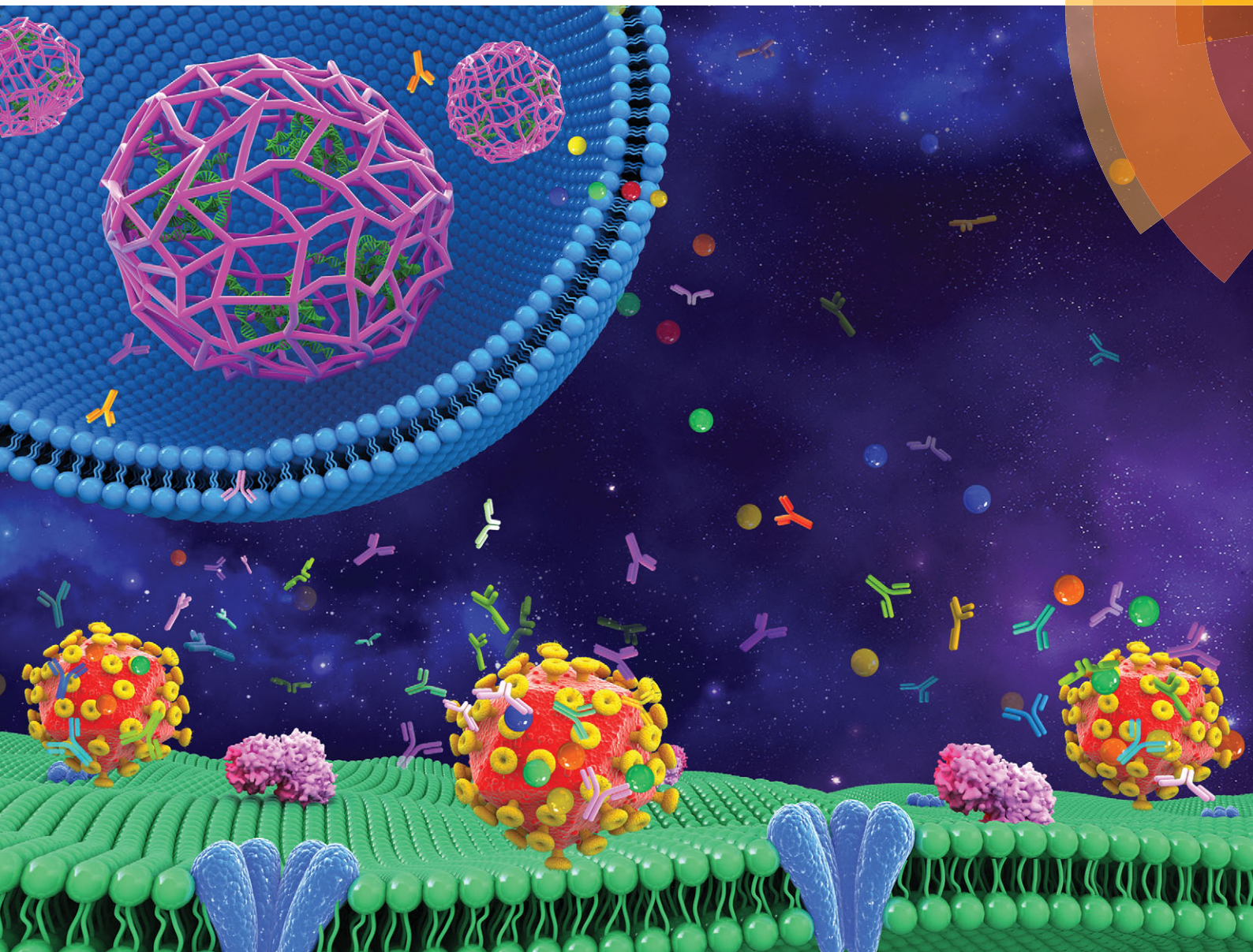


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ISSN 2055-6756



COMMUNICATION

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Cite this: *Nanoscale Horiz.*, 2016, 1, 135

Received 14th September 2015,
Accepted 14th December 2015

DOI: 10.1039/c5nh00064e

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In situ formation of peptidic nanofibers can fundamentally optimize the quality of immune responses against HIV vaccine†

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Herein, we report that the *in situ* formed peptidic nanofibers facilitate the induction of multiple crucial immunities against HIV DNA vaccine, including polyfunctional T cell response, broad IgG subclasses response, and V1/V2 loop-specific antibody response, all of which can hardly be triggered by HIV DNA vaccine alone. Such novel *in situ* formation fundamentally overcomes the big hurdle for the applications of such nanofibers, which previously can only trigger these crucial immune responses *via* adding exogenous alkaline phosphatase. Such robustness of peptidic nanofibers for inducing crucial immune responses may allow better inhibition against HIV than reported materials.

Since the first case of human immunodeficiency virus/HIV discovered in 1983, this deadly virus has caused great harm to human health worldwide. Vaccination is recognized as one of the most effective and low-cost ways to control HIV spread worldwide.¹ Up to now, a major goal for HIV vaccines is to induce good-quality, not only high-quantity, immune responses to achieve high protection efficacy of HIV vaccines.² For example, a strong, mono-functional T cell response (a single T cell simultaneously secretes only one or two cytokines) is limited for the

Conceptual insights

This work employs peptidic nanofibers to significantly improve the efficacy of vaccines for HIV, a virus against which human beings currently have no effective vaccine. As adjuvant, such nanofibers can both improve the quality of immune responses against HIV and provide convenience for administration. These two improvements allow a significant advancement in the field of HIV vaccine, which is far limited because of the immense difficulties in obtaining a high-quality immune response and complex protocols for administration. This work provides an exciting way to overcome such difficulties *via* safe and convenient adjuvants, the *in situ* formed peptidic nanofibers. The superior performance of nanofibers as adjuvants on fundamentally optimizing HIV vaccine is a disruptive advancement in the field of adjuvant research, and will further promote the development of next-generation vaccines against AIDS and other diseases.

clearance and control of the virus *in vivo*.³ Likewise, a potent antibody titer against HIV alone does not necessarily translate into good neutralization (blocking the infection of virus).⁴ In contrast, some unique profiles of the immune response, such as both the polyfunctional T cell response^{5–8} and the broad IgG subclasses response,^{9–12} are featured in LTNPs, and the IgG response against the crucial epitope (V1/V2 loop of HIV Env) can effectively reduce HIV infection risk.¹³ We thus wonder if there is a way to optimize such profiles of immune responses of HIV vaccines.

Nanomaterials have obtained widespread attention because of their specific bio-effects *in vivo*, and also have been applied in the field of biomedicine.^{14,15} In particular, peptidic nanofibers may be useful for vaccination. They not only act as basic building blocks to comprise the complex cytoskeleton network to maintain cellular structure, but also construct the complex extracellular matrix to mediate the interactions between cells and their surrounding microenvironment.¹⁶ In recent years, numerous studies focused on investigating the bio-effects of peptidic nanofibers formed *in vitro* by some inducers, such as enzymes, heating, change of pH and so forth.^{17,18} However, such nanofibers still face many problems for practical applications,

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c5nh00064e

because they have to be constructed *in vitro* via adding inducers before injection into the body. A latest study from the Bing Xu's group reported that peptidic nanofibers can self-assemble on the surface of cancer cells *in vitro*, using the ALP secreted by cells themselves.¹⁹ It suggested that peptidic precursors might self-assemble *in vivo* dependent on an endogenous inducer, such as ALP. In one of our recent studies, a kind of peptidic precursor named NMe [it contains four amino acids backbone (glycine-phenylalanine-phenylalanine-tyrosine), in which the N-terminal and the C-terminal are modified by naphthalene acetic acid and the methylamino group respectively] can self-assemble into nanofibers *in vitro* after the hydrolysis by ALP. Such peptidic nanofibers as adjuvants can increase the strength of the T cell and the IgG response against the HIV DNA vaccine in mice, *via* compressing HIV DNA vaccines into more compact spheres to enter antigen-presenting cells more effectively.²⁰ However, all above immunological effects were triggered by peptidic nanofibers which were assembled *in vitro* via addition of exogenous ALP. We thus wonder whether such peptidic nanofibers have similar effects when they are directly injected without exogenous ALP (a more convenient way for practical applications), considering endogenous ALP has an extensive contribution *in vivo*.

Herein, we investigated the effects of different assembly modes of peptidic nanofibers on regulating crucial immune responses induced by the HIV DNA vaccine, including the polyfunctional T cell response, the broad IgG subclasses response and the V1/V2-specific IgG response. Precursors of nanofibers that were directly injected in a body (a convenient route for application) had the same immunological effects as the nanofibers formed *in vitro* via addition of exogenous ALP, thus showing the robustness of such nanofibers. Our results not only explored multiple crucial immunological effects of peptidic nanofibers, but also cleared the hurdles for large-scale clinical applications by tremendously simplifying the route for administering such peptidic nanofibers.

Results and discussion

For enzyme-triggered peptidic nanofibers, the assembly mode directly determines their suitability as vaccine adjuvants. In most previous studies, enzyme-triggered nanofibers require exogenous enzymes to initiate the assembly process. Such operation procedure severely compromises the convenience, because users have to prepare the nanofibers every time before injection. Moreover, the safety of the vaccine will depend on the purity and safety of the enzymes, in addition to those of the nanofibers themselves. Having to include an enzyme in a vaccine will also significantly increase the cost for practical applications. We thus considered whether such nanofibers would successfully form *in situ*, employing only the endogenous enzymes *in vivo*.

To verify the possibility that our nanofibers can be assembled *in vivo* using endogenous ALP, we first investigated the formation of such fibers in the cell culture system. We herein designed three experiment groups (Fig. 1A). (i) HeLa

cells (a cancer cell line, 1×10^5 cells) cultured with the 0.1 wt% NMe precursor-added medium; (ii) medium supplemented by both ALP (60 U ml^{-1}) and 0.1 wt% NMe precursors (positive control). HeLa cells lysis solution (1×10^5 cells) has similar catalytic capability to NMe precursors as 60 U ml^{-1} ALP (Fig. S1, ESI[†]); (iii) HeLa cells (1×10^5 cells) cultured with the medium alone (negative control). After 24 hours, the nanofiber network was observed in groups (i) and (ii), but not in group (iii) (observed *via* TEM). The range of length of nanofibers was between 15 nm and 300 nm (Fig. 1A). The range of diameter of nanofibers was between 15 nm and 40 nm (Fig. 1A). These results indicated that NMe can self-assemble into nanofibers in the cell system without exogenous ALP.

We wonder whether the *in situ* formed nanofibers (without exogenous ALP) have similar immunological effects to that formed *in vitro* via adding exogenous ALP before injection. To answer this question, we designed three modes of nanofibers (Fig. 1B). (i) Mode I (*in situ* formation, without exogenous ALP), the mixture of the DNA vaccine and NMe was injected directly into the mice body; (ii) Mode II, NMe assembled into nanofibers by adding exogenous ALP before combination with the HIV DNA vaccine, then were injected into mice together; (iii) Mode III, the DNA vaccine and NMe were mixed together in the presence of ALP before injection into mice. Mice injected with the DNA vaccine alone was used as a control.

We first investigated whether the precursors of NMe can bind to the HIV DNA vaccines to understand the probable working mechanism in above three models. *Via* an isothermal titration calorimetry (ITC) assay, we quantified the intermolecular interactions between the precursors of NMe and the HIV DNA vaccine (Fig. 1C). By titrating the precursors of NMe ($250 \text{ } \mu\text{M}$) into the HIV DNA vaccine (153 nM), we obtained several critical thermodynamic parameters including binding affinity (K), enthalpy changes (ΔH) and binding stoichiometry (N). Approximately, one HIV DNA molecule will bind 13 precursors of NMe ($N = 0.0806 \pm 0.00517$ sites) to form an intermediate. There is the binding (binding affinity is $2.68 \pm 0.701 \times 10^5 \text{ M}^{-1}$) between the precursors of NMe and the HIV DNA vaccine (Fig. 1C). Such binding might help NMe to compress loose DNA into the compact structure along with the formation of the nanofiber network. The compact DNA will be more effective to enter the cells, finally causing enhanced immunity according to our previous report.²⁰

We next investigated the effects of three different formation modes of nanofibers on regulating the polyfunctional T cell response induced by the HIV DNA vaccine, because in the clinic, such immunity is inversely correlated with the disease progression in HIV infected individuals.^{6,8} In this study, the polyfunctionality of the T cell is defined as a single cell which can simultaneously secrete at least three cytokines among the following five cytokines, interleukin 2 (IL-2), interleukin 4 (IL-4), interferon gamma (IFN- γ), tumor necrosis factor alpha (TNF- α), and cluster of differentiation molecule 107a (CD107a). All these cytokines are crucial parameters used to evaluate the polyfunctionality of T cells of the HIV vaccine.^{6,21} "5-Cytokine-secreting T cells" indicate cells that simultaneously secrete all

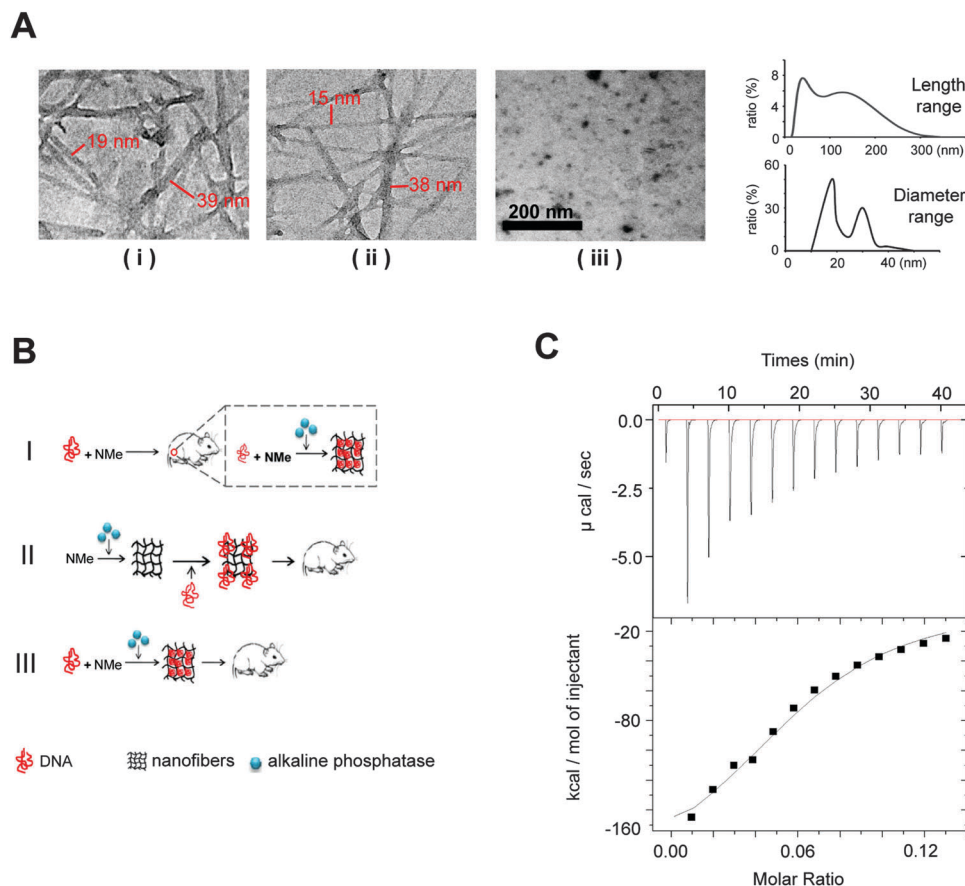


Fig. 1 The formation of peptidic nanofibers in the cell culture system, and the binding between NMe precursors and HIV DNA vaccines. (A) (i) HeLa cells (a cancer cell line, secrete ALP by themselves) cultured with the NMe precursor-added medium; (ii) medium supplemented by both ALP and NMe precursors (positive control); (iii) cells cultured with the medium alone (negative control). After 24 hours, the nanofiber network was observed by TEM in groups (i) and (ii), but not in group (iii). We randomly chose 20 fibers in each sample. By measuring the length (the distance between two nodes) and diameter of each fiber, we estimated the rough range of length and diameter of these fibers. The range of length of nanofibers was between 15 nm and 300 nm. The diameter range of nanofibers was between 15 nm and 40 nm. (B) The schematic diagram of three modes. 'I' indicates that the mixture of the DNA vaccine and NMe was injected directly into the mice body (without premixing with ALP); 'II' means that NMe assembled into nanofibers by ALP before combination with the HIV DNA vaccine; 'III' means that the DNA vaccine and NMe were mixed together in the presence of ALP before injection into the mice. (C) In the ITC test, NMe precursors (250 μM) were titrated into the solution of the HIV DNA vaccine (153 nM). The results revealed the binding between the precursors of NMe and the HIV DNA vaccine.

these five cytokines. "4-Cytokine-secreting T cells" and "3-cytokine-secreting T cells" respectively represent the cells that secrete four and three cytokines among above five cytokines. The sum of the percentage of T cells secreting 3-, 4- and 5-cytokines represents the total strength of the polyfunctional T cell response. The 5-cytokine-secreting T cell response (1.05% for Mode I, 0.93% for Mode II and 1.37% for Mode III, Fig. 2A) was effectively induced in all three modes to a level not achievable by the DNA vaccine alone. Also, significantly increased ratios of T cells expressing 4-cytokines and 3-cytokines (Mode I: 2.39% and 3.31%; Mode II: 2.61% and 3.22%; Mode III: 4.86% and 2.7%, respectively) were triggered in three modes, compared with the DNA vaccine alone (1.24% and 0.83%) (Fig. 2A). The total strength of the polyfunctional T cell response (the sum of percentage of T cells secreting 3-, 4- and 5-cytokines) was improved markedly in three modes ($P < 0.01$; Fig. 2B). However, the total strength of the polyfunctional T cell response among three modes had no significant difference in statistics (Fig. 2B).

In addition, peptidic nanofibers increased the numbers of combination ways of 3- and 4-cytokines, which respectively should be ten ($C_5^3 = 10$) and five ($C_5^4 = 5$) kinds of combination ways. *i.e.*, four types of 4-cytokine combinations and seven types of 3-cytokine combinations were triggered in three modes, which respectively included (CD107a, IFN- γ , IL-2, IL-4), (CD107a, IFN- γ , IL-4, TNF- α), (CD107a, IL-2, IL-4, TNF- α), (CD107a, IFN- γ , IL-2, TNF- α) (Fig. S2, ESI †), as well as (CD107a, IFN- γ , IL-4), (CD107a, IL-2, TNF- α), (CD107a, IL-4, TNF- α), (IFN- γ , IL-2, IL-4), (IFN- γ , IL-4, TNF- α), (CD107a, IFN- γ , TNF- α), (CD107a, IL-2, IL-4) (Fig. S3, ESI †). The HIV DNA vaccine alone triggered only two combinations of 4-cytokines (Fig. S2, ESI †) and four combinations of 3-cytokines, respectively (Fig. S3, ESI †). All together, these results indicated that peptidic nanofibers can improve the HIV-specific polyfunctional T cell response. More importantly, *in situ* formed nanofibers had the same effect as that injected with exogenous ALP on immunoregulating the polyfunctional T cell response. It thus highlighted the advantages of such

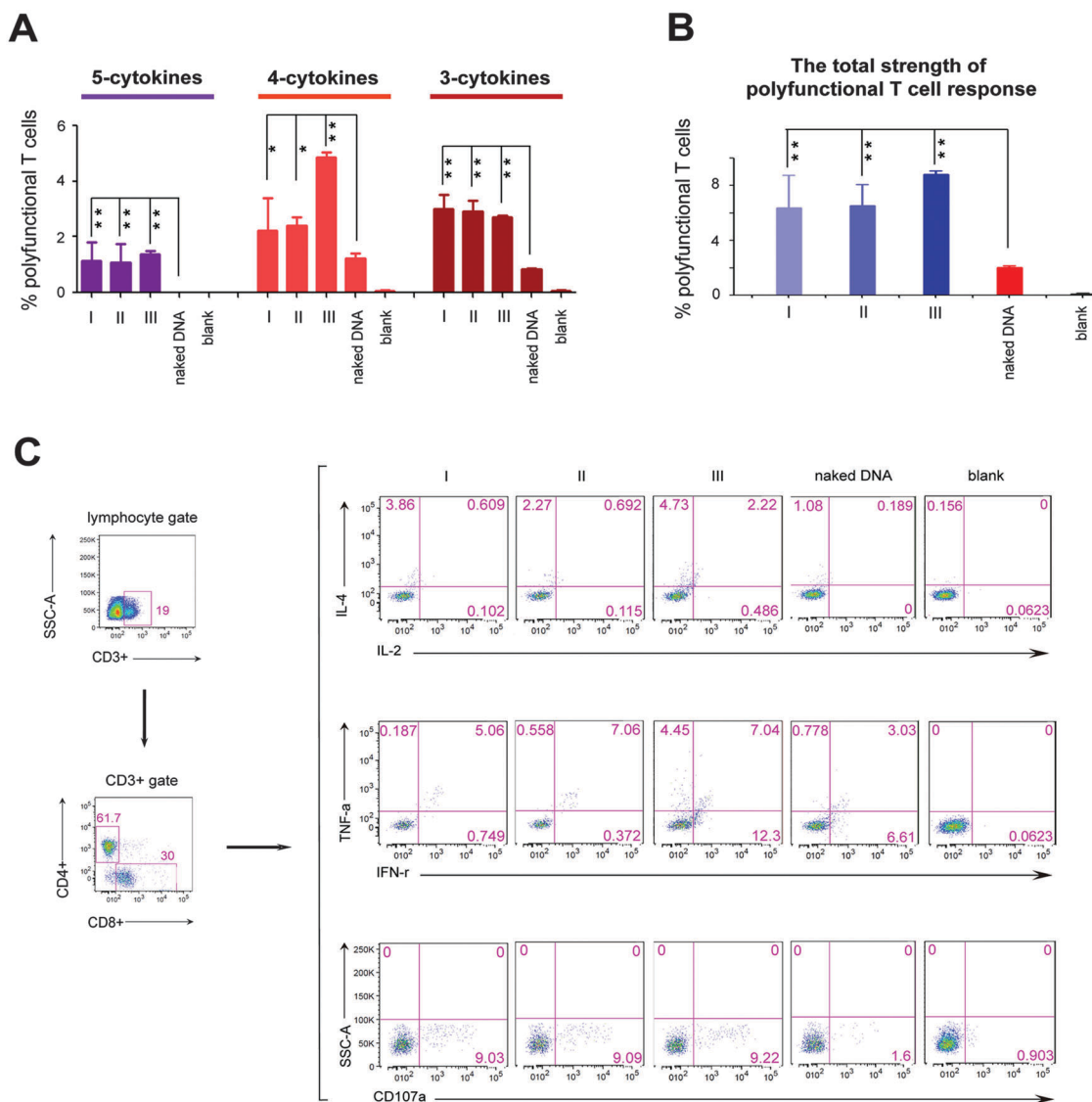


Fig. 2 Polyfunctional T cell response induced by the HIV DNA vaccine aided by peptidic nanofibers in three modes, compared with the HIV DNA vaccine alone. (A) The percentage of T cells which simultaneously secrete 5-cytokines (IL-2, IL-4, TNF- α , IFN- γ and CD107a), 4-cytokines (four cytokines among IL-2, IL-4, TNF- α , IFN- γ and CD107a), and 3-cytokines (three cytokines among IL-2, IL-4, TNF- α , IFN- γ and CD107a). (B) The total strength of the polyfunctional T cell response (the sum of proportion of T cell secreting 5-, 4- and 3-cytokines). (C) The flow cytometric analysis for four intercellular cytokines (IL-2, IL-4, TNF- α , IFN- γ) and one surface molecule (CD107a) of T cells in different mice groups. Mice were intradermally vaccinated with the DNA vaccine alone (naked DNA) and the DNA vaccine aided by peptidic nanofibers (three modes respectively). 'naked DNA': mice vaccinated with the HIV DNA vaccine alone. 'blank': mice vaccinated with an empty DNA vector aided by peptidic nanofibers. Symbols ** and * respectively mean P value < 0.01 and < 0.05.

nanofibers for practical applications on improving the polyfunctional T cell response by the HIV DNA vaccine.

Such superior function of peptidic nanofibers on optimizing the polyfunctional T cell response also implies that they can serve as candidates to substitute existing ways for enhancing vaccine-specific polyfunctionality, *e.g.*, some virus-derived vaccine vectors (vaccinia or yellow fever virus).^{22,23} Virus-derived vectors usually contain many undefined constituents that can be expressed *in vivo* and might incorrectly influence immune responses by some unknown and uncontrollable ways. For example, in the Merck phase II trials of the HIV vaccine, some unknown and *in vivo* expressed elements located in an adenoviral vector might

be the factors that caused a higher infection ratio for vaccinated individuals.²⁴ However, the components of peptidic nanofibers in this study have been identified clearly. Its safety is very good *in vitro* and *in vivo*.²⁰ It suggests that such nanofibers might be a better option for inducing the polyfunctional T cell response, compared with the virus-derived vaccine vectors.

Moreover, the result that our *in situ* formed peptidic nanofibers increased IL-4 expression encouraged us to investigate the probable effect on regulating the maturation of B cells, since IL-4 is a crucial factor on promoting the maturation of B cells.²⁵ We herein evaluated the maturation of B cells in four mice groups, which were respectively vaccinated with the HIV

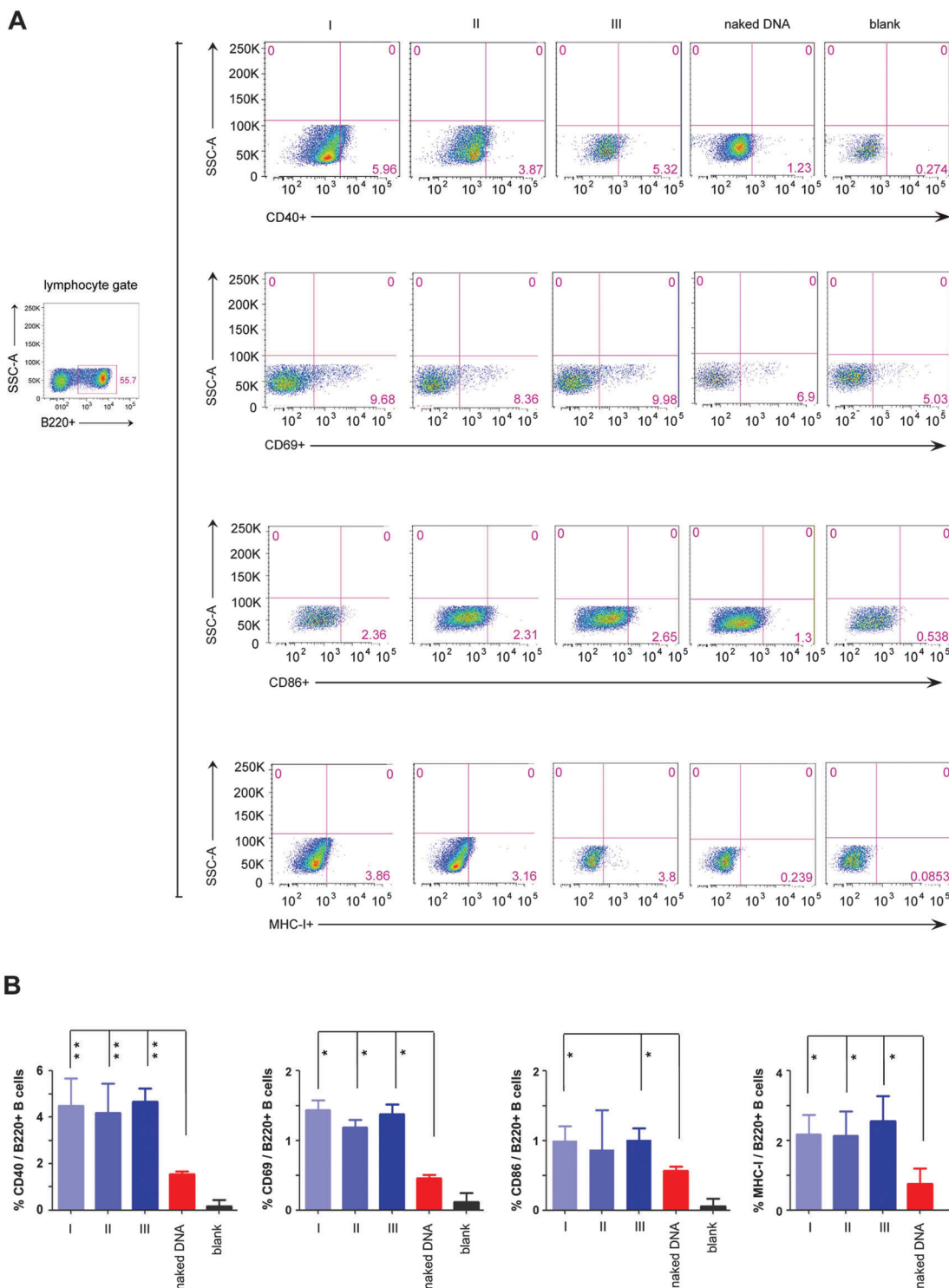


Fig. 3 The maturation of B cells stimulated by peptidic nanofibers. Flow cytometric analysis for the expressions of four surface markers indicating B cell maturation, including CD40, CD69, CD86 and MHC I. These cells were gated from B220⁺ (one specific mark of B cells) lymphocytes derived from the mice spleen. (A) Flow cytometric analysis for above surface molecules in different mice groups. (B) Histograms (the percentage of B220⁺ B cells which respectively expressed CD40, CD69, CD86 and MHC I). 'naked DNA', 'I', 'II', 'III' and 'blank' are the same as Fig. 2.

DNA vaccine alone and the HIV DNA vaccine aided by nanofibers *via* above three modes. An empty vaccine vector was used as a blank control. Three kinds of molecules on the B cell surface (CD40, CD69, CD86) were selected as markers for evaluating the

maturation of B cells.^{26,27} The ratios of B cells expressing the above three molecules increased significantly in all three mice modes vaccinated with the HIV DNA vaccine aided by peptidic nanofibers, compared with those vaccinated with the HIV DNA

vaccine alone (Fig. 3). Furthermore, the expression of the MHC molecule (the carrier of antigen epitope in cytoplasm) on the cell surface was significantly enhanced by peptidic nanofibers in three modes (Fig. 3). These results provided evidences for our above hypothesis that *in situ* formed peptidic nanofibers can promote B cell maturation, the same as that formed by adding exogenous ALP.

Next, we analyzed the effect of *in situ* formed peptidic nanofibers on the regulation of the humoral response *via* investigating antibody titers in the serum, because the facilitation of B cell maturation is consistent with the enhancement of the antibody response (B cells are in charge of secreting antibodies²⁸). Compared with DNA alone, the DNA vaccine aided by peptidic nanofibers in all three modes induced significantly higher titers of IgM and IgG (Fig. 4A and B). We analyzed the influence of peptidic nanofibers on shaping the IgG subclass response, because the broad IgG subclass response is another important clinical feature for LTNP. ^{9–12} Herein, the positive IgG subclass response is defined as the OD value (difference value between 450 nm and 630 nm) being higher than 0.1 when serum samples are diluted at 1 : 100 (log₁₀ value is 2 at least). Nanofibers in all three modes triggered positive responses against all four IgG subclasses (log₁₀ values of IgG1, IgG2a, IgG2b, IgG3 were higher than 2), but DNA alone only

induced positive IgG2a response (Fig. 4C). Therefore, *in situ* formed nanofibers, as effective as that formed *via* adding exogenous ALP, broadened the IgG subclasses response. Note-worthily, the function of such nanofibers on enhancing the IgG1 response by the HIV vaccine might further improve the value of such nanofibers. IgG1 is a main form of the neutralizing antibody in blood. Multiple broad-neutralizing monoclonal antibodies, such as F105, 2G12 and 2F5, belong to the IgG1 subclass.²⁹ Moreover, high affinity of IgG1 with the Fc- γ receptor makes IgG1 become a crucial type of antibody on inhibiting HIV *via* antibody-dependent cell-mediated cytotoxicity (ADCC),^{11,30} which is a mechanism of immune defense where ADCC-effector cells (such as natural killer cells, macrophages or neutrophils) selectively attack pathogen-infected cells *via* the affinity-dependent recognition against antibodies binding on the surface of pathogen-infected cells (mediated by the Fc- γ receptor on the surface of ADCC-effector cells).³¹

To our excitement, the peptidic nanofibers in all three modes improved the V1/V2 loop-specific IgG response (V1/V2-specific IgG2a, IgG2b and IgG3 responses; Fig. 5A and B), which was positively correlated with protection efficacy in an updated phase III HIV vaccine trial in Thailand.¹³ Such crucial immunity failed to be triggered by the HIV DNA vaccine alone (Fig. 5A). Recently, a NIH-led large-scale clinical trial (HVTN 505) was stopped,

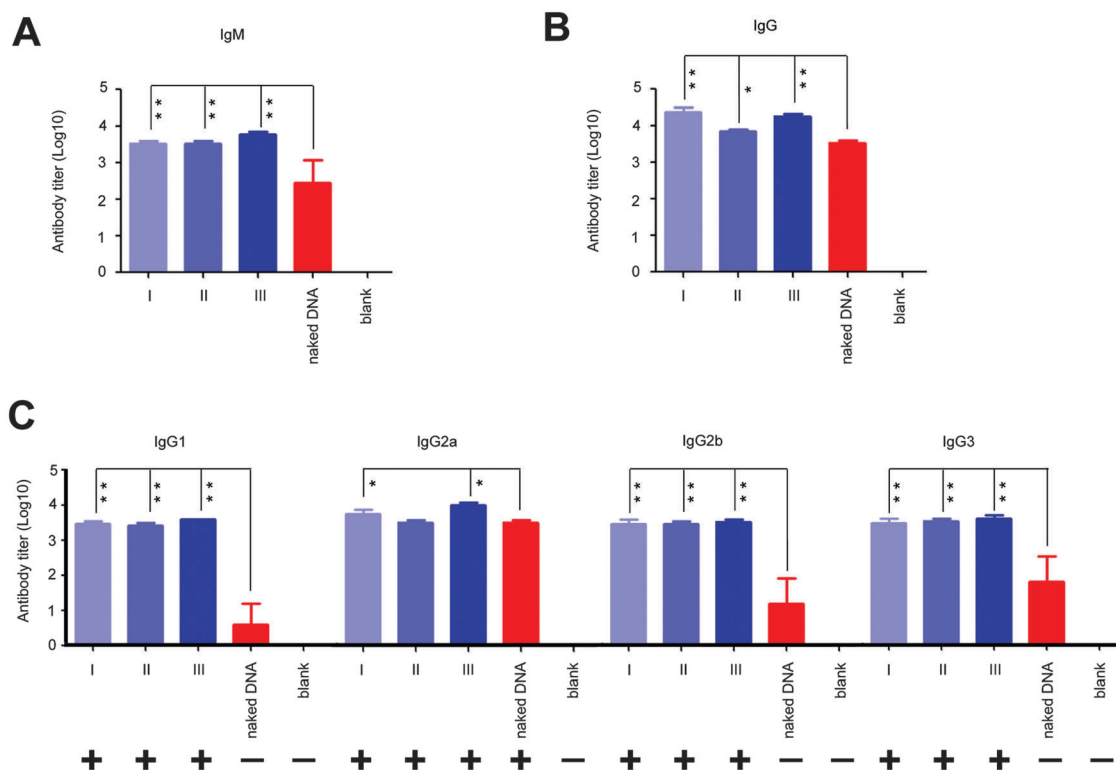


Fig. 4 The titers of IgM, IgG, and four IgG subclasses (IgG1, IgG2a, IgG2b, IgG3) against HIV Env gp145. Serum samples from vaccinated mice were harvested two weeks after the final immunization. Each mouse serum was evaluated by the standard ELISA assay (each serum sample was repeatedly detected once). The results are the mean value (log₁₀ value) of each group of mice \pm SD. (A–C) respectively indicate the titers of IgM, IgG, and four IgG subclasses (IgG1, IgG2a, IgG2b, IgG3) against the whole HIV Env gp145 in mice. '+' and '-' in part (C) respectively indicate positive and negative antibody responses. The positive value is determined: (1) OD value (difference value between 450 nm and 630 nm) > 0.1 at the dilution of 1 : 100, (2) at least 2.1 folds that of the negative control.

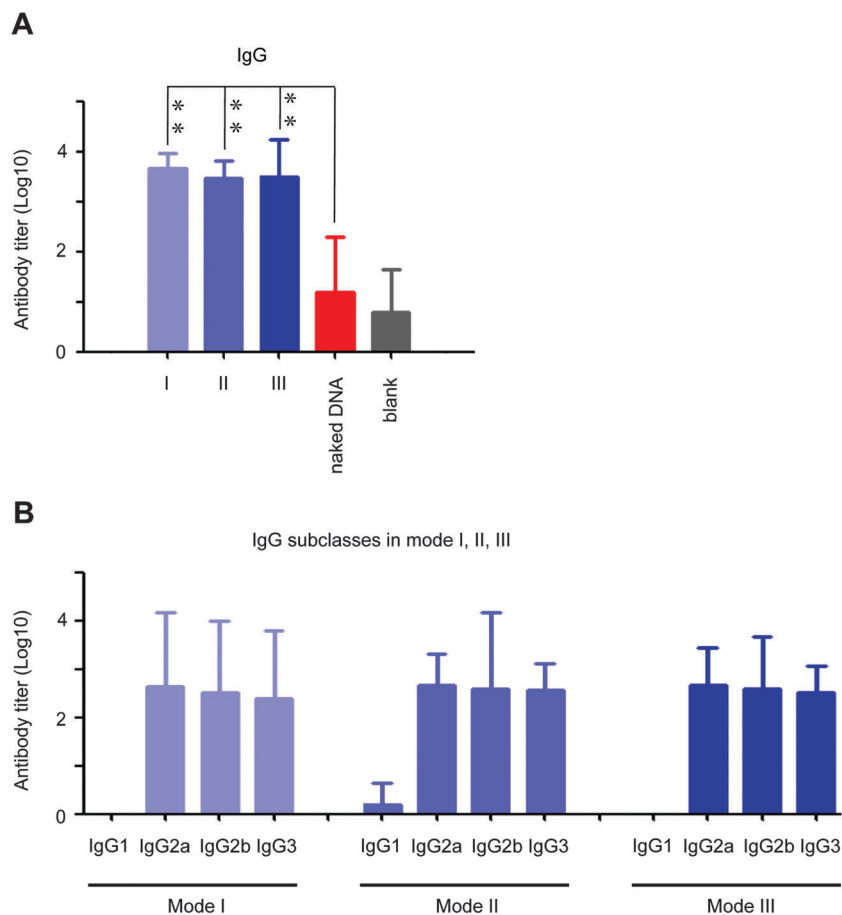


Fig. 5 IgG response against the HIV Env V1/V2 loop. The HIV Env V1/V2 loop-specific IgG response was detected by ELISA. Mice were vaccinated with 'blank', 'naked DNA', and DNA aided by peptidic nanofibers, respectively (three modes: I, II and III). The positive response is defined as that the log₁₀ value is higher than 2. (A) Significantly stronger IgG response against the HIV Env V1/V2 loop (positive response) was found in mice vaccinated with the HIV DNA vaccine aided by nanofibers in Mode I, II, and III, compared with that vaccinated with the HIV DNA vaccine alone (negative response). (B) For each IgG subclass response in Mode I, II, and III, positive IgG2a, IgG2b, IgG3 responses were triggered in mice vaccinated with the HIV DNA vaccine aided by nanofibers, but the IgG1 response was negative.

because the HIV vaccine regimen did not prevent HIV infection. One of major reasons might be that the crucial immune response, the V1/V2-specific IgG response, was not triggered.³² Considering our peptidic nanofibers can effectively optimize such immune response induced by the HIV vaccine, it thus might fundamentally optimize immunities of HIV vaccine regimens like HVTN 505. If such a clinical trial had incorporated the peptidic nanofibers, we would have predicted a better outcome.

Finally, we investigated whether NMe can still improve immunities by the HIV DNA vaccine when they are separately vaccinated. There is no binding between the HIV DNA vaccine and NMe in the separated vaccination. Such investigation is an important supplement for Mode I (injecting the HIV DNA vaccine and NMe together), and will further reveal the importance of the binding between the HIV DNA vaccine and NMe for enhancing immunities by the vaccine. One mouse group was injected with NMe precursors first, followed by the HIV DNA vaccine (the time interval is one hour). In contrast, another mouse group was injected with the HIV DNA vaccine first, followed by NMe precursors one hour later. Comparing with

the mouse group vaccinated with the HIV DNA vaccine alone, neither the 'HIV DNA vaccine first, followed by NMe' nor 'NMe first, followed by the HIV DNA vaccine' significantly improve the immunities, including the polyfunctional T cell response, IgM, IgG, IgG subclass and V1/V2 loop-specific antibody responses (Fig. S4, ESI†). It suggested that a binding between NMe precursors and the HIV DNA vaccine is necessary to improve such immunities by the HIV DNA vaccine.

Conclusion

In summary, *in situ* formed peptidic nanofibers can improve the crucial immunities of the HIV DNA vaccine, including the polyfunctional T cell response, the broad IgG subclass response, and the V1/V2-specific IgG response, all of which can effectively delay the disease process or improve the protection from the infection of HIV. Such novel *in situ* formation (triggering the formation of peptidic nanofibers without exogenous ALP) thus highlights the robustness of the nanofibers as vaccine adjuvants

for practical applications, since we can simplify the operation procedure when injecting the complex of nanofibers and the HIV DNA vaccine into the host body (without premixing with ALP). Also, such unique features will facilitate the peptidic nanofibers to be crucial components not only in next-generation HIV vaccines, but also for other infectious diseases and cancer immunotherapy vaccines, such as the yellow fever vaccine, and the lung cancer immunotherapy vaccine, considering that the importance of immunities improved by peptidic nanofibers on inhibiting the development of these diseases.

Acknowledgements

The authors wish to acknowledge Prof. Bing Xu, Prof. Yuan Gao for their assistances in experiments and valuable suggestions in this study, and the financial supports from the Ministry of Science and Technology (2011CB933201, 2012AA022703, 2012AA030308), the Ministry of Health (2012ZX10001-008), the National Science Foundation of China (21025520, 51373043, 21105018 & 81361140345), the Chinese Academy of Science (XDA09030305) and Research Supported by the CAS/SAFEA International Partnership Program for Creative Research Teams.

References

- 1 D. H. Barouch and S. G. Deeks, *Science*, 2014, **345**, 169–174.
- 2 J. McMichael, P. Borrow, G. D. Tomaras, N. Goonetilleke and B. F. Haynes, *Nat. Rev. Immunol.*, 2010, **10**, 11–23.
- 3 P. Matthieu, L. Yves and P. Giuseppe, *Curr. Opin. HIV AIDS*, 2013, **8**, 333–340.
- 4 S. W. Barnett, S. Lu, I. Srivastava, S. Cherpelis, A. Gettie, J. Blanchard, S. Wang, I. Mboudjeka, L. Leung, Y. Lian, A. Fong, C. Buckner, A. Ly, S. Hilt, J. Ulmer, C. T. Wild, J. R. Mascola and L. Stamatatos, *J. Virol.*, 2001, **75**, 5526–5540.
- 5 V. Velu, K. Titanji, B. Zhu, S. Husain, A. Pladevega, L. Lai, T. H. Vanderford, L. Chennareddi, G. Silvestri, G. J. Freeman, R. Ahmed and R. R. Amara, *Nature*, 2009, **458**, 206–210.
- 6 M. R. Betts, M. C. Nason, S. M. West, S. C. De Rosa, S. A. Migueles, J. Abraham, M. M. Lederman, J. M. Benito, P. A. Goepfert, M. Connors, M. Roederer and R. A. Koup, *Blood*, 2006, **107**, 4781–4789.
- 7 J. Liu, K. L. O'Brien, D. M. Lynch, N. L. Simmons, A. La Porte, A. M. Riggs, P. Abbink, R. T. Coffey, L. E. Grandpre, M. S. Seaman, G. Landucci, D. N. Forthal, D. C. Montefiori, A. Carville, K. G. Mansfield, M. J. Havenga, M. G. Pau, J. Goudsmit and D. H. Barouch, *Nature*, 2009, **457**, 87–91.
- 8 J. R. Almeida, D. A. Price, L. Papagno, Z. A. Arkoub, D. Sauce, E. Bornstein, T. E. Asher, A. Samri, A. Schnuriger, I. Theodorou, D. Costagliola, C. Rouzioux, H. Agut, A. G. Marcelin, D. Douek, B. Autran and V. Appay, *J. Exp. Med.*, 2007, **204**, 2473–2485.
- 9 K. Banerjee, P. J. Klasse, R. W. Sanders, F. Pereyra, E. Michael, M. Lu, B. D. Walker and J. P. Moore, *AIDS Res. Hum. Retroviruses*, 2010, **26**, 445–458.
- 10 G. D. Tomaras and B. F. Haynes, *Curr. Opin. HIV AIDS*, 2009, **4**, 373–379.
- 11 A. W. Chung, M. Ghebremichael, H. Robinson, E. Brown, I. Choi, S. Lane, A. S. Dugast, M. K. Schoen, M. Rolland, T. J. Suscovich, A. E. Mahan, L. Liao, H. Streeck, C. Andrews, S. Rerks-Ngarm, S. Nitayaphan, M. S. de Souza, J. Kaewkungwal, P. Pitisuttithum, D. Francis, N. L. Michael, J. H. Kim, C. Bailey-Kellogg, M. E. Ackerman and G. Alter, *Sci. Transl. Med.*, 2014, **6**, 228ra38.
- 12 V. A. Sundqvist, A. Linde, R. Kurth, A. Werner, E. B. Helm, M. Popovic, R. C. Gallo and B. Wahren, *J. Infect. Dis.*, 1986, **153**, 970–973.
- 13 B. F. Haynes, P. B. Gilbert, M. Juliana McElrath, S. Zolla-Pazner, G. D. Tomaras, S. M. Alam, D. T. Evans, D. C. Montefiori, C. Karnasuta, R. Sutthent, H. X. Liao, A. L. DeVico, G. K. Lewis, C. Williams, A. Pinter, Y. Fong, H. Janes, A. DeCamp, Y. Huang, M. Rao, E. Billings, N. Karasavvas, M. L. Robb, V. Ngauy, M. S. de Souza, R. Paris, G. Ferrari, R. T. Bailer, K. A. Soderberg, C. Andrews, P. W. Berman, N. Frahm, S. C. De Rosa, M. D. Alpert, N. L. Yates, X. Shen, R. A. Koup, P. Pitisuttithum, J. Kaewkungwal and S. Nitayaphan, *N. Engl. J. Med.*, 2012, **366**, 1275–1286.
- 14 Z. Zhao, Z. Zhou, J. Bao, Z. Wang, J. Hu, X. Chi, K. Ni, R. Wang, X. Chen, Z. Chen and J. Gao, *Nat. Commun.*, 2013, **4**, 2266.
- 15 N. Gu and H. Xu, *Sci. Rep.*, 2013, **3**, 2655.
- 16 Y. Chen and G. Liang, *Theranostics*, 2012, **2**, 139–147S.
- 17 P. Huang, Y. Gao, J. Lin, H. Hu, H. Liao, X. Yan, Y. Tang, A. Jin, J. Song, G. Niu, G. Zhang, F. Horkay and X. Chen, *ACS Nano*, 2015, **9**, 9517–9527.
- 18 Y. Gao, J. Shi, D. Yuan and B. Xu, *Nat. Commun.*, 2012, **3**, 1033.
- 19 Y. Kuang, J. Shi, J. Li, D. Yuan, K. A. Alberti, Q. Xu and B. Xu, *Angew. Chem., Int. Ed.*, 2014, **53**, 8104–8107.
- 20 Y. Tian, H. Wang, Y. Liu, L. Mao, W. Chen, Z. Zhu, W. Liu, W. Zheng, Y. Zhao, D. Kong, Z. Yang, W. Zhang, Y. Shao and X. Jiang, *Nano Lett.*, 2014, **14**, 1439–1445.
- 21 U. Müller, D. Pehler, W. Stenzel, G. Köhler, O. Frey, J. Held, A. Grahner, T. Richter, M. Eschke, T. Kamradt, F. Brombacher and G. Alber, *Mucosal Immunol.*, 2012, **5**, 299–310.
- 22 M. L. Precopio, M. R. Betts, J. Parrino, D. A. Price, E. Gostick, D. R. Ambrozak, T. E. Asher, D. C. Douek, A. Harari, G. Pantaleo, R. Bailer, B. S. Graham, M. Roederer and R. A. Koup, *J. Exp. Med.*, 2007, **204**, 1405–1416.
- 23 D. Gaucher, R. Therrien, N. Kettaf, B. R. Angermann, G. Boucher, A. Filali-Mouhim, J. M. Moser, R. S. Mehta, D. R. Drake III, E. Castro, R. Akondy, A. Rinfret, B. Yassine-Diab, E. A. Said, Y. Chouikh, M. J. Cameron, R. Clum, D. Kelvin, R. Somogyi, L. D. Greller, R. S. Balderas, P. Wilkinson, G. Pantaleo, J. Tartaglia, E. K. Haddad and R. Sékaly, *J. Exp. Med.*, 2008, **205**, 3119–3131.
- 24 M. J. McElrath, S. C. De Rosa, Z. Moodie, S. Dubey, L. Kierstead, H. Janes, O. D. Defawe, D. K. Carter, J. Hural, R. Akondy, S. P. Buchbinder, M. N. Robertson, D. V. Mehrotra, S. G. Self, L. Corey, J. W. Shiver, D. R. Casimiro and the Step Study Protocol Team, *Lancet*, 2008, **372**, 1894–1905.

- 25 B. Gao and T. L. Rothstein, *J. Immunol.*, 2013, **191**, 670–677.
- 26 K. Attridge, R. Kenefeck, L. Wardzinski, O. S. Qureshi, C. J. Wang, C. Manzotti, K. Okkenhaug and L. S. K. Walker, *J. Immunol.*, 2014, **192**, 2195–2201.
- 27 P. J. Nadeau, A. Roy, C. Gervais-St-Amour, M. È. Marcotte, N. Dussault and S. Néron, *Mol. Immunol.*, 2012, **49**, 582–592.
- 28 Y. Bao and X. Cao, *J. Autoimmun.*, 2014, **55**, 10–23.
- 29 T. W. Baba, V. Liska, R. Hofmann-Lehmann, J. Vlasak, W. Xu, S. Ayehunie, L. A. Cavacini, M. R. Posner, H. Katinger, G. Stiegler, B. J. Bernacky, T. A. Rizvi, R. Schmidt, L. R. Hill, M. E. Keeling, Y. Lu, J. E. Wright, T. C. Chou and R. M. Ruprecht, *Nat. Med.*, 2000, **6**, 200–206.
- 30 P. Bruhns, B. Iannascoli, P. England, D. A. Mancardi, N. Fernandez, S. Jorieux and M. Daëron, *Blood*, 2009, **113**, 3716–3725.
- 31 D. Rudnicka, A. Oszmiana, D. K. Finch, I. Strickland, D. J. Schofield, D. C. Lowe, M. A. Sleeman, D. M. Davis and HVTN 505 Study Team, *Blood*, 2013, **121**, 4694–4702.
- 32 S. M. Hammer, M. E. Sobieszczyk, H. Janes, S. T. Karuna, M. J. Mulligan, D. Grove, B. A. Koblin, S. P. Buchbinder, M. C. Keefer, G. D. Tomaras, N. Frahm, J. Hural, C. Anude, B. S. Graham, M. E. Enama, E. Adams, E. DeJesus, R. M. Novak, I. Frank, C. Bentley, S. Ramirez, R. Fu, R. A. Koup, J. R. Mascola, G. J. Nabel, D. C. Montefiori, J. Kublin, M. J. McElrath, L. Corey and P. B. Gilbert, *N. Engl. J. Med.*, 2013, **369**, 2083–2092.