

N^α-1, 3-Benzenedicarbonyl-Bis-(Amino Acid) and Dipeptide Candidates: Synthesis, Cytotoxic, Antimicrobial and Molecular Docking Investigation

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Purpose: The objective of our work was to prepare a potent and safe antimicrobial and anticancer agents, through synthesis of several peptides and examine their biological activities, namely as, cytotoxicity potent and antimicrobial and antifungal agents.

Introduction: Multidrug-resistant microbial strains have arisen against all antibiotics in clinical use. Infections caused by these bacteria threaten global public health and are associated with high mortality rates.

Methods: The main backbone structure for the novel synthesized linear peptide is N^α-1, 3-benzenedicarbonyl-bis-(Amino acids)-X, (**3–11**). A computational docking study against DNA gyrase was performed to formulate a mode of action of the small compounds as antimicrobial agents.

Results: The peptide-bearing methionine-ester (**4**) exhibited potent antimicrobial activity compared to the other synthesized compounds, while, peptide (**8**), which had methionine-hydrazide fragment was the most potent as antifungal agent against *Aspergillus niger* with 100% inhibition percent. Compounds (**6** and **7**) showed the highest potency against breast human tumor cell line “MCF-7” with 95.1% and 79.8% of cell inhibition, respectively. The nine compounds possessed weak to moderate antiproliferative effect over colon tumor cell line. The docking results suggest good fitting through different hydrogen bond interactions with the protein residues. In silico ADMET study also evaluated and suggested that these compounds had promising oral bioavailability features.

Conclusion: The tested compounds need further modification to have significant antimicrobial and antitumor efficacy compared to the reference drugs.

Keywords: anticancer, antimicrobial activity, linear-peptides, N^α-1, 3-benzenedicarbonyl-bis-peptides, molecular docking

Introduction

The application of peptide-based drugs for the treatment of different diseases has been widely developed during the last few decades. Peptides and peptide derivatives are inherently active against infectious diseases within biological systems. Moreover, antimicrobial peptides (AMPs) have been used as broad-spectrum antibiotics against several microorganisms including bacteria, fungi, and viruses.¹ The emergence of multi-resistance pathogens is still potentially problematic for the treatment of infectious diseases. Furthermore, weakening of the immune system or inflammation associated with chronic infection makes patients more vulnerable to the emergence of cancer.² Such an association between cancer and infection

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urges the identification of a new class of molecules that are dually-active as broad-spectrum antimicrobial agents and as anticancer agents.³ Promising new molecules, defined as antimicrobial peptides with anticancer properties, should present activities against broad multi-resistance pathogens and the ability to interfere with tumor growth and invasion towards healthy tissues. Physicochemical properties of peptide sequences exhibit dual-activity against pathogens and cancer. For example, the presence of positively charged or bulky hydrophobic amino acids in such sequences electrostatically interacts with negatively charged groups in the bacterial membranes. Furthermore, these amino acids provide the required characteristics to interact with tampered cancer cell membrane, through which the activity of AMPs toward cancer cells is promoted through the interactions of negatively charged phospholipids exposed in the membrane of cancer cells or by interacting with the cell membrane potential and fluidity.⁴ Despite the versatility of peptide design and synthesis to achieve such dual activities, their metabolic and conformational instabilities represent the main obstacles for the successful implementation of peptide-based drugs. This drawback urged us to overcome such obstacles by the design and synthesis of peptide derivatives with rigid conformations and specific amino acid sequences that impart, simultaneously, anticancer and antimicrobial activities. Our research group has published several studies indicating that peptide candidates are promising as potential anticancer drugs,^{5–15} anti-inflammatory and analgesic drugs,¹⁶ and as antibacterial drugs.^{17–28} The incidence of microbial infections in cancer patients increases due to the low circulating neutrophil levels in these patients in addition to the disruption of the microbial flora. Thus, the objective of the current study is to synthesize a new peptide derivative with multiple biological activities including anticancer, anti-inflammatory, and antibacterial activities to be considered as potential candidates used in the treatment of cancer.

Materials and Methods

Chemistry

These companies; Sigma (Ronkonkoma, NY, USA), Fluka (Buchs, Switzerland), and E. Merck (Hohenbrunn, Germany) were used to import all chemical and related materials. All “melting points; M.P.” uncorrected which gained from “Digital melting point apparatus; model: IA9100”. The “Micro-Analytical center” in Cairo

University of Egypt was used for measuring the following spectroscopic analysis; (i) Elemental Micro-analyses for C, N, and H, which gained through a good restriction of the theoretical values ($\pm 0.4\%$). (ii) “Potassium bromide; KBr” disks were handled for preparation tested samples to implement into the “Fourier transforms infrared spectrophotometer; IR Shimadzu; Model: with 1S affinity”. (iii) Mass spectra measured by “Shimadzu gas chromatograph–spectrometer, Model: QP2010ultra; Kyoto, Japan”. NMR-spectra in “DMSO- d_6 ” as proton and carbon were obtained from “JöEL500 MHz instrument”.

Synthesis of N^α -1, 3-Benzenedicarbonyl-Bis-[Amino Acid Methyl Ester]

First Route: The Acid Chloride Method

A DCM of compound **1** (1 mmol, 50 mL) was added dropwise while stirring to a cold (-20°C) dichloromethane solution (50 mL) of free L-alanine and L-methionine ester (2 mmol), obtained by the addition of two equivalent amounts of triethylamine (2 mmol) to the amino acid methyl ester hydrochloride and cold (-20°C) dichloromethane, 50mL, while stirring. The obtained mixture was stirred for an additional 3 hours at (-20°C), and then for 24 hours at room temperature, followed by washes with distilled water, 1 N sodium bicarbonate, 1 N potassium hydrogen sulphate, and distilled water, and then for dried for 24 hours at 0°C over anhydrous sodium sulphate. The volatile materials were evaporated until dry and were then triturated with petroleum ether (boiling point B.P. $40\text{--}60^\circ\text{C}$) to obtain residual material. The obtained precipitate was collected and then recrystallized from methanol (MeOH) to obtain compounds **3** and **4**.

Second Route: The Mixed Anhydride Method

Ethyl chloroformate (2.9 mL, 30.1 mmol), a DCM solution (50 mL) of isophthalic acid (5 g, 30.1 mmol) and triethylamine (6.6 mL, 60.2 mmol), was stirred at -20°C . The reaction mixture was stirred for an additional 30 minutes, and then the DCM solution (50 mL) free of L-alanine and L-methionine ester (8.4 g; 60.2 mmol, -20°C) was added to the dichloromethane (50 mL) under continuous stirring for 7 hours at -20°C , and subsequently for 24 hours at room temperature. The crude product was recrystallized from methanol to obtain pure compounds **3** and **4**; as determined by the melting point and TLC “thin layer chromatography” compared to the original sample prepared according to Method A.

N^α-I, 4-Benzenedicarbonyl-Bis-[L-Ala-OMe], (3)

Yield: [A]: 90; [B]: 80; m.p. 108–110°C. Rf x100 (solvent system) 55(S₂). [α]_D²⁵: - 12.04 (C, 0.02, MeOH). IR in (cm⁻¹; KBr): 3400 (NH-stretching), 3066 (CH-aromatic), 2950 (CH-aliphatic), 1738 (C=O, ester), 1644 and 1534 (C=O, amide I + II). ¹H-NMR (500 MHz, DMSO-*d*₆) δ: 9.04, 8.97 (s, 2H, CONH, D₂O, exchangeable), 8.41 (s, 1H, aromatic H, C₂), 8.15–8.04 (d, 2H, aromatic H, C_{4, 6}), 7.65–7.60 (t, 1H, aromatic H, C₅), 4.54, 4.50 (q, 2H, CHNH), 3.66 (s, 6H, OCH₃), 1.42 (d, 6H, 2CH₃, CHCH₃). ¹³C-NMR (125 MHz, δ, ppm, DMSO-*d*₆): 173.57 (2C, CO, ester), 166.33 (2C, CONH), 134.38 (2C, isophthaloyl, C_{1, 5}), 130.77 (2C, isophthaloyl, C_{2, 4}), 128.81 (1C, isophthaloyl, C₃), 127.27 (1C, isophthaloyl, C₆), 52.35 (2C, OCH₃, L-ala), 48.85 (2C, CHCH₃, L-ala), 17.15 (2C, CHCH₃, L-ala). MS (70-eV; -EI): -m/z (%) = 337 (M⁺+1, 14.75%), 336 (M⁺, 1.2%), 277 (87.52%), 234 (100%), 50 (7.13%). Molecular formula (M.wt.), C₁₆H₂₀N₂O₆ - (336.3), calculated-analysis; C, 57.14; H, 5.99; N, 8.33, found; -C 57.00, -H 6.02, -N 8.30.

N^α-I, 4-Benzenedicarbonyl-Bis-[L-Met-OMe], (4)

Yield: [A]: 85; [B]: 70; m.p. 130–132 °C. Rf x100 (solvent system) 65 (S₂). [α]_D²⁵: - 24.21 (C, 0.02, DMSO). IR in (cm⁻¹; KBr): 3421 (NH-stretching), 3015 (CH-aromatic), 2959 (CH-aliphatic), 1725 (C=O-ester), 1608 and 1523 (C=O-amide I+II). ¹H-NMR (500 MHz, DMSO-*d*₆) δ: 8.46 (s, 2H, CONH, D₂O, exchangeable), 8.20–7.69 (4H, aromatic H), 3.90 (q 2H, CHNH), 3.34 (s, 6H, OCH₃), 2.51 (t, 8H, 4CH₂, CH₂CH₂), 2.18 (s, 6H, SCH₃). ¹³C-NMR (125 MHz, δ, ppm, DMSO-*d*₆): 165.84 (2C, CO, ester), 162.00 (2C, CONH), 134.04 (2C, isophthaloyl, C_{1, 5}), 130.61–130.01 (3C, isophthaloyl, C_{2, 3, 4, 6}), 52.90–52.00 (4C, 2OCH₃, 2CHCH₂, L-met), 40.19–39.35 (4C, 2CHCH₂CH₂, L-met), 17.00 (2C, SCH₃, L-met). MS (70-eV; -EI): -m/z (%) = 457 (M⁺+1, 52.94%), 456 (M⁺, 63.73%), 442 (100%), 272 (90.20%), 233 (55.88%), 51 (66.67%). Molecular formula (M.wt.), C₂₀H₂₈N₂O₆S₂ (456.6), calculated analysis; C, 52.61; H, 6.18; N, 6.14; S, 14.05, found; C 52.56, H 6.10, N 6.09; S, 14.00.

Synthesis of N^α-I, 4-Benzenedicarbonyl-Bis-[Amino Acid]

To a stirred solution (-5°C, 20 mL) containing methyl esters of methanol (3, 4) (2 g, 5.94 mmol), sodium

hydroxide (NaOH) (1 N, 35 mL) was added dropwise and then stirred for 3 hours at -5°C, followed by stirring at room temperature for an additional 24 hours. The volatile materials were evaporated under reduced pressure and the aqueous solution was acidified (-5°C, 1 N hydrochloric acid [HCL], pH=3). The residual material was obtained and then recrystallized in an ethanol/water solution to obtain the acids 5, 6.

N^α-I, 4-Benzenedicarbonyl-Bis-[L-Ala], (5)

Yield: 75; m.p. 184–186 °C. Rf x100 (solvent system) 44 (S₁). [α]_D²⁵: - 55.15 (C, 0.06, DMSO). IR in (cm⁻¹; KBr): 3318 (NH-stretching), 3131 (CH, aromatic), 2997 (CH, aliphatic), 1715 (C=O, acid), 1643 and 1530 (C=O, amide I and amide II). ¹H-NMR (500 MHz, DMSO-*d*₆) δ: 8.93, 8.91 (s, 2H, OH), 8.82, 8.80 (s, 2H, CONH, D₂O exchangeable), 8.38 (s, 1H, aromatic H, C₂), 8.18–8.02 (d, 2H, aromatic H, C_{4, 6}), 7.66–7.57 (t, 1H, aromatic H, C₅), 4.48, 4.44 (q, 2H, CHNH), 3.66 (s, 6H, OCH₃, disappeared), 1.43, 1.41 (d, 6H, 2CH₃, CHCH₃). ¹³C-NMR (125 MHz, δ, ppm, DMSO-*d*₆): 174.62 (2C, CO, acid), 167.38, 167.10 (2C, CONH), 166.31, 165.94 (2C, isophthaloyl, C_{1, 5}), 134.76, 134, 61 (2C, isophthaloyl, C_{2, 4}), 133.85 (1C, isophthaloyl, C₃), 130.64 (1C, isophthaloyl, C₆), 52.35 (2C, OCH₃, L-ala, disappeared), 48.41 (2C, CHCH₃, L-ala), 18.25 (2C, CHCH₃, L-ala). MS (EI, 70 eV): m/z (%) = 309 (M⁺+1, 7.38%), 308 (M⁺, 3.37%), 263 (23.82%), 76 (100%), 50 (41.95%). Molecular formula (M.wt.), C₁₄H₁₆N₂O₆ (308.3), calculated analysis; C, 54.54; H, 5.23; N, 9.09, found; C 54.51, H 5.20, N 9.02.

N^α-I, 4-Benzenedicarbonyl-Bis-[L-Met], (6)

Yield: 70; m.p. 210–212 °C. Rf x100 (solvent system) 50 (S₁). [α]_D²⁵: - 38.75 (C, 0.05, DMSO). IR in (cm⁻¹; KBr): 3435 (NH-stretching), 3086 (CH-arom.), 3006 (CH-aliphatic), 1687 (C=O, acid), 1612 and 1580 (C=O, amide I and amide II). ¹H-NMR (500 MHz, DMSO-*d*₆) δ: 13.00 (br, 2H, OH), 8.48 (s, 2H, CONH, D₂O exchangeable), 8.15–7.60 (4H, aromatic H), 3.75 (q 2H, CHNH), 3.34 (s, 6H, OCH₃, disappeared), 2.50 (t, 8H, 4CH₂, CH₂CH₂), 2.16 (s, 6H, SCH₃). ¹³C-NMR (125 MHz, δ, ppm, DMSO-*d*₆): 167.03 (2C, CO, acid), 165.00 (2C, CONH), 133.81 (2C, isophthaloyl, C_{1, 5}), 131.67 (2C, isophthaloyl, C_{2, 4}), 130.43 (1C, isophthaloyl, C₃), 129.55 (1C, isophthaloyl, C₆), 52.50 (2C, 2CHCH₂, L-met), 40.47–39.52 (4C, 2CHCH₂CH₂, L-met), 15.20 (2C, SCH₃, L-met). MS (EI, 70 eV): m/z (%) = 429 (M⁺+1, 0.80%), 428 (M⁺, 0.85%),

76 (19.17%), 65 (100%), 50 (48.76%). Molecular formula (M.wt.), $C_{18}H_{24}N_2O_6S_2$ (428.5), calculated analysis; C, 50.45; H, 5.65; N, 6.54; S, 14.97, found; C 50.39, H 5.66, N 6.50; S, 14.95.

Synthesis of N^α -I, 3-Benzenedicarbonyl-Bis-[Amino Acid Hydrazides]

To a stirred solution (50 mL methanol) of esters (**3** and **4**; 1 mmol), hydrazine hydrate (0.35 mL, 10 mmol) was added. The mixture was refluxed for 3 hours; and the volatile materials were evaporated under reduced pressure. The obtained residue was triturated with ether and recrystallized from mixture of methanol/ether solution to produce the hydrazides (**7** and **8**).

N^α -I, 4-Benzenedicarbonyl-Bis-[L-Ala-NHNH₂], (**7**)

Yield: 65; m.p. 269–271 °C. Rf x100 (solvent system) 35 (S). $[\alpha]_D^{25}$: - 112.00 (C, 0.02, DMSO). IR in (cm⁻¹; KBr): 3298 (NH-stretching), 3100 (CH-arom), 2900 (CH-aliphatic), 1643 (C=O, hydrazide), 1610 and 1531 (C=O- amide I+ II). ¹H-NMR (500 MHz, DMSO-d₆) δ : 8.75 (s, 2H, CONHNH₂, D₂O exchangeable), 8.40, 8.34 (s, 2H, CONHCH, D₂O exchangeable), 8.02–7.52 (s, 4H, aromatic H), 4.49–4.47 (q, 2H, CHNH), 4.19 (s, 4H, CONHNH₂), 3.66 (s, 6H, OCH₃, disappeared), 1.35, 1.33 (d, 6H, 2CH₃, CHCH₃). ¹³C-NMR (125 MHz, δ , ppm, DMSO-*d*₆): 172.27 (2C, CO, hydrazide), 166.27 (2C, CONH), 165.00 (2C, isophthaloyl, C₁, ₅), 134.51 (2C, isophthaloyl, C₂, ₄), 130.78 (1C, isophthaloyl, C₃), 128.67 (1C, isophthaloyl, C₆), 52.35 (2C, OCH₃, L-ala, disappeared), 48.74 (2C, CHCH₃, L-ala), 17.33, 17.25 (2C, CHCH₃, L-ala). MS (EI, 70 eV): m/z (%) = 337 (M⁺+1, 1.65%), 336 (M⁺, 0.76%), 273 (100%), 76 (29.37%), 50 (7.46%). Molecular formula (M.wt.), $C_{16}H_{20}N_2O_6$ (336.3), calculated analysis; C, 57.14; H, 5.99; N, 8.33, found; C 57.11, H 5.90, N 8.31.

N^α -I, 4-Benzenedicarbonyl-Bis-[L-Met-NHNH₂], (**8**)

Yield: 70; m.p. 292–294 °C. Rf x100 (solvent system) 40 (S). $[\alpha]_D^{25}$: - 86.25 (C, 0.03, DMSO). IR in (cm⁻¹; KBr): 3288 (NH-stretching), 3155 (CH-aromatic), 3053 (CH-Aliphatic), 1661 (C=O, hydrazide), 1625 and 1525 (C=O, amide I and amide II). ¹H-NMR (500 MHz, DMSO-d₆) δ : 9.85 (br, 2H, CONHNH₂, D₂O exchangeable), 8.28 (s, 2H, CONH, D₂O exchangeable), 7.94–7.51 (4H, aromatic H), 4.00 (br, 2H, CONHNH₂), 3.17 (q, 2H, CHNH), 3.34 (s,

6H, OCH₃, disappeared), 2.51 (t, 8H, 4CH₂, CH₂CH₂), 1.75 (s, 6H, SCH₃). ¹³C-NMR (125 MHz, δ , ppm, DMSO-*d*₆): 165.91 (2C, CO, hydrazide), 160.00 (2C, CONH), 133.55 (2C, isophthaloyl, C₁, ₅), 129.88 (2C, isophthaloyl, C₂, ₄), 128.97 (1C, isophthaloyl, C₃), 126.45 (1C, isophthaloyl, C₆), 50.45 (2C, 2CHCH₂, L-met), 40.18–39.34 (4C, 2CHCH₂CH₂, L-met), 16.25 (2C, SCH₃, L-met). MS (EI, 70 eV): m/z (%) = 456 (M⁺, 3.42%), 265 (5%), 76 (100%), 64 (10.92%), 50 (81.49%). Molecular formula (M.wt.), $C_{20}H_{28}N_2O_6S_2$ (456.6), calculated analysis; C, 52.61; H, 6.18; N, 6.14; S, 14.05, found; C– 52.55, H– 6.15, N– 6.11; S– 14.00.

Synthesis of N^α -I, 3-Benzenedicarbonyl-Bis-[L-Ala-L-Phe Methyl Esters], (**9**)

Ethyl chloroformate (2.9 mL, 30.1 mmol) was added to solution of 1, 3-benzenedicarbonyl-*bis*-[L-Alanine], **5**, (-20 °C, 15.05mmol) and triethylamine (3.3 mL, 30.1mmol) in dichloromethane (50 mL). The reaction mixture was stirred for additional 30 minutes, then the two equivalents amount of free amino acid ester (L-Phe) (30.1mmol)(-20°C) in dichloromethane (50mL) was added. Then, the same procedure, mentioned in the synthesis of compound **3** using the mixed anhydride method was completed. The crude product was collected, and then purified by recrystallization from ethanol to get the corresponding ester **9**.

Yield: 85; m.p. 86–88 °C. Rf x100 (solvent system) 70 (S₂); $[\alpha]_D^{25}$: - 22.10 (C, 0.02, MeOH). IR in (cm⁻¹; KBr): 3422 (NH-stretching) 3140 (CH, aromatic) 2950 (CH, aliphatic) 1732 (C=O, ester) 1648, 1530, 1441 and 1382 (C=O, amide I, amide II, amide III and amide IV). ¹H-NMR (500 MHz, DMSO-d₆) δ : 9.09–9.01 (s, 4H, CONH, D₂O exchangeable), 8.66 (s, 1H, isophthaloyl, aromatic H, C₂), 8.48–8.40 (d, 2H, aromatic H, isophthaloyl, C₄, ₆), 7.65 (t, 1H, isophthaloyl, aromatic H, C₅), 7.50–7.21 (10H, aromatic H, L-phe), 4.56–4.51 (q, 2H, CHNH, L-phe), 4.29–4.23 (t, 2H, CHNH, L-ala), 3.64–3.59 (s, 6H, OCH₃), 3.19–2.88 (d, 4H, CH₂, L-phe), 1.41–1.03 (d, 6H, 2CH₃, CHCH₃, L-ala). ¹³C-NMR (125 MHz, δ , ppm, DMSO-*d*₆): 173.00 (2C, CO, ester), 169.00–170.25 (4C, CONH), 135.50 (1C, L-Phe, C₁), 133.20 (2C, isophthaloyl, C₁, ₅), 131.50 (2C, isophthaloyl, C₂, ₄), 128.80–124.25 (10C, the remaining aromatic carbon), 53.25 (2C, CHCH₂, L-phe), 51.50 (2C, OCH₃, L-ala), 50.00 (2C, CHCH₃, L-ala), 38.20 (2C, CH₂, L-phe), 18.00 (2C, CHCH₃, L-ala). MS (EI, 70 eV): m/z (%) = 631 (M⁺+1,

18.77%), 630 (M^+ , 23.60%), 611 (100%), 74 (2.15%), 55 (1.33%). Molecular formula (M.wt.), $C_{34}H_{38}N_4O_8$ (630.7), calculated analysis; C, 64.75; H, 6.07; N, 8.88, found; C 64.70, H 6.04, N 8.90.

Synthesis of N^α -I, 3-Benzenedicarbonyl-Bis-[L-Ala-L-Phe], (10)

To a stirred solution (-5°C , 20 mL methanol) of ester (**9**; 2 mmol), NaOH (1N, 25 mL) was added dropwise, then stirring was continuous for additional three 3 hours at -5°C . The reaction mixture was left over night at room temperature. Then complete the same procedure which was mentioned in the synthesis of compounds **5** and **6**. The precipitated crude product was filtered off and recrystallized from ethanol to get the corresponding acid **10**.

Yield: 75; m.p. 169–171 $^\circ\text{C}$. Rf x100 (solvent system) 60 (S₁); $[\alpha]_D^{25}$: -33.41 (C, 0.02, MeOH). IR in (cm^{-1} ; KBr): 3430 (NH-stretching) 3175 (CH, aromatic) 2928 (CH, aliphatic) 1643 (C=O, acid) 1529, 1480, 1421 and 1386 (C=O, amide I, amide II, amide III and amide IV). $^1\text{H-NMR}$ (500 MHz, DMSO- d_6) δ : 12.55 (br, 2H, OH), 9.19–9.09 (s, 4H, CONH, D_2O exchangeable), 8.70 (s, 1H, isophthaloyl, aromatic H, C₂), 8.40–8.33 (d, 2H, aromatic H, isophthaloyl, C₄, ₆), 7.77 (t, 1H, isophthaloyl, aromatic H, C₅), 7.47–7.18 (10H, aromatic H, L-phe), 4.61–4.49 (q, 2H, CHNH, L-phe), 4.33–4.20 (t, 2H, CHNH, L-ala), 3.64–3.59 (s, 6H, OCH₃, disappeared), 3.25–2.75 (d, 4H, CH₂, L-phe), 1.55–1.25 (d, 6H, 2CH₃, CHCH₃, L-ala). $^{13}\text{C-NMR}$ (125 MHz, δ , ppm, DMSO- d_6): 174.50 (2C, CO, acid), 170.15–172.00 (4C, CONH), 139.20 (1C, L-Phe, C₁), 136.00 (2C, isophthaloyl, C₁, ₅), 134.20 (2C, isophthaloyl, C₂, ₄), 130.50–125.00 (10C, the remaining aromatic carbon), 54.50 (2C, CHCH₂, L-phe), 51.50 (2C, OCH₃, L-ala, disappeared), 48.50 (2C, CHCH₃, L-ala), 39.70 (2C, CH₂, L-phe), 17.50 (2C, CHCH₃, L-ala). MS (EI, 70 eV): m/z (%) =603 (M^+ +1, 66.92%), 602 (M^+ , 79.23%), 484 (100%), 367 (72.31%), 159 (79.23%), 53 (64.62%). Molecular formula (M.wt.), $C_{32}H_{34}N_4O_8$ (602.6), calculated analysis; C, 63.78; H, 5.69; N, 9.30, found; C 63.70, H 5.70, N 9.32.

Synthesis of N^α -I, 3-Benzenedicarbonyl-Bis-[L-Ala-L-Phe-Hydrazide], (11)

The ester (**9**; 1 mmol) in 50 mL methanol was stirred with hydrazine hydrate (0.35 mL, 10 mmol) for 3 hours; the volatile materials were evaporated under reduced pressure. The residual product diluted with ether and recrystallized by mixture of methanol/ether to afford the corresponding hydrazide **11**.

Yield: 65; m.p. 235–237 $^\circ\text{C}$. Rf x100 (solvent system) 45 (S); $[\alpha]_D^{25}$: -133.80 (C, 0.04, DMSO). IR in (cm^{-1} ; KBr): 3424 (NH-stretching), 3273 (CH, aromatic) 3066 (CH, aliphatic) 1647 (C=O, hydrazide) 1538, 1449, 1400 and 1382 (C=O, amide I, amide II, amide III and amide IV). $^1\text{H-NMR}$ (500 MHz, DMSO- d_6) δ : 9.92 (br, 2H, CONHNH₂), 9.23–9.15 (s, 4H, CONH, D_2O exchangeable), 8.82 (s, 1H, isophthaloyl, aromatic H, C₂), 8.35–8.24 (d, 2H, aromatic H, isophthaloyl, C₄, ₆), 7.86 (t, 1H, isophthaloyl, aromatic H, C₅), 7.62–7.38 (10H, aromatic H, L-phe), 4.74–4.60 (q, 2H, CHNH, L-phe), 4.44–4.30 (t, 2H, CHNH, L-ala), 4.00 (br, 2H, CONHNH₂), 3.64–3.59 (s, 6H, OCH₃, disappeared), 3.40–2.80 (d, 4H, CH₂, L-phe), 1.90–1.60 (d, 6H, 2CH₃, CHCH₃, L-ala). $^{13}\text{C-NMR}$ (125 MHz, δ , ppm, DMSO- d_6): 176.60 (2C, CO, hydrazide), 171.25–172.50 (4C, CONH), 138.00 (1C, L-Phe, C₁), 135.20 (2C, isophthaloyl, C₁, ₅), 133.00 (2C, isophthaloyl, C₂, ₄), 128.00–123.50 (10C, the remaining aromatic carbon), 55.75 (2C, CHCH₂, L-phe), 51.50 (2C, OCH₃, L-ala, disappeared), 46.00 (2C, CHCH₃, L-ala), 36.00 (2C, CH₂, L-phe), 16.20 (2C, CHCH₃, L-ala). MS (EI, 70 eV): m/z (%) =631 (M^+ +1, 41.67%), 630 (M^+ , 69.70%), 403 (74.24%), 91 (100%), 71 (47.73%), 52 (92.42%). Molecular formula (M.wt.), $C_{32}H_{38}N_8O_6$ (630.7), calculated analysis; C, 60.94; H, 6.07; N, 17.77, found; C 60.92, H 6.02, N 17.72.

Biological Evaluation

In vitro Antimicrobial Activity

Antimicrobial efficiency for the investigated compounds (**3–11**) was evaluated using “well diffusion method” against pathogenic Gram-positive bacteria including *Streptococcus pneumonia*, *Staphylococcus aureus* ATCC25923, and *Micrococcus luteus*. Subsequently, these compounds were tested against Gram-negative bacteria “*Escherichia coli* (ATCC25922), *Pseudomonas aeruginosa* (ATCC7853) and *Proteus mirabilis*, and *Candida albicans* (ATCC1031)” as the human yeast pathogen. While antifungal activity was tested against different plant pathogens (*Rhizocotonia solani*, *Fusarium oxysporum*, and *Fusarium solani*) and human pathogen *Aspergillus niger* in terms of the percent inhibition of mycelial growth. All tested bacteria and yeast were incubated at 37 $^\circ\text{C}$ for 24 hours by inoculation into nutrient broth. The culture suspensions were prepared and adjusted at absorbance 0.5 by spectrophotometer.

Well Diffusion Method

Petri dishes with nutrient agar were inoculated with a 100 μ L suspension of each bacterial or yeast culture. Wells were generated using the agar surface around (5 mm) below the cork-borer. The tested compounds (100 μ L) were poured into the well using a sterile syringe. The bacteria seeded plate was placed in the refrigerator for 8 hours at 4°C. Next, the plates were incubated at 37 \pm 2°C for 24 hours. The inhibition zone formed around the well in the plates was measured as the diameter of the inhibition zone around the well (in mm) through the well diameter. DMSO was used to dissolve the investigated compounds (**3** to **11**; 1 mg/mL) as well as for the dilution of the compounds to be tested.²³ The average values for the three different trials (measurements were taken in three different fixed directions) are tabulated in Table 1.

Antifungal Activity

The antifungal potency for compounds **3** to **11** was evaluated as the inhibition percent for expansion mycelial (IPEM).²⁹ The test compounds **3** to **11** (100 μ L) were added to potato dextrose agar medium (PDA) plates at a concentration of 1 mg/mL. Agar discs (5 mm) were taken from 7 day-old cultures of 4 pathogenic fungi (*Aspergillus niger*, *Rhizoctonia solani*, *Fusarium oxysporum*, and *Fusarium solani*) and were placed separately in the center of the Petri plates. For the control condition, agar discs of the same size as the tested pathogen were positioned similarly on a fresh PDA plate. The incubation period for all paired cultures was performed at 25°C for 6 days. Inhibitory efficiency was measured by determining the expansion radius of the mycelium on the treated media (R2) versus the control (R1). The

IPEM were obtained using the following formula: $IPEM = \{(R1 - R2)/R1\} \times 100$.

Antioxidant Activity

Scavenging efficiency for free radicals was measured as the change the color of DPPH• radical from strong violet to yellow color as previously reported.³⁰

Anticancer Activity

Human cancer cell lines used in this study, included MCF7 (Breast carcinoma cell line), Hep-G2 (Liver carcinoma cell line), and HCT116 (Colon carcinoma cell line). They were obtained frozen in liquid nitrogen (-180°C) from Vacsera (Giza, Egypt). The tumor cell lines were maintained by serial sub-culturing. Cytotoxicity was determined using the MTT assay.

Cell Viability Assay

For the assessment of the cytotoxic activities of compounds **3** to **11**, cells were seeded in 96-well flat-bottomed microtiter plates at a density of approximately (10 \times 10³/well for MCF-7 and HepG2, while in HCT-116 20 \times 10³ cells/well), in complete RPMI-1640 Medium. Cells were incubated in a water jacketed CO₂ incubator at 37°C for 24 hours. The remaining procedures, including the MTT assays, were performed as previously reported.^{31,32}

BCL-2 Measurement

Based on previous reports,³² the effects on BCL-2 levels for the compounds under examination and the standards were tested. A contaminated biotin with antibody was added and was then streptavidin HRP. The termination of the reaction takes place by addition of acid, and the absorbance was determined at 450 nm.

Table 1 Antimicrobial Activity of the Synthesized Compounds by Well Diffusion Assay Method

Pathogenic Microbial Strains		Diameter of Inhibition Zone (mm)								
		3	4	5	6	7	8	9	10	11
Gram-positive bacteria	<i>Streptococcus pneumoniae</i>	14	-	-	-	-	-	-	-	-
	<i>Staphylococcus aureus</i> ATCC25923	-	-	-	-	-	-	-	-	-
	<i>Micrococcus luteus</i>	15	20	3.0	20	12	11	14	15	15
Gram-negative bacteria	<i>Escherichia coli</i> ATCC25922	-	14	-	13	-	-	12	12	-
	<i>Pseudomonas aeruginosa</i> ATCC7853	-	-	-	-	-	10	-	-	-
	<i>Proteus mirabilis</i>	10	22	-	15	-	15	13	20	-
Yeast	<i>Candida albicans</i> ATCC1031	-	-	-	-	-	-	-	-	-

Note: (-) No inhibition.

LC₅₀ Determination

LC₅₀ values were calculated using SPSS Statistics Base v.9 for the active extracts which showed promising values in the analysis.

Enzyme-Linked Immunosorbent Assay for Tubulin Beta (TUBB)

A total of 1.2–1.8×10,000 cells/well of MCF-7 cells were seeded in DMEM (supplemented with 10% FBS and 1% penicillin-streptomycin). Before assessment of tubulin levels, the compound under investigation was added for 18–24 hours. Next, Avidin combined with HRP was added to each microplate well and incubated. The remaining steps were performed as previously reported.³³

Human P53 Estimation

ELISA was used to measure the amount of human p53 bound to antibodies in the sample or to standards adsorbed onto the microwells. A solution containing biotin was added. The contaminated biotin with antibodies was supplemented. Then, Human p53 was measured as previously reported.³⁴

Computational Study

Preparation of Ligands and Protein

The 3-dimensional structure protein files were downloaded from the RCSB protein data bank for DNA Gyrase B (ID: 4uro),³⁵ 14 α -sterol demethylase *Candida albicans* (ID:1EA1),³ and transforming growth factor beta receptor type I (TGFBR1) (ID: 1VJY).³⁷ For mining of protein data, we used the BLAST P tool for analysis of the FASTA protein sequence. Then, the CLUSTALW⁴⁶ package was utilized for the alignment of multiple of amino acid sequences.^{38–44}

The Docking computations were completed using the MOE 2015 package.⁴⁵ Compounds **3** to **11** were built then energy minimized based on “Semi-empirical” PM3 then DFT “discrete Fourier transform” based on B3LYP/6-311G. The error correction for the structure of the catalytic sites was performed by supplementing hydrogens and partial charges using (Amber12: EHT), then minimized by utilized the same force field with RMSD= 0.100. The catalytic site was identified and analyzed using the Site Finder program. This method is based on alpha spheres as well as an energy model.⁴⁶ The catalytic zone was predicted by the MOE-Site finder.⁴⁷

Stepwise Docking Experiment

Water and inhibiting molecules were eliminated, then H-atoms were supplemented to the obtained crystal structure. The charges were designed using MMFF94x force field. The alpha-site spheres were added based on the findings of the on-site-finder module. The final energy, generated structures, and final scores were measured using procedures described previously.⁴⁸

ADMET Predictions

The in silico ADMET profile was simulated using MOE and admetSAR Lipinski criteria and ADMET properties (absorption, distribution, metabolism, excretion, and toxicity).⁴⁹

Results and Discussion

Chemistry

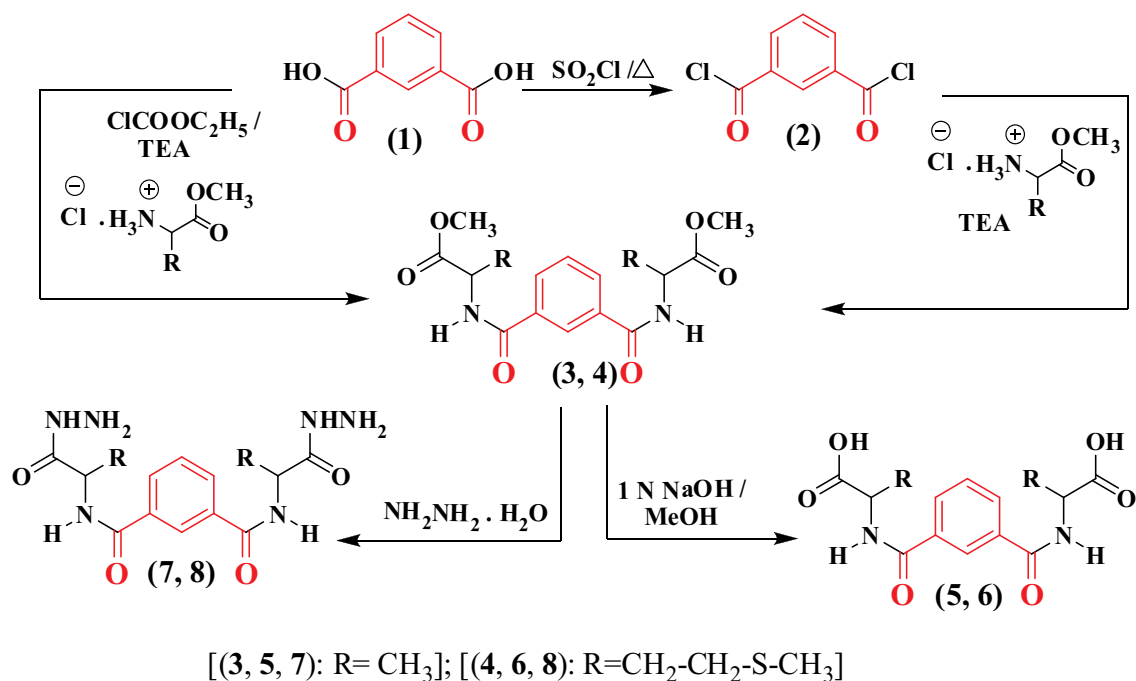
The N α -1, 3-benzenedicarbonyl-*bis*-(L-Alanine and L-Methionine methyl esters (**3**, **4**, respectively), were obtained using two synthesis routes. The first involved acylation of L-alanine and L-methionine methyl esters with 1, 3-benzenedicarbonyl dichloride **2**, which was obtained from the reaction of 1, 3-benzenedicarboxylic acid **1** with thionyl chloride. Compound **2** was then coupled with the L-alanine and L-methionine esters at a low temperature in the presence of trimethylamine. In the second synthesis route: the *bis*-esters (**3**, **4**) were prepared starting from N α -1,3-benzenedicarboxylic acid **1**, with L-alanine and L-methionine esters, in the presence of ethyl chloroformate, as a mixed anhydride partner. Hydrolysis of (**3**, **4**) with methanolic NaOH resulted in the corresponding N α -1,3-benzenedicarbonyl-*bis*-(L-alanine and L-methionine; **5**, **6**). The hydrazinolysis of **3**, **4** with alcoholic hydrazine hydrate, generated the corresponding N α -1,3-benzenedicarbonyl-*bis*-(L-alanine and L-methionine; **7**, **8**) (Scheme 1).

Treatment of **5** with free L-phenyl alanine ester generated the corresponding N α -1,3-benzenedicarbonyl-*bis*-(dipeptide)-ester (**9**). This compound was converted to the corresponding N α -1,3-benzenedicarbonyl-*bis*-(dipeptide; **10**) under hydrolysis conditions by ethanolic sodium hydroxide. Likewise, hydrazinolysis of the dipeptide methyl ester (**10**) with methanolic hydrazine hydrate generated the corresponding dipeptidehydrazide (**11**) (Scheme 2).

Biological Evaluations

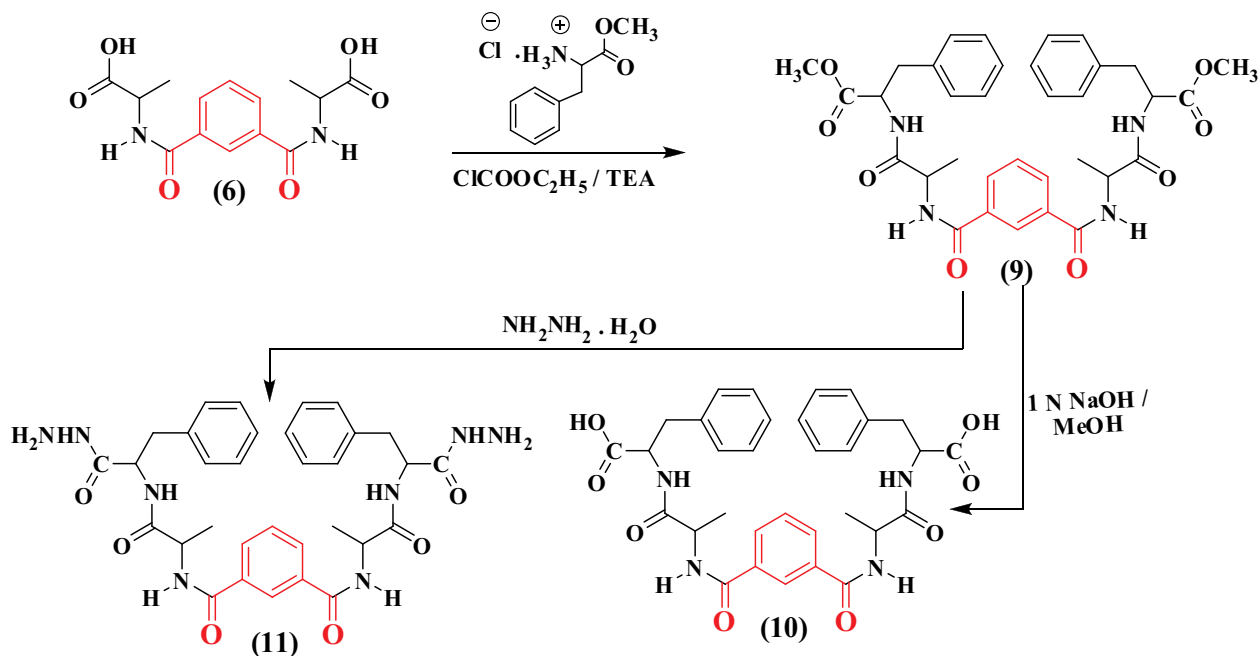
Antimicrobial Activity

The antimicrobial activity of the synthesized compounds **3** to **11** against pathogenic bacterial and fungal strains is shown in Tables 1 and 2. The tested compounds showed



- 3**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala)-OMe, **4**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Met)-OMe
5: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala)-OH, **6**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Met)-OH
7: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala)-NHNH₂, **8**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Met)-NHNH₂

Scheme 1 Synthetic routes for N^α -1, 3-benzenedicarbonyl-bis-(Amino acid methyl esters) (**3**, **4**); corresponding carboxylic acid (**5**, **6**) and the corresponding hydrazides (**7**, **8**).



- 9**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala-L-Phe)-OMe, **10**: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala-L-Phe)-OH
11: N^α -1, 3-benzenedicarbonyl-*bis*-(L-Ala-L-Phe)-NHNH₂

Scheme 2 Synthetic routes for N^α -1, 3-benzenedicarbonyl-bis-(L-Ala-L-Phe methyl esters) (**9**), corresponding carboxylic acid (**10**) and the corresponding hydrazide (**11**).

Table 2 Antifungal Activity of the Synthesized Compounds by the Percent Inhibition of Mycelial Growth

Pathogenic Fungi	The Inhibition Percent for Expansion Mycelial (IPEM)(%)								
	3	4	5	6	7	8	9	10	11
<i>Rhizocotonia solani</i>	–	30%	–	–	–	–	25%	–	40%
<i>Fusarium oxysporum</i>	–	–	11%	–	–	–	11%	–	–
<i>Fusarium solani</i>	–	–	–	–	–	–	–	–	–
<i>Aspergillus niger</i>	*	–	*	*	–	100%	–	*	No spore formation

Notes: (–) No inhibition; (*) spore color; (%>100) inhibition with sparse growth; (=100) inhibition (growth and sporulation).

different degrees of potency against pathogenic bacteria. *Micrococcus luteus* was the most sensitive bacteria against compound **6**, which showed the highest activity. Compounds **5**, **7** produced the lowest inhibitory activity and showed resistance towards all tested pathogenic microorganisms tested with the exception of *Micrococcus luteus*.

None of the tested compounds exhibited any activity against *Staphylococcus aureus* and yeast. The pathogen *Proteus sp.* was inhibited by most synthesized peptides except for compounds **6** and **11**, while *Streptococcus pneumonia* pathogen was only resistant to compound **3**, among all members of all synthesized compounds.

Most synthesized compounds showed the most significant antifungal activity against *Aspergillus niger*, except for compounds **4**, **7**, and **9**. Exposure to compounds **3**, **5**, **6**, and **10** changed the black color of *Aspergillus niger* spores to grey. Compound **8** was the most potent antifungal agent against *Aspergillus niger*, which produced 100% inhibition, while compound **11** inhibited the formation of the respective fungal spores. The inhibitory mechanism of these compounds may be attributed to a fungicidal inhibition of the THN-reductase enzyme, involved in melanin synthesis via the DHN-melanin pathway.⁵⁰ Compounds **4**, **9**, and **11** showed a moderate inhibition against *Rhizocotonia solani*, while compounds **5** and **9** showed a relatively low degree of inhibition (11%) against *Fusarium oxysporum*. Most synthesized compounds (ie, **3**, **4**, **6**, **7**, **8**, and **10**) showed no antifungal activity against *Fusarium solani*.

Antioxidant Activity

Nine compounds were screened to investigate their antioxidant potency using the free radical scavenging activity DPPH assay, and all nine compounds showed weak free radical scavenging activity (Table 3).

Table 3 Free Radical Scavenging Activity of the Synthesized Compounds on Free Radical DPPH at 50ug/mL

Samples	% DPPH Scavenging Activity
Cpd 3	6.7±1.5
Cpd 4	0.9±0.5
Cpd 5	0.6±0.2
Cpd 6	1.0±0.8
Cpd 7	7.6±0.8
Cpd 8	14.8±1.3
Cpd 9	3.6±0.7
Cpd 10	13.3±0.8
Cpd 11	0.9±0.5

Anticancer Activity

Nine samples (**3** to **11**) were preliminary screening for their antiproliferative activity against the following human carcinoma cell lines, breast MCF-7 (Figure 1), colon HCT-116 (Figure 2), and hepatocellular carcinoma HepG2 (Figure 3) cell lines. The nine compounds possessed weak to moderate antiproliferative effects in the HCT-116 cell line at concentration 10 ppm or 10µg/mL, with cytotoxicity ranging from 14% to 60.7%. While compound **9** possessed weak cytotoxic activity in the liver human tumor cell line HepG2, with cytotoxicity ranging from 2.2% to 9%. All the cytotoxicity results of the studied compounds were compared with the results of the doxorubicin reference drug, which ranged from 80% to 95% cytotoxicity for all three cell lines tested.

Conversely, compounds **6** and **7**, showed a strong cytotoxic effect on the breast cancer cell line MCF-7 with 95.1% and 79.8% of cell inhibition, respectively. Based on the promising results of compound **6** in the breast cancer cell line, it was selected for further investigation of additional anticancer activity.

The proto-oncogene coding the B-cell lymphoma 2 (Bcl-2) is known as apoptosis regulator by maintaining the equilibrium between cell death and proliferation. Low

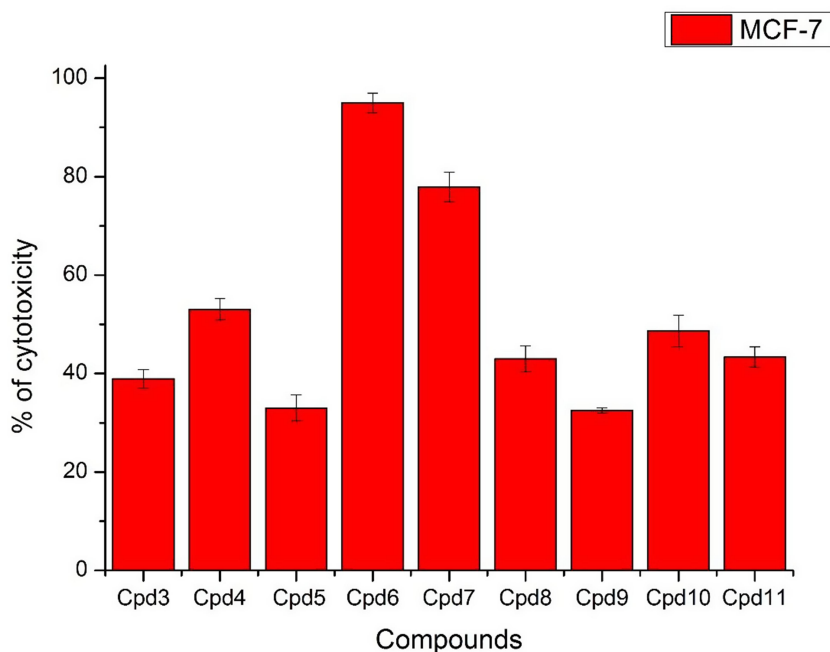


Figure 1 In vitro cytotoxicity screening for nine tested compounds against colon carcinoma (MCF-7).

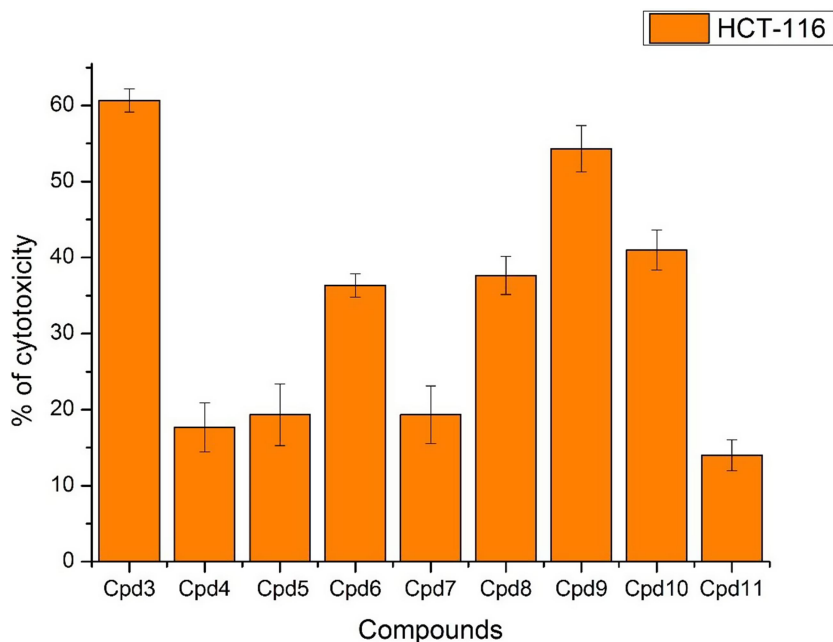


Figure 2 In vitro cytotoxicity screening for nine tested compounds via against breast carcinoma (HCT-116).

levels of Bcl-2 are associated with cell death through apoptosis.

MCF-7 cells treated with compound 6 produced BCL2 levels of 103.3 pg/mL, which lies near the borderline calibration curve of BCL2, ranging from 78.1 to 2500 pg/mL. Weak expression of Bcl2 was observed in

comparison to the reference drug doxorubicin, indicates compound 6 induced apoptosis of MCF-7 cells.

P53 is a tumor suppressor that is triggered in response to stress. Studies demonstrate that p53 is critical for the prevention of tumorigenesis and may potentially be involved in at least half of all known cancers. MCF-7

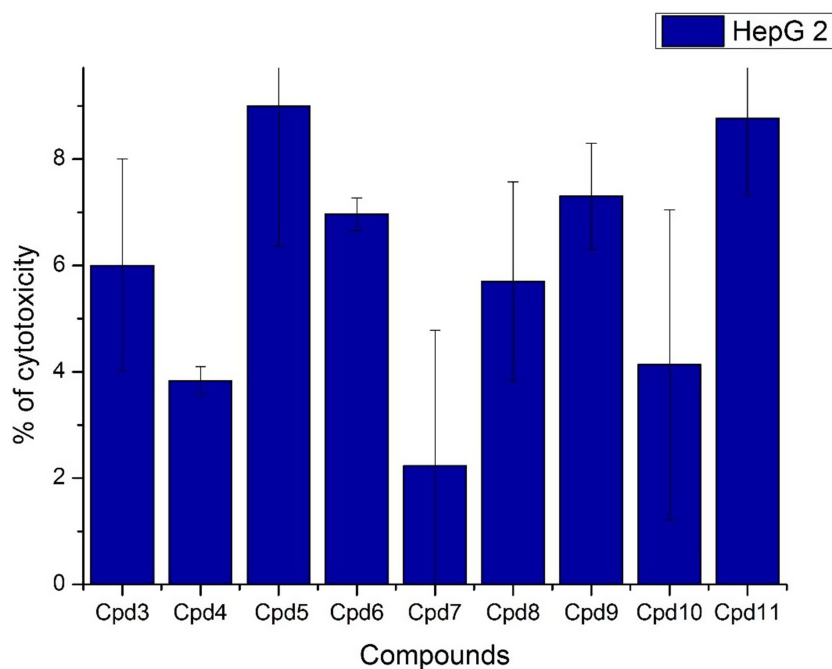


Figure 3 In vitro cytotoxicity screening for nine tested compounds via liver carcinoma (HepG 2).

treated with compound **6** showed a slight downregulation of the protein levels of p53 to 70.15 pg/mL, which suggests that compound **6**, induces apoptosis via another pathway rather than acting via p53 protein.

Microtubules are a major component of the cytoskeleton. They play a crucial role in cell division. Tubulin-binding agents are amongst the most widely used chemotherapeutic drugs in cancer treatment. We investigated the effects of compound **6** at a concentration of 10 µg/mL on TubBin MCF-7 cells. TubB levels were 0.19 pg/mL which indicated that compound **6** had no significant effect on tubulin polymerization as compared with reference drug colchicine.

Docking Studies

The docking study targeted (ID: 4uro) *DNA Gyrase B*, (ID: 1EA1) *Candida albicans* lanosterol 14- α -demethylase, and (ID: 1VJY) transforming growth factor (TGF) for β -receptor type I (β -RI), to examine the potential mode of action of the small compounds as antimicrobial and anti-tumor agents. The ligand-protein interactions were estimated based on the docking score using the functions implemented in the Molecular Operating Environment MOE 2015.10 package. All calculations of the docking experiments against crystal structures (PDB: 4uro, 1EA1 and 1VJy) are shown in Tables 4 and 5. Bacterial *DNA*-

gyrase plays a vital role in the activity of antibacterial agents and acts by breaking double-stranded DNA by catalyzing negative supercoiling which is essential for DNA replication, transcription, and recombination.⁵¹ Analysis of the co-crystallized DNA-gyrase cleavage complex with novobiocin, which is an effective antibacterial agent that acts by cleaving DNA and restricting the ATPase binding site located on the vital peptidoglycan of the bacterial cell wall (Ser 55, Ala64, Asn.65, Asp89, Thr164, Thr173, and Val79). The inhibitory effect may be a result of the distinct structure of cell wall that characterizes Gram-negative and Gram-positive bacteria. The cell wall of Gram-negative bacteria is composed of a thin peptidoglycan layer (7–8 nm) with an additional an outer membrane. The Gram-positive bacteria contain a thick peptidoglycan layer (20–80 nm) outside of the cell wall with no outer membrane. The peptidoglycan is a mesh-like polymer consisting of sugars and amino acids. A peptidoglycan layer protects microorganisms against antibacterial agents such as antibiotics, toxins, chemicals, and degradative enzymes.^{52,53} Furthermore, a docking study evaluated the lanosterol 14- α -demethylase-fluconazole complex (PDB ID: 1EA1), extracted from *Mycobacterium tuberculosis* and showed a high similarity with the lanosterol-14- α -demethylase obtained from *Candida* species, for which it is difficult to obtain X-ray crystal structure for bound

Table 4 Simulated Molecular Docking Energy in (kcal/mol) for the Tested Ligands

Cpd.	E _{d,G}	rmsd __	E _{place}	E _{conf}	E _{ele}	E _{Int.}
(ID: 4uro) DNA Gyrase B						
3	-6.02317	0.360479	50.5415	-47.7144	-5.52189	-37.2061
4	-4.67928	0.766497	43.41455	-35.2866	-5.60077	-23.5248
5	-5.31233	0.480625	-15.352	-62.0181	-6.94589	-22.5068
6	-5.65801	0.705182	-18.7582	-64.7541	-7.14161	-37.9289
7	-5.51191	0.356299	81.43585	-65.5201	-6.30798	-31.2715
8	-5.77681	0.321674	79.70246	-47.8776	-6.00393	-37.1633
9	-7.10236	3.661316	44.71742	-9.48044	-4.53216	-44.1672
10	-6.16023	2.317993	-23.5398	-54.9007	-5.2083	-34.4003
11	-7.06915	2.493655	97.9485	-86.6334	-6.06098	-51.2235
(ID: IEA1) <i>Candida albicans</i> lanosterol 14- α -demethylase						
3	-8.511	0.462	53.894	-45.302	-10.254	-42.994
4	-8.231	0.134	70.956	-34.067	-10.521	-31.988
5	-7.420	0.924	-18.645	-60.338	-13.730	-32.109
6	-7.717	0.980	-30.501	-37.080	-10.094	-42.641
7	-8.170	0.272	90.043	-63.536	-13.050	-39.811
8	-8.386	0.447	83.722	-58.822	-12.528	-45.894
9	-8.162	0.634	54.197	49.092	-6.841	-39.879
10	-8.727	0.556	-11.711	-50.456	-11.146	-41.903
11	-8.250	0.934	97.236	-5.341	-7.027	-36.096
(ID: 1VJY) transforming growth factor (TGF) for β -receptor type I (β -RI)						
3	-9.748	2.489	52.032	-61.789	-3.504	-32.149
4	-12.450	2.956	60.790	-113.823	-2.608	-13.581
5	-12.490	2.016	-18.932	-80.589	-3.453	-37.079
6	-12.863	1.361	-10.826	-102.615	-1.117	-8.393
7	-16.354	1.890	104.488	-72.455	-2.627	-6.256
8	-12.303	1.276	88.752	-81.780	0.585	-47.178
9	-11.536	2.484	44.366	-84.405	3.726	-53.218
10	-13.708	1.735	-3.721	-84.626	7.078	-24.206
11	-10.283	4.103	94.121	-62.072	7.601	-42.896

Abbreviations: E_{d,G}, final free binding energy; E_{conf}, binding energy for the ligand-conformer; E_{place}, binding energy of the ligand-receptor; E_{Int.}, ligand-receptor affinity of interaction energy; E_{ele}, ligand-receptor Electrostatic interaction; RMSD, the root mean square deviation of the pose of the docking pose compared to the co-crystal ligand position.

membrane enzyme. In addition, the crystal structure of TGF of β -RI bound with naphthyridine (ligand 460) was obtained as PDB file (ID: 1VJY) with 2.0Å resolution.

All compounds were redocked into active sites in the absence of the reference inhibitor. The ligands were successfully complexed with active sites of enzymes. The extracted docked poses of ligands were energy-minimized with a molecular mechanics (Amber12: EHT) force field, up to 0.05 kcal/mol for the gradient convergence. The poses were initially selected among those with a docking score of E_{d,G} < -7 kcal/mol). Next, poses were filtered based on lowest docking MOE score with lowest root means quart deviation (RMSD) against reference drugs.

Finally, the highest MOE scoring function for the tested compounds was applied to evaluate the binding affinities of the tested compounds (Table).

(ID: 4uro) *DNA Gyrase B*: The bispeptides **9**, **10**, and **11** exhibited the highest binding affinity and RMSD over other compounds for *DNA gyrase* (-7.1, -6.1, and -7.0 kcal/mol.), respectively, (summarized in Table 4). The other compounds showed considerable binding affinity (RMSD < 1), which was attributed to the higher stability of these compounds in the binding site (Table 4). The binding scores for compounds **3** to **8** could be arranged as 3 > 8 > 6 > 7 > 5 > 4 with a trend for RMSD between ~0.3–0.7 (Table 4). All compounds allowed H-bond

Table 5 Pharmacokinetic Features for Compounds (3–11)

Comp.	3	4	5	6	7	8	9	10	11
HBA	8	8	8	8	10	10	12	12	14
HBD	2	2	4	4	8	8	4	6	10
lip_drug like	1	1	1	1	1	1	0	0	0
lip.V	0	0	0	0	1	1	2	3	3
Log P	1.003	2.313	0.475	1.785	-1.821	-0.511	3.251	2.723	0.427
logS	-3.147	-4.880	-2.323	-4.056	-2.848	-4.581	-7.151	-6.327	-6.852
TPSA	110.8	110.8	132.8	132.8	168.44	168.44	169	191	226.64
P-GPI,II-S	NO	NO	NO	NO	NO	NO	NO	NO	NO
P-GPI,II-I	NO	NO	NO	NO	NO	NO	NO	NO	NO
Irritant	NO	NO	NO	NO	NO	NO	NO	NO	NO
CYP450-I	NO	NO	NO	NO	NO	NO	NO	NO	NO
CYP450-S	NO	NO	NO	NO	NO	NO	NO	NO	NO
Tumorigenic	NO	NO	NO	NO	NO	NO	NO	NO	NO
Carcinogens	NO	NO	NO	NO	NO	NO	NO	NO	NO
AMES.T.	NO	NO	NO	NO	NO	NO	NO	NO	NO
hERGI,II	NO	NO	NO	NO	NO	NO	NO	NO	NO
Hepatotoxicity	NO	NO	NO	NO	NO	NO	NO	NO	NO
LD50	2.639	2.951	2.222	2.278	2.486	2.465	3.075	2.483	2.465
LOAEL	0.86	0.848	1.259	1.242	2.17	2.172	2.894	3.681	4.044
Skin Sensitivity	NO	NO	NO	NO	NO	NO	NO	NO	NO

formation with important amino acid residues (Ala64, Asn.65, and Thr164) at the DNA gyrase active site ([Table S1](#), [Supplementary materials](#)). All compounds were arranged in parallel with Asn65 and Thr164, the hydrophobic portion of these compounds was considered a stabilization key in the receptor conformation. Furthermore, compounds **3**, **4**, and **6** were linked via an H₂O molecule which bound to three important backbone amino acids Val79, Ser 55, and Thr173 ([Figure 4](#)). The different ligand conformations indicated the interactions with hydrophilic amino acid backbone at the 4uro binding site ([Table S1](#), [Supplementary materials](#)). From ([Figure 4](#)), we can postulate that the hydrophobicity fragment generated in the synthesized compounds (Ala and Met) is an important pharmabiotic characteristic for binding to the DNA-gyrase pocket. Furthermore, the inhibitory efficiency of the synthesized compounds against bacterial growth may be explained by their ability to attack the naked peptidoglycan of the cell wall.^{54,55} As a result, the anti-bacterial mechanism of the synthesized compounds involves the possibility of attacking the bacterial membrane and changing its permeability. Hence, the synthesized particles may leak through sugars and proteins to deactivate hydrogen respiratory chain enzymes, and subsequently, they produce pits and gaps in the bacterial membrane (peptidoglycan layer), to induce irregular fragmentation of the bacterial cells.^{56,57} We therefore

concluded that these indicators suggested that the compounds tested would induce suitable biological functions.

(ID:1EA1) *Candida albicans* Lanosterol 14- α -Demethylase

The tested compounds **3–11** showed good binding affinity against 1EA1 ([Table 4](#), [Figure S1](#), [2](#); [Supplementary material](#)). The binding pockets of 1EA1 included amino acid residues Arg324, Arg324, Gln70, Tyr74, Arg93, Hip390, Gln70, and Arg94, which produce H-, π - π and hydrophobic interactions with the tested compounds ([Table S2](#), [Supplementary Materials](#)).

(ID: IVJY) Transforming Growth Factor (TGF) for β -Receptor Type I (β -RI)

All synthesized compounds showed higher binding affinity ([Table 4](#)) than the reference ligand 460 ($E_{d,G} = -4.81$ kcal/mol). The Ala-hydrazide derivative **7** showed the highest binding score ($E_{d,G} = -16.354$ kcal/mol). Compound **7** interacted with binding site by forming five H-bonds with the following amino acids Lys232, Ser280, and Asp290, respectively. While at the same time it formed H-bonds with H₂O which interacted with backbone amino acids Lys232 and Glu245 ([Figure S3](#); [Supplementary material](#)). The binding affinity for peptide derivatives (**4** to **6** and **8**) showed almost equally binding affinity ($E_{d,G} \approx -12$ kcal/mol). This compound occupied the binding site through similar interactions with important amino acids

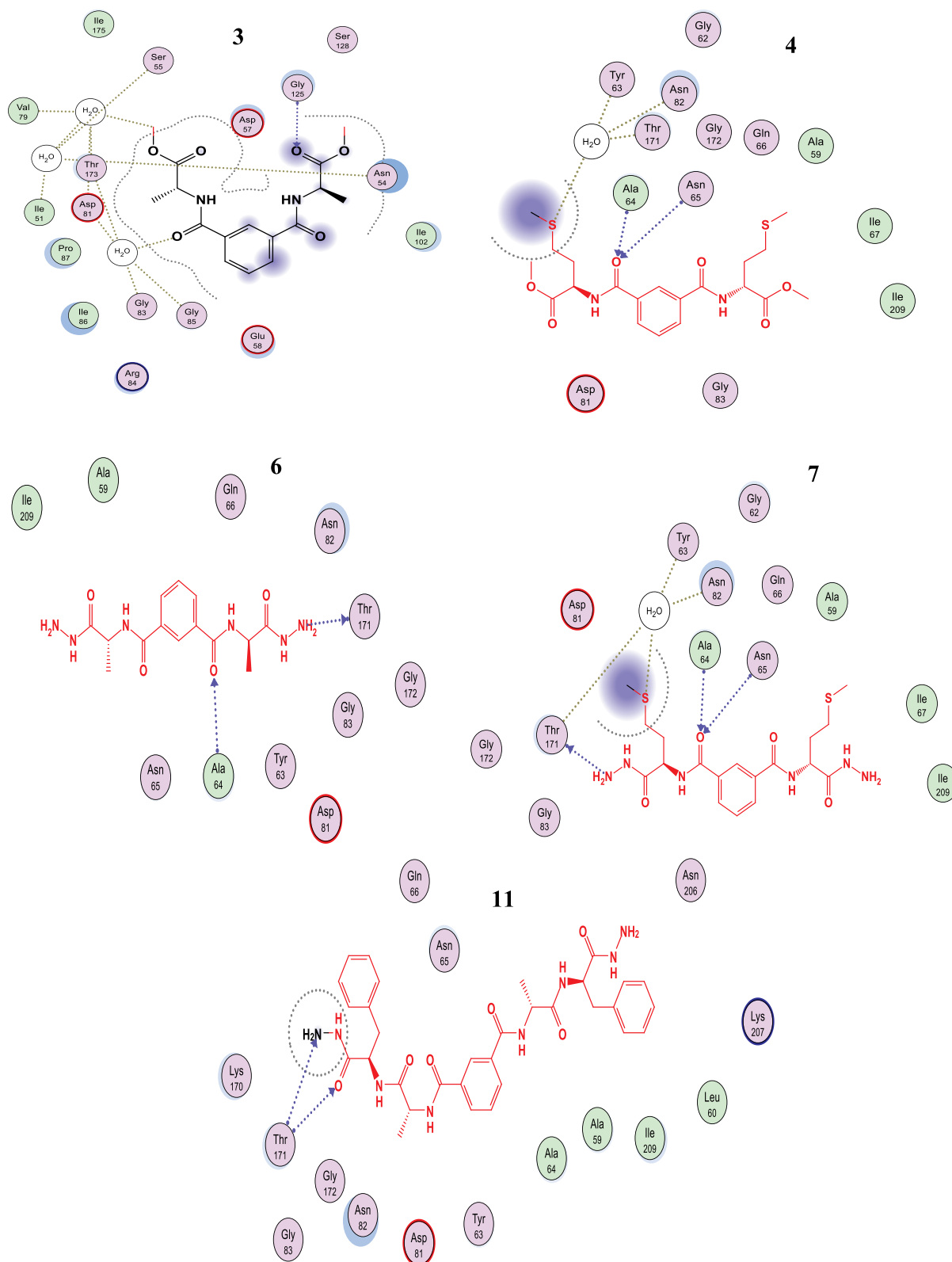


Figure 4 The 2D binding mode of 3,4,7,8 and 11 into the active site of DNA Gyrase, H-bond represented as blue dashed color.

of the binding sites for compound **3** with Gln70, Ala254, Leu319, and Arg324; for compound **5** with Arg94; for **6** with Glu255, Leu319, and Phe385; and for **8** with Ala254,

Thr258, and Cys392. The bis-peptide **9** to **11** derivatives showed lower binding affinity than the amino acid derivatives **3** to **8** (Table 4). Bis-peptide **9** formed two

H-bonds with Arg93 and two π - π interactions with Arg94. Compound **10** formed the following interactions: H-bonds with Hip390, Arg93, and π - π bonds with Cys392. While compound **11** produced two bonds with Val393 and Arg389 (Figure S3, S4; Table S3, Supplementary Material).

ADMET Profile in silico

For any bioactive molecule to be considered a potential drug should possess low toxicity with good pharmacokinetic properties. ADME/T has interfered with several agents from reaching the clinical trials. Hence, the ADME/T profile for compounds **3** to **11** was predicted using MOE and admet-SAR.[52] The pharmacokinetic descriptors in Table 5 indicate that compounds **3** to **8** fulfilled the Lipinski rules, while the other compounds **9** to **11** showed more than one violation when the Lipinski restrictions were applied. The low lipophilicity values of compound **5** (Log P < 5 mol/L) and generally for all compounds (**3–11**) predicted good permeability and absorption across the cell membrane.⁵⁷ The topological polar surface area (TPSA) is another important pharmacokinetic parameter that predicts the permeability of the membrane by polar atoms.³⁶ The passive absorption value is $TPSA > 140 \text{ \AA}^2$. Compounds (**7** to **11**) were predicted to be passively absorbed with values ranging from ~ 168 to -226 \AA^2 . The tested compounds did not act as inhibitors or substrates against the P-glycoprotein isoforms P-GPI, II-I and P-GPI, II-S, thus, it can be safely presumed that these compounds would not exert any relevant activity.^{58,59} Based on the above findings, synthesized compounds **3** to **11** may present good absorption properties.

As shown in Table 5, (i) our compounds did not show any inhibition or activation of any Cytochrome P450 (CYP450) isoform. CYP450 is an important detoxification enzyme in the body and is mainly located in liver. It oxidizes xenobiotics to facilitate their excretion. CYP450 can activate or deactivate many drugs. It is therefore important to evaluate the potential inhibitory effects against CYP450 enzymes (ie, CYP1A2, CYP2C19, CYP2C9, CYP2D6, CYP3A4) or activation of CYP2D6 or CYP3A4 for the synthesized compounds against 18,000 compounds known to inhibit or mimic these isoforms. (ii) There were no observed carcinogenic or tumorigenic effects induced by the synthesized compounds, comparing 981 known carcinogenic modules obtained from the

Carcinogenic Potency Database (CPDB).⁶⁰ (iii) The chemical structure of each of the investigated compounds was compared with 10,000 models that have been previously tested in rats to predict the lethal dose value for these compounds. Our findings showed low values of LD50 (~ 2.2 – 3.07 mg/kg) for the synthesized compounds. (iv) The LOAE was predicted by comparing the tested compounds with 576 known agents. This test is applied in chronic evaluation studies for the identification of the lowest doses of chemicals tolerable over a long period. The predictor simulated that no LOAE was detectable for any of the investigated compounds. (v) The synthesized compounds exerted no mutagenic effect on DNA of microorganisms when tested in the Ames test with over 8000 compounds. The Ames test is a widely used examination of mutagenic effects of bioactive molecules against microorganism DNA.^{61,62} (vi) A weak inhibitory action was determined against the human Ether-a-gogo-Related Gene (hERG), which can lead to long QT syndrome, for the tested compounds.^{63,64} (viii) The tested compounds did not exhibit any skin sensitivity, when tested utilizing 254 compounds able to induce skin sensitization, or hepatotoxic effects when evaluated against 531 compounds which have liver associated side effects in human. (vii) Based on the above characteristics, we can conclude that the synthesized compounds may have a good oral bioavailability without any potential for marked health adverse effects via rodent toxicity profiles.

This study introduces novel peptide derivatives based on two amino acids: alanine and methionine. Based on the activity results and docking studies, the Ala, Met, and Ala-Phe, at N ^{α} -1, 3-benzenedicarbonyl do not induce any meaningful differences in antimicrobial or antioxidant activities. The docking study showed that a binding affinity ranging, -5 to -7 kcal/mol against DNA gyrase as the model of anti-microbial activity, and against antifungal model these compounds showed higher binding score than antibacterial results (-7 to -8 kcal/mol). In addition, all the compounds were evaluated as anticancer agents in three cell lines MCF-7, HCT-116, and HepG-2. In case of MCF-7, the presence of Met.OMe (**7**) and Ala.hydrazide (**8**) in the parent compound increased the activity against all similar members of the tested compounds. Furthermore, the parent compound that carried other fragments did not present any significant variation in cytotoxic activity. In the HCT-116 cell line, introducing Ala (**3**) and Ala.Phe (**9**) fragments into the parent compound showed

highest activity against HCT116, with inhibition potency percentage (56% and 61%), respectively. These compounds showed binding scores -9.7 and -13.7 kcal/mol, respectively for compounds **3** and **9**. The presence of Met. OMe (**6**) and its derivatives from the hydrazide (**8**), in addition to the presence of Ala.Phe.OMe (**10**) showed moderate activity. The other compounds showed the lowest activity among the compounds. In the HepG-2 cell line, all compounds showed a low toxic potency in the range of 2–10%.

Conclusion

In the present study, nine novel peptide derivatives, characterized by the general stretcher, N α -1, 3-benzenedicarbonyl-*bis*-(amino acid) and dipeptide candidates were synthesized. All synthesized compounds showed potent antimicrobial activity of different degrees against different pathogenic bacteria and fungi, and only compounds **5** and **7** had the lowest inhibitory activity. These compounds showed good antimicrobial activity against important pathogens so they can be considered potent therapeutic drugs that exert biological control, warranting further investigations. Furthermore, three human cancer cell lines (MCF-7, HCT-116, and HepG-2) were used to evaluate the anticancer potency of the synthesized compounds. The novel compounds (**3–11**) showed a strong cytotoxic effect in the MCF-7 human breast cancer cell line with 95.1% and 79.8% cell inhibition, respectively. The docking results suggest good fitting through different hydrogen bond interactions with protein residues allowing eliciting anticancer activity. The ADME/T profile in silico suggested compounds **3–11** may have a good absorption property.

Sample Availability

Samples of the compounds (**1–11**) are available from Dr Gaber O. Moustafa.

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Disclosure

The authors declare no conflicts of interest in this work.

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