

Strain-engineered artificial atom as a broad-spectrum solar energy funnel

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An optoelectronic material with a spatially varying bandgap that is tunable is highly desirable for use in photovoltaics, photocatalysis and photodetection. Elastic strain has the potential to be used to achieve rapid and reversible tuning of the bandgap. However, as a result of plasticity or fracture, conventional materials cannot sustain a high enough elastic strain to create sufficient changes in their physical properties. Recently, an emergent class of materials—named ‘ultrastrength materials’—have been shown to avoid inelastic relaxation up to a significant fraction of their ideal strength. Here, we illustrate theoretically and computationally that elastic strain is a viable agent for creating a continuously varying bandgap profile in an initially homogeneous, atomically thin membrane. We propose that a photovoltaic device made from a strain-engineered MoS₂ monolayer will capture a broad range of the solar spectrum and concentrate excitons or charge carriers.

The ability to continuously control the bandgap in optoelectronic materials, in a low-cost manner, is highly desirable for a wide range of energy and sensing applications, including photovoltaics^{1–3}, photocatalysis⁴ and photodetection⁵. For example, the efficiency of photovoltaic devices comprising a single p–n junction is subject to the thermodynamic constraint of the Shockley–Queisser limit¹, as only a limited portion of solar energy can be absorbed. Although tandem solar cells consisting of multi-junctions with cascaded bandgaps can achieve higher absorption efficiencies, the increased complexity of their fabrication and the high cost of materials are major drawbacks. It would therefore be a great advantage if one could fine-tune the bandgap within a single semiconducting material. Like chemical composition, elastic strain is a continuous variable that is capable of altering many physical and chemical properties. Although elastic strain engineering is a straightforward concept, its potential in photonics remains largely under-exploited. The key hurdle is the fact that conventional bulk-scale materials cannot sustain a high enough elastic strain to induce sufficient changes to the physical properties before they deform plastically or break.

Recently, materials belonging to an emergent class termed ‘ultrastrength materials’^{6–8} have been found to avoid inelastic relaxation up to a significant fraction of their ideal strengths; that is, materials of different size and dimensionality can exhibit dramatically distinct physical and chemical properties as well as mechanical strengths. Atomically thin membranes are a notable family of materials that exhibit the property of ultrastrength^{8,9}. The elastic strain of a graphene monolayer has been predicted theoretically and also measured experimentally to be as high as 25% (refs 8,10,11), while that of bulk graphite rarely survives beyond 0.1% strain. Recently, a monolayer of molybdenum disulphide (MoS₂) was experimentally exfoliated and characterized. It demonstrated an effective in-plane strain of up to 11% (ref. 12), a direct bandgap of 1.9 eV under strain-free conditions^{13,14} and a more than 10,000-fold enhancement of its luminescence quantum yield compared to its bulk counterpart¹³. The high elastic strain limit should make it possible to control the electronic and optical properties of two-dimensional

materials through simple elastic strain engineering, which is often infeasible in their three-dimensional bulk phases. In addition, the lattice symmetry of MoS₂ also demonstrates a unique valley-selective circular dichroism^{15,16}.

Here, we illustrate theoretically and computationally that elastic strain is a viable agent for creating a continuously varying bandgap profile in an initially homogeneous, atomically thin membrane. As a proof of concept, an ‘artificial atom’ made of a nanoindented MoS₂ monolayer was investigated and predicted to be able to absorb a broad range of the solar spectrum from 2.0 eV to 1.1 eV as the biaxial strain increases from 0% to 9%. More importantly, the continuous spatial variation of electron and hole energies, together with the large exciton binding energy in the MoS₂ artificial atoms, makes it a unique solar energy funnel for conveying and concentrating the photoexcited charge carriers along the elastic strain gradient.

Introducing inhomogeneous elastic strain

When subjecting an atomically thin membrane to a simple external load, as shown in Fig. 1a, an inhomogeneous strain field can be created. The equilibrium geometry for a thin membrane is given by the Föppl–Hencky equation for membrane deformation under a stretching force at the boundaries only. For the geometry depicted in Fig. 1a, we have, for any line $\lambda(y)$, $h \int_{\lambda} dx \sigma_{\alpha} = f_{\alpha}$, where f_{α} is the stretching force on the boundary and h is the nominal membrane thickness. Owing to the force balance, the continuously narrowing membrane geometry can efficiently focus the tensile strain.

For the proposed device to function, inelastic strain relaxation as a result of dislocation plasticity or fracture must be avoided⁶. With a high-quality atomic membrane with minimal lattice defects, the edge lengths should also be minimized, as they are preferential sites for stress-aided defect nucleation¹⁷. We thus arrive at a setup similar to nanoindentation¹⁸, where a semiconducting membrane covers a circular hole on a substrate, with its rim clamped (Fig. 1b). A force probe is used to indent the suspended membrane, which will then deform towards a bottom contact. The stress gradient and hence the elastic strain gradient are maintained by the concentrated load setup with the stress scaling inversely with the radius:

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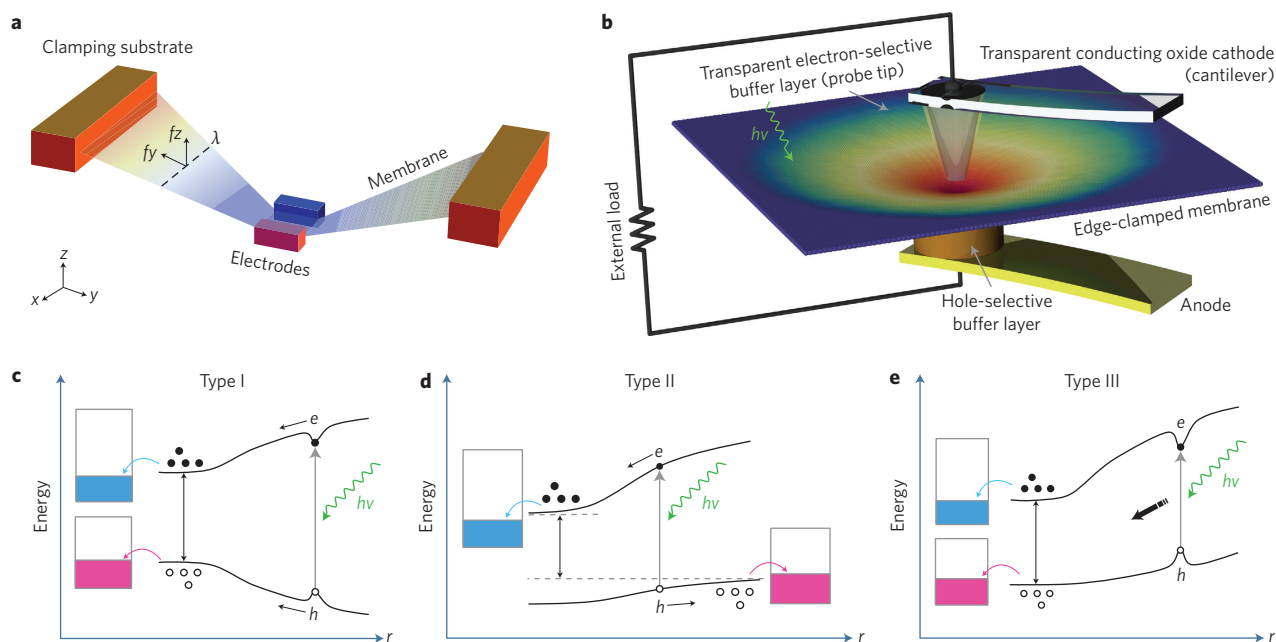


Figure 1 | Introducing inhomogeneous strain and classifying funnelling mechanisms. **a**, Schematic of an inhomogeneously strained membrane ribbon with varying width λ (dashed line). The two electrodes mechanically impose a vertical displacement on the central region of the membrane. Without the two electrodes, the equilibrium geometry of the membrane is flat. **b**, Schematic device setup for elastic strain-engineered artificial atom (not drawn to scale). Notice that although the bulk of the electrodes are metallic (for example, transparent conducting oxide), they are coated with semiconducting buffers to facilitate selective quasiparticle collection (detailed in the text). This setup completely removes the edges of the membrane ribbon in **a**, as these can serve as preferential sites for stress-aided defect nucleation. **c–e**, Three broad-spectrum solar energy funnelling mechanisms arising from a different band bending and exciton binding profile in the strain-engineered semiconducting membrane.

$\sigma_r \approx 1/r$ for $r < R$ (outer radius). Such a strain field resembles the potential field of a two-dimensional hydrogenic atom (but at a much larger length scale). It is in this sense that the strain-engineered atomic membrane can be regarded as a mesoscopic ‘artificial atom’ with electronic and optical properties that depend strongly on the strain and stress fields.

Broad-spectrum solar energy funnel

The connection between elastic strain and electronic structure in a strain-engineered artificial atom becomes apparent when the membrane is a semiconductor whose bandgap reflects a covalent bonding interaction. When stretched elastically, the covalent interaction generally weakens as the bonds lengthen, and consequently becomes a function of the imposed strain field. The stress is expected to exert significant modulation on the band energies and bandgap, analogous to band bending in a diode. The general principle here is to create a device with an electronic structure that has a continuous spatial variation due to an inhomogeneous strain field.

The scenario of continuous band bending leads us to propose the concept of a solar energy funnel. Based on the energy profiles of quasiparticles (that is, electrons and holes) and photoexcited states (that is, free electron–hole pairs or strongly bound excitons), three funnelling mechanisms are possible, as shown in Fig. 1c–e. In a Type I funnel, the energy level of photoexcited electrons continuously decreases towards the centre, while that of holes increases. Consequently, charge carriers are concentrated at the centre, as long as the lifetime of carriers is large enough for carrier migration before electron–hole recombination takes place. However, if both levels decrease with increasing strain, the result will depend on the strength of the Coulomb interaction between the quasiparticles, that is, the exciton binding energy. With weak exciton binding, electron–hole pairs will be separated by the built-in field and move in opposite directions, resulting in a Type II funnel. On the other hand, if the binding energy of an exciton is large enough to hold

the electron–hole pair together despite the built-in field, the decreasing exciton energy profile will drive the exciton to migrate towards the centre. A device based on this mechanism is classified as a Type III funnel. In all three types, the varying band/carrier energy leads to a funnelling effect. The strength of exciton binding energy determines whether Type II or III funnelling will operate in the device when the quasiparticle energy of both electron and hole decreases towards the centre of the membrane. A key advantage of such a strain-engineered device is that the performance of these devices can be adjusted to the lighting environment, as the optical bandgap is also continuously tunable.

Strain-dependent properties of MoS₂ monolayer

The structure of the MoS₂ monolayer is shown in the inset to Fig. 2d (Supplementary Fig. S1) and exhibits a remarkable trigonal prismatic geometry^{15,16,19}. On application of a biaxial strain, Poisson contraction²⁰ is expected. Owing to its hexagonal space group (*P6m2*) symmetry, the MoS₂ monolayer has isotropic in-plane elasticity, so the bandgap depends only on the two-dimensional hydrostatic strain invariant to the linear order. We therefore focus on the effect of biaxial strain on electronic structure. First-principles density functional theory^{21,22} (DFT) calculations (see Methods) indicate that both direct and indirect DFT bandgaps decrease (Fig. 2a,b), so a transition from direct bandgap to indirect bandgap occurs once the biaxial strain is applied. Other groups^{23,24} have recently reported a similar DFT bandgap reduction under biaxial tensile strain.

To acquire an accurate electronic band energy profile for the device design, we also performed many-body GW calculations^{25,26} (G, Green’s function; W, screened Coulomb interaction; see Methods). The strain-dependent bandgaps are shown in Fig. 2b, and the strain-dependent energy levels for an electron and hole at the K point in the first Brillouin zone are shown in Fig. 2d. As shown in Fig. 2b, the direct and indirect bandgaps calculated

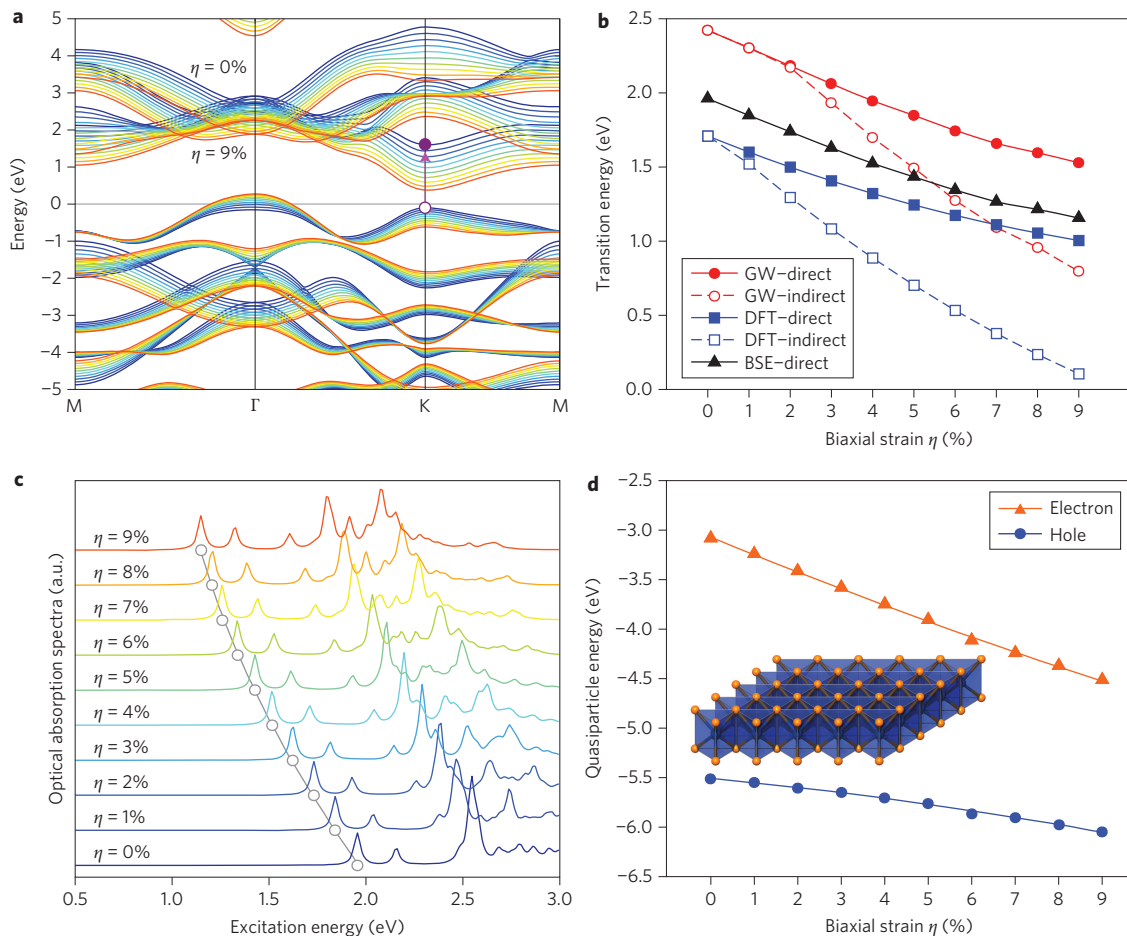


Figure 2 | Strain-dependent electronic and optical properties of MoS₂ monolayer. **a**, Electronic DFT band structure of monolayer MoS₂ under different biaxial strains from 0% (blue line) to 9% (red line). **b**, Direct and indirect bandgaps under different biaxial strains calculated using DFT and GW. The GW direct bandgap refers to the minimal direct transition energy. It is located at the K points of the hexagonal Brillouin zone, indicated by an arrow in **a**. The corresponding lowest optical excitation energy (black triangles) is calculated by BSE on top of GW quasiparticle energies with electron-hole Coulomb interaction included. **c**, Biaxial strain-dependent optical absorption spectra calculated by BSE. **d**, Biaxial strain-dependent GW quasiparticle energies for electrons and holes at the K point. Inset: atomic structure of MoS₂ monolayer.

from DFT and GW are indeed similar, except for a constant shift of 0.7 eV. Furthermore, the dielectric functions and electron energy loss spectrum of a MoS₂ monolayer are also predicted to undergo substantial changes under biaxial tensile strain (Supplementary Fig. S2).

The biaxial strain also modifies the absorption spectra and optical gap of the monolayer—the latter being another physical quantity important for device operation. As screening is generally reduced in a two-dimensional material, photoexcited quasiparticles in a MoS₂ monolayer are expected to experience stronger Coulomb attractions. We therefore go beyond the random-phase approximation by solving the Bethe–Salpeter equation (BSE)^{27,28} with GW quasiparticle energies. The calculated strain-dependent optical absorption spectra are displayed in Fig. 2c and show that the direct-to-indirect transition of the bandgap with biaxial strain does not affect the strength of the vertical optical absorption compared to the GW direct bandgap in Fig. 2b, BSE; it only shifts the whole spectrum to lower energy due to exciton binding. The lowest optical excitation in Fig. 2c originates from the doubly degenerate singlet excitons around the K points²⁹. The exciton energy of 2.0 eV at zero strain is in excellent agreement with the A peak at 1.9 eV measured using photoluminescence spectroscopy^{13,14} (the fine structure due to spin–orbit coupling discussed in ref. 29 is ignored here as it will not affect the main conclusion of strain-dependent electronic and optical energies). The excitation energy experiences substantial reduction when the strain is imposed,

decreasing to 1.1 eV at 9% biaxial strain. We then arrive at a roughly constant exciton binding energy of 0.5 eV from 0% to 9% biaxial strain. Such large constant exciton binding energy together with the reduced quasiparticle energies (Fig. 2d) makes the strain-engineered artificial MoS₂ atom a Type III solar energy funnel.

Exciton wavefunctions for the lowest excitation under zero strain are displayed in Supplementary Fig. S3. The hole state is dominated by $d_{x^2-y^2}$ and d_{xy} orbitals of molybdenum atoms and p_x and p_y orbitals of sulphur atoms, and the electron state is dominated by d_{z^2} orbitals of molybdenum atoms. The large exciton radius (~ 2 nm) and exciton binding energy (~ 0.5 eV) indicate a mixed Frenkel–Wannier–Mott exciton character. This arises from the reduced screening in two-dimensional atomically thin materials and thus stronger Coulomb attraction between electrons and holes upon photoexcitation.

Nanoindented MoS₂ monolayer as artificial atoms

To compute the morphology of the MoS₂ monolayer in the proposed device, we used a bicontinuum finite-element simulation³⁰, with a force field parameterized in the Tersoff potential form³¹ (see Supplementary Information). This coarse-grained force field method yields essentially the same membrane morphology as the analytical solution to the Föppl–Hencky equations³² (Supplementary Fig. S4). The computed morphology of the deformed membrane shows characteristic scale invariance; that is,

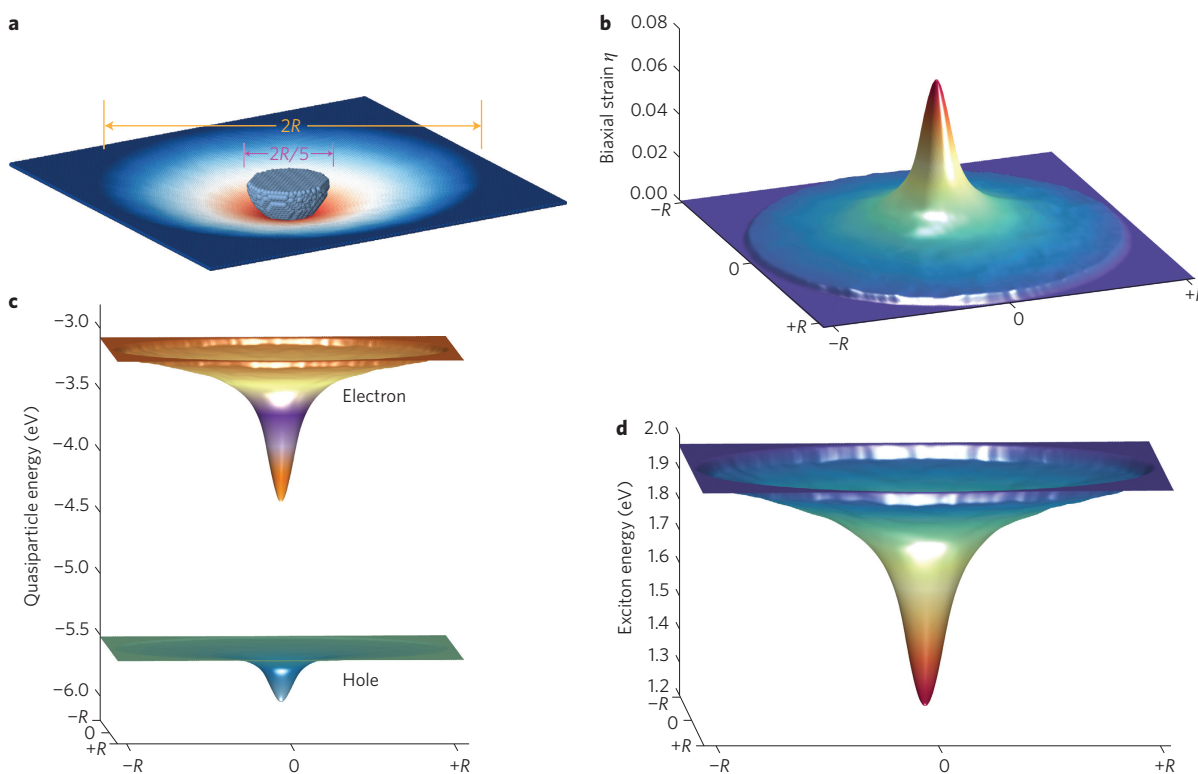


Figure 3 | Structural and physical properties of nanoindented MoS₂ monolayer. **a**, Atomic structure of circular MoS₂ membrane under indentation with a maximum vertical displacement of $R/5$ (R , circular membrane radius). **b**, Biaxial strain in the MoS₂ membrane calculated from the least-square atomic strain tensor. **c,d**, Corresponding local electron and hole energy profile (**c**) and local exciton energy profile (**d**). These are directly mapped from the profiles of homogeneously biaxial strained MoS₂ monolayer against the biaxial invariants of the strain tensor in inhomogeneously deformed MoS₂ membrane under indentation.

w/R as a function of r/R is nearly independent of R for $R > 100$ Å, where w is the vertical displacement at radius r . In practice, a force probe with a rounded tip (Fig. 1b) can be used to avoid stress singularity near the centre. A relaxed atomic structure of nanoindented MoS₂ membrane is shown in Fig. 3a, demonstrating a maximum vertical displacement of $Z = R/5$. The displacement increases gradually towards the centre of the membrane. This is expected as the boundary conditions (edge clamping and central indenting) have cylindrical symmetry, and the hexagonal lattice implies the same symmetry in the Young's modulus and Poisson ratio³³. The computed biaxial field is visualized in Fig. 3b (see Supplementary Information for calculation details). Right at the clamped rim of the suspended membrane, the biaxial strain has a small abrupt increase due to the boundary used in the simulation. As one moves inwards, the strain increases quickly, corresponding to the increase in radial stress. Because the tip of the force probe is rounded, the elastic strain does not diverge near the centre of the membrane.

With the computed strain pattern shown in Fig. 3b, we can map out the energy profiles of electronic and optical excitations on the membrane in a semiclassical sense, using those of the MoS₂ monolayer under different amounts of homogeneous biaxial strain, as presented above. The mapped band edges of electrons and holes and the energy profile of excitons for the representative membrane are shown in Fig. 3c and d, respectively. Both valence and conduction band edges are shifted downward from the rim to the centre. At the region with slowly varying strain, there is a large plateau where the positions of the band edges are almost constant, with an area occupying almost two-thirds of the circular region. The spatial variation of exciton energy in the membrane, except for the sign difference, is very similar to the strain profile, as the total

exciton energy changes almost linearly with respect to the applied biaxial strain (Fig. 2b).

The size of Type III solar energy funnel device is essentially constrained by the exciton drift length, beyond which radiative recombination will dominate and prevent the exciton from reaching the electrodes at the centre. The exciton drift length³⁴ can be estimated by

$$l_{\text{drift}} = \langle v \rangle_{\text{drift}} \tau_{1/2} = \frac{\nabla E_{\text{exc}}}{m_{\text{exc}}} \tau_{\text{dephase}} \tau_{1/2} \quad (1)$$

where $\tau_{1/2}$ is the exciton lifetime and $\langle v \rangle_{\text{drift}}$ is the average drift velocity. The latter is then estimated by the acceleration of the exciton due to the exciton energy gradient ∇E_{exc} within phase relaxation time $\langle v \rangle_{\text{drift}}$. The exciton mass is denoted m_{exc} . As we would like to have most excitons collected at the centre of the membrane, the radius of the membrane should be set to the drift length, that is, $R = l_{\text{drift}}$. Hence, the spatial gradient of the exciton energy ∇E_{exc} can be approximated by $\Delta E_{\text{exc}}/R$. We therefore have an effective membrane radius,

$$R = \sqrt{m_{\text{exc}}^{-1} \Delta E_{\text{exc}} \tau_{\text{dephase}} \tau_{1/2}} \quad (2)$$

In the plateau region, the energy variation ΔE_{exc} is ~ 0.1 eV. As we do not know exciton mass m_{exc} and phase relaxation time τ_{dephase} , we approximate them by the electron rest mass m_e and ~ 5 ps in GaAs³⁵, respectively. Fortunately, the exciton lifetime $\tau_{1/2}$ of a MoS₂ monolayer has recently been measured³⁶, with a fast photoluminescence decay component of 5 ps at temperatures of ~ 4.5 –150 K and a slow decay component of 100 ps at 270 K.

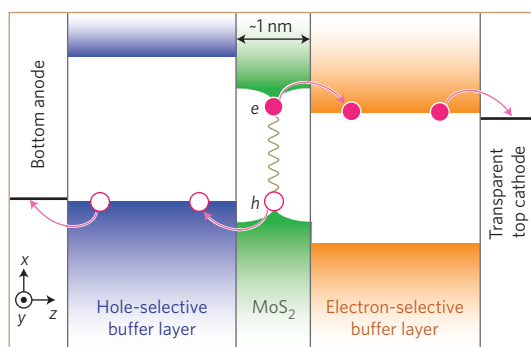


Figure 4 | Schematic of charge extraction near the centre of the atomic membrane. The atomic membrane lies on the x - y plane, and the indentation is along the z -direction. Two semiconducting buffer layers between the MoS_2 monolayer and the two electrodes serve as electron-selective and hole-selective transporters due to their specific band alignment with respect to the MoS_2 monolayer. These two buffer layers are thick enough to make direct tunnelling negligible. In addition, a transparent conducting oxide is used as the cathode so as not to block the incoming light from the top.

The estimated radius of the atomic membrane is then ~ 660 nm and $3 \mu\text{m}$ for the fast and slow decay components, respectively.

Compared to the above characteristic size of the membrane, the exciton radius of 2 nm (Supplementary Fig. S3) is indeed much smaller, meaning that the strain profile continuously and smoothly varies over a length scale comparable to the exciton size in the region of interest. The continuum strain mapping procedure is therefore justified. It is now entirely plausible that upon excitation from the ground state, the excitons will be swept by the strain-induced potential gradient and funnelled toward the centre. As both hole and electron carriers are bound in the same direction, we expect that the membrane will stay essentially charge neutral during device operation. This is advantageous, as charging might otherwise significantly reduce the efficiency of the carrier funneling. The final charge separation and harvesting of solar energy can be achieved by contacting an electron-collecting transparent semiconductor on top of the arrangement and a hole-collecting semiconductor at the bottom (Fig. 1b), with appropriate band-edge alignments to ensure one-way electron/hole flow, as in standard photovoltaic technology. As the electron-hole pairs represent energetic excited states, the membrane device works like a battery with an electropotential $\Delta V_{\text{ext}} < \Delta V_{\text{int}}$.

Discussion

What we have created in the MoS_2 membrane with the elastic strain pattern is a continuous bending of the energy levels of electrons, holes and excitons, similar to that in a photodiode, except that the photoexcited electron-hole pairs in MoS_2 are strongly bound excitons. When an electron-hole pair is generated in the plateau region following light absorption, the semiclassical potential for the quasiparticles exerts forces on the electron and hole in opposite directions. However, the large exciton binding energy will force the electron-hole pair to hold together like a molecule (despite a small polarization) and migrate towards the centre of the membrane. This is different from the case of a photodiode, where electron-hole pairs quickly dissociate, with the electrons and holes then migrating in opposite directions.

Although not central to the proposed photocarrier funneling effect, the eventual collection of solar energy is also critical. To efficiently extract charge carriers from the centre, the device design shown in the schematic in Fig. 1b was proposed, with a band alignment as shown in Fig. 4. Such a structure is similar to that of organic photovoltaic cells^{37,38}. The charge extraction process is accomplished with two carrier-collecting contacts, one of which is the

indenter itself, and the other is located beneath the indented MoS_2 membrane. The bulk of the electrodes are made of transparent conducting oxide to maximize illumination of the membrane. Key to the design are two semiconducting buffer layers, which cover the electrodes and come into direct contact with the MoS_2 monolayer. The role of these buffer layers is to selectively admit one type of charge carrier (electron or hole) while blocking the other. The carrier selectivity of these buffers arises from band alignment (detailed in Fig. 4) and follows the standard principle from photovoltaics. The nanoindebiter tip in Fig. 1b should therefore be made of, or coated with, transparent semiconductors with selected band edges. This is particularly important when building a functional funnel device such as that in Fig. 1b, because the buffer layers can help suppress direct tunnelling of carriers between anode and cathode and allow for efficient extraction of charge carriers. A layered patterning scheme is also proposed in Supplementary Fig. S6 to achieve arrays of strain-engineered artificial atoms.

The concept of a solar energy funnel achieved by engineering the electronic band energy and optical bandgap is not entirely new, and has been demonstrated experimentally in several nanostructures with graded gaps, including layer-by-layer assembled CdTe nanocrystals of different sizes³⁹ and core-shell and corrugated carbon nanotube filaments⁴⁰. Here, however, we have explored an alternative approach by generating continuously varying electronic band energy and optical bandgap in two-dimensional atomically thin membranes, by means of inhomogeneous elastic strain engineering within a single crystalline material. We would like to emphasize that it is the varying profile of band energy that leads to the direct funneling effect, while the strength of the exciton binding energy determines whether a Type II or Type III funnel device is created. The present approach is enabled by recent rapid advances in two-dimensional atomically thin sheets such as graphene, hexagonal boron-nitride, MoS_2 monolayer and so on. All of these have been exfoliated experimentally and characterized to have very high elastic strain limits. In addition, MoS_2 has some other attractive properties such as high thermal stability and chemical inertness. We therefore expect that the proposed funnel device can be verified by experiments in the near future.

In closing, we would like to make three remarks. First, photovoltaic conversion and the solar energy funnel are not the only applications of the proposed artificial atom. For example, without the electrodes, the excitons concentrated at the centre will be relaxed through radiative recombination, resulting in stronger luminescence near the centre of the membrane. This may be useful for exciton lasing, if an appropriate resonator design is devised⁴¹. The unique feature of the device, as exemplified above, is its mechanical tunability; the frequency and intensity of the emission can be tuned continuously by varying the applied strain to the atomic membrane. Second, the concept of a strain-engineered artificial atom may work with many other atomically thin sheets, including GaS, GaSe, GaTe, MX_2 -type dichalcogenides ($M=\text{Mo, Nb, Ni, Sn, Ti, Ta, Pt, V, W, Hf}$ and so on, and $X=\text{S, Se, Te}$), M_2X_3 -type trichalcogenides ($M=\text{As, Bi, Sb}$ and so on, and $X=\text{S, Se, Te}$), MPX_3 ($X=\text{S, Se}$), MAX_3 ($A=\text{Si, Ge}$, and $X=\text{S, Se, Te}$), and alloy sheets like $\text{M}_x\text{M}'_{1-x}\text{S}_2$. It can also be applied to other thin membranes based on conventional materials, such as a silicon nanomembrane⁴², which can sustain a significant tensile strain of a few percent. This level of tension is sufficient to induce the proposed funneling effect, even though the driving force is still not as large as in maximally indented atomic sheets. Finally, although we illustrate the application of the elastically deformed membrane semiconductor as a solar energy concentrator, we would also like to emphasize the more general notion of it being a tunable artificial atom. Each deformable membrane should have its own characteristic excitations (electronic and electromagnetic). Upon juxtaposition, they will interact electronically

and optically. It is our hope, then, that complex superlattice arrays of these artificial atoms can be built on a whole membrane, the properties of which will be tuned at the level of the fundamental 'atomic' building blocks by strain engineering.

Methods

Ground-state properties of MoS₂ monolayer. The ground-state atomic structure of MoS₂ monolayer under different biaxial strains was first assessed by DFT^{21,22} calculations. These were carried out using the Vienna Ab initio Simulation Package (VASP) with a plane wave basis^{43,44} and the projector-augmented wave method⁴⁵. Exchange-correlation functionals in a Perdew–Berke–Ernzerhof (PBE) form⁴⁶ within the generalized-gradient approximation (GGA)^{47,48} were used in all DFT calculations. We adopted the Monkhorst–Pack scheme⁴⁹ of k-point sampling for zone integration, and selected the number of k-points to achieve a convergence in total energy within ~1 meV/atom. The total energy of each optimized structure was calculated again with the more accurate tetrahedron zone summation with Blöchl correction⁵⁰. A qualitative electronic band structure for different biaxial tensile strain was also obtained from DFT–PBE calculations.

Excited-state properties of MoS₂ monolayer. Two physical quantities related to this design of solar energy funnel are the electronic excitation energy (quasiparticle band energy for electron and hole) and optical excitation energy (exciton energy). However, neither of these can be obtained accurately from DFT Kohn–Sham eigen energies using the GGA exchange–correlation function. We therefore computed quasiparticle energies for electrons and holes using many-body perturbation theory calculations within Hedin's G₀W₀ approximation^{25,26}, while the optical absorption spectra and exciton energies were further obtained by solving a Bethe–Salpeter equation^{27,28} on top of G₀W₀ quasiparticle energies within the Tamm–Dancoff approximation, which explicitly takes into account Coulomb interactions between quasi-electrons and quasi-holes. Here, an energy cutoff of 350 eV and Monkhorst–Pack k-point sampling of 18 × 18 × 1 were chosen. Both GW and BSE calculations require much more computational effort and memory storage than DFT calculations, because the GW method needs to evaluate frequency-dependent non-local self-energy and the BSE method is essentially solving two-particle Green's function, whereas DFT with GGA functions is intrinsically local and frequency-independent. Because an exciton corresponds to the excitation of electron–hole pairs, its wavefunction is a six-dimensional quantity, $\Psi(\mathbf{r}_e, \mathbf{r}_h) = \sum_{\mathbf{k}, \nu} A_{\nu\mathbf{k}} \phi_{\mathbf{k}}(\mathbf{r}_e) \phi_{\mathbf{k}}^*(\mathbf{r}_h)$, where $\phi_{\mathbf{k}}(\mathbf{r}_e)$ and $\phi_{\mathbf{k}}^*(\mathbf{r}_h)$ are electron and hole wavefunctions. We have reconstructed exciton wavefunctions and plotted out the corresponding electron (hole) wavefunction by holding its counterpart hole (electron) at a specific location (Supplementary Fig. S3). Spin–orbit coupling is not included as it will not affect the main conclusion of strain-dependent electronic and optical bandgaps of MoS₂ monolayer.

Geometry of the MoS₂ membrane under nanoindentation. Owing to the large size of the atomic membrane under consideration here, it is unrealistic for us to relax the nanoindented membrane by first-principles DFT calculation directly. We therefore fitted a set of Tersoff potential parameters for MoS₂, and then performed force-field calculations to obtain the geometry of the MoS₂ membrane under indentation. More details are provided in the Supplementary Information.

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Author contributions

J.L. designed the project. J.F., X.F.Q. and C.W.H. carried out the calculations and the modelling. J.F., X.F.Q. and J.L. wrote the paper. All authors contributed to discussions of the results.

Additional information

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Competing financial interests

The authors declare no competing financial interests.