

Research Article

The Antidepressant Effect of *Chlorella vulgaris* on Female Wistar Rats (*Rattus norvegicus* Berkenhout, 1769) with Chronic Unpredictable Mild Stress Treatment

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ABSTRACT

Depression is a disabling mental disorder, predicted to become the world's number 2 disability by 2020 by the World Health Organization (WHO, 2018). Chronic stress is one of the triggers for depression, causing an imbalance in brain chemicals and antioxidants levels. Although antidepressant is a common treatment, discomforting side effects has compromised its efficacy, prompting the search for alternative medicines. *Chlorella vulgaris* is a microalgae famous for its excellent protein and antioxidant content. In this study, *C. vulgaris* (360 mg/kg p.o.) potency of antidepressant in chronic unpredictable mild stress (CUMS) model of depression in female rats was evaluated compared to amitriptyline (2,25 mg/kg p.o.) for 14 days. Two types of *C. vulgaris* namely cultivation sourced and commercially-sold, were used. Sucrose preference test, forced swim test (FST) and open field test (OFT) were used as depression-like behaviour test to validate *C. vulgaris* effect. Adrenal glands were observed to further understand its effect on the stress organ. The CUMS method produced rats with depressive-like behaviour evidently by reduced body weight, sucrose preference, exploring behaviour in OFT, and increased immobility duration in FST. Furthermore, an increase in adrenal weight, fasciculata zone, and reticularis zone was observed. Both *C. vulgaris* significantly ($p < 0,05$) reversed depressive-like behaviour in rats subjected to CUMS, but not the size of adrenal glands. This finding indicated both types of *C. vulgaris* has the potential to be an alternative antidepressant but because of the short duration of treatment, it's speculated that *C. vulgaris* may not have exhibited enough difference structurally yet.

INTRODUCTION

In modern times, depression has become a disability that is common in Indonesian society. Research by the Ministry of Health in 2018 states that 6.1% of people in Indonesia are depressed. Unfortunately, from all recorded patients, only 9% were treated by professionals (Ministry of Health, 2018). One of the reasons being depression treatment which is still limited to antidepressant drugs that often bring uncomfortable side effects to the body. The common side effects include heart palpitations, insomnia, agitation, headache, and even in extreme cases, stroke (Hedaya, 2011).

One of the alleged cause of depression is chronic stress (Hammen, 2005). Chronic stress can

cause a decrease in serotonin and noradrenaline (Natarajan *et al.*, 2015; Goddard *et al.*, 2009). It can also lead to oxidative stress which causes low levels of antioxidants in the body (Srivastava & Kumar, 2015). In the human body, stress is responded by the adrenal gland. The medulla adrenal will response through the sympatho-adrenomedullary pathway while the adrenal cortex response through cortisol production from the fasciculata zone via the hypothalamus-pituitary-adrenal axis. Cortisol belongs to the glucocorticoid group which increases blood sugar levels in the body for the provision of instant energy. This energy is used to fight or flee from the stressor (fight or flight response) (Chung *et al.*, 2011). Stress response process generates reactive

oxygen species (ROS) as a byproduct. A high ROS concentration in the body can result in various poor health conditions, one of them is lipid peroxidation in brain which inhibits the binding of serotonin (Scapagnini, *et al.*, 2012). ROS effects can be reduced by antioxidant.

Serotonin is synthesized from the amino acid tryptophan, while noradrenaline uses the amino acid tyrosine (Hemat, 2004). Both of these amino acids can be obtained from food, one of which is microalgae such as *Spirulina* sp. and *Chlorella* sp. (Santhanam, 2015; Gershwin & Belay, 2008). According to Bewicke & Potter (2009), *Chlorella vulgaris* contains tryptophan and tyrosine in high content. In addition, *C. vulgaris* is also superior in chlorophyll content. Chlorophyll has been studied to have excellent antioxidant activity (Hsu *et al.*, 2013) so that the high chlorophyll in *C. vulgaris* can give an insight of its potential as a source of antioxidants for humans.

Chlorella vulgaris has long been used as health supplements for humans and are safe for consumption (Klamczynska & Mooney, 2017). However, it is not yet popular, because of the expensive price due to complicated commercial processing, which is the process of removing cell walls from *Chlorella* to make it easier to digest (Zheng *et al.*, 2011). Production with traditional cultivation methods omits said process, providing a *Chlorella* alternative at a more affordable price, but there is no research to determine the difference in performance between the two yet. Seeing the potential for the discovery of alternative medicine to reduce depression from a natural, affordable source, we conducted this study. Two types of *Chlorella vulgaris* were used (cultivated and commercially sold) to further determine the differences that may exist between the two.

MATERIALS AND METHODS

Materials

Twenty-five female Wistar rats (*R. norvegicus*) aged 1-2 months with an interval bodyweight of 100-150 grams were used as animal models and were obtained from Laboratory of Advanced Research and Testing Unit IV, University Gadjah Mada, Yogyakarta. *C. vulgaris* powdered extract was obtained from Blue Green Microalga Technology. Commercially sold *C. vulgaris* tablets were obtained from CNI Sun Chlorella. The antidepressant drug used as a control was amitriptyline (Indofarma). Supporting materials include sucrose powder, ice cubes, distilled water (CV Progo Mulyo), and wood shavings bedding. Animal feed was given in the form of AD II pellets and *reverse osmosis* (RO) water was

given as drinking water, both *ad libitum*, meaning the animals can access these freely. Animal models were kept in groups in a cage with each cage containing 5 rats. Rice husk as standard bedding was changed every 2-3 days. Ketamine (Ketalar): xylazine (Interchemie) cocktail with ratio 1: 1 was used as an anaesthetic and euthanasian agent. Materials needed for making histological slides include neutral buffered formalin (NBF) 10% (CV Progo Mulyo), graded ethanol (CV Progo Mulyo), physiological salt solution (0.9% NaCl), Ehrlich Hematoxylin dye, Eosin Y 1%, distilled water, xylol, toluol, paraffin temperature 57-60°C, Meyer's albumin, object-glass, and cover glass.

Animal models

Animal models were adapted in cages and research environment. Acclimation was carried out for 3 days to minimize stress in animals before the research was conducted so the data has good validity. This stage was carried out in the research room II Laboratory of Advanced Research and Testing Unit IV, University Gadjah Mada. Animal models were weighed once a week individually using semi-analytic scales. The weighing was also done before blood sampling for the determination of the anaesthetic dose and before euthanizing for the calculation of the adrenosomatic index.

Blood sampling was carried out for blood glucose level measurement at the beginning of the study after the acclimation process, after stress induction and after the treatment period. Before sampling, animals were anaesthetized first with ketamine (50 mg/kg BW). Blood sampling was done by inserting one end of the micro hematocrit inward *medial canthus* of the animal's eye, (forming a 30-degree angle to the nose) and was then stored in 1 mL microtube. Blood glucose measurements were made with a glucose meter (EasyTouch) with corresponding single use test strips.

Stress Induction by Chronic Unpredictable Mild Stress (CUMS) Method

Animal models stress was induced by the method CUMS (Hu *et al.*, 2017; Zhang *et al.*, 2014). The stressors given were described as follows:

- a. Treatment of cold water, animal models were put in a water tank with a water depth of 25 cm and a water temperature of 5°C for 3 minutes. After treatment, animals were allowed to dry themselves inside a dry cage with fresh wood-shavings bedding while illuminated with a lamp to keep the animals warm.
- b. Treatment of warm water, animals were put in a water tank with a water depth of 25 cm and a water temperature of 45° C for 3 minutes. After

treatment for this stressors is the same as cold water treatment.

- c. Treatment of wet cages, animals were placed in cages with wet wood shavings bedding as high as 5 cm. This treatment was given for 24 hours. After treatment, the wet bedding was removed and replaced with dry, standard bedding.
- d. Dark-light cycle reversal, animals were exposed to light at night (18.00-06.00) and in a dark place during the daytime (06.00-18.00).
- e. Sound wave exposure, animals were exposed to the ultrasonic sound for 12 hours.
- f. Treatment of tilted cage, animal's cages were tilted 45° for 24 hours.
- g. Food and water deprivation, for 24 hours animals were not fed and the water bottles were removed from the cage. The bedding was replaced beforehand to remove food burrowed in it by the animals. After treatment, feeding was continued normally and water bottles were given back.

These stressors were carried out to all groups except control for 42 days, by giving 1 type of stressor per day. The order of stressors was maintained so that the same stressors did not occur for 2 consecutive days. On day 42, animals underwent behavioural tests to determine their stress state.

Sucrose Preference Test (SPT)

The test was carried out at the beginning of the study, after stress induction and after the treatment period. The test was started by habituation of animals to two drinking bottles for 3 days. After 3 days, one of the bottles was filled with 2% sucrose solution and the other was filled with RO water. Animal cages were then not equipped with drinking bottles for 12 hours, after which both bottles were provided together. The position of the bottle was exchanged every 12 hours and the consumption of sucrose solution and RO water was measured after 24 hours. Sucrose preference was then calculated based on the following formula: $\text{Sucrose preference (\%)} = (\text{Sucrose solution consumption} / \text{Total water and solution consumption}) \text{ mL} \times 100\%$ (Shukkoor *et al.*, 2016)

Open Field Test (OFT)

The test was done after stress induction and after the treatment period and refers to the method used by Shukkoor *et al.* (2016) with changes. Animals were placed in the arena of the lidless opaque plastic box (30 cm x 60 cm with height 40 cm) without bedding individually for 10 minutes. The movements and behaviours of the animals were recorded using a camera during this period. After 10 minutes, the animal was removed from the arena.

Forced Swim Test

This test was done immediately after OFT was done. This protocol follows the literature from Slattery & Cryan (2012). This test consists of two sessions separated by a 24 hour period. At each session, animals were placed in a water tank made of clear acrylic (diameter 20 cm and height 50 cm), which was then filled with water at 26°C. Water filled the cylinder to a depth of 25 cm so that the entire body of the animal can be submerged and to prevent the animal's front limbs from reaching the edge of the container and running away.

The day before the test, 15 minutes of the pretest was carried out to habituate animals to the test environment. The next day, the animal was individually put on a water tank to swim for 5 minutes and immobility time was observed. The animal was then transferred to a dry cage containing fresh bedding and illuminated by the lamp as warmer. After the animal's body was relatively dry, it was returned to their respective cages. Faecal boli that may be present in the water tank were removed, and water was replaced every 3 test or if necessary.

Per oral Treatment

This study was designed randomly with 5 treatment groups. The treatment group consisted of 1 control group, 1 stress control group, 1 positive control group treated with 2,25mg/kg BW Amitriptyline and 2 groups of variations in treatment, namely treatment of cultivated *C. vulgaris* and treatment of commercial *C. vulgaris*, both given at 360mg/kg BW dosage. Each group consists of 5 replications. The duration of treatment is 14 days after stress induction. Treatment was given every morning between 9:00 and 11:00.

Euthanasia and surgery

On day 56 (or after the treatment period), the animals were terminated by the exsanguination method. Adrenal glands were cleaned and used to find out the adenosomatic index according to the formula: $\text{adenosomatic index} = (\text{total organ weight} / \text{total body weight}) \text{ gram} \times 100\%$ (Avinashe, 2013). Adrenal glands were then moved to neutral buffer phosphate (NBF) 10% solution to fixate it. Then the histological preparations were made with Hematoxylin-Eosin staining.

Data Analysis

Obtained data were analysed using one-way ANOVA with a confidence level of 95% ($\alpha < 0,05$) and Duncan Multiple Range Test as post-hoc test. Statistical analysis was done with SPSS 23 program and graphs needed were made using Microsoft

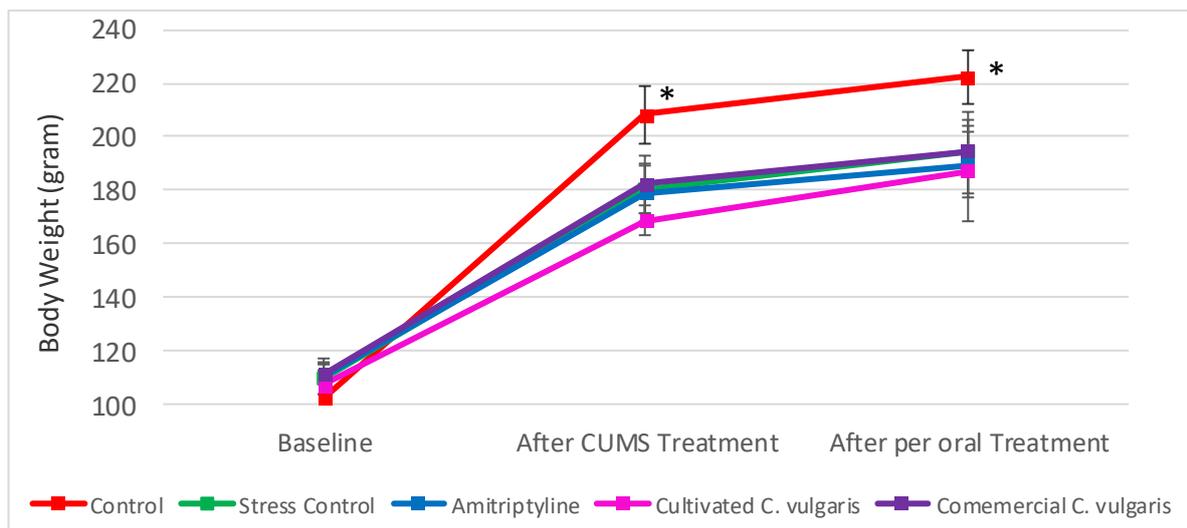


Figure 1. Weight gain in female Wistar rats during the study. * $p < 0.01$, when compared with CUMS, treated groups; one-way ANOVA followed by Duncan *post-hoc* test.

Excel. Qualitative data in the form of roaming videos were analysed using idTracker software (created by Perez-Escudero *et al.* (2014) and can be downloaded free via the web <http://www.idtracker.es/>) to see trajectory patterns made by the animal during the test period.

RESULTS AND DISCUSSION

This study has passed the required ethical feasibility test by the Animal Ethics Committee of Universitas Gadjah Mada before the study was conducted with a certificate number: 00042/04/ LPPT/VI/2018

Body Weight

Bodyweight is data that can be linked to animal physiological conditions. In this study, animal body weight was weighed at the beginning of the study, after CUMS treatment and after *per oral* treatment. Changes that might occur were observed.

Figure 1 showed that all group experienced weight gain from the beginning of the study until the end of CUMS treatment, with control group having a significant difference ($F=8.504$; $df=4.18$; $p < 0.01$) compared to all other groups. This can be caused by stress in CUMS treated groups which caused the tendency to consume fewer amounts of food (hypophagia) (Rabasa & Dickson, 2016). Stress will cause the body to release various stress hormones like CRH, ACTH, and cortisol which in turn will prepare the body for the "flight or fight" condition. This condition includes increased blood pressure, gluconeogenesis rate, and blood supply to the muscles, heart, and brain. Therefore, organ activities that require energy but not related to this response such as digestive and reproductive organs will be temporarily inhibited (Yau & Potenza, 2014). *Per*

oral treatment did not cause a difference afterwards compared to the stress control group (Figure 1). This could be due to the short period of treatment which did not provide enough time for structural change to develop. Furthermore, the adaptation to stressors in the form of increased calorie use efficiency in the stress control group (Rabasa & Dickson, 2016) and late age factor could also play a role in means of a slower metabolism, which could explain the minimum weight gain observed (Barnett, 2017).

Blood Glucose Levels

Blood glucose level can describe stress status in rat. Figure 2 showed that the stress treatment given did not have a significant effect on blood glucose.

During the initial period of the study until stress conditions were reached, all groups experienced elevated blood glucose levels with insignificant differences. After going through *per oral* treatment according to the research design, the control and stress control group had a decreased blood glucose value whereas the remaining treatment group experienced an increase.

In theory, stress can increase glucose levels acutely through the work of the epinephrine and cortisol hormones which stimulate liver glycogenolysis and gluconeogenesis. These processes will return to normal conditions (homeostasis) after the stressor ends (Preiser, 2016) so that an increase in blood glucose is also possible but when the measurement was done, the animal has been able to restore its blood glucose level to its initial state.

Besides that, it was found that microalgae treatments can increase blood glucose levels, seen from the elevated value that happened after *per oral*

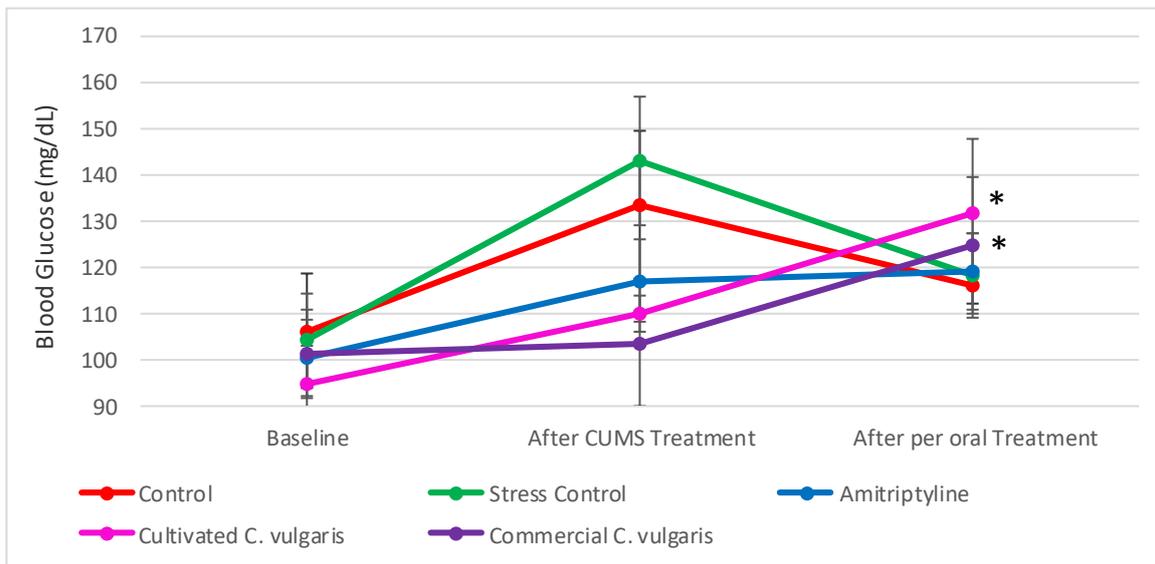


Figure 2. Blood glucose level of female Wistar rats at the beginning of the study, after CUMS treatment and after *per oral* treatment. * $p < 0.05$ when compared to its respective group on its previous period (Baseline and after CUMS treatment).

period ($F=10.829$; $df=2.12$; $p < 0.05$). According to Safi *et al.* (2014), the carbohydrate content of *C. vulgaris* is quite high, at around 55% of its dry weight, with the predominance of carbohydrates in the form of sugar as a constituent of its cell walls like galactose, glucose, xylose, arabinose, and mannose. Cultivated *C. vulgaris* provide a higher increase than commercial *C. vulgaris* because cultivated *C. vulgaris* are still intact with the cell wall, whereas in the commercial one which has been factory processed, this cell wall has been removed.

Sucrose Preference

The preference of animals against sucrose was measured using the sucrose preference test (SPT). Low preference reflects anhedonia behaviour, while high preference is generated by model animals under normal conditions.

On Figure 3, it can be seen that the sucrose preference of the control group stayed relatively high, unlike the other four groups which experienced a decline. This shows the anhedonia behaviour on groups with CUMS treatment which is one of the signs of depression.

After going through *per oral* treatment, the sucrose preference of animal in the treatment group of both amitriptyline and microalgae return to its original state and approached sucrose preferences of the control group. The animal in the stress control group had its sucrose preference decreased and differed significantly from other groups and even that group itself in the previous week ($F=7.33$; $df=4.18$; $p < 0.01$), indicating that sucrose preference in this group continues to decline. Rats naturally have a tendency to consume sweet drinks, so that a decrease in sucrose preference can be assumed as

anhedonic behaviour (Serchov *et al.*, 2016).

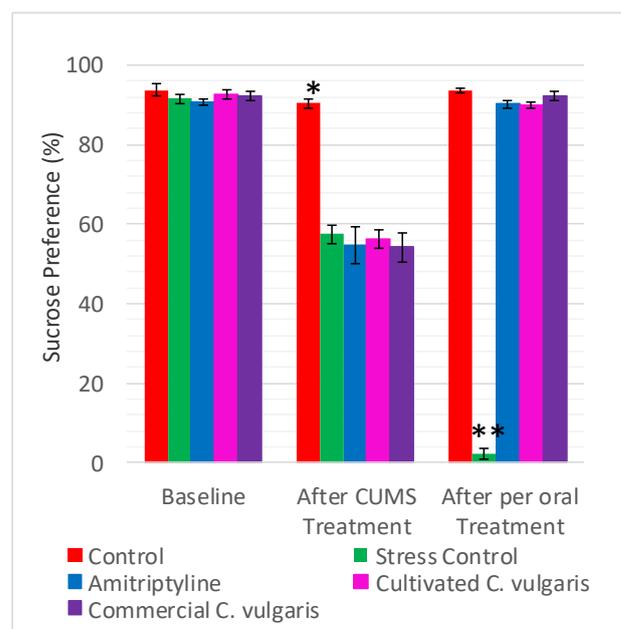


Figure 3. Sucrose preference of female Wistar rats at the beginning of the study, after CUMS treatment and after *per oral* treatment. * $p < 0.01$ compared with CUMS treated groups; ** $p < 0.01$ compared with control and *per oral* groups; one-way ANOVA followed by Duncan *post-hoc* test.

Amitriptyline was chosen because of its efficiency compared to other types of antidepressant drugs in treating depression (Cipriani *et al.*, 2018). Amitriptyline works by increasing the concentration of serotonin and norepinephrine neurotransmitters at synapses (Gupta *et al.*, 2015). When compared with groups treated with cultivated *C. vulgaris* and commercial *C. vulgaris*, group treated with cultivated *C. vulgaris* had a similar preference while the group

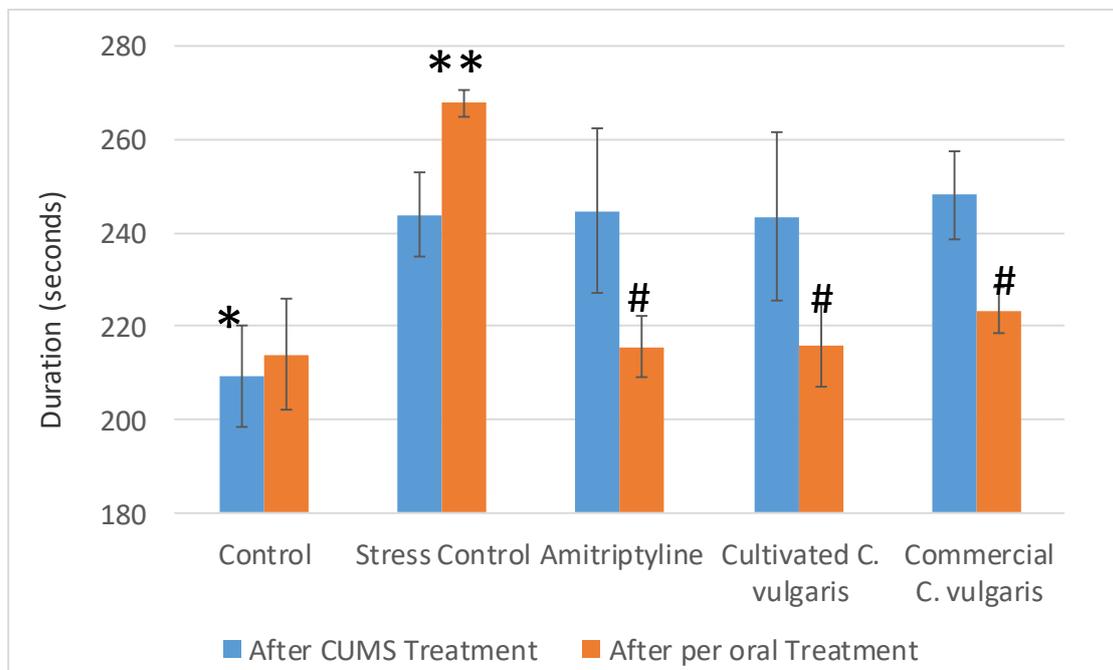


Figure 4. Immobile duration during FST on female Wistar rats after CUMS treatment and *per oral* treatment. * $p < 0.01$ compared with stress control and *per oral* treated groups; ** $p < 0.01$ compared with control and *per oral* treated groups # $p < 0.01$ compared with the same group respectively after CUMS treatment; one-way ANOVA followed by Duncan *post-hoc* test.

treated with commercial *C. vulgaris* had a higher preference value. This indicates that both type *C. vulgaris* can reduce anhedonia behaviour in animal models.

Forced Swim Test

Forced Swim Test (FST) is a common behavioural test on rodents, conducted with the aim of knowing whether a substance or a treatment has the antidepressant effect.

Based on the results on Figure 4, it was discovered that the control group had significantly ($F=5.415$; $df=4.18$; $p<0.01$) shorter duration of immobility compared with other groups. This duration has not differed much on either stress period or treatment period, which showed the consistency of animal conditions in this group throughout the study.

The negative control group had an increased immobile duration from the stress period to the treatment period. This was different from the group with *per oral* treatment, all of which showed a decline ($F=28.576$; $df=4.18$; $p<0.01$). Immobile behaviour on FST indicates despair in rodents when it realized its effort to escape from the tank is in vain, and this behaviour constitutes synonymous behaviour observed in depressed humans (Belovicova *et al.*, 2017). Amitriptyline treatment as an antidepressant caused a decrease in the duration of immobility. The same decline was observed in the group with microalgae treatment. These results indicate both

type *C. vulgaris* has the potential to reduce the duration of immobility behaviour on FST.

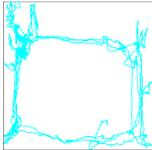
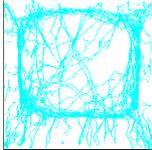
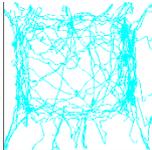
Open Field Test

Open Field Test (OFT) has been used extensively to measure anxiety behaviour in rodents. The principle of this test involves experimental animals in an unknown environment and observing the response of animals to this condition. Rodents generally choose to be on the edge and sides of the arena, avoiding the centre, unless given a substance with an anxiolytic or similar effect. This test was carried out in this research, in addition to give a more complete picture of the psychological condition of experimental animals, also because the symptoms of anxiety and depression are often times similar to each other (Tiller, 2013). The roaming of the animals was projected into a pattern and each pattern was quantified following the scale provided in Table 1.

From Figure 5 it was stated that control group has a lower roaming value than other groups, significantly in stress conditions ($F=24.657$; $df=4.18$; $p<0.01$) but not significant for animals with amitriptyline treatment at the end of study ($F=15.933$; $df=4.18$; $p<0.01$).

Chronic stress generally decreases the level of animal activity in the new environment, and this symptom can be eliminated by using antidepressants (Wang *et al.*, 2014). Group treated with microalgae produce more complex roaming patterns compared to the stress control group, but not significant

Table 1. Quantification of the trajectory pattern of rats in the OFT

Scale	Pattern	Interpretation
5		The pattern that is formed is thick on the edge and empty in the middle
4		The pattern that is formed is thick on the edge and slightly in the middle
3		The pattern formed is relatively balanced between the middle and the edges
2		The pattern that formed has many lines in the middle, on the edge is not too thick
1		Thick pattern throughout the image

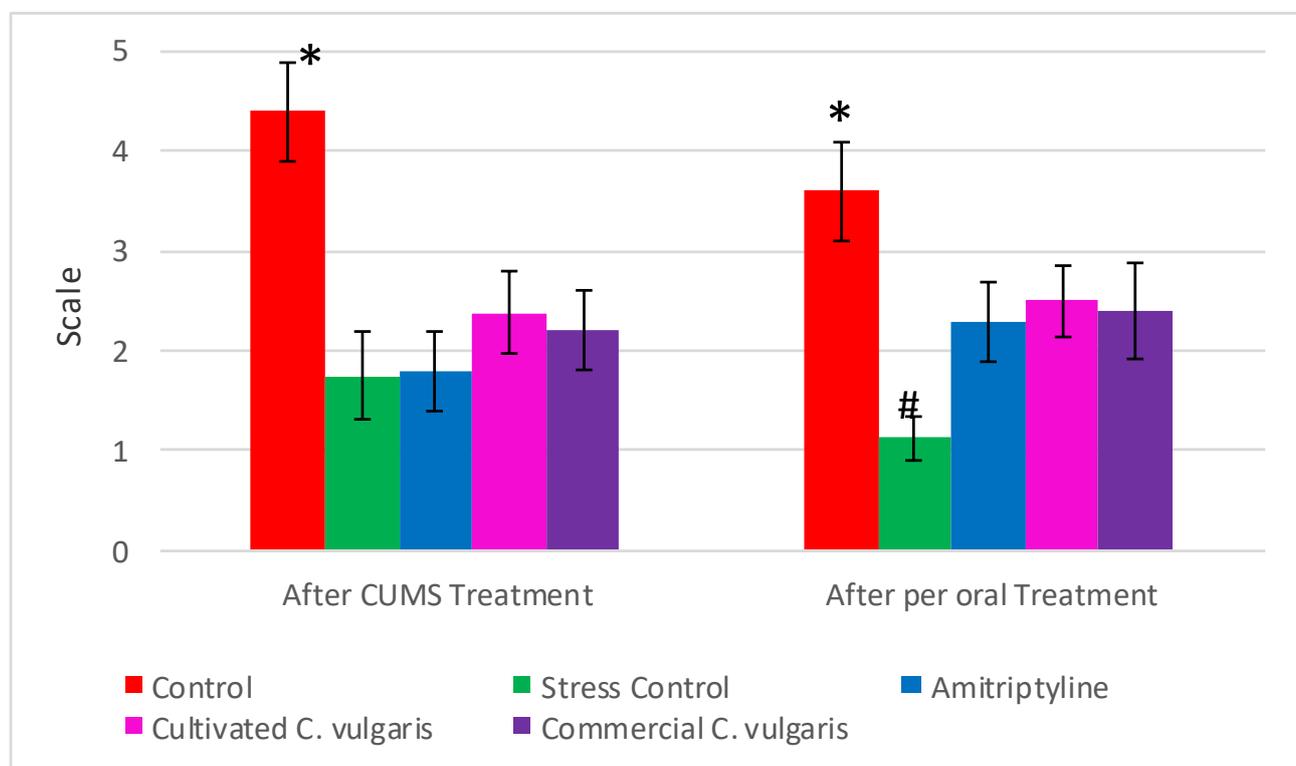


Figure 5. Comparison of the roaming behaviour of female Wistar rats in OFT after CUMS treatment and per oral treatment. * $p < 0.01$ compared with stress control and *per oral* treated groups; # $p < 0.01$ compared with control and *per oral* treated groups.

Table 2. Adrenal gland of female Wistar rats after 56 days of study.

Parameters	Control	Stress Control	Amitriptyline	Cultivated <i>C. vulgaris</i>	Commercial <i>C. vulgaris</i>
Adrenal Weight (grams)	0.045±0.012 ^x	0.067±0.005 ^z	0.052±0.004 ^{xy}	0.055±0.01 ^{xyz}	0.063±0.008 ^{yz}
Adrenosomatic Index (%)	0.020±0.006 ^x	0.035±0.004 ^y	0.028±0.003 ^y	0.029±0.002 ^y	0.032±0.004 ^y
Cortex					
Zona Glomerulosa	11.76±0.91 ^{yz}	12.40±1.71 ^z	10.53±1.33 ^{xy}	10.34±0.55 ^{xy}	9.66±0.75 ^x
Zona Fasciculata	70.66±2.3 ^x	75.80±4.18 ^y	77.27±2.59 ^y	78.49±2.25 ^y	76.87±2.37 ^y
Zona Reticularis	17.64±2.02 ^y	11,66±1,07 ^x	11.30±1.01 ^x	10.45±1.09 ^x	12.57±1.18 ^x
Cortex Medulla Ratio	48.23±5.28 ^x	51.73±7.18 ^x	47.45±6.1 ^x	51.73±4.15 ^x	52.76±4.83 ^x

x, y, z: different notation indicates significant difference at $p < 0.05$

compared to the amitriptyline treatment group. This could mean that microalgae treatment both cultivated and commercial has the potential to increase motor activity in the OFT.

and adrenal hypertrophy, reflecting the overwork of this gland (Ulrich-Lai *et al.*, 2006).

Table 2 stated that the adrenosomatic index of animals in the control group was significantly lower than the other groups ($F=3.987$; $df=4.18$; $p < 0.05$). The negative control group has the largest index, however, not significant with *per oral* treated group. No significant results between stress control groups and the amitriptyline treatment group were possible because the drug had not taken effect structurally. Zoladz *et al.* (2013) found a decreased adrenal weight compared to the stressed rat after 1 month period of amitriptyline administration. The same fact could possibly be the reason why both type *C. vulgaris* failed to give a significant effect.

Based on histological observations, it was found that the percentage of zona fasciculata in the group with CUMS treatment was significantly greater ($F=4.473$; $df=4.18$; $p < 0.05$) than the control group, but zona reticularis ($F=18.05$; $df=4.18$; $p < 0.05$) was significantly smaller (Table 2, Figure 6). Zona fasciculata is where the hormone cortisol formed, which is stimulated by the ACTH hormone from the anterior pituitary. Cortisol plays a role in stress responses such as increasing blood pressure and energy metabolism. Chronic stress causes an increase in cortisol production so that the fasciculata zone enlarges (Ulrich-Lai *et al.*, 2006). Zona reticularis plays a role in reproduction system by producing sex hormone. Stressful events may lead to the inhibition of bodily systems which do not directly support survivability, causing energy supply to these systems to reduce until the stress has passed away (Yau & Potenza, 2014). The ratio between the cortex and medulla between groups is not significantly different. This is presumably due to enlargement of the cortex followed by enlargement of the adrenal medulla. The adrenal medulla is a place for the conversion of tyrosine amino acids into catecholamines such as epinephrine and norepinephrine. These catecholamines function in the fight or flight response and cause reactions in the body such as increased heart rate, blood pressure, narrowing of blood vessels in the skin and the gastrointestinal system and increased metabolism.

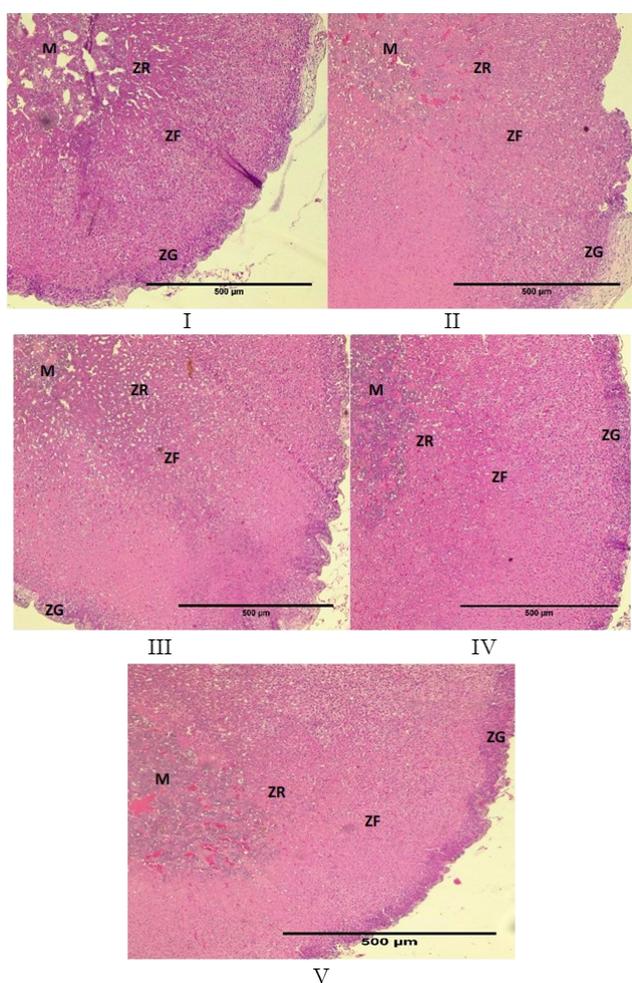


Figure 6. Histological observation of adrenal gland (100 magnification, HE stain). (I: control; II: stress control; III: amitriptyline; IV: cultivated *C. vulgaris*; V: commercial *C. vulgaris*. M: medulla; ZR: zona reticularis; ZF: zona fasciculata; ZG: zona glomerulosa).

Adrenal Glands

Stress experienced by animals can increase its adrenal gland weight due to the role of this gland in producing the hormone ACTH. This increase in weight is caused by hyperplasia

Prolonged stress will cause over-activation of the system and can cause an increase in mass in the adrenals (Ulrich-Lai *et al.*, 2006).

Brain cells are prone to damage from lipid peroxidation because most of the neuronal cell membranes are composed of unsaturated fatty acids which are the main substrates of ROS. Endogenous antioxidant defence can naturally reduce the production of ROS but chronic stress from the CUMS method is able to damage the endogenous antioxidant defences. This caused an increase in lipid peroxidation and has been studied to be one of the contributing factors to depression (Che *et al.*, 2015). *C. vulgaris* has been studied to have high antioxidant content (Saranya *et al.*, 2014) and is thought to be the cause of the antidepressant effect observed in this study. Furthermore, *C. vulgaris* is rich in tryptophan and tyrosine (Santhanam, 2015), the precursor to the neurotransmitter serotonin and noradrenaline (Hemat, 2004). These neurotransmitter's low concentration in the brain has been linked to depression as well (Lopez-Munoz & Alamo, 2011; Moret & Briley, 2011)

CONCLUSIONS

The conclusion of this study is the administration of *C. vulgaris* 360 mg/kgBW *p.o.* in female Wistar rats for 14 days after the CUMS period for 42 days was able to reduce depression-like behaviours, but was not able to revert adrenal glands size back to normal. *C. vulgaris* source did not make a notable difference to the antidepressant effects in this study. The potential of *C. vulgaris* as an antidepressant was thought to originate from its high antioxidant content which able to reduce oxidative stress in brain cells.

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