

SERCA1a	900 a.a.	ALSVLVTI E MCN
SERCA1b	900 a.a.	ALSVLVTI E MCN
SERCA1c	775 a.a.	ALSVLVTI E MCN
SERCA2a	899 a.a.	ALSVLVTI E MCN
SERCA2b	899 a.a.	ALSVLVTI E MCN
SERCA3a	900 a.a.	ALSVLVTI E MCN
SERCA3b	900 a.a.	ALSVLVTI E MCN
SERCA3c	900 a.a.	ALSVLVTI E MCN
SERCA3d	900 a.a.	ALSVLVTI E MCN
SERCA3e	900 a.a.	ALSVLVTI E MCN
SERCA3f	900 a.a.	ALSVLVTI E MCN

A

<i>Homo sapiens</i>	900 a.a.	ALSVLVTI E MCN
<i>Rattus norvegicus</i>	900 a.a.	ALSVLVTI E MCN
<i>Mus musculus</i>	900 a.a.	ALSVLVTI E MCN

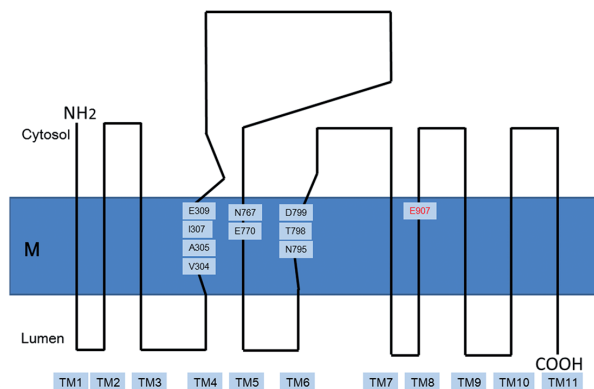


Fig. S1. Sequence alignments around the missense mutation, and the structure of SERCA2b. The sequence alignment of the SERCA family of *Homo sapiens* and the SERCA2b of mammals (A). Glu907 is in red. The glutamic acid is conserved among the SERCA family and diverse species. The structure of SERCA2b (B). There are 11 transmembrane domains (TM). Calcium-binding residues V304, A305, I307, E309 (TM4), N767, E770 (TM5), N795, T798, D799 (TM6) and E907 (TM8), located approximately in the middle of the TM. The novel mutation alters E907, indicated in red font. M indicates the membrane.