

81989_Auto_Edited.docx

Antihepatoma peptide, scolopentide, derived from the centipede scolopendra subspinipes mutilans

Hu YX *et al.* An antihepatoma peptide derived from centipede

INTRODUCTION

Centipedes¹⁴ have been widely used in traditional Chinese medicine to treat many diseases, including convulsions, strokes, rheumatic joint diseases, snake bites and tumors^[1]. It is a major challenge to isolate and characterize the effective components of centipedes due to limited peptide purification technologies for animal-derived medicines. Currently, the main extraction technologies are as follows: Aqueous extraction^[2], organic solvent extraction^[2-5], enzymatic hydrolysis^[6], ultrasonic extraction^[2], and freeze-thaw extraction^[2]. To obtain low-weight peptides, extracted proteins require further purification. At present, the main purification technologies of animal proteins are as follows: Sephadex column gel chromatography (GC)^[3,7,8], HPLC/RP-HPLC^[3,5-8], ion exchange chromatography^[7,9], and graded ultrafiltration^[7]. SDS-polyacrylamide gel electrophoresis (SDS-PAGE) is a traditional separation and identification technique^[10], and mass spectrometry (MS)⁵⁷ is the most widely used technique for identifying peptides for use in animal-derived medicines^[5-7]. To date, there is no single method that can extract a single peptide from complex mixtures (Table 1). Compound extraction plans are mostly used, which are costly and complicated. Consequently, animal-derived medicines have not been developed as herbal medicines. Currently, the main extracts are still crude peptides, and they have some disadvantages, such as a low extraction ratio, undefined composition, unstable properties and susceptibility^[11,12]. These disadvantages may be prevented by identifying and characterizing the effective compositions of animal-derived medicines and synthesizing them *in vitro*.

Some ancient Chinese medical studies have reported that centipedes can be used to treat tumors. “Compendium of Materia Medica”^[13], a famous ancient Chinese pharmaceutical work, recorded that centipedes can be used to treat tumors in the chest and abdomen. It is also recorded in the modern Chinese medical literature “Integrating Chinese and Western Medicine”^[14] that centipede liquor can be used to treat esophageal cancer with excellent clinical effects. Moreover, the Shendan Sanjie capsule, a medicine containing centipedes, has already been applied to treat cancers and has been proven to inhibit colitis-associated cancer^[15]. Although the antitumor theory and clinical efficacy of centipedes have been verified for hundreds of years in China, current experimental studies still focus on antimicrobial and anticoagulation agents. Only a few studies have observed the antitumor effects of centipedes (Table 1). The potential mechanism is also poorly understood. The tumor types focused on have been gliomas^[3], cervical cancer^[16], leukemia^[17], epidermal cancer^[4], and melanoma^[18], rather than tumors in the chest and abdomen. Purified centipede peptides can suppress tumor cells by inducing caspase-related apoptosis through a mitochondria-dependent pathway^[3,4,16,18], inducing cell cycle arrest^[3,18] and necrosis through a specific interaction with phosphatidylserine (Table 1)^[17]. Additionally, our previous studies showed that crude centipede peptides could induce apoptosis in HepG2 cells and lung cancer A549 cells by arresting cells at the G2/M phase^[19,20].

Apoptosis has been widely used as a target in cancer treatment^[21]. There are two main apoptosis signaling pathways: Mitochondria-dependent and mitochondria-independent pathways. The mitochondria-dependent pathway is controlled by Bcl-2 family proteins and is activated by DNA damage caused by chemotherapy and radiotherapy through activation of the tumor suppressor gene p53^[22]. However, the loss of p53 is common in the clinic, contributing to the resistance of conventional chemotherapy and the development of most cancers^[23,24]. Thus, the mitochondria-independent pathway, which is independent of p53, has become a promising target^[25]. Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) is part of the mitochondria-independent pathway. TRAIL can selectively initiate apoptosis of cancer

cells but has no obvious toxic effect on normal cells^[26-28], which makes it a promising chemotherapy drug for many cancers. Nonetheless, recent studies have shown that some cancers, including liver cancer and esophageal cancer, are resistant to TRAIL-mediated apoptosis, which greatly reduces its clinical efficacy^[29,30]. Current research focuses on finding suitable TRAIL sensitizers and developing TRAIL receptor agonists to maximize the cell-killing effect. However, current studies mainly focus on sensitizers rather than receptor agonists, and the clinical and laboratory results are not ideal^[26,28]. Therefore, for research and clinical applications, it would be useful to find stable and efficient TRAIL receptor agonists. Recently, ³ the regulatory effect of Chinese herbal extracts on the TRAIL apoptosis pathway was gradually discovered. Chinese herbal extracts such as luteolin and artemisinin derivatives can induce ¹⁸ apoptosis of liver cancer cells by upregulating the expression of death receptor 4 (DR4) or DR5, which are TRAIL receptors on the cell surface^[31,32].

In this study, we isolated a novel low-molecular-weight antihepatoma peptide, designated scolopentide, from *Scolopendra subspinipes mutilans* L. Koch and synthesized it *in vitro*. We observed that synthetic scolopentide inhibited tumor proliferation and was more toxic to HepG2 human liver cancer cells than to L02 human embryonic hepatocytes. Mechanistically, scolopentide activated TRAIL receptors (DR4 and DR5), ²⁰ upregulated the expression of fas-associated death domain protein (FADD), caspase-8 and caspase-3 through a mitochondria-independent pathway and finally induced tumor cell apoptosis.

³ **MATERIALS AND METHODS**

Chemicals and reagents

Dulbecco's modified Eagle medium (DMEM) was purchased from Gibco (Thermo Fisher Scientific Inc., United States) and HyClone (Logan, United States). Fetal bovine serum (FBS) ⁶ was purchased from Gibco and Invitrogen (Thermo Fisher Scientific Inc., United States). DMSO was purchased from Amresco (Solon, United States). PBS was purchased from Boster (Wuhan, Hubei Province, China) and WellBio (Shanghai, China).

Trypsin was purchased from Invitrogen. Penicillin-streptomycin was purchased from Beyotime (Shanghai, China). Hoechst was purchased from SolarBio (Beijing, China).

Cell culture

The Bel-7402 Liver cancer cell line, Eca-109 esophageal cancer cell line and A549 Lung cancer cell line were purchased from the Cell Resource Center of Shanghai University of Biology. The HepG2 Liver cancer cell line and L02 human embryonic hepatocyte cell line were purchased from The Cell Center of Xiangya Medical College, Central South University. The SMMC-7721 liver cancer cell line was purchased from the BeNa Culture Collection (Suzhou, Jiangsu Province, China). Cells were placed in culture flasks (Corning Inc., United States) and cultured in a constant temperature incubator (SHELLAB, Hitachi, United States) at 37 °C and 5% CO₂ with saturated humidity. The medium comprised 10% FBS, low-glucose DMEM and penicillin-streptomycin (100 U/mL). Cells were observed under an inverted microscope (Olympus, Japan).

Cell viability assay

The viability of HepG2, A549, Eca-109 and Bel7402 cells treated with different centipede extracts was determined by a Cell Counting Kit-8 (Dojindo, Japan). The viability of HepG2 and L02 cells treated with synthetic scolopentide was also determined. Groups and treatments were as follows: (1) Vehicle group: 100 µL cell suspension + 100 µL complete medium; (2) zero group: 200 µL of complete medium; and (3) scolopentide group: Cells were treated with different centipede extracts/synthetic scolopentide. Three replicate wells were used for each group.

Cells at the logarithmic growth stage were collected, and the density was adjusted to 1×10^4 cells/mL. The cell suspension (100 µL) was plated into a 96-well plate (Corning Inc., United States). Four hours later, drugs were added after the cells had adhered to the well. Twelve hours later, 100 µL culture solution containing 10% CCK8 was added to each well. Cells were incubated for 2 h at 37 °C with 5% CO₂. Then, the absorbance (OD) at 450 nm was analyzed by a microplate reader, and the mean was calculated. Cell

viability (%) = scolopentide group OD/vehicle group OD × 100%. The cell suppression ratio (IR) was determined with the following formula: IR (%) = (vehicle group OD - scolopentide group OD)/vehicle group OD × 100%. The zero group OD was subtracted from the OD of all groups for correction. The 50% maximum inhibitory concentration (IC₅₀) was determined by constructing dose-response curves.

Extraction of Crude Peptides from Scolopendra subspinipes mutilans

Crude centipede peptides were obtained by enzymolysis combined with acetone precipitation. The “Pharmacopoeia of the People’s Republic of China (2020)” references the use of the whole dried body of *Scolopendra subspinipes mutilans*^[1]. Centipedes were purchased from the pharmacy department of the First Affiliated Hospital of Hunan University of Chinese Medicine. They were identified as *Scolopendra subspinipes mutilans* L. Koch by Chinese herbal experts. Then, the whole bodies of the centipedes were ground into ultrafine powder by low-temperature ultrafine grinding in the Superfine Powder Engineering Research Center of the Ministry of Education of China. First, 100 g of centipede was dried at 50-60 °C. The superfine powder (particle size 1-75 μm) was crushed by a BFM-6 Bailey superfine powder machine (amplitude 100 Hz, grinding temperature 0-10 °C, grinding time 10-15 min). Microscopic image analysis and laser light scattering were used in combination to assess the particle size, and the particle size of more than 95% of the powder was smaller than 75 μm. Finally, the powder was stored at room temperature until subsequent experiments.

We chose an optimized enzymatic hydrolysis method. The extraction process was as follows: 2 g of centipede superfine powder was added to 10 mL of double distilled water, mixed with trypsin, and incubated in water at 55 °C for 4 h. The trypsin was inactivated in a 99 °C water bath for 10 min. Then, the samples were centrifuged at 3500 r/min for 5 min (low temperature ultracentrifuge, Pharmacia Inc., Sweden), and the supernatant was collected. The samples were centrifuged at 5000 r/min at 4 °C for 10 min. The supernatant was collected again. Two volumes of precooled acetone were added to precipitate the former supernatant, which was centrifuged at 5000 r/min at 4

°C for 10 min. The precipitate was dissolved in double steaming water, and the process was repeated twice. Finally, 2 volumes of precooled acetone and petroleum ether (1:1) were added to precipitate again, which was centrifuged at 5000 r/min at 4 °C for 10 min, and the precipitate was collected. Finally, the precipitate was freeze-dried (FD-1 Freeze Dryer, Rikakika Co., Ltd., Tokyo, Japan), weighed, and stored at -20 °C, and the protein content was determined for subsequent experiments.

Optimization of enzymolysis extraction methods

In the process of preparing crude centipede peptides, we tested the influence of single factors on the extraction rate of the crude extract. The five factors included protease type (trypsin, pepsin), water-centipede ratio, protease dose, enzymolysis temperature and enzymolysis time. Then, an L₉(3⁴) orthogonal design was adopted to investigate the effects of enzymolysis time, enzymolysis temperature and trypsin dose on crude extracts, which were the three most influential factors in single-factor experiments (Table 2). In addition to the extraction ratio, a CCK8 assay was used to detect the suppression effect of different crude extracts on HepG2 cell viability to determine the optimal technological conditions. We also compared the cytotoxicity of the crude extracts obtained from two methods, freeze-thawing with liquid nitrogen and enzymolysis with acetone precipitation, under optimal technological conditions to determine the significance of this improvement.

Purification and identification of an antihepatoma peptide from crude centipede peptides

Gel chromatography: For the first step, a Sephadex G-25 column was used to isolate the component with the strongest antitumor effect. With 0.1% trifluoroacetic acid-acetonitrile as the mobile phase, gradient elution was performed using an AKTA protein purification system (Amersham Bioscience, United States). In detail, a crude centipede peptide solution (50 mg/mL) was slowly added to a Sephadex G-25 column. Chromatographic columns were eluted with 0.1% trifluoroacetic acid-acetonitrile (flow

rate 0.5 mL/min). The resulting columns (3 mL/tube) were collected by an automatic collector. After isolation, tubes with the same peak are collected together, according to the peaks displayed on the computer ultraviolet spectrum. The protein contents of different peaks were determined by a fluorescence microplate reader (TECAN, Austria). The CCK8 assay was used to assess the antitumor activity of different peaks and to select the strongest peak for subsequent experiments.

High-performance liquid chromatography: For further purification, high-performance liquid chromatography (HPLC) was used as the next step. Initially, we used a conventional C18 column, but the results showed that the sample was hydrophilic. Thus, a hydrophilic XAmide column (5 μ m, 100 Å, 4.6 mm \times 250.0 mm) was chosen. The sample solution (1 mg/mL) was filtered through a 0.45 μ m microporous membrane and eluted with acetonitrile-water-100 mmol/L ammonium formate (pH 3.5) (flow rate 4 mL/min, detection wavelength 260 nm, injection volume 20 μ L). Different fractions were collected according to the HPLC chromatogram peaks. The CCK8 assay was used to select the fraction with the strongest antitumor activity. Then, the above HPLC and CCK8 assays were repeated. The final centipede peptide was concentrated, purified, freeze dried and designated scolopentide.

Determination of the molecular weight and peptide sequence of scolopentide

Quadrupole time of flight mass spectrometry (UPLC-QTOF MS 6530, SHIMADZU, Japan) was used to measure molecular weight. The ion source parameters were as follows: Electrospray ion source, positive ion mode, temperature 300 °C, gas flow 11 L/min, and spray pressure 15 psi. Scolopentide did not pass through the liquid column but went directly through the mobile phase to the QTOF system. The Mascot search engine from Matrix Science was used to match peptide sequences. Based on the protein sequence library of Mascot, the molecular weight of scolopentide was compared with that of known centipede proteins. The amino acid sequence that best matched the molecular weight was selected.

Solid-phase peptide synthesis

Scolopentides that were characterized by QTOF MS and Mascot were synthesized using solid-phase peptide synthesis methods. Peptide synthesis was performed by Genscript (Nanjing, China) as follows: (1) A total of 400 mg chloromethyl resin was selected to synthesize the polypeptides, 20% piperidine/¹⁹N, n-dimethylformamide was added to the reactor, and the reactor was shaken for 20 min; (2) the solvent was removed by filtration, ¹⁹N, n-dimethylformamide was added to the system, and the reactor was shaken for 1 min, followed by filtering to remove the liquid, which was carried out three times; (3) a total of 150 μ L ninhydrin and a small amount of resin were added to the detection tube, which was placed at 100 °C for 20 s, after which the resin's color was checked for change, which indicated that Fmoc was removed successfully; (4) the prepared amino acid solution was added to the reactor, 1 mL of N,N'-diisopropyl carbon diimine/1-hydroxybenzotriazole was added, and the reactor was shaken for 1 h; (5) step 3 was repeated, and if the color did not change, the coupling was considered successful; (6) step 2 was repeated to wash the resin; and (7) the above operation was repeated, and the corresponding amino acids were added until polypeptide synthesis was completed. Then, HPLC was used to test the purity of the synthetic scolopentide, and MS was used to test the molecular weight. Finally, the peptides ⁵were freeze-dried and stored at -20 °C until they were used.

²¹*Flow cytometry*

The apoptosis of HepG2 cells treated with extracted scolopentide was determined by Annexin V-Alexa Fluor660/7-AAD (BB20121, BestBio, China). The cells were digested and collected by trypsin. Approximately 5×10^5 cells were collected after washing ⁶²with PBS twice and centrifuged at 2000 rpm ⁵for 5 min. Then, 100 μ L of Annexin V binding ⁶³buffer was added to resuspend the cells, and the concentration was adjusted to 3×10^6 cells/mL. Five microliters ¹of annexin V-FITC and 5 μ L of propidium iodide were added ⁴⁶to the cell suspension, mixed and incubated for 10 min at 4 °C in the dark. Then, 400 μ L

of PBS was added to the cell suspension and mixed. Within 1 h, flow CytoFlex (A00-1-1102, Beckman, United States) was utilized to detect cell apoptosis.

Molecular docking

To investigate the antihepatoma mechanism of scolopentide, Discovery Studio 2020 Client and AutoDock-Vina 1.1.2 were used to predict the binding affinity between scolopentide and DR4 and DR5. BIOVIA Discovery Studio 2020 was applied to process and visualize the results. To obtain molecular structure files, scolopentide structures were constructed with Discovery Studio (Small Molecules) according to the sequence RAQNHYCK. Molecular structure files of DR4 (PDB code 5CIR) and DR5 (PDB code 419X) were retrieved from the RCSB PDB database (website: <https://www.rcsb.org/>). The DR4 and DR5 structures were modified by Discovery Studio for water deletion, binding pocket prediction, hydrogenation, and conformation optimization. Then, AutoDock-Vina was used to perform molecular docking and calculate the affinity potential energy of the protein molecule potential binding modes between scolopentide and DR4 and DR5.

Hoechst staining assay

HepG2 cells were inoculated into 6-well plates (1×10^6 cells per well), and 100 µg/mL synthetic scolopentide was added. The cell slides were removed and washed with PBS 2 or 3 times after incubation at 37 °C and 5% CO₂ with saturated humidity for different durations (6 h, 12 h, and 24 h). Then, the slides were fixed with 4% paraformaldehyde for 30 min and rinsed with PBS 3 times. Hoechst (working concentration: 25 µM) was added to the slides, which were incubated for 3 min and rinsed with PBS 3 times. Finally, the slides were sealed with glycerin buffer, stored away from light and observed under a fluorescence microscope (Zeiss, Germany).

Animal and tumor xenograft experiments

Male 5-week-old BALB/c nude mice (Shanghai SLAC Laboratory Animal Co., Ltd., Shanghai, China) were chosen for the formation of subcutaneous xenograft tumors. SMMC-7721 cells (0.2 mL/mouse, 1×10^7 cells/mL) were subcutaneously injected into the right back. The tumor diameter reached 1 cm at 3 wk. Qualified model mice were randomly divided into 2 groups ($n = 6$ per group): The scolopentide group (intraperitoneal injection of synthetic scolopentide, 500 mg/kg/d) and the vehicle group (same volume of normal saline). The length and width of tumors were measured every 2 d, and volume was calculated as the maximum tumor length \times width² \times 0.5. After 14 d of treatment, all mice were euthanized by cervical dislocation, and tumor weights were measured. All operations followed the United Kingdom Animals (Scientific Procedures) Act, 1986. All laboratory animals were carefully handled, and the study was reviewed and approved by the Ethical Review Committee of Experimental Animal Welfare at Hunan University of Chinese Medicine (Approval No. LL2021040705).

Quantitative real-time polymerase chain reaction

Total RNA was extracted from tumor tissues based on the instructions of the RNA Simple Total RNA Kit (Novoprotein Scientific Inc., Suzhou, China), and the RNA concentration was measured by a microplate reader. Then, NovoScript®Plus (Novoprotein Scientific Inc., Suzhou, China) was utilized to reverse-transcribe the RNA into cDNA, which was stored at -20 °C until subsequent use. According to the instructions of NovoStart®SYBR (Novoprotein Scientific Inc., Suzhou, Jiangsu Province, China), 10 µL SuperMix Plus, 7 µL RNase Freewater, 2 µL cDNA, 0.5 µL forward primers and 0.5 reverse primers (Genecreate Inc., Wuhan, Hubei Province, China) were mixed. GAPDH was used as the internal reference. A LightCycler®96 (Roche, Switzerland) was used for quantitative real-time polymerase chain reaction (qRT-PCR). The primer sequences are shown in Table 3. The qRT-PCR conditions were as follows: Predenaturation for 60 s at 95 °C; 45 cycles of denaturation for 20 s at 95 °C, 20 s at 60 °C, and 30 s at 72 °C; 1 cycle of melting for 10 s at 95 °C, 60 s at 65 °C, and 1 s at 97 °C;

and cooling for 30 s at 37 °C. Data were individually normalized to the mean of the relative expression of GAPDH. The fold change was calculated using the $2^{-\Delta\Delta CT}$ method. The experiment was repeated at least three times, and the results that were the most representative are shown.

Western blotting

Tumor tissue specimens were prepared and suspended in ice-cold RIPA lysis buffer (CWBIO Co., Ltd, Beijing, China) and incubated for 20 min. The supernatant was removed after centrifugation. The protein levels of tumor tissues were determined by the BCA assay using a BCA Protein Assay Kit (CWBIO Co., Ltd, Beijing, China). Proteins (50 µg) were mixed with pure water and SDS loading buffer and denatured at 100 °C for 10 min. Proteins were resolved by sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to PVDF membranes (Merck Millipore Ltd., Germany). Then, PVDF membranes were blocked for 1 h in blocking buffer (5% milk in TBST). The membranes were removed and washed with PBST (PBS + 0.1% Tween 20) 3 times, incubated with primary antibodies overnight at 4 °C, washed with TBST 3 times, incubated with Proteintech HRP Goat Anti-Rabbit IgG (H + L) (SA00001-2, diluted 1:6000) for 90 min at room temperature and washed with TBST 3 times. Protein signals were visualized by enhanced chemiluminescence using an eECL western blot kit (CWBIO Co., Ltd, Beijing, China). Membrane images were collected by an Amersham Imager 600 (GE Healthcare, United States) and qualified by Quantity One software. All experiments were repeated three times. Primary antibodies against DR5 (ab199357, diluted 1:1000), FADD (ab108601, diluted 1:1000), cleaved caspase-3 (ab32042, diluted 1:1000), and cytochrome c (Cyto-C) (ab133504, diluted 1:5000) were obtained from Abcam (Cambridge, England). Antibodies against GAPDH (10494-1-AP, diluted 1:5000) and DR4 (24063-1-AP, diluted 1:2000) were obtained from Proteintech Group, Inc. (Chicago, United States). The antibody against cleaved caspase-8 (AF5267, diluted 1:1000) was obtained from Affinity Biosciences, Inc. (Jiangsu, China).

⁷ *Analysis of intracellular reactive oxygen species accumulation*

¹
Reactive oxygen species (ROS) ¹² were measured using a ROS Assay Kit (E004-1-1 Jiancheng, Nanjing, China) according to the manufacturer's instructions. Briefly, the tumor tissue was prepared into a single cell suspension by enzymatic hydrolysis. Three groups of tubes were set up as follows: (1) Negative vehicle tube: cells without any treatment ¹² were resuspended in 0.01 M PBS; (2) positive vehicle tube: cells were resuspended in diluted 2,7-dichlorofluorescein diacetate (DCFH-DA), and an ROS ¹² hydrogen donor (concentration: 30 μ M) was added to induce the cells; and (3) sample ²¹ tube: cells were resuspended in diluted DCFH-DA (cell density: 1×10^6 - 2×10^7 /mL). Then, all cells were incubated for 50 min at 37 °C. After labeling with the probe, the single-cell suspensions were subsequently collected and centrifuged for 5-10 min. The precipitates were collected and washed with 0.01 M PBS 1-2 times. The precipitates were centrifuged and collected again. Finally, the cell precipitates were resuspended in PBS and measured on a fluorescence microplate reader (optimum excitation wavelength 500 nm, optimum emission wavelength ¹² 525 nm). All results are expressed as fluorescence values.

³² *Statistical analysis*

³²
SPSS 25.0 and GraphPad Prism 8.3 software were used for all statistical analyses. Unless otherwise stated, $P < 0.05$ was considered statistically significant. ⁶⁵ All data are expressed as the mean \pm SD. For the cell viability assay, ²⁷ ANOVA was used to compare the means of multiple groups, Student's t test was used for pairwise comparisons, and Fisher's ⁶⁸ exact test was used for ratio comparisons. For tumor weight, ⁵⁸ qRT-PCR and WB, Student's t test was used when normality and ANOVA were satisfied. For ROS assessment, the Mann-Whitney rank-sum test was used due to unsatisfactory normality. Two-way ANOVA was used for tumor volume analysis.

RESULTS

Crude centipede peptides extracted by optimal enzymolysis hydrolysis extraction ratio of crude peptides under different conditions

Based on the extract ratio and protein content, we carried out five single-factor experiments and finally obtained the optimum conditions: the protease was trypsin, the water-centipede ratio was 5:1, the protease dose was 0.1 g (300 U/g, with a centipede-trypsin ratio of 20:1), the enzymolysis temperature was 46 °C, and the enzymolysis time was 4 h. The three most influential factors were enzymolysis time, enzymolysis temperature and trypsin dose. Therefore, an $L_9(3^4)$ orthogonal design was adopted to further investigate the optimal group of three factors on the extract ratio, as shown in Table 4. In conclusion, Group 4 (A2B1C2), Group 2 (A1B2C2), and Group 9 (A3B3C2) had higher extract ratios.

Cytotoxicity of crude peptides under different conditions

The $L_9(3^4)$ orthogonal design was adopted to further investigate the optimal group of the three factors for cytotoxicity in HepG2 cells, as shown in Table 5. A CCK8 assay was used to evaluate HepG2 cell viability. The results indicated that the 9 extracts induced cell death in a concentration-dependent manner ($P < 0.05$). Extracts from Groups 4, 2, and 9 showed stronger cytotoxicity, and Groups 1, 3, 5, 6, 7, and 8 showed moderate cytotoxicity. In the $L_9(3^4)$ orthogonal experiment, Group 2 (5.86%, 0.310 mg/mL), which used the optimal process of the single-factor experiment, showed little difference from the nonoptimal process of Group 9 (5.94% and 0.278 mg/mL). Indeed, Group 2 required less time and a lower trypsin dose. Based on the IC_{50} , extract ratio and suppression ratio shown in Table 5, Group 2 was ultimately determined to have the optimal extraction process. The optimal extraction conditions were as follows: Trypsin at 0.1 g (300 U/g, centipede-trypsin ratio of 20:1), enzymolysis temperature of 46 °C, and enzymolysis time of 4 h.

Cytotoxicity of crude peptides from the optimal enzymolysis hydrolysis and freeze-thawing with liquid nitrogen

HepG2 cells were treated with crude peptides extracted using two extraction methods. The results showed that extracts obtained from the two methods significantly inhibited HepG2 cells in a concentration-dependent manner. Optimal enzymatic hydrolysis (IC_{50} : 0.31 mg/mL) was superior to freeze-thawing with liquid nitrogen (IC_{50} : 3.07 mg/mL), and the cytotoxicity of extracts increased nearly 10 times (Figure 1G).

Purification of scolopentide from crude centipede peptides

We purified and screened crude centipede peptides three times and obtained a low-molecular-weight peptide with the strongest antihepatoma effect, which was designated scolopentide.

The first purification step (Part A): Separation by a sephadex g-25 column

Sephadex G-25 gel chromatography was used, and the chromatogram obtained showed 3 peaks (Figure 1A). Part A samples (sample ID nos. A1 to A3) were then collected based on 3 peaks and assessed by a microplate reader. The protein and polypeptide contents of A1, A2 and A3 were 55, 40 and 20 μ g/mL, indicating that samples A1-3 were mainly composed of protein. Since the separation range of the Sephadex G-25 column was 1-5 kDa, we speculated that A1 contained proteins > 5 kDa and that A2 and A3 were composed of low-molecular-weight polypeptides (1-5 kDa). Subsequently, a CCK8 assay was used to detect the suppressive effects of A1, A2 and A3 on HepG2, Bel-7402 and A549 cells (Figure 1B-D). The CCK8 assay showed the following: (1) A2 showed the most significant inhibition of proliferation of the three tumor cell lines among samples A1-3 ($P < 0.05$); (2) A2 showed the strongest inhibitory effect on HepG2 cells: The IC_{50} values of A2 against the three tumor cell lines were 50.1 mg/mL (HepG2), 132.8 mg/mL (Bel-7402) and 154.5 mg/mL (A549); and (3) A1-3 showed an inhibitory effect on the three tumor cell lines in a concentration-dependent manner.

The second purification step (Part B): Purification by HPLC

In Part B, A2, which had the best antineoplastic effect *in vitro* among samples A1-3, was used. HepG2 cells, which were the most strongly inhibited among the three tumor cell lines, were used. First, tricine-SDS-PAGE^[10] was performed to further purify this sample in Part B, but the peptide separation was not obvious, and the band patterns were light and diffuse. The reason may be that the molecular weight of the sample was too low for separation by tricine-SDS-PAGE. However, satisfactory results were obtained from HPLC. Therefore, HPLC was adopted for further purification. The chromatogram indicated that the sample from Part B was composed of a variety of peptides with very similar and low molecular weights in 6 peaks (Figure 1A; part 1). Subsequently, the Part B sample was collected in 6 parts (sample ID nos. B1 to B6) based on 6 peaks. A CCK8⁴² assay was used to assess the suppressive effect of B1-6 on HepG2 cells cultured for 48 h, which were the tumor cells that were most sensitive to the centipede extracts. The CCK8 assay showed that B1-4 hardly inhibited the proliferation of HepG2 cells, while B5 and B6 inhibited proliferation ($P < 0.05$), with B5 having the strongest effect (Figure 1E).

The third purification step (Part C): Purification by HPLC

In Part C, the B5 sample, which had the best antihepatoma effect *in vitro* among B1-6, was used. HPLC was adopted for purification, and the chromatogram showed 4 peaks (Figure 1A; part 2). Then, the sample from Part C was collected in 4 parts (C1-4) based on the 4 peaks. The CCK8 assay showed that C3 had the strongest inhibitory effect on HepG2 cells cultured for 48 h, while C1, C2, and C4 showed relatively weak inhibition of proliferation (Figure 1F). Finally, we concentrated and purified C3 and designated it scolopentide.

Characterization of scolopentide

The peptide sequence of the extracted scolopentide is RAQNHYCK, and the mass spectrum (Figure 2A) showed that its molecular weight was 1018.997 Da (the highest peak). Mascot was used to determine the specific peptide sequence of scolopentide, and

the final matched sequence was RAQNH⁵⁹YCK, which conformed to the restriction site of the protease. The molecular structure of scolopentide (C₄₂H₆₆N₁₆O₁₂S) is shown in Figure 2A (the asterisk).

Synthesis and detection of scolopentide

Scolopentide was synthesized *in vitro* according to the peptide sequence RAQNH⁵⁹YCK, the length of which is 8 AA. Then, MS was used to detect the molecular weight. The observed molecular weight of synthetic scolopentide was 1018.8 Da (Figure 2B), which was similar to that of the extract (1018.997 Da). HPLC was used to assess the purity. The chromatogram is shown in Figure 2C, and the peak proportion is shown in Figure 2D. The synthetic scolopentide corresponds to peak 3; thus, its purity was inferred to be 98.014% according to the area fraction of peak 3. Consequently, the purity and molecular weight of synthetic scolopentide met the requirements.

Antineoplastic effect of extract scolopentide in vitro

Eca-109, HepG2, and A549 cells were treated with the extracted scolopentide. A CCK8 assay showed that the IC₅₀ values in the three cell types were 76.27 µg/mL (Eca-109), 22.06 µg/mL (HepG2), and 35.13 µg/mL (A549) (Figure 3A). Morphological changes in the three cell lines showed that the A549 and HepG2 cells changed considerably (Figure 3B). Based on the CCK8 assay and morphological changes, the following conclusions were drawn: extracted scolopentide still has antineoplastic effects *in vitro*, which inhibited the proliferation of three tumor cell lines, especially HepG2 cells.

Antihepatoma effect of synthetic scolopentide in vitro and in vivo

We wanted to determine whether synthetic scolopentide can exert an antihepatoma effect. Our experiments suggested that synthetic scolopentide had antihepatoma effects ²² *in vitro* and *in vivo*, which may be related to apoptosis induction. *In vitro*, the CCK8 assay suggested that synthetic scolopentide ² inhibited the proliferation of HepG2 cells in a concentration-dependent manner. The inhibitory effects were the strongest at 12 h,

64 and there was no significant difference between 24 h and 48 h (Figure 3C). Moreover, a Hoechst staining assay was used to observe the morphological changes in HepG2 cells treated with scolopentide (100 µg/mL) (6 h, 12 h, and 24 h). The results showed that cytoplasmic staining and nuclear pyknosis occurred after treatment for 12 h and 24 h, which indicated apoptosis. Apoptosis was most clearly observed at 12 h (IC₅₀: 208.11 µg/mL), consistent with the CCK8 assay (Figure 3D). *In vivo*, the tumor xenograft experiment suggested that the mean tumor volume of the two groups increased gradually, but that of the scolopentide group grew slower. The tumor weight of the scolopentide group was lower than that of the vehicle group (Figure 3E-G).

Cytotoxicity of synthetic scolopentide to L02 cells and HepG2 cells

CCK8 assays and morphological changes were used to compare the cytotoxicity of synthetic scolopentide to liver cancer cells and normal liver cells. After treatment with synthetic scolopentide for 12 h, the CCK8 assay suggested that the cytotoxicity in L02 cells was significantly lower than that in HepG2 cells (100, 150, and 200 µg/mL) (Figure 4A). Morphological changes in the scolopentide group (100 µg/mL) and vehicle group (0 µg/mL) are shown in Figure 4B. Compared to the vehicle group, most HepG2 cells in the scolopentide group died, while some L02 cells survived. The CCK8 assay and morphological changes indicated that synthetic scolopentide was cytotoxic and was significantly stronger in HepG2 cells than in L02 cells.

Molecular docking of scolopentide with DR4 and DR5

56 Scolopentide was more cytotoxic in HepG2 cells than in L02 cells (Figure 4), which is similar to the activity of TRAIL, which selectively initiates apoptosis in cancer cells without significant toxicity to normal cells. Among the known TRAIL receptors, only DR4 and DR5 are able to induce apoptosis^[33]. In addition, Hoechst staining indicated that HepG2 cells underwent apoptosis after treatment with synthetic scolopentide (Figure 3D). Therefore, we hypothesized that scolopentide could induce apoptosis by activating DR4/DR5. The binding free energies of scolopentide to DR4 and DR5 were-

55
10.4 kcal/mol and -7.1 kcal/mol, respectively. Molecular docking suggested that scolopentide tightly bound to DR4 (Figure 5A) and DR5 (Figure 5B), which supported our hypothesis.

Apoptosis of HepG2 cells treated with extracted scolopentide

Flow cytometry suggested that apoptosis occurred in HepG2 cells after treatment with extracted scolopentide *in vitro*. The apoptosis ratios were 5.01% (0 $\mu\text{g/mL}$), 12.13% (10 $\mu\text{g/mL}$), 16.52% (20 $\mu\text{g/mL}$), and 23.2% (40 $\mu\text{g/mL}$), which indicates concentration-dependence (Figure 6A).

The antihepatoma mechanism of scolopentide was determined to be induction of apoptosis by activating the TRAIL pathway

Animal experiments were used to further verify our hypothesis that scolopentide can stimulate the TRAIL pathway to induce apoptosis. The results of qRT-PCR and western blotting suggested that scolopentide activated DR4 (Figure 6B and C) and DR5 (Figure 6H and I), which promoted the expression of FADD (Figure 6E and F), activated the apoptosis promoter caspase-8 (Figure 6E and F) and the executor caspase-3 (Figure 6C and D), and finally induced apoptosis in SMMC-7721 Liver cancer cells. This form of apoptosis appears to be more related to the mitochondria-independent pathway, as Cyto-C and Bcl-2-associated X protein/B-cell lymphoma-2 (Bax/Bcl-2), which are key indicators in the mitochondria-dependent pathway, showed only slight upregulation and insignificant differences (Figure 6I-K). In addition, ROS levels were upregulated (Figure 6L). Cellular fas-associated death domain-like interleukin-1 β converting enzyme inhibitory protein (c-FLIP), an inhibitory protein of caspase-8, was downregulated (Figure 6M), and x-chromosome linked inhibitor-of-apoptosis protein (XIAP), an inhibitory protein of caspase-3, was insignificantly upregulated (Figure 6N).

DISCUSSION

Peptides from centipedes have potential antitumor activity that should not be ignored. We isolated and purified crude centipede peptides 3 times and finally obtained a low molecular weight peptide (scolopentide) with the strongest antihepatoma effect. We further demonstrated that mechanistically, scolopentide induced apoptosis in liver cancer cells by activating DR4 and DR5 and promoting the upregulation of FADD, caspase-8 and caspase-3.

Centipedes, an important part of animal Chinese medicine, have played an important role in clinical treatment^[1]. The remarkable activities and novel structure also give bioactive components of centipedes enormous potential to be exploited and modified as biological drugs^[34]. However, pharmacological studies of centipedes are far behind those of other animals, such as snakes and scorpions^[34,35]. Similar to animal-derived medicines, most centipede extracts are still crude. These crude extracts may contain a large number of histamines, polypeptide toxins and other biologically active substances^[34], leading to allergic reactions such as itching or serious adverse reactions such as acute myocardial infarction, arrhythmia, tissue necrosis, respiratory depression and hemolysis, which hinder the clinical use of centipedes^[11,36]. Small molecular substances from centipedes may not cause adverse reactions and may play a critical role with precise effects.

We tried a variety of ways to obtain a higher extraction rate and more accurate activity. For protein extraction, the optimal conditions were determined: 0.1 g trypsin (300 U/g, centipede-trypsin ratio of 20:1), enzymolysis temperature of 46 °C and enzymolysis time of 4 h. For peptide purification, GC^[37-39] and HPLC^[3,5,6], widely used techniques, were adopted to further purify crude centipede peptides. Mass spectrometry^[5-7], the most widely used technique for peptide characterization, was used to identify scolopentide. The object of extraction and method used here are also noteworthy. The whole body of the centipede was chosen for extraction as recorded in Chinese works^[14], even though most current studies focus on the venom^[12,34]. In addition, enzymatic hydrolysis^[6], a special extract technique, was chosen for the initial extraction to obtain crude centipede peptides. We aimed to simulate the environment of

Chinese medicine being digested by various proteinases in the stomach after oral administration.

Currently, studies on the antihepatoma components of centipedes are rare (Table 1). Thus, we isolated, purified, and screened the crude centipede peptides three times. Finally, we obtained scolopentide (1018.997 Da), which had the strongest antihepatoma activity. Moreover, the peptide sequence RAQNHYCK of scolopentide was matched with the Mascot search engine, which helped to achieve scolopentide synthesis *in vitro*. Mascot also revealed that scolopentide was supposed to be a polypeptide from centipede venom proteins located in Kappa-scoloptoxin(07)-Ssm²a OS. It is worth noting that the tumor types used for screening were tumors in the chest and abdomen, such as liver cancer and esophageal cancer, as recorded in ancient Chinese works^[13]. Liver cancer cells were more sensitive than other tumor cells and were chosen for subsequent mechanistic exploration (Figure 3A).

Antimicrobial activity is the most common pharmacological characteristic known among isolated centipede peptides at present (Table 1). Centipede peptides mainly destroy the integrity of cell membranes, which leads to the death of microbes^[40-42]. Inducing apoptosis is the second major mechanism by which centipede peptides induce resistance to microorganisms (Table 1). Two antimicrobial centipede peptides, scolopendin and scolopendin 1, were obtained by RNA sequencing^[41,43]. They have been shown to cause ROS accumulation in *Candida albicans*, which leads to mitochondrial depolarization, thus releasing Cyto-C into the cytoplasm from mitochondria and resulting in an increase in Ca²⁺ in the cytoplasm and mitochondria, ultimately inducing caspase-related apoptosis^[43,44]. Some centipede peptides can also block the cell cycle and affect the expression of genes related to DNA replication and repair^[45]. Additionally, some scholars have shown that a few antimicrobial centipede peptides also have antitumor activity^[17,26]. Distinct from the antimicrobial mechanism, the antitumor activities are mainly related to apoptosis, especially the mitochondrial-dependent pathway^[3,4,18]. To date, there have been no studies related to the mitochondrial-independent pathway of centipede effects (Table 1).

DR4 and DR5 are involved in the mitochondria-independent apoptosis pathway, which can receive extracellular death signals and induce intracellular apoptosis^[46]. FADD receives apoptosis signals transmitted from DR4/DR5 and forms the death-inducing signaling complex TRAIL-DR4/DR5-FADD-procaspase. Then, caspase-8 is activated to initiate two apoptotic pathways: (1) ²⁴ The mitochondria-independent pathway, in which ⁴¹ caspase-8 directly activates caspase-3; and (2) the mitochondria-dependent pathway, in which caspase-8 cleaves Bid, resulting in tBid that triggers the release of Cyto-C. Then, Cyto-C successively activates the promoter caspase-9 and the executor caspase-3. ROS accumulation can cause mitochondrial permeability transition pore opening, also leading to Cyto-C release. Apoptosis is the common end of these two pathways (Figure 7)^[28,47-49].

We found that scolopentide may activate DR4 and DR5 and induce apoptosis. Flow cytometry suggested that apoptosis occurred ²⁵ in HepG2 cells after treatment with extracted scolopentide *in vitro* (Figure 6A). Hoechst staining showed the occurrence of apoptosis ²⁵ in HepG2 cells after treatment with synthetic scolopentide (Figure 3D). The CCK8 assay showed that scolopentide was significantly more cytotoxic ²⁵ in HepG2 cells than in L02 cells (Figure 4), which was similar to the effect of TRAIL, which selectively induced tumor cell apoptosis. Molecular docking also indicated that scolopentide binds well to TRAIL receptors (DR4 and DR5) (Figure 5). In our previous study, crude centipede peptides were also found to inhibit STAT3 protein phosphorylation^[19], while some studies have suggested that dovitinib and sorafenib could overcome TRAIL resistance in HCC by inhibiting STAT3^[50,51]. Consequently, animal experiments were used to further verify our hypothesis that scolopentide can stimulate the TRAIL pathway to induce apoptosis. In animal experiments, after treatment with synthetic scolopentide, the expression of DR4 and DR5, which are upstream ² of the TRAIL pathway, was clearly upregulated, especially the expression of DR4. The expression of FADD, caspase-8, and caspase-3, which are downstream of the TRAIL pathway, was also upregulated (Figure 6B-I). The animal experiments further confirmed the viewpoint that scolopentide induces apoptosis by activating DR4 and DR5.

This process may be more related to the mitochondria-independent pathway, as Cyto-C and Bax/Bcl-2 showed only slight upregulation and insignificant differences (Figure 6I-K), which are key indicators in the mitochondria-dependent pathway^[52]. ROS were significantly upregulated but could not cause Cyto-C release (Figure 6I, J, and L). Thus, we hypothesized that ROS mainly acted as signal transduction factors of the TRAIL pathway rather than the mitochondrial-dependent pathway^[49]. Additionally, c-FLIP was inhibited, but XIAP was slightly upregulated (Figure 6M and N). A negative feedback mechanism of apoptosis may contribute to preventing excessive apoptosis in the body. Overall, our findings suggested that scolopentide may induce apoptosis in tumor cells, especially gastrointestinal tumors, by activating DR4 and DR5 and leading to the caspase cascade.

CONCLUSION

In summary, we obtained a small molecule polypeptide from *Scolopendra subspinipes mutilans* L. Koch with the strongest antitumor activity, especially for liver tumors. Mechanistically, scolopentide may induce tumor cell apoptosis by activating DR4 and DR5. Scolopentide is considered ² to be a promising drug candidate for cancer treatment, especially treatment of gastrointestinal tumors. Our studies are also expected to provide a reference for the extraction, purification and characterization of effective components in animal-derived medicines. Both extracted and synthesized scolopentide had antihepatoma activity. When HepG2 cells were cultured with two scolopentides for 48 h, a CCK8 assay showed that the IC₅₀ values were 22.06 µg/mL (extracted scolopentide) and 237.726 µg/mL (synthesized scolopentide), which indicates that the antihepatoma activity of synthesized scolopentide was weaker than that of the extracted scolopentide. This may be due to the lack of dimensional folding configurations in the synthesized peptide during synthesis. Additionally, the TRAIL pathway was activated but decreased incrementally. Key areas of future research include investigating methods of modifying the spatial architecture of synthetic scolopentide, fully activating the TRAIL pathway and improving its antihepatoma activity.

Figure Legends

Figure 1 Purification of scolopentide from crude centipede peptides. A: After the first purification, the Sephadex G-25 chromatogram showed 3 peaks, and A2 was chosen for further isolation; after the second purification, the high-performance liquid chromatography (HPLC) chromatogram showed 6 parts, and B5 was chosen for further purification (part 1); after the third purification, the HPLC chromatogram showed 4 parts, and C3 was chosen for further purification (part 2); B-D: Relative cell viability of HepG2 (B), A549 (C), and Bel7402 (D) cells treated with A1-3 at different concentrations (mg/mL). A2 showed stronger suppression than A1 and A3; E: The CCK8 assay showed that B5 (50 µg/mL) had the strongest suppression of HepG2 cells among B1-6; F: The CCK8 assay showed that C3 (20 µg/mL) had the strongest suppression of HepG2 cells among C1-4; G: The CCK8 assay showed the cytotoxicity of extracts from two methods, and optimal enzymatic hydrolysis was superior to freeze-thawing with liquid nitrogen. ^a*P* < 0.05, ^b*P* < 0.01, ^c*P* < 0.001, and ^d*P* < 0.0001.

Figure 2 Characterization of extracted scolopentide and detection of synthetic scolopentide. A: Mass spectrum of extracted scolopentide; the highest peak indicates the active peptide (scolopentide). The observed molecular weight was 1018.997 Da; the asterisk "*" means molecular structure of scolopentide; B: Mass spectrum of synthetic scolopentide. The observed molecular weight was 1018.8 Da; C and D: HPLC chromatogram of synthetic scolopentide (C); the highest peak (peak 3) indicates the active peptide, and the area % of peak 3 indicates the purity of synthetic scolopentide (98.014%) (D).

Figure 3 Antihepatoma effect of scolopentide. A: The CCK8 assay showed the suppression ratio of Eca-109, HepG2, and A549 cells treated with extracted scolopentide at different concentrations; B: Morphological changes in Eca-109, HepG2, and A549 cells under a light microscope ($\times 40$); after treatment with extracted scolopentide, three cells were morphologically changed, especially HepG2 cells; C: The CCK8 assay showed the suppression ratio of HepG2 cells treated with synthetic scolopentide at different times (6 h, 12 h, 24 h, and 48 h) and different concentrations (50 $\mu\text{g/mL}$, 100 $\mu\text{g/mL}$, 150 $\mu\text{g/mL}$, and 200 $\mu\text{g/mL}$); D: Hoechst 33342 staining ($\times 400$) of HepG2 cells. After treatment with synthetic scolopentide for 12 h and 24 h, cytoplasmic highlight staining and nuclear pyknosis occurred; E-G: Tumor volume and weight of the scolopentide group (synthetic scolopentide 500 mg/kg/d) and vehicle group (constant volume of normal saline). $n = 6$, $^bP < 0.01$, $^cP < 0.001$.

Figure 4 Cytotoxicity of synthetic scolopentide to L02 cells and HepG2 cells. A: The CCK8 assay showed that cytotoxicity to L02 cells was significantly lower than that to HepG2 cells after treatment with synthetic scolopentide for 12 h (100 $\mu\text{g/mL}$, 150 $\mu\text{g/mL}$, and 200 $\mu\text{g/mL}$); B: Morphological changes in HepG2 and L02 cells under a light microscope after treatment with synthetic scolopentide for 12 h ($\times 100$). Compared to cells in the vehicle group (0 $\mu\text{g/mL}$), most HepG2 cells in the scolopentide group (100 $\mu\text{g/mL}$) died, while some L02 cells survived. $^dP < 0.0001$.

Figure 5 Molecular docking of scolopentide and death receptor 4 and death receptor 5. A: Stereograms of the molecular docking of scolopentide and death receptor 4. Hydrogen bonds formed with amino acid residues of the receptor include LYS145, ALA201, PRO219, and CYS190; B: Stereograms of molecular docking of scolopentide and death receptor 5. Hydrogen bonds formed with amino acid residues of the receptor are TRP74, SER73, ER149, and TYR148.

Figure 6 Antihepatoma mechanism of scolopentide. A: Flow cytometry suggested that apoptosis occurred in HepG2 cells after treatment with extracted scolopentide *in vitro*. B-I: Quantitative real-time polymerase chain reaction (qRT-PCR) and western blotting showed that the expression of DR4 (B and C), DR5 (H and I), FADD (E and F), caspase-8 (F and G), and caspase-3 (C and D) was significantly upregulated in the scolopentide group. DR4 was the most considerably upregulated; I-K: qRT-PCR and western blotting showed that the expression of Cyto-C (I and J) and Bax/Bcl-2 (K) was not upregulated, which are key indicators of mitochondria dependence; L: Tumor ROS levels in the scolopentide group were higher than those in the vehicle group; M: c-FLIP, an inhibitory protein of caspase-8, was significantly downregulated in qRT-PCR; N: XIAP, an inhibitory protein of caspase-3, was insignificantly upregulated in qRT-PCR. DR4: Death receptor 4; DR5: Death receptor 5; FADD: Fas-associated death domain protein. $n = 4$ per group in qRT-PCR, $n = 3$ per group in Western blotting, $n = 4$ per group in ROS; $^aP < 0.05$, $^bP < 0.01$.

Figure 7 Activation of the mitochondria-independent and mitochondria-dependent apoptosis pathways by tumor necrosis factor-related apoptosis-inducing ligand. The mitochondria-independent pathway (engaged through death receptors, activated caspase family directly) and mitochondria-dependent pathway (triggered through the Bcl-2 gene superfamily) are represented. TRAIL: Tumor necrosis factor-related apoptosis-inducing ligand; DR4: Death receptor 4; DR5: Death receptor 5; FADD: Fas-associated death domain protein; Cyto-C: Cytochrome c; Bax: Bcl-2-associated X protein; Bcl-2: B-cell lymphoma-2; ROS: Reactive oxygen species; c-FLIP: Cellular Fas associated death domain-like interleukin-1 β converting enzyme inhibitory protein; XIAP: X-chromosome linked inhibitor-of-apoptosis protein.

Table 1 Small molecular extracts of centipede

Name	Extract methods	Function	Ref.
Scolopendrin I	Cation-exchange chromatography and two steps RP-HPLC	Antimicrobial (G ⁺ , G ⁻ , and fungi)	[53]
Scolopin 1	(1) Sephadex Chromatography; (2) RP-HPLC; and (3) characterized by MALDI-TOF MS	Antimicrobial (G ⁺ , G ⁻ , fungi, drug-resistant bacteria)	[37]
Scolopin 2	(1) Sephadex Chromatography; (2) RP-HPLC; and (3) Scolopin2-NH2 was amidation modification from scolopin 2	(1) Antimicrobial (G ⁺ , G ⁻ , fungi, and drug-resistant bacteria); and (2) inhibit proliferation of cervical cancer HeLa cells	[16,37,45]
LBLP	Transcriptomic analysis, peptide synthesis	Anti-fungi	[54]
Scolopendrasin II, V, III, X	Next-generation sequencing	(1) Antimicrobial (G ⁺ , G ⁻ , drug-resistant bacteria, fungi); and (2) regulating immune system to antimicrobial indirectly	[40,55-57]
Scolopendrasin IX	Next-generation sequencing	(1) Regulating immune system to antimicrobial	[58]

		indirectly; and (2) treating autoimmune arthritis
Scolopendrasin VII	Next-generation sequencing	(1) Antimicrobial [17,59,60] (G ⁺ , G ⁻ , drug- resistant bacteria, and fungi); (2) regulating immune system to antimicrobial indirectly; and (3) inhibiting leukemia U937 and jurkat cells
Scolopendin, 1, 2	RNA sequencing	Antimicrobial (G ⁺ , [41-44] G ⁻ , drug-resistant bacteria, and fungi)
SerGln-Leu (SQL)	(1) Ultrafiltration, Sephadex G-50 column, Source 15Q anion exchange Column; (2) RP- HPLC C18 column; and (3) characterized by ESI- MS	Anticoagulation and [7] antithrombotic
Thr-Asn-Gly- Tyr-Thr, (TNGYT)	(1) Sephadex G-50 column; (2) RP-HPLC C18 column; and (3) Characterized by ESI-MS	Anticoagulation and [38] antithrombotic
SsTx-R12A	Red-headed centipede's venom cDNA arsenal	Treating [61] autoimmune

			diseases mediated by T cells	
SsmTX-I	Red-headed centipede's venom cDNA arsenal	Analgesic		[62]
SsTx	Sephadex G-50 Gel Chromatography and RP- HPLC	Blocking potassium channels leads to cardiovascular, respiratory, muscular and nervous system damage	KCNQ	[8]
Ssm6a	Sephadex Gel Chromatography and RP- HPLC	Analgesic		[39]
Haemolymph and tissue extracts	(1) Enzymatic hydrolysis; (2) HPLC; and (3) characterized by MS	Antimicrobial (G ⁺ , G ⁻ , fungi, viruses, and parasites)		[6]
Lacrain	RP-HPLC and characterized by MS	Antibacterial (G ⁻)		[63]
Pinipessin	(1) Organic solvent extraction; (2) HPLC Jupiter® C18 column; and (3) Characterized by MS	Antimicrobial		[5]
AECS	Organic solvent extraction (ethanol)	Inducing apoptosis of human epidermal carcinoma A431 cells and human melanoma HEK293/EGFR, A375 cells		[4,18]

Two isoquinoline alkaloids	(1) Organic solvent extraction (ethanol); (2) sephadex G-25 Gel chromatography; and (3) HPLC	Inducing apoptosis of brain glioma U87 cells [3]
Centipede water extract, alcohol extract	(1) Water extraction; freeze-thawing with liquid nitrogen, ultrasonic extraction; and (3) organic solvent extraction (ethanol)	No tumour [2] inhibition in S180 and H22 Transplanted tumour mice
CE, CA	Organic solvent extraction	Inhibiting cervical tumours <i>in vivo</i> [64]
Centipede chloroform extract	Organic solvent extraction (hexane, chloroform, ethanol)	Reducing central nervous system symptoms of Alzheimer's disease [65]
Scom 5	(1) Superdex 75 gel filtration; (2) RESOURCE S ion chromatography; and (3) characterized by LC-MSMS, MALDI-TOF/TOF and protein sequencing techniques	Human allergens [11]

29

HPLC: High performance liquid chromatography; RP-HPLC: Reversed-phase high-performance liquid chromatography; MS: Mass spectrum.

Table 2 L₉(3⁴) levels of orthogonal experimental design

Level	Trypsin (g)	Time (h)	Temperature (°C)
1	0.1	3	37
2	0.2	4	46
3	0.3	5	55

28

Table 3 Primer sequences used for quantitative real-time polymerase chain reaction

Gene	Sequence (5' to 3')	
GAPDH	F:	R:
	AGAAGGCTGGGGCTCATTT	AGGGGCCATCCACAGTCTT
	G	C
DR4	F:	R:
	CCAGGCAGCATTGAAGTCA	GTTTCCAGCATCACCAGGG
	G	T
DR5	F:	R:
	CCCTGGAGTGACATCGAAT	CAGCCACAATCAAGACTAC
	G	GG
FADD	F:	R:
	TGCGGGAGTCACTGAGAAT	GGAGGTAGATGCGTCTGAG
	C	TTC
Caspase-8	F:	R:
	GAGAAGCAGCAGCCTTGA	GACAGTATCCCCGAGGTTT
	AG	G
Caspase-3	F:	R:
	CCCTCCTCAGCATCTTATCC	TGGTACAGTCAGAGCCAAC
		CT
Cyto-C	F:	R:
	CACAAGACTGGGCCAAAT	CCAGGGATGTACTTCTTGG
	CTC	GA

XIAP	F: CTATGCTCACCTAACCCCA AG	R: TTCTGACCAGGCACGATCA C
c-FLIP	F: GGAACCCTCACCTTGTTTC G	R: CTCCTTGCTTATCTTGCCTC G
Bcl-2	F: CCTTCTTTGAGTTCGGTGGG	R: CGGTTTCAGGTACTCAGTCAT CC
Bax	F: CACCAGCTCTGAGCAGATC A	R: TGTTACTGTCCAGTTCGTCC C

¹³ DR4: Death receptor 4; DR5: Death receptor 5; FADD: Fas-associated death domain protein; Cyto-C: Cytochrome C; ¹¹ XIAP: X-chromosome linked inhibitor-of-apoptosis protein; c-FLIP: Cellular Fas associated death domain-like interleukin-1 β converting enzyme inhibitory protein; ⁶ Bcl-2: B-cell lymphoma-2; Bax: Bcl-2-associated X protein.

Table 4 Extract ratio analysis of L₉ (3⁴) orthogonal experimental design

Run	A (mg, trypsin)	B (min, time)	C (°C, temperature)	D (blank)	Extract ratio (%)
1	1	1	1	1	4.55
2	1	2	2	2	5.86
3	1	3	3	3	4.32
4	2	1	2	3	5.63
5	2	2	3	1	4.21
6	2	3	1	2	3.98
7	3	1	3	2	3.34
8	3	2	1	3	3.68
9	3	3	2	1	5.94
K1	14.73	13.52	12.21	14.70	-
K2	13.82	16.01	14.83	13.18	-
K3	12.96	11.64	11.87	13.63	-
R	1.77	4.37	2.96	1.52	-

³¹ K_{ij} parameters in a first level of the first i-j-average quality index; *I* = 1, 2, 3, representative of 3 levels; *j* = 1, 2, 3; representing three factors.

Table 5 Inhibitory effects of crude centipede peptides on HepG2 cells in different conditions

Concentr ation (mg/mL)	Suppression ratio (%)								
	Group	Grou	Grou	Grou	Grou	Grou	Grou	Grou	Grou
	1	p 2	p 3	p 4	p 5	p 6	p 7	p 8	p 9
0.039	12.510	6.880	12.07	5.394	3.389	3.213	0.570	1.641	7.094
		0							
0.078	22.270	11.24	13.06	16.43	12.89	4.709	2.280	4.540	9.883
		0	0	0	0				
0.156	45.770	36.43	45.68	52.29	18.61	16.62	26.11	11.54	33.17
		0	0	0	0	0	0	0	0
0.3125	85.510	85.04	88.54	94.36	68.11	63.66	58.50	64.17	76.61
		0	0	0	0	0	0	0	0
0.625	89.090	86.06	92.44	94.52	94.17	94.96	83.20	95.18	94.57
		0	0	0	0		0	0	0
1.250	91.840	87.49	93.05	94.77	96.78	95.73	94.66	95.30	96.00
		0	0	0	0	0	0	0	0
2.500	92.070	89.47	93.43	94.69	96.94	96.67	95.54	95.67	96.92
		0	0	0	0	0	0	0	0
5.000	92.750	92.00	95.80	96.68	97.72	97.78	95.88	95.79	97.65
		0	0	0	0	0	0	0	0
IC ₅₀ (mg/mL)	0.457	0.310	0.480	0.360	0.584	0.681	0.812	0.757	0.278
Extract ratio (%)	4.550	5.860	4.320	5.630	4.210	3.980	3.340	3.680	5.940

18%

SIMILARITY INDEX

PRIMARY SOURCES

1	www.frontiersin.org Internet	161 words — 2%
2	www.science.gov Internet	145 words — 2%
3	www.mdpi.com Internet	113 words — 1%
4	www.nature.com Internet	80 words — 1%
5	mdpi-res.com Internet	63 words — 1%
6	www.spandidos-publications.com Internet	54 words — 1%
7	link.springer.com Internet	51 words — 1%
8	www.ncbi.nlm.nih.gov Internet	49 words — 1%
9	Ning Dimin, Deng Zhe, Wu Yongrong, Mei Si, Teng Yongjie, Zhou Qing, Tian Xuefei. "Niu Huang (Bovis Calculus)-Shexiang (Moschus) combination induces apoptosis	48 words — 1%

and inhibits proliferation in hepatocellular carcinoma via
PI3K/AKT/mTOR pathway", Digital Chinese Medicine, 2022

Crossref

10 f6publishing.blob.core.windows.net 38 words — < 1 %
Internet

11 tessera.spandidos-publications.com 38 words — < 1 %
Internet

12 [Xiumei Xu, Huiqing Zhang, Lin Li, Runan Yang, Guilin Li, Shuangmei Liu, Günther Schmalzing, Hong Nie, Shangdong Liang. "Study of the Involvement of the P2Y12 Receptor in Chronic Itching in Type 2 Diabetes Mellitus", Molecular Neurobiology, 2022](#) 36 words — < 1 %
Crossref

13 exp-oncology.com.ua 28 words — < 1 %
Internet

14 www.hindawi.com 28 words — < 1 %
Internet

15 pericles.pericles-prod.literatumonline.com 27 words — < 1 %
Internet

16 www.dovepress.com 26 words — < 1 %
Internet

17 www.wjgnet.com 25 words — < 1 %
Internet

18 [Sahdeo Prasad, Ji Hye Kim, Subash C. Gupta, Bharat B. Aggarwal. "Targeting death receptors for TRAIL by agents designed by Mother Nature", Trends in Pharmacological Sciences, 2014](#) 23 words — < 1 %
Crossref

19 Thayyath S. Anirudhan, Sheeba Alexander. "Synthesis and characterization of vinyl-functionalized multiwalled carbon nanotubes based molecular imprinted polymer for the separation of chlorpyrifos from aqueous solutions", Journal of Chemical Technology & Biotechnology, 2013

23 words — < 1%

Crossref

20 insights.ovid.com

Internet

22 words — < 1%

21 repository.charlotte.edu

Internet

21 words — < 1%

22 www.researchgate.net

Internet

21 words — < 1%

23 www.scribd.com

Internet

20 words — < 1%

24 Jin Na Shin, Young Woo Seo, Moonil Kim, Sun-Young Park, Mi-Ja Lee, Byung Rai Lee, Jae-Wook Oh, Dai-Wu Seol, Tae-Hyoung Kim. " Cisplatin Inactivation of Caspases Inhibits Death Ligand-induced Cell Death and Fulminant Liver Damage in Mice ", Journal of Biological Chemistry, 2005

18 words — < 1%

Crossref

25 Libin Wei. "Different apoptotic effects of wogonin via induction of H₂O₂ generation and Ca²⁺ overload in malignant hepatoma and normal hepatic cells", Journal of Cellular Biochemistry, 12/15/2010

18 words — < 1%

Crossref

26 Xiao Yuan, Wang Xiang, Kailin Yang, Huiping Liu, Guomin Zhang, Qi He, Jiapeng Fan. "Systematic Pharmacology-Based Strategy to Explore the Molecular

16 words — < 1%

Network Mechanism of Modified Taohong Siwu Decoction in the Treatment of Premature Ovarian Failure", Evidence-Based Complementary and Alternative Medicine, 2022

Crossref

-
- | | | |
|-----------|--|-----------------|
| 27 | stemcellres.biomedcentral.com
<small>Internet</small> | 16 words — < 1% |
|-----------|--|-----------------|
-
- | | | |
|-----------|---|-----------------|
| 28 | www.e-sciencecentral.org
<small>Internet</small> | 16 words — < 1% |
|-----------|---|-----------------|
-
- | | | |
|-----------|--|-----------------|
| 29 | dspace.vutbr.cz
<small>Internet</small> | 15 words — < 1% |
|-----------|--|-----------------|
-
- | | | |
|-----------|---|-----------------|
| 30 | Hui Yang, Qun Ren, Zhaojie Zhang. "Chromosome or chromatin condensation leads to meiosis or apoptosis in stationary yeast () cells ", FEMS Yeast Research, 2006
<small>Crossref</small> | 14 words — < 1% |
|-----------|---|-----------------|
-
- | | | |
|-----------|---|-----------------|
| 31 | Sun, Bao Shou, Yi Min Deng, Bo Qin Gu, and Xiao Diao Huang. "Optimization of Process Parameters for Warpage Minimization on Injection Molding Thin-Walled Parts", Applied Mechanics and Materials, 2011.
<small>Crossref</small> | 14 words — < 1% |
|-----------|---|-----------------|
-
- | | | |
|-----------|--|-----------------|
| 32 | jeccr.biomedcentral.com
<small>Internet</small> | 14 words — < 1% |
|-----------|--|-----------------|
-
- | | | |
|-----------|---|-----------------|
| 33 | koreascience.kr
<small>Internet</small> | 13 words — < 1% |
|-----------|---|-----------------|
-
- | | | |
|-----------|--|-----------------|
| 34 | onlinelibrary.wiley.com
<small>Internet</small> | 13 words — < 1% |
|-----------|--|-----------------|
-
- | | | |
|-----------|--|-----------------|
| 35 | researchmap.jp
<small>Internet</small> | 13 words — < 1% |
|-----------|--|-----------------|
-

36	search.bvsalud.org Internet	13 words — < 1%
37	pubs.rsc.org Internet	11 words — < 1%
38	watermark.silverchair.com Internet	11 words — < 1%
39	www.tandfonline.com Internet	11 words — < 1%
40	www.vet.cmu.ac.th Internet	11 words — < 1%
41	Erica J. Peterson, Vijay R. Menon, Laura Gatti, Ralph Kipping et al. "Nucleolar Targeting by Platinum: p53-Independent Apoptosis Follows rRNA Inhibition, Cell-Cycle Arrest, and DNA Compaction", Molecular Pharmaceutics, 2014 Crossref	10 words — < 1%
42	Guo, Qinglong. "VI-16, a newly synthesized flavonoid, induces apoptosis through the mitochondrial pathway in human hepatoma cells", Oncology Reports, 2011. Crossref	10 words — < 1%
43	N. Hoti. "Triphenyl Tin Benzimidazolethiol, a Novel Antitumor Agent, Induces Mitochondrial-Mediated Apoptosis in Human Cervical Cancer Cells via Suppression of HPV-18 Encoded E6", Journal of Biochemistry, 10/01/2003 Crossref	10 words — < 1%
44	datospdf.com Internet	10 words — < 1%

45	ijppr.humanjournals.com Internet	10 words — < 1 %
46	scholarbank.nus.edu.sg Internet	10 words — < 1 %
47	spandidos-publications.com Internet	10 words — < 1 %
48	www.alliedacademies.org Internet	10 words — < 1 %
49	www.pierre-fabre.co.za Internet	10 words — < 1 %
50	Hyun-Ju Kwon, Eun-Woo Lee, Yun-Kyung Hong, Hee-Jung Yun, Byung-Woo Kim. "Widdrol from <i>Juniperus chinensis</i> induces apoptosis in human colon adenocarcinoma HT29 Cells", Biotechnology and Bioprocess Engineering, 2010 Crossref	9 words — < 1 %
51	Ji - Min Dai, Xue - Qin Zhang, Jia - Jia Zhang, Wei - Jie Yang, Xiang - Min Yang, Huijie Bian, Zhi - Nan Chen. "Blockade of mIL - 6R alleviated lipopolysaccharide - induced systemic inflammatory response syndrome by suppressing NF - κB - mediated Ccl2 expression and inflammasome activation", MedComm, 2022 Crossref	9 words — < 1 %
52	Shao, Jingwei, Yongchao Dai, Wenna Zhao, Jingjing Xie, Jinping Xue, Jianhui Ye, and Lee Jia. "Intracellular distribution and mechanisms of actions of photosensitizer Zinc(II)-phthalocyanine solubilized in Cremophor EL against human hepatocellular carcinoma HepG2 cells", Cancer Letters, 2012.	9 words — < 1 %

53 Xing Chen, Hongbing Zhang, Wenqing Du, Liying Qian, Ying Xu, Yange Huang, Qingping Xiong, Hailun Li, Jun Yuan. "Comparison of different extraction methods for polysaccharides from *Crataegus pinnatifida* Bunge", International Journal of Biological Macromolecules, 2019
9 words — < 1%
Crossref

54 cmjournal.biomedcentral.com
Internet
9 words — < 1%

55 escholarship.org
Internet
9 words — < 1%

56 veterinaryresearch.biomedcentral.com
Internet
9 words — < 1%

57 www.prf.upol.cz
Internet
9 words — < 1%

58 www.scielo.br
Internet
9 words — < 1%

59 Adam A. A. Sulaiman, Naike Casagrande, Cinzia Borghese, Giuseppe Corona et al. "Design, Synthesis, and Preclinical Activity in Ovarian Cancer Models of New Phosphanegold(I)-N-heterocyclic Carbene Complexes", Journal of Medicinal Chemistry, 2022
8 words — < 1%
Crossref

60 Shengwen Cheng, Senrui Liu, Bowen Chen, Chengcheng Du, Pengcheng Xiao, Xuefeng Luo, Li Wei, Yiting Lei, Chen Zhao, Wei Huang. "Psoralidin inhibits osteosarcoma function by downregulating ITGB1 expression via

61 Takuo Kono, Masahiko Watanabe, Kotaro Koyama, Taketoshi Kishimoto, Shoji Fukushima, Takashi Sugimura, Keiji Wakabayashi. "Cytotoxic activity of pierisin, from the cabbage butterfly, *Pieris rapae*, in various human cancer cell lines", Cancer Letters, 1999

Crossref

62 Xing Lin, Yan Li, Xiaoling Zhang, Yuanyuan Wei, Shujuan Wen, Zhongpeng Lu, Quanfang Huang, Jinbin Wei. " Tormentic acid inhibits hepatic stellate cells activation via blocking /Akt/ and signalling pathways ", Cell Biochemistry and Function, 2020

Crossref

63 atvb.ahajournals.org 8 words — < 1 %

Internet

64 biblio.unibe.ch 8 words — < 1 %

Internet

65 bsdwebstorage.blob.core.windows.net 8 words — < 1 %

Internet

66 cancerci.biomedcentral.com 8 words — < 1 %

Internet

67 cellandbioscience.biomedcentral.com 8 words — < 1 %

Internet

68 rep.bioscientifica.com 8 words — < 1 %

Internet

69 www.ijbs.com

Internet

8 words — < 1%

70 www.researchsquare.com
Internet

8 words — < 1%

71 Bremer, E.. "Targeted induction of apoptosis for cancer therapy: current progress and prospects", Trends in Molecular Medicine, 200608
Crossref

7 words — < 1%

72 Hua LIU, Wei CHEN, Rui-Ting LI. "Research on complex enzyme system to extract Hibiscus Sabdariffa L reducing sugar", Food Science and Technology, 2021
Crossref

7 words — < 1%

73 Jun Long, Chenyang Gu, Qiankun Zhang, Jiale Liu et al. "Extracellular vesicles from medicated plasma of Buyang Huanwu decoction-preconditioned neural stem cells accelerate neurological recovery following ischemic stroke", Frontiers in Cell and Developmental Biology, 2023
Crossref

6 words — < 1%

EXCLUDE QUOTES OFF
EXCLUDE BIBLIOGRAPHY OFF

EXCLUDE SOURCES OFF
EXCLUDE MATCHES OFF