Synthesis, Cytotoxic Activity and 2D-QSAR Study of Some Imidazoquinazoline Derivatives

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Abstract: A novel series of 4-substituted amino-7,8-dimethoxy-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one derivatives was designed, synthesized and tested for their antitumor activity against a human mammary carcinoma cell line (MCF7). Compound 5a was found to be the most active derivative. Physico-chemical parameters were also determined and revealed that most of the compounds obeyed the “rule of five” properties with good absorption percentages. 2D-QSAR studies revealed a well predictive and statistically significant and cross validated QSAR model that helps to explore some expectedly potent compounds.

Keywords: imidazolinone; imidazo[1,5-a]quinazoline; antitumor activity; Lipinski’s parameters; 2D-QSAR

1. Introduction

Struggling efforts for the treatment and eradication of cancer have grown tremendously in the last few years led to the decrease in cancer death rates by 1.8% per year in men and by 1.5% per year in women [1]. Cancer is characterized by a deregulation of the cell cycle, resulting in progressive loss of the cellular differentiation and non-controlled cellular growth [2,3]. The control of disseminated tumor growth by systemically active chemotherapeutic agents remains a major challenge for cancer chemotherapy, despite decades of focused efforts. Although there are some notable successes with certain forms of cancer [4], the discovery and development of novel therapeutic agents for the treatment of cancer has a vital importance [5–11].

Imidazole, a small bioactive molecule, is a prominent structural motif found in numerous biologically active compounds. Interestingly, imidazolinones such as I [12] (Figure 1) and methoxyquinazoline
derivatives II, gefitinib III and IV [13–16] (Figure 1) all display significant antitumor activity. Hybrid molecules combining the imidazole and quinazoline moieties in either linear or angular imidazoquinazolines were designed and demonstrated promising antitumor activity [17–19]. Furthermore, incorporation of other heterocyclic moieties such as piperazine [20,21], piperidine [22,23], imidazole [24] and triazine [25] moieties was reported to enhance the antitumor activity too.

Figure 1. Structures of some active antitumor agents from the literature.

Inspired by the aforementioned facts, it was deemed of interest to use the hybridized 7,8-dimethoxyimidazoquinazoline ring system as a scaffold for the design of new molecules with potential antitumor activity. The target compounds incorporate different heterocyclic moieties such as piperazine, piperidine, or imidazole. The novel compounds were tested for their antitumor activity. Physico-chemical parameters were also calculated to obtain their “rule of five” properties. Moreover, 2D-QSAR studies were also applied to correlate between the structures of the synthesized compounds and their pharmacological activities.

2. Results and Discussion

2.1. Chemistry

The synthetic strategy for the preparation of the target compounds 2a,b, 3a,b, 4, 5a,b, 6a–d, 7a–d, 8 and 9 is illustrated in Schemes 1–3. Imidazolinone derivatives 2a,b were obtained through the reaction of oxazolone derivatives 1a,b, previously synthesized according to the literature method [26,27] with methyl 2-amino-4,5-dimethoxybenzoate in glacial acetic acid in the presence of sodium acetate. This reaction proceeded through open intermediates which were recyclized in the presence of glacial acetic acid to afford the imidazolinones [28]. Reaction of 2a,b with hydrazine hydrate afforded either the benzohydrazide derivatives 3a,b or imidazo[1,5-a]quinazolinone derivative 4 depending on the reaction conditions.
Scheme 1. Synthesis of the target compounds 1–4.

Reagents and conditions: (i) methyl 2-amino-4,5-dimethoxybenzoate, CH₃COOH, CH₃COONa; (ii) Hydrazine hydrate, methanol, 2 h; (iii) Hydrazine hydrate, 80% ethanol, 24 h.

Scheme 2. Synthesis of the target compounds 5–8.

Reagents and conditions: (i) Cl(CH₂)ₙCOCl, dry DMF; (ii) Piperidine or N-methylpiperazine, acetonitrile, K₂CO₃; (iii) 2/3-chloro-1-(piperidin-1-yl/4-methylpiperazin-1-yl)ethanone/propan-1-one, dry DMF, triethylamine; (iv) 2-chloroacetamide, dry DMF.
Scheme 3. Synthesis of the target compound 9.

Reagents and conditions: (i) KCNO, acetic acid.

Refluxing 2a, b with equimolar amounts of hydrazine hydrate in methanol for 2 h yielded the benzohydrazide derivatives 3a, b, whereas, the imidazoquinazoline derivative 4 was obtained according to the reported reaction conditions [29] using excess hydrazine hydrate in aqueous ethanol 80% at reflux for 24 h. (Scheme 1). On the other hand, the reaction of compound 4 with chloroacetyl chloride in DMF led to the formation of 2-chloro-N-(7,8-dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl)acetamide (5a). Similarly, the reaction with chloropropionyl chloride gave 5b. Reaction of 5a, b with secondary amines (namely, piperidine and N-methylpiperazine) in dry acetonitrile in the presence of potassium carbonate yielded compounds 6a–d. Ethylamino or propylamino-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one derivatives 7a–d were obtained by reacting compound 4 with 2/3-chloro-1-(piperidin-1-yl/4-methylpiperazin-1-yl)ethanone/propan-1-one in dry DMF in the presence of triethylamine. Moreover, formation of the oxotriazine ring was achieved via the reaction of the N-amino derivative 4 with chloroacetamide in DMF to yield 11,12-dimethoxy-8-phenyl-3,4-dihydro-2H-imidazo[1,5-a][1,2,4]triazino[2,3-c]quinazolin-2-one (8) as depicted in Scheme 2. As shown in Scheme 3, the cyclized product 9 was obtained starting with the chloroacetamide derivative 5a. Reaction with potassium cyanate yielded the 1-(7,8-dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-4(5H)-yl)-1H-imidazole-2,5-dione(9). The newly synthesized compounds were confirmed via elemental analyses and spectral data (IR, 1H-NMR, 13C-NMR and mass spectra).

2.2. Preliminary in-Vitro Antitumor Screening

All the synthesized compounds were tested for their antitumor activity against human mammary carcinoma cell line (MCF7) at the National Cancer Institute, Cairo University [30]. From the observed antitumor data (Table 1), it was noticed that imidazolinone derivative 3a and imidazoquinazoline derivatives 5a, 6a, 7a and 7b showed IC50 values of 27.1, 10.6, 26.4, 21.6 and 26.8 µM, respectively. Also, compounds 5a and 7a revealed remarkable activity comparable to that of the reference drug mitoxantrone.

Structure-activity correlations, based on the human mammary carcinoma cell line (MCF7), revealed that the chloroacetyl derivative 5a was more active than its chloropropionyl analogue 5b. Regarding the effect of the substituent on 5a, b, the results of 6a–d and 7a–d showed that a piperidinyl moiety was more active than a N-methylpiperazinyl one, and an acetyl moiety was more active than its corresponding propionyl moiety. Also, derivatives in which the carbonyl moiety was directly attached
to a piperidinyl/piperazinyl (compounds 7a–d) were more active than those in which the carbonyl was attached to an N-aminoimidazoquinazoline (compounds 6a–d).

Table 1. Cytotoxic activity of the newly synthesized compounds against breast cancer MCF7 cell line.

| Compd. No | IC_{50} µg/mL | IC_{50} µM |
|-----------|----------------|------------|
| 2a        | 18.0           | 40.6       |
| 2b        | 21.8           | 46.1       |
| 3a        | 12.0           | 27.1       |
| 3b        | 42.4           | 89.7       |
| 4         | 12.2           | 36.2       |
| 5a        | 4.4            | 10.6       |
| 5b        | 14.0           | 32.8       |
| 6a        | 12.2           | 26.4       |
| 6b        | 17.2           | 36.0       |
| 6c        | 19.0           | 39.9       |
| 6d        | 21.4           | 43.6       |
| 7a        | 10.0           | 21.6       |
| 7b        | 12.8           | 26.8       |
| 7c        | 17.4           | 36.5       |
| 7d        | 18.4           | 37.5       |
| 8         | 21             | 56.1       |
| 9         | 19.0           | 45.5       |
| Mitoxantrone | 10.1       | 22.7       |

2.3. Physico-Chemical Parameters

Computational study for the prediction of physico-chemical parameters of all synthesized compounds has been executed. The computed molecular properties are shown in Table 2. Numbers of rotatable bonds, log P, molecular polar surface area (PSA), number of hydrogen bond accepter and donor atoms of Lipinski’s “rule of five” [31] were calculated using the Molinspiration online property calculation toolkit [32]. The degree of absorption is expressed by the percentage of absorption (%ABS), that was calculated by using \(\%\text{ABS} = 109 - (0.345 \times \text{TPSA})\) [33]. From all these parameters, it was found that, all the synthesized derivatives exhibited %ABS ranging from 61.41% to 84.91%. Furthermore, most of the compounds obeyed the “rule of five” properties.

2.4. 2D-QSAR Studies

A 2D-QSAR study was performed in order to find a mathematical correlation between the structures of the newly synthesized compounds and their antitumor activity [34] that expressed as \(-\log IC_{50}\) values, using Molecular Operating Environment (MOE) software package.
Table 2. Calculated absorption (%ABS), polar surface area (PSA) and Lipinski parameters of the newly synthesized compounds.

| Compd. No | %ABS  | TPSA  | nrotb | nON ≤ 10 | nOHNH ≤ 5 | miLogP ≤ 5 | MW ≤ 500 | n violations ≤ 1 |
|-----------|-------|-------|-------|----------|-----------|------------|----------|-----------------|
| 2a        | 81.51 | 79.67 | 7     | 7        | 0         | 4.38       | 444.15   | 0               |
| 2b        | 78.32 | 88.90 | 8     | 8        | 0         | 4.43       | 472.16   | 0               |
| 3a        | 71.57 | 108.48| 6     | 8        | 3         | 2.93       | 442.16   | 0               |
| 3b        | 68.38 | 117.72| 7     | 9        | 3         | 2.98       | 472.49   | 0               |
| 4         | 79.08 | 83.80 | 3     | 7        | 2         | 2.34       | 336.35   | 0               |
| 5a        | 79.02 | 86.87 | 5     | 8        | 1         | 2.64       | 412.82   | 0               |
| 5b        | 79.02 | 86.87 | 6     | 8        | 1         | 2.91       | 426.10   | 0               |
| 6a        | 77.91 | 90.11 | 6     | 9        | 1         | 2.97       | 461.51   | 0               |
| 6b        | 76.79 | 93.35 | 6     | 10       | 1         | 1.96       | 476.52   | 0               |
| 6c        | 77.91 | 90.11 | 7     | 9        | 1         | 3.24       | 475.53   | 0               |
| 6d        | 76.79 | 93.35 | 7     | 10       | 1         | 2.23       | 490.55   | 0               |
| 7a        | 77.91 | 90.11 | 6     | 9        | 1         | 2.97       | 461.51   | 0               |
| 7b        | 76.79 | 93.35 | 6     | 10       | 1         | 1.96       | 476.52   | 0               |
| 7c        | 77.91 | 90.11 | 7     | 9        | 1         | 3.24       | 475.53   | 0               |
| 7d        | 76.79 | 93.35 | 7     | 10       | 1         | 2.23       | 490.55   | 0               |
| 8         | 80.65 | 82.17 | 3     | 8        | 1         | 2.61       | 375.38   | 0               |
| 9         | 71.13 | 109.74| 4     | 10       | 0         | 1.67       | 417.37   | 0               |

The most relevant descriptors derived for modeling the antitumor activity are listed in Table 3 and the relation between the obtained results of experimentally observed and predicted values of antitumor activity were presented in Table 4. 2D-QSAR models were validated with the leave one out (LOO) method. The best derived QSAR model for the newly synthesized derivatives was presented by the following triparametric equation with correlation coefficient ($R^2$) = 0.92:

$$[IC_{50} = -617.62254 + 757.93323 \times PEOE_{VSA \_FHYD} + 306.47775 \times glob - 4.31115 \times MNDO_{dipole}]$$

(1)

where $n = 9$, RMSE = 3.4641, $R^2 = 0.92722$, $n$: number of compounds used for construction of model; RMSE: Root mean square error; $R^2$: correlation coefficient.

From this model, it was observed that PEOE_VSA_FHYD (Fractional hydrophobic Van der Waal surface area) and glob (Globularity, or inverse condition number (smallest eigenvalue divided by the largest eigen value) of the covariance matrix of atomic coordinates. A value of 1 indicates a perfect sphere while a value of 0 indicates a two- or one-dimensional object) were positively correlated with antitumor activity but MNDO_dipole (The dipole moment calculated using the MNDO Hamiltonian [MOPAC]) was negatively correlated with activity indicated that increasing in the values PEOE_VSA_FHYD is beneficial for antitumor activity and the experimental IC$_{50}$ was matched with the expected IC$_{50}$. 


Table 3. The most relevant descriptors derived for modeling the antitumor activity.

| Compd. No | $P \text{IC}_{50}$ | $\text{PEOE}_\text{VS}$ | $\text{Glob}$ | $\text{MNDO}_{\text{di}}$ |
|-----------|---------------------|-------------------------|----------------|-------------------------|
| 4         | 36.2                | 0.8547                  | 0.0796         | 3.7779                  |
| 5a        | 10.6                | 0.8258                  | 0.0670         | 4.2328                  |
| 5b        | 32.8                | 0.8332                  | 0.0725         | 4.3450                  |
| 6a        | 26.4                | 0.8517                  | 0.0514         | 1.6865                  |
| 6b        | 36.0                | 0.8602                  | 0.0519         | 0.0000                  |
| 6c        | 39.9                | 0.8571                  | 0.1028         | 5.6151                  |
| 6d        | 43.6                | 0.8650                  | 0.0387         | 1.9895                  |
| 7a        | 21.6                | 0.8517                  | 0.0389         | 5.1133                  |
| 7b        | 26.8                | 0.8602                  | 0.0591         | 5.0499                  |
| 7c        | 36.5                | 0.8517                  | 0.0690         | 3.9824                  |

Table 4. Experimentally observed, predicted activity of the new compounds and their residual.

| Compd. No | Activity | $P \text{IC}_{50}$ | SPRED | SRES  |
|-----------|----------|---------------------|-------|-------|
| 4         | 36.2     | 38.2548             | −2.0548 |
| 5a        | 10.6     | 10.5372             | 0.0628 |
| 5b        | 32.8     |                     |       |
| 6a        | 26.4     | 36.3854             | −9.9854 |
| 6b        | 36.0     | 50.2617             | −14.2617 |
| 6c        | 39.9     | 39.2977             | 0.6023 |
| 6d        | 43.6     | 41.2839             | 2.3161 |
| 7a        | 21.6     | 17.7661             | 3.8339 |
| 7b        | 26.8     | 30.6762             | −3.8762 |
| 7c        | 36.5     | 35.9551             | 0.5448 |

3. Experimental

3.1. General Information

Unless specified all chemicals were of commercial grade, obtained from Aldrich Chemical Co. (Milwaukee, WI, USA), and used without further purification. Melting points were carried out by the open capillary tube method using a Gallenkamp digital melting point Griffin apparatus 1901 and they were uncorrected. Elemental microanalyses were recorded at the Regional Center of Mycology and Biotechnology, Al-Azhar University. Infrared Spectra were recorded on Shimadzu infrared spectrophotometer IR Affinity-1 (FTIR-8400S, Kyoto, Japan) and expressed in wave number (cm$^{-1}$), using potassium bromide discs. NMR spectra were recorded on a Varian Mercury VX 300 spectrometer ($^1$H: 300, $^{13}$C APT: 75 MHz) or a 400 MHz high performance digital FT-NMR spectrophotometer using TMS as an internal standard. The exchangeable protons (NH, and OH) were exchanged by D$_2$O. Mass Spectra was recorded on Shimadzu QP-2010 plus. All reactions were monitored by thin layer chromatography. Silica gel/TLC-cards DC-Alufolien-Kieselgel with fluorescent indicator 254 nm; layer thickness 0.2 mm; 20 × 20 cm aluminum cards were used. Petroleum ether/ethyl acetate (1:1) or (1:2) was the adopted solvent system. Compounds 1a,b [26,27] were prepared according to reported procedures.
3.2. Chemistry

3.2.1. General Procedure for the Preparation of 2a,b

A mixture of 1a,b (10 mmol) and methyl 2-amino-4,5-dimethoxybenzoate (12 mmol, 2.52 g), freshly prepared fused sodium acetate (0.3 g) was heated in a boiling water bath in glacial acetic acid (10 mL) for 3 h. The crystalline product separated on cooling was filtered, washed with water, dried, and recrystallized from ethanol.

Methyl 2-(4-benzylidene-5-oxo-2-phenyl-4,5-dihydroimidazol-1-yl)-4,5-dimethoxybenzoate (2a). Yield: 81%; mp 221–223 °C; IR (KBr, cm⁻¹): 3086 (CH aromatic), 2958, 2935 (CH aliphatic), 1693, 1674 (C=O); ¹H-NMR (DMSO-d₆): δ 3.65 (s, 3H, COOCH₃), 3.77 (s, 3H, OCH₃), 3.88 (s, 3H, OCH₃), 7.36–7.43 (m, 4H, benzyl. 3,4,5-CHs + benzo.6-CH) 7.54–7.63 (m, 6H, phenyl CHs + benzo.3-CH), 8.04 (d, 2H, J = 6.9, benzyl. 2,6-CHs), 8.57 (s, 1H, CH); ¹³C-NMR (DMSO): 52.4, 56.09, 56.1, 103.4, 107.1, 112.6, 128.4, 128.7, 129.1, 129.7, 130.0, 130.4, 131.9, 132.3, 134.0, 137.1, 144.2, 154.0, 163.9, 167.4, 167.8; MS, m/z (%): M⁺, 442 (23.54%), 105 (100%). Anal. Calcd. for C₂₆H₂₂N₂O₅ (442.46): C, 70.58; H, 5.01; N, 6.33. Found: C, 70.32; H, 5.12; N, 6.44%.

Methyl 4,5-dimethoxy-2-(4-(4-methoxybenzylidene)-5-oxo-4,5-dihydro-1H-imidazol-1-yl)-benzoate (2b). Yield: 84%; mp 211–212 °C; IR (KBr, cm⁻¹): 3062 (CH aromatic), 2939, 2839 (CH aliphatic), 1720, 1678 (C=O); ¹H-NMR (DMSO-d₆): δ 3.63 (s, 3H, COOCH₃), 3.84 (s, 6H, OCH₃), 3.86 (s, 3H, OCH₃), 6.95 (d, 2H, J = 9, benzyl. 3, 5-CHs), 7.39 (s, 1H, benzo.6-CH) 7.54–7.63 (m, 6H, phenyl CHs + benzo.3-CH), 8.04 (d, 2H, J = 6.9, benzyl. 2,6-CHs), 8.58 (s, 1H, CH); ¹³C-NMR (DMSO): 51.8, 55.5, 55.7, 56.1, 102.7, 106.4, 112.5, 114.4, 126.0, 127.6, 128.2, 130.5, 131.4, 131.7, 133.5, 134.2, 136.7, 143.5, 153.4, 160.0, 163.5, 166.8, 167.2; MS, m/z (%): M⁺, 472 (68.44%), 105 (100%). Anal. Calcd. for C₂₇H₂₄N₂O₆ (472.48): C, 68.63; H, 5.12; N, 5.93. Found: C, 68.85; H, 5.00; N, 6.01%.

3.2.2. General Procedure for the Preparation of 3a,b

A solution of 2a,b (10 mmol), hydrazine hydrate (12 mmol, 0.60 g) in methanol (50 mL) was refluxed for 2 h. The mixture was concentrated under vacuum, and water was added to the residue, which was filtered, dried, and recrystallized from ethanol.

2-(4-Benzylidene-5-oxo-2-phenyl-4,5-dihydroimidazol-1-yl)-4,5-dimethoxybenzo- hydrazide (3a). Yield: 71%; mp 151–153 °C; IR (KBr, cm⁻¹): 3456, 3356, 3305 (NH₂, NH), 3059 (CH aromatic), 2997, 2951 (CH aliphatic), 1674 (C=Os); ¹H-NMR (DMSO-d₆): δ 3.73 (s, 3H, OCH₃), 3.79 (s, 3H, OCH₃), 6.45 (s, 2H, NH₂), 7.11–7.38 (m, 4H, benzyl. 3,4,5-CHs + benzo.6-CH), 7.42–7.49 (m, 6H, phenyl CHs + benzo.3-CH), 7.79 (d, 2H, J = 6.9, benzyl. 2,6-CHs), 8.59 (s, 1H, CH), 9.33 (s, 1H, NH); ¹³C-NMR (DMSO): 55.1, 55.9, 99.0, 112.2, 126.1, 127.3, 127.9, 128.0, 129.0, 131.1, 133.9, 138.2, 139.2, 143.5, 148.2, 154.7, 166.0, 167.2, 170.5; MS, m/z (%): M⁺, 442 (0.25%), 50 (100%). Anal. Calcd. for C₂₈H₂₂N₄O₄ (442.46): C, 67.86; H, 5.01; N, 12.66. Found: C, 67.86; H, 5.12; N, 12.93%.
2-(4-(4-Methoxybenzylidene)-5-oxo-2-phenyl-4,5-dihydroimidazol-1-yl)-4,5-dimethoxybenzohydrazide (3b). Yield: 74%; mp 201–203 °C; IR (KBr, cm\(^{-1}\)): 3441, 3244, 3200 (NH, NH), 3050 (CH aromatic), 2924, 2835 (CH aliphatic), 1674, 1658 (C=O); \(^1\)H-NMR (DMSO-\(d_6\)): \(\delta\) 3.71 (s, 3H, OCH\(_3\)), 3.80 (s, 3H, OCH\(_3\)), 3.86 (s, 3H, OCH\(_3\)), 6.88 (s, 2H, NH\(_2\)), 6.98 (d, 2H, \(J = 8.7\), benzyl. 3, 5-CH\(_s\)), 7.39 (s, 1H, benzo.6-CH), 7.55–7.63 (m, 6H, phenyl CH\(_s\) + benzo.3-CH), 8.04 (d, 2H, \(J = 6.9\), benzyl. 2,6-CH\(_s\)), 8.60 (s, 1H, CH), 10.30 (s, 1H, NH); \(^13\)C-NMR (DMSO): 55.7, 55.9, 105.0, 107.9, 112.0, 127.1, 128.0, 128.3, 129.5, 135.1, 143.9, 148.7, 154.5, 160.3, 166.3; MS, \(m/z\) (%): M\(^+\) + 1, 337 (0.25%), 196 (100%). Anal. Calcd. for C\(_{18}\)H\(_{16}\)N\(_4\)O\(_3\) (336.34): C, 64.28; H, 4.79; N, 16.66. Found: C, 64.37; H, 4.86; N, 16.92%.

3.2.3. Synthesis of 4-Amino-7,8-dimethoxy-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one (4)

To a suspension of the appropriate 2a,b (10 mmol) in aqueous ethanol (80%), hydrazine hydrate (30 mmol, 1.50 g) was added. The mixture was refluxed for 24 h. The resulting solution was distilled under vacuum and the residue was triturated with ice water. The separated solid was filtered, washed with water, and crystallized from ethanol. Yield: 64% from 2a, 72% from 2b; mp 281–283 °C; IR (KBr, cm\(^{-1}\)): 3317, 3267 (NH\(_2\)), 3074 (CH aromatic), 2927, 2839 (CH aliphatic), 1680 (C=O); \(^1\)H-NMR (DMSO-\(d_6\)): \(\delta\) 3.89 (s, 3H, OCH\(_3\)), 3.90 (s, 3H, OCH\(_3\)), 4.42 (s, 2H, NH\(_2\)), 7.19 (s, 1H, imidazo.9-CH), 7.45–8.20 (m, 8H, Ar-H); \(^13\)C-NMR (DMSO): 55.7, 55.9, 105.0, 107.9, 112.0, 127.1, 128.0, 128.3, 129.5, 135.1, 143.9, 148.7, 154.5, 160.3, 166.3; MS, \(m/z\) (%): M\(^+\), 472 (0.38%), 57 (100%). Anal. Calcd. for C\(_{26}\)H\(_{24}\)N\(_4\)O\(_5\) (472.49): C, 66.09; H, 5.12; N, 11.86. Found: C, 66.34; H, 5.45; N, 12.24%.

3.2.4. General Procedure for the Preparation of 5a,b

A solution of compound 4 (5 mmol, 1.68 g) in dry DMF (5 mL) containing chloroacetyl chloride or chloropropionyl chloride (5.5 mmol) was stirred at room temperature (25–30 °C) for 24 h. The solution was poured onto crushed ice and the resulting solid was filtered, washed and crystallized from ethanol. 2-Chloro-N-(7,8-dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl) acetamide (5a). Yield: 82%; mp 176–178 °C; IR (KBr, cm\(^{-1}\)): 3224 (NH), 3035 (CH aromatic), 2947, 2854 (CH aliphatic), 1674 (C=O); \(^1\)H-NMR (DMSO-\(d_6\)): \(\delta\) 3.81 (s, 3H, OCH\(_3\)), 3.86 (s, 3H, OCH\(_3\)), 4.44 (s, 2H, CH\(_2\)), 7.40–8.20 (m, 8H, Ar-H), 11.47 (s, 1H, NH); \(^13\)C-NMR (DMSO): 43.3, 55.4, 55.7, 103.3, 107.6, 112.0, 127.3, 128.0, 128.2, 129.0, 135.0, 144.1, 144.2, 153.1, 153.4, 164.9, 167.0; MS, \(m/z\) (%): M\(^+\) + 2, 414 (0.31%), 412 (0.91%), 105 (100%). Anal. Calcd. for C\(_{20}\)H\(_{17}\)ClN\(_4\)O\(_4\) (412.82): C, 58.19; H, 4.15; N, 13.57. Found: C, 58.34; H, 4.01; N, 13.78%.

3-Chloro-N-(7,8-dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl) propamide (5b). Yield: 75%; mp 209–210 °C; IR (KBr, cm\(^{-1}\)): 3248 (NH), 3055 (CH aromatic), 2951, 2824 (CH aliphatic), 1670 (C=O); \(^1\)H-NMR (DMSO-\(d_6\)): \(\delta\) 2.88 (t, 2H, CH\(_2\)), \(J = 7.8\), 3.79 (t, 2H, CH\(_2\)), \(J = 7.8\), 3.90 (s, 3H, OCH\(_3\)), 3.92 (s, 3H, OCH\(_3\)), 7.15–8.60 (m, 8H, Ar-H), 10.7 (s, 1H, NH); \(^13\)C-NMR (DMSO): 36.0, 52.2, 54.4, 55.7, 103.9, 108.2, 112.9, 127.1, 128.0, 128.2, 128.9,131.5, 133.3, 135.0,
3.2.5. General Procedure for the Preparation of 6a–d

A mixture of equimolar amounts of the appropriate 5a,b and the corresponding secondary amine (2 mmol) in dry acetonitrile (30 mL) containing potassium carbonate (4 mmol, 0.55 g) was refluxed for 12 h and the reaction mixture was filtered hot. The solid which was separated upon storing the clear reaction mixture at room temperature overnight was collected and crystallized from ethanol.

**N-(7,8-Dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl)-2-(piperidin-1-yl)acetamide (6a).** Yield: 77%; mp 186–189 °C; IR (KBr, cm⁻¹): 3217 (NH), 3016 (CH aromatic), 2943, 2879 (CH aliphatic), 1693, 1670 (C=O); ¹H-NMR (DMSO-d₆): δ 1.42–1.43 (m, 2H, CH₂), 1.63–1.68 (m, 4H, 2CH₂), 2.46 (t, 4H, 2CH₂, J = 8.7), 3.07 (s, 2H, CH₂), 3.80 (s, 3H, OCH₃), 3.86 (s, 3H, OCH₃), 7.40–8.40 (m, 8H, Ar-H), 11.8 (s, 1H, NH); ¹³C-NMR (DMSO): 23.4, 25.2, 52.2, 55.4, 55.4, 62.9, 102.9, 106.8, 112.1, 127.4, 128.1, 128.3, 129.0, 135.9, 144.3, 144.4, 153.1, 153.4, 166.5, 169.6; MS, m/z (%): M⁺, 461 (47.5%), 254 (100%). Anal. Calcd. for C₂₅H₂₇N₅O₄ (461.51): C, 65.06; H, 5.90; N, 15.17. Found: C, 65.38; H, 5.93; N, 15.42%.

**N-(7,8-Dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl)-2-(4-methylpiperazin-1-yl)acetamide (6b).** Yield: 81%; mp 167–168 °C; IR (KBr, cm⁻¹): 3205 (NH), 3030 (CH aromatic), 2985, 2947, 2816 (CH aliphatic), 1689, 1670 (C=O); ¹H-NMR (DMSO-d₆): δ 2.19 (s, 3H, CH₃), 2.46–2.53 (m, 8H, 4CH₂), 3.75 (s, 2H, CH₂), 3.80 (s, 3H, OCH₃), 3.86 (s, 3H, OCH₃), 7.18–8.41 (m, 8H, Ar-H), 11.76 (s, 1H, NH); ¹³C-NMR (DMSO): 46.2, 52.5, 53.3, 54.7, 54.9, 56.0, 56.1, 62.5, 103.6, 107.5, 112.7, 127.9, 128.6, 129.05, 136.4, 144.0, 153.7, 167.0, 169.8; MS, m/z (%): M⁺, 476 (19.9%), 55 (100%). Anal. Calcd. for C₂₅H₂₈N₆O₄ (476.52): C, 63.01; H, 5.92; N, 17.64. Found: C, 63.13; H, 6.12; N, 17.79%.

**N-(7,8-Dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl)-3-(piperidin-1-yl)propamide (6c).** Yield: 66%; mp 105–108 °C; IR (KBr, cm⁻¹): 3263 (NH), 3010 (CH aromatic), 2931, 2894 (CH aliphatic), 1674 (br., C=O); ¹H-NMR (DMSO-d₆): δ 1.37–1.48 (m, 6H, 3CH₂), 2.53–2.55 (m, 4H, 2CH₂), 2.61 (t, 2H, 2CH₂, J = 6.3), 3.69 (t, 2H, 2CH₂, J = 9.3), 3.90 (s, 3H, OCH₃), 3.91 (s, 3H, OCH₃), 7.20–8.10 (m, 8H, Ar-H), 10.85 (s, 1H, NH); ¹³C-NMR (DMSO): 22.2, 25.6, 35.2, 52.6, 53.7, 54.4, 55.9, 56.0, 56.1, 104.6, 108.4, 112.6, 127.4, 127.9, 128.2, 128.6, 128.7, 129.1, 136.3, 144.2, 153.6, 167.6, 170.7; MS, m/z (%): M⁺, 475 (0.5%), 98 (100%). Anal. Calcd. for C₂₆H₂₉N₅O₄ (475.53): C, 65.67; H, 6.15; N, 14.73. Found: C, 65.46; H, 5.84; N, 14.50%.

**N-(7,8-Dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinazolin-5(4H)-yl)-3-(4-methylpiperazin-1-yl)-propamide (6d).** Yield: 69%; mp 130–132 °C; IR (KBr, cm⁻¹): 3217 (NH), 3059 (CH aliphatic), 2939, 2835 (CH aliphatic), 1674 (br., C=O); ¹H-NMR (DMSO-d₆): δ 2.12 (s, 3H, CH₃), 2.28–2.45 (m, 8H, 4CH₂), 2.62 (t, 2H, 2CH₂, J = 6.3), 3.72 (t, 2H, CH₂, J = 7.8), 3.90 (s, 3H, OCH₃), 3.92 (s, 3H, OCH₃), 7.19–8.24 (m, 8H, Ar-H), 10.84 (s, 1H, NH); ¹³C-NMR (DMSO): 34.4, 44.5, 51.1, 52.8, 54.3, 55.3, 103.9, 106.2, 111.7, 127.1, 128.3, 128.8, 129.5, 135.1, 143.5, 144.4, 152.7, 153.4, 167.0, 170.4; MS,
m/z (%): M+, 490 (0.11%), 70 (100%). Anal. Calcd. for C_{26}H_{30}N_{6}O_{4} (490.55): C, 63.66; H, 6.16; N, 17.13. Found: C, 64.13; H, 6.28; N, 17.46%.

3.2.6. General Procedure for the Preparation of 7a–d

A mixture of equimolar amounts of compound 4 and the corresponding 2/3-chloro-1-(piperidin-1-yl/4-methylpiperazin-1-yl) ethanone/propan-1-one (2 mmol) in dry DMF (10 mL) containing triethylamine (0.5 mL) was refluxed for 10 h. The resulting solution was cooled and poured onto crushed ice. The separated solid was filtered, dried, and crystallized from methanol.

7,8-Dimethoxy-4-(2-oxo-2-piperidin-1-yl)ethylamino)-1-phenylimidazo[1,5-a] quinazolin-5(4H)-one (7a).
Yield: 77%; mp 142–143 °C; IR (KBr, cm⁻¹): 3232 (NH), 3056 (CH aromatic), 2923, 2854 (CH aliphatic), 1670, 1654 (C=O); ¹H-NMR (DMSO-d₆): δ 1.27–1.38 (m, 2H, CH₂), 1.42–1.56 (m, 4H, 2CH₂), 3.11–3.28 (m, 4H, 2CH₂), 3.73 (s, 2H, CH₂), 3.89 (s, 3H, OCH₃), 3.91 (s, 3H, OCH₃), 6.45 (s, 1H, NH), 7.17–8.87 (m, 8H, Ar-H); ¹³C-NMR (DMSO): 21.3, 21.8, 45.2, 51.0, 55.1, 55.7, 98.9, 99.6, 112.0, 127.2, 128.1, 128.2, 128.8, 131.5, 139.2, 147.7, 153.4, 154.6, 167.4; MS, m/z (%): M⁺, 461 (0.51%), 196 (100%). Anal. Calcd. for C_{25}H_{27}N_{5}O_{4} (461.51): C, 65.06; H, 5.90; N, 15.17. Found: C, 65.11; H, 5.97; N, 15.24%.

7,8-Dimethoxy-4-(2-(4-methylpiperazin-1-yl)-2-oxo-ethylamino)-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one (7b).
Yield: 67%; mp 160–161 °C; IR (KBr, cm⁻¹): 3305 (NH), 3059 (CH aromatic), 2924, 2854 (CH aliphatic), 1670, 1661 (C=O); ¹H-NMR (DMSO-d₆): δ 2.22–2.34 (m, 7H, CH₃ and 2CH₂), 3.16 (t, 4H, 2CH₂, J= 6.3), 3.507 (s, 2H, CH₂), 3.89 (s, 3H, OCH₃), 3.90 (s, 3H, OCH₃), 6.44 (s, 1H, NH), 7.09–8.09 (m, 8H, Ar-H); ¹³C-NMR (DMSO): 44.5, 51.2, 52.1, 53.5, 54.3, 55.3, 103.9, 106.4, 111.8, 127.2, 128.3, 128.8, 129.3, 130.1, 135.2, 143.6, 144.7, 152.7, 153.4, 167.0, 170.4; MS, m/z (%): M⁺, 476 (0.22%), 105 (100%). Anal. Calcd. for C_{25}H_{28}N_{6}O_{4} (476.52): C, 63.01; H, 5.92; N, 17.64. Found: C, 62.98; H, 6.23; N, 17.81%.

7,8-Dimethoxy-4-(3-oxo-2-piperidin-1-yl)propylamino)-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one (7c).
Yield: 69%; mp 98–100 °C; IR (KBr, cm⁻¹): 3255 (NH), 3028 (CH aromatic), 2927, 2854 (CH aliphatic), 1672, 1654 (C=O); ¹H-NMR (DMSO-d₆): δ 1.19–1.27 (m, 2H, CH₂), 1.44–1.56 (m, 4H, 2CH₂), 2.72 (t, 2H, CH₂, J= 11.1), 2.97 (t, 2H, CH₂, J= 11.4), 3.43–3.69 (t, 4H, 2CH₂, J= 6.6), 3.81 (s, 3H, OCH₃), 3.91 (s, 3H, OCH₃), 4.64 (s, 1H, NH), 7.13–8.08 (m, 8H, Ar-H); ¹³C-NMR (DMSO): 23.8, 24.6, 25.9, 44.5, 51.0, 55.1, 55.7, 98.9, 99.6, 112.0, 127.1, 128.1, 128.2, 129.0, 133.2, 139.2, 147.7, 153.4, 154.6, 162.8, 167.4; MS, m/z (%): M⁺, 475 (0.52%), 207 (100%). Anal. Calcd. for C_{26}H_{29}N_{5}O_{4} (475.53): C, 65.67; H, 6.15; N, 14.73. Found: C, 65.78; H, 6.21; N, 14.92%.

7,8-Dimethoxy-4-(3-(4-methylpiperazin-1-yl)-3-oxo-propylamino)-1-phenylimidazo[1,5-a]quinazolin-5(4H)-one (7d).
Yield: 69% mp 210–211 °C; IR (KBr, cm⁻¹): 3217 (NH), 3056 (CH aromatic), 2924, 2850 (CH aliphatic), 1674, 1658 (C=O); ¹H-NMR (DMSO-d₆): δ 2.12–2.43 (m, 7H, CH₃ and 2CH₂), 2.72 (t, 2H, CH₂, J= 11.1), 2.88 (t, 2H, CH₂, J= 11.4), 3.05 (t, 4H, 2CH₂, J= 7.2), 3.89 (s, 3H, OCH₃), 3.90 (s, 3H, OCH₃), 6.45 (s, 1H, NH), 7.19–8.14 (m, 8H, Ar-H); ¹³C-NMR (DMSO): 31.1, 36.1, 45.8, 51.4, 55.6, 56.4, 99.5, 100.1, 112.8, 127.8, 127.9, 128.6, 128.9, 129.3, 139.8, 148.7, 155.3,
162.8, 167.8; MS, m/z (%): M⁺, 490 (22.1%), 70 (100%). Anal. Calcd. for C₂₆H₃₀N₆O₄ (490.55): C, 63.66; H, 6.16; N, 17.13. Found: C, 63.81; H, 6.30; N, 17.36%.

3.2.7. Synthesis of 11,12-Dimethoxy-8-phenyl-3,4-dihydro-2H-imidazo[1,5-a][1,2,4]triazino[2,3-c]quinazolin-2-one (8)

A mixture of 4 (10 mmol, 3.36 g) and 2-chloracetamide (12 mmol, 1.12 g) in dry DMF (15 mL) was refluxed for 24 h. The desired product was obtained through pouring the reaction solution onto ice water and the precipitate was collected, washed, dried, and then crystallized from DMF/water. Yield: 61%; mp 190–192 °C; IR (KBr, cm⁻¹): 3210 (NH), 3011 (CH aromatic), 2958, 2924 (CH aliphatic), 1666 (C=O); ¹H-NMR (DMSO-d₆): δ 3.74 (s, 2H, CH₂), 3.89 (s, 3H, OCH₃), 3.93 (s, 3H, OCH₃), 5.04 (s, 1H, NH), 7.29–8.31 (m, 8H, Ar-H); ¹³C-NMR (DMSO): 51.5, 55.6, 56.4, 101.4, 112.8, 126.1, 127.9, 128.1, 128.6, 129.2, 129.5, 131.9, 134.2, 138.4, 146.8, 155.3, 168.0; MS, m/z (%): M⁺–1, 374 (8.6%), 50 (100%). Anal. Calcd. for C₂₀H₁₇N₅O₃ (375.38): C, 63.99; H, 4.56; N, 18.66. Found: C, 63.81; H, 4.78; N, 18.24%.

3.2.8. Synthesis of 1-(7,8-Dimethoxy-5-oxo-1-phenylimidazo[1,5-a]quinoxalin-4(5H)-yl)-1H-imidazole-2,5-dione (9)

A suspension of 5a (5 mmol, 2.06 g) and potassium cyanate (10 mmol, 0.81 g) in acetic acid (10 mL) was refluxed for 12 h. The mixture was concentrated under vacuum; water was added to the residue, filtered, dried, and crystallized from ethanol. Yield: 82%; mp 202–203 °C; IR (KBr, cm⁻¹): 3030 (CH aromatic), 2985, 2927 (CH aliphatic), 1670 (C=O); ¹H-NMR (DMSO-d₆): δ 3.90 (s, 3H, OCH₃), 3.92 (s, 3H, OCH₃), 7.50–8.35 (m, 9H, Ar-H); ¹³C-NMR (DMSO): 52.3, 55.5, 103.5, 107.8, 112.0, 127.9, 128.1, 128.3, 129.5, 135.0, 144.1, 148.7, 153.1, 154.7, 164.9, 167.0, 167.1; MS, m/z (%): M⁺, 417 (3.05%), 55 (100%). Anal. Calcd. for C₂₁H₁₅N₅O₅ (417.37): C, 60.43; H, 3.62; N, 16.78. Found: C, 60.21; H, 3.71; N, 16.47%.

3.3. In Vitro Antitumor Activity Measurement against Human Mammary Carcinoma Cell Line (MCF7)

Cells were plated in 96-multiwell plate (104 cells/well) for 24 h before treatment with the compounds to allow attachment of cell to the wall of the plate. Different concentrations of the compound under test (0, 1, 2.5, 5 and 10 µg/mL) were added to the cell monolayer. Triplicate wells were prepared for each individual dose. Monolayer cells were then incubated with the compounds for 48 h at 37 °C and in atmosphere of 5% CO₂. After this time, cells were fixed, washed, and stained with Sulforhodamine B stain. Excess stain was washed with acetic acid and attached stain was recovered with Tris EDTA buffer. The color intensity was measured in an ELISA reader. Finally, the relation between surviving fraction and drug concentration was plotted to get the survival curve of the tumor cell line after the specific compound [30]. The optical density measured is linear to the cell number of the surviving fraction. Therefore, the assay is a sensitive measure of compound induced cytotoxicity with the best signal to noise ratio. The assay also, provides a colorimetric end point that is nondestructive, indefinitely stable and visible to naked eye.
3.4. 2D-QSAR Study

3.4.1. Data Set

All the molecular modeling calculations and docking simulation studies were performed using Molecular Operating Environment (MOE®) version 10.2010, Chemical Computing Group Inc., Montreal, Canada. The computational software operated under Windows XP installed on an Intel Pentium IV PC with a 1.6 GHz processor and 512 MB memory. All the interaction energies and different calculations were automatically calculated. The biological activity values IC\textsubscript{50} were converted to negative logarithmic (P IC\textsubscript{50}) and used as the dependant variable for the QSAR analysis. Thirteen different molecular descriptors (independent variables) were selected and calculated for the submitted structures aiming cover a wide range of different electronic, hydrophobic and topological characters. The correlation matrix was calculated to avoid multicollinearity between the calculated descriptors. The correlation matrix indicated that some of the descriptors used are highly correlated which suggests avoiding the combinations between such intercorrelated descriptors (|r| ≥ 0.80, where r is the simple linear coefficient). QSAR model was then constructed after ensuring reasonable correlation of antitumor activity with individual descriptor and minimum inter-correlation among the descriptors used in derived model.

3.4.2. Statistical Analysis

Stepwise linear regression analysis (SLRA) technique was used to test the best structural predictors for activity. For the current dataset of the new compounds, the QSAR model development was restricted to a maximum of three variables in accordance to the general accepted rule for the compounds: descriptors ratio to be around 5:1. The developed QSAR models are evaluated using the following statistical measures as root mean square error (RMSE) and correlation coefficient (R\textsuperscript{2}) and validated by the leave-out technique (LOO technique), where each object of the data set is taken away, one at a time. In this case, given n objects, n reduced models are developed. The predicted activities (p IC\textsubscript{50}) for the tested compounds calculated using multi-linear regression (SPred) technique.

4. Conclusions

Various substituted imidazo[1,5-\textalpha]quinazoline derivatives were synthesized. The structure of the newly synthesized compounds was elucidated by elemental analyses and spectral data. All compounds were tested, \textit{in vitro}, for their antitumor activity. Compound 5a is the most active against the human mammary carcinoma cell line (MCF7). Physico-chemical parameters revealed that most of the compounds obeyed the “rule of five” properties with good absorption percentages. 2D-QSAR studies helped to explore some expectedly potent compounds.

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Conflicts of Interest

The authors declare no conflict of interest.

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*Sample Availability:* Samples of the newly synthesized compounds are available from the author.

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