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Chronic administration of 13-*cis*-retinoic acid does not alter the number of serotonergic neurons in the mouse raphe nuclei

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Abstract

The synthetic retinoid 13-*cis*-retinoic acid (13-*cis*-RA), prescribed for the treatment of severe nodular acne, has been linked to an increased incidence of depression. Chronic treatment studies in rodents have shown that 13-*cis*-RA induces an increase in depression-related behaviours and a functional uncoupling of the hippocampus and dorsal raphe nucleus (DRN). Changes in the number of serotonergic neurons in the DRN have been reported in depressed human patients. Given that retinoids have apoptotic effects, we hypothesised that a decrease in the number of serotonergic neurons in the DRN or median raphe nucleus (MRN) would lead to decreased serotonergic tone and in turn to the behavioural changes seen with 13-*cis*-RA administration. Here, we used immunolabelling and unbiased stereological methods to estimate the number of serotonin (5-hydroxytryptamine, 5-HT) neurons in the MRN and DRN of vehicle control and 13-*cis*-RA-treated adult mice. In the MRN, the number of 5-HT immunolabelled cells was 1815 ± 194 in control, compared with 1954 ± 111 in 13-*cis*-RA treated tissues. The number of 5-HT immunolabelled cells was much higher in the DRN, with 7148 ± 377 cells in the control, compared with 7578 ± 424 in the 13-*cis*-RA treated group. Further analysis of the DRN revealed that there were no changes in the number of 5-HT neurons within distinct subregions of the DRN. Similarly, changes in the density of serotonergic neurons or in the volume of the MRN or DRN were not observed in 13-*cis*-RA treated animals. These data show that apoptotic actions of 13-*cis*-RA do not occur *in vivo* at drug concentrations that induce changes in depression-related behaviour and functional uncoupling of the DRN and hippocampus. The potential pro-depressant behavioural and molecular effects associated with chronic administration of 13-*cis*-RA may result from changes in serotonergic activity rather than changes in the number of serotonergic neurons.

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Keywords

Retinoids; dorsal raphe nucleus; depression; stereology; 5-hydroxytryptamine; tryptophan hydroxylase

INTRODUCTION

Vitamin A and related retinoids are increasingly recognized to play an important role in adult brain function (Lane and Bailey, 2005). Studies have shown that retinoid signalling influences learning and memory (Etchamendy et al., 2001, Cocco et al., 2002), sleep (Maret et al., 2005, Kitaoka et al., 2007), locomotor activity (Krezel et al., 1998) and mood (O'Donnell, 2004, Bremner and McCaffery, 2008, O'Reilly et al., 2008). The synthetic retinoid 13-*cis*-retinoic acid (13-*cis*-RA, isotretinoin, Accutane) is prescribed for the treatment of severe nodular acne (Zouboulis and Piquero-Martin, 2003). Controversially, the use of 13-*cis*-RA has been reported to increase the incidence of depressive illness in approximately 1-10% of patients receiving the drug (Hull and D'Arcy, 2005, Bremner and McCaffery, 2008). In animal studies chronic administration of 1 mg/kg 13-*cis*-RA has been shown to increase depression-related behaviours in both mice and rats (O'Reilly et al., 2006, Trent et al., 2009) but not in all studies of the adult rat (Ferguson et al., 2005, 2007). However, the mechanism by which 13-*cis*-RA treatment can influence depression-related behavioural changes remains poorly understood.

We have recently shown that the administration of 13-*cis*-RA in mice for six weeks leads to metabolic changes in the inferior rostral linear nucleus of the raphe and a functional uncoupling between the raphe nuclei and the hippocampus (O'Reilly et al., 2009). One explanation for this is that 13-*cis*-RA treatment adversely affects serotonergic input to forebrain regions. Retinoid signalling plays a crucial role in neuronal proliferation, differentiation and apoptosis during development of the nervous system (Maden, 2007). Similar actions of retinoid signalling have been proposed to occur in the mature brain. In the hippocampus of young adult mice treated with 13-*cis*-RA, adult neurogenesis was suppressed (Crandall et al., 2004). In cultured hippocampal neurons, high concentrations of 13-*cis*-RA reduced the number of neurons and the degree of dendritic branching (Liu et al., 2008). A similar effect on the morphology of serotonergic neurons was observed in slice cultures treated with high concentrations of 13-*cis*-RA (Ishikawa et al., 2008). Taken together these data suggest that long-term treatment with 13-*cis*-RA may lead to reduced serotonergic input to the brain regions which mediate depression-related behaviours.

A large body of evidence supports the involvement of serotonin (5-hydroxytryptamine, 5-HT) in the pathogenesis of depression (Mann, 1999, Neumeister et al., 2004). Acute tryptophan depletion transiently reduces 5-HT synthesis and induces relapse in depressed patients (Delgado et al., 1991, Bell et al., 2001). Activation of somatodendritic G_i/G_o-protein coupled 5HT_{1A} autoreceptors inhibits electrical activity in serotonergic neurons, 5-HT synthesis and 5-HT release from nerve endings (Blier and de Montigny, 1987, Sharp et al., 1989). Both increased and decreased expression of the 5-HT_{1A} receptor have been reported in depressed and suicidal patients (Neumeister et al., 2004). In addition, polymorphisms in the promoter of the human 5-HT_{1A} receptor have been linked with depression in some (Lemonde et al., 2003) but not all (Arias et al., 2002) patients. The 5-HT transporter (5-HTT) acts as a critical regulator of serotonin signalling via reuptake of 5-HT from the synaptic space (Blakely et al., 1991). Abnormalities in the expression or function of the 5-HTT may also contribute to depression. One 5-HTT polymorphism is associated with lower 5-HTT expression and anxiety related traits (Lesch et al., 1996) and with increased rates of depression and suicidality (Caspi et al., 2003). Additionally, the antidepressant

selective serotonin reuptake inhibitors (SSRIs) act by blockade of 5-HTT to enhance serotonergic neurotransmission (Blier and de Montigny, 1994). Previous work by our group, *in vitro*, has shown that 13-*cis*-RA administration leads to an increase in protein levels of the 5-HT_{1A} receptor and the 5-HTT, as well as increasing intracellular 5-HT content (O'Reilly et al., 2007). Such changes are likely to influence the activity of serotonergic neurons and hence serotonergic input to the forebrain.

The serotonin system in the brain originates in the raphe nuclei and projects throughout the brain to innervate forebrain regions including the hippocampus and frontal cortex (Jacobs and Azmitia, 1992, Barnes and Sharp, 1999, Lechin et al., 2006). The median raphe nucleus (MRN) innervates the dorsal hippocampus, medial septum, nucleus accumbens core and hypothalamus. The dorsal raphe nucleus (DRN) innervates the fronto-parietal cortex, amygdala, lateral septum, nucleus accumbens shell, ventral hippocampus and hypothalamus. The DRN thus provides the main source of serotonergic input to the cortico-limbic structures that are associated with mood disorders (Michelsen et al., 2007, Drevets et al., 2008). A decrease in the overall numbers of serotonergic neurons in the DRN of depressed patients has been reported (Baumann et al., 2002) although others have reported increased numbers of serotonergic neurons in suicide patients (Underwood et al., 1999). We hypothesised that chronic treatment with 13-*cis*-RA may cause a decrease in the number of serotonergic cells emanating from the DRN, thus reducing serotonergic input to the cortico-limbic structures implicated in mediating depression-related behaviours. In this study we have used an unbiased stereological procedure to estimate the volume and number of serotonergic neurons in the MRN and DRN of young adult mice with or without 13-*cis*-RA chronic treatment. To the best of our knowledge this is the first report of a detailed stereological analysis of the adult mouse raphe nuclei.

EXPERIMENTAL PROCEDURES

Animals

Experiments were conducted with DBA/2J young adult male mice obtained from Jackson Laboratories (Bar Harbor, ME) and Charles River Ltd (Margate, UK). Mice were received at postnatal day 21 and treatment began at postnatal day 28 and continued for six weeks. All mice were housed in groups of four per cage with *ad libitum* access to food and water and maintained on a 12:12 h light dark cycle. All of the procedures performed were approved by the University of Texas IACUC, protocol number 04100403, according to the National Institutes of Health Guide for the Care and Use of Laboratory Animals and in the UK performed under a project licence held under the Animals (Scientific Procedures) Act 1986 and in accordance with the UK Home Office guidelines.

Drug Treatment

Animals were randomly assigned to control or drug treatment groups and allowed to acclimatize to their environment for one week before commencing treatment. Drug treatment was essentially as described previously (O'Reilly et al., 2006). All mice received an intra-peritoneal injection of either vehicle (0.9% w/v sodium chloride/ dimethyl sulphoxide (DMSO) at a ratio of 1:1 v/v) or 1mg/kg 13-*cis*-RA, dissolved in vehicle, daily for six weeks. This treatment regime has been shown to induce an increase in depression-related behaviours in young adult mice (O'Reilly et al., 2006). This dose of 13-*cis*-RA is in the range of doses widely used to treat acne in patients (0.5 to 2 mg/kg/day) and achieves plasma levels similar to those seen in patients (Kerr et al., 1982, O'Reilly et al., 2006).

Tissue preparation

At the end of treatment, mice were anaesthetized and perfused transcardially with 4% paraformaldehyde. The brains were immediately removed intact and postfixed in 4% paraformaldehyde overnight followed by storage in PBS with 0.1% azide at 4°C. Brains were cut in the caudal-rostral direction across the region containing both the MRN and DRN (−5.20 to −4.04 mm from Bregma, (Paxinos and Franklin, 2001) using a vibrotome to achieve 40 µm coronal sections that were stored in PBS with 0.1% azide at 4°C until immunohistochemistry was performed.

5-HT Immunohistochemistry

For immunohistochemistry every third section across the region containing both the MRN and DRN was processed. All washes were done in PBS at room temperature unless otherwise stated. Sections were washed and then blocked for endogenous peroxides (3:1 methanol: 3% hydrogen peroxide) followed by incubation in SuperBlock (Pierce, Rockford, IL) for 15 min. Subsequently, sections were incubated in blocking solution (BS) consisting of PBS containing 10% SuperBlock, 10% normal goat serum, 2% bovine serum albumin, and 0.2% Triton X-100 with a 1:10,000 dilution of the rabbit polyclonal primary anti-5-HT antibody (Sigma, Saint Louis, MO) for 24 h. The sections were then washed and incubated for 1 h in biotinylated goat anti-rabbit secondary antibody (Sigma, Saint Louis, MO) at a 1:600 dilution in BS. After washing, immunolabelling was detected using avidin-conjugated horseradish peroxidase with diaminobenzidine as substrate using the Vectastain ABC kit (Vector Labs, Burlingame, CA) according to the manufacturer's protocols. Sections were mounted onto slides and allowed to air dry prior to Nissl counterstaining and dehydration through alcohols (75% to 100%), cleared in xylene and coverslipped with Permount mounting solution (Fisher Scientific).

Stereological analysis of 5-HT immunolabelled neurons

Using Nissl counterstained sections the MRN and DRN were outlined at a low magnification (4X objective) with the aid of a mouse atlas (Paxinos and Franklin, 2001). To demark the MRN, the anterior or ventral tegmental nucleus was used to determine the most dorsal region of the MRN, while the paramedian raphe nuclei and the tectospinal tract were used to determine the edges along the dorsal-ventral axis. The most ventral region of the MRN was identified by locating the pericentral reticulotegmental nucleus of the pons. The slices of the MRN used for the stereological analysis started at approximately −4.04 mm from Bregma and continued posterior to −4.96 mm from Bregma. The landmark used to help identify the most ventral region of the DRN was the superior cerebellar peduncle and the caudal linear nucleus of the raphe. The aqueduct was used as a landmark to determine the dorsal boundary of the DRN and sections used for stereological analysis were identified at −4.04 mm from Bregma and continued until −5.02 mm from Bregma.

The stereological analysis performed was similar to that described in (Chakraborty et al., 2003). The stereological analyses were performed using a computer-assisted morphometry system consisting of an Olympus BX61 photomicroscope (Center Valley, PA) and StereoInvestigator morphometry and stereology software (MicroBrightField, Inc., Colchester, VT). Stereological microscope analyses were performed at high magnification (100X oil objective, N.A. 1.35) and approximately nine slices were analyzed per subject for the MRN and 10 slices were analyzed per subject for the DRN. The observer was blinded as to which treatment group the subject belonged to during the stereological analysis.

The StereoInvestigator software placed dissector frames using a systematic random design within each contour outlining each region on a 90 × 90-µm² grid for the MRN and a 90 × 130-µm² grid for the dorsal raphe nuclei. 5HT labeled cells were counted within 45 × 45

μm optical dissector frames on the x-/y- axis and the final post-processing thickness of each section was measured. The average height of the tissue was $15.65 \mu\text{m}$ and 5HT labeled cells were counted over the whole thickness. The optical fractionator method (Gundersen, 1986, Chakraborty et al., 2003) was used to estimate the total number of 5-HT immunolabelled neurons in the MRN or DRN which were counted as two separate categories. The neuronal number estimates did not depend on a direct measurement of the volume of reference of the region considered and therefore tissue shrinkage during histological processing would not influence the neuronal number estimates. The total cell number is determined as the number of cells counted $\times$ sampling probability⁻¹ where the sampling probability⁻¹ is $(\text{ssf} \times \text{asf} \times \text{tsf})^{-1}$ (ssf is section sampling fraction (0.33); asf is area sampling fraction; tsf is the thickness fraction determined as the height, H, of the unbiased virtual counting spaces divided by t, the thickness of the slice after histological processing, H/t). Coefficient of error (CE) values for the cell number estimates were determined using the Gundersen method incorporated in the StereoInvestigator software and then averaged for each group.

The volumes of the MRN and DRN were measured by drawing the contour plot of each region at 10X magnification and then multiplying the contour area by the total thickness of the interslice distance (number of sections between each contour multiplied by slice thickness i.e. $3 \times 40 \mu\text{m}$). Volume estimates were calculated based on postprocessing tissues that have shrunk in all three directions. No attempts were made to correct for shrinkage because it probably differs in the z and x-y directions.

Tryptophan hydroxylase Immunohistochemistry and Microscopy

To identify whether subregions of the DRN might be affected by 13-*cis*-RA treatment we used tryptophan hydroxylase 2 (TPH-2) immunolabelling. Tissue sections from -4.84 to -4.24 mm from Bregma (Paxinos and Franklin, 2001) were selected for TPH-2 staining. All washes were performed in PBS at room temperature. Sections were washed and blocked in 3% bovine serum albumin for 1 h. Sections were then incubated with rabbit polyclonal anti-TPH2 antibody (Millipore, Billerica, MA, USA, 1:200) overnight at 4°C . Subsequently, sections were washed in PBS and incubated in donkey anti- rabbit alexafluor 568 secondary antibody (Invitrogen, 1:1000) for 2 h at room temperature. After washing, sections were mounted and coverslipped using ProLong Gold (Invitrogen) and stored at 4°C prior to imaging. Images were captured using a confocal laser scanning microscope (laser λ 488nm, Zeiss LSM510, Carl Zeiss Ltd., UK) with a 10x objective. The aqueduct and the medial longitudinal fasciculus were used as anatomical borders to demarcate the DRN (Paxinos and Franklin, 2001). All TPH-2 positive cells were counted in alternate sections, in a total of 8 sections per mouse, and the estimated neuronal number reported as the total in all 8 sections counted. The subregions examined were the DRN ventrolateral part (DRVl), the DRN dorsal part (DRD) and the DRN ventral part (DRV) as identified previously by TPH-2 immunoreactivity (Lowry et al., 2008). The observer was blinded to the treatment of each group during cell imaging and counting.

Statistical analysis

Data are reported as the mean $\pm$ S.E.M of n animals and were subject to a two-tailed *t*-test to analyse the differences between vehicle treated control, and 13-*cis*-RA treated, mouse brains. The n number for each region investigated varies because of variability in the tissue quality of some of the sections.

RESULTS

To identify whether chronic treatment with 13-*cis*-RA altered the number of serotonergic cells in the raphe nuclei, we performed a stereological analysis of immunohistochemically

labelled sections from vehicle control and drug treated adult mouse brains. Qualitative examination revealed that no gross anatomical or histochemical abnormalities were observed in 13-*cis*-RA treated brains compared with vehicle treated controls. 5-HT-immunolabelled cells were easily identified in tissue sections through the MRN and DRN (Fig. 1A-D). For the DRN, the average number of cells counted per mouse was 419.8 ± 47.2 and 458.3 ± 35.6 for control and treated groups respectively ($n=7-10$, $P=0.50$). For the MRN, the average number of cells counted per mouse was 153.8 ± 21.3 and 160 ± 35.6 for control and treated groups respectively ($n=7-10$, $P=0.76$). There were no differences between vehicle control and 13-*cis*-RA treated brains in the estimated number of 5-HT immunolabelled neurons in the MRN or DRN (Fig. 1E). In the MRN, the estimated number of 5-HT immunolabelled cells was 1815 ± 194 in control tissues compared with 1954 ± 111 in the 13-*cis*-RA treated group ($P=0.51$, $n=8$ control, $n=10$ treated), estimated with an average CE of 0.098 ± 0.006 and 0.099 ± 0.006 respectively. The number of 5-HT immunolabelled cells was much higher in the DRN when compared to the MRN with an estimated cell number of 7148 ± 377 in the control group and 7578 ± 424 in the 13-*cis*-RA treated group ($P=0.53$, $n=7$ control, $n=10$ treated), estimated with a CE of 0.081 ± 0.011 and 0.073 ± 0.004 respectively.

Further investigation of the volume of the MRN and DRN revealed no significant differences between vehicle control and 13-*cis*-RA treated mice (Fig. 2). The volume of the MRN was $0.14 \pm 0.02 \text{ mm}^3$ in the control group and $0.14 \pm 0.01 \text{ mm}^3$ in the 13-*cis*-RA treated group ($P=0.99$, $n=8$ control, $n=10$ treated), estimated with an average CE of 0.043 ± 0.005 and 0.027 ± 0.003 respectively. Similarly, in the larger DRN, the volume of the nucleus in control mice was $0.28 \pm 0.02 \text{ mm}^3$ compared with $0.29 \pm 0.01 \text{ mm}^3$ in 13-*cis*-RA treated mice ($P=0.42$, $n=7$ control, $n=10$ treated), estimated with an average CE of 0.049 ± 0.004 and 0.041 ± 0.004 respectively (Fig. 2A). Furthermore there was no difference in the density of cells contained within the MRN or DRN between control and treated groups (Fig 2B, $P=0.34$ for MRN, $P=0.91$ for DRN, $n=7-10$ per group).

To exclude the possibility that subregional differences in the number of serotonergic neurons could be obscured in the stereological analysis, we also examined the numbers of serotonergic cells within subregions of the DRN using TPH-2 immunoreactivity (Fig. 3). The number of TPH-2 immunolabelled cells in the DRV, DRD and DRVL was not significantly different when vehicle control sections were compared with 13-*cis*-RA treated brains (Fig. 3, $P=0.69$ DRV; $P=0.21$ DRD, $P=0.35$ DRVL, $n=6$).

Taken together these data show that chronic administration of 13-*cis*-RA does not decrease serotonergic cell number in the raphe nuclei of young adult mice.

DISCUSSION

In this study we have shown that chronic 13-*cis*-RA treatment has no effect on the number of serotonergic neurons in the raphe nuclei of young adult mice compared with controls. Stereological analysis of 5-HT immunolabelled neurons revealed that the volume, number of serotonergic neurons and neuronal density of the MRN and DRN was not affected by 13-*cis*-RA treatment. Furthermore, analysis of DRN subregions showed that there was not a significant difference in the number of serotonergic neurons in the DRV, DRD and DRVL following 13-*cis*-RA treatment. We have previously shown that chronic treatment with 13-*cis*-RA induces an increase in depression-related behaviours (O'Reilly et al., 2006, Trent et al., 2009). In addition, we have shown that in mice treated with 13-*cis*-RA there is a functional uncoupling of the dorsal raphe nuclei and the hippocampus (O'Reilly et al., 2009). However, the data in the present study suggest that these effects of 13-*cis*-RA are not due to neuronal loss in the MRN or DRN.

To our knowledge, this study represents the first quantitative estimate of the number of serotonergic neurons in the DRN and MRN of the adult mouse obtained by unbiased stereological methods. The total number of 5-HT immunolabelled neurons was estimated to be approximately 7500 in the DRN and 1900 in the MRN of DBA/2J adult mice. During development, at postnatal day 7 in the mouse, similar methods have shown that the number of 5-HT cell bodies in the DRN is 5641 ± 456 (Donovan et al., 2002) which is consistent with our estimates in the adult mouse. Stereological estimates of serotonergic neuronal number in the adult DRN range from 15,000 in rats (Casu et al., 2004, Strackx et al., 2008), 95,000 in dogs (Bernedo et al., 2009) through 51,000 up to 165,000 in humans (Underwood et al., 1999, Bielau et al., 2005). This variability in the number of serotonergic neurons likely reflects the interspecies variation in size.

Increases, decreases and no change in 5-HT cell number have all been reported in depressed human patients (Underwood et al., 1999, Baumann et al., 2002, Hendricksen et al., 2004). Our original hypothesis, that chronic treatment with 13-*cis*-RA would decrease serotonergic cell number, was largely based on evidence that 13-*cis*-RA and other retinoids have been shown to induce apoptosis (Sun et al., 2004, Guruvayoorappan et al., 2008, Liu et al., 2009). The serotonergic neurons of the DRN project to most forebrain areas, including those crucial for emotional behaviours, such as the amygdala and paraventricular nucleus of the hypothalamus, as well as limbic structures, such as the hippocampus (Azmitia and Segal, 1978, Imai et al., 1986, Mamounas et al., 1991, Petrov et al., 1992). An increase in cell death in the DRN in response to retinoid treatment could therefore lead to a reduction in serotonergic input to these forebrain regions involved in mediating responses to stress. Such a decrease could account for the increase in depression-related behaviours, and the functional uncoupling of the DRN and hippocampus, observed following 13-*cis*-RA treatment (O'Reilly et al., 2006, O'Reilly et al., 2009, Trent et al., 2009). Given the proposed trophic role of serotonin (Whitaker-Azmitia, 2001), a decline in serotonergic input to the hippocampus could also account for the reported decreases in adult neurogenesis following 13-*cis*-RA treatment, an effect which may be pro-depressive (Crandall et al., 2004). Ishikawa et al. (2008) used slice cultures from embryonic day 20 rats to show that high concentrations of 13-*cis*-RA (100 μ M) decreased serotonergic cell number. While these effects were blocked by nanomolar concentrations of RAR/RXR antagonists, indicating a receptor-mediated effect, concentrations of 13-*cis*-RA as high as 100 μ M have not been achieved *in vivo*. Indeed, the chronic retinoid treatment regime we used here gives rise to plasma retinoid levels of 1.5 ± 0.4 μ g/ml (5.1 μ M; O'Reilly et al., 2006), well below the high concentrations used to show a decrease in serotonergic cell number *in vitro* (Ishikawa et al., 2008). These lower plasma retinoid levels induce depression-related behaviour (O'Reilly et al., 2006) but did not reduce the number of serotonergic neurons in the raphe nucleus suggesting that some other, non-apoptotic, mechanism accounts for the change in depression-related behaviour.

Although the total number of serotonergic cells arising in the raphe nuclei remain unchanged, it is still possible that the serotonergic input to the hippocampus is impaired by chronic 13-*cis*-RA treatment. It is known that the dendritic and cellular morphology of cultured cells is affected by high levels of 13-*cis*-RA (Ishikawa et al., 2008, Liu et al., 2008) and if this is also true for the *in vivo* situation there would be a loss of serotonergic synaptic contacts occurring within the hippocampus and a reduced serotonergic drive into further hippocampal contacts. Not only may this directly account for the behavioural changes previously reported (O'Reilly et al., 2006, Trent et al., 2009) but it may also account for the functional uncoupling seen between the DRN and the hippocampus (O'Reilly et al., 2009). However, such changes in morphology are unlikely to occur at the plasma retinoid levels that are able to induce an increase in depression-related behaviours, as discussed above.

An alternative explanation for the functional uncoupling of the DRN from the hippocampus could be altered activity of serotonergic neurons following 13-*cis*-RA treatment. Interestingly, mice subjected to chronic stress, a well established paradigm for inducing depressive-like behaviours, show reduced firing of serotonergic neurons in the DRN (Bambico et al., 2009). The firing rate of serotonergic neurons from the DRN can be regulated by 5-HT_{1A} autoreceptors, where a decrease in receptor number leads to increased basal firing activity (Richer et al., 2002) and 5-HT_{1A} receptor stimulation by 5-HT_{1A} agonists or increased intra-cellular 5-HT conversely decreases the firing activity of these neurons (Blier et al., 1998). 13-*Cis*-RA, at 2.5 and 10 μ M, increased 5-HT_{1A} receptor and 5-HTT protein levels and intracellular 5-HT *in vitro* (O'Reilly et al., 2007). If these changes are recapitulated following 13-*cis*-RA treatment *in vivo* one would expect the activity of serotonergic projections from the DRN to be diminished, potentially leading to an increase in depression-related behaviour.

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Abbreviations

5-HT	5-hydroxytryptamine
5-HTT	5-hydroxytryptamine reuptake transporter
13-<i>cis</i>-RA	13- <i>cis</i> -retinoic acid
DRN	dorsal raphe nucleus
MRN	median raphe nucleus
PBS	phosphate buffered saline
SSRIs	selective serotonin reuptake inhibitors
TPH2	tryptophan hydroxylase 2

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Figure 1.

Immunohistochemical analysis of serotonergic neurons in the brain of vehicle control and 13-*cis*-RA treated young adult mice. Illustrative photomicrographs of coronal sections from control mouse brains showing that 5-HT immunolabelled neurons were readily detected in the raphe nuclei in Nissl counter-stained tissue sections at 2x (A) and 4x (B) magnification from which contour plots were drawn. Higher magnification images, 40x (C) and 100x (D), were used for cell counting and show the representative density of 5-HT immunolabelled neurons. The number of serotonergic neurons in the median (MRN) and dorsal (DRN) raphe nuclei were counted (E). Data shown are mean $\pm$ S.E.M.. For MRN control $n=8$, 13-*cis*-RA $n=10$; for DRN control $n=7$, 13-*cis*-RA $n=10$.

Figure 2.

Quantitative analysis of the volume of the raphe nuclei in vehicle control and 13-*cis*-RA treated young adult mice. Chronic 13-*cis*-RA treatment did not alter the estimated volume (A) or the estimated density of serotonergic neurons (B) in either the median (MRN) or dorsal (DRN) raphe nuclei compared with vehicle treated controls. Data shown are mean $\pm$ S.E.M. For MRN control n=8, 13-*cis*-RA n= 10 ; for DRN control n=7, 13-*cis*-RA n=10.

Figure 3.

Analysis of TPH-2 immunolabelling in the dorsal raphe nucleus from vehicle control and 13-*cis*-RA treated adult mice. (A) Illustrative confocal micrograph showing the dorsal raphe nucleus, bounded by the aqueduct (Aq) and medial longitudinal fasciculus (mlf), and the subregions of the dorsal raphe nucleus: the dorsal raphe nucleus dorsal part (DRD), ventral part (DRV) and ventrolateral part (DRVL). Scale bar: 40 μ m (B) TPH-2 immunolabelled cells were counted to estimate the number of serotonergic neurons in each dorsal raphe subregion for both vehicle control and 13-*cis*-RA treated mice. Data shown are mean $\pm$ S.E.M., n= 6 per group.