Defensive proclivity of bacoside A and bromelain against oxidative stress and AChE gene expression induced by dichlorvos in the brain of Mus musculus

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The objective of current study was to evaluate the neuroprotective effects of bacoside A and bromelain against dichlorvos induced toxicity. The healthy, 6–8 weeks old male Swiss mice were administered in separate groups subacute doses of dichlorvos (40 mg/kg bw), bacoside A (5 mg/kg bw) and bromelain (70 mg/kg bw). In order to determination of oxidative stress in different groups, thiobarbituric acid reactive substances (TBARS) and protein carbonyl content (PCC) were studied in the present investigation. Moreover, for toxic manifestation at molecular level the site-specific gene amplification of acetylcholinesterase (AChE) gene was studied in the brain. Nonetheless, the protective effects of bacoside A and bromelain were also evaluated on the TBARS, PCC and AChE gene. The exposure of dichlorvos leads to significant increase in TBARS level (p < 0.01, p < 0.001) and PCC. Besides, the decline in DNA yield, expression of amplified products of AChE gene was observed in the brain of dichlorvos treated group. The bacoside A and bromelain treatments significantly decreased the level of TBARS (p < 0.05, p < 0.01) and PCC whereas, increase in the DNA yield and expression of amplified AChE gene products were observed in the brain compared to only dichlorvos treated mice. The overall picture which emerged after critical evaluation of results indicated that the dichlorvos induced oxidative stress and alteration in AChE gene expression showed significant improvement owing to the treatments of bacoside A and bromelain. Thus, bacoside A and bromelain are very effective in alleviating neurotoxicity induced by dichlorvos.

An oxidation plays a key role in the energy metabolism for sustaining the life processes of the organisms. However, in the process the molecular oxygen is produced which in turn has the aptitude to un-pair and leave highly reactive free radicals destabilising the cellular equilibrium. Thus, the molecular reactive oxygen species (ROS) are being continuously produced during metabolism. The ROS production increases manifold under the influence of certain toxicants. Dichlorvos, a well known pesticide has been reported to cause a strong DNA alkylation effects on the brain causing hypoplasia1. Furthermore, the toxicity of dichlorvos has also been reported to reveal alterations in DNA replication, mutations2 and cellular hyperproliferation3–5. Exposure of dichlorvos has also been reported to impede the mitochondrial bioenergetics by disrupting cytochrome oxidase and electron transport system and causing oligonucleosomal DNA fragmentation of neurons6. Nevertheless, in the same study, mitochondrial DNA damage and oligonucleosomal DNA fragmentation, apoptotic neuronal degeneration and formation of 8 hydroxydeoxyguanosine are reported6,7. Moreover, in other studies human cells exposed to low concentrations of dichlorvos for short duration has been reported to damage the DNA8–10. The presence of a vinyl chloride group in the dichlorvos molecule is held responsible for the ability to induce point mutations in Salmonella and Streptomyces11. The intraperitoneal injection of dichlorvos has also been reported to cause methylation of DNA in the tissues of mice12. Moreover, dichlorvos has also been reported to increase the sister chromatid exchanges and chromosomal aberrations in chinese hamster ovary (CHO) cells, lung fibroblasts and mouse lymphoma cells13–15. The semiconservative DNA synthesis leading to low reparative synthesis is also

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affected by dichlorvos in human lymphocytes. Thus, it is quite evident from the studies that dichlorvos damages DNA at cellular level.

Bacosides are isolated from brahmi (Bacopa monniera) and have antioxidant action on the mammalian brain. Bacosides are considered to be responsible for markedly improving brain functioning. Bacoside A has been reported to repairs the damage caused in neurons by enhancing neuronal activity, protein kinase, synaptic activity and nerve impulse transmission.

Bromelain is a mixture of enzymes which is found in the pineapple juice (Ananas comosus). The target molecules for bromelain’s proteolytic enzymes are those peptidases which cause degradation to neurokinins. Bromelain has antioxidant property due to its potential to scavenge the free radicals and elevate the level of enzymatic and nonenzymatic antioxidants. It has also been reported reverse the neurotransmitters level and the enzymes such as AChE, butyrylcholinesterase (BChE), γ-amino butyric acid (GABA) and serotonin in the kidneys after the dichlorvos exposure. Now it is quite established from the earlier studies that dichlorvos impairs neuronal functions and damage DNA due to oxidative stress. Bacoside A and bromelain are well known antioxidants and preferentially act on brain functioning. Therefore, in the present investigation neuroprotective potential of bacoside A and bromelain is evaluated against dichlorvos induced toxicity with reference to oxidative stress and AChE gene expression, hitherto unreported in the literature.

Results

Study of oxidative responses. TBARS (Fig. 1). Level of TBARS increased in dichlorvos treated (group II) mice compared to control mice (group I). In bacoside A treated (group III) mice, level of TBARS decreased compared to control, dichlorvos treated and bromelain treated (group IV) mice. Meanwhile, level of TBARS increased in bacoside A treated mice compared to those mice which were treated with a combination of bacoside A and bromelain (p < 0.01) (group V) and bacoside A and bromelain along with dichlorvos (group VI). TBARS level was found to be decreased in bromelain treated mice compared to dichlorvos treated group. Increased level of TBARS found in group that was given bromelain solely, compared to bacoside A treated group, the group that was exposed to a combination of both bacoside A and bromelain (p < 0.01) and in which along with bacoside A and bromelain dichlorvos was also administered (p < 0.05). In the mice, which were exposed to a combination of both antioxidants, level of TBARS significantly decreased compared to control (p < 0.01), dichlorvos treated (p < 0.001), bacoside A treated (p < 0.01), bromelain treated (p < 0.01) and the mice which were exposed to antioxidants along with dichlorvos. Level of TBARS significantly decreased in mice that were treated with bacoside A and bromelain along with dichlorvos compared to control (p < 0.05), dichlorvos treated (p < 0.01), bacoside A treated and bromelain treated mice (p < 0.05) and it increased compared to group that was exposed to a combination of bacoside A and bromelain.
PCC (Fig. 2). In dichlorvos treated mice, PCC activity was elevated compared to control mice whereas activity of PCC was decreased in bacoside A treated mice compared to control. In bromelain treated mice, PCC activity was declined compared to all other groups. PCC level was found maximum in group of mice that was exposed to a combination of bacoside A and bromelain compared to all other groups. In the mice which were administered with bacoside A and bromelain along with dichlorvos, level of PCC was declined compared to control.

Activity of PCC was decreased in bacoside A treated mice compared to dichlorvos exposed mice. When compared with dichlorvos treatment group, level of PCC was declined in the mice which were administered bacoside A and bromelain along with dichlorvos.

In comparison of bacoside A treated mice, PCC activity was declined in dichlorvos administered mice. PCC activity was decreased in the mice which were administered individually dichlorvos and a combination of bacoside A and bromelain along with dichlorvos compared to bacoside A exposed mice.

DNA isolation and its quantitation (Fig. 3A, B). Genomic DNA was isolated from the brains of different treated mice. Isolated DNA was run on the agarose gel (1%) against Lambda DNA Eco R1 Hind III double digest ladder. Neat bands of genomic DNA near about 19,200 bp for different groups were observed on gel (Fig. 3A).

Figure 3B represents the change in the yield of DNA from different treated groups. In dichlorvos treated mice, DNA yield was decreased as compared to all among groups. Yield of DNA increased in bacoside A treated mice compared to control mice, dichlorvos treated, bromelain treated and mice that were treated with a combination of bacoside A and bromelain, whereas it was recorded to be lesser compared to mice which were administered with bacoside A and bromelain along with dichlorvos. Yield of DNA was increased in groups that were treated with bromelain and a combination of bacoside A and bromelain as compared to dichlorvos treated mice whereas it declined as compared to the group that was treated with bacoside A and bromelain along with dichlorvos. Mice that were treated with both antioxidants along with dichlorvos, DNA yield increased among all groups.

Site specific amplification of AChE gene (Fig. 4A, B). Amplification with primer 1. Figure 4A shows the expression of products of site-specific amplification of AChE gene along with the reference gene (B2M). Size of B2M is 875 bp. Compared with B2M, the position of bands from lane 1 to 6 clearly depicts the size of product i.e. 1000 bp.

Intensity of amplified product of AChE in dichlorvos treated mice decreased compared to control mice, bacoside A and bromelain treated mice, whereas increased compared to mice which received concomitant exposure of bacoside A, bromelain and dichlorvos. In bacoside treated mice, intensity of amplified product of AChE was found maximum among all groups. In groups that were exposed to purely bromelain and a combination of bacoside A and bromelain, an elevation in the intensity of amplified product of AChE is recorded as compared to control and dichlorvos treated mice. However, it diminished as compared to bacoside A treated group. Minimum
intensity of amplified product of AChE was found in mice that were given concomitant exposure of bacoside A and bromelain along with dichlorvos as among all groups (Fig. 4B).

Amplification with primer 2 (Fig. 5A, B). Figure 5A shows the expression of products of site specific amplification of AChE gene along with the reference gene (B2M). Size of B2M is 875 bp. Compared with B2M, the position of bands from lane 1 to 6 clearly depicts the size of product i.e. 879 bp.

Exposure of dichlorvos reduced the intensity of amplified product of AChE which is recorded to be minimum among all groups. Intensity of amplified product of AChE was found increased in mice which were administered bacoside A, bromelain and a combination of both bacoside A and bromelain as compared to mice of control group, dichlorvos treated group and group that was administered both antioxidants along with dichlorvos. In mice, receiving concomitant exposure of bacoside A and bromelain along with dichlorvos, intensity of amplified

Figure 3. A,B Isolated DNA and their yield from the brains of differently treated mice. M—marker, Lane 1—group I, Lane 2—group II, Lane 3—group III, Lane 4—group IV, Lane 5—group V, Lane 6—group VI.
product of AChE increased as compared to control and dichlorvos treated mice whereas it alleviated as compared to groups exposed with bacoside A, bromelain and a combination of bacoside A and bromelain (Fig. 5B).

**Amplification with primer 3 (Fig. 6A,B).** Figure 6A shows the expression of products of site specific amplification of AChE gene along with the reference gene (B2M). Size of B2M is 875 bp. Compared with B2M, the position of bands from lane 1 to 6 clearly depicts the size of product i.e. 606 bp.

Intensity of amplified product of AChE declined in dichlorvos treated mice compared to all other groups. In bacoside A treated mice, intensity of amplified product of AChE increased as compared to mice that were subjected to dichlorvos and simultaneous exposure of bacoside A, bromelain along with dichlorvos whereas declined as compared to control and bromelain treated mice. Maximum intensity of amplified product of AChE was revealed in bromelain exposed mice among all groups. In groups that were administered a combination of bacoside A and bromelain, and concomitantly with antioxidants and dichlorvos, intensity of amplified product of AChE was decreased as compared to control, dichlorvos treated, bacoside A treated and bromelain treated mice (Fig. 6B).
Discussion

Any injury or stress by extraneous factors where target organ is brain may lead to its malfunctioning. The continuous exposure to such deleterious factors in turn may be responsible for neuropsychiatric and neurodegenerative disorders such as schizophrenia, depression, Alzheimer’s disease, cerebrovascular impairment and seizure disorder. Dichlorvos is one such pesticide which crosses the blood brain barrier and consequently causing brain dysfunctioning. Dichlorvos is used to increase the productivity of the crops in agricultural fields by killing the target pests. Besides killing pests, it disseminates in the environment and accumulates at different trophic levels in the food chain. Owing to persistent nature of dichlorvos, cellular defense system is believed to be under oxidative stress besides influencing cholinergic pathway.

The level of AChE was reported to have been increased due to bacoside A and bromelain exposure in mice. These findings are in concurrence with the findings of current investigation. It seems that there is an improvement in AChE gene expression due to bacoside A and bromelain. Moreover, the bacoside A and bromelain ameliorated the toxicity induced by dichlorvos due to their antioxidant nature and having the ability to increase permeability to other antioxidants and drugs.

The substances which confer protection to cells from the damage caused by deleterious action of free radicals are known as antioxidants. The different types of antioxidants are key agents in the treatment of various

Figure 5. A,B Products of expression of site specific amplification of AChE gene with primer 2 and change in their intensity pattern in different groups. M—marker, Lane 1—group I, Lane 2—group II, Lane 3—group III, Lane 4—group IV, Lane 5—group V, Lane 6—group VI, Lane 7—B2M.
neurodegenerative disorders. In the present study, oxidative stress has been found to be increased with the simultaneous alleviation of endogenous antioxidants in the brain and cholinesterases level after dichlorvos intoxication. Earlier studies have revealed that the administration of dichlorvos resulted in the enhanced generation of free radicals. Thus, our results are in the agreement with previous report. Consequently, the overall picture which emerges after toxic treatment, leads to disturbance in the equilibrium between formation and elimination of ROS. This imbalance has been studied in current research by evaluating the concentrations of TBARS and PCC as potential indicators of oxidative stress. Brain is considered to be a major target organ of dichlorvos toxicity due to oxidative damage which has been assumed to be due to the high oxygen utilization, high level of iron and poly unsaturated fatty acids (PUFAs). Furthermore, it has been reported bio membranes rich in PUFAs lead to lipid peroxidation (LPO). In the present investigation, TBARS and PCC levels were found to be increased after dichlorvos treatment reflecting oxidative stress thus the results are in agreement with earlier findings. Other studies have also reported induction in LPO resulting in oxidative stress after dichlorvos treatment. Production of peroxides and other free radicals in the normal redox state of cell can lead to toxic effects which may damage all the components of the cells. In current study, levels of TBARS and PCC were decreased after bacoside A exposure.

Bacoside A is a triterpenoid saponin which serves as a potent nerve tonic. Bacoside A has potential of memory enhancement and found to be significant in the treatment of cardiac, respiratory and neurological disorders.
disorders such as stress, insomnia, depression, insanity, psychosis and epilepsy. Previous studies have also revealed that bacoside A markedly reduces the free radicals’ generation which in turn diminishes the oxidative stress at the cytoplasmic and mitochondrial levels. Furthermore, in earlier studies the Bacopa monniera have been shown to avert LPO in vitro and in vivo by quenching super oxide, hydroxyl and nitric oxide radicals. Bacoside A is reported to cause decline in the lipid peroxidation, thus establishing its antilipid peroxidative property. Thus, the results of earlier and present studies clearly depict that bacoside A has the potential to reduce the oxidative stress.

In the present study, bromelain exposure diminished the level of TBARS and PCC contents. In earlier reports, bromelain has been shown to reduce the generation of NO in the primary microglia of the rats. In our study, TBARS and PCC level were found to be decreased by concomitant exposure of dichlorvos, bacoside A and bromelain. Bacoside A and bromelain both have antioxidant property therefore both these agents are playing role in the reduction of ROS and also reducing the protein oxidation. The results of the present investigation are in agreement with earlier study that bromelain diminishes the levels of TBARS, NO and PCC.

In the present study, the yield of DNA and intensity of amplified product of AChE gene with primer 1, 2 and 3 was decreased by dichlorvos exposure. It is proposed that this may be due to ROS which break the DNA strands and lead to the formation of DNA adducts. The deoxy ribose sugar and base moieties in the DNA are likely to be degraded by ROS and causes oxidation of bases and cross linking to protein. DNA-MDA adducts is the most attribute feature of nucleic acid oxidation. 4-hydroxyl, 2-deoxyguanosine is the oxidative marker of DNA oxidation. The previous study suggested that dichlorvos hampers both extraction and amplification of mitochondrial and nuclear DNA.

In present investigation, Bacoside A and bromelain each separately, in combination and in an another group along with dichlorvos increased the yield of DNA and intensity of amplified product of AChE gene with primers 1, 2 and 3. It may be due to scavenging of ROS by bacoside A and bromelain at cellular level which consequently prevents the formation of the DNA adducts. Earlier, it was reported that extract of Bacopa monniera may alter the expression of AChE gene in the hippocampus of rat brain. On the basis these findings, it may be concluded that bacoside A and bromelain confer neuroprotection against dichlorvos exposure via reducing oxidative stress and improving AChE gene expression in mice brain.

Methods

Animals monitoring. Healthy mice (6–8 weeks old) were used in the investigation. Rearing of mice was done in accordance with the Committee for the Purpose of Control and Supervision on Experimentation on Animals (CPCSEA). The study on animals was carried out in compliance with the ARRIVE guidelines.

Experimental design. For ascertaining the action of bacoside A and bromelain on brain acetylcholinesterase gene expression, mice were allocated into six groups. Each group consisted a minimum number of six animals. All doses (dichlorvos, bacoside and bromelain) were prepared separately in water. Mice were exposed to bromelain by oral gavage whereas bacoside and dichlorvos were administered intraperitoneally. First group of mice was treated as control and received normal saline as a vehicle. Mice of second group were injected dichlorvos (40 mg/kg bw) to create neurotoxicity. Mice of groups III and IV were administered with bacoside A (5 mg/kg bw) and bromelain (70 mg/kg bw). Concomitant exposure of bacoside A and bromelain was given to mice of group V. In group VI, mice were exposed with dichlorvos and both the antioxidants under study.

After completion of exposure duration (i.e., 21 days in current study) mice of different groups were sacrificed. Brains were immediately isolated to make different homogenates as per their requirement in various methods.

Study of oxidative responses. TBARS and PCC serves as the marker of oxidative stress. TBARS level was measured by Ohkawa et al. (1979) method. MDA is formed as a result of peroxidation of lipids which reacts with 2-thiobarbituric acid (TBA) and forms adduct which gives pink coloured complex and can be observed spectrophotometrically at 532 nm. 100 µl of tissue sample (2%), 200 µl of 0.8% SDS, 1.5 ml of acetate, 1.5 ml of TBA solution were added to make reaction mixture. Samples were incubated at 90° C for 60 min and cooled at room temperature. 5 ml of butanol and pyridine were added in reaction mixture in the ratio of 15:1. Then samples were centrifuged at 3000 rpm for 10 min at room temperature. Upper layer was taken and read at 532 nm.

PCC level was measured by Levine et al. (1990) method. To the 100 µl of sample, 1 ml of buffer, 100 µl of TCA was added. The samples were incubated for 15 min and then centrifuged for 8000 rpm for 10 min at room temperature. After centrifugation, supernatant was carefully decanted and pellet was resuspended in 500 µl of DNPH and incubated for 1 h at room temperature. Again 500 µl of 20% TCA was added in the samples and centrifuged at 5000 rpm for 10 min at room temperature. Pellet was washed with 1.5 ml of ethanol: ethyl acetate for 3 times. This step was used to remove excess of DNPH and centrifuged at 8000 rpm for 10 min. The final pellet was dissolved in 2 ml of 6 M guanidine HCl and read at 370 nm.

Expression of AChE gene. DNA isolation. DNA was isolated by using DNA isolation kit (tissue, Nucleospin 740,952.250). Genomic DNA was amplified with the specific site of primer. Brain tissue 600 µl (10%) was mixed with lysis buffer and incubated at 55 °C for 60 min. Equal volume of buffer saturated phenol was added and centrifuged for 5 min at 12,000 rpm at 4 °C. Upper layer was added with equal volume of phenol and centrifuged at 12,000 rpm for 10 min at 4 °C (this step was repeated twice). Equal volume of chloroform was added in upper aqueous layer and centrifuged at 12,000 rpm for 10 min. Aqueous layer was taken and isopropanol was added in the ratio of 1:1 and then centrifuged at 12,000 rpm for 15 min at 4 °C. The pellet was dried and dissolved in 50 µl of TE buffer. The yield of isolated DNA was checked on 0.8% agarose gel electrophoresis.
BT/383/13-14). Approval taken from Institutional Animal Ethical Committee of Banasthali University, Banasthali (Ref no. BU/...Compliance with ethical standards.

Maintenance and treatment of animals were done in accordance with Committee for the Purpose of Control and Supervision of Experimentation on Animals (CPCSEA). Ethical approval taken from Institutional Animal Ethical Committee of Banasthali University, Banasthali (Ref no. BU/BT/383/13-14).

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Table 1. PCR program for amplification of AChE gene with specific primer.

| Program          | Steps    | Temperature (°C) | Time (min) | Cycles |
|------------------|----------|------------------|------------|--------|
| Initial denaturation | –       | 95               | 5          | 1      |
| PCR cycle        | Denaturation | 94               | 1          |        |
|                  | Annealing       | 50–60            | 1          | 32     |
|                  | Extension       | 72               | 2          |        |
|                  | Extension       | –                | 72         | 10     |
|                  | Cooling         | –                | 4          | Forever |

Table 2. List of primers for sequence specific amplification of AChE gene.

| Primer | Forward primer | Reverse primer | Annealing temp (°C) |
|--------|----------------|----------------|---------------------|
| 1      | TGAGTGTAGTGGTGTGCTGGG | AGATTGAGAGATGCCAGGACACGACACGAG | 52 |
| 2      | GGTGCGGTGTGAGTGAAGG | TGGGCTCTGGTGCGCATAAA | 54 |
| 3      | CAGATACCCCCACACCAAGG | GCAAGATTGAGTAAAGGGGGAGG | 57 |
| B2M    | CACCGAGGAGAAATAGGAAATG | ACACGAAACAGTATGCAC | 52 |

**Prime designing.** The AChE gene sequence of Mus Musculus was collected from ENSMUSG00000023328; MGI: 87,876. Sequence was used to design sequence specific primers with amplicon size of near about 600–1000 bp. Primers were designed using Gene runner and Berkeley calculator.

**Amplification of isolated DNA (Table 1).** The isolated DNA was used as a template for PCR reaction. Different primers needed for amplification of DNA are mentioned in Table 2. The amplification of DNA was done on PCR machine (Primer 96 advanced thermal cycler). Beta-2 microglobulin (B2M) was used as a house-keeping gene. The band intensity of amplified product was measured by NIH Image J Software.
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Author contributions

R.B.: Preparation and evaluation of manuscript, data interpretation and analysis, calculations of all data preparation of graphs, observation and validation of entire wet lab working. B.C.: Wet lab work was done, writing of manuscript, analysis of data. D.K.B.: Data interpretation and analysis, editing and evaluation of manuscript. All the authors read and approved the final manuscript.

Competing interests

The authors declare no competing interests.

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