## **Supporting Information**

## Enhancing the osmotic energy conversion of a nanoporous membrane: influence of pore density, pH, and temperature

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Pore-pore distance $a_1$ (nm)	Pore density (pores/cm <sup>2</sup> )
200	2.5×10 <sup>9</sup>
320	9.8×10 <sup>8</sup>
450	$5.0 \times 10^{8}$
600	$2.8 \times 10^{8}$
800	$1.6 \times 10^{8}$
1000	$1.0 \times 10^{8}$
2000	$2.5 \times 10^{7}$
3200	1.0×10 <sup>7</sup>
10000	$1.0 \times 10^{6}$

Table S1. Dependence of pore density on pore-pore distance.



**Fig. S1** Variation of diffusion current with pore density for a multi-pore system with  $a_0=5 \mu m$  (infinite reservoir). Parameters used: L=1000 nm, d=10 nm,  $\sigma_s=-0.06 \text{ C/m}^2$ ,  $C_H/C_L=100 \text{ mM/1 mM}$ , and T=298 K, n=4, and  $H_{res}=2000 \text{ nm}$ . The pore density is adjusted by the pore-pore distance  $a_1$  (Table S1).



Fig. S2 (a) Schematic representation of a single-pore system, where a cylindrical nanopore connects two large, identical reservoirs with  $a_0$  being half width. The boundary marked red has the surface charge density  $\sigma_s$ . (b) Variation of osmotic current with pore-pore distance; both the results for single-pore and those for multi-pore are presented. Temporal variation of osmotic current, (c), and Cl<sup>-</sup> concentration, (d), after a  $\Delta P$ =-1 bar is applied for the case of  $a_1$ =200 nm. The parameters used in (b)-(d) are *L*=50 nm, *d*=10 nm, pH 8,  $C_{\rm H}/C_{\rm L}$ =100 mM/1 mM, *T*=298 K,  $H_{\rm res}$ =2000 nm, and  $a_0$ = $a_1/2$ .



**Fig. S3** (a) Variation of averaged surface charge density of the wall  $\bar{\sigma}_s$  with pH for different bulk concentrations. (b) Variation of  $\bar{\sigma}_s$  with temperature for two pH levels. The parameters used in (b) are *L*=600 nm, *d*=10 nm, *C*<sub>H</sub>/*C*<sub>L</sub>=1000 mM/1 mM, *H*<sub>res</sub>=2000 nm, and  $a_0=a_1/2=100$  nm.

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Pore-pore distance	Pore density	Porosity (%)
$a_1$ (nm)	(pores/cm <sup>2</sup> )	when d=10 nm
50	$4.0 \times 10^{10}$	3.142
100	$1.0 \times 10^{10}$	0.785
140	5.1×10 <sup>9</sup>	0.401
200	2.5×10 <sup>9</sup>	0.196

 Table S2. Dependence of pore density on pore-pore distance and the corresponding porosity.



**Fig. S4** Variation of the net power  $P_{net}$ , (a) and (c), and overall power increment OPI and power increment PI, (b) and (d), with pore length. In (a) and (c), green dashed curve: power of PV work  $P_{PV}$  of  $\Delta P$ =-0.5 bar; wine dash-dotted curve: power of PV work  $P_{PV}$  of  $\Delta P$ =-1 bar. In (b) and (d), gray dashed curve: OPI=0. In (d), dash-dotted curve:  $\Delta P$ =-0.5 bar; solid curve:  $\Delta P$ =-1 bar. Parameters used are *d*=10 nm,  $C_{\rm H}/C_{\rm L}$ =1000 mM/1 mM, *T*=323 K, pH 11,  $H_{\rm res}$ =2000 nm, and  $a_0$ = $a_1/2$ . In (a) and (b),  $a_1$ =100 nm. In (c) and (d),  $a_1$ =50 nm.