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Evaluation of the Genotoxic and Antigenotoxic Effects of Chios Mastic Water by the *In Vitro* Micronucleus Test on Human Lymphocytes and the *In Vivo* Wing Somatic Test on *Drosophila*

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Abstract

Chios mastic gum, a plant-derived product obtained by the Mediterranean bush *Pistacia lentiscus* (L.) var. *chia* (Duham), has generated considerable interest because of its antimicrobial, anticancer, antioxidant and other beneficial properties. Its aqueous extract, called Chios mastic water (CMW), contains the authentic mastic scent and all the water soluble components of mastic. In the present study, the potential genotoxic activity of CMW, as well as its antigenotoxic properties against the mutagenic agent mitomycin-C (MMC), was evaluated by employing the *in vitro* Cytokinesis Block MicroNucleus (CBMN) assay and the *in vivo* Somatic Mutation And Recombination Test (SMART). In the former assay, lymphocytes were treated with 1, 2 and 5% (v/v) of CMW with or without MMC at concentrations 0.05 and 0.50 µg/ml. No significant micronucleus induction was observed by CMW, while co-treatment with MMC led to a decrease of the MMC-induced micronuclei, which ranged between 22.8 and 44.7%. For SMART, larvae were treated with 50 and 100% (v/v) CMW with or without MMC at concentrations 1.00, 2.50 and 5.00 µg/ml. It was shown that CMW alone did not modify the spontaneous frequencies of spots indicating lack of genotoxic activity. The simultaneous administration of MMC with 100% CMW led to considerable alterations of the frequencies of MMC-induced wing spots with the total mutant clones showing reduction between 53.5 and 74.4%. Our data clearly show a protective role of CMW against the MMC-induced genotoxicity and further research on the beneficial properties of this product is suggested.

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Introduction

Pistacia lentiscus (L.) var. *chia* (Duham) is an evergreen bush, uniquely cultivated in the Greek island Chios [1–3]. It produces a white semitransparent resin, which is generally known as Chios mastic gum. This product as well as its essential oil, Chios mastic oil, has been extensively used as food/beverages flavoring additives in confectionery, in perfume industry and as an ingredient of cosmetics and health products [3–5]. Their beneficial biological properties have been well documented by a number of studies showing their antibacterial, antimicrobial, anti-inflammatory and antioxidant activity [5–13] and they have been proposed for many clinical applications [14–21]. Recently, their anticancer properties against a number of human malignancies have been reported [22–28]. Despite the great number of reports analyzing the biological activities of mastic gum and mastic oil, such studies are scarce for

the very closely related commercially available product, known as Chios mastic water (CMW).

CMW is a flavoring obtained in large quantities together with mastic oil during the steam distillation of mastic resin. It is a 100% natural aqueous extract that contains all the water soluble components of mastic gum as well as a small amount (0.5–1% v/v) of mastic oil [data from Chios Mastiha Growers' Association, CMGA]. Its major identified compounds are verbenone, α -terpineol, trans-p-menth-2-ene-1,8-diol, cis-p-menth-2-ene-1,8-diol, linalool, β -phellandrenol and trans-pinocarveol [29]. With the exception of a recent study on its chemical composition and its antimicrobial activities against *Escherichia coli*, *Staphylococcus aureus* and *Candida* spp. [29], data on the biological properties of this low-cost product of mastic resin do not exist.

In an effort to evaluate the safety of use of CMW, the possible genotoxic and recombinogenic effects of this mastic product were

studied here. To further explore its biological properties, the potential protective effects of CMW against the mutagenic and recombinogenic effects of mitomycin-C (MMC) were also investigated. Both genotoxic and antigenotoxic potential activities of CMW were assessed employing the cytokinesis block micronucleus (CBMN) assay and the somatic mutation and recombination test (SMART). The former is an *in vitro* assay applied in cultured human lymphocytes for the detection of micronuclei (MN) in the cytoplasm of interphase cells. MN may originate from acentric chromosome fragments or whole chromosomes that are unable to migrate to the poles during the anaphase stage of cell division. Thus, this assay detects the potential clastogenic and aneugenic activity of chemicals in cells that have undergone cell division after exposure to the test chemical [30,31]. The simplicity, rapidity and sensitivity of the CBMN assay make it a valuable tool for genotoxicity screening. Moreover, the SMART test in *Drosophila melanogaster* (Meigen) used here, is a sensitive, low-cost, rapid eukaryotic *in vivo* assay able to detect the potential mutagenic and recombinogenic effects as well as the antigenotoxic ability of chemicals. Thus, a wide spectrum of genetic end points such as point mutations, deletions, certain types of chromosome aberrations, as well as mitotic recombination and gene conversion can be detected [32,33]. The extensive knowledge on the genetics of *D. melanogaster* and the high homology between fly and human genes [34–38] have made this organism unique in mutation research and genetic toxicology.

Since MN formation and recombinogenic events are found to be associated with carcinogenesis [39,40], our results are expected to contribute to the establishment of the safety status of this commercially available mastic product. Moreover, its potential antigenotoxic activity against mutagens could contribute to the development of chemopreventive agents capable of modulating the cellular responses to mutagens (or of phytopharmaceutical molecules of interest).

Materials and Methods

Chemicals

The CMW was supplied by CMGA (Chios, Greece). MMC and cytochalasin-B (Cyt-B) were purchased from Sigma (St. Louis, MO, USA). Ham's F-10 medium, foetal bovine serum and phytohaemagglutinin were commercially supplied (Gibco, UK). Faure's solution was prepared by mixing 100 g distilled water (H₂O), 100 g chloral hydrate (C₂HCl₃O.H₂O), 40 g glycerine (C₃H₈O₃) and 60 g arabic gum. All other chemicals and solvents were of the highest grade commercially available. Stocks of the compounds and solutions were stored at 4°C until use.

Ethics Statement

The study was approved by the Ethical Committee of the University of Patras. After informed consent two healthy, non-smoking male individuals (less than 30 years), were used as blood donors to establish whole blood lymphocyte cultures. According to the donors' declaration, they were not exposed to radiation, drug treatment or any viral infection in the recent past.

CBMN Assay in Human Lymphocytes *in vitro*

Blood samples were kept under sterile conditions in heparinized tubes. Whole blood (0.5 ml) was added to 6.5 ml Ham's F-10 medium, 1.5 ml foetal bovine serum and 0.3 ml phytohaemagglutinin to stimulate cell division.

CMW was added to final concentrations of 1, 2 and 5% (v/v) in culture volume either alone or in combination with 0.05 and 0.50 µg/ml of MMC. The MMC concentrations used in the

present study have been previously used as positive control in the particular assay and cell type [41]. The appropriate volumes were added 24 h after culture initiation. Cyt-B at final concentration of 6 µg/ml was added to the culture medium 44 h after its initiation and 20 h after the addition of the CMW, MMC or their mixtures. This concentration of Cyt-B was selected in order to obtain a higher percentage of binucleated (BN) cells and a lower baseline MN frequency [42]. Cultures were incubated at 37°C in a humidified atmosphere of 5% CO₂ for 72 h. 72 h after the initiation of culture, cells were harvested and collected by centrifugation. A mild hypotonic treatment with 3:1 solution of Ham's medium and milli-q H₂O was left for 3 min at room temperature which was followed by 10 min fixation (for at least 3 times) with a fresh 5:1 solution of methanol/acetic acid. Cells were stained with 7% Giemsa [43–45].

In total, 2000 BN cells with preserved cytoplasm were scored per experimental point. Standard criteria were used for scoring MN [46,47] and the scoring of micronuclei was performed manually and by (at least) two, independently working, experienced researchers. In order to determine possible cytotoxic effects, the cytokinesis block proliferation index (CBPI) was calculated by counting at least 1000 cells for each experimental point (500 cells per culture of each donor). CBPI is given by the equation: $CBPI = M_1 + 2M_2 + 3(M_3 + M_4)/N$ where M_1 , M_2 , M_3 and M_4 correspond to the numbers of cells with one, two, three and four nuclei and N is the total number of cells [48].

Somatic Mutation and Recombination Test (SMART)

Two *D. melanogaster* strains carrying visible wing genetic markers on the left arm of the third chromosome were used: (i) flare (*flr*³, 3–38.8) with genetic constitution *ywco/y wco; flr3 se/TM2 Ubx¹³⁰ se e* and (ii) multiple wing hairs (*mwh*, 3–0.3) with genetic constitution *f5(1)K10 w/Y; mwh se e/mwh se e* [49,50]. More detailed information on the genetic symbols and descriptions is provided by Lindsley and Zimm [49]. Insects were maintained at 24±1°C, at a photoperiod 16:8 (light:dark) on a yeast–glucose medium. The experiments were carried out following the principles and the basic procedures presented by Graf et al. [32,33]. Thus, eggs obtained by parental crosses between *flr*³ virgin females and *mwh* males were collected during a 6-hour period in culture bottles with an agar-agar base (4% w/v) topped with a thick layer of live yeast supplemented with sucrose. Three days after egg laying, larvae in the third stage of embryonic development were washed out of the bottles with Ringer's solution and collected in a stainless steel strainer. Series of 30 larvae were transferred for chronic feeding to treatment vials containing 0.85 g of *Drosophila* Instant Medium (Carolina Biological Supply, Burlington, NC, USA) rehydrated with 4 ml of 50 and 100% (v/v) CMW alone or in combination with MMC at final concentrations of 1.00, 2.50 and 5.00 µg/ml. The above concentrations of MMC were also used as positive control. Larvae were fed on these culture media for the rest of their larval life (approximately 48 h). The hatched adult flies were collected from the treatment vials and stored in 70% v/v ethanol/glycerol (1:1, v/v). The wings of the trans-heterozygous (*mwh flr+/mwh+flr³*) female flies [32,50,51], distinguished by their wild-type body color, were removed under a stereomicroscope with a pair of entomological tweezers, mounted in Faure's solution and scored at 400× magnification for the presence of mosaic spots. The rest individuals were excluded from analysis, because in the *mwh/TM2* females and the males recombinational events are suppressed [32,50,51]. The spots observed on the wings of the trans-heterozygous females were grouped into four categories based on the size, number and type of cells showing malformed wing hairs as: (i) small single spots (with one or two affected cells, either

mwh or *flr³*), (ii) large single spots (with three or more affected cells, either *mwh* or *flr³*), (iii) twin spots (consisting of both *mwh* and *flr³* subclones), and (iv) total spots [32]. Single spots (*mwh* or *flr³*) are produced by various genetic events including somatic point mutations, deletions and other types of structural rearrangements as well as by mitotic recombination between the two marker genes, while twin spots (*mwh* and *flr³*) are produced exclusively by mitotic recombination occurring between the proximal marker *flr³* and the chromosome 3 centromere [32]. For comparative analysis, parallel experiments using distilled water were carried out as the negative controls. Ten replicates per treatment were performed. Since no considerable difference in survival rates of hatched flies from independent experiments was observed, approx. 50 wing samples per treatment were randomly selected for genotoxic analysis. All experiments were performed at $24 \pm 1^\circ\text{C}$ and 60% RH. A total of about 600 wings were scored in this study.

Statistical Analysis

All results of the CBMN assay are expressed as the mean frequency \pm standard error (MF \pm se). The G-test for independence on 2×2 tables was used to perform the statistical analysis of the MN data. The chi-square test (χ^2 test) was used for the analysis of CBPI among each treatment. Differences at $p < 0.05$ were considered significant. The statistical software used for data analysis was the Origin 7.0 (OriginLab Corporation, Northampton, MA, USA), the Minitab statistical software (Minitab Inc., PA, USA) and the Statistical Package for Social Sciences (SPSS) for Windows, version 17.0.

Statistical analysis of the data derived by the SMART assay was done using the multiple-decision procedure [52,53] which is based on the conditional binomial test and the chi-squared test (K. Pearson's criterion) [54,55]. A significance level of 5% was used. For the statistical assessment of antigenotoxicity, the frequencies of each type of spots per fly were compared in pairs (negative control versus CMW; MMC versus MMC+CMW), using the non-parametric Mann-Whitney U-test [56]. Based on clone formation per 10^5 cells the percentages of CMW inhibition were calculated as follows: [(MMC - MMC combined with CMW)/MMC] \times 100 [57].

Results

Genotoxicity and Antigenotoxicity Tested with CBMN Assay

Chios mastic water was studied for genotoxicity at three different doses i.e. 1, 2 and 5% (v/v) of the total culture volume and the same doses were tested combined with different MMC concentrations (0.05 and 0.50 $\mu\text{g/ml}$) in order to identify the antigenotoxic effect of CMW against the genotoxic damage induced by MMC. A treatment with 1, 2 and 5% (v/v) of CMW doses did not induce MN at significant level as compared to control. Treatments with 0.05 and 0.50 $\mu\text{g/ml}$ of MMC induced significant MN frequencies (57.0 ± 6.0 and 177.5 ± 16.5) as compared to control. A significant decrease in MN frequencies was observed when 1, 2 and 5% (v/v) of CMW treatments were given along with both tested concentrations of MMC (Table 1). To summarize, the concentrations of CMW used in the present study were not genotoxic themselves, while they reduced the genotoxic effect of MMC.

Figure 1 shows the reduction of MMC-induced MN frequencies (%) in the presence of different concentrations of CMW. In the tested concentrations of MMC (0.05 and 0.50 $\mu\text{g/ml}$) the decrease of the MN frequencies ranges from 22.8 to 44.7%. The comparative distribution of MN frequency induced by CMW,

MMC and their combination is indicated in Figure 2. A similar pattern is shown in both MMC concentrations. In particular, a decrease in MN frequency induction is observed in co-treatment with CMW and MMC in comparison to MMC alone in both concentrations, with slightly greater decrease of the induction frequency in the CMW and 0.05 $\mu\text{g/ml}$ MMC mixtures.

The cytotoxic effect of CMW, MMC and their mixtures was evaluated by the determination of CBPI. Regarding the cytotoxic index, statistically significant differences on CBPI were detected between control cultures and the 2 and 5% (v/v) doses of CMW. The decrease of the CBPI index with some fluctuations remains in the case of MMC as well as in the mixtures of CMW with MMC (Table 1).

Genotoxicity and Antigenotoxicity Tested with SMART Assay

In a pilot experiment, the toxicity of CMW was evaluated. No toxicity of this product could be determined in *Drosophila* (data not shown). Therefore, CMW was applied at concentrations 50 and 100% (v/v) for the genotoxicity experiments. MMC was supplied to *D. melanogaster* larvae at doses of 1.00, 2.50 and 5.00 $\mu\text{g/ml}$, since preliminary experiments showed that lower concentrations did not exert strong genotoxic effects under our experimental conditions (data not shown). The antigenotoxic effect of CMW against the genotoxic damage induced by MMC was accomplished by co-treatment of the above doses of both compounds.

Table 2 summarizes the results together with the negative control experiment. No significant differences in any of the three spot categories were observed after chronic treatment of *Drosophila* larvae with CMW, compared to those of their respective negative controls, indicating that CMW was not genotoxic under our experimental conditions. On the other hand, treatment of the larvae with MMC at concentrations 1.00, 2.50 and 5.00 $\mu\text{g/ml}$ evoked a statistically significant rise in all spot categories. The correlation between the dose and the frequency of the induced total spots indicates the dose dependent genotoxic activity of MMC. Moreover, the positive effect of twin spots at the high concentrations used clearly indicates the recombinogenic activity of this agent.

After co-treatment of MMC with 50% (v/v) CMW, a reduction of the induced total wing spot frequency was observed, which, however, was not found to be statistically significant ($U = 238.5$, $p = 0.296$, $U = 230.0$, $p = 0.158$, $U = 210.0$, $p = 0.071$, for 1.00, 2.50 and 5.00 $\mu\text{g/ml}$ MMC, respectively) (Figure 3). On the contrary, a more pronounced decrease of MMC-induced total spots was provoked by 100% (v/v) CMW. This overall inhibition was 57.4%, ($U = 189.0$, $p = 0.032$), 74.4% ($U = 92.5$, $p = 0.000$) and 53.5% ($U = 129.0$, $p = 0.000$) in the case of 1.00, 2.50 and 5.00 $\mu\text{g/ml}$ MMC, respectively (Figure 3). It should be noted that when 100% (v/v) CMW is supplied along with 1.00 and 2.50 $\mu\text{g/ml}$ MMC, the frequencies of total spots are similar to those observed in the negative control meaning that CMW is able to inhibit completely the genotoxic activity of MMC at these concentrations (Table 2). On the other hand, when 100% (v/v) CMW is combined with 5 $\mu\text{g/ml}$ MMC, even though the reduction of total spots is over 50% and statistically significant ($U = 129.0$, $p = 0.000$), the genotoxic result remains positive. Considering spot sub-categories, 100% (v/v) CMW reduced significantly small single and total spots induced by 1.00 $\mu\text{g/ml}$ MMC ($U = 193.0$, $p = 0.037$) and all spot categories induced by 2.50 and 5.00 $\mu\text{g/ml}$ MMC ($U = 115.0$ – 191.0 , $p = 0.000$ – 0.031) indicating both antigenotoxic and antirecombinogenic activity.

Table 1. Frequencies of BNMN and MN as well as CBPI values in cultured human lymphocytes treated with CMW, MMC (0.05 and 0.50 µg/ml) and their mixture.

Treatment	BNMN MF (%) ± se	MN MF (%) ± se	CBPI MF (%) ± se
Control	4.5±0.5	5.0±1.0	1.89±0.04
1% (v/v) CMW	5.0±2.0	5.5±2.5	1.87±0.04
2% (v/v) CMW	4.5±0.5	4.5±0.5	1.74±0.04 ²
5% (v/v) CMW	9.0±1.0	9.0±1.0	1.64±0.03 ³
MMC (0.05 µg/ml)	55.5±6.5 ³	57.0±6.0 ³	1.67±0.02 ³
1% (v/v) CMW+MMC (0.05 µg/ml)	30.5±0.5 ^{3,c}	31.5±0.5 ^{3,c}	1.69±0.02 ^{3,a}
2% (v/v) CMW+MMC (0.05 µg/ml)	32.5±2.5 ^{3,c}	34.0±2.0 ^{3,c}	1.76±0.02 ^{1,a}
5% (v/v) CMW+MMC (0.05 µg/ml)	36.0±2.0 ^{3,b}	36.5±2.5 ^{3,b}	1.68±0.05 ^{3,a}
MMC (0.50 µg/ml)	166.0±15.0 ³	177.5±16.5 ³	1.50±0.04 ³
1% (v/v) CMW+MMC (0.50 µg/ml)	126.5±1.5 ^{3,c}	137.0±1.0 ^{3,c}	1.48±0.00 ³
2% (v/v) CMW+MMC (0.50 µg/ml)	128.5±2.5 ^{3,c}	134.5±1.5 ^{3,c}	1.50±0.03 ³
5% (v/v) CMW+MMC (0.50 µg/ml)	128.5±26.5 ^{3,c}	134.5±29.5 ^{3,c}	1.40±0.01 ^{3,c}

BN: binucleated cells; BNMN: micronucleated binucleated cells; MN: micronuclei; CBPI: Cytokinesis Block Proliferation Index; CMW: Chios Mastic Water; MMC: Mitomycin C; MF (%) ± se, mean frequencies (%) ± standard error; MN were scored in 2000 binucleated lymphocytes per experimental point;

^{1,2,3}significant difference in relation to control at $p<0.05$, $p<0.01$ and $p<0.001$ respectively;

^{a,b,c}significant difference in relation to MMC at $p<0.05$, $p<0.01$ and $p<0.001$ respectively [G-test for BNMN and MN; χ^2 for CBPI].

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Discussion

In recent years, mastic gum, a natural resin obtained from the plant *Pistacia lentiscus* var. *chia*, its essential oil (i.e. mastic oil), as well as some of their constituents [e.g. linalool, verbenone, α -terpineol, *trans*-pinocarveol] have received much attention as potentially useful bioactive compounds, with particular emphasis being given to their antioxidant, antimicrobial, anti-inflammatory or antitumor properties [5–29,58–65]. Based on the increasing international interest for mastic products, in the present study, the CMW was evaluated, for the first time, for its potential genotoxic effect as well as its antigenotoxic activity against the DNA damage induced by MMC.

For this purpose, the *in vitro* CBMN assay in cultured human lymphocytes and the *in vivo* SMART test in *D. melanogaster* were applied. Both assays are short-term genotoxicity tests able to evaluate several genetic endpoints during the cell cycle or special developmental stages [30–32].

In our testing systems, CMW was not found to be genotoxic, mutagenic or recombinogenic, as it did not induce increased frequencies of micronuclei or wing spots within a wide range of concentrations (Tables 1–2). To our knowledge there is no data on the genotoxic activity of CMW; nevertheless, one of its major constituents, linalool, with contribution 7.29% [29] was found not to exhibit genotoxic or recombinogenic activity [65]. Moreover, borneol which has low contribution (0.99%) in CMW [29] has been reported not to be genotoxic at low concentrations [61]. Since the organic fraction of CMW is a complex mixture of many constituents [29], the absence of genotoxicity found in the present study could be attributed either to its major constituents or to synergistic and/or antagonistic phenomena that may exist among its constituents [66,67]. Our results are also in line with previous reports showing that extracts of *P. lentiscus* did not exert any genotoxic effects [68,69].

As it can be seen in Table 1, a significant decrease of CBPI values were noticed at concentrations 2 and 5% (v/v) of CMW.

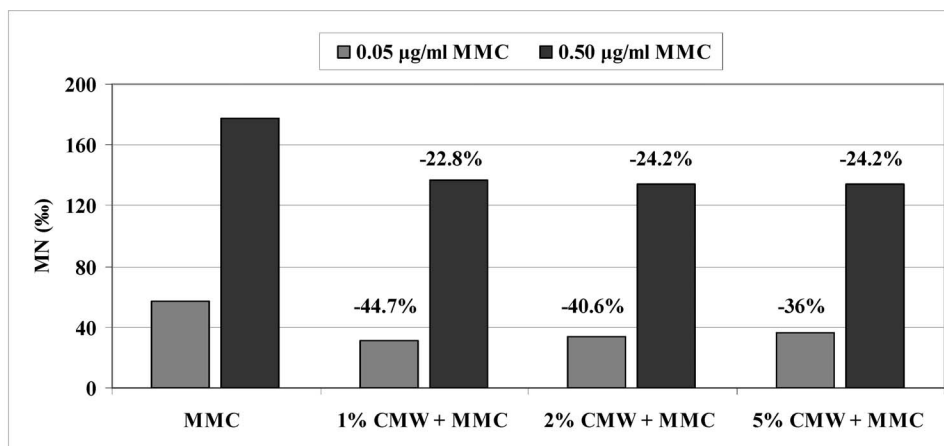


Figure 1. Reduction (%) of MN frequency induced by MMC (0.05 and 0.50 µg/ml) in presence of CMW (1, 2 and 5% v/v).

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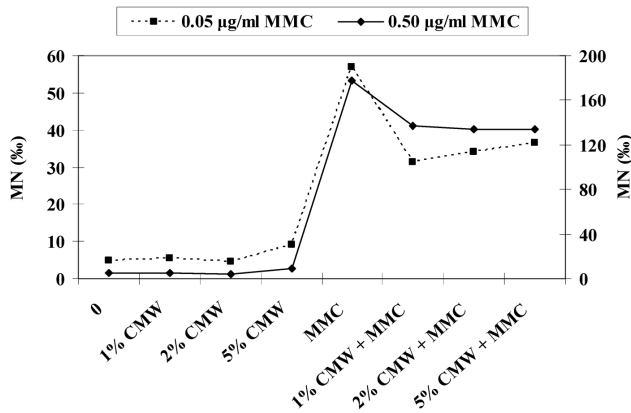


Figure 2. Comparative distribution of MN frequency (%) induced by CMW, MMC and their combination. The dotted line is read on the left hand Y-axis and the solid line on the right hand Y-axis.

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The above observation is related with the CMW constituents and is supported by literature data demonstrating that some of these constituents, namely verbenone, α -terpineol, linalol, α -phellandrenol, myrtenol/myrtenal, terpinen-4-ol and borneol, have been shown to exhibit cytotoxic activity [59–61,63,64].

Since no genotoxic activity was detected at any concentration tested, the potential antigenotoxic activity of CMW was evaluated, herein. For this purpose, MMC was used as a mutagenic inducer, similarly to a number of other antigenotoxicity studies [70–74]. MMC is an alkylating, antibiotic compound that has a range of genotoxic effects including the inhibition of DNA synthesis, mutagenesis and clastogenesis. It was found to be genotoxic in all *in vitro* and *in vivo* test systems in mammalian cells and animals and was clearly demonstrated as carcinogenic agent [75]. Consistent with previous studies [70–77], MMC was found to be

genotoxic in both our testing assays, inducing significant MN and wing spot frequencies at concentrations 0.05 and 0.50 $\mu\text{g/ml}$ in the CBMN assay and over 1.00 $\mu\text{g/ml}$ in the SMART test (Tables 1, 2). In addition, the significant induction of twin spots in SMART test indicated the recombinogenic activity of this agent (Table 2).

The co-treatment of human lymphocytes and *D. melanogaster* larvae with CMW and MMC demonstrated that CMW could afford protection against the used mutagen indicating its antigenotoxic activity under both our *in vitro* and *in vivo* testing conditions. More precisely, the frequency of micronuclei and wing spots was statistically decreased when MMC was combined with CMW in comparison to the micronuclei and spot frequencies induced by MMC alone (Tables 1–2, Figures 1, 2, 3). In the CBMN assay, the decrease of the frequency of MN induction ranged between 36.0% and 44.7% and between 22.8% and 24.2% for 0.05 and 0.50 $\mu\text{g/ml}$ of MMC, respectively. The antigenotoxic capacity of CMW was further supported by the results of the SMART test, which showed a profound reduction of total mutant spots on the wing blade after co-treatment with MMC and 100% (v/v) CMW compared to MMC alone. The decrease of the total spots was found to be 57.4, 74.4 and 53.5% for 1.00, 2.50 and 5.00 $\mu\text{g/ml}$ of MMC, respectively. The significant reduction of the frequencies of single (small and large) and twin spots indicate that 100% (v/v) CMW has a protective effect against the MMC's genotoxic and recombinogenic action (Table 2, Figure 3).

It is of note that in the CBMN assay the most profound decrease of MMC-induced MN (i.e. 44.7%, Figure 1) was observed when the lowest examined CMW concentration (i.e. 1% v/v) was co-administered to 0.05 $\mu\text{g/ml}$ of MMC. These results are in accordance with those of Kim and Neophytou [62], who observed that the lowest dose of mastic oil (0.02% v/v) had a stronger influence than the highest one (0.2% v/v) with respect to the decrease of the symptoms of clinical colitis in mice. Moreover, recent studies reported that beneficial effects of mastic gum/extracts could be achieved at low doses [22–24,26,58]. These findings lead to a possible assumption that low concentrations of

Table 2. Summary of the results obtained in the Somatic Mutation And Recombination Test (SMART) on *Drosophila melanogaster*, after larvae treatment with CMW, MMC (1.00, 2.50 and 5.00 $\mu\text{g/ml}$) and their mixture.

Treatment	Number of wings	Frequency of spots per wing and diagnosis ¹			
		Small single spots $m=2.0$	Large single spots $m=5.0$	Twin spots $m=5.0$	Total spots $m=2.0$
Control	50	0.460 (23)	0.040 (2)	0.020 (1)	0.520 (26)
50% (v/v) CMW	50	0.460 (23) –	0.000 (0) –	0.100 (5) i	0.560 (28) –
100% (v/v) CMW	50	0.320 (16) –	0.020 (1) i	0.020 (1) i	0.360 (18) –
MMC (1.00 $\mu\text{g/ml}$)	47	0.745 (35) +	0.234 (11) +	0.043 (2) i	1.021 (48) +
50% (v/v) CMW+MMC (1.00 $\mu\text{g/ml}$)	46	0.500 (23) –	0.152 (7) +	0.022 (1) i	0.740 (31) i
100% (v/v) CMW+MMC (1.00 $\mu\text{g/ml}$)	46	0.348 (16) – ^a	0.087 (4) i	0.000 (0) i	0.435 (20) – ^a
MMC (2.50 $\mu\text{g/ml}$)	50	1.540 (77) +	0.720 (36) +	0.400 (20) +	2.660 (133) +
50% (v/v) CMW+MMC (2.50 $\mu\text{g/ml}$)	48	1.021 (49) +	0.667 (32) +	0.333 (16) +	2.021 (97) +
100% (v/v) CMW+MMC (2.50 $\mu\text{g/ml}$)	47	0.426 (20) – ^b	0.255 (12)+ ^a	0.000 (0) i ^c	0.681 (32) i ^c
MMC (5.00 $\mu\text{g/ml}$)	50	2.580 (129) +	2.040 (102) +	0.540 (27) +	5.160 (258) +
50% (v/v) CMW+MMC (5.00 $\mu\text{g/ml}$)	49	2.102 (103) +	1.612 (79) +	0.429 (21) +	4.143 (203) +
100% (v/v) CMW+MMC (5.00 $\mu\text{g/ml}$)	50	1.200 (60)+ ^b	0.880 (44)+ ^b	0.320 (16)+ ^b	2.400 (120)+ ^c

Symbols next to values signify the following: +, positive mutagenic effect; –, no mutagenic effect; w, weakly positive effect; i, inconclusive effect; m is the multiplication factor¹ ($p=0.05$); ^a, ^b, ^c is significant difference in relation to MMC at $p<0.05$, $p<0.01$ and $p<0.001$, respectively (U-test).

¹The number of mutant spots is given in parenthesis. Statistical diagnosis according to Frei and Würgler [53].

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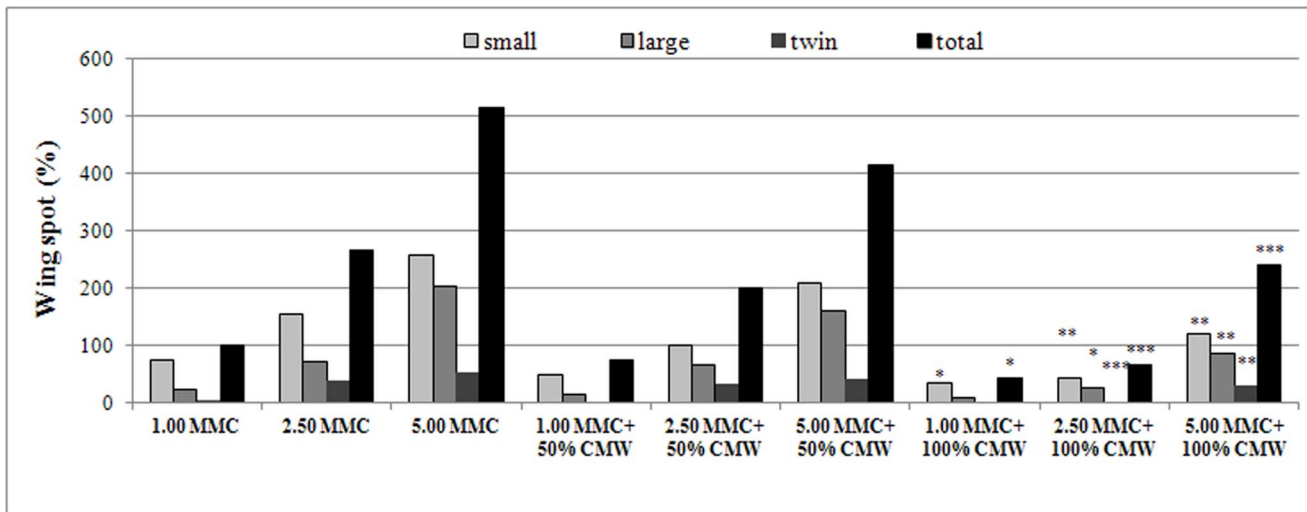


Figure 3. Wing spot frequency (%) induced by MMC (1.00, 2.50 and 5.00 µg/ml) in presence of CMW (50 and 100% v/v). [$*p < 0.05$, $**p < 0.01$, $***p < 0.001$, U-test].

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mastic's drastic constituents display the desirable effects in both antigenotoxic and therapeutic levels. The above is corroborated by a study of Doi et al. [78], according to which Chios mastic gum at high doses enhances the induction of preneoplastic lesions in rat liver. Thus, as proposed for other substances [79–82], some of CMW's constituents could potentially act as free radical scavengers at low concentrations and as pro-oxidants at higher concentrations. However, in our testing systems, the antigenotoxic activity of CMW was accomplished by quite different concentrations. Thus, even though in the CBMN assay CMW could afford protection against the used mutagen at low concentrations (1, 2 and 5% v/v), in the SMART test this was obtained by 100% (v/v) CMW (Table 2). In the latter assay 50% (v/v) CMW was unable to significantly reduce the MMC-induced mutagenic effects while 100% (v/v) CMW was capable to abolish completely the mutagenic effects induced by low concentrations of MMC (1.00 and 2.50 µg/ml) (Table 2). A number of factors may influence the observed differences in the *in vivo* and *in vitro* assays, such as compound absorption, rate and distribution of biotransportation, availability at the target site and cell permeability [83].

The antigenotoxic effects of CMW found, here, are supported by literature data demonstrating that some of its constituents, namely linalool [84], borneol [61] and perillyl alcohol [85] have been shown to exhibit antigenotoxic activity. Despite the low content of CMW in borneol and perillyl-alcohol, linalool is detected in a percentage of 7.29%, constituting one of CMW's major compounds [29]. Nevertheless, it should not be overseen that the observed protective effects of CMW would most likely be attributed to the additive/synergistic interaction of many major or minor constituents or to the combination of more than one biological activities [26,69]. Our data are consistent with the previously reported antigenotoxic and antioxidant properties of *P. lentiscus* extracts [68,69] and the antioxidant properties [5,9,12] as well as the anticancer effects of Chios mastic gum against a number of malignancies [22–28].

MMC is used in clinical cancer chemotherapy against a variety of solid neoplasms. However, due to its mutagenic and/or carcinogenic ability, secondary cancers are generated, which became a serious problem of chemotherapy. Thus, identifying new non-toxic phytochemicals capable of preventing DNA damage of

MMC is very important in developing novel nutraceuticals. Our results clearly show that CMW prevents or reduces DNA damage induced by MMC. Even though the mechanism of interaction between MMC and CMW is not known, the co-treatment protocol, used in the present study, cannot rule out the possibility of CMW acting as a desmutagen and interacting with the active groups of MMC [86]. Since CMW was present at the time of MMC exposure, it could inhibit the cytosol flavoreductases that activate MMC [87] and, thus, could block its activation and the subsequent DNA damage [88]. On the other hand, the observed antigenotoxic activity of CMW could also be ascribed to the mastic's antioxidant effects [5,9,12], as antioxidants are related with inhibition of mutagenesis [89]. This is further supported by the previously reported development of oxidative stress by MMC [74]. In any case these assumptions are not to be overestimated and further experiments are required to elucidate the mechanism by which CMW exerts its beneficial activity. Since this is the first report of anti-genotoxic activities of the CMW, further confirmation of these results could contribute to the development of herbal remedies containing natural active principles capable of compensating DNA damage and its subsequent outcomes such as cancer, accelerated ageing or degenerative conditions [90–93].

In conclusion, our work provides novel evidence that CMW does not exhibit any genotoxic or recombinogenic activity under our *in vitro* and *in vivo* experimental conditions. Moreover, CMW was found to possess antigenotoxic activity against the alkylating mutagen, MMC. The absence of the genotoxicity and the promising antigenotoxic activity of CMW suggest that this extract may contain phytochemical molecules of interest that could be used in a range of prospective applications in human healthcare. Although these results highlight the potential antigenotoxic properties of CMW, further studies are needed to delineate its pharmacological properties and its potential usefulness as a natural nontoxic dietary product.

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Author Contributions

Conceived and designed the experiments: DV ED PMT. Performed the experiments: DV DM ED IE TC CP MA EC PMT. Analyzed the data:

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