Structural Identification and Coagulation Effect of *Flammulina Velutipes* Polysaccharides

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Abstract: Two polysaccharides were isolated successfully from *Flammulina velutipes* and identified as CHFVP-1 (24.44 kDa) and CHFVP-2 (1497 kDa). Based on the results of Fourier transform-infrared spectroscopy (FT-IR), gas chromatography (GC), gas chromatography–mass spectrometry (GC–MS), and nuclear magnetic resonance (NMR) spectroscopy regarding the structure of CHFVP-1 and CHFVP-2, CHFVP-1 was constructed with the backbone of (→6)-α-D-Galp-(1→ and the branch of Galp by an (→3,6)-α-D-Manp-(1→ attached with T-β-D-GlcP or T-α-L-FucP side chains. Meanwhile, the CHFVP-2 was a glucan with the construction of (→6)-β-D-GlcP-(1→ and T-β-D-GlcP. Moreover, the coagulant activity in vitro of CHFVP-1 and CHFVP-2 was evaluated, and the results showed that CHFVP-1 exerts procoagulant activity by shortening the activated partial thromboplastin time (APTT) and thrombin time (TT), while CHFVP-2 did not reveal a definite coagulant activity. The finding would benefit the further application of *F. velutipes* in the field of medicine.

Keywords: polysaccharides; *flammulina velutipes*; identification; coagulation effect

1. Introduction

*Flammulina velutipes*, one of the edible mushrooms of Flammulina, is popular in Asia, North America, and Europe, and its production and consumption have increased rapidly due to its rich nutritional values and industrialized cultivation [1]. According to the finding of many researchers, polysaccharides, well-known key functional compounds in *F. velutipes* fruit bodies, have a variety of bioactivities, such as immunomodulatory [2], antitumor [3], antioxidant [4], anti-proliferation [5], anti-Hepatitis B Virus(HBV) activity [6], hepatocyte protection [7], renoprotective effects [8], and so on.

In our previous study, we found that several polysaccharides have coagulation or anticoagulation activities [9–12]. Up to date, there is no research on the coagulation effect of the *F. velutipes* polysaccharides. In this study, the crude polysaccharides, CHFVP-1 and CHFVP-2, were extracted, isolated, and purified from *F. velutipes*. Then, the relative molecular weight, physical and chemical properties, monosaccharide composition, residue segments, linkages, and the possible primary structure of the two *F. velutipes* polysaccharides were analyzed with corresponding methods. Meanwhile, the coagulation activity of the purified *F. velutipes* polysaccharides was studied in vitro.
2. Materials and Methods

2.1. Materials

*F. velutipes* sample were dried, ground, and collected from Henan Lonfon Industrial Corporation Limited (Puyang, China) in 2016. Male New Zealand rabbits, 2–3 kg, fed with adequate food and water for 12 weeks (temperature 20 ± 5 °C, humidity 45–65%) were used. Breviscapine was purchased from Yunnan Bio Valley Pharmaceutical Co., Ltd., Kunming 650503, China. Enzyme-linked immunosorbent assay (ELISA) kits of prothrombin time (PT, 20180502), activated partial thromboplastin time (APTT, 20180515T), and thrombin time (TT, 20180416) were all purchased from Shenzhen Dulei Biological science Co., Ltd., Shenzhen 518067, China. All other materials were purchased with the grade of the corresponding methods.

2.2. Extraction and Purification of Polysaccharides from *F. Velutipes*

The polysaccharides of *F. velutipes* were extracted and purified by the modified method of Zhang [11] and Wang [12]. During the method, lipids, pigments, and small molecular compounds were all washed out from *F. velutipes* firstly. Then, the residue was extracted with distilled water three times. The supernatant was filtered, combined, and concentrated. After that, the crude polysaccharides were precipitated by adding ethanol and washed successively with anhydrous ethanol, acetone, and petroleum ether. Then, Sevag reagent and dialysis bags were used to deproteinize the crude polysaccharides and remove some salts. Finally, crude *F. velutipes* polysaccharides (FVP) were obtained after the freeze-dried processing.

First, we dissolved the crude FVP in distilled water and separated it into fractions by a Diethylaminoethyl cellulose-52 column (DEAE-52 cellulose column). Then, the fractions were eluted by solvent gradient chromatography with 0, 0.05, 0.1, 0.2, and 0.5 M NaCl (1.3 mL/min). The phenol-sulfuric acid method was used to monitor the factions of crude FVP, and some completely separated fractions were collected. After dialysis, concentration, and purification with a Sephadex G-100 column, two main fractions, CHFVP-1 and CHFVP-2, were obtained.

2.3. Ultraviolet Spectrum Scan

CHFVP-1 and CHFVP-2 solutions (1 mg/mL) were analyzed by UV spectroscopy at a wavelengths range of 200–800 nm [13].

2.4. Infrared Spectrum Analysis

The infrared spectrum of CHFVP-1 and CHFVP-2 was measured by a Fourier-transform infrared (FT-IR) spectrometer with the KBr pellet method in the range of 4000–400 cm⁻¹.

2.5. Molecular Weight Determination Analysis

The relative molecular weight of CHFVP-1 and CHFVP-2 were commissioned for Beijing Physical and Chemical Analysis and Testing Center and determined according to the 2015 edition of the Pharmacopoeia of the People’s Republic of China.

2.6. Monosaccharide Composition Analysis

Gas chromatography (GC) in reference [12], a popular method for monosaccharide composition, was adopted in this paper. Briefly, hydrolyze sample with trifluoroacetic acid (TFA) and get the hydrolysates. Mix it with hydroxylamine hydrochloride, inositol hexyl acetate, and pyridine, add acetic anhydride, and bathe the mixture in hot water for 40 min. After that, dissolve it in chloroform. Filter the reactants and analyze it by gas chromatography fitted with a Thermo TG-Waxms capillary column (30 m × 0.32 mm × 0.5 μm). The analysis situation of GC was the same as Wang’s research [12]. The related
monosaccharide (glucose, arabinose, xylose, ribose, fucose, mannose, fructose, galactose, and rhamnose) was derived with the above method.

2.7. Methylation Analysis

In this part, the analysis of CHFV-1 and CHFV-2 was consistent with the method of Ciucanu et al. [14]. Briefly, dissolve the sample in the dimethyl sulfoxide (DMSO); then, add NaOH dry powder and CH3I to methylate it. During this step, the absorption (3200–3700 cm⁻¹) on infrared spectroscopy was used to judge the success of methylation. The sample after methylation was hydrolyzed at 100 °C by adding Trifluoroacetic Acid (TFA). After cooling, NaOH solution and NaBH₄ were added to the sample for reaction, and next, acetic acid was used to neutralize the sample. Then, we added pyridine and acetic anhydride for acetylation before analysis with the GC-Mass Spectrophotometer (MS) system with a J&W DB-5 ms capillary column (30 m × 250 μm × 0.25 μm). The GC situation was that the temperature increased from 110 to 190 °C at the rate of 5 °C/min, to 210 °C at 3 °C/min and finally to 230 °C at 5 °C/min.

2.8. Nuclear Magnetic Resonance (NMR) Spectroscopy

Dissolve 15 mg of CHFV-1 and CHFV-2 in 0.5 mL of D₂O (the internal standard was Tetramethylsilane, TMS), respectively and load them into the NMR tube. NMR experiments were recorded on a Bruker-AV-500 spectrometer at 25 °C. The measured spectra included ¹H, ¹³C, H/H correlation spectroscopy (COSY), heteronuclear single quantum coherence spectroscopy (HSQC), and heteronuclear multiple bond correlation spectroscopy (HMBC).

2.9. Coagulation Activities In Vitro

CHFV-1 and CHFV-2 were separately dissolved in saline, and 5 mg/mL of sample solutions were obtained. Then, we mixed them together for the next step. Breviscapine was dissolved in saline to get 13.3 mg/mL solution. The preparation method of Yunnan Baiyao was similar to Breviscapine [15]. Rabbits were used as experimental animals, and then PT, APTT, and TT detection kits were used for determination. Sodium citrate in saline was taken into an eppendorf (EP) tube, rabbit ear venous blood was added, and plasma was prepared after being centrifuged at 3000 g/min for 15 min (TGL-16gR, Shanghai Anting Scientific Instrument Co., Shanghai, China). CHFV-1 and CHFV-2 were transferred to plasma, and APTT was added. After preheating and incubating it for 5 min, steel balls and CaCl₂ solution were added successively and pre-warmed at 37 °C. Finally, the experimental results were recorded. For PT analysis, CHFV-1 and CHFV-2 and serum were incubated with the coagulation and minimum steel ball cell at 37 °C for 3 min. After that, PT reagent was added at 37 °C. TT analysis was determined according to the instruction. During the determination, according to many researchers, breviscapine and Yunnan Baiyao were taken as anti-coagulated [16] and pro-coagulated [17] reagents, respectively, while the normal saline was taken in the blank control group. Record the thrombus time with the RAC-030 automatic coagulation analyzer (Shenzhen Leidu Life Science Co., Ltd., Shenzhen 518067, China). The software SPSS 22.0 was used for data analysis.

3. Results and Discussion

3.1. Preparation of Polysaccharides from F. Velutipes

After the extraction of polysaccharides from F. velutipes, the crude polysaccharides were obtained, and the yield was about 1.33% (on dry; g/g). After the DEAE-52 cellulose column isolation, FVP was separated into five components (Figure 1a). In this study, FVP-1 and FVP-2 were selected for the next step of purification and structural study. Then, the purified yields of FVP-1 and FVP-2 were 8.4% and 36.2% respectively and pu-
rified with the Sephadex G-100 column to obtain the pure subfracion CHFVP-1 (Figure 1b) and CHFVP-2 (Figure 1c) with a yield of 88.4% and 78.9% (by dry weight).

Figure 1. Isolation and purification of crude polysaccharide from *F. velutipes*. (a) The elution curve of *F. velutipes* polysaccharides (FVP); (b) The elution curve of FVP-1; (c) The elution curve of FVP-2.
3.2. Spectroscopic Characteristics

The sample of FVP, CHFVP-1, and CHFVP-2 were scanned through ultraviolet spectrum, and the characterization of their absorption spectrum is shown in Figure 2. It could be found that there was no ultraviolet absorption at 260 and 280 nm, which means that FVP, CHFVP-1, and CHFVP-2 did not contain free protein and nucleic acid.

The infrared spectra of CHFVP-1 and CHFVP-2 at the range of 400–4000 cm\(^{-1}\) were scanned and are shown in Figure 3. The two fractions revealed the hydroxyl group (-OH) for the broad and strong absorption peak at around 3400 cm\(^{-1}\) [18]. The peaks at 2935 and 2923 cm\(^{-1}\) indicated the C, H and aliphatic C-H bonds, respectively [19]. Stretching peaks at 1642 and 1653 cm\(^{-1}\) corresponded to free carboxyl groups [20]. The broad absorption bands with intense intensities around 1420–1300 cm\(^{-1}\) could be owing to stretching vibrations of the C-H bond [21]. The absorption band at 1000–1200 cm\(^{-1}\) suggested that the two polysaccharides contained pyranose monomers in their structures [22]. The absorption bands at 882 and 888 cm\(^{-1}\) indicated the β-pyranoside linkage in CHFVP-1 and CHFVP-2 [23].

![Figure 2. Wavelength scanning curve of CHFVP-1, CHFVP-2, CHFVP-3, CHFVP-4, and FVP.](image1)

![Figure 3. Infrared spectra of CHFVP-1 and CHFVP-2.](image2)
Figure 3. (a) The Fourier-transform infrared (FT-IR) spectroscopy of CHFVP-1; (b) The FT-IR spectroscopy of CHFVP-2.

3.3. Molecular Weight and Monosaccharides Composition of CHFVP-1 and CHFVP-2

The average molecular masses of CHFVP-1 and CHFVP-2 were 24.44 and 1497 kDa, respectively (Table 1). The GC result of standard monosaccharides is shown in Figure 4a. According to the comparison between Figure 4a,b, it was found that CHFVP-1 was composed of fucose (Fuc), mannose (Man), glucose (Glu), and galactose (Gal) with a molar ratio of 1.51:3.42:15.28:10.00, respectively. CHFVP-2 was composed of arabinose (Ara), xylose (Xyl), ribose (Rib), Man, Glu, and Gal with a molar ratio of 0.85:0.47:0.42:1.79:18.46:1.00 (Figure 4c). Glu contributed to the main part of monosaccharides composition, indicating that glucose might be the main structure unit of CHFVP-1 and CHFVP-2, especially in CHFVP-2.

Table 1. Molecular weight of polysaccharides from *F. velutipes*.

| Polysaccharides | Mn   | Mp    | Mw    | Mz    | Mw/Mn |
|-----------------|------|-------|-------|-------|-------|
| CHFVP-1         | 23440| 22460 | 24440 | 25850 | 1.042 |
| CHFVP-2         | 1462000 | 1600000 | 1497000 | 1537000 | 1.024 |

Note: The number average molecular weight (Mn, g/mol) is the statistical average molecular weight of all polymeric chains in the polysaccharides; the weight average molecular weight (Mw, g/mol) is the statistical average of all molecular weights in the polysaccharides; Mp (g/mol) is defined as the molecular weight of the highest peak; Mz (g/mol) is defined as Z-average molecular mass, which is calculated according to the precipitation equilibrium method to determine the chain length; the ratio of Mw/Mn is called the polydispersity coefficient, which is used to describe the molecular mass distribution of polysaccharides.
Figure 4. The monosaccharide composition results of (a) reference; (b) CHFVP-1 and (c) CHFVP-2. 
(1: Rha; 2: Ara; 3: Fuc; 4: Xyl; 5: Ribose; 6: Man; 7: Glc; 8: Gal; 9: internal standard).

3.4. Methylation Analysis of CHFVP-1 and CHFVP-2

The glycosidic linkage and relative percentage of each sugar residue derivative according to the gas chromatography peak area determination from CHFVP-1 and CHFVP-2 were performed by methylation analysis (Table 2). The highest proportion of the sugar residue was the 1,6-Linked-Gal (33.25%) in CHFVP-1, followed by T-Glc (28.23%), 1,6-Linked-Glc (19.00%), 1,3,6-Linked-Man (10.18%), and 1,2,6-Linked-Man (9.24%). The residue related to the glucose, including 1, 6-Linked-Glc and T-Glc, accounted for almost half of all sugar residues. The results showed that glucose is composed of the main chain of CHFVP-1.

According to the results in Table 3, it could be found that CHFVP-2 is a glucan with T-Glc (67.01%) and 1,6-Linked-Glc (32.99%).

Table 2. GC-MS data for methylation analysis of CHFVP-1.

| Retention Time (min) | PMAA     | Linkages Types         | Percentage (%) |
|----------------------|----------|------------------------|----------------|
| 20.26                | 2,3,4,6-Me-Glc | T-Glc                 | 28.23          |
| 22.33                | 2,4-Me-Man  | 1,3,6-Linked-Man       | 10.18          |
| 23.13                | 2,3,4-Me-Glc | 1,6-Linked-Glc        | 19.00          |
| 24.14                | 2,3,4-Me-Gal | 1,6-Linked-Gal        | 33.25          |
| 26.07                | 3,4-Me-Man  | 1,2,6-Linked-Man       | 9.24           |
Table 3. GC-MS date for methylation analysis of CHFVP-2.

| Retention Time (min) | PMAA       | Linkages Types   | Percentage (%) |
|----------------------|------------|------------------|----------------|
| 20.28                | 2,3,4,6-Me-Glc | T-Glc           | 67.01          |
| 23.15                | 2,3,4-Me-Glc  | 1,6-Linked-Glc  | 32.99          |

Note: PMAA is the sugar alcohol acetate derivatives.

3.5. NMR Analysis of CHFVP-1 and CHFVP-2

The NMR spectra of CHFVP-1 are shown in Figure 5, which were the basis for inferring monosaccharide, glycosidic linkages, and chemical shifts. In Figure 5a,b, most of the proton and carbon chemical shifts of CHFVP-1 come from the regions ranging from δ 3.0 to 5.50 and δ 60 to 110, respectively, which were the typical chemical shifts of the polysaccharide [24]. Furthermore, four anomeric proton and carbon chemical shifts at δ 4.53/106.29, δ 5.001/100.79, δ 5.11/105.25, and δ 5.06/104.44 were identified in the 1H NMR, 13C NMR, and HSQC in Figure 5 and indicated as A, B, C, and D in CHFVP-1.

According to the composition of monosaccharides, methylation results, and literature comparison, the carbon signals at δ 18.52 in the 13C NMR and the proton shift at δ 1.23 in the 1H NMR indicated the existence of terminal-fucose (T-fuc) in CHFVP-1 [25,26]. The peak at δ 105.25 indicated the presence of 13C of α-1,3,6-linked D-mannopyranose units [27–29]. The peak at δ 100.79 meant the existence of C1 of the α-D-galactose [30]. The signal peak appearing at δ 64.0 suggested the presence of the C-6 galactose in the polysaccharide sugar chain structure [31]. The peaks in the region of δ 63.83–80.37 indicated the typical shift of carbohydrate. The glycosidic linkage of CHFVP-1 determined by its 1H NMR spectra in the region of δ 4.53–5.11 revealed that there were both α- and β-configurations in it [32].

![Image](attachment:image.png)
According to the anomeric chemical shift at δ 4.53/106.29 and literature comparison, the β-configuration of residue A was determined [33]. From the COSY spectrum (Figure 5c), the chemical shifts of H2-H6 can be deduced to δ 3.47, δ 3.72, δ 3.96, δ 4.36, and δ 3.76, respectively. The 13C chemical shifts of residue A were obtained from the HSQC spectrum by the chemical shift of 1H (Figure 5d), and the results are summarized in Table 4. Furthermore, according to the related literature and the chemical shifts of all 1H and 13C of residue A [25,34], residue A should be terminal-β-D-Glcp.

According to the anomeric chemical shift at δ 5.00/100.79 and literature comparison, the α-configuration of residue B was determined. The other chemical shifts of H-2, H-3, H-4, H-5, H-6 and H-6a at δ 3.89, δ 4.21, δ 4.07, δ 3.79, δ 3.91 and δ 3.70 were found from the cross-peaks in the COSY spectra (Figure 5c), and the corresponding carbon signals could be attributed to the correlations of C-H signals in the HSQC spectrum (Figure 5d), which were δ 72.43, δ 71.56, δ 71.21, δ 76.21, and δ 69.45 for C-2, C-3, C-4, C-5, and C-6, respectively. Regarding the complete 1H and 13C chemical shifts of residue B as well as previous work [35], residue B was →6)-α-D-Galp-(1→.
The heterotopic hydrogen and carbon signals of H-1 and C-1 were δ 5.11/105.25. According to the literature, residue C was inferred to be an α-configuration residue. Other signals from H-2 to H-6a were identified from the results of COSY spectrum (Figure 5c), including H-2 (δ 3.91), H-3 (δ 3.99), H-4 (δ 4.07), H-5 (δ 4.22), and H-6a (δ 4.00). In addition, the matching 13C chemical shifts were C-2 (δ 69.75), C-3 (δ 80.37), C-4 (δ 71.21), C-5 (δ 71.61), and C-6 (δ 74.25) in the HSQC spectrum (Figure 5d). The chemical shifts of all 1H and 13C of residue C were basically consistent with the literature [29], which indicated that residue C was →(3,6)-α-D-Manp-(1→.

Based on the terminal 13C and 1H shift at δ5.06/104.44, the α-configuration of the D residue was inferred. In the HSQC spectrum (Figure 5d), the cross-peak at δ1.23/18.525 indicated the H6/C6 of T-α-L-Fucp. The remaining hydrogen chemical shifts of D come from the COSY (Figure 5c) spectrum, and the 13C chemical shifts associated with it were determined in the HSQC spectrum (Figure 5d). Compared with the literature, the 1H and 13C chemical shifts for residue D were assigned [26]. The NMR chemical shifts and significant connectivity are summarized in Table 4 and Table 5.

### Table 4. 1H and 13C NMR chemical shifts of CHFVP-1 fraction recorded in D2O at 291K.

| Glycosyl residue | Chemical shift δ H/C (ppm) | H1/C1 | H2/C2 | H3/C3b | H4/C4 | H5/C5 | H6a/C6 |
|------------------|-----------------------------|-------|-------|--------|-------|-------|--------|
| A                | H                           | 4.53  | 3.35  | 3.49   | 3.66  | 3.86  | 3.77/3.9 |
| T-β-D-GlcN      | C                           | 106.29| 75.97 | 78.78/72.3 | 72.83 | 71.08 | 63.832 |
| B                | H                           | 5.00  | 3.89  | 4.21   | 4.07  | 3.79  | 3.91/3.7 |
| →6)-α-D-Galp-(1→ | C                           | 100.79| 72.43 | 71.56  | 71.21 | 76.21 | 69.45 |
| C                | H                           | 5.11  | 3.91  | 3.99   | 4.07  | 4.22  | 4.00 |
| →3,6)-α-D-Manp-(1→ | C                           | 105.25| 69.75 | 80.37  | 71.21 | 71.61 | 74.25 |
| D                | H                           | 5.06  | 3.86  | 3.91   | 4.074 | 4.17  | 1.23 |
| T-α-L-Fucp      | C                           | 104.44| 71.35 | 71.56  | 71.21 | 71.77 | 18.52 |

### Table 5. Significant connectivity observed in HMBC spectrum for the terminal 1H/13C of the residue fractions of CHFVP-1.

| Sugar Residue | Sugar Linkage | δH/δC | δH/δC | Residue | Atom |
|---------------|---------------|-------|-------|---------|------|
| A             | T-β-D-GlcN    | 4.53  | 71.77 | D       | C-5  |
| B             | →6)-α-D-Galp-(1→ | 106.29| 3.35  | A       | H-2  |
| C             | →3,6)-α-D-Manp-(1→ | 5.001| 71.77 | D       | C-5  |
| D             | T-α-L-Fucp    | 100.79| 71.35 | 71.56   | 71.77 | 18.52 |

Due to the large molecular weight and viscosity of CHFVP-1 and the wide distribution of branched chains, the NMR data of the sample were incomplete, and only partial sugar residue fragments were inferred. According to the correlation between the peaks of 13C and 1H from the HMBC spectrum (Figure 5e), the glycosides of the CHFVP-1 residues were linked as follows: AC-1/AH-2; AH-1/DC-5; BH-1/DC-5; CC-1/AH-6; CH-1/BC-5/CC-6/3; DH-1/B/C/DC-4; DC-1/AH-2. Hence, considering the aforementioned results, the most probable repeat unit of CHFVP-1 has an →6)-α-D-Galp-(1→ backbone
that is branched at O-5 of Galp by an →3,6)-α-D-Manp-(1→ attached with T-β-D-Glcp or T-α-L-Fucp side-chains.

CHFVP-2 was analyzed by the same way as that used for CHFVP-1. In Figure 6, anomeric proton and carbon signals at δ 4.53/105.88 were determined in the 1H and 13C NMR (Table 6 and Table 7) and HSQC spectra in Figure 6 and denoted as A and B in CHFVP-2 [34]. All the 1H and 13C signals were assigned by 1H/1H correlation spectroscopy (COSY), heteronuclear single quantum coherence (HSQC), and heteronuclear multiple-bond correlation (HMBC), as shown in Figure 6, indicating that residue A was T-β-D-Glcp and B was ←6)-β-D-Glcp-(1→.

The structure of CHFVP-2 was proposed based on the above analysis in the same way used for CHFVP-1, and the possible partial structure of CHFVP-2 information was shown as follows:

\[ \rightarrow [6)-β-D-Glcp-(1→6)-β-D-Glcp-(1→6)-β-D-Glcp-(1)_{6}\rightarrow \]

\[
\begin{array}{cccc}
2 & 4 & 5 \\
\uparrow & \uparrow & \uparrow \\
1 & 1 & 1 \\
\end{array}
\]

\[
\begin{array}{ccc}
β-D-Glcp & β-D-Glcp & β-D-Glcp \\
\end{array}
\]

(a)  

(b)
Figure 6. $^1$H spectrum (a); $^{13}$C spectrum (b); COSY spectrum (c); HSQC spectrum (d); HMBC spectrum (e).

Table 6. $^1$H and $^{13}$C NMR chemical shifts of CHFVP-2 fraction recorded in D$_2$O at 291K.

| Glycosyl Residue | Chemical Shift δ H/C (ppm) |
|-----------------|--------------------------|
|                 | H1/C1 | H2/C2 | H3/C3b | H4/C4 | H5/C5 | H6/C6b |
| A               | H     | 4.53  | 3.35   | 3.52  | 3.77  | 3.89   | 3.67   |
| T-β-D-Glcp     | C     | 105.88| 75.93  | 72.38  | 71.71  | 77.78  | 78.44/63.5|
| B               | H     | 4.53  | 3.35   | 3.52  | 3.77  | 3.89   | 4.23   |
| →6)-β-D-Glcp-1 | C     | 105.88| 75.93  | 72.38  | 71.71  | 77.78  | 71.71  |
Table 7. Significant connectivity observed in HMBC spectrum for the terminal $^1$H /$^13$C of the residue fractions of CHFVP-2.

| Sugar Residue | Sugar Linkage       | H-1/C-1 (ppm) Connectivities | δ Ab/δ C | δ Ab/δ C | Residue | Atom |
|--------------|---------------------|-----------------------------|---------|---------|---------|------|
| A            | T-β-D-Glcp          |                             | 4.53    | 71.71   | B       | C-6  |
|              |                     |                             | 105.88  | 3.89    | A/B    | C-4  |
|              |                     |                             |         |         | A/B    | H-5  |
|              |                     |                             |         |         | A/B    | H-2  |
| B            | →6)- β-D- Glcp-(1→6) |                             | 4.53    | 71.71   | B       | C-6  |
|              |                     |                             | 105.88  | 3.89    | A/B    | C-4  |
|              |                     |                             |         |         | A/B    | H-5  |      |

3.6. Effects of CHFVP-1 and CHFVP-2 on Parameter Coagulation Parameters In Vitro

Four kinds of coagulation kits were used to detect the coagulation activity of polysaccharide components, including APTT, PT, and TT in vitro.

In Table 8, CHFVP-2 prolonged PT significantly compared with both the blank group (P < 0.001) and breviscapine group (P < 0.01), which revealed the extremely significant anticoagulant activity of CHFVP-2. The PT of CHFVP-1 had a slight change but not significant difference. Compared with the blank group, CHFVP-1 significantly shortened APTT, meaning its significant procoagulant activity (P < 0.001). CHFVP-2 also had significant procoagulant activity in terms of APTT (P < 0.01), and the activity was stronger than that of Yunnan Baiyao. Regarding the effect of TT, CHFVP-1 significantly shortened the TT compared with the blank group, (P < 0.01) which had very significant procoagulant activity, while CHFVP-2 also had certain activity.

Table 8. Coagulation effects of CHFVP-1 and CHFVP-2 in vitro.

| Group             | PT(s)  | APTT(s) | TT(s)  |
|-------------------|--------|---------|--------|
| Control           | 13.77 ± 0.30 | 15.37 ± 0.15 | 13.97 ± 0.23 |
| Yunnan Baiyao    | 12.20 ± 0.15 | 14.47 ± 0.35 | 12.57 ± 0.11 |
| Breviscapine      | 14.77 ± 0.55 | 16.50 ± 0.36 | 16.20 ± 0.53 |
| CHFVP-1           | 13.57 ± 0.05 | 9.60 ± 0.42 | 13.07 ± 0.30 |
| CHFVP-2           | 15.77 ± 0.32 | 13.95 ± 0.78 | 13.17 ± 0.49 |

Comparing with Control, ***P < 0.001, **P < 0.01, *P < 0.05. Compared with Yunnan Baiyao, ***P < 0.001, **P < 0.01, *P < 0.05. Compared with breviscapine, #&P < 0.001, #&P < 0.01, #&P < 0.05.

Based on the above results, CHFVP-1 significantly shortened APTT and TT (P < 0.01). Thus, CHFVP-1 could activate coagulation factors and promote the condensing of fibrinogen into insoluble fibrin, indicating that CHFVP-1 promoted blood clotting through the endogenous coagulation pathway. CHFVP-2 shortened APTT but increased PT. It is temporarily unable to predict its coagulation activity in vitro.

4. Conclusions

This study isolated two polysaccharides in *F. velutipes*, CHFVP-1 (24.44 kDa) and CHFVP-2 (1497 kDa), and identified the structural characterization and coagulation activity of them. The CHFVP-1 was constructed with →6)-α-D-Galp -(1→ residues as backbone, while CHFVP-2 was constructed with →6)-β-D-Galp -(1→ residues. Moreover, the branch chain of CHFVP-1 might be inferred as →3,6)-α-D-Manp -(1→ attached with T-β-D-Glcp or t-α-L-Fucp, and that of CHFVP-2 was T-β-D-Glcp. Based on the coagulation assay results of APTT and TT of CHFVP-1, the polysaccharide exerted promote blood clotting primarily through endogenous coagulation pathways.
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Institutional Review Board Statement: The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the Ethics Committee of College of Medical, Henan University (HUSOM-2019-035, 2019).

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Ethical conduct of research: The study obtained ethical clearance from the Ethics Committee of College of Medical, Henan University (HUSOM-2019-035). The rabbit was treated as the guidelines on the care and use of animals for scientific purposes. All the animal procedures were approved by the Ethical Committee of Henan province, China and conducted in accordance with “Institute ethical committee guidelines” for animal experimentation and care.

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