Production of Antibodies for Selective Detection of Malachite Green and the Related Triphenylmethane Dyes in Fish and Fishpond Water

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This study provides a practical method for production of the antibodies against malachite green (MG) and its primary metabolite leucomalachite green (LMG). Two ELISA kits are constructed with the MG and LMG antibodies for detection of the residual MG and LMG in fish muscle and fishpond water. The detection limit is established at the level of 0.05 𝜇g/L for both MG and LMG. Our ELISA kits show the advantages of good specificity, high sensitivity, and convenience in rapid screening of MG and LMG residues. The sample of fishpond water, without extraction or prior preparation, is directly assayed by the ELISA kit. More than 80 fish samples can be simultaneously tested in a kit. The toxic crystal violet and its metabolite leucocrystal violet of illegal use in aquaculture are detected by our prepared MG and LMG antibodies, whereas the antibodies do not cross-react with common antibiotics, sulfonamides, and benzene derivatives.

KEYWORDS: Malachite green; leucomalachite green; antibody; ELISA; fish

INTRODUCTION

Malachite green (MG, see Figure 1) (1), a dye of triphenylmethane skeleton, has been extensively used in aquaculture for prevention and treatment of external fungal and parasitic infections in fish. MG is easily absorbed by fish during waterborne exposure and is rapidly metabolized into leucomalachite green (LMG, see Figure 1). The reduction derivative LMG is recognized as a major metabolite of MG in fish (2, 3) and will store in fish muscle and tissues for months. Like other triphenylmethane dyes, MG and LMG cause carcinogenesis, mutagenesis, chromosomal fractures, teratogenesis, and respiratory toxicity in animals (4–7). Use of MG in aquatic food animals is highly restricted or banned in several countries because of toxicological considerations. However, illegal use of MG continues worldwide in aquaculture due to its low cost and ready availability (8, 9).

Surveillance of MG and LMG in aquaculture products is a necessary means to protect human health. According to the European Commission and U.S. Food & Drug Administration (10), methods that can be used for the determination of MG residues in fish muscles should meet a minimum required performance limit of 2 µg/kg for the sum of MG and LMG. High-performance liquid chromatography (HPLC) and liquid chromatography–mass spectrometry (LC-MS) are two general techniques for the quantitative analysis of MG and LMG in fish tissues (11–29). These techniques rely on expensive instruments operated by well-trained analysts, and prior preparation of samples is time-consuming and is not ideal for screening large number of samples. Alternatively, enzyme-linked immunosorbent assay (ELISA) is a rapid, specific, and sensitive method that is applicable to the on-site examination of a large number of samples. We thus generated two types of polyclonal antibodies against MG and LMG, and further developed the indirect competitive ELISA.

MATERIALS AND METHODS

All solvents and reagents were reagent-grade and were used without further purification. Crystal violet (CV, see Figure 1) chloride and malachite green oxalate were purchased from Merck (Rahway, NJ). Leucomalachite green, leucocrystal violet, Freund’s complete/incomplete adjuvant, bovine serum albumin (BSA), ovalbumin (OVA), and goat anti-rabbit IgG-HRP were purchased from Sigma-Aldrich (St. Louis, MO). Melting points are uncorrected. Infrared (IR) spectra were recorded on a Nicolet Magna 550-II spectrometer. Proton NMR (1H NMR) and carbon NMR (13C NMR and DEPT) spectra were recorded on Varian Unity Plus-400 (400 MHz) and Bruker Avance-400 FT-NMR spectrometers; chemical shifts are reported in unit δ relative to tetramethylsilane (TMS) with residual protons in the solvent as an internal standard.
standard: CDCl3, δ 7.24 (for 1H NMR) and δ 77.0 (for 13C NMR and DEPT). Mass spectra (MS) and high-resolution mass spectra (HRMS) were measured using a JEOL JMS-HX 110 spectrometer. Optical density (OD) of ELISA was measured using an ELISA reader (THERMOmax, Molecular Device, USA).

All experiments requiring anhydrous conditions were performed under an atmosphere of nitrogen. Reactions were monitored by thin-layer chromatography (TLC) using slides precoated with a 0.25 mm layer of silica gel containing a fluorescent indicator. Column chromatography was carried out on Kieselgel 60 (40–63 μm).

**Synthesis of Carboxymalachite Green (CMG, 3). Method A.** According to the previously described procedure (30), a mixture of 4-formylbenzoic acid (900 mg, 6.0 mmol), freshly distilled 1,8-dimethylaniline (2.4 mL, 19 mmol), and anhydrous ZnCl2 (2.4 g, 18 mmol) in absolute ethanol (60 mL) was heated at reflux for 24 h under an atmosphere of nitrogen. The mixture was cooled, and methanol (30 mL) and aqueous HCl (1 M) were added until pH = 5 to give slightly greenish crystals. The crystals were collected by filtration, rinsed with water, and dried over KOH in vacuum to give 1.8 g of 4-[bis(4-dimethylaminophenyl)methyl] benzoic acid, CMG (30%–34%), in 80% yield.

**Method B.** Under an atmosphere of nitrogen, a mixture of methyl 4-formylbenzoate (0.55 g, 3.4 mmol), freshly distilled N,N-dimethylaniline (1.5 mL, 11.8 mmol) and conc H2SO4 (0.25 mL of 96% solution) was heated at reflux for 24 h under an atmosphere of nitrogen. The mixture was cooled, and methanol (30 mL) and aqueous HCl (1 M) were added until pH = 5 to give slightly greenish crystals. The crystals were collected by filtration, rinsed with water, and dried over KOH in vacuum to give 1.8 g of 4-[bis(4-dimethylaminophenyl)methyl] benzoic acid, CMG (30%–34%), in 80% yield.

**CLMG.** Light greenish solid, mp = 250 °C (decomposed); IR (KBr) 3417, 2886, 1682, 1611, 1519, 1343, 1279, 1166 cm−1; 1H NMR (CDCl3, 400 MHz) δ 7.98 (2 H, d, J = 8.4 Hz), 7.22 (2 H, d, J = 8.4 Hz), 6.95–6.93 (4 H, m), 6.66 (4 H, d, J = 8.4 Hz), 5.42 (1 H, s), 2.90 (12 H, s); 13C NMR (CDCl3, 100 MHz) δ 171.01, 151.25 (2x), 148.44 (2x), 131.62, 129.68 (2x), 129.51 (4x), 129.05 (2x), 126.61, 112.54 (4x), 55.23, 41.08 (2x); HRMS (ESI) calculated for C24H27N2O2: 375.2073, found: m/z 375.2022 [M+H]+.

**CLMG Methyl Ester.** White solid, mp = 128–129 °C; TLC (EtOAc/hexane (1:9)) Rf = 0.10; IR (KBr) 2800, 1720, 1612, 1519, 1441, 1343, 1279, 1110, 1102 cm−1; 1H NMR (CDCl3, 400 MHz) δ 7.90 (2 H, d, J = 8.4 Hz), 7.19 (2 H, d, J = 8.4 Hz), 6.95–6.93 (4 H, m), 6.64 (4 H, d, J = 8.8 Hz), 5.40 (1 H, s), 3.87 (3 H, s), 2.91 (12 H, s); 13C NMR (CDCl3, 100 MHz) δ 166.45, 150.35 (2x), 148.37 (2x), 131.50, 129.46 (4x), 129.01 (2x), 128.92 (2x), 127.28, 112.27 (4x), 55.11, 52.11, 40.95 (4x); HRMS (ESI) calculated for C25H29N2O2: 389.2229, found: m/z 389.2259 [M+H]+.

**Synthesis of Carboxymalachite Green (CMG, 4). Method A.** According to the previously described procedure (30–34), a mixture of CLMG (375 mg, 1 mmol), chloranil (295 mg, 1.2 mmol), and glacial acetic acid (0.75 mL) in CHCl3 (45 mL) was stirred at 25 °C for 1.5 h. The solids were collected by filtration, rinsed with CHCl3/CCl4 (1:1), and dried over KOH in vacuum to give crude CMG (360 mg), which was used without further purification for coupling with protein.

**Method B.** According to the previously described procedure (34), a mixture of CMG (150 mg, 0.60 mmol) and PbO2 (0.63 mmol) in aqueous HCl (1.5 mL of 2 M solution) was stirred at 25 °C for 18 h. The mixture was diluted with MeOH (30 mL) and filtered through a Celite bed. The filtrate was concentrated in vacuum to give green solids of CMG (246 mg), which were used without further purification for coupling with protein. By a similar procedure, CMG methyl ester was treated with PbO2 in aqueous HCl to give CMG methyl ester. Complete oxidation of CLMG using both methods was indicated by disappearance of the methine signal at δ 5.42 in the 1H NMR spectrum of the CMG product. The structure of CMG was supported by an exact mass measurement, giving m/z 373.1895 for the parent ion [C24H25N2O2]+. The prepared CMG was estimated at ~90% purity, and was good enough for the antibody production as shown in the inhibition assay without interference of any impurity (see Figure 5a).
**Characterization of MG and LMG Antibodies.** The 96-well plates (Corning, NY) were coated with MG-BSA or LMG-BSA overnight at 4 °C. The plates were washed 3 times with PBST, and then blocked with 0.1% skim milk at 37 °C for 1 h. The plates were again washed with PBST, and incubated with animal sera (100 µL/well) diluted in 0.1% BSA at 37 °C for 1 h. After incubation, plates were washed with PBST three times. Goat anti-rabbit IgG antibody conjugated with HRP was diluted (1/2000) in 0.1% BSA, and 100 µL of the resultant solution was added to each well. The plates were incubated at 37 °C for 1 h, and washed again with PBST three times. The substrate solution TMB (100 µL) was added to each well and incubated at 37 °C for 10 min. The reaction was stopped by addition of aqueous HCl (50 µL of 2 M solution). Optical density at a wavelength of 450 nm in each well was read using an ELISA reader.

The MG- and LMG-specific antibodies were purified from high-titer rabbit serum by affinity chromatography on a Protein A-Sepharose column (GE Healthcare, UK). The antibodies were concentrated using Centriprep (Amicon Ultra, 5000 MWCO; Millipore, USA). Antibody concentrations was measured with Bio-Rad protein assay (Bio-Rad Laboratories, Hercules, CA).

The specificity of MG and LMG antibodies was determined by indirect competitive ELISA using MG, LMG, CV, and leucocrystal violet (LCV) as inhibitors. The inhibitors were diluted in 0.02 M HCl with 0.15 M NaCl. The indirect competitive ELISA was performed on MG-BSA or LMG-BSA coated 96-well plates. Serially diluted inhibitors were incubated with 0.5 µg/mL of MG or 1.8 µg/mL of LMG antibodies in the antigen-coated 96-well plates at 37 °C for 30 min. Antibodies bound to the MG-BSA or LMG-BSA were detected by goat anti-rabbit IgG-HRP, followed by treatment with TMB. The optical density at a wavelength of 450 nm was recorded using an ELISA reader. The cross-reactivity of MG and LMG antibodies was expressed as the ratio of MG or LMG concentration to the cross-reactant concentration that will show 50% of B/B0 %.

**Preparation of Fish Samples for ELISA.** Fish samples were obtained from the local market including *Oreochomis* sp. (Tilapia and Taiwan Tilapia), *Chanos chanos* (milkfish), *Epinephelus* sp. (groupers), *Lateolabrax japonicus* (Japanese sea perch), *Micropterus salmoides* (California bass), and *Pagrus major* (Red sea bream). The fish was filleted, the skin and bones were removed, and the muscles were minced and frozen before being analyzed. Accurately weighed 1.0 g of homogenized fish muscle was put into a 15 mL centrifuge tube, which can be spiked with appropriate amounts of MG or LMG, and 1 mL of McIlvaine buffer (pH = 3.0) and 6 mL of acetonitrile were added. The mixture was vigorously vortexed for 3 min, and centrifuged at 3500 rpm for 10 min. The supernatant was transferred into a centrifuge tube with 1.5 mL of CH3Cl, and the sample was vortex-mixed followed by centrifuging at 3500 rpm for 10 min. An aliquot of the upper organic phase (0.5 µL) was transferred into a tube (1.5 mL), and treated with strong anion exchanger AG 1 resin (0.1 mL, Bio-Rad Laboratories, Hercules, CA) at room temperature for 30 min to reduce interference from the extracts. The supernatant (0.4 µL) was dried by evaporation under a stream of nitrogen at 50 °C. The sample was reconstituted with 0.4 mL of sample diluent (0.01 M HCl) and detected by an MG or LMG ELISA.

**MG ELISA of Fish Samples.** MG-BSA was coated on a 96-well ELISA plate and reacted with fish samples and MG antibodies in equal volume at 37 °C for 30 min. Unbound MG antibodies were removed by washing. Goat anti-rabbit IgG-HRP was added and treated with TMB. The quantity of MG antibodies bound on the MG ELISA plate was deduced from the OD at 450 nm. The percent binding (B/B0 %) for each standard or sample was calculated by the following equation, and the corresponding concentration of MG was interpolated from the standard curve. The sample concentration was corrected for dilution (7-fold).

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\text{[OD (standard or sample)/OD (blank)]} \times 100\% = \text{B/B0 %}
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**LMG ELISA of Fish Samples.** For LMG ELISA, an ELISA plate was coated with LMG-BSA. The fish samples and LMG antibodies in equal volume were subjected to indirect competitive ELISA, and the concentration of LMG was calculated by a procedure similar to that for MG ELISA.

**Preparation of Immunogens by Coupling of CMG and CLMG with Proteins.** In this study, OVA was used as a carrier for MG and LMG immunogens, and BSA was used as a carrier for the coated antigens in ELISA. To 0.85 mL of CMG or CLMG solution at 1 mg/mL in 0.1 M MES (pH 6.0) with 50% Me2SO were added antigens in ELISA. To 0.85 mL of CMG or CLMG solution at 1 mg/mL, 200 µL of 3.0% -hydroxysuccinimide (NHS, 0.26 mg, Pierce, USA). The mixture was stirred for 30 min at room temperature, and added dropwise to 5 mL of 1 mg/mL protein solution. The mixture was agitated for 2 h at room temperature. The conjugates were dialyzed against 0.1 M phosphate buffer (pH 7.4) to remove free reagents.

**Immunization.** Two groups of New Zealand rabbits (in duplicate) were immunized by sc injection with CMG-OVA or CLMG-OVA. Primary immunizations were composed of 500 µL of PBS containing 100 µg of antigen emulsified in 500 µl of Freund’s complete adjuvant. Subsequent immunizations, at 2–5 week intervals, were of the same volume, with complete adjuvant replaced by incomplete adjuvant. Rabbits were bled after each injection. The sera titer was detected by an indirect ELISA.

**Detection of Triphenylmethane Dyes in Fish and Fishpond Water.** The sample was extracted by homogenization of homogenized fish muscle was put into a 15 mL centrifuge tube, which can be spiked with appropriate amounts of MG or LMG, and 1 mL of McIlvaine buffer (pH = 3.0) and 6 mL of acetonitrile were added. The mixture was vigorously vortexed for 3 min, and centrifuged at 3500 rpm for 10 min. The supernatant was transferred into a centrifuge tube with 1.5 mL of CH3Cl, and the sample was vortex-mixed followed by centrifuging at 3500 rpm for 10 min. An aliquot of the upper organic phase (0.5 µL) was transferred into a tube (1.5 mL), and treated with strong anion exchanger AG 1 resin (0.1 mL, Bio-Rad Laboratories, Hercules, CA) at room temperature for 30 min to reduce interference from the extracts. The supernatant (0.4 µL) was dried by evaporation under a stream of nitrogen at 50 °C. The sample was reconstituted with 0.4 mL of sample diluent (0.01 M HCl) and detected by an MG or LMG ELISA.

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**LMG ELISA of Fish Samples.** For LMG ELISA, an ELISA plate was coated with LMG-BSA. The fish samples and LMG antibodies in equal volume were subjected to indirect competitive ELISA, and the concentration of LMG was calculated by a procedure similar to that for MG ELISA.
MG ELISA of Fishpond Water. The water sample collected from fishponds was diluted with 3 volumes of 0.02 M HCl, with or without spiking of MG at 0.5 or 2 µg/L. The sample was centrifuged at 3500 rpm for 10 min, and an aliquot of supernatant (100 µL) was taken for the MG ELISA.

RESULTS AND DISCUSSION

The compounds CLMG (3) and CMG (4) bearing a carboxyl group on the phenyl ring were designed as the appropriate analogues of MG and LMG for linkage with carrier proteins (Scheme 1). The synthesis of CLMG and CMG is straightforward by using known methods with minor modification (30–34).

LCMG and CMG were attached to the carrier proteins BSA and OVA via amide bond formation using NHS and EDCI as the condensation agents. Two groups of New Zealand rabbits (in duplicate) were immunized by subcutaneous injection with the MG- and LMG-OVA immunogens prepared as such to generate the specific antibodies, respectively, using the standard procedures. The results of the MG- and LMG-ELISA produced linear ranges from 0.05 to 0.5 µg/L of MG (Figure 2a) and from 0.05 to 2 µg/L of LMG (Figure 2b). The detection limit of this assay was 0.05 µg/L for both MG and LMG, lower than the regulatory limit by the European Commission and U.S. Food & Drug Administration (10).

Both our MG- and LMG-ELISA kits use a cutoff level of 1 µg/kg to distinguish positive from negative samples. A total of 53 fish samples were detected by MG ELISA. The MG concentrations in fish muscle were spiked with 0.5 or 2 µg/kg of MG, followed by 7-fold dilution, in appropriate cases for the ELISA experiments. There was 98% agreement between the results obtained by ELISA and theoretical concentration. The result shows 100% sensitivity and 96% specificity (Table 1). The recovery of MG-spiked samples was varied from 71% to 108%. A total of 115 fish samples were detected by LMG ELISA. The LMG concentrations in fish muscle were spiked with 0.5, 0.75, 1.5, or 2 µg/kg of LMG in appropriate cases for analysis. There was 90% agreement between the results obtained by ELISA and those by theoretical concentration. The result shows 83% sensitivity and 98% specificity (Table 1). The recovery of LMG-spiked samples was varied from 62% to 105%. It was noted that fatty samples might cause lower recovery of LMG, and thus inferior sensitivity in ELISA. Presumably due to the lipophilic nature of LMG, the extraction from fatty samples might become less efficient.

| Table 1. Comparison of MG and LMG Concentrations Determined by ELISA and Theoretical Concentrations in Fish Muscle (n = 53 for MG and n = 115 for LMG) |
|---------------|---------------|---------------|---------------|
| ELISA         | MG            | LMG           |
| +             | 26            | 1             | 50            |
| –             | 0             | 26            | 10            |
| sensitivity   | 100%          | 96%           | 83%           |
| specificity   |               |               | 98%           |

| Table 2. Comparison of MG Concentrations Determined by ELISA and Theoretical Concentrations in Water Samples (n = 27) |
|---------------|---------------|
| ELISA         | MG            |
| +             | 9             |
| –             | 0             |
| sensitivity   | 100%          |
| specificity   | 100%          |

MG ELISA was also used for fishpond water detection. The samples were diluted and directly analyzed by MG ELISA without the need of extraction or prior preparation. A total of 27 water samples were tested. The result showed that sensitivity and specificity are both 100% (Table 2).

The MG and LMG antibodies were insensitive to common antibiotics, sulfonamides, and benzene derivatives. No cross-reactivity (<0.005%) in the ELISA was observed in the 16 test compounds, including sulfadiazine, sulfamonomethoxine, sulfamethazine, sulfamethoxypyridazine, sulfadimethoxine, sulfathiazole, sulfaguanidine, Sulfamethoxazole, penicillin G, gentamicin, oxytetracycline, tetracycline, chloramphenicol, aniline, dimethylaniline and benzaldehyde, up to a concentration of 1000 µg/L.

The specificity of MG and LMG antibodies for the triphenylmethane dyes MG and CV as well as their metabolites LMG and LCV was investigated by indirect competitive ELISA. The MG antibody showed 100% cross-reactivity to CV, but was...
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insensitive to LMG or LCV (Figure 33a). This result might be attributable to the similar structures between MG and CV having 4-(dimethylamino)phenyl groups surround the methylium center of planar shape (Figure 11). Because LCV exhibited a methine center of tetrahedral shape, as that of LMG, the LMG antibody also reacted strongly with LCV (200% cross-reactivity), but not with CV (Figure 3b). A slight cross-reactivity of the LMG antibody with MG (~3%) was also observed. The reason is unclear, though we speculate that a small amount of MG may be derived from air oxidation of LMG during the process of immunogen preparation and immunization.

To our knowledge, using the commercially available ELISA kits, e.g., MaxSignal Malachite Green ELISA Test Kit of Bioo Scientific Co. (Texas, USA), for detection of LMG in fish samples requires a prior oxidation of LMG to MG. On the other hand, our LMG ELISA is directly applied to detection of LMG in fish samples. Though a side-by-side comparison experiment was not carried out, our MG ELISA shows a detection limit of 0.05 µg/L, better than the BIOO specification (0.5 µg/L).

In conclusion, we have successfully utilized CMG-OVA and CLMG-OVA as immunogens to produce, respectively, a high titer of rabbit polyclonal antibodies against MG and LMG. Detection of MG and LMG residues in spiked fish muscle with the ELISA kits showed 96% specificity and 100% sensitivity for the MG antibody, as well as 98% specificity and 83% sensitivity for the LMG antibody. The diluted fishpond water, without extraction or prior preparation, was directly subjected to our MG ELISA, which shows a detection limit of 0.05 µg/L, better than the BIOO specification (0.5 µg/L).

ABBREVIATIONS USED

BSA, bovine serum albumin; CLMG, carboxyleucomalachite green; CMG, carboxymalachite green; CV, crystal violet; DEPT, distorsionless enhancement by polarization transfer; EDCl, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide; ELISA, enzyme-linked immunosorbent assay; ESI, electrospray ionization; FT-NMR spectrometer, Fourier transform nuclear magnetic resonance spectrometer; HPLC, high performance liquid chromatography; HRMS, high-resolution mass spectra; HRP, horseradish peroxidase; IgG, immunoglobin G; IR spectra, infrared spectra; LC-MS, liquid chromatography–mass spectrometry; LCV, leuco metabolites; LMG, leucomalachite green; MG, malachite green; MES, 2-(N-morpholino)ethanesulfonic acid; MS, mass spectrometry or mass spectra; NHS, N-hydroxysuccinimide; NMR spectra, nuclear magnetic resonance spectra; OD, optical density; OVA, ovalbumin; PBS, phosphate-buffered saline; PBST, phosphate-buffered saline with 0.05% Tween 20; TMB, 3,3',5,5'-tetrathylbenzidine; TLC, thin-layer chromatography; TMS, tetramethylsilane.

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