

Supporting Information

Multifunctional inorganic nanomaterials for energy applications

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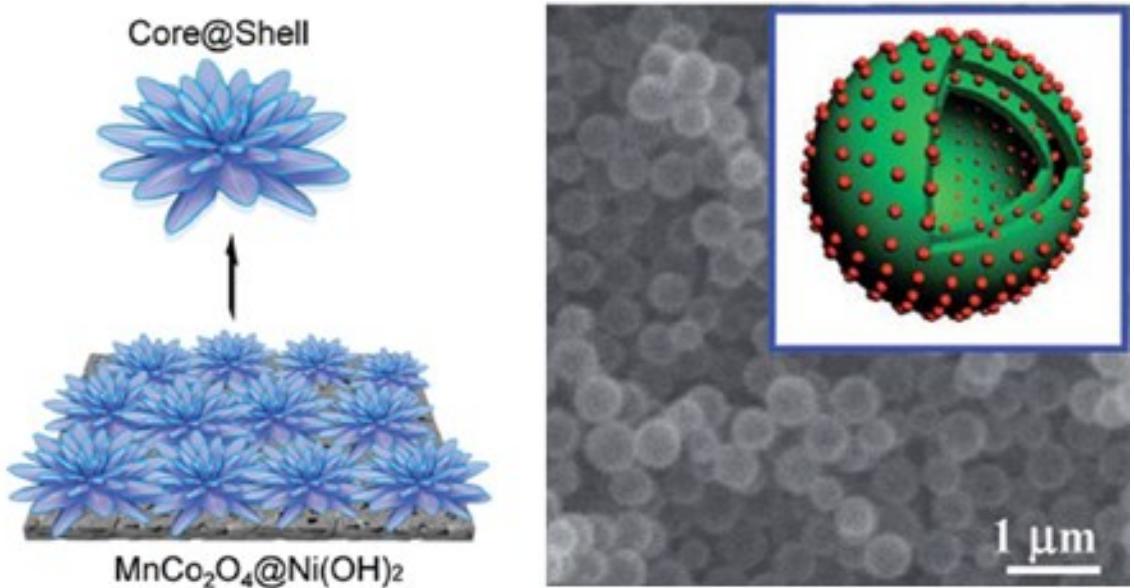


Fig. S1 MnCo₂O₄@Ni(OH)₂ core-shell flowers for asymmetric supercapacitor materials with ultrahigh specific capacitance and V₂O₅-SnO₂ double-shelled nanospheres.^{1,2}

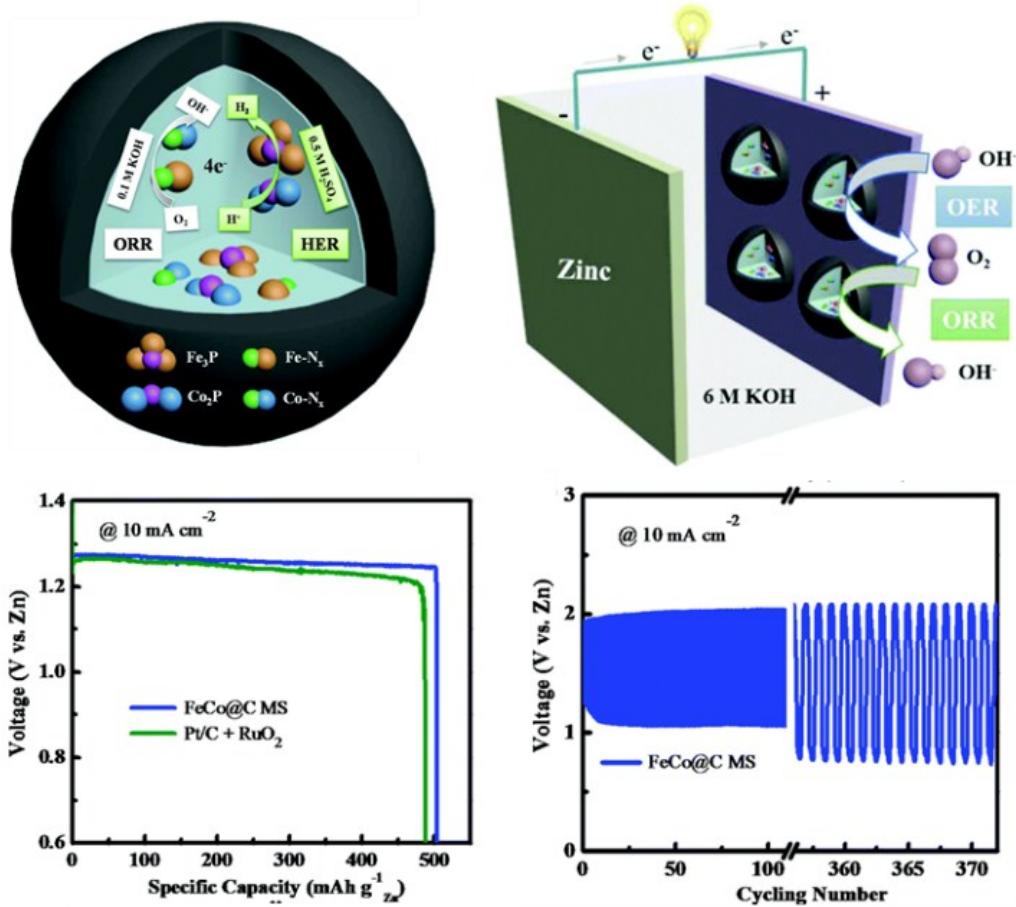


Fig. S2 (a) Schematic illustration of ORR and HER processes of FeCo@C MS. (b) Illustration of the rechargeable Zn-air battery. (c) Electrochemical performance of rechargeable Zn-air batteries using FeCo@C MS as catalysts in 6 M KOH . Galvanostatic discharge curves of primary Zn-air batteries. The specific capacity is normalized by the mass of the consumed Zn anode. (d) Pulse cycling performance of the rechargeable Zn-air batteries using FeCo@C MS at 10 mA cm^{-2} with 5 min discharge and 5 min charge.³

Table S1 Summaries of performance and preparation of multifunctional inorganic materials in various energy applications

| | Material | Morphology | Application | Performance | Preparation | Reference |
|-------------------|---|-----------------|----------------|---|------------------------------|-----------|
| Energy generation | SnSe | Nanosheet | Thermoelectric | Low thermal conductivity High ZT of 0.055 at 501 K | Thermal evaporation | 4 |
| | Cd-doped SnSe | Nano-crystal | Thermoelectric | Low thermal conductivity High power factor High hole carrier concentration High ZT approximate to 1.7 at 823 K | Solvothermal | 5 |
| | Sn _{0.99} Na _{0.01} Se-STSe | Nanoprecipitate | Thermoelectric | Ultralow lattice thermal conductivity of below 0.3 W m ⁻¹ K ⁻¹ High peak ZT of 1.33 at 773 K | High temperature melting | 6 |
| | PbTe-4% InSb | Nanocomposite | Thermoelectric | High ZT of 1.83 at 773 K | High pressure heat treatment | 7 |
| | Cu ₅ FeS ₄ | Core-shell | Thermoelectric | High power factor Low thermal conductivity Enhanced ZT value of ≈0.62 at 710 K | Colloidal synthesis | 8 |
| | Cu ₂ Se/CNT | Nanocomposite | Thermoelectric | Low lattice thermal conductivity Record-high ZT of 2.4 at 1000 K | High energy ball-milling | 9 |
| | C ₆₀ /TiS ₂ | Nanocomposite | Thermoelectric | Flexible High Seebeck coefficient High power factor | Printing | 10 |

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| | | | High ZT approximate to 0.3 at 400 K | | |
| Kevlar microfiber-ZnO | Nanowire | Piezoelectric | High flexibility Robustness and durability Open-circuit voltage and short-circuit current are 1.8 mV and 4.8 pA, respectively. | Solvothermal | 11 |
| S-treated MoS ₂ | Nanosheet | Piezoelectric | Output peak current and voltage are increased by more than 3 times (100 pA) and 2 times (22 mV), respectively. | Chemical vapor deposition | 12 |
| PVDF-AgNW | Nanofiber | Triboelectric | Excellent mechanical stability High output performances | Electrospinning | 13 |
| P(VDF-TrFE) | Nanocomposite | Triboelectric | Boosting power-generating performance is achieved for 1130 V of output voltage and 1.5 mA of output current | Spin-coating heat treatment | 14 |
| Al/PTFE | Micro-nanocomposite | Triboelectric | Excellent high-temperature tolerance (temperature ring of -30 to 550 °C) Wear-resisting ability High hardness (rockwell hardness: 63 HRM) High output voltage and current of 221 V and 27.9 μA cm ⁻² | Ball-milling | 15 |
| Energy conversion | ZnO-SnO ₂ | Core-shell | Solar cell PCE : 14.35% Good match between the energy level of the SnO ₂ shell High electron mobility of the core ZnO nanoparticles | Coprecipitation | 16 |
| | TiO ₂ -SiO ₂ | Core-shell | Solar cell PCE : 7.00% Anti-reflective coating decrease the reflectance by producing a refractive index gradient | E-beam evaporation and chemical bath deposition | 17 |

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|-----------------------------------|---------------------------|------------|---|--------------------------------------|----|
| n-ZnO/p-SnS | Core–shell nanowire array | Solar cell | Relative conversion efficiency : 37.3% The piezo-phototronic effect was effectively applied to improve The relative conversion efficiency | Rf magnetron sputtering | 18 |
| PbS–CdS | Core–shell quantum dot | Solar cell | PCE : 2.5% Improved passivation of the PbS Surface by the CdS shell, leading to a lower electron trap density | Hot injection | 19 |
| WO ₃ –TiO ₂ | Core–shell | Solar cell | PCE : 11.24% WO ₃ nanosheet arrays yield significantly enhanced photovoltaic performance as compared to nanoparticles and nanorod arrays Due to good perovskite absorber infiltration in the porous scaffold and more rapid carrier transport. | Electrospraying and hydrothermal | 20 |
| PbSe-PbS | Core-shell quantum dot | Solar cell | PCE: 6.5% PbS shell formation on the PbSe core mitigates the trade-off relationship between the open circuit voltage and the short circuit current density. | Precipitation | 21 |
| Pt–C | Core–shell | Fuel cell | A negligible voltage loss after 30000 cycles The robustness of the carbon shells that secure the Pt nanoparticles. | Ultrasonic mixing and heat treatment | 22 |
| IrNi@PdIr/C | Core–shell | Fuel cell | Specific activity is 2.2 times that of commercial Pt/C PdIr shell inhibits oxide formation. | Replacement reaction | 23 |
| Pd/PtCu | Core–shell | Fuel cell | ORR and MOR mass activities were 8.3 and 3.3 times higher than those of commercial Pt, respectively | One-pot epitaxial growth | 24 |

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| | | | Lower the catalyst cost but also improve the catalytic activity and stability | | |
| Pt-Rh _x S _y | Core-shell | Fuel cell | Electrochemical active surface area/mass of core-shell catalyst is 7 times that of commercial Rh _x S _y catalyst. | Precipitate and thermal treatment | 25 |
| CNT/Ni | □ | Fuel cell | The functionalized redox nanomaterial exhibits reversible electrocatalytic activity for the H ₂ /2H ⁺ interconversion from pH 0 to 9, with catalytic preference for H ₂ oxidation at all pH values. | - | 26 |
| Co ₉ S ₈ /MoS ₂ | Core-shell | Catalysis | HER: 97 mV (10 mA cm ⁻²) Tafel slope: 71 mV dec ⁻¹ Precisely tune Tensile surface strain | S vapor-assisted | 27 |
| CoP@PS | Core-shell | Catalysis | Her: 80 mV (10 mA cm ⁻²) The significant enhancing effect from nanometer thick amorphous PS layer on the turnover process | Wet chemical process and phosphidation | 28 |
| PtPb/Pt | Core-shell | Catalysis | 50,000 voltage cycles with negligible activity decay | Nonaqueous conditions | 29 |
| Energy storage | crystallineFe ₂ O ₃ / amorphous Fe ₂ O ₃ | Supercapacitors | Capacitance of 701 F g ⁻¹ The tunable amorphous Layer facilitates the Li ⁺ diffusion while introduced oxygen defects in Fe ₂ O ₃ can be effectively tuned to improve electronic conductivity | Hydrothermal and heat treatment | 30 |
| | P-doped Ni(OH) ₂ -MnO ₂ | Supercapacitors | Areal capacitance of 5.75 F cm ⁻² Without capacitance loss at after 10 000 cycles | Hydrothermal | 31 |

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| | | | Large surface area, good mechanical stability High ion diffusivity and numerous electroactive sites | | |
| Zinc–nickel–cobalt oxide @Ni(OH) ₂ | Core–shell nanowire arrays | Supercapacitors | Specific capacitance of 2847.5 F cm ⁻³ (10.678 F cm ⁻²) The increased electrical conductivity of the core electrode | Hydrothermal and calcination | 32 |
| Co ₃ O ₄ @NiCo ₂ O ₄ | Core-shell | Supercapacitors | Specific capacitance of 1450 F g ⁻¹ Cycling stability (~4.2% loss after 6000 cycles) Rational design of nico ₂ o ₄ nanoflakes adhering on Co ₃ O ₄ nanowires, which promotes two electroactive materials utilizing the synergistic effect to supply more pathways for accelerating fast electron and ion transfer | Hydrothermal | 33 |
| Ni-Co@Ni-Co LDH | Core-shell | Supercapacitors | Capacitance of 2200 F g ⁻¹ 98.9 % after 5000 cycles A novel hierarchical nanotube array with a massive layered top and discretely separated nanotubes in a core–shell structure | Template assisted electrodeposition | 34 |
| MnCo ₂ O ₄ @Ni(OH) ₂ | Core–shell flowers | Supercapacitors | Capacitance of 2154 F g ⁻¹ Hierarchical structure can not only efficiently ensure the synergistic effect of the two pseudocapacitive materials but also promote the diffusion and migration of electrolyte ions during the rapid charge/discharge process | Hydrothermal | 1 |
| ErOOH | Colloid | Supercapacitors | Capacitance of 1811 F g ⁻¹ Short ion diffusion and electron transfer length to enable the fast and | In situ coprecipitation | 35 |

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| | | | reversible faradaic reactions | | |
| V ₂ O ₅ –SnO ₂ | Core-shell | Lithium-ion batteries | Anode: 947 mA h g ⁻¹ Cathode: 174 mA h g ⁻¹ The current nanoarchitecture provides short li ion pathways and high electronic and ionic conductivity, and the hollow architecture is able to accommodate large volume variations | Coprecipitation | 2 |
| Mn _{1-x} Fe _x P | Solid solution phosphide | Lithium-ion batteries | A capacity of 360 mA h g ⁻¹ after 100 cycles at a high current density of 2 A g ⁻¹ | High energy mechanical milling | 36 |
| Cu ₃ Si-Si@C@G | Core-shell | Lithium-ion batteries | Good rate performance and delivers reversible capacity of 483 mA h g ⁻¹ after 500 cycles with capacity retention of about 80% at high current density of 4 A g ⁻¹ | Sol-gel coating | 37 |
| H-MoP@rGO | Core-shell | Sodium-ion batteries | The specific capacity is as high as 353.8 mA h g ⁻¹ at 1 A g ⁻¹ after 600 cycles Extraordinary rate performance of 183.4 mA h g ⁻¹ at an ultrahigh current density of 10 A g ⁻¹ even after 3000 cycles. | Hydrothermal followed by phosphorization procedure | 38 |
| FeS ₂ | Foam-like nanostructure | Sodium-ion batteries | High electrical conductivity Good ion diffusion kinetics High inhibition capacity of volume expansion. High capacity of 823 mA h g ⁻¹ at 0.1 A g ⁻¹ , very close to the theoretical capacity of FeS ₂ Good rate capability of 581 mA h g ⁻¹ at 5.0 A g ⁻¹ Good cycle ability of 754 mA h g ⁻¹ at 0.2 A g ⁻¹ with 97% retention after | Solution combustion followed by heat treatment | 39 |

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| | | | 80 cycles | | |
| VS ₄ | 3D self-assembled nanoarchitectures | Sodium-ion batteries | High reversible capacity of 412 mA h g ⁻¹ at 0.2 A g ⁻¹ after 230 cycles Capacity of 345 and 293 mA h g ⁻¹ even at 1.0 and 2.0 A g ⁻¹ , respectively | Hydrothermal | 40 |
| Ti ₂ Nb ₂ O ₉ | Nanosheet | Sodium-ion batteries | High reversible capacity of 250 mA h g ⁻¹ at 50 mA g ⁻¹ at a suitable average voltage of approximate to 0.7 V. | Liquid exfoliation combined with thermal dehydration | 41 |
| Ni/PCNFO | Nanofiber | Lithium-sulfur batteries | High electrical conductivity High specific capacity of 1320 mA h g ⁻¹ Excellent rate capability of 780 mA h g ⁻¹ Long cycling stability of 910 mA h g ⁻¹ after 500 cycles at 0.2 C | Electrospinning | 42 |
| 3DNG/TiN | Composite | Lithium-sulfur batteries | Ultrahigh areal capacity of 12.0 mA h cm ⁻² at a high current density of 8.03 mA cm ⁻² | Hydrothermal | 43 |
| rGO/GC/S | Composite | Lithium-sulfur batteries | High capacity of 524 mA h g ⁻¹ after 100 cycles at a current rate of 0.2 C | Low-temperature wet spinning | 44 |
| Fe ₃ C@N-C | Core-shell | Lithium-sulfur batteries | High capacity of 1351 mA h g ⁻¹ at 0.1 C Outstanding rate capability and cycling stability | Thermal treatment | 45 |
| Ni-doped CoO | Nanosheet | zinc-air batteries | Excellent performance with a record-high discharge peak power density of 377 mW cm ⁻² , and works stable for >400 h at 5 mA cm ⁻² | Hydrothermal | 46 |
| FeCo@C MS | Core-shell | zinc-air batteries | High discharge voltage of 1.27 V High specific capacity of 503 mA h g ⁻¹ | Solvothermal followed by two-step | 3 |

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| | | | | Energy density of 639 W h kg^{-1} | carbonization | |
| Energy saving | WO_3 | Tunneled phosphorus-doped film | Smart window | 6.1 s for the colouration speed and 2.5 s for the bleaching speed Colouration efficiency $55.9 \text{ cm}^2 \text{ C}^{-1}$ Higher retention (91.5%) of transmittance modulation after 1000 electrochromic (EC) cycles | Precipitation, anneal, red P ignition | 47 |
| | WO_3 | Mesoporous film | Smart window | $T_c = 1.4 \text{ s}$ $T_b = 1.1 \text{ s}$ 93.1% of the original value after 1000 cycles and 74.6% after 2500 cycles CE value of $79.7 \text{ cm}^2 \text{ C}^{-1}$ | Dip-coating sol-gel | 48 |
| | Cs_xWO_3 | Nanorod | Smart window | Visible light transmittance reaching 78.22% NIR shielding rate being 97.36% Transparent insulation index attaining 175.58 | Solvothermal, heat-treat, citric acid thermolysis | 49 |
| | Pt-doped K_xWO_3 | Nanorod | Smart window | Visible light transmittance reaching 39.58% NIR shielding rate being 94.75% | Solvothermal | 50 |
| | $\text{M}_x\text{WO}_3/\text{ZnO}$ | Nanorod | Smart window | Visible light transmittance reaching 92 % High NIR shielding rate 50% of toxic no gas could be decomposed by composite and relevant film under the irradiation of UV light | Hydrothermal | 51 |
| | BN | Nanosheet | Cooling liquids | 2.39 W mK^{-1} at 24 vol% loading | Molten alkali-assisted liquid | 52 |
| | Graphene | Nanoplatelet | Coolants | The maximum thermal conductivity enhancement of 22.92% is | Sonication | 53 |

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| | | | | attained for the sample containing 0.1 wt% at a fluid temperature of 45 °C The dynamic viscosity and density are close to those for DI water High stability | | |
| | GO-TiO ₂ | Nanosheet | Nanolubricant | The film thickness of lubricant with GO-TiO ₂ was 28.07 nm | Solvothermal | 54 |
| Energy transmission | Al/InAs | Nanowire | Superconductivity | Biassing the junction to $\phi \approx \pi$ reduces the critical field at which the zero-bias peak appears, with respect to $\phi = 0$ | Electron beam lithography | 55 |
| | Nb/InAs, Al/InAs | Nanowire | Superconductivity | Al/InAs-nanowire/Al junctions comprising an Al electrode separation of 220 nm With no clear Josephson supercurrent was achieved, the measured voltage signal becomes nonlinear around zero bias current | Mbe(molecular beam epitaxy) | 56 |
| | InAs/V | Nanowire | Superconductivity | The nanoscale superconducting vanadium had a high out-of-plane critical field BC = 1.8 T, far exceeding the bulk value Devices fabricated from hybrid InAs/vanadium nanowires showed superconducting transitions at T < 4 K | Mbe(molecular beam epitaxy) | 57 |
| | YBa ₂ Cu _{3-x} Ni _x O _{7-δ} | Nanowire | Superconductivity | The T _c values range between 70 and 93.2 K for samples with x = 0.04 to x = 0.00, respectively, where the substitution of Cu with Ni results in a lower T _c | Electrospinning | 58 |
| | ABC-TLG/hBN | Hall bar geometry | Superconductivity | 1/4-filling Mott insulator with D = -0.54 V nm ⁻¹ and n = -5.4 × 10 ¹¹ cm ⁻² . Estimated superconducting T _c of 0.65 K | Dry transfer | 59 |

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| | | | At $D = -0.17 \text{ V nm}^{-1}$, where only the 1/2-filling Mott state exists, the phase diagram shows no superconductivity even at base temperature | | |
| Graphene/ α -Mo ₂ C | Nonlayered | Superconductivity | The extracted value of the interface transparency is $T \approx 0.7\text{--}0.8$, indicating a highly transparent high-quality graphene-2D α -Mo ₂ C interface in comparison with previous work on graphene hybrid/heterostructure devices with either deposited or stacked superconducting contacts | Chemical vapor deposition | 60 |
| NbSe ₂ | Nanoplate | Superconductivity | Superconductivity transition is observed within the temperature range of 4.60–5.40 K The T_c at zero magnetic field is 5.05 K, slightly lower than that of bulk NbSe ₂ (~ 7.2 K) A magnetic field of 3.0 T completely suppresses the superconductivity down to the lowest measured temperature. $T_{\text{BKT}} \approx 4.83$ K | Chemical vapor deposition | 61 |
| NbC | 3D nanowire | Superconductivity | FIBID planar nanowires become superconducting at $T_c \approx 5$ K The critical temperature of free-standing 3D nanowires is as high as $T_c \approx 11$ K, which is close to the value of bulk NbC. | Focused ion beam direct writing | 62 |
| WC | 3D hollow nanowire | Superconductivity | The nanowires become superconducting at 6.4 K and show large critical magnetic field and critical current density resulting from their quasi-one-dimensional superconducting character. | Focused electron beam induced deposition | 63 |
| ZnO | Flower-shaped particle | Insulating | The conductivity decreased with increasing amount (0.1 to 3 wt%) of coated ZnO nanoparticles The addition of 3 wt% C ₈ -coated ZnO nanoparticles reduced the | Heat treatment | 64 |

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| | | | conductivity of the LDPE by 2–3 orders of magnitude | | |
| MgO | Hexagonal platelet | Insulating | The lowest volume conductivity was ca. 7×10^{16} S m ⁻¹ for 3 wt% surface coated nanoparticles Uniformly dispersed MgO nanoparticles up to contents as high as 9 wt%, with maintained 10–100 times reduced volume conductivity | Liquid precipitation and heat treatment | 65 |

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