

Fig. S1. Extra- and intracellular concentrations of K^+ at different temperatures and cooling rates. (A) Extracellular $[K^+]$ measured at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment. A significant interaction was found between cooling rate and temperature using a Two-way ANOVA. The means flanked by different letters within a temperature group are statistically different at $P < 0.05$. (B) Intracellular $[K^+]$ measured at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment.

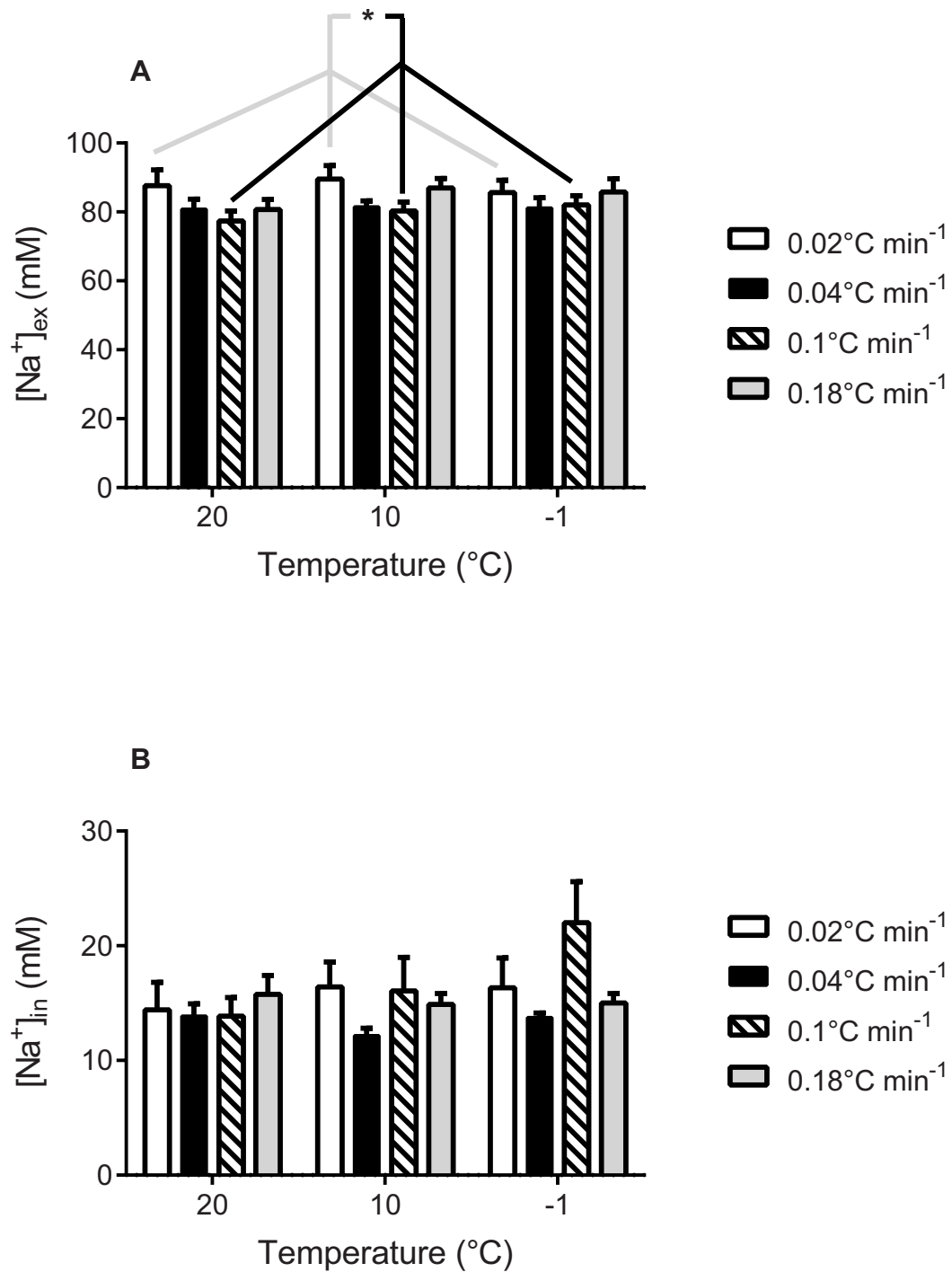


Fig. S2. Extra- and intracellular concentrations of Na⁺ at different temperatures and cooling rates. (A) Extracellular [Na⁺] measured at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment. A significant effect of cooling rate was found using a Two-Way ANOVA. Statistical differences ($P < 0.05$) between cooling rates are indicated by *. (B) Intracellular [Na⁺] measured at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment.

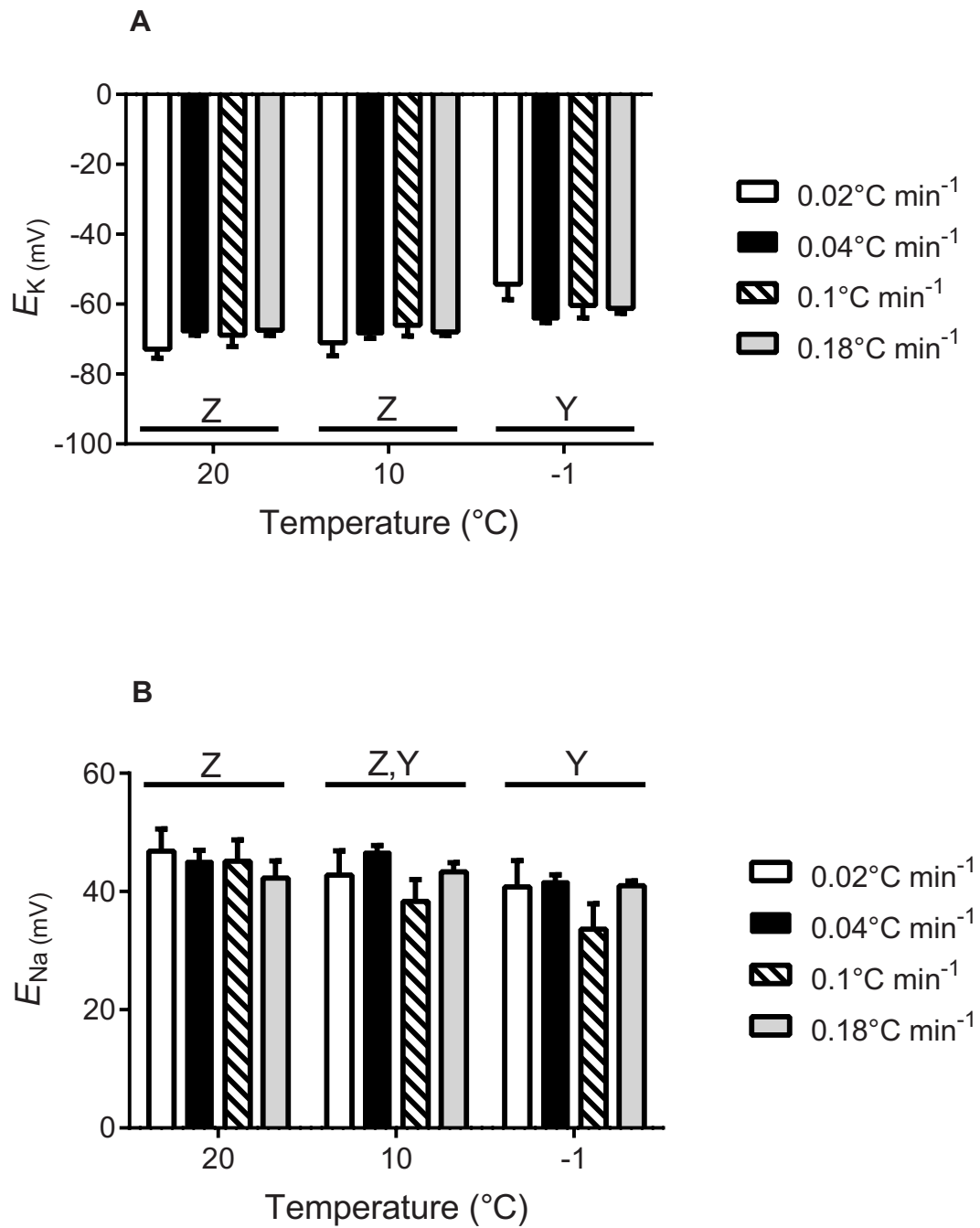


Fig. S3. Nernst potential for K^+ and Na^+ at different temperatures and cooling rates. (A) Nernst potential for K^+ (E_K) calculated from the intra- and extracellular $[\text{K}^+]$ at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment. A significant effect of temperature was found using a Two-way ANOVA. Dissimilar letter indicate groups that differ significantly ($P < 0.05$). (B) Nernst potential for Na^+ (E_{Na}) calculated from the intra- and extracellular $[\text{Na}^+]$ at three temperatures during different cooling rates. Bars represent means \pm s.e.m. of 8-10 samples in each treatment. A significant effect of temperature was found using a Two-way ANOVA. Dissimilar letter indicate groups that differ significantly ($P < 0.05$).