Venus YFP was coelectroporated with control vector into the chick neural tube in ovo at stages 12–15. The electroporation procedure was essentially done as described previously (Liu et al., 2004). Embryos were isolated at stage 23 and examined for YFP fluorescence. The spinal cord showing green fluorescence was isolated as an open-book preparation and the whole mount immunostaining was carried by using the chicken commissural axon marker, anti-axonin-1 antibody. The merged image shows commissural axons project toward the floor plate with most of them arriving at or crossing over the floor plate. For more information, see the article by Liu et al. in this issue (957–968).
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Activity of Ventral Medial Thalamic Neurons during Absence Seizures and Modulation of Cortical Paroxysms by the Nigrothalamic Pathway
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Correction: The address for Jack Favor in the article “Molecular Interaction between Projection Neuron Precursors and Invading Interneurons via Stromal-Derived Factor 1 (CXCL12)/CXCR4 Signaling in the Cortical Subventricular Zone/Intermediate Zone” by Marie-Catherine Tiveron, Mireille Rossel, Barbara Moepps, Yong Li Zhang, Ralph Seidenfaden, Jack Favor, Norbert König, and Harold Cremer, which appears on pages 13273–13278 of the December 20, 2006 issue was listed incorrectly. The correct address for Jack Favor should have been listed as Institute of Human Genetics, GSF-National Center for Environment and Health, 85764, Neuherberg, Germany.

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