Harmonine, a defence compound from the harlequin ladybird, inhibits mycobacterial growth and demonstrates multi-stage antimalarial activity

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The harlequin ladybird beetle Harmonia axyridis has been introduced in many countries as a biological control agent, but has become an invasive species threatening the biodiversity of native ladybirds. Its invasive success has been attributed to its vigorous resistance against diverse pathogens. This study demonstrates that harmonine ((17R,9Z)-1,17-diaminooctadec-9-ene), which is present in Harmonia axyridis haemolymph, displays broad-spectrum antimicrobial activity that includes human pathogens. Antibacterial activity is most pronounced against fast-growing mycobacteria and Mycobacterium tuberculosis, and the growth of both chloroquine-sensitive and -resistant Plasmodium falciparum strains is inhibited. Harmonine displays gametocytocidal activity, and inhibits the exflagellation of microgametocytes and zygote formation. In an Anopheles stephensi mosquito feeding model, harmonine displays transmission-blocking activity.

Keywords: Harmonia axyridis; insect immunity; harmonine; antimicrobial activity

1. INTRODUCTION
Harmonia axyridis, known as the Asian lady beetle or the harlequin ladybird, is a ladybird beetle native to continental, temperate and subtropical parts of East and Central Asia. Since the beginning of the twentieth century, this species has been introduced as a biological control agent against aphid and/or coccid pests into North America, Europe and the Soviet Union. In addition, H. axyridis has been commercially available as a biological control agent for greenhouses and urban ecosystems since the mid-1990s. Over the last two decades, H. axyridis has become an invasive species in many countries. In Europe, H. axyridis populations have been growing rapidly since the turn of the millennium, threatening populations of native ladybird species [1]. Its invasive success has been attributed to its enduring resistance against diverse pathogens, which allows it to outperform and therefore dominate the most abundant native European ladybirds, Coccinella septempunctata and Adalia bipunctata [2]. Besides antimicrobial peptides encoded by small genes and synthesized on ribosomes [3], many insects synthesize low-molecular mass defence compounds, or sequester such compounds from their diet. Ladybirds exude droplets of haemolymph containing deterrent alkaloids through their leg joints when threatened or attacked, a behaviour known as reflex bleeding [4]. In the present study, harmonine ((17R,9Z)-1,17-diaminooctadec-9-ene) was identified as the principal antimicrobial compound of H. axyridis haemolymph.

2. MATERIAL AND METHODS

(a) Origin and rearing of ladybirds
Adults of H. axyridis subsequently used for captive breeding were collected in and around Gießen and Ober-Mörlen, Germany. Adults of the seven-spot ladybird (C. septempunctata) and eggs of the two-spot ladybird (A. bipunctata) were obtained from Kath Biotech AG (Baruth, Germany). All ladybird species were reared in cages at 26°C and 60 per cent relative humidity under a 16:8 photoperiod. Bean plants (Phaseolus vulgaris) infested by pea aphids (Acyrthosiphon pisum) were provided as a food source.

(b) Purification, structure determination and synthesis of harmonine
Haemolymph released by reflex bleeding was collected from 500 H. axyridis beetles. Groups of five beetles were vortexed for 10 s in 0.2 ml water in a 1.5 ml tube, and the combined liquid was heated to 95°C for 1 h and the precipitated material removed by centrifugation. The supernatant was supplemented with acetonitrile to a final concentration of 20 per cent (v/v) and passed over a strong anion exchange solid-phase extraction cartridge (ISOLUTE SAX 100 mg/3 ml, Biotage). The flow-through was loaded onto a strong cation exchange column (Mono S 5 ml, Biotage). The flow-through was loaded onto a strong cation exchange column (Mono S 5/50 GL, GE Healthcare) and eluted with a linear gradient of NaCl (0–1 M in water containing 20% acetonitrile). Fractions containing active compounds from the radial diffusion assay eluted at approximately 700 mM NaCl. After removal of excess acetonitrile by vacuum evaporation, final purification was achieved by chromatography on a reversed-phase column (Acclaim 120, C18, 3 µm, 4.6 x 150 mm; Dionex) by applying a gradient of 8–80% acetonitrile in water containing 1% per cent formic acid. The activity was recovered at approximately 45 per cent acetonitrile. Structure determination was performed on a microOTOF-Q II mass spectrometer (Bruker Daltonics). Harmonine was synthesized following the protocol of Enders & Bartzen [5].

(c) Antibacterial activity
For radial diffusion assays, beetles were homogenized in 20 per cent acetonitrile (10 µl mg-1 beetle weight), and 5 µl of the supernatant was applied to yeast extract and tryptone agar test plates (well diameter 3 mm) containing Escherichia coli DE 31. Minimal inhibitory concentration (MIC) values were determined in triplicate with 1 : 2 serial dilutions. Activity against Mycobacterium tuberculosis was determined using the BACTEC MGIT 960 system (Becton Dickinson).

(d) Antimalarial activity
Synchronized cultures containing Plasmodium falciparum ring forms were plated in 96-well plates at a parasitemia of 1 per cent in the presence of 1 : 2 serial dilutions of harmonine. After incubating the plates for 72 h, the viability of the parasites was assessed using the Malstat assay [6]. Gametocytocidal activity was determined by plating stage II P. falciparum gametocytes in triplicate in 24-well plates in the presence of harmonine [7]. The cultures were incubated with harmonine for 48 h and then for another 5 days without the compound. The numbers of stage IV and V gametocytes in 1000 red blood cells were counted. Inhibition of microgametogenesis was determined by adding harmonine to mature gametocyte cultures for 15 min at 37°C prior to activation with 100 µM xanthurenic acid. After another 15 min, the numbers of exflagellation centres were

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counted in 30 optical fields using a Leica DMLS microscope (400× magnification). Inhibition of zygote formation was determined by adding harmonine to mature gametocyte cultures prior to activation. The activated cultures were incubated at room temperature for 20 h. Zygotes that had subsequently formed were highlighted using anti-Pfs25 antibodies (ATCC). The numbers of zygotes were counted in 90 optical fields at 400× magnification. Transmission-blocking activity was determined by feeding female Anopheles stephensi mosquitoes on harmonine-containing gametocyte cultures. The mosquito midguts were dissected 12 days post-infection and inspected for oocysts by mercurochrome staining.

(e) Antiproliferative and cytotoxic activity

Cells (HUVEC (ATCC CRL-17230), K-562 (DSMZ ACC 10) and MCF-7 (DSMZ ACC 115), S9 (Promega), High Five (Invitrogen)) were plated together with harmonine in 96-well plates and incubated for 72 h. Cell viability was determined using the CellTiter-Blue (Promega) or 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. Subconfluent monolayers of HeLa cells (DSMZ ACC 57) were incubated with harmonine for 72 h before staining with methylene blue.

3. RESULTS

(a) Bioactivity guided purification of harmonine

In initial experiments, the antimicrobial activity of haemolymph from ladybirds was assessed using a radial diffusion assay. Harmonia axyridis haemolymph generated an inhibition zone that was not observed with haemolymph from C. septempunctata or A. bipunctata (figure 1a). The active compound was purified by collecting haemolymph released by reflex bleeding from 500 H. axyridis beetles. Applying the radial diffusion assay for the detection of active fractions, a purification procedure by heat denaturation, anion exchange chromatography (activity in flow-through), cation exchange chromatography and reversed-phase high-performance liquid chromatography resulted in a single compound, which was identified as harmonine ((17R,9Z)-1,17-diaminooctadec-9-ene) (figure 1b) by electrospray ionization quadrupole time-of-flight (ESI-Qq-TOF) mass spectrometry. Using synthetic harmonine as standard, the amount of harmonine contained in an individual H. axyridis beetle was determined to be 160 ± 44 mg. With an estimated haemolymph volume of 21 ± 4 μl (determined by weight loss on drying), this corresponds to a harmonine concentration of approximately 27 mM. Harmonine was neither detectable in C. septempunctata nor in A. bipunctata haemolymph (figure 1c).

(b) Antibacterial activity

Synthetic harmonine displayed activity against 12 bacterial strains and the yeast Candida albicans with MIC

Figure 1. Detection of harmonine as the principal antimicrobial compound in H. axyridis haemolymph. (a) Haemolymph from three individual beetles from each of the three ladybird species were tested in radial diffusion assays against Escherichia coli. The activity of the H. axyridis samples corresponds to 499, 473 and 535 μg ml−1 gentamycin, respectively. (b) Chemical structure of harmonine. (c) The haemolymph samples used for the radial diffusion assays were analysed for the presence of harmonine by liquid chromatography–mass spectrometry. Extracted ion chromatograms were recorded at m/z 283 ± 0.5. Synthetic harmonine was analysed as a control.
values in the range between 5.5 and 354 μM (table 1). Harmonine proved to be equally active against a drug-sensitive \textit{Staphylococcus aureus} and an multi-resistant \textit{S. aureus} (MRSA) strain. Pronounced activity was observed against four strains of fast-growing mycobacteria (MIC values of 5.5 μM against four strains of fast-growing mycobacteria was also observed against \textit{Mycobacterium fortuitum} and \textit{Mycobacterium smegmatis}). Comparable activity was also observed against \textit{Mycobacterium vaccae} and \textit{Candida albicans}.

(d) \textbf{Antiproliferative and cytotoxic activity}

Proliferation of the human cell lines HUVEC, K-562 and MCF-7 was inhibited by harmonine with IC$_{50}$ values of 21.3 ± 2.1, 18.5 ± 0.6 and 38.0 ± 5.6 μM, respectively. The IC$_{50}$ values for the lepidopteran cell lines SF9 and High Five were 57 ± 8 and 53 ± 18 μM, respectively. Cytotoxicity for HeLa cells was observed at 37.0 ± 1.7 μM, causing 50 per cent destruction of the cell monolayer.

4. DISCUSSION

Harmonine had previously been isolated by Braconier \textit{et al.} \cite{8} based on its reactivity with Dragendorff’s reagent. The same group showed that the compound acted as a feeding deterrent against the common red ant, \textit{Myrmica rubra} \cite{9}. Alam \textit{et al.} \cite{10} reported cytotoxicity against five human solid tumour cell lines and moderate inhibition of the enzymes acetylcholinesterase, prolyl endopeptidase and neuraminidase. The broad-spectrum antimicrobial activity observed in the present study demonstrates that harmonine is an important factor in beetle immunity and may explain the invasive success of \textit{H. axyridis}. Although harmonine was less active than standard antibiotics, the MIC values were significantly below the estimated harmonine concentration in \textit{H. axyridis} haemolymph. At antibacterial concentrations, harmonine also displayed antiproliferative and cytotoxic activity against human and lepidopteran cell lines. How \textit{H. axyridis} is able to resist the cytotoxic potential of harmonine remains unknown. The pronounced activity of harmonine against mycobacteria together with activity against an MRSA strain is indicative for a novel mode of action, which might be exploited by the development of derivatives less toxic to human cells. At remarkably low concentrations, harmonine inhibited the growth of the malaria parasite \textit{Plasmodium falciparum} and prevented transmission of sexual parasite stages to the mosquito. Therefore, further studies may provide a base for the development of novel anti-malarial drugs with both parasitocidal and transmission-blocking activities.
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