Figure EV1. Borneol has no effect on DRG neurons from TRPM8<sup>−/−</sup> mice. Averaged intracellular Ca<sup>2+</sup> increases in TRPM8<sup>−/−</sup> DRG neurons in response to consecutive applications of 200 μM borneol, 200 μM menthol, and 67 mM KCl. A total of 1,127 KCl-responsive neurons from four mice were included in the analysis, and none of the neurons responded to borneol.

Figure EV2. The borneol effect on mock-transfected HEK 293 cells and the menthol effect on TRPM8-expressing cells.

A Representative intracellular Ca<sup>2+</sup> signals in HEK 293 cells transfected with empty vector in response to borneol and subsequent applied Ca<sup>2+</sup> ionophore ionomycin (n = 6).

B Representative intracellular Ca<sup>2+</sup> signals in HEK 293 cells expressing hTRPM8 in response to different concentrations of menthol and the subsequently applied Ca<sup>2+</sup> ionophore ionomycin.
Figure EV3. Borneol activates TRPM8.

A Quantification of consecutively applied 100 μM menthol- and 600 μM borneol-induced hTRPM8 currents. Currents were normalized to 100 μM menthol-induced currents at +80 mV (n = 6).

B Representative intracellular Ca\textsuperscript{2+} signals in HEK 293 cells expressing mouse TRPM8 in response to different concentrations of borneol.

C Dose–response curves of borneol-induced increase in intracellular Ca\textsuperscript{2+} in mouse TRPM8-expressing HEK 293 cells. The smooth curve is a fit to the Hill equation with an EC\textsubscript{50} of 116 μM (n = 12). The data were normalized to ionomycin-induced intracellular Ca\textsuperscript{2+} increases.

D Time course of menthol- and subsequently applied borneol-induced whole-cell currents in mouse TRPM8-expressing HEK 293 cells (n = 5).

Data information: All the data are presented as the mean ± standard error of the mean (SEM).
Figure EV4. Intrathecal naloxone antagonizes menthol- or morphine-induced analgesia.

A Quantification of the effect of intrathecal injection of naloxone on menthol-induced analgesia in TRPM8−/− mice. After control saline or naloxone was intrathecally injected in TRPM8−/− mice, 15% menthol was applied to a hindpaw for a total of three times. After 10 min, 100 µM Cap was injected into the paw, and paw licking and lifting time was measured within 5 min.

B Quantification of 100 µM Cap-induced nociceptive responses in WT mice after intrathecal injection of morphine with or without naloxone.

Data information: The number of mice is indicated on top of each bar. Statistical significance was evaluated using two-tailed t-test. **P < 0.01; ***P < 0.001; the exact P-values are indicated in Appendix Table S1. All the data are presented as the mean ± standard error of the mean (SEM).

Figure EV5. Menthol causes cold hypersensitivity in TRPM8−/− mice.

Ethanol or menthol was applied to both hindpaws of TRPM8−/− mice, and the nociceptive response latencies were measured in a cold plate test (0°C). Statistical significance was evaluated using two-tailed t-test. ***P < 0.001; the exact P-value is indicated in Appendix Table S1. The number of mice is indicated on top of each bar. All the data are presented as the mean ± standard error of the mean (SEM).