Functional properties of edible insects: a systematic review

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Abstract

Consumption of edible insects has been widely suggested as an environmentally sustainable substitute for meat to reduce greenhouse gas emissions. However, the novel research field for edible insects relies on the content of bioactive ingredients and on the ability to induce a functional effect in humans. The goal of this manuscript is to review the available body of evidence on the properties of edible insects in modulating oxidative and inflammatory stress, platelet aggregation, lipid and glucose metabolism and weight control. A search for literature investigating the functional role of edible insects was carried out in the PubMed database using specific keywords. A total of 55 studies, meeting inclusion criteria after screening, were divided on the basis of the experimental approach: *in vitro* studies, cellular models/*ex vivo* studies or *in vivo* studies. In the majority of the studies, insects demonstrated the ability to reduce oxidative stress, modulate antioxidant status, restore the impaired activity of antioxidant enzymes and reduce markers of oxidative damage. Edible insects displayed anti-inflammatory activity reducing cytokines and modulating specific transcription factors. Results from animal studies suggest that edible insects can modulate lipid and glucose metabolism. The limited number of studies focused on the assessment of anti-coagulation activity of edible insects makes it difficult to draw conclusions. More evidence from dietary intervention studies in humans is needed to support the promising evidence from *in vitro* and animal models about the functional role of edible insect consumption.

Key words: Antioxidants: Edible insects: Functional foods: Health: Inflammation

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Introduction

The increasing global food demand, strictly linked to the growing world population, expected to reach almost 10 billion in

2050, has brought a new global interest on the human consumption of edible insects⁽¹⁾. The consumption of insects by humans is not a novel phenomenon, having been practiced since early in human evolution, and is nowadays part of the usual diet in many

Abbreviations: 8-OHdG, 8-hydroxy-2'-deoxyguanosine; ABCA1, ATP-binding cassette transporter A1; ABCG1, ATP-binding cassette transporter G1; ABTS, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid); ACAT2, acetyl-CoA acetyltransferase 2; ACE, angiotensin-converting enzyme; ADP, adenosine diphosphate; ADRP, adipose differentiation-related protein; AGPAT1, 1-acylglycerol-3-phosphate O-acyltransferase 1; AMPK- α , adenosine monophosphate-activated protein kinase- α ; aPTT, activated partial thromboplastin time; AUC, area under the curve; C/EBP α , CCAAT/enhancer-binding protein α ; CA, chromosome aberration; BW, body weight; BWG, body weight gain; CAT, catalase; CD36, cluster of differentiation 36; ChREBP, carbohydrate-response element-binding protein; CLP, caecal ligation and puncture; CP, compound; CYP7A1, cholesterol 7 α hydroxylase; DGAT1, diacylglycerol O-acyltransferase 1; D-HMVECs, diabetic type 2 microvascular endothelial cells; DPPH, 1,1-diphenyl-2-picrylhydrazyl; DPP-IV, dipeptidyl peptidase-4; ELOVL2, fatty acid elongase 2; ELOVL5, fatty acid elongase 5; ERK, extracellular signal-regulated kinases; ET-1, endothelin-1; F4/80+ KCs, F4/80-positive Kupffer cells; FABP1, fatty acid-binding protein 1; FADS2, fatty acid desaturase 2; FAS, fatty acid synthase; FATP5, fatty acid transport protein 5; FRAP, ferric reducing antioxidant power; FXa, factor Xa; GHG, greenhouse gas; GI, glucose index; GM-CSF, granulocyte-macrophage colony-stimulating factor; GPAT1, glycerol-3-phosphate acyltransferase 1; GPAT4, glycerol-3-phosphate acyltransferase 4; GPx, glutathione peroxidase; GR, glutathione reductase; GST, glutathione S-transferase; HbA1c, glycosylated haemoglobin concentration; HDL, high-density lipoprotein; HMGCR, 3-hydroxy-3-methylglutaryl-CoA reductase; HOMA-IR, homeostasis model assessment of insulin resistance; HUVECs, human umbilical vein endothelial cells; ICAM-1, intercellular adhesion molecule-1; IFN α , interferon α ; IFN- γ , interferon γ ; IgA, immunoglobulin A; IgE, immunoglobulin E; IgG, immunoglobulin G; IgM, immunoglobulin M; IL-10, interleukin-10; IL-12, interleukin-12; IL-13, interleukin-13; IL-1 α , interleukin-1 α ; IL-1 β , interleukin-1 β ; IL-2, interleukin-2; IL-4, interleukin-4; IL-5, interleukin-5; IL-6, interleukin-6; IL-7, interleukin-7; IL-8, interleukin-8; ISI, insulin sensitivity index; ITT, insulin tolerance test; JNK, c-Jun N-terminal kinase; LDL, low-density lipoprotein; LPL, lipoprotein lipase; LPS, lipopolysaccharide; LS, lipo-soluble extract; LXR α , liver X receptor; MARCKS, myristoylated alanine-rich C-kinase substrate; MDA, malondialdehyde; MN, micronucleus; MUFA, monounsaturated fatty acids; NEFA, non-esterified fatty acid; NF- κ B, nuclear factor- κ B; NO, nitric oxide; Nrf2, nuclear factor erythroid 2-related factor; OGTT, oral glucose tolerance test; ORAC, oxygen radical absorbance capacity; PAI-1, plasminogen activator inhibitor-1; PAP1, phosphatidate phosphatase 1; PH, protein hydrolysates; p-IkB- α , phosphorylated-inhibitor of nuclear factor-kappa B-alpha; PPAR γ , peroxisome proliferator-activated receptor γ ; PT, prothrombin time; PUFA, polyunsaturated fatty acids; RNS, reactive nitrogen species; ROS, reactive oxygen species; SAHR, scavenging activity on hydroxyl radicals; SCD1, stearoyl-CoA desaturase-1; SCE, sister chromatid exchange; SCFA, short-chain fatty acids; SFA, saturated fatty acids; SOD, superoxide dismutase; SREBP-1c, sterol regulatory element-binding protein 1c; SREBP-2, sterol regulatory element-binding protein 2; SRSC, superoxide radical scavenging capacity; TAC, total antioxidant capacity; TBA, total bile acid; TC, total cholesterol; TEAC, Trolox equivalent antioxidant capacity; TG, triacylglycerol; TLR4, toll-like receptor 4; TNF- α , tumour necrosis factor- α ; TOS, total oxidant status; tPA, tissue plasminogen activator; TT, thrombin time; VCAM-1, vascular cell adhesion molecule-1; VEGF, vascular endothelial growth factor; WS, water-soluble extract.

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countries⁽²⁾. Edible insects have positive environmental and economic implications and are characterised by high feed-conversion efficiency and low greenhouse gas (GHG) emissions, land use and environmental contamination⁽¹⁾. Recently, consumption of edible insects has been widely suggested as an environmentally sustainable substitute for meat and animal products to reduce their extremely high global intake, responsible for massive GHG emission⁽¹⁾.

From a nutritional point of view, edible insects are a good source of nutrients characterised by high levels of essential amino acids, fibre, vitamin B12, iron, zinc, omega-3 and omega-6 fatty acids, and polysaccharides and might contribute to a balanced gut microbiota⁽³⁾. However, the innovative aspect of edible insects is related to their content of bioactive ingredients and their role as functional foods, defined as foods providing a health benefit beyond basic nutrition.

Throughout human history, entomotherapy, the medicinal use of edible insects, has been used in traditional medicine in wound healing and as curative therapy for respiratory disorders and stomachache⁽⁴⁾. The available body of evidence on the functional role of edible insects is mainly focused on their role as antioxidant ingredients *in vitro* or in cellular models, as recently reviewed by D'Antonio *et al.*⁽⁵⁾. Moreover, there is scattered evidence on the role of edible insects in platelet aggregation, as anti-inflammatory agents⁽⁶⁾ or as modulators of lipid⁽⁷⁾ and glucose metabolism⁽⁸⁾. Furthermore, the available studies tested a wide range of different insects, extracts or protein fractions, making it difficult to draw clear conclusions about their functional properties without a review of the available evidence. Recently, findings related to the nutritional, functional and health properties as well as consumer acceptance of edible insects have been thoroughly reviewed^(9–13). However, none of the manuscripts provided a comprehensive and systematic review of the available evidence on the functional roles of edible insects.

The goal of this work is to review the available body of evidence on the properties of edible insects in modulating oxidative and inflammatory stress, platelet aggregation, weight control, and lipid and glucose metabolism.

Material and methods

The search strategy of the study is shown in Figure 1. We first systematically searched PubMed database (National Library of Medicine, Bethesda, MD; <https://pubmed.ncbi.nlm.nih.gov>) using the following keywords: (edible insect) AND (functional) OR cardiovascular) OR platelet aggregation) OR oxidative) OR antioxidant) OR glycemia) OR diabetes) OR inflammation) OR immune) OR cholesterol) OR hypolipidemic) OR hypocholesterolemic) OR thrombosis) OR hepatoprotective) OR liver) OR adipose tissue) OR body composition) OR hypoglycaemic). The search, carried out in November 2020, with no limit ranges for the year of publication and with the 'English' filter activated, yielded 620 results (Figure 1). Reviews, systematic reviews and meta-analyses ($n = 78$) were excluded. Articles were screened to exclude those not relevant to the topics such as allergenic properties, nutrient composition, technological functionality, safety assessment, rearing, non-edible insects or other

functional effects ($n = 492$). The references of the selected papers were screened, and two additional papers were identified. Furthermore, to include papers without edible insect as key word, we performed a search for the following insect names: *Bombyx mori*, *Tenebrio molitor*, cricket and grasshoppers, identifying three more papers. Fifty-five papers, focusing on oxidative and inflammatory stress, platelet aggregation, weight control, lipid and glucose metabolism, were identified.

Results are presented in Tables 1–6: *in vitro* studies (Tables 1–3), cellular models/*ex vivo* studies (Tables 4 and 5) and *in vivo* (Table 6) studies. They were further grouped by topic: antioxidant activity *in vitro* (Table 1), *in vitro* effects on platelet aggregation (Table 2), other *in vitro* effects (Table 3), antioxidant activity in cellular models or *ex vivo* (Table 4), and other effects in cellular models/*ex vivo* (Table 5). For the *in vivo* studies, results were divided into six categories: body and organ weight and composition, inflammatory status, oxidative/antioxidant status, lipid status, glycaemia/insulin status and coagulation markers.

Results

Antioxidant activity *in vitro*

Table 1 summarises the results of twenty-two studies^(6,14–34) on the *in vitro* antioxidant activity of different extracts from edible insects. The studies involved thirty-one species of insects, among which the most studied was *Tenebrio molitor*, tested in ten papers^(14,15,20–23,25,26,28,30). *Acheta domesticus* and *Gryllobates sigillatus* were cited respectively in four and^(14,25,26,30) three papers^(6,15,34), while *Alphitobius diaperinus*^(26,29), *Bombyx mori*^(26,31), *Hermetia illucens*^(16,17) and *Lebocerus indicus*^(26,27) were investigated in two papers. Water-soluble fractions were investigated in fourteen research articles^(6,19–22,24–32), while protein hydrolysates were reported in six papers^(14–17,23,34) and liposoluble fractions were tested in two papers only^(26,33). Among the several methods taken into account, the most used was the 1,1-diphenyl-2-picrylhydrazyl (DPPH) method, cited in sixteen studies^(6,14,15,18,20–22,24,25,27,28,30,32–35), while 2,2'-azo-bis(3-ethylbenzothiazoline-6-sulfonic acid (ABTS) and ferric reducing antioxidant power (FRAP) methods were performed in thirteen^(6,15,16,19–21,23,25,26,28,29,34,35) and nine papers^(17,21,22,25,26,28,31,32,34), respectively. Seven studies investigated the scavenging activity of selected fractions against different radicals^(16,17,22,24,28,29,32), three involving metal ion chelating activity^(15,25,34) and one utilising β -carotene and linolenic acid bleaching tests⁽³⁶⁾. All tested fractions showed significant antioxidant activity, the only exception being the protein hydrolysate from *Gryllobates sigillatus* in FRAP assay⁽³⁴⁾. Only the study conducted by Di Mattia *et al.*⁽²⁶⁾ provided a comparison between antioxidant activity of water and lipo-soluble fractions of edible insect and food extracts. More specifically, when water-soluble extracts of edible insects were analysed, *Calliptamus italicus*, *Bombyx mori* and *Acheta domesticus* crickets showed an antioxidant capacity, expressed as Trolox equivalent antioxidant capacity (TEAC), five-fold higher than fresh orange juice. Furthermore, water-soluble extracts of *Calliptamus italicus*, *Imbrasia oyemensis* and *Acheta domesticus* displayed a

Fig. 1. Flow diagram for search strategy.

reducing power (FRAP) double that of fresh orange juice. As regards the liposoluble fraction, *Bombyx mori*, *Tanna japonensis* and *Imbrasia oyemensis* showed a TEAC twice that of olive oil. Regarding endogenous antioxidants, Dutta *et al.* (2016) showed that an aqueous extract of *Vespa affinis* was able to increase the activity of catalase (CAT) and glutathione *S*-transferase (GST)⁽²⁴⁾.

Anti-platelet aggregation activity in vitro

Seven edible insects were tested for their effects on coagulation markers as described in Table 2. In the study by Pyo and colleagues (2020)⁽²⁸⁾, ethanol extract of the edible insect *Teleogryllus emma* was able to prolong thrombin time (TT) and activated partial thromboplastin time (aPTT), but not prothrombin time (PT); also relative PT, TT and aPTT were increased. In the same study, *Allomyrina dichotoma*, *Apis mellifera*, *Gryllus bimaculatus*, *Protaetia brevitarsis* and *Tenebrio molitor* did not exert the same effect. However, all above-mentioned insects, with the exception of *Apis mellifera*, were able to increase platelet aggregation; conversely, haemolytic activity was observed only in *Apis mellifera* and *Teleogryllus emma*. *N*-acetyl dopamine dimers obtained from *Oxya chinensis sinuosa* showed the ability of inhibit factor Xa (FXa) and to reduce platelet aggregation induced by adenosine diphosphate (ADP)

or U46619, but not by thrombin. However, they did not inhibit thrombin, trypsin, elastase, plasmin, streptokinase, tissue plasminogen activator (tPA) or urokinase⁽³⁷⁾. Indole alkaloids isolated from *Protaetia brevitarsis seulensis* affected coagulation by inhibiting platelet aggregation, myristoylated alanine-rich C-kinase substrate (MARCKS) phosphorylation and reduced Ca²⁺ peak in platelets. Moreover, they were able to prolong aPTT and PT and to inhibit thrombin and FXa production⁽³⁸⁾. Finally, two compounds (1, cyclo(L-Pro-L-Tyr) and 2, *N*-acetyltyramine) extracted from *Tenebrio molitor* prolonged aPTT and reduced platelet aggregation induced by ADP, collagen or U46619, but not by thrombin. Moreover, FXa production and activity were inhibited, but not thrombin, trypsin, plasmin, activated protein C or tPA. These compounds also inhibited MARCKS phosphorylation and expression of P-selectin and PAC-1, and reduced Ca²⁺ peak in platelets⁽³⁹⁾.

Other activities in vitro

Table 3 presents seven studies^(19,29,30,34,40–42) evaluating other *in vitro* effects for seven different insects. Proteins from *Bombyx mori*, *Tenebrio molitor* and *Gryllus bimaculatus*⁽⁴⁰⁾, *Alphitobius diaperinus*⁽²⁹⁾ and *Gryllobius sigillatus*^(34,41) were able to inhibit activity of angiotensin-converting enzyme (ACE). These proteins also showed potential anti-diabetic

Table 1. *In vitro* antioxidant activity of edible insects

Sample	Concentrations	Method	Antioxidant activity	References
<i>Acheta domesticus</i> , <i>Tenebrio molitor</i> – PH	0.05–5.0 mg/ml	DPPH	+	Messina <i>et al.</i> , 2019 ⁽¹⁴⁾
<i>Acheta domesticus</i> , <i>Rhynchophorus ferrugineus</i> , <i>Tenebrio molitor</i> , <i>Zophobas morio</i>	–	ABTS, DPPH, FRAP, Fe ²⁺ chelating activity	+	Botella-Martinez <i>et al.</i> , 2020 ⁽²⁵⁾
<i>Allomyrina dichotoma</i> , <i>Apis mellifera</i> , <i>Gryllus bimaculatus</i> , <i>Protaetia brevitarsis</i> , <i>Teleogryllus emma</i> , <i>Tenebrio molitor</i>	500 µg/ml 200 µg/ml	ABTS, DPPH, FRAP Nitric scavenging activity	+	Pyo <i>et al.</i> , 2020 ⁽²⁸⁾
<i>Alphitobius diaperinus</i> – PH	–	ABTS, ORAC	+	Sousa <i>et al.</i> , 2020 ⁽²⁹⁾
<i>Acheta domesticus</i> , <i>Tenebrio molitor</i>	10 mg/ml	DPPH	+	Navarro del Hierro <i>et al.</i> , 2020 ⁽³⁰⁾
<i>Bombyx mori</i>	10 g/50 ml	DPPH, ABTS, FRAP	+	Anuduang <i>et al.</i> , 2020 ⁽³¹⁾
<i>Brachytrupes orientalis</i>	0.25–6.25 mg/ml 1.25–12.5 mg/ml	DPPH FRAP, SAHR, SRSC	+	Dutta <i>et al.</i> , 2017 ⁽³²⁾
<i>Clanis bilineata</i>	10–200 µg/ml 0.1–4.0 mg/ml	DPPH β-carotene and linolenic acid bleaching test	+	Sun <i>et al.</i> , 2018 ⁽³³⁾
<i>Gryllosides sigillatus</i> – PH	1 mg/ml	ABTS, DPPH, metal ion chelating activity FRAP	+	Hall <i>et al.</i> , 2018 ⁽³⁴⁾
<i>Gryllosides sigillatus</i> , <i>Schistocerca gregaria</i> , <i>Tenebrio molitor</i> – PH	–	ABTS, DPPH, Fe ²⁺ chelating activity	+	Zielinska <i>et al.</i> , 2017 ⁽¹⁵⁾
<i>Gryllus bimaculatus</i>	–	ABTS DPPH	+	Hwang <i>et al.</i> , 2019 ⁽⁶⁾
<i>Hermetia illucens</i> – PH	14 g/l	SAHR	+	Mintah <i>et al.</i> , 2019a ⁽¹⁷⁾
<i>Hermetia illucens</i> – PH	2 mg/ml 4 mg/ml	ABTS FRAP, SRSC	+	Mintah <i>et al.</i> , 2019b ⁽¹⁶⁾
Honey-bee brood	0.0025 g/ml	DPPH	+	Haber <i>et al.</i> , 2019 ⁽¹⁸⁾
<i>Pachymerus nucleorum</i>	1 g/100 ml	ABTS	+	Alves <i>et al.</i> , 2016 ⁽¹⁹⁾
<i>Tenebrio molitor</i>	–	ABTS, DPPH	+	Son <i>et al.</i> , 2020 ⁽²⁰⁾
<i>Tenebrio molitor</i>	3 g/10 ml	ABTS, DPPH, FRAP	+	Mancini <i>et al.</i> , 2019 ⁽²¹⁾
<i>Tenebrio molitor</i>	0.625–5.0 mg/ml	DPPH, FRAP, ORAC, SAHR, hydrogen peroxide radical scavenging activity	+	Tang <i>et al.</i> , 2018 ⁽²²⁾
<i>Tenebrio molitor</i> , <i>Uromoides dermestoides</i> – PH	0.1–1.0 mg/ml	ABTS	+	Flores <i>et al.</i> , 2020 ⁽²³⁾
<i>Vespa affinis</i> L.	0.25–6.25 µg/µl 1.25–15.0 µg/µl 1.25–10.0 µg/µl	DPPH SAHR, SRSC Activities of CAT and GST enzymes	+	Dutta <i>et al.</i> , 2016 ⁽²⁴⁾
Various (<i>Acheta domesticus</i> , <i>Alphitobius diaperinus</i> , <i>Bombyx mori</i> , <i>Calliptamus italicus</i> , <i>Imbrasias oyemensis</i> , <i>Lasius niger</i> , <i>Lethocerus indicus</i> , <i>Rhynchophorus ferrugineus</i> , <i>Scolopendra</i> , <i>Tanna japonensis</i> , <i>Tenebrio molitor</i> , <i>Haplopelma albostratum</i> , <i>Pandinus imperator</i>)	–	ABTS hydro, ABTS lipo, FRAP	+	Di Mattia <i>et al.</i> , 2019 ⁽²⁶⁾
Various (<i>Crocothemis servilia</i> , <i>Cybister tripunctatus</i> , <i>Hydrophilus olivaceus</i> , <i>Laccotrephes maculatus</i> , <i>Lethocerus indicus</i>)	1–500 mg/ml	DPPH	+	Shantibala <i>et al.</i> , 2014 ⁽²⁷⁾

PH, protein hydrolysates; DPPH, 2,2-diphenyl-1-picrylhydrazyl; ABTS, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid); FRAP, ferric reducing antioxidant power; ORAC, oxygen radical absorbance capacity; SAHR, scavenging activity on hydroxyl radicals; RSC, superoxide radical scavenging capacity; CAT, catalase; GST, glutathione S-transferase.

Table 2. *In vitro* anti-platelet aggregation activity of edible insects

Sample	Concentrations	Activity	Result	References
<i>Allomyrina dichotoma</i> , <i>Apis mellifera</i> , <i>Gryllus bimaculatus</i> , <i>Protaetia brevitarsis</i> , <i>Tenebrio molitor</i> – WS	5.0 mg/ml	Prolonged PT, TT, aPTT	–	Pyo <i>et al.</i> , 2020 ⁽²⁸⁾
	0.25 mg/ml	Platelet aggregation	+	
	1.0 mg/ml	Haemolytic activity	– (A-M–)	
<i>Teleogryllus emma</i> – WS	5.0 mg/ml	Prolonged relative TT and aPTT	– (A-M+)	
	5.0 mg/ml	Prolonged PT	+	
	5.0, 6.0, 7.0 mg/ml	Relative PTT, TT, aPTT	–	
	0.25 mg/ml	Platelet aggregation	+	
	1.0 mg/ml	Haemolytic activity	+	
<i>Oxya chinensis sinuosa</i> – 5 CPs (<i>N</i> -acetyldopamine dimers)	–	Inhibition of thrombin, trypsin, elastase, plasmin, protein Ca, streptokinase, tPA and urokinase	–	Lee <i>et al.</i> , 2017b ⁽³⁷⁾
	1, 2, 5 µM	Reduction of platelet aggregation induced by ADP or U46619	+ (5–)	
	1,2,5 µM	Reduction of platelet aggregation induced by thrombin	–	
	2, 5 µM (c1–4)	Inhibition of FXa	+	
	5–50 µM	Inhibition of platelet aggregation induced by U46619 or collagen	+	
<i>Protaetia brevitarsis seoulensis</i> – 2 CPs (Indole alkaloids)	50 µM	Inhibition of ADP- and U46619-induced MARCKS phosphorylation (platelets)	+	Lee <i>et al.</i> , 2017a ⁽³⁸⁾
	10, 25, 50 µM	Reduction of Ca ²⁺ peak (platelets)	+	
	5–50 µM	Prolonged aPTT and PT	+	
	Various	Inhibition of thrombin and FXa production	+	
	1, 5, 10 µM	Prolonged aPTT	+	
<i>Tenebrio molitor</i> – 2 CPs (1, cyclo(L-Pro-L-Tyr) and 2, <i>N</i> -acetyltyramine)	1, 5, 10 µM	Reduction of platelet aggregation induced by ADP, collagen or U46619	+	Lee <i>et al.</i> , 2017c ⁽³⁹⁾
	1, 5, 10 µM	Reduction of platelet aggregation induced by thrombin	–	
	1–10 µM	Inhibition of FXa production	+	
	5, 10 µM	Inhibition of FXa activity	+	
	–	Inhibition of thrombin, trypsin, plasmin, activated protein C, and tPA	–	
	10 µM	Inhibition of ADP- and U46619-induced MARCKS phosphorylation (platelets)	+	
	1, 5, 10 µM	Reduction of Ca ²⁺ peak (platelets)	+	
	5, 10 µM	Reduction of ADP- or U46619-induced expression of P-selectin and PAC-1 (platelets)	+	

WS, water-soluble extract; PT, prothrombin time; TT, thrombin time; aPTT, activated partial thromboplastin time; CP, compound; tPA, tissue plasminogen activator; ADP, adenosine diphosphate; FXa, factor Xa; MARCKS, myristoylated alanine-rich C-kinase substrate.

Table 3. Other *in vitro* activity of edible insects

Sample	Concentrations	Activity	Result	References
<i>Acheta domestica</i> , <i>Tenebrio molitor</i> – WS	–	Pancreatic lipase inhibition	+	Navarro del Hierro <i>et al.</i> , 2020 ⁽³⁰⁾
<i>Alphitobius diaperinus</i> – PH	–	ACE inhibition	+	Sousa <i>et al.</i> , 2020 ⁽²⁹⁾
<i>Bombyx mori</i> , <i>Gryllus bimaculatus</i> , <i>Tenebrio molitor</i> – PH	–	α -glucosidase inhibition	+	Yoon <i>et al.</i> , 2019 ⁽⁴⁰⁾
<i>Gryllodes sigillatus</i> – PH	1–25 mg/ml	ACE inhibition	+	Hall <i>et al.</i> , 2018 ⁽³⁴⁾
<i>Gryllodes sigillatus</i> – PH	–	DPP-IV inhibition	+	Hall <i>et al.</i> , 2020 ⁽⁴¹⁾
<i>Gryllodes sigillatus</i> – PH	–	ACE inhibition	+	Alves <i>et al.</i> , 2016 ⁽¹⁹⁾
<i>Pachymerus nucleorurn</i> – LS	–	α -amylase inhibition	+	Alves <i>et al.</i> , 2016 ⁽¹⁹⁾
<i>Tenebrio molitor</i>	1 % (w/v)	α -glucosidase inhibition	+	De Carvahlo <i>et al.</i> , 2019 ⁽⁴²⁾
		ACE inhibition	+	
		Tryptic activity	+	
		Anti-tryptic and anti-chymotryptic activities	+	
		<i>Lactobacillus</i> , <i>Bifidobacterium</i> : viability, SCFA production, viability in nutritive stress conditions	+	

WS, water-soluble extract; PH, protein hydrolysates; ACE, angiotensin converting enzyme; DPP-IV, dipeptidyl peptidase-4; LS, lipo-soluble extract; SCFA, short-chain fatty acids.

activity by inhibition of dipeptidyl peptidase-4 (DPP-IV)⁽³⁴⁾ and α -glucosidase^(40,41). Moreover, extracts from *Acheta domestica* and *Tenebrio molitor* were able to inhibit pancreatic lipase⁽³⁰⁾, while *Pachymerus nucleorurn* larvae showed tryptic activity together with absence of anti-nutritional factors as well as anti-tryptic and chymotryptic activity⁽¹⁹⁾. Lastly, flour from *Tenebrio molitor* affected the growth of *Lactobacillus* and *Bifidobacterium*, improving short-chain fatty acid (SCFA) production and viability in nutritive stress conditions⁽⁴²⁾.

Antioxidant activity in cellular models

Nine studies^(6,20,24,32,39,40,43–45) investigating the antioxidant activity in cellular model of ten species of insects are described in Table 4. *Gryllus bimaculatus* and *Tenebrio molitor* were the most utilised insects, examined in three studies. Different fractions of the insects, including water-soluble fraction ($n = 6$), compounds ($n = 2$), protein hydrolysates ($n = 1$) and lipo-soluble fraction ($n = 1$), were tested. In three different studies, aqueous extracts of *Gryllus bimaculatus*⁽⁶⁾, methanolic extract of defatted powder and unsaponifiable lipids, obtained by *Tenebrio molitor*⁽²⁰⁾, and *Bombyx mori* protein hydrolysates showed the ability to reduce lipopolysaccharide-induced nitric oxide (NO) production in the murine macrophage cell line RAW 264-7. A similar effect was exerted by 1, cyclo(L-Pro-L-Tyr) and 2, *N*-acetyltyramine isolated from *Tenebrio molitor* in human umbilical vein endothelial cells (HUVECs)⁽³⁹⁾ and by glycosaminoglycan from *Gryllus bimaculatus* in diabetic type 2 microvascular endothelial cells (D-HMVECs)⁽⁴⁵⁾. According to a study by Yoon and co-workers⁽⁴⁰⁾, protein hydrolysates of *Tenebrio molitor* and *Gryllus bimaculatus* did not exert any effect on NO release. Water-soluble extract of dung beetles of *Onitis* sp., mole crickets of *Gryllotalpa* sp., grasshopper of *Caelifera* sp.⁽⁴³⁾, *Oryctes boas* and *Zonocerus variegatus*⁽⁴⁴⁾ were tested in human peripheral blood lymphocytes to evaluate whether they affected oxidative status. Results showed that at lower concentrations (10–40 ppm) the insects display an antioxidant effect; however, at higher concentrations (2000 ppm) they exhibited a pro-oxidant effect. According to the results obtained in a cell-free system⁽²⁴⁾, the aqueous extract of *Vespa affinis* was able to increase the activity of both GST and CAT also in THP-1 human monocytes and human plasma; moreover, it reduced reactive oxygen species (ROS) formation in THP-1. Finally, the hydro-alcoholic extract of *Brachytrupes orientalis* was able to restore nuclear factor erythroid 2-related factor (Nrf2) and GST protein expression, reducing radical and malondialdehyde (MDA) levels in C2C12, a murine myotube cell line, following high glucose stress⁽³²⁾.

Other activity in cellular models or ex vivo

The effect of edible insects on coagulation markers^(37–39), inflammatory status^(41,46) and lipid metabolism^(7,47–49) are described in Table 5. A total of eleven studies involving eight insects were reported: *Protaetia brevitarsis* was cited in three research articles, whilst *Oxya chinensis*, *Gryllus bimaculatus* and *Tenebrio molitor* were investigated in two studies each. Compounds (indole alkaloids) isolated from *Protaetia brevitarsis seulensis* were able to reduce prothrombin-produced thrombin, plasminogen

Table 4. Antioxidant activity of edible insects in cellular models or *ex vivo*

Sample	Cell type	Concentration	Antioxidant/Oxidant Marker	References
<i>Bombyx mori</i> – PH	RAW264-7	0.1, 0.3, 0.5 mg/ml	NO ↓	Yoon <i>et al.</i> , 2019 ⁽⁴⁰⁾
<i>Gryllus bimaculatus</i> – PH			NO ↔	
<i>Tenebrio molitor</i> – PH			NO ↔	
<i>Brachytrupes orientalis</i> – WS	C2C12	7.5, 10*, 12.5* mg/ml	Lipid peroxidation: MDA ↓*; ROS ↓*; GST, Nrf-2 ↑*	Dutta <i>et al.</i> , 2017 ⁽³²⁾
<i>Caelifera</i> sp. – WS	hPBL	5–2000 ppm	TOS ↔ (↑ 2000 ppm); TAC ↔ (↓ 2000 ppm)	Koc <i>et al.</i> , 2014 ⁽⁴³⁾
<i>Gryllotalpa</i> sp. – WS			TOS ↔ (↑ 2000 ppm); TAC ↔ (↑ 10 ppm, ↓ 1000, 2000 ppm)	
<i>Onitis</i> sp. – WS			TOS ↔ (↑ 2000 ppm) TAC ↔ (↑ 15 ppm, ↓ 1000, 2000 ppm)	
<i>Gryllus bimaculatus</i> – CP (glycosaminoglycan)	D-HMVECs	5, 10 mg/ml	NO ↓	Ahn <i>et al.</i> , 2020 ⁽⁴⁵⁾
<i>Gryllus bimaculatus</i> – WS	RAW264-7	20–100 µg/ml	NO ↓	Hwang <i>et al.</i> , 2019 ⁽⁶⁾
<i>Oryctes boas</i> – WS	hPBL	5–2000 ppm	TAC ↑ (10–40 ppm), ↓ (2000 ppm) TOS ↑ (1000, 2000 ppm)	Memis <i>et al.</i> , 2013 ⁽⁴⁴⁾
<i>Zonocerus variegatus</i> – WS			TAC ↑ (10–25 ppm), ↓ (500–2000 ppm) TOS ↑ (200–2000 ppm)	
<i>Tenebrio molitor</i> – 2 CPs	HUVECs	5, 10 µM	NO ↓	Lee <i>et al.</i> , 2017c ⁽³⁹⁾
<i>Tenebrio molitor</i> – WS, LS	RAW264-7	WS: 25–500 µg/ml LS: 0.05–5.0 µg/ml	NO ↓	Son <i>et al.</i> , 2020 ⁽²⁰⁾
<i>Vespa affinis</i> – WS	THP-1	0.4, 0.8*, 1.2* µg/µl	GST, CAT ↑*	Dutta <i>et al.</i> , 2016 ⁽²⁴⁾
	THP-1	0.8 µg/µl	ROS ↓	
	hPlasma	1.25–10.00 µg/µl	GST ↑ (except for 1.25 and 2.50 µg/µl) CAT ↑ (except for 1.25 µg/µl)	

PH, protein hydrolysates; NO, nitric oxide; WS, water-soluble extract; MDA, malondialdehyde; ROS, reactive oxygen species; GST, glutathione *S*-transferase; Nrf2, nuclear factor erythroid 2-related factor; TOS, total oxidant status; TAC, total antioxidant capacity; CP, compound; D-HMVECs, diabetic type 2 microvascular endothelial cells; hPBL, human peripheral blood lymphocytes; HUVECs, human umbilical vein endothelial cells; LS, lipo-soluble extract; CAT, catalase.

Table 5. Other activity in cellular models or *ex vivo*

Source	Cells	Dose	Results	References
<i>Allomyrina dichotoma</i> – CPs (Tetrahydroquinolines)	HUVECs	1, 2, 5*, 10* μ M	Cell viability $\leftrightarrow$; permeability $\downarrow$ *, **, VCAM-1*, **, ICAM-1** $\downarrow$; adherence of monocytes to HUVEC monolayers $\downarrow$ *, **, migration of human neutrophils through HUVEC monolayers $\downarrow$ *, **	Park <i>et al.</i> , 2020 ⁽⁴⁶⁾
<i>Bombyx mori</i> – PUFAs	HUVECs L-02	10 μ M 100–1600 μ g/ml	NF- κ B p65 activity; TNF- α and IL-1 β production, expression of phospho-p38 $\downarrow$ Cell viability $\leftrightarrow$ ($\downarrow$ 1600 μ g/ml)	Luo <i>et al.</i> , 2020 ⁽⁷⁾
<i>Bombyx mori</i> – α -linolenic acid		200, 400*, 800** μ g/ml 2.5–40* μ g/ml	Lipid accumulation, TC $\downarrow$ *, **, TBA $\uparrow$; mRNA and protein expression (cholesterol metabolism related): LXR α **, PPAR γ *, **, ABCA1 *, **, ABCG1, CYP7A1 *, ** $\uparrow$ Cell viability: $\leftrightarrow$ ($\downarrow$ 40 μ g/ml)	
<i>Gryllos sigillatus</i> – PH	RAW 264.7	5, 10*, 20** μ g/ml 0.5–3.0 μ g/ml	Lipid accumulation, TC $\downarrow$ *, **, TBA $\uparrow$; mRNA and protein expression (chol-metabolism related): LXR α , PPAR γ , ABCA1, ABCG1, CYP7A1 $\uparrow$ ** Viability $\leftrightarrow$; NF- κ B $\downarrow$	Hall <i>et al.</i> , 2020 ⁽⁴¹⁾
<i>Gryllus bimaculatus</i> – CP (glycosaminoglycan)	D- HMVECs	5, 10* mg/ml	Laminin $\leftrightarrow$; VEGF $\uparrow$ *	Ahn <i>et al.</i> , 2020 ⁽⁴⁵⁾
<i>Gryllus bimaculatus</i> , <i>Oxya chinensis sinuosa</i> , <i>Protaetia brevitarsis seulensis</i> – WS	HepG2	10–200 μ g/ml 10, 50, 100 μ g/ml	Cell viability $\leftrightarrow$ Intracellular lipid accumulation, TG $\downarrow$	Im <i>et al.</i> , 2018 ⁽⁴⁷⁾
<i>Locusta migratoria</i> – WS	hPBL	1–1000 mg/l	MN/1000 cells, SCEs, CAs $\leftrightarrow$	Tukez <i>et al.</i> , 2014 ⁽⁵⁰⁾
<i>Oxya chinensis sinuosa</i> – 5 CPs (<i>N</i> -acetyldopamine dimers)	Mice plasma	1, 2, 5 μ M	aPTT $\uparrow$; PT $\leftrightarrow$	Lee <i>et al.</i> , 2017b ⁽³⁷⁾
<i>Oxya chinensis sinuosa</i> – 1–4 CPs (<i>N</i> -acetyldopamine dimers)	HUVECs	0.5–5.0 μ M	FXa $\downarrow$ (except for 0.5 μ M)	
<i>Protaetia brevitarsis seulensis</i> – 5 CPs (results are referred to CPs 1 and 2) (Indole alkaloids)	HUVECs	5–50 μ M 0.5–5.0 μ M	Prothrombin-produced thrombin, PAI 1 $\downarrow$ (except for cp5) tPA $\leftrightarrow$ FXa $\downarrow$ (except for 0.5 μ M)	Lee <i>et al.</i> , 2017a ⁽³⁸⁾
	Mice platelet-rich plasma	10, 25, 50 μ M	Aggregation % $\downarrow$; aPTT, PT $\uparrow$	
<i>Protaetia brevitarsis</i> – WS	3T3-L1	0.1, 0.5*, 1.0* mg/ml	Adipogenesis-related genes (mRNA expression): C/EBP α , aP2, FAS $\downarrow$ *	Ahn <i>et al.</i> , 2019 ⁽⁴⁹⁾
<i>Tenebrio molitor</i> – 2 CPs (1, cyclo(L-Pro-L-Tyr) and 2, <i>N</i> -acetyltyramine)	Mice blood	1, 5, 10 μ M	aPTT, clotting time $\uparrow$	Lee <i>et al.</i> , 2017c ⁽³⁹⁾
<i>Tenebrio molitor</i>	HUVECs	5, 10 μ M	ET-1 $\uparrow$	
	3T3-L1	0.1, 1*, 2*, 3* mg/ml	Lipid droplet formation, TG $\downarrow$ *; adipogenic differentiation (mRNA expression): PPAR γ , C/EBP- α $\downarrow$ *; lipogenesis-specific genes mRNA expression: SREBP-1c, LPL, SCD1, FAS $\downarrow$ *(mRNA); p-ERK $\downarrow$, p-AMPK- α , p-p38 $\uparrow$, p-JNK $\leftrightarrow$	Seo <i>et al.</i> , 2017 ⁽⁴⁸⁾

HUVECs, human umbilical vein endothelial cells; VCAM-1, vascular cell adhesion molecule-1; ICAM-1, intercellular adhesion molecule-1; NF- κ B, nuclear factor- κ B; TNF- α , tumour necrosis factor- α ; IL-1 β , interleukin-1 β ; PUFA, polyunsaturated fatty acids; TC, total cholesterol; TBA, total bile acid; LXR α , liver X receptor; PPAR γ , peroxisome proliferator-activated receptor γ ; ABCA1, ATP-binding cassette transporter A1; ABCG1, ATP-binding cassette transporter G1; CYP7A1, cholesterol 7 α hydroxylase; PH, protein hydrolysates; D-HMVECs, diabetic type 2 microvascular endothelial cells; VEGF, vascular endothelial growth factor; WS, water-soluble extract; TG, triacylglycerol; MN, micronucleus; SCE, sister chromatid exchange; CA, chromosome aberration; aPTT, activated partial thromboplastin time; PT, prothrombin time; FXa, factor Xa; PAI-1, plasminogen activator inhibitor-1; tPA, tissue plasminogen activator; C/EBP α , CCAAT/enhancer-binding protein α ; FAS, fatty acid synthase; ET-1, endothelin-1; SREBP-1c, sterol regulatory element-binding protein 1c; LPL, lipoprotein lipase; SCD1, stearoyl-CoA desaturase-1; ERK, extracellular signal-regulated kinases; AMPK- α , adenosine monophosphate-activated protein kinase- α ; JNK, c-Jun *N*-terminal kinase.

Table 6. Effect of edible insects on body weight and composition, inflammation, redox, lipid and glycaemia/insulin status and coagulation markers in animal and human studies

Animal studies				Results						References
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	
<i>Acheta domesticus</i>	Malnourished weanling Sprague-Dawley rats	283.0 g/kg	14 d	Body weight gain, organs weight ↑ Relative biological value ↑ Bone mineral content, lean mass, ↑ Fat mass ↑						Agbemafle <i>et al.</i> , 2019 ⁽⁶¹⁾
	Malnourished weanling Sprague-Dawley rats	154.3 g/kg	14 d	Body weight gain, organ weight ↑ Relative biological value ↔ Bone mineral content, lean mass ↑ Fat mass ↔						
<i>Rhynchophorus phoenicis fabricius</i>	Malnourished weanling Sprague-Dawley rats	403.0 g/kg	14 d	Body weight gain, organ weight ↑ Relative biological value ↑ Bone mineral content, lean mass ↑ Fat mass ↔						Park <i>et al.</i> , 2020 ⁽⁴⁶⁾
<i>Allomyrina dichotoma</i> – 4 CPs (results are referred to as CPs 1, 2 and 3) (Tetrahydroquinolines)	C57BL/6 mice, LPS inflammation	0.13, 0.26* μM per mouse	Acute		Peritoneum: leucocyte migration, vascular permeability ↓ *					
	CLP-induced septic C57BL/6 mice	0.26 μM per mouse	Acute	Survival rate ↑	Lungs: interstitial oedema, tissue damage ↔					Ryu, 2014 ⁽⁵⁶⁾
<i>Bombyx mori</i>	Sprague-Dawley rats, healthy	200 g/kg	5 weeks	Body weight ↔			TG, TC ↓; HDL ↔			
<i>Bombyx mori</i> – LS	Wistar rats, hypercholesterolaemia	1*, 2**, 4*** mL/kg/d	6 weeks	Organ size: liver*** ↓; kidney ↓; heart* ↓; spleen** ↑		Serum: TAC**, SOD **, *** GPx **, ** ↑; MDA ↓ Liver: TAC, SOD ↑; GPx ↔; MDA ↓	TC, LDL, LDL/HDL ↓; TG, HDL ↔			Zou <i>et al.</i> , 2017 ⁽⁶⁴⁾

Table 6. (Continued)

Animal studies				Results						
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	References
<i>Clanis bilineata</i> – CPs (chitooligosaccharides)	Sprague-Dawley rats, high-fat diet	5 % (w/w)	6 weeks	Body weight gain, food efficiency ratio ↓; food intake ↔			TG, TC, LDL ↓; HDL ↑ Faecal content: TG, TC ↑			Xia <i>et al.</i> , 2013 ⁽⁶³⁾
<i>Caelifera</i> sp.	Wistar rats, healthy	10, 20 g/kg 1*, 3** %	Acute 28/90 d	Body weight ↔; toxicity signs: no; mortality: no Organs and tissues weight ↔			TG, TC, HDL ↔	Insulin ↓ (**, 90 d); plasma glucose ↑*, ↓** (28 d)		Ochiai <i>et al.</i> , 2020 ⁽⁵⁷⁾
Grain larvae – WS	Sprague-Dawley rats, healthy	5.0, 7.0, 9.0 mg	4 weeks	Body weight gain ↔; weight: liver, kidney ↔; thymus, spleen ↑; abdominal fat ↓	IgG, IgA, IgM ↓		SCFA ↓*, pH ↔ TG, TC, LDL ↓	Blood glucose ↓		Park & Park 2015 ⁽⁵⁸⁾
Green cocoon shell of <i>Bombyx mori</i> – WS	ICR mice, type 2 diabetes	150, 250*, 350** mg/kg	7 weeks	Body weight ↔	Liver: NF-κB, IL6, TNF-α ↓*, ** Liver: inflammatory infiltrations, oedema ↓	Liver: GPx, SOD ↑; MDA, 8-OHdG ↓*, **	TC, HDL ↔; TG** ↓; LDL*, ** ↓ Liver: lipid droplet accumulation ↓	OGTT: blood glucose reduction rate ↑, AUC ↓ ITT: blood glucose ↓, AUC ↓ Fasting blood glucose ↓; fasting blood insulin ↓; HbA1c*, **, HOMA-IR ↓; ISI ↑		Zhao <i>et al.</i> , 2019 ⁽⁸⁾

Table 6. (Continued)

Animal studies										
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Results				References
						Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	
<i>Gryllobates sigillatus</i>	Mice, malnourished	–	6 weeks	Body weight ↑ (versus malnourished)	TLR4, TNFα		TG ↓; leptin, adiponectin ↔ (versus control)	Pancreas: islet structure and area, pancreatic β cells, insulin secreted by β cells ↑		Bergmans <i>et al.</i> , 2020 ⁽⁶²⁾
<i>Gryllus bimaculatus</i>	Sprague-Dawley rats, varicocele	1.63, 6.5 mg/kg	42 d	Body and organ weights ↔	IL-1β, IFNγ, IL-4 in spleen tissue ↔ (versus control)	TNF-α, IL-6 ↓	Testicular tissues			Karna <i>et al.</i> , 2020 ⁽⁵²⁾
<i>Gryllus bimaculatus</i> – WS	Wistar rats, obesity	100*, 200** mg/kg	2 months	Body weight ↔; fat weight: abdominal*, total ↓, epididymal ↔; death/toxicity: no	IL-10* ↓	MDA, ROS/RNS ↓ SOD, GPx, CAT ↑ Blood protein carbonyl content ↓** (2m) CAT ↔ Liver: MDA ↓ Serum: uric acid ↔ ↓ (1m), ↑ (2m)	TG ↓ (100 mg/kg-1m); TC, HDL, LDL ↔; abdominal fat composition: polyunsaturated ↑, monounsaturated ↓	Glucose ↓ (1m*), ↔ (2m);		Ahn <i>et al.</i> , 2015 ⁽⁵¹⁾
<i>Gryllus bimaculatus</i> – WS	C57BL/6J mice, alcoholic liver damage	200 mg/kg	2 weeks		F4/80+ KCs, IL-1β ↓	Liver: 8-OHdG, MDA ↓ Small intestine: 8-OHdG ↓	Liver: lipid droplet accumulation ↓			Hwang <i>et al.</i> , 2019 ⁽⁶⁾
<i>Gryllus bimaculatus</i> – CPs (glycosaminoglycan)	BKS.Cg-m+/+Leprdb mice, diabetes	5 mg/kg	1 month	Body weight ↔, abdominal fat ↔		Carbonyl content: blood ↓, liver ↔, GST ↔;	Adipocyte density in lungs, liver and kidneys ↓ pancreas ↔	Blood glucose level ↔; non-fasting blood glucose ↔ (1 week ↓)		Ahn <i>et al.</i> , 2020 ⁽⁴⁵⁾

Table 6. (Continued)

Animal studies				Results						References
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	
						CAT, GPx ↑		pancreatic islet status, liver tissue status ↑		
<i>Gryllus bimaculatus</i> – WS	C57BL/6Jmice, non-alcoholic fatty liver disease	100*, 200** mg/kg/d	14 weeks	Body weight gain, liver weight ↓, adipose tissue: abdominal, kidney ↔; intestinal, epididymal ↓	Liver and adipose tissue: TNF-α, IL-1β, IL-6↓ (mRNA); TNF-α, IL-1β ↓ (expression) adiponectin↑↔#		TG*, TC*, LDL, HDL*, NEFA↓ Liver: lipid droplet accumulation ↓			Im <i>et al.</i> , 2018 ⁽⁴⁷⁾
<i>Oxya chinensis sinuosa</i> – WS							TG*, TC**, LDL, HDL**, NEFA↓ Liver: lipid droplet accumulation ↓	Fasting blood glucose** ↓		
<i>Protaetia brevitarsis seulensis</i> – WS							TG ↔; TC ↓*; LDL, HDL ↔; NEFA ↓ Liver: lipid droplet accumulation ↓	Fasting blood glucose ↓		
<i>Hermetia illucens</i>	Zebrafish, healthy	25, 50, 75*, 100** %	57 d	Specific growth rate ↑*,**; survival ↔	Intestine: no inflammatory events (histological analysis) mRNA expression (intestine): IL-1β, IL-10, TNF-α ↑*,**		Liver: steatosis, fat (%) ↑*,** SFA, n-6 ↑; MUFA, PUFA, n-3, n-9 ↓ Lipid metabolism-related genes: elovl5 ↑*,**; elovl2, fads2 ↔			Zarantonello <i>et al.</i> , 2020 ⁽⁵⁹⁾
<i>Oxya chinensis sinuosa</i> – 4 CPs (N-acetyldopamine dimers)	C57BL/6 mice, healthy	1, 2, 5 μM	4 d						Blood clotting time ↑	Lee <i>et al.</i> , 2017b ⁽³⁷⁾
	C57BL/6 mice, healthy	10, 20, 30 μM	Acute						Tail bleeding time ↔	
	C57BL/6 mice, arterial thrombosis	1, 2*, 5** μM	Acute						Thrombi formation time ↑ thrombi size ↓ (c1; c2: *,**; c3: *,**; c4: **)	
	C57BL/6 mice, pulmonary thrombosis	1, 2, 5 μM	Acute						Mortality ↓	
<i>Oxya chinensis sinuosa</i> – WS	Hos/HR-1 mice, UV irradiation	100 mg / kg/d	12 weeks		Skin: IL1-β, IL-6, TNF-α ↓	Skin: SOD, CAT ↑				Im <i>et al.</i> , 2019 ⁽⁵⁵⁾
<i>Protaetia brevitarsis</i> – WS	C57BL/6 mice, obesity	100, 200* mg/kg/d	7 weeks	Body weight gain, organ weight: liver ↓, spleen ↔; fat weight:		Liver: GPx ↑; CAT ↑*	TG, TC, LDL ↓			Ahn <i>et al.</i> , 2019 ⁽⁴⁹⁾

Table 6. (Continued)

Animal studies				Results						References
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	
				epididymal, sub-cutaneous ↓						
<i>Samia ricinii</i> – LS	Weanling Wistar NIN rats, healthy	10 % (w/w)	18 weeks	Body weight gain, nutrient retention, organ weight ↔. Toxicity: no			Liver: lipid droplet accumulation ↓ TG, TC ↓, LDL ↑			Longvah <i>et al.</i> , 2012 ⁽³⁶⁾
<i>Protaetia brevitarsis seulensis</i> – 5 CPs (indole alkaloids) (results are referred to CPs 1 and 2)	C57BL/6 mice, arterial thrombosis	10, 25, 50 µM	Acute				Liver: TG ↔		Arterial thrombi formation rate and size ↓ Mortality ↓	Lee <i>et al.</i> , 2017a ⁽³⁸⁾
	C57BL/6 mice, pulmonary thrombosis model	10, 25, 50 µM	Acute							
	C57BL/6 mice, healthy	c1: 7-5, 14-9 µg/mouse; c2: 11-5, 23-0 µg/mouse	Acute						Tail bleeding time ↑	
<i>Tenebrio molitor</i> – 2 CPs (1, cyclo(L-Pro-L-Tyr) and 2, <i>N</i> -acetyltyramine)	C57BL/6 mice, arterial thrombosis model	1, 5, 10 µM	Acute						Thrombi formation time ↑ thrombi size ↓ (↔ c2: 1 µM)	Lee <i>et al.</i> , 2017c ⁽³⁹⁾
	C57BL/6 mice, pulmonary thrombosis model	1, 5, 10 µM	Acute						Mortality ↓; thrombi size ↓ (↔ c2: 1 µM)	
<i>Tenebrio molitor</i>	BALB/c mice, obesity	100, 3000* mg/kg/d	6 weeks	Body weight gain; fat weight: epididymal cell volume*, abdominal-to-peripheral adipose tissue volume; epididymal white adipose tissue volume ↓			Liver: lipid droplet accumulation ↓			Seo <i>et al.</i> , 2017 ⁽⁴⁸⁾

Table 6. (Continued)

Animal studies				Results						References
Sample	Animal/disease	Dose	Duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	
<i>Tenebrio molitor</i> – WS (fermented)	Sprague-Dawley rats, alcoholic liver disease	50*, 100**, 200*** mg/kg/d	8 weeks	Liver weight ↓*,**	Serum: TNF-α ↓; IL-6 ↔; Liver: p-IkB-a, TNF-α, ** ↓, IL-6 ↔	SOD, CAT ↔; GPx*, GR, Tot GPx, reduced GPx ↑	Liver: lipid accumulation, focal necrosis fibrosis ↓ Liver: TG, TC, NEFA ↓*** Hepatic genes: lipid synthesis transcription factors; ChREBP**, ***, SREBP-1c**, ***, SREBP-2 ↓; fatty acid uptake and transport-related genes: CD36, FATP5, FABP1 ↔ Triacylglycerol synthesis-related: GPAT1*, ***, GPAT4 **, AGPAT1, PAP1**, DGAT1**, ADRP ↓ Cholesterol synthesis and esterification-related: HMGCR, ACAT2 ↓***			Choi <i>et al.</i> , 2020 ⁽⁵³⁾
<i>Tenebrio molitor</i> – WS (fermented)	Sprague-Dawley rats, alcoholic liver disease	50, 100, 200 mg/kg/d	8 weeks			Liver: β-oxidation ↑				Choi <i>et al.</i> , 2020 ⁽⁵⁴⁾
<i>Zophobas atratus</i> – WS	SPF Sprague-Dawley rats, healthy	1250, 2500* and 5000** mg/kg	2 weeks 13 weeks	Body weight, organs weight ↔. Toxicity signs: no; mortality: no.	IgE: ↔					Kim <i>et al.</i> , 2020 ⁽⁶⁰⁾

Table 6. (Continued)

Human studies										
Edible insects-extract	Animal/disease	Dose	Treatment duration	Body and organ weight and composition	Inflammatory status	Antioxidant/oxidant markers	Lipid status	Glycaemia/insulin status	Coagulation markers	References
<i>Bombyx mori</i>	Humans, healthy	Noodles: wheat flour 93 g + 0.83 g of <i>Bombyx mori</i> powder	Acute					Post-prandial blood glucose, glucose peak, IAUC glucose, GI value ↓		Suk <i>et al.</i> , 2016 ⁽⁶⁵⁾
<i>Cricket</i>	Humans, healthy	25 g/d, in a muffin and a smoothie	14 d	Gastro-intestinal functionality ↔	slgA ↔, TNF-α ↓, GM-CSF, IFNα, IL-1α, IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-10, IL-12 (p70), IL-13↔		Faecal content: butyrate ↔, acetate, propionate ↓; bile acids, TG ↓	Glucose ↔		Stull <i>et al.</i> , 2018 ⁽⁶⁶⁾

CP, compound; LPS, lipopolysaccharide; CLP, caecal ligation and puncture; TG, triacylglycerol; TC, total cholesterol; HDL, high-density lipoprotein; LS, lipo-soluble extract; TAC, total antioxidant capacity; SOD, superoxide dismutase; GPx, glutathione peroxidase; MDA, malondialdehyde; LDL, low-density lipoprotein; WS, water-soluble extract; SCFA, short-chain fatty acids; IgA, immunoglobulin A; IgG, immunoglobulin G; IgM, immunoglobulin M; NF-κB, nuclear factor-κB; IL-6, interleukin-6; TNF-α, tumour necrosis factor-α; 8-OHdG, 8-hydroxy-2'-deoxyguanosine; OGTT, oral glucose tolerance test; AUC, area under curve; ITT, insulin tolerance test; HbA1c, glycosylated haemoglobin concentration; HOMA-IR, homeostasis model assessment of insulin resistance; ISI, insulin sensitivity index; TLR4, toll-like receptor 4; IL-1β, interleukin-1β; IFN-γ, interferon gamma; IL-4, interleukin-4; ROS, reactive oxygen species; RNS, reactive nitrogen species; GST, glutathione S-transferase; CAT, catalase; IL-10, interleukin-10; F4/80+ KCs, F4/80-positive Kupffer cells; SFA, saturated fatty acids; MUFA, monounsaturated fatty acids; PUFA, polyunsaturated fatty acids; ELOVL5, fatty acid elongase 5; ELOVL2, fatty acid elongase 2; FADS2, fatty acid desaturase 2; p-IkB-α, phosphorylated-inhibitor of nuclear factor-kappa B-alpha; GR, glutathione reductase; NEFA, non-esterified fatty acid; ChREBP, carbohydrate-response element-binding protein; SREBP-1c, sterol regulatory element-binding protein 1c; SREBP-2, sterol regulatory element-binding protein 2; CD36, cluster of differentiation 36; FATP5, fatty acid transport protein 5; FABP1, fatty acid-binding protein 1; GPAT1, glycerol-3-phosphate acyltransferase 1; GPAT4, glycerol-3-phosphate acyltransferase 4; AGPAT1, 1-acylglycerol-3-phosphate O-acyltransferase 1; PAP1, phosphatidate phosphatase 1; DGAT1, diacylglycerol O-acyltransferase 1; ADRP, adipose differentiation-related protein; HMGCR, 3-hydroxy-3-methylglutaryl-CoA reductase; ACAT2, acetyl-CoA acetyltransferase 2; SPF, specific pathogen-free; IgE, immunoglobulin E; GI, glucose index; slgA, secretory immunoglobulin A; GM-CSF, granulocyte-macrophage colony-stimulating factor; IFNα, interferon alpha; IL-1α, interleukin-1 α; IL-2, interleukin-2; IL-5, interleukin-5; IL-7, interleukin-7; IL-8, interleukin-8; IL-12, interleukin-12; IL-13, interleukin-13.

activator inhibitor (PAI-1) and FXa, but not tPA, in HUVECs. Moreover, in platelet-rich plasma (*ex vivo*) from mouse, they were able to reduce aggregation percentage and increase aPTT and PT⁽³⁸⁾. Compounds (*N*-acetyl dopamine dimers) extracted from *Oxya chinensis sinuosa* were able to increase aPTT but did not affect PT in mice plasma (*ex vivo*); in the same study, they also showed the ability to reduce FXa in HUVECs⁽³⁷⁾. Lastly, two compounds (1, cyclo(L-Pro-L-Tyr) and 2, *N*-acetyltyramine) from *Tenebrio molitor* were able to increase aPPT and clotting time in mice blood (*ex vivo*) and increase endothelin-1 (ET-1) production – a vasoconstrictor – in HUVECs⁽³⁹⁾.

In four studies^(7,47–49), the ability of edible insects to modulate lipid pattern in cellular models was investigated through evaluation of lipid content or the expression of genes related to lipid metabolism. Ethanol extract of *Gryllus bimaculatus*, *Oxya chinensis sinuosa* and *Protaetia brevitarsis seulensis* diminished intracellular lipid accumulation and triacylglycerol (TG) in HepG2, a human liver cancer cellular line⁽⁴⁷⁾. Moreover, lipid accumulation, together with total cholesterol (TC) levels, was reduced in L-02 cells, a human fetal hepatocyte line, by polyunsaturated fatty acids (PUFAs) and α -linolenic acid from *Bombyx mori*. As reported in the same paper, the extracts showed the ability to increase the mRNA and protein expression of cholesterol metabolism-related genes⁽⁷⁾. Incubation with *Tenebrio molitor* larvae decreased lipid droplet formation and TG levels in 3T3-L1 cells (murine pre-adipocytes) and mRNA expression of genes related to adipogenic differentiation and lipogenesis. Additionally, the treatment decreased phosphorylation of extracellular signal-regulated kinase (ERK) and increased phosphorylation of adenosine monophosphate-activated protein kinase- α (AMPK- α) and p-p38 but did not affect phosphorylation of c-Jun *N*-terminal kinase (JNK)⁽⁴⁸⁾. In the same cellular model, ethanol extract of *Protaetia brevitarsis* larvae decreased the mRNA expression of genes related to adipogenesis⁽⁴⁹⁾.

Two studies investigated the effect of edible insects on inflammatory response in cellular lines. In particular, Park and colleagues⁽⁴⁶⁾ evaluated how three tetrahydroquinolines from *Allomyrma dichotoma* affected vascular inflammatory responses in HUVECs: they reduced vascular cell adhesion molecule-1 (VCAM-1) and intercellular adhesion molecule-1 (ICAM-1) levels, adherence of monocytes to HUVECs monolayers and migration of human neutrophils; moreover, they decreased nuclear factor- κ B (NF- κ B) p65 activity, tumour necrosis factor- α (TNF- α) and interleukin-1 β (IL-1 β) production and expression of phospho-p38. Furthermore, protein hydrolysates from *Gryllodes sigillatus* reduced NF- κ B in RAW 264.7 cells⁽⁴¹⁾. Aqueous extract of *Locusta migratoria*, tested on human peripheral blood lymphocytes, did not affect the ratio between micronucleus (MN) and cells, sister chromatid exchange (SCE) and chromosome aberration (CA) formation⁽⁵⁰⁾. Finally, in D-HMVECs glycosaminoglycan from *Gryllus bimaculatus* increased vascular endothelial growth factor (VEGF), but not laminin⁽⁴⁵⁾.

Effect of edible insects in animal and human studies

A total of twenty-five intervention studies in animal models and two in humans have been published, as described in Table 6. Fifteen different edible insects were investigated: five papers

involved *Gryllus bimaculatus*^(6,45,47,51,52) and four *Tenebrio molitor*^(39,48,53,54), while the use of *Oxya chinensis sinuosa*^(37,47,55) and *Protaetia brevitarsis seulensis*^(38,47,49) was reported in three interventions each. One study investigated the effects of both acute and chronic settings, while four were only acute (i.e. single dose) and twenty were chronic (medium to long term) interventions. Six studies^(36,56–60) were conducted on animals that were not affected by particular stresses. Among these, Kim and colleagues investigated any toxic effects of a skimmed powder obtained from *Zophobas atratus* on specific pathogen-free (SPF) Sprague-Dawley rats for a 2-week repeated-dose toxicity study. Neither toxicological lesions nor mortality was observed, nor was body and organ weight influenced. The same outcomes were observed in a 13-week repeated-dose toxicity study; moreover, in macroscopic or histopathological examinations, no test-substance-related toxicological lesions were observed⁽⁶⁰⁾. No toxicity signs were observed in healthy Wistar rats after the acute ingestion of locust (*Caelifera* sp.) powder, which was therefore administered to the same animal model for 28 or 90 d; organ and tissue weight, as well as TG, TC and high-density lipoprotein (HDL) cholesterol plasma levels, were not affected. However, TG and TC liver contents were increased after 90 d of supplementation, as faecal crude lipids. As regards caecum content, TG and TC concentration did not change, while SCFA and insulin levels were decreased⁽⁵⁷⁾. The oil extracted from *Samia ricinii* was administered to healthy weanling Wistar National Institute of Nutrition (NIN) rats for 18 weeks, causing no variation in body weight gain or organ weight. Serum levels of TG and TC were reduced, while low-density lipoprotein (LDL) level was increased; TG content of liver was not affected⁽³⁶⁾. Effects on lipid status were observed also in a study by Ryu⁽⁵⁶⁾: 5 weeks of a diet comprising powder of whole *Bombyx mori* did not affect their body weight or HDL levels, but reduced TG and TC, of healthy Sprague-Dawley rats⁽⁵⁶⁾. Neither body weight gain nor liver and kidney weight were altered in healthy Sprague-Dawley rats after 4 weeks of supplementation with grain larvae. However, their thymus and spleen weight increased, while abdominal fat was reduced. A reduced level of immunoglobulins (IgA, IgM, IgG), blood glucose and TG, TC and LDL was reported. Furthermore, this treatment modified the composition of caecal organic acids, increasing their total quantity and the acetic and propionic acid level, whilst reducing butyric, isobutyric, valeric and isovaleric acid contents⁽⁵⁸⁾.

Zebrafish fed with diet containing different percentages of *Hermetia illucens* (75–100 % of total meal) showed a significant increase in the specific growth rate when compared with the control, while survival was not affected. Considering mRNA expression of IL-1 β , interleukin 10 (IL-10) and TNF- α , groups fed with this percentages of insect powder showed a significant up-regulation with respect to control; furthermore, this condition induced a severe degree of steatosis. As concerns lipid metabolism-related gene expression, no significant differences in fatty acid elongase 2 (ELOVL2) or fatty acid desaturase 2 (FADS2) were detected, while fatty acid elongase 5 (ELOVL5) was up-regulated. Conversely, lower doses (25–50 %) did not significantly affect the previously reported parameters. However, all the doses were able to increase the percentage of lipid categories such as saturated fatty acids (SFA) and *n*-6 and to decrease

monounsaturated fatty acids (MUFA), PUFA, *n*-3 and *n*-9 in zebrafish; histological analyses of fish intestine did not show any morphological alteration or inflammatory event⁽⁵⁹⁾.

Two studies^(61,62) were conducted on malnourished animals. In particular, powder of whole *Acheta domesticus* or *Rhynchophorus phoenicis fabricius* was included in the diet of malnourished weanling Sprague-Dawley rats, increasing their body weight gain, organ (spleen, right kidney, brain, liver) weight, bone mineral content and lean mass. Fat mass was increased only by *Acheta domesticus*; moreover, the relative biological value of the diet was improved by the insect addition⁽⁶¹⁾. Six weeks of a diet based on *Gryllos sigillatus* helped malnourished mice to recover by increasing their body weight; compared with the control group, whose mice did not suffer from malnutrition, the treated mice did not show changes in levels of several inflammatory markers, such as toll-like receptor 4 (TLR4), TNF- α , IL-1 β and interferon γ (IFN- γ), and anti-inflammatory markers, such as interleukin 4 (IL-4), in spleen tissue. Similarly, leptin and adiponectin levels were not affected, while triacylglycerols were reduced⁽⁶²⁾.

The effect of edible insect consumption on animals fed a high-fat diet was evaluated in six studies^(47–49,51,63,64). Chitoooligosaccharides from *Clanis bilineata* administered for 6 weeks improved the lipid status of Sprague-Dawley rats fed a high-fat diet by reducing TC, TG and LDL and increasing HDL levels. Faecal excretion of TG and TC was increased, while food efficiency ratio was decreased; although food intake remained stable, body weight gain decreased⁽⁶³⁾.

The ethanolic extracts of *Protaetia brevitarsis* larvae, administered with a high-fat diet for 7 weeks, decreased body weight gain, epididymal and subcutaneous fat weight and liver – but not spleen – weight of obese C57BL/6J mice; the treatment increased glutathione peroxidase (GPx) and CAT and reduced lipid droplet accumulation in mouse liver, as well as TG, TC and LDL levels⁽⁴⁹⁾. Ethanolic extract of *Gryllus bimaculatus*, added to a high-fat diet, did not affect body weight or epididymal fat in obese rats, but it reduced total and abdominal fat weight. Moreover, it changed the composition of abdominal fat, reducing MUFA and increasing PUFA content. The supplementation did not affect TC, HDL or LDL levels, but reduced TG, IL-10 and glucose levels. Even though serum CAT was not affected, the prolonged treatment with ethanolic extract of *Gryllus bimaculatus* reduced protein and lipid oxidative damage caused by high-fat diet in both liver and blood, where serum uric acid and carbonyl concentrations were reduced⁽⁵¹⁾.

In a study by Im and colleagues⁽⁴⁷⁾, the ethanol extracts of three different insects, that is, *Gryllus bimaculatus*, *Oxya chinensis sinuosa* and *Protaetia brevitarsis seulensis*, were administered for 14 weeks to C57BL/6J mice subjected to a high-fat diet to counteract the effects of non-alcoholic fatty liver disease. All the three supplementations reduced body weight gain and liver weight, together with intestinal and epididymal adipose tissue, but no significant effect was described for abdominal and kidney adipose tissue. Moreover, when inflammation markers were measured in liver and adipose tissue, mRNA levels of TNF- α , IL-1 β and interleukin 6 (IL-6) were found to be decreased. *Oxya chinensis sinuosa* and *Gryllus bimaculatus* reduced lipid droplet accumulation in liver, TG, TC, LDL, HDL and non-

esterified fatty acid (NEFA) levels, together with fasting blood glucose, while among these markers *Protaetia brevitarsis seulensis* reduced only lipid droplet accumulation, TC, NEFA and fasting blood glucose⁽⁴⁷⁾.

Zou *et al.*⁽⁶⁴⁾ reported that, in Wistar rats with hypercholesterolaemia, *Bombyx mori* pupae oil supplementation was able to counteract the impairments induced by a high-cholesterol diet: lipid status was improved through a reduction of TC, LDL and LDL/HDL ratio without affecting TG or HDL levels. Moreover, the size of liver, kidney and heart was reduced, while that of spleen was increased. The supplementation improved antioxidant status, restoring superoxide dismutase (SOD) levels, increasing TAC levels, decreasing MDA in liver and serum and restoring the activity of GPx in rats' liver⁽⁶⁴⁾. Finally, 6 weeks of treatment of obese BALB/c mice with whole powder of *Tenebrio molitor* resulted in a decrease in body weight gain and epididymal and abdominal-to-peripheral adipose cell volume, as well as a decrease in lipid accumulation in liver⁽⁴⁸⁾.

Two studies^(8,45) investigated whether the consumption of edible insects was able to counteract the effect of diabetes in mice. The supplementation with ethanol extract of the sericin layer from the green cocoon shell of *Bombyx mori* improved glycaemic status in obese mice with type 2 diabetes, increasing blood glucose reduction rate and reducing blood glucose during tolerance tests, fasting blood glucose and insulin, glycosylated haemoglobin (HbA1c) and homeostasis model assessment of insulin resistance (HOMA-IR), while improving insulin sensitivity index (ISI). Pancreas functionality was restored: islet structure and area of pancreatic beta cells and the insulin quantity secreted by them was increased. Treatment also improved antioxidant status, increasing the activity of liver GPx and SOD and reducing the liver content of MDA and 8-hydroxy-2'-deoxyguanosine (8-OHdG). Inflammation markers in liver, as well as infiltrations and oedema and NF- κ B, IL-6 and TNF- α levels, were also reduced, while body weight was not affected⁽⁸⁾. Glycosaminoglycan extracted from *Gryllus bimaculatus* and administered for 1 month to BKS.Cg-m+/+Lepr^{db} diabetic mice, despite not affecting body weight and abdominal fat, reduced circulating levels of carbonyl. As regards antioxidant enzymes, this extract did not affect GST, but improved activity of CAT and GPx. It also reduced adipocyte density in lungs, liver and kidneys, but not in pancreas. However, the treatment improved the status of pancreas islet and liver tissue that was damaged by the diabetic condition. Blood glucose levels were not affected, whilst non-fasting blood glucose was reduced only after 1 week of treatment⁽⁴⁵⁾.

The effect of edible insects against stress induced by alcohol consumption was evaluated in three different studies^(6,53,54). The first reported positive action of the aqueous extract of *Gryllus bimaculatus* in restoring the normal physiological levels of 8-OHdG levels and MDA content in liver and small intestine of C57BL/6J mice with liver damage caused by acute alcohol exposure; liver droplet accumulation was also reduced, as well as levels of inflammation markers (F4/80-positive Kupffer cells (F4/80+ KCs) and IL-1 β)⁽⁶⁾. The treatment with fermented *Tenebrio molitor* powder of Sprague-Dawley rats, impaired by a chronic alcohol diet, reduced liver weight and increased activity of glutathione reductase (GR) and total and reduced GPx hepatic content, but no effect was observed on SOD and CAT

levels. Moreover, phosphorylated-inhibitor of nuclear factor-kappa B- α (p-I κ B- α) levels were reduced, as well as serum and liver content of TNF- α , but not of IL-6. As regards lipid status, hepatic levels of TG, NEFA and TC were also reduced, as alcohol-induced hepatic lipid accumulation, focal necrosis and mild fibrosis were attenuated. Also, expression of lipid synthesis transcription factors, triacylglycerol synthesis-related genes and cholesterol synthesis and esterification-related genes was reduced, while expression of fatty acid uptake and transport-related genes was not affected.⁽⁵³⁾ The treatment with fermented defatted *Tenebrio molitor* powder of Sprague-Dawley rats fed with a chronic alcohol diet dose-dependently increased hepatic β -oxidation⁽⁵⁴⁾.

Concerning the effect on coagulation markers^(37–39), compounds extracted from *Protaetia brevitarsis seulensis*⁽³⁸⁾, *Oxya chinensis sinuosa*⁽³⁷⁾ and *Tenebrio molitor*⁽³⁹⁾ decreased arterial thrombi formation rate and size in mice with arterial thrombosis and decreased mortality in mice with pulmonary thrombosis; in this model, 1, cyclo(L-Pro-L-Tyr) and 2, N-acetyltyramine from *Tenebrio molitor* were also able to reduce thrombi size⁽³⁹⁾. Indole alkaloids and N-acetyldopamine dimers extracted respectively from *Protaetia brevitarsis seulensis*⁽³⁸⁾ and *Oxya chinensis sinuosa*⁽³⁷⁾ increased tail bleeding time in healthy mice.

Edible insects were also able to improve antioxidant and inflammatory status that was previously impaired by different stressing agents such as sepsis⁽⁴⁶⁾, varicocele⁽⁵²⁾ or UV radiation⁽⁵⁵⁾. Four tetrahydroquinolines from *Allomyrina dichotoma* were administered twice to caecal ligation and puncture (CLP)-induced septic C57BL/6 mice, resulting in an increase in the survival rate. There were no significant differences between the lungs of the treated and untreated mice: interstitial oedema was observed, and the pulmonary architecture was severely damaged. Moreover, to evaluate effect of compounds against inflammation, lipopolysaccharide (LPS)-induced vascular permeability in mice was inhibited by the tetrahydroquinolines^(1–3), which reduced LPS-induced leucocyte migration into the murine peritoneal cavities⁽⁴⁶⁾. Moreover, treatment with *Gryllus bimaculatus* improved antioxidant and anti-inflammatory status in testicular tissue of Sprague-Dawley rats affected by varicocele. Furthermore, the increased levels of MDA, ROS and reactive nitrogen species (RNS) were significantly reduced⁽⁵²⁾. Finally, the extract of *Oxya chinensis sinuosa*, administered to Hos/HR-1 hairless mice for 12 weeks, reduced the damage induced by UV irradiation as well as inflammation markers (IL-1- β , IL-6 and TNF- α), while the activity of SOD and CAT was increased⁽⁵⁵⁾.

Evidence in humans is available only for two dietary intervention trials. Wheat noodles enriched with *Bombyx mori* powder, were provided to thirteen healthy humans, fasting in the morning, following a cross-over acute ingestion design. The *Bombyx mori* noodles significantly reduced post-prandial blood glucose, glucose peak, area under the curve (AUC) of glucose and glucose index (GI) compared with control noodles⁽⁶⁵⁾. In a recent chronic intervention study, 25 g/d of dried roasted cricket powder, included in a muffin and in a dry mix shake, was given to twenty healthy humans for breakfast for 14 d following a cross-over design. The treatment did not modify intestinal microbiota, gastrointestinal functionality, glycaemia or IgA levels of

the subjects. However, TNF- α plasma levels decreased, as did the acetate, propionate, bile acids and TG content of faeces⁽⁶⁶⁾.

Discussion

In this work, we summarised the body of evidence on selected functional properties of edible insects in modulating oxidative and inflammatory stress, platelet aggregation, lipid and glucose metabolism and weight control in different experimental models. Concerning the considered species, among a total of forty-three edible insects, the Gryllidae family and *Tenebrio molitor* were the most investigated, respectively in seventeen and sixteen studies. Both species are included in the list of insects for human consumption from the European Food Safety Authority and are also widely present in the market. Regarding the different aspects, antioxidant properties were investigated in thirty-six different species (*Tenebrio molitor* and Gryllidae in fourteen and twelve studies, respectively); eleven species were tested for their ability to affect lipid status (*Gryllus bimaculatus* and *Bombyx mori* in four studies and *Tenebrio molitor* in three studies). As regards the anti-inflammatory properties, the effects of nine insects were investigated: *Gryllus bimaculatus* was cited in five research articles, whilst *Grillodes sigillatus* and *Bombyx mori* were tested in two studies each. *Gryllus bimaculatus* was investigated for its ability to modulate glucose metabolism in four different studies, followed by *Grillodes sigillatus* with two studies. Lastly, the studies involving coagulation markers investigated seven insects, of which *Tenebrio molitor* and *Protaetia brevitarsis seulensis* were evaluated twice each. It is noticeable that the Gryllidae family, represented mainly by *Gryllus bimaculatus* and *Grillodes sigillatus*, is the most cited in all the topics, the only exception being the effect on coagulation. Moreover, one of the two studies on human subjects involved crickets of the Gryllidae family, while the other study was focused on *Bombyx mori*.

The antioxidant properties of edible insects were investigated in thirty-six out of fifty-five studies involving animals ($n = 11$), cell cultures ($n = 9$) or *in vitro* ($n = 22$). Results clearly show that all insects tested *in vitro* and in cellular models displayed radical scavenging or metal ion chelation properties, as well as the ability to modulate glutathione S-transferase and catalase, with activity depending on the utilised concentration. Also, the findings in animal models were consistent with *in vitro* results, supporting the antioxidant properties of edible insects. According to the different studies, the effect was evident in serum, liver and skin, with a mechanism of action ranging from the increase in antioxidant capacity up to the modulation of endogenous antioxidant enzymes. In the majority of the studies, the antioxidant effect was more evident when specific stressors such as high-fat diet, oxidative stress, obesity, alcoholic liver disease or UV irradiation were present. Although the antioxidant properties have been widely investigated *in vitro* and in cellular and animal models, studies on human subjects are lacking. Overall, the evidence from the available studies clearly showed that all the tested insects, with varying ability, were able to reduce an induced oxidative stress, modulating redox status of cellular and body fluids and restoring the impaired activity of antioxidant enzymes.

The effect on lipid markers was evaluated in twenty-one research articles; the studies were carried out mostly in animal models ($n = 18$), while only few were in cellular models ($n = 4$) or *in vitro* ($n = 1$). When markers of dyslipidaemia were evaluated in animal studies, edible insects were able to reduce TG (10/12), TC (7/11) and LDL (6/8) levels, without affecting HDL (5/7), in body fluids and tissues. Moreover, the reduction of TG was detected also in hepatocytes and adipocytes, while a study reported a reduction of TC in hepatocytes. Reduction of lipid droplets in liver was detected in animal models (7/7) and in two studies on cellular model of hepatocytes, suggesting an effect on liver steatosis. A reduction in fat tissue weight or volume was detected in five out of seven studies, together with a change in fat composition (2/2). Furthermore, edible insects positively modulated lipid metabolism and fat accumulation in animal ($n = 2$) and cellular models ($n = 3$). The only *in vitro* study evaluating the ability of edible insects to affect lipid metabolism reported an inhibition of pancreatic lipase, with the possible consequence of preventing the breakdown of triacylglycerol and delaying the absorption of fatty acids⁽⁶⁷⁾. Furthermore, the effect of consumption of edible insects on weight control, expressed as body weight (BW) or body weight gain (BWG) was evaluated in fifteen studies; considering both parameters, either an increase (2/15) or a decrease (4/15) was reported, but in most of the cases no significative effect was observed (9/15). However, if the results are classified on the basis of the type of diet or the body weight at the baseline, the supplementation with edible insects induced a decrease in BW or BWG in four out of five cases in obese animals or following a high-fat diet, while an increase of BW was reported when malnourished animals were considered ($n = 2$). Based on the evidence from animal studies, dietary intervention with edible insects reduced TC, TG and LDL, while the effect on BW was dependent on whether animals were obese or malnourished.

The anti-inflammatory properties of edible insects were evaluated in sixteen papers, twelve *in vivo* and five in cellular models. Concentrations of cytokines were evaluated in nine *in vivo* studies, leading to a reduction in circulating levels increased by different stressors (8/9). An increase in cytokine levels was observed only when high doses of *Hermetia illucens* (75–100 % of the total diet) were given to healthy zebrafish⁽⁵⁹⁾, although no inflammatory events were observed through histological analysis. An effect on reducing circulating levels of TNF- α was shown in humans, although we need to consider that the study was conducted on healthy subjects, who may have low or scarce levels of inflammation, and that the number of subjects enrolled was low⁽⁶⁶⁾. Effects on immunoglobulins were evaluated in two papers: IgE levels – as an identifier of allergic reactions in rats – were not detected, whilst an increase in IgG, IgA and IgM was revealed, considered by the authors to be a bifidogenic effect from the antimicrobial peptides in the used extract. NF- κ B levels, transcription-factor-regulating genes involved in inflammatory responses, were decreased in cellular models, HUVECs and RAW 264-7, and in an animal study. However, TLR4 levels, whose stimulation leads to NF- κ B activation, were not affected. Concerning the effect on specific organs, reduction

of inflammatory infiltrations and oedema in liver, leucocyte migration and vascular permeability in peritoneum were observed. Finally, three studies showed activity in reducing NO production in macrophages, a radical involved in the modulation of inflammation and immunity⁽⁶⁸⁾. Therefore, the evidence from cellular and animal models supports an effect on reducing inflammatory cytokines thorough the modulation of NF- κ B levels, without affecting immunoglobulins.

Effects of edible insects on glucose and/or insulin status were evaluated in nine studies; six out of the total were *in vivo* interventions, and three were *in vitro* studies. Most of the *in vivo* studies reported improvements in diabetes markers, such as reduced blood glucose (4/6) and serum insulin (2/2), as well as an enhancement of pancreas structure and functionality (2/2). Moreover, a reduction in glycated haemoglobin (HbA1c), a tool in both routine management and diagnosis of diabetes⁽⁶⁹⁾, was detected. However, only two interventions out of six involved diabetic mice. All the *in vitro* studies outlined a positive effect through the inhibition of DPP-IV and α -glucosidase, suggesting a role in modulating glucose homeostasis⁽⁷⁰⁾. The results *in vitro* and in animal models are supported by evidence in humans showing an effect of noodles enriched with *Bombyx mori* powder on reducing glycaemic index⁽⁶⁵⁾. Although further evidence is needed, results suggest that edible insects can modulate glycaemia/insulin homeostasis with potential effect on the management of glycaemic index, a key aspect for diabetes prevention.

Anti-coagulation properties of edible insects have been investigated in only four different works: three out of four studies were performed by the same research group, investigating compounds extracted from one insect for each, using either *in vitro*, *ex vivo*, cellular or animal models *vivo* with acute interventions, while the other study investigated *in vitro* the properties of seven different insects. The selected edible insects showed anti-coagulant properties *in vitro* and *ex vivo*, by reducing aPTT and PT – both measuring speed of blood clotting – and affecting the coagulation cascade by modulation of factor X and thrombin. Moreover, a reduction in PAI-1 was recorded, whose inhibition may reduce the incidence of thrombotic events⁽⁷¹⁾. PAI-1 is also an important physiological inhibitor of tPA⁽⁷²⁾: this molecule, whose main function within the vascular system is the removal of fibrin⁽⁷³⁾, was not affected by *in vitro* treatment with compounds extracted from edible insects. When compounds derived from edible insects were administered to mice in acute interventions, a decrease of thrombi formation and consequent mortality in mice models of arterial or pulmonary thrombosis was observed; moreover, these compounds prolonged tail bleeding time in healthy mice. Overall, the available literature is scarce and from one research group only, making it hard to draw any conclusion on the anti-platelet aggregation properties of edible insects.

In terms of research quality of the results, the evidence from *in vitro*, cellular and animal studies is solid because the markers and biomarkers utilised are appropriate and the animal studies are properly designed. Human studies involved a low number of subjects, short period of dietary intervention and lack of a

dietary assessment of the subjects during the studies, making it difficult to draw any conclusions about the clinical relevance of the findings.

A few more considerations are needed: first of all, different species display similar activities such as antioxidants and anti-inflammatory properties, despite being characterised by different genes, habitat, physiological needs and eating habits, making it difficult to associate the specific functional role to the different species. Studies assessing the functional properties of different species characterised by dietary habits and environment, as well as the effect of different feedings, will help us to understand the variables influencing the functional properties of the insects.

Secondly, the majority of the studies tested extracts without identifying the compounds responsible for the functional effect or without testing sub-fractions with different chemical composition. However, although functional properties of the edible insects might be related to their variegated molecular composition, as occurs for foods, the identification of the single molecules is a necessary step for nutraceutical or pharmacological purposes.

Thirdly, the evidence from *in vitro*, cellular and animal models on the functional properties reviewed in this manuscript is based on robust and reliable markers with a statistical significance. However, the scarce evidence in humans highlights the urgent need for long-term dietary intervention trials to endorse the role of edible insects as functional foods. The step of human consumption is critical because there is no information on the bioavailability of functional molecules from edible insects in body fluids following consumption. Moreover, it would be interesting to understand the effect of intestinal microbiota on molecules from the digestion processes of edible insects and if the arising metabolites are endowed with functional properties. In this view, it would be interesting to develop observational or epidemiological studies assessing biomarkers related to antioxidant, inflammatory and immune status as well as microbiota analyses in populations where insects are a part of the usual diet.

The inclusion of edible insects in the category of functional food might open a new scenario for food industries through the enrichment of food with proteins, extracts or flour obtained from edible insects, representing a novel aspect with scientific, commercial and social impacts on society in the following years. However, for such an approach to be efficacious, it is important to investigate consumers' acceptance, mainly in Western countries, of insect-based foods⁽⁷⁴⁾. This is a process that requires time and campaigning of information for consumers, focusing on the importance of reducing the ecological impact of animal-based products with more sustainable and functional foods.

Conclusions

In conclusion, based on the available body of evidence, mainly *in vitro*, cellular and animal studies, edible insects are a promising source of bioactive ingredients endowed with antioxidant and anti-inflammatory properties, involved in the modulation of glucose and lipid metabolism, potentially leading to health benefits. However, dietary intervention trials are urgently needed to confirm and define the efficacy of edible insects as

functional foods in humans, confirming the promising evidence from *in vitro* and animal models. We think that this review could promote research on the functional properties of edible insects, amplifying our knowledge on this topic and ensuring that correct information is relayed from the media to consumers.

Author contributions

M.S. conceived the topic of the review and supervised the work. V.D.A. wrote the initial draft. N.B., G.S. and C.D.M. revised the work. All authors approved the submitted version.

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Conflict of interest

None.

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