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Multicomponent synthesis of novel functionalized spiroindenopyridotriazine-4*H*-pyrans†

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A very convenient one-pot method for the synthesis of new biologically active compounds, spiro-4*H*-pyran derivatives, has been developed. This approach, enables the incorporation of three pharmacophoric cores: pyran, pyridine and triazine scaffolds, within a single molecular skeleton using an efficient sequential multicomponent reaction. The starting materials include ninhydrin, cyanoacetohydrazide, ethyl cyanoacetate, aromatic aldehydes, pyrazolone and malononitrile. In this process, spiropyran derivatives were produced with high efficiency through three consecutive reactions. The first step involves the synthesis of 1,6-diaminopyridinone via a three-component approach using readily available starting materials. The second step, the synthesis of pyridotriazine, occurs in a domino way without the need to separate the intermediate product of the first step. Finally, the third step involves the synthesis of new spiro-4*H*-pyran derivatives through a one-pot three-component reaction. This method offers several advantages, including fast and simple purification of products, the use of safe solvents, no need for metallic and toxic catalysts, high atomic economy and excellent chemoselectivity of the reactions.

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Introduction

One of the goals of organic synthesis is the search for efficient conversion of simple raw materials into complex functional products using methods that combine environmental and economic aspects. In this regard, the use of multicomponent reactions (MCRs) has been developed as a powerful synthetic tool to achieve this purpose. MCRs have significant advantages over linear multistep processes. In these reactions, by changing each component, a large number of products can be created around a common framework. Also, unlike linear processes, there is no need for purification after each transformation. Therefore, due to the saving of time, cost and consumption of many chemical solvents, multicomponent reactions have been introduced as one of the most attractive topics in green chemistry.^{1–9} Indeed, performing multicomponent reactions in which cyclization is possible as a process step, is a powerful strategy for the facile synthesis of diverse heterocyclic frameworks. Increasing attention to diversity-oriented synthesis, which is easily achievable in multicomponent reactions, has increased interest to this approach.^{10–14}

The compounds containing the spirocyclic moiety have prominent medicinal activities and are present in the structure of many natural molecules with diverse biological effects. The presence of a spiro center has a significant impact on biological

activities due to the rigidity of the molecule structure. The three-dimensional nature of spiro compounds and natural bioactive structures (*e.g.* proteins) makes such compounds have more effective interactions with binding sites in proteins, which is an important feature in the drug development process.^{15–18}

Meanwhile, spiro-4*H*-pyrans exhibit various biological properties, including: antibacterial, anti-anaphylactic, anti-fungal, antitumor and antirheumatic activities.^{19–21} The structure of some synthetic products with biological properties is shown in Fig. 1.^{20,22–25}

Some spiro structures have also been found to act as anti-oxidants (AO), compounds that prevent damage from reactive oxygen and nitrogen species (ROS and RNS). These types can be produced naturally in living cells or caused by some biological disorders such as anxiety. AOs are free radical scavengers and therefore protect proteins, DNA and lipids from oxidative degradation.²⁵ Among the compounds whose antioxidant properties have been investigated is spiro-4*H*-pyran (**VII**), which is very similar to the products synthesized in this work.²⁶

In publications, there are several reports on the reaction of cyclic compounds containing active carbonyl moiety (isatin, ninhydrin, acenaphthoquinone) with malononitrile (or ethyl/methyl cyanoacetate) and various CH-acids in order to synthesize fused spiro-4*H*-pyrans through multicomponent reactions (Fig. 2(A–C)).^{27–34}

In recent years, novel carbonyl structures have been synthesized as a starting material for the preparation of new spiro-4*H*-pyrans. As shown in Fig. 2, these compounds include indenopyridazine (**D**),¹⁸ indenoquinoxaline (**E**)^{35–40} and indoloquinazoline (**F**).^{24,41,42} Recently, we used indenopyridotriazine as

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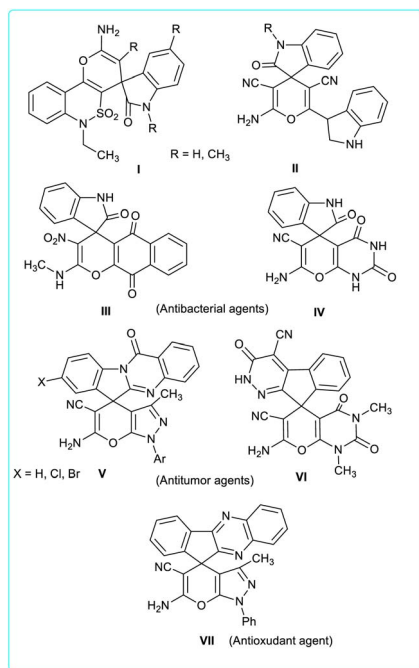


Fig. 1 Bioactive synthetic spiro-4H-pyrans.

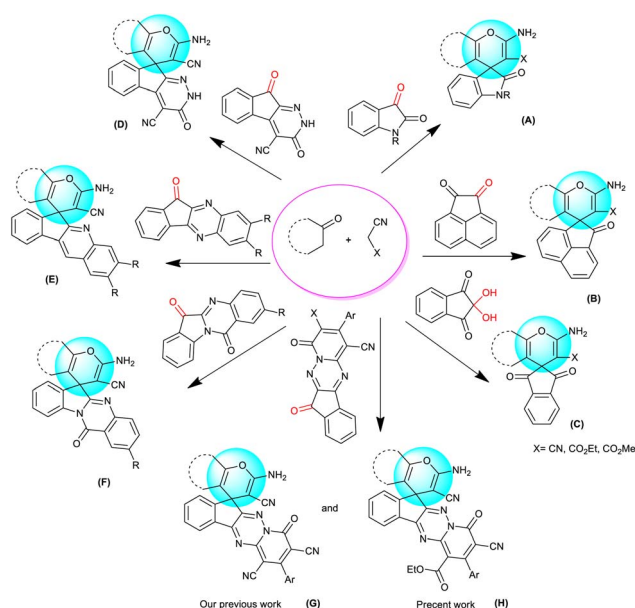


Fig. 2 Reports on the synthesis of various spiro-4H-pyrans.

the active carbonyl component for multicomponent synthesis of polysubstituted fused spiro-4H-pyran derivatives (G,H).⁴³ Most of the methods reported in the synthesis of spiro-4H-pyrans are associated with disadvantages such as the use of toxic solvents, expensive and metallic catalysts and the difficulty of the synthetic steps.^{27,30,36,44}

In this research, we succeeded in synthesizing new products from the spiro-4H-pyrans family by using multicomponent reactions compatible with green chemistry. In these structures, the frameworks of spiropyran and pyridotriazine are connected

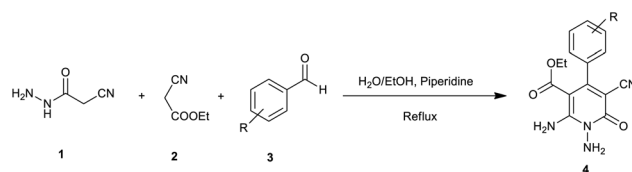
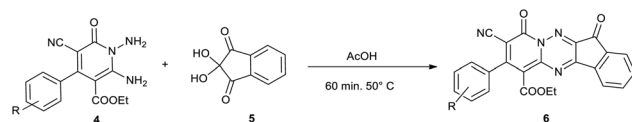
to each other, which according to the available references, each one is a part of the biologically privileged cores.

Results and discussion

First, the pyridinone products **4** were synthesized from the reaction of cyanoacetohydrazide **1**, ethyl cyanoacetate **2** and benzaldehyde derivatives **3** according to the previously reported method (Scheme 1).⁴⁵ Subsequently, pyridotriazine **6** was obtained by adding ninhydrin **5** in acetic acid to the pyridinone intermediate (Scheme 2).⁴⁶ Finally, the target spiropyran compounds **9**, were prepared by investigating the reaction of pyridotriazine **6** with malononitrile **7** and pyrazolone **8**.

Optimization of the conditions

To find the optimal conditions for these reactions, dihydroindeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-1-carboxylate **6a**, malononitrile **7** and 3-methyl-1-phenyl-1H-pyrazol-5(4H)-one **8** were used as model substrates under different conditions. As

Scheme 1 Synthetic scheme for the formation of pyridinones **4**.Scheme 2 Synthetic scheme for the formation of pyridotriazines **6**.Table 1 Optimization of reaction conditions for the generation of **9a**^a

Entry	Solvent	Catalyst	Time (h)	Temp (°C)	Yield ^b (%)
1	H ₂ O	—	10	100	No reaction
2	MeOH	—	8	64	25
3	MeCN	—	6	82	33
4	DMF	—	12	120	No reaction
5	EtOH	—	10	25	No reaction
6	EtOH	—	6	78	86
7	EtOH	Piperidine	6	78	45
8	EtOH	NEt ₃	6	78	35
9	EtOH	HOAc	6	78	Trace

^a Reagents and conditions: **6a**, **7**, **8** (1 mmol), catalyst (0.1 mmol), solvent (10 mL). ^b Isolated yield based on **9a**.



shown in Table 1, performing the reaction in water did not yield the desired product (entry 1, Table 1). Similarly, using organic solvents such as methanol, acetonitrile and dimethylformamide resulted in either low efficiency or no formation of spiro-4*H*-pyran product **9a** (entries 2–4 respectively, Table 1). No product was obtained when ethanol was used at room temperature (entry 5, Table 1). However, at 78 °C within 6 hours, product **9a** was synthesized with 86% yield (entry 6, Table 1). The use of acidic or basic catalysts under these condition reduced the efficiency of the final product (entry 7–9, Table 1). A detailed analysis of these reactions showed that the TLC of the final mixture, revealed a spot corresponding to pyridone. This observation suggests that some triazines decompose into their constituent components, probably due to the presence of the catalyst.

Based on the information obtained from Table 1, we successfully synthesized a series of new spiro-4*H*-pyran derivatives **9a–g** using triazines **6a–g**, malononitrile **7** and pyrazolone **8** as starting materials in ethanol at reflux conditions without the use of any catalyst (Scheme 3). The target products, spiroindenopyridotriazine-pyranopyrazole **9a–g**, were obtained with high efficiency (78–90%) and their information is given in Table 2.

Structure determination

The structure of the synthesized products **9a–g** is clearly were confirmed by evaluating their spectral data, including IR, ¹H NMR, ¹³C NMR and mass spectroscopic analyses (see the ESI†).

As a representative example, we were investigated the proton and carbon spectrum data of compound **9a** (Fig. 3). In the ¹H NMR spectrum of **9a**, the NH₂ group appears at δ 7.96 ppm (this peak is exchangeable with D₂O). Two methyl groups appear at δ 1.07 and 1.40 ppm (triplet and singlet respectively). A quartet signal at δ 4.15–4.20 ppm corresponds to the methylene group. Other signals in the range of 7.40 to 8.16 are related to the aromatic protons. In the ¹H-decoupled ¹³C NMR spectrum of **9a**, 32 distinct resonances confirm the proposed structure. The key signals include two methyl carbons (2CH₃, at δ 12.3, 13.7 ppm), the spiro carbon (C_{spiro}, at δ 47.7 ppm), the C=CNH₂ and CH₂ carbons (at δ 56.4 and 61.6 ppm) respectively. The carbonyl groups are observed at δ 163.9 and 160.1 ppm. Two signals at δ 95.5, 96.9 are assigned to C–COO and C=C–O (Fig. 3). In the FT-IR spectrum, absorption bands of **9a** appeared at 3365, 2199, 1727, 1682 cm^{−1} due to NH₂, CN and 2CO groups respectively. The mass spectrum of **9a** did not display the molecular-ion peak

(*m/z* 678) but the observation of the [M – 29]⁺ peak and the fragmentation pattern confirm the proposed structure.

Scope and limitations

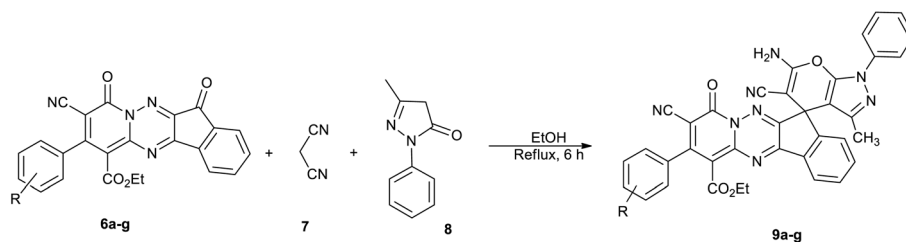
To further expand the scope of the reaction, we also tested various aromatic aldehydes (bearing electron-donating or electron-withdrawing groups) and 2-cyanoacetamide, ethyl-2-cyanoacetate, and malononitrile³⁸ under the optimized conditions. However, reactions with aldehydes such as 2-nitro and 2-hydroxybenzaldehyde did not result the desired product. With these derivatives, the synthesis of primary pyridone, in pure form was unsuccessful. TLC analysis indicated, the formation of two-component product likely resulting from the reaction of cyanoacetohydrazide with aldehyde. This outcome suggests that the amide group is hydrolyzed in ethanol under the reaction conditions. When malononitrile was used as the substrate spirane products were obtained with good to high yields.

With the established conditions in hand, we are continuing to explore the reaction's scope. Our ongoing research focuses on synthesizing diverse spirane products using other CH-acids, alkyl-2-cyanoacetate and pyrazolone derivatives. This work aims to enhance method for creating various complex spiro compounds.

We began our synthesis (of model substrates) with the preparation of a series of spiroindenopyridotriazine-4*H*-pyrans in just three steps on gram scale. This simple and scalable method significantly improves the ability to obtain useful quantities of these molecules for further research and potential applications.

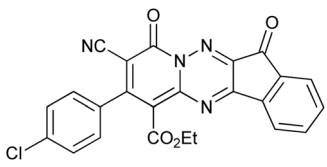
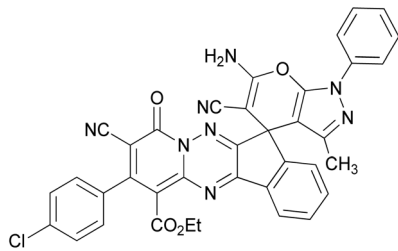
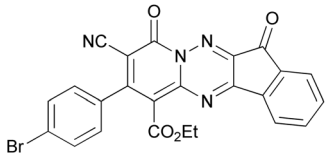
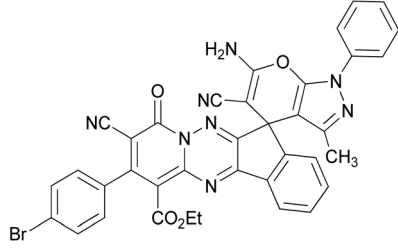
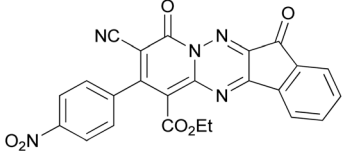
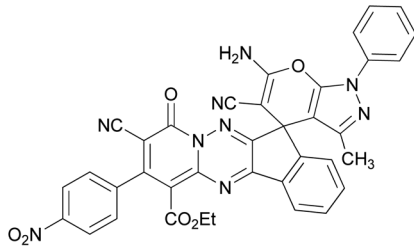
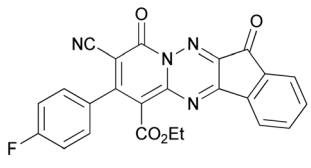
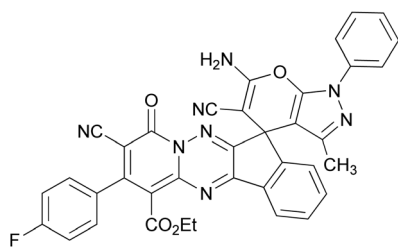
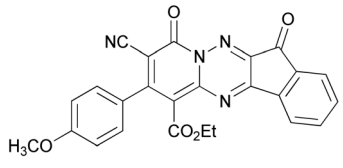
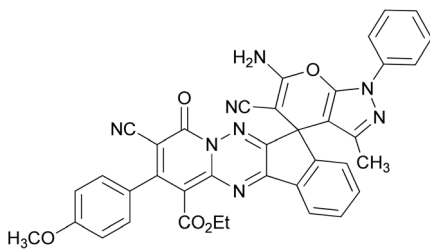
Proposed mechanism

A plausible mechanism for the formation of spiropyrans **9** (as an example **9a**) is shown in Scheme 4. At first, from the reaction of ninhydrin **5** and pyridone **4**, during two sequential condensation steps between the amino groups and the carbonyl groups, the 6-membered triazine ring is closed. It is clear that, the amine group attached to nitrogen participates first due to its higher nucleophilicity and attacks to the middle carbonyl group of ninhydrin, which is more electrophile. Then, the intermediate **10** is formed through the Knoevenagel condensation between pyridotriazine **6** and malononitrile **7**. The Michael addition of pyrazolone **8** to adduct **10**, generates intermediate **11**, which undergoes intramolecular cyclization *via* the nucleophilic attack of oxygen on the nitrile group to form

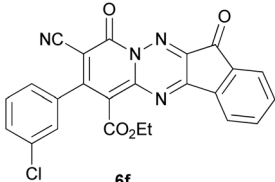
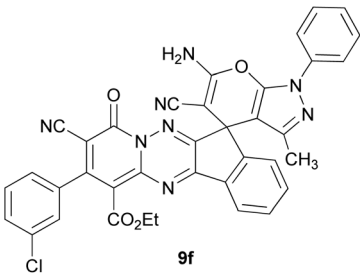
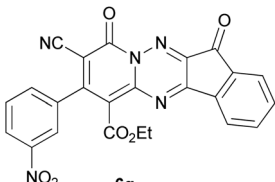
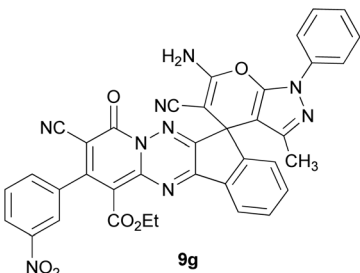


Scheme 3 Synthesis of products **9a–g**.

Table 2 Compounds 6a–g^a

Entry	Triazine	Product	Yield (%)	M.P. (°C)
1	 6a	 9a	86	253–255
2	 6b	 9b	82	266–268
3	 6c	 9c	90	260–262
4	 6d	 9d	78	275–278
5	 6e	 9e	80	295–297



Entry	Triazine	Product	Yield (%)	M.P. (°C)
6	 6f	 9f	82	245–247
7	 6g	 9g	85	282–284

intermediate **12**. Finally, intermediate **12** undergoes imine-enamine tautomerization to afford the target product **9a**.^{23,43}

Materials

Typical procedure for the synthesis of pyridotriazines (6a–g)

Firstly, 1,6-diaminopyridine-2-one compounds **4** were easily prepared *via* a three-component reaction between cyanoacetohydrazide **1**, benzaldehyde derivatives **3** and ethyl cyanoacetate **2** according to reported method in our previous work.⁴⁰ Then, a stoichiometric ratio (1 mmol) of ninhydrin **5** and acetic acid (5 mL) was added to the above mixture in a 25 mL round-bottomed flask connected to a reflux condenser, in a one-pot cascade manner. The resulting mixture was magnetically stirred in an oil-bath (50 °C, for 60 min). The completion time of the reaction was monitored by TLC using EtOAc/*n*-hexane (1 :

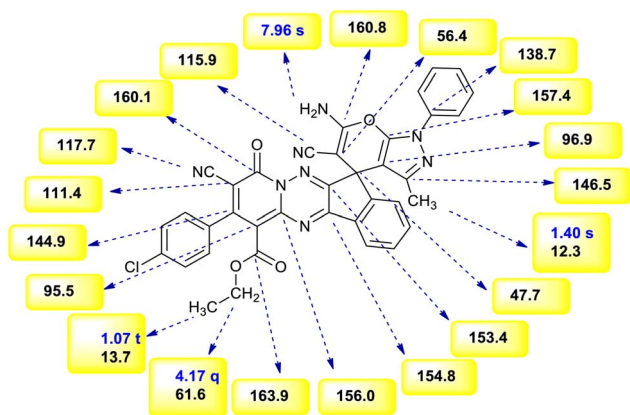
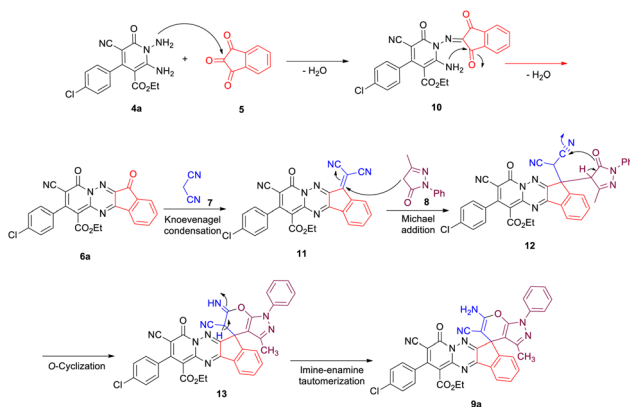


Fig. 3 ^1H and ^{13}C NMR chemical shifts of 9a.



Scheme 4 Proposed mechanism for the synthesis of products **9**.

1) as the eluent. The reaction mixture was cooled to room temperature. As a result, an orange solid was obtained. The solid was separated, washed with water (2×10 mL) and subsequently with Et₂O (2×10 mL) to give the pure pyrido[1,2,4]triazines products **6**.

Synthetic procedure of spiroindenopyridotriazine-4H-pyrans (9a–g)

Indenopyrido[1,2,4]triazine **6** (1 mmol), malononitrile **7** (1 mmol), pyrazolone **8** (1 mmol) and 10 mL EtOH were placed in a 25 mL round-bottomed flask mounted on a magnetic stirrer-heater. The reaction mixture was refluxed for 6 hours. After completion of the reaction as determined by TLC, the cooled insoluble crude product was collected by simple filtration and subsequently washed with warm ethanol to give the novel spiro-4H-pyrans **9** without the need for chromatography column or recrystallization.

Ethyl 1,6-diamino-4-(4-chlorophenyl)-3-cyano-2-oxo-1,2-dihydropyridine-5-carboxylate (4a). White solid; 87%; mp: 245–247 °C; IR (KBr): 3387, 3226, 2984, 2923, 2215, 1686, 1651, 1585, 1473, 1207, 1092, 839, 618 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 0.57 (3H, t, ³*J*_{HH} = 7.2 Hz, CH₃), 3.73 (2H, q, ³*J*_{HH} = 7.2 Hz, CH₂), 5.67 (2H, s, N-NH₂), 7.25 (2H, d, ³*J*_{HH} = 8.4 Hz, ArH), 7.49 (2H, d, ³*J*_{HH} = 8.4 Hz, ArH), 8.42 (1H, brs, NH₂), 8.72 (1H, brs, NH₂); ¹³C{¹H} NMR (75.4 MHz, DMSO-*d*₆): δ : 13.2 (CH₃), 60.5 (O-CH₂), 88.1 (C-CN), 91.4 (C-CO₂Et), 117.0 (CN), 128.4, 129.2, 133.4, 138.1 (Ar), 156.6 (C-4), 158.6 (C-NH₂), 159.2 (C=O), 166.1 (CO₂Et); MS (EI, 70 eV): *m/z* (%) = 334 (35) [M + 2]⁺, 333 (21) [M + 1]⁺, 332 (100) [M]⁺, 303 (5), 286 (13), 251 (13), 229 (21), 187 (5), 162 (12), 138 (7), 111 (4), 75 (4), 58 (3), 43 (5).

Ethyl 3-cyano-2-(4-nitrophenyl)-4,7-dioxo-4,7-dihydroindeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-1-carboxylate (6c). Orange solid; 85%; mp: 305–307 °C; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 1.04 (3H, t, ³*J*_{HH} = 6 Hz, CH₃), 4.13 (2H, q, ³*J*_{HH} = 6 Hz, CH₂), 7.83 (2H, d, ³*J*_{HH} = 9 Hz, ArH), 8.07–8.22 (4H, m, ArH), 8.45 (2H, d, ³*J*_{HH} = 9 Hz, ArH); ¹³C{¹H} NMR (125.7 MHz, DMSO-*d*₆): δ : 13.6 (CH₃), 61.8 (OCH₂), 99.0 (C-COO), 111.9 (C-CN), 115.0 (CN), 123.8, 124.8, 125.0, 129.5, 136.0, 137.0, 137.8, 140.2, 140.4 (Ar), 144.9 (C=N), 148.5 (C=N), 154.9 (C=Ar-NO₂), 155.8 (N-C-N), 157.9 (N-C=O), 163.9 (CO₂Et), 183.5 (C=O).

Ethyl 6'-amino-2-(4-chlorophenyl)-3,5'-dicyano-3'-methyl-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-7,4'-pyrano[2,3-*c*]pyrazole]-1-carboxylate (9a). Brown powder, 86%, mp: 253–255 °C; IR (KBr): 3365, 3196, 2960, 2199, 1727, 1682, 1594, 1525, 1463, 1399, 1092, 761 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 1.07 (3H, t, ³*J*_{HH} = 6.0 Hz, CH₃CH₂), 1.40 (3H, s, Me), 4.18 (2H, q, ³*J*_{HH} = 6.0 Hz, CH₂), 7.40–7.87 (11H, m, ArH), 7.96 (2H, s, NH₂), 8.00 (1H, d, ³*J*_{HH} = 6.0 Hz, ArH), 8.15 (1H, d, ³*J*_{HH} = 6.0 Hz, ArH); ¹³C{¹H} NMR (75.4 MHz, DMSO-*d*₆): δ : 12.3 (CH₃), 13.7 (CH₃CH₂), 47.7 (C_{Spiro}), 56.4 (C=CNH₂), 61.6 (OCH₂), 95.5 (C-COO), 96.9 (C=C-O), 111.4 (C-CN), 115.9 (CN), 117.7 (CN), 120.5, 125.3, 125.6, 126.9, 128.9, 129.5, 129.7, 131.4, 133.5, 135.0, 137.1, 138.7, 144.4 (Ar), 144.9 (C-C₆H₅Cl), 146.5 (Me-C=N), 153.4 (C=N-N), 154.8 (C=N), 156.0 (N-C-N), 157.4 (O-C-N), 160.1 (N-C=O), 160.8 (CNH₂), 163.9 (CO₂Et); MS (EI, 70 eV): *m/z* (%) = 649

(0.6) [M - 29]⁺, 621 (0.1), 553 (0.4), 523 (0.5), 495 (2), 461 (2), 409 (4), 352 (4), 324 (3), 293 (4), 239 (14), 189 (28), 137 (18), 109 (81), 81 (52), 43 (100), 29 (22).

Ethyl 6'-amino-2-(4-bromophenyl)-3,5'-dicyano-3'-methyl-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-7,4'-pyrano[2,3-*c*]pyrazole]-1-carboxylate (9b). Brown powder, 82%, mp: 266–268 °C; IR (KBr): 3347, 3195, 2960, 2199, 1732, 1683, 1591, 1525, 1461, 1399, 1265, 1071, 754 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 1.07 (3H, t, ³*J*_{HH} = 6.0 Hz, CH₃CH₂), 1.41 (3H, s, Me), 4.17 (2H, m, CH₂), 7.37–7.58 (5H, m, ArH), 7.77–7.87 (6H, m, ArH), 7.97 (2H, s, NH₂), 7.98–8.01 (1H, m, ArH), 8.15 (1H, d, ³*J*_{HH} = 9.0 Hz, ArH); ¹³C{¹H} NMR (75.4 MHz, DMSO-*d*₆): δ : 12.3 (CH₃), 13.7 (CH₃CH₂), 47.7 (C_{Spiro}), 56.4 (C=CNH₂), 61.7 (OCH₂), 95.5 (C-COO), 96.8 (C=C-O), 111.3 (C-CN), 115.9 (CN), 117.7 (CN), 120.5, 123.8, 124.2, 126.8, 129.5, 129.9, 131.4, 131.9, 133.6, 133.8, 137.1, 138.2, 144.4 (Ar), 144.9 (C-C₆H₅Br), 146.5 (Me-C=N), 153.4 (C=N-N), 154.8 (C=N), 156.0 (N-C-N), 157.4 (O-C-N), 160.1 (N-C=O), 160.8 (CNH₂), 163.9 (CO₂Et); *m/z* (%) = 647 (0.03), 503 (0.01), 459 (0.03), 414 (0.4), 399 (0.1), 355 (0.4), 315 (4), 269 (2), 224 (13), 178 (15), 135 (57), 91 (100), 43 (60), 15 (6).

Ethyl 6'-amino-3,5'-dicyano-3'-methyl-2-(4-nitrophenyl)-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-7,4'-pyrano[2,3-*c*]pyrazole]-1-carboxylate (9c). Red brown powder, 90%, mp: 260–262 °C; IR (KBr): 3361, 3193, 2960, 2204, 1727, 1688, 1597, 1524, 1462, 1348, 1127, 1073, 755 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 1.07 (3H, m, CH₃CH₂), 1.42 (3H, s, Me), 4.15 (2H, m, CH₂), 7.40–7.58 (3H, m, ArH), 7.75–7.87 (7H, m, ArH), 7.99 (2H, s, NH₂), 8.15 (1H, d, ³*J*_{HH} = 9.0 Hz, ArH), 8.44–8.47 (2H, m, ArH); ¹³C{¹H} NMR (75.4 MHz, DMSO-*d*₆): δ : 12.3 (CH₃), 13.7 (CH₃CH₂), 47.7 (C_{Spiro}), 56.3 (C=CNH₂), 61.8 (OCH₂), 95.5 (C-COO), 96.7 (C=C-O), 111.0 (C-CN), 115.7 (CN), 117.8 (CN), 120.5, 124.0, 124.3, 126.9, 129.5, 131.4, 131.5, 137.1, 137.8, 138.4, 141.4, 144.4 (Ar), 145.0 (C-C₆H₅NO₂), 146.9 (Me-C=N), 148.4 (C-NO₂), 153.5 (C=N-N), 154.0 (C=N), 156.0 (N-C-N), 157.6 (O-C-N), 160.3 (N-C=O), 160.9 (CNH₂), 163.7 (CO₂Et); MS (EI, 70 eV): *m/z* (%) = 578 (0.08), 551 (0.2), 523 (0.2), 495 (0.1), 467 (0.09), 416 (0.8), 368 (1), 343 (2), 313 (1), 287 (11), 261 (29), 230 (48), 204 (56), 174 (9), 127 (52), 78 (81), 43 (100), 18 (38).

Ethyl 6'-amino-3,5'-dicyano-2-(4-fluorophenyl)-3'-methyl-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-*e*]pyrido[1,2-*b*][1,2,4]triazine-7,4'-pyrano[2,3-*c*]pyrazole]-1-carboxylate (9d). Orange powder, 78%, mp: 275–278 °C; IR (KBr): 3301, 3188, 2965, 2195, 1730, 1684, 1599, 1514, 1463, 1400, 1333, 1161, 1074, 764 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆): δ : 1.06 (3H, t, ³*J*_{HH} = 6.0 Hz, CH₃CH₂), 1.41 (3H, s, Me), 4.14 (2H, q, ³*J*_{HH} = 6.0 Hz, CH₂), 7.37–7.58 (7H, m, ArH), 7.77–7.87 (5H, m, ArH), 7.97 (2H, s, NH₂), 8.15 (1H, d, ³*J*_{HH} = 9.0 Hz, ArH); ¹³C{¹H} NMR (75.4 MHz, DMSO-*d*₆): δ : 12.3 (CH₃), 13.7 (CH₃CH₂), 18.62 (EtOH), 47.7 (C_{Spiro}), 56.09 (EtOH), 56.5 (H₂N-C=C-CN), 61.6 (OCH₂), 95.5 (C-COO), 97.1 (C=C-O), 111.6 (C-CN), 115.8 (CN), 116.1 (d, ²*J*_{CF} = 21.8 Hz), 117.8 (CN), 120.4, 124.2, 126.8, 126.9, 129.5, 130.3 (d, ³*J*_{CF} = 8.3 Hz), 130.4 (d, ⁴*J*_{CF} = 6.0 Hz), 131.4, 137.1, 138.2, 144.4 (Ar), 145.0 (C-C₆H₅F), 146.4 (Me-C=N), 153.4 (C=N-N), 155.0 (C=N), 156.1 (N-C-N), 157.4 (O-C-N), 160.1 (N-C=O), 160.8 (CNH₂), 161.4 (d, ¹*J*_{CF} = 240.5 Hz), 164.0 (CO₂Et); MS (EI,



70 eV): m/z (%) = 596 (0.06), 550 (0.1), 523 (0.1), 488 (0.4), 443 (0.2), 416 (0.6), 388 (0.4), 346 (0.2), 313 (0.1), 301 (3), 255 (4), 212 (1), 174 (21), 145 (7), 105 (16), 77 (100), 51 (42), 28 (23).

Ethyl 6'-amino-3,5'-dicyano-2-(4-methoxyphenyl)-3'-methyl-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-e]pyrido[1,2-b][1,2,4]triazine-7,4'-pyrano[2,3-c]pyrazole]-1-carboxylate (9e). Dark brown powder, 80%, mp: 295–298 °C; IR (KBr): 3365, 3196, 2960, 2201, 1724, 1655, 1463, 1250, 1025, 762 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ : 1.10 (3H, t, $^3J_{\text{HH}} = 9.0$ Hz, CH_3CH_2), 1.40 (3H, s, Me), 3.84 (3H, s, OCH_3), 4.18 (2H, q, $^3J_{\text{HH}} = 9.0$ Hz, CH_2), 7.12–7.15 (3H, m, ArH), 7.40–7.43 (2H, m, ArH), 7.53–7.58 (3H, m, ArH), 7.79–7.87 (4H, m, ArH), 7.90 (2H, s, NH_2), 8.13 (1H, d, $^3J_{\text{HH}} = 9.0$ Hz, ArH); $^{13}\text{C}\{^1\text{H}\}$ NMR (125.7 MHz, $\text{DMSO}-d_6$) δ : 12.2 (CH_3), 13.7 (CH_3CH_2), 47.6 (C_{Spiro}), 55.3 (OCH_3), 56.6 ($\text{C}=\text{CNH}_2$), 61.4 (OCH_2), 95.5 ($\text{C}-\text{COO}$), 96.9 ($\text{C}=\text{C}-\text{O}$), 111.6 ($\text{C}-\text{CN}$), 114.1 (CN), 116.1 (CN), 114.6, 120.4, 124.0, 126.5, 126.7, 126.8, 129.4, 131.3, 131.4, 137.0, 138.0, 144.3, 144.9 (Ar), 145.9 ($\text{Me}-\text{C}=\text{N}$), 153.2 ($\text{C}=\text{N}-\text{N}$), 155.7 ($\text{C}=\text{N}$), 156.1 ($\text{N}-\text{C}-\text{N}$), 157.1 ($\text{O}-\text{C}-\text{N}$), 159.7 ($\text{C}_{\text{Ar}}-\text{OMe}$), 160.6 ($\text{N}-\text{C}=\text{O}$), 160.7 (CNH_2), 164.1 (CO_2Et).

Ethyl 6'-amino-2-(3-chlorophenyl)-3,5'-dicyano-3'-methyl-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-e]pyrido[1,2-b][1,2,4]triazine-7,4'-pyrano[2,3-c]pyrazole]-1-carboxylate (9f). Brown powder, 82%, mp: 245–248 °C; IR (KBr): 3192, 2198, 1738, 1683, 1462, 1126, 1017, 762 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ : 1.05 (3H, t, $^3J_{\text{HH}} = 6.0$ Hz, CH_3CH_2), 1.41 (3H, s, Me), 4.17 (2H, q, $^3J_{\text{HH}} = 6.0$ Hz, CH_2), 7.39–7.69 (9H, m, ArH), 7.80–7.87 (3H, m, ArH), 7.98 (2H, s, NH_2), 8.15 (1H, d, $^3J_{\text{HH}} = 9.0$ Hz, ArH); $^{13}\text{C}\{^1\text{H}\}$ NMR (75.4 MHz, $\text{DMSO}-d_6$) δ : 12.2 (CH_3), 13.7 (CH_3CH_2), 47.7 (C_{Spiro}), 56.5 ($\text{C}=\text{CNH}_2$), 61.6 (OCH_2), 95.5 ($\text{C}-\text{COO}$), 97.0 ($\text{C}=\text{C}-\text{O}$), 111.3 ($\text{C}-\text{CN}$), 115.7 (CN), 117.7 (CN), 120.5, 124.2, 126.6, 126.8, 127.5, 129.5, 130.0, 130.8, 131.3, 133.3, 136.5, 137.1, 138.3, 144.4 (Ar), 144.9 ($\text{C}-\text{C}_6\text{H}_5\text{Cl}$), 146.5 ($\text{Me}-\text{C}=\text{N}$), 153.4 ($\text{C}=\text{N}-\text{N}$), 154.3 ($\text{C}=\text{N}$), 156.0 ($\text{N}-\text{C}-\text{N}$), 157.4 ($\text{O}-\text{C}-\text{N}$), 160.2 ($\text{N}-\text{C}=\text{O}$), 160.8 (CNH_2), 163.8 (CO_2Et).

Ethyl 6'-amino-3,5'-dicyano-3'-methyl-2-(3-nitrophenyl)-4-oxo-1'-phenyl-1'H,4H-spiro[indeno[1,2-e]pyrido[1,2-b][1,2,4]triazine-7,4'-pyrano[2,3-c]pyrazole]-1-carboxylate (9g). Brown powder, 85%, mp: 283–285 °C; IR (KBr): 3351, 3196, 2223, 1732, 1686, 1532, 1464, 1348, 1125, 1019, 755 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ : 1.00–1.07 (3H, m, CH_3CH_2), 1.41 (3H, s, Me), 4.13–4.17 (2H, m, CH_2), 7.40–7.42 (1H, m, ArH), 7.55 (1H, d, $^3J_{\text{HH}} = 6.0$ Hz, ArH), 7.81–7.93 (5H, m, ArH), 7.98 (2H, s, NH_2), 7.99–8.03 (2H, m, ArH), 8.18 (1H, d, $^3J_{\text{HH}} = 9.0$ Hz, ArH), 8.30–8.47 (3H, m, ArH); $^{13}\text{C}\{^1\text{H}\}$ NMR (75.4 MHz, $\text{DMSO}-d_6$) δ : 12.3 (CH_3), 13.8 (CH_3CH_2), 47.7 (C_{Spiro}), 56.5 ($\text{H}_2\text{N}-\text{C}=\text{C}-\text{CN}$), 61.6 (OCH_2), 95.6 ($\text{C}-\text{COO}$), 97.1 ($\text{C}=\text{C}-\text{O}$), 111.7 ($\text{C}-\text{CN}$), 115.8 (CN), 116.1, 117.8 (CN), 120.5, 124.2, 126.7, 126.9, 129.6, 130.3, 130.4, 131.4, 137.1, 138.3, 144.5, 145.0 (Ar), 146.4 ($\text{Me}-\text{C}=\text{N}$), 153.4 ($\text{C}=\text{N}-\text{N}$), 155.1 ($\text{C}=\text{N}$), 156.1 ($\text{N}-\text{C}-\text{N}$), 157.4 ($\text{O}-\text{C}-\text{N}$), 160.1 ($\text{N}-\text{C}=\text{O}$), 160.8 (CNH_2), 161.4, 164.0 (CO_2Et).

Conclusion

In summary, we have developed an attractive and green approach for the synthesis of novel highly functionalized fused spiro-2-amino-4H-pyrans by sequential multicomponent

reaction between pyridotriazines, pyrazolone as cyclic CH-acid and malononitrile in ethanol under mild conditions. To reach the target products, 1,6-diaminopyridine-2-one compounds were firstly prepared, then their reaction was done with ninhydrin. As a result of this condensation, pyridotriazine derivatives were synthesized as starting material for the main reaction. These two processes were carried out as a four-component cascade reaction. This method not only provides high efficiency and reaction speed, but also prevent the use of dangerous solvents or catalysts. Other promising points for the introduced synthesis include: clean reaction profile, ease of product purification, good to high yields and high atom economy.

Data availability

The data supporting this article are provided in the ESI.†

Conflicts of interest

The authors declare no competing financial interest.

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Notes and references

- Q. Guan, L. L. Zhou and Y. B. Dong, *J. Am. Chem. Soc.*, 2023, **145**, 1475.
- S. E. John, S. Gulati and N. Shankaraiah, *Org. Chem. Front.*, 2021, **8**, 4237.
- Y. E. Ryzhkova, M. N. Elinson, O. I. Maslov and A. N. Fakhrutdinov, *Molecules*, 2021, **26**, 6839.
- A. Shaabani, R. Mohammadian, R. Afshari, S. E. Hooshmand, M. T. Nazeri and S. Javanbakht, *Mol. Divers.*, 2021, **25**, 1145.
- J. D. Sunderhaus and S. F. Martin, *Chem. Eur. J.*, 2009, **15**, 1300.
- M. O. Rodrigues, M. N. Eberlin and B. A. D. Neto, *Chem. Rec.*, 2021, **10**, 2762.
- L. R. Wen, Z. R. Li, M. Li and H. Cao, *Green Chem.*, 2012, **14**, 707.
- M. Karami, A. Hasaninejad, H. Mahdavi, A. Iraj, S. Mojtavavi, M. A. Faramarzi and M. Mahdavi, *Mol. Divers.*, 2021, **26**, 2393.
- C. Nicolaou, D. J. Edmonds and P. G. Bulger, *Angew. Chem. Int. Ed.*, 2006, **45**, 7134.
- S. Sharma, V. Vaishali, N. Devi and V. Singh, *Chem.-Asian J.*, 2024, **20**, e202400862.
- D. Singh, S. Sharma, R. K. Thakur, V. Vaishali, S. Nain, Jyoti, C. C. Malakar and V. Singh, *Tetrahedron*, 2024, **152**, 133809.
- S. Sharma, C. C. Malakar and V. Singh, *Chem.-Asian J.*, 2020, **9**, 1857.



- 13 S. Sharma, D. Singh, S. Kumar, V. Vaishali, R. Jamra, N. Banyal, K. Deepika, C. C. Malakar and V. Singh, *Beilstein J. Org. Chem.*, 2023, **19**, 231.
- 14 S. S. Mritunjay, P. Singh, A. Gupta, V. Singh, L. Singh and A. Kumar, *ChemistrySelect*, 2024, **9**, e202402133.
- 15 V. F. Batista, D. C. Pinto and A. M. S. Silva, *Expert Opin. Drug Discovery*, 2022, **17**, 603.
- 16 K. Hiesinger, D. Darin, E. Proschak and M. Krasavin, *J. Med. Chem.*, 2021, **64**, 150.
- 17 Y. J. Zheng and C. M. Tice, *Expert Opin. Drug Discovery*, 2016, **11**, 831.
- 18 Y. Zheng, C. M. Tice and S. B. Singh, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 3673.
- 19 L. Emami, L. Moezi, L. Amiri-Zirtol, *et al.*, *Mol. Divers.*, 2022, **26**, 3129.
- 20 M. Saigal, P. Irfan, M. Khana and M. M. Khan, *ACS Omega*, 2019, **4**, 16794.
- 21 D. C. Wang, Y. M. Xie and C. Fan, *Chin. Chem. Lett.*, 2014, **25**, 1011.
- 22 R. X. Liu, Y. N. Liang, X. X. Ren, Q. Q. Wu, *et al.*, *Curr. Org. Synth.*, 2023, **20**, 870.
- 23 F. Safari, H. Hosseini, M. Bayat and A. Ranjbar, *RSC Adv.*, 2019, **9**, 24843.
- 24 Z. Sadeghian, M. Bayat and F. Safari, *J. Mol. Struct.*, 2022, **1250**, 131759.
- 25 M. Valko, D. Leibfritz, J. Moncol, M. T. D. Cronin, M. Mazur and J. Telser, *Int. J. Biochem. Cell Biol.*, 2007, **39**, 44.
- 26 K. Acosta-Quiroga, C. Rojas-Peña, L. S. Nerio, M. Gutiérrez and E. Polo-Cuadrado, *RSC Adv.*, 2021, **11**, 21926.
- 27 L. Jalili-Baleh, N. Mohammadi, M. Khoobi, L. Mamani, A. Foroumadi and A. Shafiee, *Helv. Chim. Acta*, 2013, **96**, 1601.
- 28 P. Saluja, K. Aggarwal and J. M. Khurana, *Synth. Commun.*, 2013, **43**, 3239.
- 29 L. Mohammadi, M. M. Heravi, A. Saljooqi and P. Mohammadi, *Sci. Rep.*, 2022, **12**, 22281.
- 30 C. B. Li, L. S. Huang, R. S. Wu and D. Z. Xu, *ChemistrySelect*, 2019, **4**, 1635.
- 31 M. M. Li, C. S. Duan, Y. Q. Yu and D. Z. Xu, *Dyes Pigm.*, 2018, **150**, 202.
- 32 K. Nikoofar and S. M. Dizgarani, *Monatsh. Chem.*, 2015, **8**, 1161.
- 33 S. Mal and M. Jana, *Arkivoc*, 2022, **2022**, 361.
- 34 D. Suven, *RSC Adv.*, 2020, **10**, 18875.
- 35 R. Singh, D. Bhardwaj and M. R. Saini, *RSC Adv.*, 2021, **11**, 4760.
- 36 P. Khanna, L. Khanna, S. J. Thomas, A. M. Asiri and S. S. Panda, *Curr. Org. Chem.*, 2018, **22**, 67.
- 37 S. F. Hojati, A. Amiri and E. Fardi, *Appl. Organomet. Chem.*, 2020, **34**, 1.
- 38 M. R. Poor Heravi and F. Norouzy, *Res. Chem. Intermed.*, 2017, **43**, 4265.
- 39 G. Mohammadi Ziarani, Z. Kheilkordi and F. Mohajer, *J. Iran. Chem. Soc.*, 2020, **17**, 247.
- 40 N. H. Nasab and J. Safari, *Polyhedron*, 2019, **164**, 74.
- 41 Z. Sadeghian and M. Bayat, *Med. Chem. Res.*, 2022, **31**, 497.
- 42 Z. Sadeghian, M. Bayat, M. Eskandari and F. Safari, *Results Chem.*, 2024, **9**, 101636.
- 43 M. Rezaei and M. Bayat, *RSC Adv.*, 2023, **13**, 31488.
- 44 D. M. Patel, P. J. Patel and H. M. Patel, *Eur. J. Org. Chem.*, 2022, **46**, e202201119.
- 45 H. Hosseini and M. Bayat, *RSC Adv.*, 2018, **8**, 27131.
- 46 M. Shokoohian, N. Hazeri, M. T. Maghsoodlou and M. Lashkari, *Polycyclic Aromat. Compd.*, 2022, **42**, 1.

