



Cadmium Chloride Induced Cognitive Decline in Female Wistar Rats Exposed to Chronic Restraint Stress

**Gbenga Opeyemi Owolabi ^{a*}, Toluwalase Oyenike Oyewale ^a,
Richard Adedamola Ajike ^a, Innocent Effiom Offiong ^b
and Aliyat Nafiu Olanrewaju ^a**

^a Department of Physiology, Faculty of Basic Medical Sciences, Ladoke Akintola University of Technology, Ogbomoso, Oyo State, Nigeria.

^b Department of Anatomy, Faculty of Basic Medical Sciences, Ladoke Akintola University of Technology, Ogbomoso, Oyo State, Nigeria.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: <https://doi.org/10.9734/ajrimps/2024/v13i4279>

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/126535>

Original Research Article

Received: 15/09/2024

Accepted: 18/11/2024

Published: 22/11/2024

ABSTRACT

Cadmium (Cd) is known to have some adverse effects on different biochemical and physiological functions causing neurotoxicity leading to neurodegeneration and increasing the risk factor for neurodegenerative disorders. Restraint stress is also associated with changes in behavioral, neuroendocrine function, and brain morphology. The study aimed to evaluate the effects of cadmium chloride administration and restraint stress exposure on cognitive function of female

*Corresponding author: E-mail: owolabiphysiology@yahoo.com;

Cite as: Owolabi, Gbenga Opeyemi, Toluwalase Oyenike Oyewale, Richard Adedamola Ajike, Innocent Effiom Offiong, and Aliyat Nafiu Olanrewaju. 2024. "Cadmium Chloride Induced Cognitive Decline in Female Wistar Rats Exposed to Chronic Restraint Stress". *Asian Journal of Research in Medical and Pharmaceutical Sciences* 13 (4):118-27. <https://doi.org/10.9734/ajrimps/2024/v13i4279>.

Wistar rats. 24 female Wistar rats (180-220g) were randomly divided into 4 groups (n=6 each): Control (CTL), Restraint stress alone (RSS), Cadmium alone (CCC), Cadmium + Restraint stress (RSC). The experimental groups were subjected to cadmium chloride 100mg/kg orally and restraint stress for 30 minutes using wire mesh. Prior to the animal sacrifice, behavioral tests were carried out to assess the effects of cadmium chloride and restraint stress on cognitive performance of rats. 24 hours post last cadmium administration and restraint stress exposure, all animals were anesthetized and sacrificed. The brain was excised, weighed and homogenized for biochemical analysis (Serotonin and acetylcholinesterase activity). Results showed that there was significant ($p<0.05$) decrease in serotonin level in Cadmium alone group when compared to the control group. The restraint stress + cadmium group showed a significant ($p<0.05$) increase in acetylcholine esterase level when compared control, cadmium alone and restraint stress alone groups. The findings also revealed that Cadmium exposure led to a significant ($p<0.05$) decrease on number of entries in open arms of elevated plus maze. Furthermore, spontaneous alteration (Y maze) was significantly ($p<0.05$) decrease in restraint stress alone, cadmium alone and restraint stress + cadmium groups when compared to the control group. In conclusion, cadmium exposure and restraint stress altered neurotransmission, increased anxiety-like behavior, decreased cognitive abilities, increased alteration in hippocampal architecture and neuronal depletion as revealed in the histological evaluation resulting in cognitive deficits.

Keywords: Cadmium chloride; restraint stress; neurodegeneration; spontaneous alteration; cognitive deficit.

1. INTRODUCTION

Cadmium (Cd) is a highly toxic environmental pollutant absorbed into the human body via the respiratory system, gastrointestinal tract and skin [1]. Exposure to cadmium can be through food, water, cigarette smoke, and air contamination [2]. Its environmental half-life ranges between 25-30 years and accumulation of cadmium can result in a long term toxicological effect on multiple systems [3]. There are various sources of cadmium, it can be found naturally in the earth's crust, agriculture, and produced as a byproduct of mining (zinc, lead and copper) [4,5]. Bioavailability of this toxic heavy metal in the environment is due to increased domestic and industrial indiscriminate disposal into the water bodies and the soil [6]. In this way cadmium gains access into the aquatic animals and crops, this is a secondary route of exposure. The nervous system is particularly vulnerable to cadmium toxicity, as prolonged exposure to cadmium disrupts the normal biochemical and physiological process in the nervous system [7]. The neurotoxic effect of Cd leads to alteration or disruption in various neurological functions such as memory and learning abilities [8,9]. Furthermore, cadmium exposure can impair the synthesis, release and reuptake of neurotransmitters such as serotonin and acetylcholine which play a key role in cognitive process [10].

Stress is any physical or psychological stimuli that disrupt homeostasis [11]. Restraint stress is

a commonly used experimental model for inducing stress response syndromes in animals [12,13]. Various studies have reported that restraint stress is associated with changes in behavioral, neuroendocrine function, and brain morphology, this makes it useful for researching the underlying mechanisms in stress-related neurological disorders.

Cognitive function refers to range mental processes which include memory, learning, attention, decision making and language abilities [14]. Serotonin and acetylcholine are neurotransmitters involved in the modulation of cognitive functions [15]. Exposure to both acute and chronic stress is associated with cognitive impairment [16]. Serotonin and Acetylcholine are neurotransmitters that are involved in the modulation of cognitive function [17]. Alteration in the levels of these neurotransmitters has a significant impact on cognitive function. Nevertheless, studies on the combined effect of cadmium chloride and restraint stress on the cognitive function in female Wistar rats are limited. Therefore, this study aimed to evaluate the effect of cadmium chloride administration and restraint stress on the cognitive function in female Wistar rats.

2. MATERIALS AND METHODS

Chemical and Compounds: Cadmium chloride (Kermel, China), chloroform, Normal Saline, distilled water, was purchased from department of pure and applied science laboratory,

LAUTECH, Oyo State, Nigeria, Buffered formalin was purchased from the department of Anatomy, FBMS, LAUTECH, Oyo State, phosphate buffer saline was purchased from department of science laboratory, LAUTECH, Oyo, Nigeria.

2.1 Experimental Planning and Animals

Twenty-four (24) female rats (180-220g) were used for the study. The rats were kept in a standardized laboratory environment. The rats were acclimatized for two weeks and had free access to clean water and food. The rats were housed in standardized plastic cage maintained between 12-hour light and dark cycle. After acclimatization, the rats were divided randomly into four groups with six (6) rats in each group and the experiment lasted for 21 days. Group I represent the control group while groups II, III, IV served as the experimental groups. The groups designate are: I= Control group (CTL), II= Restraint Stress Alone (RSS), III= Cadmium Alone (CCC) and IV= Cadmium+ Restraint stress (RSC).

2.2 Collection and Preparation of Samples

On the last day of restraint stress induction or/cadmium chloride administration, behavioral assessments using Elevated plus Maze (Guedri et al., 2017) and Y maze (Yoshizak et al., 2020) were carried out. Twenty- four hours after the last treatment regimen, rats were euthanized by placing them in desiccator with chloroform soaked cotton wool. Blood samples were collected via cardiac puncture in the heparinized tubes and then centrifuged at 1500 rpm for 10 minutes. Brain samples were excised, then rinsed in PBS and homogenized over ice in 0.1 M cold sodium phosphate buffer (pH 7.4). The homogenate was centrifuged at 4°C for 10

minutes at 10,000 rpm. The supernatant obtained was aliquot for subsequent biochemical analysis.

2.3 Behavioural Assessments

On the 21st day of cadmium chloride or/ restraint stress conduction, behavioural assessment with Elevated plus maze and Y maze were performed.

Elevated plus maze test: Elevated plus maze was used to measure anxiogenic behavior. The maze was built with two open and two closed arms enclosed with 30cm high walls, which is 10cm wide and mounted 50cm above the ground. Rats were placed at the junction of the open and closed arms, facing the open arm opposite to where the experimenter and their behavior (the number of entries made in the open) was recorded for six (6) minutes. The maze was cleaned with an alcoholic solution followed by wet and dry paper towels following each test (Guedri et al., 2017).

Y maze: Y maze was used to assess the state of working memory of the rats. The arm length of Y maze is about 40cm; arm bottom width is 3cm, arm upper width 13cm, height of the wall 15cm. Each rat was placed in the central area and the number of entries into the arms and alteration were recorded for 10 minutes. The working memory was calculated as number of spontaneous alterations/ number of the total new arm entries (Yoshizak et al., 2020).

2.4 Biochemical Assays

Serotonin (SRO) and Acetylcholinesterase (AChE) activities, were assayed using commercial kits and standardized methods.

Table 1. Animal grouping and experimental procedures

Groups	Administration
Control (CTL)	Rats were given only animal feed and water <i>ad libitum</i> for 21 days.
Restraint Stress Alone (RSS)	Rats were subjected to restraint stress using wire mesh for 30 minutes daily for 21 days.
Cadmium Alone (CCC)	Rats received cadmium chloride (100mg/kg/b.w) orally for 21 days.
Cadmium+ Restraint stress (RSC)	Rats received cadmium chloride (100mg/kg/b.w) daily orally and were subjected to restraint stress using wire mesh for 30 minutes for 21 days.

Statistical Analysis: SPSS (version 16.0) was used for all statistical analysis. All results obtained are expressed as Mean ± Standard Error of the Mean (SEM). Data were analyzed using one-way ANOVA and Duncan's posthoc test for multiple comparisons. P value < 0.05 was considered to be statistically significant.

3. RESULTS AND DISCUSSION

Results

Table 2. Effect of restraint stress and cadmium chloride administration on brain weight in female Wistar rats

Group	CTL	RSS	CCC	RSC
Mean ± SEM	1.33±0.03 ^a	1.48±0.12 ^a	1.32±0.03 ^a	1.20±0.06 ^a

Values are expressed as mean± SEM (n=6). Groups with superscript of different letters are significantly (p<0.05) different from each other. Groups with superscript of same letters are not significantly different from each other. In female Wistar rats exposed to cadmium chloride and restraint stress, there was no significant difference across all groups

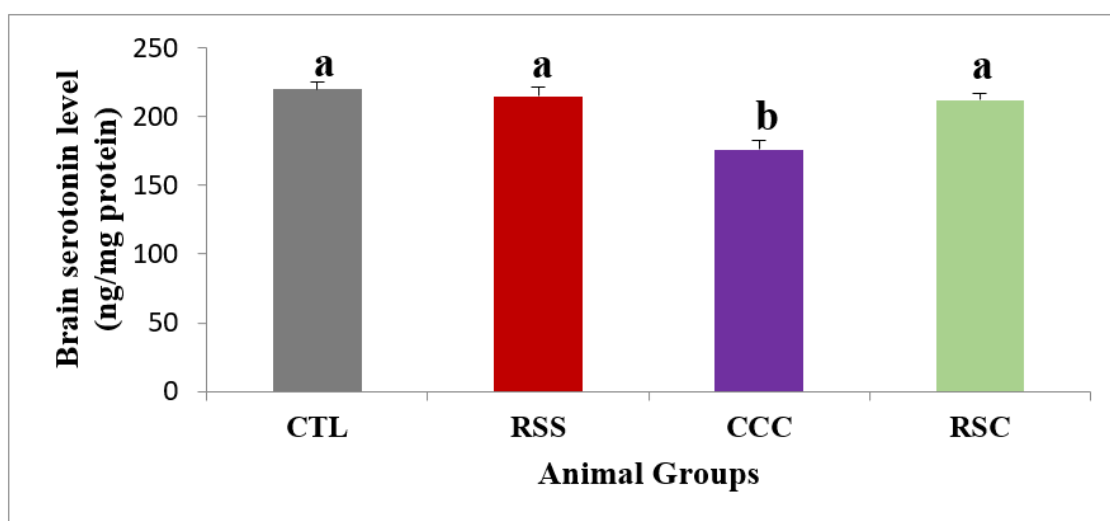


Fig. 1. Effect of restraint stress and cadmium chloride administration on brain serotonin in female Wistar rats

Values are expressed as mean ±SEM (n= 6). Groups with superscript of different letters are significantly (p<0.05) different from each other. Groups with superscript of same letters are not significantly different from each other. Result showed that there was significant (p<0.05) decrease in brain serotonin in CCC group when compared to CTL. There was no significant difference in RSS when compared to CTL. There was no statistical significant difference in RSC when compared to RSC and CCC

Table 3. Effect of restraint stress and cadmium chloride administration on cerebral acetylcholine esterase in female Wistar rats

Group (µmol/ gtissue)	CTL	RSS	CCC	RSC
Mean ± SEM	0.05±0.003 ^a	0.06±0.007 ^a	0.08±0.003 ^a	0.14±0.009 ^b

Values are expressed as mean± SEM (n=6). Groups with superscript of different letters are significantly (p<0.05) different from each other. Groups with superscript of same letters are not significantly different from each other. There was significant (p<0.05) increase in acetylcholine esterase in RSC group when compared to RSS and CCC. There was no significant difference in RSS and CCC when compared to CTL

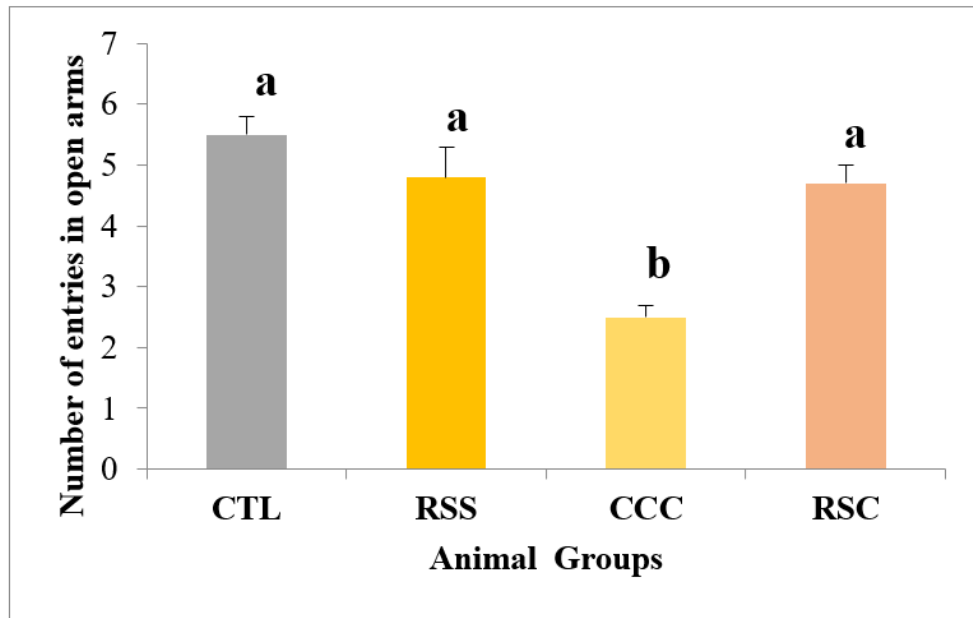


Fig. 2. Effect of restraint stress and cadmium chloride administration on number of entries in open arms of an Elevated Plus Maze in female Wistar rats

Values are expressed as mean \pm SEM ($n=6$). Groups with superscript of different letters are significantly ($p<0.05$) different from each other. Groups with superscript of same letters are not significantly different from each other.

There was significant ($p<0.05$) decrease in the numbers of entries in open arms in CCC group when compared to CTL. However there was no significant difference in RSS when compared to CTL. Also there was no significant difference in RSC when compared to RSS and CCC.

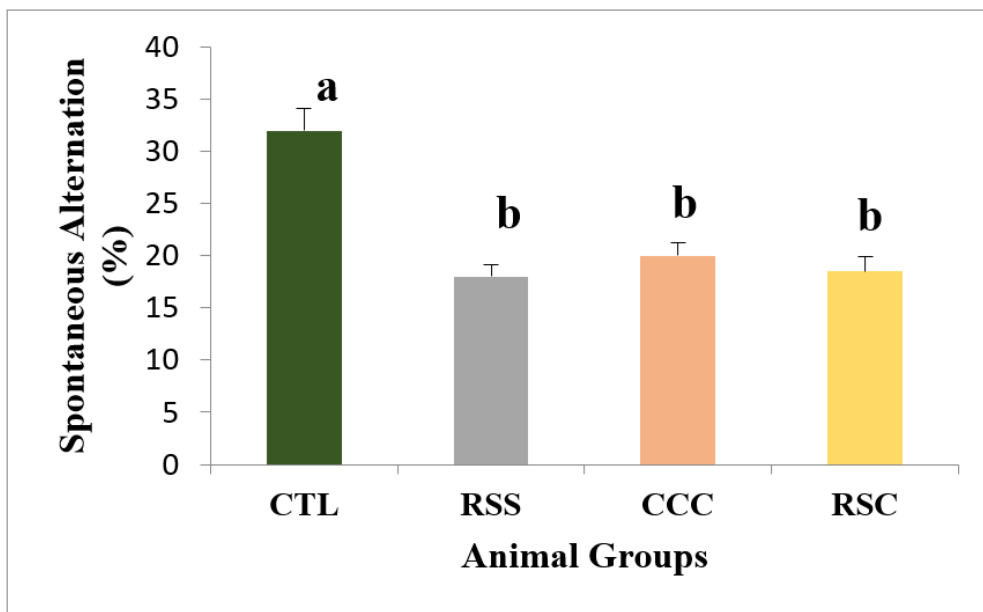


Fig. 3. Effect of restraint stress and cadmium chloride administration on spontaneous alteration (Y maze) in female Wistar rats

Values are expressed as mean \pm SEM ($n=6$). Groups with superscript of different letters are significantly ($p<0.05$) different from each other. Groups with superscript of same letters are not significantly different from each other.

There was significant ($p<0.05$) decrease in spontaneous alteration in RSS, CCC, and RSC group when compared to CTL. However there was no significant difference in RSC when compared to RSS and CCC.

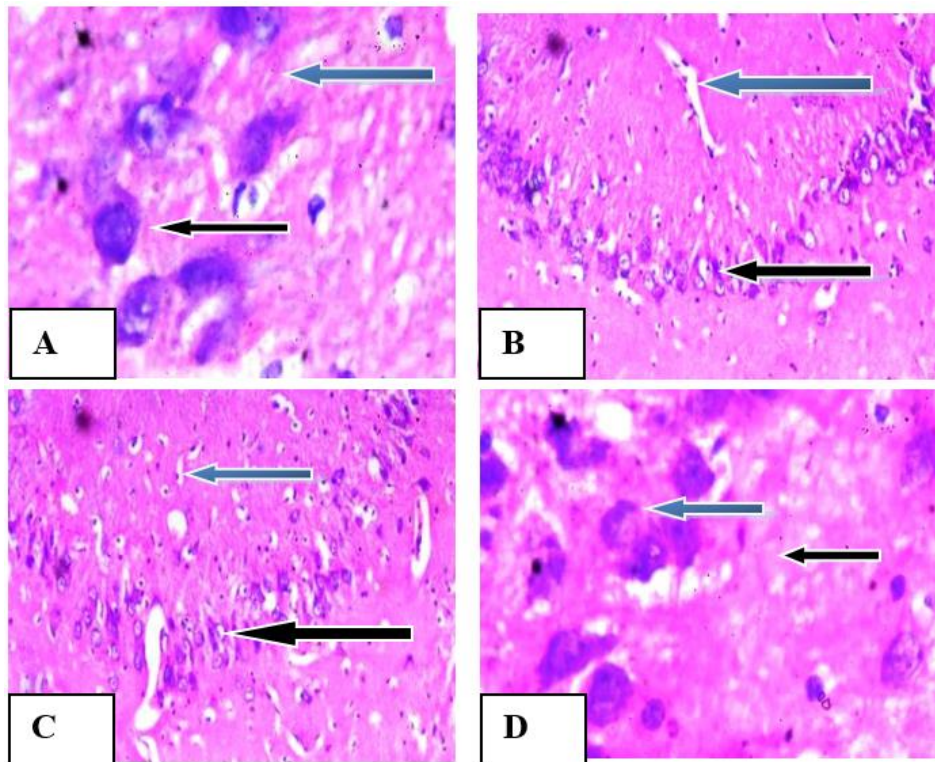


Fig.4. Effect of cadmium chloride administration and restraint stress on hippocampal histology in female Wistar rats

Haematoxylin and Eosin stained micrographs of the hippocampus of control and experimental rats. Histological sections of CTL rats (A) and CCC rats (B) showed a normal structural architecture (blue arrow) with normal neuronal cells (black arrow) while in the RSS (C) rats the hippocampus is fat filled with degenerated neuronal cells (black arrow) and poor structural organization (blue arrow). In the RSC (D), hippocampus with depleted neuronal cells (black arrow) and disorganized structural organization (blue arrow) was seen (H & E, $\times 400$).

Discussion: The toxicological effects of cadmium on the nervous system in both *in vitro* and *In vivo* have been shown extensively in previous studies [18,19]. The brain is one of the organs vulnerable to cadmium toxicity [20]. The hippocampus is the major organ associated with mood and cognition. Chronic stress has detrimental impact on cognition [21].

Organ weight changes are key indicators of toxin induced organ damage [22]. In this study, the brain weight showed no significant difference across all groups. Result observed in restraint stress alone group could be due to integrated stress response (ISR), a cellular defense mechanism that functions to help cells adapt to acute stress by modulating protein synthesis [23]. ISR pathway also plays a crucial role in synaptic plasticity, learning and memory [24]. The combined effects of cadmium and restraint stress exposure did not have significant effect on brain weight, indicating that the study might have been conducted over a short duration for a

significant change to be observed. Serotonin is one of the major monoamine neurotransmitters involved in learning and memory consolidation. In Fig. 1, results observed in the cadmium only group is consistent with previous study of Ojo et al. [25], where there was significant ($p < 0.05$) decrease in brain serotonin following cadmium exposure when compared to control. Reduction in serotonin level leads to disruption in memory consolidation [26]. Decrease in serotonin level may be due to disruption in tryptophan hydroxylase, an enzyme that plays a crucial role in converting tryptophan to serotonin (5-HT), resulting to decreased serotonin synthesis (Rasha et al., 2015). In the restraint stress alone group, there was relative decrease in serotonin level when compared to control group. This result is partially in-line with the study of Oh et al. [18] who reported that restraint stress significantly ($p < 0.05$) decreased serotonin level indicating that, elevation in stress hormone level as a result of hyperactivation of hypothalamic pituitary adrenal (HPA) axis can

disrupt the catecholaminergic and monoaminergic systems leading to a reduction in serotonin level. The study observed the combination of restraint stress and cadmium when compared to cadmium and restraint stress alone group showed no significant difference in serotonin level suggests that the level serotonin might have been reduced to a threshold by either cadmium alone or stress alone.

Acetylcholinesterase (AChE) is an enzyme known to play a major role in cholinergic neurotransmission. It hydrolyzed acetylcholine (a neurotransmitter involved in memory process) in the synaptic cleft of cholinergic synapses and neuromuscular junction. Cadmium toxicity has been implicated in cholinergic neurotransmission disruption [27]. In Table 3, the result observed in the cadmium and restraint stress showed a significant ($p < 0.05$) increase in acetylcholinesterase level when compared to the cadmium alone and restraint stress alone. The combination of cadmium and stress resulted in an increase in acetylcholinesterase level signifying a decrease in acetylcholine. The decrease in acetylcholine disrupts cholinergic signaling, which affect synaptic plasticity and result in cognitive deficit.

According to this study, in Fig. 2 there was significant ($p < 0.05$) decrease in number of entries in open arm of elevated plus maze in cadmium alone group when compared to control group, this is consistent with the previous study of Adeniyi et al. [28]. Decrease in the number of entries in open arm suggests an increase in anxiety level. The neurotoxic effect of cadmium can results in depletion in serotonin level leading to alteration in neurotransmission, as serotonin is a key modulator in anxiety regulation. The combined effect of cadmium and stress showed no statistical significant difference in the restraint stress and cadmium chloride group when compared to the cadmium and restraint stress alone group. However, histological evaluation showed neuronal apoptosis and altered structural organization indicating that if the study had been carried out for a longer duration significant changes might have been observed. Also, this could be as a result of the rats adapting to stress, thereby reducing anxiogenic behavior. This suggests that adaptation to stress might have reduced the synergetic effect of cadmium when combined with restraint stress causing a reduction in anxiety-like behavior.

Spontaneous alteration depends on the natural ability of an animal to explore the novel arm of the Y maze rather than revisiting the previously explored arm. In Fig. 3, the observed statistical significant ($p < 0.05$) decrease in spontaneous alteration in restraint stress alone group when compared to the control group is consistent with the previous studies of Amin et al. [29] and Thongrong et al. [30], suggesting that hyperactivation of the HPA axis as a result of stress could have induced hippocampal degeneration and neuronal apoptosis which result in learning and memory impairment. Result observed in the cadmium alone group is consistent with the findings of Lamtai et al. [31] where spontaneous alteration were significantly decrease following cadmium intoxication in treated groups when compared to their relative controls. Cadmium interferes with serotonergic system resulting in spatial working memory impairment this is evident in result observed in the brain serotonin level. The combined effect of cadmium and stress showed no significant difference in spontaneous alteration when compared to cadmium alone group and restraint stress alone group. This suggests that the individual effect of cadmium or stress might have decrease the spontaneous alteration to a threshold in which the combination of cadmium and stress may not cause further impairment in cognitive behavior this could be attributed to converging mechanism of cadmium and stress.

In this present work, examination of section of hippocampus, by light microscope, did not show notable differences among rats in control and cadmium alone group. On the other hand, Restraint stress induced histological changes in the structural organization and neuronal cell degeneration. This is consistent with the findings of Elfakharany et al. [32]. Furthermore, the neurotoxic effects of cadmium and restraint stress were evident in the altered morphological structure and neuronal depletion of the hippocampus.

4. CONCLUSION

In this study, Cadmium exposure and restraint stress altered neurotransmission, increased anxiety-like behavior, decreased cognitive abilities, increased alteration in hippocampal architecture, increased neuronal degeneration and depletion as revealed in the histological evaluation resulting in cognitive deficits. Continuous bioavailability to cadmium poses

threat globally due to increased environmental pollution and increase in psychological stress intensified by socio-economic crisis, academic and work-related demand and other environmental factor.

CONSENT

It is not applicable.

ETHICAL APPROVAL

All protocols and treatment procedures were done according to the Institutional Animal Care and Use Committee (IACUC) guidelines, in strict compliance with the National Institutes of Health (NIH) guideline for the care and use of laboratory animals.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or editing of this manuscript.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Das SC, Al-Naemi HA. Cadmium toxicity: Oxidative Stress, Inflammation and Tissue Injury. *Occupational Diseases and Environmental Medicine*. 2019;7:144-163.
2. Hayat MT, Nauman M, Nazir N, Ali S, Bangash N. Environmental hazards of cadmium: past, present, and future. In *Cadmium toxicity and tolerance in plants*. 2019;163-183.
3. Genchi G, Sinicropi MS, Lauria G, Carocci A, Catalano A. The effects of cadmium toxicity. *International Journal of Environmental Research and Public Health*. 2020;17(11):3782.
4. Tomza-Marciniak A, Pilarczyk B, Marciniak A, Udała J, Bąkowska M, Pilarczyk R. Cadmium, Cd. mammals and birds as bioindicators of trace element contaminations in terrestrial environments: An Ecotoxicological Assessment of the Northern Hemisphere. 2019;483-532.
5. Haider FU, Coulter JA, Liqun C, Hussain S, Cheema SA, Wu J, et al. An overview on biochar production, its implications, and mechanisms of biochar-induced amelioration of soil and plant characteristics. *Pedosphere*. 2022;32:107–130.
DOI: 10.1016/S1002-0160(20)60094-7
6. Mishra S, Bharagava RN, More N, Yadav A, Zainith S, Mani S, Chowdhary P. Heavy metal contamination: An alarming threat to environment and human health. *Environmental biotechnology: For sustainable future*. 2019;103-125.
7. Tsentsevitsky AN, Petrov AM. Synaptic mechanisms of cadmium neurotoxicity. *Neural Regeneration Research*. 2021;16(9):1762-1763.
8. Elemile OO, Gana AJ, Ejigboye PO, Ibitogbe EM, Olajide OS, Ibitoye OO. Analysis of potentially toxic elements from selected mechanical workshops using the geo-accumulation index and principle component analysis in Omu-Aran Communities, Nigeria. *Environmental Monitoring and Assessment*. 2023;195(2):276.
9. Rezaei K, Mastali G, Abbasgholinejad E, Bafrani MA, Shahmohammadi A, Sadri Z, Zahed MA. Cadmium neurotoxicity: Insights into behavioral effect and neurodegenerative diseases. *Chemosphere*. 2024;143180.
10. Gonçalves JF, Dressler VL, Assmann CE, Morsch VMM, Schetinger MRC. Cadmium neurotoxicity: From its analytical aspects to neuronal impairment. In *Advances in neurotoxicology* (Vol. 5, pp. 81-113). Academic Press; 2021.
11. Chu B, Marwaha K, Sanvictores T, Awosika AO, Ayers D. Physiology, stress reaction. In *StatPearls* [Internet]. StatPearls Publishing; 2024.
12. Atrooz F, Alkadhi KA, Salim S. Understanding stress: Insights from rodent models. *Current Research in Neurobiology*. 2021;2:100013.
13. Şahin Z, Özkürkçüler A, Koç A, Solak H, Koca RÖ, Çakan P, Görmüş ZIS, Kutlu S. An evaluation of the effects of two chronic immobilization stress protocols on depression/anxiety-related behavior in male rats. *Acıbadem Üniversitesi Sağlık Bilimleri Dergisi*. 2019;(3):535-541.

14. Zhang J. Cognitive functions of the brain: perception, attention and memory. arXiv preprint arXiv: 2019;1907.02863.
15. Slater C, Liu Y, Weiss E, Yu K, Wang Q. The neuromodulatory role of the noradrenergic and cholinergic systems and their interplay in cognitive functions: A focused review. *Brain Sciences*. 2022;12(7):890.
16. Forghani N, Hosseinian S, Akhoond-Ali Z, Gholami AA, Assaran-Darban R, Vafae F. Effect of acute and chronic stress on memory impairment and hippocampal oxidative stress following global cerebral ischemia in adult male rats. *Research in Pharmaceutical Sciences*. 2024;19(4):436-446.
17. Handra C, Coman OA, Coman LAURENȚIU, Enache T, Stoleru S, Sorescu AM, Ghită I, Fulga I. The connection between different neurotransmitters involved in cognitive processes. *Farmacologia*. 2019;67(2):193-201.
18. Oh DR, Yoo JS, Kim Y, Kang H, Lee H, Lm SJ, Choi EJ, Jung MA, Bae D, Oh KN, Hong JA. *Vaccinium bracteatum* leaf extract reverses chronic restraint stress-induced depression-like behavior in mice: regulation of hypothalamic-pituitary-adrenal axis, serotonin turnover systems, and ERK/Akt phosphorylation. *Frontiers in pharmacology*. 2018;9:604.
19. Subaraja M, Arokiyaraj S, Mathew P. Neuroprotective effect of huperzine-A against cadmium chloride-induced Huntington's disease in *Drosophila melanogaster* model. *Journal of King Saud University-Science*. 2024;36(8):103319.
20. Arruebarrena MA, Hawe CT, Lee YM, Branco RC. Mechanisms of cadmium neurotoxicity. *International Journal Molecular Sciences*. 2023;24:16558.
21. McEwen BS. Neurobiological and systemic effects of chronic stress. *Chronic stress*. 2017;1:2470547017692328.
22. Lazic SE, Semenova E, Williams DP. Determining organ weight toxicity with Bayesian causal models: Improving on the analysis of relative organ weights. *Scientific Reports*. 2020;10(1):6625.
23. Lawrence RE, Shoemaker SR, Deal A, Sangwan S, Anand AA, Wang L, Marqusee S, Walter P. A helical fulcrum in eIF2B coordinates allosteric regulation of stress signaling. *Nature Chemical Biology*. 2024;20(4):422-431.
24. Helseth AR, Hernandez-Martinez R, Hall VL, Oliver ML, Turner BD, Caffall ZF, Rittiner JE, Shipman MK, King CS, Gradinaru V, Gerfen C. Cholinergic neurons engage the integrated stress response for dopamine modulation and skill learning. *Science (New York, NY)*. 2021;372(6540).
25. Ojo OA, Rotimi DE, Ojo AB, Ogunlakin AD, Ajiboye BO. Gallic acid abates cadmium chloride toxicity via alteration of neurotransmitters and modulation of inflammatory markers in Wistar rats. *Scientific Reports*. 2023;13:1577.
26. Coray R, Quednow BB. The role of serotonin in declarative memory: A systematic review of animal and human research. *Neuroscience and Biobehavioral Reviews*. 2022;139:104729.
27. Gupta R, Shukla RK, Chandravanshi LP, Srivastava P, Dhuriya YK, Shanker J, Singh MP, Pant AB, Khanna VK. Protective role of quercetin in cadmium-induced cholinergic dysfunctions in rat brain by modulating mitochondrial integrity and MAP kinase signaling. *Molecular neurobiology*. 2017;54:4560-4583.
28. Adeniyi PA, Olatunji BP, Ishola AO, Ajonijebu DC, Ogundele OM. Cadmium increases the sensitivity of adolescent female mice to nicotine-related behavioral deficits. *Behavioural Neurology*. 2014;2014(1):360978.
29. Amin SN, El-Aidi AA, Ali MM, Attia YM, Rashed LA. Modification of hippocampal markers of synaptic plasticity by memantine in animal models of acute and repeated restraint stress: implications for memory and behavior. *Neuromolecular Medicine*. 2015;17:121-136.
30. Thongrong S, Surapinit S, Promsrisuk T, Jittiwat J, Kongsui R. Pinostrobin alleviates chronic restraint stress-induced cognitive impairment by modulating oxidative stress and the function of astrocytes in the hippocampus of rats. *Biomedical Reports*. 2023;18(3):1-10.
31. Lamtai M, Azirar S, Zghari O, Ouakki S, El Hessni A, Mesfioui A, Ouichou A, Melatonin ameliorates cadmium-induced affective and cognitive impairments and hippocampal oxidative stress in rat.

- Biological Trace Element Research. 2021;199:1445-1455.
32. Elfakharany SA, Eskaros SS, Azhary NME, Abdelmonsif DA, Zeitoun TM, Ammar GA, Hatem YA. Neuroprotective role of selenium nanoparticles against behavioral, neurobiochemical and histological alterations in rats subjected to chronic restraint stress. *Molecular Neurobiology*. 2024;1-23.

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of the publisher and/or the editor(s). This publisher and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.

© Copyright (2024): Author(s). The licensee is the journal publisher. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/126535>