## SUPPORTING INFORMATION

## S-Acylation controls functional coupling of BK channel pore-forming α-subunits and β1-subunits

Peter J Duncan, Danlei Bi, Heather McClafferty, Lie Chen, Lijun Tian & Michael J Shipston

## LIST OF MATERIALS

Figure S1

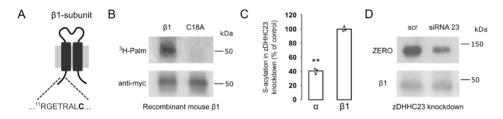


Figure S1: β1-subunit is S-acylated at Cys18 in HEK293 cells

(A) Schematic of BK channel  $\beta$ 1-subunit indicating cysteine 18 (Cys18) at the intracellular interface of the first transmembrane domain. (B) Metabolic incorporation of <sup>3</sup>[H]-Palmitate in wild-type  $\beta$ 1-subunit but not in C18A mutant  $\beta$ 1-subunits expressed in HEK293 cells. (C) siRNA knockdown of zDHHC23 reduces S-acylation of  $\alpha$ -subunit but not  $\beta$ 1-subunit in HEK293 cells. (D) Representative blot showing reduced incorporation of <sup>3</sup>[H]-Palmitate in  $\alpha$ -subunits, but not  $\beta$ 1-subunits, of zDHHC23 siRNA-treated cells (siRNA 23) compared to scrambled siRNA (scr). Data are Means ± SE (n = 3), \*\* p < 0.01 compared to scrambled siRNA control (t-test).