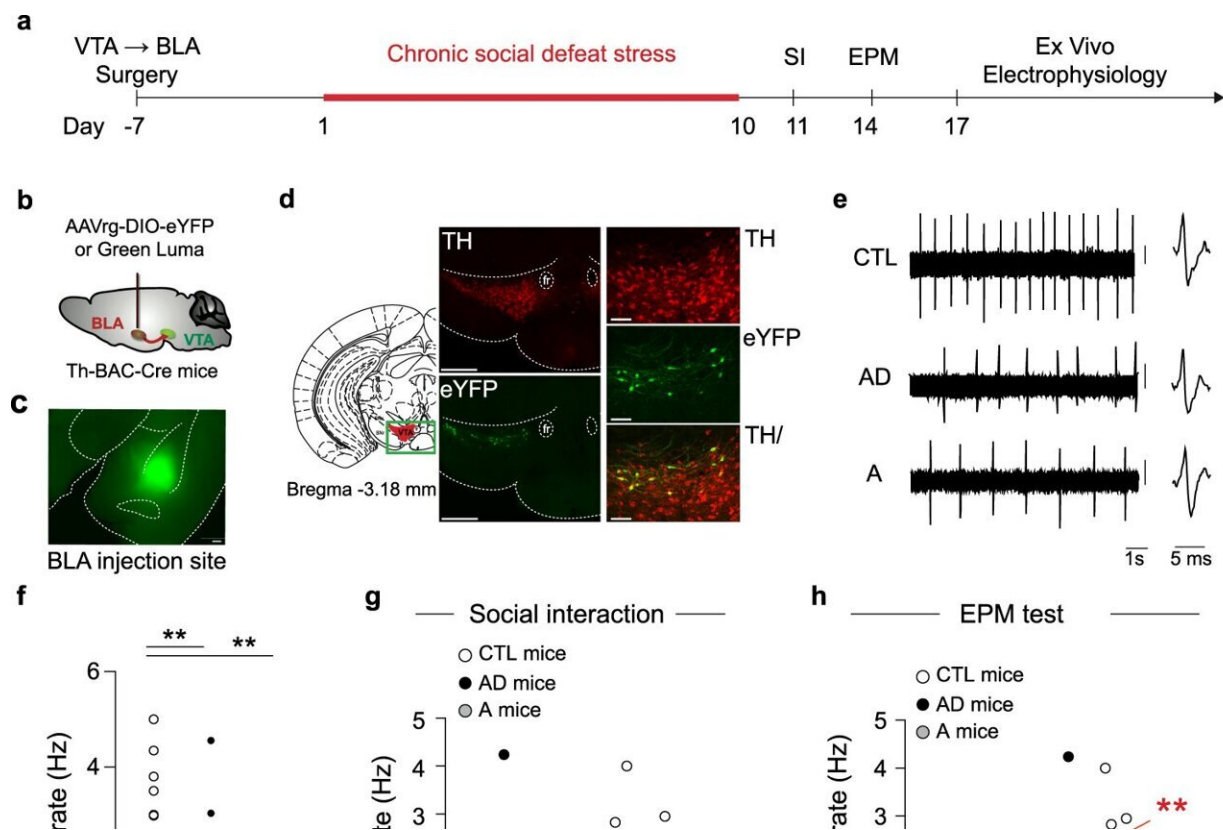


Midbrain projection to the basolateral amygdala encodes anxiety-like behaviors

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Anxiety-like behavior correlates with the hypoactivity of VTA → BLA dopamine neurons. **a** Experimental timeline. **b** Schematic of the brain surgery to dissect VTA → BLA circuit. **c** BLA surgery injection site (scale bar=500 μm). **d** Morphological validation showing the targeted VTA → BLA dopamine neurons in TH-BAC-Cre mice injected with AAVrg-DIO-eYFP (scale bar = 500 and 100 μm, representative images of the 23 recorded mice). **e** Sample traces of ex vivo cell-attached recordings from CTL, AD, and A mice (scale bar = 0.2 mV). **f** Spontaneous firing activity of VTA → BLA dopamine neurons in AD and A

mice compared to control mice (mean \pm s.e.m., ANOVA, $F_{(2, 104)} = 6.750$ $p = 0.0018$; post hoc test, $t = 3.48$ $p = 0.002$; $t = 3.50$ $p = 0.003$, $n = 30, 31, 45$ neurons, $n = 23$ combined C57BL6/J and TH-BAC-Cre mice injected with AAVrg-DIO-eYFP and Green Luma, respectively). g Pearson correlation analyses of VTA \rightarrow BLA dopamine neuron firing with the social interaction behavior after CSDS ($p = 0.59$, 3–7 neurons per mouse, $n = 23$ combined C57BL6/J and TH-BAC-Cre mice). h Pearson correlation analyses of VTA \rightarrow BLA dopamine neuron firing activity with the time in EPM open arms ($p = 0.0015$, 3–7 neurons per mouse, $n = 23$ combined C57BL6/J and TH-BAC-Cre mice). i Sample traces of ex vivo whole-cell recordings from CTL, AD, and A mice at a 40 pA step current injection. j VTA \rightarrow BLA dopamine neurons excitability in AD and A mice compared to CTL mice following incremental steps in currents injections (20–280 pA; mean \pm s.e.m., RM two-way ANOVA: group effect: $F_{(2, 33)} = 3.818$ $p = 0.021$; Interaction $F_{(28, 434)} = 3.164$ $p = 1.08e-07$; post hoc tests: $t = 2.41$ $p = 0.04$; $t = 2.53$ $p = 0.04$; $t = 1.95$ $p = 0.04$; $t = 2.63$ $p = 0.04$; $t = 1.64$ $p = 0.04$; $t = 2.52$ $p = 0.04$; $t = 1.72$ $p = 0.04$; $t = 2.25$ $p = 0.04$; $n = 11, 12, 14$ neurons/4, 5, 6 TH-BAC-Cre mice). k VTA \rightarrow BLA dopamine neurons rheobase in AD and A mice compared to CTL mice (mean \pm s.e.m., ANOVA: Group effect: $F_{(2, 33)} = 4.016$ $p = 0.013$; post hoc tests $t = 2.43$ $p = 0.04$; $t = 2.85$ $p = 0.02$; $n = 11, 13, 14$ neurons/4, 5, 6 TH-BAC-Cre mice). l VTA \rightarrow BLA dopamine neurons hyperpolarization-activated current, i.e., I_h current in AD and A mice compared to CTL mice following incremental voltage steps (mean \pm s.e.m., RM two-way ANOVA: group effect: $F_{(2, 33)} = 4.194$ $p = 0.017$; interaction $F_{(10, 175)} = 3.393$ $p = 9.7e-06$; post hoc tests $t = 2.22$ $p = 0.04$; $t = 2.71$ $p = 0.025$; $n = 11, 13, 14$ neurons/4, 5, 6 TH-BAC-Cre mice). m VTA \rightarrow BLA dopamine neurons sag ratio in AD and A mice compared to CTL mice (mean \pm s.e.m., ANOVA: group effect: $F_{(2, 32)} = 7.225$ $p = 0.001$; $t = 3.04$ $p = 0.009$; $t = 3.79$ $p = 0.002$, $n = 11, 13, 14$ neurons/4–6 TH-BAC-Cre mice). In all panels, two-sided statistical analyses post hoc corrected tests were performed, * p

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