

- ④ 三好 耕, 宮崎育子, 浅沼幹人: Pericentrin は発達期大脳皮質の神経一次繊毛の基部に局在する. 第 29 回日本神経科学大会, 京都, 2006, 7, 21.
- ⑤ Miyoshi K., Miyazaki, I. and Asanuma, M.: Involvement of pericentrin in the formation of neuronal primary cilia. 第 28 回日本生物学的精神医学会・第 36 回日本神経精神薬理学会・第 49 回日本神経化学会大会合同年会, 名古屋, 2006, 9, 14.
- ⑥ 宮崎育子, 浅沼幹人, 小川紀雄: Dopamine quinone-related neurotoxicity and potential neuroprotective agents. 第 28 回日本生物学的精神医学会・第 36 回日本神経精神薬理学会・第 49 回日本神経化学会大会合同年会, 名古屋, 2006, 9, 15.
- ⑦ 浅沼幹人, 喜多大三, 日名俊行, 宮崎育子, 小川紀雄, 北村佳久, 千堂年昭, 五味田裕: Effect of methylphenidate on excess extra-vesicular dopamine-induced dopaminergic neurotoxicity. 第 28 回日本生物学的精神医学会・第 36 回日本神経精神薬理学会・第 49 回日本神経化学会合同年会, 名古屋, 2006, 9, 16.
- ⑧ 浅沼幹人, 宮崎育子, 北市清幸: メタンフェタミン急性神経毒性におけるドパミンキノン体生成の関与とキノン消去による阻止. シンポジウム II 覚せい剤による神経傷害の生化学, 臨床画像, そして修復・治療, 第 18 回日本アルコール精神医学会・第 9 回ニコチン・薬物依存研究フォーラム平成 18 年度合同学術総会, 千葉, 2006, 9, 29.
- ⑨ Miyazaki, I., Asanuma, M., Diaz-Corrales, F.J. and Ogawa, N.: Protective effects of a novel anti-parkinsonian agent zonisamide on dopamine quinone-related neurotoxicity. 10th International Congress of Parkinson's Disease and Movement Disorders, Kyoto,

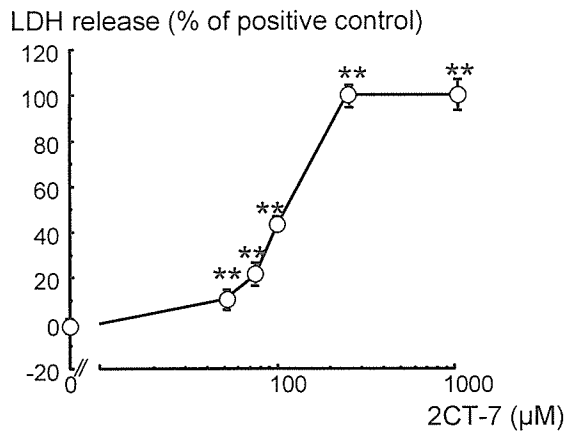
2006, 10, 31.

- ⑩ Asanuma, M., Miyazaki, I., Diaz-Corrales, F.J. and Ogawa, N.: A novel anti-parkinsonian agent zonisamide increases glutathione levels in the basal ganglia. 10th International Congress of Parkinson's Disease and Movement Disorders, Kyoto, 2006, 10, 31.

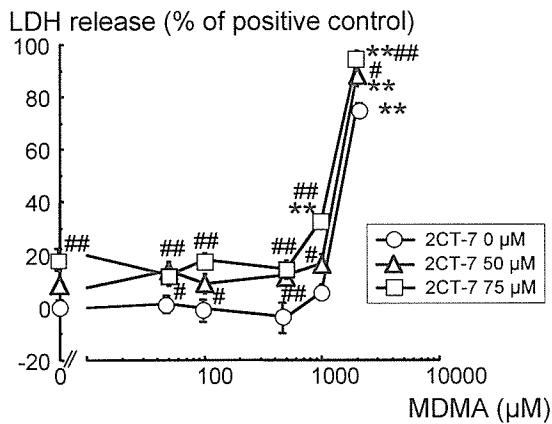
H. 知的財産権の出願・登録状況

特許取得
特になし
実用新案登録
特になし
その他
特になし

A CATH.a cells



B CATH.a cells



C CATH.a cells

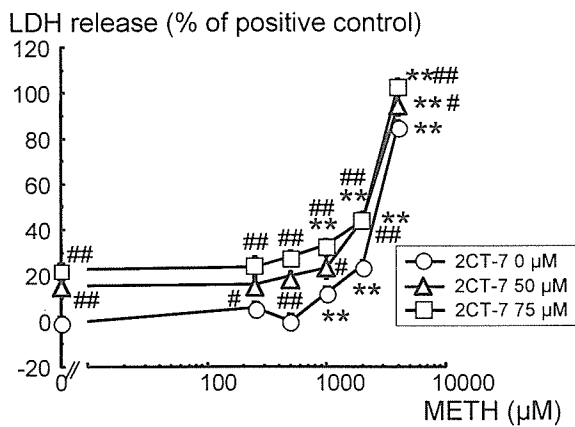


Fig. 1. Changes in released LDH from dopaminergic CATH.a cells after exposure to 2CT-7 (A), 2CT-7+MDMA (B) or 2CT-7+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. ** p <0.01 vs. each control group without MDMA or METH. # p <0.05, ## p <0.01 vs. MDMA/METH-dose-matched control group without 2CT-7.

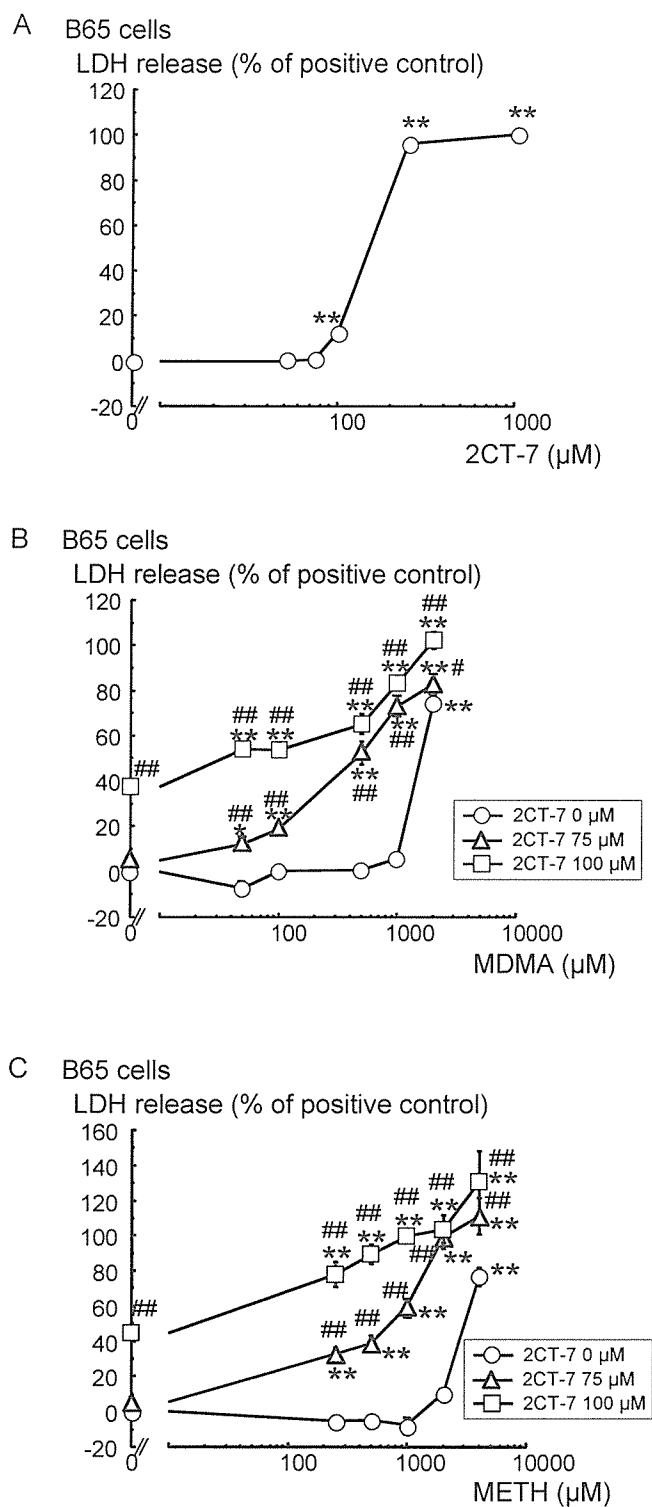


Fig. 2. Changes in released LDH from dopaminergic B65 cells after exposure to 2CT-7 (A), 2CT-7+MDMA (B) or 2CT-7+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. * $p < 0.05$, ** $p < 0.01$ vs. each control group without MDMA or METH. # $p < 0.05$, ## $p < 0.01$ vs. MDMA/METH-dose-matched control group without 2CT-7.

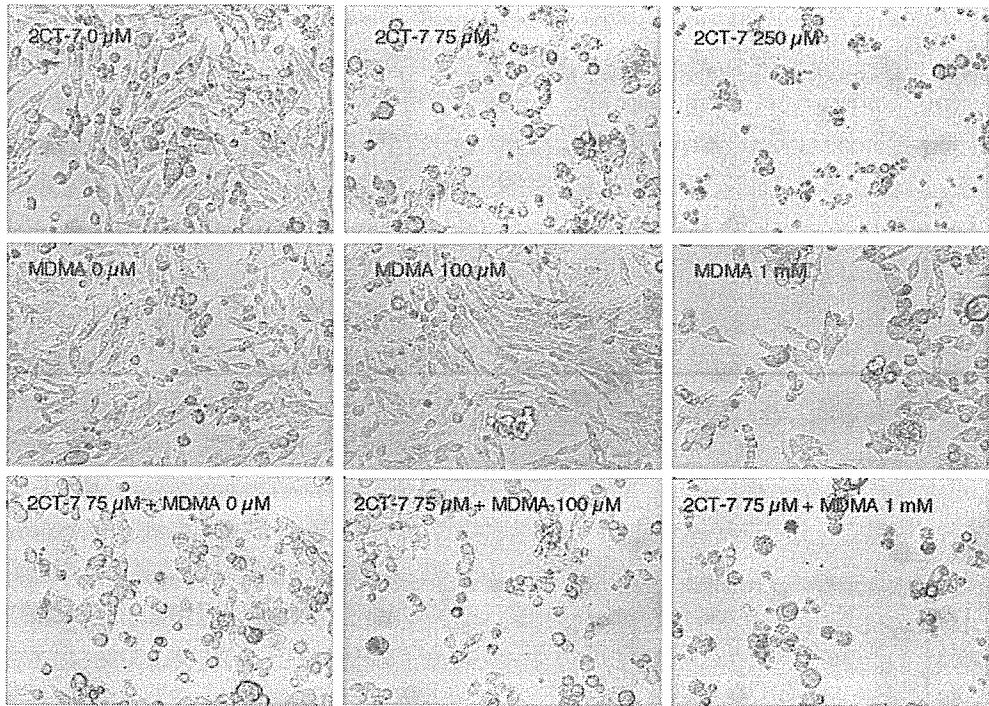


Fig. 3. Photographs of B65 cells treated with 2CT-7 and/or MDMA (final concentration: 0, 100 μM, 1 mM) for 24 hours.

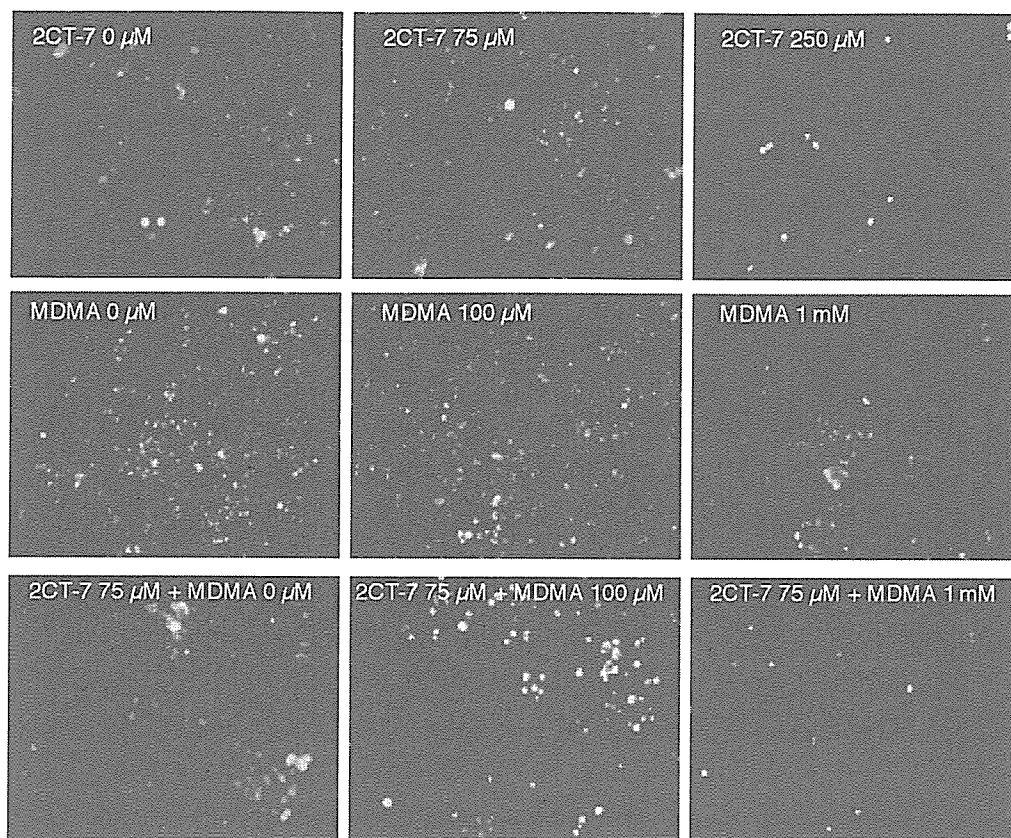


Fig. 4. Nuclear staining of B65 cells treated with 2CT-7 and/or MDMA (final concentration: 0, 100 μ M, 1 mM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

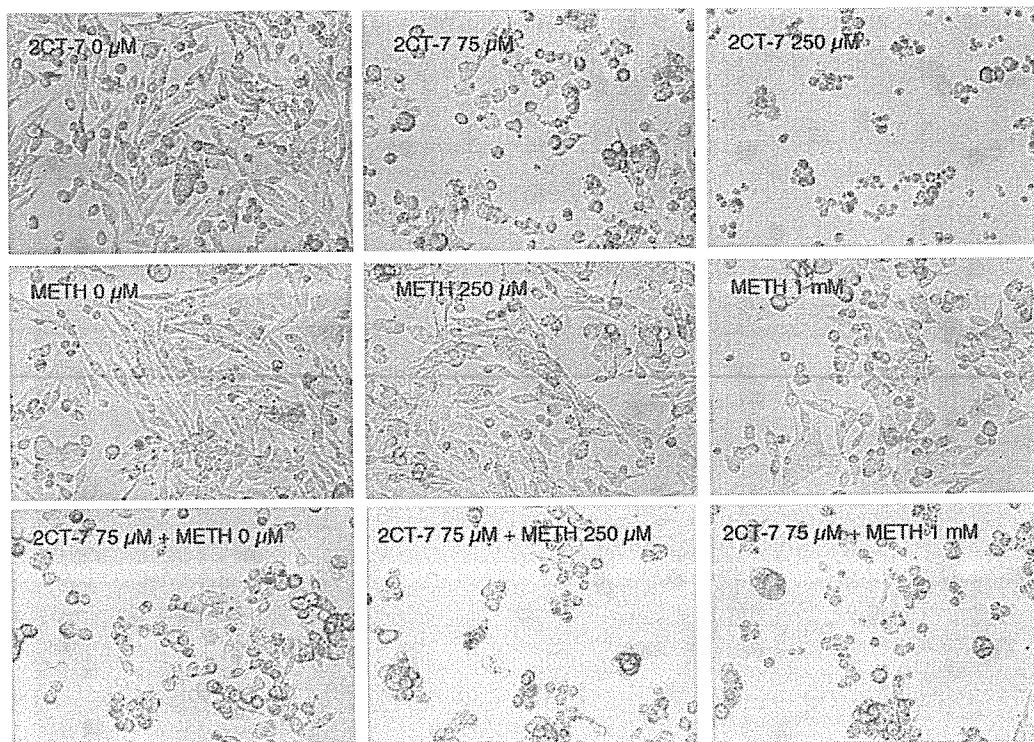


Fig. 5. Photographs of B65 cells treated with 2CT-7 and/or METH (final concentration: 0, 250 μM, 1 mM) for 24 hours.

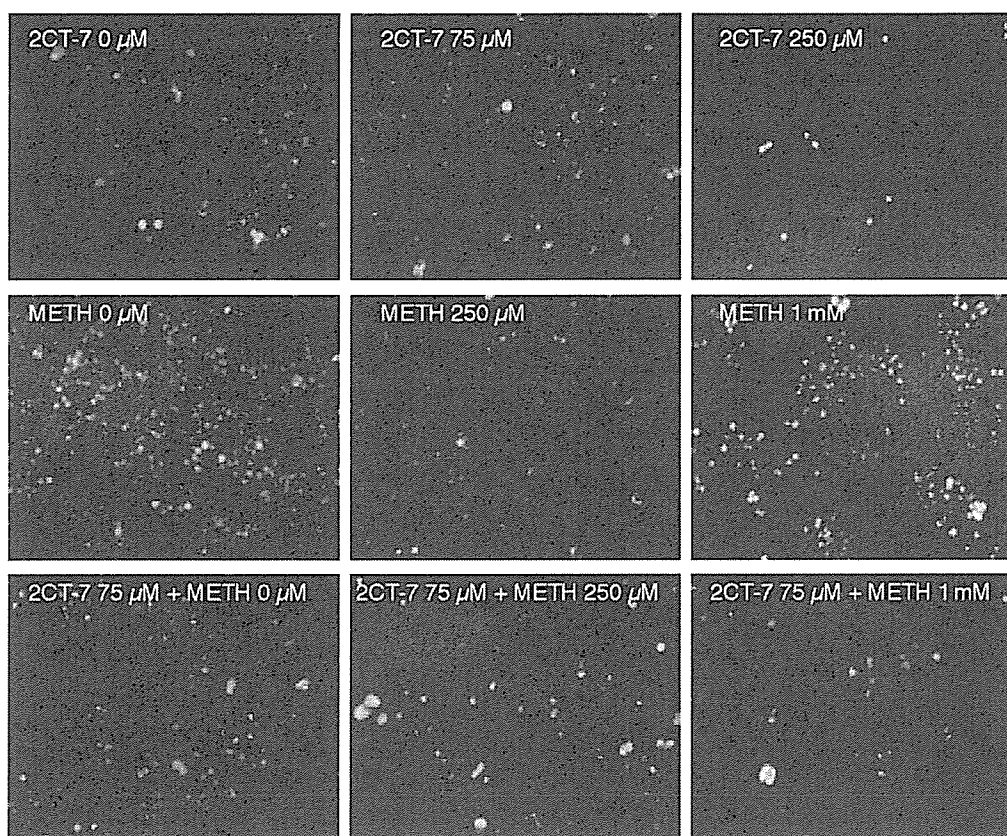
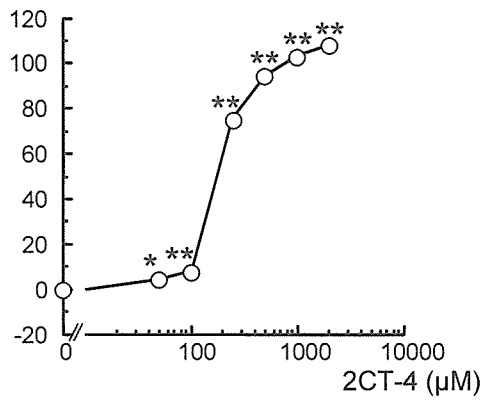


Fig. 6. Nuclear staining of B65 cells treated with 2CT-7 and/or METH (final concentration: 0, 250 μM , 1 mM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

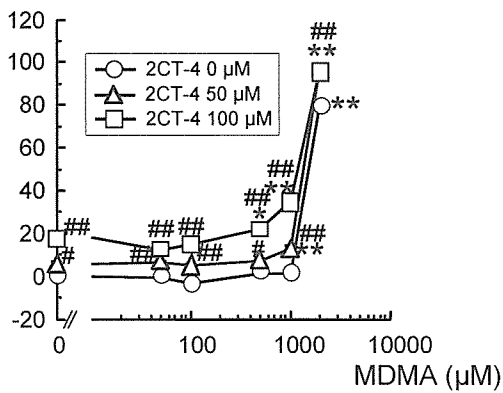
A CATH.a cells

LDH release (% of positive control)



B CATH.a cells

LDH release (% of positive control)



C CATH.a cells

LDH release (% of positive control)

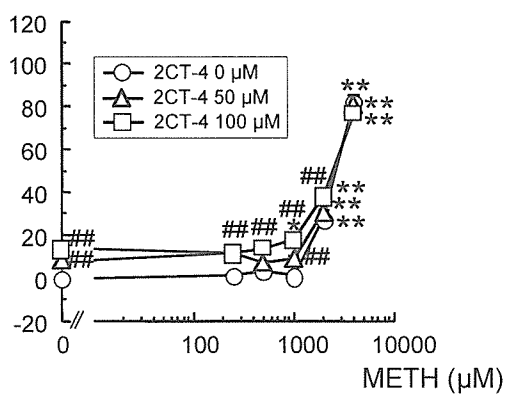
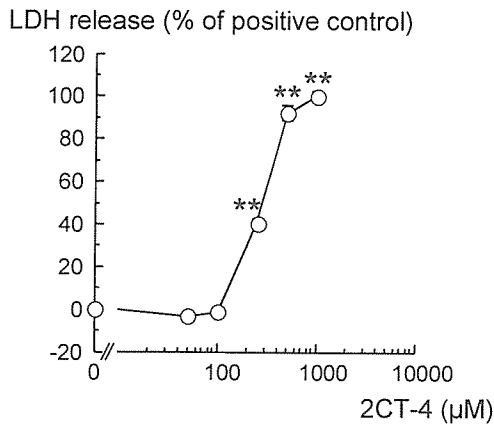
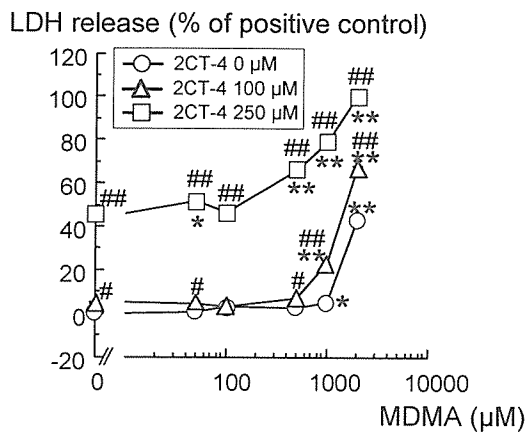


Fig. 7. Changes in released LDH from dopaminergic CATH.a cells after exposure to 2CT-4 (A), 2CT-4+MDMA (B) or 2CT-4+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. * p <0.05, ** p <0.001 vs. each control group without MDMA or METH. # p <0.01, ## p <0.001 vs. MDMA/METH-dose-matched control group without 2CT-4.

A B65 cells



B B65 cells



C B65 cells

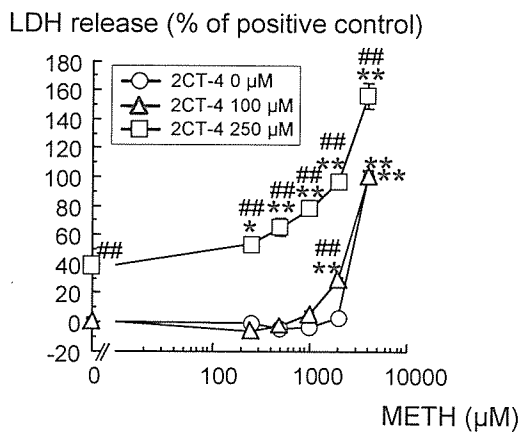


Fig. 8. Changes in released LDH from dopaminergic B65 cells after exposure to 2CT-4 (A), 2CT-4+MDMA (B) or 2CT-4+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. * p <0.05, ** p <0.001 vs. each control group without MDMA or METH. # p <0.05, ## p <0.001 vs. MDMA/METH-dose-matched control group without 2CT-4.

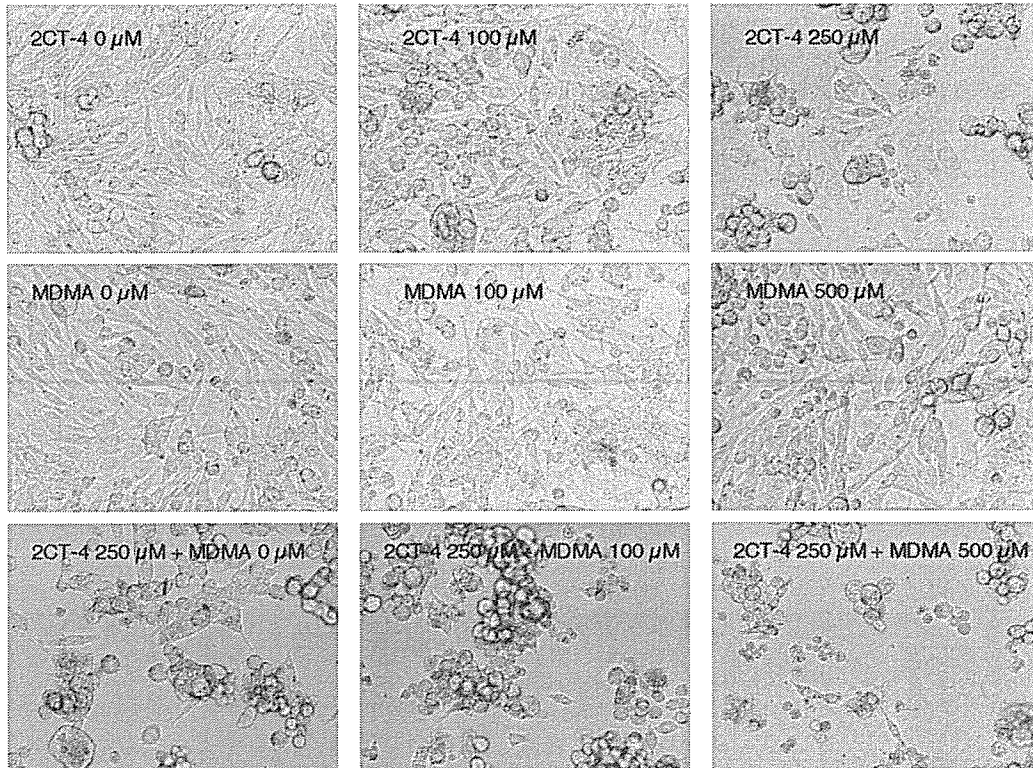


Fig. 9. Photographs of B65 cells treated with 2CT-4 and/or MDMA (final concentration: 0, 100 μM , 500 μM) for 24 hours.

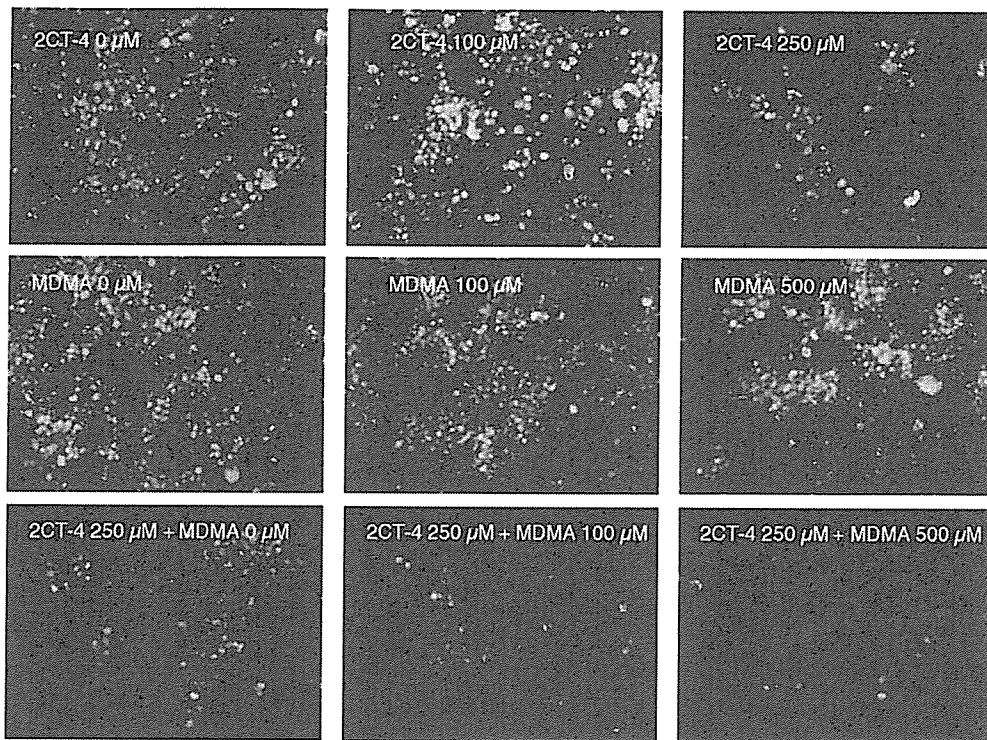


Fig. 10. Nuclear staining of B65 cells treated with 2CT-4 and/or MDMA (final concentration: 0, 100 μM , 500 μM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

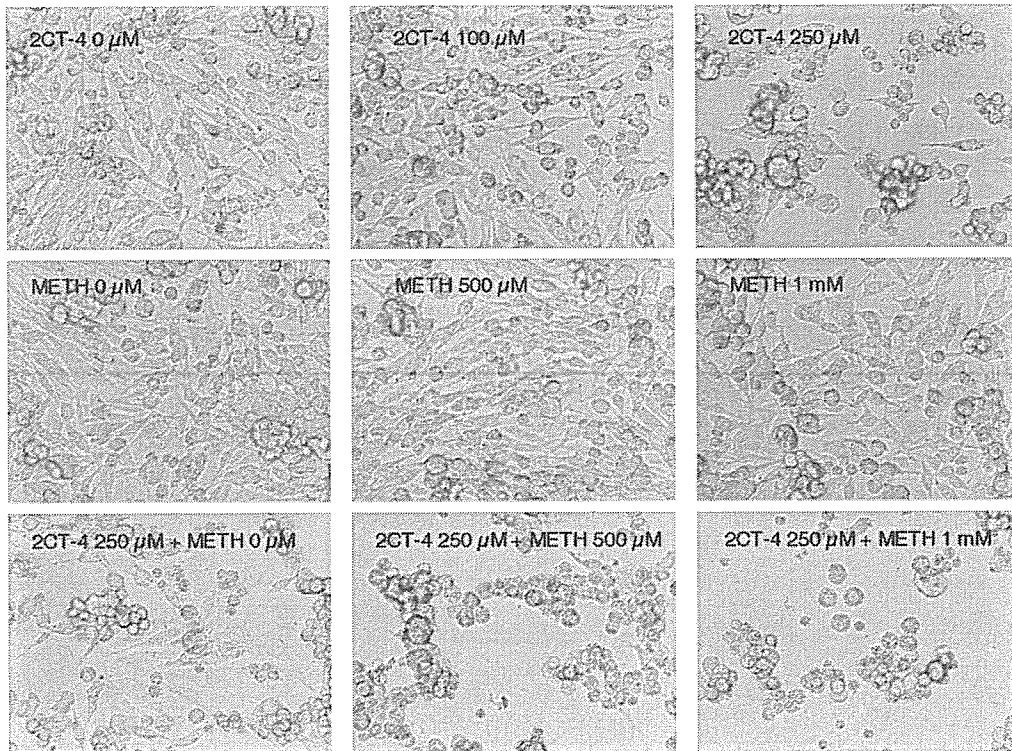


Fig. 11. Photographs of B65 cells treated with 2CT-4 and/or METH (final concentration: 0, 500 μ M, 1 mM) for 24 hours.

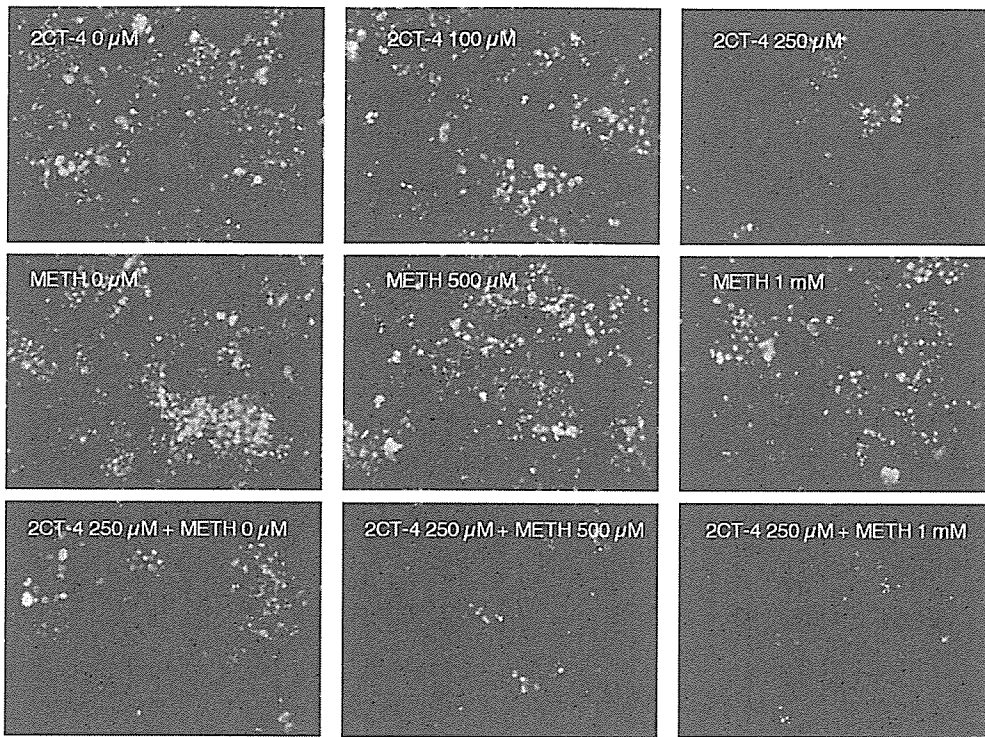
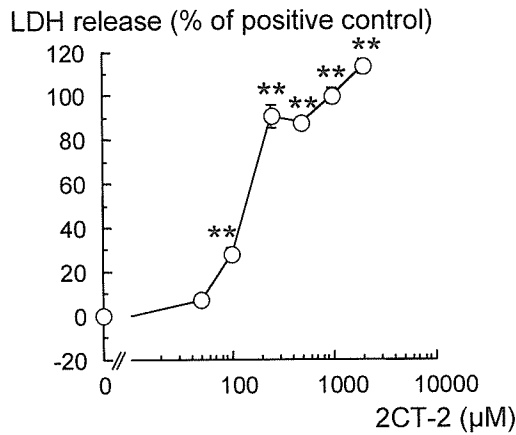
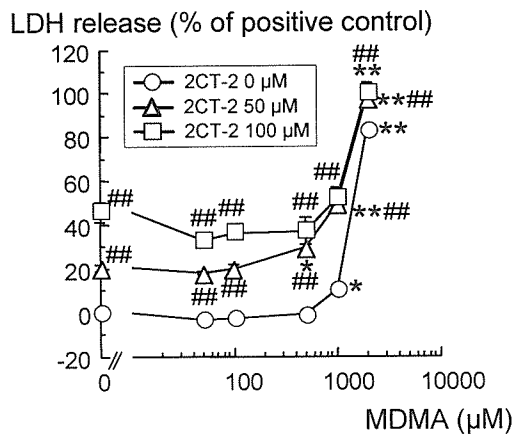


Fig. 12. Nuclear staining of B65 cells treated with 2CT-4 and/or METH (final concentration: 0, 500 μM , 1 mM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

A CATH.a cells



B CATH.a cells



C CATH.a cells

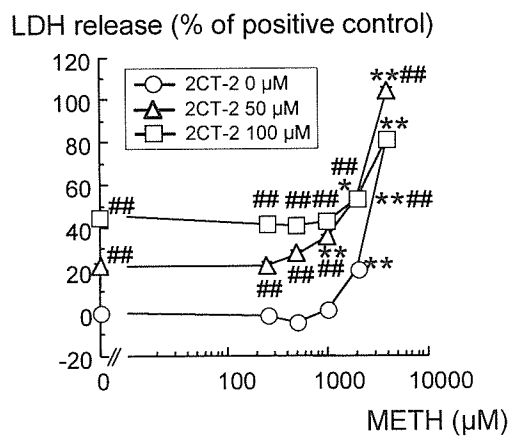


Fig. 13. Changes in released LDH from dopaminergic CATH.a cells after exposure to 2CT-2 (A), 2CT-2+MDMA (B) or 2CT-2+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. * $p < 0.05$, ** $p < 0.001$ vs. each control group without MDMA or METH. ## $p < 0.001$ vs. MDMA/METH-dose-matched control group without 2CT-2.

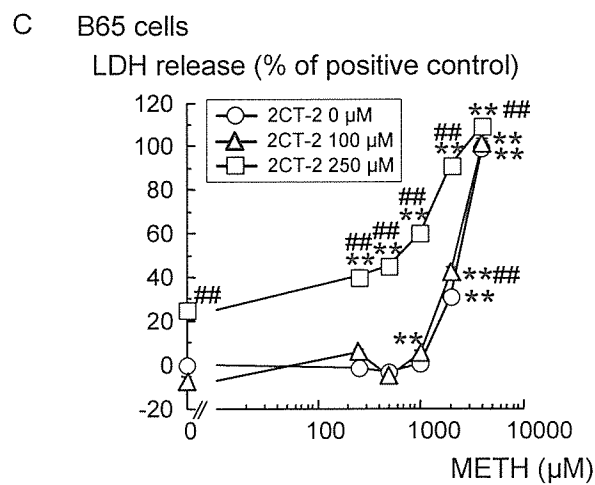
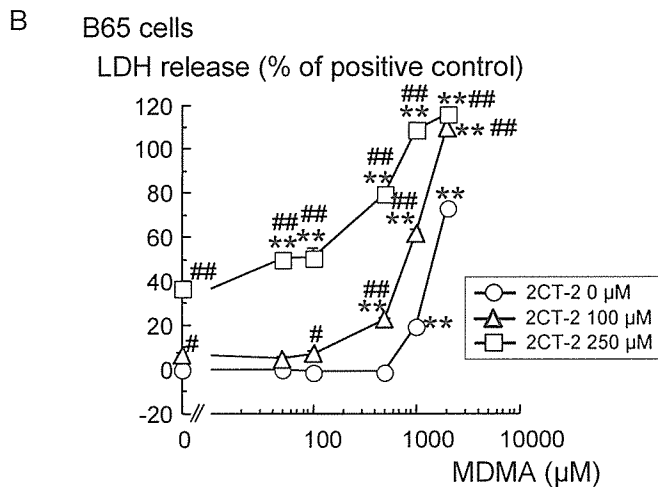
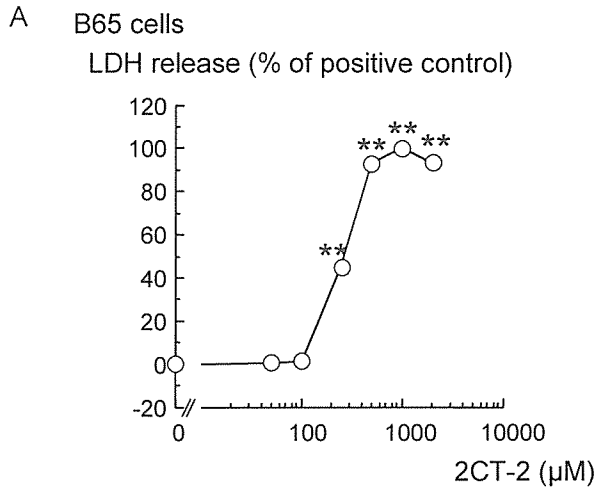


Fig. 14. Changes in released LDH from dopaminergic B65 cells after exposure to 2CT-2 (A), 2CT-2+MDMA (B) or 2CT-2+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. ** $p < 0.001$ vs. each control group without MDMA or METH. # $p < 0.05$, ## $p < 0.001$ vs. MDMA/METH-dose-matched control group without 2CT-2.

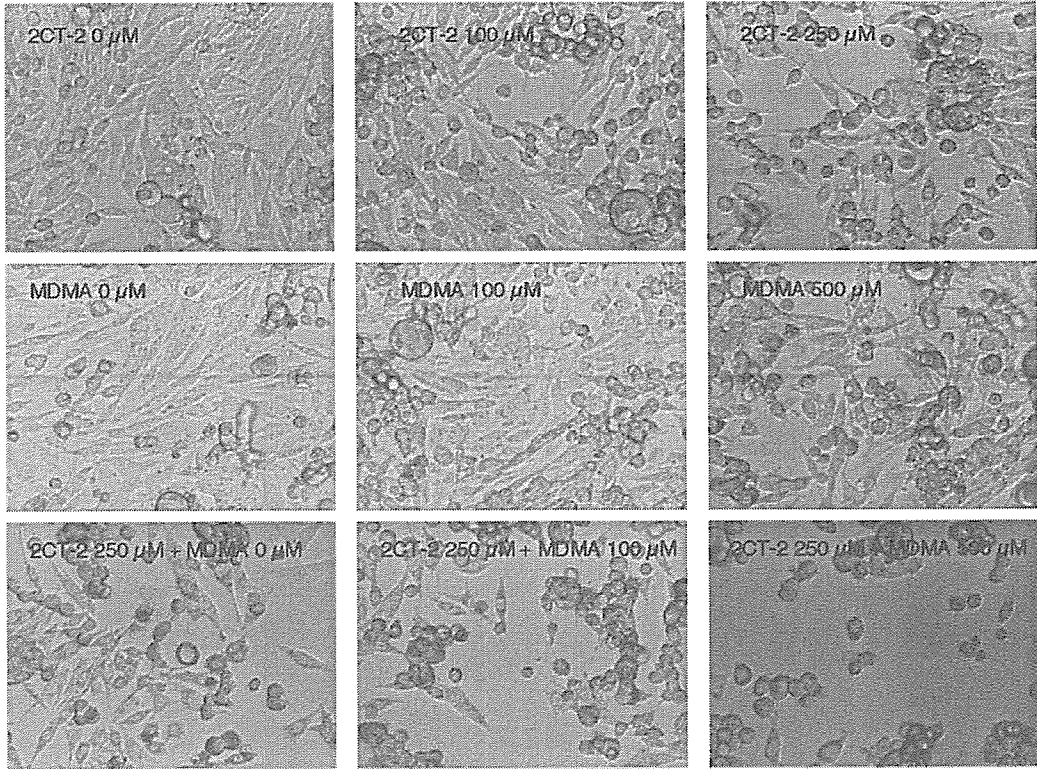


Fig. 15. Photographs of B65 cells treated with 2CT-2 and/or MDMA (final concentration: 0, 100 μM , 500 μM) for 24 hours.

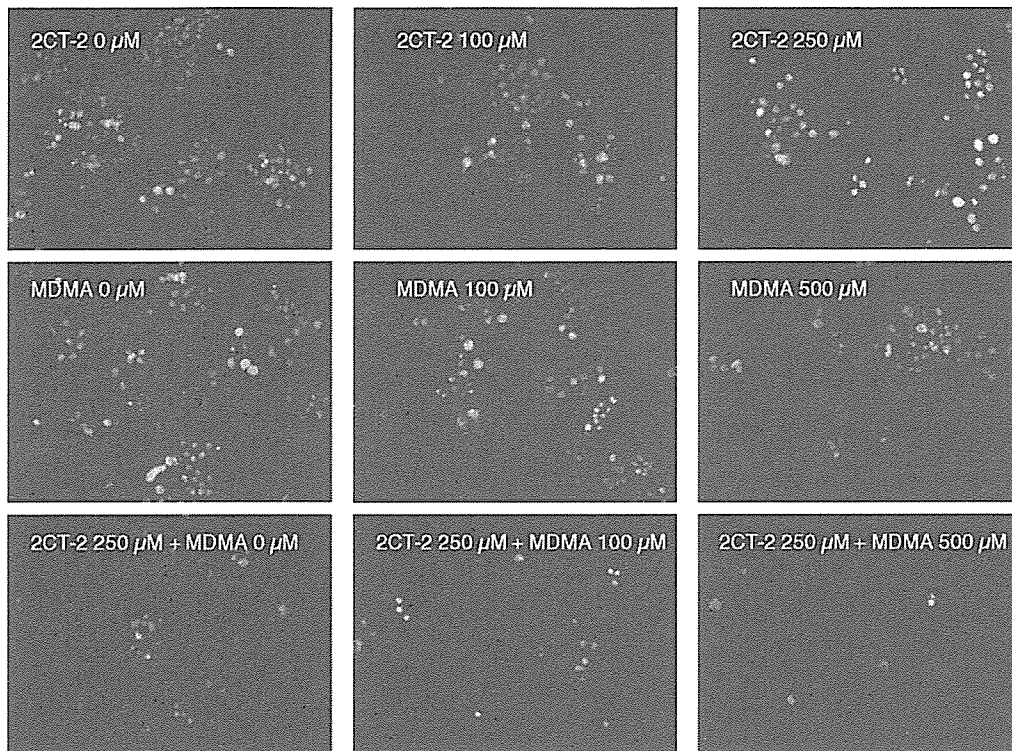


Fig. 16. Nuclear staining of B65 cells treated with 2CT-2 and/or MDMA (final concentration: 0, 100 μM , 500 μM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

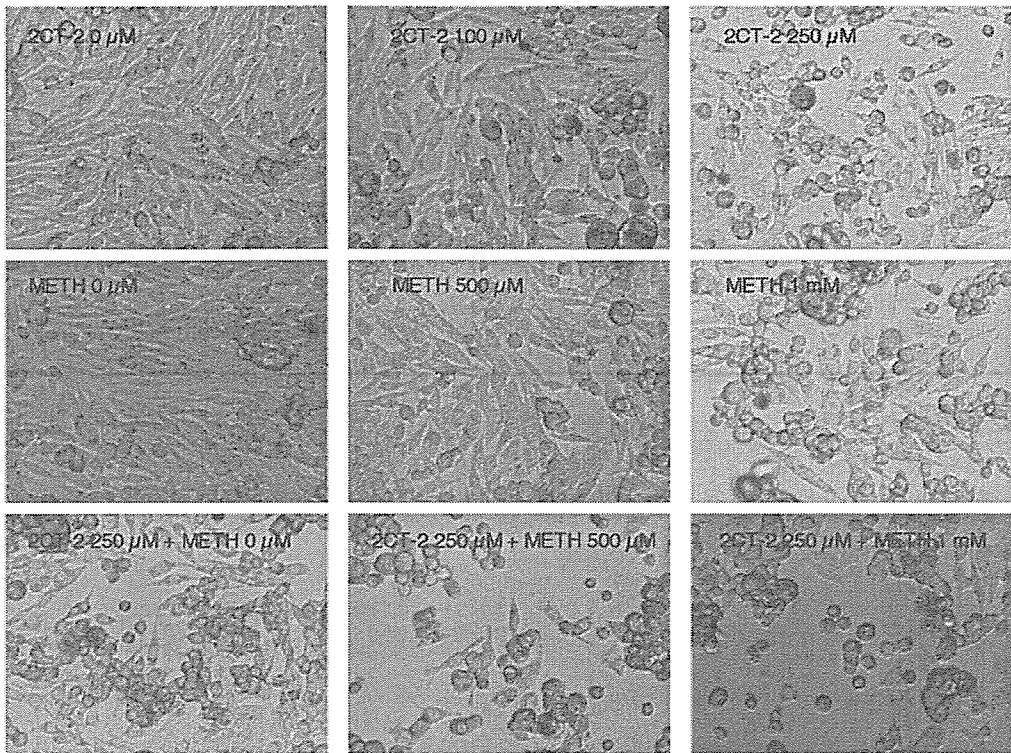


Fig. 17. Photographs of B65 cells treated with 2CT-2 and/or METH (final concentration: 0, 500 μ M, 1 mM) for 24 hours.

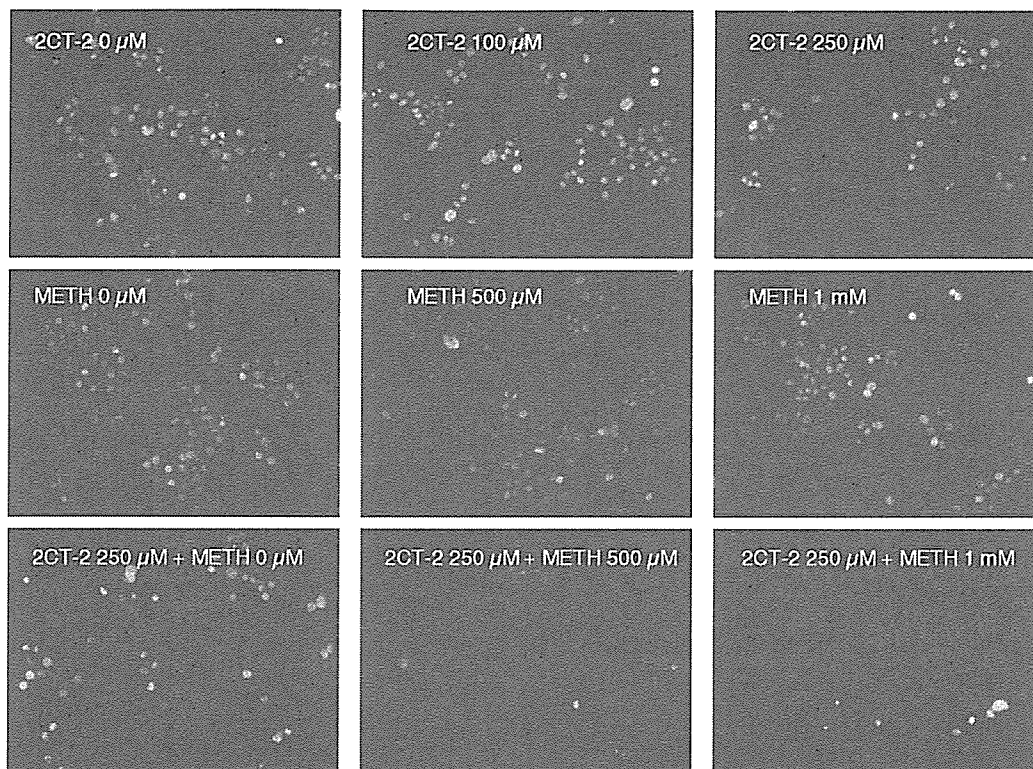
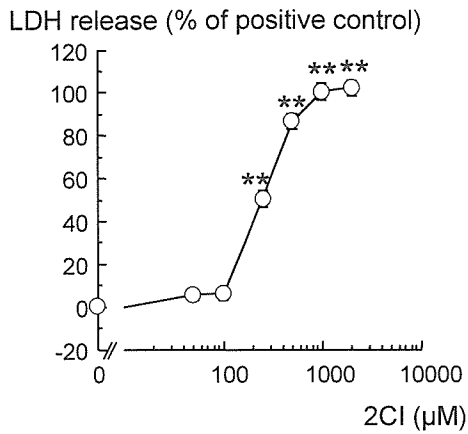
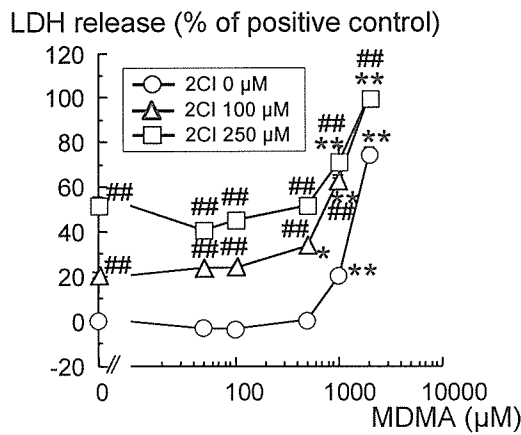


Fig. 18. Nuclear staining of B65 cells treated with 2CT-2 and/or METH (final concentration: 0, 500 μM , 1 mM) for 24 hours. Nuclei were visualized by incubation with Hoechst33342 dye.

A CATH.a cells



B CATH.a cells



C CATH.a cells

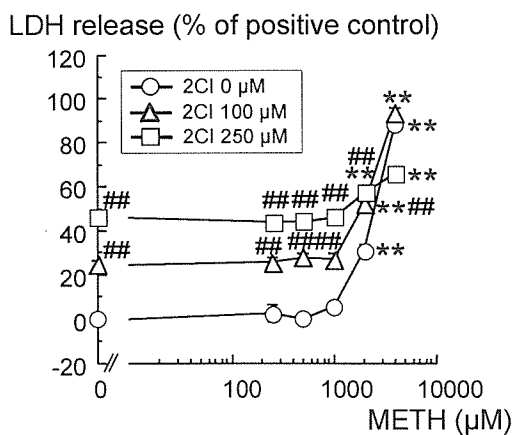


Fig. 19. Changes in released LDH from dopaminergic CATH.a cells after exposure to 2CI (A), 2CI+MDMA (B) or 2CI+METH (C) for 24 hours. Each value mean \pm SEM of released LDH expressed as percentage of Tween-20-treated positive control. * $p < 0.01$, ** $p < 0.001$ vs. each control group without MDMA or METH. ## $p < 0.001$ vs. MDMA/METH-dose-matched control group without 2CI.