

Supporting information

Berbamine reduces chloroquine-induced itch in mice through inhibition of MrgprX1

Kunhi Ryu, Yunkyung Heo, Yechan Lee, Hyejin Jeon and Wan Namkung *

¹College of Pharmacy, Yonsei Institute of Pharmaceutical Sciences, Yonsei University, Incheon 21983, Korea.

Correspondence to: Wan Namkung, Ph.D. College of Pharmacy, Yonsei Institute of Pharmaceutical Sciences, Yonsei University, Incheon 21983, Korea; Tel: +82 32 749 4519; Fax: +82 32 749 4109; E-mail: wnamkung@yonsei.ac.kr

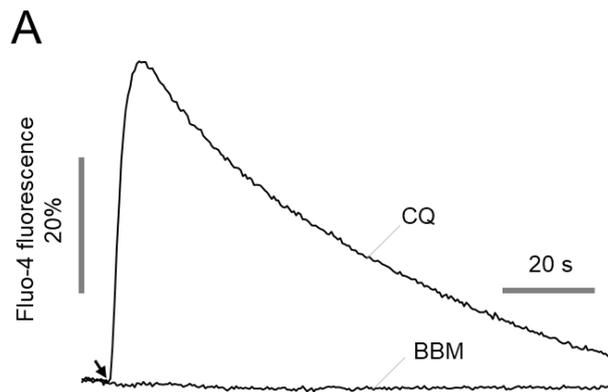


Figure S1. Effect of berbamine on MrgprX1 activation. (A) Intracellular calcium levels were measured using Fluo-4 NW in HEK293T cells expressing MrgprX1. The cells were treated with 30 μ M berbamine (BBM) or 1 mM of chloroquine (CQ).

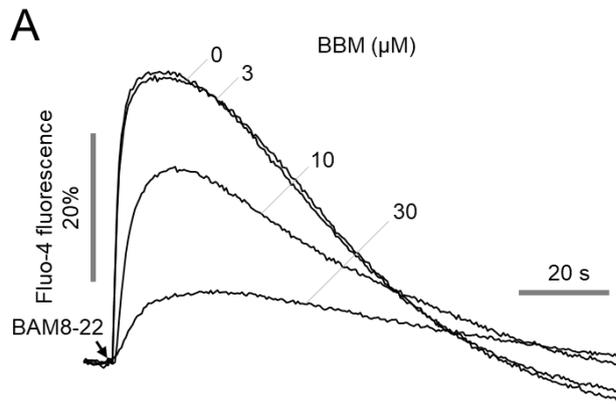


Figure S2. Effect of berbamine on BAM8-22 induced MrgprX1 activation. (A) Intracellular calcium levels were measured using Fluo-4 NW in HEK293T cells expressing MrgprX1. The cells were pretreated with indicated concentrations of berbamine (BBM) for 10 min prior to the treatment of 500 nM BAM8-22.

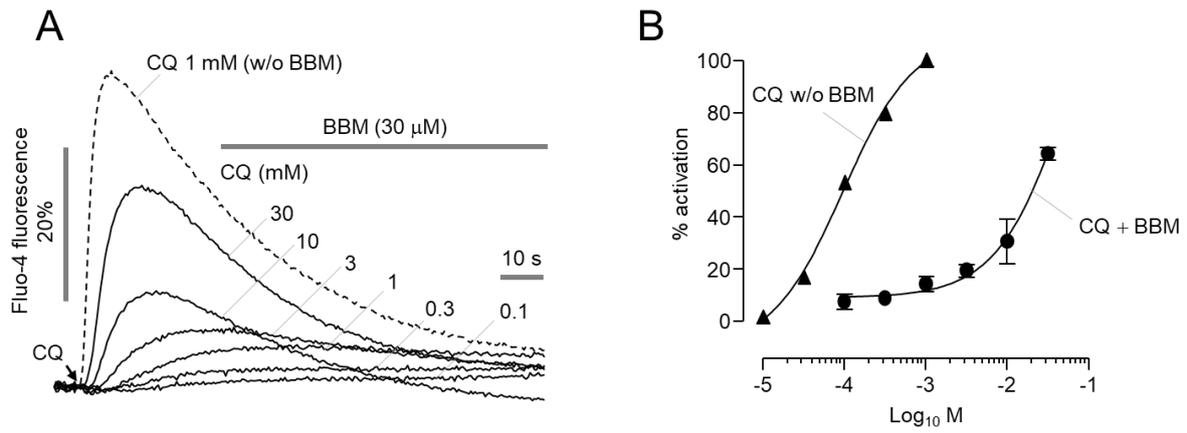


Figure S3. Dose response of CQ-induced MrgprX1 activation in the presence or absence of berbamine. (A) Intracellular calcium levels were measured using Fluo-4 NW in HEK293T cells expressing MrgprX1. The cells were pretreated with 30 μM of berbamine (BBM) and applied with indicated concentrations of CQ. (B) Summary of dose-response (mean \pm S.E., $n = 4$). The dose response data of CQ in the absence of BBM was obtained from Figure 1B.