



Dorsal Hippocampus ERK2 Signaling Mediates Anxiolytic-Related Behavior in Male Rats

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Abstract

Background: Anxiety disorders are the most common neuropathologies worldwide, but the precise neuronal mechanisms that underlie these disorders remain unknown. The hippocampus plays a role in mediating anxiety-related responses, which can be modeled in rodents using behavioral assays, such as the elevated plus maze. Yet, the molecular markers that underlie affect-related behavior on the elevated plus maze are not well understood.

Methods: We used herpes simplex virus vector delivery to overexpress extracellular signal-regulated kinase-2, a signaling molecule known to be involved in depression and anxiety, within the dorsal hippocampus of adult Sprague-Dawley male rats. Three days post virus delivery, we assessed anxiety-like responses on the elevated plus maze or general locomotor activity on the open field test.

Results: When compared to controls, rats overexpressing extracellular signal-regulated kinase-2 in the dorsal hippocampus displayed an anxiolytic-like phenotype, per increases in time spent in the open arms, and less time in the closed arms, of the elevated plus maze. Furthermore, no changes in locomotor activity as a function of virus infusion were observed on the open field test between the experimental groups.

Conclusion: This investigation demonstrates that virus-mediated increases of extracellular signal-regulated kinase-2 signaling, within the hippocampus, plays a critical role in decreasing anxiogenic responses on the rat elevated plus maze. As such, our data provide construct validity, at least in part, to the molecular mechanisms that mediate anxiolytic-like behavior in rodent models for the study of anxiety.

Keywords

anxiety, construct validity, ERK, elevated plus maze, MAPK1

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Introduction

Anxiety disorders constitute the most common forms of psychiatric illnesses worldwide.^{1,2} These disorders are associated with increased health-care costs and exhibit high comorbidity with other mental diseases, such as depression and posttraumatic stress disorder.^{3,4} Despite their high prevalence, the specific neurobiological mechanisms underlying these affect-related neuropathologies remain to be fully understood.

Members of the mitogen-activated protein kinases (MAPKs) function by integrating extracellular signals received by membrane receptors and transferring them through a phosphorylation cascade into the nucleus,

thus regulating gene expression.⁵ Although they are ubiquitously expressed, MAPKs have been found to play critical functions in the mammalian brain, ranging from synaptic plasticity, long-term potentiation, and

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neuronal differentiation.⁶ Within the MAPK superfamily, the actions of members of the subfamily of extracellular signal-regulated kinase (ERK), particularly the ERK1 and ERK2 isoforms, have been extensively characterized in the central nervous system, and it is becoming increasingly clear that they play a prominent role in several psychiatric diseases, including depression and anxiety disorders.^{7–10}

Preclinical pharmacological experiments have been useful in establishing a link between anxiety-related behavior and the ERK signaling pathway.^{11–13} However, despite significant efforts, there are currently no pharmacological agents that directly target individual and/or specific ERK isoforms for activation or inhibition,^{14,15} which in turn, limit our understanding on the role that each MAPK isoform plays in the regulation of mood-related behavior. Addressing this issue, we have previously shown that the upregulation of ERK2, via viral vector delivery in the dorsal hippocampus, reduces despair-related behavior in male rats, as determined in the forced swim test.¹⁶ Conversely, stress-induced decreases of ERK-related signaling within this brain region has been linked to enhanced anxiety-like behavior,¹⁷ which can be successfully studied in rodents using the elevated plus maze (EPM)—a behavioral paradigm with high face and predictive validity.^{18–20} Yet, to date, the precise role that ERK2 plays in the regulation of anxiety-related behavior has not been clearly delineated. Therefore, in order to determine if the upregulation of hippocampal ERK2 signaling is involved in the modulation of anxiety-like behavior, we combined a targeted molecular approach of ERK2 overexpression in the rat dorsal hippocampus, with behavioral assessment using the EPM and open field tests.

Materials and Methods

Animals

Male Sprague-Dawley rats (250–275 g) were obtained from Charles River Laboratories (Hollister, CA). Rats were maintained in a humidity- and temperature-controlled animal facility with a 12-h light/dark cycle and were subjected to a one-week acclimation period before surgery and the start of experiments. Rats were housed in pairs in standard laboratory polycarbonate cages (Ancare, model R20, Bellmore, NY) along with sani-chip bedding (P.J. Murphey San-Chips®, Murville, NJ) and had access to food and water *ad libitum*. Animals were cared for in accordance with the National Institutes of Health *Guide for the Care and Use of Laboratory Animals*²¹ and with approval from the Institutional Animal Care and Use Committee.

Virus Vectors

Overexpression of ERK2 was achieved by microinjecting herpes simplex virus (HSV) vectors encoding green fluorescent protein (GFP)-alone or a GFP-wild-type ERK2 (wtERK2) in the dorsal hippocampus. Virus was packaged in a p1005 plasmid with expression of the target gene (i.e., ERK2) driven by an HSV IE 4/5 promoter, while GFP expression was driven by a CMV promoter.²² All viruses were diluted in phosphate-buffered saline (PBS) + 10% sucrose and 20 mM HEPES (pH 7.3). For further specifics on vector design, construction, neurotropic effects, as well as *in vitro* and *in vivo* validation, see previously published work.^{23,24} Highest levels of transgene expression by the HSV vectors were reached three days after surgery. Therefore, behavioral experiments began after this time period (see Figure 1(a) for experimental design).^{25,26} Transgene expression was restricted to an area of ~1 mm² surrounding injection site, as previously demonstrated.^{27,28} Behavioral tests (described below) were recorded using a video tracking system (Ethovision XT, Noldus, Leesburg, VA).

Animal Surgery

Stereotaxic surgeries were conducted for selective delivery of the HSV vectors. Rats were anesthetized with an intramuscular injection of ketamine/xylazine cocktail (80/10 mg/kg), with subcutaneous atropine (0.25 mg/kg) administered to reduce bronchial secretions; bilateral microinjections of either GFP or GFP-wtERK2 vectors were then administered (1 µl per hemisphere over a 10-min period) into the dorsal hippocampus (coordinates from Bregma: anteroposterior: –3.8, lateral: –2, dorsoventral: –3.2 mm below dura) using a 32-gauge Hamilton syringe.^{16,29} Postsurgical discomfort was minimized by applying the local anesthetic bupivacaine.

Elevated Plus Maze

The EPM was used to evaluate anxiety-like behavior.³⁰ The maze consisted of two perpendicular, intersecting runways (12 cm wide × 100 cm long): one with tall, closed arms (30 cm), and the other with open arms, connected by a central area. The maze was elevated 1 m from the floor in a behavioral testing room under controlled lighting (~90 lux) conditions. Rats were placed initially in the central area, facing one of the open arms, and grooming counts, as well as the cumulative time spent in the closed and open arms was recorded for 5 min.³¹

Locomotor Activity

Locomotor activity of GFP-alone or GFP-wtERK2 rats was assessed as distance traveled (cm) in an open field apparatus (63 × 63 × 26 cm) for 5 min.

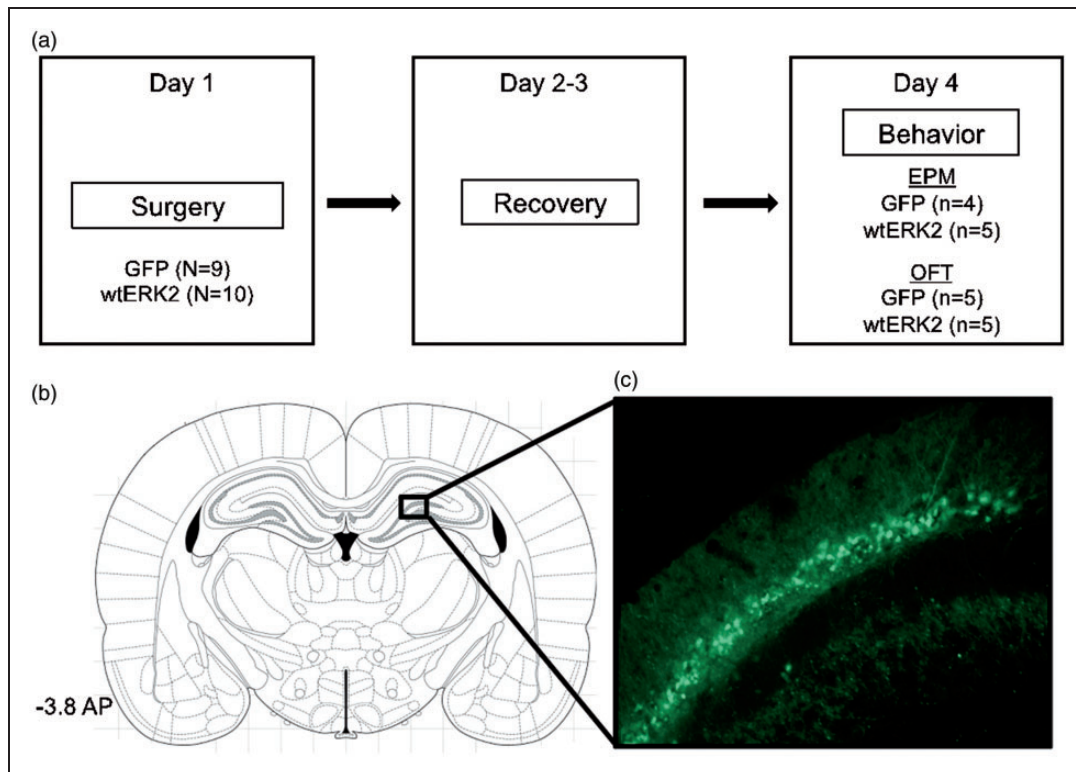


Figure 1. Experimental design and virus-mediated overexpression of ERK2 in the rat dorsal hippocampus. (a) Male rats were infused with HSV vectors overexpressing GFP-alone or a GFP-wtERK2 on day 1 (surgery) and allowed to rest for 48 h (days 2–3). On day 4, rats were tested on the EPM or the OFT. (b) Region of the hippocampus to which microinjections of HSV vectors was targeted (AP: -3.8 , lateral: -2 , dorsoventral: -3.2 mm below dura). Adapted from *The Rat Brain in Stereotaxic Coordinates*.⁵³ (c) Cells expressing GFP-wtERK2 (green, cyanine 2) fluorescence (magnification, $\times 400$). GFP: green fluorescent protein; EPM: elevated plus maze; OFT: open field test; AP: anteroposterior; wtERK2: wild-type extracellular signal-regulated kinase-2.

Histology and Transgene Detection

The sites of HSV vector injection were confirmed in all rats. Animals were euthanized with an overdose of sodium pentobarbital administered 1 h after behavioral assessment, followed by transcardial perfusion with 0.9% saline and 4% paraformaldehyde (PFA). Brains were excised and postfixed by overnight immersion in 4% PFA and stored in 20% glycerol. Hippocampal coronal sections ($45\ \mu\text{m}$) were obtained on a microtome and stored in 0.1 M sodium phosphate buffer containing 0.05% sodium azide. Tissue sections were used to examine GFP expression within the hippocampus as previously described.³² Hippocampal free-floating coronal sections were blocked in 3% normal donkey serum (NDS), followed by overnight incubation with primary antibody (rabbit anti-GFP antibody, 1:1000; Abcam, Cambridge, MA) along with 0.3% Triton X-100 and 1% NDS. Tissue was then incubated with anti-rabbit secondary antibody (1:1000; Jackson ImmunoResearch, West Grove, PA) for 2 h at room temperature. Immunostained sections were mounted on microscope slides, dehydrated in ethanol and Citrosolv, and coverslipped with DPX (Sigma, St. Louis, MO).

Slides were visualized and photographed using a confocal fluorescence microscope equipped with a digital camera.

Statistical Analysis

Rats were randomly assigned to receive either HSV-GFP-alone or HSV-GFP-wtERK2 vectors, three days prior to behavioral testing. Statistical testing was carried out using two-tailed Student's *t* test. Data are expressed as mean \pm standard error of the mean. A value of $p < 0.05$ was adopted to determine statistical significance.

Results

Figure 1(b) depicts the hippocampal region targeted for microinjection delivery of HSV vectors. As observed in Figure 1(c), immunofluorescence confocal microscopy revealed that ERK2 was successfully overexpressed in the dorsal hippocampus as previously shown.^{16,27}

Using the EPM test to determine anxiety-like behavioral responses to GFP-wtERK2 overexpression in the dorsal hippocampus, we found that, when compared to

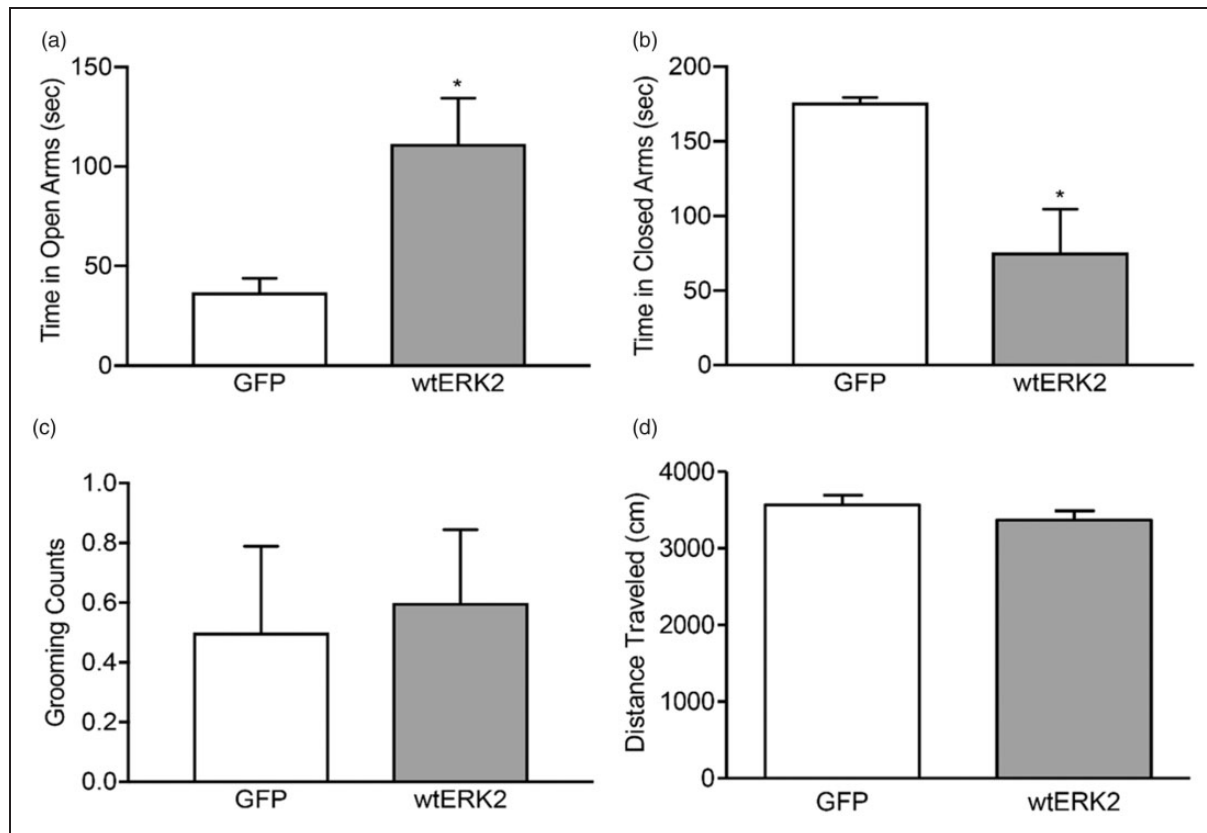


Figure 2. Herpes simplex virus overexpression of ERK2 (wtERK2) mediates an anxiolytic-like effect in the elevated plus maze. When compared to controls (GFP, $n = 4$), male rats overexpressing wtERK2 ($n = 5$) in the dorsal hippocampus spent higher time in the open arms (a), along with decreased time in the closed arms (b), of the maze. No differences in grooming counts (c) or distance traveled (d, $n = 5$ per group) were noted between the experimental groups. * $p < 0.05$. GFP: green fluorescent protein; wtERK2: wild-type extracellular signal-regulated kinase-2.

their GFP-only counterparts ($n = 4$), GFP-wtERK2 ($n = 5$) rats spent significantly more time in the open arms ($t_7 = 2.78$, $p < 0.05$; Figure 2(a)) and less time in the closed arms ($t_7 = 3.03$, $p < 0.05$; Figure 2(b)) of the EPM. No changes were observed in grooming counts ($t_7 = 0.26$, $p > 0.05$; Figure 2(c)). Lastly, Figure 2(d) shows that hippocampal overexpression of ERK2 ($n = 5$) did not influence locomotor activity as measured in the open field test when compared to GFP-only rats ($n = 5$; $t_8 = 1.12$, $p > 0.05$).

Discussion

In this study, we aimed to determine the behavioral effect of viral vector-mediated ERK2 overexpression in the dorsal hippocampus of adult male Sprague-Dawley rats, using the EPM as a tool to assess anxiety-related behavior.³⁰ Our results indicate that rats overexpressing GFP-wtERK2 in the dorsal hippocampus spent more time in the open arms and less time in the closed arms of the EPM, without overall changes in locomotor activity. These behavioral responses are consistent with an

anxiolytic effect,¹⁸ a behavioral outcome that is commonly reported after acute benzodiazepine (BDZ) exposure,^{33,34} or chronic administration of selective serotonin reuptake inhibitors (SSRIs).^{31,35} Of note, these two classes of drugs, which are used clinically to treat anxiety-related disorders,³⁶ have been reported to mediate their therapeutic effects via neuronal ERK signaling regulation. Specifically, treatment with the BDZ drug diazepam has been shown to increase the levels of ERK1/2 in the rat hippocampus,³⁷ and the administration of the SSRI fluoxetine similarly enhances ERK1/2 signaling across different brain regions,³⁸ including the hippocampus³⁹; brain circuitry that is activated during both stress and anxiety responses.⁴⁰ Likewise, activation of the ERK pathway via treatment with brain-derived neurotrophic factor results in elevated levels of activated ERK2 as well as reduced anxiety-like behavior in mice.⁴¹ Conversely, pharmacological inhibition of hippocampal ERK signaling in rats has been shown to induce anxiety-like behavior in the EPM.¹¹ However, we must note that the role of ERKs, as a molecular link between stress and anxiety, is equivocal across the literature.^{42–45}

While previous work has suggested that BZD and SSRI medications mediate their therapeutic effects via increases in hippocampal ERK1/2 signaling, our experimental approach confirms that the upregulation of ERK2 specifically, in the stress-naïve rat's dorsal hippocampus, is a contributor in the modulation of anxiety (Figure 2) and despair-like behavior.¹⁶ Importantly, we do so by using a genetically targeted approach, providing construct and mechanistic validity to the male rat EPM test. Furthermore, our findings expand on recent work indicating that the ventral hippocampus also modulates behavioral responses in the EPM^{46,47}—however, whether anxiolytic behavior in this paradigm is ERK2-dependent, within the ventral portion of the hippocampus, remains to be explored. Unfortunately, a limitation of the current investigation is that we did not include female rats in our experimental design, reducing the interpretability of our findings to the clinical setting, where anxiety-related disorders are more frequently diagnosed in the female, versus male, population.⁴⁸ Lastly, given that ERK1 and ERK2 share over 80% homology, it will be important to evaluate how selective regulation of ERK1 influences responses on the EPM, since ERK1-alone has been found to modulate behavior differentially in other tasks.^{49–51}

Conclusion

The neurobiological correlates between stress and anxiety are not well understood. Previous animal and human postmortem studies have shown that ERKs, across different brain regions,^{10,16,23,52} play a crucial role in modulating affect-related responses. Here, we expand this work to include a role for ERK2 specifically, within the dorsal hippocampus of rats, in mediating anxiolytic behavior when assessed on the EPM—a classic model used to screen for anxiolytic or anxiogenic compounds. Indeed, virus-mediated increases of ERK2 resulted in animals spending more time in the open arms, and lower time in the closed arms, of the EPM; an anxiolytic response mimicking the effects of traditional pharmaceutical agents used for the management of anxiety (i.e., SSRIs and BDZs). As such, our data may prove insight for the development of pharmaceutical agents, or alternative strategies, to treat/prevent the development of anxiety-related disorders.

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Declaration of Conflicting Interests

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References

1. Canuto A, Weber K, Baertschi M, et al. Anxiety disorders in old age: psychiatric comorbidities, quality of life, and prevalence according to age, gender, and country. *Am J Geriatr Psychiatry*. 2018; 26: 174–185.
2. Kessler RC, Berglund P, Demler O, Jin R, Merikangas KR, Walters EE. Lifetime prevalence and age-of-onset distributions of DSM-IV disorders in the National Comorbidity Survey Replication. *Arch Gen Psychiatry*. 2005; 62: 593–602.
3. Tiller JW. Depression and anxiety. *Med J Aust*. 2013; 199: S28–S31.
4. Bandelow B, Michaelis S. Epidemiology of anxiety disorders in the 21st century. *Dialog Clin Neurosci*. 2015; 17: 327–335.
5. Pearson G, Robinson F, Beers Gibson T, et al. Mitogen-activated protein (MAP) kinase pathways: regulation and physiological functions. *Endocr Rev*. 2001; 22: 153–183.
6. Mao LM, Wang JQ. Synaptically localized mitogen-activated protein kinases: local substrates and regulation. *Mol Neurobiol*. 2016; 53: 6309–6315.
7. Dwivedi Y, Rizavi HS, Roberts RC, Conley RC, Tamminga CA, Pandey GN. Reduced activation and expression of ERK1/2 MAP kinase in the post-mortem brain of depressed suicide subjects. *J Neurochem*. 2001; 77: 916–928.
8. Wang JQ, Fibuch EE, Mao L. Regulation of mitogen-activated protein kinases by glutamate receptors. *J Neurochem*. 2007; 100: 1–11.
9. Wang JQ, Mao L. The ERK pathway: molecular mechanisms and treatment of depression. *Mol Neurobiol*. 2019; 56: 6197–6205.

10. Duric V, Banasr M, Licznernski P, et al. A negative regulator of MAP kinase causes depressive behavior. *Nat Med*. 2010; 16: 1328–1332.
11. Qi X, Lin W, Wang D, Pan Y, Wang W, Sun M. A role for the extracellular signal-regulated kinase signal pathway in depressive-like behavior. *Behav Brain Res*. 2009; 199: 203–209.
12. Al Rahim M, Rimando AM, Silistrelis K, El-Alfy AT. Anxiolytic action of pterostilbene: involvement of hippocampal ERK phosphorylation. *Planta Med*. 2013; 79: 723–730.
13. Anchan D, Clark S, Pollard K, Vasudevan N. GPR30 activation decreases anxiety in the open field test but not in the elevated plus maze test in female mice. *Brain Behav*. 2014; 4: 51–59.
14. Di Benedetto B, Wefers B, Wurst W, Kuhn R. Local knockdown of ERK2 in the adult mouse brain via adeno-associated virus-mediated RNA interference. *Mol Biotechnol*. 2009; 41: 263–269.
15. Shin M, Franks CE, Hsu KL. Isoform-selective activity-based profiling of ERK signaling. *Chem Sci*. 2018; 9: 2419–2431.
16. Iñiguez SD, Parise LF, Lobo MK, et al. Upregulation of hippocampal extracellular signal-regulated kinase (ERK)-2 induces antidepressant-like behavior in the rat forced swim test. *Behav Neurosci*. 2019; 133: 225–231.
17. Kim EJ, Pellman B, Kim JJ. Stress effects on the hippocampus: a critical review. *Learn Mem*. 2015; 22: 411–416.
18. Walf AA, Frye CA. The use of the elevated plus maze as an assay of anxiety-related behavior in rodents. *Nat Protoc*. 2007; 2: 322–328.
19. Barkus C, McHugh SB, Sprengel R, Seeburg PH, Rawlins JN, Bannerman DM. Hippocampal NMDA receptors and anxiety: at the interface between cognition and emotion. *Eur J Pharmacol*. 2010; 626: 49–56.
20. Cominski TP, Jiao X, Catuzzi JE, Stewart AL, Pang KC. The role of the hippocampus in avoidance learning and anxiety vulnerability. *Front Behav Neurosci*. 2014; 8: 273.
21. Council NR. *Guidelines for the Care and Use of Mammals in Neuroscience and Behavioral Research*. Washington, DC: National Academy Press; 2003.
22. Neve RL, Neve KA, Nestler EJ, Carlezon WA Jr. Use of herpes virus amplicon vectors to study brain disorders. *BioTechniques*. 2005; 39: 381–391.
23. Iñiguez SD, Vialou V, Warren BL, et al. Extracellular signal-regulated kinase-2 within the ventral tegmental area regulates responses to stress. *J Neurosci*. 2010; 30: 7652–7663.
24. Neve RL, Howe JR, Hong S, Kalb RG. Introduction of the glutamate receptor subunit 1 into motor neurons in vitro and in vivo using a recombinant herpes simplex virus. *Neuroscience*. 1997; 79: 435–447.
25. Barrot M, Olivier JD, Perrotti LI, et al. CREB activity in the nucleus accumbens shell controls gating of behavioral responses to emotional stimuli. *Proc Natl Acad Sci*. 2002; 99: 11435–11440.
26. Carlezon WA Jr, Nestler EJ, Neve RL. Herpes simplex virus-mediated gene transfer as a tool for neuropsychiatric research. *Crit Rev Neurobiol*. 2000; 14: 47–67.
27. Iñiguez SD, Alcantara LF, Warren BL, et al. Fluoxetine exposure during adolescence alters responses to aversive stimuli in adulthood. *J Neurosci*. 2014; 34: 1007–1021.
28. Iñiguez SD, Warren BL, Neve RL, Nestler EJ, Russo SJ, Bolaños-Guzmán CA. Insulin receptor substrate-2 in the ventral tegmental area regulates behavioral responses to cocaine. *Behav Neurosci*. 2008; 122: 1172–1177.
29. Chen AC, Shirayama Y, Shin KH, Neve RL, Duman RS. Expression of the cAMP response element binding protein (CREB) in hippocampus produces an antidepressant effect. *Biol Psychiatry*. 2001; 49: 753–762.
30. Montgomery KC. The relation between fear induced by novel stimulation and exploratory behavior. *J Comp Physiol Psychol*. 1955; 48: 254–260.
31. Iñiguez SD, Warren BL, Bolaños-Guzmán CA. Short- and long-term functional consequences of fluoxetine exposure during adolescence in male rats. *Biol Psychiatry*. 2010; 67: 1057–1066.
32. Warren BL, Iñiguez SD, Alcantara LF, et al. Juvenile administration of concomitant methylphenidate and fluoxetine alters behavioral reactivity to reward- and mood-related stimuli and disrupts ventral tegmental area gene expression in adulthood. *J Neurosci*. 2011; 31: 10347–10358.
33. Rex A, Stephens DN, Fink H. “Anxiolytic” action of diazepam and abecarnil in a modified open field test. *Pharmacol Biochem Behav*. 1996; 53: 1005–1011.
34. Barbalho CA, Nunes-de-Souza RL, Canto-de-Souza A. Similar anxiolytic-like effects following intra-amygdala infusions of benzodiazepine receptor agonist and antagonist: evidence for the release of an endogenous benzodiazepine inverse agonist in mice exposed to elevated plus-maze test. *Brain Res*. 2009; 1267: 65–76.
35. Farhan M, Haleem DJ. Anxiolytic profile of fluoxetine as monitored following repeated administration in animal rat model of chronic mild stress. *Saudi Pharm J*. 2016; 24: 571–578.
36. Sartori SB, Singewald N. Novel pharmacological targets in drug development for the treatment of anxiety and anxiety-related disorders. *Pharmacol Ther*. 2019; 204: 107402.
37. Sevastre-Berghian AC, Fagarasan V, Toma VA, et al. Curcumin reverses the diazepam-induced cognitive impairment by modulation of oxidative stress and ERK 1/2/NF-kappaB pathway in brain. *Oxid Med Cell Longev*. 2017; 2017: 3037876.
38. Di Benedetto B, Radecke J, Schmidt MV, Rupprecht R. Acute antidepressant treatment differently modulates ERK/MAPK activation in neurons and astrocytes of the adult mouse prefrontal cortex. *Neuroscience*. 2013; 232: 161–168.
39. Qi X, Lin W, Li J, Ji H, Wang W, Wang D, Sun M. Fluoxetine increases the activity of the ERK-CREB signal system and alleviates the depressive-like behavior in rats exposed to chronic forced swim stress. *Neurobiol Dis*. 2008; 2: 278–285.
40. Takagi Y, Sakai Y, Abe Y, et al. A common brain network among state, trait, and pathological anxiety from whole-brain functional connectivity. *NeuroImage*. 2018; 172: 506–516.

41. Schmidt HD, Duman RS. Peripheral BDNF produces antidepressant-like effects in cellular and behavioral models. *Neuropsychopharmacology*. 2010; 35: 2378–2391.
42. Xiang M, Jiang Y, Hu Z, Yang Y, Botchway BOA, Fang M. Stimulation of anxiety-like behavior via ERK pathway by competitive serotonin receptors 2A and 1A in post-traumatic stress disordered mice. *Neurosignals*. 2017; 25: 39–53.
43. Xiang M, Jiang Y, Hu Z, et al. Serotonin receptors 2A and 1A modulate anxiety-like behavior in post-traumatic stress disordered mice. *Am J Transl Res*. 2019; 11: 2288–2303.
44. Wefers B, Hitz C, Holter SM, et al. MAPK signaling determines anxiety in the juvenile mouse brain but depression-like behavior in adults. *PLoS One*. 2012; 7: e35035.
45. Satoh Y, Endo S, Nakata T, et al. ERK2 contributes to the control of social behaviors in mice. *J Neurosci*. 2011; 31: 11953–11967.
46. Mastrodonato A, Martinez R, Pavlova IP, et al. Ventral CA3 activation mediates prophylactic ketamine efficacy against stress-induced depressive-like behavior. *Biol Psychiatry*. 2018; 84: 846–856.
47. Wang C, Zhang Y, Shao S, Cui S, Wan Y, Yi M. Ventral hippocampus modulates anxiety-like behavior in male but not female C57BL/6J mice. *Neuroscience*. 2019; 418: 50–58.
48. McLean CP, Asnaani A, Litz BT, Hofmann SG. Gender differences in anxiety disorders: prevalence, course of illness, comorbidity and burden of illness. *J Psychiatr Res*. 2011; 45: 1027–1035.
49. Selcher JC, Nekrasova T, Paylor R, Landreth GE, Sweatt JD. Mice lacking the ERK1 isoform of MAP kinase are unimpaired in emotional learning. *Learn Mem*. 2001; 8: 11–19.
50. Dwivedi Y, Zhang H. Altered ERK1/2 signaling in the brain of learned helpless rats: relevance in vulnerability to developing stress-induced depression. *Neural Plast*. 2016; 2016: 7383724.
51. Mazzucchelli C, Vantaggiato C, Ciamei A, et al. Knockout of ERK1 MAP kinase enhances synaptic plasticity in the striatum and facilitates striatal-mediated learning and memory. *Neuron*. 2002; 34: 807–820.
52. Labonte B, Engmann O, Purushothaman I, et al. Sex-specific transcriptional signatures in human depression. *Nat Med*. 2017; 23: 1102–1111.
53. Paxinos G, Watson C. *The Rat Brain in Stereotaxic Coordinates*. 6th ed. Amsterdam, the Netherlands: Elsevier; 2007.