

Supporting Information

Tin Doped PrBaFe₂O_{5+δ} Anode Material for Solid Oxide Fuel Cells

Guohui Dong, Chunyang Yang, Fei He, Yanmei Jiang, Chunlei Ren, Yun Gan, Myongjin Lee, and Xingjian Xue *

Department of Mechanical Engineering, University of South Carolina, Columbia, SC 29208, USA

Table S1. Element molar ratios of as-prepared PrBaFe_(2-x)Sn_xO_{5+δ} (x=0, 0.05, 0.1, 0.15, 0.2, 0.3) powders obtained from ICP–OES experiment and analysis.

X value of Sn	Nominal compositions	ICP-OES tested results			
		Tested compositions	Pr/Ba	Fe/Ba	Sn/Ba
0	PrBaFe ₂ O _{5+δ}	Pr _{0.97} Ba _{0.99} Fe _{2.00} O _{5+δ}	0.98	2.02	0
0.05	PrBaFe _{1.95} Sn _{0.05} O _{5+δ}	Pr _{0.99} Ba _{0.99} Fe _{1.93} Sn _{0.04} O _{5+δ}	1.00	1.95	0.04
0.1	PrBaFe _{1.9} Sn _{0.1} O _{5+δ}	Pr _{0.98} Ba _{0.99} Fe _{1.88} Sn _{0.09} O _{5+δ}	0.99	1.90	0.09
0.15	PrBaFe _{1.85} Sn _{0.15} O _{5+δ}	Pr _{0.99} Ba _{1.01} Fe _{1.86} Sn _{0.15} O _{5+δ}	0.98	1.84	0.15
0.2	PrBaFe _{1.8} Sn _{0.2} O _{5+δ}	Pr _{0.98} Ba _{0.98} Fe _{1.77} Sn _{0.20} O _{5+δ}	1.00	1.81	0.20
0.3	PrBaFe _{1.7} Sn _{0.3} O _{5+δ}	Pr _{0.99} Ba _{1.01} Fe _{1.74} Sn _{0.29} O _{5+δ}	0.98	1.72	0.29

Table S2. The oxygen content (5+δ) in the as-prepared PrBaFe_(2-x)Sn_xO_{5+δ} (x=0, 0.05, 0.1, 0.15, 0.2, 0.3) and after reducing treatment in humidified gas mixture of 10 % H₂ + 90% N₂ at 800 °C for 24h.

x value of Sn	Oxygen content (5+δ)	
	As-prepared	After reducing
0	5.7688	5.5853
0.05	5.8211	5.5571
0.1	5.8216	5.496
0.15	5.8386	5.4417
0.2	5.7605	5.4305
0.3	5.7458	5.3571

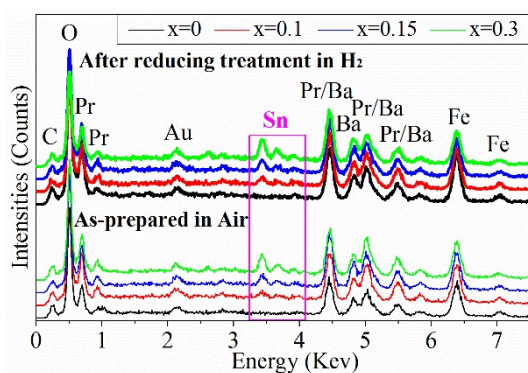


Figure S1. The EDS spectra of as-prepared $\text{PrBaFe}_{(2-x)}\text{Sn}_x\text{O}_{5+\delta}$ ($x=0, 0.1, 0.15, 0.3$) powders and after reducing treatment (humidified gas mixture of 10% H_2 + 90% N_2) at 800 °C for 24h.

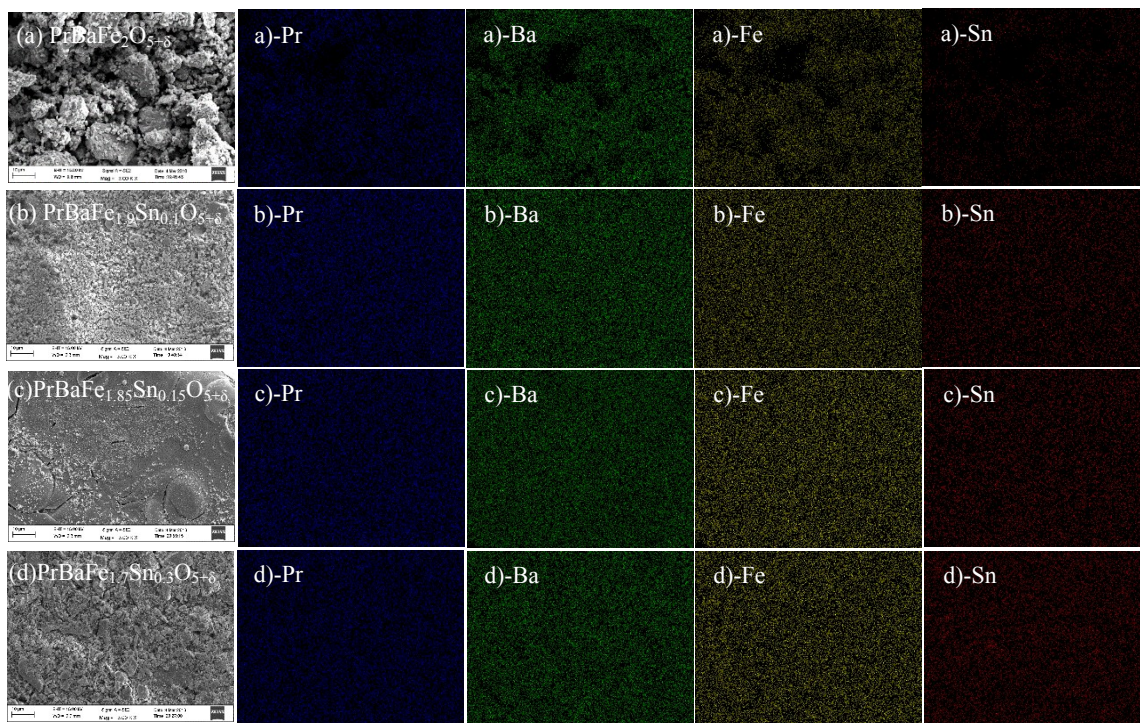


Figure S2. SEM images and EDS element mappings of as-prepared $\text{PrBaFe}_{(2-x)}\text{Sn}_x\text{O}_{5+\delta}$ ($x=0, 0.1, 0.15, 0.3$) powders. (a) $x=0$; (b) $x=0.1$; (c) $x=0.15$; (d) $x=0.3$. The scale length in all SEM images is 10 μm .

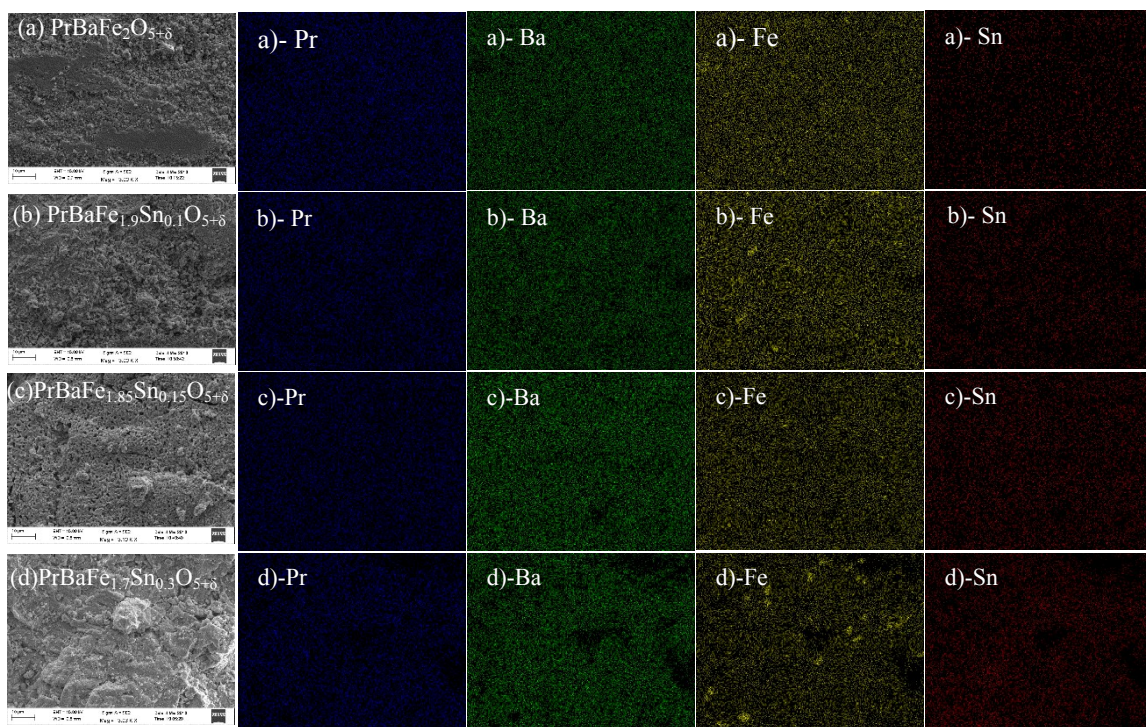


Figure S3. SEM images and EDS element mappings of $\text{PrBaFe}_{(2-x)}\text{Sn}_x\text{O}_{5+\delta}$ ($x=0, 0.1, 0.15, 0.3$) powders after reducing treatment at $800\text{ }^\circ\text{C}$ for 24 h in the humidified gas mixture of $10\% \text{H}_2 + 90\% \text{N}_2$. (a) $x=0$; (b) $x=0.1$; (c) $x=0.15$; (d) $x=0.3$. The scale length in all SEM images are $10\text{ }\mu\text{m}$.

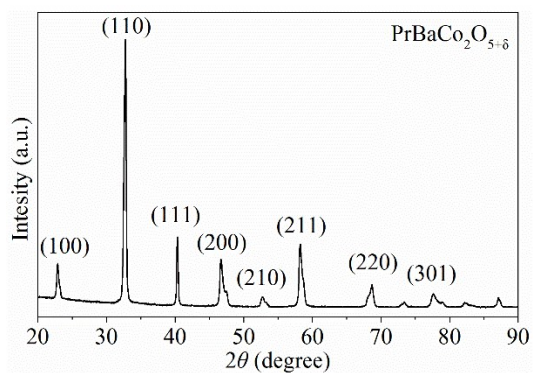


Figure S4. The XRD patterns of as-synthesized $\text{PrBaCo}_2\text{O}_{5+\delta}$ powder samples calcinated at $950\text{ }^\circ\text{C}$ in air for 5 h.

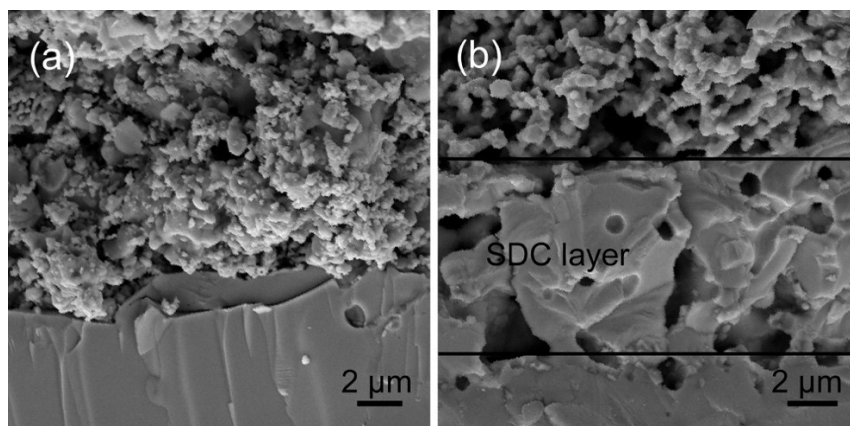


Figure S5. SEM micrographs of the single cell with configuration of $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ | LSGM | SDC | PBCO after testing: $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ anode (a) and SDC buffer layer and PBCO cathode (b).

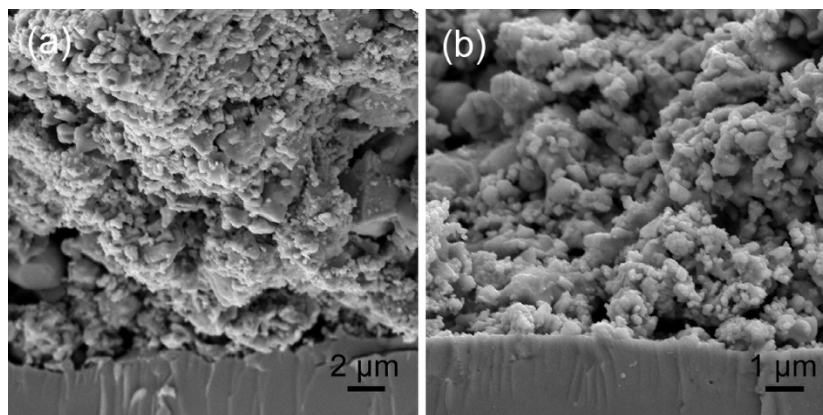


Figure S6. SEM micrographs of the symmetrical cell with configuration of $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ | LSGM | $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ after testing: $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ anode (a) and $\text{PrBaFe}_{1.9}\text{Sn}_{0.1}\text{O}_{5+\delta}$ cathode (a).