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Exploring the antitumor potential of novel quinoline derivatives *via* tubulin polymerization inhibition in breast cancer; design, synthesis and molecular docking†

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A series of quinoline derivatives was designed and synthesized as novel tubulin inhibitors targeting the colchicine binding site. All the rationalized compounds **3a–e**, **4a–e**, **5a–e**, and **6a–e** have been chosen for screening their cytotoxic activity against 60 cell lines by NCI. Compounds **3b**, **3c**, **4c**, **5c** and **6c** demonstrated the most notable antitumor activity against almost all cell lines. Compound **4c** emerged as the most potent compound as an antiproliferative agent. This compound was subsequently chosen for five-dose testing and it exhibited remarkable broad-spectrum efficacy with strong antitumor activity against several cell lines. Compound **4c** significantly induced cell cycle arrest in MDA-MB-231 cells at G2 and M phases where the cell population increased dramatically to 22.84% compared to the untreated cells at 10.42%. It also increased the population in MDA-MB-231 cells at both early and late stages of apoptosis. Compound **4c** can successfully inhibit tubulin polymerization with an IC₅₀ value of 17 ± 0.3 μM. The β-tubulin mRNA levels were notably reduced in MDA-MB-231 cells treated with compound **4c** which is similar to the effect observed with colchicine treatment. Docking studies revealed that compound **4c** interacted well with crucial amino acids in the active site.

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1. Introduction

Cancer is the development of abnormal cells that proliferate uncontrollably and affect human health and it is one of the leading causes of death globally.^{1–3} The latest data indicate that breast cancer is one of the most prevalently diagnosed cancers among women with an estimated 2.3 million new cases (11.7%) and the 5th cause of cancer-related deaths with an estimated 6.9%.^{1–3} Currently, chemotherapy is the main approach for cancer treatment; since microtubules are essential for cell viability, especially for the fast division of cancer cells, drugs that interfere with the dynamics of microtubule/tubulin have become essential therapeutics.^{4,5} Most of these agents act by

binding to the tubulin, an α/β heterodimer protein that forms the microtubule which is a major component of the eukaryotic cytoskeleton.⁶

Microtubule targeting agents (MTA) are also named antimetabolic agents which bind to the tubulin in the microtubules and prohibit the proliferation of the cells. There are two distinct categories of microtubule targeting agents; those that bind to the binding site of paclitaxel, such as paclitaxel which are identified as microtubule stabilizing agents or tubulin promoters.⁷ Conversely, agents that bind to the binding site of colchicine, such as colchicine, combretastatin A-4 (CA-4), and podophyllotoxin or to the binding site of vinca alkaloid, such as vincristine, are identified as tubulin inhibitors or destabilizing agents (Fig. 1).⁸

The colchicine binding site (CBS), which sits at the interface between α and β-tubulin heterodimers, has been the subject of much research to identify potential anticancer drugs. Colchicine was the first tubulin destabilizing agent. It was extracted from the poisonous meadow saffron *Colchicum autumnale* and it binds with high affinity to β-tubulin subunit, interfering with microtubule polymerization and causing mitosis.^{5,9,10}

Pharmacophoric patterns identified in studies of colchicine binding site inhibitors (CBSIs) consistently displayed specific structural characteristics and recurring interactions between tubulin and ligands at pharmacophoric points. These studies

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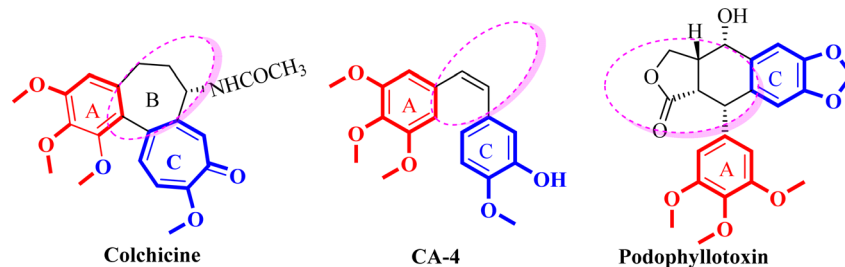


Fig. 1 Structure of tubulin destabilizing agents.

proposed that the different structural classes of CBSIs can be linked by a six to seven-point pharmacophore consisting of three hydrogen bond acceptors (HBA), one hydrogen bond donor (HBD), one or two hydrophobic center (HY), and/or one planar group.¹¹ Ten top-scored hypothetical pharmacophores were generated in another study using the HypoGen algorithm, the training set consisted of 26 drugs with tubulin inhibitory activity ranging from 0.52 to 13.800 nM. One hydrogen-bond acceptor (HBA), one hydrogen-bond donor (HBD), one ring aromatic feature (RA), one hydrophobic feature (HY), and three excluded volumes (EV) made up the greatest hypothetical pharmacophore.¹² Another training set for the 3D QSAR pharmacophore model used the structures of 21 different drugs defined four different feature types: the hydrophobic feature (HY), hydrophobic aromatic group (HY-AR), hydrogen bond acceptor (HBA), and hydrogen bond donor (HBD) (Fig. 2).^{13–15} The previously mentioned features are necessary to ensure stable and effective binding to the CBS.¹⁶ The presence of many hydrophobic and aromatic groups ensures a fit to the hydrophobic pocket of the β -tubulin subunit, while the HBA and HBD groups extend the inhibitors towards the hydrophilic α -tubulin subunit.

SAR analysis of **colchicine** revealed that the important trimethoxyphenyl group is oriented within β -tubulin. The

interaction between ring A and CBS provides the strength of **colchicine** binding to tubulin. On the other hand, interactions between the CBS and the oxygen atoms on ring C control the inhibition. It is suggested that within the binding locus, ring A anchors and keeps the B and C rings oriented correctly.^{5,9,10}

Moreover, **colchicine** showed its anticancer effect on apoptotic genes of human breast cancer cell lines.^{10,17,18} **CA-4** binds at the CBS in a manner akin to that of **colchicine**.¹⁰ Nevertheless, the instability of the *cis*-double bond, which might change into an inactive *trans*-conformation, has hindered the development of **CA-4**.¹⁹ In recent years, various conformationally restricted analogs of **CA-4** and colchicine-bearing triazolopyrimidine **I**, benzimidazole **II**, azetidine **III**, pyridine **IV** and **VI**, pyrazole **V**, pyrimidine **VII**, imidazopyridine **VIII** have been reported as potent tubulin inhibitors (Fig. 3).^{20–26}

Quinoline is a privileged scaffold in the expansion of anti-cancer drugs as they have exhibited potent antiproliferative activity *via* different mechanisms of action including cell cycle arrest, induction of apoptosis, and inhibition of angiogenesis, and cell migration disruption. Many tubulin inhibitors **IX–XIV** also, possess quinoline motifs (Fig. 4), these compounds exhibited their antiproliferative activity because of tubulin polymerization inhibition and the disruption of microtubule assembly.^{27–32}

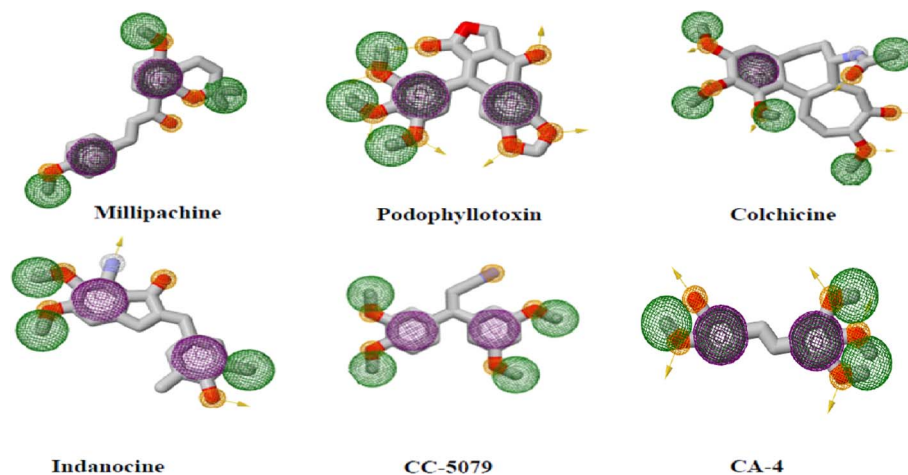


Fig. 2 Two-dimensional structures of reported CBSI's with pharmacophoric features necessary to impart depolymerization upon tubulin where purple spheres represent aromatic pharmacophoric points (AR), the orange spheres with and without arrow denote the functional groups that act as hydrogen bond donor (HBD) and acceptor (HBA), respectively. Finally green spheres represent hydrophobic centers (HY).

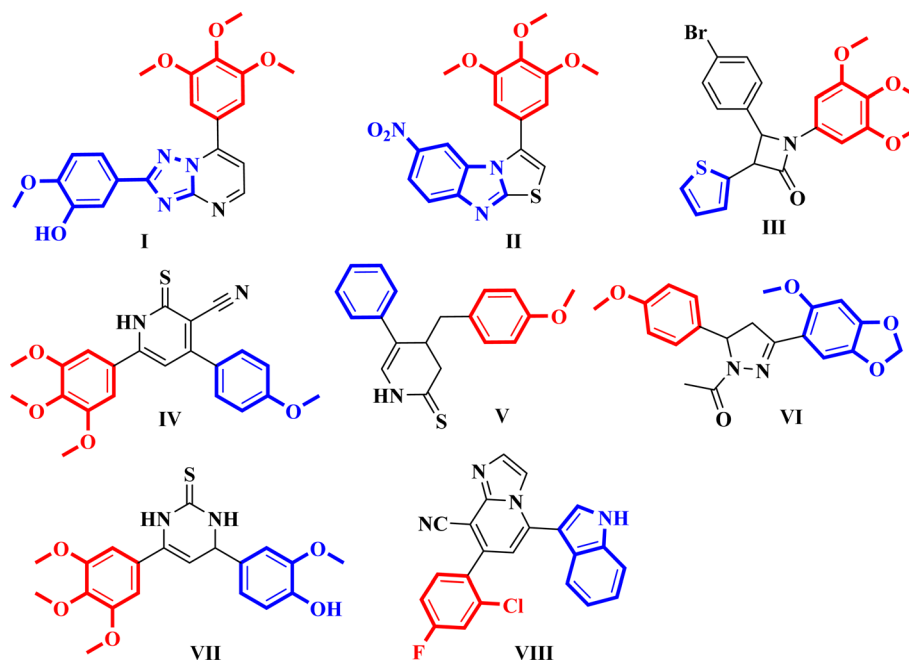


Fig. 3 Chemical structure of reported CA-4 analogs with cytotoxic activity.

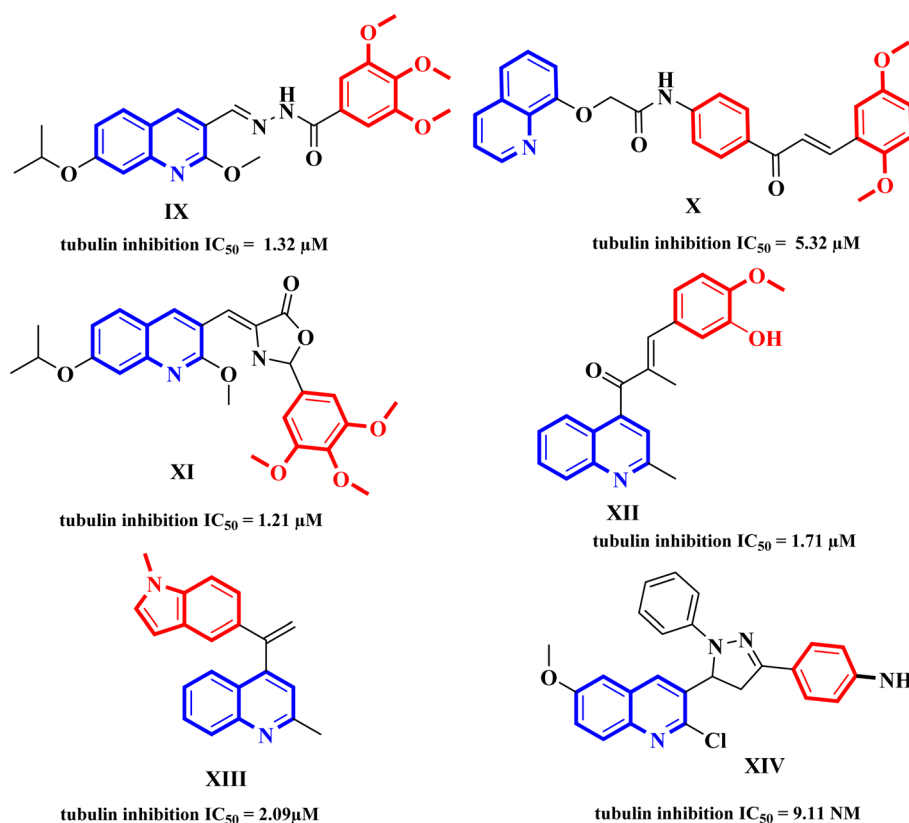


Fig. 4 Some quinoline-containing antimitotic agents and tubulin polymerization inhibitors.

In light of the aforementioned findings, we aimed to synthesize novel molecules with promising tubulin polymerization inhibition activity *via* structure optimization of lead compounds CA-4 and **colchicine**. As shown in our design

strategy (Fig. 5), bioisosteric replacement of ring C in **colchicine** and CA-4 was performed with a quinoline ring. 3,4,5-Trimethoxyphenyl moiety (ring A) in **colchicine** and CA-4 was retained in certain synthesized derivatives, while others were



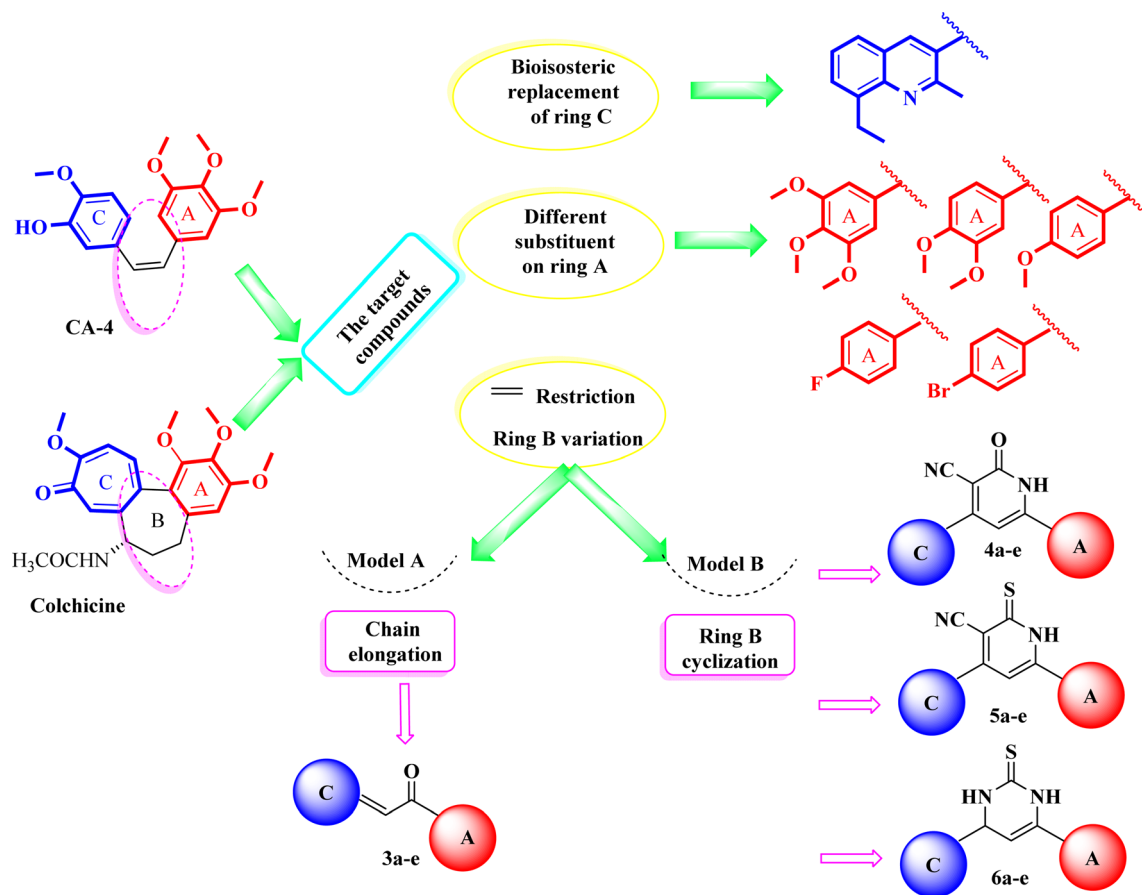


Fig. 5 Design strategy of the rationalized compounds 3a–e, 4a–e, 5a–e, and 6a–e.

embellished with 3,4-dimethoxyphenyl, 4-methoxyphenyl, 4-fluorophenyl or 4-bromophenyl moieties. Chain elongation in compounds 3a–e by the introduction of 2-propen-1-one instead of an olefinic bond aimed to increase the rigidity of the structure to overcome the drawbacks of CA-4. For more rigidification, cyclization with rigid core ring B was performed by 2-pyridinone 4a–e, pyridinethione 5a–e and pyrimidinethione ring 6a–e. Our designed plan was achieved according to Scheme 1 (Fig. 6). The novel compounds were also fully characterized for their cytotoxic activity on NCI 60-cell lines and additional mechanistic biochemical tests were conducted for the molecule with the highest potency. Moreover, docking studies were conducted in an attempt to interpret the outcomes of biological tests.

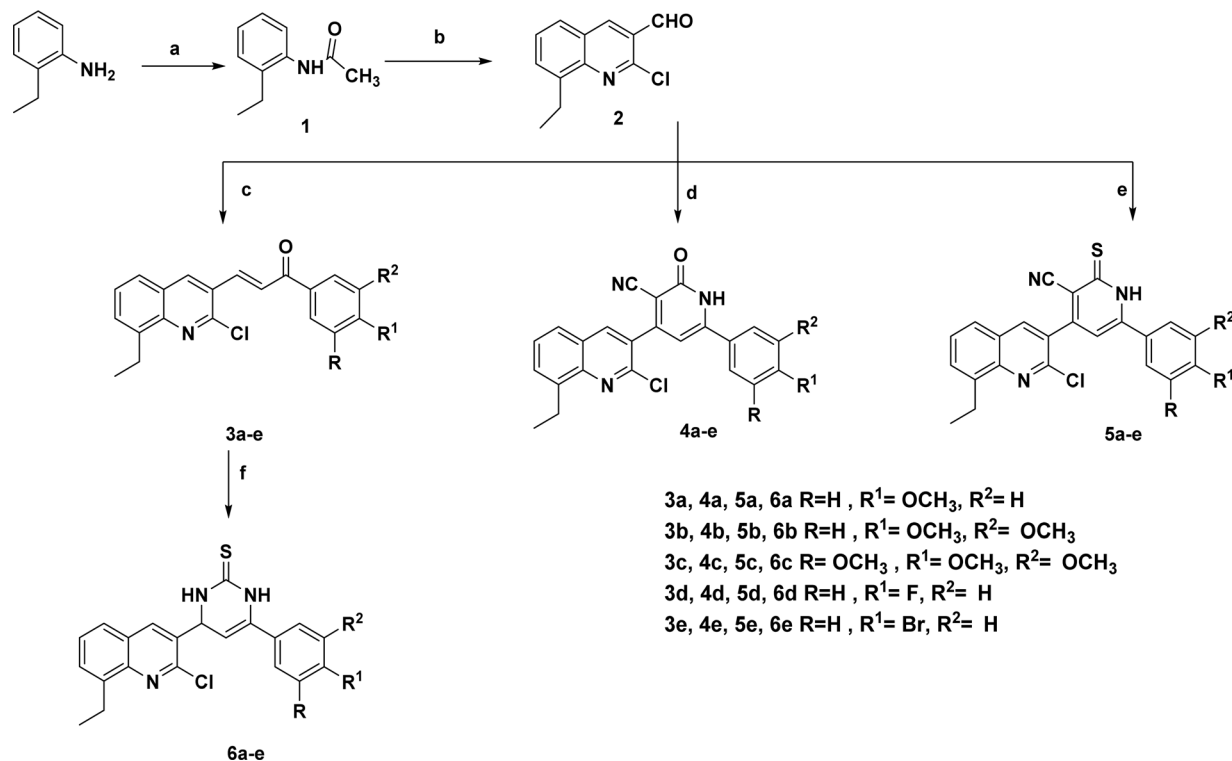
2. Results and discussion

2.1. Chemistry

The present study described the synthetic plan of novel compounds 3a–e, 4a–e, 5a–e, and 6a–e as illustrated in Scheme 1. Compound 1 and the key intermediate 2 were prepared in excellent yields as described in the reported literature.^{33–36} Chalcone derivatives 3a–e were prepared following the Claisen–Schmidt condensation reaction by reacting a mixture of the carbalddehyde derivative 2 with the appropriate substituted acetophenone in the presence of 50% aqueous potassium

hydroxide. The IR spectrum of each synthesized compound exhibited a strong absorption band corresponding to the carbonyl (C=O) functional group, which was observed in the range of 1600–1656 cm^{-1} . ^1H NMR charts showed signals corresponding to α , β alkene protons, as a doublet signal at δ 8.12–8.20 ppm referred to the α -CH alkene proton while another doublet appeared at δ 7.64–8.11 ppm corresponding to β -CH alkene proton. On the other hand, their ^{13}C NMR spectra showed a peak referring to the carbonyl (C=O) functional group at δ 186.8–188.0 ppm. Refluxing the 8-ethylquinoline-3-carbaldehyde derivative 2 with different acetophenones and either ethyl cyanoacetate or 2-cyanothioacetamide in the presence of ammonium acetate afforded the target pyridine derivatives 4a–e and 5a–e, respectively. The IR spectra of the carbonitrile derivatives 4a–e and 5a–e showed a peak at 2218–2239 cm^{-1} of the added CN group and a band of NH appeared at 3398–3433 cm^{-1} . ^1H NMR spectra of 2-oxo-1,2-dihydropyridine derivatives 4a–e showed a singlet at δ 6.85–7.16 ppm corresponding to the proton of 1,2-dihydropyridine ring, while ^{13}C NMR spectra revealed signals at δ 116.4–118.2 and 156.0–159.6 ppm for nitrile and carbonyl groups, respectively. ^1H NMR spectra of the isosteric analog; 2-thioxo-1,2-dihydropyridine derivatives 5a–e showed similar singlet at δ 6.78–7.17 ppm corresponding to the proton of 1,2-dihydropyridine ring, although, their ^{13}C NMR spectra revealed signals at δ 116.0–





Scheme 1 Synthesis of the target compounds 3a–e, 4a–e, 5a–e, and 6a–e.

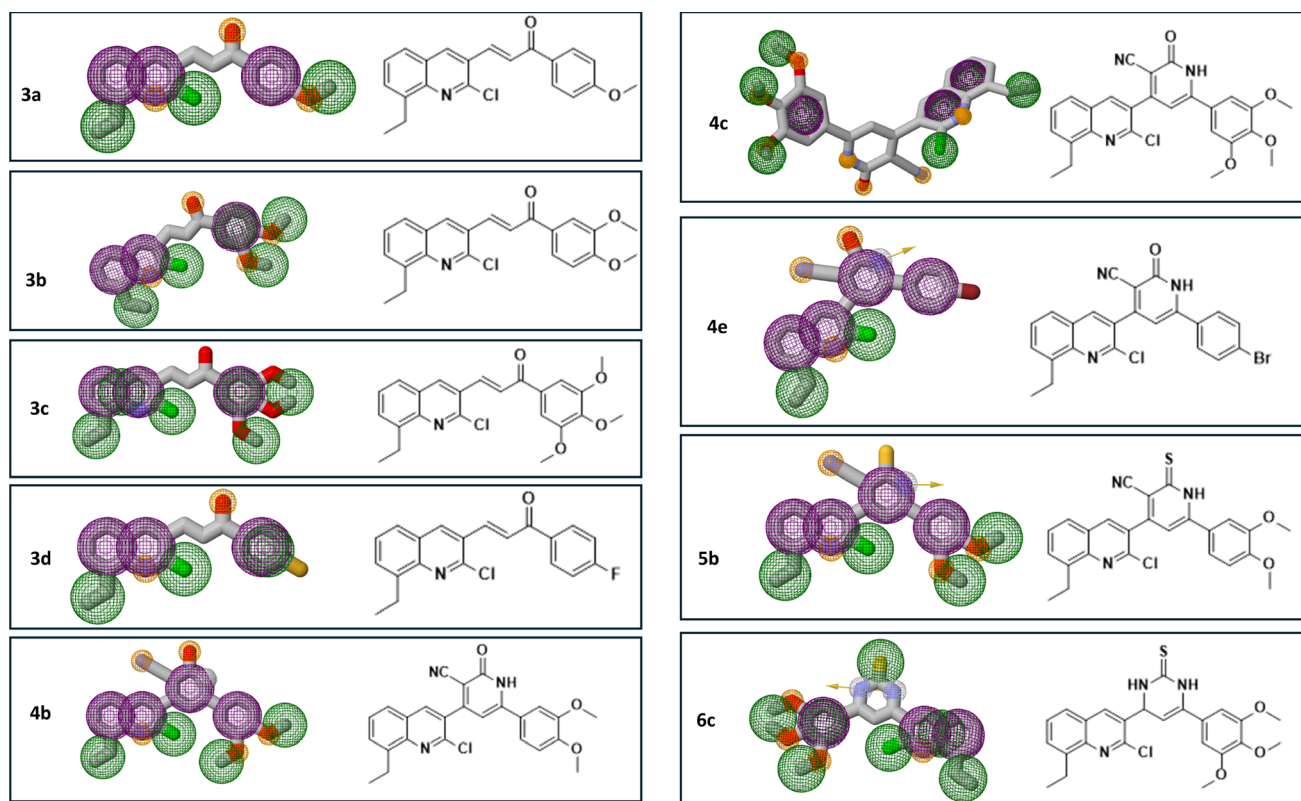


Fig. 6 Some of the designed titled compounds with common structural features to reported tubulin depolymerization agents.



117.0 and 156.5–162.1 ppm for nitrile and thiocarbonyl groups, respectively. Additionally, mass spectrometric analysis was conducted to further confirm the structure of the synthesized 1,2-dihydropyridine derivative compounds, designated as **4a–e** and **5a–e**. The mass spectrometric data showed the presence of molecular ion peaks that corresponded to the expected molecular weights of the respective synthesized compounds. 3,4-Dihydropyrimidinethione derivatives **6a–e** were synthesized by refluxing the corresponding chalcone **3a–e** with thiourea in an alkaline medium. The afforded compounds were prepared with an overall good yield of 70–74%. The IR spectra confirmed that the carbonyl functional group present in the starting materials **3a–e** was no longer detected in the final synthesized compounds but an absorption band appeared at 3162–3169 cm^{-1} corresponding to NH. ^1H NMR of **6a–e** showed two doublets at δ 5.39–5.61 ppm which is characteristic to the dihydropyrimidine ring protons along with the exchangeable protons of 2NH that appeared at δ 9.10–9.10 and 10.04–10.20 ppm. Additionally, ^{13}C NMR showed a peak at δ 96.5–99.7 ppm pointing to C5 of the pyrimidine ring and a peak at δ 174.0–177.0 ppm referring to C=S. Finally, it is worth mentioning that all our novel compounds were subjected to elemental analyses for further authentication of the synthesized compounds.

2.2. Biology

2.2.1. In vitro cytotoxic activity screening. In order to determine the cytotoxic activity of 20 recently synthesized quinoline derivatives, their activity against a comprehensive panel of 60 cancer cell lines, including those representing leukemia, melanoma, lung, colon, central nervous system (CNS), ovary, kidney, prostate, and breast cancers, was screened at the National Cancer Institute (NCI) in Bethesda, MD, USA. This screening involved administering a single dose of 10 μM and the data were reported as percent growth ($G\%$) of the treated cell lines (Fig. S68–S87[†]) and then expressed as percentage growth inhibition (GI%) (Tables S1 and S2[†]). The heatmap shown in Fig. 7 illustrates the anti-proliferative activity (cells viability percentage) of all quinoline derivatives tested against all 60 cancer cell lines. Based on the findings, the

majority of the newly created compounds exhibited cytotoxic effects ranging from moderate to high, as indicated by their growth inhibition percentages (GI%). Some of the compounds demonstrated only mild cytotoxic activity. Chalcone derivative **3a** (methoxy-substituted) showed selective remarkable cytotoxic activity on non-small cell lung cancer (NCI-H226) with GI value of 79.88%. Chalcone derivative **3b** (dimethoxy-substituted) demonstrated potent cytotoxic activity against leukemia (SR), CNS cancer (SF-539) with GI value of 80.08 and 99.38%, respectively. Additionally, it showed lethal effect on non-small cell lung cancer (NCI-H226) and CNS cancer (SNB-75). Chalcone derivative **3c** (trimethoxy-substituted) displayed potent antitumor activity against leukemia (CCRF-CEM) with GI value of 87.78%, non-small cell lung cancer (NCI-H522) with GI values of 80.68%, colon cancer (KM12 and HT29) with GI value of 80.66 and 77.4%, respectively and ovarian cancer (OVCAR-8) with GI value of 84%. Other chalcones **3d** (fluoro-substituted) and **3e** (bromo-substituted) exhibited low cytotoxic activity across all tested cell lines.

Pyridin-2-one **4c** (trimethoxy-substituted) demonstrated the highest level of cytotoxic activity among the synthesized compounds. It possessed potent cytotoxic activity against wide range of tumor cells as leukemia (CCRF-CEM, MOLT-4, RPMI-8226, and SR) with GI values of 80.39, 76.43, 74.51 and 77.32%, respectively, non-small cell lung cancer (A549/ATCC and HOP-62) with GI values of 88.16 and 87.86%, respectively, CNS cancer (SF-295 and SNB-19) with GI values of 96.38 and 89.58%, respectively, melanoma (LOX IMVI, MALME-3M and SK-MEL-5) with GI values of 91.58, 89.41, 74.51 and 78.58%, respectively, ovarian cancer (OVCAR-8) with GI value of 92.11%, renal cancer (ACHN and CAKI-1) with GI values of 75.69 and 95.06%, respectively, prostate cancer (DU-145) with GI value of 80.55% and breast cancer (T-47D) with GI value of 91.56%. Its lethal effect was evident across a wide range of tumor cells including non-small cell lung cancer (HOP-92, NCI-H226 and NCI-H522), CNS cancer (SF-539, SNB-75 and U251), melanoma (UACC-62), ovarian cancer (OVCAR-3 and OVCAR-4), renal cancer (786-0, A498, RXF 393, SN12C, TK-10 and UO-31) and breast cancer (MDA-MB-231/ATCC and HS 578T). Other

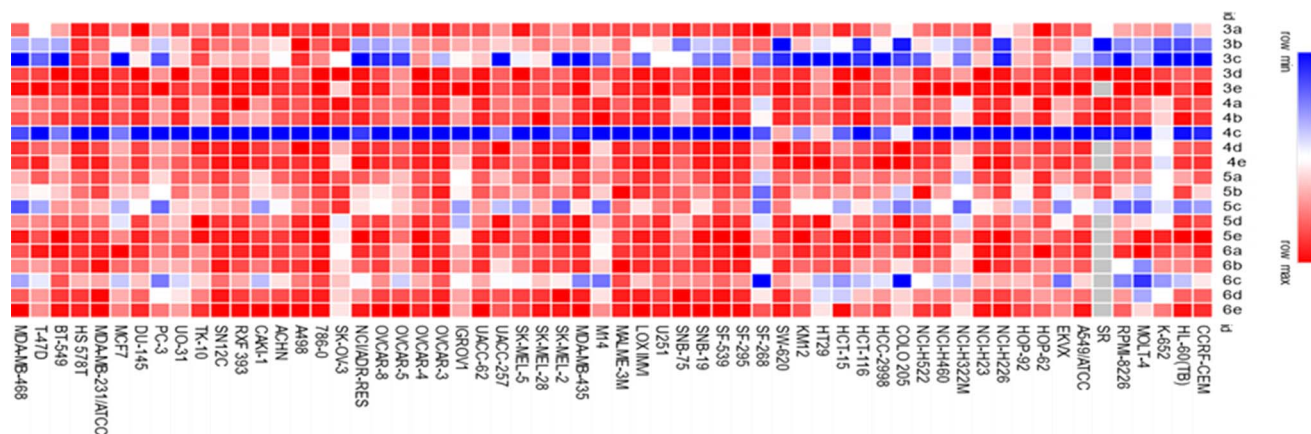


Fig. 7 Heatmap of all the synthesized quinoline derivatives **3a–e**, **4a–e**, **5a–e** and **6a–e** demonstrating their effect on tumor cells viability of 60 different cancer cell lines. Red color indicates higher tumor cells viability, while blue color indicates less tumor cells viability.



derivatives in the series including pyridin-2-one derivative **4a** (methoxy-substituted) showed moderate activity over CNS cancer (SNB-75) with GI value of 54.69%, where, compound **4b** (dimethoxy-substituted), **4d** (fluoro-substituted) and **4e** (bromo-substituted) possessed mild cytotoxic activity over multiple cell lines.

Pyridinethione derivative **5c** displayed significant cytotoxic activity on non-small cell lung cancer (HOP-92) with GI value of 77.38%, while, its moderate cytotoxic activity appeared over numerous cell lines as leukemia (CCRF-CEM, HL-60(TB), MOLT-4 and RPMI-8226) with GI values of 57.63, 51.76, 61.16 and 63.06%, respectively, non-small cell lung cancer (NCI-H522) with GI value of 65.68%, renal cancer (CAKI-1) with GI value of 69.81%, and breast cancer (MDA-MB-231/ATCC, T-47D and MDA-MB-468) with GI values of 52.51, 60.59 and 50.55%, respectively. Additional derivatives bearing pyridinethione namely, **5a** (methoxy-substituted), **5b** (dimethoxy-substituted),

5d (fluoro-substituted) and **5e** (bromo-substituted) showed a slight cytotoxic activity over multiple cell lines.

Finally, pyrimidine derivative **6c** (trimethoxy-substituted) possessed moderate cytotoxic effect over leukemia (MOLT-4 and RPMI-8226) with GI values of 67.65 and 57.59%, respectively, non-small cell lung cancer (NCI-H226 and NCI-H522) with GI values of 57.97 and 50.3%, respectively and renal cancer (CAKI-1 and UO-31) with GI values of 64.11 and 67.70%, respectively. Pyrimidine derivative **6b** (dimethoxy-substituted) possessed moderate cytotoxic effect over leukemia (MOLT-4) with GI value of 53.3%. Other derivatives bearing pyrimidine moiety specially, **6a** (methoxy-substituted), **6d** (fluoro-substituted) and **6e** (bromo-substituted) showed limited cytotoxic activity over multiple cell lines.

In summary, upon comparing all the newly synthesized compounds (Fig. 7), clearly shows that **4c** > **3c** > **3b** > **5c** > **6c** derivatives demonstrated the highest antitumor activity against

Table 1 Median growth inhibitory (GI₅₀, μM), total growth inhibitory (TGI, μM) and median lethal (LC₅₀) concentrations of compound **4c**

Subpanel tumor cell lines	Activity			Subpanel tumor cell lines	Activity		
	GI ₅₀	TGI	LC ₅₀		GI ₅₀	TGI	LC ₅₀
Leukemia				Melanoma			
CCRF-CEM	12.00	>100	>100	M14	29.70	>100	>100
HL-60(TB)	13.40	>100	>100	MDA-MB-435	32.60	>100	>100
K-562	7.72	43.80	>100	SK-MEL-2	39.80	>100	>100
MOLT-4	8.17	79.90	>100	SK-MEL-28	21.10	>100	>100
RPMI-8226	5.16	>100	>100	SK-MEL-5	17.50	90.20	>100
SR	5.70	>100	>100	UACC-257	42.60	>100	>100
				UACC-62	17.50	60.10	>100
Non-small cell lung cancer				Ovarian cancer			
A549/ATCC	12.20	48.70	>100	IGROV1	12.20	46.30	>100
EKVX	28.60	>100	>100	OVCAR-3	14.20	29.80	62.30
HOP-62	20.70	78.90	>100	OVCAR-4	17.20	>100	>100
HOP-92	2.37	9.68	>100	OVCAR-5	48.40	>100	>100
NCI-H226	10.90	32.60	97.70	OVCAR-8	18.50	93.60	>100
NCI-H23	3.20	>100	>100	NCI/ADR-RES	44.90	>100	>100
NCI-H322M	10.50	>100	>100	SK-OV-3	26.00	>100	>100
NCI-H460	16.10	41.80	>100				
NCI-H522	19.10	>100	>100	Renal cancer			
Colon cancer				786-0	11.70	26.70	61.10
COLO 205	45.60	>100	>100	A498	15.80	45.20	>100
HCC-2998	35.70	>100	>100	ACHN	15.60	31.00	61.80
HCT-116	14.10	33.30	78.40	CAKI-1	10.70	22.90	49.20
HCT-15	27.60	>100	>100	RXF 393	2.21	6.69	35.40
HT29	32.10	>100	>100	SN12C	10.30	25.20	61.80
KM12	33.60	>100	>100	TK-10	17.50	34.50	>100
SW-620	27.50	>100	>100	UO-31	12.60	>100	>100
CNS cancer				Prostate cancer			
SF-268	13.80	>100	>100	PC-3	40.30	>100	>100
SF-295	19.40	>100	>100	DU-145	15.50	38.00	93.50
SF-539	12.00	24.80	51.30				
SNB-19	9.25	50.90	>100	Breast cancer			
SNB-75	2.38	16.90	52.60	MCF7	24.00	>100	>100
U251	10.80	28.40	74.90	MDA-MB-231/ATCC	14.50	31.80	69.80
Melanoma				HS 578T	2.38	7.26	>100
LOX IMVI	13.50	33.20	81.30	BT-549	4.11	28.10	>100
MALME-3M	10.10	40.00	>100	T-47D	12.20	64.10	>100
				MDA-MB-468	12.70	61.60	>100



almost all cell lines. Notably, compound **4c** being the most potent compound as anti-proliferative agent. Interestingly, **4c** derivative along with the other compounds with the highest activity (*i.e.*, **3c**, **5c**, **6c**) belonged to the **c** derivatives series which is characterized by trimethoxy group substitution in the main ring. Hence, the trimethoxy group substitution is shown to be superior to all other modifications in the main ring (methoxy, dimethoxy, fluorine, and bromine). On the other hand, cyclization with pyridine-2-one cyclization is more effective than (open chain > pyridinethione > pyrimidine). This suggests that trimethoxy group and pyridin-2-one are responsible for the compounds' high antitumor activity.

2.2.2. Five doses testing of compound 4c. Since pyridin-2-one derivative **4c** exhibited the highest level of cytotoxic activity against almost all tumor cell lines, it has progressed to the full 5-dose assay for the determination of median growth inhibitory (GI_{50} , μM), total growth inhibitory (TGI, μM) and median lethal (LC_{50}) concentrations (Table 1). The dose response curves of **4c** derivative against all different cancer cell

lines are shown in (Fig. 8). Compound **4c** showed remarkable cytotoxic activity across various cell lines. In particular, it demonstrated notable activity on leukemia (K-562, MOLT-4, RPMI-8226 and SR) with GI_{50} values of 7.72, 8.17, 5.16 and 5.70 μM , respectively, non-small cell lung cancer (HOP-92 and NCI-H23) with GI_{50} values of 2.37 and 3.20 μM , respectively, CNS cancer (SNB-75) with GI_{50} value of 2.38 μM , renal cancer (RXF 393) with GI_{50} value of 2.21 μM and breast cancer (HS 578T, BT-549) with GI_{50} values of 2.38 and 4.11 μM , respectively. Additionally, compound **4c** also disclosed moderate cytotoxic activity across leukemia (CCRF-CEM and HL-60(TB)) with GI_{50} values of, 12.00 and 13.40 μM , respectively, non-small cell lung cancer (A549/ATCC and NCI-H226, NCI-H322M) with GI_{50} values of 12.20, 10.90 and 10.50 μM , respectively, colon cancer (HCT-116) with GI_{50} value of 14.10 μM , CNS cancer (SF-268, SF-539, SNB-19, U251) with GI_{50} values of 13.80, 12.00, 9.25 and 10.80 μM , respectively, ovarian cancer (IGROV1 and OVCAR-3) with GI_{50} values of 12.20 and 14.20 μM , respectively, renal cancer (786-0, CAKI-1, SN12C and UO-31) with GI_{50} values of

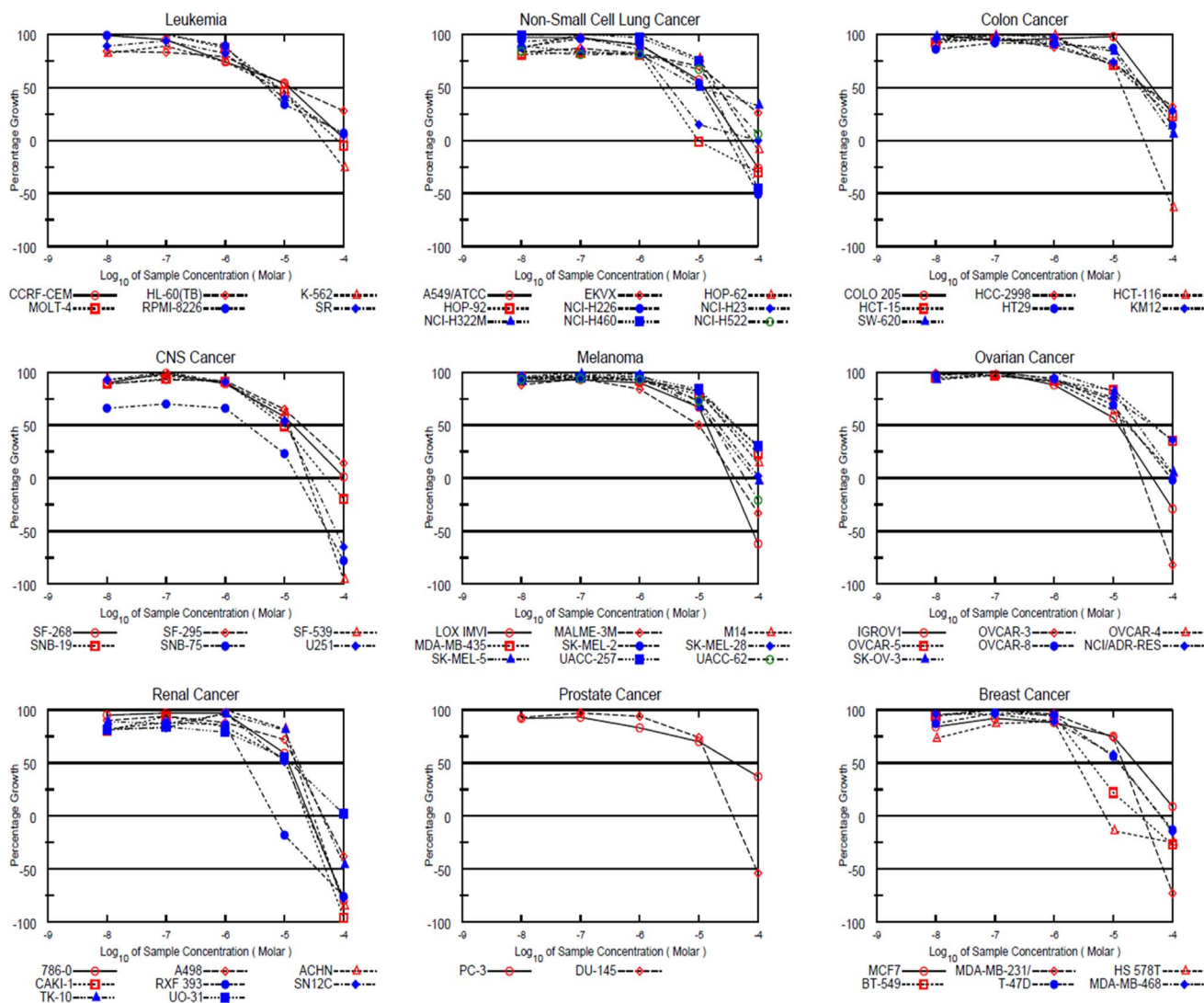


Fig. 8 Dose response curves of **4c** derivative against all 60 cell lines.



11.70, 10.70, 10.30, and 12.60 μM , respectively, and breast cancer (HS 578T) with GI_{50} value of 14.50 μM .

2.2.3. Cell cycle analysis by flow cytometry. Cell cycle is a well-coordinated cellular division process which is composed of several phases. The mitotic phase (M phase) is a central phase which is characterized by the segregation of the duplicate DNA content into two daughter cells.³⁷ Accurate spindle assembly is required for normal cell division and interfering with the formation of microtubules, which is composed of α -tubulin and β -tubulin subunits, hinders cell division leading to cell cycle arrest.³⁸ Therefore, several tubulin polymerization inhibitors have been developed.³⁹ Since we developed our quinoline derivatives to act as tubulin polymerization inhibitors, their ability to induce cell cycle arrest at G2 and M phases was evaluated. Hence, we focused on elucidating the cellular mechanisms underlying its potent cytotoxic activity against MDA-MB-231. Breast cancer cell lines are among the tumor cells against which pyridin-2-one derivative **4c** showed high cytotoxic activity. Breast cancer is one of the most diagnosed cancers among women with 2.26 million new cases in 2020.⁴⁰ It has several subtypes such as luminal breast cancer, Her-2/neu positive type, and triple-negative breast cancer.⁴¹ MDA-MB-231 cell line is one example of the cell lines used to study triple-

negative breast cancer. It is used to identify pathways that regulate metastasis when used for xenografts.⁴² Therefore, MDA-MB-231 cell line is considered a valuable tool for developing new antiproliferative drugs against metastatic breast cancer.⁴³

Hence, we investigated the distribution of cells at the different phases of cell cycle after MDA-MB-231 cells treatment with the newly synthesized **4c** derivative and the positive control **colchicine** which is a well-established tubulin polymerization inhibitor. Fig. 9A shows the histograms of the control non-treated cells in addition to **4c**, and colchicine-treated cells demonstrating propidium iodide PI signals at each phase. Both **4c** and **colchicine** histograms showed an increase in signal intensity at G2/M phase. Moreover, the distribution of cells in each phase of the cell cycle was measured. As illustrated in Fig. 9B, **4c** and **colchicine** significantly induced cell cycle arrest at G2 and M phases where the cells population increased dramatically to 22.84% and 26.51%, respectively compared with control non-treated cells which were 10.42%. There was no significant difference between **4c** and **colchicine** in inducing cell cycle arrest at G2 and M phases. Hence, **4c** derivative exerts its cytotoxic action against breast cancer *via* inhibiting tubulin polymerization causing cell cycle arrest at G2 and M phases.

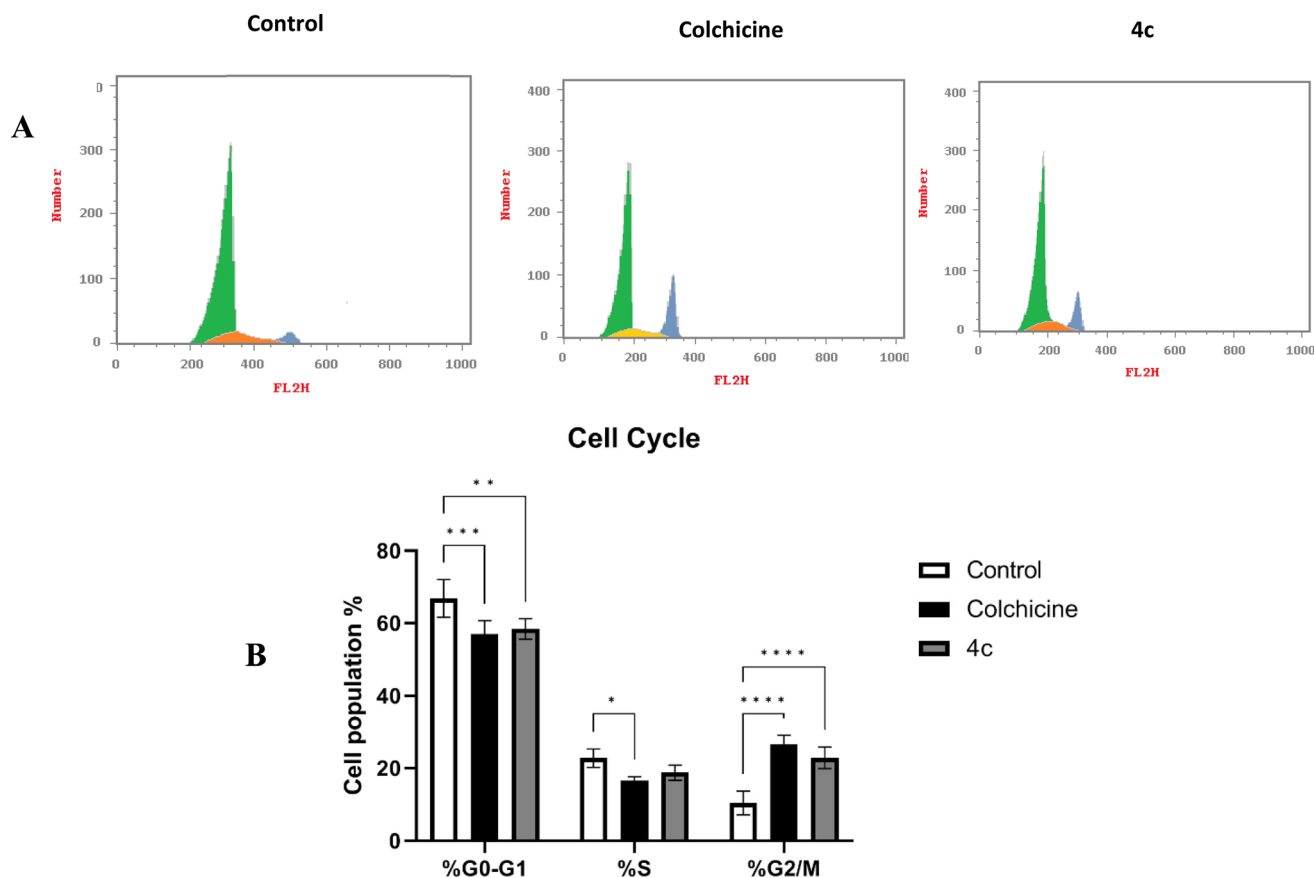


Fig. 9 (A) Histograms of cell cycle phase distribution of control, colchicine, and **4c** treated cells using PI staining for FACS analysis. (B) Percentages of cells accumulation at G0–G1, S, and G2/M cell cycle phases induced by control, colchicine, and **4c**. Data are represented as the mean \pm SD of three independent experiments. Statistical analysis was conducted using two-way ANOVA followed by Tukey's multiple comparison test; * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, **** $p < 0.0001$ compared to the control.



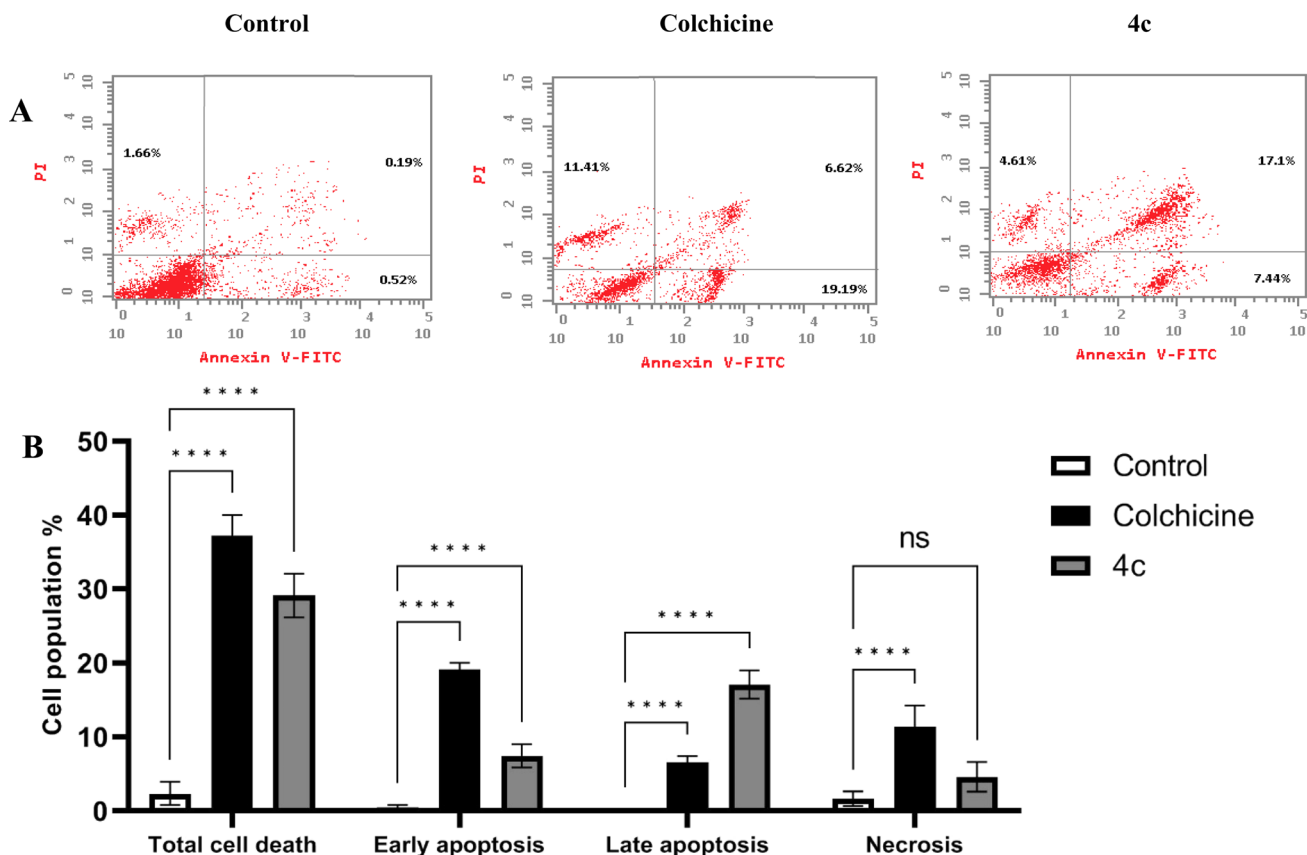


Fig. 10 (A) Dot blots of apoptotic cells populations using PI/annexin V-FITC staining for FACS analysis. (B) Percentages of early apoptosis, late apoptosis and necrosis induced by 4c and colchicine in MDA-MB-231 cells compared with non-treated control cells. Data are represented as mean \pm SD of three individual experiments. Statistical analysis was done by applying either two-way ANOVA followed by with Dunnett's multiple comparison test; * p < 0.05, ** p < 0.01, *** p < 0.001, **** p < 0.0001 compared to the control.

2.2.4. Cell apoptosis analysis. Tubulin polymerization inhibitors can trigger cellular apoptosis due to cell cycle progression hindrance at M phase and the accumulation of mitotic cells.⁴⁴ Hence, we examined the ability of compound 4c to induce cellular apoptosis in MDA-MB-231 cells. Following the reported protocol by staining MDA-MB-231 cells stained with both PI and annexin V-FITC. Fig. 10A shows the dot blots of the control non-treated cells, 4c and colchicine treated cells. The percentages of apoptotic cells significantly increased after treatment with 4c and colchicine treatment compared with non-treated cells. Both 4c and colchicine significantly increased the percentages of cells at both early and late apoptosis compared with the control. Table S3† shows the percentages of the cells at

each apoptotic phase. As displayed in (Fig. 10B) colchicine significantly induced high levels of early apoptosis, while 4c induced higher levels of late apoptosis. This difference indicates that 4c acts with a faster rate in promoting apoptosis, thus reaching the late apoptosis phase earlier compared with colchicine. Additionally, there was no significant difference in

Table 2 IC₅₀ values of 4c, colchicine and CA-4 for tubulin polymerization inhibition

Compound	IC ₅₀ (μM) for tubulin polymerization inhibition
4c	17 \pm 0.3
Colchicine	7.48 \pm 0.11
CA-4	4.647 \pm 0.06

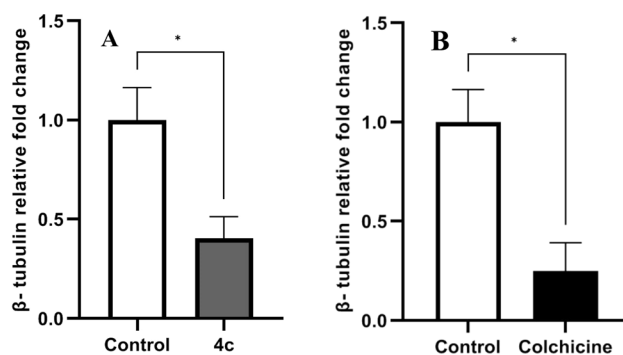


Fig. 11 Relative gene expression of β-tubulin induced by (A) 4c and (B) colchicine compared with the control. Data are represented as mean \pm SD of three individual experiments. Statistical analysis was done by applying Mann-Whitney test; * p < 0.05, ** p < 0.01, *** p < 0.001, **** p < 0.0001 compared to the control.



necrosis levels between **4c** treated and non-treated cells. This indicates the higher safety levels of compound **4c** compared with **colchicine** which significantly increased necrosis levels to 11.41%. This is in alignment with other studies that shows the

ability of tubulin polymerization inhibitors to cause apoptosis.^{45,46}

2.2.5. Tubulin polymerization inhibition assay. To verify whether the newly synthesized compound **4c** can successfully

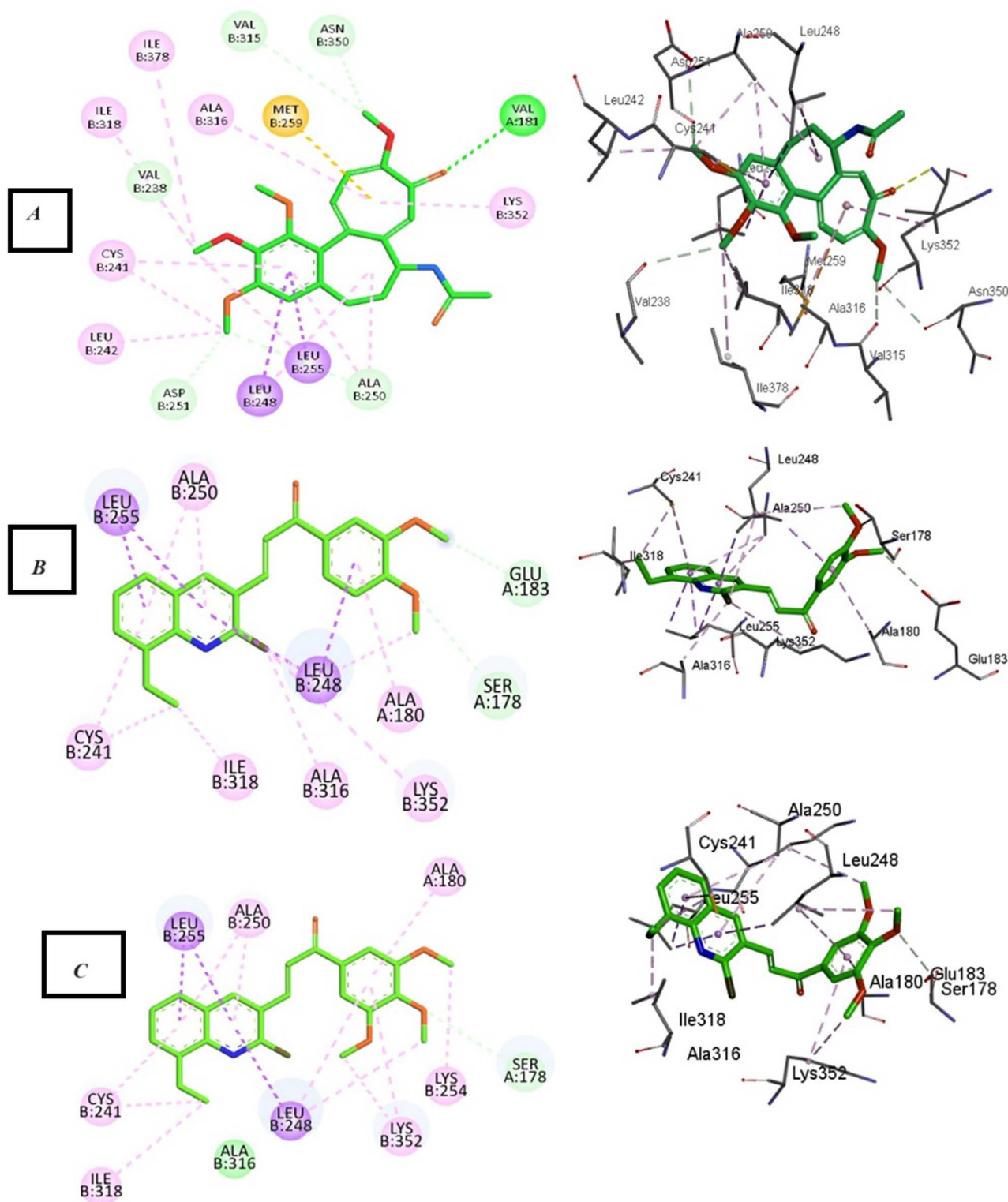


Fig. 12 2D and 3D interaction diagrams of (A) colchicine, (B) compound **3b** and (C) compound **3c** into the colchicine binding site of tubulin enzyme.



inhibit tubulin polymerization, the enzymatic IC_{50} for inhibiting tubulin polymerization was measured. Table 2 shows the IC_{50} for tubulin polymerization inhibition of compound **4c**, **colchicine**, and combretastatin also, graphs of tubulin polymerization inhibition assay data are represented in Fig. S88 and S90.† All compounds were able to inhibit tubulin polymerization with CA-4 being the most active with the lowest IC_{50} . This is in alignment with several studies that showed the ability of different quinoline derivatives to inhibit tubulin polymerization.^{28,47,48}

2.2.6. Tubulin expression assay. α -Tubulin and β -tubulin subunits are the main building blocks of microtubules.⁶ Several tubulin polymerization inhibitors interfere with microtubule assembly.⁴⁴ However, such inhibitors can additionally suppress β -tubulin expression. Thus, we investigated the ability of compound **4c** to decrease β -tubulin mRNA levels. MDA-MB-231 cells were treated with either compound **4c** or **colchicine** and the expression of β -tubulin was determined by real-time PCR. As demonstrated in Fig. 11A and B, both compound **4c** and **colchicine** significantly suppressed β -tubulin transcription compared with non-treated cells.

2.3. In silico studies

2.3.1. Docking studies. Discovery Studio client software was used for the analysis of the possible binding modes between our titled compounds and CBS of tubulin. According to reports, the **colchicine** binding site comprises two hydrophilic

sites partially located at the α -tubulin subunit that are prone to form additional hydrogen bonds with ligands that add stability to the interactions^{49,50} and three hydrophobic pockets located at the β -tubulin subunit that serve as the main locations for ligand interactions. α Thr179, α Val181, and β Leu248 are residues that engage in hydrophilic interactions, whereas the hydrophobic pockets, which are the primary target for compounds in the “deep binding mode” category, include essential amino acids as α Ala180, β Cys241, β Ala250, β Asn258, β Ala316, β Ile318, and β Lys352.⁵⁰

In the current simulation, redocking of **colchicine** was done which revealed the reported binding mode of the compound.^{51,52} After validation of our docking procedure, the titled compounds **3b**, **3c**, **4c**, and **5c** displayed correct binding modes into the CBS. Chalcone derivatives **3b** and **3c** which were lethal against non-small cell lung cancer cells; NCI-H226, CNS cancer cells; SF-539 and SNB-75 and ovarian cancer cells; OVCAR-3, showed a high binding interaction within the hydrophobic pocket of the β -tubulin subunit with the correct pose equal to -9.0 and -8.0 kcal mol⁻¹, respectively. As can be deduced from (Fig. 12) both compounds revealed identical binding modes where they were deeply buried in the **colchicine** site at the α , β intradimer interface and formed mainly hydrophobic contacts with several residues of β -tubulin where the 2-chloro-8-ethylquinoline ring bonded through several π -alkyl and π -sigma interactions to β Cys241, β Leu248, β Ala250, β Leu255, β Ala316, β Ile318 and β Lys352 (Table S4†). The

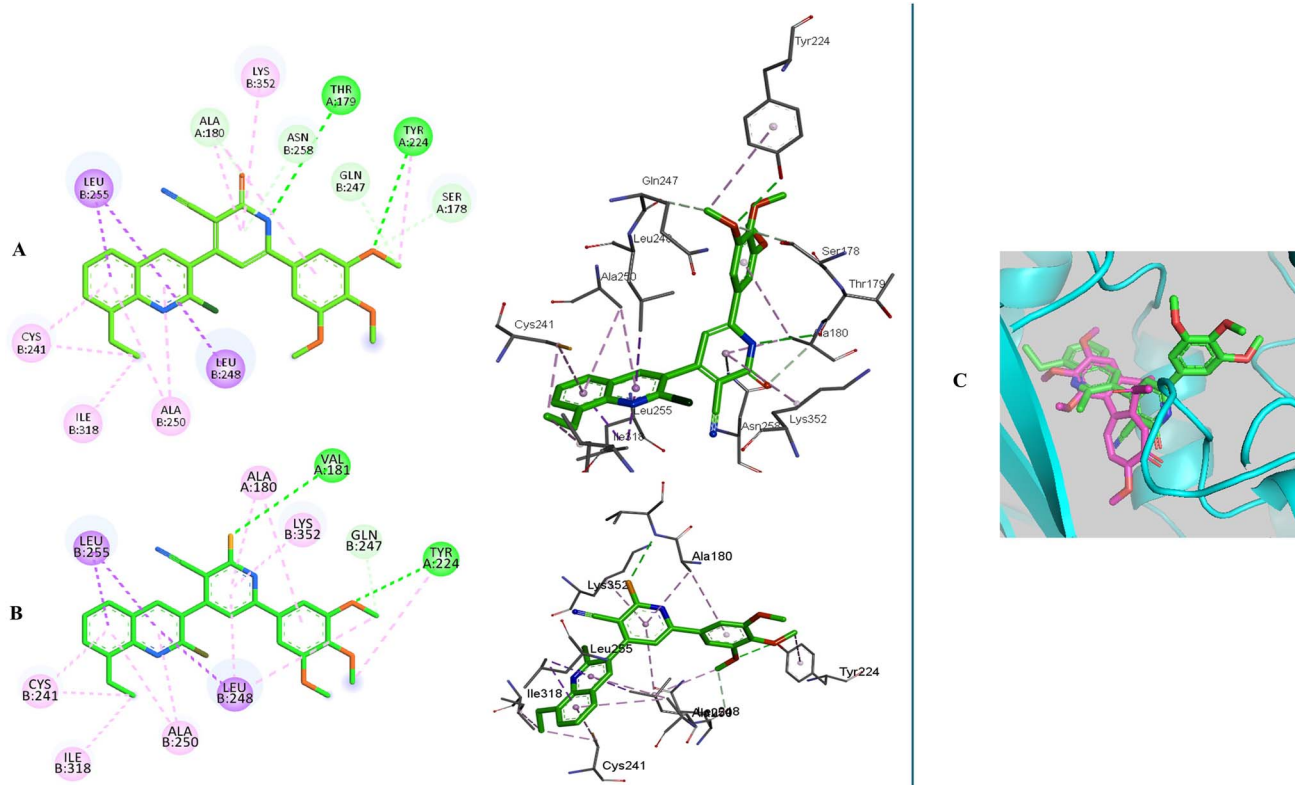


Fig. 13 2D and 3D interaction diagrams of (A) compound **4c**, (B) compound **5c** into the colchicine binding site of tubulin enzyme and (C) aligned docking pose of **colchicine** (pink) and **4c** into the CBS.



dimethoxy and trimethoxyphenyl rings of **3b** and **3c**, respectively made H-bond interactions with α Ser178 (3.33 and 3.29 Å, respectively) acting as hydrogen acceptors, both rings made π -alkyl interactions with α Ala180. The dimethoxy group in **3b** formed an extra H-bond with α Glu183 (3.48 Å) acting as a hydrogen donor while the trimethoxy group in **3c** extended towards β Lys254 forming π -alkyl interaction with the residue. The 2-oxo-1,2-dihydropyridine derivative **4c** was the most cytotoxic synthesized compound against 60 cell lines. It inhibited the growth of most of the tested cells and was lethal towards many non-small cell lung cancer cell lines, CNS cancer cells, renal and breast cancer cells, while its isosteric derivative 2-thioxo-1,2-dihydropyridine containing compound **5c** revealed less cytotoxic effect than **4c** and exhibited reasonable

cytotoxicity against most of the tested cell lines. Compound **4c** revealed the highest predicted binding interaction with CBS of tubulin compared with **5c** and **colchicine** equal to -11.5 , -10.9 and -9.8 kcal mol $^{-1}$, respectively, which support the results of the biological assay. Compounds **4c** and **5c** fitted comfortably into the active space of the CBS of tubulin, their mode of binding was almost identical and was like that of **colchicine** as revealed by superimposing both compounds (Fig. 13). The 2-chloro-8-ethylquinoline ring oriented itself towards the hydrophobic pocket of the CBS of tubulin forming multiple π -alkyl and π -sigma interactions with β Cys241, β Leu248, β Ala250, β Leu255 and β Ile318. The trimethoxy phenyl ring in both compounds faced the hydrophilic region of CBS of tubulin. Both compounds are linked to α Tyr224 and β Gln247. Compound **4c** formed hydrogen bond interactions with both residues (3.28 and 3.26 Å, respectively) acting as hydrogen acceptor and donor, respectively. Compound **5c** bonded to the same two residues with π -alkyl and π -sigma interactions, respectively. Moreover, **4c** made an additional hydrogen bond through its trimethoxy group with α Ser178 (3.34 Å) acting as a hydrogen acceptor. π -Alkyl interaction occurred between β Lys352 and the 2-oxo-1,2-dihydropyridine ring in compound **4c**, as well as with the 2-thioxo-1,2-dihydropyridine ring in compound **5c**. 2-Oxo-1,2-dihydropyridine moiety in **4c** linked by three hydrogen bonds to α Thr179, α Ala180 and β Asn258 (3.37,

Table 3 CDocker energy of the tested compounds **3b**, **3c**, **4c** and **5c** and colchicine

Compound	CDocker energy (kcal mol $^{-1}$), tubulin enzyme (PDB ID: 4O2B)
3b	-9.0
3c	-8.0
4c	-11.5
5c	-10.9
Colchicine	-9.8

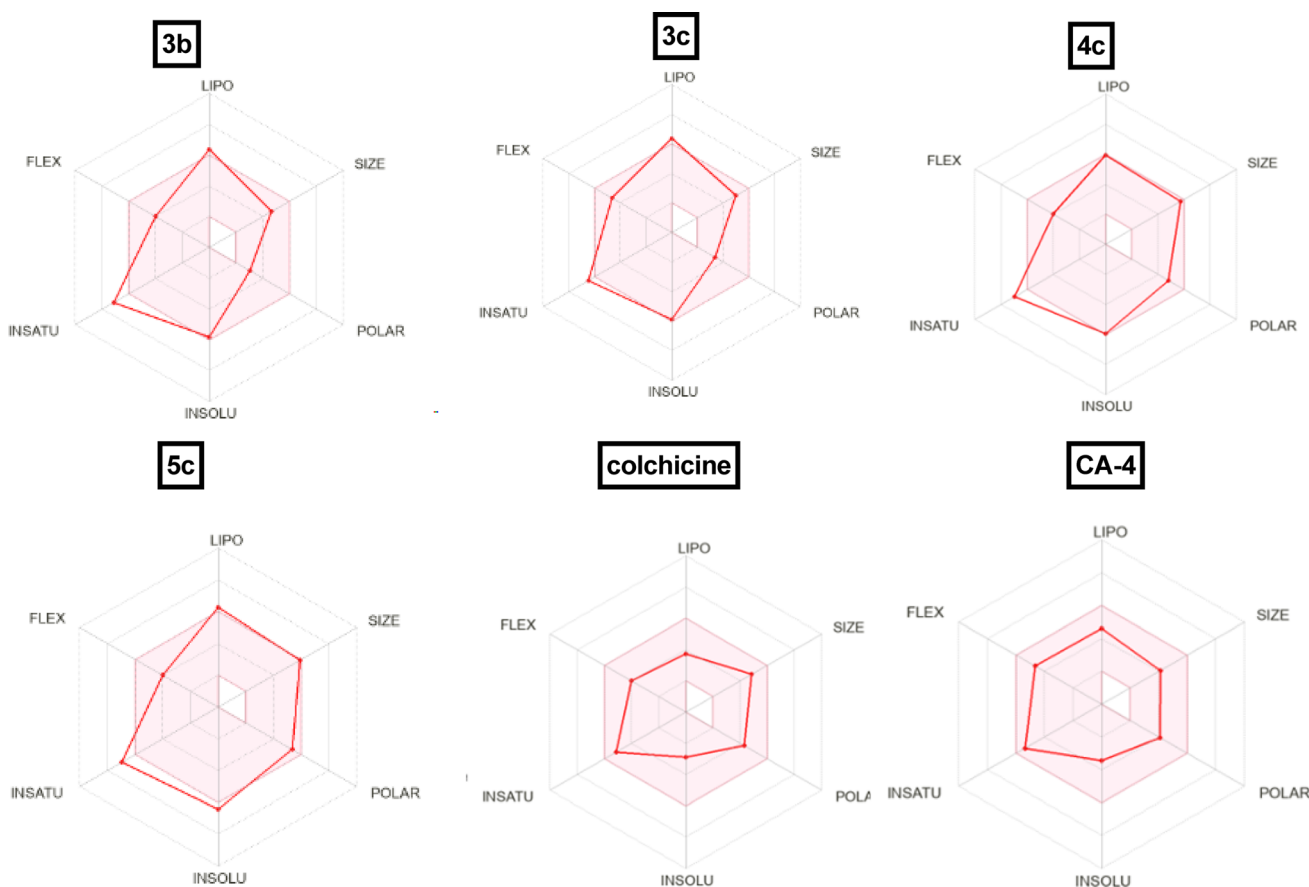


Fig. 14 Radar chart showing six predicted physicochemical properties of the tested compound **3b**, **3c**, **4c**, **5c**, colchicine and CA-4.



3.29, 4.12 Å, respectively) acting as hydrogen donor, acceptor and donor, respectively. Conversely, in compound **5c** the 2-thioxo-1,2-dihydropyridine ring in **5c** formed a hydrogen bond with α Val181 (3.65 Å) acting as hydrogen acceptor. Close analysis of the binding mode of **4c** revealed that it formed many hydrogen bond interactions with the hydrophilic part of CBS of tubulin in addition to the hydrophobic interactions with the β -tubulin subunit. These interactions support the promising activity of compound **4c** towards tubulin, as well as its cytotoxicity against numerous cancer cell lines (Table 3).

2.3.2. Physicochemical properties. The physicochemical features of the most promising cytotoxic compounds **3b**, **3c**, **4c** and **5c** against various tested cell lines were investigated compared to reference drugs **colchicine** and **CA-4**. The results of the ADME revealed six physicochemical properties: lipophilicity, size, polarity, solubility, saturation, and flexibility.

Compounds **3b**, **3c** and **5c** showed a slight deviation in the saturation and solubility properties from the validated range compared to **colchicine** and **CA-4** while compound **4c** showed a slight deviation in saturation only (Fig. 14) and compound **6c** showed no deviation in all properties. Other physicochemical parameters include topological polar surface area, pan assay interference structures, lipophilicity parameter WLOGP, number of rotatable bonds, number of hydrogen bond acceptors, number of hydrogen bond donors, gastrointestinal absorption blood–brain barrier permeability and drug likeness are shown in Table 4. Moreover, compounds **3b**, **3c** and **CA-4** could not be good substrates for *P*-glycoprotein (Pgp) while compounds **4c**, **5c** and **colchicine** could be good substrates for Pgp. In addition, compound **4c** and **colchicine** showed high GI absorption as they were located in the Boiled-Egg's white but compound **5c** showed low GI absorption as it wasn't located in

Table 4 The predicted pharmacokinetic profile for the investigated compounds **3b**, **3c**, **4c**, **5c**, **colchicine** and **CA-4**

Comp. no.	TPSA ^a	PAINS ^b	WLOGP ^c	NRB ^d	HBD ^e	HBA ^f	GI absorption ^g	BBB permeability ^h	Lipinski ⁱ
3b	48.42	0	5.26	6	0	4	High	Yes	0 violation
3c	57.65	0	5.26	7	0	5	High	Yes	0 violation
4c	97.23	0	5.37	6	1	6	High	No	0 violation
5c	112.25	0	6.74	6	1	5	Low	No	0 violation
Colchicine	83.09	0	2.55	6	1	6	High	No	0 violation
CA-4	77.38	0	2.38	7	2	6	High	Yes	0 violation

^a Topological polar surface area. ^b Pan assay interference structures. ^c Lipophilicity parameter WLOGP. ^d Number of rotatable bonds. ^e Number of hydrogen bond acceptors. ^f Number of hydrogen bond donor. ^g Gastrointestinal absorption. ^h Blood–brain barrier permeability. ⁱ Lipinski (drug likeness).

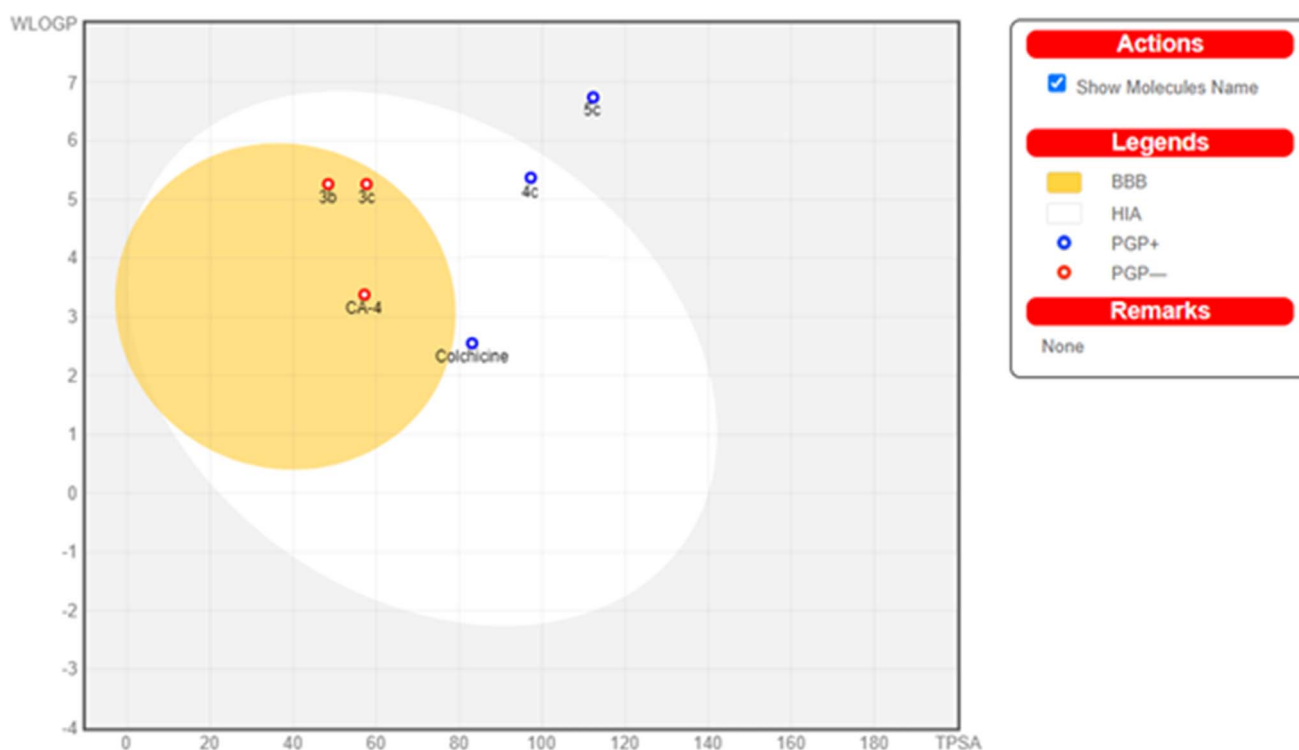


Fig. 15 Boiled egg model of the tested compounds **3b**, **3c**, **4c**, **5c**, **colchicine** and **CA-4**.



the Boiled-Egg's white. Finally, **3b**, **3c** and **CA-4** which were located in the Boiled-Egg yolk could penetrate the blood–brain barrier (Fig. 15).

3. Conclusion

To sum up, a group of novel quinoline derivatives **3a–e**, **4a–e**, **5a–e**, and **6a–e** were designed and synthesized as tubulin polymerization inhibitors targeting the **colchicine** binding site. Their cytotoxic activity against 60 cell lines by NCI were evaluated. Compounds **3b**, **3c**, **4c**, **5c** and **6c** possessed the most remarkable antitumor activity against almost all cell lines, especially compound **4c** (pyridin-2-one ring substituted with 3,4,5-trimethoxy groups) which was the most potent compound as an antiproliferative agent. Compound **4c** significantly induced cell cycle arrest at G2 and M phases and encouraged apoptosis, as evidenced by a rise in the number of cells in both early and late phases of apoptosis. Compound **4c** successfully inhibited tubulin polymerization with IC₅₀ value 17 ± 0.3 μM. The β-tubulin mRNA levels were markedly reduced in MDA-MB-231 cells treated with compound **4c**, resembling the effect observed with **colchicine** treatment. Molecular docking exhibited the interaction mode of compound **4c** with tubulin including formation of hydrogen bonds and hydrophobic interactions, which demonstrated energy score −11.5 kcal mol^{−1} in comparison to **colchicine** with energy score −9.8 kcal mol^{−1}, and it interacted with essential amino acids in the active sites. Compound **4c** could have high GI absorption, drug-like properties and the ability to penetrate the blood–brain barrier. Taken together, compound **4c** is a highly promising candidate for further preclinical studies against breast cancer.

4. Experimental

4.1. Chemistry

4.1.1. Procedure for synthesis of N-(2-ethylphenyl)acetamide (1).^{33,34,36} To a solution of 2-ethyl aniline (20 mmol, 2.42 mL) in glacial acetic acid (5 mL), acetic anhydride (20 mmol, 2.04 mL) was added, and the mixture was stirred in ice bath for 2 h. The mixture was poured onto ice-cooled water after completion of the reaction, the formed precipitate was collected by filtration and then dried to be used in the next step.

4.1.2. Procedure for synthesis of 2-chloro-8-ethylquinoline-3-carbaldehyde (2).^{35,36} DMF (62.5 mmol, 4.55 mL) in an ice bath (0–5 °C) was mixed continuously while phosphorus oxychloride (175 mmol, 26.82 mL) was added dropwise. Acetanilide **1** (25 mmol, 4.08 g) was then added portion-wise. The reaction mixture was heated in a water bath for 16 h, at 70 to 90 °C, and after that, it was poured onto ice-cold water and stirred until a precipitate was produced. The mixture was let to stand overnight before being filtered, dried, and crystallized from ethyl acetate to produce compound **2**.

4.1.3. General procedure for synthesis of 3-(2-chloro-8-ethylquinolin-3-yl)-1-(substituted phenyl)prop-2-en-1-one (3a–e). To a mixture of 2-chloro-8-ethylquinoline-3-carbaldehyde (**2**) (10 mmol, 2.19 g) and the appropriate substituted

acetophenone (10 mmol) in absolute ethanol (30 mL), an aqueous solution of potassium hydroxide (50%, 5 mL) was added while stirring. Then the reaction mixture was stirred for an additional 3 h at room temperature. The separated product was collected by filtration and crystallized from methanol to give compounds **3a–e**.

3-(2-Chloro-8-ethylquinolin-3-yl)-1-(4-methoxyphenyl)prop-2-en-1-one (3a). Yellow powder, yield 85%, mp 138–140 °C. IR (KBr, cm^{−1}): 3022 (CH aromatic), 2960 (CH aliphatic), 1656 (C=O), 1568 (C=N). ¹H NMR (400 MHz, DMSO-*d*₆), δ ppm: 1.30 (t, 3H, *J* = 8.0 Hz, CH₂CH₃), 3.11–3.16 (q, 2H, *J* = 8.0 Hz, CH₂CH₃), 3.89 (s, 3H, OCH₃), 7.12 (d, 2H, *J* = 8.0 Hz, Ar-H), 7.64 (t, 1H, *J* = 8.0 Hz, Ar-H), 7.71 (d, 1H, *J* = 8.0 Hz, Ar-H), 7.91 (d, 1H, *J* = 8.0 Hz, Ar-H), 8.02 (d, 1H, *J* = 16 Hz, CH alkene β proton), 8.16 (d, 1H, *J* = 16 Hz, CH alkene α proton), 8.21 (d, 2H, *J* = 8.0 Hz, Ar-H), 9.21 (s, 1H, Ar-H). ¹³C NMR (100 MHz, DMSO-*d*₆), δ ppm: 15.2, 24.0, 56.0, 114.5 (2C), 126.2, 126.9, 127.0, 127.4, 128.1, 130.4, 130.6, 131.5 (2C), 137.3, 138.0, 141.6, 145.9, 149.1, 163.9, 187.2. Anal. calcd for C₂₁H₁₈ClNO₂ (351.83): C, 71.69; H, 5.16; N 3.98; found: C, 71.85; H, 5.33; N, 4.21.

3-(2-Chloro-8-ethylquinolin-3-yl)-1-(3,4-dimethoxyphenyl)prop-2-en-1-one (3b). Yellow powder, yield 84%, mp 135–137 °C. IR (KBr, cm^{−1}): 3060 (CH aromatic), 2964 (CH aliphatic), 1653 (C=O), 1577 (C=N). ¹H NMR (400 MHz, DMSO-*d*₆), δ ppm: 1.29 (t, 3H, *J* = 8.0 Hz, CH₂CH₃), 3.09–3.15 (q, 2H, *J* = 8.0 Hz, CH₂CH₃), 3.87 (s, 3H, OCH₃), 3.89 (s, 3H, OCH₃), 7.13 (d, 1H, *J* = 8.0 Hz, Ar-H), 7.62 (t, 2H, *J* = 8.0 Hz, Ar-H), 7.70 (d, 1H, *J* = 8.0 Hz, Ar-H), 7.89–7.95 (m, 2H, Ar-H), 8.00 (d, 1H, *J* = 16 Hz, CH alkene β proton), 8.13 (d, 1H, *J* = 16 Hz, CH alkene α proton), 9.15 (s, 1H, Ar-H). ¹³C NMR (100 MHz, DMSO-*d*₆), δ ppm: 15.3, 24.0, 56.1, 56.3, 111.2, 111.3, 124.3, 126.5, 127.0, 127.3, 127.5, 128.3, 130.5, 130.8, 137.4, 138.2, 141.7, 146.0, 149.2, 149.3, 154.0, 187.3. Anal. calcd for C₂₂H₂₀ClNO₃ (381.86): C, 69.20; H, 5.28; N 3.67; found: C, 69.43; H, 5.40; N, 3.85.

3-(2-Chloro-8-ethylquinolin-3-yl)-1-(3,4,5-trimethoxyphenyl)prop-2-en-1-one (3c). White powder, yield 86%, mp 140–142 °C. IR (KBr, cm^{−1}): 3064 (CH aromatic), 2958 (CH aliphatic), 1639 (C=O), 1583 (C=N). ¹H NMR (400 MHz, DMSO-*d*₆), δ ppm: 1.28 (t, 3H, *J* = 8.0 Hz, CH₂CH₃), 3.06–3.15 (q, 2H, *J* = 8.0 Hz, CH₂CH₃), 3.79 (s, 3H, OCH₃), 3.92 (s, 6H, 2 × OCH₃), 7.46 (s, 2H, Ar-H), 7.64 (t, 1H, *J* = 8.0 Hz, Ar-H), 7.72 (d, 1H, *J* = 8.0 Hz, Ar-H), 7.93 (d, 1H, *J* = 8.0 Hz, Ar-H), 8.04 (d, 1H, *J* = 16 Hz, CH alkene β proton), 8.12 (d, 1H, *J* = 12 Hz, CH alkene α proton), 9.14 (s, 1H, Ar-H). ¹³C NMR (100 MHz, DMSO-*d*₆), δ ppm: 15.3, 24.0, 56.7, 60.7 (2C), 106.8 (2C), 126.1, 126.2, 127.0, 127.4, 130.8, 132.8, 138.2, 138.3, 141.7, 142.7, 145.9, 149.1, 150.7, 153.3 (2C), 188.0. Anal. calcd for C₂₃H₂₂ClNO₄ (411.88): C, 67.07; H, 5.38; N 3.40; found: C, 67.41; H, 5.45; N, 3.62.

3-(2-Chloro-8-ethylquinolin-3-yl)-1-(4-fluorophenyl)prop-2-en-1-one (3d). White powder, yield 83%, mp 215–217 °C. IR (KBr, cm^{−1}): 3078 (CH aromatic), 2964 (CH aliphatic), 1643 (C=O), 1598 (C=N). ¹H NMR (400 MHz, DMSO-*d*₆), δ ppm: 1.31 (t, 3H, *J* = 8.0 Hz, CH₂CH₃), 3.11–3.17 (q, 2H, *J* = 8.0 Hz, CH₂CH₃), 7.46 (t, 2H, *J* = 8.0 Hz, Ar-H), 7.60 (d, 1H, *J* = 7.2 Hz, Ar-H), 7.64 (d, 1H, *J* = 16 Hz, CH alkene β proton), 7.74 (t, 1H, *J* = 8.0 Hz, Ar-H), 8.06–8.09 (m, 3H, Ar-H), 8.20 (d, 1H, *J* = 16 Hz, CH alkene α proton), 9.29 (s, 1H, Ar-H). ¹³C NMR (100 MHz, DMSO-*d*₆),



δ ppm: 15.5, 23.6, 115.3 (2C), 126.1, 126.9, 127.0, 127.5, 128.1, 130.4, 130.6, 131.6 (2C), 137.4, 138.1, 141.6, 146.0, 149.2, 167.3, 187.5. Anal. calcd for $C_{20}H_{15}ClFNO$ (339.79): C, 70.70; H, 4.45; N, 4.12; found: C, 70.59; H, 4.62; N, 4.37.

1-(4-Bromophenyl)-3-(2-chloro-8-ethylquinolin-3-yl)prop-2-en-1-one (3e). White powder, yield 83%, mp 210–212 °C. IR (KBr, cm^{-1}): 3101 (CH aromatic), 2958 (CH aliphatic), 1600 (C=O), 1583 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.31 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.06–3.15 (q, 2H, $J = 8.0$ Hz, CH_2CH_3), 7.06 (d, 1H, $J = 8$ Hz, Ar-H), 7.36 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.61 (m, 1H, Ar-H), 7.73 (d, 2H, $J = 8.4$ Hz, Ar-H), 7.91 (d, 2H, $J = 8.4$ Hz, Ar-H), 8.11 (d, 1H, $J = 15.2$ Hz, CH alkene β proton), 8.17 (d, 1H, $J = 16$ Hz, CH alkene α proton), 8.50 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.9, 23.6, 126.2, 126.8, 127.1, 127.5, 128.2, 128.5 (2C), 129.1, 130.3, 130.6, 131.7 (2C), 137.9, 139.9, 142.3, 146.0, 150.3, 186.8. Anal. calcd for $C_{20}H_{15}ClBrNO$ (400.70): C, 59.95; H, 3.77; N 3.50; found: C, 60.23; H, 3.91; N, 3.64.

4.1.4. General procedure for synthesis of 4-(2-chloro-8-ethylquinolin-3-yl)-6-(substituted phenyl)-2-oxo-1,2-dihydropyridine-3-carbonitrile (4a–e).

2-Chloro-8-ethylquinoline-3-carbaldehyde (**2**) (1 mmol, 0.219 g) was heated under reflux for 4–6 h with the appropriate substituted acetophenone (1 mmol), ethyl cyanoacetate (1 mmol, 0.11 g), and ammonium acetate (8 mmol, 0.62 g) in absolute ethanol (10 mL). After cooling the formed solid was filtered, dried and crystallized from methanol to give compounds **4a–e**.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-methoxyphenyl)-2-oxo-1,2-dihydropyridine-3-carbonitrile (4a). Yellow powder, yield 79%, mp 270–272 °C. IR (KBr, cm^{-1}): 3421 (NH), 3095 (CH aromatic), 2962 (CH aliphatic), 2218 (CN), 1647 (C=O), 1514 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.34 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.14–3.20 (m, 2H, CH_2CH_3 + s, 1H, NH, D_2O exchangeable), 3.84 (s, 3H, OCH_3), 7.00 (s, 1H, CH pyridine), 7.08 (d, 2H, $J = 8.8$ Hz, Ar-H), 7.69 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.80 (d, 1H, $J = 6.8$ Hz, Ar-H), 7.92 (d, 2H, $J = 8.8$ Hz, Ar-H), 7.96 (d, 1H, $J = 8.0$ Hz, Ar-H), 8.66 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.3, 23.9, 55.7, 106.4, 115.0 (2C), 116.4, 124.8, 126.1, 126.8, 126.9, 127.8, 127.9, 128.6, 129.8, 129.9 (2C), 131.0, 140.3, 141.9, 145.9, 152.9, 156.7, 162.4. MS m/z (%): 417.40 (M + 2, 5.89), 415.49 (M $^+$, 13.79), 125.14 (100.00). Anal. calcd for $C_{24}H_{18}ClN_3O_2$ (415.88): C, 69.31; H, 4.36; N 10.10; found: C, 69.08; H, 4.50; N, 10.29.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(3,4-dimethoxyphenyl)-2-oxo-1,2-dihydropyridine-3-carbonitrile (4b). Yellow powder, yield 71%, mp 280–282 °C. IR (KBr, cm^{-1}): 3431 (NH), 3040 (CH aromatic), 2962 (CH aliphatic), 2239 (CN), 1641 (C=O), 1517 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.34 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 2.52 (s, 1H, NH, D_2O exchangeable), 3.17–3.26 (m, 2H, CH_2CH_3), 3.80 (s, 3H, OCH_3), 3.83 (s, 3H, OCH_3), 6.85 (s, 1H, CH pyridine), 7.00 (d, 1H, $J = 8.4$ Hz, Ar-H), 7.60–7.67 (m, 3H, Ar-H), 7.75 (d, 1H, $J = 6.8$ Hz, Ar-H), 7.95 (d, 1H, $J = 8.0$ Hz, Ar-H), 8.52 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.5, 22.6, 56.0, 56.1, 105.8, 110.5, 110.8, 111.0, 111.8, 181.2, 121.0, 123.7, 126.7, 127.1, 128.1, 130.4, 130.9, 140.0, 141.9, 145.8, 146.8, 149.3, 151.3, 156.1, 173.9. MS m/z (%): 447.88 (M + 2, 3.22), 445.74 (M $^+$, 9.87), 313.27 (100.00). Anal. calcd for

$C_{25}H_{20}ClN_3O_3$ (445.90): C, 67.34; H, 4.52; N 9.42; found: C, 67.53; H, 4.61; N, 9.68.

4-(2-Chloro-8-ethylquinolin-3-yl)-2-oxo-6-(3,4,5-trimethoxyphenyl)-1,2-dihydropyridine-3-carbonitrile (4c). Yellow powder, yield 72%, mp 297–299 °C. IR (KBr, cm^{-1}): 3446 (NH), 3140 (CH aromatic), 2953 (CH aliphatic), 2220 (CN), 1645 (C=O), 1581 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.33 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.14–3.23 (m, 3H (2H, CH_2CH_3 + 1H, NH, D_2O exchangeable)), 3.73 (s, 3H, OCH_3), 3.86 (s, 6H, $2OCH_3$), 7.16 (s, 1H, CH pyridine), 7.26 (s, 2H, Ar-H), 7.69 (t 1H, $J = 8.0$ Hz, Ar-H), 7.80 (d, 1H, $J = 8$ Hz, Ar-H), 7.98 (d, 1H, $J = 8$ Hz, Ar-H), 8.66 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 16.2, 23.9, 56.7, 60.8 (2C), 105.6 (2C), 107.1, 116.3, 116.7, 126.9, 127.6, 128.0, 128.6, 129.8, 130.7, 140.3, 140.5, 141.8, 145.9, 146.1, 152.6, 153.6 (2C), 156.7, 162.6. MS m/z (%): 477.68 (M + 2, 11.23), 475.05 (M $^+$, 34.00), 42.89 (100.00). Anal. calcd for $C_{26}H_{22}ClN_3O_4$ (475.93): C, 65.62; H, 4.66; N 8.83; found: C, 65.49; H, 4.85; N, 9.07.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-fluorophenyl)-2-oxo-1,2-dihydropyridine-3-carbonitrile (4d). White powder, yield 77%, mp 260–262 °C. IR (KBr, cm^{-1}): 3431 (NH), 3047 (CH aromatic), 2966 (CH aliphatic), 2222 (CN), 1656 (C=O), 1535 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.34 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.11–3.21 (m, 2H, CH_2CH_3 + s, 1H, NH, D_2O exchangeable), 7.11 (s, 1H, CH pyridine), 7.39 (t, 2H, $J = 8.0$ Hz, Ar-H), 7.63–7.71 (m, 1H, Ar-H), 7.80 (d, 1H, $J = 6.8$ Hz, Ar-H), 7.96–7.803 (m, 3H, Ar-H), 8.68 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 16.6, 23.3, 106.2, 114.6 (2C), 117.6, 119.5, 120.6, 122.5, 124.2, 125.9, 127.2 (2C), 127.9, 129.2, 129.9, 131.3, 144.6, 147.3, 153.6, 159.7, 161.7, 162.4. MS m/z (%): 405.62 (M + 2, 4.18), 403.80 (M $^+$, 11.20), 376.34 (100.00). Anal. calcd for $C_{23}H_{15}ClFN_3O$ (403.84): C, 68.41; H, 3.74; N, 10.41; found: C, 68.70; H, 3.86; N 10.64.

6-(4-Bromophenyl)-4-(2-chloro-8-ethylquinolin-3-yl)-2-oxo-1,2-dihydropyridine-3-carbonitrile (4e). White powder, yield 78%, mp 290–262 °C. IR (KBr, cm^{-1}): 3419 (NH), 3034 (CH aromatic), 2964 (CH aliphatic), 2222 (CN), 1647 (C=O), 1589 (C=N). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.33 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 1.87 (s, 1H, NH, D_2O exchangeable), 3.06–3.15 (m, 2H, CH_2CH_3), 7.00 (s, 1H, CH pyridine), 7.69–7.70 (m, 3H, Ar-H), 7.79 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.91–7.96 (m, 3H, Ar-H), 8.61 (s, 1H, Ar-H). MS m/z (%): 466.63 (M + 2, 29.98), 465.61 (M + 1, 54.96), 464.52 (M $^+$, 48.74), 353.35 (100.00). Anal. calcd for $C_{23}H_{15}BrClN_3O$ (464.75): C, 59.44; H, 3.25; N 9.04; found: C, 59.32; H, 3.41; N, 9.29.

4.1.5. General procedure for synthesis of 4-(2-chloro-8-ethylquinolin-3-yl)-6-(substituted phenyl)-2-thio-1,2-dihydropyridine-3-carbonitrile (5a–e). A mixture of 2-chloro-8-ethylquinoline-3-carbaldehyde (**2**). (1 mmol, 0.219 g), required acetophenones (1 mmol), 2-cyanothioacetamide (1 mmol, 0.1 g), and ammonium acetate (8 mmol, 0.62 g) in absolute ethanol (10 mL) was heated under reflux for 6–8 h. After the reaction mixture was cooled to room temperature, the separated solid was filtered, dried and crystallized from methanol to afford compounds **5a–e**.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-methoxyphenyl)-2-thio-1,2-dihydropyridine-3-carbonitrile (5a). Yellow powder, yield



71%, mp 290–292 °C. IR (KBr, cm^{-1}): 3433 (NH), 3070 (CH aromatic), 2962 (CH aliphatic), 2218 (CN), 1577 (C=N), 1147 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.33 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.15–3.21 (q, 2H, $J = 8.0$ Hz, $\text{CH}_2\text{CH}_3 + \text{s}$, 1H, NH, D_2O exchangeable), 3.83 (s, 3H, OCH_3), 6.99 (s, 1H, CH pyridine), 7.08 (d, 2H, $J = 9.3$ Hz, Ar-H), 7.68 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.79 (d, 1H, $J = 7.2$ Hz, Ar-H), 7.92 (d, 2H, $J = 8.8$ Hz, Ar-H), 7.96 (d, 1H, $J = 7.6$ Hz, Ar-H), 8.65 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.3, 24.3, 56.3, 106.7, 114.9 (2C), 117.0, 120.7, 121.4, 124.8, 125.4, 126.7, 126.9, 128.6, 130.0 (2C), 131.0, 140.3, 142.0, 145.8, 152.8, 156.5, 160.1, 162.4. MS m/z (%): 433.93 (M + 2, 16.24), 432.21 (M + 1, 40.09), 431.34 (M^+ , 50.68), 423.98 (100.00). Anal. calcd for $\text{C}_{24}\text{H}_{18}\text{ClN}_3\text{OS}$ (431.94): C, 66.74; H, 4.20; N 9.73; found: C, 66.83; H, 4.34; N, 9.91.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(3,4-dimethoxyphenyl)-2-thioxo-1,2-dihydropyridine-3-carbonitrile (**5b**). Yellow powder, yield 71%, mp 280–282 °C. IR (KBr, cm^{-1}): 3446 (NH), 3151 (CH aromatic), 2964 (CH aliphatic), 2218 (CN), 1577 (C=N), 1145 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.33 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 1.88 (s, 1H, NH, D_2O exchangeable), 3.20 (s, 2H, CH_2CH_3), 3.83 (s, 3H, OCH_3), 3.84 (s, 3H, OCH_3), 7.02 (s, 1H, CH pyridine), 7.08 (d, 1H, $J = 8.4$ Hz, Ar-H), 7.33–7.61 (m, 2H, Ar-H), 7.68 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.79 (d, 1H, $J = 6.8$ Hz, Ar-H), 7.96 (d, 1H, $J = 7.6$ Hz, Ar-H), 8.63 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.3, 24.3, 56.1, 56.2, 100.0, 106.5, 110.9, 112.3, 116.4, 121.6, 124.6, 124.7, 126.8, 126.9, 128.3, 129.9, 130.8, 140.3, 141.8, 145.9, 146.1, 149.3, 152.1, 156.7, 162.5. MS m/z (%): 463.93 (M + 2, 2.78), 462.63 (M + 1, 9.64), 461.68 (M^+ , 6.26), 43.21 (100.00). Anal. calcd for $\text{C}_{25}\text{H}_{20}\text{ClN}_3\text{O}_2\text{S}$ (461.96): C, 65.00; H, 4.36; N 9.10; found: C, 64.88; H, 4.51; N, 9.32.

4-(2-Chloro-8-ethylquinolin-3-yl)-2-thioxo-6-(3,4,5-trimethoxyphenyl)-1,2-dihydropyridine-3-carbonitrile (**5c**). Yellow powder, yield 69%, mp 279–281 °C. IR (KBr, cm^{-1}): 3446 (NH), 3051 (CH aromatic), 2966 (CH aliphatic), 2220 (CN), 1585 (C=N), 1128 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.31 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.18–3.22 (m, 2H, $\text{CH}_2\text{CH}_3 + \text{s}$, 1H, NH, D_2O exchangeable), 3.73 (s, 3H, OCH_3), 3.86 (s, 6H, $2 \times \text{OCH}_3$), 7.17 (s, 1H, CH pyridine), 7.25 (s, 2H, Ar-H), 7.67 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.80 (d, 1H, $J = 8$ Hz, Ar-H), 7.98 (d, 1H, $J = 8$ Hz, Ar-H), 8.69 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 14.9, 23.7, 56.8 (2C), 60.5, 101.8, 106.4, 109.2, 116.0 (2C), 126.1, 126.9, 127.4, 128.1, 129.6, 130.8, 132.7, 138.0, 138.4, 146.1, 149.3, 153.2 (2C), 159.5, 173.7, 175.8. MS m/z (%): 493.53 (M + 2, 26.83), 491.95 (M^+ , 72.17), 441.90 (100.00). Anal. calcd for $\text{C}_{26}\text{H}_{22}\text{ClN}_3\text{O}_3\text{S}$ (491.99): C, 63.47; H, 4.51; N 8.54; found: C, 63.60; H, 4.63; N, 8.70.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-fluorophenyl)-2-thioxo-1,2-dihydropyridine-3-carbonitrile (**5d**). Yellow powder, yield 76%, mp 235–237 °C. IR (KBr, cm^{-1}): 3398 (NH), 3049 (CH aromatic), 2966 (CH aliphatic), 2222 (CN), 1508 (C=N), 1165 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.34 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.11–3.22 (m, 2H, $\text{CH}_2\text{CH}_3 + \text{s}$, 1H, NH, D_2O exchangeable), 6.78 (s, 1H, CH pyridine), 7.23 (t, 2H, Ar-H), 7.65 (d, 1H, $J = 8.0$ Hz, Ar-H), 7.73–7.77 (m, 1H, Ar-H), 7.94 (d, 1H, $J = 8$ Hz, Ar-H), 8.08 (d, 2H, $J = 7.6$ Hz, Ar-H), 8.49 (s, 1H, Ar-H). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.0, 22.0, 105.5, 115.9, 116.1

(2C), 118.3, 126.1, 126.7, 128.1, 128.3 (2C), 128.5, 129.7, 130.0, 133.9, 141.5, 145.2, 145.7, 149.7, 162.2, 166.4, 173.6. MS m/z (%): 421.12 (M + 2, 9.84), 419.78 (M^+ , 27.88), 64.08 (100.00). Anal. calcd for $\text{C}_{23}\text{H}_{15}\text{ClFN}_3\text{S}$ (419.90): C, 65.79; H, 3.60; N 10.01; found: C, 66.05; H, 3.74; N, 10.29.

6-(4-Bromophenyl)-4-(2-chloro-8-ethylquinolin-3-yl)-2-thioxo-1,2-dihydropyridine-3-carbonitrile (**5e**). Buff powder, yield 79%, mp charring 250 °C. IR (KBr, cm^{-1}): 3446 (NH), 3126 (CH aromatic), 2910 (CH aliphatic), 2218 (CN), 1564 (C=N), 1195 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.33 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.17–3.26 (m, 2H, $\text{CH}_2\text{CH}_3 + \text{s}$, 1H, NH, D_2O exchangeable), 6.99 (s, 1H, CH pyridine), 7.05 (d, 2H, $J = 9.2$ Hz, Ar-H), 7.66 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.77 (d, 1H, $J = 7.2$ Hz, Ar-H), 7.91–7.97 (m, 3H, Ar-H), 8.67 (s, 1H, Ar-H). MS m/z (%): 482.73 (M + 2, 19.45), 481.83 (M + 1, 17.34), 480.23 (M^+ , 41.55), 474.31 (100.00). Anal. calcd for $\text{C}_{23}\text{H}_{15}\text{BrClN}_3\text{S}$ (480.81): C, 57.46; H, 3.14; N 8.74; found: C, 57.70; H, 3.23; N, 8.97.

4.1.6. General procedure for synthesis of 4-(2-chloro-8-ethylquinolin-3-yl)-6-(substituted phenyl)-3,4-dihydropyrimidine-2(1H)-thione (6a-e). The corresponding chalcone **3a-e** (10 mmol) and thiourea (10 mmol, 0.76 g) were added successively to a solution of sodium ethoxide (10 mmol) in absolute ethanol (15 mL). The reaction mixture was heated under reflux for 2 h. After completion of the reaction, the solvent was concentrated under reduced pressure then, poured into ice-cooled water and acidified with acetic acid (2 mL). The obtained precipitate was filtered, washed with water, dried and crystallized from ethanol to afford pyrimidine derivatives **6a-e**.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-methoxyphenyl)-3,4-dihydropyrimidine-2(1H)-thione (**6a**). White powder, yield 71%, mp above 300 °C. IR (KBr, cm^{-1}): 3165 (NH), 3016 (CH aromatic), 2964 (CH aliphatic), 1130 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.28 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.12–3.19 (q, 2H, $J = 8.0$ Hz, CH_2CH_3), 3.76 (s, 3H, OCH_3), 5.39 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 5.57 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 6.91 (d, 2H, $J = 8.8$ Hz, Ar-H), 7.45 (d, 2H, $J = 8.8$ Hz, Ar-H), 7.60 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.69 (d, 1H, $J = 7.6$ Hz, Ar-H), 7.89 (d, 1H, $J = 7.6$ Hz, Ar-H), 8.21 (s, 1H, Ar-H), 9.16 (s, 1H, NH, D_2O exchangeable), 10.04 (s, 1H, NH, D_2O exchangeable). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 16.3, 24.1, 55.3, 58.3, 99.7, 120.7 (2C), 124.1, 125.8, 126.5, 126.8, 128.8, 130.2, 131.9 (2C), 136.3, 139.3, 141.7, 145.8, 148.5, 159.4, 175.6. Anal. calcd for $\text{C}_{22}\text{H}_{20}\text{ClN}_3\text{OS}$: C, 64.46; H, 4.92; N 10.25; found: C, 64.32; H, 5.11; N, 10.49.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(3,4-dimethoxyphenyl)-3,4-dihydropyrimidine-2(1H)-thione (**6b**). Orange powder, yield 74%, mp 237–239 °C. IR (KBr, cm^{-1}): IR (KBr, cm^{-1}): 3196 (NH), 3080 (CH aromatic), 2960 (CH aliphatic), 1172 (C=S). ^1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.29 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.12–3.16 (m, 2H, CH_2CH_3), 3.75 (s, 3H, OCH_3), 3.78 (s, 3H, OCH_3), 5.42 (d, 1H, CH pyrimidine, $J = 4$ Hz), 5.58 (d, 1H, CH pyrimidine, $J = 4$ Hz), 6.92 (d, 1H, $J = 8$ Hz, Ar-H), 7.05–7.08 (m, 2H, Ar-H), 7.61 (d, 1H, $J = 7.6$ Hz, Ar-H), 7.69 (d, 1H, $J = 7.2$ Hz, Ar-H), 7.90 (d, 1H, $J = 8$ Hz, Ar-H), 8.22 (s, 1H, Ar-H), 9.10 (s, 1H, NH, D_2O exchangeable), 10.16 (s, 1H, NH, D_2O exchangeable). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.0, 24.1, 56.1, 56.2, 56.2, 97.5, 111.3, 124.0, 124.5, 126.3, 126.7, 127.1, 127.5, 128.2, 130.4,



130.8, 137.5, 146.0, 149.0, 149.4, 151.5, 154.0, 174.3. Anal. calcd for $C_{23}H_{22}ClN_3O_2S$: C, 62.79; H, 5.04; N 9.55; found: C, 62.51; H, 5.25; N, 9.76.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(3,4,5-trimethoxyphenyl)-3,4-dihydropyrimidine-2(1H)-thione (**6c**). Yellow powder, yield 72%, mp 235–237 °C. IR (KBr, cm^{-1}): 3165 (NH), 3099 (CH aromatic), 2964 (CH aliphatic), 1130 (C=S). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.29 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.12–3.18 (q, 2H, $J = 8.0$ Hz, CH_2CH_3), 3.65 (s, 3H, OCH_3), 3.80 (s, 6H, $2OCH_3$), 5.51 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 5.61 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 6.81 (s, 2H, Ar-H), 7.60 (t, 1H, $J = 8.0$ Hz, Ar-H), 7.69 (d, 1H, $J = 8$ Hz, Ar-H), 7.92 (d, 1H, $J = 8$ Hz, Ar-H), 8.31 (s, 1H, Ar-H), 9.10 (s, 1H, NH, D_2O exchangeable), 10.16 (s, 1H, NH, D_2O exchangeable). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.2, 24.1, 53.3, 56.0 (2C), 60.7, 99.4, 103.4 (2C), 126.3, 127.6, 128.2, 128.9, 129.8, 134.6, 135.7, 137.9, 138.6, 141.6, 145.2, 147.3, 153.0 (2C), 177.0. Anal. calcd for $C_{24}H_{24}ClN_3O_3S$: C, 61.33; H, 5.15; N 8.94; found: C, 61.08; H, 5.32; N, 9.09.

4-(2-Chloro-8-ethylquinolin-3-yl)-6-(4-fluorophenyl)-3,4-dihydropyrimidine-2(1H)-thione (**6d**). Orange powder, yield 70%, mp 220–222 °C. IR (KBr, cm^{-1}): 3167 (NH), 3016 (CH aromatic), 2960 (CH aliphatic), 1197 (C=S). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.29 (t, 3H, $J = 8.0$ Hz CH_2CH_3), 3.12–3.17 (q, 2H, $J = 8.0$ Hz, CH_2CH_3), 5.44 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 5.59 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 7.19 (t, 2H, $J = 8.0$ Hz, Ar-H), 7.53–7.62 (m, 3H, Ar-H), 7.69 (d, 1H, $J = 6.8$ Hz, Ar-H), 7.90 (d, 1H, $J = 8$ Hz, Ar-H), 8.22 (s, 1H, Ar-H), 9.19 (s, 1H, NH, D_2O exchangeable), 10.17 (s, 1H, NH, D_2O exchangeable). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 16.0, 24.1, 59.2, 97.3, 115.8 (2C), 125.0, 126.6, 127.0, 128.2, 128.7, 129.9, 130.4 (2C), 131.1, 136.5, 145.5, 146.8, 149.7, 162.2, 174.0. Anal. calcd for $C_{21}H_{17}ClFN_3S$: C, 63.39; H, 4.31; N 10.56; found: C, 63.54; H, 4.45; N, 10.78.

6-(4-Bromophenyl)-4-(2-chloro-8-ethylquinolin-3-yl)-3,4-dihydropyrimidine-2(1H)-thione (**6e**). Orange powder, yield 74%, mp 248–240 °C. IR (KBr, cm^{-1}): 3162 (NH), 3018 (CH aromatic), 2964 (CH aliphatic), 1197 (C=S). 1H NMR (400 MHz, DMSO- d_6), δ ppm: 1.29 (t, 3H, $J = 8.0$ Hz, CH_2CH_3), 3.12–3.19 (m, 2H, CH_2CH_3), 5.50 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 5.60 (d, 1H, CH pyrimidine, $J = 4.4$ Hz), 7.46 (d, 2H, $J = 8.4$ Hz, Ar-H), 7.54–7.62 (m, 3H, Ar-H), 7.69 (d, 1H, $J = 7.2$ Hz, Ar-H), 7.90 (d, 1H, $J = 7.6$ Hz, Ar-H), 8.25 (s, 1H, Ar-H), 9.18 (s, 1H, NH, D_2O exchangeable), 10.20 (s, 1H, NH, D_2O exchangeable). ^{13}C NMR (100 MHz, DMSO- d_6), δ ppm: 15.0, 24.5, 57.9, 96.5, 123.1, 123.8, 124.5, 129.1 (2C), 129.8, 130.5, 132.6 (2C), 133.3, 137.2, 139.7, 140.8, 148.9, 152.8, 153.5, 174.4. Anal. calcd for $C_{21}H_{17}BrClN_3S$: C, 54.98; H, 3.73; N 9.16; found: C, 54.87; H, 3.90; N, 9.35.

4.2. Biology

4.2.1. In vitro cytotoxic activity screening. The cytotoxic activity of the synthesized quinoline derivatives was determined by the national cancer institute (NCI, Bethesda, Maryland, USA) against 60 different cancer cell lines at a single high dose (10^{-5} M) in the full NCI 60 cell panel using the sulforhodamine B assay for cell viability evaluation. In summary, 5×10^3 cells were in 96-well plate and incubated in complete media for 24 h. Cells were treated with 100 μ L of media containing each newly

synthesized compound. After drug exposure, cells were fixed by replacing media with 150 μ L of 10% TCA and incubated at 4 °C for 1 h. The TCA solution was removed, and the cells were washed 5 times with distilled water. Aliquots of 70 μ L SRB solution (0.4% w/v) were added and incubated in a dark place at room temperature for 10 min. The plates were washed 3 times with 1% acetic acid and allowed to air-dry overnight. Then, 150 μ L of TRIS (10 mM) was added to dissolve protein-bound SRB stain. The absorbance was measured at 540 nm.

$$\% \text{ Cytotoxicity} = [1 - (AV_x/AV_{NC})] \times 100$$

where AV_x denotes the average absorbance of the sample well and AV_{NC} denotes the average absorbance of the negative control.

4.2.1.1. Five doses testing of compound 4c. The methodology of the NCI anticancer screening has been described in details at (<http://www.dtp.nci.nih.gov>). The anticancer assay was performed in full NCI 60 cell lines derived from nine tumor subpanels, including leukemia, lung, colon, CNS, melanoma, ovarian, renal, prostate, and breast cancer cell lines.

The cell lines were first cultured for 48 h, after being exposed to a single dosage of the compound. Using the protein-binding dye sulforhodamine B dye, cell proliferation and viability were evaluated (SRB assay). The results were presented for each chemical as GI%. Compound that met NCI criteria for substantial activity were further evaluated using a five-dose experiment. The NCI60 cell lines were treated with the chemicals in this assay at five different concentration levels (0.01, 0.1, 1, 10, and 100 μ M). Following cell line incubation, the results were depicted using three parameters: GI₅₀, which signifies the molar concentration resulting in a 50% reduction in all cell count after incubation; TGI, indicating the molar concentration leading to total growth inhibition; and LC₅₀, representing the molar concentration causing a 50% loss of the initial cells after incubation.

4.2.2. Cell cycle analysis by flow cytometry. Propidium Iodide Flow Cytometry Kit for Cell Cycle Analysis (Abcam) was used according to manufacturer's instructions. Briefly, MDA-MB 231 cells (2×10^5 cells) were seeded in 6-well plates and cultured for 24 h. The cells were treated with IC₅₀ of the synthesized compound for 24 h. Cells were collected, washed with PBS and centrifuged. Subsequently, the cells were re-suspended and fixed in 66% ethanol on ice for at least 2 h. The cells were stained with a working solution containing propidium iodide (PI) in the dark at 37 °C for 30 min. DNA content of the cells were measured by flow cytometry using FACSCalibur (BD Biosciences, Mountain View, CA). The data were analyzed using the CellQuest software (BD Biosciences).²¹

4.2.3. Cell apoptosis analysis. Annexin-V-FITC Apoptosis Detection Kit (Biovision, USA) was used to determine apoptotic cells according to manufacturer's instructions. Briefly, MDA-MB 231 cells (2×10^5 cells) were seeded in 6-well plates and cultured for 24 h. The cells were treated with IC₅₀ of the synthesized compound for 24 h. Cells were collected, washed with PBS and centrifuged. The cells were re-suspended in 500 μ L of 1 \times binding buffer. 5 μ L of annexin V-FITC and 5 μ L of



propidium iodide (PI 50 mg mL⁻¹) were added and the cells were incubated in the dark at room temperature for 5 min. The fluorescence of the stained cells was analyzed by flow cytometry using FACSCalibur (BD Biosciences, Mountain View, CA). The data were analyzed using the CellQuest software (BD Biosciences).^{53,54}

4.2.4. Tubulin polymerization inhibition assay. Tubulin polymerization inhibition assay was performed using Tubulin Polymerization Assay Kit (Cytoskeleton, USA). 2 mg mL⁻¹ tubulin was dissolved in 80 mM PIPES pH 6.9, 2.0 mM MgCl₂, 0.5 mM EGTA, 1.0 mM GTP and 15% glycerol. A 96-well plate was warmed to 37 °C for 10 minutes. Serial dilutions of the test compounds were prepared. The assay components were mixed. Polymerization was monitored using a plate reader at 360 nm and emission at 420 nm at 37 °C every 1 min intervals for 60 min. A dose–response curve was generated and the IC₅₀ for tubulin polymerization inhibition was calculated.

4.2.5. Tubulin expression assay

4.2.5.1. RNA isolation. The cells were seeded in a six-well plate at a concentration of 3 × 10⁵ cells per well. After 24 h. incubation, the IC₅₀ dose of compound **4c** and the positive control **colchicine** was applied to each well for 48 h. The cells were washed, collected and centrifuged at 2500 rpm for 15 min at 4 °C. The total RNA isolation kit (RNeasy extraction kit – QIAGEN) was used to isolate RNA according to the manufacturer's instructions.

4.2.5.2. Gene expression analysis by quantitative real-time PCR. Gene expression of β-tubulin was determined using iScript One-Step RT-PCR kit with SYBR Green (BIORAD) according to the manufacturer protocol. Primers sequences used for qPCR are illustrated in (Table 5). The genes expression was normalized to β-actin as the housekeeping gene. The protocol used was cDNA synthesis for 10 min at 50 °C, iScript Reverse transcriptase inactivation for 5 min at 95 °C and PCR cycling and detection (30 to 45 cycles): 10 s at 95 °C and 30 s at 55 °C to 60 °C.

4.3. In silico studies

4.3.1. Docking studies. The docking simulation was performed to ascertain the fit of the titled compounds **3b**, **3c**, **4c**, and **5c** in the active site of the tubulin receptor. Docking was performed on AutoDock software using the Vina plugin.^{55,56} Protein structure was downloaded from PDB (PDB ID: 4O2B with a resolution of 2.30 Å). The simulation started with the preparation of protein through the removal of all the subunits



Fig. 16 Alignment between the conformation of the co-crystallized ligand (colchicine) colored in grey and the best-docked pose (colored in blue) into α–β tubulin enzyme with RMSD value equal to 0.17 Å.

except chain A (α-tubulin subunit) and chain B (β-tubulin subunit) having the **colchicine** binding site (CBS), water was removed, protonation, and addition of Kollman charges was performed, finally, the prepared enzyme, co-crystallized ligand (**colchicine**, PDB entry: LOC), and titled compounds were saved in pdbqt format. Grid box coordinates were kept at 17.30 × 66.41 × 42.90 Å, while exhaustiveness was kept at 8. The internal ligand of the enzyme was redocked to validate the method and the RMSD value was reported using the DockRMSD server.⁵⁷ The result revealed that the re-docked **colchicine** showed an RMSD value of 0.17 Å, while the pose of the redocked ligand aligned perfectly with the native ligand (Fig. 16) which revealed a validated docking procedure. After docking completion 2D and 3D images were visualized using DS client software.⁵⁸ The key amino acid residues, bonded atom or group and the interaction energies in kcal mol⁻¹ of the new compounds as well as for the co-crystallized ligand were summarized (Table S5†).

4.3.2. Physicochemical properties. The SwissADME web tool was used to estimate physicochemical and pharmacokinetics characteristics *in silico* (SwissADME: <http://www.swissadme.ch/>).⁵⁹

Table 5 Primers sequences of different genes

Gene	Primer sequence
β-Tubulin	F: 5'-TCAGCGTCTACTACAACGAGGC-3' R: 5'-GCCTGAAGAGATGTCCAAGGC-3'
β-Actin	F: 5'-CATTGCTGACAGGATGCAGAAGG-3' R: 5'-TGCTGGAAGGTGGACAGTGAGG-3'

Data availability

The authors of this article declare that and data sharing should be upon request.

Conflicts of interest

There are no conflicts to declare.



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