Introduction

Vaccination has been one of the most effective ways to control pathogens and prevent disease in the history of medicine. Essentially all successful prophylactic vaccines work by inducing neutralizing or protective antibodies against pathogens (polio, small pox, tetanus) [1–3]. The first attempts to vaccinate were dominated by live vaccines as e.g., vaccinating against small pox using first small pox virus itself followed by vaccinia virus. A constant search for safer vaccines has led to the development of virus-like-particles (VLPs) derived from the RNA bacteriophage AP205 for epitope-based vaccines.

Methodology/Principal Findings: Peptides of angiotensin II, S.typhi outer membrane protein (D2), CXCR4 receptor, HIV1 Nef, gonadotropin releasing hormone (GnRH), Influenza A M2-protein were fused to either N- or C-terminus of AP205 coat protein. The A205-peptide fusions assembled into VLPs, and peptides displayed on the VLP were highly immunogenic in mice. GnRH fused to the C-terminus of AP205 induced a strong antibody response that inhibited GnRH function in vivo. Exposure of the M2-protein peptide at the N-terminus of AP205 resulted in a strong M2-specific antibody response upon immunization, protecting 100% of mice from a lethal influenza infection.

Conclusions/Significance: AP205 VLPs are therefore a very efficient and new vaccine system, suitable for complex and long epitopes, of up to at least 55 amino acid residues in length. AP205 VLPs confer a high immunogenicity to displayed epitopes, as shown by inhibition of endogenous GnRH and protective immunity against influenza infection.
structure on the epitope. In addition, some of the viral carriers face major challenges for regulatory approval due to infectivity or presence of DNA and resistance genes.

Recently, we have described new vaccines based on a VLP derived from the RNA bacteriophage QB, made from 180 subunits assembling into an icosahedral capsid in E. coli. Epitopes conjugated chemically to these VLPs are displayed in a highly repetitive and organized manner, inducing strong immune responses in all species tested [14]. Remarkably, the highly repetitive display allows to elicit strong antibody responses even against self-antigens [15–20]. Indeed, vaccines based on epitopes coupled to QB have shown clinical efficacy in smoking cessation and the treatment of hypertension [21–24].

In this report, we describe a new RNA phage based VLP-vaccine carrier allowing both N- or C-terminal fusion of epitopes. The VLP is formed from 180 copies of the coat protein. We show that long (up to at least 55 amino acids) or complex epitopes (containing multiple cysteine residues) can be fused to this AP205-derived VLP-carrier. One vaccine successfully induced antibodies against GnRH, a self-antigen involved in fertility and prostate cancer. A second epitope fused to AP205 VLPs was the extracellular domain of the Influenza A M2 protein. Vaccination of mice with M2-AP205 resulted in 100% protection from lethal infection with influenza virus.

Results

AP205 has been recently identified as a new RNA bacteriophage infecting the Gram-negative bacteria Acinetobacter sp. Sequence alignment of its coat protein with other RNA phage coat proteins is difficult since only 5 amino acids (aa) are conserved among all RNA phages isolated so far, even when using structural information [25,26]. The N- and C-termini of RNA bacteriophage coat proteins come close together at the three-fold axis of their icosahedral shell and are often involved in interactions with each other, or shielded from the surface. We generated four AP205 genes, allowing rapid insertion of oligonucleotides or genes encoding a desired epitope to both the N- and C-terminus of AP205 coat protein, using either a short (3 or 4) or long (8 or 11) aa spacer to ensure optimal display of the epitope (Fig. 1). Whilst the shorter spacers were combinations of G and S residues, the long spacer sequences used were modifications of a linker sequence present in the so called A1 extension of QB coat protein [27].

Table 1 presents a list of epitopes vs. VLP vector and spacer variants used for the construction of model vaccines. In a first set of experiments, a short peptide, angiotensin II (Ang II), was fused to the N- and C-terminus of the coat protein, using all four vectors. The resulting fusion proteins were expressed in E. coli, and spontaneously assembled into VLPs as detected by electron microscopy (Fig. 2). VLPs were purified by gel filtration and tested by Coomassie staining after LDS-PAGE (Fig. 3a). Western blot analysis of the fused proteins using an antiserum against AP205 (Fig. 3b) or Ang II (Fig. 3c) confirmed presence of the angiotensin peptide. In the absence of a crystal structure of AP205 VLP (Tars et al., in progress), the accessibility of epitopes fused at any one of the coat protein termini can only be tested experimentally. To this end, we assessed the ability of the various AP205 VLPs displaying Ang II to inhibit binding of an Ang II-specific antiserum to an Ang II-conjugate coated on an ELISA plate (Fig. 4a and b). VLPs were preincubated with the antiserum specific for Ang II and remaining free antibodies were detected by ELISA. As seen in Fig. 4a and 4b, both the VLPs with Ang II fused to the C-terminus, or the N-terminus of AP205 coat protein, inhibited binding of the anti-Ang II antiserum. Wild type AP205 VLP did not inhibit antibody binding, confirming the specificity of the inhibition and thereby the accessibility of the peptides on the surface of the VLPs.

The display of the fused epitopes on the surface of the AP205-VLP would predict that these epitopes should be highly immunogenic. In order to test immunogenicity and at the same

Table 1. Epitopes and AP205 VLPs referred to in the manuscript.

| Epitope | Designation | Source | Length, aa | C-terminal fusions | N-terminal fusions |
|---------|-------------|--------|------------|-------------------|-------------------|
|         |             |        | short linker | long linker | short linker | long linker |
| Ang II  | Angiotensin II, aa 1–8 | 8 | + | + | + | + |
| D2      | S.lyphi outer membrane protein, aa 266–280 | 15 | + | + | + | + |
| CXCR4   | CXCR4, N-terminal extracellular part, aa 1–39 | 39 | – | – | + | – |
| Nef55   | HIV Nef, aa 66–100, 132–151 | 55 | – | + | – | – |
| GnRH    | Gonadotropin releasing hormone, aa 1–10 | 10 | – | + | – | – |
| M2      | M2, N-terminal ectodomain, aa 2–24 | 24 | – | – | + | – |

AP205 VLPs are designated by the linker used to fuse the corresponding epitope. Combinations described in the manuscript are referred to with a “+”.

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time to assess whether AP205 coat protein could accommodate epitopes larger than the 8 aa peptide Ang II, we fused a sequence encoding the Salmonella typhi outer membrane protein derived 15 aa peptide D2 [28] to AP205 coat protein, using our four vectors. The resulting coat proteins were expressed in E.coli and the VLPs purified. All four coat-protein fusions assembled into VLPs, as was tested by size exclusion chromatography (not shown) and electron microscopy (Fig. S1). After purification of chimeric AP205-D2 VLPs by gel filtration to reliable purity confirmed by LDS-PAGE (Fig. S2), competitive ELISA showed that the D2 peptides are displayed on the outer VLP surface to the same extent as the Ang II peptide in the corresponding VLPs (not shown). In order to further test accessibility of the peptide on the VLP and to test the immunogenicity of the particles, we immunized Balb/c mice at fortnightly interval with 25 μg of D2-VLPs in the absence of adjuvant. As shown in Fig. 5, all 4 VLPs were highly immunogenic, and a robust titer was measured after two injections. The response is specific for the D2 peptide, as AP205 linked to a control peptide induces antibodies that do not bind to D2 (data not shown). Display of the D2 peptide with a long linker at the N-terminus of AP205 VLP seemed, however, to yield less immunogenic VLPs than using a short linker at the N-terminus, or displaying the peptide at the C-terminus of the VLP carrier. These data independently confirm the accessibility of fused epitopes on the surface of AP205 VLP.

In order to investigate whether peptides of greater length can be displayed on AP205 VLP, we fused the N-terminal 39 aa extracellular part of CXCR4 to the N-terminus, and a 55 aa HIV polyepitope containing Nef consensus T-cell epitopes to the C-terminus of AP205 VLP. Both fusion coat proteins assembled into VLPs and displayed their epitopes, as was confirmed by electron microscopy (Fig. S3), SDS-PAGE and Western blot (not shown). Thus, peptides as large as 55 aa in length can be fused to AP205 VLP.

We wanted next to see whether the high immunogenicity of the AP205-VLP vaccines could generate antibodies neutralizing their target and having a useful therapeutic effect in vivo. To this end, we selected GnRH as a target for neutralisation in mice. This provided additionally the opportunity to test whether AP205 VLP based vaccines could break B cell unresponsiveness towards self-
antigens. This is important, as a number of attractive molecules currently targeted for immunotherapy are self-antigens. Antibodies neutralising GnRH have the potential to act as androgen-deprivation therapy used to date to treat prostate cancer [29–31], since prostate cancer cells depend on testosterone for growth [32]. Fertility management may be an additional potential use of such a vaccine [33]. Therapeutically meaningful induction of neutralising antibodies against GnRH should lead to a decrease in LH and FSH production and subsequently to reduced levels of androstenedione and gonadal steroids like estrogens and testosterone. This in turn will cause testis atrophy that can be monitored as testis weight reduction.

We therefore fused murine GnRH to the C-terminus of AP205 coat protein using a long spacer. The fused coat protein assembled to VLPs in E. coli as determined by electron microscopy of purified VLPs. Presence of the peptide in AP205-VLPs was confirmed by western blot. Male adult mice were immunized once at 9 weeks of age with 50 μg of AP205-GnRH and boosted three weeks later with the same dose. Neutralization of GnRH activity was assessed by measuring testis weight 70 days after initial immunisation. As seen in Fig. 6a, immunisation with AP205-GnRH led to a 38% decrease in testis weight (p = 0.003, t-test), demonstrating the neutralization activity of raised antibodies. Antibody titers and testosterone levels were monitored on day 0, 21, 28, 45 and 70. A high antibody titer was already achieved after the first immunisation in all animals on day 21, and was further boosted after the second injection (Fig. 6b), further confirming that

Figure 3. AP205-Ang II VLPs analyzed by dodecyl sulphate PAGE and Western blot. After purification by gel filtration, AP205 VLPs carrying Ang II peptide fused to the C-terminus (AP441 and AP442) or N-terminus (AP446 and AP447) of AP205 coat protein were heated at 95°C in sample buffer for 5 minutes prior to loading. (a) Coomassie-stained LDS-PAGE of corresponding chimeric proteins; (b) Western blot with an AP205 coat protein-specific rabbit serum; (c) Western blot with an Ang II-specific mouse serum. The migration of all four AP205 coat proteins carrying Ang II is retarded in comparison to wt AP205 coat protein (130 aa). Accordingly, migration of coat protein fused to AP205 coat protein using a long spacer (AP442 and AP447) is slower than the migration of coat proteins containing a short spacer (AP441 and AP446).

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Figure 4. Display of Ang II on AP205 VLP assessed by inhibition ELISA. Ang II was conjugated to RNAse, and coated on ELISA plates. The anti-Ang II serum was pre-incubated with serial dilutions of the 4 VLPs displaying Ang II or wt AP205, and subsequently added to the wells coated with RNAse-Ang II. Signals shown on the Figure are normalized and are given as % binding. The VLP concentration [μg/ml] was log transformed. (a) VLPs displaying Ang II at the C-terminus of AP205 coat protein using a short (AP441) or long spacer (AP442) or wt AP205 VLP (AP205). (b) VLPs displaying Ang II at the N-terminus of AP205 coat protein using a short (AP446) or long spacer (AP447) or wt AP205 VLP (AP205).

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the immunisation had efficiently overcome B-cell unresponsiveness. The response against GnRH is specific, as antibodies generated against AP205 alone do not recognize GnRH (data not shown). As expected, GnRH neutralization led to a decrease in testosterone levels, and we observed a reduction in testosterone starting from day 28, coincidentally with reaching high antibody titers after the booster immunisation (Fig. 6, b and c). The median of the testosterone level, averaged between day 28 and 70, was 81% lower in the AP205-GnRH treated animals when compared to untreated animals (p = 0.008, Mann-Whitney test).

The experiments above demonstrate that AP205 VLP induced rapidly a high antibody response in all animals after a single injection. This is particularly advantageous in cases where a rapid immune response has to be mounted against a pathogen, for example in the case of an influenza pandemic. Currently, influenza vaccines contain as antigens hemagglutinin and neuraminidase from three strains selected every year by the WHO. Use of a conserved antigen not subject to yearly variation would be a considerable improvement. The N-terminal M2 epitope of influenza is such a conserved epitope [34–36]. Antibodies raised against M2 do not neutralise influenza virions, but rather act through antibody dependent cytotoxicity (ADCC) to limit and clear viral infection [35]. In order to test whether an AP205 VLP based vaccine may elicit a high enough antibody responses to protect against influenza virus-infection, we fused the N-terminal ectodomain of Influenza A M2 protein to the N-terminus of AP205 VLP using a short spacer. The coat proteins fused to M2 assembled into VLPs, as was assessed by electron microscopy. The protective effect of the M2-AP205 VLPs against a challenge with the influenza PR 8 strain was assessed in C57BL/6 mice. The animals were immunized twice with M2-AP205 or AP205 only, and challenged intranasally (i.n.) 13 days after the last injection with a lethal dose of influenza virus. The mice mounted a strong immune response against M2 as well as against AP205 already after the first injection, and this response was boosted after the second injection (Fig. 7a). The mice immunized with control AP205 produced high antibody titers against AP205 but not against M2. All mice immunized with M2-AP205 were protected against the influenza challenge, while for the control group, all mice succumbed to the lethal infection (Fig. 7b). While control animals experienced severe weight and temperature loss M2-AP205 vaccinated animals were largely protected from symptoms of disease.
Discussion

VLPs are very potent inducers of antibody responses not only against themselves but also against essentially any antigen displayed on them. Here we describe a new RNA-phage derived VLP that allows genetic fusion of peptide epitopes for display on their surface. Using various antigens, including a self-peptide, we demonstrate that this VLP-epitope display platform is able to induce potent antibody responses neutralizing GnRH in vivo and protecting mice from lethal infection with influenza virus. Moreover, the antibody response generated with AP205 VLPs is largely comparable in magnitude and IgG-subtype pattern (Th1) to what is obtained using RNA phage Qβ (unpublished), whose immunogenicity has been demonstrated in humans [22–24].

Remarkably, these AP205-VLPs were very receptive in accepting foreign sequences, and self-assembled even when an epitope with multiple cysteines such as the M2 epitope was fused to the coat protein. The longest epitope we have fused so far to the N-terminus of AP205 coat protein was 39 aa long, while to the C-terminus it was 55 aa in length. Chemical peptide synthesis becomes challenging with peptide length beyond 40 aa and this new technology therefore widens the range of epitopes available for vaccination. The ability to fuse to both ends of the AP205 VLP-subunit allows better display of N-or C- terminal epitopes which is often difficult for other VLPs. Indeed, VLPs such as those derived from the Hepatitis B core antigen (HBcAg), the bacteriophage FQ or filamentous phage particles only allow fusion at one terminus of the subunit [37,38]. The availability of VLPs suitable for fusing epitopes at either end is important, since for a number of epitopes, the protective immune response raised, is specific for their N- or C-terminal part (unpublished and [21]). In addition, this will allow to more exhaustively test epitopes when screening a whole pathogen genome or “antigenome” for antigenicity as recently described [39].

A number of epitope-based vaccines targeting self-antigens are in development targeting e.g. Ang II, GIP, Aβ or GnRH [4]. We have shown in this study that AP205 VLP fused to a self-antigen is able to overcome B-cell unresponsiveness. In addition, the antibodies neutralized their target, GnRH, as shown by testosterone level reduction and testis atrophy. An anti-GnRH vaccine may find application as a therapeutic vaccine for prostate cancer. Several vaccines are already in development for this indication.

Similarly, an influenza-pandemic vaccine with a rapid onset of action is highly desirable. The M2 response we obtained is remarkable, since not only was mortality avoided, but also morbidity greatly reduced. Moreover, a 100% protection was achieved even with the high dose of virus used in the challenge. High antibody titres were already reached by day 14 in all animals, demonstrating the rapid onset of the antibody response when immunizing with AP205 RNA phage VLP. RNA bacteriophage VLPs are produced recombinantly in E. coli, achieving high yield at a low cost of goods. An additional advantage of using bacteria rather than eggs or eukaryotic cells for production is speed. These features of the technology are highly relevant to the design and production of vaccines for a pandemic. They are also advantageous for the design of vaccines against emerging pathogens or for biodefense applications.

Material and Methods

Cloning of coat protein fusions

Four intermediary plasmids for convenient subcloning of fragments coding for epitopes by cassette mutagenesis were prepared by PCR with pAP283 as template (Gielens et al., unpublished). Plasmids pAP378 and pAP382 were designed to introduce epitopes with a short (GSGG), respectively a long amino acid spacer (GSGTAGGGSGS) at the N-terminus, while plasmids pAP409 and pAP405 were designed for introducing epitopes with a short (GSG), respectively a long amino acid spacer (GTAGGGSG), at the C-terminus of AP205 coat protein. Oligodeoxynucleotide pairs were the following: p2.561 (5’-TGGGATCCATGG-
GATCCGGAGGGCCAAAATAGCAGCAACC-3′ and p1.46 (5′-TGAAGCTTAAAGCAGCATATGAGCAGCATGG-3′) for pAP378, p2.589 (5′-TGGCATGCTTGTCGCAATTAA-GCCAAAATGCCGGAACCCGCGGGCTGAAGCTTAAAGCAGCATATGAGCAGCATGG-3′) and p1.46 (5′-TGAAGCTTAAAGCAGCATATGAGCAGCATGG-3′) for pAP382, p1.35 (5′-TGTCAGGGAGGCTGGAAGCCTGGGAACCGCCTCCTGCGGTTCCAGCAGTAGTATC-3′) and p2.587 (5′-TGATGATCCTCCTGGGATCGAGCAACCGTGAGTTGAGCGATACG-3′) were subcloned into plasmid pAP405 (41). In pAP378 asparagine 14 is mutated to aspartate. Fragments coding for epitopes to be fused at the N-terminus were cloned as a NcoI cassette (Fig. 1). The primer pairs for the plasmids coding an epitope fused to AP205 coat protein N- or C-terminus are described in the supplementary material and methods (Text S1).

Expression and purification of AP205-fusion VLPs

Plasmids were transformed into E.coli JM109. A seed culture was prepared by inoculating a single colony into LB medium containing 20 mg/l Ampicillin and growing the culture overnight at 37°C without shaking. For expression, the overnight culture was diluted in M9 medium supplemented with casaminoacids (Difco) and containing 20 mg/l Ampicillin and growth of the culture carried out at 37°C with vigorous aeration for 14–20 hours.

Cells were resuspended in TEN buffer (20 mM Tris-HCl, 5mM EDTA, 150 mM NaCl, pH 7.8) containing 1 mg/ml lysozyme and 0.1% Tween 20, and lysed by three freeze thaw cycles followed by ultrasonification. Clarified lysates were purified by one or more sequential gel filtration steps using CL-4B, CL-2B and Sepharose 6B columns (GE Healthcare). Some VLPs were additionally purified by ultracentrifugation over a CsCl gradient. Concentrations of the purified proteins were determined by the Bradford test. For the AP205-GnRH vaccine, two bands reactive with an anti-GnRH antiserum were observed when using vector pAP405 for expression. This wasn’t observed in any other construct. Subcloning into a pK2K222 derivative however led to a single band. The immunogenicity of both constructs was equivalent.

Electron microscopy: Negative staining

Suspensions of VLPs were adsorbed on carbon-formvar coated grids and stained with 1% phosphotungstic acid (pH 6.8). The grids were examined with a JEM 100C electron microscope (Jeol Ltd., Tokyo, Japan) at an accelerating voltage of 80 kV. Photographic records (negatives) were performed on Kodak electron image film and electron micrographs were obtained by scanning of negatives with EPSON 2480 PHOTO scanner.

Anti-peptide antibody response measured by ELISA

Peptides modified to contain a cysteine at their N- or C-termini, and in some cases additionally a diglycine spacer, were conjugated to ribonuclease A (RNase) using the cross-linker SPDP (Pierce, Rockford, IL, USA). The resulting conjugate was coated overnight at 4°C on maxisorb plates (Nunc, Denmark), at a concentration of 10 μg/ml. Microtiter plates were blocked with 2% bovine serum albumin in PBS supplemented with 0.05% Tween 20. Binding of the sera was detected with a Horseradish-peroxidase goat anti-mouse IgG conjugate. Binding of VLPs to anti-peptide sera was assessed in an inhibition ELISA assay. Serial VLP dilutions were pre-incubated with anti-peptide sera on blocked microtiter plates. Remaining free antibody was detected by ELISA using an RNase-peptide conjugate as described above. The sera specific for Ang II were raised using Ang II conjugated at its N-terminus to carrier Qb.

Immunization of mice

Animal experiments were conducted according to guidelines set by the Swiss Federal Veterinary Office (BVET) and were approved by the Kantonale Veterináramt of Zurich. Balb/c mice (n = 3) were immunized subcutaneously on day 0 and 14 with 25 μg proteins of AP410, AP420, AP421 and AP422. The proteins were diluted to a final volume of 200 μl in PBS, and 100 μl were injected in the left and right inguinal region of each animal. Animals were bled on day 14 and 21 for measurement of antibody titre.

Anti-GnRH immunization

Male C57 BL/6 mice (n = 5) were either immunized subcutaneously with 50 μg of AP205-GnRH on day 0 and 21, or untreated (control animals). Animals were additionally bled on day 28 and 45, and were sacrificed on day 70. Testes were sampled, weighed and testosterone was measured in serum.

Anti-M2 immunization and influenza challenge

C57BL/6 mice were kept in our animal facility, free of specific pathogens. Influenza Virus PR8 (A/PuertoRico8/34, H1N1) was originally provided by J. Pavlovic, University Zurich. C57BL/6 mice (n = 6) were immunized on day 0 and 18 with 50 μg of M2-AP205 or AP205. On day 31, mice were infected intranasally (i.n.) with 4×LD50 (4×103 p.f.u./ml) of PR8 virus (H1N1) in endotoxin-free phosphate buffer saline (PBS) under isoflurane anaesthesia. Mice were monitored for another 22 days thereafter, when body weight and rectal body temperature were determined daily.

Supporting Information

Text S1 Supplementary Material and Methods. Found at: doi:10.1371/journal.pone.0009809.s001 (0.04 MB DOC)

Figure S1 Electron micrographs of AP205-D2 VLPs displaying the D2 peptide. The D2 peptide was fused to the C-terminus (AP418 - short linker, AP420 - long linker) or the N-terminus (AP421 - short linker, AP422 - long linker) of AP205 coat protein. Found at: doi:10.1371/journal.pone.0009809.s002 (5.78 MB TIF)

Figure S2 Coomassie-stained LDS-PAGE of AP205-D2 VLPs. AP205 VLPs carrying D2 peptide fused to the C-terminus (AP418 and AP420) or N-terminus (AP421 and AP422) of AP205 coat protein, analyzed by LDS PAGE. Lane 1 is the Marker, lane 2 AP205 VLP conjugated chemically to D2 peptide at an epitope density of 1 peptide per AP205, lane 3 AP205 conjugated chemically to 2 D2 peptides at an epitope density of 2 peptides per AP205. The bands corresponding to 1 peptide per AP205 and 2 peptides per AP205 are marked by a star, the band corresponding to 3 peptides by two stars, and the band corresponding to 3 peptides by 3 stars. Found at: doi:10.1371/journal.pone.0009809.s003 (0.89 MB TIF)

Figure S3 Electron micrographs of chimeric AP205 VLPs displaying longer foreign epitopes. Electron micrographs of AP205 VLPs with Nef55 epitope fused to the C-terminus via short linker (AP418), CXCR4 epitope fused to the N-terminus via long linker (AP459), CXCR4 epitope fused to the N-terminus via short linker (AP543), and M2 epitope fused to the N-terminus via short linker (AP551). Found at: doi:10.1371/journal.pone.0009809.s004 (3.14 MB TIF)
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Author Contributions

Conceived and designed the experiments: ACT RR IC EM VO GTJ PS PP MFB. Performed the experiments: RR IC EM VO. Analyzed the data: ACT RR IC EM VO GTJ PS MFB. Wrote the paper: ACT PP MFB. Preformed all the work on Influenza Virus: NS.