Designer DNA architecture offers precise and multivalent spatial pattern-recognition for viral sensing and inhibition

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DNA, when folded into nanostructures with a specific shape, is capable of spacing and arranging binding sites into a complex geometric pattern with nanometre precision. Here we demonstrate a designer DNA nanostructure that can act as a template to display multiple binding motifs with precise spatial pattern-recognition properties, and that this approach can confer exceptional sensing and potential viral inhibitory capabilities. A star-shaped DNA architecture, carrying five molecular beacon-like motifs, was constructed to display ten dengue envelope protein domain III (ED3)-targeting aptamers into a two-dimensional pattern precisely matching the spatial arrangement of ED3 clusters on the dengue (DENV) viral surface. The resulting multivalent interactions provide high DENV-binding avidity. We show that this structure is a potent viral inhibitor and that it can act as a sensor by including a fluorescent output to report binding. Our molecular-platform design strategy could be adapted to detect and combat other disease-causing pathogens by generating the requisite ligand patterns on customized DNA nanoarchitectures.

Infectious diseases, such as viruses, bacteria and toxins, present unique spatial patterns of antigens on their surfaces1–3. Existing weakly binding ligands (called ‘binders’ herein) that interact with these epitopes can be linked to a scaffold to improve multivalent binding avidity4–6. However, some viruses, such as dengue (DENV), present complex geometric patterns that cannot be addressed by existing scaffolds because they are not as precise in ligand spacing or provide limited control over the scaffold shape and ligand valency7–12. Furthermore, previous synthetic scaffolds (polymers, dendrimers, nanofibres, nanoparticles and lipid nanoemulsions) have shown some toxicity. Designer DNA nanostructures13–17 are dendrimers, nanofibres, nanoparticles and lipid nanoemulsions) eventually removed by the liver21 and kidney28,29. These nanostructures have exhibited blood circulation times of up to 24 h (ref. 21) but are eventually removed by the liver21 and kidney28,29.

We demonstrate our strategy by directly targeting DENV through a star-shaped DNA scaffold. This architecture was strategically designed to multivalently arrange binders in a 2D pattern mirroring the complex spatial distribution of dengue epitopes with nanometre precision. Although previous studies have shown that optimizing bivalent interactions is important in protein binding18–20, cell signalling21 and antibody spatial tolerance using DNA nanostructures22, to our knowledge the importance of multivalent, 2D spatial pattern-recognition properties, and that this approach can confer exceptional sensing and potential viral inhibitory capabilities. A star-shaped DNA architecture, carrying five molecular beacon-like motifs, was constructed to display ten dengue envelope protein domain III (ED3)-targeting aptamers into a two-dimensional pattern precisely matching the spatial arrangement of ED3 clusters on the dengue (DENV) viral surface. The resulting multivalent interactions provide high DENV-binding avidity. We show that this structure is a potent viral inhibitor and that it can act as a sensor by including a fluorescent output to report binding. Our molecular-platform design strategy could be adapted to detect and combat other disease-causing pathogens by generating the requisite ligand patterns on customized DNA nanoarchitectures.

Results

Design and characterization of the DNA star. The envelope protein domain III (ED3) viral epitope on the DENV surface is organized into a complex polyhedral pattern23 with alternating clusters of trivalent or pentavalent ED3 sites. By connecting the clusters of ED3 sites linearly, we determined that a star shape, consisting of an interior pentagon connected to five exterior triangles, would provide an optimal multivalent scaffold (Fig. 1a). The viral diameter of

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DENV is 50 nm (ref. 39) and vector distances were measured using PyMOL. The distance between adjacent trivalent–trivalent clusters was 15.3 nm and 14.3 nm, respectively. The DNA star scaffold is designed to match the pattern and spacing of ED3 clusters. Each DNA strand is represented by an arrow going from the 5′ end to the 3′ end. Non-hybridizing polyT regions are for angle formation between the edges. A 3D representation of the scaffold is also shown. A schematic of the partial or complete star complexes (left) and characterization of the DNA star structure using non-denaturing PAGE (right). Yields of the 1-triangle, 2-triangle, 3-triangle, 4-triangle, star and unclosed star complexes are 100%, 98.2%, 81.2%, 86.3%, 88.1% and 98.3%, respectively. Experiments were repeated independently three times, with similar results.

Single-stranded regions of the scaffold strands form stem-loops. AFM confirmed the formation. Experiments were repeated independently three times, with similar results.

The formation of the complete DNA star (lane ‘Star’, Fig. 1c) was first characterized by 4% non-denaturing polyacrylamide gel electrophoresis (PAGE), with one of each unique component strand and partial star complex included for reference. Atomic force microscopy (AFM) imaging further confirmed formation of the star (Fig. 1d). For individual DNA nanostructure assemblies, non-denaturing gel electrophoresis is a valid and more reliable method for characterizing structural formation and yield (for example, see previous studies41–43). Our non-denaturing PAGE showed a distinct, major band corresponding to each of the 2D architectures, confirming that DNA star structures were formed with high efficiency (Fig. 1c). It is also worth noting that the AFM probe can damage or displace the thin DNA tile-based structures during the scan (Supplementary Fig. 1). Additionally, AFM may display the overall
shape of DNA star but cannot fully resolve the details of the structure at the atomic level due to the limited resolution.

**DNA star-based DENV sensing.** The DNA star was functionalized for DENV detection and inhibition by hybridizing a well-characterized, ED3-binding aptamer\(^\text{11}\) at each of the 10 vertices of the DNA star to form a star–aptamer complex that geometrically matched and targeted ED3 clusters (Fig. 2a and Supplementary Fig. 2). The aptamer exhibited weak binding to DENV, as determined by surface plasmon resonance (Supplementary Fig. 3). However, the binding strength increased as aptamers were placed onto scaffolds that increasingly matched the pattern of ED3 sites on DENV with the DNA star showing the greatest binding avidity (Supplementary Fig. 3). 6-FAM (fluorophore) and BHQ-1 (quencher) carrying strands were hybridized to each inner edge, flanking the hairpin to turn the DNA star–aptamer complex into a viral sensor. In the absence of DENV, FAM and BHQ-1 molecules would be brought together by canonical Watson–Crick base pairing in the stem, much like a molecular beacon\(^\text{12}\). However, unlike a molecular beacon, which gives a fluorescent readout of target nucleic acid hybridization events, here the hairpins are pulled apart and converted to ssDNA as a result of flexible linear DNA scaffold (abbreviated as linear). In both scaffold–aptamer complexes, the strand(s) used to create the DNA star were reused by a flexible linear DNA scaffold (abbreviated as linear). In both scaffold–aptamer complexes, the strand(s) used to create the DNA star were reused.

The DNA star–aptamer sensor was able to directly detect DENV virions with high sensitivity, affording a limit of detection (LoD) of \(1 \times 10^3\) p.f.u. ml\(^{-1}\) and \(1 \times 10^5\) p.f.u. ml\(^{-1}\), respectively, in human serum and plasma (Fig. 2c). To exclude the possibility that the series of DENV-mixed DNA star sensor signals obtained within 1–2 min resulted from an automatic (non-target-triggered) separation of the fluorophores from quenchers, we measured fluorescent readouts over time at different temperatures. Fluorescent measurements over time showed that the FAM fluorophores remained quenched during the duration required for sensing and functioned at different temperatures (Supplementary Fig. 4). To examine the specificity of our sensor, we tested it against another virus available to us, the adenovirus. If the star–aptamer complex does not recognize the virus through proper binding, the complex will remain quenched (Fig. 2b).

The experimental results verified that the star–aptamer complex designed for DENV detection is not able to sense the adenovirus, even at high concentrations (Fig. 2d).

To make a comparison between the DNA star-based and gold standard methods under similar conditions, we carried out RT–qPCR and ELISA assays to respectively detect DENV RNA and NS1 antigen (Supplementary Fig. 5). The more sensitive method, RT–qPCR, showed a LoD of \(1 \times 10^3\) p.f.u. ml\(^{-1}\) while the heptagon could not detect the virus even at a DENV serum sample concentration of \(1 \times 10^5\) p.f.u. ml\(^{-1}\) when standardized with the same fluorophore–quencher pair concentrations.

The positive sensing provided by the heptagon was expected, because when we analysed DENV surface ED3 patterns, we also recognized a hexagon-shaped distribution of ED3 clusters when centring the DENV virion on a trivalent cluster (Fig. 3b); this additional pattern differed from the pentavalent cluster-centred star shape (Fig. 1a). The fact that the hexagon can serve as a good positive control sensor proves the need for pattern matching in detection. The sensitivity of the hexagon sensor was tested, providing a LoD of \(1 \times 10^3\) p.f.u. ml\(^{-1}\) (Fig. 3d). Although the heptagon carried more aptamers than the hexagon, the sensor resulted in poor to no detection. This comparison is particularly interesting as it demonstrates that a small mismatch of 2D geometry, under the same bivalently correct spacing and with higher ligand valency and density (as compared to the hexagon), leads to dramatically reduced sensing capabilities.

**In vitro DENV inhibition.** Encouraged by the previous results, we investigated the in vitro inhibition of DENV through standard antiviral, plaque-forming half-maximum effective concentration (EC\(_{50}\)) assays. Briefly, the DENV viral particles were incubated with different concentrations of each inhibitor in human serum and the remaining infectivity was determined by a plaque reduction assay (Fig. 4a). The dose-dependent inhibition of DENV for the monovalent aptamer and each of the scaffold–aptamer complexes was examined (Fig. 4b). The EC\(_{50}\) value of the DNA star–aptamer complex for DENV infection inhibition was 2 nM, whereas the EC\(_{50}\) value for the monovalent aptamer was 15\(\mu\)M (Fig. 4c,d). These results demonstrate that the DNA star–aptamer multivalent inhibitor was 7.5 \(\times\) 10\(^5\)-fold more effective than the monovalent aptamer for the in vitro inhibition of DENV infection in human serum.

We also examined the EC\(_{50}\) values of the other control scaffolds, where a lower EC\(_{50}\) value demonstrates more potent inhibition (Fig. 4b–d). In correlation with our DENV sensing data, the hexagon scaffold provided the second most potent inhibitory effect, with an EC\(_{50}\) value of 10 nM, while the heptagon scaffold exhibited poor inhibition, with an EC\(_{50}\) value of 440 nM. The flexible linear scaffold provided better inhibition, at an EC\(_{50}\) of 90 nM, than the bivalent scaffold, at an EC\(_{50}\) of 130 nM. Additionally, as previous studies predicted, potentially matching the spacing could increase the inhibitory effect as all the structures proved to be more potent than the monovalent aptamer.
These results point out that stronger inhibition was achieved when the spatial pattern was properly matched. Interestingly, the 2D heptagon scaffold exhibited worse inhibitory potency than the 1D bivalent and flexible linear scaffolds. A mismatch of a fixed geometric pattern, while keeping the same initial spacing, leads to reduced inhibition potency when compared to flexibly matching...
Fig. 3 | Evaluation of control sensors. a, Schematic showing the design of bivalent, flexible, linear, hexagon-centred and heptagon-centred control sensors. AFM imaging verified formation of the 2D scaffolds. AFM experiments were repeated independently three times, with similar results. b, Schematic demonstrating that ED3 clusters show a hexagonal-shaped pattern when a trivalent ED3 cluster is centred. The pattern-matching hexagon scaffold allows aptamers to strongly bind the ED3 domains, enabling a fluorescent readout. c, At a DENV concentration of $1 \times 10^5$ p.f.u. ml$^{-1}$, the DNA star and hexagon sensors exhibit good detection ability. The linear sensor exhibits poor detection ability. Heptagon and bivalent sensors show little to no sensing. Data are presented as mean ± s.d., $n = 3$ biologically independent samples. A two-sided t-test was performed to test significance against the background ($^{***}P < 0.0001$). Individual data points below background are not shown but involved in error calculation. d, A comparison of the DNA star and hexagon sensors in DENV sensing capabilities. The hexagon sensor exhibits a LoD of $10^3$ p.f.u. ml$^{-1}$. Data are presented as mean ± s.d., $n = 3$ biologically independent samples. A two-sided t-test was performed to test significance against the background ($^{*}P < 0.05; ^{**}P < 0.005; ^{***}P < 0.0005$). Individual data points below background are not shown but were involved in error calculation.
the spacing. In summary, the more the binders match the 2D pattern of binding sites, the stronger the inhibition as aptamers bound to the star scaffold show superior performance compared to the other scaffold shapes.

Confocal imaging. Time-lapsed, live confocal imaging demonstrated that DENV virions lose their cell internalization ability after being bound by the DNA star–aptamer complex (Fig. 5). For the assay, the HepG2 line was chosen as it is derived from the liver, an organ DENV infects, and was cultured in medium containing human serum. Membrane-bound structures, including the cell membrane, were stained with Dil (yellow) while cell nuclei (blue) were stained with Hoechst. DENV virions were labelled with DiD (red) and the DNA star–aptamer complex was labelled with the 6-FAM fluorophore (green), quenched in the absence of DENV. When the green DNA stars and the red virions co-localize, the merged signals appear yellow. The Dil stain and 6-FAM dye also exhibit fluorescence spillover due to slightly overlapping excitation and emission spectra. The finer details between labelled structures can be resolved when viewing the merged image as separate fluorescence channels (Supplementary Fig. 8).

Unbound or DNA star-bound virions were introduced to the labelled cells (Fig. 5, Supplementary Fig. 9 and Supplementary Videos 1 and 2). In the unbound condition, DENV accumulated...
**Fig. 5 | Confocal imaging.**

**a.** DENV virions (red), membrane bound structures (yellow) and cell nuclei (blue) were labelled with DiD, Dil and Hoescht stains, respectively. Due to the combined effects of fluorescence spillover and the merged red and green signals, co-localized DNA star-bound virions appear yellow. Representative viral particles are indicated with white arrows. Top row: viral accumulation over time for the no-inhibitor treated condition. Bottom row: viral entry inhibition over time during DNA star treatment. Each panel is accompanied by a cross-section of the same image (along the dotted lines in the main images). An eye symbol orients the viewing direction for the cross-sections reconstructed from Z stacks, with images taken at different focal planes.

**b.** A volume reconstruction of a close-up confocal view shows unbound DENV particles (red spheres) accumulating in the cell (top panels) and a DNA star-bound DENV particle (green sphere) inhibited from cell entry (bottom panels). Each confocal experiment was performed in singlicate.
into the cells over time. In the DNA star-bound condition, DENV accumulation was drastically reduced and virions were generally electrostatically repelled from cells. Even if some DNA star-bound virions reached the cell membrane, they were unable to enter cells over the time course of an hour.

A cross-section (indicated by the white dotted line in Fig. 5a) accompanies each panel, with an eye symbol denoting the viewing direction. The view was reconstructed by images taken from different planes (Z stacks) during imaging. Volume reconstruction using Imaris software confirmed viral accumulation in the unbound condition and viral entry inhibition when bound to the DNA star (Fig. 5b).

**Discussion**

By targeting a challenging exemplar platform, the DENV flavivirus, we have demonstrated the importance of integrating a structurally defined DNA nanoarchitecture with precise, multivalent spatial pattern-recognizing properties. Sensors and inhibitors recognizing the DENV pattern identity allow them to work effectively. When aptamers are arranged into an optimal 2D shape, they can be induced to fit the correct DENV pattern. In contrast, linear complexes, such as the bivalent–aptamer complex, initially have neither optimal shape nor optimal spacing, so they have weak affinity to ED3 sites. Moreover, for those that bind transiently, they have trouble staying on because the hairpin region also has affinity to the base pair. Without the optimal shape identity to keep these hairpin interactions from forming, sensing and inhibitory abilities suffer.

We also observed that an incorrect shape will be more detrimental than having no shape at all, as observed with the heptagon scaffold. We speculate that the heptagon–aptamer complex can bind, in the best-case scenario, bivalently to two ED3 clusters, but that leaves certain sites unbound and capable of cell internalization (Supplementary Fig. 7). In addition, a bound heptagon–aptamer complex would sterically prevent more heptagon–aptamer complexes from binding while linear scaffolds do not experience a steric block to such a degree.

Our DNA star sensor shows superiority over current gold standard DENV detection methods. The reduced sensitivity of RT-qPCR can result from the low amount of starting material (one genome copy per viral particle), RNA extraction process and instability of the extracted RNA. The star sensor, on the other hand, has direct access to the unprocessed viral sample. Up to two stars can specifically bind to a single viral particle, translating to a 10-fluorophore label (five per star). Additionally, the immunoglobulin-M (IgM) ELISA or IgM rapid test cannot provide early viral sensing as antibody production in the body requires several days after initial infection. So, no gold standard method for DENV detection can achieve the same sensing capacity in terms of cost, ease, sensitivity and speed (see Supplementary Fig. 6, Supplementary Table 1, and Note for Supplementary Table 1 for a detailed comparison). As is
standard with viral infection screening, secondary confirmation of infection by some of the standard methods (for example, viral isolation or nucleic acid sensing) should still be employed after the initial precautionary screening in order to make a full documentation and evaluation of an epidemic outbreak.

The performance of the DNA star during in vitro DENV inhibition shows promising therapeutic potential. As is the case for various infectious diseases (for example, the malaria parasite, DENV, Zika and influenza A), interactions with negatively charged glycosaminoglycans (GAGs) are required for host cell invasion. Thus, the DNA architecture not only relies on specific multivalent interactions for binding DENV but also electrostatically traps and isolates virions from the host cell plasma membrane and GAGs via the negative charges of the scaffold. Moreover, other monovalent aptamer combinations have been used as antiviral agents for human viruses. Thus, our scaffold-based strategy can be used to arrange these same aptamers into arrangements that increase specific, on-target binding while reducing off-target binding based on pattern identity.

As binders specific to other binding sites can be evolved or synthesized, our strategy could be widely applicable, and such scaffolds could be used to efficiently detect and inhibit a wide range of pathogens (Fig. 6), modulate cell signalling by analysing the distribution of cellular receptors, or organize previous drugs into more potent configurations. Although DNA nanostructures have shown certain stability in vivo, the component DNA strands can be chemically modified to further improve its stability.

**Reporting Summary.** Further information on research design is available in the Nature Research Reporting Summary linked to this article.

**Data availability**

All data supporting the findings of this work are available within this paper (figures, videos and description) and its Supplementary Information. All other data are available from the corresponding authors upon request.

**Code availability**

The SEQUIN program used in this study is available from the corresponding authors upon reasonable request.

**Online content**

Any Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41557-019-0369-8.

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Author contributions
P.S.K., S.-J.K., J.S.D., R.J.L. and X.W. conceived the original 2D spatial pattern-matching strategy. P.S.K., S.-J.K., R.J.L., J.C. and X.W. conceived and designed the experiments. S.R. and J.C. designed the sequences of the DNA scaffolds and performed non-denaturing PAGE. S.R., M.X., D.Z., J.C., F. Zhou and N.C.S. performed the AFM experiments. P.S.K., S.R. and M.E.K. purified and folded the DNA. L.K. and L.D.K. prepared viruses, conducted RT–qPCR detection and performed the plaque forming assays. F. Zhang performed SPR analysis. P.S.K., S.-J.K. and D.K. performed confocal microscopy. P.S.K. performed all other experiments. K.F. analysed the aptamer binding pattern of DENV. P.S.K., S.-J.K. and X.W. performed data analysis. P.S.K., S.-J.K., R.J.L. and X.W. led the preparation of the manuscript with contributions from all authors. P.S.K., S.R., S.-J.K., M.E.K. and L.K. contributed equally to this work.

Competing interests
The authors declare no competing interests.

Additional information
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Data collection

- Gel image was collected by GeneSys Software for G: BOX F3 (Syngene); AFM images were collected by PeakForce QNM Software, ScanAsyst-HR accessory for Nanoscope Multimode AFM; SPR data were collected using BIAevaluation software for BIAcore 3000 SPR spectrometer; Confocal microscopy images were collected using embedded software for Zesis LSM 510 META confocal; Time-lapsed imaging assay was done using embedded software for Leica TCS SP8 STED microscope.

Data analysis

- Gel images were analyzed by GeneTools Software for G: BOX F3 (Syngene); AFM images were analyzed by PeakForce QNM Software, ScanAsyst-HR accessory for Nanoscope Multimode AFM; SPR data were automatically analysed using BIAevaluation software for BIAcore 3000 SPR spectrometer; Confocal images were analyzed with Imaris software (Ver. 9.2); GraphPad Prism (Ver. 6.01) was used to make the plots for EC50 assay and other statistical analyses; PyMOL (version 2.1.1) was used to analyze the dimensions and pattern distribution of a dengue surface epitope; Nanoengineer-1 (Ver. 1.1.1) was used to make DNA models; Adobe Illustrator (Ver. 16.0.0) was used to generate figures in the paper; ChemDraw (Ver. 16.0) was used to generate a part of Figure 2; Biorender (biorender.com) was used to generate parts of Figure 6.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:
- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

All data supporting the findings of this work are available within this paper (figures, movies, and description) and its supplementary information documents. All
other data are available from the corresponding author upon request.

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

- Life sciences
- Behavioural & social sciences
- Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size

The sample size was selected for the eight assay used were those typically used for the chemical/physical measurements described. With the exception of confocal spectroscopy all samples were analyzed in triplicate.

Data exclusions

No data were excluded. All data obtained in analyses are presented.

Replication

AFM, gel analysis, qPCR, fluorimetry, SPR and ELISA were analyzed in triplicate. Plaque assay was performed in biological triplicate with singlicate analysis. Confocal was performed in singlicate.

Randomization

The eight analytical methods (AFM, confocal, gel analysis, plaque assay, qPCR, fluorimetry, SPR and ELISA) used in this study were chemical/physical measurements and unlike in vivo assays do not routinely use randomization. The necessary positive and negative controls were used in all the experiments where appropriate.

Blinding

Four of our analytical methods (AFM, confocal, qPCR, and fluorimetry) used in this study were chemical/physical measurements and unlike biological or in vivo assays do not routinely use blinded samples. The gel analysis, plaque assay, SPR and ELISA were essentially blinded as the samples were prepared in one lab and assayed/analyzed in a second lab.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

- n/a Involved in the study
- Antibodies
- Eukaryotic cell lines
- Palaeontology
- Animals and other organisms
- Human research participants
- Clinical data

Methods

- n/a Involved in the study
- ChIP-seq
- Flow cytometry
- MRI-based neuroimaging

Eukaryotic cell lines

Policy information about cell lines

Cell line source(s)

Vero and HepG2 cells were obtained from ATCC.

Authentication

All cell lines used in this study were obtained from ATCC. For all cell lines, ATCC uses morphology, karyotyping, and PCR based approaches to confirm the identity of human cell lines and to rule out both intra- and interspecies contamination. These include an assay to detect species specific variants of the cytochrome C oxidase I gene (COI) analysis to rule out inter-species contamination and short tandem repeat (STR) profiling to distinguish between individual human cell lines and rule out intra-species contamination.

Mycoplasma contamination

No. All cells are fully authenticated by ATCC and routinely tested for every 2 weeks.

Commonly misidentified lines

None.