



Cite this: DOI: 10.1039/d4nj00824c

Novel fused pyran derivatives induce apoptosis and target cell cycle progression in anticancer efficacy against multiple cell lines†

 K. Fabitha,^a Anoop Kallingal,^b Natalia Maciejewska,^b C. G. Arya,^a Munugala Chandrakanth,^a Neethu Mariam Thomas,^a Yupeng Li,^c Ramesh Gondru,^d Manne Munikumar^d and Janardhan Banothu^{*a}

Nitrogen-based heterocycles such as pyrazole, imidazole, 1,2,4-triazole, benzimidazole, and benzotriazole substituted fused pyran derivatives (**6a–e**, **8a–e**, **10a–e**, **12a–e**, & **14a–e**) have been synthesized and tested for their *in vitro* anticancer efficacies against MCF7, A549, and HCT116 cancer cell lines. Among the compounds, **6e**, **14b**, and **8c** were identified as the most potent against MCF7, A549, and HCT116, with IC₅₀ values of 12.46 ± 2.72 μM, 0.23 ± 0.12 μM, and 7.58 ± 1.01 μM, respectively. Further studies demonstrated that these compounds can change cellular and nuclear morphology and inhibit colony formation in the tested cancer cells. They also remarkably block/inhibit the cell cycle progression of cancer cells at various phases. DNA damage analysis and apoptosis studies revealed that these compounds have the potential to induce DNA double-strand breaks and apoptosis. *In silico* absorption, distribution, metabolism, excretion, and toxicity (ADMET) properties of the potent compounds were assessed, revealing that all the compounds exhibited favorable pharmacokinetic and toxicological properties. The potent compounds identified from this study can be considered as a lead for further drug design and development.

 Received 20th February 2024,
Accepted 16th March 2024

DOI: 10.1039/d4nj00824c

rsc.li/njc

1. Introduction

Cancer, a multifaceted and relentless disease, continues to pose a significant global health challenge, affecting millions of lives worldwide.^{1–3} With its diverse range of manifestations and inherent heterogeneity, cancer remains a major focus of scientific investigation, necessitating a comprehensive understanding of its underlying mechanisms and the development of effective therapeutic strategies.^{1–4} One of the hallmarks of cancer is uncontrolled cell growth, driven by alterations in crucial cellular pathways that regulate proliferation and apoptosis. Genetic mutations disrupt the delicate balance between

cell division and cell death, leading to the uncontrolled proliferation of abnormal cells.^{5–7}

Fused pyran derivatives have emerged as promising candidates in cancer therapeutics due to their ability to target the cellular pathways, resulting in the inhibition of cancer cell growth, induction of apoptosis, and disruption of essential cellular processes that are crucial for tumor progression.^{8–20} These moieties have also been reported to exhibit a wide array of pharmacological activities that include antimicrobial,^{21–27} antitubercular,²¹ antioxidant,^{24,28} analgesic,²⁵ antileishmanial,²⁹ antiplatelet,³⁰ formyl peptide receptor 1 (FPR1) antagonist,³¹ and anticonvulsant activities.³² In addition, they were also identified as potent selective estrogen receptor modulators (SERMs),³³ inhibitors of enzymes such as acetylcholinesterase,²⁸ monoamine oxidase,^{34,35} topoisomerase 2,³⁶ c-Src kinase,³⁷ xanthine oxidase,³⁸ and aldehyde reductase 2 (ALR2).³⁹ Structures and pharmacological applications of a few fused pyran derivatives are summarized in Fig. 1. Of these compounds, the chromene analog EPC2407 (Crolibulin, **I**) is the most promising vascular disrupting agent and apoptosis inducer for treating advanced solid tumors.^{10,14,40,41} In phase I clinical trials in patients with advanced thoracic and abdominal tumors, cell swelling and decreased tumor perfusion monitored by functional MRI imaging modalities 2–3 days

^a Department of Chemistry, National Institute of Technology Calicut, Kozhikode 673601, Kerala, India. E-mail: janardhan@nitc.ac.in; Tel: +91 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 Pharmaceutical Sciences, School of Pharmacy and Border Biomedical Research Center, The University of Texas at El Paso, El Paso, TX 79968, USA

^d Indian Council of Medical Research, National Institute of Nutrition, Hyderabad 500007, Telangana, India

 † Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4nj00824c>

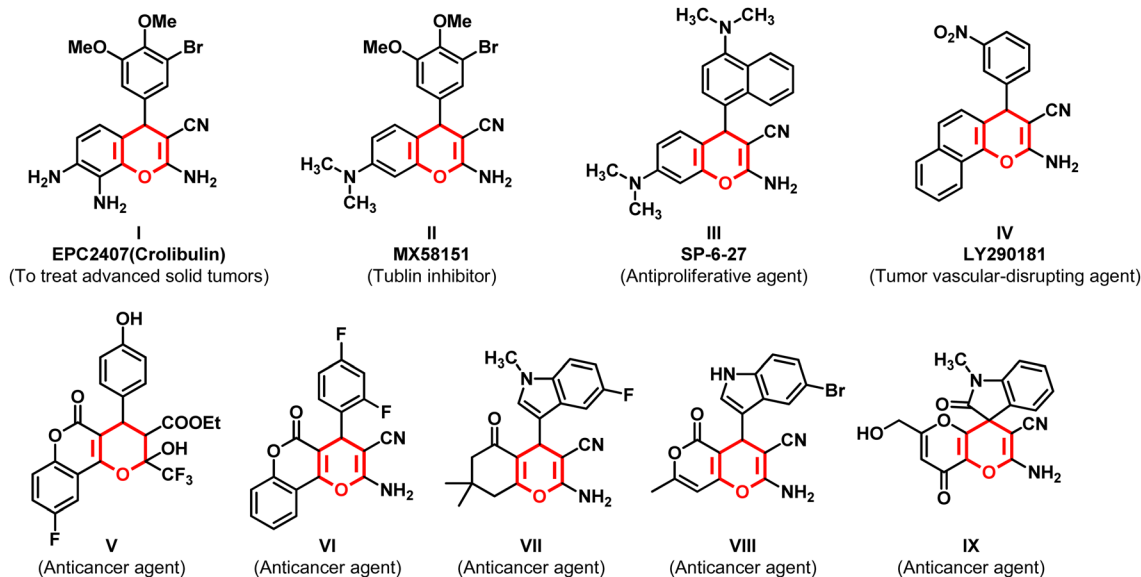



Fig. 1 Structures of potential chemotherapeutic pyran derivatives available in the clinic (I–IV) and several reported pyran derivatives as anticancer agents (V–IX).

post-treatment with crolibulin were confirmed.⁴¹ It is now under phase I/II clinical trials in combination with cisplatin for anaplastic thyroid cancer (ATC).^{14,40} Similarly, MX58151 (II) was reported as a caspase activator and tubulin inhibitor,^{15,16} and SP-6-27 (III) displayed the highest potency towards glioma, melanoma, and prostate cancer cell lines, and is known for its high antiproliferative activity.¹⁴ Another chromene analog LY290181 (IV) has been identified as a potent tumor vascular-disrupting agent and acts as an inhibitor for mitosis and microtubules.^{18,42,43} Likewise, fluorine-containing pyrano-chromenes (V)²⁰ and (VI),¹⁹ indole-substituted tetrahydro-chromene (VII),¹⁴ and indole-tethered pyrano-pyrans (VIII)⁴⁴ and (IX)⁴⁵ have been reported as promising anticancer agents. Despite the progress in the development of chemotherapeutic agents, several new target-specific and improved anticancer agents with minimum/no side effects are essential to tackle drug resistance.^{46–48}

According to the U.S. Food and Drug Administration (FDA) database, nearly 60% of small-molecule drugs are nitrogen-containing heterocycles.⁴⁹ Incorporation of nitrogen atoms or N-based heterocycles in a pharmacophore such as a fused pyran may not only increase its water solubility but also enhance the binding to a variety of biological targets such as enzymes and receptors.^{49,50} The structural versatility of fused pyran derivatives enables the synthesis of compounds with tailored properties, optimizing their anticancer efficacy and minimizing off-target effects. By introducing diverse functional groups and modifying specific regions of the fused pyran scaffold, researchers have enhanced their potency and improved their pharmacokinetic profiles, allowing for improved bioavailability and target specificity.^{14–20} Therefore, we aimed to synthesize a series of novel fused pyran derivatives containing N-based heterocycles to examine their anticancer efficacies against different cancer cell lines, and study their mechanisms of

action and *in silico* ADMET predictions to evaluate their drug-likeness properties.

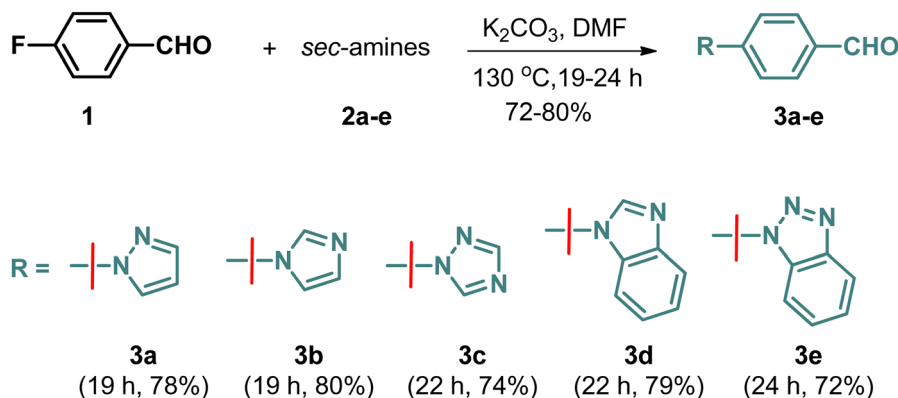
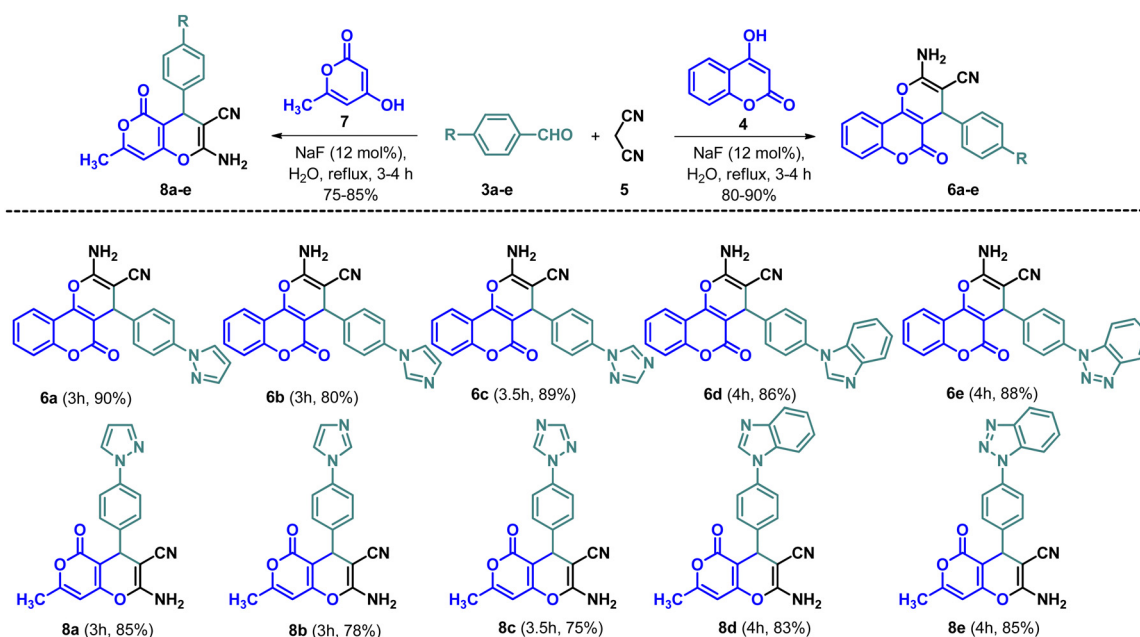
2. Results and discussion

2.1 Chemical synthesis of pyran/chromene derivatives

A multicomponent Knoevenagel–Michael reaction of α -naphthol/ β -naphthol/4-hydroxycoumarin with aryl/heteroaryl aldehydes and malononitrile in H₂O in the presence of sodium fluoride (NaF) catalyst under microwave irradiation resulted in the formation of diverse pyran/chromene derivatives.⁵¹ In continuation, we have synthesized a variety of N-based heterocycles substituted fused pyran derivatives using the strategy shown in Schemes 2 and 3. The intermediates, N-based heterocycles substituted aryl aldehydes (3a–e) were obtained by the S_NAr reactions of 4-fluorobenzaldehyde (1) with secondary amines such as pyrazole (2a), imidazole (2b), 1,2,4-triazole (2c), benzimidazole (2d), benzotriazole (2e) in the presence of potassium carbonate in DMF at 130 °C (Scheme 1). The desired products (6a–e, 8a–e, 10a–e, 12a–e & 14a–e) were obtained by treating 4-hydroxycoumarin (4), 4-hydroxy-6-methyl-2H-pyran-2-one (7), 5-hydroxy-2-(hydroxymethyl)-4H-pyran-4-one (9), cyclohexane-1,3-dione (11) and 5,5-dimethylcyclohexane-1,3-dione (13) individually with N-based heterocycles substituted benzaldehydes (3a–e) and malononitrile (5) in H₂O utilizing 12 mol% of NaF as a catalyst (Schemes 2 and 3). The reaction was carried out in H₂O at the reflux temperature. All the compounds were obtained in 75–90% of the yields within 3–4 hours.

Structures of all the target molecules 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 4.30–4.90 ppm confirms the presence of the pyran-C4 proton. Moreover, the presence of a peak at 30–39 ppm in



Scheme 1 Synthesis of N-based heterocycles substituted aryl aldehydes (**3a-e**).Scheme 2 Synthesis of novel fused pyran derivatives (**6a-e** & **8a-e**).

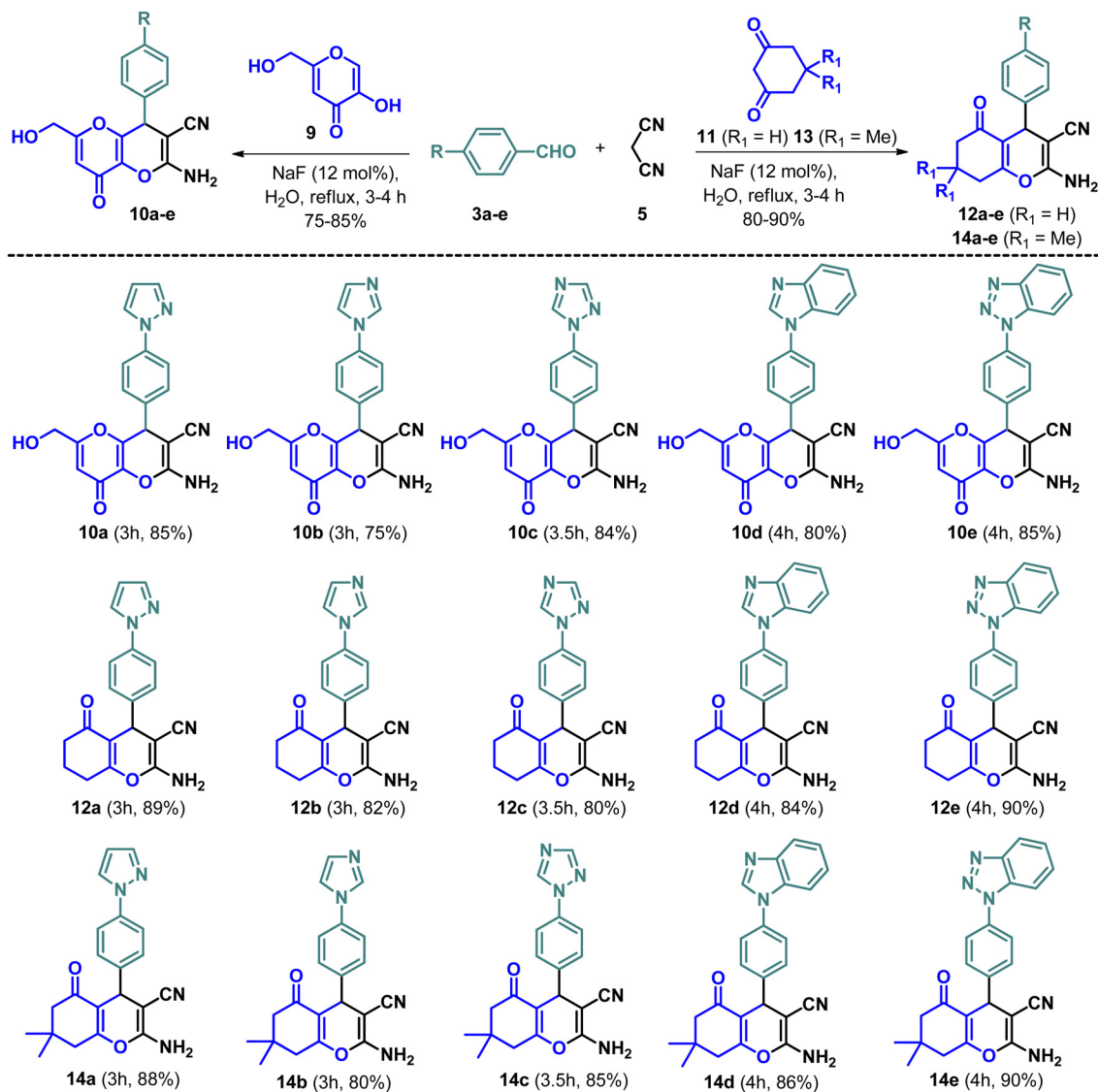
the ^{13}C NMR spectra corresponding to the pyran-C4 carbon provided additional evidence of the formation of the pyran ring. Furthermore, the molecular ion peak from the mass spectra also confirmed the product formation. The purity of all tested compounds was greater than 99.5% determined by LC-HRMS.

2.2 Impact of the fused pyran derivatives on the viability of cancer cells

To assess the impact of the synthesized fused pyran derivatives on the viability of cancer cells, an MTT ((3-[4,5-dimethylthiazol-2-yl]-2,5 diphenyl tetrazolium bromide) assay was conducted in three different cancer cell lines MCF7 (breast adenocarcinoma), HCT116 (colorectal carcinoma), and A549 (lung adenocarcinoma) and the results were compared with standard drug cisplatin. The results, presented in Table 1, provide valuable

insights into the compounds' half-maximal inhibitory concentration (IC_{50}) values for each tested cell line. Among the compounds tested, compounds **6e**, **12a**, and **14b** exhibited promising results in inhibiting cellular growth in MCF7 cells (Fig. 2). These compounds demonstrated significant inhibitory effects on the proliferation of MCF7 cells, indicating their potential in treating breast adenocarcinoma compared to cisplatin. Similarly, compounds **12a**, **12c**, **12d**, **14b**, **14c**, and **14d** exhibited notable anti-proliferation activity on A549 cells, and compounds **8c** and **8e** displayed inhibitory effects specifically on HCT116 cells. They may serve as potential candidates for further optimization to treat lung adenocarcinoma and colorectal carcinoma, respectively. It is also noted that some compounds such as **12a** and **14b** showed cell toxicity in both MCF7 and A549 cells, indicating their broad-spectrum potential against multiple cancer cell lines.





Scheme 3 Synthesis of novel fused pyran derivatives (**10a–e**, **12a–e** & **14a–e**).

2.3 Effect of the fused pyran derivatives on cancer cell morphology and colony formation

To evaluate the changes caused by the most toxic fused pyran derivatives in cancer cells and nuclear morphology, the morphological analysis using microscopic techniques was performed with the results shown in Fig. 3. After a 48-hour incubation with the tested compounds at their corresponding IC_{50} concentrations, all compounds induced cytoplasmic morphological alterations (Fig. 3(A)–(C)), suggesting their cytotoxic effect on cellular structures and functions in the cancer cells. Interestingly, it was observed that all compounds, except **14b** in MCF7, induced the formation of micronuclei and apoptotic-like nuclear bodies (indicated by red arrows in Fig. 3(A)–(C)). This phenomenon indicates that the compounds have the potential to cause nucleic acid damage and potentially activate the pathways leading to cell death in the tested cell lines. Furthermore, compound **14b** exhibited notable variations in its effects

across two different cell lines. In MCF7 cells, compound **14b** did not induce the formation of apoptotic-like nuclear bodies and micronuclei. However, in the A549 cell line, exposure to **14b** resulted in a significant increase in apoptotic nuclear abnormalities. Interestingly, these nuclear changes were accompanied by a distinct phenomenon of cytoplasmic shrinking, suggesting a comprehensive cellular response specific to the A549 cell line upon exposure to compound **14b**.

The colony formation analysis showed a significant decrease in colony formation after treatment with the fused pyran derivatives (Fig. 3(D)). Notably, compound **8c** exhibited the highest inhibition rate ($70.0 \pm 4.96\%$) against HCT116 cells when treated with its IC_{50} concentration. Additionally, compounds **6e** and **14b** showed considerable inhibitions in colony formation with values of $66.4 \pm 4.50\%$ and $71.6 \pm 2.01\%$ for the MCF7 and A549 cell lines respectively (Fig. 3(E)). These results suggest that these molecules possess the ability to hinder the



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

Cell lines (IC ₅₀ in μM)			
Compound	MCF 7	A549	HCT116
6a	> 50	> 50	> 50
6b	> 50	> 50	> 50
6c	> 50	> 50	> 50
6d	> 50	> 50	> 50
6e	12.46 ± 2.72	> 50	> 50
8a	> 50	> 50	> 50
8b	> 50	> 50	> 50
8c	> 50	> 50	7.58 ± 1.01
8d	> 50	> 50	> 50
8e	> 50	> 50	21.43 ± 1.23
10a	> 50	> 50	> 50
10b	> 50	> 50	> 50
10c	> 50	> 50	> 50
10d	> 50	> 50	> 50
10e	> 50	> 50	> 50
12a	21.51 ± 4.83	2.25 ± 0.67	> 50
12b	> 50	> 50	> 50
12c	> 50	31.12 ± 0.45	> 50
12d	> 50	26.58 ± 2.05	> 50
12e	> 50	> 50	> 50
14a	> 50	> 50	> 50
14b	32.14 ± 0.83	0.23 ± 0.12	> 50
14c	> 50	15.05 ± 0.11	> 50
14d	> 50	4.99 ± 3.33	> 50
14e	> 50	> 50	> 50
Cisplatin	30.56 ± 2.54	28.54 ± 1.64	22.61 ± 2.03

formation of cell colonies, indicating potential anti-proliferative effects. Specifically, compound **8c** displayed remarkable inhibition against HCT116 cells, while compounds **6e** and **14b** demonstrated notable reductions in colony formation for the MCF7 and A549 cell lines. These results highlight the potential of these compounds as promising candidates for further investigation as new anticancer agents.

2.4 Effect of the fused pyran derivatives on cell cycle

Regulation of cell cycle is a crucial process for maintaining normal cell growth and division. Dysregulation of the cell cycle is one of the hallmarks of cancer, which has emerged as a promising targeting strategy for cancer therapy. We thus evaluated the fused pyran derivatives for their effects on the cell cycle progression in MCF7, A549, and HCT116 cells. As shown in Fig. 4, nearly all tested compounds possess remarkable abilities to arrest the cell cycle progression of the three cancer cells at various phases, including G0/G1, S, and G2/M (Fig. 4). For example, in the A549 cell line (Fig. 4(A)), compound **14b** exhibited a strong inhibition of 57.35 ± 3.18% in the G0/G1 phase after 24 h of treatment at its IC₅₀ concentration, suggesting its potential to prevent the cancer cells from entering the DNA synthesis phase and thereby impeding cancer cell proliferation. Similarly, in the MCF7 cell line (Fig. 4(B)), compound **6e** showed a robust block in the G0/G1 phase (59.10 ± 0.24%). The G0/G1 phase is critical for regulating cell growth and

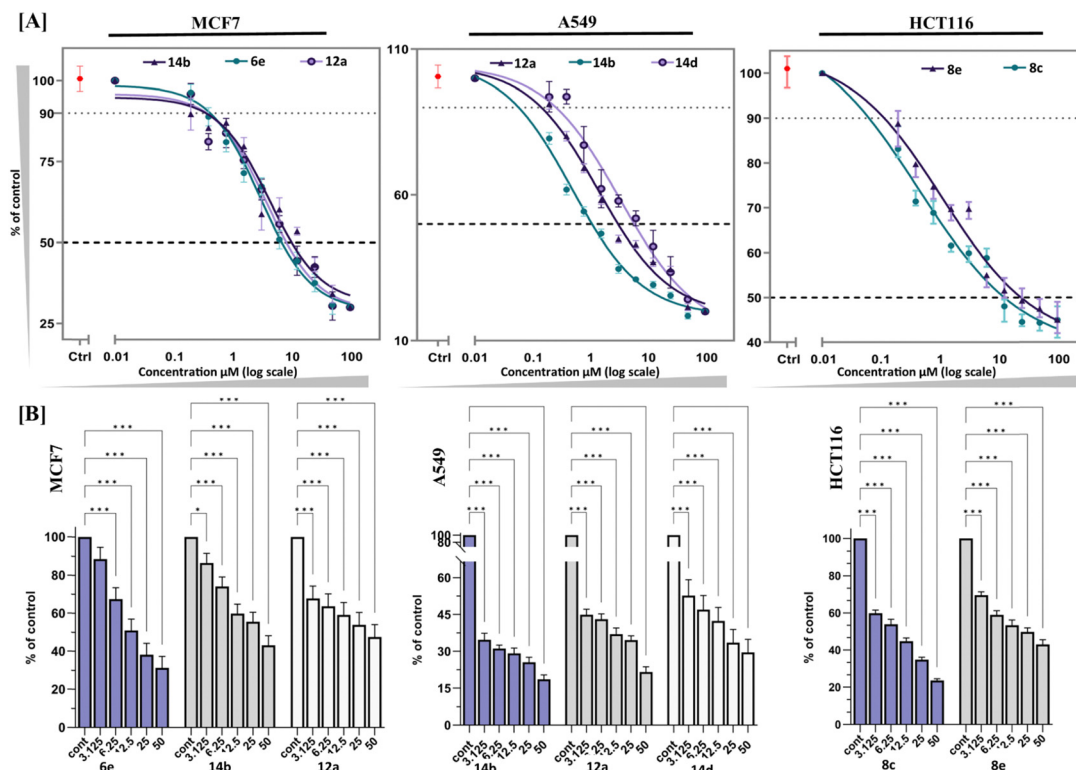


Fig. 2 (A) IC₅₀ comparison graph of top selected compounds. (B) Dose-dependent inhibition of the viability of cancer cells under compound treatment at various concentrations (3.125–50 μM). The statistical analysis was conducted by one-way ANOVA and *post hoc* Dunnett's test. The data represent mean ± SD (*n* = 3), * *P* ≤ 0.01, ** *P* ≤ 0.001, *** *P* ≤ 0.0001, and **** *P* ≤ 0.00001.



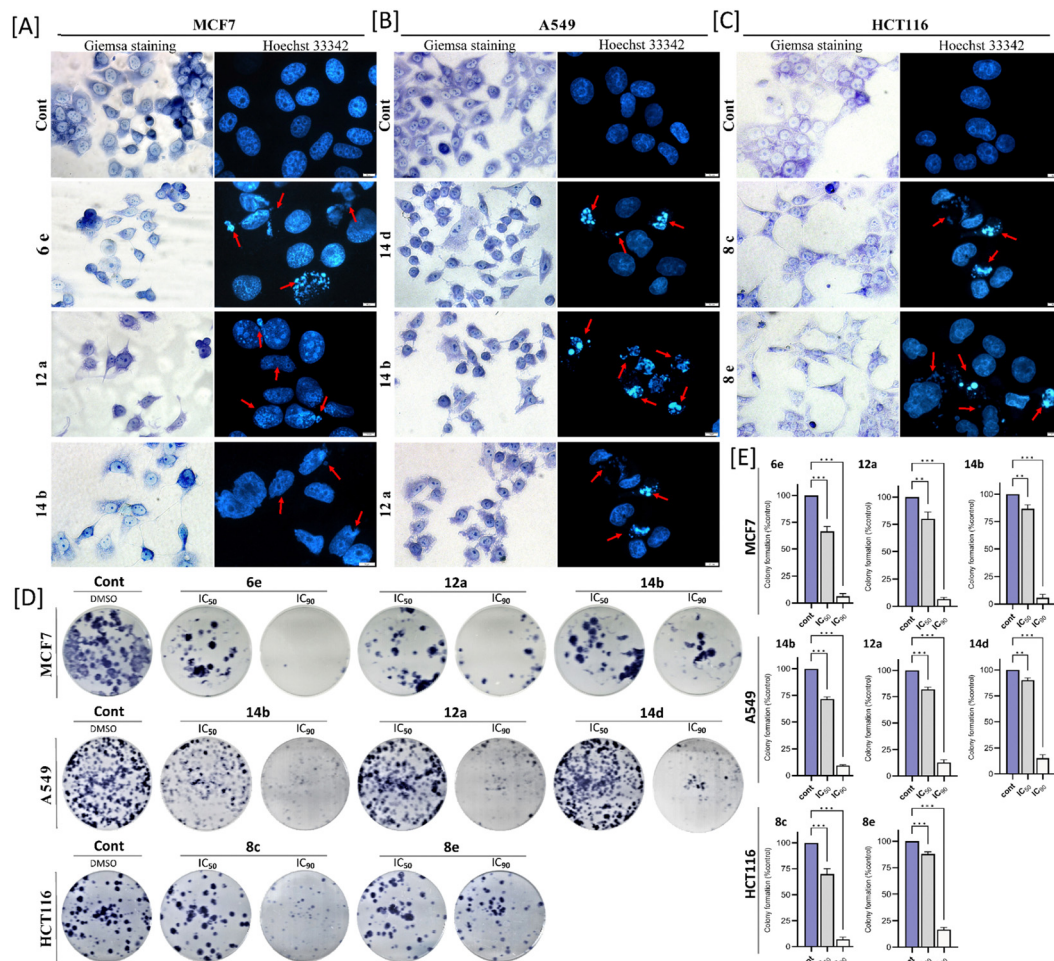


Fig. 3 (A)–(C) Morphological and nuclear changes of MCF7, A549, and HCT116 cells after exposure to the selected fused pyran derivatives concerning the non-treated control cells. (D) Representative image of colony formation assay after compound treatment with IC₅₀ and IC₉₀ concentrations. (E) Quantitative analysis of colony formation ability after compound treatment with IC₅₀ and IC₉₀ concentrations. The data represent mean \pm SD ($n = 3$). The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

deciding whether cells will undergo division or remain quiescent. Therefore, the significant inhibition observed with compound **6e** suggests its potential to halt the cell division of MCF7 cancer cells. Furthermore, compounds **14b** and **12a** exhibited inhibition to the S phase of MCF7 cells, with inhibition rates of $33.60 \pm 2.97\%$ and $35.15 \pm 7.57\%$, respectively. This indicated their activity of interfering with DNA replication and halting subsequent cell division. In the HCT116 cell line (Fig. 4(C)), compound **8c** showed substantial inhibition to the G₂/M phase ($60.40 \pm 0.99\%$) after 48 h incubation. The G₂/M phase is responsible for ensuring accurate chromosome segregation and cell division. The notable block observed with compound **8c** suggests its potential to disrupt these essential processes, thereby inhibiting the growth of HCT116 cancer cells. Additionally, compound **8e** exhibited a significant inhibition in the G₀/G₁ phase ($48.15 \pm 2.89\%$) compared to the untreated cells (Fig. 4(C)) after 48 h incubation. These results highlight the potential of fused pyran derivatives as potential agents for disrupting the cell cycle events in cancer cells. However, to

provide a more robust assessment, it is essential to further investigate and compare the cell cycle effects of the fused pyran derivatives with those of the drugs employed in clinical settings.

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

DNA damage analysis conducted by using γ H2AX and 7AAD staining has provided significant insights into the capacity of the tested compounds to induce DNA double-strand breaks (DSBs) in A549, MCF7, and HCT116 cells (Fig. 5). In the MCF7 cell line, compound **6e** induced a significantly high level of DNA damage of $72.63 \pm 2.14\%$ after 48 h incubation. The other two compounds **12a** and **14b** also exhibited significant DNA damage of 42.53 ± 2.93 and $36.93 \pm 3.14\%$, respectively (Fig. 5(A)) at the 48 h treatment timepoint. Similarly, in the A549 cell line, compound **14b** was found to induce a significant amount of DSBs, accounting for $35.17 \pm 2.787\%$ of the analyzed cells after 48 h incubation. Compounds **12a** and **14d** were



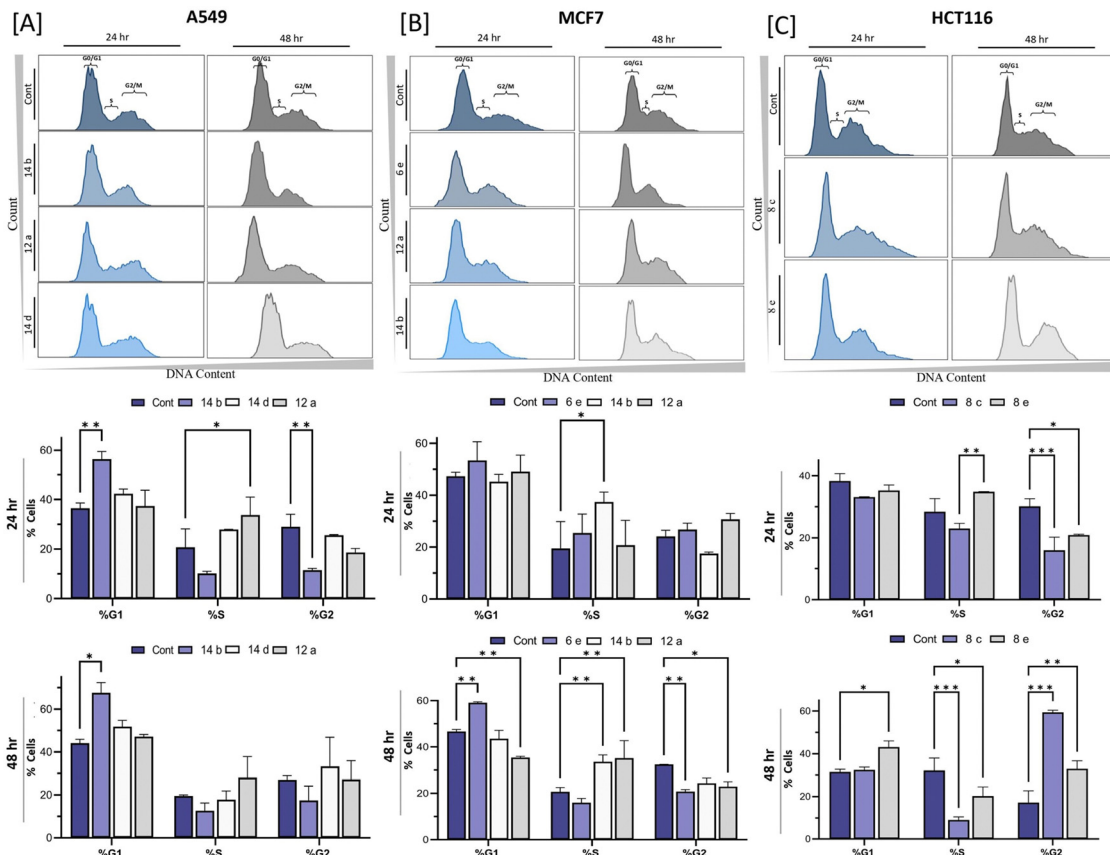


Fig. 4 (A)–(C) The effects of selected fused pyran derivatives on cell cycle distribution in MCF7, A549, and HCT116 cells and its quantitative analysis after compound treatment for 24 and 48 h at IC_{50} concentration. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

slightly weaker than 14b, inducing moderate levels of $23.67 \pm 1.97\%$ and $19.47 \pm 1.87\%$ of DNA damage, respectively (Fig. 5(B)). In the context of the HCT116 cell line, compounds **8c** and **8e** were observed to induce DNA damage as evidenced by γ H2AX detection. Notably, 48 h treatment of compound **8c** resulted in $23.05 \pm 5.162\%$ of DSBs in the cells, while compound **8e** led to $20.75 \pm 3.65\%$ γ H2AX-positive cells (Fig. 5(C)).

The confocal microscopy analysis provided valuable insights into the presence of significant amounts of γ H2AX foci in the cells treated with the tested fused pyran compounds when compared to the untreated cells. Particularly, among the compounds tested, compound **6e** exhibited the highest susceptibility for inducing DNA damage in the MCF7 cell line, indicating its promising efficacy as a potential anticancer agent. Similarly, compound **14b** showed notable induction of DSBs in the A549 cell line, suggesting its potential therapeutic value against breast and lung cancers. Additionally, compound **8c** displayed discernible staining of γ H2AX in the HCT116 cell line, further supporting its capacity to induce DNA damage. Identifying the DNA-damaging effects caused by the fused pyran derivatives holds significant implications as they shed light on the potential mechanisms by which the tested compounds can elicit apoptotic pathways and subsequent cell death in cancer cells. By inducing DSBs, the compounds have

the ability to disrupt DNA integrity, triggering cellular responses that can ultimately lead to the demise of cancerous cells.

2.6 Ability of the fused pyran derivatives to induce apoptosis

Apoptotic assessment, conducted *via* annexin V binding and subsequent flow cytometry analysis, aimed to investigate the potential of tested compounds in inducing apoptotic bodies within the cell cultures. Our results indicated that all tested compounds exhibited a time-dependent capacity to induce apoptosis. Specifically, in the A549 cell line, compound **14b** displayed a remarkable apoptotic activity, with a substantial increase of apoptotic bodies at $34.48 \pm 4.79\%$ and $47.20 \pm 0.28\%$ after 24 and 48 h incubation, respectively. Notably, the standard drug Mitoxantrone exhibited a lower percentage of cell death at $27.60 \pm 3.56\%$ after 24 h incubation (Fig. 6(A)).

Furthermore, when evaluating the apoptotic effect of compound **6e** on the MCF7 cell line, a significant apoptotic induction ability was observed, resulting in $8.55 \pm 3.71\%$ and $57.44 \pm 4.16\%$ of apoptotic cells after 24 and 48 h incubation, respectively. In contrast, Mitoxantrone exhibited a lower apoptotic rate of $15.08 \pm 2.86\%$ after 24 hours of incubation (Fig. 6(B)). In the HCT116 cell line, compound **8c** demonstrated the highest percentage of apoptotic cells, with $11.45 \pm 1.14\%$ and $29.20 \pm 3.29\%$ observed after 24 and 48 hours of incubation,



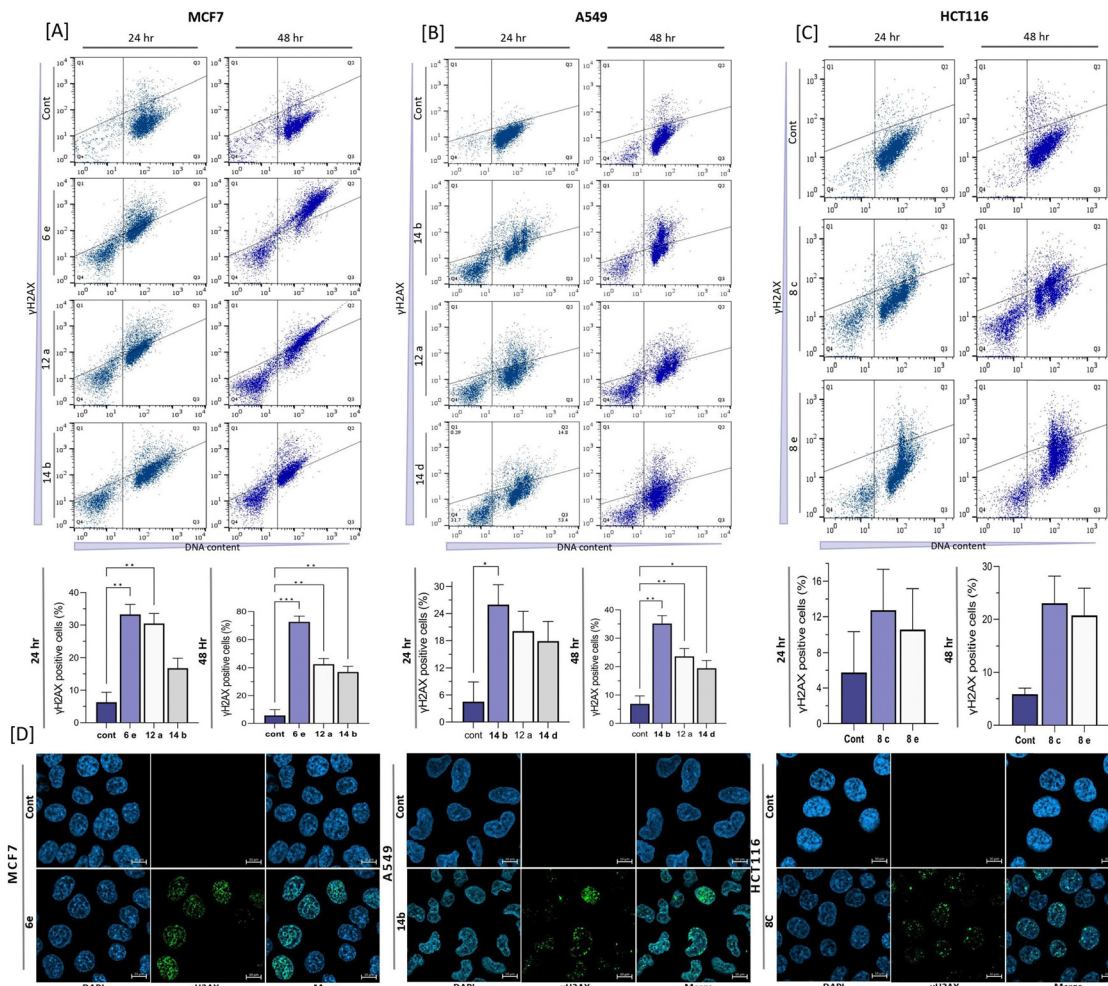


Fig. 5 Formation of DSBs caused by the fused pyran derivatives on MCF7, A549, and HCT116. (A)–(C) Flow cytometric analysis of γ H2AX detection on different cell lines and their quantification after compound treatment for 24 and 48 h at IC_{50} concentrations. (D) Confocal microscopic images of γ H2AX detection in the cells. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

respectively. Comparatively, Mitoxantrone induced a cell death rate of $35.25 \pm 1.90\%$ after 24 h incubation (Fig. 6(C)). These results highlight the apoptotic potential of the tested compounds in different cell lines, with compound **14b** showing significant activity in A549 cells, compound **6e** demonstrating remarkable effects in MCF7 cells, and compound **8c** displaying the most prominent apoptotic induction ability in HCT116 cells. The observed variations in apoptotic responses among the compounds and cell lines merit further investigation, emphasizing the need for comprehensive analysis to elucidate the underlying mechanisms of their action.

2.7 ADMET predictions

Drug-likeness is a qualitative parameter that distinguishes drug candidates from other chemical entities. Approximately 40% of drug failures were allegedly due to their poor pharmacokinetics (ADME) and toxicity properties.⁵² To become an efficient drug, any promising compound should possess higher activity at minimum concentrations with low toxicity and be stable and

active until the desired biological response occurs. Recent developments in combinatorial chemistry and high-throughput screenings (HTS) have significantly increased the capacity of traditional drug discovery programs by early predictions of physicochemical, pharmacokinetic, metabolism, and potential toxicity of a clinical candidate, thereby reducing the late-stage attritions.⁵³

An *in silico* evaluation encompassing the realms of absorption, distribution, metabolism, excretion, and toxicity (ADMET) was conducted to predict the drug-likeness of potent compounds. We delved into the ADMET attributes of the aforementioned compounds, employing the ADMET Protocol housed within the Discovery Studio software (Accelrys, situated in San Diego, California, USA). These taxations were exclusively predicated upon the molecular structural composition of each constituent. A multitude of parameters were computed, including the two-dimensional Fast Polar Surface Area (2D_FPASA), Atom-based Log P98 ($A\text{Log}P98$), the integrity of the Blood–Brain Barrier (BBB), the presence and influence of Cytochrome P4502D6 (CYP2D6), and the propensity for Hepatotoxicity (HEPATOX).^{54–56}



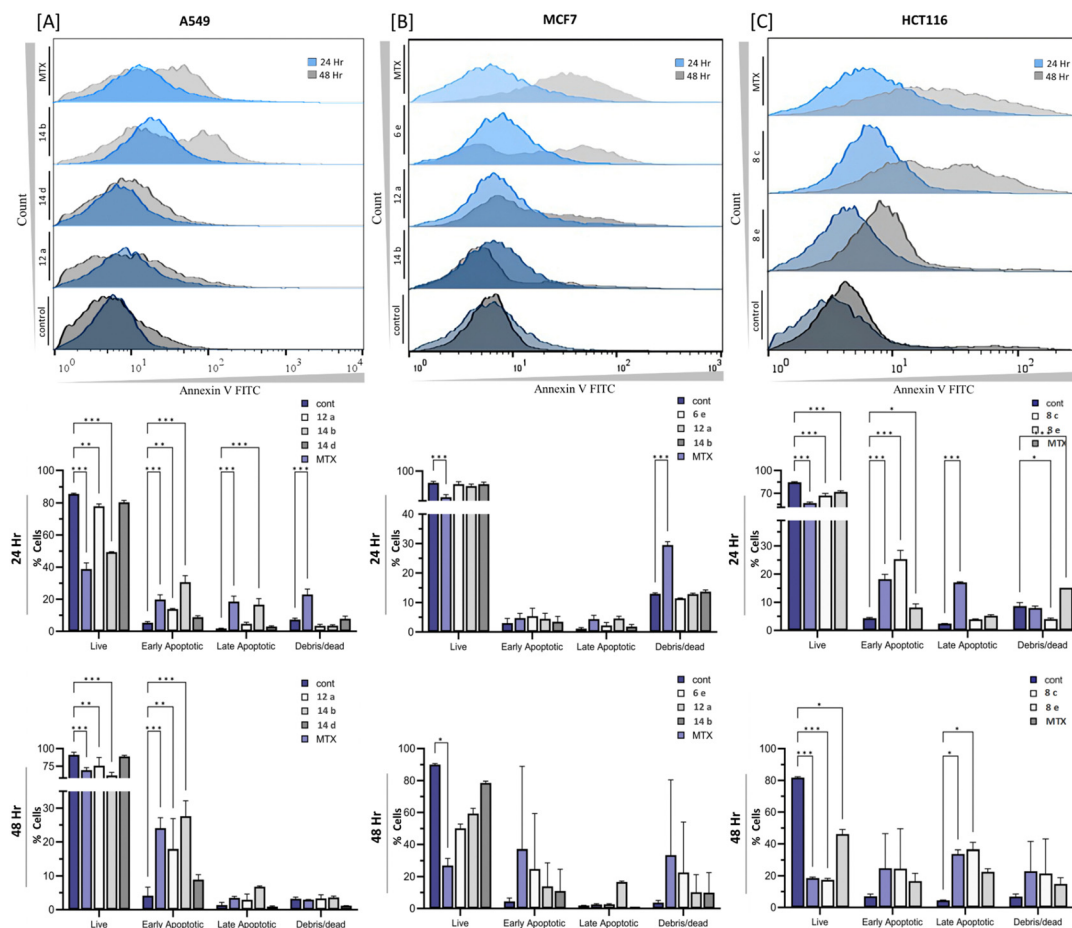


Fig. 6 The apoptotic ability of the selected fused pyran derivatives on different cancer cell lines was detected *via* flow cytometric analysis and its quantification. (A) A549 cell line, (B) MCF7 cell line and (C) HCT116 cell line. The statistical analysis was performed using a two-way ANOVA and Tukey's test. The data represent mean \pm SD ($n = 3$), ns $P > 0.05$, * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

From Tables 2 and 3, it can be concluded that all the potent and obeyed Lipinski's "Rule of Five" and "Veber Rule" and compounds displayed appropriate physicochemical properties hence, they have drug-likeness and can be considered as probable

Table 2 *In silico* physicochemical properties and pharmacokinetic (ADME) predictions of potent compounds

Comp. no.	Lipinski's rule-of-five (RO5)					Veber rule		ADME			
	$A \text{ Log } P$	M_w	HBD	HBA	No. of violations	PSA_2D	RB	Solubility	BBB	CYP2D6	HIA
	≤ 5	≤ 500	≤ 5	≤ 10	≤ 1	≤ 140	≤ 10				
6e	3.98	433.42	1	6	0	112.51	2	1	4	False	1
8c	1.08	347.33	1	6	0	112.51	2	3	4	False	0
8e	2.92	397.39	1	6	0	112.51	2	2	4	False	0
12a	2.08	332.36	1	4	0	92.32	2	2	3	False	0
12c	1.40	333.34	1	5	0	103.58	2	3	3	False	0
12d	3.15	382.42	1	4	0	92.32	2	2	3	False	0
14b	1.98	360.41	1	4	0	92.32	2	2	3	False	0
14c	1.85	361.40	1	5	0	103.58	2	2	3	False	0
14d	3.61	410.47	1	4	0	92.315	2	2	2	False	0

$A \text{ Log } P$ = Ghose-Crippen-Viswanadhan octanol-water partition coefficient, M_w = molecular weight, HBD = hydrogen bond donar, HBA = hydrogen bond acceptor, HT = hepatotoxic, HIA = human intestinal absorption, NI = non-inhibitor, NT = non-toxic. Solubility level: 0 (extremely low), 1 (very low), 2 (low), 3 (good), 4 (optimal), 5 (too soluble), 5 (molecules with one or more $A \text{ Log } P$ 98 types). Blood-brain barrier penetration level (BBB): 0 (very high penetrant), 1 (high), 2 (medium), 3 (low), 4 (undefined), 5 (molecules with one or more $A \text{ Log } P$ calculation). Cytochrome P450 2D6 (CYP2D6) level: 0 (non-inhibitor), 1 (Inhibitor). Hepatotoxic level and prediction: 0 and False (non-toxic), 1 and True (toxic). Human intestinal absorption level (HIA): 0 (good absorption), 1 (moderate), 2 (poor), 3 (very poor). Plasma protein binding (PPB) level and prediction: 0 (binding is $< 90\%$), 1 (binding is $\geq 90\%$), 2 (binding is $\geq 95\%$), and true (binding), false (not-binding).



Table 3 TOPKAT analysis of potent compounds

Comp. no.	NTP rat		Ames mutagen	Fathead minnow LC ₅₀	Rat oral LD ₅₀	Carcinogenic potency _{mouse} TD ₅₀	Daphnia EC ₅₀
	Male	Female					
6e	NC	C	NM	0.000322	0.205353	3.782	0.109699
8c	NC	NC	NM	0.0139834	0.188742	16.9529	2.28863
8e	NC	NC	NM	0.00110458	0.172766	11.575	0.63856
12a	NC	NC	NM	0.0674315	0.0890379	21.0331	1.4037
12c	NC	NC	NM	0.0988727	0.0922284	15.1738	1.26637
12d	NC	NC	NM	0.00806484	0.241895	10.8802	0.40211
14b	NC	NC	NM	0.233863	0.201545	9.78652	4.9743
14c	NC	NC	NM	0.186816	0.162781	9.84066	3.43744
14d	NC	NC	NM	0.0150864	0.422685	6.98587	1.08061

C = carcinogen, NC = non-carcinogen, NM = non-mutagen, LC₅₀ (g L⁻¹), LD₅₀ (g kg⁻¹ body weight), TD₅₀ (mg kg⁻¹ body weight day⁻¹), EC₅₀ (mg L⁻¹)

lead compounds for the development of anticancer drugs. Pharmacokinetic properties like aqueous solubility play a significant role in the drug molecules' bioavailability; from Table 2, the compounds **8c** and **12c** have good aqueous solubility levels (level 3). All others have low-level aqueous solubility levels except for ligand **6e**, having a very low level (level 2), of solubility (level 1). Table 2 shows that the majority of the compounds have low values for BBB penetration levels (level 3) except for compound **14d**, which has a moderate value (level 2), and compounds **6e**, **8c**, and **8e** have undefined BBB penetration levels (level 4). Also, the results indicate that all potent ligands are non-inhibitors concerning CYP2D6 liver, which suggests they are well metabolized in Phase-I and all ligands have good intestinal absorption levels (level 1).

Similarly, the toxicological properties of the 2D molecular structures 9 potent ligands were predicted *in silico* by TOPKAT wizard that utilizes a quantitative structure-toxicity relationship (QSTR) model to assess specific toxicological endpoints such as fathead minnow (LC₅₀), rat oral (LD₅₀), rat (TD₅₀), and Daphnia (EC₅₀) are summarized in Table 3. All potent ligands were depicted to be non-carcinogen and non-mutagen towards NTP rats (male and female) and ames mutagen, except the compound **6e** found to be carcinogenic against female NTP rats (Table 3).

3. Conclusion

Twenty-five new fused pyran derivatives containing nitrogen-based heterocycles such as pyrazole, imidazole, 1,2,4-triazole, benzimidazole, and benzotriazole (**6a-e**, **8a-e**, **10a-e**, **12a-e** & **14a-e**) have been synthesized *via* a three-component reaction involving 4-hydroxycoumarin (**4**)/4-hydroxy-6-methyl-2H-pyran-2-one (**7**)/5-hydroxy-2-(hydroxymethyl)-4H-pyran-4-one (**9**)/cyclohexane-1,3-dione (**11**)/dimedone (**13**), N-based heterocycles substituted benzaldehydes (**3a-e**) and malononitrile (**5**). All the compounds were screened for their *in vitro* anticancer potentials against three cancer cell lines MCF7, A549, and HCT116. Among them, compounds **6e**, **14b**, and **8c** were identified as the most potent against MCF7, A549, and HCT116 with IC₅₀ values of 12.46 ± 2.72 μM, 0.23 ± 0.12 μM, and 7.58 ± 1.01 μM, respectively. The top active compounds

showed strong activities to induce cytoplasmic morphological alterations and inhibit colony formation in the tested cancer cells. They also remarkably inhibited the cell cycle progression of cancer cells at various phases. DNA damage analysis and apoptosis studies revealed that these compounds have the potential to induce DNA double-strand breaks and apoptosis. The *in silico* ADMET studies also suggest that these compounds can be considered lead compounds for further development of anticancer drugs. These data together suggest a good potential of the fused pyran derivatives as promising candidates for further investigation as anticancer agents.

4. Experimental section

Chemistry

All the reagents and solvents required for the synthesis were obtained commercially with the highest quality and were used directly without treatment unless otherwise indicated. The melting point of all the compounds was determined using the Stuart-SMP20 capillary melting point apparatus and is uncorrected. IR spectra of the compounds were analyzed by dispersing the compounds in potassium bromide pellets on a JASCO FT-IR 4700 spectrophotometer. ¹H NMR and ¹³C NMR spectra were obtained on a Jeol spectrometer (500 MHz) in CDCl₃/DMSO-d₆ solvent using TMS as an internal standard. Coupling constant (*J*) values are expressed in Hertz (Hz) and the chemical shift (δ) values are reported in (ppm). LC-HRMS of all the compounds were recorded on Agilent 6530 ESI-QTOF mass instrument (mass accuracy of 5 ppm) operating in positive ion acquisition mode using reverse phase C18 column and all the compounds were confirmed to be greater than 98% pure. Thin-layer chromatography (TLC) was analyzed using a mixture of (4:1 v/v) hexanes and ethyl acetate. All the compounds were purified through column chromatography using silica gel (Merck 100–200 mesh) as a stationary phase and a mixture of hexanes and ethyl acetate as a mobile phase.

4-(1H-Pyrazol-1-yl)benzaldehyde (3a). To a solution of 4-fluorobenzaldehyde (248.22 mg, 2 mmol) in dimethylformamide (5 mL) was added 1H-pyrazole (136.15 mg, 2 mmol) and anhydrous potassium carbonate (829.23 mg, 6 mmol). The reaction mixture was heated at 130 °C for 19 h. The reaction



mixture was cooled to room temperature and poured into water (50 mL). The resulting mixture was extracted with ethyl acetate (20 mL) three times and the combined organic layer was dried over Na_2SO_4 , concentrated under reduced pressure, and purified by column chromatography using 20% ethyl acetate in hexanes to obtain the pure product. Yield: 78%. Pale yellow solid. M.p.: 83–85 °C. IR (KBr, ν_{max} , cm^{-1}): 3126 (C–H), 1692 (C=O), 1604 (C=C), 1520 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.94 (s, 1H), 7.96 (d, $J = 2.5$ Hz, 1H), 7.92–7.90 (m, 2H), 7.83–7.81 (m, 2H), 7.71 (d, $J = 1.5$ Hz, 1H), 6.46 (t, $J = 2.2$ Hz, 1H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.898, 144.248, 142.281, 134.064, 131.290, 126.942, 118.774, 108.810. MS (ESI-QTOF) for $\text{C}_{10}\text{H}_9\text{N}_2\text{O}$ $[\text{M} + \text{H}]^+$ calculated 173.0709, found 173.0712.

Compounds 3b–e were synthesized by following the same procedure

4-(1H-Imidazol-1-yl)benzaldehyde (3b). 1H-Imidazole (136.15 mg, 2 mmol) was used as a reagent, and the reaction time was 19 h. Yield: 80%. Pale yellow solid. M.p.: 145–147 °C. IR (KBr, ν_{max} , cm^{-1}): 3112 (C–H), 1689 (C=O), 1609 (C=C), 1515 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 10.01 (s, 1H), 8.00–7.95 (m, 3H), 7.56–7.54 (m, 2H), 7.33–7.36 (m, 1H), 7.21 (s, 1H). ^{13}C NMR (126 MHz, CDCl_3): δ 191.1, 162.6, 142.3, 134.1, 131.3, 127.0, 118.8, 108.9. MS (ESI-QTOF) for $\text{C}_{10}\text{H}_9\text{N}_2\text{O}$ $[\text{M} + \text{H}]^+$ calculated 173.0709, found 173.0712.

4-(1H-1,2,4-Triazol-1-yl)benzaldehyde (3c). 1H-1,2,4-Triazole (138.13 mg, 2 mmol) was used as a reagent, and the reaction time was 22 h. Yield: 74%. White solid. M.p.: 145–147 °C. IR (KBr, ν_{max} , cm^{-1}): 3099 (C–H), 1693 (C=O), 1604 (C=C), 1520 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.99 (s, 1H), 8.63 (s, 1H), 8.09 (s, 1H), 7.98 (d, $J = 8.4$ Hz, 2H), 7.84 (d, $J = 8.6$ Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.562, 153.156, 141.177, 141.033, 135.523, 131.405, 119.839. MS (ESI-QTOF) for $\text{C}_9\text{H}_8\text{N}_3\text{O}$ $[\text{M} + \text{H}]^+$ calculated 174.0662, found 174.0658.

4-(1H-Benzo[d]imidazol-1-yl)benzaldehyde (3d). 1H-Benzo[d]imidazole (236.27 mg, 2 mmol) was used as a reagent, and the reaction time was 22 h. Yield: 79%. White solid. M.p.: 108–111 °C. IR (KBr, ν_{max} , cm^{-1}): 3143 (C–H), 1698 (C=O), 1604 (C=C), 1511 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 10.03 (s, 1H), 8.15 (s, 1H), 8.04 (d, $J = 8.6$ Hz, 2H), 7.84–7.81 (m, 1H), 7.68–7.65 (m, 2H), 7.56–7.53 (m, 1H), 7.33–7.27 (m, 2H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.620, 144.085, 141.724, 141.225, 135.293, 132.912, 131.607, 124.322, 123.717, 123.486, 120.885, 110.442. MS (ESI-QTOF) for $\text{C}_{14}\text{H}_{11}\text{N}_2\text{O}$ $[\text{M} + \text{H}]^+$ calculated 223.0866, found 223.0871.

4-(1H-Benzo[d][1,2,3]triazol-1-yl)benzaldehyde (3e). 1H-Benzo[d][1,2,3]triazole (238.25 mg, 2 mmol) was used as a reagent, and the reaction time was 24 h. Yield: 72%. White solid. M.p.: 186–190 °C. IR (KBr, ν_{max} , cm^{-1}): 3148 (C–H), 1693 (C=O), 1604 (C=C), 1515 (C=N). ^1H NMR (500 MHz, CDCl_3): δ 9.94 (s, 1H), 7.96 (d, $J = 2.5$ Hz, 1H), 7.93–7.89 (m, 2H), 7.84–7.81 (m, 2H), 7.71 (d, $J = 1.5$ Hz, 1H), 7.19 (s, 1H), 6.46–6.45 (m, 1H). ^{13}C NMR (126 MHz, CDCl_3): δ 190.898, 144.248, 142.281, 134.064, 131.290, 126.942, 118.774, 108.810. MS (ESI-QTOF) for $\text{C}_{13}\text{H}_{10}\text{N}_3\text{O}$ $[\text{M} + \text{H}]^+$ calculated 224.0818, found 224.0818.

4-(4-(1H-Pyrazol-1-yl)phenyl)-2-amino-5-oxo-4,5-dihydro-pyrano[3,2-c]chromene-3-carbonitrile (6a). To a solution of 4-(1H-pyrazol-1-yl)benzaldehyde (3a, 172.18 mg, 1 mmol) in water (2 mL) was added 4-hydroxycoumarin (4, 162.14 mg, 1 mmol), malononitrile (5, 66.06 mg, 1 mmol) and sodium fluoride (5 mg, 12 mol%). The reaction mixture was refluxed for 3 h. 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 obtain the pure product. Yield: 90%. Pale yellow solid. M.p.: 248–253 °C. IR (ν_{max} , cm^{-1}): 3367, 3255 (–NH₂), 3173 (C–H), 2211 (CN), 1710 (C=O), 1652 (C=C), 1200 (C–O–C). ^1H NMR (500 MHz, DMSO-d_6): δ 8.45 (d, $J = 2.5$ Hz, 1H), 7.93–7.91 (m, 1H), 7.77–7.71 (m, 4H), 7.53–7.38 (m, 6H), 6.53–6.52 (m, 1H), 4.53 (s, 1H). ^{13}C NMR (126 MHz, DMSO-d_6): δ 159.589, 157.948, 153.475, 152.179, 141.285, 140.939, 138.770, 132.991, 128.864, 127.750, 124.708, 122.529, 119.217, 118.622, 116.606, 113.307, 107.785, 103.696, 57.690, 36.467. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{14}\text{N}_4\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 383.1139, found 383.1141.

4-(4-(1H-Imidazol-1-yl)phenyl)-2-amino-5-oxo-4,5-dihydro-pyrano[3,2-c]chromene-3-carbonitrile (6b). Yield: 80%. White solid. M.p.: 210–214 °C. IR (ν_{max} , cm^{-1}): 3460, 3178 (–NH₂), 3125 (C–H), 2196 (CN), 1710 (C=O), 1671 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO-d_6): δ 8.32 (s, 1H), 7.92 (d, $J = 7.1$ Hz, 1H), 7.7–7.72 (m, 2H), 7.59 (d, $J = 8.4$ Hz, 2H), 7.53–7.44 (m, 6H), 7.16 (s, 1H), 4.56 (s, 1H). ^{13}C NMR (126 MHz, DMSO-d_6): δ 159.608, 157.996, 153.580, 152.198, 142.379, 135.688, 135.266, 133.010, 129.248, 129.133, 124.660, 122.557, 120.945, 119.179, 118.526, 116.549, 112.997, 103.571, 57.584, 36.438. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{15}\text{N}_4\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 383.1139, found 383.1137.

4-(4-(1H-1,2,4-Triazol-1-yl)phenyl)-2-amino-5-oxo-4,5-dihydro-pyrano[3,2-c]chromene-3-carbonitrile (6c). Yield: 89%. Off-white solid. M.p.: 282–285 °C. IR (ν_{max} , cm^{-1}): 3348, 3241 (–NH₂), 3100 (C–H), 2186 (CN), 1715 (C=O), 1671 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO-d_6): δ 9.24 (s, 1H), 8.20 (s, 1H), 7.91–7.90 (m, 1H), 7.78 (d, $J = 8.6$ Hz, 2H), 7.73–7.69 (m, 1H), 7.51–7.44 (m, 6H), 4.55 (s, 1H). ^{13}C NMR (126 MHz, DMSO-d_6): δ 159.541, 157.929, 153.571, 152.342, 152.189, 142.983, 142.312, 135.679, 132.982, 129.094, 124.660, 122.519, 119.697, 119.083, 116.577, 112.968, 103.466, 57.546, 36.544. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{13}\text{N}_5\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 384.1091, found 384.1091.

4-(4-(1H-Benzo[d]imidazol-1-yl)phenyl)-2-amino-5-oxo-4,5-dihydro-pyrano[3,2-c]chromene-3-carbonitrile (6d). Yield: 86%. Off-white solid. M.p.: 245–248 °C. IR (ν_{max} , cm^{-1}): 3430, 3294 (–NH₂), 3153 (C–H), 2201 (CN), 1691 (C=O), 1603 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO-d_6): δ 8.55 (s, 1H), 7.92 (d, $J = 6.7$ Hz, 1H), 7.76–7.70 (m, 2H), 7.64–7.46 (m, 9H), 7.31–7.27 (m, 2H), 4.60 (s, 1H). ^{13}C NMR (126 MHz, DMSO-d_6): δ 159.599, 158.101, 153.609, 152.179, 143.617, 143.262, 142.830, 134.834, 133.010, 129.286, 124.698, 123.690, 123.441, 122.500, 122.433, 119.831, 119.179, 116.587, 112.968, 110.703, 103.677, 57.574, 36.553. MS (ESI-QTOF) for $\text{C}_{26}\text{H}_{16}\text{N}_4\text{O}_3$ $[\text{M} + \text{H}]^+$ calculated 433.1295, found 433.1292.



4-(4-(1*H*-Benzo[*d*][1,2,3]triazol-1-yl)phenyl)-2-amino-5-oxo-4,5-dihydropyrano[3,2-*c*]chromene-3-carbonitrile (6e). Yield: 88%. Yellow solid. M.p.: 220–225 °C. IR (ν_{\max} , cm^{-1}): 3387, 3251 (–NH₂), 3073 (=C–H), 2196 (CN), 1715 (C=O), 1667 (C=C), 1214 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.60 (s, 1H), 8.50 (d, *J* = 7.6 Hz, 1H), 8.23–8.16 (m, 3H), 8.00–7.88 (m, 3H), 7.69–7.47 (m, 6H), 4.58 (s, 1H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 160.424, 160.136, 158.552, 154.204, 152.764, 145.421, 144.970, 139.067, 133.595, 132.876, 131.772, 129.823, 129.084, 128.201, 125.254, 123.075, 121.424, 121.012, 118.948, 118.948, 117.172, 113.505, 103.945, 57.948, 37.205. MS (ESI-QTOF) for C₂₅H₁₅N₅O₃ [M + H]⁺ calculated 434.1248, found 434.1248.

4-(4-(1*H*-Pyrazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4,5-dihydropyrano[4,3-*b*]pyran-3-carbonitrile (8a). Yield: 85%. Mustard yellow solid. M.p.: 232–238 °C. IR (ν_{\max} , cm^{-1}): 3465, 3300 (–NH₂), 3129 (=C–H), 2186 (CN), 1725 (C=O), 1671 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.41 (s, 1H), 7.76–7.72 (m, 3H), 7.31 (d, *J* = 8.2 Hz, 2H), 7.16 (s, 2H), 6.52 (s, 1H), 6.27 (s, 1H), 4.36 (s, 1H), 2.23 (s, 3H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 163.006, 161.365, 158.197, 158.053, 141.544, 140.843, 138.635, 128.806, 128.566, 127.875, 127.578, 119.284, 118.603, 107.718, 100.442, 98.004, 57.622, 35.766, 19.295. MS (ESI-QTOF) for C₂₆H₁₆N₄O₃ [M + H]⁺ calculated 347.1139, found 347.1143.

4-(4-(1*H*-Imidazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4,5-dihydropyrano[4,3-*b*]pyran-3-carbonitrile (8b). Yield: 78%. Light brown solid. M.p.: 213–217 °C. IR (ν_{\max} , cm^{-1}): 3367, 3125 (–NH₂), 3086 (=C–H), 2196 (CN), 1695 (C=O), 1622 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.25 (s, 1H), 7.72 (t, *J* = 1.1 Hz, 1H), 7.58 (d, *J* = 8.4 Hz, 2H), 7.34–7.32 (m, 2H), 7.26 (s, 2H), 7.12 (s, 1H), 6.30 (s, 1H), 4.37 (s, 1H), 2.23 (s, 3H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 163.083, 161.355, 158.255, 158.101, 142.427, 135.717, 128.969, 128.864, 120.763, 119.265, 118.161, 100.356, 98.023, 96.305, 57.498, 35.747, 19.228. MS (ESI-QTOF) for C₁₉H₁₄N₄O₃ [M + H]⁺ calculated 347.1139, found 347.1139.

4-(4-(1*H*-1,2,4-Triazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4,5-dihydropyrano[4,3-*b*]pyran-3-carbonitrile (8c). Yield: 75%. Off-white solid. M.p.: 260–265 °C. IR (ν_{\max} , cm^{-1}): 3363, 3144 (–NH₂), 3105 (=C–H), 2192 (CN), 1706 (C=O), 1671 (C=C), 1209 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 9.25 (s, 1H), 8.22 (s, 1H), 7.79 (d, *J* = 8.6 Hz, 2H), 7.39 (d, *J* = 8.6 Hz, 2H), 7.26 (s, 2H), 6.30 (s, 1H), 4.40 (s, 1H), 2.24 (s, 3H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 163.102, 161.346, 158.284, 158.063, 152.294, 143.271, 142.331, 135.612, 128.960, 119.697, 119.217, 100.231, 98.004, 57.440, 35.833, 19.304. MS (ESI-QTOF) for C₁₈H₁₃N₅O₃ [M + H]⁺ calculated 348.1091, found 348.1095.

4-(4-(1*H*-Benzo[*d*]imidazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4,5-dihydropyrano[4,3-*b*]pyran-3-carbonitrile (8d). Yield: 83%. Dark yellow solid. M.p.: 255–258 °C. IR (ν_{\max} , cm^{-1}): 3353, 3309 (–NH₂), 3110 (=C–H), 2206 (CN), 1715 (C=O), 1667 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.59 (s, 1H), 7.79–7.77 (m, 1H), 7.67–7.63 (m, 3H), 7.46 (d, *J* = 8.4 Hz, 2H), 7.33–7.31 (m, 4H), 6.33 (s, 1H), 4.44 (s, 1H), 2.25 (s, 3H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 163.141, 161.403, 158.341, 158.226, 143.751, 143.444, 143.099, 134.777, 132.962, 129.219,

123.681, 122.509, 119.860, 119.323, 110.770, 100.442, 97.975, 57.478, 35.929, 19.295. MS (ESI-QTOF) for C₂₃H₁₆N₄O₃ [M + H]⁺ calculated 397.1295, found 397.1299.

4-(4-(1*H*-Benzo[*d*][1,2,3]triazol-1-yl)phenyl)-2-amino-7-methyl-5-oxo-4,5-dihydropyrano[4,3-*b*]pyran-3-carbonitrile (8e). Yield: 85%. Dark brown solid. M.p.: 223–226 °C. IR (ν_{\max} , cm^{-1}): 3455, 3319 (–NH₂), 3129 (=C–H), 2192 (CN), 1710 (C=O), 1671 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.45 (d, *J* = 2.5 Hz, 1H), 7.77–7.73 (m, 4H), 7.31 (d, *J* = 8.6 Hz, 2H), 7.25 (s, 2H), 6.53 (t, *J* = 2.1 Hz, 1H), 6.30 (s, 1H), 4.35 (s, 1H), 2.23 (s, 3H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 163.016, 161.365, 158.197, 158.053, 141.544, 140.843, 138.635, 128.797, 128.566, 127.885, 127.587, 119.284, 118.612, 107.718, 100.442, 97.946, 57.613, 35.766, 19.285. MS (ESI-QTOF) for C₂₂H₁₅N₅O₃ [M + H]⁺ calculated 398.1248, found 398.1241.

4-(4-(1*H*-Pyrazol-1-yl)phenyl)-2-amino-6-(hydroxymethyl)-8-oxo-4,8-dihydropyrano[3,2-*b*]pyran-3-carbonitrile (10a). Yield: 85%. Dark brown solid. M.p.: 223–227 °C. IR (ν_{\max} , cm^{-1}): 3411 (OH), 3304, 3197 (–NH₂), 3129 (=C–H), 2181 (CN), 1652 (C=O), 1613 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.50 (d, *J* = 1.9 Hz, 1H), 7.86 (d, *J* = 8.4 Hz, 2H), 7.75 (s, 1H), 7.42 (d, *J* = 8.4 Hz, 2H), 7.28 (s, 2H), 6.55 (s, 1H), 6.35 (s, 1H), 5.69 (t, *J* = 5.5 Hz, 1H), 4.89 (s, 1H), 4.24–4.12 (m, 2H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 169.601, 168.257, 159.224, 148.743, 141.054, 139.250, 138.635, 136.399, 129.037, 128.950, 127.952, 127.674, 119.246, 118.929, 111.433, 107.967, 107.900, 59.091, 55.491. MS (ESI-QTOF) for C₁₉H₁₄N₄O₄ [M + H]⁺ calculated 363.1088, found 363.1087.

4-(4-(1*H*-Imidazol-1-yl)phenyl)-2-amino-6-(hydroxymethyl)-8-oxo-4,8-dihydropyrano[3,2-*b*]pyran-3-carbonitrile (10b). Yield: 75%. Grey solid. M.p.: 248–252 °C. IR (ν_{\max} , cm^{-1}): 3411 (OH), 3304, 3197 (–NH₂), 3129 (=C–H), 2181 (CN), 1652 (C=O), 1613 (C=C), 1200 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 8.26 (s, 1H), 7.75 (s, 1H), 7.67 (d, *J* = 8.6 Hz, 2H), 7.44 (d, *J* = 8.4 Hz, 2H), 7.29 (s, 2H), 7.11 (s, 1H), 6.35 (s, 1H), 5.70 (t, *J* = 6.1 Hz, 1H), 4.90 (s, 1H), 4.24–4.12 (m, 2H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 169.581, 168.247, 159.263, 148.628, 139.451, 136.456, 135.324, 129.651, 129.238, 120.964, 119.227, 118.094, 118.017, 118.0, 111.394, 59.101, 55.415. MS (ESI-QTOF) for C₁₉H₁₄N₄O₄ [M + H]⁺ calculated 363.1088, found 363.1090.

4-(4-(1*H*-1,2,4-triazol-1-yl)phenyl)-2-amino-6-(hydroxymethyl)-8-oxo-4,8-dihydropyrano[3,2-*b*]pyran-3-carbonitrile (10c). Yield: 84%. Yellowish-white solid. M.p.: 290–294 °C. IR (ν_{\max} , cm^{-1}): 3363 (OH), 3319, 3164 (–NH₂), 3095 (=C–H), 2206 (CN), 1667 (C=O), 1608 (C=C), 1209 (C–O–C). ¹H NMR (500 MHz, DMSO-*d*₆): δ 9.30 (s, 1H), 8.25 (s, 1H), 7.89 (d, *J* = 8.6 Hz, 2H), 7.50 (d, *J* = 8.6 Hz, 2H), 7.30 (s, 2H), 6.35 (s, 1H), 5.69 (t, *J* = 6.2 Hz, 1H), 4.94 (s, 1H), 4.24–4.12 (m, 2H). ¹³C NMR (126 MHz, DMSO-*d*₆): δ 169.610, 168.276, 159.253, 152.640, 148.512, 142.436, 140.373, 136.485, 136.284, 129.248, 120.052, 119.208, 111.394, 59.101, 55.357. MS (ESI-QTOF) for C₁₈H₁₃N₅O₄ [M + H]⁺ calculated 364.1040, found 364.1039.

4-(4-(1*H*-Benzo[*d*]imidazol-1-yl)phenyl)-2-amino-6-(hydroxymethyl)-8-oxo-4,8-dihydropyrano[3,2-*b*]pyran-3-carbonitrile (10d). Yield: 80%. Dark brown solid. M.p.: 210–213 °C. IR (ν_{\max} , cm^{-1}): 3416 (OH), 3333, 3178 (–NH₂), 3119 (=C–H), 2181 (CN), 1657



(C=O), 1583 (C=C), 1224 (C-O-C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.60 (s, 1H), 7.79–7.73 (m, 3H), 7.68–7.66 (m, 1H), 7.56 (d, J = 8.4 Hz, 2H), 7.34–7.30 (m, 4H), 6.37 (s, 1H), 5.73 (t, J = 6.0 Hz, 1H), 4.98 (s, 1H), 4.28–4.17 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 170.138, 168.861, 159.953, 149.222, 144.356, 143.837, 140.775, 137.070, 136.081, 133.423, 129.958, 124.601, 124.054, 123.066, 120.503, 119.821, 112.018, 111.317, 59.676, 55.923. MS (ESI-QTOF) for $\text{C}_{23}\text{H}_{16}\text{N}_4\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 413.1244, found 413.1252.

4-(4-(1H-Benzo[d][1,2,3]triazol-1-yl)phenyl)-2-amino-6-(hydroxymethyl)-8-oxo-4,8-dihydropyran[3,2-*b*]pyran-3-carbonitrile (10e). Yield: 85%. Dark brown solid. M.p.: 268–270 °C. IR (ν_{max} , cm^{-1}): 3382 (OH) 3294, 3178 (–NH₂), 3071 (=C–H), 2196 (CN), 1628 (C=O), 1583 (C=C), 1214 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.35 (d, J = 8.6 Hz, 2H), 8.05–8.04 (m, 2H), 7.61 (d, J = 8.8 Hz, 2H), 7.53 (q, J = 3.2 Hz, 2H), 7.34 (s, 2H), 6.36 (s, 1H), 5.69 (t, J = 6.1 Hz, 1H), 5.01 (s, 1H), 4.25–4.13 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 170.138, 168.861, 159.886, 148.963, 145.027, 142.599, 139.604, 137.070, 132.885, 131.781, 129.986, 128.335, 121.405, 119.725, 118.957, 118.794, 111.998, 59.628, 55.769. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{15}\text{N}_5\text{O}_4$ $[\text{M} + \text{H}]^+$ calculated 414.1197, found 414.1195.

4-(4-(1H-Pyrazol-1-yl)phenyl)-2-amino-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (12a). Yield: 89%. Off-white solid. M.p.: 207–210 °C. IR (ν_{max} , cm^{-1}): 3363, 3304 (–NH₂), 3110 (=C–H), 2196 (CN), 1671 (C=O), 1594 (C=C), 1200 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.39 (s, 1H), 7.69 (d, J = 7.6 Hz, 3H), 7.24 (d, J = 8.4 Hz, 2H), 7.02 (s, 2H), 6.48 (s, 1H), 4.21 (s, 1H), 2.63–2.57 (m, 2H), 2.32–2.20 (m, 2H), 1.96–1.86 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 196.448, 165.079, 158.994, 143.348, 141.361, 138.865, 128.796, 128.230, 120.263, 119.121, 114.072, 108.207, 58.418, 36.860, 35.554, 27.031, 20.340. MS (ESI-QTOF) for $\text{C}_{19}\text{H}_{16}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 333.1346, found 333.1351.

4-(4-(1H-Imidazol-1-yl)phenyl)-2-amino-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (12b). Yield: 82%. Yellow solid. M.p.: 209–211 °C. IR (ν_{max} , cm^{-1}): 3391, 3324 (–NH₂), 3105 (=C–H), 2192 (CN), 1681 (C=O), 1642 (C=C), 1205 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.39 (s, 1H), 7.69 (d, J = 8.4 Hz, 3H), 7.23 (d, J = 8.4 Hz, 2H), 7.01 (s, 2H), 6.48 (s, 1H), 4.21 (s, 1H), 2.60–2.53 (m, 2H), 2.30–2.19 (m, 2H), 1.95–1.85 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 195.920, 164.705, 158.409, 143.934, 135.218, 128.902, 128.537, 120.887, 119.678, 119.361, 118.516, 113.391, 57.843, 36.304, 35.046, 27.914, 26.494. MS (ESI-QTOF) for $\text{C}_{19}\text{H}_{16}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 333.1346, found 333.1351.

4-(4-(1H-1,2,4-Triazol-1-yl)phenyl)-2-amino-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (12c). Yield: 80%. Off-white solid. M.p.: 231–233 °C. IR (ν_{max} , cm^{-1}): 3402, 3275 (–NH₂), 3134 (=C–H), 2186 (CN), 1676 (C=O), 1608 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 9.23 (s, 1H), 8.22 (s, 1H), 7.76 (d, J = 8.4 Hz, 2H), 7.35 (d, J = 8.4 Hz, 2H), 7.08 (s, 2H), 4.28 (s, 1H), 2.68–2.58 (m, 2H), 2.36–2.23 (m, 2H), 2.00–1.86 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 195.920, 164.686, 158.447, 152.448, 152.217, 144.548, 142.302, 135.295, 128.537, 119.649, 113.343, 57.699, 36.304, 35.152, 26.503,

19.775. MS (ESI-QTOF) for $\text{C}_{18}\text{H}_{15}\text{N}_5\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 334.1299, found 334.1302.

4-(4-(1H-Benzo[d]imidazol-1-yl)phenyl)-2-amino-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (12d). Yield: 84%. Yellowish-white solid. M.p.: 259–261 °C. IR (ν_{max} , cm^{-1}): 3391, 3300 (–NH₂), 3086 (=C–H), 2181 (CN), 1691 (C=O), 1652 (C=C), 1200 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.57 (s, 1H), 7.79–7.76 (m, 1H), 7.64–7.61 (m, 3H), 7.42 (d, J = 8.4 Hz, 2H), 7.33–7.29 (m, 2H), 7.11 (s, 2H), 4.33 (s, 1H), 2.70–2.61 (m, 2H), 2.37–2.28 (m, 2H), 2.03–1.89 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 195.968, 164.811, 158.601, 144.375, 143.607, 143.444, 143.195, 134.383, 132.991, 128.739, 123.690, 123.594, 119.889, 119.755, 113.496, 110.732, 57.766, 36.323, 35.113, 26.503, 19.784. MS (ESI-QTOF) for $\text{C}_{23}\text{H}_{18}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 383.1503, found 383.1502.

4-(4-(1H-Benzo[d][1,2,3]triazol-1-yl)phenyl)-2-amino-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (12e). Yield: 90%. Off-white solid. M.p.: 255–257 °C. IR (ν_{max} , cm^{-1}): 3445, 3387 (–NH₂), 3076 (=C–H), 2202 (CN), 1700 (C=O), 1642 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.66 (s, 1H), 8.55 (d, J = 8.4 Hz, 1H), 8.25–8.21 (m, 2H), 8.07–8.03 (m, 2H), 7.58–7.51 (m, 2H), 7.46 (d, J = 8.6 Hz, 1H), 7.13 (s, 1H), 4.34 (s, 1H), 2.70–2.60 (m, 2H), 2.38–2.26 (m, 2H), 2.02–1.92 (m, 2H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 196.496, 165.310, 159.070, 146.755, 144.951, 138.683, 129.276, 128.143, 120.916, 120.186, 118.716, 113.832, 58.073, 36.841, 35.852, 27.069, 20.312. MS (ESI-QTOF) for $\text{C}_{22}\text{H}_{17}\text{N}_5\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 384.1455, found 384.1456.

4-(4-(1H-Pyrazol-1-yl)phenyl)-2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (14a). Yield: 88%. White solid. M.p.: 271–273 °C. IR (ν_{max} , cm^{-1}): 3372, 3309 (–NH₂), 3178 (=C–H), 2181 (CN), 1691 (C=O), 1652 (C=C), 1214 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.44 (s, 1H), 7.75–7.72 (m, 3H), 7.26 (d, J = 8.2 Hz, 2H), 7.06 (s, 2H), 6.52 (s, 1H), 4.24 (s, 1H), 2.53 (s, 2H), 2.27 (d, J = 16.0 Hz, 1H), 2.12 (d, J = 16.0 Hz, 1H), 1.04 (s, 3H), 0.96 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 195.709, 162.526, 158.485, 142.734, 140.891, 140.766, 138.319, 128.413, 128.144, 127.808, 127.530, 119.668, 118.584, 112.498, 107.651, 57.949, 49.963, 35.113, 31.811, 28.366, 26.791. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{20}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 361.1659, found 361.1664.

4-(4-(1H-Imidazol-1-yl)phenyl)-2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (14b). Yield: 80%. Pale yellow solid. M.p.: 245–247 °C. IR (ν_{max} , cm^{-1}): 3440, 3129 (–NH₂), 2950 (=C–H), 2177 (CN), 1681 (C=O), 1598 (C=C), 1209 (C–O–C). ^1H NMR (500 MHz, DMSO- d_6): δ 8.18 (s, 1H), 7.67 (s, 1H), 7.54 (d, J = 8.4 Hz, 2H), 7.28 (d, J = 8.4 Hz, 2H), 7.09 (s, 1H), 6.98 (s, 2H), 4.26 (s, 1H), 2.53 (s, 2H), 2.26 (d, J = 16.0 Hz, 1H), 2.13 (d, J = 16.0 Hz, 1H), 1.05 (s, 3H), 0.98 (s, 3H). ^{13}C NMR (126 MHz, DMSO- d_6): δ 195.488, 162.536, 158.418, 143.377, 135.343, 129.363, 128.537, 128.288, 120.494, 119.342, 117.979, 112.335, 57.987, 49.963, 35.056, 31.648, 28.183, 26.811. MS (ESI-QTOF) for $\text{C}_{21}\text{H}_{20}\text{N}_4\text{O}_2$ $[\text{M} + \text{H}]^+$ calculated 361.1659, found 361.1663.

4-(4-(1H-1,2,4-Triazol-1-yl)phenyl)-2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (14c). Yield: 85%.



White solid. M.p.: 254–256 °C. IR (ν_{\max} , cm^{-1}): 3440, 3265 ($-\text{NH}_2$), 3105 ($=\text{C}-\text{H}$), 2181 (CN), 1681 ($\text{C}=\text{O}$), 1594 ($\text{C}=\text{C}$), 1224 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 9.23 (s, 1H), 8.22 (s, 1H), 7.77 (d, $J = 8.4$ Hz, 2H), 7.33 (d, $J = 8.6$ Hz, 2H), 7.09 (s, 2H), 4.27 (s, 1H), 2.54 (s, 2H), 2.27 (d, $J = 16.0$ Hz, 1H), 2.12 (d, $J = 16.0$ Hz, 1H), 1.05 (s, 3H), 0.97 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 195.719, 162.670, 158.476, 152.438, 152.217, 144.481, 142.283, 135.189, 128.547, 119.601, 112.296, 57.786, 49.943, 35.229, 31.821, 28.346, 26.839. MS (ESI-QTOF) for $\text{C}_{20}\text{H}_{19}\text{N}_5\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 361.1612, found 362.1613.

4-(4-(1H-Benzod[*d*]imidazol-1-yl)phenyl)-2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (14d). Yield: 86%. Off-white solid. M.p.: 249–251 °C. IR (ν_{\max} , cm^{-1}): 3353, 3290 ($-\text{NH}_2$), 3086 ($=\text{C}-\text{H}$), 2196 (CN), 1691 ($\text{C}=\text{O}$), 1613 ($\text{C}=\text{C}$), 1214 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.56 (s, 1H), 7.77 (d, $J = 7.2$ Hz, 1H), 7.63–7.56 (m, 3H), 7.41–7.29 (m, 4H), 7.11 (s, 2H), 4.32 (s, 1H), 2.56 (s, 2H), 2.29 (d, $J = 16.0$ Hz, 1H), 2.17 (d, $J = 16.0$ Hz, 1H), 1.06 (s, 3H), 1.01 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 195.805, 162.834, 158.591, 144.289, 143.751, 143.195, 134.441, 132.982, 128.806, 128.720, 123.575, 119.937, 119.707, 112.402, 110.722, 57.910, 49.991, 35.257, 31.859, 28.298, 27.031. MS (ESI-QTOF) for $\text{C}_{25}\text{H}_{22}\text{N}_4\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 411.1816, found 411.1818.

4-(4-(1H-benzod[*d*]1,2,3-triazol-1-yl)phenyl)-2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (14e). Yield: 90%. Off-white solid. M.p.: 266–268 °C. IR (ν_{\max} , cm^{-1}): 3387, 3304 ($-\text{NH}_2$), 3168 ($=\text{C}-\text{H}$), 2181 (CN), 1681 ($\text{C}=\text{O}$), 1642 ($\text{C}=\text{C}$), 1205 ($\text{C}-\text{O}-\text{C}$). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 8.19 (d, $J = 7.1$ Hz, 2H), 7.97 (s, 2H), 7.46–7.38 (m, 4H), 7.08 (s, 2H), 4.27 (s, 1H), 2.49 (s, 2H), 2.22 (d, $J = 16.0$ Hz, 1H), 2.08 (d, $J = 16.0$ Hz, 1H), 0.99 (s, 3H), 0.92 (s, 3H). ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$): δ 195.738, 162.718, 158.562, 146.161, 144.394, 138.136, 128.710, 127.645, 120.503, 120.254, 119.591, 118.353, 118.219, 117.988, 112.277, 57.613, 49.924, 35.392, 31.898, 28.289, 26.868. MS (ESI-QTOF) for $\text{C}_{24}\text{H}_{21}\text{N}_5\text{O}_2$ [$\text{M} + \text{H}$] $^+$ calculated 412.1768, found 412.1771.

Cell culturing and maintenance

There are 3 different cancer cell lines were used in this study named MCF 7 (Breast Adenocarcinoma, obtained from ATCC, HTB-22) A549 (Lung Carcinoma, obtained from ATCC, CRM-CCL-185), and HCT116 (Colorectal Carcinoma, obtained from ATCC, CCL-247EMT). The HCT116 was cultured in McCoy's 5A medium supplemented with 10% fetal bovine serum, 2 mM L-glutamine, and antibiotics (penicillin 62.6 $\mu\text{g mL}^{-1}$ and streptomycin 40 $\mu\text{g mL}^{-1}$). The MCF7 and A549 cells were cultured in RPMI-1640 medium supplemented with 10% fetal bovine serum, 2 mM L-glutamine, and antibiotics (penicillin 62.6 $\mu\text{g mL}^{-1}$ and streptomycin 40 $\mu\text{g mL}^{-1}$). All the culturing experiments were performed with 5% CO_2 under controlled humidity.

Determination of cell viability

Cell viability was evaluated using the MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) (Sigma-Aldrich) assay. To perform this assay, cells were seeded into 96-well

culture plates and allowed to adhere overnight to ensure proper attachment. Subsequently, the cells were exposed to various concentrations of the synthesized compounds for a duration of 72 h. DMSO (Merck) was employed as a control group. After the incubation period, each well was treated with an MTT solution at a final concentration of 0.4 mg mL^{-1} and incubated for an additional 2–3 hours at 37 °C. Following the completion of incubation, the culture medium was replaced with 100 μL of DMSO. The absorbance of the formazan product was measured at 450 nm using an ASYS UVM340 microplate reader. The IC_{50} value for each compound was determined using GraphPad Prism 9 software. The IC_{50} values were estimated based on the dose–response curves obtained from survival as a function of dose. The data were derived from three independent experiments and averaged to ensure the robustness and reliability of the results.

Morphological evaluation

Cell lines HCT116, A549, and MCF7 were cultured in appropriate culture plates and allowed to adhere overnight. Following this, the cells were exposed to the experimental compounds for 24 hours. After the incubation period, the cells were stained using Hoechst 33542 (Sigma-Aldrich) and subsequently observed under a fluorescence microscope (Olympus BX53). Additionally, cellular morphology was assessed using a modified Giemsa Stain solution under a brightfield microscope.

Colony formation assay

In this study, we employed well-established cell lines, HCT-116, MCF7, and A549, each seeded at a density of 170, 140, and 120 cells per well, respectively, in 12-well plates. Before treatment, the cells were allowed to attach and subsequently subjected to various concentrations of test compounds for 24 hours. Following this incubation period, the culture medium was replaced, and the cells were maintained in culture for an additional 8 days. Upon completion of the 8-day culture period, the cells were subjected to a series of preparatory steps. Firstly, thorough washing with PBS was performed to remove any residual compounds. Next, the cells were fixed using 99.5% methanol for 20 minutes, ensuring cellular preservation. Subsequently, a 0.5% crystal violet solution was used to stain the cells for a further 20 minutes. The stained plates were air-dried to facilitate the visualization of colonies. To quantitatively assess the effects of the compounds on cell viability, the visible colonies were meticulously counted using a colony counter. The percentage of viable cells was then calculated by comparing the results to the control.

Cell cycle analysis

Cell lines HCT116, A549, and MCF7 were cultured in appropriate culture plates and allowed to adhere overnight. Subsequently, the cells were exposed to the test compounds for 24 and 48 hours. After the incubation, the cells were fixed using ice-cold ethanol (75%) and stored at a temperature of -20 °C overnight. The cells were then subjected to two washes with PBS and centrifuged. After centrifugation, the cells were



stained with a solution containing propidium iodide (PI) at a concentration of $20 \mu\text{g } \mu\text{L}^{-1}$ and RNase A at a concentration of $50 \mu\text{g } \mu\text{L}^{-1}$ in PBS. This staining process was carried out for 30 minutes at room temperature. Subsequently, the cell cycle distribution was analyzed using a Guava easyCyte 8 flow cytometer (Merck Millipore) in conjunction with FlowJo v10 software.

Evaluation of DNA DSBs induction

The cell lines HCT116, A549, and MCF7 were cultured and seeded onto tissue culture plates, allowing them to adhere overnight. The following day, the cells were exposed to the experimental compounds for durations of 24 and 48 h. Mitoxantrone ($1 \mu\text{M}$) was utilized as a reference compound. Subsequently, the cells were harvested through trypsinization, fixed in 75% ethanol, and stored at $-20 \text{ }^\circ\text{C}$ until further analysis. To prepare the cells for analysis, they were rehydrated with PBS while placed on ice for 5 minutes. Following this, the cells were permeabilized with a solution consisting of 0.2% Triton X-100 in PBS and incubated at room temperature for 15 minutes. For labeling purposes, the cells were treated with Alexa Fluor 488-conjugated mouse anti-p- γH2AX (Ser139) antibody (#613406, BioLegend, USA) at a dilution of 1:200, and incubated at $37 \text{ }^\circ\text{C}$ for 1.5 hours. Afterwards, the cells were washed with PBS and stained with propidium iodide ($20 \mu\text{g } \mu\text{L}^{-1}$) and RNase ($50 \mu\text{g } \mu\text{L}^{-1}$) from Thermo Fisher Scientific for 20 minutes. Analysis of the labeled cells was analyzed using a Guava EasyCyte 8 cell sorter (Merck Millipore, USA) and FlowJo v10 software (BD Life Sciences, USA).

Immunofluorescence

The cell lines HCT116, A549, and MCF7 were cultured and seeded onto tissue culture plates, allowing them to adhere overnight. On the subsequent day, the cells were exposed to the tested compounds at their IC_{90} concentration for the specified duration. After exposure, the cells were washed with PBS and fixed with 4% paraformaldehyde (PFA) in PBS for 15 minutes at room temperature. Subsequently, the cells were permeabilized with 0.25% Triton X-100 in PBS for 15 minutes. To minimize non-specific binding, the cells were then blocked with 3% bovine serum albumin (BSA) in PBS for 1 hour at RT. The cells were incubated with Alexa Fluor 488-conjugated mouse anti-H2AX (pS139) antibody (#560445, diluted 1:200) from BD Pharmingen for 1.5 h at $37 \text{ }^\circ\text{C}$ in a humidified chamber. Following antibody incubation, the cells were washed three times for 10 minutes each with PBS-T. For nuclear staining, the cells were treated with $0.25 \mu\text{g } \text{mL}^{-1}$ 4',6-diamidino-2-phenylindole (DAPI) for 15 minutes and then mounted onto slides using a mounting medium consisting of PBS-glycerol (90%) containing 2.5% (w/v) 4-diazobicyclo-(2,2,2-octane) (DABCO). The coverslips were washed twice with PBS-T and stained again with $0.25 \mu\text{g } \text{mL}^{-1}$ DAPI. Finally, images were acquired using an LSM 800 inverted laser scanning confocal microscope (Carl Zeiss, Dresden, Germany) equipped with an Airyscan detector and a $\times 63$ 1.4 NA Plan Achromat objective (Carl Zeiss).

Apoptosis detection

To assess the proapoptotic activity of the compounds, cellular experiments were conducted following standard protocols. Initially, cells were seeded onto suitable Petri dishes and allowed to attach overnight. Subsequently, the cells were treated with the compounds at their respective IC_{50} concentrations for 24 and 48 hours. To establish a baseline, DMSO at a concentration of 1% (Merck) and Mitoxantrone at $1 \mu\text{M}$ (Sigma-Aldrich) were utilized as reference samples. After the treatment period, the cells were harvested and subjected to staining with FITC-Annexin V (Thermo Fisher, V13242) as per the manufacturer's instructions. To discriminate between viable and non-viable cells, propidium iodide (Thermo Fisher) staining was subsequently performed. The stained cells were then analyzed using a Guava easyCyte 8 cell sorter (Merck Millipore). The acquired data were further processed and analyzed using FlowJo v10 software.

Statistical analyses

The data are presented 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.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

FK and ACG thank the Ministry of Education (MoE) for providing the fellowships. MC is thankful to the DST-INSPIRE (IF190417) for the fellowship. JB sincerely acknowledges the DST-SERB for a research grant (file name: EEQ/2020/000303) and the Director, National Institute of Technology Calicut for providing research facilities. NMT is grateful to DST-SERB for providing fellowship from the project (EEQ/2020/000303). All the authors are thankful to CMC, and NIT Calicut for the characterization data.

References

- 1 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).
- 2 J. Ferlay, M. Colombet, I. Soerjomataram, D. M. Parkin, M. Piñeros, A. Znaor and F. Bray, *Int. J. Cancer*, 2021, **149**, 778–789, DOI: [10.1002/ijc.33588](https://doi.org/10.1002/ijc.33588).
- 3 C. de Martel, D. Georges, F. Bray, J. Ferlay and G. M. Clifford, *Lancet Global Health*, 2020, **8**, e180–e190, DOI: [10.1016/S2214-109X\(19\)30488-7](https://doi.org/10.1016/S2214-109X(19)30488-7).



- 4 J. Fares, M. Y. Fares, H. H. Khachfe, H. A. Salhab and Y. Fares, *Signal Transduction Targeted Ther.*, 2020, **5**, 28, DOI: [10.1038/s41392-020-0134-x](https://doi.org/10.1038/s41392-020-0134-x).
- 5 American Cancer Society. Learn About Cancer. <https://www.cancer.org/cancer.html>.
- 6 American Society of Clinical Oncology. Stages of Cancer. <https://www.cancer.net/navigating-cancer-care/diagnosing-cancer/stages-cancer>.
- 7 National Cancer Institute. What is Cancer? <https://www.cancer.gov/about-cancer/understanding/what-is-cancer>.
- 8 G. Feuer, in *Progress in Medicinal Chemistry*, ed. G. P. Ellis and G. B. West, North-Holland Publishing Company, New York, 1st edn, 1974, vol. 10, p.85.
- 9 F. M. Dean, *Naturally Occurring Oxygen Ring Compounds*, Butterworth-Heinemann, London, 1963, pp. 176–220.
- 10 S. A. Patil, R. Patil, L. M. Pfeffer and D. D. Miller, *Future Med. Chem.*, 2013, **5**(14), 1647–1660, DOI: [10.4155/fmc.13.126](https://doi.org/10.4155/fmc.13.126).
- 11 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).
- 12 M. K. Katiyar, G. K. Dhakad, Shivani, S. Arora, S. Bhagat, T. Arora and R. Kumar, *J. Mol. Struct.*, 2022, **1263**, 133012, DOI: [10.1016/j.molstruc.2022.133012](https://doi.org/10.1016/j.molstruc.2022.133012).
- 13 B. Borah, K. D. Dwivedi and L. R. Chowhan, *Polycyclic Aromat. Compd.*, 2022, **42**, 5893–5937, DOI: [10.1080/10406638.2021.1962923](https://doi.org/10.1080/10406638.2021.1962923).
- 14 M. S. Malik, H. Ather, S. M. Asif Ansari, A. Siddiqua, Q. M. S. Jamal, A. H. Alharbi, M. M. Al-Rooqi, R. S. Jassas, E. M. Hussein, Z. Moussa, R. J. Obaid and S. A. Ahmed, *Pharmaceuticals*, 2023, **16**, 1–13, DOI: [10.3390/ph16030333](https://doi.org/10.3390/ph16030333).
- 15 W. Kemnitzer, J. Drewe, S. Jiang, H. Zhang, C. Crogan-Grundy, D. Labreque, M. Bubenick, G. Attardo, R. Denis, S. Lamothe, H. Gourdeau, B. Tseng, S. Kasibhatla and X. C. Sui, *J. Med. Chem.*, 2008, **51**, 417–423, DOI: [10.1021/jm7010657](https://doi.org/10.1021/jm7010657).
- 16 W. Kemnitzer, S. Kasibhatla, S. Jiang, H. Zhang, J. Zhao, S. Jia, L. Xu, C. Crogan-Grundy, R. Denis, N. Barriault, L. Vaillancourt, S. Charron, J. Dodd, G. Attardo, D. Labreque, S. Lamothe, H. Gourdeau, B. Tseng, J. Drewe and S. X. Cai, *Bioorg. Med. Chem. Lett.*, 2005, **15**, 4745–4751, DOI: [10.1016/j.bmcl.2005.07.066](https://doi.org/10.1016/j.bmcl.2005.07.066).
- 17 J. M. Doshi, D. Tian and C. Xing, *J. Med. Chem.*, 2006, **49**, 7731–7739, DOI: [10.1021/jm060968r](https://doi.org/10.1021/jm060968r).
- 18 D. A. N. L. Wood, D. Panda, T. R. Wiernicki, L. Wilson, M. A. N. N. Jordan, J. A. I. P. A. L. Singh and D. L. W. Indiana, *Mol. Pharmacol.*, 1997, **52**, 437–444, DOI: [10.1124/mol.52.3.437](https://doi.org/10.1124/mol.52.3.437).
- 19 A. M. El-Agrody, A. M. Fouda, M. A. Assiri, A. Mora, T. E. Ali, M. M. Alam and M. Y. Alfaiif, *Med. Chem. Res.*, 2020, **29**, 617–629, DOI: [10.1007/s00044-019-02494-3](https://doi.org/10.1007/s00044-019-02494-3).
- 20 N. R. Emmadi, K. Atmakur, G. K. Chityal, S. Pombala and J. B. Nanubolu, *Bioorg. Med. Chem. Lett.*, 2012, **22**, 7261–7264, DOI: [10.1016/j.bmcl.2012.09.018](https://doi.org/10.1016/j.bmcl.2012.09.018).
- 21 N. R. Kamdar, D. D. Haveliwala, P. T. Mistry and S. K. Patel, *Med. Chem. Res.*, 2011, **20**, 854–864, DOI: [10.1007/s00044-010-9399-x](https://doi.org/10.1007/s00044-010-9399-x).
- 22 G. Singh, A. Sharma, H. Kaur and M. P. S. Ishar, *Chem. Biol. Drug Des.*, 2016, **87**, 213–223, DOI: [10.1111/cbdd.12653](https://doi.org/10.1111/cbdd.12653).
- 23 C. Bingi, E. Narender Reddy, M. Chennapuram, Y. Poornachandra, C. G. Kumar, N. Jagadeesh Babu and K. Atmakur, *Bioorg. Med. Chem. Lett.*, 2015, **25**, 1915–1919, DOI: [10.1016/j.bmcl.2015.03.034](https://doi.org/10.1016/j.bmcl.2015.03.034).
- 24 N. D. Vala, H. H. Jardosh and M. P. Patel, *Chin. Chem. Lett.*, 2016, **27**, 168–172, DOI: [10.1016/j.cclet.2015.09.020](https://doi.org/10.1016/j.cclet.2015.09.020).
- 25 E. Rajanarendar, M. Nagi Reddy, S. Rama Krishna, K. Rama Murthy, Y. N. Reddy and M. V. Rajam, *Eur. J. Med. Chem.*, 2012, **55**, 273–283, DOI: [10.1016/j.ejmech.2012.07.029](https://doi.org/10.1016/j.ejmech.2012.07.029).
- 26 N. C. Lazzara, R. J. Rosano, P. P. Vagadia, M. T. Giovine, M. W. Bezpalko, N. A. Piro, W. S. Kassel, W. J. Boyko, D. L. Zubris, K. K. Schrader, D. E. Wedge, S. O. Duke and R. M. Giuliano, *J. Org. Chem.*, 2019, **84**, 666–678, DOI: [10.1021/acs.joc.8b02490](https://doi.org/10.1021/acs.joc.8b02490).
- 27 J. H. Roireau, R. J. Rosano, N. C. Lazzara, T. Chen, J. Bajsa-Hirschel, K. K. Schrader, S. O. Duke, D. Wykoff and R. M. Giuliano, *J. Agric. Food Chem.*, 2020, **68**, 9906–9916, DOI: [10.1021/acs.jafc.0c02564](https://doi.org/10.1021/acs.jafc.0c02564).
- 28 Y. Dgachi, O. M. Bautista-Aguilera, M. Bencheqroun, H. Martin, A. Bonet, D. Knez, J. Godyń, B. Malawska, S. Gobec, M. Chioua, J. Janockova, O. Soukup, F. Chabchoub, J. Marco-Contelles and L. Ismaili, *Molecules*, 2016, **21**, 1–15, DOI: [10.3390/molecules21050634](https://doi.org/10.3390/molecules21050634).
- 29 A. Foroumadi, S. Emami, M. Sorkhi, M. Nakhjiri, Z. Nazarian, S. Heydari, S. K. Ardestani, F. Poorrajab and A. Shafiee, *Chem. Biol. Drug Des.*, 2010, **75**, 590–596, DOI: [10.1111/j.1747-0285.2010.00959.x](https://doi.org/10.1111/j.1747-0285.2010.00959.x).
- 30 K. S. Lee, L. Y. Khil, S. H. Chae, D. Kim, B. H. Lee, G. S. Hwang, C. H. Moon, T. S. Chang and C. K. Moon, *Life Sci.*, 2006, **78**, 1091–1097, DOI: [10.1016/j.lfs.2005.06.017](https://doi.org/10.1016/j.lfs.2005.06.017).
- 31 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).
- 32 V. T. Angelova, Y. Voynikov, P. Andreeva-Gateva, S. Surcheva, N. Vassilev, T. Pencheva and J. Tchekalarova, *Med. Chem. Res.*, 2014, **26**, 1884–1896, DOI: [10.1007/s00044-017-1902-1](https://doi.org/10.1007/s00044-017-1902-1).
- 33 N. Jain, R. M. Kanojia, J. Xu, G. Jian-Zhong, E. Pacia, M. T. Lai, F. Du, A. Musto, G. Allan, D. W. Hahn, S. Lundeen and Z. Sui, *J. Med. Chem.*, 2006, **49**, 3056–3059, DOI: [10.1021/jm060353u](https://doi.org/10.1021/jm060353u).
- 34 K. Takao, H. Yahagi, Y. Uesawa and Y. Sugita, *Bioorg. Chem.*, 2018, **77**, 436–442, DOI: [10.1016/j.bioorg.2018.01.036](https://doi.org/10.1016/j.bioorg.2018.01.036).
- 35 K. Takao, S. U. 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).
- 36 A. P. Sarkate, V. S. Dofe, S. V. Tiwari, D. K. Lokwani, K. S. Karnik, D. D. Kamble, M. H. S. H. Ansari, S. Dodamani, S. S. Jalalpure, J. N. Sangshetti, R. Azad, P. V. L. S. Burra and S. V. Bhandari, *Bioorg. Med. Chem. Lett.*, 2021, **40**, 127916, DOI: [10.1016/j.bmcl.2021.127916](https://doi.org/10.1016/j.bmcl.2021.127916).
- 37 H. E. A. Ahmed, M. A. A. El-Nassag, A. H. Hassan, H. M. Mohamed, A. H. Halawa, R. M. Okasha, S. Ihmaid, S. M. Abd El-Gilil, E. S. A. E. H. Khattab, A. M. Fouda, A. M. El-Agrody, A. Aljuhani and T. H. Afifi, *J. Mol. Struct.*, 2019, **1186**, 212–223, DOI: [10.1016/j.molstruc.2019.03.012](https://doi.org/10.1016/j.molstruc.2019.03.012).
- 38 R. Kaur, F. Naaz, S. Sharma, S. Mehndiratta, M. K. Gupta, P. M. S. Bedi and K. Nepali, *Med. Chem. Res.*, 2015, **24**, 3334–3349, DOI: [10.1007/s00044-015-1382-0](https://doi.org/10.1007/s00044-015-1382-0).



- 39 G. Gopinath, V. Sankeshi, S. Perugu, M. D. Alaparathi, S. Bandaru, V. K. Pasala, P. R. Chittineni, G. L. D. Krupadanam and S. R. Sagurthi, *Eur. J. Med. Chem.*, 2016, **124**, 750–762, DOI: [10.1016/j.ejmech.2016.08.070](https://doi.org/10.1016/j.ejmech.2016.08.070).
- 40 Z. Zhang, C. Wang, L. Ma, X. Jiang, C. Wu, Y. Wang, Y. Jiang, W. Zheng, Y. Yang, Y. Ma and J. Yang, *Biochem. Biophys. Res. Commun.*, 2019, **511**, 381–386, DOI: [10.1016/j.bbrc.2019.02.064](https://doi.org/10.1016/j.bbrc.2019.02.064).
- 41 A. M. A. Lorza, H. Ravi, R. C. Philip, J. P. Galons, T. P. Trouard, N. A. Parra, D. D. Von Hoff, W. L. Read, R. Tibes, R. L. Korn and N. Raghunand, *Sci. Rep.*, 2020, **10**, 1–12, DOI: [10.1038/s41598-020-71246-w](https://doi.org/10.1038/s41598-020-71246-w).
- 42 D. Panda, J. P. Singh and L. Wilson, *J. Biol. Chem.*, 1997, **272**, 7681–7687, DOI: [10.1074/jbc.272.12.7681](https://doi.org/10.1074/jbc.272.12.7681).
- 43 F. Schmitt, M. Gold, M. Rothmund, I. Andronache, B. Biersack, R. Schobert and T. Mueller, *Eur. J. Med. Chem.*, 2019, **163**, 160–168, DOI: [10.1016/j.ejmech.2018.11.055](https://doi.org/10.1016/j.ejmech.2018.11.055).
- 44 D. Rajguru, B. S. Keshwal and S. Jain, *Med. Chem. Res.*, 2013, **22**, 5934–5939, DOI: [10.1007/s00044-013-0586-4](https://doi.org/10.1007/s00044-013-0586-4).
- 45 K. Parthasarathy, C. Praveen, C. Balachandran, P. Senthil Kumar, S. Ignacimuthu and P. T. Perumal, *Bioorg. Med. Chem. Lett.*, 2013, **23**, 2708–2713, DOI: [10.1016/j.bmcl.2013.02.086](https://doi.org/10.1016/j.bmcl.2013.02.086).
- 46 L. Braconi, E. Teodori, C. Riganti, M. Coronello, A. Nocentini, G. Bartolucci, M. Pallecchi, M. Contino, D. Manetti, M. N. Romanelli, C. T. Supuran and S. Dei, *J. Med. Chem.*, 2022, **65**, 14655–14672, DOI: [10.1021/acs.jmedchem.2c01175](https://doi.org/10.1021/acs.jmedchem.2c01175).
- 47 S. Wang, S. Q. Wang, Q. Teng, L. Yang, Z. Lei, X. Yuan, J. Huo, X. Chen, M. Wang, B. Yu, Z. Chen and H. Liu, *J. Med. Chem.*, 2020, **63**, 15979–15996, DOI: [10.1021/acs.jmedchem.0c01741](https://doi.org/10.1021/acs.jmedchem.0c01741).
- 48 D. J. Hong, S. H. Jung, J. Kim, D. Jung, Y. G. Ahn, K. H. Suh and K. H. Min, *J. Enzyme Inhib. Med. Chem.*, 2020, **35**, 227–234, DOI: [10.1080/14756366.2019.1693555](https://doi.org/10.1080/14756366.2019.1693555).
- 49 M. M. Heravi and V. Zadsirjan, *RSC Adv.*, 2020, **10**, 44247–44311, DOI: [10.1039/d0ra09198g](https://doi.org/10.1039/d0ra09198g).
- 50 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).
- 51 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).
- 52 H. van de Waterbeemd and E. Gifford, *Nat. Rev. Drug Discovery*, 2003, **2**, 192–204, DOI: [10.1038/nrd1032](https://doi.org/10.1038/nrd1032).
- 53 M. J. Waring, J. Arrowsmith, A. R. Leach, P. D. Leeson, S. Mandrell, R. M. Owen, G. Pairaudeau, W. D. Pennie, S. D. Pickett, J. Wang, O. Wallace and A. Weir, *Nat. Rev. Drug Discovery*, 2015, **14**, 475–486, DOI: [10.1038/nrd4609](https://doi.org/10.1038/nrd4609).
- 54 Y. Sivamani, D. Shanmugarajan, T. Durai Ananda Kumar, S. Faizan, B. Channappa, N. L. Naishima and B. R. Prashantha Kumar, *Comput. Biol. Chem.*, 2021, **95**, 107600, DOI: [10.1016/j.compbiolchem.2021.107600](https://doi.org/10.1016/j.compbiolchem.2021.107600).
- 55 N. Kang, S. Y. Heo, S. H. Cha, G. Ahn and S. J. Heo, *Mar. Drugs*, 2022, **20**, 1–14, DOI: [10.3390/md20060399](https://doi.org/10.3390/md20060399).
- 56 M. Szumilak, W. Lewgowd and A. Stańczak, *Acta Pol. Pharm.*, 2016, **73**, 1191–1200.

