



ORIGINAL ARTICLE

α -glucosidase inhibitory, antioxidant activity, and GC/MS analysis of *Descurainia sophia* methanolic extract: *In vitro*, *in vivo*, and *in silico* studies



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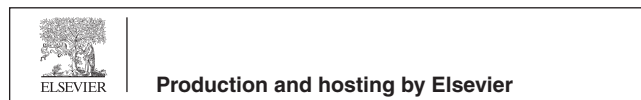
α -glucosidase;
Descurainia sophia;
Desolphosinigrin;
Molecular docking;
Molecular dynamics
simulation

Abstract Due to the presence of various phenolic compounds in *D.sophia*, this plant may have an inhibitory effect on α -Glc and ultimately diabetes control. Therefore, this work aims to scrutinize total phenolic, flavonoid contents, antioxidant capacity, and α -Glc inhibitory activity in aerial parts of methanolic *D.sophia* extract. The methanolic flower extracts were selected from among aerial parts for the experimental study of anti-diabetic effects by α -Glc inhibitory assays. The flower extracts were also studied by GC/MS to detect the compounds. The total phenolic and flavonoid contents were 21.38 ± 0.93 GAE/g and 96.2 ± 0.20 QE/g, respectively. The IC_{50} value of flower extract for α -Glc inhibition with mixed (Competitive/non-competitive) mode was found to be 20.34 ± 0.11 mg/ml. Furthermore, *in-vivo* studies showed that the blood glucose level reduced after consumption of flower extract compared to the control group. Twenty-one compounds were identified by GC/MS technique. These compounds were assessed for high docking scores against α -Glc *in silico*. Docking score calculations exhibited that the DES- α -Glc complex had a significantly higher binding energy (-6.13 Kcal/mol) than other compounds. The DES- α -Glc complex which displayed a higher docking energy value than the ACR was subjected to MDs studies. The findings

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of this study suggest that the flower extract of *D.sophia* can be used as a suitable additive in syrups or foods with anti-diabetic capacity.

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1. Introduction

DM is a metabolic disease characterized by high blood glucose levels (hyperglycemia) and lack or insufficient insulin secretion. Hyperglycemia is related to atherosclerotic disease and enhanced occurrence of microvascular complications in diabetic people. Chronic hyperglycemia can activate the non-enzymatic glycosylation and the diabetic complication (Shim et al., 2003, Becerra-Jiménez and Andrade-Cetto 2012). Furthermore, evidence indicated that long-time exposure to chronic high glucose levels stimulates the generation of ROS that has a pivotal role in diabetic complications. Moreover, diabetes also changes the activity of antioxidant enzymes in some tissues, such as the liver, aorta, and kidney (Fatehi-Hassanabad et al., 2010, Anichini et al., 2020).

Controlling blood glucose levels is an effective method to delay hyperglycemia. This is correlated to the breakdown of carbohydrates including α -Glc (Cakar et al., 2017, Ćakar et al., 2018). The α -Glc enzyme, a membrane-bound enzyme, is present in the brush border of the small intestine, which catalyses the hydrolysis of 1,4- α bonds of oligo and disaccharides and converts them into absorbable monosaccharides (Han et al., 2017, Ni et al., 2019, Sadeghi and Zarei 2020). Therefore, α -Glc inhibition can be an effective way to delay absorption and control blood glucose levels. Several AGIs such as metformin, ACR, thiazolidinediones, meglitinides, and sulfonylureas have been advanced to reduce postprandial blood glucose levels. However, continued use of these AGIs leads to numerous side effects including diarrhea, flatulence, and abdominal pain. (Cai et al., 2020, Xu et al., 2020). Thus, the use of natural resources such as plant-derived compounds with low side effects can be an interesting approach to reducing the blood glucose level in diabetic patients.

Some biologically active compounds derived from plants, foods, and fruits such as flavonoids, phenolic acids, stilbenoids, C-glycosylflavonoids, and terpenoids have been considered important sources of α -glucosidase inhibitors (Đorđević et al., 2018, Fettach et al., 2019, Aalim et al., 2021). Numerous pharmaceutical herbs have been used for several years to control diabetes in different countries. Iran is one of the countries with a great diversity in plant species, which grow across different regions (Sadat-Hosseini et al., 2017). The *Descurainia sophia* (L.) Webb ex Prantl (*D.sophia*) belongs to the Brassicaceae family in The Plant List (<https://www.theplantlist.org/>), a commonly used herbal medicine in Iranian folk medicine, is known as Khakshir (khak-e shir) (Amiri and Joharchi 2013, Zarshenas et al., 2013). Some reports have shown that the secondary metabolites of *D.sophia* include phenolics, cardiac glycosides, flavonoids, and sulfur glycosides (Amiri and Joharchi 2013). In Iran, Khakshir with other complementary syrups is used as a beverage (Mohammadinia et al., 2012, Pasalar et al., 2013).

The α -Glc inhibitory activity hexane extract of *D.sophia* has been reported (Zarei and Poursharifi 2015) but the hexane extract is not used as part of Khakshir. Due to the abundance of *D.sophia* in the Kurdistan region of Iran, it will be very interesting to investigate the inhibitory effects of its relevant extracts on α -Glc and the subsequent reduction in blood glucose levels. Hence, the present study aimed to determine phenolic and flavonoid contents, antioxidant activities, and α -Glc inhibitory of the aerial parts of *D.sophia*, as well as to characterize the potent extracts, *in vivo* investigation, GC/MS analysis, followed by molecular docking and molecular dynamics simulation for compounds with potential inhibitory effect.

2. Materials and methods

2.1. Chemicals

p-Nitrophenyl α -D-glucopyranoside (*p*NPG), *Saccharomyces cerevisiae* α -Glc, ACR, DPPH, sodium carbonate, and BSA, was purchased from Sigma Aldrich Chemical Co. London, UK. Alloxan, Aluminium chloride, quercetin, ascorbic acid, gallic acid, ferric chloride, Folin-Ciocalteu's reagent, TCA, and potassium acetate were purchased from Merck Co. Darmstadt, Germany.

2.2. Plant materials

The plant's aerial parts were collected from the central areas of Kurdistan Province, Iran, from June to July 2018. Voucher specimens have been deposited in the Herbarium of the Kurdistan Agricultural and Natural Resources Research & Education Center (HKC), Sanandaj, Iran. Their flowers, leaves, and stems were separated from each other. Separated aerial parts were dried and grinded by an electric grinder (Dolatabadi et al., 2011, Khan et al., 2018).

2.3. Preparation of crude extract

To determine the enzyme inhibitory and antioxidant activities of the crude extracts, 40 g of each ground powder was soaked in 200 ml methanol for 24 h with sporadic shaking. The soaked plant materials were filtered through a Whatman filter paper (grade No, 42). The solvent was evaporated with the rotary evaporator. The residues were air-dried under the chemical hood and finally collected in small microfuge tubes. A stock solution (200 mg/ml) was prepared, then diluted to different concentrations.

2.4. Determination of total phenolics

The total phenolics were measured according to Shahidi and Nacz (Shahidi and Nacz 1995). Briefly, 0.5 ml of 100 ppm the extract was mixed with 0.5 ml of Folin-Ciocalteu reagent and incubated for 2 min, followed by adding 10 ml Na₂CO₃ (7:100, v/v). Subsequently, the absorbance of samples was measured at 765 nm after heating at 45 °C water for 60 min. The total Phenolic contents were expressed as mg of Gallic acid equivalents per g of dry weight.

2.5. Determination of total flavonoid

The content of flavonoid compounds was measured by the Chang-Yang method and according to the aluminium chloride colorimetric method (Chang et al., 2002). Briefly, 0.5 ml of diluted samples were mixed with 0.1 ml of 10% AlCl₃ methanol solution, 0.1 ml potassium acetate 1 M, and 2.8 ml of dis-

tilled water. After incubating at RT for 30 min, the absorbance was measured at 415 nm. The flavonoid content was calculated from a Quercetin standard curve and expressed in mg Quercetin equivalent per g of dry mass.

2.6. DPPH free radical-scavenging capacity

Free radical scavenging activity was calculated using the method described by Prakash et al with a few modifications (Prakash et al., 2007). Briefly, a methanolic solution (0.1 ml) of the sample at different concentrations was added to 0.1 ml of DPPH (0.1 mM) solution. The reaction mixture was incubated at RT for 30 min with gentle shaking, the absorbance was measured at 517 nm. The percentage (%) of RSA was determined using the following formula:

$$\% \text{ RSA} = \frac{\text{Absorbance}(\text{Control}) - \text{Absorbance}(\text{Extract})}{\text{Absorbance}(\text{Control})}$$

The Ascorbic acid concentrations of 0.031, 0.25, 0.5, 1.0, 2.0, 5.0, 8.0, and 12.0 $\mu\text{g/ml}$ were used as positive controls and also the DPPH solution was used as a negative control. EC50 values indicate the concentration of the sample needed to scavenge 50% of the DPPH free radicals.

2.7. Determination of reducing power

The reduction power of the extracts was measured according to the method of Yen et al (Yen and Chen 1995). Briefly, 2.5 ml of phosphate buffer (0.2 M, pH 6.6) and 2.5 ml of potassium ferricyanide (1% v/v) were added to 2.5 ml extract solutions (50–800 $\mu\text{g/ml}$). After 30 min incubation at 45 °C, 2.5 ml of 10% trichloroacetic acid was added to the mixture and centrifuged for 10 min at 2300 g. Thereafter, 2.5 ml of the supernatant was mixed with 2.5 ml distilled water and 0.5 ml FeCl_3 (0.1%), and finally, absorbance was measured at 700 nm.

2.8. α -Glc inhibition assay

α -Glc inhibitory activity was assayed according to Pistia-Brueggeman et al, with a few changes (Pistia-Brueggeman and Hollingsworth 2003). Briefly, 20 μl of each plant extract dilution (5.0, 10.0, 15.0, 20.0, 25.0, 50.0, 100.0, and 200.0 mg/ml), 50 μl of phosphate buffer (50 mM; pH 6.8), 10 μl of α -Glc (1 U/ml) were mixed together in a well of 96-well microplate, then incubated at 37 °C for 5 min. subsequently, 20 μl of pNPG (1 mM) substrate was added to the mixture and after incubation at 37 °C for 30 min, the reaction was stopped by the addition of 50 μl of 0.1 M Na_2CO_3 . ACR was used as the positive control. Each experiment was performed in three replicates with appropriate blanks. The percentage of inhibition of α -Glc was obtained as follows:

$$\% \text{ inhibition} = \frac{\text{Absorbance}(\text{Control}) - \text{Absorbance}(\text{Extract})}{\text{Absorbance}(\text{Control})}$$

The IC_{50} was estimated using nonlinear regression analysis using SigmaPlot software. All values are expressed as Mean \pm Standard Deviation.

2.9. Kinetic analysis

The inhibition kinetics of plant extracts were analysed by the same method as described above for the α -Glc activity assay. The inhibitory method was considered by the effects of seven variant concentrations (5.0, 15.0, 20.0, 25.0, 50.0, 100.0, and 200 mg/ml) of plant extracts solutions. The maximum velocity (V_{max}) and Michaelis–Menten constant (K_m) were calculated by the Lineweaver–Burk (LB) curve in equation (1) (Lin et al., 2015). Equations (2) and (3) were used to find the inhibition type of the extract and the kinetic constants (K_i^k and K_i^i), respectively (Yang et al., 2013).

$$\frac{1}{V_0} = \frac{K_m}{V_{\text{max}}} \times \frac{1}{[S_0]} + \frac{1}{V_{\text{max}}} \quad (1)$$

$$K_{m1} = K_m \left(1 + \frac{[I_0]}{K_i^k} \right) \quad (2)$$

$$V_{\text{max}1} = \frac{V_{\text{max}}}{1 + \frac{[I_0]}{K_i^i}} \quad (3)$$

where $V_{\text{max}1}$ and K_{m1} are the maximum velocities of the enzyme and the Michaelis–Menten constant in the presence of inhibitor, respectively. $[S_0]$ and $[I_0]$ are the concentrations of the pNPG and Extract, respectively.

2.10. Animals

Male Wistar rats weighing 190 to 230 g were purchased from the Isfahan Royan Institute. Animals were housed in standard conditions with free access to water, food, and adequate light. All animal protocols were approved by the AJA University of medical sciences Animal Ethics Committee (ethical certificate: IR.AJAUMS.REC.1401.035). The rats became diabetic with an oral administration of alloxan (Rats 130 mg/kg body weight) and definite by blood glucose level (> 210 mg/dl) (Kalaivani et al., 2012).

2.11. Induced diabetes

Induction of diabetes with ALX was performed by previous studies with few modifications (Macdonald Ighodaro et al., 2017). The diabetes was induced by oral administration of ALX 130 mg/kg. Five groups of rats were selected ($n = 5$). Group I: As negative group, during the test period received 0.5 ml of 60% in NaCl (5 g/l) and did not receive any drug treatment. Group II: As diabetic group, received oral administration ALX 130 mg/kg dissolved at 60% in NaCl (5 g/l), and during the trial period they did not receive any drug treatment. Group III: As under treatment with oral administration of acarbose (10 mg/kg) and ALX (130 mg/kg) dissolved at 60% in NaCl (5 g/l). Group IV: As under treatment with oral administration of extract (2.25 g/kg) and ALX (130 mg/kg) dissolved at 60% in NaCl (5 g/l). Group V: As under treatment with oral administration of extract (4.50 g/kg) and ALX (130 mg/kg) dissolved at 60% in NaCl (5 g/l).

2.12. GC/MS study

To determine natural components of *D.sophia* flower, the considered extract was analyzed by GC/MS using an Agilent gas chromatograph model GC 7890A coupled with a mass spectrometer model MS 5973A (Schlawis et al., 2018). The column conditions were as follows: Elite-1 Capillary Column, 3 m × 0.25 mm I.D × 0.25 μm, working in electron influence method at 60 eV. A fixed stream of 1.32 ml/min and an additional volume of 1 μl of helium gas was used as carrier gas. The ion source and injector temperatures were adjusted to 210 °C and 250 °C, respectively. The control of the oven temperature was as follows: First, the oven temperature started from 60 °C with a rise of 20 °C/min to 310 °C/min and then finished with isotherm for 10 min at 310 °C. MS spectra were recorded at 60 eV, with a scan value of 30–1100 *m/z*.

2.13. Molecular docking

The *S.cerevisiae* α-Glc 3A4A (PDB ID) was downloaded from Protein Data Bank (<https://www.rcsb.org/structure/3A4A>) and the pattern quality and validation were checked by the PROCHECK server (Laskowski et al., 2006). Then the desired pattern was selected as the docking input file. The structures of

the ACR and compounds in the *D.sophia* flower were downloaded from PubChem (<https://pubchem.ncbi.nlm.nih.gov/>). All the water molecules were deleted and added the hydrogen bond, compute Gasteiger and atoms assign AD4 type with the AutoDock tools software (version 4.2). All docking parameters including Grid box (110 × 115 × 110), algorithm type, and runs (1000 runs) were calculated. Then, the docking score was calculated for each compound. Finally, Discovery Studio (DS version 3.5) software was used to analyze the inhibitor/enzyme interactions.

2.14. MD simulation

MD simulation with Gromacs was used to obtain further comprehensive binding information. The AMBER99SB-ILDN protein and the general amber force field were added to charge α-Glc and ligand, respectively. Ligand topology was created by ProdrG. The complex was solvated in SPC/E Water Models (Mark and Nilsson 2001). After adding the water, the complex was neutralized by adding thirty Na⁺ and ten Cl⁻. The covalent contacts between the atoms were limited by short energy minimization. Then the system was equilibrated in two stages. The first stage contained a fixed number of particles, volume, and temperature (NVT). The second stage included a constant number of particles, pressure, and temperature (NPT). Finally, the simulation was performed at 300 K and 1 bar pressure for 100 ns. MD simulation protocol between receptor-ligand complex was analysed according to the previous reports (Ni et al., 2020). Finally, the stability of the system was computed the RMSD, RMSF, Rg, and hydrogen bonding analysis.

2.15. Statistical analysis

The data and results of the experiments were analyzed using SigmaPlot software, version 12. ANOVA was used to assess the significant differences between groups; A value of

Table 1 Total phenolic and flavonoid content of different aerial parts of the *D.sophia*. The results are determined as mean ± SD.

Name	Aerial part	Total phenolic content (μg/ml)	Total flavonoid content (μg/ml)
<i>D. sophia</i>	Flower	21.38 ± 0.93	35.85 ± 0.73
	Leaf	2.96 ± 0.68	12.81 ± 0.87
	Stem	0.89 ± 0.07	4.53 ± 0.46

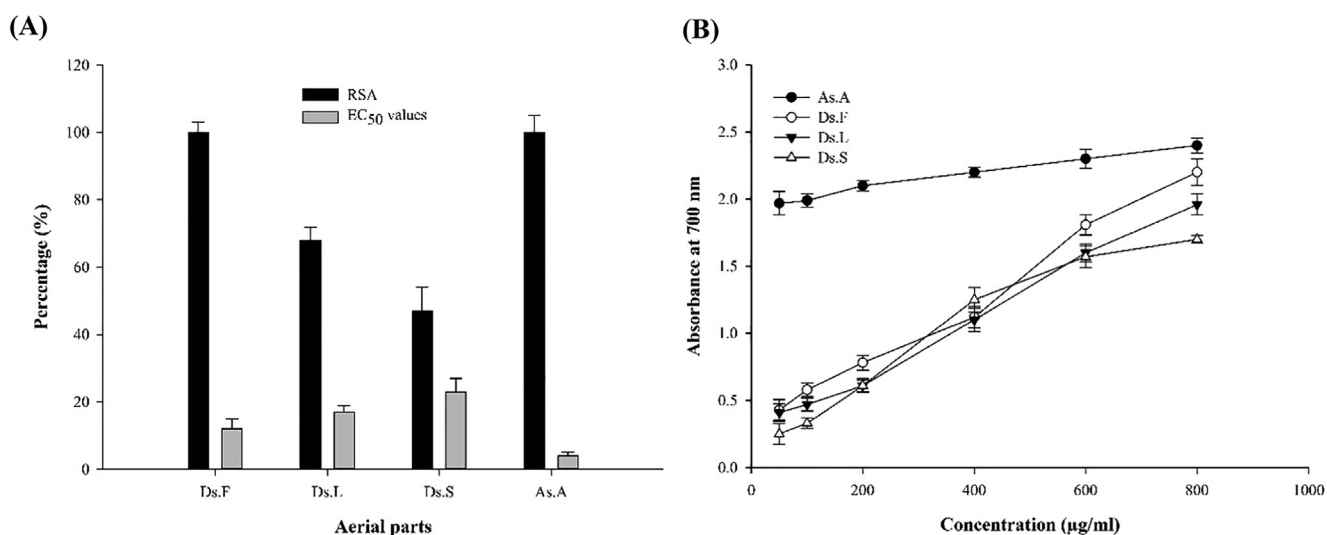


Fig. 1 Antioxidant activity results. (A) RSA and EC₅₀ values for different aerial parts of extracts (B) Reducing power graph of aerial parts of extracts. Results are expressed as mean ± SD in triplicate measurements.

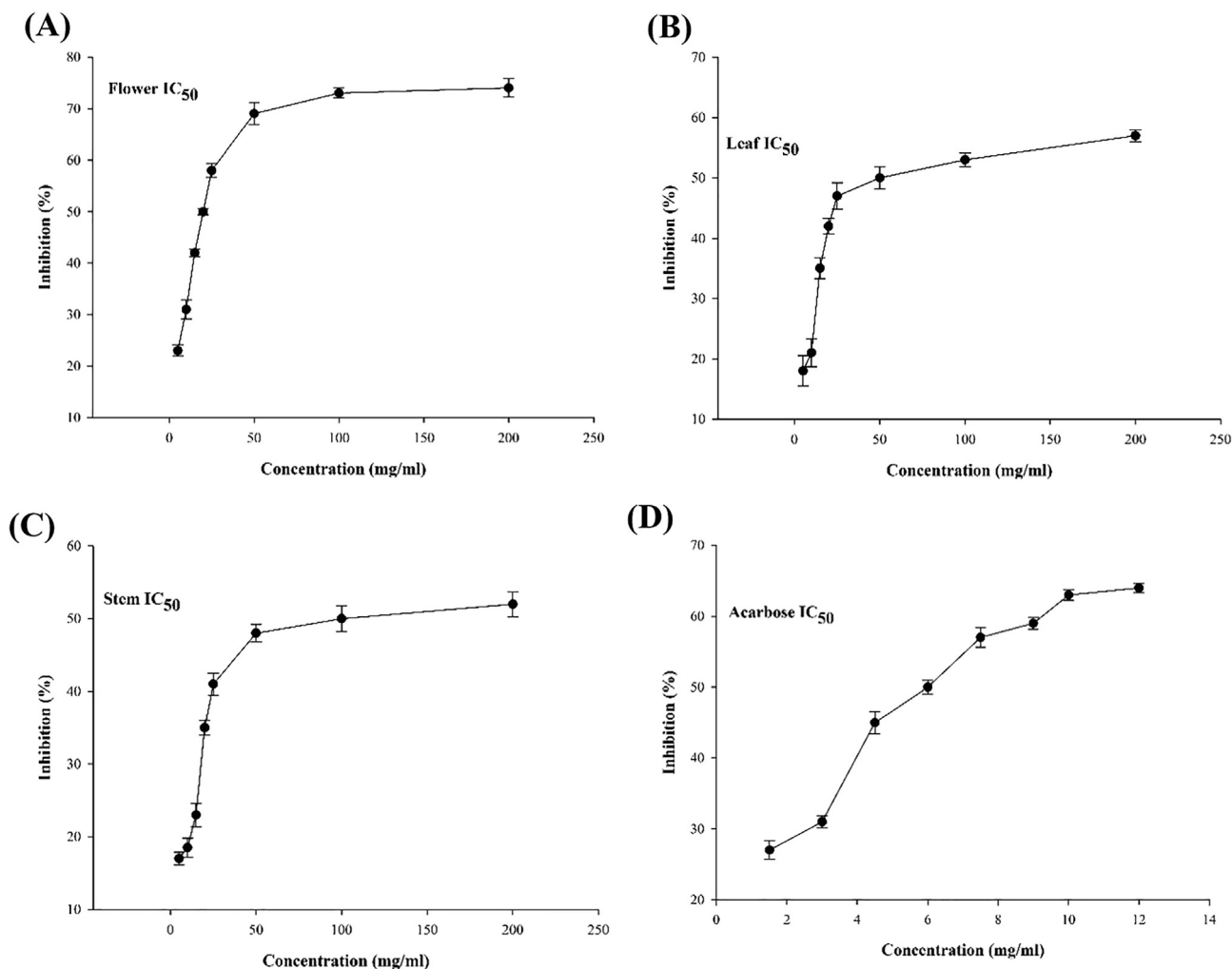


Fig. 2 The α -Glc inhibitory activity of aerial part extracts. (A) flower, (B) leaf, (C) stem, (D) ACR, positive control was ACR ($IC_{50} = 6.12 \pm 0.13$ mg/ml). The results are presented as IC_{50} versus concentration and determined as mean \pm SD.

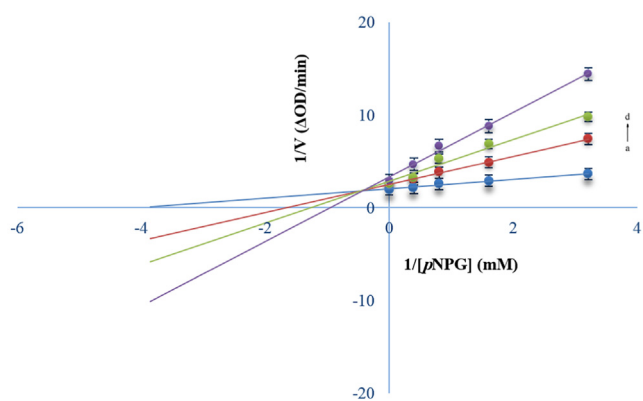


Fig. 3 LB plots. $c(\alpha\text{-glucosidase}) = 3.28 \times 10^{-7}$ mol/L, and $c(\text{flower extract}) = 0, 50.0, 100.0,$ and 200.0 mg/ml for Curves $a \rightarrow d$, respectively.

$p < 0.05$ were considered significant. And all the data from the experiment were replicated and their values were reported as Mean \pm Standard Deviation.

Table 2 Effect of different concentrations of flower extract on V_{max} , K_m , and ratios of the K_i^y to K_i^k .

$[I_0]$ (Extract)/mg. ml ⁻¹	K_m (mg)	V_{max} (mg.ml ⁻¹ . min ⁻¹)	K_i^y / K_i^k
0	2.63 ± 0.08	5.31 ± 0.14	–
50	5.93 ± 0.79	3.01 ± 1.03	2.3
100	6.71 ± 0.67	3.64 ± 1.06	3.14
200	8.03 ± 0.93	2.19 ± 0.58	3.79

3. Results and discussion

3.1. Phenolic and flavonoid contents

Flavonoids and other Phenolic are secondary metabolites found in some plants with antioxidant properties. The antioxidant activity of these compounds is due to the presence of hydroxyl groups in the aromatic ring structure and also using non-coupled electrons around the ring to eliminate free radicals (Alam et al., 2007). Previous studies have shown a positive

correlation between phenolic and flavonoids content and their relative anti-diabetic activities (Ahmed et al., 2014). The amount of total phenolic and flavonoid compounds in the aerial parts of the *D.sophia* was calculated (Table 1). The highest levels of Phenolic and flavonoid contents were quantified in flower part of *D.sophia* as 21.38 ± 0.93 mg GAE/g and 35.85 ± 0.73 QE/g, respectively. Therefore, due to the relationship between phenolic and flavonoid contents with α -Glc inhibition (Dirar et al., 2019), it is expected that this part will show more inhibition.

Added report corroborated the presence of flavonoid and phenolic complexes in the methanol extract of *D.sophia* as 41.77 ± 0.17 QE/g and 25.44 ± 0.53 mg GA/g, respectively (Mahomoodally et al., 2018). In this work, the *D.sophia* flower extract exhibited a low amount of flavonoid and phenolic compounds than the previous studies. This variation is possibly due to aspects such as environmental settings, solubility degree in the solvents, degree of polymerization of flavonoid and phenolic compounds, seasonal differences, etc (Medini et al., 2014). Most previous studies of plant extracts containing high flavonoid and phenolic compounds revealed good antioxidant capacity and AGI ability (Chen et al., 2019, Sarikurku et al., 2019). Thus, the present work was attentive on the antioxidant capacity and AGI identification from *D.sophia*.

3.2. Antioxidant activity

To evaluate the antioxidant activity, two common methods including free RSA and reducing power of plant extracts were employed. The first approach is to trap the DPPH radicals by the ability of hydrogen donation (Đorđević et al., 2018). In the second method, the reducing power of the material in the extract is assessed by reducing iron III into II. Iron reducing is often used as an electron-donating potential measurement, which is a suitable method for evaluating the antioxidant activity of phenolic compounds (Franco et al., 2018). RSA and EC₅₀ values were determined for aerial parts of *D.sophia* (Fig. 1A). The flower extract has the highest RSA and the lowest EC₅₀, which is similar to ascorbic acid. The results of the reducing power test is displayed in Fig. 1B. The range of reducing power in flower extract was 0.43 to 1.78, while it was lower in other aerial parts. Therefore, the antioxidant results showed that flower extract may have potential antioxidant activity due to higher RSA and power reducing levels as well as lower EC₅₀. Given the association between antioxidant potency and α -Glc inhibition (Girish et al., 2012), it is expected that flower extract may show more inhibition.

3.3. In vitro α -Glc inhibitory ability

The AGI abilities of aerial parts methanol extracts were signified in Fig. 2. The methanolic flower extract displayed good ability with an IC₅₀ value of 20.34 ± 0.11 mg/ml. The AGI ability of flower extract exhibited the highest ability of the other aerial parts. Further, the control ACR (IC₅₀ = 6.12 ± 0.13 mg/ml) revealed a very good inhibitory ability of α -Glc which are significantly better than flower extract. The purity of compounds could minor role in their inhibitory effects. But it depends simply on the nature of the compounds. If one compound in the crude extract owns a strong affinity to the active center, with a proper concentration the extract could

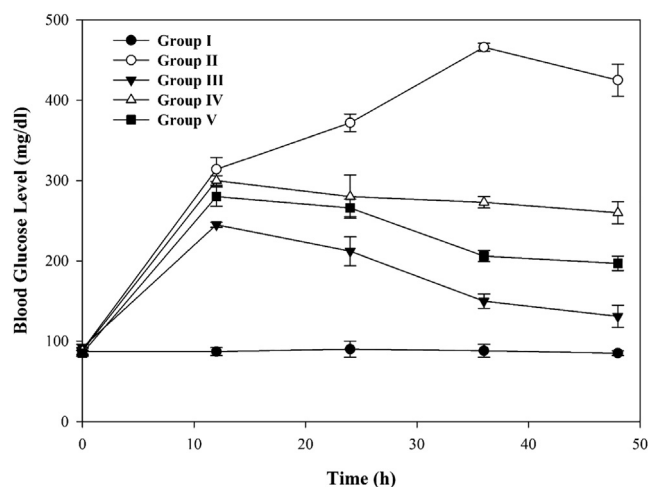


Fig. 4 Effect of *D.sophia* flower extract on blood glucose level in ALX diabetic rats. Thirty rats were divided into five groups: Negative Control (Group I), ALX diabetic (Group II), under treatment with ALX and ACR (Group III), under treatment with ALX and EXT 2.25 g/kg (Group IV), and treatment with ALX and EXT 4.50 g/kg (Group V). BGL was measured at 0, 12, 24, 36, and 48 h after administration of ALX according to the injection ALX test method described in the text. Oral administration dose; ALX at 130 mg/kg body weight, EXT at 2.25 g/kg body weight, EXT at 4.50 g/kg body weight, and ACR at 10 mg/kg body weight. Each dot signifies mean \pm S.D.

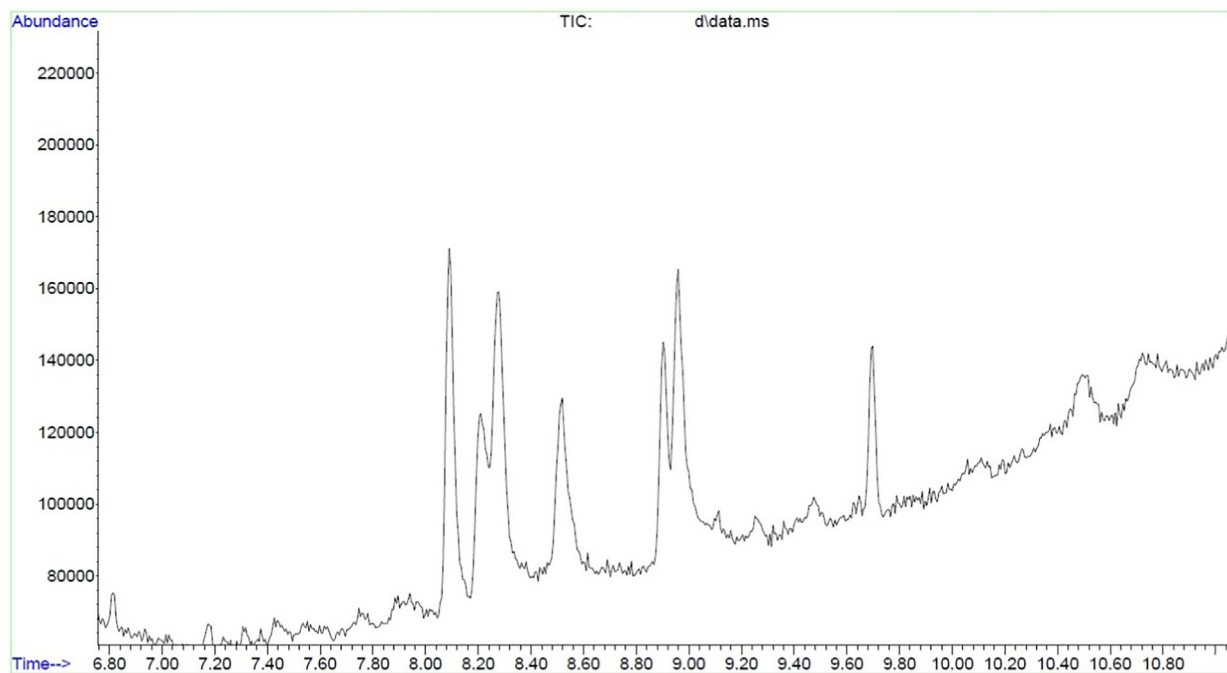
have a low IC₅₀ despite its complexity. There are numerous studies on the AGI ability of the aerial part extracts from other plants. The methanolic extract from *Ipomoea alba* has shown AGI ability with an IC₅₀ value of 17.12 ± 0.18 mg/ml against the α -Glc enzyme (Rosas-Ramirez et al., 2020). The flower and leaf extracts from *Psychotria malayana* revealed AGI ability with IC₅₀ values of 14.23 ± 0.16 and 21.16 ± 0.12 mg/ml respectively (Nipun et al., 2021).

3.4. Inhibition kinetics mediated by flower extract

Due to the high phenolic and flavonoid contents, antioxidant capacity, and a significant amount of α -Glc inhibitors in the *D.sophia* flower extract, the inhibition type of the α -Glc enzyme was determined for this part. The LB diagram was used to evaluate the inhibition type and kinetic parameters (Xiong et al., 2018). Fig. 3 shows the α -Glc activity in the presence of various concentrations of substrate and different amounts of flower extract. The lines intersect at a common point indicating a mixed (competitive/non-competitive) mode of inhibition (Beedessee et al., 2013).

The amount of K_m and V_{max} increases and decreases respectively, calculated by equation (1) with increasing of $[I_0]$, indicating a mixed inhibition by the flower extract (Table 2). In addition, the inhibition type of flower extract was further checked by Equation (2) and Equation (3) according to Johnson's theory (Johnson and Goody 2011). K_i^y into K_i^k (K_i^y/K_i^k) ratios can also be used to determine the type of inhibition. If this value is between 2.0 and 5.0, it indicates the mixed mode of inhibition (Ni et al., 2020). The mentioned ratios were 2.03 to 3.79, which signifies this type of inhibition on α -Glc.

(A)



(B)

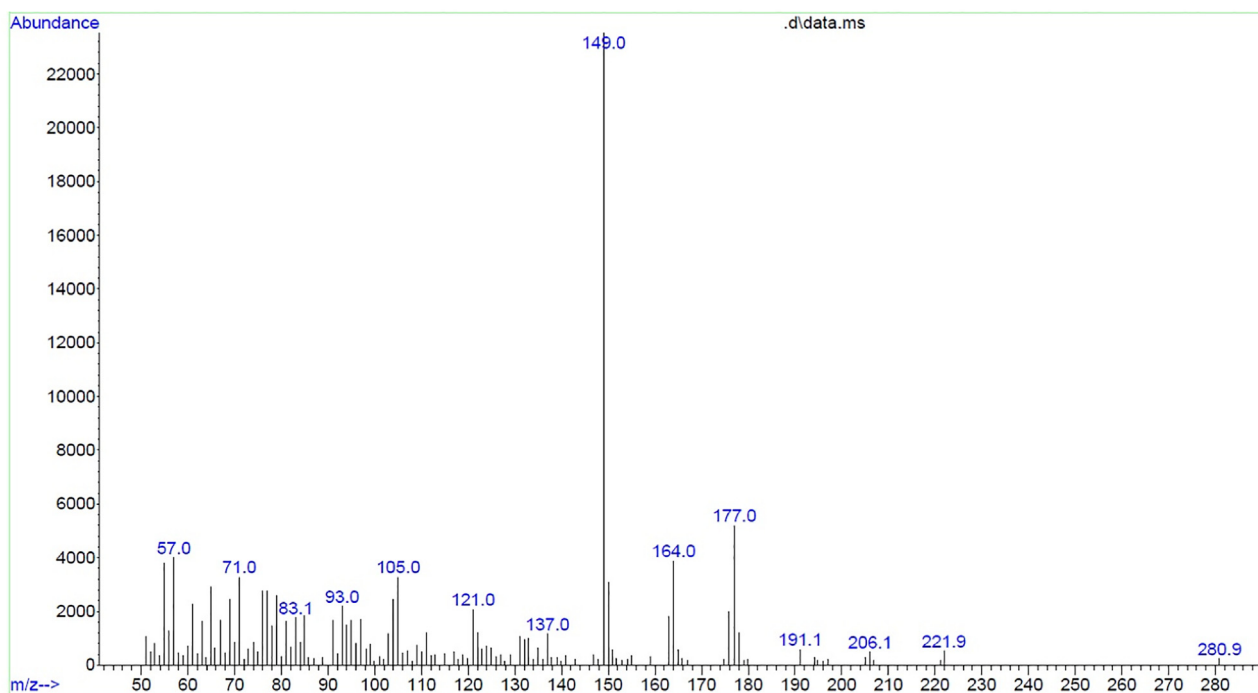


Fig. 5 GC/MS of flower extract of *D.sophia*. (A) Chromatogram of abundance versus RT (min), (B) Chromatogram of abundance versus m/z .

3.5. *In vivo* study of blood glucose level

Among the aerial parts of *D.sophia*, methanolic flower extract had more α -Glc inhibitory and antioxidant capacity. Therefore, in the present work, the hypoglycemic effect mediated

by *D.sophia* flower extract was investigated in diabetic rats. BGL was in the range of 85–94 mg/dl in all rat groups at the time of 0 h. The oral administration of the alloxan (130 mg/kg body weight) was utilized to heighten BGL; thirty-six hours after administration, it reached a maximum value of 466 mg/dl

Table 3 Phytochemicals recognized by GC/MS technique.

S.No.	RT	Peak area (%)	Compounds	Formula	PubChem CID	Mw (g/mol)
1.	6.82	0.63	Descurainin	C ₁₆ H ₁₈ O ₆	101857593	306.31
2.	7.16	1.78	Desulphosinigrin	C ₁₀ H ₁₇ NO ₆ S	9601716	279.31
3.	–	1.73	9-Octadecen-12-ynoic acid, methyl ester	C ₁₉ H ₃₂ O ₂	5363161	292.5
4.	7.88	2.36	5-Bromoadamantan-2-one	C ₁₀ H ₁₃ BrO	590905	229.11
5.	–	2.31	1-Methyl-1-hydroxymethyladamantane	C ₁₂ H ₂₀ O	590910	180.29
6.	8.00	0.51	Cinnamylaldehyde	C ₉ H ₈ O	637511	132.16
7.	8.20	42.13	Benzylidenemalonaldehide	C ₁₀ H ₈ O ₂	583063	160.17
8.	–	42.17	Ethyl 3-(3-pyridyl) propenoate	C ₁₀ H ₁₁ NO ₂	5371888	177.2
9.	8.33	14.17	Bicyclohexyliden-2-one	C ₁₂ H ₁₈ O	13892	178.27
10.	8.50	11.18	Quercetin	C ₁₅ H ₁₀ O ₇	5280343	302.23
11.	–	11.14	8,11,14-Eicosatrienoic acid, (Z,Z,Z)-	C ₂₀ H ₃₄ O ₂	5282826	306.5
12.	–	12.19	Tricyclo[4.3.1.1(3,8)]undecane-3-carboxylic acid	C ₁₂ H ₁₈ O ₂	541957	194.27
13.	8.11	3.45	Isovanillic acid	C ₈ H ₈ O ₄	12575	168.15
14.	8.17	9.14	Isorhamnetine	C ₁₆ H ₁₂ O ₇	5281654	316.26
15.	8.90	7.66	(E)-Isoeugenol	C ₁₀ H ₁₂ O ₂	853433	164.2
16.	9.11	15.43	Bergapten	C ₁₂ H ₈ O ₄	2355	216.19
17.	–	16.05	Phthalic acid, butyl undecyl ester	C ₂₃ H ₃₆ O ₄	6423450	376.5
18.	–	16.88	Phthalic acid, hexyl tetradecyl ester	C ₂₈ H ₄₆ O ₄	6423379	446.7
19.	9.46	6.48	4,7-Octadecadiynoic acid, methyl ester	C ₁₉ H ₃₀ O ₂	569159	290.4
20.	9.69	3.66	Phen-1,4-diol, 2,3-dimethyl-5-trifluoromethyl	C ₉ H ₉ F ₃ O ₂	590850	206.16
21.	10.5	9.14	Pterin-6-carboxylic acid	C ₇ H ₅ N ₅ O ₃	135403803	207.15

Table 4 Docking score compounds in the active site of α -Glc.

S. No.	Compounds	Binding energy (kcal/mol)
1.	Descurainin	-4.13
2.	Desulphosinigrin	-6.13
3.	9-Octadecen-12-ynoic acid, methyl ester	-3.98
4.	5-Bromoadamantan-2-one	-4.17
5.	1-Methyl-1-hydroxymethyladamantane	-4.33
6.	Cinnamylaldehyde	-5.03
7.	Benzylidenemalonaldehide	-4.99
8.	Ethyl 3-(3-pyridyl) propenoate	-4.17
9.	Bicyclohexyliden-2-one	-4.13
10.	Quercetin	-5.14
11.	8,11,14-Eicosatrienoic acid, (Z,Z,Z)-	-4.76
12.	Tricyclo[4.3.1.1(3,8)]undecane-3-carboxylic acid	-3.19
13.	Isovanillic acid	-4.97
14.	Isorhamnetine	-5.08
15.	(E)-Isoeugenol	-5.11
16.	Bergapten	-3.78
17.	Phthalic acid, butyl undecyl ester	-4.08
18.	Phthalic acid, hexyl tetradecyl ester	-3.66
19.	4,7-Octadecadiynoic acid, methyl ester	-3.17
20.	Phen-1,4-diol, 2,3-dimethyl-5-trifluoromethyl	-4.19
21.	Pterin-6-carboxylic acid	-3.77
22.	ACR (control)	-5.23

(Fig. 4). The rats that received ACR (oral administration 10 mg/kg body weight) exhibited a reduction of the BGL in comparison with the alloxan group. Rats given *D.sophia* flower extract (oral administration 2.25 g/kg body weight) showed a

considerable reduction in BGL up to 273 mg/dl at 36 h post-administration. Moreover, intake of *D.sophia* flower extract (oral administration 4.50 g/kg body weight) in rats reduced the BGL up to 206 mg/dl at 36 h post-administration. Accordingly, the ACR (10 mg/kg body weight) and flower extract (4.5 g/kg body weight) groups showed the lowest BGL in comparison with the diabetic group (p-value < 0.05). Therefore, our results propose that ACR and *D.sophia* flower extract can prevent the hydrolysis of the disaccharide and reduce the BGL. These findings may be due to structural similarity between ACR and compounds of *D.sophia* flower extract in α -Glc inhibition.

3.6. Phytochemistry

Among the aerial parts of *D.sophia*, methanolic flower extract had more α -Glc inhibitory potency and antioxidant properties. Therefore, the *D.sophia* flower compounds were identified through GC/MS technique. The different compounds were recognized by analyzing the spectral data obtained from GC/MS (Fig. 5A, 5B) and comparing them with the information of the National Institute of Standard and Technology (NIST). The compound's name, RT, formula, peak area, Mw, PubChem CID, and the structure of the flower extract were shown (Table 3). Most plants and their components can lower the blood glucose amount due to their diversity of flavonoid, phenolic, tannins, and terpenoids compounds (Osadebe et al., 2010). Natural compounds described by GC/MS study were found to have potential hypoglycemic effects. Numerous healing compounds were recognized through GC/MS by many scientists. These compounds possess antifungal, antibacterial, antioxidant, anti-diabetic, anticancer, and anti-inflammatory properties (Kalaivani et al., 2012, Gopinath et al., 2013). From the prior reports, it was established that quercetin (Kim et al., 2017), isoeugenol (Topal 2019), isorhamnetine (Shibano et al., 2008), and Isovanillic acid (Choi et al., 2020) are important

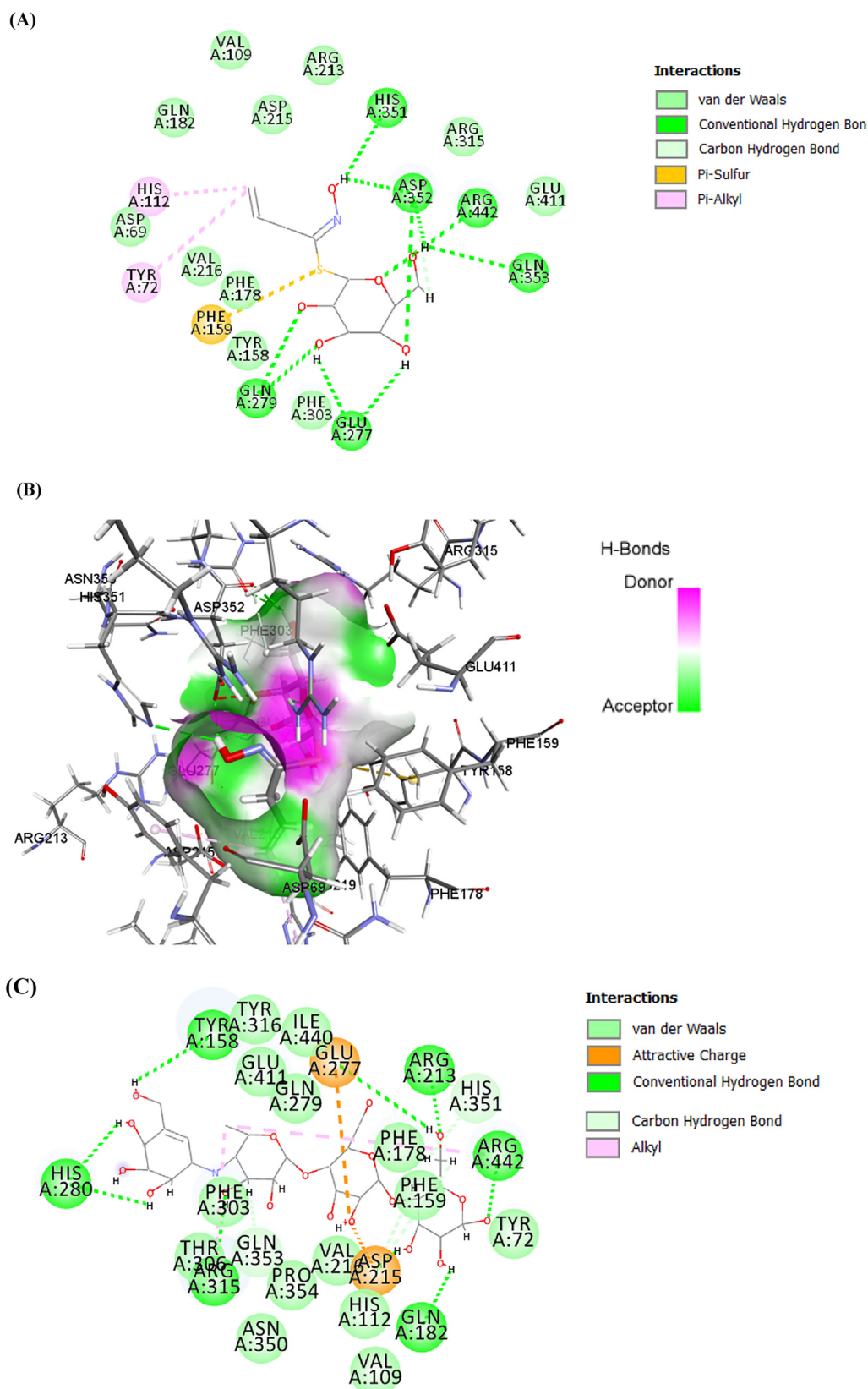


Fig. 6 Interaction results. (A, B) 2D and 3D interaction between α -Glc-DES complex, respectively. (C, D) 2D and 3D interaction between α -Glc-ACR complex, respectively.

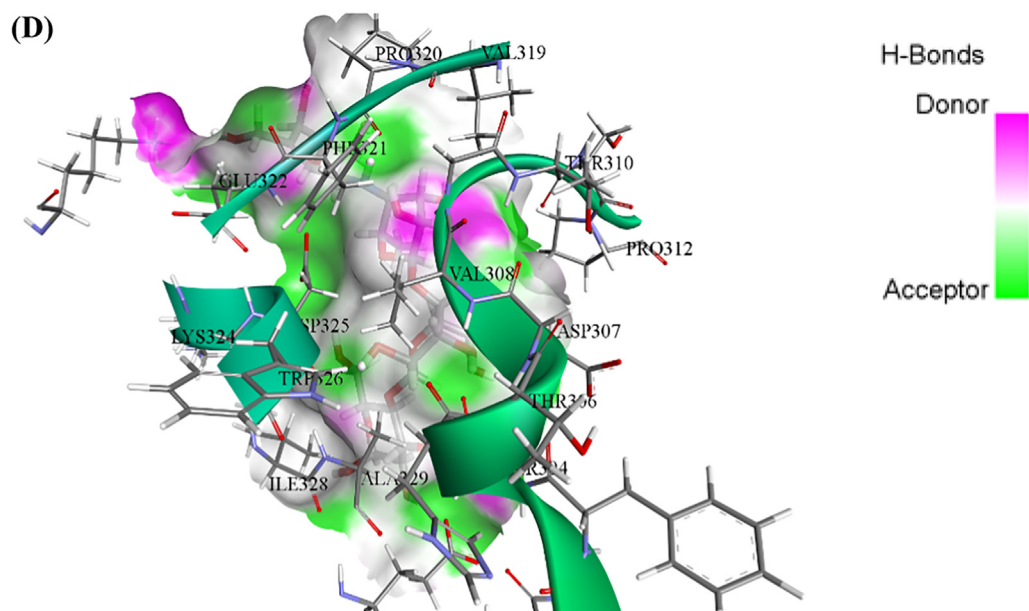


Fig. 6 (continued)

natural compounds with antioxidant and α -Glc inhibitory activity. GC/MS investigation of dichloromethane extract of *Ferulago bracteata* exhibited the presence of bergapten, 4,7-Octadecadiynoic acid, methyl ester, Cinnamylaldehyde and other compounds with α -Glc inhibitory capacity (Karakaya et al., 2018). Hence, in consistent with the previous studies, our findings demonstrated the presence of natural compounds with α -Glc inhibitory activity in the *D.sophia* flower methanolic extract. According to GC/MS study, there are twenty-one compounds. In next step, molecular docking was performed to bind affinity each of these compounds to the α -Glc enzyme.

3.7. Molecular docking studies

The molecular binding approach has been widely used to gain a better understanding of the receptor-ligand interactions (Şöhretoğlu et al., 2018, Li et al., 2022). Moreover, the position of *D.sophia* flower compounds into the active site of α -Glc was prophesied by this method. In this respect, the lowest docking score between α -Glc with *D.sophia* flower compounds was selected to carry out MDs in the next stage. The docking score was regarded as the comparison criterion and ACR (Standard drug) was considered as a control compound. The docking score of ACR with α -Glc was -5.23 kcal/mol (Table 4). However, among the twenty-one *D.sophia* flower compounds, DES showed a better docking score than ACR.

Although no collected work is existing on the docking score of this compound, the docking energy of DES with the α -Glc enzyme is very promising. If the DES retains effective docking energy against human α -Glc similar to this work, there would be a prospect to improve this compound into a new AGI. Many compounds such as quercetin, isorhamnetine, bergapten, etc. have been reported and isolated in the Brassicaceae family for diverse biological actions including the antidiabetic ability (Mahomoodally et al., 2018). Nonetheless, a detailed study on the anti-diabetic ability of isolated compounds from *D.sophia* is lacking. In this work, the α -Glc inhi-

biton potential of the *D.sophia* flower was assessed using experimental and computational studies signifying that there are active inhibitors present in the *D.sophia* flower extract. This work would be a motivation for more comprehensive studies on these identified compounds to expand them as natural AGI.

ASP69, ASP215, PHE303, ARG315, HIS351, and ARG442 were found at the active site of the α -Glc enzyme (Ning et al., 2019). The α -Glc-DES docked complex exhibited the minimum docking score (-6.13 kcal/mol). The interaction type of enzyme residues with the ligand can be divided into four groups: van der Waal, Conventional H-bonds, carbon-hydrogen, and pi-alkyl. Among the residues, GLU277, GLN279, HIS351, ASP352, GLN353, HIS351, and ARG442 were observed to interact through a conventional hydrogen bond. VAL109, GLN182, ARG213, ASP215, ARG315, and GLU411 residues formed van der Waals bond. ASP69, TYR158, and PHE303 residues can interact with DES by carbon-hydrogen bond formation. Furthermore, TYR72 and HIS112 formed pi alkyl bonds with DES (Fig. 6A, 6B). In the case of α -Glc-ACR complex, TYR158, GLN182, ARG213, HIS280, ARG315, and ARG442 residues were seen to interact through a conventional hydrogen bond. HIS351 and GLN353 formed carbon-hydrogen bonds and GLU277 and ASP215 can be interacted with ACR by Pi anion bond formation. Fourteen residues involved in the formation of van der Waals bonds can be seen too (Fig. 6C, 6D). It is observed that the conventional hydrogen bond number in the α -Glc-DES is more than α -Glc-ACR complex. Therefore, considering the importance of conventional hydrogen bonds compared to other bonds, it is expected that the α -Glc-DES docking score is better than the α -Glc-ACR complex.

3.8. Molecular dynamics simulation analysis

MDs are used to study the binding of a ligand to an enzyme over some time at the atomic level. Several factors, such as

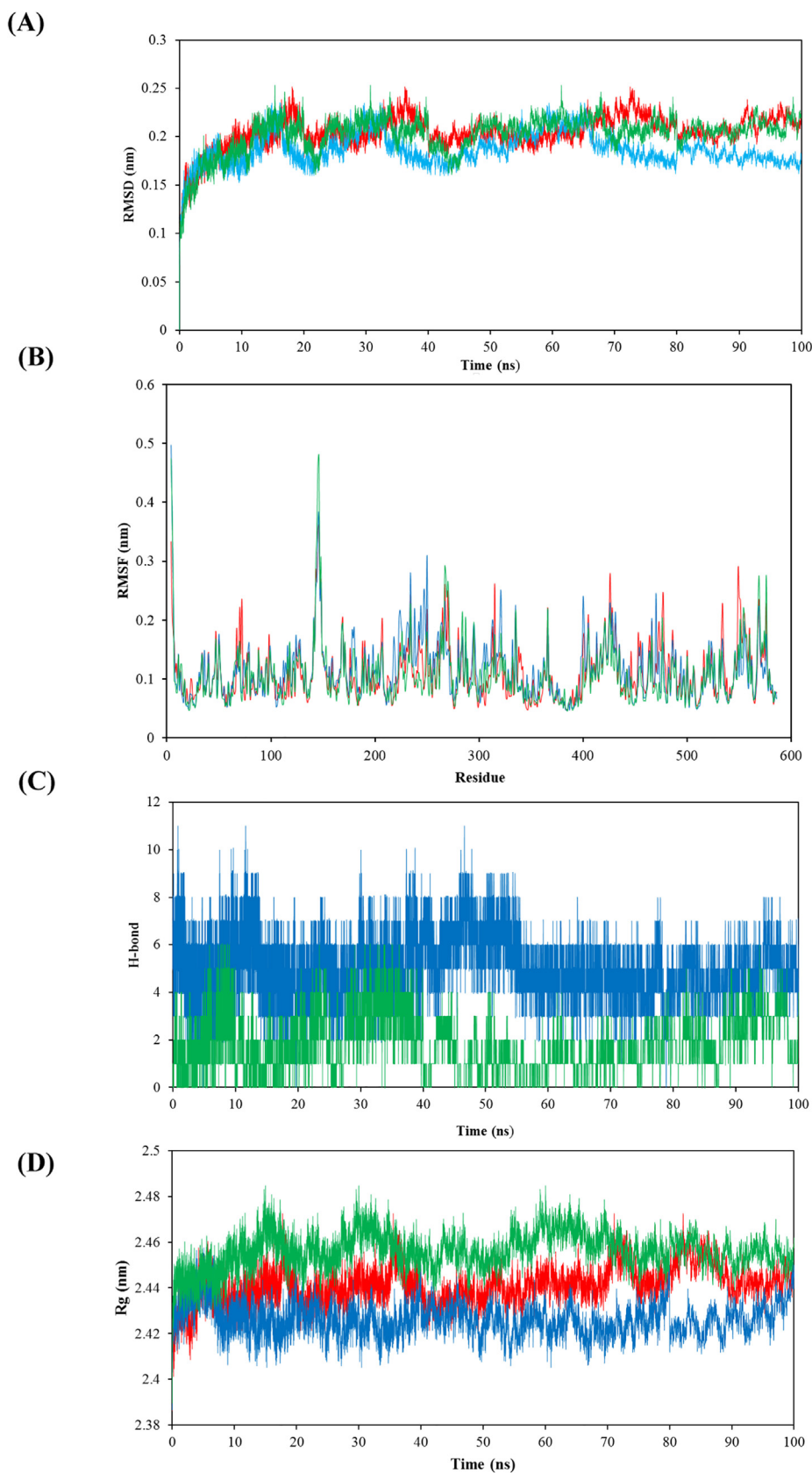


Fig. 7 Molecular dynamics simulation results. (A) RMSD values for α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex during 100 ns molecular dynamics simulation. (B) RMSF of residues of α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex all amino acids. (C) The number of hydrogen bonds between α -Glc-DES complex, and α -Glc-ACR complex for 100 ns. (D) Rg values for the α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex during 100 ns MD simulation. The red, blue, and green colors represent α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex, respectively.

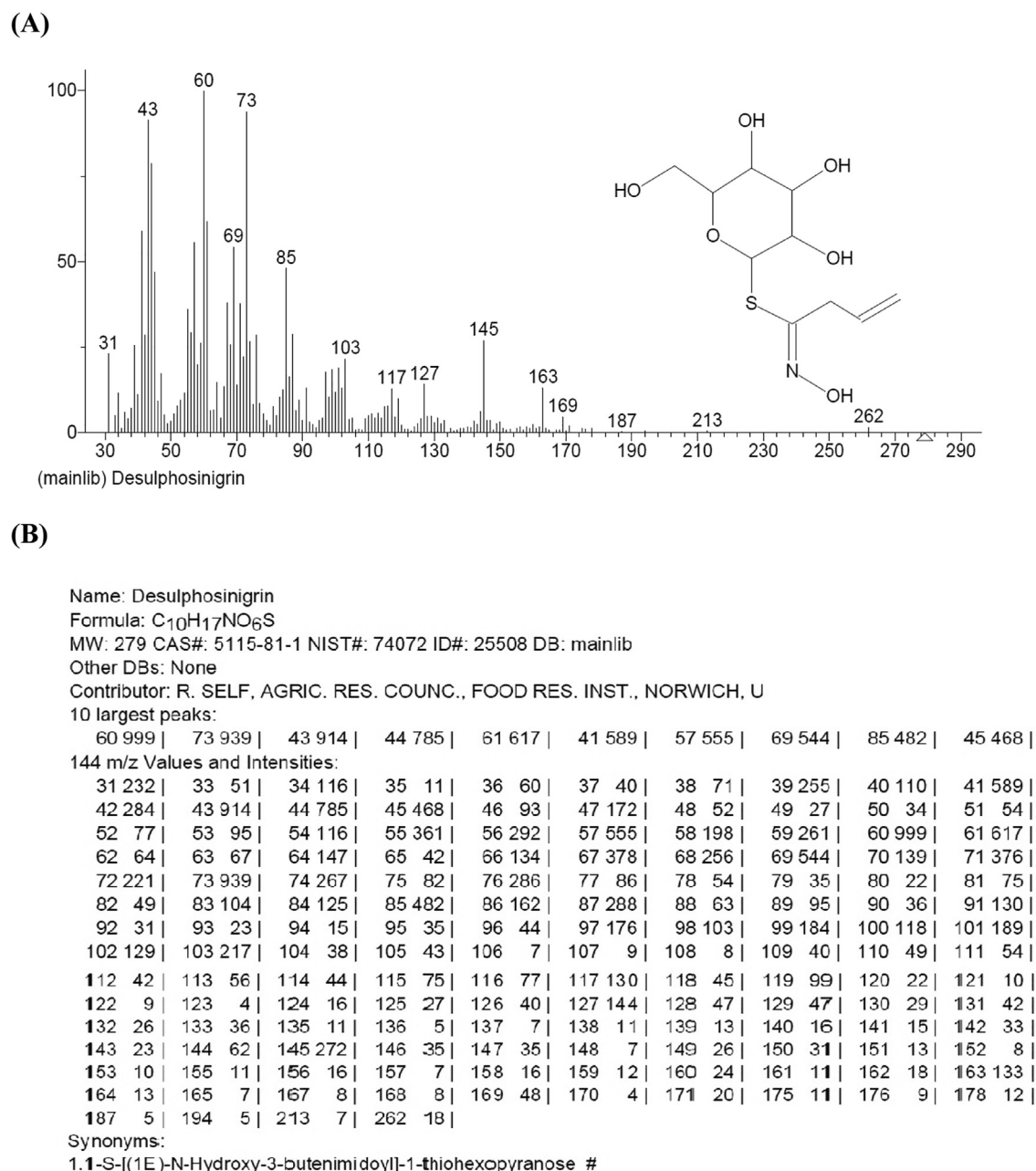


Fig. 8 Mass Analysis. (A) Mass Spectra of Desulphosinigrin, (B) Mass data of Desulphosinigrin.

RMSD, RMSF, Rg, and hydrogen bonds, give more information about how the enzyme and ligand interact (Sneha and Doss 2016). The RMSD shows a broad view of the stability of a structural complex. Therefore, an RMSD study for α -Glc was performed in the presence of DES, ACR, and free. According to our results, it can be seen that both complexes (α -Glc-DES complex and α -Glc-ACR complex) have maintained their equilibrium concerning the single enzyme and show a slight deviation during 100 ns (Fig. 7A). The RMSD level in α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex fluctuated around the mean values of 0.1719 ± 0.0112 , 0.1791 ± 0.0141 , and 0.1856 ± 0.0174 respectively, which are almost similar. The lower RMSD level of the α -Glc-DES com-

plex shows more stability than that of the α -Glc-ACR complex.

RMSF studies the flexibility among the α -Glc in the presence of DES and ACR. Fig. 7B displayed the RMSF level of the α -Glc free, α -Glc-DES complex, and α -Glc-ACR complex. From the RMSF level, it was observed that the α -Glc-DES complex displayed a lesser fluctuation than the α -Glc-ACR complex illustrating the restricted movements during the simulation.

Hydrogen bonds play an important role in the stability of the enzyme structure. Analysis of hydrogen bonds was performed for both complexes (α -Glc-DES complex and α -Glc-ACR complex) and the results are shown in Fig. 7C. Our stud-

ies showed that both complexes contained a lot of hydrogen bonds during the simulation period, but the α -Glc-DES complex had a higher number of hydrogen bonds. Therefore, the α -Glc-DES complex is more stable in the structure of the enzyme.

The Rg value indicates the dimensions of the system and the amount of compactness (Sneha and Doss 2016). From the Rg, it can be seen that the protein is compacted or enlarged during the simulation. As shown in Fig. 7D, the value of the Rg level of the α -Glc-ACR complex started at 2.38 and continued up to 2.49, while the α -Glc-DES complex started at 2.38 and continued up to 2.45. Therefore, fewer fluctuations have taken place in the α -Glc-DES complex. It is clear from the figure that the α -Glc-DES complex shows a lower Rg level than the α -Glc-ACR complex, indicating slight structural changes during the simulation period.

3.9. Desulphosinigrin data

DES is a thiohexopyranose compound found naturally in plants such as *Hybanthus enneaspermus*, *Arabidopsis thaliana*, *Vitis vinifera*, *Melastoma malabathricum*, *Euphorbia lathyris*, and *Annona reticulate* (Andersson et al., 2009, Rout and Kar 2014, Kadhim et al., 2017). Antibacterial, antiasthmatic, and anticancer properties of DES have been reported. It is also known as an inhibitor of Cyclin-dependent kinases and urease enzymes (Nile et al., 2021). However, the inhibition of this compound has not been investigated on the α -Glc enzyme. Due to the better docking score and the optimal parameters of the α -Glc-DES complex simulation, it is expected that by performing *in vitro* and *in vivo* tests can be introduced as an anti-diabetic compound. Mass Spectra and data of Desulphosinigrin are given in Fig. 8.

4. Conclusions

Our present work confirmed the phenolic and flavonoid contents, α -Glc inhibitory effect, and antioxidant capacity of methanolic flower extract of *D.sophia*. The results displayed that flower extract of *D.sophia* was an effective α -Glc inhibitor with the IC₅₀ was 20.34 \pm 0.11 mg/ml, and this inhibiting process was mediated through a mixed inhibition pattern. Moreover, *In vivo* investigation exhibited that flower extract of *D.sophia* has a significant effect on decreasing the blood glucose level in diabetic rats. The GC/MS analysis identified twenty-one compounds. Molecular docking of these compounds displayed that the DES compound has a better docking score comparing other compounds. The analysis of MD simulations illustrated that the interaction of DES to α -Glc was even more favorable than ACR. Therefore, *in vivo* and *in vitro* studies may be used as the next step to understand its practical application in diabetes management. Furthermore, this study proposed the flower extract of *D.sophia* might be introduced as a suitable additive in syrups or foods with anti-diabetic capacity.

Declaration of Competing Interest

The authors declare that there are no conflicts of interest.

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