell

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**Fig. 4** Understanding and optimizing the cy(LLFFF) ligand through strategic chemical modifications. (A) Sandwich ELISA results of an alanine scan of cy(LLFFF), which evaluates the affinity of alanine-substituted cy(LLFFF) analogues to full-length MrkA protein. (B) Plot of the isocratic point versus affinity to MrkA protein, relative to cy(LLFFF), for various cy(LLFFF) analogues in which residues were either substituted or removed, or an N-terminal peptide tag was appended. Modifications generally reduced isocratic point but also reduced target binding, with the exception of an N-terminal hexa-arginine tag ("cy(LLFFF)+R6"). This compound also bound MrkA protein in solutions of 10 and 100% mouse sera, albeit at lower levels versus in 0% serum (inset). The relative affinities plotted in (B) are the ratio of ELISA signals yielded by the analogue and those produced by a cy(LLFFF) reference on the same ELISA plate. All ELISA data are background corrected to signal from an otherwise identical measurement except with biotin-peg6 conjugated instead of the PCC. Error bars are standard deviations. Each PCC used here had an N-terminal Biotin-peg group.
ELISA data in Fig. 5C show strong signals from the two strains of *K. pneumoniae* (BAA 1705 and BAA 2146) but only weak signals from *E. coli* or *S. Typhimurium* cells. The same bacterial strains treated instead with the control compound, biotinylated cy(LLFFA), produced low signals that were comparable to secondary antibody controls ("No PCC", Fig. 5C), establishing low binding. Strong binding of cy(LLFFF) towards *K. pneumoniae* cells demonstrate excellent binding selectivity of cy(LLFFF) towards MrkA-expressing *K. pneumoniae* cells versus non-target bacteria. The compound cy(LLFFF)+R6, which was chemically optimized for increased hydrophilicity versus cy(LLFFF), exhibited similarly high selectivity towards MrkA-expressing cells (Fig. 5C). Importantly, though not explicitly tested here, *E. coli* and *S. Typhimurium* are known to express fimbrae under growth conditions similar to those used here, suggesting selectivity of cy(LLFFF) to type 3 fimbiae of *K. pneumoniae*. Moreover, salt aggregation tests reveal that the cell surfaces of the *K. pneumoniae* strains used here have similar or lower hydrophobicities than *E. coli* or *S. Typhimurium* (Fig. S8 and S9†), indicating that binding selectivity cannot be explained simply by differences in surface hydrophobicity across these bacteria.

**AR-PCC-driven opsonization of *K. pneumoniae***

We next sought to use cy(LLFFF) to promote opsonization by appending an antibody-recruiting (AR) handle, the hapten 2,4-dinitrophenyl (DNP), to form an AR-PCC. DNP is a hapten that is bound by 1% of endogenous human antibodies, and has been employed in several immune-recruiting therapeutics to recruit antibodies to various pathogens and cancer cells. We hypothesized that, once the AR-PCC binds to the *K. pneumoniae* surface, the AR handle should recruit antibodies to the pathogen. PCCs were tagged with a DNP group via conjugation of a DNP-modified lysine residue, in which a DNP moiety is covalently attached at the terminal sidechain amine. Both DNP-conjugated cy(LLFFF) and cy(HNGPT) PCCs, referred to respectively as cy(LLFFF)-DNP and cy(HNGPT)-DNP, showed absorbances at 360 nm and 420 nm (Fig. S10†) that are characteristic of DNP-modified lysine and indicate successful labeling of the PCC.

Flow cytometry was used to determine whether the cy(LLFFF)-DNP AR-PCC promoted opsonization of resistant *K. pneumoniae* by anti-DNP antibodies. *K. pneumoniae* cells (strain BAA 2146) were first treated with cy(LLFFF)-DNP and then Alexafluor 488 fluorophore-labeled anti-DNP antibodies (Fig. 6A), so that fluorescence served as a proxy for opsonization. The cytometry data in Fig. 6B show weak fluorescence (predominantly <0.2 × 10^4 intensity) from *K. pneumoniae* cells that were either unstained or incubated with fluorescent anti-DNP only, and slightly higher fluorescence (predominantly <1.0 × 10^4 intensity) was observed for cells exposed to control AR-PCC cy(HNGPT)-DNP at 50 μM plus anti-DNP, indicating slight cross-reactivity. Fluorescence counts for all these samples, however, are substantially less than for cells treated with much lower concentrations (1 or 5 μM) of cy(LLFFF)-DNP plus fluorescent anti-DNP (Fig. 6B, light and dark green, respectively). This demonstrates that the cy(LLFFF)-DNP AR-PCC promotes the opsonization of *K. pneumoniae* cells with anti-DNP. The same behaviors were observed on a separate resistant *K. pneumoniae* strain, BAA 1705, (Fig. S11†) and by using whole-cell ELISA methods that were sensitive to anti-DNP binding (Fig. S12†). Paired with the recruitment of SAv-HRP to cells by biotin-cy(LLFFF) (Fig. 5B and C), these results lend to the versatility of AR-PCCs to recruit diverse and specified biomolecules to bacterial surfaces.

**AR-PCC-driven phagocytosis and opsonophagocytic killing (OPK) of *K. pneumoniae***

Given that the cy(LLFFF)-DNP AR-PCC recruits antibodies to *K. pneumoniae* surfaces, we posited that *K. pneumoniae* opsonized in this manner would be more susceptible to phagocytosis by macrophages. AR-PCC-driven phagocytosis and OPK were

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**Fig. 5** Binding specificity of cy(LLFFF) towards MrkA-expressing drug-resistant *K. pneumoniae* cells. (A) Schematic illustration of a whole-cell ELISA affinity test in which *K. pneumoniae* was incubated with PCCs conjugated with biotin, opsonized with streptavidin-HRP, and then detected colorimetrically. (B) Whole-cell ELISA results show that biotinylated cy(LLFFF) binds wild type MrkA-expressing KPPR1 cells (solid orange bars), but not MrkA-knockout KPPR1 (dashed orange bars) cells that do not produce MrkA. Much lower binding is observed from the highly homologous control compound cy(LLFFA) (solid and dashed blue bars). Western blots of wild type and MrkA-knockout KPPR1 cells are shown in the inset. (C) Cell-binding ELISA results show that biotinylated cy(LLFFF) at 5 μM binds two strains of MrkA-expressing *K. pneumoniae* cells with excellent selectivity versus *E. coli* and *S. Typhimurium*. By comparison a biotinylated control ligand cy(LLFFA) shows no or minimal binding above baseline. Asterisks (∗) indicate ELISA data for target *K. pneumoniae* strains, and error bars reflect standard deviations.
quantified by a gentamicin protection assay, as depicted in Fig. 7A. For this assay, *K. pneumoniae* cells (strain BAA 1705) were treated with cy(LLLFF)-DNP, opsonized by anti-DNP antibodies, and then incubated with murine bone-marrow-derived macrophages (BMDMs). BMDMs were subsequently exposed to a solution containing gentamicin, to which *K. pneumoniae* strain BAA 1705 is susceptible, to kill any extracellular bacteria. Phagocytosed bacteria remain viable inside BMDMs for at least 1 h, yet are rendered inviable after 24 h by OPK (Fig. S13†). Macrophages harvested at 1 h and 24 h postinfection were lysed and plated to generate bacterial colonies, which were enumerated to quantify phagocytosis and OPK.

Gentamicin protection assays were carried out using the cy(LLLFF)-DNP AR-PCC on a strain of *K. pneumoniae* (BAA 1705) that harbored high antimicrobial resistance, including all tested carbapenems, but is susceptible to gentamicin. As shown in Fig. 7B, BMDMs exposed to untreated *K. pneumoniae* cells for 1 h (*"K. pneumoniae* only", black bar) yielded counts of ~39 000 CFU per million cells, establishing a basal level of phagocytosis (dashed line in Fig. 7B). Similar signals were also observed in all control samples (Fig. 7B, leftmost six samples, black bars) harvested at the 1 h time point. The controls were untreated bacteria or bacteria treated either with anti-DNP antibody only, cy(LLLFF)-DNP only, cy(LLFFA)-DNP only, cy(LLFFF)+R6-DNP only or cy(LLFFA)-DNP plus anti-DNP. None of these treatments elicited phagocytosis above the basal level. Much greater counts of 89 000 CFU (indicated by the solid blue arrow, Fig. 7B) were observed from samples in which *K. pneumoniae* was treated with cy(LLLFF)-DNP plus anti-DNP, which compared well with the phagocytosis-promoting performances of the *K. pneumoniae* antiserum and the anti-MrkA antibody (Fig. 7B, rightmost two samples). When tagged with DNP, the optimized compound cy(LLLFF)+R6 also enhanced phagocytosis, as shown by the dashed blue arrow in Fig. 7B. In fact, the phagocytosis-promoting activity of DNP-tagged cy(LLLFF)-R6 was comparable to that of cy(LLLFF)-DNP. Statistical analyses (T-tests comparing phagocytosed CFUs after one hour of treatment) indicated that cy(LLLFF)-DNP plus anti-DNP promoted phagocytosis of *K. pneumoniae* above the basal level of the controls (*p*-value <0.01, Fig. S14, ESI†). Nonetheless, the level of phagocytosis induced by cy(LLLFF)-DNP plus anti-DNP was about 13% lower than that induced by *K. pneumoniae* antiserum and anti-MrkA treatment (positive controls: *p* <0.001). At 24 h of incubation, the cy(LLLFF)-DNP plus anti-DNP sample (and all other samples) showed little to no cell counts (Fig. 7B, red bars) indicating near complete OPK of phagocytosed *K. pneumoniae* bacteria. Thus, AR-PCCs demonstrably enhance the OPK of a highly resistant *K. pneumoniae* bacterium, presumably by engaging Fc-receptor-mediated phagocytosis pathways.

**Conclusions**

We demonstrated, *in vitro*, a new concept for targeted antibiotics called antibody-recruiting protein-catalyzed capture agents (AR-PCCs). AR-PCCs were designed to exhibit specific *in vitro* antimicrobial activities against highly-resistant *Klebsiella pneumoniae* bacteria. AR-PCC molecules comprise of two molecular motifs: a peptide ligand that binds a specific surface protein epitope on the pathogen, and a hapten that recruits antibodies. Combined multi-omic data and bioinformatic analyses provided an algorithm for selecting a highly abundant surface protein and epitopes on *K. pneumoniae* as targets for AR-PCCs. A single-generation protein-catalyzed capture agent combinatorial screen was then used to rapidly identify macrocyclic peptide ligands to the chosen epitope targets. The lead AR-PCC ligand, cy(LLLFF), exhibited strong binding to full-length MrkA (EC50 ~ 50 nM) and one of the highly conserved target epitopes, as well as high binding specificity towards MrkA-expressing *K. pneumoniae* versus other MrkA-deficient bacterial species. Further, the lead AR-PCC ligand conjugated with 2,4-dinitrophenyl (DNP) moieties recruited anti-DNP antibodies to the *K. pneumoniae* surface, which led to increased levels of phagocytosis by macrophages and ultimately greater opsonophagocytic killing. Chemical modifications to the cy(LLLFF) suggest that the macrocycle scaffold can be optimized for desired *in vivo* PK characteristics. Such mouse model experiments are currently underway. While *K. pneumoniae* served as the target in this study, the approaches used here should be adaptable to other antibiotic resistant extracellular pathogens. This versatility also makes feasible the development of cocktails of AR-PCCs that simultaneously target several conserved surface epitopes on a single pathogen, to facilitate complete clearance of bacterial populations that exhibit
heterogeneous surface protein expression. Overall, AR-PCCs are a promising all-synthetic molecular platform that can be rapidly designed, built and deployed against resistant microbes.

Materials and Methods

Gene expression analysis

Transcriptional profiles (normalized read counts) of *Klebsiella pneumoniae* strain CH1034 in stationary phase, exponential phase, and biofilms (7 h, 13 h and detached cells) generated by Guilhen and collaborators were downloaded from the GEO database (accession number: GSE71754). The downloaded dataset included 5146 genes and triplicates for each condition. Although additional transcriptomes are publicly available for *K. pneumoniae*, we restricted our analysis to a single dataset (with key stages of *K. pneumoniae* life cycle) due to: (i) the high number of *K. pneumoniae* strains studied by different research groups. This means available transcriptomic data involve multiple strains and have been collected on multiple platforms. (ii) The large size of *K. pneumoniae* pan-genome makes it possible that transcript levels of any given gene may be present in only some of the available transcriptional profiles. Inspired by the superior performance of rank-combined predictions that integrate multiple network inference methods/predictions (with respect to single ones), we ranked all genes in each replicate based on their normalized read counts. Then, we computed the average ranking position of each gene along the 15 replicates. This approach identified genes with consistently high transcript levels along the sampled conditions and, for downstream analyses, we focused on the set of genes that were in the top 10% of the ranking (515 genes).

Identification of highly expressed genes encoding outer membrane or extracellular proteins

We mined available literature to identify *K. pneumoniae* proteins that localize in the outer membrane or extracellular space. We found 54 proteins that localize to the regions of interest. Then, we evaluated the overlap between this set of proteins and the group of 515 highly expressed genes defined in the “Gene Expression Analysis” section. To compare the two sets, we first downloaded the genome annotation of *Klebsiella pneumoniae* strain CH1034 from the NCBI website in May 2018. We used the genome annotation to convert the locus names used as gene ID in the analyzed transcriptomics data to standard gene names (*e.g.* CH1034_10002 corresponds to *phnV*). Finally, we manually reviewed available information about genes present in both sets to select a target, prioritizing gene targets that yielded virulence-associated proteins that were either membrane-spanning or oriented on the extracellular side of the outer membrane.

MrkA sequence alignment

First we performed a Blastn (megablast) in the NCBI BLAST website using as query the nucleotide sequence of *mrkA* in the *K. pneumoniae* ATCC BAA-1705 strain. We restricted the search to the *K. pneumoniae* taxa and set the maximum number of allowed target sequences to 500. Other parameters were kept as default. We downloaded all hits (402). Then, we removed any hit from plasmid sequences or duplicated sequences. Nucleotide hit sequences were then translated using EMBOSS Translate. Finally, MrkA protein sequences were aligned using MAFFT and visualized in Jalview.

Bioinformatics analysis

Predictions for surface exposure and surface antigenicity were performed using NetsurfP 2.0 (ref. 57) and Bepipred2.0 (ref. 58) software (using default parameters), with the primary sequence of MrkA as an input, which assign either a value of 1 or 0 to each residue. The resulting prediction values were then averaged over 14-residue epitopes, converted to a percent value, and plotted. The uniqueness of each 14-residue epitope of MrkA among all epitopes in the human proteome was determined by performing a BlastP 2.0 search with the parameters defined in the heuristic string method by Berglund *et al.* Many matches
were obtained for each epitope, and the match with greatest homology with the MrkA epitope was extracted, the corresponding homology converted to a percent value, and plotted as “Maximum homology.” All results from these bioinformatic analyses are tabulated in Table S1.†

Peptide synthesis and purification

All peptides were synthesized by using standard Fmoc solid-phase peptide synthesis procedures, using either Rink Amide resin (Aapptech, RRZ005), Sieber Amide resin (Aapptech, RST001), or Tentagel S NH₂ resin (Rapp Polymere, S30902). N-methylpyrrolidone (NMP, Alfa Aesar, 43894) was used as a solvent for all synthesis procedures, except the coupling of biotin (AK Scientific, C820) which used a 50/50 mixture of dimethylsulfoxide (DMSO, Fisher, D128-4) and NMP for solubility. All standard Fmoc-protected amino acids were purchased from ChemPep. Fmoc deprotection was achieved with 20% piperidine (Alfa Aesar, A12442) in NMP, and coupling reactions employed O-(7-aza-benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexa-fluorophosphate (HATU, Chem-Impex Intl Inc., 12881) as an activator and N-disopropylethylamine (DIEA, Alfa Aesar, A11801) as the base. An Aapptech Titan 357 instrument was used to couple all Fmoc-protected standard amino acids and non-natural residues with click handles, i.e. propargylglycine (Pra, ChemPep, 180710) and azidolysine (Az4, ChemPep, 101227). Fmoc- NH-peg5-CH₂CH₂OH (ChemPep, 280110), Fmoc-NH-peg10-CH₂CH₂OH (ChemPep, 280113), and Fmoc-Lys(DNP)-OH (ApexBio, A7926) were coupled overnight with molar excesses of 2, 2, and 6 for the Fmoc-protected compound, HATU, and DIEA, respectively (excesses are with respect to the reactive groups on bead surface). The Cu-catalyzed azide–alkyne click reaction was conducted by incubating beads overnight with shaking in a solution of 1.5× molar excess of Cu(i) (Millipore Sigma, 818311) and 5× molar excess of l-ascorbic acid (Sigma, A0278) in 20% piperidine in NMP (excesses are with respect to the reactive groups on resin surface). The resin was then washed 5 × 1 minute with 5–8 mL of NMP, after which extra copper was removed by incubating beads with shaking for 5 minutes in a solution of 5 v/v% DIEA, 5 w/w% Sodium diethyldithiocarbamate trihydrate (Chem-Impex) in NMP. The resin is washed extensively with NMP (3× without shaking, 5 × 5 min or more with shaking) until the bead color turned white to light yellow and remained constant. After synthesis was complete, the resin was dried in dichloromethane (DCM, Acros Organics, 40692-0040) for at least 15 min on a vacuum manifold. Peptides were cleaved from dried resin by mixing with 10 mL solution of Trifluoroacetic acid (TFA, Alfa Aesar, L06374) : Triethylsilane (TES, Millipore Sigma, 230197) Millipore Water (H₂O) at volumetric ratios of 95 : 2.5 : 2.5, respectively, and vigorously stirring for 2 h. Cleavage under these acidic conditions also removed all acid-labile sidechain protecting groups. The resulting solution was added to 40 mL of diethyl ether (Acros Organics, 615080-0040) and stored overnight at −20 °C to precipitate the peptide product. The product was then pelleted by centrifugation, dried in air, and then resuspended in an aqueous solution of 30% acetonitrile (Fisher, A955-4) (aq.) with 0.1% of either TFA or formic acid (Fisher, A117-50) before purification by liquid-chromatography-mass spectrometry.

PCC compounds were purified on a Waters Autopurification system, which isolated compounds based on MS peaks corresponding to protonated [M + 1H]⁺ and [M + 2H]²⁺, and/or sodium adducts [M + Na]⁺ and [M + 2Na]²⁺. A representative HPLC chromatogram from the Waters Autopurification system with an analytical HPLC run of the corresponding product is shown in Fig. S15.† The isotopic points of PCCs were determined based on the elution of the compound from a C18 prep-scale column. Synthetic epitopes were either purified on the Waters Autopurification System, or by semi-preparative HPLC and then using matrix-assisted laser desorption/ionization mass spectroscopy to identify fractions with the desired m/z ratio. The resulting peptides were lyophilized, the yield determined by mass difference or UV-visible absorbance, and resuspended at a concentration of up to 10 nM peptide in DMSO. Peptides were stored at −80 °C before thawing for each use.

Standard Fmoc-protected synthesis and split-and-mix procedures were used to synthesize a one-bead one-compound library on Tentagel S NH₂ beads with the structure NH₂-Pra(80%)/Gly(20%)-X₇-X₆-X₅-X₄-X₃-Az4-M-Resin, where Xᵢ indicates one of 16 natural amino acids (excluding Methionine, Cysteine, Glutamine and Isoleucine). The coupling solution for the N-terminal residue included 80% Fmoc-propargylglycine-OH and 20% Fmoc-glycine-OH. The resulting library was clicked under copper-catalyzed conditions as described above to yield on each bead ~80% of the cyclic product cy(Pra-X₇-X₆-X₅-X₄-Az4) and 20% of the linear product Gly-X₇-X₆-X₅-X₄-Az4, which facilitates identification by tandem mass spectroscopy. A final Pra was coupled onto the library to enable PCC combinatorial screening. The library was then incubated with a solution containing volumetric ratios of 95 : 2.5 : 2.5 of TFA : H₂O : TES under vigorous stirring for 2 h to remove acid-labile sidechain protecting groups, and the washed 3 × 5 min in H₂O, NMP, methanol (Fisher, A454-1), and then DCM.

Combinatorial PCC screening

Combinatorial in situ click screens were performed as described previously.⁴⁴ Briefly, 500 mg (representing ~1.4 copies of approximately 1 000 000 different compounds) of a combinatorial one-bead one-compound libraries was incubated overnight with shaking in TBS buffer (25 mM Tris HCl, 150 mM NaCl, pH 7.6). A preclear to remove beads that bound SAv-AP was performed as follows. Unless otherwise noted, all steps were performed at room temperature and all incubation and washing steps were conducted with 4 mL of the stated solution with shaking. Beads were blocked overnight by incubation in blocking buffer (TBS buffer with 1% BSA and 0.05% Tween-20, pH 7.6), rinsed with blocking buffer 3 × 3 min, incubated for 1 h with 1 : 10 000 streptavidin-Alkaline phosphatase (SAV-AP) (Thermofisher Scientific, SA1008) in 5 mL of blocking buffer, and then washed with the following: 3 × 5 min in TBS, 3 × 5 min in 0.1 M glycine (pH 2.8), 3 × 5 min in TBS buffer, 3 × 5 min of alkaline phosphatase (AP) buffer (100 mM Tris–HCl, 150 mM NaCl, 1 mM MgCl₂, pH 9.0). The beads were then split and transferred into two separate Petri dishes by using AP buffer, such that there was a total of 16 mL of AP buffer per dish. Separately,
10 mL of BCIP/NBT development buffer was prepared from the Promega BCIP/NBT Color Development substrate kit (Promega, S3771) by adding 66 μL of the NBT solution to 10 mL of AP buffer, mixing by hand, then adding 33 μL of the BCIP solution followed by vortexing. Four mL of the BCIP/NBT development buffer was added to each plate and then each plate was gently swirled for 30 seconds to ensure the development buffer was well-mixed. The reaction was quenched after 25 minutes by adding 4 mL of 5.0 N HCl (aq.) to each plate and swirling to homogenize. The beads were transferred to a new SPPS tube by using Millipore water and then were washed copiously with Millipore water (10× without shaking, then 10 × 1 minute with shaking). The beads were resuspended in 0.05 N HCl (aq.), returned to a Petri dish, and the purple beads removed by using a 10 μL pipette.

After all of the purple beads were removed, the library was incubated overnight in NMP to decolorize the residual purple coloration on the remaining beads. The library was then washed 5 × 1 minute in Millipore water and incubated overnight. The in situ click screen was then performed with the MrkA SynEpss. The library was collected and rinsed in TBS buffer (3 × 5 min), incubated with a TBS buffer containing 20 μM of each purified MrkA SynEp for 6 h. The library was then subjected to the following treatments: washed with TBS buffer (10 × 1 min), incubated for 1 h in 7.5 M Guanidine-Hydrochloride (pH 2.0), washed with TBS (10 × 1 min), incubated for 2 h with blocking buffer, washed with blocking buffer (3 × 1 min), and incubated for 1 h with 1 × 10 000 SAv-AP in 5 mL of blocking buffer. The library was then washed as follows: 3 × 5 min in TBS, 3 × 5 min in 0.1 M glycine (pH 2.8), 3 × 5 min in TBS buffer, 3 × 5 min of AP buffer. The library was then split into two Petri dishes and developed as described above. The reaction was quenched by the addition of 4 mL of 5.0 N HCl (aq.), then the beads were transferred to a new SPPS tube, washed copiously with Millipore water (10× without shaking, then 10 × 1 minute with shaking), resuspended in 0.05 N HCl, and then returned to Petri dishes. Dark and medium-dark colored beads were picked as hits and collected into Corning Costar Spin-X centrifuge tube filters (Sigma-Aldrich, CLS8170). Hit beads were then rinsed 10 × 30 s (7000 RPM, top ETF centrifuge), decolorized by overnight incubation in NMP, rinsed in Millipore water (10 × 30 s), resuspended in TBS, and stored at 4 °C.

Individual beads were transferred to wells in a 96 well plate and subjected to cyanogen-bromide cleavage, prepared, and sequenced as described previously.44,45

In vitro ELISA assays
ELISA assays for MrkA-binding and epitope selectivity were performed on clear NeutrAvidin Coated High Capacity Plates (ThermoFisher, 15507). The buffer used for all washes and to dissolve all compounds was TBST + 0.1% BSA (TBST + 0.05% Tween 20 + 0.1% BSA, pH 7.3) and, unless otherwise stated, steps were conducted at room temperature. Each wash involved a brief 20 s agitation, and all conjugation, blocking, and incubation steps were performed with gentle agitation over the entire stated time period. The general procedure for an ELISA assay was: wash 3 × 200 μL per well, conjugate with 2 μM of biotinylated compound for 2 h at room temperature (100 μL per well), wash 3 × 200 μL per well, blocked overnight in 5 wt% milk at 4 °C, wash 3 × 200 μL per well with wash buffer, incubate with titrated compound (e.g., MrkA or PCC) for 1 h, 3 × 200 μL per well, incubate with primary antibody (either or) for 1 h (100 μL per well), wash 3 × 200 μL per well, incubate with 1 : 2000 anti-rabbit secondary antibody-horseradish peroxidase conjugate (Cell signaling Technologies, 7074S) for 1 h, 3 × 200 μL per well, develop with the Microwell Peroxidase Substrate System (2 C) (SeraCare, 5120-0047) using 100 μL per well for 1–40 minutes, and quench using 1 M H2SO4 (aq.) at 100 μL per well. For MrkA binding assays, biotinylated PCCs were conjugated to the well surface, recombinant MrkA with a N-terminal 6xHis-SUMO-tag (My Biosource, MBS1248970) was titrated at the desired concentration, and the primary antibody was His-tag antibody, pAb, Rabbit (Genscript, A00174) at a 1 : 5000 dilution. For epitope selectivity assays, biotinylated SynEpss were conjugated, DNPs-conjugated AR-PCCs were titrated at a desired concentration, and the primary antibody was anti-DNP antibody produced in Rabbit (Sigma-Aldrich, D9656) at a 1 : 8000 dilution. The PCCs and SynEpss used for plate-based ELISA assays had a peg5 linker between the N-terminus of the peptide and the tag, which was either biotin or DNP-modified lysine. For binding tests in the context of mouse sera, the “titrated compound” described in the steps above was dispersed in a solution containing the desired concentration of mouse serum.

Cell culture
Bacteria were grown overnight from glycerol stocks by inoculation into either minimal media containing 1% glycerol and 0.3% casamino acids (G-CAA) or Lysogeny Broth (LB) with shaking at 37 °C. G-CAA media was used to promote the expression of MrkA, while LB was used as a control medium for binding tests that involved K. pneumoniae that do not produce MrkA. An immortalised mouse bone marrow derived macrophage cell line (iBMDDM, a kind gift from Dr Eicke Latz) was used in this study. Cells were maintained in RPMI containing 10% serum at 37 °C in a humidified atmosphere with 5% CO2. K. pneumoniae strains BAA 1705 and BAA 2146 were obtained from ATCC, while wild type and MrkA/B/C knockout versions of K. pneumoniae ATCC 43816 were kindly provided by Prof. Matthew Wargo.

Salt aggregation tests
Bacteria were cultured overnight in G-CAA media, washed once in 0.02 M phosphate buffer (0.01 M Na2HPO4, 0.01 M NaH2PO4, pH 8), and then resuspended to an O.D.600 of 0.95. The bacterial solutions were arrayed onto a single glass slide in 10 μL spots, and then each spot was mixed with an equal volume of phosphate buffer containing (NH4)2SO4. The glass slide was gently agitated for 2 m. Images were recorded at 10 and 30 m following agitation, and image acquisition of all the spots took less than 1 m. Measurements were conducted on both live bacteria and heat-killed bacteria that were treated for 10 minutes at 90 °C.

Whole-cell ELISAs
Detection of the binding of biotinylated PCCs to bacterial surface was performed as follows. 108 bacteria from a culture grown...
overnight were used for each test. Bacteria were incubated with PBS containing 1% BSA (PBS-BSA) for 1 h at 37 °C, washed once with PBS, and then incubated with various concentrations of the biotinylated PCC in PBS-BSA for 1 h at 37 °C. After washing thrice with PBS to remove unbound PCCs, bacteria were incubated with Streptavidin-HRP (1 : 1000) for 1 h at 37 °C. Bacteria were washed thoroughly and the TMB reagent was added until visible coloration was observed. The reaction was quenched using 2 N H2SO4 and absorbance was measured at 450 nm. The biotinylated PCCs used for cell-based ELISA assays had a peg5-peg5 linker between the N-terminus of the PCC and the biotin tag.

The protocol for detecting anti-DNP recruitment to bacterial cell surfaces is as follows. Bacteria were incubated with the desired concentration of DNP-conjugated PCCs in PBS-BSA for 1 h at 37 °C. Residual PCCs were washed off by using PBS-BSA and bacteria were incubated with anti-DNP antibody (1 : 1000) for 1 h at 37 °C. After washing, bacteria were incubated with HRP conjugated secondary antibody (Bio-Rad) at a dilution of 1 : 10 000 for 1 h at 37 °C. Unbound antibody was washed off by using PBS-BSA and the cells were then developed by using TMB reagent. The reaction was quenched using 2 N H2SO4 and absorbance was measured at 450 nm. Anti-MrkA antibody was procured from Biorbyt (orb51318) and used at a concentration of 5 μg mL⁻¹. K. pneumoniae antiserum was obtained from abcam (ab20947) and used at a concentration of 5 μg mL⁻¹. The DNP-conjugated PCCs used for anti-DNP recruitment assays had a peg5-peg5 linker between the N-terminus of the PCC and the DNP tag.

Opsonophagocytic killing (OPK) assays

*Klebsiella pneumoniae* BAA1705 was treated with DNP-conjugated PCCs followed by incubation with anti-DNP antibody, as described above in the Cell-Based ELISAs section. These opsonized bacteria were then used to infect BMDMs at a multiplicity of infection of 50 for 0.5 h at 37 °C. Bacteria were then washed by using RPMI 1640 medium and BMDMs were left in cell culture medium containing gentamicin (100 μg mL⁻¹). After 1 h and 24 h, BMDMs were washed to remove gentamicin and intracellular bacteria were harvested by lysing BMDMs in RPMI medium containing 0.2% Triton X 100. Bacterial CFUs were enumerated by plating onto LB agar. Bacterial counts at 1 h indicated the degree of opsonization while those at 24 h served as a measure of microbialic activity of macrophages. For these measurements, anti-MrkA antibody and *K. pneumoniae* antiserum were used at dilutions of 5 μg mL⁻¹ each. The DNP-conjugated PCCs used for the phagocytosis and OPK assays had a peg5-peg5 linker between the N-terminus of the PCC and the DNP-modified lysine.

Flow cytometry

Cytometry measurements were used to quantify AR-PCC-driven opsonization. For these measurements, *K. pneumoniae* cells were cultured in G-CAA medium overnight, washed, incubated with DNP-conjugated PCC at a desired concentration, washed, incubated with anti-DNP antibody conjugated with AlexaFlour488, washed, then resuspended in PBS media. The samples were stored at 4 °C for 2 d before performing cytometry measurements. The cytometer was calibrated by using Rainbow fluorescent beads (3.5 μm diameter) from BD (559123), which aided identification of single *K. pneumoniae* cells in subsequent measurements. Samples were excited with 488 nm light and the fluorescence emission at 530 nm was measured. Sample with no AlexaFlour488 stain was used as fluorescent minus one (FMO) control for gating of DNP+ population. FACS data was analyzed on Flowjo v10 software, and the resulting histograms each include fluorescence data from >18 000 cells for strain BAA 2146 (Fig. 6B) and >4000 cells for strain BAA 1705 (Fig. S11, ESI†). PCCs used for flow cytometry assays had a peg5-peg5 linker between the N-terminus of the PCC and the DNP tag.

Conflicts of interest

J.R.H is a co-founder and B.T.L is an employee of Indi Molecular Inc., which seeks to commercialize PCC technology.

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Notes and references


62 Private communication with Indi Molecular Inc.


