

# Organocatalytic Discrimination of Non-Directing Aryl and Heteroaryl Groups: Enantioselective Synthesis of Bioactive Indole-Containing Triarylmethanes

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**Abstract:** Despite the enormous developments of asymmetric catalysis, the basis for asymmetric induction is largely limited to spatial interaction between substrate and catalyst. Consequently, asymmetric discrimination between two sterically similar groups remains a challenge. This is particularly formidable for enantiodifferentiation between aryl and heteroaryl groups without a directing group or electronic manipulation. Here we address this challenge by a robust organocatalytic system leading to excellent enantioselection between aryl and heteroaryl groups. With the versatile 2-indole imine methide as platform, an excellent combination of a superb chiral phosphoric acid and the optimal hydride source provided

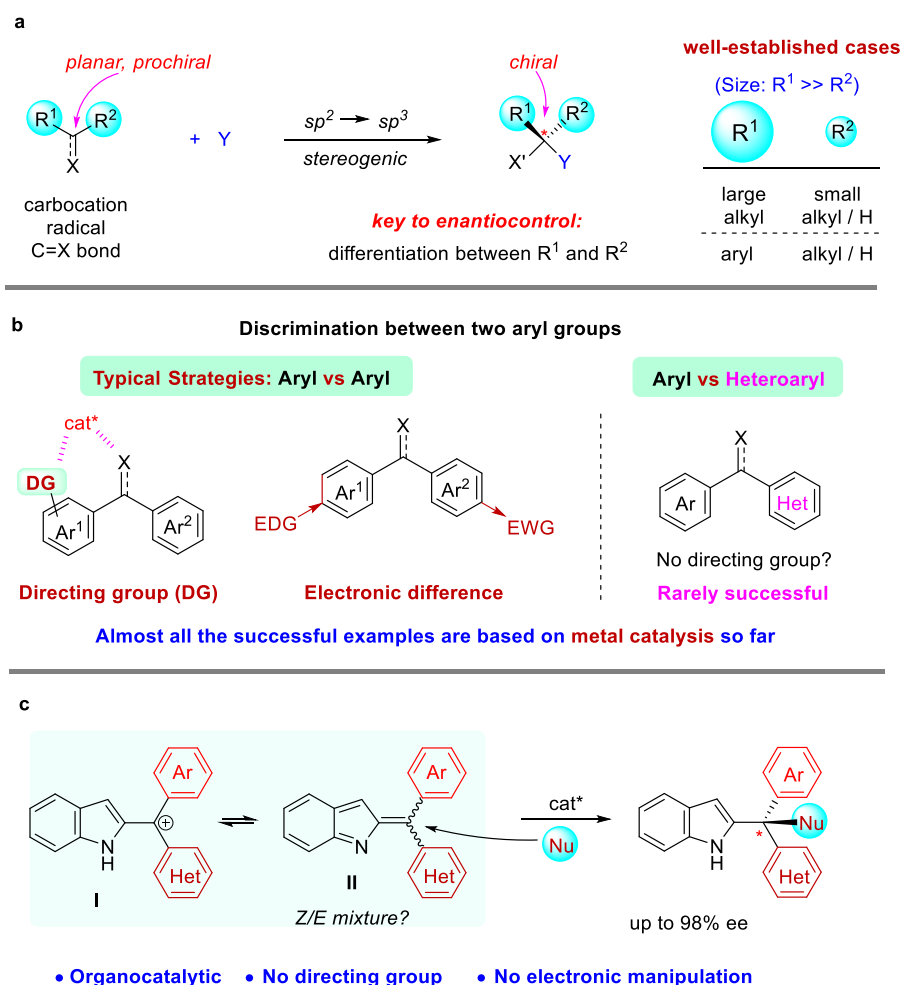
**efficient access to a range of highly enantioenriched indole-containing triarylmethanes. Control experiments and kinetic studies provided important insights into the mechanism. DFT calculations also indicated that, while hydrogen bonding is important for activation, the key interaction for discrimination of the two aryl groups is mainly  $\pi$ - $\pi$  stacking. Preliminary biological studies also demonstrated the great potential of these triarylmethanes for anticancer and antiviral drug development.**

Asymmetric catalysis has evolved arguably into the most powerful method for the synthesis of enantioenriched molecules.<sup>1</sup> It features high efficiency and atom-economy in principle as compared to other approaches such as chiral resolution and auxiliary-based asymmetric synthesis, thereby enabling increasing applications in industrial synthesis.<sup>2</sup> In the past few decades, a wide range of chiral catalytic systems with diverse activation modes have been developed. However, the fundamental basis for enantiocontrol remains essentially unchanged, i.e., spatial interaction between substrate and catalyst is still the key to success.<sup>1, 2</sup> For example, in the construction of a tetrahedral  $C(sp^3)$ -chiral center from a prochiral  $C(sp^2)$ -based planar substrate (e.g., carbocation, radical, carbonyl, olefin), a chiral catalyst typically provides enantiodifferentiation by blocking one face of the plane and directing the reaction partner (Y) to approach towards the other face (Fig 1a). To achieve this, the catalyst must be able to effectively discriminate between the two substituents (R1 and R2) on the prochiral carbon. Obviously, the larger the difference of these two substituents is, the better enantioselectivity will be expected. Consequently, it has been well-established to achieve high enantioselectivity for cases bearing two sterically different groups (e.g., alkyl/aryl vs. H, large alkyl vs. small alkyl). In contrast, cases bearing two substituents of similar size remain challenging.<sup>1</sup>

1,1-Diarylmethinyl stereocenters is a widely prevalent structural motif in various natural products and biologically important molecules.<sup>3-4</sup> Asymmetric addition to the 1,1-diaryl  $C=C$  and  $C=X$  (X = heteroatom) bonds represents one of the most direct approaches for the construction of this unit.<sup>5-21</sup> However, this requires effective discrimination between two (often) sterically similar aryl groups, which represents a notable challenge in asymmetric catalysis (Fig 1b).<sup>5</sup> So far, success has mainly relied on the use of a directing group<sup>6-14</sup> in one aryl group to allow catalyst recognition (e.g., by coordination) or electronic difference by incorporating electron-donating/withdrawing groups<sup>15, 16</sup>.

Notably, the effective enantiodifferentiation between aryl and heteroaryl groups still remains challenging, particularly in the absence of a directing group or electronic manipulation.<sup>17-21</sup> Currently, successful differentiation between aryl and heteroaryl mostly involves a pyridine heterocycle due to its special coordinating ability.<sup>17-21</sup> Moreover, despite the above-mentioned important progress, it is worth noting that almost all these examples relied on metal catalysis, and little success has been achieved by organocatalysis. In this context, here we describe an organocatalytic discrimination of non-directing aryl and heteroaryl groups with excellent enantioselectivity.

Indole imine methides (IIMs) have recently emerged as versatile intermediates for the asymmetric synthesis of enantioenriched indole derivatives, a family of useful units in medicinal chemistry.<sup>22-31</sup> In particular, those with the methide motif adorned in the 2-position of indole are particularly useful to construct indole-fused polyheterocycles via asymmetric annulation processes, as pioneered by Shi and co-workers.<sup>22-28</sup> However, the majority of these examples either involve straightforward discrimination (e.g., aryl vs. H) or do not involve asymmetric control in the 2-benzylic position (e.g., with two same substituents).<sup>22-31</sup> In continuation of our efforts in this area,<sup>14, 29-31</sup> we envisioned that this type of intermediates would be a good platform to study the power of organocatalysis for the challenging discrimination between aryl and heteroaryl groups lacking a directing group (Scheme 1c). However, additional challenges should be expected since this intermediate **II** is likely generated as a *Z/E* mixture, typically in equilibrium with carbocation **I**. Therefore, the equilibrium should be made in synergy with the nucleophilic addition step to allow dynamic asymmetric control in order to achieve high enantioselectivity.



**Fig. 1. Introduction to asymmetric differentiation in  $C(sp^2)$ -prochiral centers.** **a** Asymmetric differentiation in prochiral planar substrates. **b** Challenges in discrimination between two aryl groups. **c** Our reaction design and the challenges.

## Results

**Reaction development.** To test our hypothesis, we employed racemic tertiary alcohol **1a** as the model precursor to the 2-indole imine methide intermediate. Notably, no directing group is incorporated in the two aryl groups (phenyl and thienyl) to be discriminated by catalyst. Despite the above-mentioned substantial challenges on this asymmetric control, considerable efforts were devoted to condition optimization and ultimately led to excellent reaction efficiency and enantiocontrol (Table 1). Specifically, among the broad range of chiral acid catalysts evaluated, the SPINOL-derived chiral phosphoric acid **C1** was identified as the optimal catalyst.<sup>32-37</sup> With benzothiazoline **2a** as the hydride source,<sup>38,30</sup> the asymmetric reduction proceeded smoothly to form indole-containing triarylmethane **3a** under mild conditions in essentially quantitative yield and 95%

ee (entry 1, Table 1). For comparison, other conditions typically led to inferior results. For example, other SPINOL-based chiral phosphoric acids gave lower enantioselectivity (entries 2-5). In particular, the previously well-known STRIP catalyst **C2** resulted in only 16% ee (entry 2). In addition, the catalyst chiral backbones were compared with the same 3,5-bis(trifluoromethyl)phenyl substituent in the 3,3'-positions (entries 3-5). While the SPINOL-based catalyst **C3** gave 81% ee, in sharp contrast, the BINOL- and [H<sub>8</sub>]BINOL-derived analogues **A** and **B** did not show any asymmetric induction (<2% ee). This result not only highlighted the superiority of the spirocyclic skeleton, but also corroborated the elusive stereocontrol in this case. Other hydride sources were also examined (entries 6-8). Benzothiazolines **2b** and **2c**, bearing a different aryl substituent, led to drastically low chemoselectivity and enantioselectivity (entries 6-7). More surprisingly, Hantzsch ester **2d**, the previously well-established hydride source,<sup>39</sup> gave drastically low enantiocontrol (entry 8). Other solvents than DCM did not provide a better result either (entries 9-11). The reaction was very sensitive to coordinating solvents, such as ether and ethyl acetate, which completely shut down the reaction, presumably due to competing binding with the acid catalyst. Decreasing the reaction temperature to 0 °C maintained high enantioselectivity, but moderately affected the reaction rate (entry 12). Finally, at a higher concentration, slightly lower enantioselectivity was observed (entry 13).

**Table 1.** Evaluation of reaction conditions.<sup>a</sup>

**1a (racemic)** + **2a** (1.1 equiv)  $\xrightarrow[\text{DCM (0.05 M), RT, 12 h}]{(R)\text{-C1 (5 mol\%)}}$  **3a**

"standard conditions"

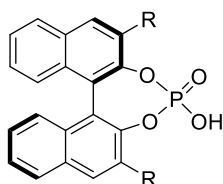
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Ar = 2-naphthyl (**2a**)  
Ar = 1-naphthyl (**2b**)  
Ar = 9-phenanthracenyl (**2c**)

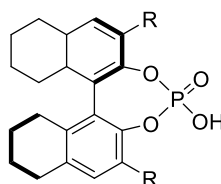
**2d**

Entry	Deviation from the "standard conditions"	Yield (%) <sup>b</sup>	ee(%) <sup>b</sup>
1	none	> 95	95
2	( <i>R</i> )- <b>C2</b> instead of ( <i>R</i> )- <b>C1</b>	> 95	16
3	( <i>R</i> )- <b>C3</b> instead of ( <i>R</i> )- <b>C1</b>	> 95	81
4	( <i>R</i> )- <b>A</b> instead of ( <i>R</i> )- <b>C1</b>	> 95	< 2
5	( <i>R</i> )- <b>B</b> instead of ( <i>R</i> )- <b>C1</b>	> 95	< 2

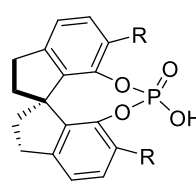
6	<b>2b</b> instead of <b>2a</b>	11 <sup>c</sup>	80
7	<b>2c</b> instead of <b>2a</b>	15 <sup>c</sup>	55
8	<b>2d</b> instead of <b>2a</b>	78 <sup>d</sup>	−9
9	Et <sub>2</sub> O as solvent	< 5 <sup>e</sup>	—
10	Toluene as solvent	87	89
11	EtOAc as solvent	< 5 <sup>e</sup>	—
12	Run at 0 °C	84 <sup>d</sup>	96
13	<i>c</i> = 0.2 M	> 95	93



(*R*)-**A**  
R = 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>



(*R*)-**B**  
R = 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>



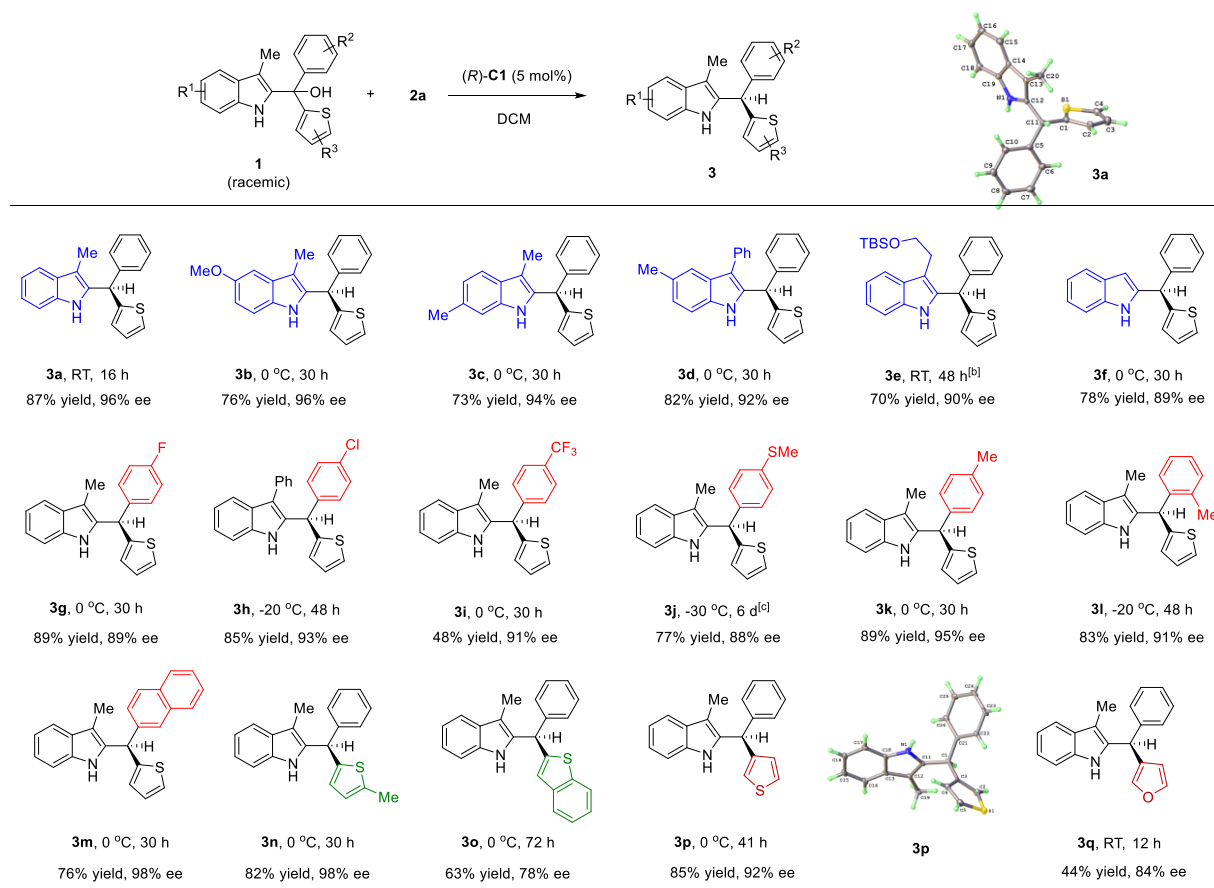
(*R*)-**C1**: R = 3,5-(<sup>t</sup>Bu)<sub>2</sub>-4-(OMe)C<sub>6</sub>H<sub>2</sub>  
(*R*)-**C2**: R = 2,4,6-(<sup>i</sup>Pr)<sub>3</sub>C<sub>6</sub>H<sub>2</sub>  
(*R*)-**C3**: R = 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>

<sup>a</sup>Reaction conditions: **1a** (25 μmol), hydride source (27.5 μmol), catalyst (2.5 μmol), solvent (0.5 mL). <sup>b</sup>Yield was determined by analysis of the <sup>1</sup>H NMR spectrum of the crude reaction mixture with CH<sub>2</sub>Br<sub>2</sub> as internal standard. Ee was determined by HPLC analysis on a chiral stationary phase. <sup>c</sup>A mixture of unidentifiable products was formed. <sup>d</sup>Clean conversion. The starting material accounts for the remainder of the mass balance. <sup>e</sup>Conversion < 5%.

**Substrates scope exploration.** With the optimized conditions, we examined the reaction scope with variously substituted indole-derived tertiary alcohol substrates (Table 2). In general, this protocol provided efficient access to a wide range of highly enantioenriched indole-containing triarylmethanes. Substrates bearing electron-withdrawing and electron-donating groups at different positions were all suitable. The presence of a substituent at the 3-position of indole is not necessary (**3f**), although this position is nucleophilic and can potentially serve as a competitive intermolecular nucleophile. In addition to substitution at the 2-position of the thiophene ring in most cases, it is worth noting that substitution at the 3-position provided equally high enantioselectivity (**3p**). Finally, it is worth noting that, other than those thiophene-containing examples, the discrimination between benzene and furan ring is also possible, leading to good enantiocontrol (**3q**). In all these cases, no directing group to provide additional interaction (e.g., hydrogen bonding) with catalyst is needed in order to achieve high enantiocontrol. Steric perturbation on enantioselectivity by increasing the size of either of the two aryl groups (e.g., **3l-m**) was not obvious. Notably, the structure and absolute

configuration of products **3a** and **3p** were confirmed by X-ray crystallography.

**Table 2.** Reaction scope.<sup>a</sup>

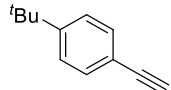
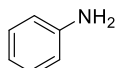

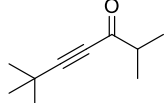
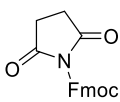
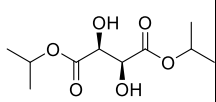
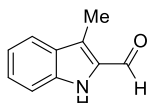
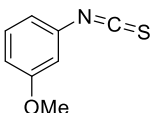
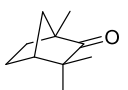
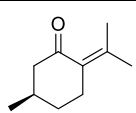


<sup>a</sup>Reaction scale: **1** (0.4 mmol), **2** (0.44 mmol), (*R*)-**C1** (5 mol%), DCM (8.0 mL). <sup>b</sup>Run with 10 mol% of catalyst.

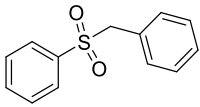
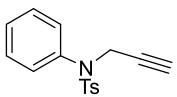
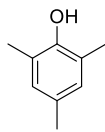
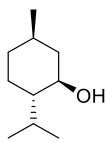
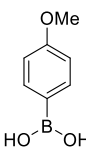
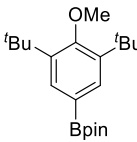
The robustness of this protocol was further examined by stoichiometric adulteration of various additives bearing different functional groups.<sup>40</sup> As shown in Table 3, in most cases, the excellent chemical efficiency and enantioselectivity were not obviously affected by the additives in the standard reaction of **1a** and **2a**. Many of these additives contain highly polar and reactive functionalities that are typical strong hydrogen-bonding partners, such as primary amine, thiol, alcohol, carbonyl, sulfone, and boronic acid. This is particularly remarkable in view of the fact that hydrogen bonding is a key catalyst-substrate interaction in the process. Notably, from a different point of view, the little influence on enantiocontrol by polar additives might also imply that it is not hydrogen bonding, but other interactions such as  $\pi$ - $\pi$  stacking, that provide the basis for asymmetric

discrimination (*vide infra*). Nevertheless, these results clearly illustrated the excellent functional group tolerance as well as the robust enantiodifferentiation ability of this mild but powerful catalytic system.

**Table 3.** Compatibility of external functional groups.<sup>a</sup>

$\mathbf{1a} + \mathbf{2a} + \text{additive} \xrightarrow[\text{DCM, rt, 12 h}]{\text{standard conditions}} \mathbf{3a}$					
Entry	Additive	Recovered additive (%)	Conversion of <b>1a</b> (%)	Yield of <b>3a</b> (%)	Ee of <b>3a</b> (%)
1		90	100	95	91
2		82	70	49	94
3		95	100	88	96
4		85	100	99	94
5		95	100	87	94
6	Ph <sub>3</sub> CSH	66	100	83	89
7		61	100	77	87
8		67	100	73	94
9		78	100	64	90
10		83	100	75	94
11	PPh <sub>3</sub>	80	100	95	94
12		67	100	57	94

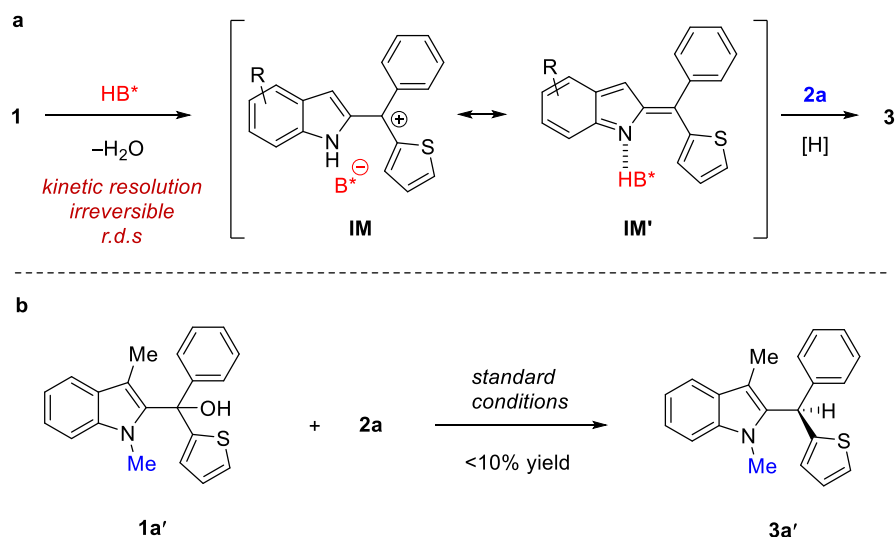


13		89	100	98	94
14		90	100	92	95
15		87	100	80	94
16		80	100	86	94
17		50	100	85	92
18		78	100	73	94

<sup>a</sup>Reaction scale: **1a** (25  $\mu$ mol, **2a** (27.5  $\mu$ mol), additive (25  $\mu$ mol), (*R*)-**C1** (5 mol%), DCM (0.5 mL).

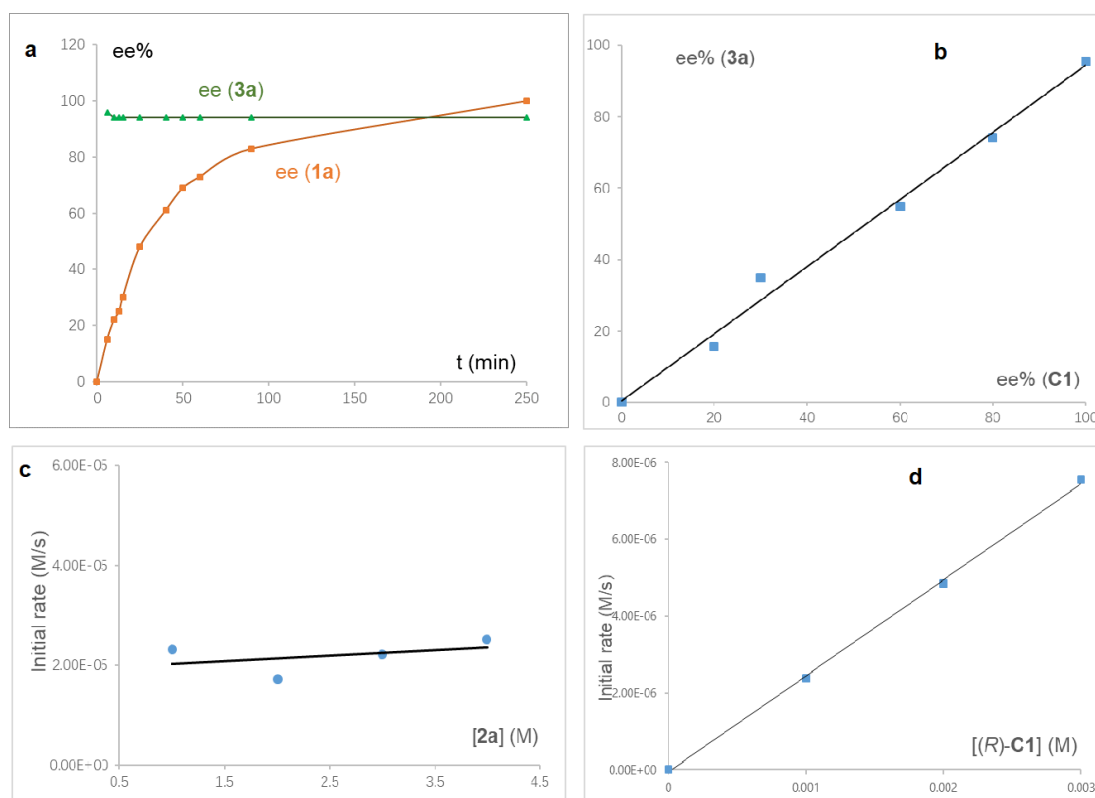
Yield was determined by analysis of the <sup>1</sup>H NMR spectrum of the crude reaction mixture with CH<sub>2</sub>Br<sub>2</sub> as internal standard. Ee was determined by HPLC analysis on a chiral stationary phase.

**Mechanistic studies.** A possible mechanism is proposed in Fig 2. We believe that this reaction begins with acid-catalyzed dehydration to form indolyl cation **IM**, paired with a phosphate counter anion. This ion pair might be in equilibrium (or pseudo resonance) with the activated indole imine methide form **IM'**. Subsequently, the hydride source approaches the benzylic carbon to deliver the product **3**.



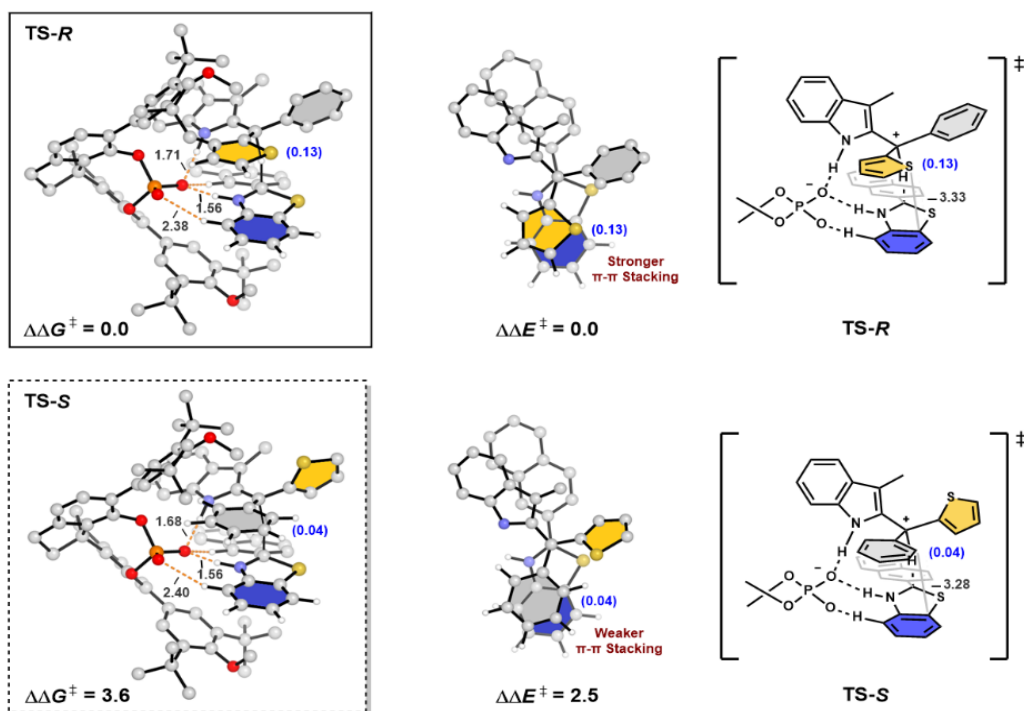
**Fig. 2.** Proposed mechanism and a control experiment. **a** Proposed mechanism **b** Control experiments with *N*-substituted substrate.

We have carried out a series of control experiments. First of all, under the standard conditions, the reaction with *N*-methylated substrate **1a'** did not proceed to form the desired product **3a'** (Scheme 3b). This result suggests that the free N–H motif in the indole moiety is essential to the observed reactivity, which is consistent with the intermediacy of 2-indole imine methide **IM'**, as this intermediate cannot be formed from **1a'**. Next, the enantiomeric excess (ee) values of substrate and product were both monitored during the reaction progress (Fig 3a). The product ee remained constant (95% ee) during the entire reaction, but substrate ee gradually increased over time. This enantioconvergent feature agrees with the initial formation of an achiral 2-indole imine methide intermediate followed by stereodefined asymmetric addition of nucleophile. The observation of substrate enantioenrichment is indicative of kinetic resolution during the first step, which is likely irreversible. Taken together, a direct  $\text{S}_{\text{N}}2$  mechanism could be excluded. Furthermore, this reaction did not exhibit non-linear effects, suggesting that the enantiodetermining transition state likely involves only one catalyst molecule. Finally, kinetic studies indicated that this reaction exhibits zeroth order in nucleophile and first order in catalyst, which further confirmed that the first step is rate-determining and irreversible.



**Fig. 3.** Mechanistic studies. **a** Time-dependence of substrate and product ee values. **b** Absence of non-linear effects. **c** Zeroth order in nucleophile. **d** First order in catalyst.

**Density functional theory studies.** The rate-determining step of this reaction is the chiral phosphoric acid-catalyzed dehydration of tertiary alcohols **1a** to obtain the indolyl cation **IM**. However, the stereo-determining step for this reaction is the nucleophilic addition of the hydride **2a** on the indolyl cation **IM**. To evaluate the origins of stereoselectivity, density functional theory (DFT) calculations were performed. We first explored the stereo-determining transition structures (Fig. 4). Geometry optimizations were conducted at B3LYP-D3(BJ) functional with the def2svp basis set in the Gaussian 16 package.<sup>41-43</sup> Single-point energies and solvent effects in dichloromethane were calculated with the CPCM solvation model at the M06-2X/def2tzvpp level of theory.<sup>44-47</sup> Conformational searches were conducted using the CREST conformer-rotamer ensemble sampling tool version 2.10.2 with xtb version 6.3.3.<sup>48-52</sup> Based on the results of conformational searches, the lowest energy conformers are depicted in Figure 2. The computed transition state **TS-R** delivering the preferred R-product is 3.6 kcal/mol lower than that of **TS-S** generating the S-product, in good accord with the 96% ee observed experimentally.



**Fig. 4.** DFT-optimized stereo-determining transition structures. The distances are given in Ångstroms, and energies are given in kcal/mol. Coloured rings: grey, phenyl; yellow, thienyl; blue, benzo group on the benzothiazoline. Numbers in parentheses are the total Hirshfeld charges on the aryl groups.

To gain further insights into the factors that impact the enantioselectivity, the geometries of transition states **TS-R** and **TS-S** were compared. No obvious steric clashes and hydrogen-bonding interaction difference between the catalyst and substrates are detected in these two competing transition states. Computational studies of the total Hirshfeld charges on the aryl groups show that the key interaction for discrimination of the two aryl groups is mainly  $\pi$ - $\pi$  stacking. Thienyl is a better donor than phenyl so it donates more electrons to the C<sup>+</sup>. In major **TS-R**, the electron-deficient thienyl (0.13 e) is in closer contact with the electron-rich benzo ring of the benzothiazoline. By contrast, in minor **TS-S**, the phenyl group (0.04 e) forms a slip-stacked configuration with the benzo group on the hydride. As a result, the stronger  $\pi$ - $\pi$  stacking stabilizes **TS-R** more than the weaker  $\pi$ - $\pi$  stacking stabilizes **TS-S**. This conclusion rather than some interaction of transition state with catalyst was tested by calculations on the fixed transition state formed by removing the catalyst, Single-point  $\Delta\Delta E^\ddagger$  without optimization show a 2.5 kcal/mol advantage for the stronger attractive  $\pi$ - $\pi$  stacking in **TS-R**. This is the significant contribution to the 3.6 kcal/mol preference for the

formation of R-product. Therefore, attractive  $\pi$ - $\pi$  stacking plays a major role in the selectivity.

**Biological Studies.** To investigate the potential anticancer activity of the enantioenriched indole-containing triarylmethanes, we examined the cytotoxicity of the representative product **3d** towards human cervical adenocarcinoma (HeLa), ovarian carcinoma (A2780), breast adenocarcinoma (MCF-7), colorectal carcinoma (HCT116), and lung carcinoma (A549) cells. A widely used anticancer drug, doxorubicin, was used as the control. As shown in Table 4, **3d** exhibited significant cytotoxicity, with the half-maximal inhibitory concentration (IC<sub>50</sub>) values ranging from 5.6 to 18.2  $\mu$ M. A549 lung carcinoma cells were found to be the most sensitive cell line towards **3d**, and a normal cell line from the same origin (MRC-5 lung fibroblasts) was used to evaluate the cancer cell selectivity of **3d**. The IC<sub>50</sub> value in MRC-5 cells is 27.1  $\mu$ M, corresponding to a selectivity index (SI; cytotoxicity in the normal cells / cytotoxicity in the cancer cells) of 4.8; whereas the SI value for doxorubicin is only 2.6. These preliminary results suggest that this class of compounds have promising potential for further development as anticancer drug candidates.

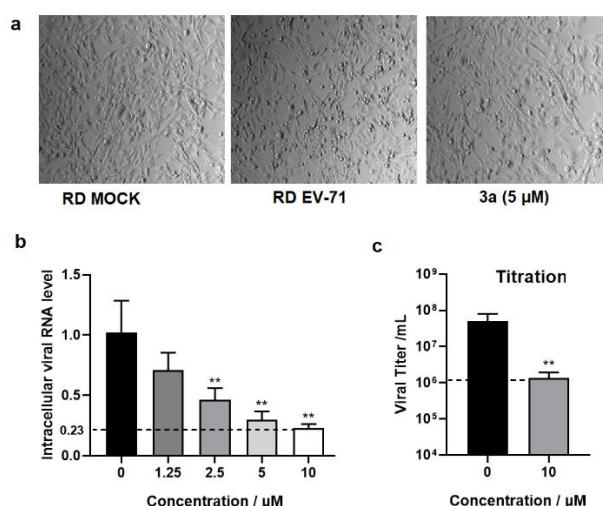
**Table 4.** The cytotoxicity of **3d** in various human cell lines.

Cell line	IC <sub>50</sub> value ( $\mu$ M) <sup>a</sup>					
	HeLa	MCF-7	A2780	A549	HCT116	MRC-5
Doxorubicin	1.4 $\pm$ 0.4	0.55 $\pm$ 0.11	0.36 $\pm$ 0.07	0.28 $\pm$ 0.06	1.4 $\pm$ 0.3	0.72 $\pm$ 0.14
<b>3d</b>	18.2 $\pm$ 2.9	15.3 $\pm$ 2.3	8.9 $\pm$ 1.7	5.6 $\pm$ 0.9	9.7 $\pm$ 1.3	27.1 $\pm$ 3.4

<sup>a</sup>The half inhibitory concentration (IC<sub>50</sub>) values were determined by MTT assay in 72 h. The error bars were obtained as the standard deviation from the mean value based on three independent experiments.

We also tested the antiviral activity of another representative product **3a** with enterovirus A71 (EV-A71) using Rhabdomyosarcoma (RD) cell line. The cytopathic effect (CPE) and intracellular viral RNA level were measured to reflect the antiviral effects. Indeed, a strong CPE was observed after EV-A71 infection at the multiplicity of infection (MOI) of 0.01 for 36 hours. The morphology of RD cells changed from flat to unhealthy round and death. As shown in Fig. 5a, the CPE induced by EV-A71 infection was significantly reduced upon treatment with **3a**. Moreover, the intracellular viral RNA level was decreased by 80-90% after treating with **3a** at the concentration of 5-10  $\mu$ M

compared with untreated EV-A71 infected cells (Fig. 5b). The strong antiviral effect of **3a** was also confirmed by viral titration. The virus titer was decreased by 35 folds upon treatment with **3a** (Fig. 5c). Moreover, this compound showed low cytotoxicity according to the MTT assay (Table 5), thus indicating a high selectivity index and suggesting a great potential of such molecules for antiviral drug development.



**Fig. 5. The antiviral effects of 3a shown by CPE assay and intracellular viral RNA level.** **a** RD cells were first treated with compounds at different concentration and then infected with EV-A71 at MOI of 0.01 after 2 hours. Cell morphology was observed 36 h post-infection. **b** Relative intracellular EV-A71 genome RNA level was determined by RT-qPCR. **c** The EV-A71 viral titer in the supernatant was measured by TCID<sub>50</sub> assay. Data are represented as mean ± SD (n = 3). \*\*p < 0.01, compared with no infected group.

**Table 5. Cytotoxicity concentration (CC<sub>50</sub>) and antiviral activity IC<sub>50</sub>.<sup>a</sup>**

Compound	CC <sub>50</sub> (μM)	IC <sub>50</sub> (μM)	Selectivity index
<b>3a</b>	55.46	2.27	24.43

<sup>a</sup> CC<sub>50</sub>, 50% cytotoxic concentration tested by the viability assay with no viral infection. IC<sub>50</sub>, viral RNA copies decreased 50% compared with the control group (without compound treatment) in the secreted virions. A selectivity index (CC<sub>50</sub>/IC<sub>50</sub>) >10 is assumed as a potential candidate for further research analysis.

## Conclusion

Despite the longstanding challenge on asymmetric discrimination between two sterically similar diaryl groups and the dominant role of metal catalysis in the limited previous studies, here we have demonstrated a rare organocatalytic example with excellent efficiency and enantiocontrol. The versatile 2-indole imine methide bearing aryl and heteroaryl groups without a directing group was used as a platform for this study. The combined use of a superb chiral phosphoric acid catalyst and a benzothioazoline hydride source is critically important to the success. This protocol provided efficient access to a wide range of highly enantioenriched indole-containing triarylmethanes from the corresponding racemic tertiary alcohols. Mechanistic experiments, including control reactions and kinetic studies, provided important insights into the mechanism, which involves initial rate-determining dehydration (with concomitant substrate kinetic resolution) and subsequent enantioconvergent nucleophilic addition. Further DFT studies suggested that it is  $\pi$ - $\pi$  stacking, but not hydrogen bonding, that provides the key interaction for asymmetric discrimination between the benzene and thiophene rings. This is also consistent with the robust enantiocontrol in the presence of various polar functional groups that are likely hydrogen-bond destroyers. Preliminary biological studies also demonstrated the great potential of these triarylmethanes for anticancer and antiviral drug development.

## Methods

### General procedure for the synthesis of enantioenriched triarylmethanes 3.

At 0 °C, to an oven-dried 30-mL vial were added the tertiary alcohol **1** (0.40 mmol), the hydride source **2a** (115.9 mg, 0.44 mmol), (*R*)-**C1** (15.04 mg, 0.02 mmol, 5.0 mol%) and DCM (8.0 mL). The mixture was stirred at the same temperature for 30 h and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography to afford the desired product.

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### **Author contributions**

Q.Y. conceived the project, performed the synthesis experiments, and wrote the paper. M.D. and P.Y. performed DFT calculations. C.C., Z.D. and M.W. performed the biological experiments. M.H. and G.Z. directed the biological experiments and data analysis. K.N.H. directed the DFT calculations and mechanism analysis. J.S. conceived and directed the project and wrote the paper. All the authors discussed the results and commented on the manuscript.

### **Competing interests**

The authors declare no competing interests.