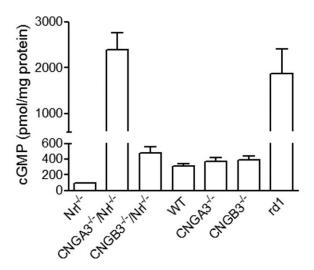
Supplemental Figures

Supplemental Figure 1. Elevation of retinal cGMP levels in $CNGA3^{-/-}/Nrl^{-/-}$ and $CNGB3^{-/-}/Nrl^{-/-}$ mice. ELISA (Assay Designs Inc., Ann Arbor, MI) was performed using light-adapted mouse retina (P30). Compared to the levels in $Nrl^{-/-}$ mice, $CNGA3^{-/-}/Nrl^{-/-}$ mice exhibited a remarkable elevation in retinal cGMP levels while $CNGB3^{-/-}/Nrl^{-/-}$ mice showed a moderate increase. The cGMP level was about 25-fold higher in $CNGA3^{-/-}/Nrl^{-/-}$ retina than that in $Nrl^{-/-}$ retina. No such increase was clearly detected in $CNGA3^{-/-}$ and $CNGB3^{-/-}$ mice, compared to the wild type (WT). This is likely because of the small proportion of cones in the rod-dominant retina. In addition, cGMP levels in the WT retina were significantly higher than the levels in $Nrl^{-/-}$ retinas. Retinas of rdl mice were included as an assay control. Data are represented as means \pm SEM of measurements from four independent experiments using retinas from 8-10 mice.



Supplemental Figure 2. Expression of CNGA3 in $CNGB3^{-/-}/Nrl^{-/-}$ retinas. Western blot was performed to detect retinal expression of CNGA3 in $CNGB3^{-/-}/Nrl^{-/-}$ and $Nrl^{-/-}$ mice (P30). Shown are representative images (*left panel*) and the correlating densitometric analysis (*right panel*). Actin was probed as a loading control. Data are represented as means \pm SEM of measurements from three independent experiments using retinas from 4-5 mice. Unpaired student's t test was used for determination of the significance (*, P < 0.05).

