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In situ visualization of ozone in the brains of mice with depression phenotypes by using a new near-infrared fluorescence probe†

Ping Li,‡ Jijuan Wang,‡ Xin Wang, Qi Ding, Xiaoyi Bai, Yandi Zhang, Di Su, Wei Zhang, Wen Zhang and Bo Tang *

Ozone (O₃), one of the reactive oxygen species (ROS), is deeply involved in diseases including depression. However, the lack of appropriate *in situ* detection methods suitable for the complex biological context of brain impedes uncovering the exact relationship between depression and changes in the O₃ level. Therefore, we developed a near-infrared (NIR) fluorescent probe (ACy7) for the direct visualization of O₃ in mice brains. The specific cycloaddition reaction between O₃ and the terminal double bond of the butenyl group extends the conjugation of the "pre-" heptamethine cyanine system, which emits NIR fluorescence of heptamethine cyanine. This makes the ACy7 specific, highly sensitive and able to deeply penetrate tissue. Using ACy7, we found that under glutamate stimulation, the O₃ content in PC12 cells was significantly higher than that in control cells. By imaging analysis on the brains of mice, we revealed for the first time that the levels of O₃ in mice with depression phenotypes were markedly higher than that in control mice. Intriguingly, experimental results unravelled that excess O₃ promoted high expression of the pro-inflammatory cytokine interleukin-8 (IL-8), which ultimately induced depression phenotypes. Our work demonstrates the pivotal role of elevated O₃ in depression and provides a fresh entry point for exploring oxidative stress contributing to depression.

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Introduction

Depression is a mental illness affecting people throughout the world, causing high mortality and disability rates.¹ However, the aetiology of depression is still unclear. In recent years, the immunoinflammatory hypothesis of depression has received increasing attention. This hypothesis suggests that inflammatory mediators such as cytokines play an important role in the development of depression and that the immunoinflammatory reaction may be an essential cause of the poor efficacy of conventional antidepressant treatments.^{2,3}

As one of the reactive oxygen species (ROS), ozone (O₃) has attracted widespread attention due to its key role in mental disorders,^{4–7} such as depression. Unfortunately, there are two different perspectives on the relationship between O₃ and depression. Some researchers have shown that exogenous O₃ stimulation induces depressive behaviors.^{8–11} Others have

shown that ozonation in conjunction with autologous blood transfusion could be used as a new, fast acting antidepressant treatment.¹² O₃ is endogenously produced in the inflammation and antibacterial response of the immune system,^{13,14} which causes the production of cytokines, including the pro-inflammatory cytokine interleukin-8 (IL-8), amplifying the inflammatory cascade.¹⁵ So, we speculate that during depression, the level of endogenous O₃ in the brain is increased, resulting in the overexpression of IL-8, which is the ultimate cause of depression. To confirm our hypothesis and clarify the role of O₃ in depression, the development of appropriate *in vivo* detection methods to *in situ* visually monitor changes in O₃ in brains of mice with depression phenotypes is of special note.

Fluorescent probes are characterized by high sensitivity, relatively simple preparation, and good biocompatibility, and therefore they have been used to observe target molecules in living organisms.^{16–20} In particular, optical imaging associated with the near-infrared (NIR) fluorescent probe has the advantages of being suitable for deep tissue penetration and low autofluorescence background, making this technique more applicable for *in vivo* detection.²¹ In recent years, fluorescent probes for monitoring O₃ have been developed, but these probes are mostly used for detecting O₃ at the environmental or the cellular level.^{22–26} Due to the complex biological environment in the brain and the blood–brain barrier, probes for measuring O₃ in living brains have not been reported.

College of Chemistry, Chemical Engineering and Materials Science, Key Laboratory of Molecular and Nano Probes, Ministry of Education, Collaborative Innovation Center of Functionalized Probes for Chemical Imaging in Universities of Shandong, Institutes of Biomedical Sciences, Shandong Normal University, Jinan 250014, People's Republic of China. E-mail: tangb@sdsu.edu.cn

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‡ These authors contributed equally to this work.

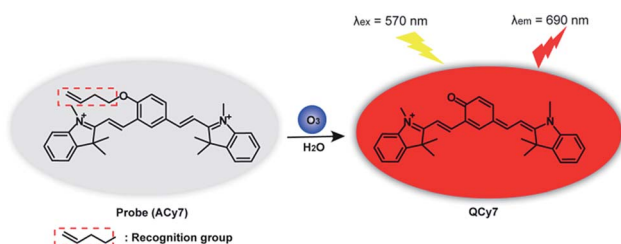


Thus, we designed and synthesized an “off-on” NIR small-molecule fluorescent probe (ACy7) for high sensitivity fluorescence imaging of O₃ in the brains of mice with depression phenotypes. Our probe uses a Cy7-like molecule as the precursor of the fluorophore and 3-butenyl as the recognition group.²⁶ O₃ can undergo a specific cycloaddition reaction with the terminal olefin of the 3-butenyl moiety to convert ACy7 to the quinone derivative QCy7, extending the degree of conjugation. This reaction triggers emitting bright, NIR fluorescence of heptamethine cyanine (Scheme 1), allowing the specific and highly sensitive detection of O₃. We first explored the applicability of ACy7 in macrophages and PC12 cells. Then, we applied ACy7 to the *in situ* fluorescence imaging of O₃ in the brains of mice with depression phenotypes. In addition, we investigated the molecular mechanisms of depression associated with O₃ regulation of IL-8 involvement.

Results and discussion

In vitro optical properties of ACy7 to O₃

ACy7 was synthesized by a two-step reaction (Scheme S1, ESI[†]), purified by thin-layer chromatography and column chromatography, and then characterized by ¹H NMR, ¹³C NMR and MS. To understand the effect of O₃ on the photophysical properties of ACy7, we first measured the absorption spectra of the probe in the absence and presence of O₃. As shown in Fig. S1a (ESI[†]), the absorption maximum of the probe was at approximately 420 nm in the absence of O₃. When O₃ was added to the system, the probe exhibited a new absorption band at approximately 570 nm, and the intensity of the absorption at 570 nm enhanced with increasing O₃ concentration. Subsequently, we studied the fluorescence properties of QCy7. Fig. S1b (ESI[†]) shows that the maximum excitation and emission peaks of QCy7 are at 570 nm and 690 nm, respectively, with a large Stokes shift. We examined the fluorescence response of ACy7 to O₃ and optimized the reaction conditions (Fig. S2–S5, ESI[†]). We measured the fluorescence emission intensities of ACy7 before and after the addition of O₃ (Fig. 1A). The figure shows that when the reaction solution did not contain O₃, the fluorescence emission of the probe at 690 nm was negligible. When O₃ solution is added to the mixture, the fluorescence emission intensity of the probe at 690 nm is significantly enhanced, increasing by more than 80-fold. This indicates that the probe has a high signal to noise ratio. This result is consistent with the design of our probe, tuning fluorescence from “off” to “on” by extending the conjugation of the “pre-” heptamethine cyanine system. As depicted



Scheme 1 “Off-on” type NIR fluorescent probe ACy7 reaction with O₃.

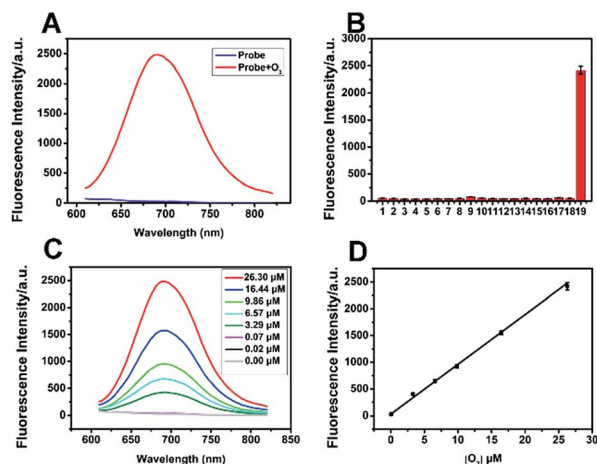


Fig. 1 Fluorescence spectral properties of ACy7. (A) Fluorescence emission spectrum of 10 μM ACy7 before and after reaction with O₃ (26.30 μM). (B) Fluorescence responses of 10 μM ACy7 to some biological interfering substances. (C) Fluorescence response of 10 μM ACy7 after the addition of serial concentrations of O₃. (D) A linear relationship between the fluorescence emission intensity of the ACy7 solution and the O₃ concentration ($\lambda_{\text{ex}}/\lambda_{\text{em}} = 570/690 \text{ nm}$). Error bars are \pm SD ($n = 3$). (Interfering substances: (1) blank, (2) H₂O₂ (0.1 mM), (3) O₂^{•-} (125 μM), (4) ¹O₂ (50 μM), (5) \cdot OH (50 μM), (6) *t*-BuOOH (30 μM), (7) NaClO (50 μM), (8) NO (50 μM), (9) ONO⁻ (6 μM), (10) GSH (5 mM), (11) vitamin C (0.3 mM), (12) Na⁺ (0.5 mM), (13) K⁺ (0.5 mM), (14) Zn²⁺ (0.5 mM), (15) Ca²⁺ (0.5 mM), (16) Na₂S₂O₃ (0.5 mM), (17) Fe³⁺ (0.5 mM), (18) citric acid (50 μM), and (19) O₃ (26.30 μM).)

in Scheme S2 (ESI[†]), through a (3 + 2) cycloaddition reaction, the terminal olefin of probe 1 reacts with O₃ to form molecular ozonide 2, which further generates aldehyde 3, HCHO and H₂O₂ in the aqueous solution. Then, aldehyde 3 undergoes β-elimination to produce CH₂=CHCHO and compound 4. Phenate compound 4 undergoes electron rearrangement to form a fluorescent quinone derivative QCy7.²⁷ This specific reaction mechanism can endow the probe with outstanding selective and sensitive response to O₃.

Under the optimal conditions, we added various concentrations of O₃ to the ACy7 reaction solution. As expected, the fluorescence emission intensity of QCy7 at 690 nm gradually increased with increasing O₃ concentration. The O₃ concentration and fluorescence emission intensity (ΔF) showed good linearity in the range of 0.02–26.30 μM (Fig. 1C and D). The linear correlation equation was $\Delta F = 93.23[\text{O}_3] (\mu\text{M}) + 29.50$ ($R^2 = 0.9964$). The limit of detection ($3S/k$, in which S is the standard deviation of the blank measurements, $n = 11$, and k was the slope of the linear equation) was 10 nM. To further confirm that the turn-on fluorescence is based on the probe reacting with ozone, we carried out RP-HPLC analysis combined with MS to detect the conversion of ACy7 before and after reaction with an equimolar amount of ozone (Fig. S6, ESI[†]). The results of HPLC-MS verify that 33% of ACy7 transformed to QCy7 with an equimolar amount of ozone, causing a notable increase in fluorescence. These findings demonstrate that the probe can measure low concentration of O₃ with high sensitivity.

To evaluate the specificity of the reaction, we examined some interfering substances that are common in the biological



environment, such as ROS, reactive nitrogen species (RNS), bio-antioxidants and metal ions, under the same conditions (Fig. 1B). When these interfering substances were added, the fluorescence intensity of the probe was extremely low. The addition of O_3 brought about obvious fluorescence enhancement, indicating that the probe has good specificity and can specifically detect O_3 . In addition, we examined the reaction kinetics and photostability of ACy7 to O_3 . As shown in Fig. S7 (ESI[†]), when O_3 was added, the emission intensity of the ACy7 solution reached a plateau at approximately 40 min, and the intensity remained basically unchanged over the next 25 min. The fluorescence emission intensity of ACy7 basically did not change with time. Therefore, ACy7 can quickly react with O_3 and has good photostability. These results indicate that the probe (ACy7) has desirable properties, including emission located in the NIR region, high sensitivity and high specificity. Therefore, ACy7 is expected to be suitable for *in situ* detection and bio-imaging of O_3 in biological samples.

Endogenous O_3 imaging in live cells with ACy7

The above studies on the optical properties of ACy7 *in vitro* proved that ACy7 was well suited for the detection of O_3 under simulated physiological conditions. To test the feasibility of the probe ACy7 as an imaging reagent, its potential toxicity to cells was evaluated using a standard MTT [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide] assay (Fig. S8, ESI[†]).²⁸ The results showed that ACy7 had an IC_{50} value of 0.322 mM, meaning low cytotoxicity.

Then, the applicability of ACy7 for the imaging of RAW 264.7 macrophages was investigated. We firstly tried to visualize exogenous O_3 within living macrophages. Macrophages were incubated directly in 5 μ M of O_3 solution, and then fluorescence brightness was examined (Fig. 2B). The images showed that the fluorescence intensity within these cells was 1.5-fold stronger than that of the control (Fig. 2A). This result indicates that the probe ACy7 can be used for imaging of exogenous O_3 at the cellular level. To investigate whether the probe ACy7 is capable of monitoring endogenous O_3 in cells, phorbol 12-myristate 13-acetate (PMA) was used to stimulate macrophages for producing O_3 (Fig. 2C).²⁹ As expected, intracellular fluorescence intensity was 1.9-fold higher than that of the control. To verify that the

enhancement of fluorescence intensity was caused by an increase in O_3 , we added ethyl 4-vinylbenzoate to eliminate intracellular O_3 (Fig. 2D).¹⁵ We observed that the fluorescence intensity of these cells was significantly reduced (by 1.9-fold) compared to that of the PMA-stimulated group. This experiment further proves that the endogenous O_3 produced in the cells undergoes a cleavage reaction with ACy7 to release QCy7 to emit light instead of reacting with other ROS. The above results demonstrate that our probe has sufficient membrane permeability and can be used for imaging changes in intracellular O_3 .

To investigate the feasibility of ACy7 in nerve cell applications, we visualized the changes in endogenous O_3 levels inside PC12 cells. Considering that glutamate can cause nerve cells to undergo oxidative stress with concomitant ROS production,³⁰ high concentration glutamate was added to the cells (Fig. 3C). We found that the fluorescence signal within these cells significantly increased (8.6-fold) compared with that of the control. This result indicates that excess of O_3 is produced in glutamate-stimulated cells, and ACy7 can be used to image endogenous O_3 in PC12 cells. To further demonstrate that the change in the fluorescence intensity was caused by O_3 , we pre-incubated the cells with ethyl 4-vinylbenzoate (Fig. 3B and D), and found that the fluorescence intensity of these cells was greatly lower than that of the control. The imaging of PC12 cells showed that ACy7 can effectively distinguish between control cells and cells under oxidative stress. Overall, the results of the cell imaging experiments demonstrate that ACy7 has the advantages of membrane permeability and biocompatibility.

Fluorescence imaging of O_3 in the brains of mice with depression phenotypes

Since the emission of the probe is in the NIR region, it causes only slight light damage to biological samples. In particular, it could penetrate deeper tissue with weak background fluorescence interference. These merits make ACy7 well suited for *in vivo* imaging experiments. Therefore, we explored *in vivo* fluorescence imaging of O_3 with the probe. First, we established a chronic unpredictable mild stress (CUMS) mouse model (Fig. S9–S11, ESI[†]).³¹ After successful modelling, both control mice and mice with depression-like behaviors were injected intraperitoneally with 0.15 mg kg^{-1} of ACy7. Subsequently, *in*

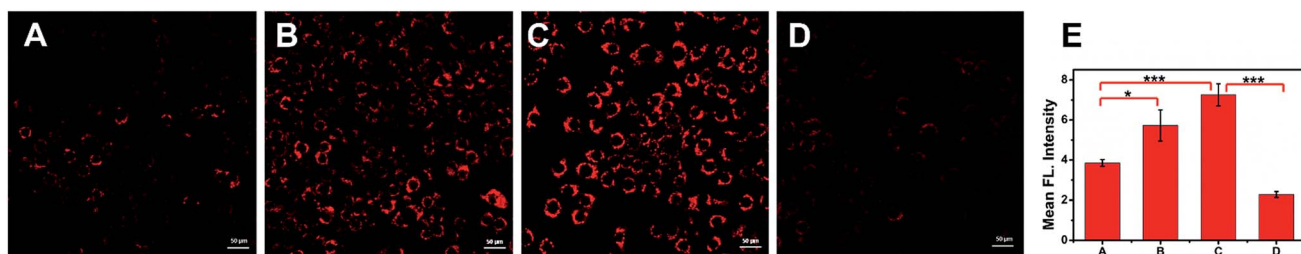


Fig. 2 Laser confocal imaging of O_3 in RAW 264.7 macrophages. (A) Cells incubated with 10 μ M ACy7 for 15 min at 37 $^{\circ}$ C. (B) ACy7-loaded cells incubated with 5 μ M O_3 for 10 min. (C) Cells incubated with PMA (100 ng mL^{-1}) for 20 min and then incubated with 10 μ M ACy7 at 37 $^{\circ}$ C for 15 min. (D) Cells incubated with PMA (100 ng mL^{-1}) for 20 min, and then with 0.05 mM ethyl 4-vinylbenzoate for 30 min, and then with 10 μ M ACy7 for 15 min. (E) The data output of the graphs in (A)–(D). All cells were washed 3 times with 0.01 M PBS prior to imaging. Emissions from 640 nm to 740 nm were collected using a 561 nm laser. The values are the mean \pm s.d. for $n = 3$, * $p < 0.05$, and *** $p < 0.001$.



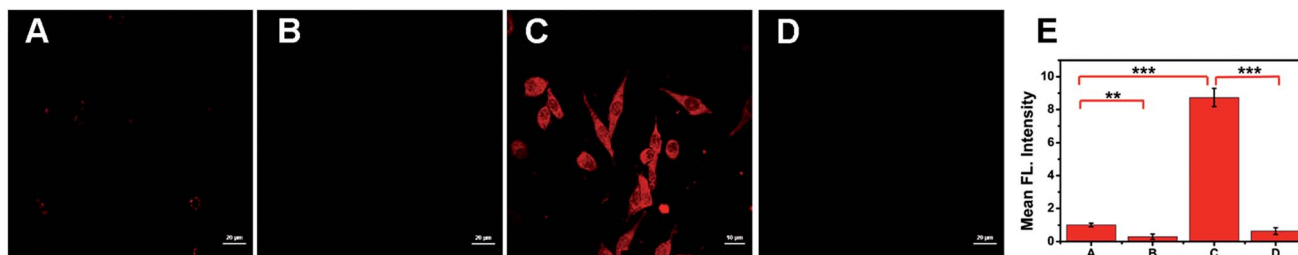


Fig. 3 Laser confocal imaging of O_3 in PC12 cells. (A) Cells incubated with 10 μM ACy7 for 15 min at 37 $^\circ\text{C}$. (B) Cells incubated with 0.05 mM ethyl 4-vinylbenzoate and then treated according to (A). (C) Cells incubated with 10 mM glutamate for 17 h and then incubated with ACy7 for 15 min at 37 $^\circ\text{C}$. (D) Cells incubated with 10 mM glutamate for 17 h, and then with 0.05 mM ethyl 4-vinylbenzoate for 30 min and then with ACy7 for 15 min. (E) The data output of the graphs in (A)–(D). All cells were washed 3 times with 0.01 M PBS prior to imaging. Emissions from 640 nm to 740 nm were collected using a 561 nm laser. The values are the mean \pm s.d. for $n = 3$, ** $p < 0.01$, and *** $p < 0.001$.

situ fluorescence imaging of the brain of each mouse was performed using an IVIS Lumina III (Fig. 4). The results showed that the brain of mice with depression-like behaviors exhibited brighter fluorescence, about 1.8-times higher than that of control mice. This indicates that the O_3 contents in the brains of mice with depression-like behaviors are higher than the levels in control mice. Overall, based on the superior *in vivo* fluorescence imaging performance of the probe ACy7, we revealed the elevation of O_3 in the brains of mice with depression phenotypes for the first time.

Study of the signalling pathway of O_3 -induced depression phenotypes in mice

We have demonstrated that the level of O_3 in the brain of mice is positively correlated with depression phenotypes. We wondered why the flux of O_3 is associated with depression-like behaviors. Studies have shown that O_3 as one of the important ROS is deeply involved in inflammation.^{32–34} In the meantime, there is a close relationship between inflammation and depression.^{35,36} So, we speculated that IL-8 served as a bridge between O_3 and depression. To verify this assumption, we analysed IL-8 in the brains of mice with depression-like behaviors. First, we used lipopolysaccharide (LPS) to establish a mouse model of acute inflammation-induced depression (Fig. S12–S14, ESI[†]),³⁷ and

divided the mice into a control group, an LPS group and an LPS + Lim group. We next injected 0.15 mg kg^{-1} of ACy7 into the mice by intraperitoneal injection and performed *in situ* fluorescence imaging of the mouse brains using an IVIS Lumina III (Fig. 5A and B). The images showed higher fluorescence intensity in the brain tissue of the mice with depression-like behaviors, 1.9-fold greater than that of the control group. Similar results were also obtained by utilizing sulfo-ACy7 (Fig. S15, ESI[†]). This means the obvious increase in O_3 in depression caused by inflammation.

Next, to investigate how IL-8 levels changed in the brains of mice, we measured the IL-8 content in the brain of each mouse by enzyme linked immunosorbent assay (ELISA) (Fig. 5C). The results showed that the IL-8 contents in mice with depression-like behaviors were 1.7-fold greater than those in the control group, indicating that the increase in IL-8 in the brains of mice with depression-like behaviors is caused by inflammation. To prove that the changes in the IL-8 contents in the brains of mice are ascribed to O_3 , we used limonene (Lim, a natural O_3 scavenger) to establish an LPS + Lim mouse model.³⁸ The LPS + Lim group of mice was given 50 mg kg^{-1} Lim by gavage 7 days in advance. Next, we injected 0.83 mg kg^{-1} LPS into the mice by

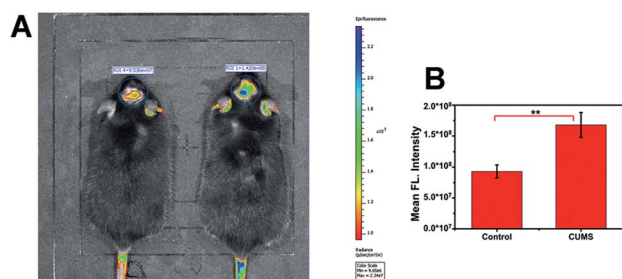


Fig. 4 Visual imaging of O_3 in the brains of mice. (A) Fluorescence imaging (pseudocolor) of ACy7 in control mice (left) and mice with depression-like behaviors (right). (B) The output of the fluorescence intensity in image (A). The excitation wavelength filter is 570 nm, and the emission wavelength filter is 690 nm. The values are the mean \pm s.d. for $n = 3$, and *** $p < 0.01$.

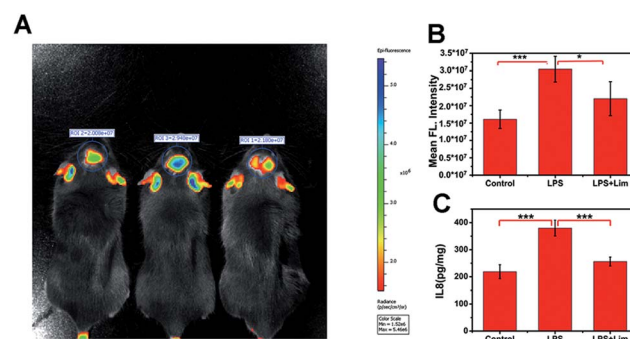


Fig. 5 Imaging of O_3 and detection of IL-8 contents in the brains of mice. (A) Fluorescence imaging (pseudocolor) in the control (left), LPS (middle), and LPS + Lim (right) groups. (B) The output of fluorescence intensity in image (A). (C) Analysis of the IL-8 content in mouse brain tissue by ELISA. The excitation wavelength filter is 570 nm, and the emission wavelength filter is 690 nm. The values are the mean \pm s.d. for $n = 5$, * $p < 0.05$ and *** $p < 0.001$.



intraperitoneal injection to induce depression-like behaviors. The images showed that the fluorescence in the brain of each mouse in the LPS + Lim group was significantly lower than those of the mice in the LPS group. These results indicated that the O₃ in the brains of the LPS + Lim mice was decreased. At the same time, by using DepressionScan (Clever Sys. Inc.), we found that the depression-like behaviors of mice in the LPS + Lim group was relieved. Importantly, compared with mice in the LPS group, the IL-8 level in the brains of the LPS + Lim group was also significantly decreased, which is consistent with our previous speculation. Altogether, the above results fully demonstrate that over-generated O₃ in the mouse brain promotes the production of excess IL-8, which ultimately induces depression.

Conclusions

In summary, we developed a new NIR fluorescent probe (ACy7) to trace O₃ in the brain tissue of small animals. ACy7 was proved to be extremely sensitive and selective to O₃, as well as having good biocompatibility. Relying on these merits, we successfully applied this probe in the visualization of over produced O₃ in PC12 cells under oxidative stress induced by glutamate. Importantly, we realized the *in situ* imaging observation of O₃ in the brain tissue of mice with depression phenotypes for the first time, revealing the elevated O₃ levels in mice with depression phenotypes. Furthermore, we uncovered that O₃ in the brain induces mouse depression by producing excessive IL-8. This work provides strong direct evidence of a positive correlation between O₃ and depression and will help to more fully elucidate the pathogenesis of depression associated with oxidative stress.

Live subject statement

All the animal experiments were carried out in accordance with the relevant laws and guidelines issued by the Ethical Committee of Shandong University and were in agreement with the guidelines of the Institutional Animal Care and Use Committee.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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