



## Synthesis and biological activity evaluation of 3-(hetero) arylideneindolin-2-ones as potential c-Src inhibitors

Salvatore Princiotta, Loana Musso, Fabrizio Manetti, Valentina Marcellini, Giovanni Maga, Emmanuele Crespan, Cecilia Perini, Nadia Zaffaroni, Giovanni Luca Beretta & Sabrina Dallavalle

To cite this article: Salvatore Princiotta, Loana Musso, Fabrizio Manetti, Valentina Marcellini, Giovanni Maga, Emmanuele Crespan, Cecilia Perini, Nadia Zaffaroni, Giovanni Luca Beretta & Sabrina Dallavalle (2022) Synthesis and biological activity evaluation of 3-(hetero) arylideneindolin-2-ones as potential c-Src inhibitors, Journal of Enzyme Inhibition and Medicinal Chemistry, 37:1, 2382-2394, DOI: [10.1080/14756366.2022.2117317](https://doi.org/10.1080/14756366.2022.2117317)

To link to this article: <https://doi.org/10.1080/14756366.2022.2117317>



© 2022 The Author(s). Published by Informa UK Limited, trading as Taylor & Francis Group.



[View supplementary material](#)



Published online: 01 Sep 2022.



[Submit your article to this journal](#)



Article views: 38



[View related articles](#)




[View Crossmark data](#)

RESEARCH PAPER



## Synthesis and biological activity evaluation of 3-(hetero) arylideneindolin-2-ones as potential c-Src inhibitors

Salvatore Princiotta<sup>a</sup>, Loana Musso<sup>a</sup>, Fabrizio Manetti<sup>b</sup>, Valentina Marcellini<sup>b</sup>, Giovanni Maga<sup>c</sup>, Emmanuele Crespan<sup>c</sup>, Cecilia Perini<sup>c</sup>, Nadia Zaffaroni<sup>d</sup>, Giovanni Luca Beretta<sup>d</sup> and Sabrina Dallavalle<sup>a</sup> 

<sup>a</sup>Department of Food, Environmental and Nutritional Sciences (DeFENS), University of Milan, Milan, Italy; <sup>b</sup>Dipartimento di Biotecnologie, Chimica e Farmacia, Università di Siena, Siena, Italy; <sup>c</sup>Institute of Molecular Genetics IGM, CNR “Luigi Luca Cavalli-Sforza”, Pavia, Italy; <sup>d</sup>Molecular Pharmacology Unit, Department of Applied Research and Technological Development, Fondazione IRCCS Istituto Nazionale Tumori, Milan, Italy

### ABSTRACT

Inhibition of c-Src is considered one of the most studied approaches to cancer treatment, with several heterocyclic compounds approved during the last 15 years as chemotherapeutic agents. Starting from the biological evaluation of an *in-house* collection of small molecules, indolinone was selected as the most promising scaffold. In this work, several functionalised indolinones were synthesised and their inhibitory potency and cytotoxic activity were assayed. The pharmacological profile of the most active compounds, supported by molecular modelling studies, revealed that the presence of an amino group increased the affinity towards the ATP-binding site of c-Src. At the same time, bulkier derivatizations seemed to improve the interactions within the enzymatic pocket. Overall, these data represent an early stage towards the optimisation of new, easy-to-be functionalised indolinones as potential c-Src inhibitors.

### ARTICLE HISTORY

Received 15 June 2022  
Revised 9 August 2022  
Accepted 21 August 2022

### KEYWORDS

c-Src; indolinone;  
Knoevenagel reaction;  
molecular docking

## 1. Introduction



c-Src is the first discovered and most studied member of the Src kinase family<sup>1</sup>. It is a cytoplasmic non-receptor protein kinase<sup>2</sup>, highly involved in intracellular signal transmission for the regulation of proliferation, migration, adhesion, invasion, and drug resistance of cancer cells<sup>3,4</sup>. c-Src is a 60-kDa protein anchored to the plasmatic and endosomal membranes, as well as in the nuclear compartment, through its N-terminal tail, in which basic residues are particularly abundant<sup>5</sup>. The protein structure consists of four domains: SH1, the kinase region in which the autophosphorylation site is present; SH2, which hosts the phosphorylated Tyr530 (in the C-terminal tail) and binds to PDGFR; SH3, responsible for the closed and inactive conformation of the protein; SH4, which presents the myristoylation site, essential for the membrane localisation<sup>6</sup>. c-Src is constitutively inactive, with a close conformation in which SH2 and SH3 protect a phosphorylated Tyr530, preventing interactions within the catalytic site. Dephosphorylation of Tyr530 triggers the activation of c-Src, whose open conformation allows Tyr419 phosphorylation and full availability to ligand-protein interactions<sup>7,8</sup> (Figure 1).


Overexpression and/or high kinase activity of c-Src is associated with cancer progression and metastasis processes, mainly because of the reduced cell-cell adhesion<sup>9</sup> and their increased motility, migration, and invasiveness<sup>10</sup>. Moreover, interaction with other overexpressed receptor tyrosine kinases (such as EGFR<sup>11</sup>, PDGFR<sup>12</sup>, ERBB<sup>13</sup>, FGFR<sup>12</sup>, and HGF<sup>14</sup>) promotes the disruption of the closed conformation and exacerbates the activation of c-Src; such a

condition is widely observed in glioblastoma<sup>15</sup>, breast<sup>16</sup>, lung<sup>17</sup>, thyroid<sup>18</sup>, bones<sup>19,20</sup>, colon<sup>21</sup>, pancreas<sup>22</sup>, and prostate cancers<sup>23</sup>.

In this context, many efforts have been made to design and discover new molecules acting as c-Src inhibitors, blocking its downstream pathways and impairing tumour progression<sup>24–27</sup>. To date, five molecules targeting Src have been approved by FDA (Figure 2), and several others are in clinical trials for the treatment of solid tumours and leukemias, often in association with other cytotoxic agents. In particular, Dasatinib and Bosutinib have been approved for the treatment of chronic myeloid leukaemia (CML) in adult and paediatric patients<sup>28,29</sup>. Saracatinib, characterised by a more selective profile in terms of Src family inhibition<sup>30</sup>, is in clinical trials for the treatment of several solid tumours<sup>24</sup>. Vandetanib and Ponatinib are multi-tyrosine kinase inhibitors, approved by FDA to treat CML, thyroid carcinoma, and Philadelphia chromosome-positive acute lymphoblastic leukaemia in adults<sup>31,32</sup>. Other promising c-Src inhibitors (in phase I or II, (Figure 2) are DGY-06–116<sup>33</sup>, eCF506<sup>34</sup>, Elzovantinib (TPX-0022)<sup>35</sup>, and Tirbanibulin, the latter targeting the Src substrate binding site<sup>36</sup> and recently approved for the topical treatment of actinic keratosis<sup>37</sup>.

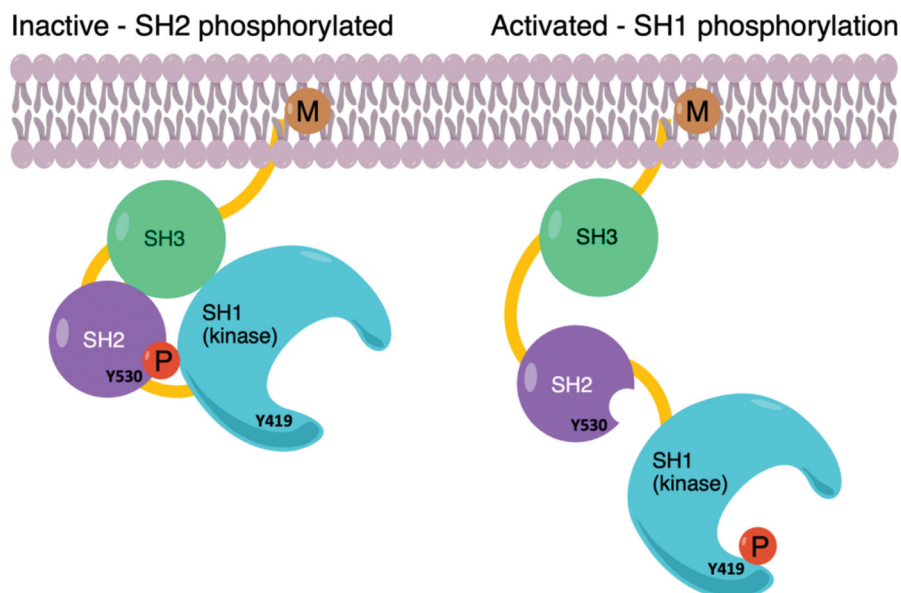
As part of an ongoing research program aimed at identifying new chemical entities for anticancer chemotherapy, we focussed our efforts on the search of c-Src inhibitors. For this purpose, we initially screened an *in-house* collection library of small molecules characterised by great structural diversity. Analogues of the most promising compounds, containing the indolinone scaffold, were synthesised to highlight the structural determinants for the

**CONTACT** Salvatore Princiotta  [salvatore.princiotta@unimi.it](mailto:salvatore.princiotta@unimi.it)  Department of Food, Environmental and Nutritional Sciences (DeFENS), University of Milan, Via Celoria 2, Milan, I-20133, Italy

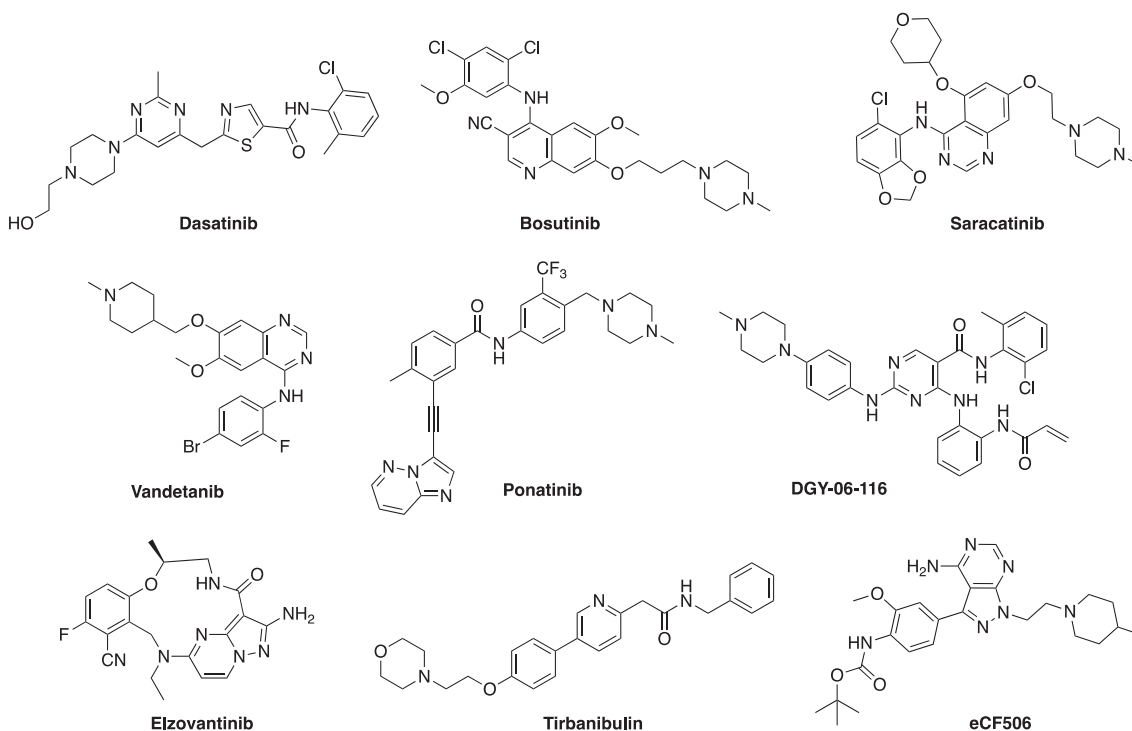
 Supplemental data for this article is available online at <https://doi.org/10.1080/14756366.2022.2117317>.

© 2022 The Author(s). Published by Informa UK Limited, trading as Taylor & Francis Group.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.



**Figure 1.** (a) Inactive conformation of c-Src shows phosphorylated Tyr530 on the SH2 domain; (b) Phosphorylation of Tyr419 on the kinase domain SH1 allows to activate of the enzyme.



**Figure 2.** Structures of the molecules approved by the FDA or in clinical trials as Src-family inhibitors.

activity. Molecular modelling studies were also performed to propose a binding mode within the c-Src binding site.

## 2. Experimental

### 2.1. Computational details

Docking studies were performed on the three-dimensional structure of the complex between c-Src and **AP23464** (entry 2BDJ of the protein data bank, 2.5 Å resolution). The structure of c-Src was prepared using the Protein Preparation Wizard of the Maestro Software Package (Schrödinger suite, version 19.2). The software

corrects bond orders and partial charges of the ligands and adds hydrogens to all atoms. Water molecules were removed, except those useful for stable binding interactions with the protein or ligand. Prime software was applied to define the correct conformation of missed side chains. Pre-processed protein was optimised with PROPKA and then minimised with an OPLS3 force field (convergence of heavy atoms set to a rmsd 0.3 Å).

The 2D structures of all synthesised compounds and native ligand **AP23464** were designed using the sketcher module of the Maestro suite and prepared with the Ligprep tool, which used the Epik module to find protonation and tautomerization states of compounds at pH of  $7.4 \pm 0.5$ . To optimise the three-dimensional

structures of the ligands, the Minimisation tool of the Macro model package was applied, using OPLS3 force field, while the other parameters were set to default values. Glide's receptor grid generation wizard was used to generate a three-dimensional grid with default settings. Then, flexible docking was performed with standard precision docking mode.

## 2.2. Biological assays

N-Terminal His6-tagged Src was purchased from Merck-Millipore (cat. 14–326). The reaction was performed according to the manufacturer's instructions with minor modifications: 500  $\mu$ M Src-peptide (KVEKIGEGTYGVVYK), 100  $\mu$ M ATP, 0.00087% NP-40, 10 ng of the enzyme, 10% DMSO in 10  $\mu$ L at 30 °C for 10 min. To avoid peptide adsorption to the plastic surface, protein low-binding tubes were used. ADP-Glo Kinase Assay (Promega) was then used to detect kinase activity accordingly with manufacturer's instruction with minor modifications. In details, reactions were transferred to white 384 well-plates and stopped by adding 10  $\mu$ L of ADP-Glo Reagent (Promega) for 50 min at rt. 20  $\mu$ L of Detection Reagent (Promega) was then added for 30 min and the reaction read using GloMax Discover microplate reader (Promega). Data were plotted using GraphPad Prism 5.0. ID<sub>50</sub> values were obtained according to Equation (1)

$$v = V / \{1 + (I/ID_{50})\} \quad (1)$$

where  $v$  is the measured reaction velocity,  $V$  is the apparent maximal velocity in the absence of an inhibitor,  $I$  is the inhibitor concentration, and ID<sub>50</sub> is the 50% inhibitory dose. Compounds tested were assumed to act as fully ATP-competitive inhibitors. Therefore,  $K_i$  values were calculated accordingly to Equation (2).

$$K_i = ID_{50} / (1 + K_m/[S]) \quad (2)$$

where  $K_i$  is the affinity of the inhibitor to the enzyme,  $[S]$  is the substrate (namely, ATP) concentration, and  $K_m$  is the affinity of ATP calculated accordingly to Michaelis-Menten equation.

Cytotoxicity experiments were carried out using the human MCF7 breast cancer cell line (ATCC HTB-22). Cells were cultured in RPMI1640 plus 10% foetal bovine serum at 37 °C and 5% CO<sub>2</sub>. Compound potency was determined by a growth inhibition assay (CellTiter 96<sup>®</sup> Aqueous One Solution Cell Proliferation Assay MTS, Promega). 24 h after seeding in a 96-well plate, cells were exposed to the compounds (concentration range 1–100  $\mu$ M) and 72 h later, 20  $\mu$ L of 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium salt (MTS) was added to each well. The absorbance was measured using a FLUOstar OPTIMA plate reader (BMG Labtech GmbH, Offenburg, Germany) at 492 nm after 4 h of incubation at 37 °C in 5% CO<sub>2</sub>. The IC<sub>50</sub> is the drug concentration causing 50% cell growth inhibition determined by the dose-response curves. Experiments were performed in triplicate.

## 2.3. Chemistry

All reagents and solvents were reagent grade or were purified by standard methods before use. Melting points were determined in open capillaries by an SMP3 apparatus and are uncorrected. <sup>1</sup>H and <sup>13</sup>C NMR spectra were recorded on Bruker AV600 spectrophotometer at 600 and 150 MHz, respectively. Chemical shifts are reported in ppm ( $\delta$ ) relative to TMS. The coupling constants,  $J$  are reported in Hertz (Hz). All compounds were routinely checked by thin layer chromatography (TLC) using precoated silica gel 60 F<sub>254</sub>, aluminium foil and the spots were detected under UV light at

254 nm and 365 nm or were revealed by spraying with 10% phosphomolybdic acid in ethanol.

Compounds **1**, **2**, **4**<sup>38</sup>, **13**, **14**<sup>39</sup>, **15**, **16**<sup>40</sup>, **7**, **8**, **23**, **24**<sup>41</sup>, **9**<sup>42</sup>, **10**, **11**<sup>43</sup>, **17**<sup>44</sup>, **18**<sup>45</sup>, **19**<sup>45</sup>, **25**<sup>46</sup>, **28**<sup>47</sup> are part of an in-house library. The synthetic procedures relative to the above-mentioned products have been already described and reported in the cited references.

## 2.4. General procedure for preparation of compounds **22**, **27**, **29**, **30**, **32–38**, **43–50**

To a solution of the benzaldehyde (0.9 mmol) in EtOH (9 ml), indolinone (100 mg, 0.6 mmol) was added, followed by piperidine (0.06 mmol). After heating at reflux for 4 h, the crude was concentrated under reduced pressure and purified by flash chromatography.

**1-hydroxy-3-(3-phenoxybenzylidene)indolin-2-one (22)** was prepared from 1-hydroxy-3H-indol-2-one<sup>41</sup> and 3-phenoxybenzaldehyde. Purification by column chromatography (CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 97:3) gave the title compound in 85% yield, as a yellow oil. <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 10.90 (1H, s); 7.67 (1H, s); 7.57–7.53–7.73 (1H, m); 7.48–7.41 (2H, m); 7.40–7.37 (1H, m); 7.33–7.7.28 (1H, m); 7.26 (1H, s); 7.21–7.15 (2H, m); 7.14–7.11 (2H, m); 6.95 (1H, d,  $J$  = 7.6 Hz); 6.88–6.82 (1H, m). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 162.9, 157.3, 156.1, 142.4, 136.0, 135.9, 130.6, 130.4, 130.2 ( $\times$  2 C), 130.1, 124.6, 124.1, 122.1, 121.8, 119.9, 119.4 ( $\times$  2 C), 118.6, 118.3, 107.5.

**3-((5-phenylthiophen-2-yl)methylene)indolin-2-one (27)** was prepared from commercially available 2-oxindole and 5-phenylthiophene-2-carbaldehyde. Purification by column chromatography (CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 95:5) gave the title compound in 95% yield, as an orange solid (m.p. 257–258 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 10.63 (1H, s); 8.06 (1H, s); 7.90 (1H, d,  $J$  = 3.8 Hz); 7.79–7.73 (2H, m); 7.68 (1H, d,  $J$  = 7.7 Hz); 7.64 (1H, d,  $J$  = 3.8 Hz); 7.50–7.43 (2H, m); 7.41–7.35 (1H, m); 7.24–7.16 (1H, m); 7.02–6.97 (1H, m); 6.86 (1H, d,  $J$  = 7.6 Hz). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 167.4, 150.2, 140.5, 139.0, 136.9, 133.5, 129.3 ( $\times$  2 C), 128.6, 128.5, 128.0, 125.8 ( $\times$  2 C), 124.4, 124.2, 121.6, 121.0, 119.4, 109.5.

**3-(3,5-dibromo-4-hydroxybenzylidene)-5,6-dimethoxyindolin-2-one (29)** was prepared from commercially available 5,6-dimethoxy-2-oxindole and 3,5-dibromo-4-hydroxybenzaldehyde. Purification by column chromatography (CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 95:5) gave the title compound in 52% yield, as a yellow solid (291–292 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>, *E/Z* 76/24,  $\delta$ : 10.52 (1H, bs); 10.36 (1H isomer *Z*, bs); 10.32 (1H isomer *E*, bs); 8.75–8.69 (2H isomer *Z*, m); 7.95–7.89 (2H isomer *E*, m); 7.52 (1H isomer *Z*, s); 7.34 (1H isomer *Z*, s); 7.28 (1H isomer *E*, s); 7.17 (1H isomer *E*, s); 6.51 (1H isomer *E*, s); 6.45 (1H isomer *Z*, s); 3.79 (3H isomer *E*, s); 3.77 (3H isomer *Z*, s); 3.76 (3H isomer *Z*, s); 3.64 (3H isomer *E*, s). <sup>13</sup>C-NMR (150 MHz, DMSO-*d*<sub>6</sub>, isomer *E*+*Z*,  $\delta$ : 169.1, 167.8, 151.2 ( $\times$  2 C), 144.2 ( $\times$  2 C), 143.2 ( $\times$  2 C), 139.3, 135.6 ( $\times$  2 C), 133.1 ( $\times$  4 C), 131.5, 129.6, 128.5, 116.0, 111.8 ( $\times$  2 C), 111.4, 111.3, 107.0, 105.1, 95.5 ( $\times$  2 C), 95.0 ( $\times$  2 C), 56.3 ( $\times$  2 C), 55.7 ( $\times$  2 C).

**3-((1,1'-biphenyl)-4-ylmethylene)-5,6-dimethoxyindolin-2-one (30)** was prepared from 5,6-dimethoxy-2-oxindole and (1,1'-biphenyl)-4-carbaldehyde. Purification by column chromatography (CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 95:5) gave the title compound an 83% yield, as a yellow solid (m.p. 231–232 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 10.36 (1H, bs); 7.87–7.78 (4H, m); 7.78–7.72 (2H, m); 7.54–7.45 (4H, m); 7.43–7.36 (1H, m); 7.23 (1H, s); 3.79 (3H, s); 3.58 (3H, s). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>, major isomer (*E*),  $\delta$ : 169.3, 151.4, 143.2, 140.9, 139.3, 138.3, 133.8, 131.8, 130.0 ( $\times$  2 C), 129.0

( $\times 2$ C), 127.9 ( $\times 2$ C), 126.8 ( $\times 2$ C), 126.7 ( $\times 2$ C), 111.7, 107.9, 95.5.

**4-((6-chloro-2-oxoindolin-3-ylidene)methyl)benzonitrile (32)** was prepared from **31** and 4-cyanobenzaldehyde. Purification by column chromatography (hexane/EtOAc 90:10  $\rightarrow$  20:80) gave the title compound in 45% yield, as a yellow solid (m.p. 310 °C, dec.).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), *E/Z* 73/27,  $\delta$ : 10.88 (1H isomer Z, s); 10.86 (1H isomer E, s); 8.48–8.42 (2H isomer Z, m); 8.04–7.98 (2H isomer E, m); 7.96–7.92 (3H isomer Z); 7.91–7.86 (2H isomer E, m); 7.80–7.75 (1H isomer Z, d,  $J=8.3$  Hz); 7.70 (1H isomer E, s); 7.37 (1H isomer E, d,  $J=8.9$  Hz); 7.10 (1H isomer Z, dd,  $J=8.3$  Hz, 1.8 Hz); 6.96–6.90 (2H isomer E, m); 6.87 (1H isomer Z, d,  $J=1.8$  Hz).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 168.2, 144.7, 139.2, 134.8, 134.3, 132.7 ( $\times 2$ C), 132.0, 130.0 ( $\times 2$ C), 128.6, 124.0, 121.1, 119.3, 118.5, 110.3.

**6-chloro-3-(4-nitrobenzylidene)indolin-2-one (33)** was prepared from **31** and 4-nitrobenzaldehyde. Purification by column chromatography ( $\text{CH}_2\text{Cl}_2 + 0.1\%$  Et $_3$ N,  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  99:1  $\rightarrow$  80:20, EtOAc/ $\text{CH}_3\text{OH}$  80:20) gave the title compound in 45% yield, as a red solid (m.p. 322–323 °C, dec.).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 10.89 (1H, s); 8.40–8.33 (2H, m); 8.00–7.92 (2H, m); 7.74 (1H, s); 7.40 (1H, d,  $J=8.2$  Hz); 6.95–6.90 (2H, m).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 168.1, 147.6, 144.8, 141.1, 135.0, 133.8, 130.5 ( $\times 2$ C), 129.0, 124.0 ( $\times 2$ C), 123.2, 121.2, 119.2, 110.3.

**6-chloro-3-(4-(dimethylamino)benzylidene)indolin-2-one (34)** was prepared from **31** and 4-(dimethylamino)benzaldehyde. Purification by column chromatography (petroleum ether/EtOAc 95:5  $\rightarrow$  20:80,  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  95:5) gave the title compound an 84% yield, as an orange solid (m.p. 253–254 °C).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 10.62 (bs, 1H); 7.78 (1H d,  $J=8.3$  Hz); 7.68–7.64 (2H, m); 7.57 (1H, s); 6.96 (1H, dd,  $J=8.3$  Hz, 1.6 Hz); 6.88 (1H, d,  $J=1.6$  Hz); 6.86–6.82 (2H, m); 3.02 (6H, s).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ),  $\delta$ : 169.8, 152.0, 143.8, 138.7, 139.9, 132.6 ( $\times 2$ C), 123.2, 121.3 ( $\times 2$ C), 121.2, 121.0, 112.1 ( $\times 2$ C), 110.1, 40.1 ( $\times 2$ C).

**6-chloro-3-(4-hydroxybenzylidene)indolin-2-one (35)** was prepared from **31** and 4-hydroxybenzaldehyde. Purification by column chromatography (hexane/EtOAc 90:10,  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  95:5  $\rightarrow$  90:10) gave the title compound in 94% yield, as a yellow solid (m.p. 330–331 °C, dec.).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 10.70 (1H, bs); 10.27 (1H, bs); 7.69 (1H, d,  $J=8.2$  Hz); 7.66–7.60 (2H, m); 7.59 (1H, s); 6.97–6.91 (3H, m); 6.90–6.99 (1H, m).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), isomer *E+Z*,  $\delta$ : 169.0, 167.3, 160.5, 159.6, 143.8, 141.1, 138.5, 137.5, 135.0, 133.3, 132.0 ( $\times 4$ C), 125.4, 124.7, 124.6, 123.4, 123.2, 121.6, 120.7, 120.5, 120.4, 120.3, 115.7 ( $\times 2$ C), 115.3 ( $\times 2$ C), 109.8, 109.0.

**4-((6-chloro-2-oxoindolin-3-ylidene)methyl)benzoic acid (36)** was prepared from **31** and 4-formylbenzoic acid. Purification by column chromatography (hexane/EtOAc 90:10  $\rightarrow$  75:25,  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  95:5  $\rightarrow$  80:20, EtOAc/ $\text{CH}_3\text{OH}$  80:20) gave the title compound in 22% yield, as a yellow solid (m.p. 271–272 °C, dec.).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 10.86 (1H, bs); 8.10–8.03 (2H, m); 7.81–7.73 (2H, m); 7.71 (1H, s); 7.48 (1H, d,  $J=7.48$  Hz); 6.96–6.89 (2H, m).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 168.4, 168.0, 144.5, 137.5, 135.6, 134.4, 131.6, 129.7 ( $\times 2$ C), 129.1 ( $\times 2$ C), 127.5, 123.8, 121.0, 119.6, 110.2.

**3-(4-(benzyloxy)benzylidene)-6-chloroindolin-2-one (37)** was prepared from **31** and 4-benzyloxybenzaldehyde. Purification by column chromatography (hexane/EtOAc 90:10  $\rightarrow$  75:25) gave the title compound in 50% yield, as a yellow solid (m.p. 239–240 °C, dec.).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), *E/Z* 75/25,  $\delta$ : 10.74 (1H isomer *E* + 1H isomer *Z*, s); 8.48 (2H isomer *Z*, m); 7.82 (1H isomer *Z*, s); 7.78–7.69 (2H isomer *E* + 1H isomer *Z*, m); 7.68–7.61

(2H, m); 7.54–7.48 (2H isomer *E*, m); 7.47–7.40 (2H isomer *E* + 2H isomer *Z*, m); 7.39–7.34 (2H isomer *Z*, m); 7.22–7.16 (2H isomer *E*, m); 7.16–7.11 (2H isomer *Z*, m); 7.07–7.02 (1H isomer *Z*, m); 6.95 (1H isomer *E*, dd,  $J=8.4$  Hz, 2.1 Hz); 6.90 (1H isomer *E*, d,  $J=2.1$  Hz); 6.86–6.83 (1H isomer *Z*, m); 5.21 (2H isomer *E* + 2H isomer *Z*, s).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), isomer *E+Z*,  $\delta$ : 168.8, 167.2, 160.51, 59.8, 144.0, 141.3, 137.9, 136.9, 136.6, 134.6, 133.6, 132.2, 131.7 ( $\times 2$ C), 128.5 ( $\times 6$ C), 128.0 ( $\times 2$ C), 127.8 ( $\times 4$ C), 127.0, 126.6, 124.5, 124.3, 123.3, 122.9, 120.8, 120.7 ( $\times 2$ C), 120.1, 115.1 ( $\times 2$ C), 114.6 ( $\times 2$ C), 110.0, 109.1.

**6-chloro-3-(3-phenoxybenzylidene)indolin-2-one (38)** was prepared from **31** and 3-phenoxybenzaldehyde. Purification by column chromatography (hexane/EtOAc 90:10 + 0.1% Et $_3$ N, hexane/EtOAc 80:20  $\rightarrow$  50:50,  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  99:1  $\rightarrow$  80:20) gave the title compound in 19% yield, as a yellow solid (m.p. 188–189 °C).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 10.78 (1H, bs); 7.63 (1H, s); 7.58–7.52 (1H, m); 7.49–7.41 (3H, m); 7.40–7.35 (1H, m); 7.25–7.19 (2H, m); 7.18–7.10 (3H, m); 6.91–6.82 (1H, m).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), major isomer (E),  $\delta$ : 168.4, 157.3, 156.0, 144.4, 136.0, 135.7, 134.2, 130.6, 130.2 ( $\times 2$ C), 126.9, 124.4, 124.1, 123.7, 120.8, 119.8, 119.5 ( $\times 2$ C), 119.4, 118.2, 110.1.

#### 2.4.1. Ethyl 2-(4-((6-chloro-2-oxoindolin-3-ylidene)methyl)phenoxy)acetate (43)

To a solution of 4-hydroxybenzaldehyde (100 mg, 0.82 mmol) and bromoethyl acetate (0.14 ml, 1.23 mmol) in dry acetone (3 ml),  $\text{K}_2\text{CO}_3$  (170 mg, 1.64 mmol) was added, and the reaction was left stirring at reflux. After 4 h, the crude was concentrated under reduced pressure, treated with EtOAc and washed with water. The resulting organic phase was washed with brine and dried over  $\text{Na}_2\text{SO}_4$ , then evaporated under reduced pressure. Purification by flash chromatography (petroleum ether/EtOAc 90:10  $\rightarrow$  85:15) afforded ethyl 2-(4-formylphenoxy)acetate **40** (150 mg, 88% yield) as a yellow oil.  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ),  $\delta$ : 9.93 (1H, s); 7.91–7.85 (2H, m); 7.07–7.02 (2H, m); 4.73 (2H, s); 4.31 (2H, q,  $J=6.6$  Hz); 1.33 (3H, t,  $J=6.6$  Hz).

Compound **43** was prepared from **31** and the above compound. Purification by column chromatography (hexane/EtOAc 90:10  $\rightarrow$  75:25) gave the title compound in 64% yield, as a yellow solid (m.p. 158–159 °C).  $^1\text{H}$  NMR (600 MHz, DMSO- $d_6$ ), *E/Z* 90/10,  $\delta$ : 10.74 (1H isomer *E*, bs); 8.49–8.44 (2H isomer *Z*, m); 7.82 (1H isomer *Z*, s); 7.75–7.68 (2H isomer *E*, m); 7.76–7.58 (2H isomer *E*, m); 7.13–7.07 (2H isomer *E*, m); 7.06–7.03 (1H isomer *Z*, m); 6.94 (1H isomer *E*, dd,  $J=1.9$  Hz, 8.4 Hz); 6.90 (1H isomer *E*, d,  $J=1.9$  Hz); 6.84 (1H isomer *Z*, d,  $J=1.8$  Hz); 4.90 (2H isomer *E* + 2H isomer *Z*, s); 4.21 (2H isomer *E* + 2H isomer *Z*, q,  $J=6.9$  Hz); 1.24 (3H isomer *E* + 3H isomer *Z*, t,  $J=6.9$  Hz).  $^{13}\text{C}$  NMR (150 MHz, DMSO- $d_6$ ), isomer *E+Z*,  $\delta$ : 168.8, 168.4 ( $\times 2$ C), 167.2, 159.6, 159.0, 144.0, ( $\times 2$ C), 141.4, 137.7, 136.7 ( $\times 2$ C), 134.4 ( $\times 2$ C), 133.7, 132.3, 131.5 ( $\times 2$ C), 127.4, 127.0, 124.8, 124.3, 123.3 ( $\times 2$ C), 120.8 ( $\times 3$ C), 120.0, 114.9 ( $\times 2$ C), 114.3 ( $\times 2$ C), 64.7, 64.6, 60.7 ( $\times 2$ C), 14.0 ( $\times 2$ C).

#### 2.4.2. Methyl 4-((4-((6-chloro-2-oxoindolin-3-ylidene)methyl)phenoxy)methyl)benzoate (44)

To a solution of 4-hydroxybenzaldehyde (100 mg, 0.82 mmol) and methyl 4-bromomethyl benzoate (205 mg, 0.9 mmol) in dry DMF (3 ml),  $\text{Cs}_2\text{CO}_3$  (200 mg, 0.57 mmol) was added, and the reaction was left stirring at rt. After 16 h, the crude was concentrated under reduced pressure, treated with EtOAc and washed with water. The resulting organic phase was washed with brine and dried over  $\text{Na}_2\text{SO}_4$ , then evaporated under reduced pressure.

Purification by flash chromatography (petroleum ether/EtOAc 90:10 → 75:25) afforded methyl 4-((4-formylphenoxy)methyl)benzoate **41** (175 mg, 79% yield) as a white solid. <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), δ: 9.93 (1H, s); 8.14–8.08 (2H, m); 7.91–7.84 (2H, m); 7.56–7.50 (2H, m); 7.14–7.07 (2H, m); 5.24 (2H, s); 3.95 (3H, s).

Compound **44** was prepared from **31** and the above compound. Purification by column chromatography (hexane/EtOAc 90:10 → 50:50) gave the title compound in 93% yield, as a yellow solid (m.p. 201–202 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), *E/Z* 70/30, δ: 10.73 (1H isomer *Z*, bs); 10.71 (1H isomer *E*, bs); 8.49–8.43 (2H, isomer *Z*, m); 8.05–7.96 (2H isomer *E*, m); 7.79 (1H isomer *Z*, s); 7.74–7.66 (2H isomer *E*+2H isomer *Z*, m); 7.65–7.58 (4H isomer *E*+3H isomer *Z*, m); 7.19–7.15 (2H isomer *E*, m); 7.14–7.11 (2H isomer *Z*, m); 7.02 (1H isomer *Z*, dd, *J* = 1.6 Hz, 8.2 Hz); 6.92 (1H isomer *E*, dd, *J* = 1.6 Hz, 8.3 Hz); 6.87 (1H isomer *E*, d, *J* = 1.6 Hz); 6.81 (1H isomer *Z*, d, *J* = 1.6 Hz); 5.29 (2H isomer *E*+2H isomer *Z*, s); 3.85 (3H isomer *E*+2H isomer *Z*, s). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), isomer *E*+*Z*, δ: 168.8, 167.2, 166.0 (× 2C), 160.2, 159.6, 144.0, 142.2 (× 2C), 141.4, 137.8, 136.7, 134.6, 133.6, 131.7, 131.6 (× 2C), 129.4 (× 6C), 129.1 (× 2C), 127.2 (× 4C), 127.2, 126.8, 124.6, 124.3, 123.3, 123.1, 120.8 (× 3C), 120.1, 115.2 (× 2C), 114.6 (× 2C), 110.0, 109.1, 68.8 (× 2C), 52.1 (× 2C).

#### 2.4.3. 6-Chloro-3-((4-(cyclopropylmethoxy)benzylidene)indolin-2-one (45)

To a solution of 4-hydroxybenzaldehyde (100 mg, 0.82 mmol) and bromomethyl cyclopropane (0.16 ml, 1.64 mmol) in dry acetone (3 ml), K<sub>2</sub>CO<sub>3</sub> (450 mg, 3.28 mmol) was added, and the reaction was left stirring at reflux. After 16 h, the crude was concentrated under reduced pressure, treated with EtOAc and washed with water. The resulting organic phase was washed with brine and dried over Na<sub>2</sub>SO<sub>4</sub>, then evaporated under reduced pressure. Purification by flash chromatography (petroleum ether/EtOAc 95:5 → 90:10) afforded compound **42** (110 mg, 76% yield) as a colourless oil. <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), δ: 9.90 (1H, s); 7.93–7.81 (2H, m); 7.06–6.95 (2H, m); 3.92 (2H, d, *J* = 6.9 Hz); 1.37–1.26 (1H, m); 0.77–0.62 (2H, m); 0.50–0.34 (2H, m).

Compound **45** was prepared from **31** and the above compound. Purification by column chromatography (hexane/EtOAc 90:10 → 70:30) gave the title compound in 47% yield, as a yellow solid (m.p. 231–232 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), isomer *E*, δ: 10.74 (1H, bs), 8.42–8.50 (2H, m); 7.81 (1H, s); 7.71 (1H, d, *J* = 8.1 Hz); 7.10–6.99 (3H, m), 6.81 (1H, s); 3.93 (2H, d, *J* = 6.7 Hz); 1.32–1.22 (1H, m); 0.67–0.53 (2H, m); 0.42–0.31 (2H, m). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 167.2, 160.9, 141.3, 138.0, 134.6 (× 2C), 132.1, 131.6, 126.7, 124.4, 122.7, 120.6, 114.2 (× 2C), 109.1, 72.3, 10.0, 3.1 (× 2C).

#### 2.4.4. 2-((4-((6-Chloro-2-oxoindolin-3-ylidene)methyl)phenoxy)acetic acid (46)

To a solution of compound **43** (60 mg, 0.17 mmol) in THF/water 2:1 (6 ml), LiOH monohydrate (18 mg, 0.42 mmol) was added and the reaction was left stirring at rt. After 16 h, the mixture was concentrated under reduced pressure and extracted twice with Et<sub>2</sub>O. The resulting aqueous phase was treated with HCl 37% until pH = 2, then extracted three times with EtOAc, washed with brine, dried over Na<sub>2</sub>SO<sub>4</sub>, then evaporated under reduced pressure to afford 40 mg of **46** in 71% yield, as an orange solid (m.p. 197 °C, dec.). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), *E/Z* 74/26, δ: 13.08 (1H isomer *E*+1H isomer *Z*, bs); 10.75 (1H isomer *Z*, bs); 10.73 (1H isomer *E*, bs); 7.90–7.86 (2H isomer *Z*, m); 7.82 (1H isomer *Z*, s); 7.74–7.68 (2H, isomer *Z*, m); 7.66–7.61

(1H isomer *E*+1H isomer *Z*, m); 7.48 (1H, d, *J* = 8.3 Hz); 7.11–7.15 (2H isomer *Z*, m); 7.10–7.06 (2H isomer *E*+1H isomer *Z*, m); 6.97–6.93 (1H isomer *E*+1H isomer *Z*); 6.90 (1H isomer *E*, d, *J* = 2.2 Hz); 4.84 (2H isomer *Z*, s); 4.80 (2H isomer *E*, s). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 169.9, 168.8, 159.2, 144.0, 136.7, 133.7, 131.6 (× 2C), 126.8, 124.6, 123.3, 120.8, 120.0, 114.8, 110.0, 64.6.

**4-((4-((6-chloro-2-oxoindolin-3-ylidene)methyl)phenoxy)methyl)benzoic acid (47)**. To a solution of compound **44** (70 mg, 0.17 mmol) in THF/water 2:1 (6 ml), LiOH monohydrate (18 mg, 0.42 mmol) was added and the reaction was left stirring at rt. After 16 h, the mixture was concentrated under reduced pressure and extracted twice with Et<sub>2</sub>O. The resulting aqueous phase was treated with HCl 37% until pH = 2, then extracted three times with EtOAc, washed with brine, dried over Na<sub>2</sub>SO<sub>4</sub>, then evaporated under reduced pressure to afford 60 mg of **47** in 87% yield, as a yellow solid (m.p. 279–280 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 13.08 (1H, bs); 10.73 (1H, bs); 8.04–7.96 (2H, m); 7.77–7.71 (2H, m); 7.66–7.56 (4H, m); 7.22–7.17 (2H, m); 6.94 (1H, dd, *J* = 1.4 Hz, 8.6 Hz); 6.90 (1H, d, *J* = 1.4 Hz); 5.32 (2H, s). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 168.8, 167.0, 159.6, 144.0, 141.7, 136.8, 133.6, 131.8 (× 2C), 130.3, 129.5 (× 2C), 127.5 (× 2C), 126.7, 124.6, 123.3, 120.8, 120.1, 110.0, 69.0.

**6-chloro-3-((5-phenylthiophen-2-yl)methylene)indolin-2-one (48)** was prepared from **31** and 5-phenylthiophene-2-carbaldehyde. Purification by column chromatography (hexane/EtOAc 90:10, CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 95:5 → 90:10) gave the title compound 98% yield, as a red solid (m.p. 233–234 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 10.80 (1H, bs); 8.18 (1H, s); 7.93 (1H, d, *J* = 3.4 Hz); 7.81–7.76 (2H, m); 7.72 (1H, d, *J* = 8.3 Hz); 7.68 (1H, d, *J* = 3.4 Hz); 7.53–7.47 (2H, m); 7.44–7.38 (1H, m); 7.07 (1H, dd, *J* = 8.3 Hz, 1.9 Hz); 6.89 (1H, d, *J* = 1.9 Hz). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 167.2, 150.7, 141.5, 139.5, 136.7, 133.4, 132.4, 129.3 (× 2C), 129.0, 128.7, 125.8 (× 2C), 124.3, 123.4, 120.8 (× 2C), 120.3, 109.4.

**3-((2-bromothiazol-5-yl)methylene)-6-chloroindolin-2-one (49)** was prepared from **31** and 2-bromo-5-formylthiazole. Purification by column chromatography (petroleum ether/EtOAc 95:5 → 10:90, CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 95:5 → 85:15) gave the title compound an 81% yield, as a red solid (m.p. 262–263 °C, dec.). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 10.99 (bs, 1H); 8.23 (2H, s); 7.71 (1H, d, *J* = 8.4 Hz); 7.11 (1H, dd, *J* = 8.4 Hz, 1.9 Hz); 6.92 (1H, d, *J* = 1.9 Hz). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 167.4, 151.2, 143.7, 142.1, 135.6, 133.5, 124.8, 123.0, 122.3, 121.5, 121.3, 109.9.

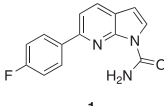
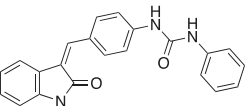
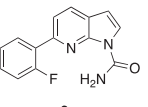
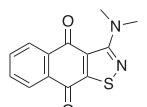
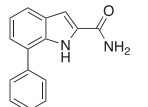
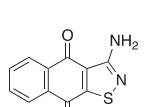
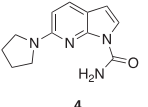
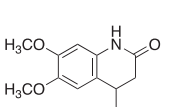
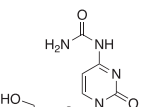
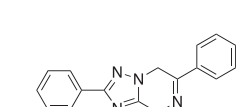
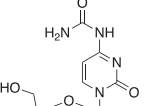
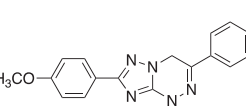
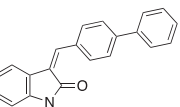
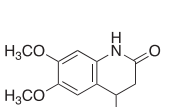
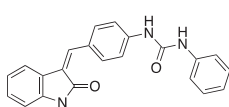
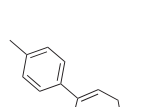
**3-((1H-indol-5-yl)methylene)-6-chloroindolin-2-one (50)** was prepared from **31** and 1H-indole-5-carbaldehyde. Purification by column chromatography (hexane/EtOAc 95:5 → 20:80, CH<sub>2</sub>Cl<sub>2</sub>:CH<sub>3</sub>OH 95:5 → 90:10) gave the title compound in 34% yield, as a yellow solid (m.p. 238–239 °C). <sup>1</sup>H NMR (600 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 11.46 (1H, bs); 10.70 (1H, s); 7.98 (1H, s); 7.82 (1H, s); 7.77 (1H, d, *J* = 8.1 Hz); 7.55 (1H, d, *J* = 8.6 Hz); 7.51 (1H, d, *J* = 8.6 Hz); 7.49–7.46 (1H, m); 6.96–6.92 (1H, m); 6.92–6.88 (1H, m). <sup>13</sup>C NMR (150 MHz, DMSO-*d*<sub>6</sub>), major isomer (*E*), δ: 169.1, 152.5, 143.8, 139.6, 136.8, 133.2, 127.8, 127.0124.8, 123.4, 122.9 (× 3C), 120.7, 120.5, 111.8, 109.8, 102.1.

## 3. Results and discussion

### 3.1. Synthesis and biological activity evaluation

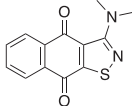
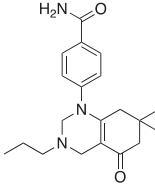
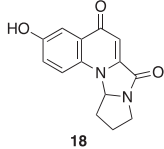
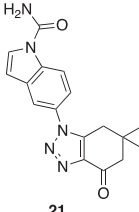
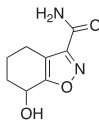
A small collection of molecules containing highly diverse chemical scaffolds was initially screened on c-Src to evaluate their inhibiting activity at a 100 μM concentration (Table 1). Triazolo-triazine **13**

**Table 1.** Evaluation of the inhibitory activity of c-Src by the *in-house* prepared collection of small molecules.

Compound	% Inhibition Src (100 $\mu$ M)	ID <sub>50</sub>	Compound	% Inhibition Src (100 $\mu$ M)	ID <sub>50</sub>
	58.1%			84.0%	
	28.6%			43.5	
	34.8%			2.2%	
	4.0%			40.0%	
	1.8%			90.0%	50.0 $\mu$ M
	10.3%			n.d.	
	77.3%			7.8%	
	83.2%	12.5 $\mu$ M		0.6%	

(continued)

Table 1. Continued.

Compound	% Inhibition Src (100 $\mu$ M)	ID <sub>50</sub>	Compound	% Inhibition Src (100 $\mu$ M)	ID <sub>50</sub>
	57.6%			7.7%	
<b>10</b>			<b>20</b>		
	15.5%			n.d.	
<b>18</b>			<b>21</b>		
	12.8%		<b>Dasatinib</b>	100.0%	1.6 nM
<b>19</b>					

Dasatinib was used as reference compound.

showed the best inhibition potency (90%). However, its closely related analogue **14** did not affect the enzyme activity. Indolinone and *N*-hydroxyl indolinone derivatives **7–9** exhibited a significant activity (percent inhibition ranging from 77 to 84%). A modest inhibition profile was highlighted for azaindole **1** and kynurenic derivative **17** (about 58%), followed by thiazolybenzoquinone **10** (43%) and dihydroquinolin-2-one **12** (40%), whereas all the other entries of the library showed low to no activity towards c-Src. ID<sub>50</sub> of both triazolo-triazine **13** and indolinone **8** were in the two-digit micromolar concentration (50 and 12.5  $\mu$ M, respectively).

These results prompted us to focus on the indolinone core for further investigations. This skeleton can be considered a very attractive target for biological activity evaluation and SAR studies, offering the possibility of easy chemical accessibility and several points for chemical decoration. We thus prepared and tested a second-generation set of compounds containing the indolinone scaffold substituted at positions 5 and/or 6 and bearing an arylidene side chain appended to position 3. We also prepared **24** to evaluate the role of the heteroaromatic system and both **22** and **23** to investigate the effect of the OH group on the nitrogen atom. All the compounds showed a good dose-dependent activity at 100  $\mu$ M and 10  $\mu$ M concentrations, with **26** having the best inhibitory potency of the series (95 and 67% at 100 and 10  $\mu$ M, respectively) (Table 2).

Compound **26** was prioritised and its structure was used as the starting point to prepare new analogues, maintaining unaltered the 6-chloro oxindole moiety and introducing decorations on the aromatic ring connected by a methylene bridge to C3. Knoevenagel condensation conditions were exploited to react 6-chloro oxindole **31** with substituted aromatic/heteroaromatic aldehydes (Scheme 1), obtaining 15 new variously substituted derivatives.

All the analogues contained a phenyl ring connected by a conjugated vinyl fragment to C3. Various substituents with different

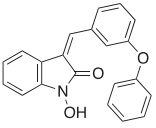
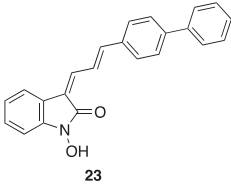
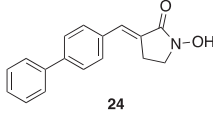
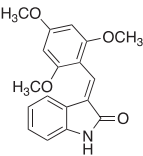
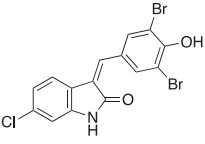
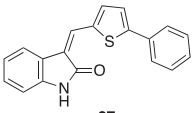
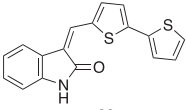
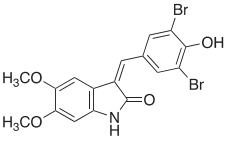
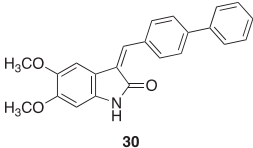
stereoelectronic properties were introduced in the para position of the ring: cyano (**32**), nitro (**33**), dimethylamino (**34**), phenolic (**35**) and carboxylic (**36**) groups. Commercially available *p*-benzyloxy benzaldehyde and *m*-phenoxy benzaldehyde were employed for the preparation of **37** and **38**. Additionally, alkylation of 4-hydroxy benzaldehyde **39** as in **43–47** gave the opportunity to increase the overall compound length and eventually fill some empty regions of the cavity, either in presence of aromatic or aliphatic moieties, as well as with a terminal acidic or ester portion. Finally, compounds containing heterocyclic substituents were synthesised (**48–50**) to investigate their potential role as hydrogen bond acceptors/donors.

All the obtained compounds were characterised by NMR analysis, showing the presence of isomeric mixtures (from 9:1 to 7:3 *E/Z* isomers). The relative ratios of the *E* and *Z* stereoisomers within this series (**32–50**) were assigned based on the chemical shifts of protons at the C2' and C6' positions<sup>41</sup>. Previous investigations on 3-arylideneoxindoles have reported the *E/Z* isomerisation to be solvent-, temperature-, time-, and light-dependent<sup>48–50</sup>. For this reason, the compounds were tested as mixtures of the two isomers.

The percentage of inhibition at the concentrations of 100  $\mu$ M and 10  $\mu$ M, ID<sub>50</sub> of the most active compounds, and their *K<sub>i</sub>* were evaluated and reported in Table 3. Compounds **32–34** showed a good inhibitory activity towards the enzyme, with **34** having the best profile of inhibition (85% at 10  $\mu$ M). Conversely, **36**, with a carboxylic group directly linked to the phenyl ring, had very low activity. However, the percentage of inhibition was enhanced by increasing the length of the spacer between the aromatic ring and the acidic group (**46**, 70% at 100  $\mu$ M, **47** 94% at 100  $\mu$ M), with the corresponding esters (**43** and **44**) showing a good activity as well. Replacement of the acidic group of **46** with a hydrophobic cyclopropyl substituent led to an almost inactive compound (**45**). Moreover, the phenolic derivative **35** and the corresponding



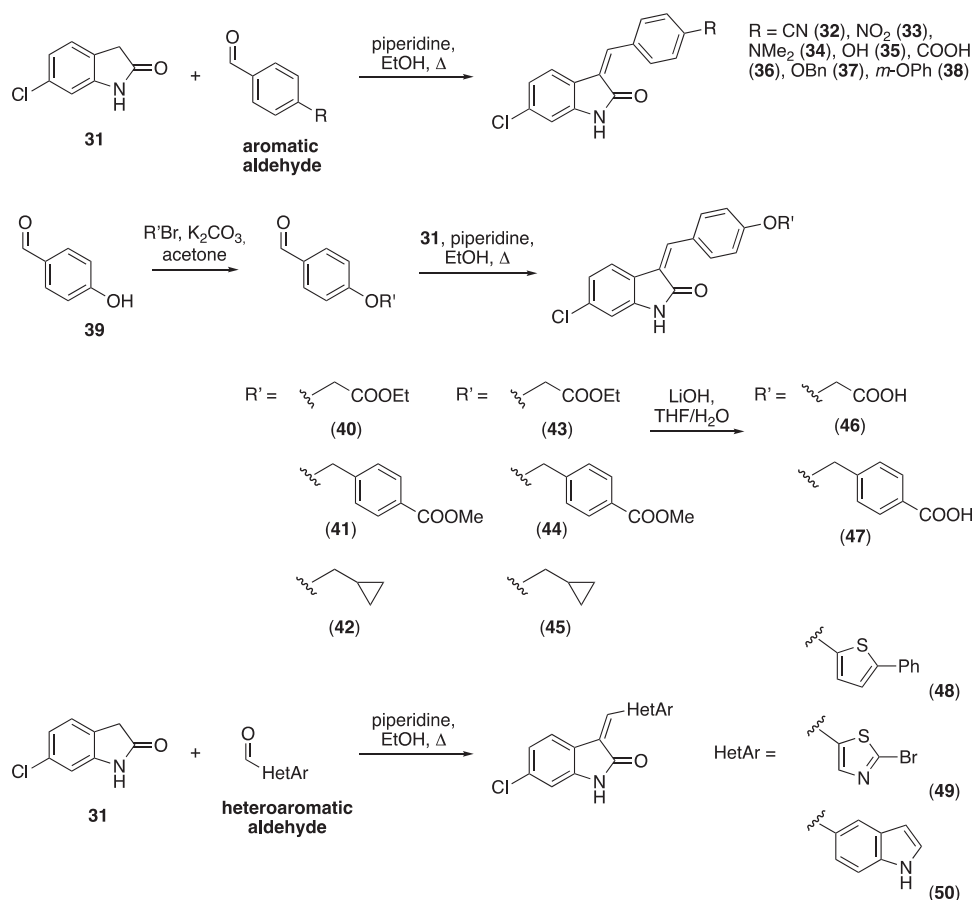
**Table 2.** Inhibitory activity evaluation of indolinones 22–30.

Compound	% Inhibition		ID <sub>50</sub> (μM)	K <sub>i</sub> (ATP competitive)
	100 μM	10 μM		
<b>Dasatinib</b>	100	100	1.60 nM ± 0.22 nM	0.80 nM ± 0.11 nM
 <b>22</b>	64.61 ± 3.80	32.06 ± 3.53		
 <b>23</b>	57.44 ± 6.31	22.77 ± 2.94		
 <b>24</b>	87.46 ± 7.41	34.18 ± 0.60		
 <b>25</b>	35.79 ± 3.69	23.15 ± 5.55		
 <b>26</b>	94.70 ± 0.30	67.03 ± 0.10	5.63 μM ± 2.46 μM	3.80 μM ± 0.63 μM
 <b>27</b>	45.58 ± 7.85	25.30 ± 9.71		
 <b>28</b>	55.15 ± 0.91	30.69 ± 10.16		
 <b>29</b>	78.56 ± 1.32	39.81 ± 3.16		
 <b>30</b>	42.03 ± 8.91	9.39 ± 19.22		

Dasatinib was used as reference compound.

benzyl ether **37** maintained high inhibitory activity at the highest dose, which significantly decreased at 10 μM. The shift of a lipophilic aromatic ring to *meta* position on the phenyl group (**38**) did not influence the activity. Among compounds containing

heterocyclic rings, **50** showed the best activity at both concentrations, whereas the thiophene- and thiazole-containing compounds **48** and **49**, respectively, evidenced a drop in activity, in particular at the lowest concentration.



**Scheme 1.** General synthesis by Knoevenagel condensation starting from 6-chloro oxindole and different aromatic and heteroaromatic compounds.

Compound **34** resulted to be the best of the series, showing significant inhibition of the enzyme at  $10 \mu\text{M}$  (85%), with  $\text{ID}_{50}$  and  $K_i$  in the sub-micromolar range (0.71 and  $0.48 \mu\text{M}$ , respectively). These findings suggest that the presence of an anilino nitrogen could have an important role in the inhibitory activity of the compounds.

With these results in our hands, the cytotoxicity test of selected molecules was carried out on the MCF-7 breast cancer cell line and compared to the *c*-Src inhibitor Dasatinib used as standard (Table 4). Thiazole **49** showed potency comparable to Dasatinib. Ester **43** and indole derivative **50** were able to exert some relevant but not excellent cytotoxic activity. It is worth noting that **34**, the best inhibitor in the cell-free assay, did not show any cytotoxic activity, probably due to pharmacokinetic issues.

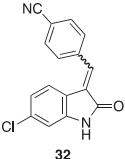
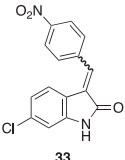
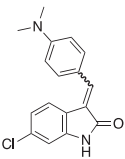
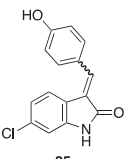
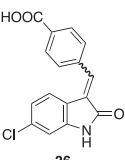
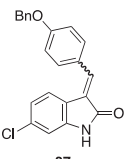
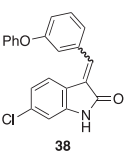
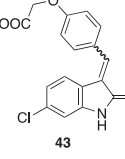
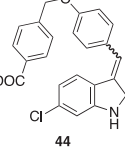
### 3.2. Molecular docking studies

In the attempt to find a putative binding mode of the new indolinone derivatives within the ATP-binding site of *c*-Src, molecular docking simulations were performed using the software Glide. For this purpose, the three-dimensional structure of the co-crystallized complex between *c*-Src and **AP23464** (entry 2BDJ of the protein data bank,  $2.50 \text{ \AA}$  resolution) was used for calculations. Preliminary validation of the docking protocol was done by re-docking **AP23464** into the binding site. The resulting rmsd =  $0.903 \text{ \AA}$ , which was calculated on the atomic coordinates of the inhibitor, confirmed the reliability of the docking protocol. Two highly conserved hydrogen bonds were found between N7 and aniline nitrogen of **AP23464** with the amide nitrogen and the carbonyl of the

backbone of the Met341, respectively (Figure 3). The 3-hydroxyphenylethyl group in position N9 on the purine scaffold protruded into the hydrophobic pocket and mediated a series of interactions. Specifically, the hydroxyl substituent formed hydrogen bonds with the carboxyl group of the Glu310 side chain and the backbone NH group of Asp404.

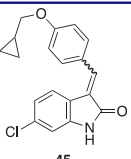
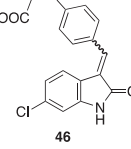
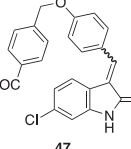
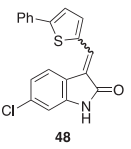
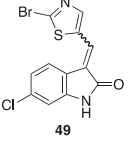
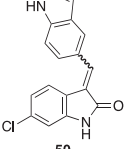
Considering that the new compounds were assayed as *E/Z* isomers, isomerisation was allowed during calculations. Docking scores showed that for each compound, the *E*-isomer was preferred to the *Z*-isomer. Overall, the indolinone derivatives showed a common binding pose with the lactam moiety that mimicked the hydrogen bond acceptor-donor motif represented by the N9 and the anilino NH group of **AP23464**. They served as hydrogen bond acceptor-donor system to give the classical interactions with Met341 of the hinge region (Figure 3). Moreover, the chlorophenyl moiety was superimposable to the cyclopentyl ring of **AP23464**. Finally, the aryl appendage at C3 could mimic the phenol portion of the co-crystallized inhibitor. As an example, the predicted binding mode of **33**, which showed single-digit micromolar  $\text{ID}_{50}$  and  $K_i$ , was able to maintain the two hydrogen bonds with Met341 in its best-scored pose (Figure 3). In addition, one of the oxygen atoms of the nitro group matched the phenol oxygen of **AP23464**, thus mimicking its ability to make an additional hydrogen bond with Asp404. A very similar binding mode was found for **50**, where the NH group of the pendant indole nucleus was able to build a hydrogen bond with the side chain of Glu310, in addition to the classical interactions with Met341 (Figure S1). On the other hand, molecular docking and scoring were not able to justify the binding affinity of **34**, whose peripheral dimethylanilino

**Table 3.** Evaluation of the inhibitory activity of the novel indolinone derivatives.

Compound	% Inhibition		ID <sub>50</sub>	K <sub>i</sub> (ATP competitive)
	100 μM	10 μM		
<b>Dasatinib</b>	100	100	1.60 nM ± 0.22 nM	0.80 nM ± 0.11 nM
	88.96 ± 1.35	35.74 ± 9.66	>10 μM	–
	69.94 ± 0.19	53.03 ± 18.08	2.43 μM ± 1.03 μM	1.64 μM ± 0.69 μM
	75.72 ± 7.63	85.35 ± 6.98	0.71 μM ± 0.43 μM	0.48 μM ± 0.29 μM
	86.86 ± 2.50	37.90 ± 0.35	>10 μM	–
	47.19 ± 9.33	4.15 ± 1.28	–	–
	84.27 ± 0.82	16.86 ± 13.95	–	–
	53.62 ± 12.91	31.48 ± 3.16	–	–
	80.84 ± 6.47	71.67 ± 2.27	3.24 μM ± 0.43 μM	2.19 μM ± 1.03 μM
	92.13 ± 3.84	41.05 ± 10.30	–	–

(continued)

Table 3. Continued.

Compound	% Inhibition		ID <sub>50</sub>	K <sub>i</sub> (ATP competitive)
	100 μM	10 μM		
 45	34.40 ± 13.26	14.44 ± 2.09	–	–
 46	70.56 ± 4.80	0	–	–
 47	93.88 ± 0.58	63.39 ± 1.06	5.31 μM ± 0.47 μM	3.58 μM
 48	87.71 ± 3.04	12.83 ± 1.32	–	–
 49	76.82 ± 9.22	53.85 ± 26.57	>10 μM	–
 50	96.08 ± 1.42	41.20 ± 1.12	9.14 μM ± 1.74 μM	6.17 μM

Dasatinib was used as reference compound.

Table 4. Cytotoxicity evaluation of selected compounds on human MCF-7 breast cancer cell line.

Compound	MCF-7 (IC <sub>50</sub> , μM)
Dasatinib	27 ± 1
33	>100
34	>100
43	85 ± 2
49	33 ± 2
50	68 ± 2

24 h after seeding, cells were exposed for 72 h to the compounds and cytotoxicity was measured using MTS assay. Experiments were performed in triplicate and data represent mean values ± SD. Dasatinib was used as reference compound.

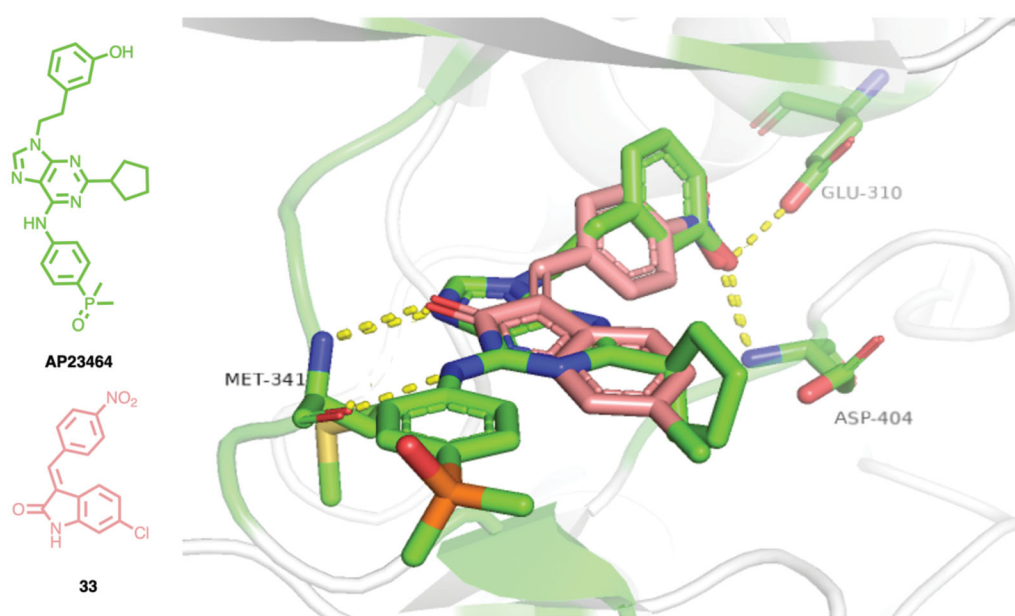
moiety gave hydrophobic interactions with Ala403, whereas other interactions were not highlighted, in addition to the classical contacts with Met341 (Figure S2).

Increasing the length and bulkiness of the substituent at the pendant phenyl ring forced the terminal molecular edge outside the hydrophobic side pocket where the phenol ring of **AP23464** was accommodated. As an example, the benzoic carboxyl group

of the long side chain of **47**, which retained the two hydrogen bonds with Met341, made two additional hydrogen bonds with Cys277 and Phe278. On the contrary, the docking pose of the corresponding methyl ester **44** lacked the fundamental anchor points on the binding site to avoid steric clashes due to the terminal methyl group of the ester function.

#### 4. Conclusion

An *in-house* library of structurally diverse small molecules was screened with the aim of finding new c-Src inhibitors. It emerged that the most active compounds contained an indolinone core, which was selected as a promising scaffold for further investigations. Various substituted 3-(hetero)arylideneindolin-2-ones were designed and synthesised to highlight the structural determinants for the activity. Most of the compounds showed a percentage of inhibition of the enzyme >70% at 100 μM and some of them also had a significant activity at 10 μM. Selected molecules were assayed on the human MCF-7 breast cancer cell line, showing



**Figure 3.** Graphical representation of the best-scored binding pose of **33** (magenta) within the c-Src ATP binding site, in comparison to the co-crystallized inhibitor AP23464 (green). Yellow dashed lines represent hydrogen bonds between Met341, Glu310, Asp404, and both ligands.

moderate antiproliferative activity. Among the tested compounds, **49** demonstrated cytotoxicity comparable to that of the reference compound Dasatinib, but lacked any significant activity in the cell-based assay, indicating the possibility that some off-target effect could take place. Conversely, **34**, which showed the best inhibition profile towards the isolated enzyme, resulted ineffective ( $IC_{50} > 100 \mu M$ ) on the cell line. This finding likely depends on pharmacokinetic issues probably related to the difficulty of the compound to reach the intracellular target. Therefore, further studies will be necessary to improve the pharmacological profile of this class of compounds. The results of molecular docking studies, which were performed to propose a binding mode within the c-Src binding site, will guide and complement future activities.

### Acknowledgements

We thank AIRC Foundation for Cancer Research in Italy (Fondazione AIRC per la Ricerca sul Cancro, AIRC): Grant IG-2017, project code 20762, PI Giovanni Maga iii) Grant IG-2020, project code 24448, PI Emmanuele Crespan; E.C. wishes to thank Cariplo Foundation, project 2019–1836.

### Disclosure statement

No potential conflict of interest was reported by the author(s).

### Funding

This work was supported by MIUR (PRIN 2017, project code 2017SA5837).

### ORCID

Sabrina Dallavalle  <http://orcid.org/0000-0002-3340-5165>

### References

1. Simatou A, Simatos G, Goulielmaki M, et al. Historical retrospective of the SRC oncogene and new perspectives. *Mol Clin Oncol* 2020;13:1–10.
2. Parsons SJ, Parsons JT. Src family kinases, key regulators of signal transduction. *Oncogene* 2004;23:7906–9.
3. Brown MT, Cooper JA. Regulation, substrates and functions of src. *Biochimica et Biophysica Acta (BBA) – Reviews on Cancer* 1996;1287:121–49.
4. Playford MP, Schaller MD. The interplay between Src and integrins in normal and tumor biology. *Oncogene* 2004;23:7928–46.
5. Sigal CT, Zhou W, Buser CA, et al. Amino-terminal basic residues of Src mediate membrane binding through electrostatic interaction with acidic phospholipids. *Proc Natl Acad Sci USA* 1994;91:12253–7.
6. Boggon TJ, Eck MJ. Structure and regulation of Src family kinases. *Oncogene* 2004;23:7918–27.
7. Amata I, Maffei M, Pons M. Phosphorylation of unique domains of Src family kinases. *Front Genet* 2014;5:181.
8. Roskoski R. Src kinase regulation by phosphorylation and dephosphorylation. *Biochem Biophys Res Commun* 2005;331:1–14.
9. McLachlan RW, Kraemer A, Helwani FM, et al. E-cadherin adhesion activates c-Src signaling at cell–cell contacts. *Mol Biol Cell* 2007;18:3214–23.
10. Patel A, Sabbineni H, Clarke A, Somanath PR. Novel roles of Src in cancer cell epithelial-to-mesenchymal transition, vascular permeability, microinvasion and metastasis. *Life Sci* 2016;157:52–61.
11. Maa MC, Leu TH, McCarley DJ, et al. Potentiation of epidermal growth factor receptor-mediated oncogenesis by c-Src: implications for the etiology of multiple human cancers. *Proc Natl Acad Sci* 1995;92:6981–5.
12. Shah K, Vincent F. Divergent roles of c-src in controlling platelet-derived growth factor-dependent signaling in fibroblasts. *Mol Biol Cell* 2005;16:5418–32.

13. Ishizawar RC, Miyake T, Parsons SJ. c-Src modulates ErbB2 and ErbB3 heterocomplex formation and function. *Oncogene* **2007**; 26:3503–10.
14. Rahimi N, Hung W, Tremblay E, et al. c-Src kinase activity is required for hepatocyte growth factor-induced motility and anchorage-independent growth of mammary carcinoma cells\*. *J Biol Chem* **1998**;273:33714–21.
15. Calgani A, Vignaroli G, Zamperini C, et al. Suppression of SRC signaling is effective in reducing synergy between glioblastoma and stromal cells. *Mol Cancer Ther* **2016**;15:1535–44.
16. González L, Agulló-Ortuño MT, García-Martínez JM, et al. Role of c-Src in human MCF7 breast cancer cell tumorigenesis\*. *J Biol Chem* **2006**;281:20851–64.
17. Zhang J, Kalyankrishna S, Wislez M, et al. Src-family kinases are activated in non-small cell lung cancer and promote the survival of epidermal growth factor receptor-dependent cell lines. *Am J Pathol* **2007**;170:366–76.
18. Schweppe RE, Kerege AA, French JD, et al. Inhibition of Src with AZD0530 reveals the Src-focal adhesion kinase complex as a novel therapeutic target in papillary and anaplastic thyroid cancer. *J Clin Endocrinol Metabolism* **2009**;94:2199–203.
19. Lowe C, Yoneda T, Boyce BF, et al. Osteopetrosis in Src-deficient mice is due to an autonomous defect of osteoclasts. *Proc Natl Acad Sci* **1993**;90:4485–9.
20. Rucci N, Susa M, Teti A. Inhibition of protein kinase c-Src as a therapeutic approach for cancer and bone metastases. *Anti-Cancer Agents Med Chem* **2008**;8:342–9.
21. Jin W. Regulation of Src family kinases during colorectal cancer development and its clinical implications. *Cancers* **2020**;12:1339.
22. Lutz MP, Eßer IBS, Flossmann-Kast BBM, et al. Overexpression and activation of the tyrosine kinase Src in human pancreatic carcinoma. *Biochem Biophys Res Commun* **1998**;243:503–8.
23. Varkaris A, Katsiampoura AD, Araujo JC, et al. Src signaling pathways in prostate cancer. *Cancer Metastasis Rev* **2014**;33:595–606.
24. Belli S, Esposito D, Servetto A, et al. c-Src and EGFR inhibition in molecular cancer therapy: what else can we improve? *Cancers*. **2020**;12:1489.
25. Rivera-Torres J, San José E. Src tyrosine kinase inhibitors: new perspectives on their immune, antiviral, and senotherapeutic potential. *Front Pharmacol* **2019**;10:1011.
26. Musumeci F, Brullo C, Grossi G, et al. Identification of new pyrrolo[2,3-d]pyrimidines as Src tyrosine kinase inhibitors in vitro active against Glioblastoma. *Eur J Med Chem* **2017**;127:369–78.
27. Ceccherini E, Indovina P, Zamperini C, et al. SRC family kinase inhibition through a new pyrazolo[3,4-d] pyrimidine derivative as a feasible approach for glioblastoma treatment. *J Cell Biochem* **2015**;116:856–63.
28. Gnoni A, Marech I, Silvestris N, et al. Dasatinib: an anti-tumour agent via Src inhibition. *Current Drug Targets* **2011**;12:563–78.
29. Steinbach A, Clark SM, Clemmons AB. Bosutinib: a novel src/abl kinase inhibitor for chronic myelogenous leukemia. *J Adv Pract Oncol* **2013**;4:451–5.
30. Hennequin LF, Allen J, Breed J, et al. N-(5-Chloro-1,3-benzodioxol-4-yl)-7-[2-(4-methylpiperazin-1-yl)ethoxy]-5-(tetrahydro-2H-pyran-4-yloxy)quinazolin-4-amine, a novel, highly selective, orally available, dual-specific c-Src/Abl kinase inhibitor. *J Med Chem* **2006**;49:6465–88.
31. Cooper MR, Yi SY, Alghamdi W, et al. Vandetanib for the treatment of medullary thyroid carcinoma. *Ann Pharmacother* **2014**;48:387–94.
32. Tanimura K, Yamasaki K, Okuhiro Y, et al. Monitoring ponatinib in a child with philadelphia chromosome-positive acute lymphoblastic leukemia. *Case Rep Oncol* **2021**;14:24–8.
33. Gurbani D, Du G, Henning NJ, et al. Structure and characterization of a covalent inhibitor of Src kinase. *Front Mol Biosci* **2020**;7:81–9.
34. Temps C, Lietha D, Webb ER, et al. A conformation selective mode of inhibiting SRC improves drug efficacy and tolerability. *Cancer Res* **2021**;81:5438–50.
35. Hong DS, Sen S, Park H, et al. A phase I, open-label, multicenter, first-in-human study of the safety, tolerability, pharmacokinetics, and antitumor activity of TPX-0022, a novel MET/CSF1R/SRC inhibitor, in patients with advanced solid tumors harboring genetic alterations in MET. *J Clin Oncol* **2020**;38:TPS3663–TPS3663.
36. Wang X, Ren R, Xu Z, et al. Tirbanibulin attenuates pulmonary fibrosis by modulating Src/STAT3 signaling. *Front Pharmacol* **2021**;12:693906.
37. Markham A, Duggan S. Tirbanibulin: first approval. *Drugs* **2021**;81:509–13.
38. Cincinelli R, Musso L, Merlini L, et al. 7-Azaindole-1-carboxamides as a new class of PARP-1 inhibitors. *Bioorg Med Chem* **2014**;22:1089–103.
39. Aiwale TS, Dallavalle S. Efficient synthesis of 3,7-Diaryl-1,4-dihydro[1,2,4]triazolo[5,1-c][1,2,4]triazines. *Synthesis* **2012**;44:3055–8.
40. Cincinelli R, Cassinelli G, Dallavalle S, et al. Synthesis, modeling, and RET protein kinase inhibitory activity of 3- and 4-Substituted  $\beta$ -Carbolin-1-ones. *J Med Chem* **2008**;51:7777–87.
41. Musso L, Cincinelli R, Zuco V, et al. 3-Arylidene-N-hydroxyoxindoles: a new class of compounds endowed with antitumor activity. *ChemMedChem* **2016**;11:1700–4.
42. Rizzi E, Cassinelli G, Dallavalle S, et al. Synthesis and RET protein kinase inhibitory activity of 3-arylidenebenzylideneindolin-2-ones. *Bioorg Med Chem Lett* **2007**;17:3962–8.
43. Cautain B, Castillo F, Musso L, et al. Discovery of a novel, isothiazolonaphthoquinone-based small molecule activator of FOXO nuclear-cytoplasmic shuttling. *PLOS One* **2016**;11:e0167491.
44. Cincinelli R, Musso L, Beretta G, Dallavalle S. 4-Quinolone fused heterocyclic ring systems by intramolecular reactions of 4-quinolone-2-carboxamides. *Tetrahedron* **2014**;70:9797–804.
45. Musso L, Cincinelli R, Giannini G, et al. Synthesis of 5,6-dihydro-4H-benzo[d]isoxazol-7-one and 5,6-dihydro-4H-isoxazolo[5,4-c]pyridin-7-one Derivatives as Potential Hsp90 Inhibitors. *Chemical Biology & Drug Design* **2015**;86:1030–5.
46. Höttecke N, Liebeck M, Baumann K, et al. Inhibition of  $\gamma$ -secretase by the CK1 inhibitor IC261 does not depend on CK1 $\delta$ . *Bioorg Med Chem Lett* **2010**;20:2958–63.
47. la Pietra V, Sartini S, Botta L, et al. Challenging clinically unresponsive medullary thyroid cancer: discovery and pharmacological activity of novel RET inhibitors. *Eur J Med Chem* **2018**;150:491–505.
48. Chu W, Zhou D, Gaba V, et al. Design, synthesis, and characterization of 3-(Benzylidene)indolin-2-one derivatives as ligands for  $\alpha$ -synuclein fibrils. *J Med Chem* **2015**;58:6002–17.
49. Ngai MH, So CL, Sullivan MB, et al. Photoinduced isomerization and hepatotoxicities of semaxanib, sunitinib and related 3-substituted indolin-2-ones. *ChemMedChem* **2016**;11:72–80.
50. Chen X, Yang T, Deivasigamani A, et al. N'-Alkylaminosulfonyl analogues of 6-fluorobenzylideneindolinones with desirable physicochemical profiles and potent growth inhibitory activities on hepatocellular carcinoma. *ChemMedChem* **2015**;10:1548–58.