

1 Supporting Information

2 Green and Efficient Piezocatalytic Hydrogen Production Achieved by 3 Modifying SrBi₄Ti₄O₁₅ with CdSe

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20 **1. Experimental**

21 **1.1 Synthesis methods**

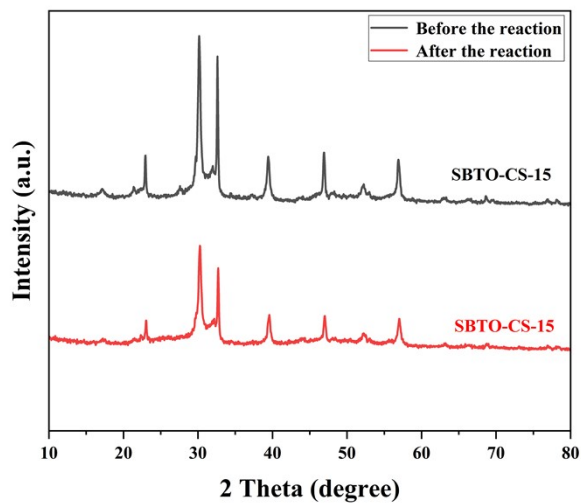
22 The CdSe QDs were synthesized by imitating previous reported work^[1]. In
23 a typical routine utilizing a three-neck flask, in order to prepare the Se
24 precursor, Na₂SO₃ (0.03 mol) and selenium powder (5.0 mmol) were added
25 into 100 mL deionized water and refluxed under a nitrogen atmosphere at
26 80 °C for 6 h. After that, CdAc·2H₂O (4 mmol) was dissolved in 50 mL of
27 deionized water and then 100 mL Se precursor was added dropwise into
28 the mixture at 500 μL/min using a peristaltic pump during reflux and
29 insulation at 80 °C for 6 h. The crimson precipitates were centrifuged and
30 washed with ethanol and deionized water repeatedly to remove the residual
31 reactants, and dried at 60 °C in a vacuum oven, labeled as CS.

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33 **2.XPS-VB calculation**

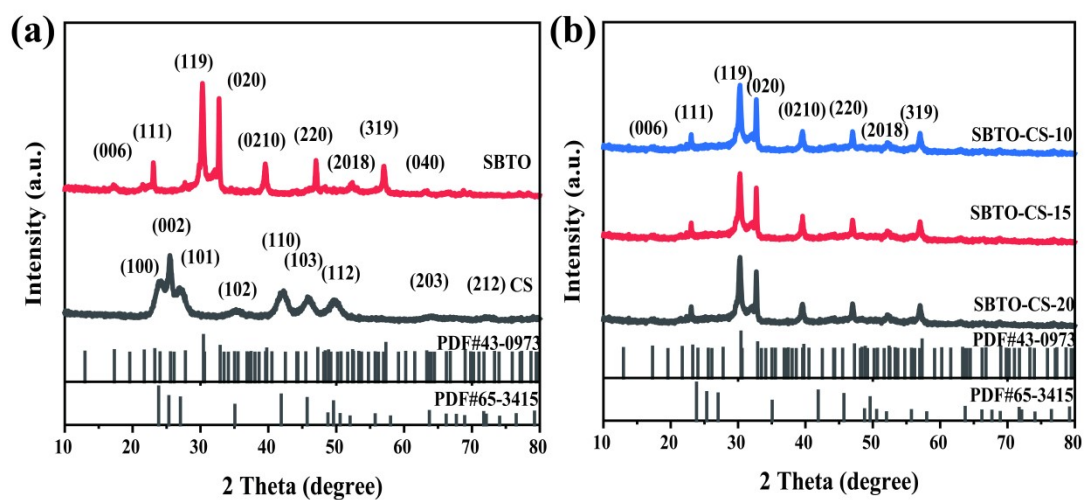
34 The small binding energy range of VB_{-XPS}^[2] is used to calculate its valence
35 band. It is reported that the work function of the sample and the contact
36 potential difference (ΔV) have the following relationship: $\Delta V = \Phi - \phi$ (Φ
37 and ϕ are the work functions of the material and the XPS analyzer,
38 respectively). So $E_{VB, NHE} = \phi - 4.44 \text{ eV} + E_{VB, XPS}$ (ϕ is the work function of
39 XPS analyzer as 4.44, where 4.44 eV is the energy difference from Fermi
40 energy level to valence band relative to the standard hydrogen electrode at
41 pH = 7, $E_{VB, NHE}$ is the position of energy band under the standard hydrogen

42 electrode, and $E_{VB, XPS}$ is the position of the intersection of slope and X-
43 axis).



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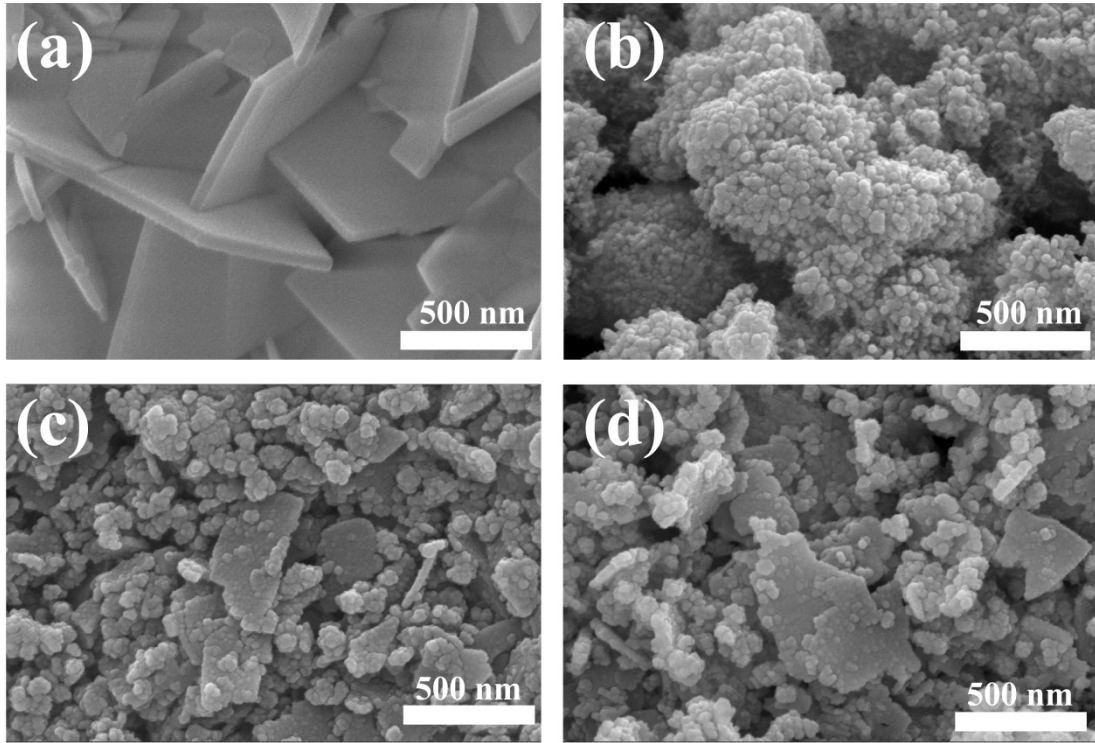
45 Fig. S1 XRD patterns of SBTO-CS-15 before and after reaction (reaction
46 time is 3h).



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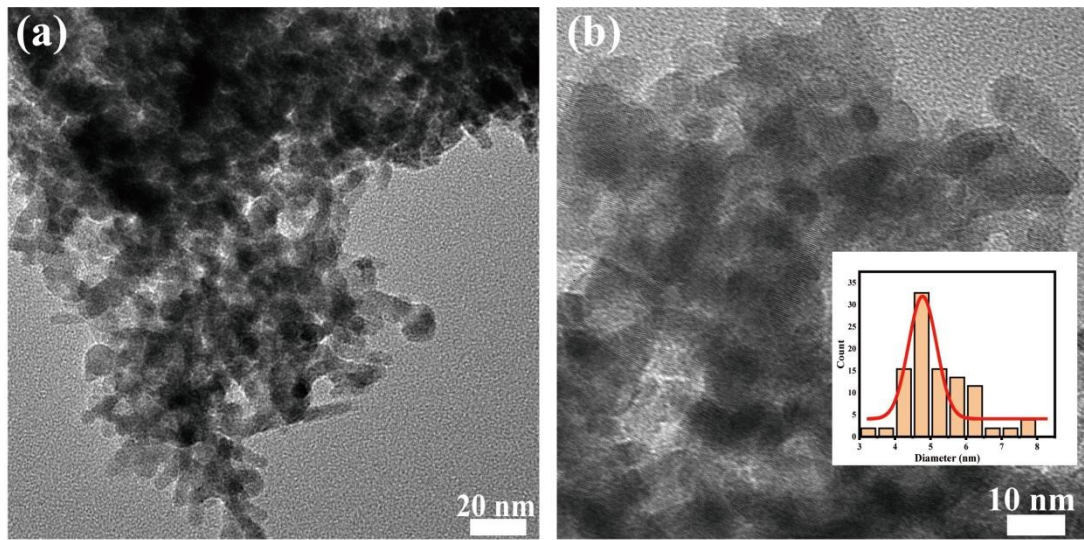
48 Fig. S2 (a) XRD patterns of pure SBTO, CS samples. (b) XRD patterns of
49 SBTO-CS-10, 15, 20 samples.

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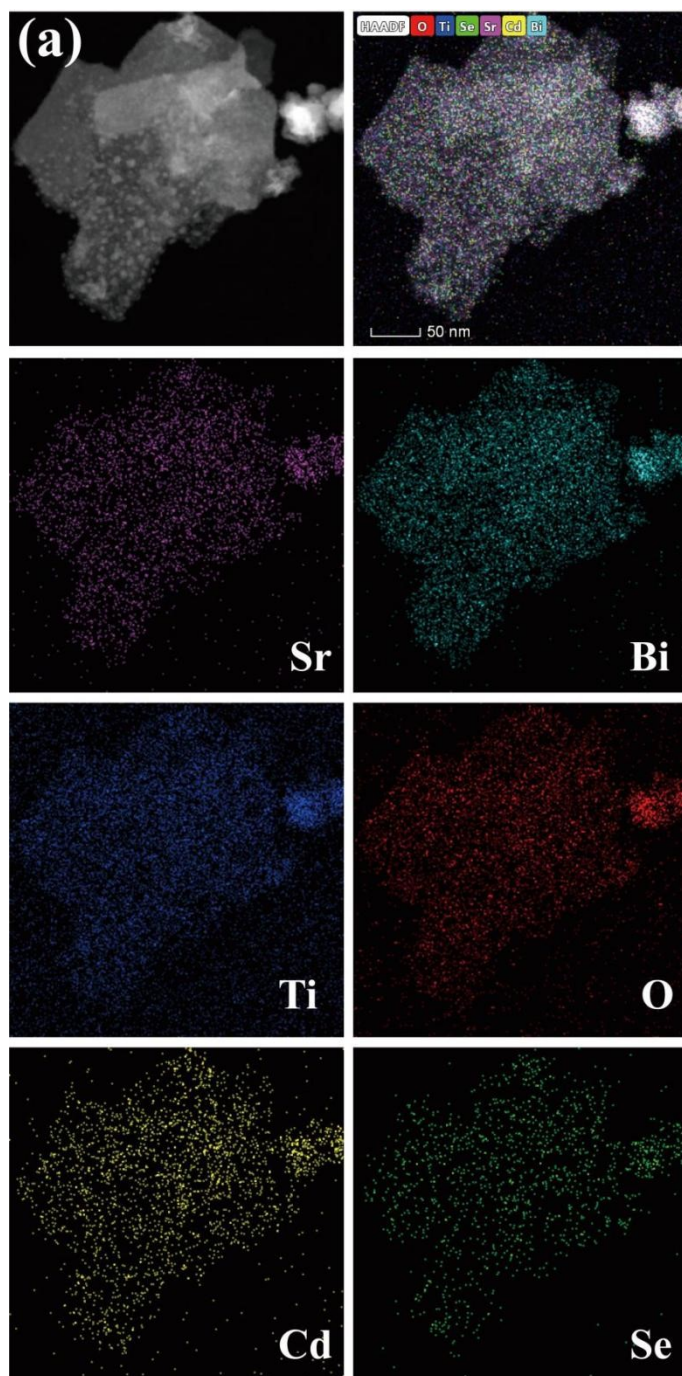
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52 Fig S3. SEM images of (a) pure SBTO and (b) CS. (c-d) SEM images of
 53 SBTO-CS-15 (at different regions).



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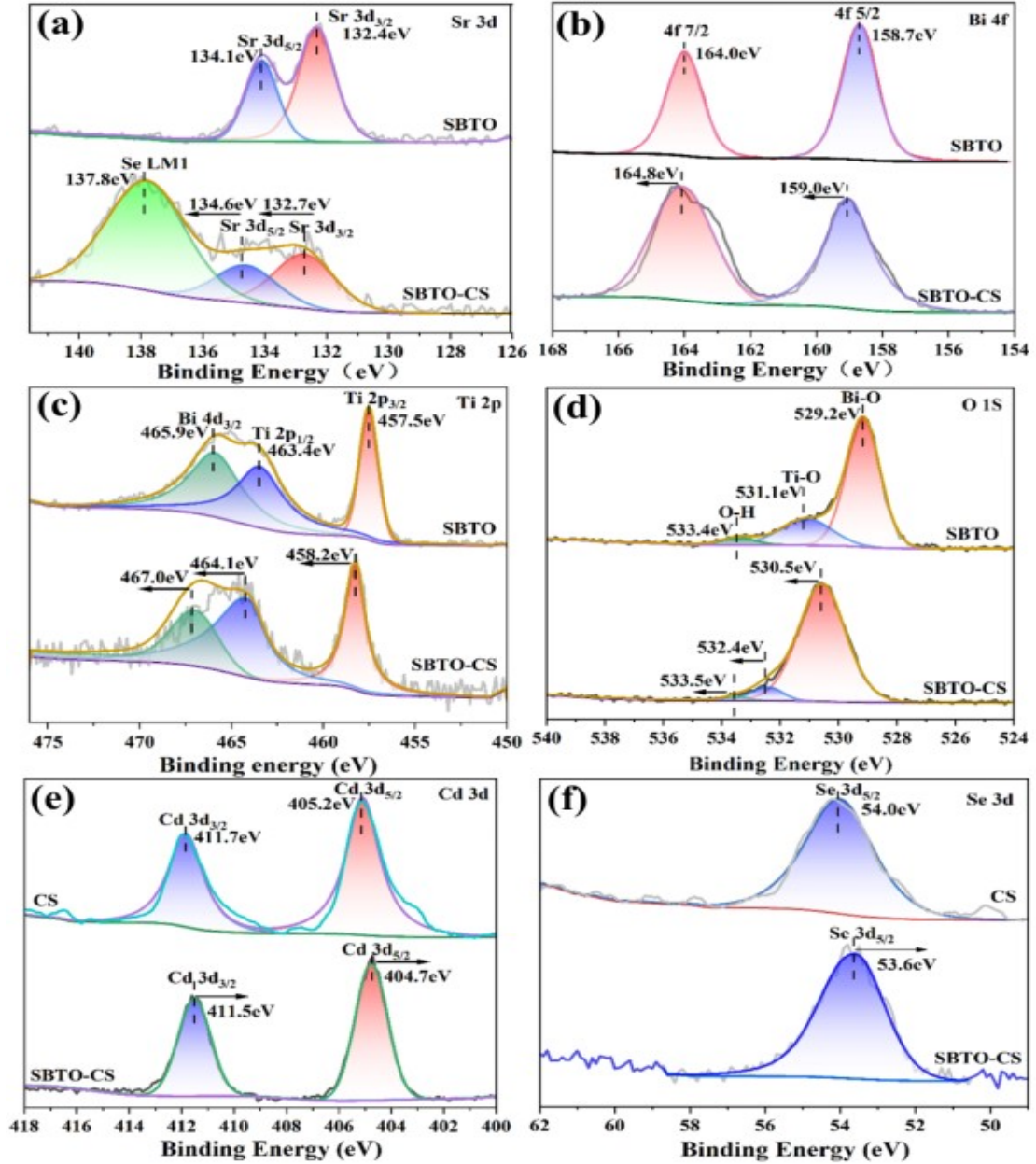
55 Fig. S4 (a-b) TEM images of CS QDs. (b) a higher magnification image of
 56 CS and the insert shows the particles size distribution.



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58 Fig. S5 EDS-mapping data of SBTO-CS-15 (Scanning for Sr, Bi, Ti, O,

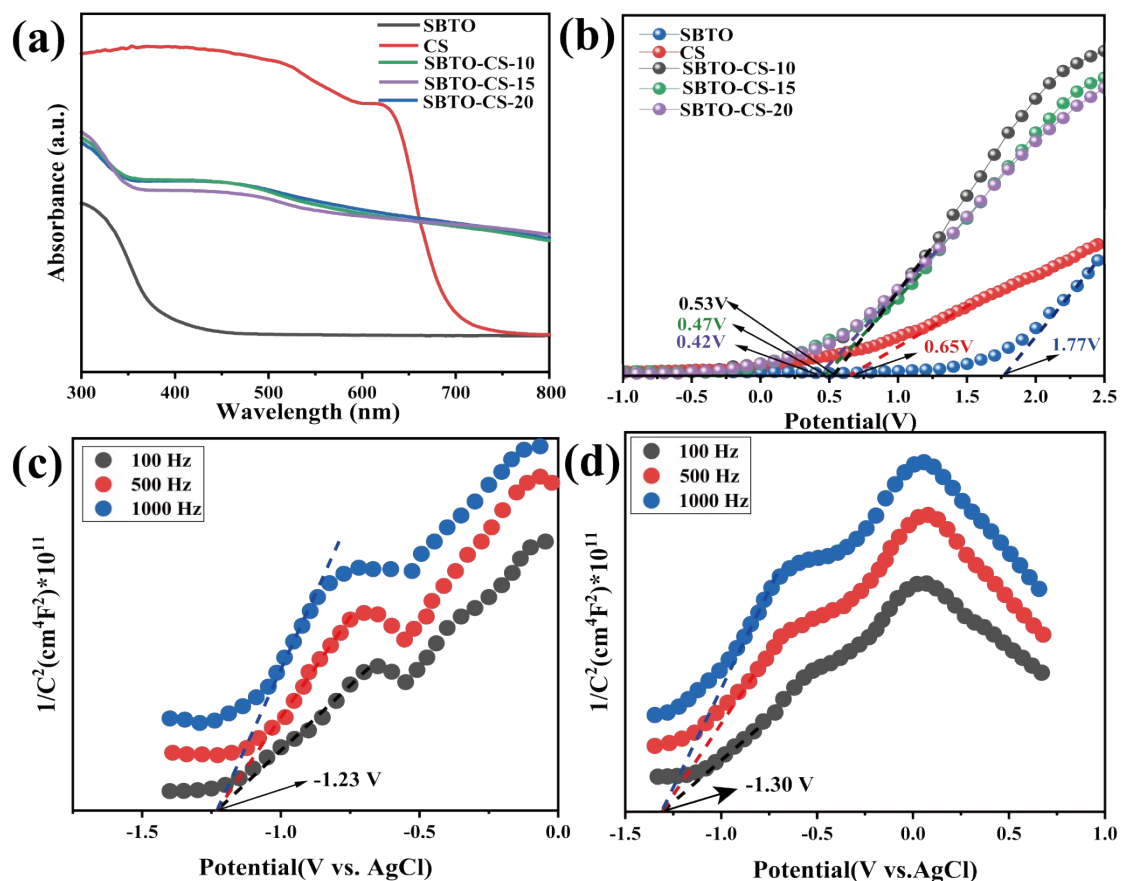
59 Cd, Se elements.)



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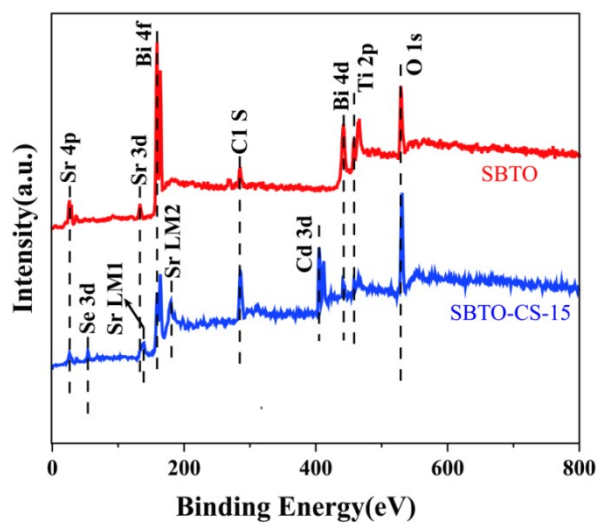
61 Fig. S6 XPS patterns of (a) Sr, (b) Bi, (c) Ti, (d) O, (e) Cd, and (f) Se
 62 elements collected from the pure SBTO, CS and SBTO-CS-15 samples
 63 (before reactions). The names of the samples are marked on these curves.

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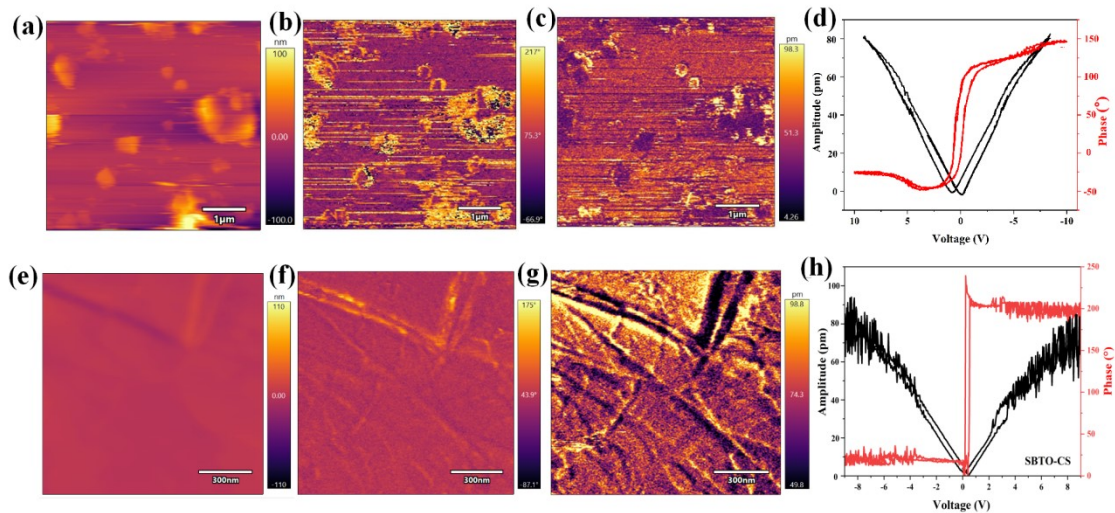
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66 Fig. S7 (a) the UV-Vis absorption spectra, (b) VB-XPS diagrams for
 67 SBTO, CS, SBTO-CS-10, 15, 20. (c-d) Mott Schottky curves of (c) SBTO-
 68 CS-10 and (d) SBTO-CS-20.



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70 Fig. S8 XPS spectra of two materials.



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72 Fig. S9 PFM images of (a–d) pristine SBTO and (e–h) SBTO-CS: (a, e)
 73 topography image; (b, f) phase image; (c, g) amplitude image and (d, f)
 74 amplitude butterfly loops (black) and phase hysteresis loops (red).

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References

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