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# Eco-friendly synthesis of new olanzapine derivatives and evaluation of their anticancer potential†

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New derivatives of the known antipsychotic drug olanzapine have been obtained as potential compounds with anticancer activity in two metabolically different breast cancer cell lines: MCF-7 and triple negative MDA-MB-231. The compounds were obtained under phase transfer catalysis (PTC) in the presence of microwave irradiation (MW) or ultrasound ("")), evaluating the effect of solvents such as dimethylformamide, water, or choline chloride/urea (natural deep eutectic solvent, NaDES). In the best option, the compounds were obtained within 2 minutes with a yield of 57–86% in MW. Two of the obtained compounds which have a naphthalimide moiety and a pentyl (7) or hexyl chain (8) show pronounced cytotoxicity. Interestingly, neither olanzapine nor desmethylolanzapine (DOLA), which was one of the substrates for the synthesis reaction, showed any significant activity in the study.

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## Introduction

Olanzapine is an atypical antipsychotic that is approved by the FDA for use in the treatment of schizophrenia and bipolar disorder, including mixed or manic episodes. In addition, olanzapine may be used with fluoxetine, a selective serotonin reuptake inhibitor (SSRI), in patients who experience depressive episodes associated with bipolar I disorder and in treatment-resistant depression.<sup>1</sup> Interestingly, olanzapine can be also used in combination with samidorphan to alleviate olanzapine-induced weight gain in schizophrenia and bipolar I disorder.<sup>2,3</sup> Moreover, it can be used in cancer patients as an anti-emetic, but it can also have a beneficial effect on enhancing the effect of chemotherapeutic agents,<sup>4,5</sup> as well as in the treatment of insomnia<sup>6–9</sup> and as an analgesic agent.<sup>10–14</sup> Additionally, it has also been tested in patients with Anorexia Nervosa.<sup>15–17</sup> These

numerous advantages of olanzapine prompted scientists to search for olanzapine derivatives, which are tested, for example, as selective ligands of 5HT<sub>2A</sub>/D<sub>2</sub> receptors for more effective treatment of schizophrenia<sup>18,19</sup> as PDE4B inhibitors. This means the derivatives can have procognitive, neuroprotective, and anti-inflammatory effects.<sup>20,21</sup>

In recent years, antipsychotic drugs have often been assessed with regard to their anticancer potential.<sup>22–32</sup> It should be noted that olanzapine, as an antipsychotic drug, has an antagonistic effect on dopamine D<sub>2</sub> receptors and serotonin 5-HT<sub>2A</sub> receptors. In many studies, it has been observed that D<sub>2</sub> receptor antagonists exhibit anticancer properties both *in vitro* and in animal models to reduce tumor growth, cause apoptosis and induce autophagy.<sup>22</sup> The D<sub>2</sub> receptor pathway is hypothesized to play a role in promoting cell death due to reduced survivin expression. Some known psychotropic drugs which include olanzapine were previously considered as potential chemotherapeutic agents for the treatment or adjuvant treatment of cancer.<sup>34</sup> In recent years, many studies have focused on discovering the link between serotonin and breast cancer. Undoubtedly, there is a connection here, because serotonin antagonists are known to prevent the development of breast cancer, but the full mechanism is still not understood<sup>33–42</sup>

It is known that such an activity profile, *i.e.* an antagonist of serotonin and dopamine receptors, is very beneficial in anti-cancer chemotherapeutics. This justifies undertaking research in the context of the use of olanzapine itself as well as its structural analogues for anti-cancer research, especially with regard to breast, lung, and ovarian cancers.

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Considering the already known evidence for the anticancer activity of olanzapine<sup>43</sup> and the recent results of studies in the context of the assessment of activity to MCF-7 human breast cancer cells,<sup>44</sup> we decided to synthesize four new olanzapine derivatives and evaluate their activity to both MCF-7 and triple negative MDA-MB-231.

The methods of obtaining olanzapine by *N*-methylation of desmethylolanzapine (2-methyl-4-piperazin-1-yl-10*H*-thieno [2,3-*b*] [1,5]benzodiazepine, DOLA) known so far in the literature are based on syntheses in the presence of sodium hydroxide or hydride, potassium carbonate, triethylamine. This is because an alkaline environment that favors the reaction needs to be created. The most common solvents used are methanol, acetone, dichloromethane, tetrahydrofuran, tert-butanol.<sup>45,46</sup> The reaction time is 2–4 hours.

Thanks to previous experience where we successfully obtained olanzapine and its derivatives in conditions in the presence of microwave irradiation in less than one minute or using ultrasound conditions, we decided to use this greener alternative.<sup>47</sup> Reactions were carried out under phase transfer catalysis conditions using dimethylformamide as solvent and more ecofriendly options like choline chloride/urea, which is a deeply eutectic solvent. Deep Eutectic Solvents (DES) and Natural Deep Eutectic Solvents (NADES) are eutectic mixtures comprised of one or more hydrogen bond donors and hydrogen bond acceptors (from natural origin in the case of NADES) and have shown great potential as solvents and catalysts for a variety of organic reactions.<sup>48–51</sup> The advantages of DES and NADES over conventional organic solvents include low vapor pressure, low production costs, usually high biodegradability and recyclability. Moreover, DES and NADES can be prepared by a 100% atom economy, simply by mixing the components in the proper ratio and heating for a few hours.

## Results

### Synthesis of olanzapine

We started our research by developing a method of olanzapine synthesis in the presence of microwave irradiation. In this case, instead of the previously used microwave oven, we used a CEM Discovery reactor. In the first reactions, we tested the participation of a phase-transfer catalyst (PTC), *i.e.* TBAB (tetrabutylammonium bromide), as before. We conducted the experiment under pressure (entries 1, 2, Table 1) and under reflux (entry 3, Table 1). The reaction progress was assessed with TLC. In the next step, we assessed the effects of the addition of DABCO (1,4-diazabicyclo [2.2.2] octane) (entry 4, Table 1), which is often recommended for *N*-alkylation reactions<sup>52,53</sup> and other PTCs (entries 5–9, Table 1), such as BTBAC (benzyltributylammonium chloride), TEAC (tetraethylammonium chloride), TMAB (tetramethylammonium borohydride), TEBA (benzyltriethylammonium chloride), CTAB (hexadecyltrimethylammonium bromide). The reactions proceeded by nucleophilic substitution, which is favored by an alkaline environment. That is why potassium carbonate or in one case DIPEA (*N,N*-diisopropylethylamine) was used in the process. DMF (dimethylformamide) was used as the solvent in the reactions, but a greener solvent, *i.e.* choline chloride/urea (NaDES), was also used for comparison (Scheme 1).

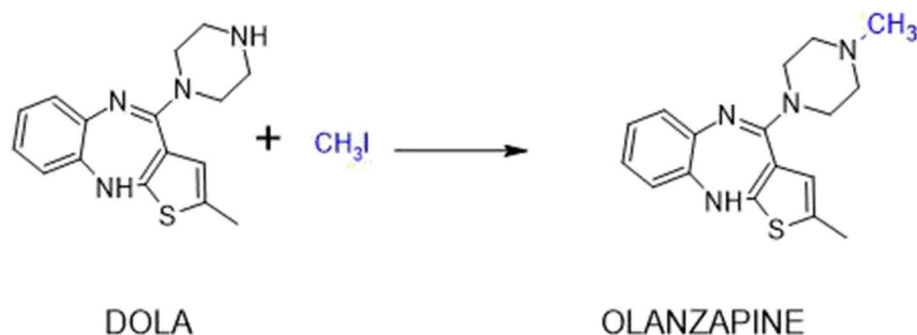
The experiments carried out proved that olanzapine could be effectively obtained using all tested catalysts (Table 1) and solvents. In the case of reactions carried out in the presence of microwave irradiation and using DMF, the highest yields were obtained for CTAB ( $Y = 95\%$ ) and TEBA ( $Y = 87\%$ ). In other cases, the efficiency was not much lower and amounted to 69–73%. When no catalyst was used (entry 10, Table 1) only 38% yield was observed after 60 seconds. In the case of using NaDES, within 2 minutes, a yield of 46% was obtained with the catalyst

**Table 1** Conditions for the synthesis of olanzapine derivatives in the reaction of DOLA (1 eq.) : CH<sub>3</sub>I (1.5 eq.) : K<sub>2</sub>CO<sub>3</sub> or DIPEA (3 eq.) : catalyst (0.01 eq.) in the presence of microwave irradiation (MW) and ultrasound ("")

No.	Conditions	Temp. <sup>a</sup> [°C]	Pressure <sup>b</sup> [bar]	Solvent/base/catalyst	Reaction time [min]	Yield [%]
1	MW	147	2.3	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	57
2	MW	150	4.7	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	1.5	69
3	MW	105	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	1.5	48
4	MW	150	3.2	DMF//K <sub>2</sub> CO <sub>3</sub> /DABCO	1	71
5	MW	150	1.9	DMF/K <sub>2</sub> CO <sub>3</sub> /BTBAC	1	69
6	MW	150	2.6	DMF/K <sub>2</sub> CO <sub>3</sub> /TEAC	1	73
7	MW	150	2.3	DMF/K <sub>2</sub> CO <sub>3</sub> /TMAB	1	72
8	MW	150	2.2	DMF/K <sub>2</sub> CO <sub>3</sub> /TEBA	1	87
9	MW	150	2.5	DMF/K <sub>2</sub> CO <sub>3</sub> /CTAB	1	95
10	MW	150	0.5	DMF/K <sub>2</sub> CO <sub>3</sub> /-	1	38
11	MW	150	0.5	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	46
12	MW	150	0.5	NaDES/K <sub>2</sub> CO <sub>3</sub> /-	2	37
13	MW	150	2.5	NaDES/Glycerine/K <sub>2</sub> CO <sub>3</sub> /TBAB	2.5	53
14	)))	45	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	40
15	)))	45	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /DABCO	120	45
16	)))	45	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /-	120	15
17	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	25
18	)))	45	Atm	NaDES/DIPEA/TBAB	120	61

<sup>a</sup> Highest observed data entered, the maximum temperature was set to 150 °C; reactions in the presence of ultrasound were carried out under reflux, due to the boiling point of CH<sub>3</sub>I (43 °C). <sup>b</sup> Highest observed data entered, the maximum pressure was set to 10 bar.





Scheme 1 Synthesis of olanzapine.

and 25% in the reaction without the catalyst. When, apart from NaDES, glycerin was also used as a solvent, the yield of olanzapine was 53%.

We decided to choose TBAB and DABCO to continue research on olanzapine synthesis in ultrasound conditions, because it has several advantages: it is eco-friendly, inexpensive, acts as a non-toxic base, and can be recycled and reused.<sup>54,55</sup> In the presence of ultrasonics in DMF, within 2 hours, the highest efficiency was obtained with DABCO, but in the case of TBAB it was also satisfactory (40%). As in the case of microwave irradiation, the lack of the catalyst caused a significant, *i.e.* more than two-fold decrease in the efficiency of obtaining olanzapine at the same time. In the case of NaDES, in the reaction in the presence of potassium carbonate, the yield was only 25%, but the exchange of the basic medium from solid to liquid increased the yield to 61%.

### Synthesis of olanzapine derivatives

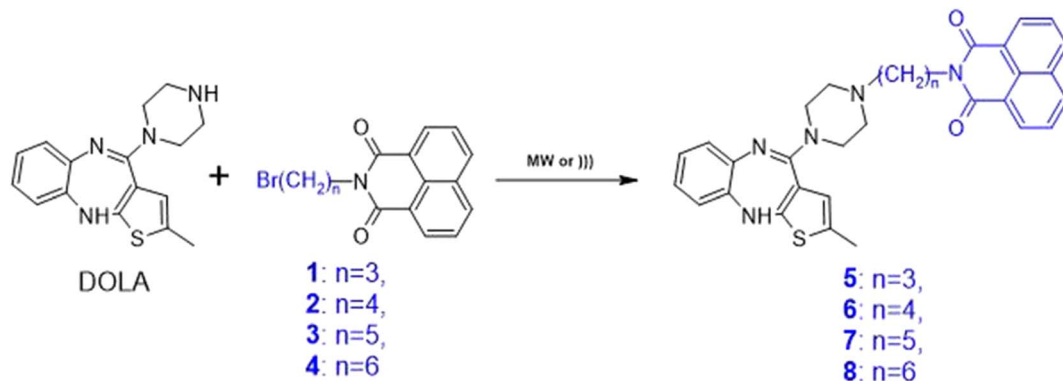
In the next stage, a study was carried out to obtain four new naphthalimidoalkyl derivatives of olanzapine (5–8), which differed only in the length of the alkyl chain (Scheme 2). Introduction of naphthalimidoalkyl moiety to DOLA core was aimed to augment the compound with structure bearing nicotinamide part of nicotinamide adenine dinucleotide (NAD<sup>+</sup>), which serves as an endogenous substrate of poly(ADP-ribose) polymerase (PARP) family of enzymes.<sup>56</sup>

Based on the experience in the synthesis of olanzapine, we decided to carry out the synthesis in the presence of microwave irradiation and ultrasound using similar reaction conditions (Table 2).

For each of the compounds, we performed syntheses in the presence of microwave irradiation using DMF and NaDES. In each case, higher yields (57–86%) were obtained with DMF, but despite the lower yields with NaDES (30–47%) or NaDES/Glycerine (5–53%), it is a very interesting synthesis option due to the non-toxic nature of the solvent. When NaDES was used in MW-assisted reactions, it was observed that the yield clearly decreases with the elongation of the linker, so the highest yields are observed for the compound with the propyl chain, and the lowest for the compound with the hexyl linker. In reactions in the presence of ultrasound, the highest yields for derivatives 5, 7 and 8 ( $Y = 27$ –51%) were obtained in the reaction with water as a solvent, in the presence of potassium carbonate and TBAB as a catalyst. In the case of derivative 6, product formation was observed only in the NaDES reaction ( $Y = 10\%$ ).

### Cytotoxicity

As a result of the synthesis reaction, four new olanzapine derivatives 5–8 were obtained, which were then purified by crystallization from methanol. Next, they were subjected to biological tests to assess their anticancer potential against breast cancer cells, *i.e.* triple negative MDA-MB-231 and



Scheme 2 Synthesis of olanzapine derivatives 5–8.



**Table 2** Conditions for the synthesis of olanzapine derivatives in the reaction of DOLA (1 eq.) : 1–4 (1 eq.) : K<sub>2</sub>CO<sub>3</sub> (3 eq.) : DABCO or TBAB (0.01 eq.) in MW or ultrasound

Entry	EE no.	Comp.	<i>n</i>	Conditions	Temp. <sup>a</sup> [°C]	Press. <sup>b</sup> [bar]	Solvent/base/catalyst	Reaction time [min]	Yield [%]
1		5	3	MW	150	2.8	DMF/K <sub>2</sub> CO <sub>3</sub> /DABCO	2	86
2		5	3	MW	150	0.3	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	30
3		5	3	MW	150	4.3	NaDES/Glycerine/K <sub>2</sub> CO <sub>3</sub> /TBAB	1.5	57
4		5	3	)))	45	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	60	16
5		5	3	)))	45	Atm	DMF/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	30
6		5	3	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /TBAB	60	0
7		5	3	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	6
8		5	3	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	60	0
9		5	3	)))	65	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	180	51
10		5	3	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	1
11		5	3	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	180	30
12		5	3	)))	45	Atm	NaDES/DIPEA/TBAB	120	10
13		6	4	MW	150	2.7	DMF/K <sub>2</sub> CO <sub>3</sub> /DABCO	2	65
14		6	4	MW	150	0.5	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	47
15		6	4	MW	150	1.5	NaDES/Glycerine/K <sub>2</sub> CO <sub>3</sub> /TBAB	3	40
16		6	4	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	120	1
17		6	4	)))	65	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	180	0
18		6	4	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	0
19		6	4	)))	45	Atm	NaDES/DIPEA/TBAB	120	10
20		7	5	MW	150	2.3	DMF/K <sub>2</sub> CO <sub>3</sub> /DABCO	2	57
21		7	5	MW	150	0.5	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	36
22		7	5	MW	150	4.2	NaDES/Glycerine/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	13
23		7	5	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	120	7
24		7	5	)))	65	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	180	27
25		7	5	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	2
26		7	5	)))	45	Atm	NaDES/DIPEA/TBAB	120	0
27		8	6	MW	150	2.6	DMF/K <sub>2</sub> CO <sub>3</sub> /DABCO	2	70
28		8	6	MW	150	0.8	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	2	31
29		8	6	MW	150	2.8	NaDES/Glycerine/K <sub>2</sub> CO <sub>3</sub> /TBAB	3	5
30		8	6	)))	45	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	120	0
31		8	6	)))	65	Atm	H <sub>2</sub> O/K <sub>2</sub> CO <sub>3</sub> /DABCO	180	31
32		8	6	)))	45	Atm	NaDES/K <sub>2</sub> CO <sub>3</sub> /TBAB	120	2
33		8	6	)))	45	Atm	NaDES/DIPEA/TBAB	120	0

<sup>a</sup> Highest observed data entered; the maximum temperature was set to 150 °C; reactions in the presence of ultrasound were carried out under reflux.

<sup>b</sup> Highest observed data entered; the maximum pressure was set to 10 bar.

whether they were hormone-dependent (both estrogen and progesterone receptor positive) MCF7. The derivatives showed dose-dependent inhibitory effect against the viability of the cultured breast cancer cells (Fig. 1). The compounds 7 and 8 showed more pronounced cytotoxicity characterized by generally lower half maximal inhibitory concentration (IC<sub>50</sub>) and higher maximal inhibition compared to other derivatives (Table 3). Interestingly, both parent DOLA and olanzapine itself elicited little to no effect on cell fitness under the conditions of our assay.

In general, MDA-MB-231 cell line was more susceptible to the generated naphthalimidoalkyl derivatives of olanzapine compared to MCF7. As the naphthalimidoalkyl moiety is supposed to affect the activity of PARP enzymes, we decided to investigate the expression of *PARP1* gene in cancer cell lines and to cross-compare the expression levels with the CRISPR-Cas9 viability screens using <https://depmap.org/portal> (Fig. 2). The gathered data indicate that MDA-MB-231 cells express more PARP1 and the MDA-MB-231 cell line is more susceptible to

*PARP1* knock-down compared to MCF7 cell line. These facts are in line with the cytotoxicity data.

### Molecular modeling

To further corroborate the effects of the obtained naphthalimidoalkyl derivatives, compounds 5–8 were evaluated in a molecular modeling procedure using the known crystal structures of three key PARP polymerases, namely TNKS2, TNKS1, and PARP1 (ref. 57–59). In the case of synthesized olanzapine derivatives, the presence of a hydrogen bond with Gly 1032 and a  $\pi$ - $\pi$  interaction with Tyr 1071 was observed in the complex with TNKS2. The 1,8-naphthalimide moiety was the most important fragment of the structure, directed deep into the binding pocket. Interactions of this type also appear in the reference complex (5*S*)-5-methyl-5-[4-(4-oxo-3,4-dihydroquinazolin-2-yl)phenyl]imidazolidine-2,4-dione with TNKS2, playing the key role in the activity. No additional stabilization was noted through the interactions in the tricyclic terminal region (Fig. 3A).





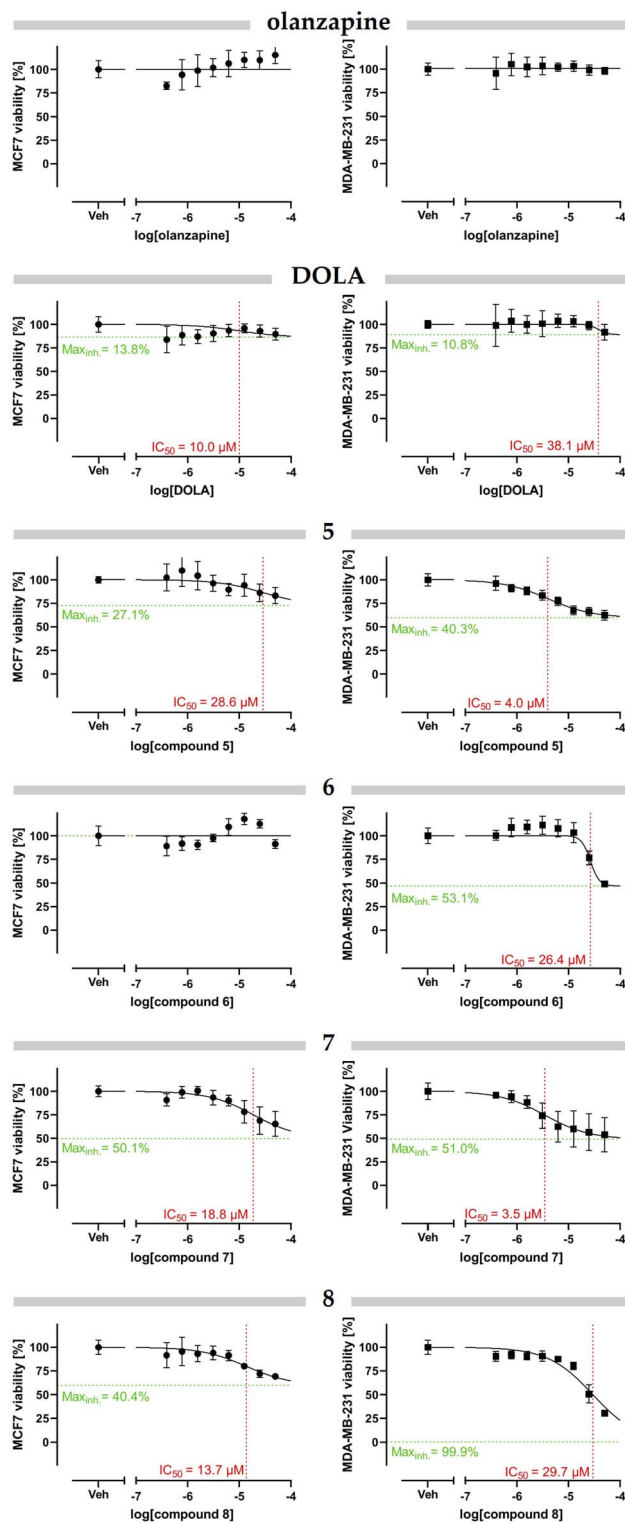


Fig. 1 Dose-response curves for olanzapine, DOLA, and its derivatives (compounds 5–8). The MCF7 (left column) and MDA-MB-231 (right column) cell lines were exposed to the increasing concentrations of our compounds of interest or vehicle (0.1% DMSO). The viability of the cells was measured using MTS assay (see Materials and methods for more details). The obtained values were normalized to vehicle control and three-parameters sigmoidal curve was fitted. Half-maximal inhibitory concentrations (IC<sub>50</sub> values, red dotted lines) and levels of maximal inhibition (Max<sub>inh</sub> values, green dotted lines) were calculated.

In conformations representing the binding mode in complexes with TNKS1, stabilizing  $\pi$ - $\pi$  interactions with Tyr 1213 and Tyr 1224 residues were observed. In the designed molecules, there was no key interaction with Gly 1185, which may suggest a lack of affinity towards the tested protein target (Fig. 3B).

In complexes with PARP1, a coherent binding mode was obtained as compared to the conformation of the reference compound. In the docked poses of the designed ligands, stabilizing hydrogen bonds with Gly 863 and  $\pi$ -stacking with Tyr907 were observed, which are key interactions for the stability of the complex. Importantly, the long-chain derivatives showed a slightly biased orientation as compared to olaparib in the central region of the binding pocket. However, in the docked poses, additional stabilizing hydrogen-type and  $\pi$ - $\pi$  interactions were observed in the terminal imide fragment (Fig. 3C).

## Discussion

In conclusion, we obtained new alkyl derivatives of olanzapine by reaction under PTC conditions. The compounds were obtained in the presence of microwave irradiation or ultrasound. The main advantage of the 5–8 synthesis method is the possibility to carry out the reaction under green chemistry conditions, *i.e.* through the use of ecological NaDES solvent and short reaction time. The obtained compounds were tested for inhibition in tests using two metabolically different breast cancer cell lines MCF-7 and triple negative MDA-MB-231. The obtained results, which were additionally justified in the procedure of molecular modeling for three key PARP polymerases (TNKS2, TNKS1, PARP1), prove that olanzapine naphthalimides with a pentyl (7) or hexyl (8) chain have a very interesting profile because they show a clear activity in both tested cell lines. The key limitation of the current study is the lack of biological screening to determine the inhibitory activity of the synthesized compounds against TNKS2, TNKS1, and PARP1. This will be addressed in a separate study and will be reported elsewhere.

## Methods

**Chemistry.** The reactions in the presence of microwave irradiation were carried out in a CEM Discover microwave reactor, while ultrasonic reactions were carried out in the PS-08 ultrasonic bath (80 W, 40 kHz). All reagents and solvents for the synthesis were purchased from Sigma-Aldrich, Merck, POCH, BIOSYNT CARBOSYNT and used as received. NaDES – choline chloride/urea – was obtained in accordance with the procedure described in 2020.<sup>47</sup>

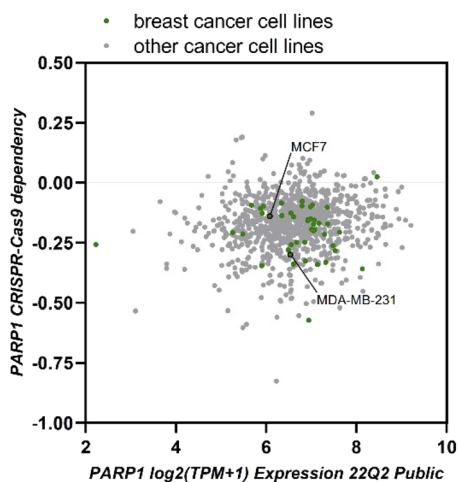
Sigma-Aldrich silica plates: 200  $\mu$ m, pore diameter: 60 Å, fluorescence index: 254 nm were used to evaluate the reaction progress with thin-layer chromatography (TLC) using 8 : 2 methanol : chloroform as eluent. Purification was carried out by column chromatography using silica gel 0.063–0.200 mm and methylene chloride : methanol 8 : 2 as eluent. UPLC-MS analyzes were performed with a Waters Acquity UPLC system – a Waters TQD (ESI-tandem quadrupole) mass spectrometer,



**Table 3** Toxicity of olanzapine derivatives towards MCF7 and MDA-MB-231 breast cancer cell lines. Half-maximal inhibitory concentrations ( $IC_{50}$ ) and maximal inhibition levels were calculated by fitting the normalized experimental values to three-parameter sigmoidal curve equation<sup>a</sup>

Entry	Compound	MCF7		MDA-MB-231	
		$IC_{50}$	Maximal inhibition	$IC_{50}$	Maximal inhibition
1	Olanzapine	n/d	n/d	n/d	n/d
2	DOLA	10.0 $\mu$ M	13.8%	38.1 $\mu$ M	10.8%
3	5	28.6 $\mu$ M	27.1%	4.0 $\mu$ M	40.3%
4	6	n/d	n/d	26.4 $\mu$ M	53.1%
5	7	18.8 $\mu$ M	50.1%	3.5 $\mu$ M	51.0%
6	8	13.7 $\mu$ M	40.4%	29.7 $\mu$ M	99.9%

<sup>a</sup> n/a, not determined due to lack of significant inhibition of cell viability.



**Fig. 2** Dependency of cancer cell lines on *PARP1* gene and the expression of *PARP1* across cell lines. Data were retrieved from the <https://depmap.org/portal>.

PDA detector, Acquity UPLC BEH C18 column, 1.7, 2.1  $\times$  100 mm (Waters Corporation, Milford, MA, USA), isocratically mobile phase: methanol : water + formic acid (4 : 6 + 0.1%, v/v). The  $^1H$  and  $^{13}C$  NMR spectra of products were recorded on a Bruker Avance 400 MHz spectrometer, using TMS as an internal reference. The detailed description of NMR spectra is provided in the ESI File.†

**Cytotoxicity assessment.** The cytotoxicity of the newly synthesized compounds was assessed using MTS assay as described before (ref. 60). The cell lines used in this study were human invasive breast ductal carcinoma MCF7 cell line (RRID:CVCL\_0031) and human breast adeno-carcinoma cell line MDA-MB-231 (RRID:CVCL\_0062). The drug exposure time was 72 h. DMSO (0.1%) was used as vehicle control.

**Datamining.** Data on gene expression and gene dependency were retrieved from DepMap Project portal available at <https://depmap.org/portal/>.

**Molecular modeling.** LigPrep 3.7 was used to prepare the three-dimensional structures. Epic. 2 was used to determine the respective ionization states at pH 7.4. The bond order and the respective ionization states of the amino acids were assigned using the protein preparation wizard. Grids were generated by centering a 35 Å grid field on the appropriate rest using the

OPLS\_2005 force field. Glide version 7.0 with flexible docking enabled was used for docking.

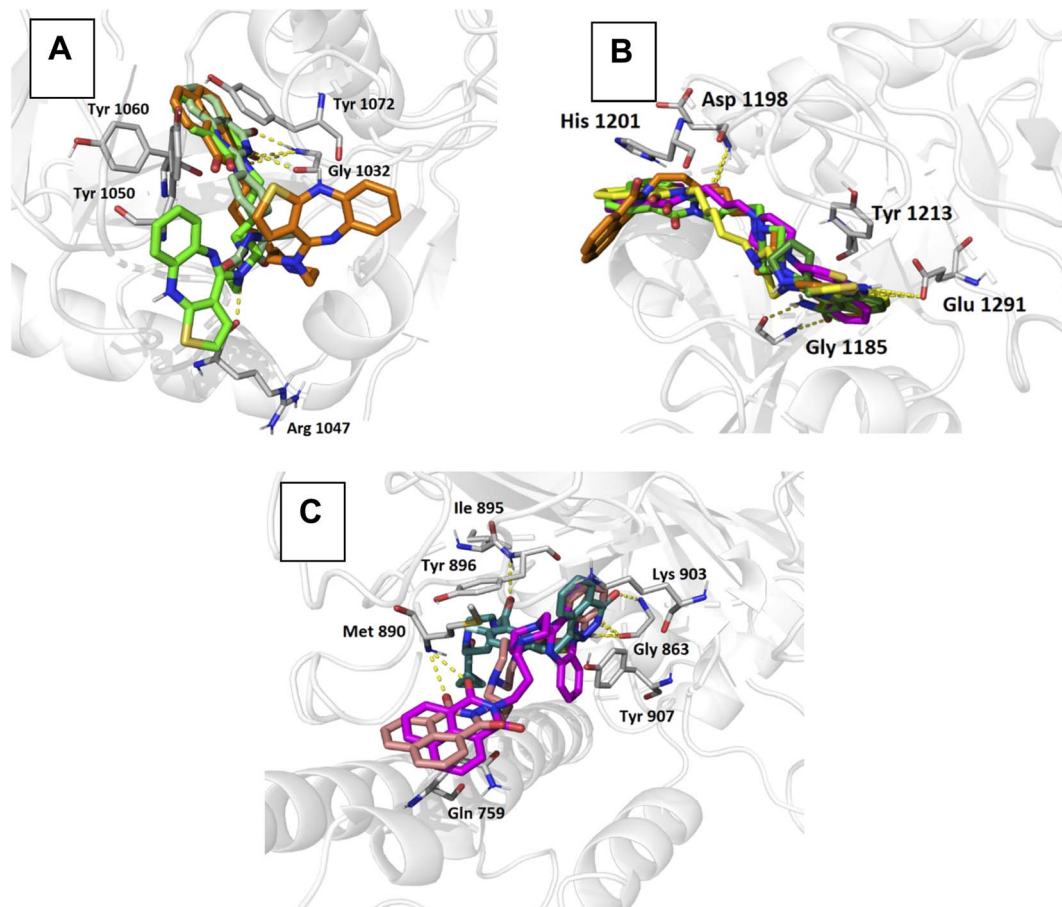
**General procedure for the synthesis of olanzapine in MW.** In a test tube equipped with magnetic stirrer bar, *N*-desmethyloanzapine (DOLA) (90 mg, 0.3 mmol, 1 equiv.), methyl iodide (28  $\mu$ L, 0.45 mmol, 1.5 equiv.), potassium carbonate (124 mg, 0.9 mmol, 3 equiv.), PTC catalyst (0.03 mmol, 0.1 equiv.), and solvent (2 mL) were mixed under microwave irradiation for 1–2 min. The reaction progress was monitored with TLC. After the reaction, 15 mL of water was added to the mixture and extracted with three portions of methylene chloride. The organic layer was evaporated to dryness, which was then crystallized from methanol or purified on a column chromatography.

**General procedure for the synthesis of olanzapine in ultrasound.** In a test tube *N*-desmethyloanzapine (DOLA) (90 mg, 0.3 mmol, 1 equiv.), methyl iodide (28  $\mu$ L, 0.45 mmol, 1.5 equiv.), potassium carbonate or DIPEA (0.9 mmol, 3 equiv.), PTC catalyst (0.03 mmol, 0.1 equiv.), and solvent (5 mL) placed in an ultrasonic bath and kept at 45–50 °C for 1–3 h. The reaction progress was monitored with TLC. After the reaction, 15 mL of water was added to the mixture and extracted with three portions of methylene chloride. The organic layer was evaporated to dryness, which was then crystallized from methanol or purified on a column chromatography.

**General procedure for the synthesis of 5–8 in MW.** In a test tube equipped with magnetic stirrer bar, *N*-desmethyloanzapine (DOLA) (90 mg, 0.3 mmol, 1 equiv.), naphthalimide alkyl derivative 1–4 (0.30 mmol, 1 equiv.), potassium carbonate or DIPEA (0.9 mmol, 3 equiv.), PTC catalyst (0.03 mmol, 0.1 equiv.), and solvent (2 mL) were mixed under microwave irradiation for 1–2 min. The reaction progress was monitored with TLC. After the reaction, 15 mL of water was added to the mixture and extracted with three portions of methylene chloride. The organic layer was evaporated to dryness, which was then crystallized from methanol or purified on a column chromatography.

**General procedure for the synthesis of 5–8 in ultrasound.** In a test tube *N*-desmethyloanzapine (DOLA) (90 mg, 0.3 mmol, 1 equiv.), naphthalimide alkyl derivative 1–4 (0.30 mmol, 1 equiv.), potassium carbonate or DIPEA (0.9 mmol, 3 equiv.), PTC catalyst (0.03 mmol, 0.1 equiv.) and solvent (5 mL) placed in an ultrasonic bath and kept at 45–50 °C for 1–3 h. The reaction progress was monitored with TLC. After the reaction,





**Fig. 3** (A) Binding modes of (5S)-5-methyl-5-[4-(4-oxo-3,4-dihydroquinazolin-2-yl)phenyl]imidazolidine-2,4-dione (pale green), 7 (green), 8 (orange) into TNKS2. The yellow dash-lines indicate the hydrogen bonds. (B) Binding modes of veliparib (dark green), 5 (magenta), 6 (green), 7 (yellow), 8 (orange) into TNKS1. The yellow dash-lines indicate the hydrogen bonds. (C) Binding modes of olaparib (cyan), 5 (magenta), 8 (orange) into PARP1. The yellow dash-lines indicate the hydrogen bonds.

15 mL of water was added to the mixture and extracted with three portions of methylene chloride. The organic layer was evaporated to dryness, which was then crystallized from methanol or purified on a column chromatography.

## Data availability

All data generated or analyzed during this study are included in this published article [and its ESI<sup>†</sup> File].

## Author contributions

A. K. D., W. T., W. B. running experiment, D. K. interpretation of NMR test results, P. Z. molecular modelling research, A. A., A. W., biological research – planning, experiments and interpretation, A. T. A. D. research concept using NaDES, manuscript revision, J. J. supervision, conceptualization, methodology, investigation, visualization, writing original draft.

## Conflicts of interest

The authors declare no competing interests.

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