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Xanthone synthetic derivatives with high anticandidal activity and positive mycostatic selectivity index values

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With the current massive increases in drug-resistant microbial infection as well as the significant role of fungal infections in the death toll of COVID-19, discovering new antifungals is extremely important. Natural and synthetic xanthenes are promising derivatives, although only few reports have demonstrated their antifungal mechanism of action in detail. Newly synthesized by us xanthone derivative 44 exhibited strong antifungal activity against reference and fluconazole resistant *C. albicans* strains. Our results indicate that the most active compounds 42 and 44 are not substrates for fungal ABC transporters (Cdr1p and Cdr2p) and Mdr1p, the main representative of the major facilitator superfamily efflux pumps, membrane proteins that are responsible for the development of resistance. Moreover, fungicidal mode of action reduces the probability of persistent or recurrent infections and resistance development. In this light, the demonstrated killing activity of the examined derivatives is their undoubted advantage. Novel synthesized compounds exhibited moderate cytotoxicity against human cell lines, although the selectivity index value for human pathogenic strains remained favourable. Our results also indicate that novel synthesized compounds 42 and 44 with antifungal activity target yeast topoisomerase II activity. In summary, further validation of xanthenes applicability as antifungals is highly valuable.

Fungal microorganisms are etiological factors of severe, often deadly, infectious diseases, especially in immunocompromised patients. The number of these patients is growing rapidly, not only because of diseases resulting in immunodeficiency, like AIDS but also as a consequence of the frequent use of therapies that affect the human immune defense system (e.g. anticancer therapy with cytostatics, steroid therapy, use of the immunosuppressive agents in transplant patients). Systemic mycoses are caused in these patients mainly by yeast-like microorganisms from the *Candida* genus, especially *Candida albicans* and *Candida glabrata*, and filamentous fungi from the *Aspergillus* genus¹. On the other hand, many fungal microorganisms are known as one of the most frequent reasons of nosocomial infections. *C. albicans* is considered the fourth most popular etiological agent of nosocomial infections worldwide. Moreover, chemotherapeutics used in clinical treatment have become factors stimulating the selection of resistant cells. A newly described pathogen, *Candida auris*, is an emerging multidrug-resistant organism that poses a global threat². Additionally, invasive fungal infections complicate the clinical course of COVID-19 and are associated with a significant increase in mortality, especially in critically ill patients admitted to an intensive care unit³. Thus, with the current massive increases in drug-resistant microbial infections as well as the significant role of fungal infections in the death toll of COVID-19, discovering new antifungal compounds is extremely important.

There are several approaches in novel drug discovery. First of all, researchers are looking for new drugs targeting old pathways (e.g., ergosterol synthesis)⁴ or cell membranes⁵, while others are trying to find out new solutions. The biosynthesis of fungal proteins, DNA, and other essential molecules is extremely important^{6,7}. As long as new targets are concerned, our group is looking for new drugs targeting fungal topoisomerases. Significant work has been done on the structure and function of topoisomerase I and II in fungi and results indicated that

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their activities are crucial for some specific strains^{8–10}. Moreover, inhibition of yeast topoisomerase II resulted in antifungal activity^{11,12} and even managed to overcome fluconazole-resistance^{13,14}.

Natural xanthone derivatives are a promising group of antifungal compounds^{15–17}. They are present in nature as metabolites of various plant, lichen, fungal, and bacterial species^{18,19}. The interesting structural scaffold and biological efficacy of those compounds lead many scientists to synthesize xanthone derivatives for the development of new prospective drug candidates as anticancer, antimicrobial, antimalarial, anti-HIV, antioxidant, anti-inflammatory, and antimalarial agents²⁰. Several articles have been published highlighting the antifungal activity of synthetic xanthone analogues^{15,21–23}, although only a few have been examined thoroughly in order to define their mechanism of action. According to previous reports, 1,2-dihydroxyxanthone is the most active compound against all fungal strains tested, showing its effect on sterol biosynthesis by reducing the amount of ergosterol detected²⁴.

Emerging from previous studies performed for xanthone analogues as potential antimicrobials^{15–24}, we have decided to analyze the antifungal activity of four new groups of compounds. As xanthone derivatives with anti-cancer activity are reported to be effective human topoisomerase inhibitors²⁰, we have also analyzed the inhibitory effect of selected derivatives on yeast topoisomerase II activity (yTOPO II).

Results and discussion

Synthesis of xanthone derivatives. The structures of xanthone derivatives synthesized in this study are divided into four groups (Fig. 1). Compounds 1–16 and 34 were prepared according to a previously published procedure^{25,26}, while 25, 26, 35–38, and 41–45 are newly synthesized analogues (Table 1). These groups (I–IV) differ in the nitro or amino substitution and the presence of a pyrazole or a benzene fused ring on the xanthone core (Fig. 1).

The list of xanthone and benzoxanthone analogues analyzed in this study is presented in Table 1.

For the synthesis of the new compounds we have used an analogous synthetic procedure, with slight alterations (Figs. 2 and 3). Briefly, the reaction of ethyl salicylate (17) or ethyl 3-hydroxy-2-naphthoate (27) with 2,4-dichloronitrobenzene (18) afforded a mixture of the isomeric diarylethers 19, 20 and 28, 29 respectively. Trituration of both mixtures with methanol resulted in pure 20 and 29, while we obtained a 1/1 mixture of 19, 20, and 28, 29, which could not be further purified. For the synthesis of the ortho-substituted amines 25, 26 and 34, each of the above mixtures was saponified under mild conditions. Without further purification, the resulting mixture of acids 21, 22, and 30, 31, was ring closed upon treatment with polyphosphoric acid (PPA) to afford nitro compounds 23, 24, and 32, 33 respectively, as inseparable mixtures. Reaction of the above mixtures with the suitable amines resulted in the aminoderivatives 2, 5, 25, 26, and 9, 34 respectively. Each aminoderivative was isolated in pure form by column chromatography and identified by means of ¹H and ¹³C spectral data, using both direct and long-range experiments (HMBC and HMQC). In order to prepare the corresponding p-substituted nitro derivatives 35–38 and 41–45, ethyl ester 29 was saponified and ring closed upon treatment with PPA to afford the nitrosubstituted benzoxanthone 32 in pure form. Consequently, the amino derivatives 35–38 were prepared upon nucleophilic substitution of the chloro group of 32 by the appropriately substituted amines. For

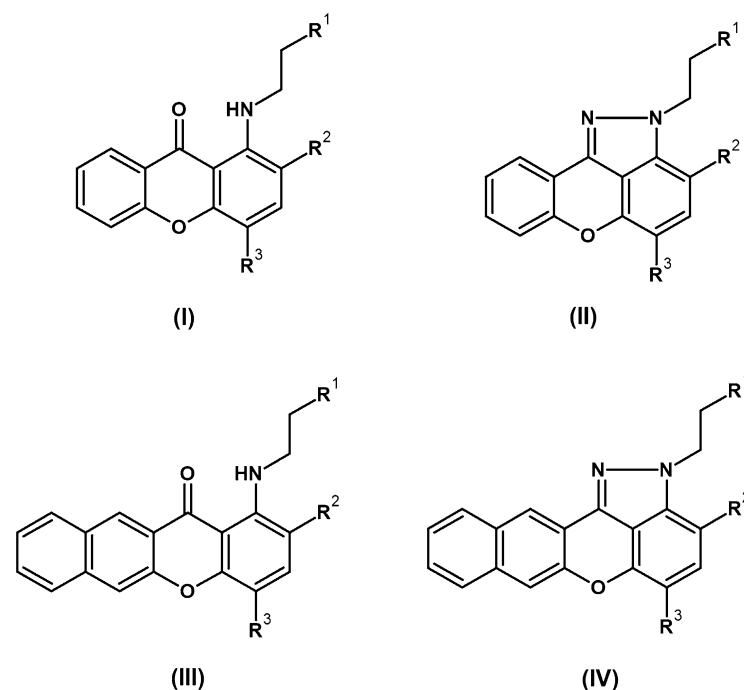


Figure 1. Overall structures of derivatives that were analysed in this study. R¹—aminosubstitution, R² or R³—NO₂ or H.








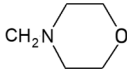
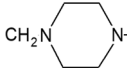
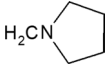

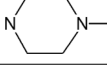



Compound	R ¹	R ²	R ³
Group (I)			
1	N(CH ₂ CH ₃) ₂	H	NO ₂
2	N(CH ₂ CH ₃) ₂	H	H
3		H	NO ₂
4		H	H
25	N(CH ₂ CH ₃) ₂	NO ₂	H
26		NO ₂	H
Group (II)			
5		H	H
6		H	NO ₂
7	N(CH ₂ CH ₃) ₂	H	NO ₂
8	N(CH ₃) ₂	H	H
Group (III)			
9		H	NO ₂
10	N(CH ₂ CH ₃) ₂	H	NO ₂
11	N(CH ₃) ₂	H	H
34		NO ₂	H
35	OCH ₂ CH ₂ OH	H	NO ₂
36	CH ₂ OH	H	NO ₂
37	OH	H	NO ₂
38	CH ₂ OCH ₃	H	NO ₂
41		H	NO ₂
42		H	NO ₂
43		H	NO ₂
44		H	NO ₂
45		H	NO ₂
Group (IV)			
12		H	H
13	N(CH ₂ CH ₃) ₂	H	NO ₂
14		H	NO ₂
15		H	H
16	N(CH ₂ CH ₃) ₂	H	NO ₂

Table 1. The list of xanthone (group I and II) and benzoxanthone (group III and IV) analogues.

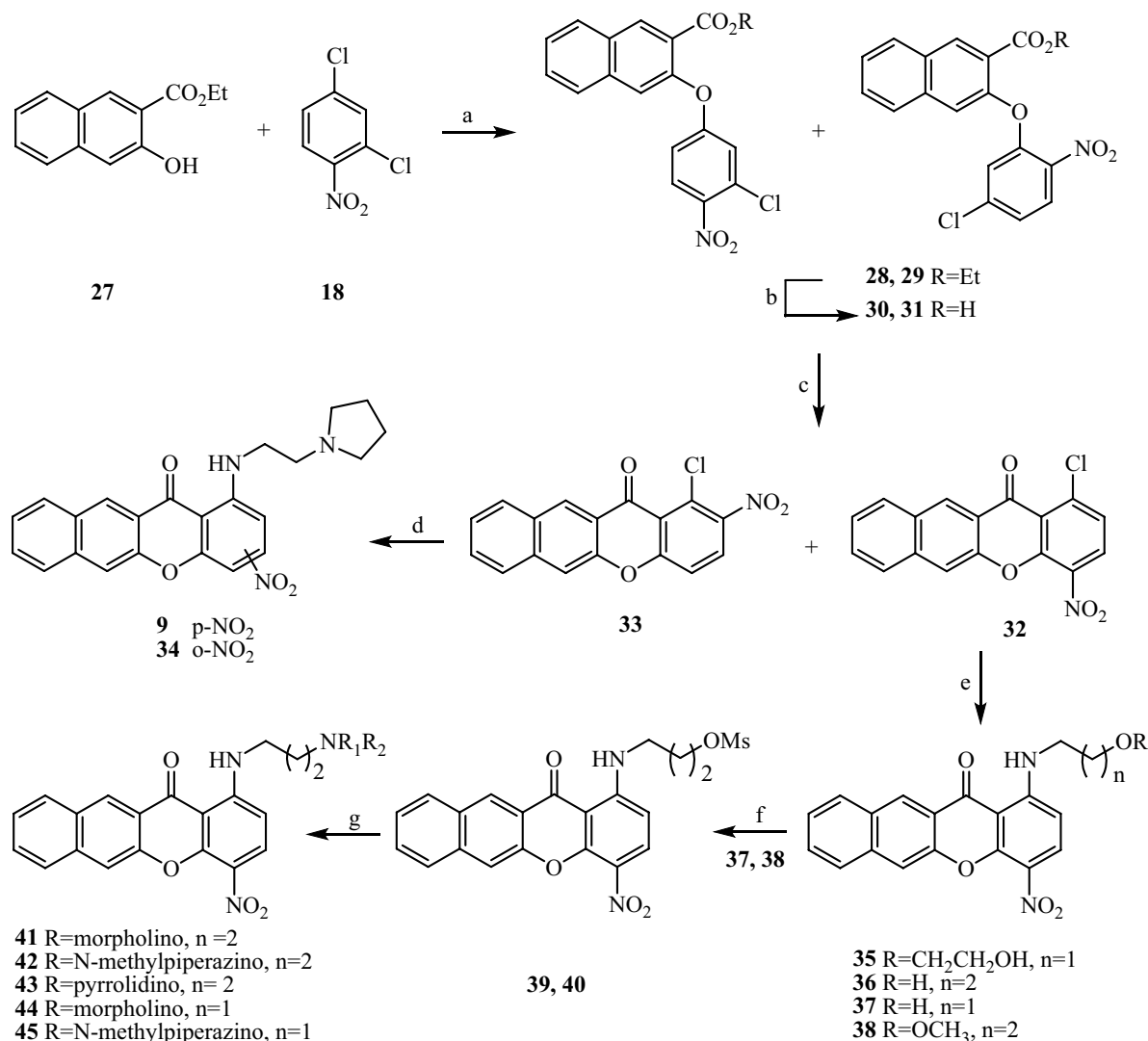


Figure 3. Reaction and conditions: (a) K₂CO₃, Cu₂O, DMF dry, 110 °C; (b) NaOH 40%, EtOH, rt; (c) PPA, 110 °C; (d) suitable amine, pyridine, reflux; (e) suitable amine, pyridine, reflux; (f) MsCl, Et₃N, THF, rt; (g) suitable amine, EtOH, reflux.

Molecular mechanism of antifungal activity. *Killing activity of selected derivatives.* To establish the possible mode of action of xanthone and benzoxanthone analogues we have analyzed the killing activity and determined the minimal fungicidal concentrations (MFCs) for selected derivatives (Table 4).

Our results indicate that the mode of action of the most active compounds is fungicidal. The higher fungicidal activity was determined for derivatives **9**, **42** and **44** from group III and **14** from group IV. The use of fungicidal therapy of invasive candidiasis and candidemia is associated with a higher probability of early therapeutic success. A decreased probability of persistent or recurrent infection and resistance development is also expected. In addition, frequent use of available fungistatic drugs like fluconazole promotes drug resistance³⁰. In this light, the demonstrated killing activity of the examined derivatives is their undoubted advantage.

Inhibition of the relaxation activity of yeast topoisomerase II in vitro. The xanthone derivatives act as anticancer agents through several mechanisms of action. The most important are the activation of caspase proteins and the inhibition of protein kinases and topoisomerases³¹. Due to known activity of xanthone derivatives as human topoisomerase II inhibitors^{20,31}, we decided to evaluate the influence of the six more active compounds on the fungal equivalent of that enzyme. The effect of selected compounds on yeast topoisomerase II-mediated relaxation activity suggests their molecular target (Table 5 and Fig. 4).

As reported in Table 5, the most effective inhibitors towards fungal topoisomerase II are compounds **42** and **44**. Surprisingly, no inhibition was observed in the tested concentration range for compound **9**, the closely related to **42** and **44** derivatives. Thus, the molecular mode of action may vary between these compounds and needs more in depth analysis.

Compound	*MIC ₉₀ µg mL ⁻¹				
	<i>Candida albicans</i> ATCC 10231	<i>Candida glabrata</i> ATCC 90030	<i>Candida krusei</i> ATCC 6258	<i>Candida parapsilosis</i> ATCC 22019	<i>Saccharomyces cerevisiae</i> ATCC 9763
1	32	32	32	>64	16
2	>64	>64	64	32	32
3	16	16	16	64	8
4	>64	>64	64	64	64
5	16	>64	32	32	32
6	16	8	16	>64	8
7	16	16	16	>64	16
8	32	64	32	32	16
9	4	4	4	8	4
10	>16	>16	>16	>16	>16
11	>64	>64	>64	>64	>64
12	8	>64	8	>64	8
13	8	4	4	16	2
14	8	4	4	16	4
15	8	32	8	64	4
16	32	16	8	32	16
25	>64	>64	>64	64	64
26	>64	>64	64	64	64
34	32	>32	8	>32	8
35	>64	>64	>64	>64	>64
36	>64	>64	>64	>64	>64
37	>64	>64	>64	>64	>64
38	>64	>64	>64	>64	>64
41	>64	>64	>64	>64	>64
42	8	4	4	8	4
43	>64	>64	>64	>64	>64
44	4	2	2	4	2
45	32	32	32	32	32
Amphotericin B	0.5	1	1	1	0.5

Table 2. Antifungal activity against reference strains. MIC₉₀, minimal inhibitory concentration—a concentration that inhibits 90% of fungal cell growth. * > Means no activity at the concentration mentioned. In this assay, the MIC₉₀ value of amphotericin B was recorded as a positive control. The experiments were performed in three replicates.

Compound	*MIC ₉₀ µg mL ⁻¹						
	<i>Candida albicans</i> ATCC 10231	<i>Candida albicans</i> B3	<i>Candida albicans</i> B4	<i>Candida albicans</i> Gu4	<i>Candida albicans</i> Gu5	<i>Candida albicans</i> F2	<i>Candida albicans</i> F5
3	16	16	32	16	32	16	32
5	16	64	64	64	64	32	64
6	16	16	64	16	64	32	32
7	16	32	64	32	32	32	32
8	32	32	32	32	32	32	32
9	4	8	8	8	8	8	8
12	8	64	64	64	64	>64	>64
13	8	16	16	16	16	16	16
14	8	16	16	8	8	8	8
15	8	8	8	8	8	8	8
42	8	8	8	8	8	8	8
44	4	4	4	4	4	4	4
45	32	64	64	64	64	64	64
Fluconazole	8	1	16	4	>64	8	>64

Table 3. Antifungal activity of selected derivatives against clinical strains in comparison with *C. albicans* ATCC 10231. * > Means no activity at the concentration mentioned. In this assay, the MIC₉₀ value of fluconazole was recorded as a positive control. The experiments were performed at least in three replicates.

Compound	*MFC $\mu\text{g mL}^{-1}$			
	<i>Candida albicans</i> ATCC 10231	<i>Candida glabrata</i> ATCC 90030	<i>Candida krusei</i> ATCC 6258	<i>Saccharomyces cerevisiae</i> ATCC 9763
1	64	>128	>128	16
2	>64	>64	>64	64
3	32	128	64	16
4	>64	>64	>64	64
5	64	>128	128	32
6	32	128	64	8
7	64	>128	128	16
8	64	>64	64	32
9	8	8	8	4
10	>16	>16	>16	>16
11	>64	>64	>64	>64
12	>64	>64	>64	>64
13	32	32	8	16
14	16	16	8	8
15	64	64	32	16
16	64	32	32	32
25	>64	>64	>64	>64
26	>64	>64	>64	>64
34	>32	>32	32	32
42	8	8	4	4
44	4	4	4	2
45	32	32	32	32
Amphotericin B	2	2	2	2

Table 4. Fungicidal activity of selected compounds and amphotericin B (MFC—minimal fungicidal concentration—a concentration that inhibits 99% of fungal cell growth). * > Means no 99% killing activity at the concentration mentioned. Higher concentrations have not been tested due to the solubility limitations of some compounds. In this assay, the MFC value of amphotericin B was recorded as a positive control. The experiments were performed at least in three replicates.

Compound	Complete inhibition [μM]	IC ₅₀ [μM]
9	>50*	–
13	>150	–
14	50	18.62 ± 1.81
15	>150	–
42	10	5.86 ± 1.07
44	10	5.36 ± 1.22

Table 5. The concentrations of selected compounds that totally inhibited yeast topoisomerase II-mediated relaxation activity. For selected compounds IC₅₀—inhibitory concentrations representing a concentration that inhibits 50% of enzyme activity were determined. The experiments were performed at least in three replicates. *Higher concentrations have not been tested due to the low solubility of the compound.

Selectivity in relation to mammalian cells. Selected compounds were screened for their in vitro anti-proliferative activity towards a human embryonic kidney cell line (HEK-293) and human liver cancer cell line (HEPG2) using colorimetric MTT assay. Our results were compared with the cytotoxic activity determined previously for human colorectal adenocarcinoma cell line (HT29)²⁶ and are presented in Table 6.

As indicated for etoposide and doxorubicin³², both chemotherapy drugs used to treat different types of cancers, results obtained for newly synthesized compounds are within acceptable cytotoxicity levels. To estimate the selectivity in relation to mammalian cells mycostatic selectivity index (MSI) for *C. albicans* ATCC 10231 was calculated as the ratio of EC₅₀ to MIC₉₀ values after converting the MIC₉₀ value to micromolar concentrations. The results are presented in Table 7.

Positive mycostatic selectivity index values (MSI = EC₅₀/MIC₉₀) were obtained for all selected derivatives, ranging from 0.105 to 0.655. Our results indicate the possibility of using xanthone derivatives as efficient anti-fungal agents. However, it is necessary to improve their selectivity. Compound 44 is the most interesting, for

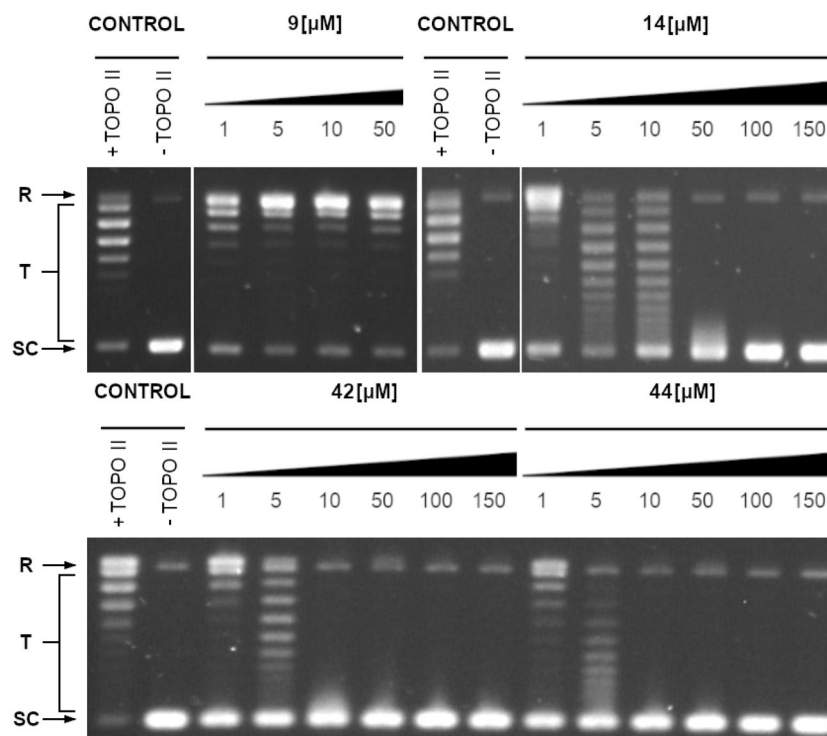


Figure 4. Inhibition of the catalytic activity of purified yeast DNA topoisomerase II by compounds **9**, **14**, **42** and **44** as measured by relaxation. Supercoiled pBR322 plasmid DNA (- TOPO II) was relaxed by purified yeast topoisomerase II in the absence (+ TOPO II) or presence of analyzed compounds **14**, **42** and **44** at 1–150 μM concentrations or 1–50 μM for **9** (higher concentrations have not been tested due to the low solubility of the compound). The resulting topological forms of DNA were separated by gel electrophoreses. SC, supercoiled DNA; R, relaxed DNA; T, DNA topoisomers. Data shown are typical of three independent experiments and one set of representative pictures are shown. Original gels are presented in Supplementary Fig. S1.

Compound	HEK293	HEPG2	HT29
9	3.64 \pm 0.23	1.94 \pm 0.67	–
13	5.73 \pm 1.64	1.92 \pm 0.85	1.85 \pm 0.21*
14	5.99 \pm 1.05	4.2 \pm 0.71	1.83 \pm 0.46*
15	8.33 \pm 0.75	2.53 \pm 0.82	10.7 \pm 3.25*
42	5.38 \pm 0.35	4.88 \pm 0.31	–
44	3.57 \pm 0.25	4.94 \pm 0.56	–
45	48.47 \pm 1.50	26.64 \pm 2.36	–
Etoposide	1.91 \pm 0.97	5.01 \pm 0.22	–
Doxorubicin	6.52 \pm 0.13**	–	–

Table 6. Determination of cytotoxic effect using MTT assay towards HEK293 and HEPG2 cell lines. In vitro growth inhibitory activity of compounds presented as an $\text{EC}_{50} \pm \text{SD}$ (μM) value—a concentration that inhibits 50% of mammalian cell growth. *Antiproliferative activity of compounds **13**–**15** published previously²⁶. **Antiproliferative activity of doxorubicin published previously³². In this assay, the EC_{50} value of etoposide was recorded as a positive control.

which a higher level of MSI using HEPG2 cells was obtained. The HEPG2 cell line, derived from human liver cancer, is the most commonly used in studies on drug metabolism and toxicity³³.

Material and methods

Chemical synthesis. *General experimental procedures.* All commercially available reagents and solvents were purchased from Alfa Aesar and used without any further purification. Melting points were determined on a Büchi apparatus and were uncorrected. All NMR spectra were recorded on 400 or 600 MHz Bruker spectrometers respectively Avance™ DRX and III instruments (Bruker BioSpin GmbH–Rheinstetten, Germany). ¹H NMR

Compound	EC ₅₀ HEK293/MIC ₉₀	EC ₅₀ HEPG2/MIC ₉₀
9	0.400	0.213
13	0.314	0.105
14	0.327	0.229
15	0.408	0.124
42	0.300	0.272
44	0.375	0.518
45	0.655	0.360

Table 7. Mycostatic selectivity index values determined for *C. albicans* ATCC 10231 in relation to mammalian cell lines HEK293 and HEPG2.

(400 and 600 MHz) and ¹³C NMR (101 and 151 MHz, recorded with complete proton decoupling) spectra were obtained with samples dissolved in CDCl₃ or DMSO-*d*₆ with the residual solvent signals used as internal references: 7.26 ppm for CHCl₃, and 2.50 ppm for (CD₃)(CD₂H)S(O) regarding ¹H NMR experiments; 77.2 ppm for CDCl₃ and 39.4 ppm for (CD₃)₂S(O) concerning ¹³C NMR experiments. Chemical shifts (δ) are given in ppm to the nearest 0.01 (1H) or 0.1 ppm (¹³C). The coupling constants (*J*) are given in Hertz (Hz). The signals are reported as follows: (s=singlet, d=doublet, t=triplet, m= multiplet, br=broad). Assignments of ¹H and ¹³C NMR signals were unambiguously achieved with the help of D/H exchange and 2D techniques: COSY, NOESY, HMQC, and HMBC experiments. Compounds **1–8**, **10–16** and **34** were synthesized according to literature and their ¹H NMR and ¹³C NMR were compared with those reported into the literature^{25,26}. Flash chromatography was performed on Merck silica gel (40–63 μm) with the indicated solvent system using gradients of increasing polarity in most cases (Merck KGaA—Darmstadt, Germany). The reactions were monitored by analytical thin-layer chromatography (Merck pre-coated silica gel 60 F254 TLC plates, 0.25-mm layer thickness). Compounds were visualized on TLC plates by both UV radiation (254 and 365 nm). All solvents for absorption and fluorescence experiments were of spectroscopic grade. Mass spectra were recorded on a UPLC Triple TOF–MS {UPLC:Acquity of Waters (USA), SCIEX Triple TOF–MS 5600+ (USA)}.

General procedure for the synthesis of amino substituted xanthenes **25** and **26**. A solution of ethyl salicylate (1.66 g, 10 mmol, **17**), 2,4-dichloronitrobenzene (1.91 g, 9.95 mmol, **18**), K₂CO₃ (1.38 g, 10 mmol) and Cu₂O (142 mg, 1 mmol) in dry DMF (10 mL) was heated at 110 °C for 12 h, under an argon atmosphere. After completion of the reaction the mixture was filtered hot, washed with EtOAc and the filtrate was vacuum evaporated. The oily residue was diluted in CH₂Cl₂, washed with water, dried over anh. Na₂SO₄ and evaporated to dryness. The obtained oily residue was triturated with hot ethanol (15 mL) with stirring and after cooling was filtrated to afford 1.0 g (36%) of **20** and 1.2 g (44%) of an oily mixture of esters **19** and **20** which was used for the next step without any further purification. To a suspension of the above mixture in ethanol (10 mL) a cold 40% NaOH solution was added and the resulting mixture was stirred at room temperature for 40 min. After completion of the reaction the mixture was poured into ice-water and acidified with 18% HCl solution. The resulting mixture of acids **21** and **22** was filtered, dried over P₂O₅ and dissolved in hot polyphosphoric acid. The resulting mixture was stirred at 110 °C for 1 h and upon cooling was poured into ice, and the precipitate was filtered and dried over P₂O₅ to afford xanthenes **23** and **24** which were used for the next step without any further purification. Thus, **23** and **24** and the suitable amine, in dry pyridine, was refluxed for 1 1/2 h. After completion of the reaction, pyridine was vacuum evaporated, the oily residue was diluted in EtOAc, washed with water, dried over anh. Na₂SO₄ and evaporated to dryness. Flash chromatography on silica gel using a mixture of CH₂Cl₂/MeOH 93/7 as the eluent afforded the title compounds **25**, **26** and their isomers **2²⁶** and **5²⁶**, respectively.

Data for 1-((2-(diethylamino)ethyl)amino)-2-nitro-9H-xanthen-9-one (**25**). Yield: 12%; M.p. 123–124 °C (EtOAc-*n*-Hexane); ¹H NMR (400 MHz, CDCl₃) δ 10.89 (s br, D₂O exchange, 1H), 8.26 (dd, *J*=7.9, 1.7 Hz, 1H), 8.09 (d, *J*=9.2 Hz, 1H), 7.69 (td, *J*=7.9, 1.7 Hz, 1H), 7.46–7.36 (m, 2H), 6.57 (d, *J*=9.3 Hz, 1H), 3.03 (q, *J*=5.7 Hz, 2H), 2.72 (t, *J*=5.7 Hz, 2H), 2.65 (q, *J*=7.1 Hz, 4H), 1.08 (t, *J*=7.1 Hz, 6H). ¹³C NMR (400 MHz, CDCl₃) δ 179.0, 161.0, 155.1, 148.2, 135.2, 134.3, 130.6, 126.8, 125.2, 122.2, 117.8, 109.8, 104.0, 52.1, 47.2, 45.4, 12.0. (+) ESI QqToF (m/z): Calcd. for C₁₉H₂₂N₃O₄⁺: [M + H]⁺ 356.1605, found 356.1614.

Data for 2-nitro-1-((2-(pyrrolidin-1-yl)ethyl)amino)-9H-xanthen-9-one (**26**). Yield: 9%; M.p. 173–175 °C (EtOAc-*n*-Hexane); ¹H NMR (400 MHz, CDCl₃) δ 10.88 (t, D₂O exchange, *J*=4.7 Hz, 1H), 8.21 (dd, *J*=7.9, 1.7 Hz, 1H), 8.06 (d, *J*=9.3 Hz, 1H), 7.68 (td, *J*=8.6, 1.7 Hz, 1H), 7.45–7.32 (m, 2H), 6.56 (d, *J*=9.3 Hz, 1H), 3.13 (q, *J*=6.3 Hz, 2H), 2.84 (t, *J*=6.3 Hz, 2H), 2.66 (m, 4H), 1.85 (m, 4H). ¹³C NMR (400 MHz, CDCl₃) δ 178.9, 160.5, 154.6, 147.6, 134.9, 133.9, 130.1, 126.3, 124.8, 121.6, 117.4, 109.2, 103.7, 54.9, 54.0, 46.0, 23.7. (+) ESI QqToF (m/z): Calcd. for C₁₉H₂₀N₃O₄⁺: [M + H]⁺ 354.1448, found 354.1440.

2-nitro-1-((2-(pyrrolidin-1-yl)ethyl)amino)-12H-benzo[*b*]xanthen-12-one (**34**). This compound was prepared by an analogous procedure as described for **26** starting from ethyl 3-hydroxy-2-naphthoate (**27**) with 2,4-dichloronitrobenzene (**18**). Flash chromatography on silica gel using a mixture of CH₂Cl₂/MeOH 94/6 as the

elouent afforded the title compound **34**, and the para nitro substituent compound **9**²⁶. Yield: 12%; M.p. > 220 °C (EtOAc); ¹H NMR (400 MHz, CDCl₃) δ 10.99 t, D₂O exchange, *J* = 6.0 Hz, 1H), 8.85 (s, 1H), 8.17 (d, *J* = 9.4 Hz, 1H), 8.08 (d, *J* = 8.3 Hz, 1H), 7.93 (d, *J* = 8.3 Hz, 1H), 7.86 (s, 1H), 7.66 (t, *J* = 8.3 Hz, 1H), 7.55 (t, *J* = 8.3 Hz, 1H), 6.73 (d, *J* = 9.4 Hz, 1H), 3.31 (q, *J* = 6.5 Hz, 2H), 3.03 (t, *J* = 6.5 Hz, 2H), 2.88 (m, 4H), 1.96 (m, 4H). ¹³C NMR (400 MHz, CDCl₃) δ 180.0, 161.2, 150.6, 148.0, 136.8, 134.5, 130.3, 130.2, 129.8, 129.6, 128.3, 127.2, 126.2, 120.6, 113.4, 108.6, 105.5, 54.0, 43.3, 23.5. (+) ESI QqToF (m/z): Calcd. for C₂₃H₂₂N₃O₄⁺: [M + H]⁺ 401.1605, found 401.1616.

General procedure for the synthesis of amino substituted benzo[*b*]xanthenes **35–38**. A solution of pure benzo[*b*]xanthone **32**, obtained from the synthesis of **34**, and the suitable amine (× 10 equiv.) in dry pyridine (5 mL) was refluxed for 2 h. After completion of the reaction, pyridine was vacuum evaporated, the oily residue was diluted in EtOAc, washed with water, dried over anh. Na₂SO₄ and evaporated to dryness. Flash chromatography on silica gel, using a mixture of CH₂Cl₂/MeOH 100/1 as the elouent provided the title compounds **35–38**.

Data for 1-((2-(2-hydroxyethoxy)ethyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**35**). Yield: 91%; Oil; ¹H NMR (600 MHz, CDCl₃) δ 10.95 (t, *J* = 5.0 Hz, 1H), 8.77 (s, 1H), 8.32 (d, *J* = 9.5 Hz, 1H), 8.01 (d, *J* = 8.3 Hz, 1H), 7.99 (s, 1H), 7.92 (d, *J* = 8.3 Hz, 1H), 7.62 (t, *J* = 8.2 Hz, 1H), 7.51 (t, *J* = 8.1 Hz, 1H), 6.42 (d, *J* = 9.5 Hz, 1H), 3.88 (m, 4H), 3.76–3.73 (m, 2H), 3.59 (q, *J* = 5.2 Hz, 2H). ¹³C NMR (600 MHz, CDCl₃) δ 180.1, 155.9, 154.0, 150.7, 136.9, 134.1, 130.4, 129.8, 129.5, 127.8, 127.5, 126.3, 125.5, 120.9, 114.3, 104.8, 103.4, 72.9, 68.6, 62.1, 42.8. (+) ESI QqToF (m/z): Calcd. for C₂₁H₁₉N₂O₆⁺: [M + H]⁺ 395.1238, found 395.1248.

Data for 1-((3-hydroxypropyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**36**). Yield: 84%; M.p. > 220 °C (EtOAc); ¹H NMR (400 MHz, CDCl₃) δ 10.78 (s, 1H), 8.77 (s, 1H), 8.33 (d, *J* = 9.6 Hz, 1H), 8.02 (d, *J* = 9.9 Hz, 2H), 7.93 (d, *J* = 8.4 Hz, 1H), 7.63 (t, *J* = 8.2 Hz, 1H), 7.52 (t, *J* = 8.2 Hz, 1H), 6.50 (d, *J* = 9.7 Hz, 1H), 3.91 (t, *J* = 5.8 Hz, 2H), 3.57 (q, *J* = 6.2 Hz, 2H), 2.07 (m, 2H). ¹³C NMR (400 MHz, CDCl₃) δ 180.0, 156.1, 154.0, 150.7, 136.8, 134.0, 130.3, 129.8, 129.5, 127.6, 127.5, 126.3, 125.3, 120.9, 114.3, 104.6, 103.4, 60.3, 40.3, 31.5. (+) ESI QqToF (m/z): Calcd. for C₂₀H₁₆N₂O₅Na⁺: [M + Na]⁺ 387.0951, found 387.0962.

Data for 1-((2-hydroxyethyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**37**). Yield: 89%; M.p. > 220 °C (EtOH-H₂O); ¹H NMR (400 MHz, CDCl₃, MeOD) δ 8.75 (s, 1H), 8.30 (d, *J* = 8.4 Hz, 1H), 8.01 (d, *J* = 8.4 Hz, 1H), 7.97 (s, 1H), 7.91 (d, *J* = 8.4 Hz, 1H), 7.61 (t, *J* = 8.4 Hz, 1H), 7.50 (t, *J* = 8.3 Hz, 1H), 6.53 (d, *J* = 9.7 Hz, 1H), 3.89 (t, *J* = 5.6 Hz, 2H), 3.54 (t, *J* = 5.6 Hz, 2H). ¹³C NMR (400 MHz, CDCl₃, MeOD) δ 180.4, 156.7, 154.5, 151.0, 137.3, 134.4, 130.8, 130.1, 129.9, 128.0, 127.9, 126.7, 125.5, 121.3, 114.6, 105.0, 104.2, 60.4, 45.7. (-) ESI QqToF (m/z): Calcd. for C₁₉H₁₃N₂O₅⁻: [M - H]⁻ 349.0830, found 349.0825.

Data for 1-((3-methoxypropyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**38**). Yield: 94%; M.p. 174–176 °C (EtOAc); ¹H NMR (400 MHz, CDCl₃) δ 10.76 (s, 1H), 8.78 (s, 1H), 8.32 (d, *J* = 9.7 Hz, 1H), 8.03 (d, *J* = 8.4 Hz, 1H), 8.00 (s, 1H), 7.93 (d, *J* = 8.4 Hz, 1H), 7.63 (t, *J* = 8.3 Hz, 1H), 7.52 (t, *J* = 8.1 Hz, 1H), 6.48 (d, *J* = 9.7 Hz, 1H), 3.57 (t, *J* = 5.7 Hz, 2H), 3.55–3.49 (m, 2H), 3.41 (s, 3H), 2.05 (m, 2H). ¹³C NMR (400 MHz, CDCl₃) δ 179.9, 156.1, 154.0, 150.7, 136.8, 134.0, 130.3, 129.8, 129.4, 127.6, 127.5, 126.2, 125.3, 121.0, 114.2, 104.6, 103.4, 69.9, 59.0, 40.6, 29.2. (+) ESI QqToF (m/z): Calcd. for C₂₁H₁₈N₂O₅Na⁺: [M + Na]⁺ 401.1108, found 401.1115.

General procedure for the synthesis of amino substituted benzo[*b*]xanthenes **41–43** and **44–45**. Methanesulfonyl chloride (82 μL, 1.05 mmol) was added dropwise at 0 °C to a suspension of **36** (1 mmol) or **37** (1 mmol) and Et₃N (279 μL, 2 mmol) in dry CH₂Cl₂ (10 mL) and the resulting mixture was stirred at room temperature for 6 h. After completion of the reaction the mixture was washed with 1N HCl (2 × 5 mL) and water (2 × 5 mL) and the organic layer was dried over Na₂SO₄ and vacuum evaporated. Without further purification, the obtained crude mesylate (**39** and **40** respectively), was dissolved in absolute ethanol (10 mL) and to this solution a suitable amine (× 10 equiv.) was added. The resulting solution was stirred under reflux for 12 h. The solvent was then vacuum evaporated and the oily residue was purified by flash chromatography on silica gel, using a mixture of CH₂Cl₂/MeOH 9/1 as the elouent to provide the title compounds **41–43** and **44–45**.

Data for 1-((3-morpholinopropyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**41**). Yield: 77%; M.p. 179–180 °C (EtOAc-*n*-Hexane); ¹H NMR (400 MHz, CDCl₃) δ 10.73 (t, D₂O exchange, *J* = 5.5 Hz, 1H), 8.73 (s, 1H), 8.30 (d, *J* = 9.6 Hz, 1H), 8.00 (d, *J* = 8.4 Hz, 1H), 7.97 (s, 1H), 7.91 (d, *J* = 8.4 Hz, 1H), 7.62 (t, *J* = 8.2 Hz, 1H), 7.50 (t, *J* = 8.2 Hz, 1H), 6.47 (d, *J* = 9.7 Hz, 1H), 3.77 (m, 4H), 3.47 (q, *J* = 6.4 Hz, 2H), 2.59–2.45 (m, 6H), 1.96 (p, *J* = 6.4 Hz, 2H). ¹³C NMR (400 MHz, CDCl₃) δ 179.7, 155.9, 153.8, 150.5, 136.6, 133.8, 130.2, 129.6, 129.3, 127.4 (2 C), 126.1, 125.1, 120.8, 114.1, 104.4, 103.3, 67.0, 56.0, 53.8, 41.3, 25.7. (+) ESI QqToF (m/z): Calcd. for C₂₄H₂₄N₃O₅⁺: [M + H]⁺ 434.1710, found 434.1717.

Data for 1-((3-(4-methylpiperazin-1-yl)propyl)amino)-4-nitro-12*H*-benzo[*b*]xanthen-12-one (**42**). Yield: 81%; Oil; ¹H NMR (400 MHz, CDCl₃) δ 10.74 (t, D₂O exchange, *J* = 5.4 Hz, 1H), 8.73 (s, 1H), 8.30 (d, *J* = 9.6 Hz, 1H), 8.06–8.00 (d, *J* = 9.6 Hz, 1H), 7.96 (s, 1H), 7.94–7.89 (d, *J* = 9.6 Hz, 1H), 7.63 (t, *J* = 8.2 Hz, 1H), 7.51 (t, *J* = 8.2 Hz, 1H), 6.46 (d, *J* = 9.6 Hz, 1H), 3.47 (q, *J* = 6.3 Hz, 2H), 2.81 (m, 4H), 2.67 (m, 4H), 2.61 (t, *J* = 6.3 Hz, 2H), 2.54 (s, 3H), 1.97 (p, *J* = 6.3 Hz, 2H). ¹³C NMR (400 MHz, CDCl₃) δ 179.5, 155.6, 153.6, 150.3, 136.5, 133.6,

130.0, 129.5, 129.2, 127.2, 127.1, 126.0, 125.0, 120.5, 114.0, 104.2, 103.1, 55.3, 54.1, 51.5, 44.7, 41.4, 25.5. (+) ESI QqToF (m/z): Calcd. for $C_{25}H_{27}N_4O_4$: $[M + H]^+$ 447.2027, found 447.2024.

Data for 4-nitro-1-((3-(pyrrolidin-1-yl)propyl)amino)-12H-benzo[*b*]xanthen-12-one (**43**). Yield: 79%; M.p. 175–176 °C (EtOAc–*n*-Hexane); 1H NMR (600 MHz, $CDCl_3$) δ 10.73 (t, D_2O exchange, $J=5.3$ Hz, 1H), 8.77 (s, 1H), 8.32 (d, $J=9.6$ Hz, 1H), 8.02 (d, $J=8.3$ Hz, 1H), 7.99 (s, 1H), 7.93 (d, $J=8.4$ Hz, 1H), 7.63 (t, $J=8.2$ Hz, 1H), 7.52 (t, $J=8.2$ Hz, 1H), 6.50 (d, $J=9.6$ Hz, 1H), 3.53 (q, $J=6.9$, 2H), 2.80 (t, $J=6.9$ Hz, 2H), 2.76 (brs, 4H), 2.10 (p, $J=7.0$ Hz, 2H), 1.91 (brs, 4H). ^{13}C NMR (600 MHz, $CDCl_3$) δ 179.8, 155.8, 153.7, 150.5, 136.7, 133.8, 130.2, 129.6, 129.3, 127.4, 127.3, 126.0, 125.2, 120.8, 114.1, 104.4, 103.3, 54.0, 53.4, 41.3, 27.4, 23.5. (+) ESI QqToF (m/z): Calcd. for $C_{24}H_{24}N_3O_4$: $[M + H]^+$ 418.1761, found 418.1764.

Data for 1-((2-morpholinoethyl)amino)-4-nitro-12H-benzo[*b*]xanthen-12-one (**44**). Yield: 82%; M.p. > 220 °C (dec) (EtOAc–*n*-Hexane); 1H NMR (400 MHz, $CDCl_3$) δ 10.87 (s, D_2O exchange, 1H), 8.84 (s, 1H), 8.35 (d, $J=9.6$ Hz, 1H), 8.06 (d, $J=8.4$ Hz, 1H), 8.02 (s, 1H), 7.94 (d, $J=8.4$ Hz, 1H), 7.68–7.61 (m, 1H), 7.58–7.48 (m, 1H), 6.45 (d, $J=9.6$ Hz, 1H), 3.84 (m, 4H), 3.49 (brs, 2H), 2.80 (brs, 2H), 2.61 (m, 4H). ^{13}C NMR (400 MHz, $CDCl_3$) δ 179.8, 155.7, 154.0, 150.7, 136.8, 134.0, 130.4, 129.8, 129.4, 127.7, 127.5, 126.2, 125.5, 121.1, 114.3, 104.8, 103.6, 67.2, 56.4, 53.6, 40.2. (+) ESI QqToF (m/z): Calcd. for $C_{23}H_{22}N_3O_5$: $[M + H]^+$ 420.1554, found 420.1560.

Data for 1-((2-(4-methylpiperazin-1-yl)ethyl)amino)-4-nitro-12H-benzo[*b*]xanthen-12-one (**45**). Yield: 68%; M.p. > 220 °C (dec) (EtOAc–*n*-Hexane); 1H NMR (600 MHz, $CDCl_3$) δ 10.83 (t, D_2O exchange, $J=4.8$ Hz, 1H), 8.81 (s, 1H), 8.34 (d, $J=9.5$ Hz, 1H), 8.05 (d, $J=8.3$ Hz, 1H), 8.01 (s, 1H), 7.94 (d, $J=8.4$ Hz, 1H), 7.64 (t, $J=8.2$ Hz, 1H), 7.53 (t, $J=8.2$ Hz, 1H), 6.43 (d, $J=9.6$ Hz, 1H), 3.48 (q, $J=5.8$ Hz, 2H), 3.02–2.74 (m, 10H), 2.57 (s, 3H). ^{13}C NMR (400 MHz, $CDCl_3$) δ 180.0, 155.9, 154.2, 151.0, 137.1, 134.2, 130.6, 130.0, 129.7, 127.8, 127.8, 126.5, 125.8, 121.3, 114.5, 105.1, 103.8, 55.7, 54.9, 51.6, 45.3, 40.6. (+) ESI QqToF (m/z): Calcd. for $C_{24}H_{25}N_4O_4$: $[M + H]^+$ 433.1870, found 433.1870.

Microorganisms strains and growth conditions. The following reference and clinical fungal strains were used: *C. albicans* ATCC 10231, *C. glabrata* ATCC 90030, *C. krusei* ATCC 6258, *C. parapsilosis* ATCC 22019, *S. cerevisiae* ATCC 9763, *C. albicans* B3, *C. albicans* B4, *C. albicans* Gu4, *C. albicans* Gu5, *C. albicans* F2, *C. albicans* F5^{27,28}. Fungal strains used in this investigation were routinely grown over 18 h at 30 °C in YPG liquid medium (1% yeast extract, 1% peptone, 2% glucose) in a shaking incubator. For growth on solid media, 1.5% agar was added to the YPG medium.

Antifungal activity assays. Antifungal in vitro activity was determined by the modified M27-A3 specified by the CLSI²⁹ by minimal inhibitory concentrations (MICs) determination as described previously¹⁴. The MIC was defined as the lowest drug concentration at which at least a 90% decrease in turbidity, in comparison to the drug-free control, was observed. Antifungal activity was determined in RPMI-1640 medium buffered to pH 7.0. The final concentration of the compound solvent (DMSO) did not exceed 2.5% volume of final suspension in each well and did not influence the growth of microorganism.

Minimum fungicidal concentrations (MFCs) were determined as described previously¹³ by spot assay. The MFC was determined as the lowest concentration of the test compound in which no recovery of microorganisms was observed.

Yeast topoisomerase II relaxation assay and inhibition. The inhibition of yeast topoisomerase II was analysed according to the relaxation assay kit from Inspiralis (Inspiralis Limited, Norwich, UK) and as described previously¹⁴. The relaxation inhibition effectivity (IC_{50}) of the analyzed compounds was determined by densitometry quantification of the transition from supercoiled to relaxed forms and was expressed in relation to the control. The gels were photographed with Gel Doc XR+ Gel Documentation System (Bio-Rad: Hercules, CA, 647 USA) and image processing was performed by GIMP 2.10.18.

Antiproliferative activity determination. The human embryonic kidney cell line (HEK-293) and human liver cancer cell line (HEPG2) were purchased from ATCC (Manassas, Virginia, USA). HEK-293 was cultured in Dulbecco's Modified Eagle Medium and HEPG2 in Minimum Essential Medium Eagle. For each cell line, the culture medium was supplemented with 10% fetal bovine serum, 2 mM L-glutamine, and antibiotics: penicillin 62.6 $\mu g mL^{-1}$ and streptomycin 40 $\mu g mL^{-1}$. The cells were cultured at the humidified atmosphere of 5% $CO_2/95\%$ air and routinely tested for *Mycoplasma* contamination. Antiproliferative activity of the compounds was determined as previously described by MTT method¹⁴. Percentage of cell viability was calculated, and EC_{50} values were determined using GraphPad Prism® software (version 8.3.1).

Conclusions

28 xanthone and benzoxanthone analogues were synthesized and evaluated for their antifungal activity. Benzo-xanthenes proved to be the best antifungal agents with positive mycostatic selectivity index values in relation to HEK293 and HEPG2 cell lines. The fused benzene ring is thus crucial for their activity. The evaluation of biological properties suggests that the mode of action of the most effective derivatives is fungicidal. The demonstrated killing activity is associated with a higher probability of early therapeutic success and a decreased probability of resistance development. Moreover, the clinical strains of *C. albicans* resistant to fluconazole, due

to the FLU-induced overexpression of Cdr1p/Cdr2p drug efflux pumps as well as Mdr1p membrane transport protein of the major facilitator superfamily (MFS), remained sensitive to our novel compounds. This finding suggests that the most active against clinical resistant strains compounds **9**, **42** and **44** are not good substrates for ABC as well as MFS efflux pumps.

Since xanthone derivatives are human topoisomerase inhibitors, the antifungal activity of analyzed compounds may be related to their inhibitory effect on the fungal equivalent of that enzyme. Our results indicate a strong relationship between the antifungal activity and the inhibitory effectiveness against yeast topoisomerase II.

Summing up, we were able to show a proof of concept that xanthone modification may result in the discovery of a new group of selective antifungal drugs affecting fungal topoisomerase II. Moreover, our results indicate the possibility of using those derivatives against resistant fungal cells. Further validation of xanthenes applicability as antifungals by designing, synthesizing and evaluating the activity of new inhibitors is highly valuable. Future investigations will be focused on improving selectivity.

Data availability

All data generated or analysed during this study are included in this published article.

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Author contributions

K.R. conducted the experiments of antifungal activity determination and topoisomerase II inhibition studies, analyzed data, wrote manuscript; R.I. conducted the synthesis of the compounds, wrote manuscript; P.M. analyzed data; N.P. analyzed data; M.O. conducted the experiments of cytotoxic activity, analyzed data; I.K.K. and I.G. conceived the experiments, analyzed data, wrote manuscript. All authors reviewed the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

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