Research Article

Wound Healing Potential and In Silico Appraisal of *Convolvulus arvensis* L. Methanolic Extract

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Received 3 August 2022; Revised 21 October 2022; Accepted 28 October 2022; Published 24 November 2022

1. Introduction

Wound healing is an intricate pathophysiological process involving parallel roleplay of various biochemical and cellular pathways to reestablish cutaneous and subcutaneous tissues. Apart from that, various enzymatic pathways also activate during this process to accomplish tissue repair [1, 2]. This multifaceted physiological process involves different limitations, including bacterial infection. It also becomes challenging in the presence of many physiological disorders, for example, rheumatoid arthritis, zinc deficiency, and diabetes [3]. The neuropathy associated with diabetes diminishes the blood flow across the site of the wound, resulting in retardation of the healing process. Wound healing in diabetic patients, particularly in the form of a diabetic foot ulcer, is an overwhelming concern for health care authorities around the world. In various populations around the world, up to 25% diabetic patients have been reported to have a diabetic foot ulcer, which can consume as high as 25% of the health care budget pledged for diabetes [4].

Plant-based natural products are the basic source of traditional herbal medicines. These substances are derived from plants with the least possible industrial processing. People are primarily dependent on this health care facility throughout the world [2]. The market for these products has
mounted up to a staggering US$ 60 billion per annum (WHO traditional medicine strategy, 2002). The occurrence of various plant-derived life-supporting constituents, with significant physiological benefits, has inspired researchers to scrutinize these plants in order to regulate potential wound healing activities [5]. Numerous phytochemicals and their corresponding plant extracts have proven promising role in wound healing and are used as an alternative due to their diverse active ingredients and limited side effects [6, 7]. Various studies have reported the wound healing properties of different phytochemicals including flavonoids, alkaloids, terpenes, and carbohydrates [8, 9].

Convolvulus arvensis L. (C. arvensis) is a yearly or occasionally perennial climber weed with a wide distribution all over the Europe and Asia belonging to family Convolvulaceae [10]. Traditionally, the flower is used as a tea infusion and also in the treatment of wounds, skin ulcers, inflammation, and fever, whereas the leaf can be helpful during the menstrual period [10, 11]. Phytochemical research on this plant exhibited the presence of saponins, flavonoids, alkaloids, and lipids [12]. Previous studies have shown the abundant presence of benzoic acid along with different phenolic compounds comprising p-hydroxybenzoic acid, vanillin acid, syringic acid, and ferulic acid in C. arvensis extract [13]. Moreover, it is reported the presence of neophytadiene, hexadecanamide, 9-octadecanamide, 2-benzendicarboxylic acid, 9-octadecanamide, stigment-5-en-3-ol, stearic acid, and hexadecanamide, 9-octadecanamide, 2-benzendicarboxylic acid [13].

2. Materials and Methods

2.1. Collection of Plant and Its Authentication. The stem portion of C. arvensis was collected (March 2018) and recognized, and its authenticity was approved by the taxonomist Dr. Mansoor Hameed (Botany Department, University of Agriculture Faisalabad), and a voucher specimen (244-2-18) was deposited in the herbarium.

2.2. Preparation of Methanol Extract. To prepare the methanol extract of C. arvensis stem (CaME), the fresh stem was shade dried and powdered by means of an electric mixer. The powder weighing 50 g was macerated in 100 mL methanol for 20 hours at standard room temperature. The mixture was then passed through a fine muslin cloth following its filtration through Whatman No. 1 filter paper. The remainder filtrate was dried on a rotary evaporator at 40°C under vacuum to evaporate the organic solvent and then freeze-dried for complete dryness [29, 30].

2.3. HPLC Analysis. For quantification of phenolic compounds, analytical HPLC system (Shimadzu Corporation, Kyoto, Japan) consisting of LC-10AT pump, UV visible detector SPD-10-AV, and CLC-ODS Shim-pack C18 column (250 mm × 4.6 mm, 5 μm) was used. The data acquisition was performed using CSW32 software. The HPLC grade water, acetic acid, and acetonitrile were used as mobile phases. The separation was carried out by using acetic acid (1 M, pH 2.3) as solvent A and acetonitrile as solvent B (0 – 15 min = 15% B, 15 – 30 min = 45% B, and 30 – 45 min = 100% B). The elution was carried out at 25°C with the flow rate of 1 mL/min. The volume injection was 10 μL, and 280 nm was selected as the detector wavelength. The peaks were identified with respect to the retention time, and the concentrations were calculated by comparing with the corresponding standard curve [31].

2.4. GC-MS Analysis. The GC-MS analysis was performed on Agilent 6890 GC system coupled with an Agilent 5973N MSD operational at 70 eV. The ion source temperature was kept 200°C along with split inoculation (1 μL injection volume, split ratio 50:1). HP-5MS-fused silica capillary column (30 m × 0.25 mmID × 0.25 μm film, Agilent J&W, USA) was used. The oven temperature was varied from 100 to 275°C at 10°C/min for 20 min. As a carrier, helium gas was used at a continuous flow rate of 1 mL/min; data acquisition was done by Agilent GC/MSD CS Version D.02.00. The identification of C. arvensis constituents was
based on the direct association of the retention times and mass spectral records with standard compounds and NIST/EPA/NIH mass spectral library stored on computer [14].

2.5. Antioxidant Assays. The 2,2-diphenyl-1-picrylhydrazyl (DPPH) radical scavenging activity, hydrogen peroxide (H$_2$O$_2$) scavenging assay, and ferric reducing power assay of CaME were performed according to the published protocols with minor modifications [29]. For a fair comparison, test extract or standard ascorbic acid solutions (1 mg/mL) were mixed with reagent solutions to achieve an absolute concentration of 0.2 mg/mL in every case.

2.5.1. DPPH Assay. Methanolic solutions of CaME test sample (1 mg/mL), ascorbic acid (1 mg/mL) as standard, and DPPH (0.3 mmol/L) were used. The test sample or ascorbic acid solution was mixed with DPPH solution (1:4) and incubated for half an hour at room temperature in darkness, and absorbance was noted at 517 nm. The control solution contained methanol and DPPH solution (1:4). The percentage of DPPH radical scavenging activity was evaluated with the following relation:

\[
\text{Inhibition of DPPH radical} = \frac{A_c - A_t}{A_c} \times 100, \tag{1}
\]

where $A_c$ is the absorbance of the control and $A_t$ is the absorbance of the test sample. The experiment was repeated triple times with results represented as mean ± S.D.

2.5.2. FRAP. The ferric reducing antioxidant potential (FRAP) of CaME was measured in a manner described by Benzie and Strain [32]. GaME and standard ascorbic acid solutions (1 mg/mL) were prepared in phosphate buffer (50 mmol/L, pH 6.7) and mixed with 1% w/v potassium ferri cyanide (1:3.5) and were centrifuged for 10 min at 3000 rpm. The supernatant was mixed with 0.1% w/v FeCl$_3$ (1:0.1) and the absorbance was measured at 593 nm. The control was prepared by adding potassium ferricyanide and FeCl$_3$ to the blank phosphate buffer in a similar way. The ferric reducing power was calculated using the following relation:

\[
\text{Ferric reducing antioxidant potential (FRAP)} = \frac{A_c - A_t}{A_c} \times 100, \tag{2}
\]

where $A_c$ describes the absorbance of the control and $A_t$ is the absorbance of the testing sample. The experiment was performed in triplicate and outcomes were represented at mean ± S.D.

2.5.3. H$_2$O$_2$ Scavenging Assay. The phosphate buffer (50 mmol/L, pH 7.3) was used to prepare 40 mmol/L H$_2$O$_2$ solution. The confirmation of concentration of the prepared solution was done by taking the absorbance at 230 nm. CaME and standard ascorbic acid solutions (1 mg/mL) were prepared in distilled water. The test sample and H$_2$O$_2$ solutions were mixed (1:4) and absorbance was determined at 230 nm after 10 min. The phosphate buffer solution minus H$_2$O$_2$ was used as black and the % of H$_2$O$_2$ scavenging was calculated using the following relation:

\[
\text{Hydrogen peroxide scavenged} = \frac{A_c - A_t}{A_c} \times 100, \tag{3}
\]

where $A_c$ is the absorbance of the control and $A_t$ is the test sample absorbance. The experiment was repeated 3 times and results were denoted at mean ± S.D.

2.6. Acute Dermal Toxicity Testing. The acute dermal toxicity trial was conducted according to the test guidelines 402 of the Organization for Economic Cooperation and Development (OECD) [33]. The adult nonpregnant nulliparous female rats of albino specie with intact skin, weighing 230-250 g, were arbitrarily selected. Animals were adjusted to the laboratory environment for at least five days before proceeding to the test in standard conditions of temperature and relative humidity. All animals were given water and feed during the experiment. The dorsal area of the trunk of the animals was shaved to remove fur at least 24 hours before the test.

The plant extract at a dose of 2000 mg/kg body weight was applied evenly over the uncontaminated clean skin of each rat. A porous gauge dressing and nonirritating tape was used to cover the area throughout the 24 h treatment exposure. The similar procedure was used for the control group treated with vehicle. Both groups were observed frequently on the first day and then a vigilant clinical analysis was done at least once every day for a total of 14 days. After the exposure time ends, the residual extract was carefully removed with water and observed for any sign of the local skin reaction. The cage side interpretations and weights of the animals were also examined.

2.7. Animal Grouping. The rats of any sex with weight ranging from 200 to 250 g were attained from the animal house of the Department of Pharmacology, Government College University Faisalabad and placed in four groups each of five rats ($n = 5$). Before the commencement of study, the test animals were familiarized with the laboratory surroundings for fourteen days and given typical diet and water ad libitum. Group 1 was served with the blank ointment formulation which served as reference control, group 2 was treated with the standard gentamycin ointment (0.1%), and groups 3 and 4 were treated with the ointment batches containing 10% and 20% CaME. Excision wound was made of 177 mm$^2$ area and 2 mm thickness with a round seal in rats pre-anesthetized with ketamine and xylazine according to previous method [34].

The experimental research was conducted after receiving approval from the Institutional Review Board (Reference No. GCUF/ERC/1981; 27-6-2018) of the Government College University, Faisalabad, Pakistan, and the experiments were conducted according to the regulations of the Institute of Laboratory Animal Resources, Commission on Life Sciences, National Research Council, Washington DC, USA (1996).
2.8. Excisional Wound Model. British Pharmacopoeia (B.P.) grade petroleum jelly was used to prepare ointment formulations of gentamycin (0.1%) and CaME (10 and 20%) while plain petroleum jelly was used as a blank. A 15 mm diameter of the round ethanolic seal was applied to the margins of the depilated central trunk for sterilization. The wound excision was performed under I.V. anesthesia of ketamine (120 mg/kg body weight). The skin was removed from marked area via inflicted to obtain a wound having approximately 177 mm² diameter. A complete hemostasis was achieved by blotting the wound with a cotton swab soaked in warm normal saline. The topical application of test and control ointments was performed at 24h intervals until the complete wound closure. The wound area was measured with a luminous paper and afterwards estimated on a graph page every 4th day up to completion of epithelialization and wound closing was measured [35].

The two parameters for wound healing are percentage wound contraction rate and epithelialization period. Percent wound contraction was measured using the following formula:

\[
\text{Percentage wound contraction} = \frac{\text{Initial wound size} - \text{Wound size on a specific day}}{\text{Initial wound size}} \times 100.
\]

As epithelialization takes days, so wound epithelialization was done for several days until there was the fall of scale with no raw wound behind. Histopathological study of the skin tissues using hematoxylin-eosin staining was performed for the further wound study [36].

2.9. Statistical Analysis. The results were stated as mean ± standard deviation (S.D.). Antioxidant activities of CaME were evaluated by one-way ANOVA. Data pertaining to percentage wound contraction were investigated by two-way ANOVA followed by Tukey's post hoc test using GraphPad Prism software version 7.0.

2.10. Molecular Modeling Studies. To discover the binding manner and molecular interactions of identified compounds with therapeutic targets, molecular docking studies were active using AutoDock Vina with default settings and parameters [37]. The X-ray-resolved crystal structures of potential targets, including TGF-β complexed with staurosporine (PDB ID: SE8W) [38] and GSK-3β complexed with N-(6-(3,4-dihydroxyphenyl)-1H-pyrazolo[3,4-b]pyridine-3-yl)acetamide; B4K) (PDB ID: 5OY4) [39], were retrieved from the PDB with good resolution and R-free factor. For β-catenin and c-myc, a complex structure of β-catenin with hTcf-4 (PDB ID: 1JDH) [40] and c-myc-max recognizing DNA (PDB ID: 1NKP) [41] crystal structures were extracted from the PDB.

For structural analysis, cocrystralized ligands were divided from the equivalent proteins and redocked another time to compute the root mean square deviation (RMSD) of docked and cocrystralized ligand for the reliability of an AutoDock Vina protocol [37]. Since no cocrystralized ligand was reported against β-catenin and c-myc, previously high-throughput screening studies were considered to study the binding pockets. For β-catenin, binding pocket residues reported to interact with Tcf-4 (PDB IDs: 1JPW and 1JDH) [42] were selected to generate a reliable docking grid, whereas the region Arg363-Ile381 along with the loop Pro382-Lys392 of c-myc was defined as the active site for ligand binding, as reported in previous molecular modeling studies [43, 44]. The respective binding sites were used for molecular docking procedure. The general docking procedure of protein preparation and its minimization and optimization have been described elsewhere [27, 45, 46]. The molecular structures of identified active compounds were produced by ChemDraw Professional (v. 16.0). AutoDockTools was used to merge nonpolar hydrogens and add Gas- teiger charges, docked complexes were inspected utilizing UCSF Chimera (v. 13) [47], and 2D molecular interactions were examined using ligplot [48].

The dynamic stability of the binding pose with respect to time was investigated via MD simulations over a period of 10 nanoseconds (ns). All MD simulations were performed using AMBER 16 simulation package [49]. After minimization and equilibration protocols, the simulated system with explicit solvent molecules (TIP3) was submitted to a production run of 10 ns under standard temperature (300 K) and pressure (1 bar). We applied the same MD simulation procedure as defined in prior studies [24]. For the trajectory analysis, the CPPTRAJ module of AMBER 16 [50] was collected with a time interval of 2 picoseconds (ps).

3. Results

3.1. HPLC Analysis. The HPLC analysis of the CaME showed seven compounds. Ferulic acid was in the highest concentration among all the compounds detected. The

| Peak no. | Compound          | Retention time (min) | Area (mV.s) | Area (%) | Amount (ppm) |
|---------|-------------------|----------------------|-------------|----------|--------------|
| 1       | Quercetin         | 2.813                | 9.807       | 0.5      | 0.51         |
| 2       | Gallic acid       | 4.967                | 34.728      | 1.7      | 1.24         |
| 3       | Caffeic acid      | 12.880               | 65.561      | 3.2      | 3.13         |
| 4       | Vanillic acid     | 13.687               | 101.001     | 4.9      | 6.26         |
| 5       | p-Coumaric acid  | 17.380               | 136.218     | 6.6      | 1.56         |
| 6       | Ferulic acid      | 22.533               | 322.329     | 15.5     | 23.73        |
| 7       | Cinnamic acid     | 24.653               | 127.464     | 6.1      | 4.45         |
Table 2: GC-MS analysis of the methanolic extract of C. arvensis stem.

| Peak no. | Compound name                              | Retention time (min) | Pharmacological use               |
|---------|--------------------------------------------|----------------------|----------------------------------|
| 1       | Isoeugenol                                 | 10.787               | Flavouring agent                 |
| 2       | 2-Hexadecyllicosan-1-ol                     | 14.635               | For the treatment of blisters or cold sores |
| 3       | 7-Hexadecenoic acid                         | 16.593               | Anti-inflammatory                 |
| 4       | 1,2-Benzenedicarboxylic acid, 1,2-bis(7-methyloctyl) ester | 19.206               | Thyroid-disrupting agent         |
| 5       | Octadecyl 3-(3,5-di-tert-butyl-4-hydroxyphenyl) propionate | 20.939               | Antioxidant                      |

Table 3: Antioxidant analysis of C. arvensis stem methanolic extract (CaME) by 2,2-diphenyl-1-picrylhydrazyl (DPPH), ferric reducing antioxidant power (FRAP), and hydrogen peroxide (H2O2) scavenging assay.

| Tested samples | DPPH (% inhibition) | FRAP (% activity) | H2O2 (% scavenged) |
|----------------|----------------------|-------------------|---------------------|
| CaME           | 21.296 ± 1.223****   | 1.053 ± 0.11      | 2.405 ± 0.127****   |
| Ascorbic acid  | 78.703 ± 4.245       | 0.923 ± 0.204     | 59.340 ± 3.377      |

The triplicate values were expressed as mean ± S.D. Results were significant at ****P < 0.0001 with respect to the control group.

Table 4: Effect of standard drug and methanolic extract from C. arvensis stem (CaME) on percentage wound contraction and epithelialization period on excision wound in rats.

| Groups   | 0 d | Wound size (mm²) at different days (% healing) | Epithelialization |
|----------|-----|-----------------------------------------------|-------------------|
|          | 0 d | 4 d 8 d 12 d 16 d 20 d                        |                   |
| Control  | 177 ± 0.00 (100%) | 117 ± 2.44 (33.89%) | 59.8 ± 1.71 (66.21%) | 35 ± 1.22 (80.22%) | 26.4 ± 0.98 (85.08%) | 11.6 ± 0.83 (93.42%) | 23.4 |
| Standard | 177 ± 0.00 (100%) | 113 ± 0.00**** (36.15%) | 54.2 ± 1.71**** (69.37%) | 30 ± 1.22**** (83.05%) | 0.54 ± 0.14**** (99.69%) | 0.00 ± 0.00**** (100%) | 17.2 |
| 10%      | 177 ± 0.00 (100%) | 139 ± 2.44**** (21.46%) | 70 ± 3.64**** (60.45%) | 35 ± 1.22**** (78.53%) | 28 ± 0.00**** (84.18%) | 7.1 ± 0.00**** (95.98%) | 21.6 |
| 20%      | 177 ± 0.00 (100%) | 119 ± 2.44* (32.76%) | 41.6 ± 1.47**** (76.49%) | 22.4 ± 0.98**** (87.34%) | 1.18 ± 0.25**** (99.33%) | 0.00 ± 0.00**** (100%) | 17.6 |

The values are expressed as mean ± standard error (n = 05) in each group. The results were significant at ****P < 0.0001 and *P < 0.05 with respect to the control group and * * * * P < 0.0001 with respect to standard therapy.

Figure 1: Pictorial representation of the wound healing process up to day 20 treated with standard, 10%, and 20% CaME ointment and control group.
compounds were in the following increasing order of concentration: ferulic acid > vanillic acid > cinnamic acid > caffeic acid > p-coumaric acid > gallic acid > quercetin (Table 1).

3.2. GC-MS Analysis. GC-MS analysis of CaME identified isoegenol (10.787 min), 2-hexadecylicosan-1-ol (14.635 min), 7-hexadecenoic acid (16.593 min), 1,2-benzenedicarboxylic acid 1,2-bis(7-methyloctyl) ester (19.206 min), and octadecyl-3-(3,5-di-tert-butyl-4-hydroxyphenyl)-propionate (20.939 min). The retention times, exact masses, and chemical formulas of all compounds are shown in Table 2.

3.3. Antioxidant Assays. The CaME exhibited notable antioxidant potential (Table 3). CaME induced 21.296% DPPH inhibition, 1.053% FRAP activity, and 2.405% H₂O₂ scavenging. Standard ascorbic acid exhibited statistically (P < 0.0001) higher DPPH inhibition and 59.34% H₂O₂ scavenging as compared with CaME. The experiments were carried out at the same final concentration of 0.2 mg/mL CaME or ascorbic acid. It is evident that CaME depicted slightly higher FRAP activity, but statistically comparable to the standard. DPPH inhibition of the extract was approximately 4 times lower than the standard and H₂O₂ scavenging was not significant.

3.4. Acute Dermal Toxicity. The results from the acute dermal toxicity trial of CaME produce no mortality during the 14-day observation period. Likewise, no deformities were noticed during gross interpretations on the skin and behavioral patterns. After 1-, 7-, and 14-day postexposure to the extract, there were no signs of edema, erythema, or any signs of toxicity on the skin. The differences in increase of body

Table 5: Binding affinities (in kcal/mol) isolated compounds from the methanolic extract of C. arvensis stem as calculated from AutoDock Vina.

| Compound                                           | c-myc | β-Catenin | TGF-β | GSK-3β |
|----------------------------------------------------|-------|-----------|-------|--------|
| Quercetin                                          | -6.8  | -6.5      | -9.2  | -8.1   |
| 1,2-Benzenedicarboxylic acid 1,2-bis(7-methyloctyl) ester | -6.3  | -6.2      | -7.4  | -6.5   |
| Cinnamic acid                                      | -5.7  | -5.9      | -6.2  | -5.6   |
| Caffeic acid                                       | -5.4  | -5.8      | -6.4  | -5.9   |
| Ferulic acid                                       | -5.2  | -5.8      | -6.7  | -5.8   |
| Isoeugenol                                         | -5.2  | -6        | -6.7  | -5.7   |
| p-Coumaric acid                                    | -5.1  | -5        | -6.2  | -5.9   |
| 7-Hexadecenoic acid                                | -5.1  | -5.1      | -6.2  | -5.4   |
| Octadecyl 3-(3,5-di-tert-butyl-4-hydroxyphenyl) propionate | -5    | -5.2      | -6.9  | -5.5   |
| Gallic acid                                        | -5    | -5.2      | -6    | -5     |
| Vanillic acid                                      | -4.9  | -5.2      | -6.9  | -5.4   |
| 2-Hexadecylicosan-1-ol                             | -4.9  | -5.8      | -6.1  | -5.1   |

Figure 2: Histopathological results of healed skin on 17th day postwounding. (a) Negative control group, (b) ointment containing 10% CaME, (c) standard gentamycin group, and (d) ointment containing 20% CaME.
Figure 3: Molecular modeling analysis of cmd1 (quercetin). (a) The favorable docked conformation of cmd1 (grey stick) together with cocrystallized staurosporine (yellow stick) (PDB ID: 5E8W) inside the binding pocket of TGF-β. (b) The same representation of quercetin complexed with GSK-3β together with a cocrystallized B4K (PDB ID: 5OY4). (c) Root mean square deviation of TGF-β/cmd1 and GSK-3β/cmd1 (d) over a period of 10 ns molecular dynamic simulations. (e) 2D-ligplot molecular interaction analysis of TGF-β/cmd1 and GSK-3β/cmd1 (f) after MD simulation together with corresponding cocrystallized complexes. The hydrogen interacting residues (labeled green) are displayed in stick and hydrophobic interacting residues (labeled black) are highlighted with red spoked arcs, and the distance for H-bonds are highlighted in Angstrom Å (default set < 3 Å).
Figure 4: Continued.
weight percentage from 0 to 14th day after exposure were also observed insignificant.

3.5. Excisional Wound Model. The results of excisional wound healing of different rat groups treated with standard gentamycin (0.1%), 10% CaME, and 20% CaME ointments in comparison with the negative control are described in Table 4. The group treated with 20% CaME had substantially accelerated wound closure resulting in the reduction of the average epithelialization period to 17.6 days compared to the 10% CaME and blank ointment treatments with an average epithelialization periods of 21.6 and 23.4 days, respectively. The percentage of wound contraction of standard and 20% CaME ointment is almost similar as shown in Figure 1. Almost 80% of healing in wound was documented through extract and groups treated by antibiotic for 12 days. Although 100% healing was obtained in the C. arvensis (20%) ointment and standard group after 16th day, which was significantly different ($P < 0.0001$) from healing percentages in the other extract-treated group, the percentage healing in the control group was significantly lower than that of standard and 20% ointment-treated group.

Moreover, a histopathology study of the 20% CaME ointment treated group exhibited significant healing as compared to the control group (Figure 2). It is visible that the skin architecture of standard gentamycin and 20% CaME treatments was similar after 17 days.

3.6. Molecular Modeling Interpretations. The most representative conformation of ligands from the highest cluster was selected as the best docking pose inside the binding pocket of β-catenin and c-myc. The predicted binding affinities of identified active compounds of C. arvensis are tabulated in Table 5. The docking study revealed fairly good binding affinities against all four targets, including GSK-3β, TGF-β, β-catenin, and c-myc ranging from -8.1 to -5.1 kcal/mol, -9.2 to -6.1 kcal/mol, -6.5 to -5 kcal/mol, and -6.8 to -4.9 kcal/mol, respectively. Among all active compounds, quercetin and 1,2-benzenedicarboxylic acid 1,2-bis(7-methyloctyl) ester designated as cmd1 and cmd2, respectively, were ranked top 2. These compounds showed significant binding affinities against GSK-3β (-8.1 and -6.5 kcal/mol), TGF-β (-9.2 and -7.4 kcal/mol), β-catenin (-8.1 and -7.8 kcal/mol), and c-myc (-7.4 and -6.7 kcal/mol). The stability of both compounds, as being the highest binding affinities, inside the binding groove of all four targets was estimated using MD simulations to analyze the RMSD over a period of 10 ns. The overall binding interaction analysis with corresponding proteins together with their cocrystallized ligands is displayed in Figures 3–6.

3.7. Binding Interactions of cmd1 (Quercetin). Overall, the binding pose of quercetin was deeply found in the binding pocket organized along parallel cocrystallized staurosporine [38] and N-(6-(3,4-dihydroxyphenyl)-1H-pyrazol[3,4-b]pyridine-3-yl) acetamide (B4K) [39] in TGF-β and GSK-3β, respectively (Figures 3(a) and 3(b)). The generated RMSD plot of TGF-β during the entire simulation period of 10 ns revealed the stability of quercetin within the binding pocket followed by small fluctuations in the start (Figure 3(c)) and Ca-backbone stability of TGF-β which also remain converged between ~1 Å, whereas quercetin showed small initial fluctuations inside the binding pocket of GSK-3β which also triggered some dynamic impact on
Figure 5: Molecular modeling analysis of cmd2 (1,2-benzenedicarboxylic acid 1,2-bis(7-methyloctyl)). (a) The favorable docked conformation of cmd2 (grey stick) together with cocrystallized staurosporine (yellow stick) (PDB ID: 5E8W) inside the binding pocket of TGF-β. (b) The same representation of cmd2 complexed with GSK-3β together with a cocrystallized B4K (PDB ID: 5OY4). (c) Root mean square deviation of TGF-β/cmd2 and GSK-3β/cmd2 (d) over a period of 10 ns molecular dynamic simulations. (e) 2D-ligplot molecular interaction analysis of TGF-β/cmd2 and GSK-3β/cmd2 (f) after MD simulation together with corresponding cocrystallized complexes. The hydrogen interacting residues (labeled green) are displayed in stick and hydrophobic interacting residues (labeled black) are highlighted with red spoked arcs, and the distance for H-bonds are highlighted in Angstrom Å (default set < 3 Å).
Figure 6: Molecular modeling analysis of cmd2 (1,2-benzenedicarboxylic acid 1,2-bis(7-methyloctyl)). (a) The favorable docked conformation of cmd1 (grey stick) inside the binding pocket of β-catenin. (b) The same representation of quercetin complexed with c-myc. (c) Root mean square deviation of β-catenin/cmd1 and c-myc/cmd1 (d) over a period of 10 ns molecular dynamic simulations. (e) 2D-ligplot molecular interaction analysis of β-catenin/cmd1 and c-myc/cmd1 (f) after MD simulation. The hydrogen interacting residues (labeled green) are displayed in stick and hydrophobic interacting residues (labeled black) are highlighted with red spoked arcs, and the distance for H-bonds are highlighted in Angstrom (Å) (default set < 3 Å).
backbone atoms of protein; however, later, the complex remained stable from 4 to 10 ns as seen in Figure 3(d). In complex with β-catenin and c-myc for the duration of 10 ns simulation period as displayed in Figure 4, quercetin stayed constant inside the binding pocket β-catenin/quercetin showing Ca-backbone stability, whereas quercetin elicited higher fluctuations in Ca-backbone of c-myc.

The 2-dimensional interaction study after 10 ns discovered the influence of hydroxyl groups of quercetin in establishing H-bonds (<3 Å distance), and most residues were interacted over hydrophobic interfaces, lining the binding position of TGF-β and GSK-3β, respectively. As seen in the binding pose of quercetin with both proteins (Figures 3(e) and 3(f)), the 3' and 4'-hydroxyl group of dihydroxyphenyl moiety established H-bonds with sidechain OE2 atom of Glu245 (2.70 Å), backbone N atom of Lys232 (2.61 Å) in TGF-β, and sidechain OD2 atom of Asp200 (2.21 Å) in GSK-3β. Moreover, tetrahydroxycromen-4-one moiety was bound deep inside the groove with the hydroxyl group hydrogen-bonded to three adjacent residues, Ser280 (2.38 Å), Asp281 (2.02 Å) and His283 (2.45 Å) of TGF-β, and Asp200 (2.29 Å) of GSK-3β. Other than H-bonds, a huge number of conserved nonhydrophilic interactions were also detected in TGF-β/cmd1 (Figure 3(e)) and GSK-3β/cmd1 (Figure 3(f)) complexes.

Similarly, quercetin established H-bonds mainly through terminal hydroxyl groups in complex β-catenin and c-myc. Figures 4(e) and 4(f) showed that 3' and a 4'-hydroxyl group of dihydroxyphenyl moiety established H-bonds with sidechain NE2 and NE atom of Gln302 (2.70 Å) and Arg342 (1.98 Å) in β-catenin and backbone O and N atom of Arg914 (2.32 Å) in c-myc, whereas tetrahydroxycromen-4-one moiety established a network of hydrogen bonds with two adjacent arginines, Asp213 (2.61 Å), Arg914 (2.88 Å), and one nearby Glu910 (1.96 Å) of c-myc. Along with H-bonds, enormous amount of hydrophobic interactions was also witnessed in β-catenin/quercetin (Figure 4(e)) and c-myc/quercetin (Figure 4(f)) complexes.

3.8. Binding Interactions with cmd2. Likewise, the cmd2 was present deep within the binding pocket along with parallel cocrystallized staurosporine [38] and B4K [39] in TGF-β and GSK-3β, respectively (Figures 5(a) and 5(b)). The RMSD trajectory plot showed the stability of cmd2 inside the binding pocket followed by the convergence of Ca-backbone stability of TGF-β between ~0.5 Å from 3 to 10 ns (Figure 5(c)). However, cmd2 complexed with GSK-3 experienced some fluctuations in proteins’ backbone atoms but converged between ~1.25 Å throughout the simulation (Figure 5(d)). The RMSD deviation in GSK-3 β/cmd2 complex was evident from the flexibility resulted in the open conformation of bis-7-methyloctyl ester (Figure 5(b)) to attain a more favorable conformation inside the pocket which triggered fluctuations in protein backbone. Likewise, in complex with β-catenin and c-myc, cmd2 remained stable internally in the binding pocket for the duration of whole simulation phase (Figures 6(a) and 6(b)) but showed slight fluctuations due to the open conformation of bis-7-methyloctyl ester (Figures 6(c) and 6(d)). β-catenin/cmd2 showed Ca-backbone stability, whereas cmd2 elicited fluctuations in c-myc Ca-backbone.

Further insight into the molecular interactions of cmd2 with both targets, it was found that the benzenedicarboxylic acid interacted mainly through H-bonds while bis-7-methyloctyl ester interacted through a large number of hydrophobic interactions (Figures 5(e) and 5(f)). The oxygen atom of benzenedicarboxylic acid established an H-bond with the backbone nitrogen atom of Ser287 (2.45 Å) in TGF-β and sulfur atoms of Cys199 (2.20 Å) in GSK-3β, whereas both residues showed hydrophobic interactions with their corresponding cocrystallized ligands [38, 39]. The long bis-7-methyloctyl ester of cmd2 established a large network of conserved hydrophobic interaction in TGF-β (Figure 5(e)) which anticipated stable conformation of cmd2 inside the binding pocket (Figure 4(c)) as compared to GSK-3β/cmd2 with lesser hydrophobic interactions and experienced open conformation (Figure 5(f)).

Likewise, in TGF-β and GSK-3β, cmd2 showed a similar interaction pattern in β-catenin and c-myc. Figures 6(e) and 6(f) show that the oxygen atoms of benzenedicarboxylic acid established H-bonds with the sidechain N and OH atoms of Lys345 (2.66 Å) and Tyr306 (2.51 Å) in complex with β-catenin, whereas the benzenedicarboxylic acid established two H-bonds with N atom of Lys939 (2.44 Å and 1.97 Å) and one H-bond with sidechain N atom of Lys918 (2.11 Å) in complex with c-myc. Along with H-bonds, the long bis-7-methyloctyl ester of cmd2 established a large network of hydrophobic interaction with β-catenin (Figure 6(e)) and c-myc (Figure 6(f)), which somewhat anticipated the stability of cmd2 in the respective binding pockets.

For both compounds, the binding site residues interacted in a similar way as seen in cocrystallized complexes, TGF-β/staurosporine (Figure 3(e)) [38] and GSK-3/B4K (Figure 3(f)) [39], and other reported cocrystallized bound inhibitors of TGF-β [51] and GSK-3 [52, 53], respectively. For example, in TGF-β/cmd1 complex, hydrogen-bonded oxygen atoms of Ser280 and Asp281 and backbone nitrogen atom of His283 were conserved as reported in TGF-β/staurosporine [38] together with other hydrophobic interactions including Gly212, Leu340, Ile211, Gly286, Ala230, Leu260, Lys232, and Val219 (Figure 3(e)), likewise in TGF-β/cmd2, which interacted mainly through hydrophobic interactions (Figure 5(e)), whereas the 1H-pyrazolo[3,4-b]pyridine rings of cocrystallized-bound ligand (B4K) of GSK-3 superimposed exactly on tetrahydroxycromen-4-one moiety of quercetin and evidenced conserved hydrophobic interactions with residues, Ala83, Tyr134, Val135 (which interacted through H-bond with B4K), and Leu188 (Figure 3(f)). Likewise, several conserved hydrophobic interactions were observed in GSK-3β/cmd2 (Figure 5(f)).

4. Discussion

The current research was aimed at determining the wound healing potential of *C. arvensis* methanolic extract. Moreover, antioxidant assays and chemical characterization were also performed. The antioxidant potential of the CaME was assessed by the bona fide DPPH inhibition, ferric reducing
antioxidant potential, and H₂O₂ scavenging assays while ascorbic acid was used as a standard. The plant extract exhibited notable antioxidant potential in contrast to ascorbic acid. These promising results could be endorsed to the presence of excessive flavonoid and phenolic contents in GaME [10, 14]. The antioxidant potential is an established therapeutic indicator for accelerating wound healing characteristics of plant extracts [54]. Different medicinal plants having promising wound healing characteristic have been found to possess significant antioxidant activity. It is evident from a wide range of literature that wound healing property of plant extracts coexist with their antioxidant potential [55].

The wound healing action of CaME could be attributed to the promising antioxidant potential of CaME. Another concomitant factor of these results is related to the antibacterial activity of C. arvensis. An ethanolic extract of this plant has been reported to possess strong antibacterial activity towards Staphylococcus aureus, Streptococcus pyogenes, and Escherichia coli [56]. These microorganisms have their proven predominant role in delaying wound healing process [57]. In the current study, CaME exhibiting wound healing potential might be due to the antibacterial action as evidenced from the skin histology slides. Additionally, 20% CaME substantially accelerated wound closure in contrast to 10% dosage of GaME and control group. The healing activity of 20% CaME was insignificantly varied from gentamicin.

Ferulic acid has proven to be effective in treating erythema as it is greatly absorbed in the skin when topically applied [58]. Quercetin may also support speed wound curing; researches have also revealed that quercetin acted on nerve tissues in skin wounds to heal the damage [59]. Gallic acid possesses strong antioxidant activity that causes the nerve tissues in skin wounds to heal the damage [59]. Vanillyl acid has proven to have relatively high permeation due to its flavor and GC-MS. In silico validation of the activity of each complex after 10 ns MD simulations was in line with the reference compounds. Dermal application of an agent can produce a wide range of undesired effects ranging from cutaneous toxicity to general physiological disorders. Plant extracts contain a large number of compounds having different polarities that can not only affect surface skin tissues but also capable of crossing different membrane barriers to reach the bloodstream, resulting in their general physiological effects. This study is the first account of the acute dermal toxicity of C. arvensis. Although the oral ingestion of this plant causes severe gastric and hepatic toxicity [72], the dermal toxicity study exhibited no mortality and toxicity till 14 days.

5. Conclusion

In this study, the presence of pharmacologically promising phytoconstituents in CaME was evident through HPLC and GC-MS. In silico validation of the affinity of these phytoconstituents towards wound healing targets and promising antioxidant activity corroborate wound healing potential of CaME. These outcomes potentiate further investigation of CaME for the treatment of diabetic wound healing. To the best of our knowledge, this is the first account of the experimental validation of the wound healing potential and acute dermal toxicity of the C. arvensis. These outcomes provide the basis of the future direction involving isolation and identification of the active phytoconstituents and their validation for wound healing potential. This study lacked mechanism-based investigation of CaME for wound healing that must be elucidated in further studies. It is suggested that further extension of this work towards validation of the pathways
involving GSK-3β, c-myc, TGF-β, and β-catenin must be carried out, which are responsible for the accelerated wound healing. These findings are economically beneficial because the use of this plant extract for wound healing by local communities will help reduce the health care cost due to reduced usage of health facilities in distant areas.

Data Availability
Authors declare that all the data supporting the findings of this study are included in the article.

Conflicts of Interest
There was no conflict of interest among authors.

Authors’ Contributions
UZ was responsible for conceptualization, visualization, review, supervision, and writing the original draft. SK, SZ, FA, MFK, and LH performed literature, experimentation, and analysis; BA, AS, and MFA were responsible for historical analysis and editing the manuscript.

Acknowledgments
The authors are grateful to Dr. Sarfraz Ahmad and Dr. Muhammad Usman Mirza for providing their services for docking study.

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