

Cite this: *RSC Med. Chem.*, 2025, 16, 3671

Improved anti-breast cancer activity through structural modification of fused pyran derivatives and mechanistic investigations†

K. Fabitha, ^a Anoop Kallingal, ^b Natalia Maciejewska, ^b Hiba Haneefa P. T., ^a Varun Thachan Kundil, ^c Majed Alharbi ^d and Janardhan Banothu ^{*a}

Novel fused pyran derivatives were synthesized and evaluated for anticancer activity against MCF7 (ATCC – HTB-22, breast), A549 (ATCC – CCL-185, lung), and HCT116 (ATCC – CCL-247, colorectal) cancer cell lines. Most compounds exhibited broad-spectrum activity. Among them, imidazole-containing derivatives **8a** and **8b** demonstrated potent anti-breast cancer activity with IC₅₀ values of 8.24 ± 0.19 μM and 4.22 ± 0.81 μM, respectively. Both compounds were nontoxic to MCR5 (ATCC – CCL-171, human diploid fibroblast) cell line. *In vitro* studies using MCF7 cells showed that **8a** and **8b** significantly reduced colony formation and inhibited spheroid progression. Cell cycle analysis revealed G1 arrest, a mechanism distinct from the standard drug etoposide. Furthermore, these compounds induced persistent DNA damage and both early and late apoptosis. In addition, *ex ovo* CAM assay revealed a significant reduction in new blood vessel formation over time compared to untreated controls. Network pharmacology identified EGFR, SRC, and GSK3B as potential targets, and molecular docking studies confirmed strong binding affinities to these proteins. This interaction is proposed to inhibit uncontrolled breast cancer cell growth. Additionally, the compounds exhibited favorable pharmacokinetic properties *in silico*. These findings suggest that novel fused pyran derivatives, particularly **8a** and **8b**, are promising candidates for further preclinical development as breast cancer therapeutics.

Received 19th February 2025,
Accepted 3rd May 2025

DOI: 10.1039/d5md00161g

rsc.li/medchem

1. Introduction

Cancer continues to be a major global health concern and a leading cause of mortality worldwide.¹ Among various cancer types, female breast cancer is the most prevalent and remains the primary cause of cancer-related deaths.² According to the GLOBOCAN 2020 report, approximately 2.3 million new breast cancer cases were diagnosed, with around 685 000 deaths attributed to the disease as of 2020.^{1–3} Despite significant advancements in treatment options, there remains an urgent need for novel therapeutic strategies, particularly for aggressive subtypes.^{4–7}

Recent advancements in drug development have highlighted fused pyran/chromene derivatives as promising antitumor agents.^{8–10} These compounds, found in various natural sources, exhibit significant anticancer activity against various malignancies.^{8–10} For instance, Tephrosin has shown efficacy against lung cancer through the autophagy pathway and against pancreatic cancer by generating reactive oxygen species.^{11,12} Calanone has demonstrated activity against cervical carcinoma by upregulating the p53 tumor suppressor gene,¹³ while Acronycine has been effective against ovarian, lung, and colon cancers by inducing DNA damage.¹⁴ Similarly, Seselin has the potential to fight skin cancer by inhibiting tumour-promoting mechanisms.¹⁵ Further studies on fused pyran/chromene derivatives have shown their ability to inhibit tubulin polymerization by binding to or near the colchicine-binding site, triggering apoptosis through caspase activation and inducing G2/M cell cycle arrest, ultimately leading to cancer cell death.¹⁶ Additionally, these molecules disrupt tumor vascular endothelial cell capillary tubules, indicating their potential as vascular-disrupting agents.¹⁷ EPC2407 (Crolibulin™), a potent microtubule-targeting agent, has progressed to phase I/II clinical trials for anaplastic thyroid cancer.^{8,18,19} Another lead compound,

^a Department of Chemistry, National Institute of Technology Calicut, Kozhikode – 673601, Kerala, India. E-mail: janardhan@nitc.ac.in; Tel: +0495 228 5324^b Department of Pharmaceutical Technology and Biochemistry, Faculty of Chemistry, Gdansk University of Technology, Narutowicza St 11/12, 80-233 Gdansk, Poland^c Department of Biotechnology and Microbiology, School of Lifesciences, Kamur University, Kannur – 670661, Kerala, India^d Department of Pharmaceutical Chemistry, Faculty of Pharmacy, King Abdulaziz University, Jeddah 21589, Saudi Arabia† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d5md00161g>

MX-58151, is a robust tubulin destabilizer, triggering cell apoptosis through caspase3/7 activation and nuclear fragmentation.^{20,21} Furthermore, recent reports indicate that fused pyran/chromene derivatives selectively inhibit formyl peptide receptor-1 (FPR-1), thereby blocking calcium ion (Ca^{2+}) flux and suppressing chemotaxis in human neutrophils, ultimately preventing their migration to inflammatory sites.²² These compounds have also been shown to specifically target triple-negative breast cancer cells, significantly inhibiting their proliferation while exerting minimal effects on hormone-responsive cells.²³ All these findings highlight the therapeutic potential of fused pyran/chromene derivatives in developing novel anticancer therapies.

A comprehensive literature review on fused pyran/chromene derivatives highlights their ability to inhibit cancer cells through diverse mechanisms of action. For instance, 3-amino-9-methoxy-1-aryl-1*H*-benzo[*f*]chromene-2-carbonitrile (A) demonstrated significant potency against MCF7/ADR cells.²⁴ It effectively inhibited P-gp expression, induced apoptosis, caused cell cycle arrest at various phases, disrupted mitochondrial transmembrane potential, and activated caspase 3/7, collectively promoting apoptotic cell death.²⁴ Similarly, 3-amino/3,5-diamino-1-aryl-1*H*-benzo[*f*]chromene-2-carbonitrile (B) showed a significant effect on multiple mechanisms, including induction of apoptosis, flow cytometry analysis, and regulation of apoptosis-related genes.^{25,26} Moreover, 3-amino-8-bromo/methoxy-1-aryl-1*H*-benzo[*f*]chromene-2-carbonitrile (C) displayed potent and selective cytotoxic activity against MCF7, HCT-116, and HepG-2 cell lines, targeting the c-Src kinase enzyme and exhibiting anti-invasion effects on MDA-MB-231 cells.²⁷ Furthermore, 2-amino-4-(3-nitrophenyl)-5-oxo-4,5-dihydropyrano[3,2-*c*]chromene-3-carbonitrile (D) showed a strong antiproliferative effect mediated through DNA interaction.²⁸ The structures of compounds A–D are presented in Fig. 1. In addition to their anticancer properties, fused pyran derivatives have been reported as potential anticonvulsant agents,²⁹ antidiabetic agents,³⁰ antitubercular agents,³¹ antimicrobial agents,³² and inhibitors of estrogen biosynthesis,³³ and monoamine oxidase (MAO).³⁴

Nitrogen-containing heterocycles are common drug fragments with immense importance in medicine, as they

are abundant in nature and serve as key components in numerous biologically active compounds, such as vitamins, hormones, antibiotics, *etc.*³⁵ Analysis of U.S. FDA-approved drugs reveals that around 60% of approved small-molecule drugs contain N-based heterocycles, highlighting their structural significance in drug design.³⁶ The prevalence of these compounds in pharmacology is primarily attributed to their stability and efficiency in the human body and their ability to form hydrogen bonds with DNA to enhance their therapeutic potential, particularly in cancer treatment.³⁷ Recognizing the therapeutic potential of fused pyran derivatives and as a part of our endeavour to investigate the anticancer potentials of heterocyclic small molecules,^{38–42} we recently reported the synthesis and anticancer evaluation of various fused pyran derivatives featuring N-based heterocycles.^{43,44} Notably, some compounds demonstrated significant inhibitory activity against lung, breast, and colon cancer cell lines.^{43,44} In addition, the potent compounds were identified to inhibit colony formation and various phases of the cell cycle, as well as induce DNA double-strand breaks and apoptosis in the tested cancer cells.⁴⁴ Encouraged by these results and to enhance the efficacy of the compounds, we have designed several new fused pyran derivatives, as illustrated in Fig. 2. All the designed molecules were synthesized and screened for their *in vitro* anticancer activity against MCF7, A549, and HCT116 cell lines, and the potent compounds were evaluated for their mechanism of action.

2. Results and discussion

2.1 Chemical synthesis

The three-component condensation reaction of 4-hydroxycoumarin/4-hydroxy-6-methyl-2*H*-pyran-2-one/5-hydroxy-2-(hydroxymethyl)-4*H*-pyran-4-one/cyclohexane-1,3-dione/5,5-dimethylcyclohexane-1,3-dione with N-based heterocycles substituted arylaldehydes and malononitrile in water, catalyzed by sodium fluoride (NaF), produced diverse fused pyran derivatives.⁴⁴ *In vitro* anticancer evaluation of all the compounds led to identifying potential anti-lung, anti-breast and anti-colon cancer agents (I–IV, Fig. 2). To enhance the anticancer efficacy, various substituted pyrazoles,

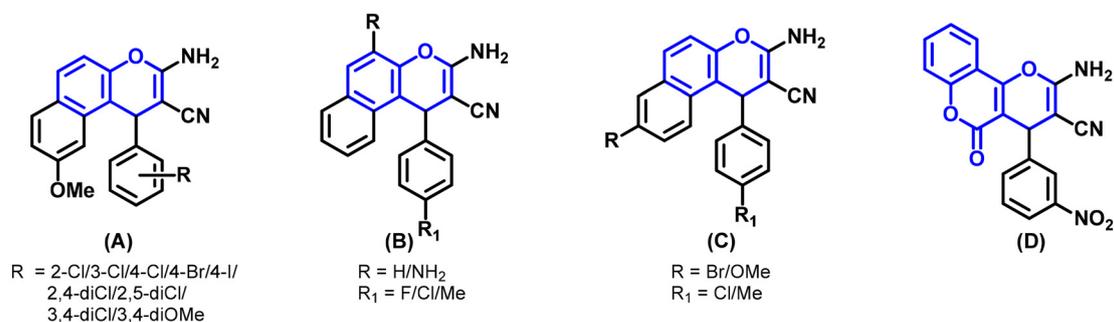


Fig. 1 Examples of fused pyran/chromene derivatives (A–D) as potential anticancer agents with multiple mechanisms of action.



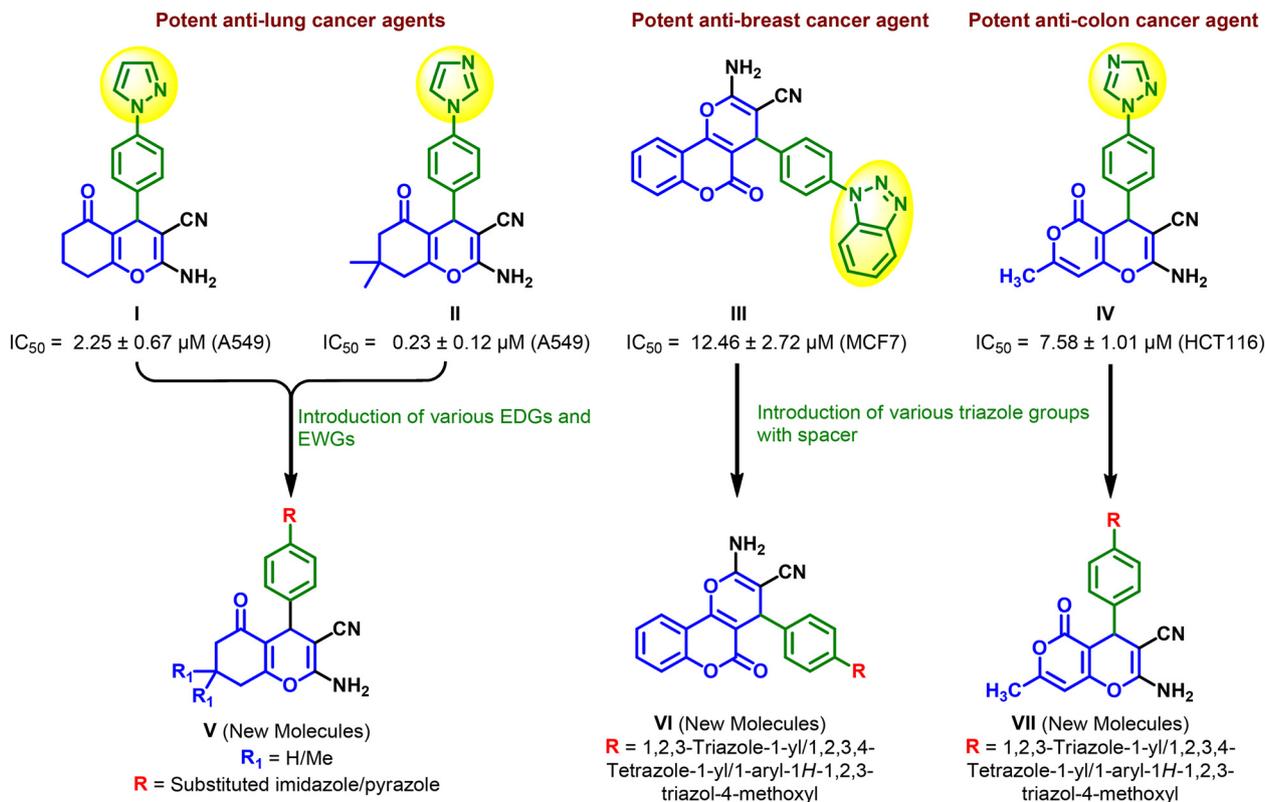
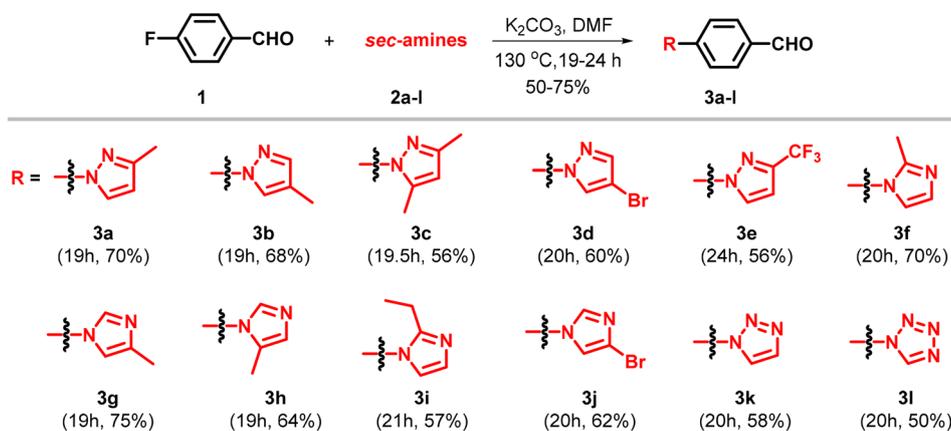


Fig. 2 The design strategy of new fused pyran derivatives from the reported potent anticancer agents.

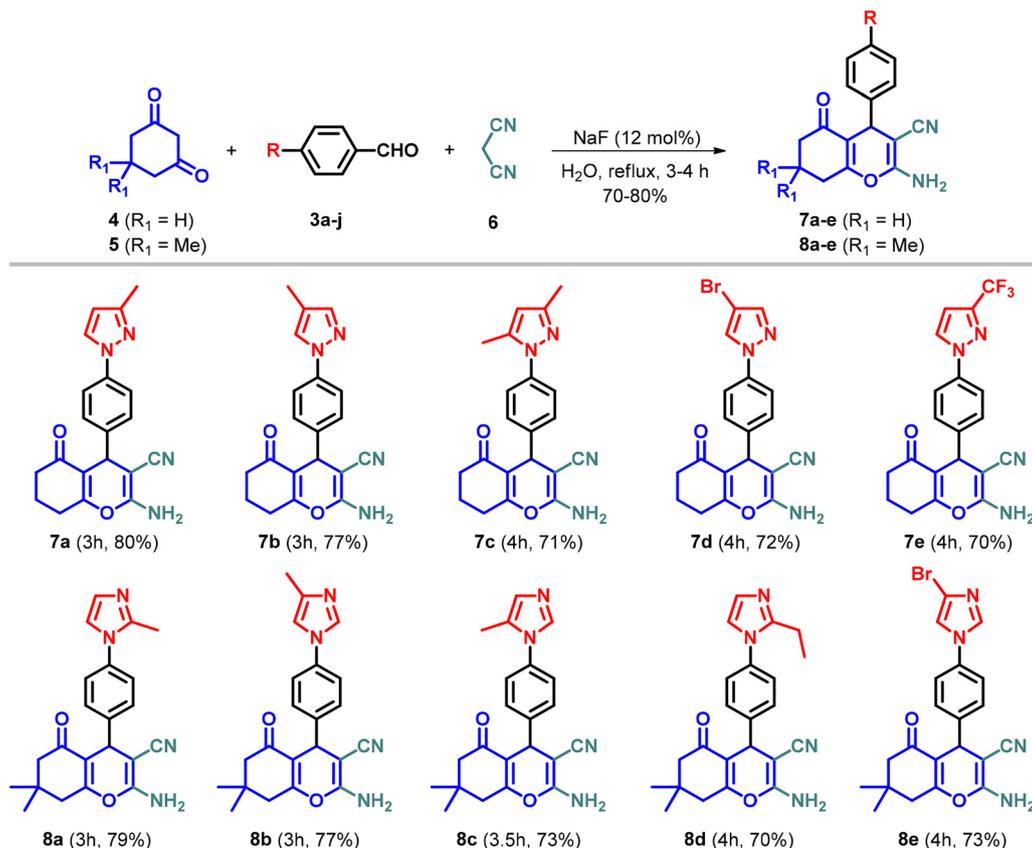
imidazoles, triazoles and tetrazoles were introduced as outlined in Schemes 2 and 4. The intermediates, N-heterocycle-substituted arylaldehydes (**3a-l**) were synthesized through S_NAr reactions of 4-fluorobenzaldehyde (**1**) with substituted secondary amines, including pyrazole (**2a-e**), imidazole (**2f-j**), 1,2,3-triazole (**2k**), and 1,2,3,4-tetrazole (**2l**), in the presence of potassium carbonate in DMF at 130 °C (Scheme 1). Substituted 1,2,3-triazole-based aldehydes (**13a-e**) were prepared by following the synthetic route depicted in Scheme 3.⁴⁵ The intermediates required, *i.e.* azide (**10**) was obtained through the classical diazotization-azidation

method. Whereas, *o*-propargylation of commercially available 4-hydroxybenzaldehyde (**11**) was achieved by treating it with propargyl bromide in the presence of anhydrous K_2CO_3 in DMF. Subsequently, a copper-catalyzed azide-alkyne cycloaddition (CuAAC) between azide (**10**) and terminal alkyne (**12**) resulted in the formation of **13a-e** in 65–75% yield.⁴⁵ Finally, the desired products (**7a-e**, **8a-e**, **15a-g**, & **17a-g**) were synthesized by reacting cyclohexane-1,3-dione (**4**), 5,5-dimethylcyclohexane-1,3-dione (**5**), 4-hydroxycoumarin (**14**), and 4-hydroxy-6-methyl-2H-pyran-2-one (**16**) individually with N-heterocycle-substituted benzaldehydes (**3a-l**)/1,2,3-

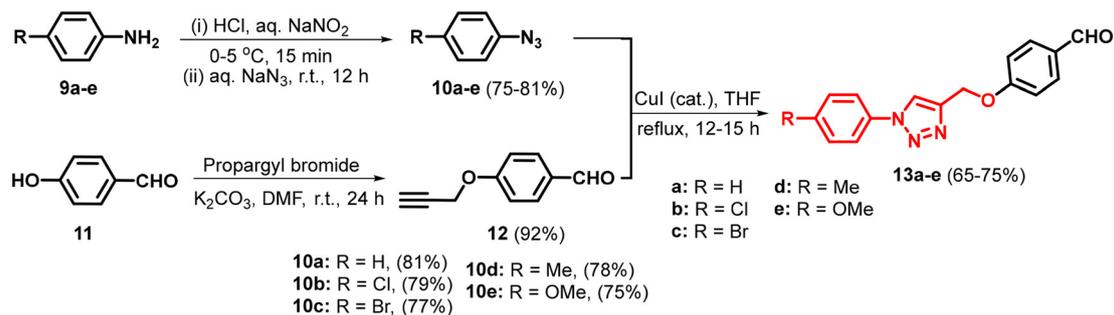


Scheme 1 Synthesis of N-based heterocycles substituted arylaldehydes (**3a-l**).





Scheme 2 Synthesis of novel N-based heterocycles embedded fused pyran derivatives (7a-e & 8a-e).



Scheme 3 Synthesis of 1,2,3-triazole substituted arylaldehydes (13a-e).

triazole substituted arylaldehydes (13a-e) and malononitrile in H₂O, using 12 mol% sodium fluoride (NaF) as a catalyst. The reactions were performed at reflux temperature, yielding 65–80% of the products within 3–4 hours.

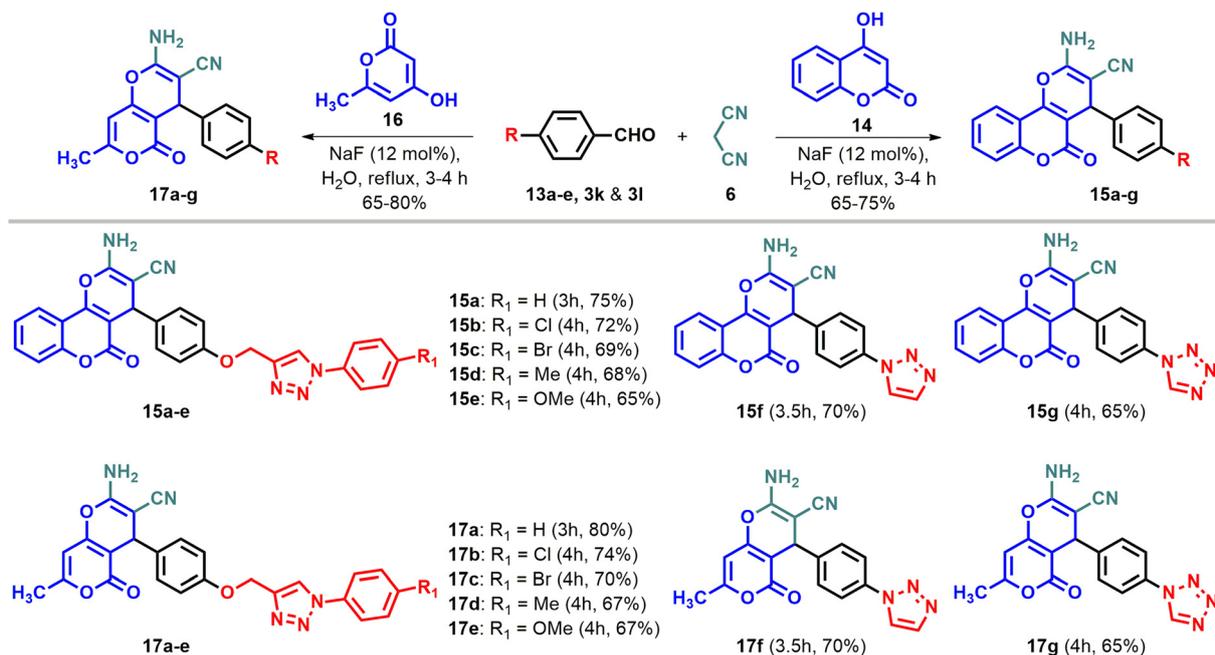
Structures of all the target compounds were confirmed by spectral studies such as FT-IR, ¹H NMR, ¹³C NMR, and LC-HRMS. From the ¹H NMR spectra, the appearance of a singlet peak at 4.10–4.60 ppm confirms the presence of the pyran-C4 proton. Moreover, the presence of a peak at 34–38 ppm in the ¹³C NMR spectra corresponding to the pyran-C4 carbon provided additional evidence of the formation of the pyran ring. Furthermore, the molecular ion peak from the mass spectra also confirmed the product formation. The

purity of all tested compounds was greater than 99.5% as determined by LC-HRMS.

2.2 Effects of fused pyran derivatives on cancer cell viability

The cytotoxic effects of all the synthesized compounds were systematically evaluated using the MTT assay across multiple cancer cell lines, such as A549 (ATCC – CCL-185, lung carcinoma), HCT116 (ATCC – CCL-247, colon carcinoma), and MCF7 (ATCC – HTB-22, breast carcinoma), and a noncancerous human diploid fibroblast cell line, MCR5 (ATCC – CCL-171). As summarized in Table 1, the most tested compounds demonstrated dose-dependent cytotoxicity





Scheme 4 Synthesis of novel fused pyran derivatives (**15a-g** & **17a-g**).

across all selected cancer cell lines, suggesting their potential as effective anti-cancer agents (Fig. 3). Determining half maximal inhibitory concentration (IC₅₀)

values revealed varying degrees of potency among the compounds. Specifically, compounds **8a** and **8b** exhibited the most significant cytotoxic activity against the MCF7

Table 1 IC₅₀ values of fused pyran derivatives against MCF7, A549, HCT116 and MRC5. Data are shown as mean ± SD, *n* = 3

Compound	Cell lines			
	MCF7 IC ₅₀ (μM)	A549	HCT116	MRC5
7a	190.47 ± 5.34	89.68 ± 4.38	139.13 ± 3.89	>200
7b	23.98 ± 2.27	147.08 ± 3.61	163.48 ± 3.14	>200
7c	22.92 ± 1.31	>200	193.79 ± 6.01	>200
7d	56.13 ± 3.79	23.56 ± 1.29	45.70 ± 0.67	>200
7e	15.03 ± 0.78	17.96 ± 0.45	72.21 ± 0.96	>200
8a	8.24 ± 0.19	111.46 ± 4.91	142.96 ± 6.35	>200
8b	4.22 ± 0.81	109.56 ± 4.94	129.16 ± 5.79	>200
8c	83.25 ± 2.11	133.38 ± 5.81	161.97 ± 6.91	>200
8d	>200	19.82 ± 0.94	136.80 ± 5.82	>200
8e	74.87 ± 0.17	30.60 ± 1.85	77.57 ± 1.58	>200
15a	>200	>200	>200	146.5 ± 0.53
15b	>200	>200	>200	>200
15c	>200	>200	>200	66.9 ± 1.6
15d	>200	>200	>200	>200
15e	>200	>200	>200	>200
15f	144.92 ± 2.75	72.86 ± 2.26	163.25 ± 5.45	156 ± 0.48
15g	>200	>200	>200	>200
17a	>200	80.51 ± 4.68	>200	>200
17b	36.06 ± 0.31	20.44 ± 1.87	47.44 ± 4.13	>200
17c	>200	>200	>200	>200
17d	>200	>200	>200	>200
17e	52.25 ± 1.34	>200	>200	>200
17f	30.32 ± 1.73	>200	>200	>200
17g	39.21 ± 2.71	164.10 ± 3.26	>200	>200
I ⁴⁴	—	2.25 ± 0.67	—	—
II ⁴⁴	—	0.23 ± 0.12	—	—
III ⁴⁴	12.46 ± 2.72	—	—	—
IV ⁴⁴	—	—	7.58 ± 1.01	—



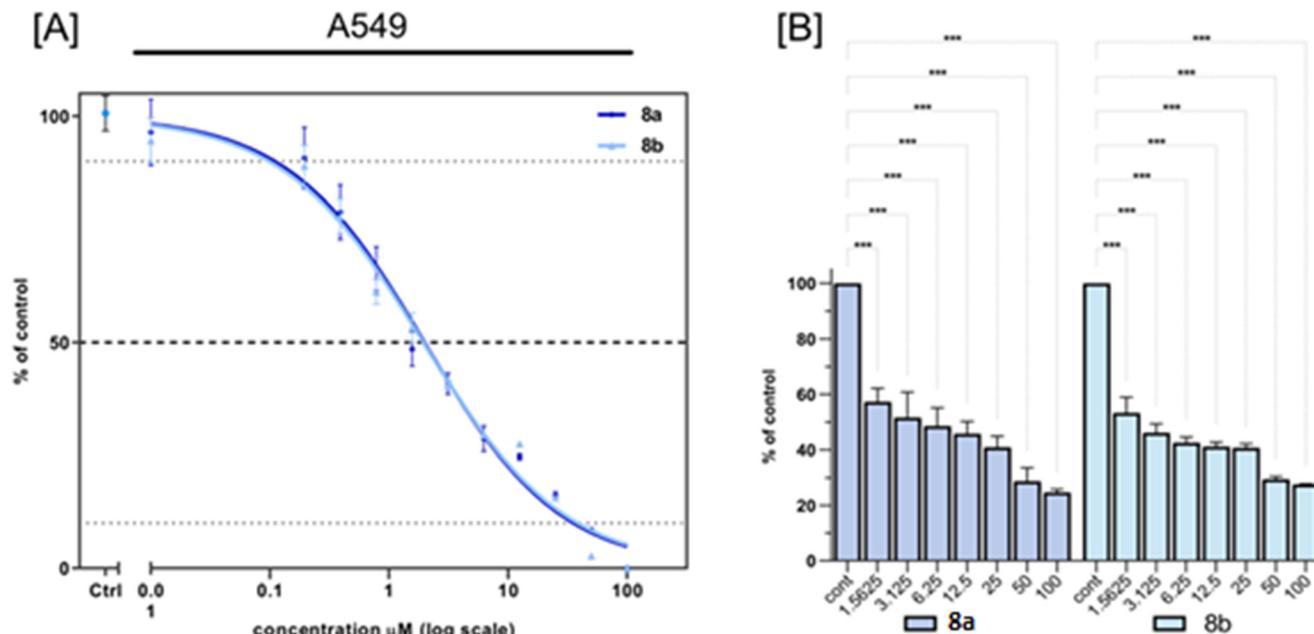


Fig. 3 IC₅₀ comparison graph of top selected compounds [A] and the dose-dependent inhibition of the viability of cancer cells under compound treatment at various concentrations (100–1.56 μM) [B]. The statistical analysis was conducted using one-way ANOVA and *post hoc* Dunnett's test. The data represent mean SD ($n = 3$), * $P \leq 0.01$, ** $P \leq 0.001$, *** $P \leq 0.0001$, and **** $P \leq 0.00001$.

cell line, with IC₅₀ values of $8.24 \pm 0.19 \mu\text{M}$ and $4.22 \pm 0.81 \mu\text{M}$, respectively. Notably, compound **8a** demonstrated a 1.5-fold higher potency, while compound **8b** exhibited a 3-fold increase in potency compared to the previously reported fused pyran derivative (**III**, Fig. 2).⁴⁴ The structure–activity relationship (SAR) study against the MCF-7 cell lines is presented in Fig. 4. These low IC₅₀ values suggest a strong inhibitory effect on MCF7 cell viability, highlighting the superior efficacy of these compounds in targeting breast cancer cells. In addition, both compounds were found to be nontoxic to MRC5 cells. The enhanced and selective cytotoxicity observed with these compounds (**8a** and **8b**) underscores their potential therapeutic relevance and justifies their selection for further mechanistic studies in MCF7 cell lines. The MTT assay confirmed the synthesized compounds' cytotoxic capabilities and provided crucial quantitative

insights into their relative effectiveness across different cancer types. These findings contribute to the ongoing efforts to develop potent anti-cancer agents and support the continued investigation of compounds **8a** and **8b** in subsequent evaluations.

2.3 Inhibitory effects of fused pyran derivatives on colony formation and spheroid formation

The data from the colony formation assays for MCF-7 cells treated with compounds **8a** and **8b** were systematically analyzed to determine the percentage of colony formation at various inhibitory concentrations (IC₁₀, IC₃₀, IC₅₀, IC₇₀, and IC₉₀). For compound **8a**, the percentage of colony formation progressively decreased from $94.3 \pm 4.04\%$ at IC₁₀ to $5.67 \pm 4.04\%$ at IC₉₀, demonstrating a clear dose-dependent response. Similarly, for compound **8b**, the percentage of

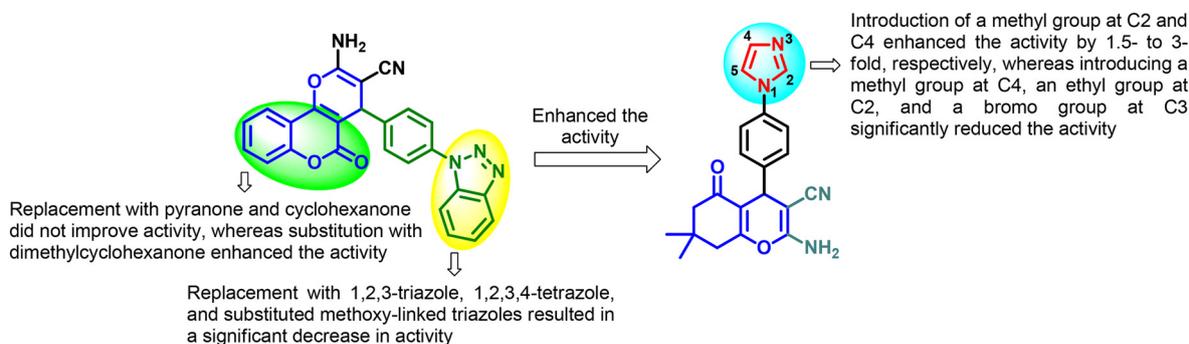


Fig. 4 Structure–activity relationship (SAR) study of fused pyran derivatives against MCF-7 cell lines.



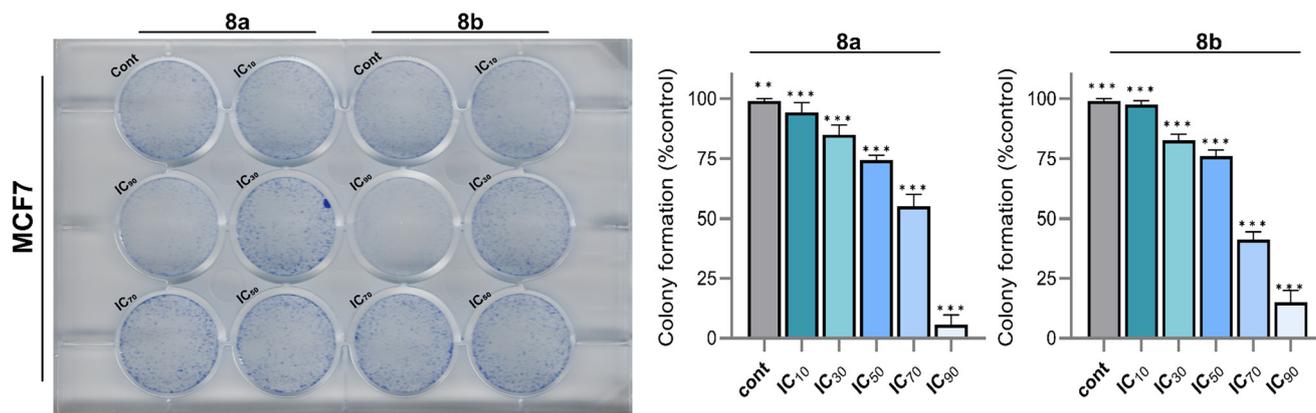


Fig. 5 Compounds **8a** and **8b** colony formation ability on MCF7. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

colony formation declined from $97.7 \pm 1.53\%$ at IC_{10} to $15 \pm 5\%$ at IC_{90} . Intermediate inhibitory concentrations also showed marked reductions, with IC_{50} values of $74.3 \pm 2.08\%$ for compound **8a** and $76 \pm 2.65\%$ for compound **8b**, reflecting their potential to impair cellular reproductive viability Fig. 5. Colony formation assays serve as a critical tool in cancer research, providing insights into cancer cells long-term survival and proliferative potential under various treatment conditions. This assay uniquely measures the ability of single cells to grow into colonies over time, making it a gold standard for assessing the cytotoxic and anti-proliferative effects of anticancer agents. By evaluating the

percentage of colony formation across a range of concentrations, the assay provides a robust quantitative assessment of a compound's efficacy in impairing clonogenic survival, a hallmark of tumorigenicity. The sharp reductions in colony formation observed with compounds **8a** and **8b** underscore their potential therapeutic relevance, as they effectively inhibit the colony-forming ability of MCF-7 cells, a widely used model for breast cancer research. The results indicate that both compounds exhibit significant anti-proliferative activity, with compound **8a** demonstrating a steeper colony formation decline at higher concentrations compared to compound **8b**. These findings are particularly

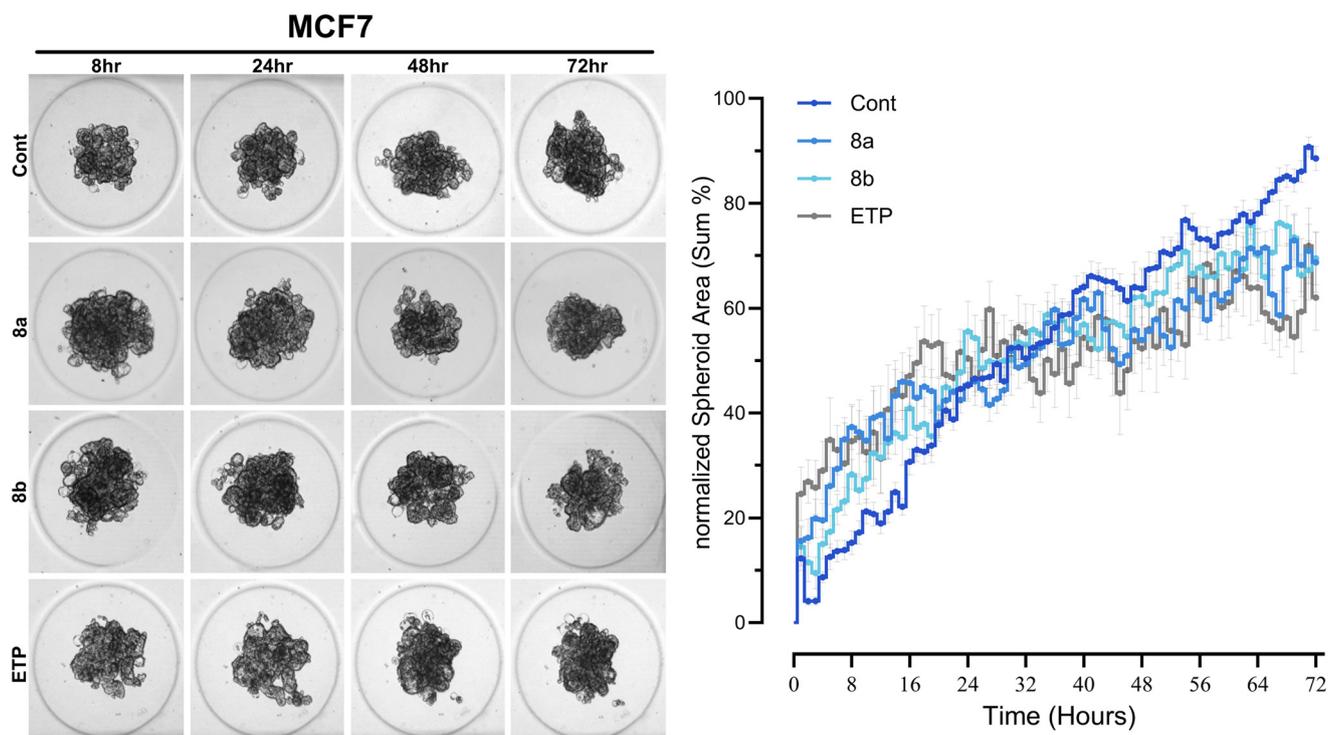


Fig. 6 Effects of **8a** and **8b** on spheroid formation and growth in MCF7 cell lines over 72 hours. Data are presented as mean \pm standard deviation from three independent experiments.



significant as the ability to inhibit colony formation reflects a compound's potential to prevent tumor regrowth and metastasis, critical factors in improving cancer treatment outcomes.

The spheroid formation analysis has emerged as a critical *in vitro* model for cancer drug discovery due to its ability to closely mimic the three-dimensional (3D) architecture and the microenvironment of solid tumors. Unlike traditional two-dimensional (2D) monolayer cultures, spheroids replicate key tumor characteristics such as cell-cell and cell-matrix interactions, oxygen and nutrient gradients, and resistance to chemotherapeutic agents. This makes spheroid models highly suitable for evaluating the efficacy of candidate compounds, as they provide a more physiologically relevant environment to assess tumor progression, cellular proliferation, and drug penetration. By monitoring spheroid dynamics, researchers can obtain valuable insights into the anti-tumor potential of new compounds and better predict their *in vivo* efficacy. In this study, compounds **8a** and **8b** were evaluated for their anti-tumor activity using MCF7 spheroids over a treatment period of 72 hours. MCF7 cells, a well-established breast cancer cell line, readily form compact spheroids that allow for robust monitoring of growth and progression under treatment conditions. The results demonstrated a significant inhibitory effect of compounds **8a** and **8b** on spheroid progression. In untreated control spheroids, a steady and measurable increase in spheroid size was observed over time, which indicates continuous cellular

proliferation and tumor-like growth. Conversely, spheroids treated with compounds **8a** and **8b** showed a marked reduction in size throughout the experimental duration, signifying an inhibition of cellular growth and potential cytotoxic effects (Fig. 6). Specifically, after 72 hours of treatment, spheroids exposed to compound **8a** exhibited a pronounced reduction in size compared to untreated controls, indicating a robust suppression of cellular proliferation and spheroid expansion. Compound **8b** produced a similar effect, demonstrating its ability to impede spheroid growth over the same period significantly. The consistent reduction in spheroid size under both treatment conditions highlights the efficacy of these compounds in targeting tumor-like structures and inhibiting growth in a 3D environment. Furthermore, the observed inhibition suggests that compounds **8a** and **8b** may disrupt key processes required for tumor progression, such as cell division, nutrient uptake, or survival signaling within the spheroid.

2.4 Impact of the fused pyran derivatives on the cancer cell cycle

The cell cycle distribution of MCF7 cells treated with compounds **8a** and **8b** at their IC_{50} concentrations was analyzed using flow cytometry following propidium iodide staining, with Etoposide (ETP) at 10 μ M as a positive control and 0.1% DMSO as the vehicle control. The results revealed significant alterations in the cell cycle profile at both 24 and

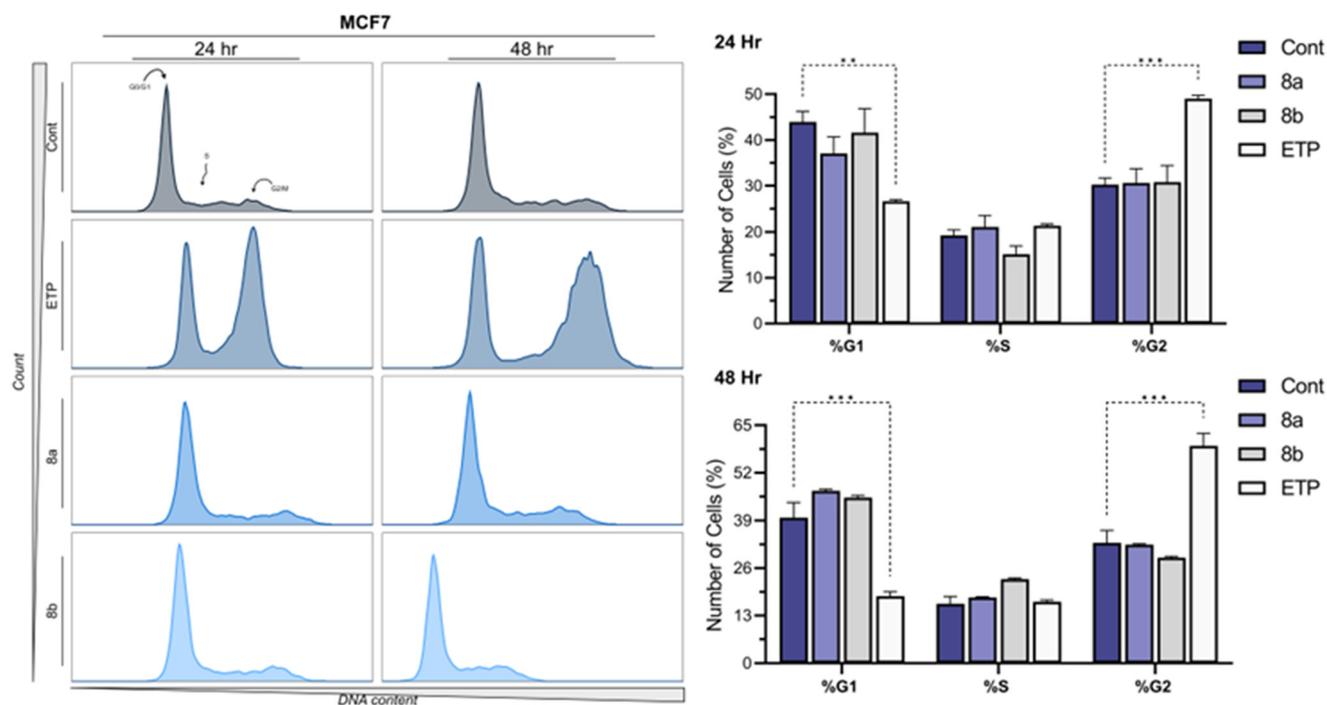


Fig. 7 Flow cytometry analysis and quantitative evaluation of cell cycle distribution after 24 and 48 hours of treatment with the compounds **8a** and **8b** in the MCF7 cell line. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, $*P \leq 0.05$, $**P \leq 0.01$, $***P \leq 0.001$.



48 hours of incubation. After 24 hours, compound **8a** induced a modest decrease in the percentage of cells in the G1 phase ($37.08 \pm 5.07\%$) compared to the control ($43.93 \pm 3.26\%$), coupled with an increase in the S phase population ($21.11 \pm 3.39\%$) and a slight shift in the G2 phase population ($30.63 \pm 4.35\%$). Similarly, compound **8b** caused a reduction in G1 phase cells ($41.59 \pm 7.44\%$) with a concurrent decrease in the S phase population ($15.11 \pm 2.59\%$) and a marginal increase in G2 phase cells ($30.79 \pm 5.11\%$). In contrast, ETP treatment resulted in a pronounced G2/M phase arrest ($49.03 \pm 0.99\%$), with a significant reduction in the G1 phase population ($26.61 \pm 0.54\%$), consistent with its known mechanism as a topoisomerase II inhibitor. After 48 hours, the impact of both compounds became more distinct. Compound **8a** prominently increased the G1 phase population ($47.08 \pm 0.6\%$) compared to the control ($39.82 \pm 5.78\%$), suggesting a potential G1 phase arrest. Meanwhile, **8b** showed a similar trend with an elevated G1 phase population (45.22

$\pm 0.82\%$) and a reduced G2 phase population ($28.83 \pm 0.52\%$). These observations contrasted sharply with ETP, which induced a robust G2/M phase arrest ($59.36 \pm 4.79\%$) and a significant reduction in the G1 phase population ($18.3 \pm 1.67\%$). The control cells maintained relatively consistent distribution across all phases at both time points, reflecting the typical cell cycle dynamics of untreated MCF7 cells Fig. 7. Studying the cell cycle is essential in cancer drug research because abnormalities in its regulation are a key feature of cancer. Targeting specific cell cycle phases allows new drugs to stop cancer cells from dividing while minimizing effects on normal cells. In this study, compounds **8a** and **8b** showed their ability to alter the cell cycle of MCF7 cells, mainly by causing G1 arrest, suggesting their potential as effective cancer treatments. Comparing these effects to those of Etoposide highlights the different ways anticancer drugs work. These results set the stage for more research to understand the molecular pathways behind these changes.

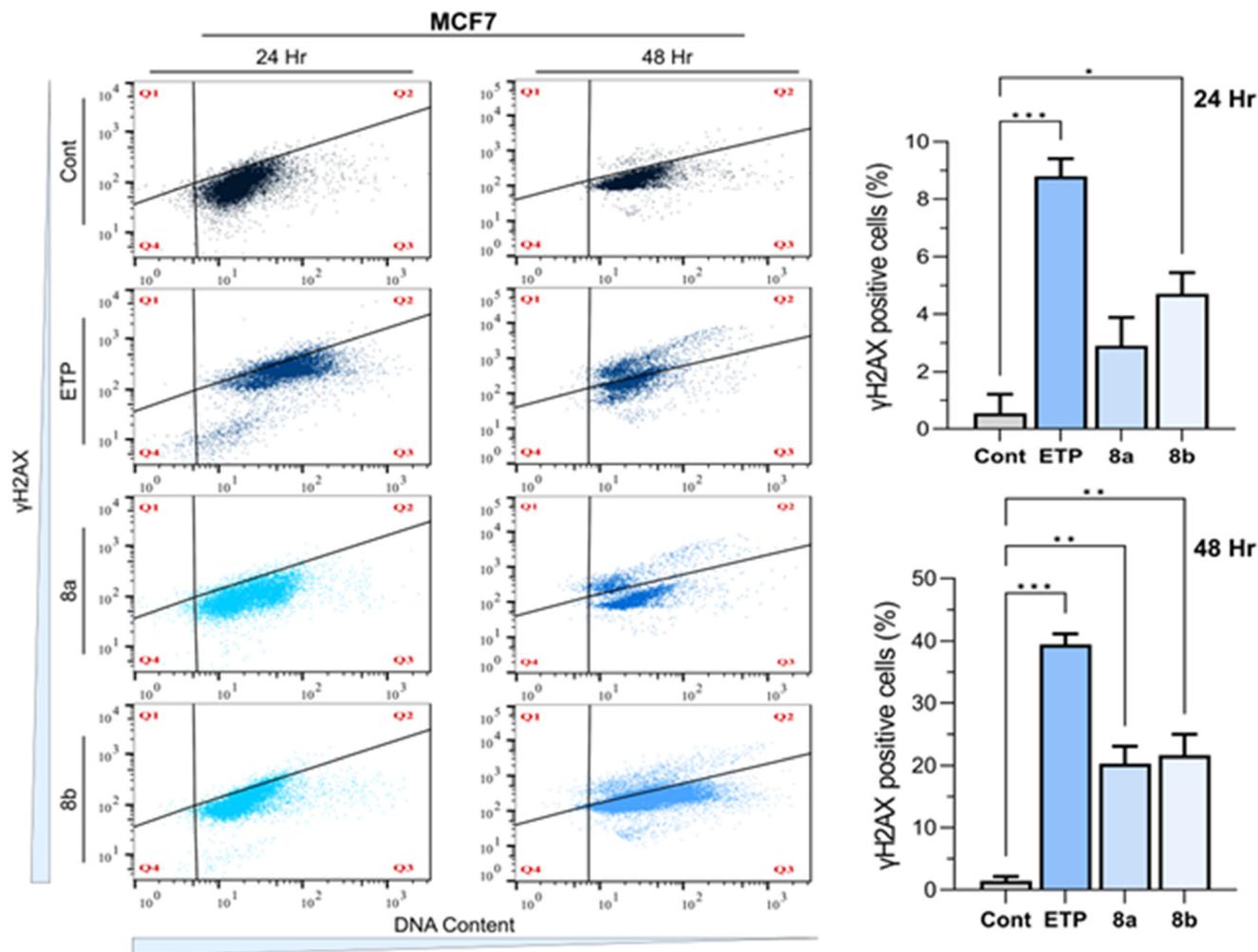


Fig. 8 Flow cytometry analysis and quantitative assessment of γ H2AX expression following 24 and 48 hours of treatment with the respective compounds in MCF7 cell lines. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.



2.5 Ability of the fused pyran derivatives to induce DNA double-strand breaks (DSBs)

γ H2AX labelling, a widely recognized marker for double-strand breaks (DSBs), pivotal role in evaluating DNA damage in cancer cells treated with anticancer compounds. In the present study, γ H2AX labelling was utilized to assess DNA damage in the MCF7 cell line following treatment with compounds **8a** and **8b** at their IC_{50} concentrations for 24 and 48 hours, alongside etoposide (ETP, a positive control at 10 μ M) and 0.1% DMSO as a vehicle control. Flow cytometry analysis revealed distinct patterns of γ H2AX expression that correlated with treatment duration and the nature of the compounds, providing insights into the potential of these compounds as DNA damage-inducing agents. At 24 hours, the control group showed minimal γ H2AX labelling (0.56 ± 0.66), indicating negligible baseline DNA damage. In contrast, etoposide treatment markedly increased γ H2AX levels (8.8 ± 0.62), consistent with its well-documented role in inducing DSBs *via* topoisomerase II inhibition. Both compounds **8a** and **8b** exhibited significant DNA damage induction compared to the control, with γ H2AX levels of 2.89 ± 0.99 and 4.71 ± 0.75 , respectively. Compound **8b** elicited a stronger γ H2AX response than compound **8a**, suggesting higher potency in inducing DSBs within the same incubation period. A more pronounced difference was observed at 48 hours, highlighting the time-dependent effects of the treatments. γ H2AX levels in the control group were slightly elevated (1.44 ± 0.68), likely due to natural cellular processes over extended incubation. Etoposide demonstrated a dramatic increase in γ H2AX expression (39.44 ± 1.69), consistent with sustained and severe DNA damage. Similarly, Compounds **8a** and **8b** showed substantial γ H2AX labelling (20.3 ± 2.74 and 21.62 ± 3.38 , respectively), further indicating

the ability of these compounds to induce persistent DNA damage. Compound **8b** again exhibited slightly higher activity than compound **8a**, though the difference was not statistically significant within the margin of experimental variability Fig. 8. The observed γ H2AX responses for compounds **8a** and **8b** suggest their potential mechanisms of action may involve the induction of DNA damage, a hallmark of effective chemotherapeutics. Furthermore, the time-dependent increase in γ H2AX expression highlights the sustained activity of these compounds, which may contribute to their cytotoxic effects against cancer cells. These results warrant further investigation into the detailed molecular pathways and potential therapeutic applications of compounds **8a** and **8b**.

2.6 Efficacy of fused pyran derivatives in inducing apoptosis in MCF7 cell lines

Apoptosis, a programmed cell death mechanism, pivotal in cancer drug discovery. It is often dysregulated in cancer cells, allowing them to evade cell death and proliferate uncontrollably. Compounds that can restore or induce apoptosis selectively in cancer cells are of significant interest in developing of anticancer therapies. This study assessed the apoptotic effects of compounds **8a** and **8b** in MCF7 breast cancer cells, using annexin V staining and flow cytometry at their IC_{50} concentrations over 24 and 48 hour incubation periods. The results were compared to etoposide (10 μ M) as a positive control and 0.1% DMSO as the untreated control. At 24 hours, etoposide significantly increased late apoptotic cells to $12.25 \pm 0.08\%$, compared to $7.58 \pm 0.03\%$ in the control group, reflecting its potent pro-apoptotic effect. Early apoptotic cells were also elevated in the etoposide group ($6.42 \pm 0.27\%$) relative to the control ($3.15 \pm 0.12\%$).

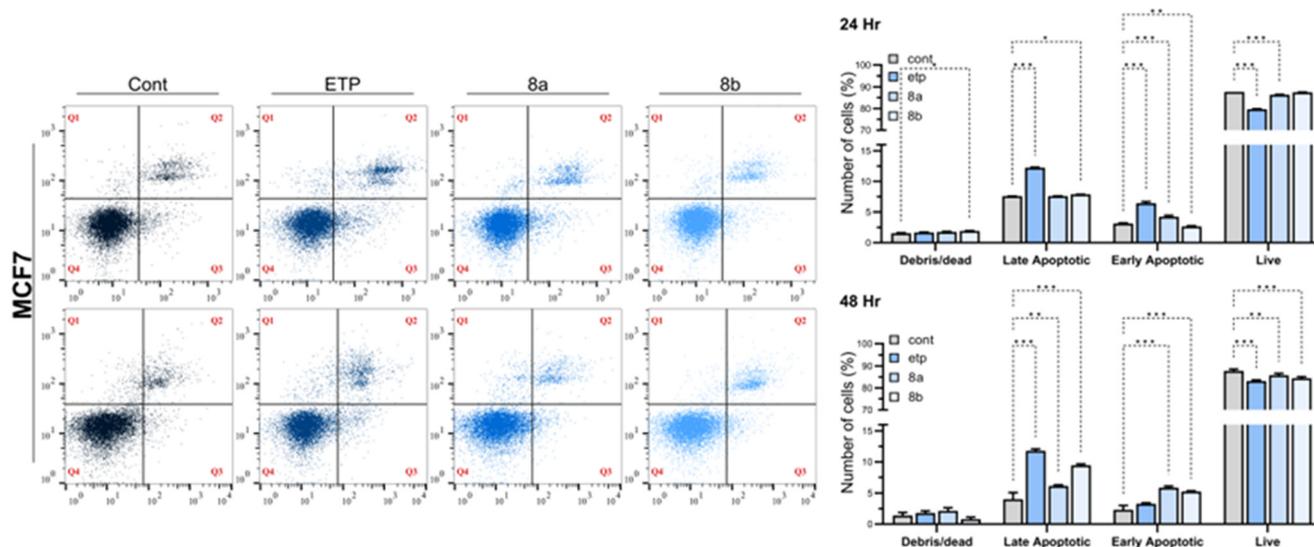


Fig. 9 Flow cytogram and quantitative assessment of apoptotic cells following 24 and 48 hours of treatment with compounds **8a** and **8b** in the MCF7 cell line. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.



Compound **8a** showed a modest increase in early apoptosis ($4.27 \pm 0.20\%$) and late apoptosis ($7.6 \pm 0.01\%$) compared to the control, while compound **8b** exhibited a slight increase in late apoptosis ($7.87 \pm 0.02\%$) but a decrease in early apoptosis ($2.66 \pm 0.12\%$). Both compounds preserved a high percentage of live cells, with $86.4 \pm 0.15\%$ and $87.55 \pm 0.08\%$ for **8a** and **8b**, respectively, compared to $87.7 \pm 1.03\%$ in the control group. After 48 hours, the apoptotic effects became more pronounced. Etoposide treatment led to a substantial increase in late apoptotic cells ($11.8 \pm 0.29\%$) and a modest rise in early apoptotic cells ($3.28 \pm 0.16\%$), with live cells reduced to $83.15 \pm 0.50\%$. Compound **8a** markedly increased early apoptosis ($5.89 \pm 0.20\%$) and late apoptosis ($6.17 \pm 0.13\%$), indicating a time-dependent enhancement of its proapoptotic activity. Similarly, compound **8b** showed an increase in both early apoptosis ($5.27 \pm 0.13\%$) and late apoptosis ($9.46 \pm 0.22\%$), with fewer live cells ($84.45 \pm 0.64\%$) compared to the 24 hour time point. Interestingly, compound **8b** also reduced debris/dead cells ($0.82 \pm 0.30\%$), suggesting

it may limit necrosis while inducing apoptosis more effectively over time Fig. 9. These results highlight the potential of compounds **8a** and **8b** to modulate apoptotic pathways in MCF7 cells. The apoptotic response elicited by both compounds was time-dependent, with increased efficacy observed at 48 hours. While etoposide demonstrated the highest apoptotic activity, the observed effects of **8a** and **8b**, particularly in increasing early apoptotic populations, underscore their therapeutic potential. Mechanistically, the ability of these compounds to induce apoptosis while maintaining a significant proportion of live cells may indicate selective targeting of cancer cells without widespread cytotoxicity.

2.7 Effect of fused pyran derivatives on angiogenesis

To evaluate the anti-angiogenic potential of compounds **8a** and **8b**, an *ex ovo* chorioallantoic membrane (CAM) assay was performed. Angiogenesis, the formation of new blood vessels

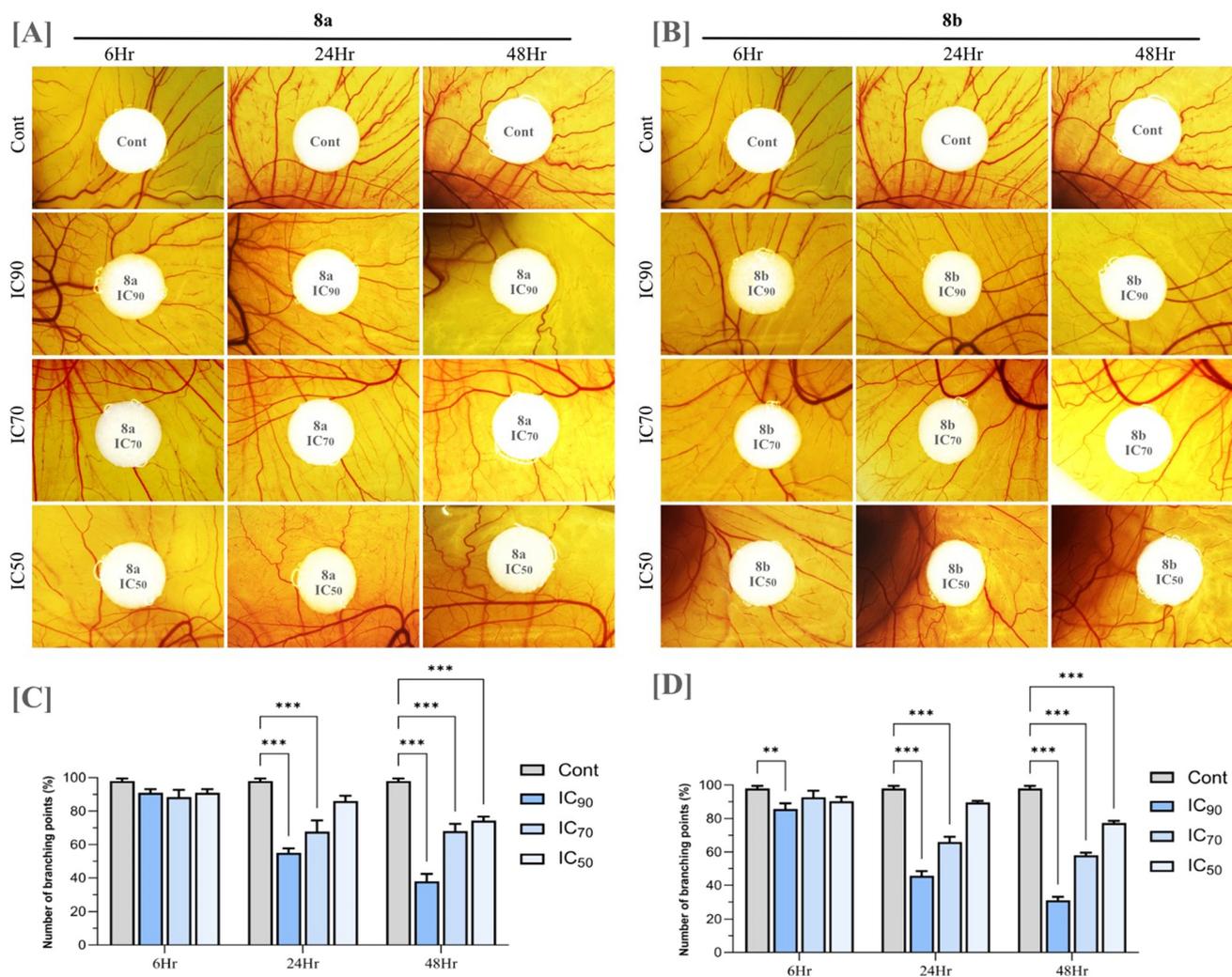


Fig. 10 Photomicrograph of CAM vascular area around the drug applied site (A) compound **8a** (B) compound **8b** (C and D) quantification of angiogenesis based on the number of vascular branching points. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 5$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.



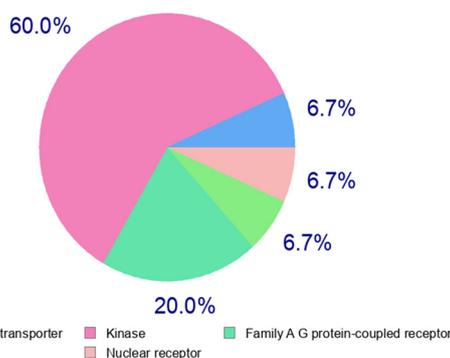
from pre-existing vasculature, is a fundamental process in both physiological and pathological conditions. In the context of cancer, angiogenesis plays a critical role in tumor development and progression by supplying oxygen and essential nutrients to rapidly growing tumor cells. It also facilitates metastatic spread by providing a conduit for tumor cells to enter the circulation. The CAM assay is a widely used *in vivo* model for studying angiogenesis due to its highly vascularized nature and ease of manipulation. The *ex ovo* version allows for direct visualization and quantification of blood vessel growth in response to experimental treatments, making it an ideal platform to assess the impact of potential anti-angiogenic compounds. Upon treatment with compounds **8a** and **8b**, a significant reduction in new blood vessel formation was observed over time compared to untreated controls. This inhibitory effect was found to be concentration-dependent, with higher concentrations resulting in a more pronounced suppression of vascular growth (Fig. 10). The treated CAMs exhibited thin, underdeveloped vasculature, suggesting that both

compounds effectively hinder angiogenic activity. While anti-angiogenic effects are not a universal characteristic of all anticancer agents and may in some cases be attributed to off-target actions, the ability to disrupt angiogenesis can be a valuable therapeutic asset. By limiting tumor vascularization, such compounds may contribute to reduced tumor growth and metastatic potential. The observed anti-angiogenic properties of compounds **8a** and **8b** enhance their overall anticancer profile and support further investigation into their mechanisms of action and therapeutic potential.

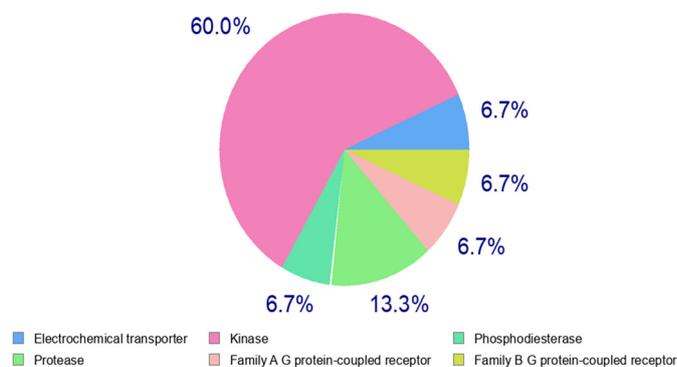
2.8 Network pharmacological analysis of fused pyran derivatives: identifying targets and biological mechanisms

Over the past decade, drug discovery has followed a “one drug, one disease” paradigm, where each drug is designed to target a specific disease. The action of drugs can be understood through the “lock and key” model, in which a drug acts like a selective key that binds to its target, or “lock”. However, drug effectiveness can improve when a

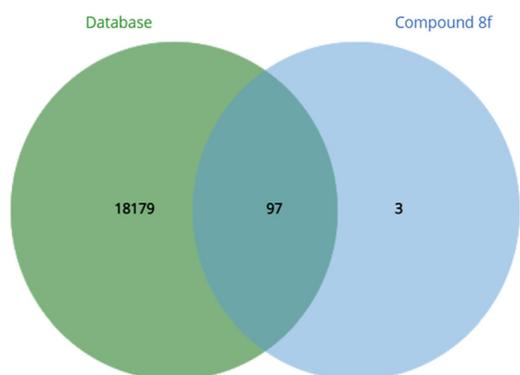
A (8a)



B (8b)



C (8a)



D (8b)

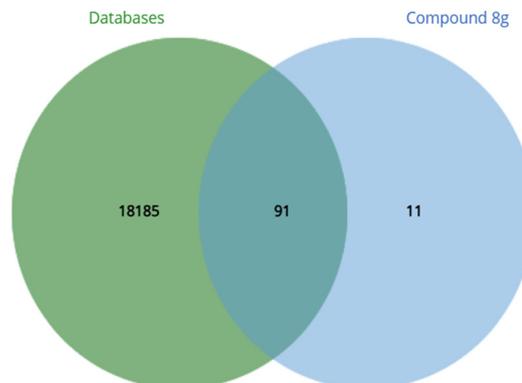


Fig. 11 A and B represent the overview of the Swiss target prediction of compounds **8a** and **8b**. C and D represent the venn diagram of common targets in each set.



single key fits multiple locks, allowing it to impact multiple targets. Viewing drug action through the lens of network biology may provide insights into how drugs bind to different targets. In 2017, Hopkins introduced a novel concept of network pharmacology to identify various drug targets using the available biomedical data in the postgenomic era, systems biology, and polypharmacology. The emergence of network pharmacology helps to examine the multi-pathway interactions in cancer (how different pathways interact and influence each other) and helps the researchers to understand drug-response signature datasets, which contain information on how cancer cells respond to various drugs and thereby decode more complex mechanism of drug action.^{46–49} Therefore, we applied a network pharmacology approach to identify the targets of fused pyran derivatives against lung, breast, and colorectal cancer. Subsequently, molecular docking was performed on these identified targets followed by *in vitro* analysis to determine whether the predicted mechanisms from the network pharmacology and molecular docking studies are accurate in real biological systems. The study is framed as a three-phase workflow stated below: (i) identification of the potential targets of compound **8a** and **8b** based on its association with breast cancer through retrieval from databases; (ii) establishment of the core targets based on the interaction relationships by network analysis; and (iii) validation of the potential targets by molecular docking verification and *in vitro* assessment.

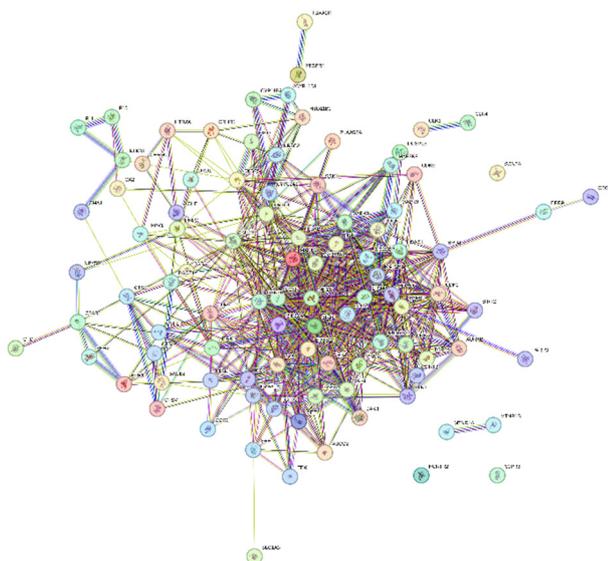
2.8.1 Identification of potential targets. Swiss target prediction was used to identify potential targets for compounds **8a** and **8b**. For compound **8a**, approximately 100 likely protein targets were predicted, with 60% being kinases, 20% family A G protein receptors, 6.7% nuclear receptors, and 6.7% electrochemical transporters Fig. 11A. For

compound **8b**, around 101 potential protein targets were identified, with 60% being kinases, 13% phosphodiesterases, 6.7% electrochemical transporters, 6.7% family B G protein receptors, 6.7% family A G protein receptors, and another 6.7% kinases Fig. 11B.

A comprehensive database search was conducted for breast cancer across multiple resources. For breast cancer, data from Malacards (18 253 records), Disginet (16 200 records), and Omim (250 records) were merged, yielding 18 276 unique gene targets after removing duplicates. To identify co-relative targets specific to each compound, a Venn diagram was constructed using the Venny website.⁵⁰ This approach helped pinpoint common targets across each compound's dataset, revealing their specificity for breast cancer. Notably, compound **8a** was found to have 97 common targets for breast cancer, while compound **8b** shared 91 common targets for breast cancer. These identified targets, considered potential anticancer candidates, were selected for further analysis (Fig. 11C and D).

2.8.2 Network interaction and its ranking. The identifiable targets of compounds associated with the respective cancer were introduced into the STRING web server to obtain the functional protein–protein association network. The protein–protein interaction (PPI) network of these anti-cancer targets was constructed. The identified 97 targets of compound **8a** and 91 targets of compound **8b** associated with breast cancer were introduced to obtain the functional protein–protein association network. For **8a**, STRING database identified 97 nodes and 578 edges, representing a network of interconnected genes (Fig. 12A). The top ten hub genes, ranked by the degree method through the CytoHubba plug-in in Cytoscape, were EGFR, GSK3B, MTOR, PTGS2, PIK3CA, PARP1, EP300, ERBB2, MAPK1, CREBBP

8a (A)



8b (B)

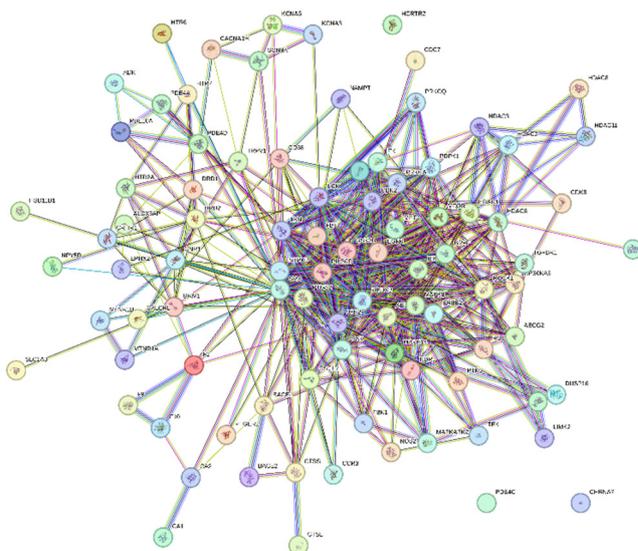


Fig. 12 STRING database analysis of compounds **8a** and **8b**.



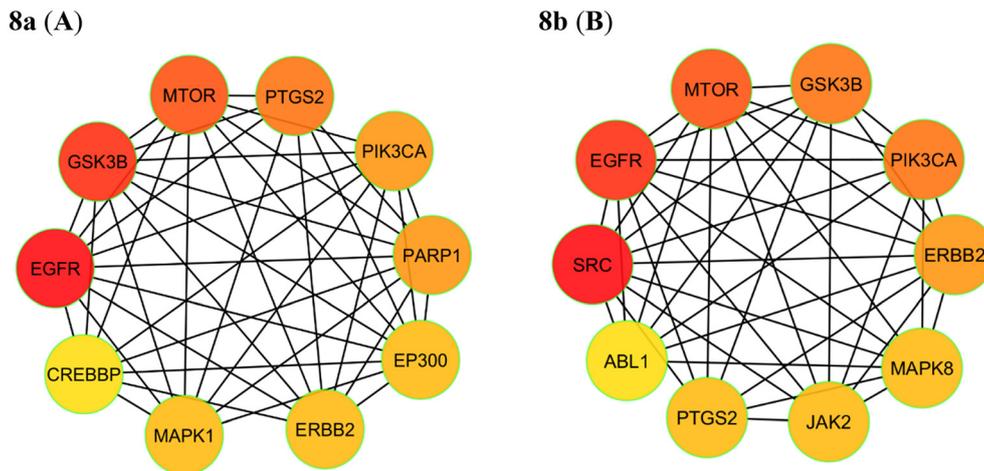


Fig. 13 Hub genes are coloured and sized according to their degree from red to yellow (A) **8a** and (B) **8b**.

(Fig. 13A). These genes, with high degree values, highlight their potential significance in the development of breast cancer. For **8b** string database identified 91 nodes and 500 edges, representing a network of interconnected genes (Fig. 12B). The top ten hub genes, ranked by the degree method through the CytoHubba plug-in in Cytoscape, were SRC, EGFR, MTOR, GSK3B, PIK3CA, ERBB2, MAPK8, JAK2, PTGS2, ABL1 (Fig. 12B).

2.8.3 Molecular docking studies. The top two hub genes identified for compound **8a** were EGFR and GSK3B, while for compound **8b**, they were SRC and EGFR. Both compounds were imported into Schrodinger Maestro and prepared to generate all possible 3D conformations. The 3D protein structures of EGFR (PDB ID: 4I23), GSK3B (PDB ID: 4ACC), and SRC (PDB ID: 3QLF) were retrieved from the PDB database and processed using the Protein Preparation Wizard. XP docking was then performed for these targets with both compounds.

EGFR is a receptor tyrosine kinase, and its crystal structure includes a co-crystallized ligand, dacomitinib. The coordinates (-0.26, -52.67, -22.5) of dacomitinib were used as the center of the grid for docking the compounds. For GSK3B, a serine/threonine protein kinase, the crystal structure is a homodimer, and chain A was selected for the study. The structure features a co-crystallized inhibitor, 7YG, with coordinates (17.46, 18.83, 10.69) used to define the grid center. Similarly, SRC, a protein tyrosine kinase, also forms a homodimer, and chain A was chosen for the analysis. The crystal structure of SRC is co-crystallized with the inhibitor pyrazolopyrimidine 5, and its coordinates (6.22, -33.92, -6.7) were used for grid generation.

The docking analysis of compounds **8a** and **8b** against various target proteins revealed differences in their binding affinities. For compound **8a**, docking with EGFR yielded a docking score of $-5.42 \text{ kcal mol}^{-1}$ and a binding energy of $-53.69 \text{ kcal mol}^{-1}$ (Fig. 14A), with interactions predominantly through hydrophobic contacts. When docked with GSK3B, **8a** showed a docking score of $-5.99 \text{ kcal mol}^{-1}$

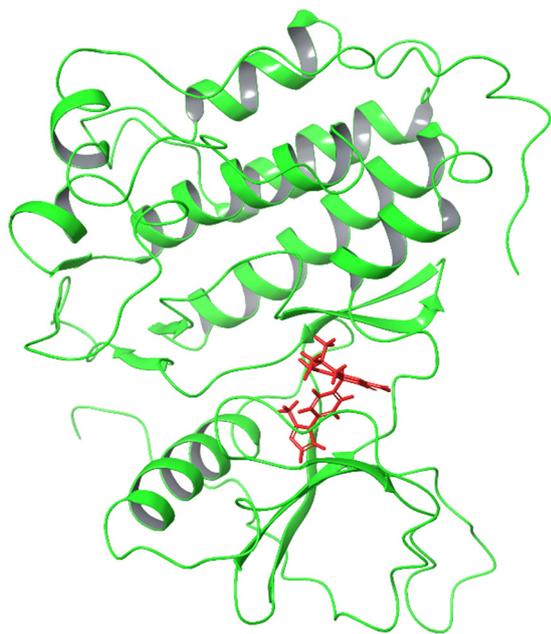
and a binding energy of $-40.05 \text{ kcal mol}^{-1}$, forming hydrogen bonds with Val135 (bond length: 2.35 Å; D-H...A angle: 177.1°) (Fig. 14B). In contrast, compound **8b** exhibited a docking score of $-5.51 \text{ kcal mol}^{-1}$ and a binding energy of $-64.77 \text{ kcal mol}^{-1}$ against SRC, forming one hydrogen bond (bond length: 2.07 Å; D-H...A angle: 141.9°) and a salt bridge (distance: 2.94 Å) with Asp348, along with a second hydrogen bond (bond length: 2.07 Å; D-H...A angle: 140.4°) with Glu339 (Fig. 15A). When docked with EGFR, **8b** showed a docking score of $-4.66 \text{ kcal mol}^{-1}$ and a binding energy of $-41.54 \text{ kcal mol}^{-1}$, forming a hydrogen bond (bond length: 2.07 Å; D-H...A angle: 129.7°) with Leu718 (Fig. 15B). The docking scores, binding energies, interacting residues and free energy (ΔG) parameters for compounds **8a** and **8b** are summarized in Tables 2 and 3.

Breast cancer is a heterogeneous disease characterized by diverse pathological features and clinical implications.⁵¹ The subtypes of breast cancer, including luminal A, luminal B, HER2-enriched, and triple-negative, exhibit distinct gene expression profiles and treatment responses.⁵² Network pharmacology was employed to identify key target proteins in breast cancer, followed by molecular docking studies to assess the binding affinity of the synthesized compounds to these targets.^{46,47} This approach resulted in the identification of epidermal growth factor receptor (EGFR), proto-oncogene tyrosine-protein kinase Src (SRC), and glycogen synthase kinase 3 beta (GSK3B) as targets for docking studies. These proteins play critical roles in breast cancer development and progression. Molecular docking simulations conducted on EGFR, SRC, and GSK3B demonstrated favorable binding affinity with fused pyran derivatives. This observation suggests that the compounds may interact with these proteins at their active sites, potentially resulting in the inhibition of their oncogenic functions.

EGFR is a receptor tyrosine kinase that initiates intracellular signaling cascades upon binding to its



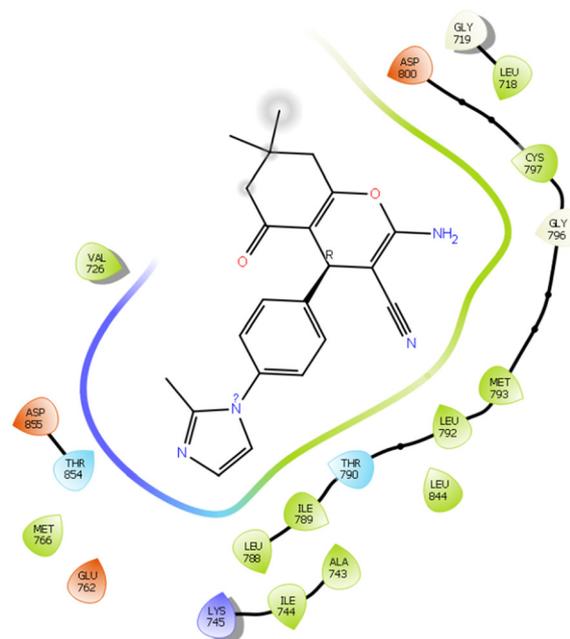
(A)



(C)



(B)



(D)

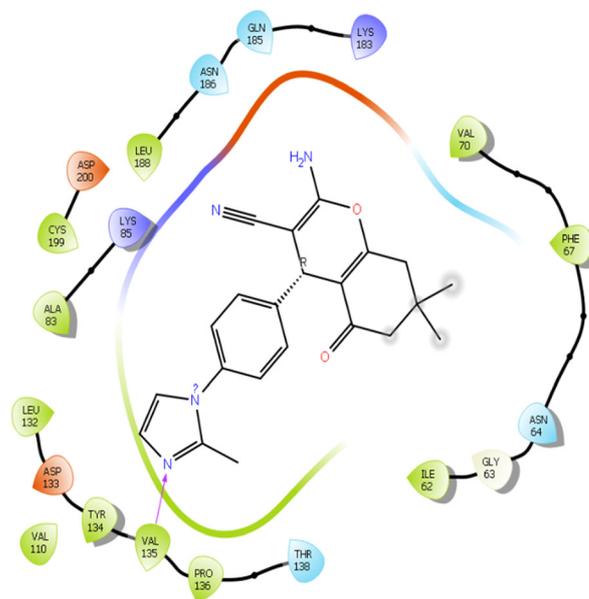


Fig. 14 Molecular docking analysis of compound **8a** against EGFR (A and B) and GSK3B (C and D) in 3D and 2D.

ligands.^{53,54} This process involves pathways including PI3K/Akt and MAPK, which regulate cell proliferation, survival, and differentiation.⁵⁵ In breast cancer, EGFR is frequently overexpressed or exhibits activating mutations, resulting in the activation of these pathways and subsequent uncontrolled cell growth.⁵⁶ EGFR signaling promotes cell cycle progression and DNA repair mechanisms. Disruption of these processes by EGFR inhibition can lead to DNA damage accumulation and trigger apoptosis.⁵⁷ SRC is a non-receptor tyrosine kinase involved in various cellular processes, including cell

growth, adhesion, migration, angiogenesis, and survival.⁵⁸ It is involved in cell survival signaling and DNA repair pathways.⁵⁸ Elevated SRC activity is observed in breast cancer, contributing to tumor progression, invasion, and metastasis.^{58,59} SRC can interact with and phosphorylate EGFR, enhancing EGFR signaling and promoting tumor growth.⁶⁰ Its inhibition by fused pyran derivatives can disrupt these processes, resulting in increased DNA damage and apoptosis. GSK3B is a serine/threonine kinase with diverse functions in cellular processes, including cell cycle regulation,⁶¹ apoptosis,⁶² and DNA repair.⁶³ Its role



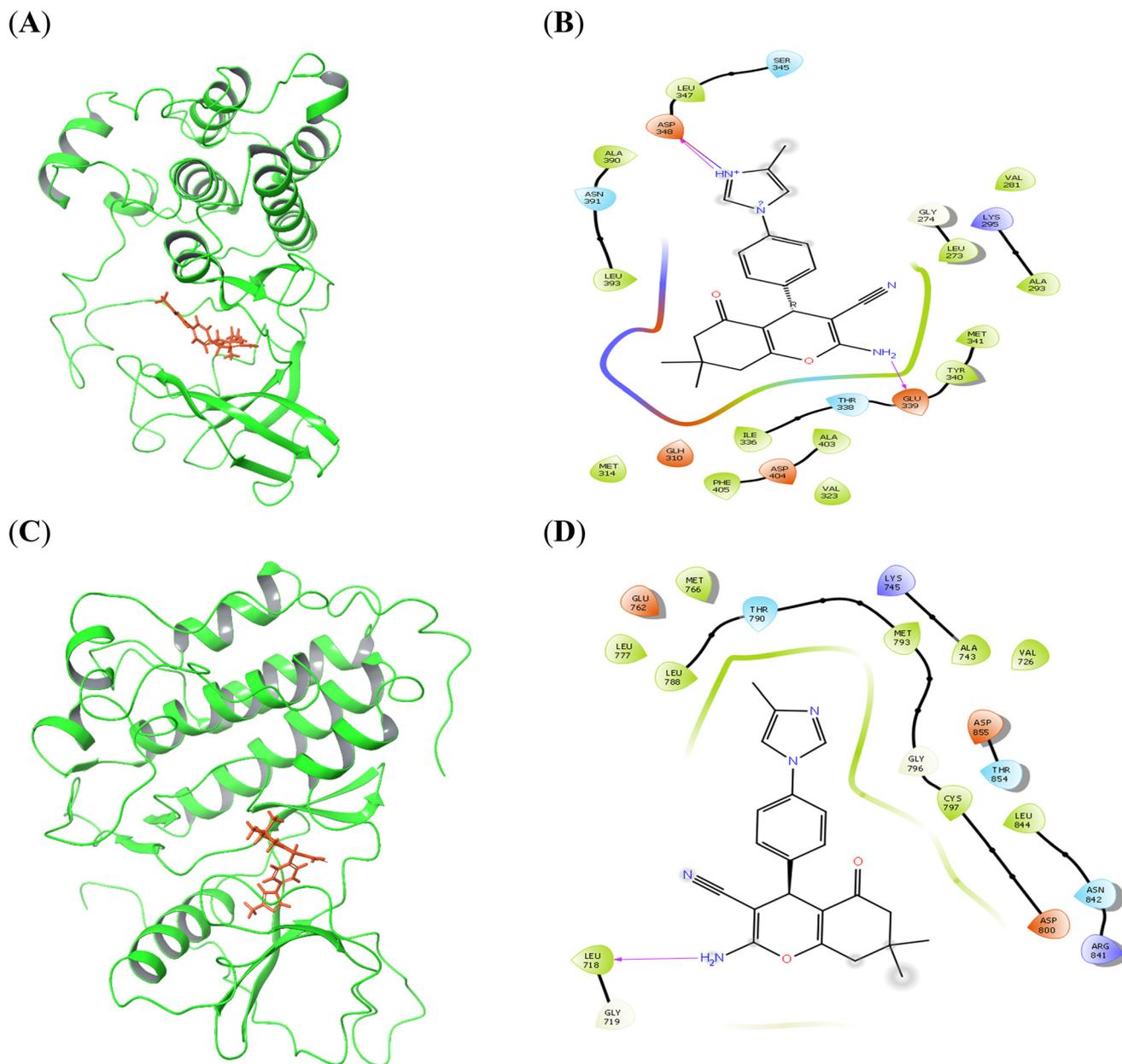


Fig. 15 Molecular docking analysis of compound **8b** against SRC (A and B) and EGFR (C and D) in 3D and 2D.

Table 2 The docking scores and binding energies of **8a** and **8b** against the target proteins

Compound	Target protein	Dock score (kcal mol ⁻¹)	Interacting residues	MMGBSA binding energy (kcal mol ⁻¹)
8a	EGFR	-5.42	—	-53.69
	GSK3B	-5.99	Val135 (H-bond)	-40.05
8b	SRC	-5.517	Asp348 (H-bond and salt bridge) Glu339 (H-bond)	-64.77
	EGFR	-4.66	Leu718 (H-bond)	-41.54

in cancer is complex and context-dependent. In breast cancer, GSK3B can be involved in both tumor-promoting and tumor-suppressing functions, contingent upon the specific context and signaling pathways involved.^{64,65} Targeting these proteins with drugs can disrupt oncogenic signaling, leading to apoptosis and DNA breaks.

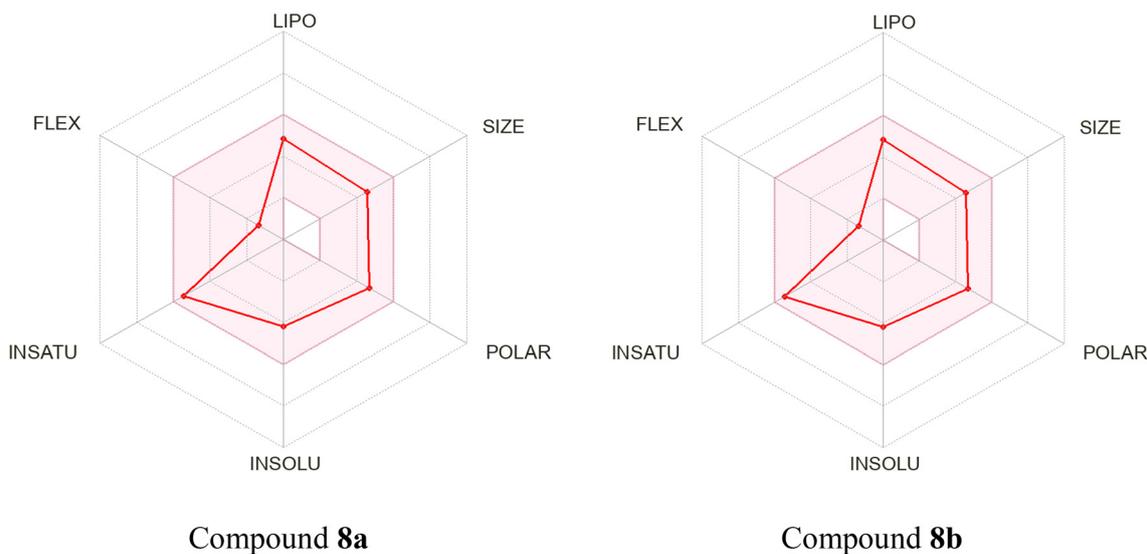
2.9 ADME prediction

The Swiss ADME webserver was utilized to calculate the physicochemical properties of the compounds. Several parameters, including lipophilicity, number of hydrogen bond donors, number of hydrogen bond acceptors, water



Table 3 The free energy (ΔG) parameters and the binding score of compounds **8a** and **8b** with its respective targets. All the energies are reported in kcal mol⁻¹

	MMGBSA, ΔG , bind	MMGBSA, ΔG , bind, Coulomb	MMGBSA, ΔG , bind, covalent	MMGBSA, ΔG , bind, H-bond	MMGBSA, ΔG , bind, lipo	MMGBSA, ΔG , bind, packing	MMGBSA, ΔG , bind, SelfCont.	MMGBSA, ΔG , bind, Solv. GB	MMGBSA, ΔG , bind, vdW
8a -EGFR	-53.69	-8.88	7.56	-0.32	-37.35	0.00	0.00	21.77	-36.48
8a -GSK3B	-40.05	-14.44	4.16	-1.12	-29.44	-0.00	0.00	31.90	-31.11
8b -SRC	-64.77	27.00	2.44	-1.05	-43.32	0.00	0.00	-15.70	-34.13
8b -EGFR	-41.54	-5.72	4.68	-0.26	-31.15	0.00	0.00	27.68	-36.76

**Fig. 16** Bioavailability radar and score prediction of compounds **8a** and **8b** were generated using SwissADME.

solubility, and drug-likeness, were calculated based on pre-trained models. A radar plot was generated for each molecule, wherein the colored zone represented a suitable physicochemical space for oral bioavailability (Fig. 16). Compounds **8a** and **8b** exhibited similar physicochemical properties, rendering them promising candidates for further investigation (Table 4). Both compounds possessed a molecular weight of 374.44 g mol⁻¹ and demonstrated moderate water solubility. Each compound contained four hydrogen bond acceptors and one donor, indicating their capacity to interact with biological molecules. Both compounds exhibited high gastrointestinal (GI) absorption, indicating good oral bioavailability, although neither demonstrates permeability of the blood–brain barrier (BBB). Furthermore, both **8a** and **8b** are substrates of P-glycoprotein (P-gp) and inhibitors of several cytochrome P450 enzymes (CYP1A2, CYP2C19, CYP2C9, and CYP3A4), which may have implications for drug metabolism and potential drug–drug interactions. They do not inhibit CYP2D6, which suggests a lower risk of interfering with the metabolism of other drugs that rely on this enzyme. Both compounds exhibited a $\log K_p$ value of -6.47 cm s^{-1} , indicating low skin permeation. In terms of drug-likeness, both compounds pass Lipinski's rule of five with no violations and meet the criteria for Ghose,

Veber, Egan, and Muegge rules, suggesting that they possess favorable characteristics for oral drug development. Their bioavailability score was 0.56, indicating moderate potential for effective absorption.

Table 4 The physicochemical properties of compounds **8a** and **8b**

Properties	8a	8b
Water solubility	Moderately soluble	Moderately soluble
Number of H-bond acceptors	4	4
Number of H-bond donors	1	1
GI absorption	High	High
BBB permeant	No	No
P-gp substrate	Yes	Yes
CYP1A2 inhibitor	Yes	Yes
CYP2C19 inhibitor	Yes	Yes
CYP2C9 inhibitor	Yes	Yes
CYP2D6 inhibitor	No	No
CYP3A4 inhibitor	Yes	Yes
$\log K_p$ (skin permeation)	-6.47 cm s^{-1}	-6.47 cm s^{-1}
Druglikeness		
Lipinski	Yes; 0 violation	Yes; 0 violation
Ghose	Yes	Yes
Veber	Yes	Yes
Egan	Yes	Yes
Muegge	Yes	Yes
Bioavailability score	0.56	0.56



3. Conclusion

In this study, we successfully designed and synthesized a series of N-heterocyclic substituted fused pyran derivatives using NaF catalyst in an aqueous solvent. All the fused pyran derivatives were evaluated for their anti-proliferative activity through *in vitro* assay on MCF7, A549 and HCT116 cell lines, and several compounds exhibited promising anticancer properties. Notably, compounds **8a** and **8b** demonstrated significant inhibitory effects on the MCF7 cell line, with IC₅₀ values of 8.24 ± 0.19 and 4.22 ± 0.81 μM, respectively. These compounds were nontoxic to the human diploid fibroblast cell line, MCR5. Colony formation assays revealed a marked reduction in the colony-forming capacity of MCF7 cells treated with compounds **8a** and **8b**, suggesting their potential to inhibit cellular proliferation and tumor progression. Spheroid formation assays in three-dimensional cultures further demonstrated consistent decreases in spheroid size, highlighting the efficacy of these compounds in targeting tumor-like structures in a 3D environment. Compounds **8a** and **8b** were shown to induce G1 phase cell cycle arrest in MCF7 cells, which is a key mechanism in limiting cancer cell proliferation. Comparisons with the standard drug, etoposide, further elucidated their therapeutic potential. Additionally, DNA damage analysis revealed high-intensity γH2AX responses for compounds **8a** and **8b**, indicating that their mechanisms of action may involve the induction of DNA damage, a characteristic of effective chemotherapeutics. Furthermore, apoptosis assays demonstrated the capacity of these compounds to modulate apoptotic pathways, particularly by increasing the early apoptotic populations in MCF7 cells. In addition, *ex ovo* CAM assay (angiogenesis analysis) revealed a significant reduction in new blood vessel formation over time compared to untreated controls. The docking studies of compounds **8a** and **8b** with targets revealed favorable binding affinities with EGFR, SRC, and GSK3β. This interaction of compounds with targets potentially disrupts critical signaling pathways that drive breast cancer development and progression. *In silico* ADME analysis indicates favorable pharmacokinetic properties. These findings collectively elucidate the therapeutic potential of compounds **8a** and **8b** as promising candidates for further development as effective anticancer agents.

4. Experimental section

4.1 Chemistry

All the solvents and reagents were purchased from Sigma Aldrich, TCI, Thermo Fischer, SRL, or Spectrochem and used without further purification unless otherwise mentioned. The melting point of all the compounds was determined using the Stuart-SMP20 capillary melting point apparatus and are uncorrected. FTIR spectra were recorded on a JASCO FT-IR 4700 spectrometer by dispersing the compounds in potassium bromide pellets. Thin layer chromatography (TLC) was performed on silica gel plates (60 F254, 0.25 mm, Merck)

using a 4:1 (v/v) mixture of hexanes and ethyl acetate and components were visualized under UV light or with iodine. Chromatographic separation was carried out on 100–200 mesh silica gel. ¹H NMR and ¹³C NMR spectra were measured using a 500 MHz JEOL spectrometer in CDCl₃/DMSO-*d*₆ solvent using TMS as an internal standard. High-resolution mass spectra (HRMS) data were obtained using an Agilent 6530 ESI-QTOF mass spectrometer (mass accuracy of 5 ppm) in positive ion mode with a reverse-phase C18 column.

Synthesis of 4-(3-methyl-1H-pyrazol-1-yl)benzaldehyde (3a). 3-Methyl-1H-pyrazole (**2a**, 164.2 mg, 2 mmol) and anhydrous potassium carbonate (829.23 mg, 6 mmol) were added to a solution of 4-fluorobenzaldehyde (**1**, 248.22 mg, 2 mmol) in dimethylformamide (5 mL). The obtained reaction mixture was heated at 130 °C for 19 h. The mixture was cooled to room temperature, poured into water (50 mL) and extracted the crude product with ethyl acetate (20 mL) thrice. The combined organic layer was dried over Na₂SO₄, concentrated under reduced pressure, and purified by column chromatography using 20% ethyl acetate in hexanes to obtain the pure product. Yield: 70%. Pale yellow solid. Melting point: 95–96 °C. IR (KBr, ν_{max}, cm⁻¹): 3136 (C–H), 2861, 2753 (C–H, aldehyde), 1703 (C=O), 1613 (C=C), 1518 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 9.97 (s, 1H), 7.94–7.90 (m, 3H), 7.82–7.80 (m, 2H), 6.30 (d, *J* = 2.3 Hz, 1H), 2.37 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 191.1, 152.1, 144.4, 133.6, 131.4, 127.7, 118.3, 109.1, 13.9. MS (ESI-QTOF) for C₁₁H₁₀N₂O [M + H]⁺ calculated 187.0866, found 187.0871.

Compounds **3b–l** were synthesized following the same procedure described for compound **3a**.

4-(4-Methyl-1H-pyrazol-1-yl)benzaldehyde (3b). 4-Methyl-1H-pyrazole (**2b**, 164.2 mg, 2 mmol) was used as a reagent. Yield: 68%. White solid. Melting point: 88–90 °C. IR (KBr, ν_{max}, cm⁻¹): 3100 (C–H), 2812, 2754 (C–H, aldehyde), 1704 (C=O), 1612 (C=C), 1512 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 9.97 (s, 1H), 7.94–7.92 (m, 2H), 7.82–7.78 (m, 3H), 7.57 (s, 1H), 2.15 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 191.0, 144.4, 143.3, 133.8, 131.4, 125.4, 119.7, 118.3, 9.0. MS (ESI-QTOF) for C₁₁H₁₀N₂O [M + H]⁺ calculated 187.0866, found 187.0867.

4-(3,5-Dimethyl-1H-pyrazol-1-yl)benzaldehyde (3c). 3,5-Dimethyl-1H-pyrazole (**2c**, 192.26 mg, 2 mmol) was used as a reagent. Yield: 56%. Yellow liquid. IR (KBr, ν_{max}, cm⁻¹): 3136 (C–H), 2831, 2721 (C–H, aldehyde), 1694 (C=O), 1604 (C=C), 1511 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 9.86 (s, 1H), 7.82–7.76 (m, 2H), 7.51 (d, *J* = 8.4 Hz, 1H), 7.02 (d, *J* = 8.4 Hz, 1H), 5.91 (s, 1H), 2.25 (s, 3H), 2.14 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 191.2, 150.2, 144.7, 139.8, 134.2, 132.1, 130.6, 123.8, 119.3, 108.8, 13.5, 13.0. MS (ESI-QTOF) for C₁₂H₁₂N₂O [M + H]⁺ calculated 201.1022, found 201.1021.

4-(4-Bromo-1H-pyrazol-1-yl)benzaldehyde (3d). 4-Bromo-1H-pyrazole (**2d**, 293.94 mg, 2 mmol) was used as a reagent. Yield: 60%. White solid. Melting point: 120–124 °C. IR (KBr, ν_{max}, cm⁻¹): 3120 (C–H), 2812, 2730 (C–H, aldehyde), 1694 (C=O), 1612 (C=C), 1502 (C=N). ¹H NMR (500



MHz, CDCl₃): δ 10.00 (s, 1H), 8.03 (s, 1H), 7.98–7.96 (m, 2H), 7.83–7.81 (m, 2H), 7.71 (s, 1H). ¹³C NMR (126 MHz, CDCl₃): δ 190.9, 143.6, 142.8, 134.6, 131.5, 127.2, 118.7, 97.2. MS (ESI-QTOF) for C₁₀H₇BrN₂O [M + H]⁺ calculated 250.9815, found 250.9819.

4-(3-(Trifluoromethyl)-1H-pyrazol-1-yl)benzaldehyde (3e). 3-(Trifluoromethyl)-1H-pyrazole (**2e**, 272.15 mg, 2 mmol) was used as a reagent. Yield: 56%. Pale yellow liquid. IR (KBr, ν_{\max} , cm⁻¹): 3138 (C–H), 2821, 2739 (C–H, aldehyde), 1690 (C=O), 1617 (C=C), 1511 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.01 (s, 1H), 8.06 (d, *J* = 1.5 Hz, 1H), 7.99–7.97 (m, 2H), 7.90–7.88 (m, 2H), 6.76 (d, *J* = 2.3 Hz, 1H). ¹³C NMR (126 MHz, CDCl₃): δ 190.9, 145.10 (q, *J* = 38.8 Hz), 143.4, 135.2, 131.4, 128.6, 119.80 (m), 119.7, 107.0. MS (ESI-QTOF) for C₁₁H₇F₃N₂O [M + H]⁺ calculated 241.0583, found 241.0581.

4-(2-Methyl-1H-imidazol-1-yl)benzaldehyde (3f). 2-Methyl-1H-imidazole (**2f**, 164.2 mg, 2 mmol) was used as a reagent. Yield: 70%. Yellow solid. Melting point: 84–86 °C. IR (KBr, ν_{\max} , cm⁻¹): 3164 (C–H), 2839, 2750 (C–H, aldehyde), 1684 (C=O), 1600 (C=C), 1511 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.04 (s, 1H), 8.00–7.98 (m, 2H), 7.46 (d, *J* = 8.4 Hz, 2H), 7.03 (s, 2H), 2.40 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 190.9, 144.6, 142.9, 135.6, 131.1, 128.5, 125.8, 120.3, 14.1. MS (ESI-QTOF) for C₁₁H₁₀N₂O [M + H]⁺ calculated 187.0866, found 187.0872.

4-(4-Methyl-1H-imidazol-1-yl)benzaldehyde (3g). 4-Methyl-1H-imidazole (**2g**, 164.2 mg, 2 mmol) was used as a reagent. Yield: 75%. Yellow liquid. IR (KBr, ν_{\max} , cm⁻¹): 3124 (C–H), 2831, 2748 (C–H, aldehyde), 1694 (C=O), 1603 (C=C), 1511 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.06 (s, 1H), 8.02–8.01 (m, 2H), 7.60 (s, 1H), 7.46 (d, *J* = 8.4 Hz, 2H), 6.92 (s, 1H), 2.22 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 190.7, 141.9, 140.6, 134.7, 134.5, 131.7, 120.6, 114.0, 13.7. MS (ESI-QTOF) for C₁₁H₁₀N₂O [M + H]⁺ calculated 187.0866, found 187.0867.

4-(5-Methyl-1H-imidazol-1-yl)benzaldehyde (3h). 5-Methyl-1H-imidazole (**2h**, 164.2 mg, 2 mmol) was used as a reagent. Yield: 64%. Yellow solid. Melting point: 80–83 °C. IR (KBr, ν_{\max} , cm⁻¹): 3156 (C–H), 2835, 2748 (C–H, aldehyde), 1703 (C=O), 1607 (C=C), 1520 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.00 (s, 1H), 7.97 (d, *J* = 8.4 Hz, 2H), 7.86 (d, *J* = 1.5 Hz, 1H), 7.51 (d, *J* = 8.4 Hz, 2H), 7.07 (s, 1H), 2.28 (s, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 190.8, 141.9, 140.6, 134.7, 134.5, 131.7, 120.6, 114.0, 13.7. MS (ESI-QTOF) for C₁₁H₁₀N₂O [M + H]⁺ calculated 187.0866, found 187.0873.

4-(2-Ethyl-1H-imidazol-1-yl)benzaldehyde (3i). 2-Ethyl-1H-imidazole (**2i**, 192.26 mg, 2 mmol) was used as a reagent. Yield: 57%. Yellow liquid. IR (KBr, ν_{\max} , cm⁻¹): 3149 (C–H), 2852, 2762 (C–H, aldehyde), 1690 (C=O), 1600 (C=C), 1514 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.03 (s, 1H), 7.99–7.97 (m, 2H), 7.45–7.43 (m, 2H), 7.04 (d, *J* = 1.5 Hz, 1H), 6.99 (d, *J* = 1.5 Hz, 1H), 2.67 (q, *J* = 7.4 Hz, 2H), 1.22 (t, *J* = 7.6 Hz, 3H). ¹³C NMR (126 MHz, CDCl₃): δ 190.9, 149.5, 142.9, 135.7, 131.0, 128.4, 126.1, 120.3, 20.8,

12.3. MS (ESI-QTOF) for C₁₂H₁₂N₂O [M + H]⁺ calculated 201.1022, found 201.1031.

4-(4-Bromo-1H-imidazol-1-yl)benzaldehyde (3j). 4-Bromo-1H-imidazole (**2j**, 293.94 mg, 2 mmol) was used as a reagent. Yield: 62%. Yellow solid. Melting point: 159–162 °C. IR (KBr, ν_{\max} , cm⁻¹): 3101 (C–H), 2808, 2735 (C–H, aldehyde), 1690 (C=O), 1612 (C=C), 1511 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.03 (s, 1H), 8.03–8.00 (m, 2H), 7.83 (d, *J* = 1.5 Hz, 1H), 7.55–7.53 (m, 2H), 7.34 (d, *J* = 1.5 Hz, 1H). ¹³C NMR (126 MHz, CDCl₃): δ 190.5, 140.9, 135.6, 135.0, 131.8, 121.3, 118.0, 117.2. MS (ESI-QTOF) for C₁₀H₇BrN₂O [M + H]⁺ calculated 250.9815, found 250.9811.

4-(1H-1,2,3-Triazol-1-yl)benzaldehyde (3k). 1H-1,2,3-Triazole (**2k**, 138.12 mg, 2 mmol) was used as a reagent. Yield: 58%. White solid. Melting point: 106–108 °C. IR (KBr, ν_{\max} , cm⁻¹): 3101 (C–H), 2854, 2744 (C–H, aldehyde), 1699 (C=O), 1617 (C=C), 1507 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.03 (s, 1H), 8.26–8.24 (m, 2H), 8.01–7.98 (m, 2H), 7.86 (s, 2H). ¹³C NMR (126 MHz, CDCl₃): δ 191.1, 143.7, 136.7, 135.1, 131.2, 119.2. MS (ESI-QTOF) for C₉H₇N₃O [M + H]⁺ calculated 174.0662, found 174.0664.

4-(1H-Tetrazol-1-yl)benzaldehyde (3l). 1H-Tetrazole (**2l**, 140.11 mg, 2 mmol) was used as a reagent. Yield: 50%. Pale grey solid. Melting point: 98–99 °C. IR (KBr, ν_{\max} , cm⁻¹): 3100 (C–H), 2862, 2733 (C–H, aldehyde), 1691 (C=O), 1604 (C=C), 1504 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 10.10 (s, 1H), 8.71 (s, 1H), 8.36 (d, *J* = 9.2 Hz, 2H), 8.10 (d, *J* = 10.7 Hz, 2H). ¹³C NMR (126 MHz, CDCl₃): δ 190.7, 153.5, 140.4, 137.0, 131.3, 120.5. MS (ESI-QTOF) for C₈H₆N₄O [M + H]⁺ calculated 175.0614, found 175.0623.

Synthesis of 4-((1-phenyl-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (13a). To a mixture of azidobenzene **10a** (238.24 mg, 5 mmol) and 4-(prop-2-yn-1-yloxy)benzaldehyde **12** (800.26 mg, 5 mmol) in anhydrous THF (10 mL) was added catalytic amount of copper iodide. Then, the reaction mixture was refluxed for 12 h. After completion of the reaction (monitored by TLC), the mixture was filtered and washed with cold THF, and the crude product present in the filtrate was extracted with ethyl acetate (3 × 20 mL) three times. The combined organic layer was washed with brine solution, dried over Na₂SO₄, and concentrated using a rotary evaporator. The product obtained was recrystallized using a mixture of chloroform and hexanes (8:2 v/v) to afford the pure product. Yield: 74%. Off-white solid. Melting point: 100–103 °C. IR (KBr, ν_{\max} , cm⁻¹): 3124 (C–H), 2808, 2735 (C–H, aldehyde), 1690 (C=O), 1617 (C=C), 1507 (C=N). ¹H NMR (500 MHz, CDCl₃): δ 9.83 (s, 1H), 8.02 (s, 1H), 7.81–7.77 (m, 2H), 7.68–7.65 (m, 2H), 7.48–7.44 (m, 2H), 7.40–7.37 (m, 1H), 7.09–7.06 (m, 2H), 5.31 (s, 2H). ¹³C NMR (126 MHz, CDCl₃): δ 190.8, 163.1, 144.0, 136.8, 132.0, 130.4, 129.8, 129.0, 121.1, 120.6, 115.1, 62.1. MS (ESI-QTOF) for C₁₆H₁₃N₃O₂ [M + H]⁺ calculated 280.1081, found 280.1082.

Compounds **13b–e** were synthesized by following the same procedure as described for compound **13a**.

4-((1-(4-Chlorophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (13b). 1-Azido-4-chlorobenzene (**10b**, 307.12



mg, 2 mmol) was used as a reagent. Yield: 75%. Pale brown solid. Melting point: 138–140 °C. IR (KBr, ν_{\max} , cm^{-1}): 3082 (C–H), 2837, 2762 (=C–H, aldehyde), 1709 (C=O), 1606 (C=C), 1504 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.87 (s, 1H), 8.06 (s, 1H), 7.84–7.82 (m, 2H), 7.69–7.67 (m, 2H), 7.50–7.47 (m, 2H), 7.13–7.11 (m, 2H), 5.35 (s, 2H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.8, 163.1, 144.3, 135.4, 135.0, 132.1, 130.6, 130.1, 121.9, 121.2, 115.2, 62.1. MS (ESI-QTOF) for $\text{C}_{16}\text{H}_{12}\text{ClN}_3\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 314.0691, found 314.0690.

4-((1-(4-Bromophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (13c). 1-Azido-4-bromobenzene (**10c**, 396.04 mg, 2 mmol) was used as a reagent. Yield: 70%. Pale yellow solid. Melting point: 159–161 °C. IR (KBr, ν_{\max} , cm^{-1}): 3065 (C–H), 2835, 2753 (C–H, aldehyde), 1685 (C=O), 1612 (C=C), 1516 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.88 (s, 1H), 8.07 (s, 1H), 7.86–7.83 (m, 2H), 7.66–7.61 (m, 4H), 7.14–7.11 (m, 2H), 5.36 (s, 2H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.9, 163.1, 135.9, 133.1, 132.1, 130.6, 122.8, 122.1, 121.2, 121.0, 115.2, 62.1. MS (ESI-QTOF) for $\text{C}_{16}\text{H}_{12}\text{BrN}_3\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 358.0186, found 358.0189.

4-((1-(*p*-Tolyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (13d). 1-Azido-4-methylbenzene (**10d**, 266 mg, 2 mmol) was used as a reagent. Yield: 69%. Off-white solid. Melting point: 134–136 °C. IR (KBr, ν_{\max} , cm^{-1}): 3133 (C–H), 2812, 2733 (C–H, aldehyde), 1699 (C=O), 1612 (C=C), 1525 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.88 (s, 1H), 8.03 (s, 1H), 7.85–7.83 (m, 2H), 7.60–7.58 (m, 2H), 7.30 (d, $J = 7.6$ Hz, 2H), 7.14–7.11 (m, 2H), 5.36 (s, 2H), 2.41 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.9, 163.2, 143.9, 139.3, 134.7, 132.1, 130.5, 130.4, 121.3, 120.6, 115.2, 62.2, 21.2. MS (ESI-QTOF) for $\text{C}_{17}\text{H}_{15}\text{N}_3\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 294.1237, found 294.1233.

4-((1-(4-Methoxyphenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (13e). 1-Azido-4-methoxybenzene (**10e**, 298 mg, 2 mmol) was used as a reagent. Yield: 65%. Pale orange solid. Melting point: 130–133 °C. IR (KBr, ν_{\max} , cm^{-1}): 3051 (C–H), 2803, 2721 (C–H, aldehyde), 1694 (C=O), 1612 (C=C), 1520 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.87 (s, 1H), 7.99 (s, 1H), 7.84–7.82 (m, 2H), 7.62–7.59 (m, 2H), 7.14–7.11 (m, 2H), 7.02–6.98 (m, 2H), 5.35 (s, 2H), 3.84 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.9, 163.2, 160.1, 143.8, 132.1, 130.5, 130.4, 122.4, 121.4, 115.2, 114.9, 62.2, 55.7. MS (ESI-QTOF) for $\text{C}_{17}\text{H}_{15}\text{N}_3\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 310.1186, found 310.1182.

Synthesis of 2-amino-4-(4-(3-methyl-1H-pyrazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (7a). To a solution of 4-(3-methyl-1H-pyrazol-1-yl)benzaldehyde (**3a**, 186.07 mg, 1 mmol) in water (2 mL) was added cyclohexane-1,3-dione (**4**, 112.05 mg, 1 mmol), malononitrile (**5**, 66.06 mg, 1 mmol), and sodium fluoride (5 mg, 12 mol%). The obtained reaction mixture was refluxed for 3 h. After completion of the reaction, the mixture was cooled to room temperature, and the solid product obtained was filtered and washed with water (10 mL) three times to remove the NaF. The crude product obtained was purified by column chromatography using 30% ethyl acetate in hexanes to get the pure product. Yield: 80%. Off-white solid. Melting point: 224–226 °C. IR (ν_{\max} , cm^{-1}): 3381, 3216 ($-\text{NH}_2$), 3134

(=C–H), 2186 (CN), 1676 (C=O), 1617 (C=C), 1222 (C–O–C). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.25 (d, $J = 2.3$ Hz, 1H), 7.64–7.61 (m, 2H), 7.20–7.19 (m, 2H), 7.01 (s, 2H), 6.27 (d, $J = 2.3$ Hz, 1H), 4.19 (s, 1H), 2.60–2.54 (m, 2H), 2.29–2.22 (m, 5H), 1.97–1.83 (m, 2H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.4, 165.0, 159.0, 149.9, 142.7, 138.9, 128.8, 128.7, 120.3, 118.6, 114.1, 108.1, 58.4, 36.9, 35.5, 27.0, 20.3, 14.0. MS (ESI-QTOF) for $\text{C}_{20}\text{H}_{18}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 347.1503, found 347.1505.

Compounds **7b–e**, **8a–e**, **15a–g** and **17a–g** were synthesized using the same procedure outlined for compound **7a**.

2-Amino-4-(4-(4-methyl-1H-pyrazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (7b). 4-(4-Methyl-1H-pyrazol-1-yl)benzaldehyde (**3b**, 186.21 mg, 1 mmol) was used as a reagent. Yield: 77%. Off-white solid. Melting point: 255–258 °C. IR (KBr, ν_{\max} , cm^{-1}): 3344, 3179 ($-\text{NH}_2$), 3032 (=C–H), 2194 (CN), 1686 (C=O), 1630 (C=C), 1236 (C–O–C). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.15 (s, 1H), 7.62 (d, $J = 8.4$ Hz, 2H), 7.49 (s, 1H), 7.20 (d, $J = 8.4$ Hz, 2H), 7.00 (s, 2H), 4.19 (s, 1H), 2.60–2.56 (m, 2H), 2.27–2.20 (m, 2H), 2.00–2.09 (m, 3H), 1.91–1.83 (m, 2H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.4, 165.1, 159.0, 142.9, 142.0, 138.9, 128.7, 126.5, 120.3, 118.6, 118.0, 114.1, 58.4, 36.9, 35.5, 27.0, 20.3, 9.3. MS (ESI-QTOF) for $\text{C}_{20}\text{H}_{18}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 347.1503, found 347.1501.

2-Amino-4-(4-(3,5-dimethyl-1H-pyrazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (7c). 4-(3,5-Dimethyl-1H-pyrazol-1-yl)benzaldehyde (**3c**, 200.24 mg, 1 mmol) was used as a reagent. Yield: 71%. Mustard yellow solid. Melting point: 220–224 °C. IR (KBr, ν_{\max} , cm^{-1}): 3335, 3183 ($-\text{NH}_2$), 2941 (=C–H), 2189 (CN), 1680 (C=O), 1635 (C=C), 1255 (C–O–C). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 7.35 (d, $J = 8.4$ Hz, 2H), 7.22 (d, $J = 8.4$ Hz, 2H), 7.02 (s, 2H), 6.00 (s, 1H), 4.22 (s, 1H), 2.61–2.57 (m, 2H), 2.28–2.23 (m, 5H), 2.12 (s, 3H), 1.98–1.87 (m, 2H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.5, 165.2, 159.1, 148.2, 143.9, 139.5, 138.7, 128.3, 124.5, 120.3, 114.1, 107.6, 58.4, 36.9, 35.6, 27.0, 20.3, 13.9, 12.7. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{20}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 361.1659, found 361.1660.

2-Amino-4-(4-(4-bromo-1H-pyrazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (7d). 4-(4-Bromo-1H-pyrazol-1-yl)benzaldehyde (**3d**, 251.08 mg, 1 mmol) was used as a reagent. Yield: 72%. Grey solid. Melting point: 220–224 °C. IR (KBr, ν_{\max} , cm^{-1}): 3310, 3188 ($-\text{NH}_2$), 3120 (=C–H), 2182 (CN), 1683 (C=O), 1642 (C=C), 1243 (C–O–C). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.91 (s, 1H), 8.67 (s, 1H), 8.48 (s, 1H), 8.04 (d, $J = 1.5$ Hz, 1H), 7.81 (s, 1H), 7.67 (d, $J = 8.4$ Hz, 1H), 7.24 (d, $J = 8.4$ Hz, 1H), 7.02 (s, 1H), 4.20 (s, 1H), 2.63–2.56 (m, 2H), 2.29–2.20 (m, 2H), 1.94–1.85 (m, 2H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.4, 165.1, 158.9, 145.0, 142.9, 142.6, 137.9, 131.1, 128.9, 120.2, 113.9, 106.7, 58.2, 36.8, 35.7, 27.0, 20.3. MS (ESI-QTOF) for $\text{C}_{19}\text{H}_{15}\text{BrN}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 411.0451, found 411.0448.

2-Amino-5-oxo-4-(4-(3-(trifluoromethyl)-1H-pyrazol-1-yl)phenyl)-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (7e). 4-(3-(Trifluoromethyl)-1H-pyrazol-1-yl)benzaldehyde (**3e**, 240.18 mg,



1 mmol) was used as a reagent. Yield: 70%. Yellow solid. Melting point: 200–204 °C. IR (KBr, ν_{\max} , cm^{-1}): 3386, 3306 ($-\text{NH}_2$), 2964 ($=\text{C}-\text{H}$), 2195 (CN), 1693 ($\text{C}=\text{O}$), 1640 ($\text{C}=\text{C}$), 1249 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.61 (s, 1H), 7.72 (d, $J = 8.4$ Hz, 2H), 7.30 (d, $J = 8.4$ Hz, 2H), 7.03 (s, 2H), 6.98 (d, $J = 2.3$ Hz, 1H), 4.24 (s, 1H), 2.62–2.57 (m, 2H), 2.30–2.20 (m, 2H), 1.96–1.84 (m, 2H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.4, 165.2, 159.0, 145.0, 142.9, 142.7, 137.9, 131.1, 129.0, 120.8, 120.2, 113.9, 106.7, 58.2, 36.8, 35.7, 27.0, 20. MS (ESI-QTOF) for $\text{C}_{20}\text{H}_{15}\text{F}_3\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 401.1220, found 401.1222.

2-Amino-7,7-dimethyl-4-(4-(2-methyl-1H-imidazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (8a). 4-(2-Methyl-1H-imidazol-1-yl)benzaldehyde (**3f**, 186.21 mg, 1 mmol) was used as a reagent. Yield: 79%. Pale yellow solid. Melting point: 229–232 °C. IR (KBr, ν_{\max} , cm^{-1}): 3381, 3312 ($-\text{NH}_2$), 2991 ($=\text{C}-\text{H}$), 2194 (CN), 1699 ($\text{C}=\text{O}$), 1630 ($\text{C}=\text{C}$), 1255 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 7.33 (d, $J = 8.4$ Hz, 2H), 7.25 (d, $J = 8.4$ Hz, 3H), 7.05 (s, 2H), 6.85 (s, 1H), 4.24 (s, 1H), 2.50 (s, 2H), 2.25–2.10 (m, 5H), 1.01 (s, 3H), 0.95 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.3, 163.4, 159.1, 144.9, 144.0, 136.6, 128.7, 127.6, 125.6, 121.5, 120.2, 112.9, 58.4, 50.5, 35.7, 32.4, 28.8, 27.5, 14.2. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{22}\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 375.1816, found 375.1810.

2-Amino-7,7-dimethyl-4-(4-(4-methyl-1H-imidazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (8b). 4-(4-Methyl-1H-imidazol-1-yl)benzaldehyde (**3g**, 186.21 mg, 1 mmol) was used as a reagent. Yield: 77%. Off-white solid. Melting point: 230–234 °C. IR (KBr, ν_{\max} , cm^{-1}): 3326, 3183 ($-\text{NH}_2$), 2941 ($=\text{C}-\text{H}$), 2194 (CN), 1690 ($\text{C}=\text{O}$), 1625 ($\text{C}=\text{C}$), 1236 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.03 (s, 1H), 7.46 (d, $J = 8.4$ Hz, 2H), 7.34 (s, 1H), 7.21 (d, $J = 8.4$ Hz, 2H), 7.02 (s, 2H), 4.20 (s, 1H), 2.49 (s, 2H), 2.24–2.06 (m, 5H), 1.00 (s, 3H), 0.93 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.3, 163.2, 159.0, 143.7, 138.7, 136.1, 135.2, 129.1, 120.6, 120.2, 114.8, 112.9, 58.5, 50.5, 35.6, 32.4, 28.9, 27.4, 14.1. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{22}\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 375.1816, found 375.1815.

2-Amino-7,7-dimethyl-4-(4-(5-methyl-1H-imidazol-1-yl)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (8c). 4-(5-Methyl-1H-imidazol-1-yl)benzaldehyde (**3h**, 272.15 mg, 1 mmol) was used as a reagent. Yield: 73%. Pale yellow solid. Melting point: 225–228 °C. IR (KBr, ν_{\max} , cm^{-1}): 3312, 3188 ($-\text{NH}_2$), 2959 ($=\text{C}-\text{H}$), 2194 (CN), 1685 ($\text{C}=\text{O}$), 1621 ($\text{C}=\text{C}$), 1245 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.05 (s, 1H), 7.46 (d, $J = 8.4$ Hz, 2H), 7.35 (s, 1H), 7.21 (d, $J = 9.2$ Hz, 2H), 7.02 (s, 2H), 4.19 (s, 1H), 2.49 (s, 2H), 2.24–2.06 (m, 5H), 1.00 (s, 3H), 0.93 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.3, 163.2, 159.0, 143.7, 138.6, 136.0, 135.2, 129.1, 120.7, 120.2, 114.8, 112.9, 58.5, 50.5, 35.6, 32.4, 28.9, 27.4, 14.1. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{22}\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 375.1816, found 375.1811.

2-Amino-4-(4-(2-ethyl-1H-imidazol-1-yl)phenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (8d). 4-(2-Ethyl-1H-imidazol-1-yl)benzaldehyde (**3i**, 200.24 mg, 1 mmol) was used as a reagent. Yield: 70%. Yellow

solid. Melting point: 230–233 °C. IR (KBr, ν_{\max} , cm^{-1}): 3358, 3321 ($-\text{NH}_2$), 3138 ($=\text{C}-\text{H}$), 2203 (CN), 1690 ($\text{C}=\text{O}$), 1653 ($\text{C}=\text{C}$), 1245 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 7.31 (d, $J = 8.4$ Hz, 2H), 7.26–7.21 (m, 3H), 7.06 (s, 2H), 6.88 (s, 1H), 4.24 (s, 1H), 2.56–2.50 (m, 4H), 2.25–2.10 (m, 2H), 1.07 (t, $J = 7.6$ Hz, 3H), 1.01 (s, 3H), 0.94 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.3, 163.4, 159.1, 148.8, 145.1, 136.4, 128.7, 127.5, 126.0, 121.5, 120.2, 112.9, 58.4, 50.5, 35.7, 32.4, 28.8, 27.5, 20.6, 12.7. MS (ESI-QTOF) for $\text{C}_{23}\text{H}_{24}\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 389.1972, found 389.1978.

2-Amino-4-(4-(4-bromo-1H-imidazol-1-yl)phenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (8e). 4-(4-Bromo-1H-imidazol-1-yl)benzaldehyde (**3j**, 251.08 mg, 1 mmol) was used as a reagent. Yield: 73%. Yellow solid. Melting point: 197–200 °C. IR (KBr, ν_{\max} , cm^{-1}): 3358, 3307 ($-\text{NH}_2$), 3147 ($=\text{C}-\text{H}$), 2185 (CN), 1691 ($\text{C}=\text{O}$), 1653 ($\text{C}=\text{C}$), 1250 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.17 (s, 1H), 7.87 (d, $J = 1.5$ Hz, 1H), 7.52 (d, $J = 8.4$ Hz, 2H), 7.24 (d, $J = 9.2$ Hz, 2H), 7.04 (s, 2H), 4.22 (s, 1H), 2.50 (s, 2H), 2.24–2.07 (m, 2H), 1.01 (s, 3H), 0.93 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 196.2, 163.3, 159.1, 144.8, 136.4, 135.3, 129.1, 121.2, 120.1, 118.1, 116.0, 112.9, 58.4, 50.5, 35.7, 32.4, 28.8, 27.5. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{19}\text{BrN}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 439.0764, found 439.0768.

2-Amino-5-oxo-4-(4-((1-phenyl-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15a). 4-((1-Phenyl-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13a**, 279.29 mg, 1 mmol) was used as a reagent. Yield: 75%. White solid. Melting point: 220–223 °C. IR (KBr, ν_{\max} , cm^{-1}): 3436, 3248 ($-\text{NH}_2$), 3178 ($=\text{C}-\text{H}$), 2181 (CN), 1708 ($\text{C}=\text{O}$), 1662 ($\text{C}=\text{C}$), 1259 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.90 (s, 1H), 7.86 (t, $J = 7.2$ Hz, 3H), 7.68–7.65 (m, 1H), 7.56 (t, $J = 8.0$ Hz, 2H), 7.47–7.41 (m, 3H), 7.35 (s, 2H), 7.17 (d, $J = 9.2$ Hz, 2H), 6.99 (d, $J = 8.0$ Hz, 2H), 5.17 (s, 2H), 4.38 (s, 1H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 158.5, 157.7, 153.7, 152.6, 144.4, 137.1, 136.4, 133.4, 130.4, 129.4, 129.3, 125.2, 123.4, 123.0, 120.7, 119.9, 117.1, 115.2, 113.5, 104.8, 61.6, 58.7, 36.7. MS (ESI-QTOF) for $\text{C}_{28}\text{H}_{19}\text{N}_5\text{O}_4$ [$\text{M} + \text{H}$] $^+$ calculated 490.1510, found 490.1511.

2-Amino-4-(4-((1-(4-chlorophenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-5-oxo-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15b). 4-((1-(4-Chlorophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13b**, 313.74 mg, 1 mmol) was used as a reagent. Yield: 72%. White solid. Melting point: 255–258 °C. IR (KBr, ν_{\max} , cm^{-1}): 3468, 3261 ($-\text{NH}_2$), 3101 ($=\text{C}-\text{H}$), 2185 (CN), 1717 ($\text{C}=\text{O}$), 1653 ($\text{C}=\text{C}$), 1250 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.93 (s, 1H), 7.92 (d, $J = 9.2$ Hz, 2H), 7.85 (d, $J = 6.9$ Hz, 1H), 7.68–7.62 (m, 3H), 7.46–7.40 (m, 2H), 7.35 (s, 2H), 7.17 (d, $J = 8.4$ Hz, 2H), 6.98 (d, $J = 8.4$ Hz, 2H), 5.17 (s, 2H), 4.38 (s, 1H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 158.5, 157.6, 153.7, 152.6, 144.6, 136.5, 135.9, 133.5, 133.4, 130.4, 129.3, 125.2, 123.4, 123.0, 122.4, 119.9, 117.1, 115.2, 113.5, 104.8, 61.6, 58.6, 36.7. MS (ESI-QTOF) for $\text{C}_{28}\text{H}_{18}\text{ClN}_5\text{O}_4$ [$\text{M} + \text{H}$] $^+$ calculated 524.1120, found 524.1115.



2-Amino-4-(4-((1-(4-bromophenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-5-oxo-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15c). 4-((1-(4-Bromophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13c**, 358.19 mg, 1 mmol) was used as a reagent. Yield: 69%. Off-white solid. Melting point: 223–225 °C. IR (KBr, ν_{\max} , cm^{-1}): 3458, 3243 ($-\text{NH}_2$), 3096 ($=\text{C}-\text{H}$), 2189 (CN), 1703 ($\text{C}=\text{O}$), 1653 ($\text{C}=\text{C}$), 1255 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.93 (s, 1H), 7.85 (d, $J = 8.4$ Hz, 3H), 7.76 (d, $J = 9.2$ Hz, 2H), 7.66 (t, $J = 7.6$ Hz, 1H), 7.46–7.40 (m, 2H), 7.35 (s, 2H), 7.17 (d, $J = 9.2$ Hz, 2H), 6.98 (d, $J = 9.2$ Hz, 2H), 5.17 (s, 2H), 4.38 (s, 1H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 158.4, 157.6, 153.7, 152.6, 144.6, 136.5, 136.2, 133.4, 133.3, 129.4, 125.2, 123.4, 123.0, 122.6, 121.9, 119.8, 117.1, 115.2, 113.4, 104.7, 61.6, 58.6, 36.7. MS (ESI-QTOF) for $\text{C}_{28}\text{H}_{18}\text{BrN}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 568.0615, found 568.0613.

2-Amino-5-oxo-4-(4-((1-(*p*-tolyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15d). 4-((1-(*p*-Tolyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13d**, 293.32 mg, 1 mmol) was used as a reagent. Yield: 68%. White solid. Melting point: 245–247 °C. IR (KBr, ν_{\max} , cm^{-1}): 3453, 3233 ($-\text{NH}_2$), 3022 ($=\text{C}-\text{H}$), 2184 (CN), 1716 ($\text{C}=\text{O}$), 1621 ($\text{C}=\text{C}$), 1236 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.84 (s, 1H), 7.85 (d, $J = 7.6$ Hz, 1H), 7.74 (d, $J = 9.2$ Hz, 2H), 7.68–7.64 (m, 1H), 7.46–7.40 (m, 2H), 7.35 (t, $J = 3.8$ Hz, 4H), 7.17 (d, $J = 9.2$ Hz, 2H), 6.98 (d, $J = 9.2$ Hz, 2H), 5.16 (s, 2H), 4.38 (s, 1H), 2.33 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 158.5, 157.7, 153.7, 152.6, 144.3, 138.9, 136.4, 134.9, 133.4, 130.7, 129.2, 125.2, 123.2, 123.0, 120.6, 119.9, 117.1, 115.2, 113.5, 104.8, 61.6, 58.7, 36.7, 21.1. MS (ESI-QTOF) for $\text{C}_{29}\text{H}_{21}\text{N}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 504.1666, found 504.1668.

2-Amino-4-(4-((1-(4-methoxyphenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-5-oxo-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15e). 4-((1-(4-Methoxyphenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13e**, 309.32 mg, 1 mmol) was used as a reagent. Yield: 65%. Off-white solid. Melting point: 253–255 °C. IR (KBr, ν_{\max} , cm^{-1}): 3408, 3275 ($-\text{NH}_2$), 3101 ($=\text{C}-\text{H}$), 2185 (CN), 1703 ($\text{C}=\text{O}$), 1667 ($\text{C}=\text{C}$), 1245 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.78 (s, 1H), 7.86–7.65 (m, 4H), 7.46–7.40 (m, 2H), 7.35 (s, 2H), 7.17 (d, $J = 8.4$ Hz, 2H), 7.09 (d, $J = 9.2$ Hz, 2H), 6.98 (d, $J = 8.4$ Hz, 2H), 5.15 (s, 2H), 4.38 (s, 1H), 3.78 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 159.8, 158.5, 157.7, 153.7, 152.6, 144.2, 136.4, 133.4, 130.5, 129.4, 125.2, 123.3, 123.0, 122.4, 119.9, 117.1, 115.4, 115.2, 113.5, 104.8, 61.6, 58.7, 56.1, 36.7. MS (ESI-QTOF) for $\text{C}_{29}\text{H}_{21}\text{N}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 520.1615, found 520.1614.

4-(4-(1H-1,2,3-Triazol-1-yl)phenyl)-2-amino-5-oxo-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15f). 4-(1H-1,2,3-Triazol-1-yl)benzaldehyde (**3k**, 173.17 mg, 1 mmol) was used as a reagent. Yield: 70%. White solid. Melting point: 257–259 °C. IR (KBr, ν_{\max} , cm^{-1}): 3345, 3212 ($-\text{NH}_2$), 3042 ($=\text{C}-\text{H}$), 2201 (CN), 1721 ($\text{C}=\text{O}$), 1630 ($\text{C}=\text{C}$), 1236 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.23–8.21 (m, 1H), 8.12–8.06 (m, 2H), 7.93 (d, $J = 8.4$ Hz, 2H), 7.88 (d, $J = 8.4$ Hz, 1H), 7.69 (t, $J = 6.9$ Hz, 1H), 7.48–7.42 (m, 5H), 4.52 (s, 1H). ^{13}C NMR (126 MHz,

$\text{DMSO}-d_6$): δ 160.1, 158.5, 154.1, 152.7, 143.4, 138.7, 138.4, 137.0, 133.6, 133.0, 129.6, 125.2, 123.1, 119.7, 119.5, 119.2, 117.2, 113.5, 104.1, 58.1, 37.1. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{13}\text{N}_5\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 384.1091, found 384.1086.

4-(4-(1H-Tetrazol-1-yl)phenyl)-2-amino-5-oxo-4H,5H-pyrano[3,2-c]chromene-3-carbonitrile (15g). 4-(1H-Tetrazol-1-yl)benzaldehyde (**3l**, 174.16 mg, 1 mmol) was used as a reagent. Yield: 65%. Yellow solid. Melting point: 240–243 °C. IR (KBr, ν_{\max} , cm^{-1}): 3371, 3261 ($-\text{NH}_2$), 3178 ($=\text{C}-\text{H}$), 2208 (CN), 1708 ($\text{C}=\text{O}$), 1648 ($\text{C}=\text{C}$), 1250 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 9.19 (s, 1H), 8.01 (d, $J = 8.4$ Hz, 2H), 7.88 (d, $J = 8.0$ Hz, 1H), 7.69 (t, $J = 7.6$ Hz, 1H), 7.56 (d, $J = 8.4$ Hz, 2H), 7.48–7.43 (m, 4H), 4.59 (s, 1H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 160.1, 158.5, 154.4, 154.3, 152.8, 146.0, 135.6, 133.6, 130.1, 125.3, 123.1, 120.8, 119.6, 117.2, 113.5, 103.8, 57.8, 37.2. MS (ESI-QTOF) for $\text{C}_{20}\text{H}_{12}\text{N}_6\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 385.1044, found 385.1046.

2-Amino-7-methyl-5-oxo-4-(4-((1-phenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-4H,5H-pyrano[4,3-*b*]pyran-3-carbonitrile (17a). 4-((1-Phenyl)-1H-1,2,3-triazol-4-yl)methoxybenzaldehyde (**13a**, 279.29 mg, 1 mmol) was used as a reagent. Yield: 80%. Brownish red solid. Melting point: 167–170 °C. IR (KBr, ν_{\max} , cm^{-1}): 3382, 3310 ($-\text{NH}_2$), 3178 ($=\text{C}-\text{H}$), 2190 (CN), 1699 ($\text{C}=\text{O}$), 1650 ($\text{C}=\text{C}$), 1239 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.93 (s, 1H), 7.90 (d, $J = 7.6$ Hz, 2H), 7.59 (t, $J = 7.9$ Hz, 2H), 7.49 (t, $J = 7.4$ Hz, 1H), 7.14–7.11 (m, 4H), 7.01 (d, $J = 8.6$ Hz, 2H), 6.25 (s, 1H), 5.20 (s, 2H), 4.23 (s, 1H), 2.20 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 162.7, 161.3, 158.0, 157.9, 157.0, 143.9, 136.6, 136.2, 129.9, 128.7, 128.6, 122.8, 120.2, 119.4, 114.6, 101.0, 97.9, 61.1, 58.1, 35.5, 19.3. MS (ESI-QTOF) for $\text{C}_{25}\text{H}_{19}\text{N}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 454.1510, found 454.1515.

2-Amino-4-(4-((1-(4-chlorophenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-7-methyl-5-oxo-4H,5H-pyrano[4,3-*b*]pyran-3-carbonitrile (17b). 4-((1-(4-Chlorophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13b**, 313.74 mg, 1 mmol) was used as a reagent. Yield: 74%. Pale orange solid. Melting point: 200–203 °C. IR (KBr, ν_{\max} , cm^{-1}): 3408, 3340 ($-\text{NH}_2$), 3120 ($=\text{C}-\text{H}$), 2194 (CN), 1708 ($\text{C}=\text{O}$), 1644 ($\text{C}=\text{C}$), 1245 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.93 (s, 1H), 7.93–7.91 (m, 2H), 7.65–7.62 (m, 2H), 7.13 (s, 2H), 7.09 (d, $J = 8.4$ Hz, 2H), 6.97 (d, $J = 8.4$ Hz, 2H), 6.22 (s, 1H), 5.17 (s, 2H), 4.20 (s, 1H), 2.17 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 163.3, 161.9, 158.6, 158.5, 157.5, 144.6, 136.7, 135.9, 133.6, 130.4, 129.2, 123.4, 122.4, 119.9, 115.1, 101.5, 98.5, 61.5, 58.6, 35.9, 19.8. MS (ESI-QTOF) for $\text{C}_{25}\text{H}_{18}\text{ClN}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 488.1120, found 488.1122.

2-Amino-4-(4-((1-(4-bromophenyl)-1H-1,2,3-triazol-4-yl)methoxy)phenyl)-7-methyl-5-oxo-4H,5H-pyrano[4,3-*b*]pyran-3-carbonitrile (17c). 4-((1-(4-Bromophenyl)-1H-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13c**, 358.19 mg, 1 mmol) was used as a reagent. Yield: 70%. Off-white solid. Melting point: 206–209 °C. IR (KBr, ν_{\max} , cm^{-1}): 3403, 3321 ($-\text{NH}_2$), 3106 ($=\text{C}-\text{H}$), 2198 (CN), 1703 ($\text{C}=\text{O}$), 1648 ($\text{C}=\text{C}$), 1236 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.93 (s, 1H), 7.86 (d, $J = 8.4$ Hz, 2H), 7.76 (d, $J = 9.2$ Hz, 2H), 7.13–7.08 (m, 4H), 6.97 (d, $J = 9.2$ Hz,



2H), 6.22 (s, 1H), 5.17 (s, 2H), 4.20 (s, 1H), 2.17 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 163.3, 161.9, 158.6, 158.5, 157.5, 144.7, 136.7, 136.3, 133.3, 129.2, 123.4, 122.6, 121.9, 119.9, 115.1, 101.5, 98.5, 61.6, 58.6, 36.0, 19.8. MS (ESI-QTOF) for $\text{C}_{25}\text{H}_{18}\text{BrN}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 532.0615, found 532.0610.

2-Amino-7-methyl-5-oxo-4-(4-((1-(*p*-tolyl)-1*H*-1,2,3-triazol-4-yl)methoxy)phenyl)-4*H*,5*H*-pyrano[4,3-*b*]pyran-3-carbonitrile (17d). 4-((1-(*p*-Tolyl)-1*H*-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13d**, 293.32 mg, 1 mmol) was used as a reagent. Yield: 67%. Pale yellow solid. Melting point: 208–211 °C. IR (KBr, ν_{max} , cm^{-1}): 3408, 3307 (–NH₂), 3179 (=C–H), 2189 (CN), 1708 (C=O), 1658 (C=C), 1255 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.85 (s, 1H), 7.75 (d, $J = 8.4$ Hz, 2H), 7.36 (d, $J = 8.4$ Hz, 2H), 7.13 (s, 2H), 7.09 (d, $J = 9.2$ Hz, 2H), 6.98 (d, $J = 8.4$ Hz, 2H), 6.22 (s, 1H), 5.16 (s, 2H), 4.20 (s, 1H), 2.34 (s, 3H), 2.17 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 163.3, 161.9, 158.6, 158.5, 157.6, 144.3, 138.9, 136.7, 134.9, 130.8, 129.2, 123.2, 120.6, 119.9, 115.1, 101.5, 98.5, 61.6, 58.6, 36.0, 21.1, 19.8. MS (ESI-QTOF) for $\text{C}_{26}\text{H}_{21}\text{N}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 468.1666, found 468.1668.

2-Amino-4-(4-((1-(4-methoxyphenyl)-1*H*-1,2,3-triazol-4-yl)methoxy)phenyl)-7-methyl-5-oxo-4*H*,5*H*-pyrano[4,3-*b*]pyran-3-carbonitrile (17e). 4-((1-(4-Methoxyphenyl)-1*H*-1,2,3-triazol-4-yl)methoxy)benzaldehyde (**13e**, 309.32 mg, 2 mmol) was used as a reagent. Yield: 67%. Pale pink solid. Melting point: 202–205 °C. IR (KBr, ν_{max} , cm^{-1}): 3330, 3211 (–NH₂), 3106 (=C–H), 2194 (CN), 1722 (C=O), 1648 (C=C), 1245 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.99 (s, 1H), 8.93 (s, 1H), 8.38 (s, 1H), 7.97–7.91 (m, 2H), 7.64 (t, $J = 7.6$ Hz, 2H), 7.30 (d, $J = 8.4$ Hz, 1H), 7.13–7.08 (m, 2H), 6.97 (d, $J = 8.4$ Hz, 1H), 6.22 (s, 1H), 5.37 (s, 2H), 5.17 (s, 3H), 4.20 (s, 1H), 2.17 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 163.3, 161.9, 159.8, 158.6, 158.5, 157.6, 144.2, 136.7, 130.5, 129.2, 123.3, 122.4, 119.9, 115.4, 115.1, 101.5, 98.5, 61.6, 58.6, 56.1, 36.0, 19.8. MS (ESI-QTOF) for $\text{C}_{26}\text{H}_{21}\text{N}_5\text{O}_5$ $[\text{M} + \text{H}]^+$ calculated 484.1615, found 484.1615.

4-(4-(1*H*-1,2,3-Triazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4*H*,5*H*-pyrano[4,3-*b*]pyran-3-carbonitrile (17f). 4-(1*H*-1,2,3-Triazol-1-yl)benzaldehyde (**3k**, 173.17 mg, 1 mmol) was used as a reagent. Yield: 70%. Off-white solid. Melting point: 232–234 °C. IR (KBr, ν_{max} , cm^{-1}): 3372, 3309 (–NH₂), 3178 (=C–H), 2203 (CN), 1700 (C=O), 1658 (C=C), 1236 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.06 (s, 2H), 7.93 (d, $J = 8.4$ Hz, 2H), 7.36 (d, $J = 9.2$ Hz, 2H), 7.23 (s, 2H), 6.26 (s, 1H), 4.35 (s, 1H), 2.19 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 163.6, 161.9, 158.8, 158.6, 143.7, 138.7, 136.9, 129.5, 119.8, 119.2, 100.8, 98.5, 58.0, 36.4, 19.9. MS (ESI-QTOF) for $\text{C}_{18}\text{H}_{13}\text{N}_5\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 348.1091, found 348.1091.

4-(4-(1*H*-Tetrazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4*H*,5*H*-pyrano[4,3-*b*]pyran-3-carbonitrile (17g). 4-(1*H*-Tetrazol-1-yl)benzaldehyde (**3l**, 174.16 mg, 1 mmol) (**2e**, 272.15 mg, 2 mmol) was used as a reagent. Yield: 65%. Brown solid. Melting point: 227–230 °C. IR (KBr, ν_{max} , cm^{-1}): 3385, 3280 (–NH₂), 3106 (=C–H), 2185 (CN), 1690 (C=O), 1644 (C=C), 1245 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 9.20 (s, 1H), 8.01 (d, $J = 8.3$ Hz, 2H), 7.47 (d, $J = 8.3$ Hz, 2H), 7.27

(s, 2H), 6.27 (s, 1H), 4.42 (s, 1H), 2.20 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 163.8, 161.9, 159.0, 158.7, 154.4, 146.3, 135.5, 129.9, 120.8, 119.7, 100.6, 98.5, 57.7, 36.6, 19.9. MS (ESI-QTOF) for $\text{C}_{17}\text{H}_{12}\text{N}_6\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 349.1044, found 349.1047.

4.2 Anticancer activity

Cell viability assessment, colony formation assay, cell cycle analysis, DNA double-strand break evaluation, and apoptosis detection were performed following our previously reported protocol.⁴⁴ Etoposide (1 μM) served as the reference compound for cell cycle analysis and DNA double-strand break assessment. Data are expressed as mean \pm standard error of the mean. Statistical differences between control and treated groups were assessed using a one-way and two-way analysis of variance (ANOVA) followed by Tukey and Dunnett *post hoc* test using GraphPad Prism 9 software. FACS data were quantified by FlowJo software.

Spheroid formation assay. Spheroid formation and growth were assessed using agar molds prepared with the 3D Petri Dish® system (Microtissues®). Agar was liquefied in normal saline, sterilized, and cast into the molds. Once solidification, the molds were rinsed with the appropriate culture medium and placed into 24-well plates. Cells were seeded onto the molds and allowed to adhere for 20 minutes. Subsequently, an additional medium containing various concentrations of the test drugs was added. Live-cell imaging was conducted using an Olympus CellVivo Inverted Microscope, and spheroid dimensions were analyzed with Olympus CellSens software.

Ex ovo CAM assay. Fertilized chicken eggs were sourced from a local supplier and incubated at 37.5 °C with a humidity of 65% and periodic rotation for three days. On the third day, the eggs were carefully opened into sterile weighing boats, each covered with a sterile lid, and placed in a cell culture-grade CO₂ incubator maintained at 37 °C and 5% CO₂ concentration. To assess the angiogenic potential of various compounds, on the seventh day, glass filter paper discs, previously dipped in designated drug concentrations and dried, were applied to the chorioallantoic membrane CAM vascular zones. The development of the vascular networks was monitored, and images were captured periodically. These images were subsequently analysed using AngioTool software to quantify vascular growth. The CAM experiments are not considered to be animal experimentation from a regulatory point of view in Europe (EU directive 2010/63/EU), European Commission (2021/2784(RSP)). Therefore, they are exempt from Institutional Animal Ethics Committee (IAEC) approval in EU countries.

4.3 Network interaction and its ranking

4.3.1 Data collection and preparation. Fused pyrazole derivatives **8a** and **8b** demonstrated good activity *in vitro* against a breast cancer cell line MCF7. Based on this key information, various databases were utilized to identify



targets related to breast cancer. In this study, the Swiss Target Prediction database⁶⁶ was employed to screen for potential targets of compounds **8a** and **8b**. Breast cancer hits were obtained from the Disgenet,⁶⁷ Malacard⁶⁸ and OMIM⁶⁹ (accessed on October 15, 2024) using the keyword searches “breast cancer”. The overlapping targets identified from both database searches for each compound set will provide a relevant anticancer target, which is taken for further analysis.

4.3.2 Establishment of STRING network and module formation. Protein–protein interaction (PPI) data for the identified targets were generated using the STRING Database (<https://stringdb.org/>, accessed on 15 October 2024).⁷⁰ A list of target genes was provided as input to STRING, which offers valuable insights into protein interactions and correlations, assigning confidence scores to evaluate their reliability. The overlapping targets for each cancer type were entered into the database, with the *Homo sapiens* category selected for network visualization. A medium confidence threshold ranging from 0.4 to 0.7 was applied to construct the interaction network. The resulting network was then imported into Cytoscape,⁷¹ and the top 10 hub genes were identified and ranked using the CytoHubba app.⁷²

4.4 Molecular docking

Molecular docking is a protocol used to simulate the binding of a ligand to target proteins. Docking was performed using Schrödinger Maestro.⁷³ The respective protein targets, selected after ranking using CytoHubba, were retrieved from the PDB database. The proteins were prepared using a standard protocol, which involved removing crystallized water molecules, adding missing loops and side chains, and performing a short minimization step. Grid generation was performed by selecting the center of the co-crystallized ligand as the reference. This was followed by extra precision docking using the Glide module. Binding free energy calculation was done for each of the docked complexes using a prime module.⁷⁴

4.5 ADME analysis

The physicochemical properties of the compounds were obtained from the SwissADME webserver.⁷⁵ SwissADME is a freely accessible online tool that evaluates the pharmacokinetics, drug-likeness, and medicinal chemistry properties of compounds. The SMILES notation of the respective compounds **8a** and **8b** was input, and an analysis was conducted to calculate various parameters, including compliance with Lipinski's rule of five, lipophilicity, and water solubility.

Data availability

The primary data supporting this work are included in the ESI.†

Author contributions

Fabitha K. synthesized all the target compounds and performed data analysis and validation. Hiba Haneefa P. T. prepared and characterized the intermediates. Anoop Kallingal and Natalia Maciejewska conducted the biological assays. Varun Thachan Kundil conducted *in silico* studies. Fabitha K. drafted the manuscript with assistance from Anoop Kallingal and Varun Thachan Kundil. Dr. Janardhan Banothu designed the project, provided support, supervised the entire study, and reviewed, and edited the manuscript for final submission. Dr. Majed Alharbi reviewed the manuscript. All the authors approved the final version of the manuscript.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgements

JB sincerely thanks the Science and Engineering Research Board (SERB) for awarding the research grant (DST No: EEQ/2020/000303) and the Centre for Innovation, Entrepreneurship & Incubation (CIEI), NIT Calicut for the financial support under Students Innovative Project 2023–24. FK sincerely acknowledges the Ministry of Education and the National Institute of Technology Calicut for the fellowship. All the authors thanks to Varun Thachan Kundil for conducting *in silico* studeis. All the authors are grateful to the CMC, NIT Calicut for NMR characterization and the Director, NIT Calicut for providing research facilities. We sincerely thank DST-FIST for providing the HRMS facility (SR/FIST/CS-I/2019/120) to the Department of Chemistry, NIT Calicut.

References

- 1 F. Bray, M. Laversanne, H. Sung, J. Ferlay, R. L. Siegel, I. Soerjomataram and A. Jemal, *Ca-Cancer J. Clin.*, 2024, **74**, 229–263, DOI: [10.3322/caac.21834](https://doi.org/10.3322/caac.21834).
- 2 L. Wilkinson and T. Gathani, *Br. J. Radiol.*, 2021, **95**, 20211033, DOI: [10.1259/bjr.20211033](https://doi.org/10.1259/bjr.20211033).
- 3 H. Sung, J. Ferlay, R. L. Siegel, M. Laversanne, I. Soerjomataram, A. Jemal and F. Bray, *Ca-Cancer J. Clin.*, 2021, **71**, 209–249, DOI: [10.3322/caac.21660](https://doi.org/10.3322/caac.21660).
- 4 N. Kerru, P. Singh, N. Koorbanally, R. Raj and V. Kumar, *Eur. J. Med. Chem.*, 2017, **142**, 179–212, DOI: [10.1016/j.ejmech.2017.07.033](https://doi.org/10.1016/j.ejmech.2017.07.033).
- 5 S. Sharmin, M. M. Rahaman, M. Martorell, J. Sastre-Serra, J. Sharif-Rad, M. Butnariu, I. C. Bagiu, R. V. Bagiu and M. T. Islam, *Cancer Cell Int.*, 2021, **21**, 1–19, DOI: [10.1186/s12935-021-02309-9](https://doi.org/10.1186/s12935-021-02309-9).
- 6 F. Ye, S. Dewanjee, Y. Li, N. K. Jha, Z.-S. Chen, A. Kumar, Vishakha, T. Behl, S. K. Jha and H. Tang, *Mol. Cancer*, 2023, **22**, 105, DOI: [10.1186/s12943-023-01805-y](https://doi.org/10.1186/s12943-023-01805-y).



- 7 J. Wang, K. Lei and F. Han, *Riv. Eur. Sci. Med. Farmacol.*, 2018, **22**, 3855–3864, DOI: [10.26355/eurrev.201806_15270](https://doi.org/10.26355/eurrev.201806_15270).
- 8 S. A. Patil, R. Patil, L. M. Pfeffer and D. D. Miller, *Future Med. Chem.*, 2013, **5**, 1647–1660, DOI: [10.4155/fmc.13.126](https://doi.org/10.4155/fmc.13.126).
- 9 A. Chaudhary, K. Singh, N. Verma, S. Kumar, D. Kumar and P. P. Sharma, *Mini-Rev. Med. Chem.*, 2022, **22**, 2736–2751, DOI: [10.2174/1389557522666220331161636](https://doi.org/10.2174/1389557522666220331161636).
- 10 M. Costa, T. A. Dias, A. Brito and F. Proença, *Eur. J. Med. Chem.*, 2016, **123**, 487–507, DOI: [10.1016/j.ejmech.2016.07.057](https://doi.org/10.1016/j.ejmech.2016.07.057).
- 11 J. Li, X. Wang, Y. Fang and C. Wang, *J. Asian Nat. Prod. Res.*, 2010, **12**, 992–1000, DOI: [10.1080/10286020.2010.513034](https://doi.org/10.1080/10286020.2010.513034).
- 12 J. Du, F. Jiang, S. Xu, Z. Huang, L. L. Chen and L. Li, *J. Cancer*, 2021, **12**, 270–280, DOI: [10.7150/jca.50360](https://doi.org/10.7150/jca.50360).
- 13 H. Ekowati, I. Astuti and Mustofa, *Indones. J. Chem.*, 2010, **10**, 247–251, DOI: [10.22146/ijc.21467](https://doi.org/10.22146/ijc.21467).
- 14 N. Guilbaud, L. Kraus-Berthier, F. Meyer-Losic, V. Malivet, C. Chacun, M. Jan, F. Tillequin, S. Michel, M. Koch, B. Pfeiffer, G. Atassi, J. Hickman and A. Pierré, *Clin. Cancer Res.*, 2001, **7**, 2573–2580.
- 15 H. Nishino, T. Okuyama, M. Takata, S. Shibata, H. Tokuda, J. Takayasu, T. Hasegawa, A. Nishino, H. Ueyama and A. Iwashima, *Carcinogenesis*, 1990, **11**, 1557–1561, DOI: [10.1093/carcin/11.9.1557](https://doi.org/10.1093/carcin/11.9.1557).
- 16 V. Raj and J. Lee, *Front. Chem.*, 2020, **8**, 1–23, DOI: [10.3389/fchem.2020.00623](https://doi.org/10.3389/fchem.2020.00623).
- 17 T. Kottha and V. Gnanasambandam, *ChemistrySelect*, 2018, **3**, 8381–8386, DOI: [10.1002/slct.201801210](https://doi.org/10.1002/slct.201801210).
- 18 W. Kemnitzer, J. Drewe, S. Jiang, H. Zhang, Y. Wang, J. Zhao, S. Jia, J. Herich, D. Labreque, R. Storer, K. Meerovitch, D. Bouffard, R. Rej, R. Denis, C. Blais, S. Lamothe, G. Attardo, H. Gourdeau, B. Tseng, S. Kasibhatla and S. X. Cai, *J. Med. Chem.*, 2004, **47**, 6299–6310, DOI: [10.1021/jm049640t](https://doi.org/10.1021/jm049640t).
- 19 Y. Liu, D. Yang, Z. Hong, S. Guo, M. Liu, D. Zuo, D. Ge, M. Qin and D. Sun, *Eur. J. Med. Chem.*, 2018, **146**, 185–193, DOI: [10.1016/j.ejmech.2018.01.052](https://doi.org/10.1016/j.ejmech.2018.01.052).
- 20 S. Kasibhatla, H. Gourdeau, K. Meerovitch, J. Drewe, S. Reddy, L. Qiu, H. Zhang, F. Bergeron, D. Bouffard, Q. Yang, J. Herich, S. Lamothe, S. X. Cai and B. Tseng, *Mol. Cancer Ther.*, 2004, **3**, 1365–1374, DOI: [10.1158/1535-7163.1365.3.11](https://doi.org/10.1158/1535-7163.1365.3.11).
- 21 H. Gourdeau, L. Leblond, B. Hamelin, C. Desputeau, K. Dong, I. Kianicka, D. Custeau, C. Boudreau, L. Geerts, S.-X. Cai, J. Drewe, D. Labrecque, S. Kasibhatla and B. Tseng, *Mol. Cancer Ther.*, 2004, **3**, 1375–1383, DOI: [10.1158/1535-7163.1375.3.11](https://doi.org/10.1158/1535-7163.1375.3.11).
- 22 I. A. Schepetkin, L. N. Kirpotina, A. I. Khlebnikov, N. Cheng, R. D. Ye and M. T. Quinn, *Biochem. Pharmacol.*, 2014, **92**, 627–641, DOI: [10.1016/j.bcp.2014.09.027](https://doi.org/10.1016/j.bcp.2014.09.027).
- 23 A. Alneyadi, Z. N. Nizami, H. E. Aburawi, S. Hisaindee, M. Nawaz, S. Attoub, G. Ramadan, N. Benhalilou, M. Al Azzani, Y. Elmahi, A. Almqbali, K. Muhammad, A. H. Eid, R. Vijayan and R. Iratni, *Cancers*, 2023, **15**, 2682, DOI: [10.3390/cancers15102682](https://doi.org/10.3390/cancers15102682).
- 24 L. M. Al-harbi, E. A. Al-Harbi, R. M. Okasha, R. A. El-Eisawy, M. A. A. El-nassag, H. M. Mohamed, A. M. Fouda, A. A. Elhenawy, A. Mora, A. M. El-Agrody and H. K. A. El-Mawgoud, *J. Enzyme Inhib. Med. Chem.*, 2023, **38**, 2155814, DOI: [10.1080/14756366.2022.2155814](https://doi.org/10.1080/14756366.2022.2155814).
- 25 T. H. Afifi, R. M. Okasha, H. E. A. Ahmed, J. Ilaš, T. Saleh and A. S. Abd-El-Aziz, *EXCLI J.*, 2017, **16**, 868–902, DOI: [10.17179/excli2017-356](https://doi.org/10.17179/excli2017-356).
- 26 M. H. Ahagh, G. Dehghan, M. Mehdipour, R. Teimuri-Mofrad, E. Payami, N. Sheibani, M. Ghaffari and M. Asadi, *Bioorg. Chem.*, 2019, **93**, 103329, DOI: [10.1016/j.bioorg.2019.103329](https://doi.org/10.1016/j.bioorg.2019.103329).
- 27 H. E. A. Ahmed, M. A. A. El-Nassag, A. H. Hassan, R. M. Okasha, S. Ihmaid, A. M. Fouda, T. H. Afifi, A. Aljuhani and A. M. El-Agrody, *J. Enzyme Inhib. Med. Chem.*, 2018, **33**, 1074–1088, DOI: [10.1080/14756366.2018.1476503](https://doi.org/10.1080/14756366.2018.1476503).
- 28 T. C. Braga, M. M. Silva, E. O. O. Nascimento, E. C. D. da Silva, Y. F. Rego, M. Mandal, Z. A. de Souza, A. L. T. G. Ruiz, J. E. de Carvalho, F. T. Martins, I. M. Figueiredo, T. M. de Aquino, C. M. da Silva, B. Mandal, G. Brahmachari, J. C. C. Santos and A. de Fatima, *Eur. J. Med. Chem. Rep.*, 2022, **4**, 100030, DOI: [10.1016/j.ejmcr.2022.100030](https://doi.org/10.1016/j.ejmcr.2022.100030).
- 29 P. Rawat and S. M. Verma, *Drug Des., Dev. Ther.*, 2016, **10**, 2779–2788, DOI: [10.2147/DDDT.S111266](https://doi.org/10.2147/DDDT.S111266).
- 30 R. Soni, S. D. Durgapal, S. S. Soman and J. J. George, *Arabian J. Chem.*, 2019, **12**, 701–708, DOI: [10.1016/j.arabjc.2016.11.011](https://doi.org/10.1016/j.arabjc.2016.11.011).
- 31 W. Zhao, B. Wang, Y. Liu, L. Fu, L. Sheng, H. Zhao, Y. Lu and D. Zhang, *Eur. J. Med. Chem.*, 2020, **189**, 112075, DOI: [10.1016/j.ejmech.2020.112075](https://doi.org/10.1016/j.ejmech.2020.112075).
- 32 M. Mashhadinezhad, M. Mamaghani, M. Rassa and F. Shirini, *ChemistrySelect*, 2019, **4**, 4920–4932, DOI: [10.1002/slct.201900405](https://doi.org/10.1002/slct.201900405).
- 33 Y. Kanbe, M. Kim, M. Nishimoto, Y. Ohtake, N. Kato, T. Tsunenari, K. Taniguchi, I. Ohizumi, S. Kaiho, K. Morikawa, J. Jo, H. Limb and H. Kimb, *Bioorg. Med. Chem.*, 2006, **14**, 4803–4819, DOI: [10.1016/j.bmc.2006.03.020](https://doi.org/10.1016/j.bmc.2006.03.020).
- 34 K. Takao, U. Shiori, H. Kamauchi and Y. Sugita, *Bioorg. Chem.*, 2019, **87**, 594–600, DOI: [10.1016/j.bioorg.2019.03.042](https://doi.org/10.1016/j.bioorg.2019.03.042).
- 35 N. Kerru, L. Gummidi, S. Maddila, K. K. Gangu and S. B. Jonnalagadda, *Molecules*, 2020, **25**, 1909, DOI: [10.3390/molecules25081909](https://doi.org/10.3390/molecules25081909).
- 36 E. Vitaku, D. T. Smith and J. T. Njardarson, *J. Med. Chem.*, 2014, **57**, 10257–10274, DOI: [10.1021/jm501100b](https://doi.org/10.1021/jm501100b).
- 37 H. B. Broughton and I. A. Watson, *J. Mol. Graphics Modell.*, 2004, **23**, 51–58, DOI: [10.1016/j.jmgn.2004.03.016](https://doi.org/10.1016/j.jmgn.2004.03.016).
- 38 P. Telukuntla, M. Chandrakanth, P. G. Amrutha, N. M. Thomas, R. Gondru, K. R. Valluru and J. Banothu, *Tetrahedron Lett.*, 2024, **146**, 155180, DOI: [10.1016/j.tetlet.2024.155180](https://doi.org/10.1016/j.tetlet.2024.155180).
- 39 R. Deshineni, C. G. Arya, J. Banothu, R. Velpula and G. Chellamella, *Chem. Data Collect.*, 2020, **30**, 100546, DOI: [10.1016/j.cdc.2020.100546](https://doi.org/10.1016/j.cdc.2020.100546).
- 40 J. Banothu, R. Gondru, R. Bavantula, Y. Velivela and P. A. Crooks, *Bioorg. Med. Chem. Lett.*, 2015, **25**, 106–112, DOI: [10.1016/j.bmcl.2014.10.100](https://doi.org/10.1016/j.bmcl.2014.10.100).
- 41 R. Gali, J. Banothu, M. Porika, R. Velpula, R. Bavantula and S. Abbagani, *RSC Adv.*, 2014, **4**, 53812–53819, DOI: [10.1039/C4RA11428K](https://doi.org/10.1039/C4RA11428K).



- 42 R. Gali, J. Banothu, M. Porika, R. Velpula, S. Hnamte, R. Bavantula, S. Abbagani and S. Busi, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 4239–4242, DOI: [10.1016/j.bmcl.2014.07.030](https://doi.org/10.1016/j.bmcl.2014.07.030).
- 43 K. Fabitha, C. G. Arya, M. Chandrakanth and J. Banothu, *Res. Chem. Intermed.*, 2023, **49**, 997–1014, DOI: [10.1007/s11164-022-04929-w](https://doi.org/10.1007/s11164-022-04929-w).
- 44 K. Fabitha, A. Kallingal, N. Maciejewska, C. G. Arya, M. Chandrakanth, N. M. Thomas, Y. Li, R. Gondru, M. Munikumar and J. Banothu, *New J. Chem.*, 2024, **48**, 8038–8054, DOI: [10.1039/d4nj00824c](https://doi.org/10.1039/d4nj00824c).
- 45 K. Fabitha, C. G. Arya, P. Shyam, R. Gondru and J. Banothu, *Polycyclic Aromat. Compd.*, 2022, **43**, 7093–7109, DOI: [10.1080/10406638.2022.2130369](https://doi.org/10.1080/10406638.2022.2130369).
- 46 S. Wang, Y. Tong, T. Ng, L. Lao, J. K. W. Lam, K. Y. Zhang, Z. Zhang and S. C. W. Sze, *Chin. Med.*, 2015, **10**, 1–12, DOI: [10.1186/s13020-015-0051-z](https://doi.org/10.1186/s13020-015-0051-z).
- 47 A. L. Hopkins, *Nat. Biotechnol.*, 2007, **25**, 1110–1111, DOI: [10.1038/nbt1007-1110](https://doi.org/10.1038/nbt1007-1110).
- 48 A. S. Azmi, *Future Med. Chem.*, 2012, **4**, 939–941, DOI: [10.4155/fmc.12.44](https://doi.org/10.4155/fmc.12.44).
- 49 X. Liang, H. Li and S. Li, *Mol. BioSyst.*, 2014, **10**, 1014–1022, DOI: [10.1039/c3mb70507b](https://doi.org/10.1039/c3mb70507b).
- 50 J. C. Oliveros, (2007–2015) Venny, An Interactive Tool for Comparing Lists with Venn's Diagrams, <https://bioinfo.pcnb.csic.es/tools/venny/index.html>.
- 51 A. Prat and C. M. Perou, *Mol. Oncol.*, 2010, **5**, 5–23, DOI: [10.1016/j.molonc.2010.11.003](https://doi.org/10.1016/j.molonc.2010.11.003).
- 52 X. Dai, T. Li, Z. Bai, Y. Yang, X. Liu, J. Zhan and B. Shi, *Am. J. Cancer Res.*, 2015, **5**, 2929–2943.
- 53 P. Wee and Z. Wang, *Cancers*, 2017, **9**, 52, DOI: [10.3390/cancers9050052](https://doi.org/10.3390/cancers9050052).
- 54 H. Masuda, D. Zhang, C. Bartholomeusz, H. Doihara, G. N. Hortobagyi and N. T. Ueno, *Breast Cancer Res. Treat.*, 2012, **136**, 331–345, DOI: [10.1007/s10549-012-2289-9](https://doi.org/10.1007/s10549-012-2289-9).
- 55 S. Jahani-Sherafat, M. Azimirad, H. Raeisi, M. A. Looha, S. Tavakkoli, H. A. Amoli, S. Moghim, M. Rostami-Nejad, A. Yadegar and M. R. Zali, *Mol. Biol. Rep.*, 2024, **51**, 265, DOI: [10.1007/s11033-024-09273-3](https://doi.org/10.1007/s11033-024-09273-3).
- 56 A. E. Maennling, M. K. Tur, M. Niebert, T. Klockenbring, F. Zeppernick, S. Gattenlöhner, I. Meinhold-Heerlein and A. F. Hussain, *Cancers*, 2019, **11**, 1826, DOI: [10.3390/cancers11121826](https://doi.org/10.3390/cancers11121826).
- 57 T. Fang, Y. Zhang, V. Y. Chang, M. Roos, C. M. Termini, L. Sigmaevskaia, M. Quarmyne, P. K. Lin, A. Pang, J. Kan, X. Yan, A. Javier, K. Pohl, L. Zhao, P. Scott, H. A. Himgurg and J. P. Chute, *Blood*, 2020, **136**, 441–454, DOI: [10.1182/blood.2020005895](https://doi.org/10.1182/blood.2020005895).
- 58 D. L. Wheeler, M. Iida and E. F. Dunn, *Oncologist*, 2009, **14**, 667–678, DOI: [10.1634/theoncologist.2009-0009](https://doi.org/10.1634/theoncologist.2009-0009).
- 59 Y. Fukumoto, M. Morii, T. Miura, S. Kubota, K. Ishibashi, T. Honda, A. Okamoto, N. Yamaguchi, A. Iwama, Y. Nakayama and N. Yamaguchi, *J. Biol. Chem.*, 2014, **289**, 12313–12329, DOI: [10.1074/jbc.M113.533752](https://doi.org/10.1074/jbc.M113.533752).
- 60 W. Wu, L. M. Graves, G. N. Gill, S. J. Parsons and J. M. Samet, *J. Biol. Chem.*, 2002, **277**, 24252–24257, DOI: [10.1074/jbc.m200437200](https://doi.org/10.1074/jbc.m200437200).
- 61 K. Park, H. Yang, S. Youn, H. Yang, I. Chae, B. Oh, M. Lee, Y. Park, Y. Choi, H. Kim and K. Walsh, *Arterioscler., Thromb., Vasc. Biol.*, 2003, **23**, 1364–1369, DOI: [10.1161/01.ATV.0000081633.53390.B4](https://doi.org/10.1161/01.ATV.0000081633.53390.B4).
- 62 E. Beurel and R. S. Jope, *Prog. Neurobiol.*, 2006, **79**, 173–189, DOI: [10.1016/j.pneurobio.2006.07.006](https://doi.org/10.1016/j.pneurobio.2006.07.006).
- 63 Y. Yang, T. Lei, S. Du, R. Tong, H. Wang, J. Yang, J. Huang, M. Sun, Y. Wang and Z. Dong, *Int. J. Oncol.*, 2018, **52**, 709–720, DOI: [10.3892/ijo.2018.4237](https://doi.org/10.3892/ijo.2018.4237).
- 64 R. He, S. Du, T. Lei, X. Xie and Y. Wang, *Oncol. Rep.*, 2020, **44**, 2373–2385, DOI: [10.3892/or.2020.7817](https://doi.org/10.3892/or.2020.7817).
- 65 J. Luo, *Cancer Lett.*, 2009, **273**, 194–200, DOI: [10.1016/j.canlet.2008.05.045](https://doi.org/10.1016/j.canlet.2008.05.045).
- 66 A. Daina, O. Michielin and V. Zoete, *Nucleic Acids Res.*, 2019, **47**, W357–W364, DOI: [10.1093/nar/gkz382](https://doi.org/10.1093/nar/gkz382).
- 67 J. Piñero, J. M. Ramírez-Anguaita, J. Saüch-Pitarch, F. Ronzano, E. Centeno, F. Sanz and L. I. Furlong, *Nucleic Acids Res.*, 2020, **48**, D845–D855, DOI: [10.1093/nar/gkz1021](https://doi.org/10.1093/nar/gkz1021).
- 68 N. Rappaport, N. Nativ, G. Stelzer, M. Twik, Y. Guan-Golan, T. I. Stein, I. Bahir, F. Belinky, C. P. Morrey, M. Safran and D. Lancet, *Database*, 2013, **2013**, bat018, DOI: [10.1093/database/bat018](https://doi.org/10.1093/database/bat018).
- 69 A. Hamosh, A. F. Scott, J. S. Amberger, C. A. Bocchini and V. A. McKusick, *Nucleic Acids Res.*, 2005, **33**, D514–D517, <https://pubmed.ncbi.nlm.nih.gov/articles/PMC539987/>.
- 70 D. Szklarczyk, R. Kirsch, M. Koutrouli, K. Nastou, F. Mehryary, R. Hachilif, A. L. Gable, T. Fang, N. T. Doncheva, S. Pyysalo, P. Bork, L. J. Jensen and C. V. Mering, *Nucleic Acids Res.*, 2023, **51**, D638–D646, <https://pubmed.ncbi.nlm.nih.gov/36370105/>.
- 71 P. Shannon, A. Markiel, O. Ozier, N. S. Baliga, J. T. Wang, D. Ramage, N. Amin, B. Schwikowski and T. Ideker, *Genome Res.*, 2003, **13**, 2498–2504, <https://pubmed.ncbi.nlm.nih.gov/articles/PMC403769/>.
- 72 C. Chin, S. Chen, H. Wu, C. Ho, M. Ko and C. Lin, *BMC Syst. Biol.*, 2014, **8**, 1–7, <https://pubmed.ncbi.nlm.nih.gov/25521941/>.
- 73 Y. Yang, K. Yao, M. P. Repasky, K. Leswing, R. Abel, B. K. Shoichet and S. V. Jerome, *J. Chem. Theory Comput.*, 2021, **17**, 7106–7119, DOI: [10.1021/acs.jctc.1c00810](https://doi.org/10.1021/acs.jctc.1c00810).
- 74 M. P. Jacobson, D. L. Pincus, C. S. Rapp, T. J. F. Day, B. Honig, D. E. Shaw and R. A. Friesner, *Proteins: Struct., Funct., Bioinf.*, 2004, **55**, 351–367, DOI: [10.1002/prot.10613](https://doi.org/10.1002/prot.10613).
- 75 A. Daina, O. Michielin and V. Zoete, *Sci. Rep.*, 2017, **7**, 42717, DOI: [10.1038/srep42717](https://doi.org/10.1038/srep42717).

