Supporting Information for

Palladium-catalyzed asymmetric allylic amination of

vinylethylene carbonate with N-heteroaromatics

Chao Xia,^a Dong-Chao Wang,^b Gui-Rong Qu^b and Hai-Ming Guo*, ^{a, b}

^aSchool of Environment, Henan Normal University, Xinxiang, Henan Province 453007, China

^bNMPA Key Laboratory for Research and Evaluation of Innovative Drug, Key Laboratory of Green Chemical Media and Reactions, Ministry of Education, Collaborative Innovation Center of Henan Province for Green Manufacturing of Fine Chemicals, School of Chemistry and Chemical Engineering, Henan Normal University, Xinxiang, Henan 453007, China.

E-mail: ghm@htu.edu.cn

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1. General information

¹H NMR and ¹³C NMR spectra were recorded on Bruker Avance III HD 600 or Avance 400 MHz spectrometer. Chemical shifts are recorded in ppm relative to tetramethylsilane with the solvent resonance as the internal standard. Data are represented as follows: chemical shift, multiplicity (s = singlet, d = doublet, t = triplet, q = quartet, dd = doublet of doublets, m = multiplet), coupling constants (*J*) are in Hertz (Hz), and integration. Enantiomer excesses were determined by chiral HPLC analysis on Chiralcel IA/AS-H/ID/OD-H/IE/IG in comparison with the authentic racemates. Chiral HPLC analysis was recorded on Thermo Scientific Dionex Ultimate 3000 and Agilent Technologies 1260 Infinity. Optical rotations were recorded on Autopol Automatic Polarimeter, and were reported as follows: $[\alpha]_D^T$ (c: g/100 mL, in CH₂Cl₂). High resolution mass spectra (HRMS) was recorded on Supernova Atlas S2 CCD detector. Melting point (m.p.) data were obtained on X-5 micro melting point apparatus. For column chromatography, silica gel (200-300 mesh) was used as the stationary phase. Unless stated otherwise, all the solvent and reagents were purchased from commercial suppliers and used without further purification.

2. The Optimization of Reaction Conditions

	$ \begin{array}{c} CI \\ N \\ N \\ N \\ H \\ 1a \end{array} $	20 or 0	Pd₂(dba)₃ (5 mmol%) L (10 mmol%) DCM, N₂, rt, 6 h	$ \begin{array}{c} CI \\ N \\ N \\ N \\ HO \\ 3a \\ \end{array} $
	PPh ₂ PPh ₂ L	$ \begin{array}{c} $	L3 V	$L4$ $\downarrow \downarrow $
entry	2	L	yield ^{b} (%)	ee^{c} (%)
1	2a	L1	37	43
2	2b	L1	NR	
3	2a	L2	50	25
4	2b	L2	48	25
5	2a	L3	28	23
6	2b	L3	35	23
7	2a	L4	trace	
8	2b	L4	NR	
9	2a	L5	42	20
10	2b	L5	45	17
11	2a	L6	88	83
12	2b	L6	88	77
13	2a	L7	85	85
14	2b	L7	80	83
15	2a	L8	90	95
16	2b	L8	86	90

Table S1 The optimization of reaction conditions for selecting substrate 2^a

^{*a*} Reaction conditions: **1a** (0.2 mmol), **2** (0.22 mmol), and solvent (2.0 mL). ^{*b*} Isolated yields. ^{*c*} Determined by chiral HPLC analysis.

Table S2 The Screening of Palladium^a

			[Pd] (x mol%) L8 (1 mol%) CH ₃ CN, N ₂ , rt, 12 h		
	1a	2		3a	
entry	[Pd] (x mol%)		yield ^{b} (%)	ee^{c} (%)	
1	Pd ₂ (dba) ₃ ·CHCl ₃ (0.5)		86	96	
2	$Pd(dba)_2(1)$		88	96	
3	[Pd(C ₃ H ₅)Cl] ₂ (0.5)		20	73	
4	$Pd(PPh_3)_4(1)$		47	20	
5	$Pd(OAc)_2(1)$		NR		
^{<i>a</i>} Reaction conditions: 1a (0.4 mmol), 2 (0.44 mmol), and solvent (4.0 mL). ^{<i>b</i>} Isolated yields. ^{<i>c</i>}					
Determined by chiral HPLC analysis.					

		Pg Me	+	Pd₂(dba)₃ (0.5 mol%) L8 (1 mol%) Solvent, N₂, 4 h, rt	Pg Me O N HO	
		4	2		5	
_	entry	4	Pg	Solvent (x mL)	yield ^b (%)	ee ^c (%)
	1	4 a	Bz	$CH_3CN(2)$	97	82
	2	4 b	Boc	$CH_3CN(2)$	93	82
	3	4 a	Bz	DCM (2)	85	66
	4	4 a	Bz	DCE (2)	83	69
	5	4 a	Bz	$CHCl_{3}(2)$	89	50
	6	4 a	Bz	EA (2)	90	76
	7	4 a	Bz	THF (2)	NR	
	8	4 a	Bz	toluene (2)	92	74
	9	4 a	Bz	CH ₃ CN (4)	97	84
	10	4 a	Bz	CH ₃ CN (6)	80	84
	11	4 a	Bz	CH ₃ CN (8)	53	90
	12	4 a	Bz	CH ₃ CN (10)	35	92
	13 ^d	4 a	Bz	CH ₃ CN (10)	35	92
	14^e	4a	Bz	CH ₃ CN (10)	95	92

Table S3 The optimization of reaction conditions for pyrimidine 4^a

^{*a*} Reaction conditions: **4** (0.2 mmol), **2** (0.22 mmol). ^{*b*} Isolated yields. ^{*c*} Determined by chiral HPLC analysis. ^{*d*} The reaction was carried out at 0 °C. ^{*e*} 3 equivalent **2** was used.

3. General procedure for the asymmetric allylic amination reactions



A reaction tube was charged with 1 (0.4 mmol), $Pd_2(dba)_3$ (1.8 mg, 0.002 mmol, 0.5 mol%) and L8 (3.2 mg, 0.004 mmol, 1 mol%). The reaction tube was placed under vacuum and backfilled with argon three times. CH₃CN (4.0 mL) followed by 2 (0.44 mmol, 1.1 equiv) were added via syringe under argon. The resulting mixture was stirred at rt for 12 h, and the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 2:1 to 1:1) to give the corresponding products **3**.



A reaction tube was charged with **4** (0.2 mmol), $Pd_2(dba)_3$ (0.9 mg, 0.001 mmol, 0.5 mol%) and **L8** (1.6 mg, 0.002 mmol, 1 mol%). The reaction tube was placed under vacuum and backfilled with argon three times. CH₃CN (10.0 mL) followed by **2** (0.6 mmol, 3 equiv) were added via syringe under argon. The resulting mixture was stirred at rt for 4 h, and the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 2:1 to 1:1) to give the corresponding annulation **5**.



A reaction tube was charged with **6** (0.4 mmol), $Pd_2(dba)_3$ (1.8 mg, 0.002 mmol, 0.5 mol%) and **L8** (3.2 mg, 0.004 mmol, 1 mol%). The reaction tube was placed under vacuum and backfilled with argon three times. CH₃CN (4.0 mL) followed by **2** (0.44 mmol, 1.1 equiv) were added via syringe under argon. The resulting mixture was stirred at rt for 6 h, and the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 4:1 to 2:1) to give the corresponding products **5**.

4. Scale-up synthesis of product 3g



In a 50 mL reaction flask, $Pd_2(dba)_3$ (18 mg, 0.02 mmol, 0.5 mol%), L8 (32 mg, 0.04 mmol, 1 mol%) and CH₃CN (40 mL) were introduced under an argon atmosphere. The resulting solution was stirred for 30 minutes. Then, 1g (1.34g, 4 mmol) and 2 (502 mg, 4.4 mmol) were added in one portion. The resulting mixture was stirred at rt for 12 h, and the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 2:1 to 1:1) to give product 3g as white solid (1.49 g, 92% yield, 95% ee).

5. Transformation



To a solution of **5a** (60 mg, 0.2 mmol) in MeOH (2.0 mL), NaOH (16 mg, 0.4 mmol) was added. The reaction was stirred at the room temperature for 12 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 1:1 to 1:2) to give product **5aa** as white solid (37.7 mg, 96% yield, 92% ee).



STEP 1: To a solution of **3g** (81 mg, 0.2 mmol) in MeOH (2.0 mL), ^{*t*}BuONa (38.4 mg, 0.4 mmol) was added. The reaction was stirred at the room temperature for 12 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 1:1 to 1:2) to give product **3ga** as white solid (58 mg, 95% yield).

STEP 2: Heated the temperature of the reaction solution that had been completed in **STEP 1** to 50 °C and stirred the reaction for other 16 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (DCM:MeOH = 10:1) to give product **3gb** as white solid (39.1 mg, 95% yield, 94% ee).



STEP 1: To a solution of **3g** (81 mg, 0.2 mmol) and Cs_2CO_3 (130 mg, 0.4 mmol) in CH₃CN (2.0 mL), allyl iodide (51 mg, 0.3 mmol) was added. The reaction was stirred at 40 °C for 12 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 4:1 to 2:1) to give product **3gc** as colorless oil (80.2 mg, 90% yield).

STEP 2: To a solution of **3gc** (89 mg, 0.2 mmol) in DCM (2.0 mL), Grubbs II catalyst (10 mol%) was added. The reaction was stirred under N₂ at 40 °C for 12 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 3:1 to 3:2) to give product **3gd** as colorless oil (73.4 mg, 88% yield, 94% ee).

STEP 3: To a solution of **3gd** (83.5 mg, 0.2 mmol) in MeOH (2.0 mL), 'BuONa (38.4 mg, 0.4 mmol) was added. The reaction was stirred at 50 °C for 16 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (DCM:MeOH = 10:1) to give product **3ge** as white solid (41.2 mg, 95% yield, 95% ee).

STEP 4: K₂OsO₂(OH)₄ (2.4 mg, 0.008 mmol, 4 mol %), (DHQ)₂PYR (17.6 mg, 0.02 mmol, 10 mol %), K₃Fe(CN)₆ (196 mg, 0.6 mmol, 3 equiv), methanesulfonamide (38 mg, 0.4 mmol, 2 equiv) and K₂CO₃ (84 mg, 0.6 mmol, 3 equiv) were suspended in a mixture of water and *tert*-butyl alcohol (1:1, 8 mL). The mixture was stirred at room temperature for 1 h and then added **3gd** (83.5 mg, 0.2 mmol). The reaction was stirred at the room temperature for 24 h and monitored by TLC until the reaction was completed. The reaction was quenched at 0 °C by addition of Na₂S₂O₃ and the mixture stirred at room temperature for 2 h. The reaction mixture was then partitioned between ethyl acetate and water. The combined organic phases were dried (Na₂SO₄), filtered and concentrated in vacuo to afford a crude oil. Purification by flash column chromatography (PE:EA =

1:1 to 1:3) to furnished **3gf** (79.3 mg, 88% yield, 93% ee and >20:1 dr) as white solid.

STEP 5: To a solution of **3gd** (90.2 mg, 0.2 mmol) in MeOH (2.0 mL), 'BuONa (38.4 mg, 0.4 mmol) was added. The reaction was stirred at 50 °C for 16 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (DCM:MeOH = 10:1 to 3:1) to give product **3gg** as white solid (45.2 mg, 90% yield).



STEP 1: To a solution of **3**g (81 mg, 0.2 mmol) and TsCl (53 mg, 0.3 mmol) in DCM (2.0 mL), Et₃N (40 mg, 0.4 mmol) was added. The reaction was stirred at the room temperature for 6 h. After the reaction was consumed (determined by TLC), the solvent was removed under vacuum. The residue was purified by flash column chromatography on silica gel (PE:EA = 4:1 to 2:1) to give product **3gh** as white solid (104 mg, 93% yield).

STEP 2: K₂OsO₂(OH)₄ (1.2 mg, 0.004 mmol, 2 mol %), (DHQ)₂PYR (17.6 mg, 0.02 mmol, 10 mol %), K₃Fe(CN)₆ (196 mg, 0.6 mmol, 3 equiv), methanesulfonamide (38 mg, 0.4 mmol, 2 equiv) and K₂CO₃ (84 mg, 0.6 mmol, 3 equiv) were suspended in a mixture of water and *tert*-butyl alcohol (1:1, 8 mL). The mixture was stirred at room temperature for 1 h and then added **3gh** (112 mg, 0.2 mmol). The reaction was stirred at 0 °C for 12 h and monitored by TLC until the reaction was completed. The reaction was quenched at 0 °C by addition of Na₂S₂O₃ and the mixture stirred at room temperature for 2 h. The reaction mixture was then partitioned between ethyl acetate and water. The combined organic phases were dried (Na₂SO₄), filtered and concentrated in vacuo to afford a crude oil. Purification by flash column chromatography (PE:EA = 1:1 to 1:3) to furnished **3gi** (109.2 mg, 92% yield, 96% ee and >20:1 dr) as white solid.

STEP 3: Warmed the temperature of the reaction solution that had been completed in **STEP 2** up to room temperature and K_2CO_3 (56 mg, 0.4 mmol, 2 equiv) was added. The reaction was stirred at the room temperature for other 12 h. The reaction was quenched at 0 °C by addition of Na₂S₂O₃ and the mixture stirred at room temperature for 2 h. The reaction mixture was then partitioned

between ethyl acetate and water. The combined organic phases were dried (Na₂SO₄), filtered and concentrated in vacuo to afford a crude oil. Purification by flash column chromatography (PE:EA = 1:1 to 1:2) to furnished **3gj** (69.6 mg, 85% yield, 94% ee and >20:1 dr) as colorless oil.

6.Characterization Data of Products

(S)-2-(6-chloro-9H-purin-9-yl)but-3-en-1-ol (3a)



White solid; m.p. 104.8-106.3 °C; 81.6 mg, 91% yield, 97% ee; $[\alpha]_D^{23} = -73.67$ (c = 0.600, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 85/15, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 16.395 min (major), 17.837 min (minor); ¹**H NMR** (600 MHz, CDCl3) δ 8.66 (s, 1H), 8.22 (s, 1H), 6.21 (ddd, *J* = 17.4, 10.8, 6.6 Hz, 1H), 5.41 (dd, *J* = 10.8, 1.2 Hz, 1H), 5.27 (dddt, *J* = 6.6, 4.8, 3.6, 1.2 Hz, 1H), 5.24 (dd, *J* = 17.4, 1.8 Hz, 1H), 4.25 (dd, *J* = 12.0, 6.0 Hz, 1H), 4.15 (dd, *J* = 12.0, 3.6 Hz, 1H), 4.01 (brs, 1H); ¹³**C NMR** (150 MHz, CDCl3) δ 151.8, 151.4, 150.9, 145.5, 132.1, 131.4, 120.4, 63.4, 60.8; **HRMS** (ESI-TOF): exact mass calcd. for C₉H₁₀ClN₄O [M+H]⁺ requires m/z 225.0538, found m/z 225.0538.

(S)-2-(9H-purin-9-yl)but-3-en-1-ol (3b)



White solid; m.p. 70.4-71.6 °C; 54.8 mg, 72% yield, 95% ee; $[\alpha]_D^{23} = -84.21$ (c = 0.585, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 13.320 min (minor), 17.232 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.88 (s, 1H), 8.79 (s, 1H), 8.19 (s, 1H), 6.21 (ddd, *J* = 17.6, 10.4, 6.4 Hz, 1H), 5.37 (d, *J* = 10.4 Hz, 1H), 5.28 – 5.23 (m, 1H), 5.20 (d, *J* = 17.2 Hz, 1H), 4.96 (brs, 1H), 4.23 (dd, *J* = 12.0, 6.0 Hz, 1H), 4.10 (dd, *J* = 12.0, 3.6 Hz, 1H); ¹³**C NMR** (100 MHz, CDCl₃) δ 152.1, 151.0, 148.3, 145.5, 133.8, 132.5, 120.0, 63.3, 60.4; **HRMS** (ESI-TOF): exact mass calcd. for C₉H₁₁N₄O [M+H]⁺ requires m/z 191.0927, found m/z 191.0924.

(S)-2-(6-methyl-9H-purin-9-yl)but-3-en-1-ol (3c)



White solid; m.p. 108.6-110.2 °C; 63.6 mg, 78% yield, 96% ee; $[\alpha]_D^{23} = -100.55$ (c = 0.610, CHCl₃);

HPLC CHIRALCEL IE, *n*-hexane/2-propanol = 85/15, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 21.165 min (major), 23.288 min (minor); ¹**H** NMR (400 MHz, CDCl₃) δ 8.64 (s, 1H), 8.09 (s, 1H), 6.20 (ddd, *J* = 17.0, 10.4, 6.4 Hz, 1H), 5.35 (d, *J* = 10.4 Hz, 1H), 5.22 – 5.09 (m, 3H), 4.28 – 4.20 (m, 1H), 4.10 (d, *J* = 12.4 Hz, 1H), 2.68 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 159.4, 151.7, 150.1, 144.1, 132.9, 132.6, 119.7, 63.4, 61.0, 19.3; **HRMS** (ESI-TOF): exact mass calcd for C₁₀H₁₃N₄O [M+H]⁺ requires m/z 205.1084, found m/z 205.1083.

(S)-2-(6-methoxy-9H-purin-9-yl)but-3-en-1-ol (3d)



White solid; m.p. 116.2-117.4 °C; 81.0 mg, 76% yield, 92% ee; $[\alpha]_D^{23} = -93.33$ (c = 0.660, CHCl₃); **HPLC** CHIRALCEL IG, *n*-hexane/2-propanol = 80/20, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 13.873 min (major), 15.707 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 8.43 (s, 1H), 7.92 (s, 1H), 6.23 (ddd, *J* = 17.2, 10.4, 6.4 Hz, 1H), 5.37 (d, *J* = 10.4 Hz, 1H), 5.28 – 5.22 (m, 1H), 5.19 (d, *J* = 17.2 Hz, 1H), 5.16 – 5.11 (m, 1H), 4.25 – 4.14 (m, 2H), 4.10 (s, 3H); ¹³**C NMR** (100 MHz, CDCl₃) δ 160.8, 151.6, 151.2, 142.5, 132.8, 121.2, 119.7, 63.6, 61.4, 54.3; **HRMS** (ESI-TOF): exact mass calcd for C₁₀H₁₃N₄O₂ [M+H]⁺ requires m/z 221.1033, found m/z 221.1032.

(S)-2-(6-(propylthio)-9H-purin-9-yl)but-3-en-1-ol (3e)



Colorless oil; 87.7 mg, 83% yield, 94% ee; $[\alpha]_D^{23} = -81.97$ (c = 0.610, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 80/20, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 10.732 min (major), 11.790 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 8.58 (s, 1H), 7.98 (s, 1H), 6.16 (ddd, *J* = 17.2, 10.4, 6.4 Hz, 1H), 5.33 (d, *J* = 10.4 Hz, 1H), 5.20 – 5.05 (m, 3H), 4.15 (s, 2H), 3.26 (t, *J* = 7.2 Hz, 2H), 1.75 (h, *J* = 7.2 Hz, 2H), 1.03 (t, *J* = 7.2 Hz, 3H); ¹³C **NMR** (100 MHz, CDCl₃) δ 161.7, 151.5, 147.7, 142.7, 132.7, 131.0, 119.6, 63.6, 60.6, 30.7, 22.9, 13.5; **HRMS** (ESI-TOF): exact mass calcd for C₁₂H₁₇N₄OS [M+H]⁺ requires m/z 265.1118, found m/z 265.1119.

(S)-2-(6-(diethylamino)-9H-purin-9-yl)but-3-en-1-ol (3f)



Colorless oil; 74.2 mg, 71% yield, 94% ee; $[\alpha]_D^{23} = -87.87$ (c = 0.500, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 85/15, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 11.492 min (major), 12.808 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 8.23 (s, 1H), 7.73 (s, 1H), 6.15 (ddd, J = 16.8, 10.4, 6.0 Hz, 1H), 5.30 (d, J = 10.4 Hz, 1H), 5.03 (d, J = 17.2 Hz, 2H), 4.14 – 4.05 (m, 2H), 3.96 (brs, 4H), 1.27 (t, J = 7.2 Hz, 6H); ¹³**C NMR** (100 MHz, CDCl₃) δ 154.0, 151.9, 149.7, 138.6, 133.5, 119.9, 118.6, 64.2, 61.5, 43.3, 13.6; **HRMS** (ESI-TOF): exact mass calcd for C₁₃H₂₀N₅O [M+H]⁺ requires m/z 262.1662, found m/z 262.1663.

(S)-2-(6-(diBocamino)-9H-purin-9-yl)but-3-en-1-ol (3g)



White solid; m.p. 59.5-61.2 °C; 142.6 mg, 88% yield, 94% ee; $[\alpha]_D^{23} = -44.95$ (c = 0.525, CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol = 90/10, flow rate = 0.7 mL/min, λ = 256 nm, retention time: 11.008 min (minor), 12.845 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.75 (s, 1H), 8.18 (s, 1H), 6.15 (ddd, *J* = 17.2, 10.4, 6.4 Hz, 1H), 5.29 (d, *J* = 10.4 Hz, 1H), 5.23 (q, J = 5.6 Hz, 1H), 5.07 (d, *J* = 17.2 Hz, 1H), 4.36 (brs, 1H), 4.12 (dd, *J* = 12.0, 6.2 Hz, 1H), 4.01 (dd, *J* = 12.0, 3.6 Hz, 1H), 1.39 (s, 18H); ¹³**C NMR** (100 MHz, CDCl₃) δ 153.1, 151.6, 150.5, 150.2, 145.2, 132.7, 128.8, 119.4, 84.0, 63.2, 60.3, 27.8; **HRMS** (ESI-TOF): exact mass calcd for C₁₉H₂₈N₅O₅ [M+H]⁺ requires m/z 406.2085, found m/z 406.2080.

(S)-2-(2-chloro-9H-purin-9-yl)but-3-en-1-ol (3h)

White solid; m.p. 89.0-90.6 °C; 68.2 mg, 76% yield, 96% ee; $[\alpha]_D^{23} = -53.55$ (c = 0.620, CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol = 90/10, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 23.547 min (minor), 25.702 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.85 (s, 1H), 8.23 (s, 1H), 6.19 (ddd, *J* = 17.2, 10.4, 6.4 Hz, 1H), 5.42 (d, *J* = 10.4 Hz, 1H), 5.31 – 5.22 (m, 2H), 4.21 (dd, *J* = 12.0, 6.2 Hz, 1H), 4.09 (dd, *J* = 12.0, 3.7 Hz, 1H), 3.60 (brs, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 154.2, 153.0, 150.0, 146.1, 133.0, 132.0, 120.5, 63.4, 59.8; **HRMS** (ESI-TOF): exact mass calcd for C₉H₁₀ClN₄O [M+H]⁺ requires m/z 225.0538, found m/z 225.0537.

(S)-2-(2,6-dichloro-9H-purin-9-yl)but-3-en-1-ol (3i)



White solid; m.p. 144.6-146.2 °C; 53.8 mg, 52% yield, 97% ee; $[\alpha]_D^{23} = -49.25$ (c = 0.555, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 90/10, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 14.182 min (major), 16.353 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 8.27 (s, 1H), 6.20 (ddd, *J* = 17.2, 10.4, 6.4 Hz, 1H), 5.45 (d, *J* = 10.4 Hz, 1H), 5.32 – 5.26 (m, 2H), 4.20 (dd, *J* = 12.0, 6.0 Hz, 1H), 4.12 (dd, *J* = 12.0, 3.6 Hz, 1H), 3.20 (brs, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 153.0, 152.9, 151.7, 146.0, 131.7, 130.6, 120.9, 63.4, 60.1; **HRMS** (ESI-TOF): exact mass calcd for C₉H₉Cl₂N₄O [M+H]⁺ requires m/z 259.0148, found m/z 259.0148.

(S)-2-(2-chloro-6-(dimethylamino)-9H-purin-9-yl)but-3-en-1-ol (3j)



White solid; m.p. 119.5-121.1 °C; 77.0 mg, 72% yield, 94% ee; $[\alpha]_D^{23} = -73.53$ (c = 0.835, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 6.885 min (major), 8.505 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 7.70 (s, 1H), 6.13 (ddd, *J* = 17.2, 10.4, 6.2 Hz, 1H), 5.34 (d, *J* = 10.4 Hz, 1H), 5.14 (d, *J* = 17.2 Hz, 1H), 5.11 – 5.05 (m, 1H), 4.28 (s, 1H), 4.15 – 4.02 (m, 2H), 3.62 (s, 3H), 3.29 (s, 3H); ¹³**C NMR** (150 MHz, CDCl₃) δ 155.0, 153.5, 151.3, 138.5, 132.9, 119.4, 118.9, 64.0, 60.3, 39.2, 38.2; **HRMS** (ESI-TOF): exact mass calcd for C₁₁H₁₅ClN₅O [M+H]⁺ requires m/z 268.0960, found m/z 268.0957.

(S)-2-(2-(diBocamino)-6-(benzyloxy)-9H-purin-9-yl)but-3-en-1-ol (3k)



White solid; m.p. 61.0-62.2 °C; 184.0 mg, 90% yield, 95% ee; $[\alpha]_D^{23} = -30.00$ (c = 0.600, CHCl₃); HPLC CHIRALCEL IG, *n*-hexane/2-propanol = 80/20, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 12.292 min (minor), 14.260 min (major); ¹H NMR (400 MHz, CDCl₃) δ 7.99 (s, 1H), 7.48 (d, J = 7.6 Hz, 2H), 7.35 – 7.25 (m, 3H), 6.10 (ddd, J = 16.8, 10.4, 6.0 Hz, 1H), 5.59 (s, 2H), 5.27 (d, J = 10.4 Hz, 1H), 5.10 (q, J = 5.6, 4.4 Hz, 1H), 5.03 (d, J = 17.2 Hz, 1H), 4.44 (t, J = 6.0 Hz, 1H), 4.10 (dt, J = 12.0, 6.0 Hz, 1H), 4.01 (dt, J = 12.0, 4.0 Hz, 1H), 1.40 (s, 18H); ¹³C NMR (100 MHz, CDCl₃) δ 160.9, 152.6, 151.4, 150.8, 143.2, 135.8, 133.1, 128.6, 128.5, 128.3, 119.7, 119.0, 83.4, 68.8, 63.4, 60.7, 27.9; **HRMS** (ESI-TOF): exact mass calcd for C₂₆H₃₄N₅O₆ [M+H]⁺ requires m/z 512.2504, found m/z 512.2503.

(S)-9-(1-hydroxybut-3-en-2-yl)-1,3-dimethyl-3,9-dihydro-1H-purine-2,6-dione (31)



White solid; m.p. 140.9-142.6 °C; 76.1 mg, 76% yield, 97% ee; $[\alpha]_D^{23} = -65.11$ (c = 0.600, CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol =60/40, flow rate = 0.4 mL/min, λ = 256 nm, retention time: 18.970 min (major), 25.622 min (minor); ¹**H NMR** (400 MHz, CDCl₃) δ 7.75 (s, 1H), 6.16 (ddd, *J* = 17.2, 10.4, 6.8 Hz, 1H), 5.41 (t, *J* = 9.2 Hz, 2H), 5.31 (d, *J* = 17.2 Hz, 1H), 4.06 (qd, *J* = 12.0, 5.2 Hz, 2H), 3.50 (s, 3H), 3.33 (s, 3H); ¹³**C NMR** (150 MHz, CDCl₃) δ 155.4, 151.5, 149.0, 141.0, 132.6, 120.5, 106.9, 64.2, 61.8, 29.9, 28.2; **HRMS** (ESI-TOF): exact mass calcd for C₁₁H₁₅N₄O₃ [M+H]⁺ requires m/z 251.1139, found m/z 251.1138.

(S)-3-benzoyl-1-(1-hydroxybut-3-en-2-yl)-5-methylpyrimidine-2,4(1H,3H)-dione (5a)

White solid; m.p. 49.5-50.8 °C; 57.0 mg, 95% yield, 92% ee; $[\alpha]_D^{23} = -1.85$ (c = 0.505, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 13.050 min (minor), 14.757 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 7.95 – 7.86 (m, 2H), 7.66 – 7.60 (m, 1H), 7.50 – 7.45 (m, 2H), 7.22 (q, *J* = 1.2 Hz, 1H), 5.88 (ddd, *J* = 17.2, 10.8, 5.6 Hz, 1H), 5.41 (dd, *J* = 10.8, 1.2 Hz, 1H), 5.33 (dd, *J* = 17.2, 1.2 Hz, 1H), 5.14 – 5.08 (m, 1H), 3.91 – 3.78 (m, 2H), 2.75 (brs, 1H), 1.92 (d, *J* = 1.2 Hz, 3H); ¹³**C NMR** (150 MHz, CDCl₃) δ 169.3, 163.0, 150.6, 138.4, 135.2, 132.0, 131.6, 130.5, 129.3, 120.6, 110.7, 62.7, 59.0, 12.6; **HRMS** (ESI-TOF): exact mass calcd for C₁₆H₁₆N₂NaO₄ [M+Na]⁺ requires m/z 323.1002, found m/z 323.0998. *tert*-butyl (*S*)-3-(1-hydroxybut-3-en-2-yl)-5-methyl-2,6-dioxo-3,6-dihydropyrimidine-1(2*H*)carboxylate (5b)

Colorless oil; 53.2 mg, 90% yield, 90% ee; $[\alpha]_D^{23} = -12.02$ (c = 0.560, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 80/20, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 11.653 min (minor), 13.247 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 7.14 (d, J = 1.2 Hz, 1H), 5.91 (ddd, J = 17.2, 10.8, 5.6 Hz, 1H), 5.42 (ddd, J = 10.8, 1.6, 0.8 Hz, 1H), 5.35 (ddd, J = 17.2, 1.6, 0.8 Hz, 1H), 5.15 – 5.10 (m, 1H), 3.91 (h, J = 6.8, 6.0 Hz, 2H), 2.80 (s, 1H), 1.89 (d, J = 1.2 Hz, 3H), 1.59 (s, 9H); ¹³**C NMR** (150 MHz, CDCl₃) δ 161.6, 149.7, 148.3, 138.2, 132.1, 120.7, 110.2, 87.0, 62.7, 59.2, 27.5, 12.6; **HRMS** (ESI-TOF): exact mass calcd for C₁₄H₂₀N₂NaO₅ [M+Na]⁺ requires m/z 319.1264, found m/z 319.1258.

(S)-3-benzoyl-1-(1-hydroxybut-3-en-2-yl)pyrimidine-2,4(1*H*,3*H*)-dione (5c)



Colorless oil; 54.9 mg, 96% yield, 89% ee; $[\alpha]_D^{23} = -6.27$ (c = 0.765, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 60/40, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 11.888 min (minor), 14.988 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.00 – 7.87 (m, 2H), 7.67 – 7.62 (m, 1H), 7.52 – 7.46 (m, 2H), 7.43 (d, J = 8.0 Hz, 1H), 5.91 (ddd, J = 17.2, 10.7, 5.5 Hz, 1H), 5.78 (d, J = 8.0 Hz, 1H), 5.47 – 5.42 (m, 1H), 5.40 – 5.34 (m, 1H), 5.18 – 5.11 (m, 1H), 3.88 (qd, J = 10.8, 10.4, 6.8 Hz, 2H), 2.68 (s, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 169.0, 162.4, 150.5, 142.9, 135.3, 131.8, 131.5, 130.6, 129.4, 121.0, 101.9, 62.7, 59.2; **HRMS** (ESI-TOF): exact mass calcd for C₁₅H₁₄N₂NaO₄ [M+Na]⁺ requires m/z 309.0846, found m/z 309.0842.

(S)-3-benzoyl-5-ethyl-1-(1-hydroxybut-3-en-2-yl)pyrimidine-2,4(1H,3H)-dione (5d)



Colorless oil; 59.6 mg, 95% yield, 93% ee; $[\alpha]_D^{23} = -1.04$ (c = 0.575, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 80/20, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 19.610 min

(minor), 21.112 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 7.91 (d, J = 7.8 Hz, 2H), 7.63 (t, J = 7.2 Hz, 1H), 7.48 (t, J = 7.8 Hz, 2H), 7.18 (s, 1H), 5.90 (ddd, J = 16.8, 10.8, 5.4 Hz, 1H), 5.42 (d, J = 10.8 Hz, 1H), 5.34 (d, J = 17.4 Hz, 1H), 5.12 (q, J = 5.4 Hz, 1H), 3.89 (dd, J = 11.4, 3.6 Hz, 1H), 3.87 – 3.81 (m, 1H), 2.63 (s, 1H), 2.36 (qd, J = 7.8, 3.0 Hz, 2H), 1.12 (t, J = 7.2 Hz, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 169.4, 162.6, 150.6, 137.7, 135.2, 132.0, 131.7, 130.5, 129.3, 120.6, 116.5, 62.8, 59.1, 20.3, 12.8; **HRMS** (ESI-TOF): exact mass calcd for C₁₇H₁₉N₂O₄ [M+H]⁺ requires m/z 315.1339, found m/z 315.1334.

(S)-3-benzoyl-1-(1-hydroxybut-3-en-2-yl)-5-methoxypyrimidine-2,4(1H,3H)-dione (5e)



Colorless oil; 58.8 mg, 93% yield, 94% ee; $[\alpha]_D^{23} = -4.31$ (c = 0.650, CHCl₃); **HPLC** CHIRALCEL AS, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 11.307 min (major), 14.690 min (minor); ¹**H NMR** (600 MHz, CDCl₃) δ 7.91 (d, *J* = 7.8 Hz, 2H), 7.64 (t, *J* = 7.8 Hz, 1H), 7.48 (t, *J* = 7.8 Hz, 2H), 7.02 (s, 1H), 5.89 (ddd, *J* = 16.8, 10.8, 5.4 Hz, 1H), 5.43 (d, *J* = 10.8 Hz, 1H), 5.35 (d, *J* = 17.8 Hz, 1H), 5.17 – 5.12 (m, 1H), 3.90 (dd, *J* = 12.0, 4.2 Hz, 1H), 3.85 (dd, *J* = 12.0, 7.2 Hz, 1H), 3.72 (s, 3H), 2.83 (s, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 168.4, 158.7, 149.2, 136.1, 135.4, 131.9, 131.4, 130.6, 129.3, 123.0, 120.7, 62.7, 59.1, 58.1; **HRMS** (ESI-TOF): exact mass calcd for C₁₆H₁₆N₂NaO₅ [M+Na]⁺ requires m/z 339.0951, found m/z 339.0946.

(S)-3-benzoyl-5-fluoro-1-(1-hydroxybut-3-en-2-yl)pyrimidine-2,4(1H,3H)-dione (5f)



Colorless oil; 57.6 mg, 95% yield, 90% ee; $[\alpha]_D^{23} = -2.35$ (c = 0.510, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 80/20, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 21.060 min (minor), 22.550 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.00 – 7.84 (m, 2H), 7.70 – 7.64 (m, 1H), 7.57 (d, *J* = 6.0 Hz, 1H), 7.54 – 7.47 (m, 2H), 5.91 (ddd, *J* = 17.6, 10.8, 5.6 Hz, 1H), 5.49 (dd, *J* = 10.8, 1.6 Hz, 1H), 5.41 (dd, *J* = 17.2, 1.6 Hz, 1H), 5.18 (dtt, *J* = 5.6, 4.0, 1.6 Hz, 1H), 3.96 (dd, *J* = 12.0, 4.0 Hz, 1H), 3.88 (dd, *J* = 12.0, 6.4 Hz, 1H), 2.45 (s, 1H); ¹³**C NMR** (100 MHz, CDCl₃) δ 167.5, 156.3 (*J*_{C-F}= 27.0 Hz), 149.1, 139.9 (*J*_{C-F}= 238.0 Hz), 135.7, 131.3, 131.1, 130.7, 129.5, 127.2

 $(J_{C-F}= 33.0 \text{ Hz})$, 121.5, 62.7, 59.0; ¹⁹F NMR (376 MHz, CDCl₃) δ -164.58 (s); HRMS (ESI-TOF): exact mass calcd for C₁₅H₁₃FN₂NaO₄ [M+Na]⁺ requires m/z 327.0752, found m/z 327.0750.

(S)-3-benzoyl-5-chloro-1-(1-hydroxybut-3-en-2-yl)pyrimidine-2,4(1H,3H)-dione (5g)

White waxy solid; m.p. 65.0-70.0 °C; 59.0 mg, 92% yield, 88% ee; $[\alpha]_D^{23} = -6.38$ (c = 0.690, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 9.748 min (minor), 10.738 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 7.92 (d, *J* = 7.8 Hz, 2H), 7.71 – 7.63 (m, 2H), 7.50 (t, *J* = 7.8 Hz, 2H), 5.93 (ddd, *J* = 16.8, 10.8, 5.4 Hz, 1H), 5.51 (d, *J* = 10.8 Hz, 1H), 5.43 (d, *J* = 17.4 Hz, 1H), 5.19 (q, *J* = 6.0 Hz, 1H), 3.97 (dt, *J* = 12.0, 3.6 Hz, 1H), 3.92 (dt, *J* = 11.4, 4.8 Hz, 1H), 2.29 (s, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 167.8, 158.3, 149.6, 139.8, 135.6, 131.4, 131.1, 130.7, 129.4, 121.7, 108.6, 62.7, 59.4; **HRMS** (ESI-TOF): exact mass calcd for C₁₅H₁₃ClN₂NaO₄ [M+Na]⁺ requires m/z 343.0456, found m/z 343.0452.

(S)-4-(diBocamino)-1-(1-hydroxybut-3-en-2-yl)pyrimidin-2(1H)-one (5h)



White solid; m.p. 118.2-119.5 °C; 72.2 mg, 95% yield, 84% ee; $[\alpha]_D^{23} = -41.10$ (c = 0.725, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 60/40, flow rate = 0.5 mL/min, λ = 256 nm, retention time: 12.065 min (minor), 13.032 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 7.69 (d, *J* = 7.2 Hz, 1H), 7.04 (d, *J* = 7.2 Hz, 1H), 5.98 (ddd, *J* = 17.2, 10.8, 5.6 Hz, 1H), 5.37 (d, *J* = 10.8 Hz, 1H), 5.31 (d, *J* = 17.2 Hz, 1H), 5.27 – 5.20 (m, 1H), 4.01 – 3.86 (m, 2H), 3.67 (s, 1H), 1.54 (s, 18H); ¹³**C NMR** (150 MHz, CDCl₃) δ 161.9, 155.9, 149.7, 147.2, 132.8, 120.3, 96.3, 85.1, 62.5, 61.1, 27.8; **HRMS** (ESI-TOF): exact mass calcd for C₁₈H₂₇N₃NaO₆ [M+Na]⁺ requires m/z 404.1792, found m/z 404.1789.

(S)-1-(1-hydroxybut-3-en-2-yl)pyridin-2(1H)-one (7a)



White solid; m.p. 88.2-90.1 °C; 55.4 mg, 84% yield, 75% ee; $[\alpha]_D^{23} = -86.90$ (c = 0.290, CHCl₃);

HPLC CHIRALCEL IG, *n*-hexane/2-propanol = 80/20, flow rate = 0.4 mL/min, λ = 240 nm, retention time: 22.160 min (minor), 23.753 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 7.37 (d, *J* = 6.6 Hz, 1H), 7.30 (t, *J* = 7.2 Hz, 1H), 6.53 (d, *J* = 9.0 Hz, 1H), 6.21 (t, *J* = 6.6 Hz, 1H), 5.97 (ddd, *J* = 16.8, 10.8, 5.4 Hz, 1H), 5.54 (q, *J* = 5.4 Hz, 1H), 5.35 (d, *J* = 10.8 Hz, 1H), 5.25 (d, *J* = 17.4 Hz, 1H), 4.10 (t, *J* = 6.0 Hz, 1H), 3.96 (dt, *J* = 12.0, 4.2 Hz, 1H), 3.89 (dt, *J* = 12.0, 6.0 Hz, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 163.3, 139.6, 135.8, 133.3, 120.6, 119.6, 106.6, 63.3, 59.4; **HRMS** (ESI-TOF): exact mass calcd for C₉H₁₁NNaO₂ [M+Na]⁺ requires m/z 188.0682, found m/z 188.0677.

(S)-1-(1-hydroxybut-3-en-2-yl)indoline-2,3-dione (7b)



Red solid; m.p. 102.7-104.5 °C; 57.2 mg, 66% yield, 96% ee; $[\alpha]_D^{23} = 58.00$ (c = 0.500, CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol = 90/10, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 17.402 min (major), 20.738 min (minor); ¹**H NMR** (600 MHz, CDCl₃) δ 7.61 (d, *J* = 7.2 Hz, 1H), 7.55 (t, *J* = 7.8 Hz, 1H), 7.12 (t, *J* = 7.8 Hz, 1H), 6.99 (d, *J* = 7.8 Hz, 1H), 6.02 (ddd, *J* = 16.6, 10.8, 5.8 Hz, 1H), 5.37 (d, *J* = 10.8 Hz, 1H), 5.33 (d, *J* = 17.4 Hz, 1H), 4.79 (q, *J* = 6.0 Hz, 1H), 4.21 (t, *J* = 10.2 Hz, 1H), 4.05 (dd, *J* = 12.0, 4.2 Hz, 1H), 2.85 (s, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 183.0, 158.9, 150.8, 138.3, 131.1, 125.7, 124.0, 119.5, 118.1, 112.1, 61.9, 58.2; **HRMS** (ESI-TOF): exact mass calcd for C₁₂H₁₁NNaO₃ [M+Na]⁺ requires m/z 240.0631, found m/z 240.0628.

(S)-2-(1-hydroxybut-3-en-2-yl)isoindoline-1,3-dione (7c)



White solid; m.p. 64.9-66.6 °C; 70.3 mg, 81% yield, 96% ee; $[\alpha]_D^{23} = -62.22$ (c = 0.585, CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 5.817 min (minor), 6.797 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 7.78 (dd, J = 5.6, 2.8 Hz, 2H), 7.70 – 7.65 (m, 2H), 6.11 (ddd, J = 17.2, 10.4, 6.8 Hz, 1H), 5.26 (dt, J = 4.8, 1.2 Hz, 1H), 5.22 (p, J = 1.2 Hz, 1H), 4.88 (dddt, J = 8.4, 7.2, 4.8, 1.2 Hz, 1H), 4.12 (dt, J = 11.6, 8.0 Hz, 1H), 3.91 (dt, J = 11.6, 4.0 Hz, 1H), 3.06 (dd, J = 8.0, 4.0 Hz, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 168.6, 134.2, 132.1, 131.8, 123.4, 118.9, 62.7, 56.0; **HRMS** (ESI-TOF): exact mass calcd for C₁₂H₁₁NNaO₃ [M+Na]⁺ requires m/z 240.0631, found m/z 240.0629.

(S)-2-(7H-pyrrolo[2,3-d]pyrimidin-7-yl)but-3-en-1-ol (7d)



Colorless oil; 59.7 mg, 79% yield, 92% ee; $[\alpha]_D^{23} = -59.48$ (c = 0.575, CHCl₃); **HPLC** CHIRALCEL IG, *n*-hexane/2-propanol = 80/20, flow rate = 0.7 mL/min, $\lambda = 256$ nm, retention time: 16.857 min (minor), 18.233 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 8.79 (s, 1H), 8.68 (s, 1H), 7.32 (d, J =3.6 Hz, 1H), 6.54 (d, J = 3.6 Hz, 1H), 6.14 (ddd, J = 16.8, 10.8, 6.0 Hz, 1H), 5.31 (p, J = 4.2 Hz, 1H), 5.27 (d, J = 10.8 Hz, 1H), 5.02 (d, J = 17.4 Hz, 2H), 4.15 (dd, J = 12.0, 6.6 Hz, 1H), 4.10 (dd, J = 12.0, 3.6 Hz, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 150.7, 150.1, 149.3, 133.8, 129.0, 119.3, 118.5, 100.0, 64.0, 60.8; **HRMS** (ESI-TOF): exact mass calcd for C₁₀H₁₂N₃O [M+H]⁺ requires m/z 190.0975, found m/z 190.0975.

(S)-2-(1-hydroxybut-3-en-2-yl)isoquinolin-1(2H)-one (7e)



Colorless oil; 55.0 mg, 64% yield, 79% ee; $[\alpha]_D^{23} = -69.70$ (c = 0.505, CHCl₃); **HPLC** CHIRALCEL IG, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 8.133 min (minor), 9.602 min (major); ¹**H NMR** (400 MHz, CDCl₃) δ 8.36 (d, J = 7.6 Hz, 1H), 7.60 (ddd, J =8.0, 6.8, 1.2 Hz, 1H), 7.50 – 7.40 (m, 2H), 7.15 (d, J = 7.6 Hz, 1H), 6.52 (d, J = 7.6 Hz, 1H), 6.00 (ddd, J = 17.2, 10.8, 5.2 Hz, 1H), 5.67 (dtt, J = 6.8, 5.2, 2.0 Hz, 1H), 5.35 (ddd, J = 10.8, 2.0, 0.8 Hz, 1H), 5.26 (ddd, J = 17.2, 2.0, 0.8 Hz, 1H), 4.04 (dd, J = 11.6, 4.8 Hz, 1H), 3.96 (t, J = 9.6 Hz, 1H), 3.62 (s, 1H); ¹³**C NMR** (100 MHz, CDCl₃) δ 163.0, 136.8, 133.5, 132.5, 129.1, 128.1, 127.0, 126.0, 125.9, 119.3, 106.7, 63.6, 59.1; **HRMS** (ESI-TOF): exact mass calcd for C₁₃H₁₄NO₂ [M+H]⁺ requires m/z 216.1019, found m/z 216.1021.

(S)-3-(1-hydroxybut-3-en-2-yl)quinazolin-4(3H)-one (7f)



White solid; m.p. 79.9-81.5 °C; 71.7 mg, 83% yield, 88% ee; $[\alpha]_D^{23} = -75.96$ (c = 0.545, CHCl₃);

HPLC CHIRALCEL IG, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 8.798 min (major), 10.423 min (minor); ¹**H** NMR (400 MHz, CDCl₃) δ 8.11 (s, 1H), 8.07 (dd, *J* = 8.0, 1.6 Hz, 1H), 7.63 (ddd, *J* = 8.4, 7.2, 1.6 Hz, 1H), 7.51 (d, *J* = 7.8 Hz, 1H), 7.35 (ddd, *J* = 8.0, 7.2, 1.2 Hz, 1H), 6.08 (ddd, *J* = 17.2, 10.8, 6.0 Hz, 1H), 5.47 – 5.39 (m, 2H), 5.36 (ddd, *J* = 17.2, 1.6, 0.8 Hz, 1H), 4.30 (s, 1H), 4.09 (qd, *J* = 12.0, 5.2 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 161.2, 147.0, 145.9, 134.5, 132.7, 127.4, 126.9, 126.8, 121.6, 120.4, 62.7, 58.6; **HRMS** (ESI-TOF): exact mass calcd for C₁₂H₁₃N₂O₂ [M+H]⁺ requires m/z 217.0972, found m/z 217.0971. (*S*)-1-(1-hydroxybut-3-en-2-yl)quinoxalin-2(1*H*)-one (7g)



White solid; m.p. 112.5-114.4 °C; 82.0 mg, 95% yield, 99% ee; $[\alpha]_D^{23} = -77.04$ (c = 0.540, CHCl₃); **HPLC** CHIRALCEL AS, *n*-hexane/2-propanol = 80/20, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 7.980 min (minor), 9.965 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 8.20 (s, 1H), 7.85 (d, *J* = 7.8 Hz, 1H), 7.53 – 7.49 (m, 2H), 7.33 (ddd, *J* = 8.4, 5.4, 2.4 Hz, 1H), 6.20 (ddd, *J* = 17.4, 10.8, 4.8 Hz, 1H), 5.71 (s, 1H), 5.35 – 5.32 (m, 1H), 5.23 – 5.19 (m, 1H), 4.31 (dt, *J* = 12.0, 6.6 Hz, 1H), 4.17 (d, *J* = 9.4 Hz, 1H), 3.51 (s, 1H); ¹³**C NMR** (150 MHz, CDCl₃) δ 155.9, 150.5, 134.2, 132.4, 132.2, 131.0, 130.8, 124.1, 118.6, 115.2, 62.4, 59.7; **HRMS** (ESI-TOF): exact mass calcd for C₁₂H₁₂N₂NaO₂ [M+Na]⁺ requires m/z 239.0791, found m/z 239.0789.

(S)-1-(1-hydroxybut-3-en-2-yl)-5-methylpyrimidine-2,4(1H,3H)-dione (5aa)



White solid; m.p. 139.5-141.2 °C; 37.7 mg, 96% yield, 92% ee; $[\alpha]_D^{23} = -2.95$ (c = 0.745, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 11.372 min (minor), 12.470 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 9.99 (s, 1H), 7.10 (d, *J* = 1.2 Hz, 1H), 5.91 (ddd, *J* = 17.4, 10.8, 5.4 Hz, 1H), 5.43 – 5.39 (m, 1H), 5.35 (dd, *J* = 17.4, 1.2 Hz, 1H), 5.21 – 5.17 (m, 1H), 4.00 (dt, *J* = 12.0, 4.8 Hz, 1H), 3.89 (dt, *J* = 12.0, 6.6 Hz, 1H), 3.71 (t, *J* = 6.0 Hz, 1H), 1.86 (d, *J* = 1.2 Hz, 3H); ¹³**C NMR** (150 MHz, CDCl₃) δ 164.4, 152.1, 138.7, 132.4, 120.4, 110.6, 62.6, 59.2, 12.6; **HRMS** (ESI-TOF): exact mass calcd for C₉H₁₂N₂NaO₃ [M+Na]⁺ requires m/z 219.0740, found m/z 219.0739. tert-butyl (S)-(9-(1-hydroxybut-3-en-2-yl)-9H-purin-6-yl)carbamate (3ga)



White solid; m.p. 118.2-119.6 °C; 58.0 mg, 95% yield; ¹H NMR (600 MHz, CDCl₃) δ 8.65 (s, 1H), 8.28 (s, 1H), 7.99 (s, 1H), 6.19 (ddd, J = 16.8, 10.8, 6.6 Hz, 1H), 5.38 (d, J = 10.8 Hz, 1H), 5.19 – 5.13 (m, 2H), 4.60 (s, 1H), 4.24 (dd, J = 12.0, 6.0 Hz, 1H), 4.15 (dd, J = 12.6, 3.0 Hz, 1H), 1.55 (s, 9H); ¹³C NMR (150 MHz, CDCl₃) δ 152.5, 150.7, 150.1, 149.9, 142.7, 132.8, 121.8, 119.7, 82.5, 63.8, 61.4, 28.3; HRMS (ESI-TOF): exact mass calcd for C₁₄H₂₀N₅O₃ [M+H]⁺ requires m/z 306.1561, found m/z 306.1557.

(S)-2-(6-amino-9H-purin-9-yl)but-3-en-1-ol (3gb)



White solid; m.p. 163.2-164.7 °C; 39.1 mg, 95% yield, 94% ee; $[\alpha]_D^{20} = -55.28$ (c = 0.615, CH₃OH); **HPLC** CHIRALCEL IG, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 17.220 min (minor), 19.225 min (major); ¹**H NMR** (600 MHz, Methanol-*d*₄) δ 8.22 (s, 1H), 8.19 (s, 1H), 6.26 (ddd, *J* = 17.4, 10.2, 6.6 Hz, 1H), 5.33 (dt, *J* = 10.8, 1.2 Hz, 1H), 5.23 – 5.20 (m, 1H), 5.17 (dt, *J* = 17.4, 1.2 Hz, 1H), 4.13 (dd, *J* = 12.0, 7.2 Hz, 1H), 3.98 (dd, *J* = 12.0, 4.2 Hz, 1H); ¹³**C NMR** (150 MHz, Methanol-*d*₄) δ 157.3, 153.5, 150.7, 142.0, 134.7, 120.0, 119.3, 64.0, 61.2; **HRMS** (ESI-TOF): exact mass calcd for C₉H₁₂N₅O [M+H]⁺ requires m/z 206.1036, found m/z 206.1035.

(S)-9-(1-(allyloxy)but-3-en-2-yl)-9H-purin-6-diBocamine (3gc)



Colorless oil; 80.2 mg, 90% yield; ¹**H NMR** (600 MHz, CDCl₃) δ 8.82 (s, 1H), 8.24 (s, 1H), 6.21 (ddd, *J* = 16.8, 10.8, 6.0 Hz, 1H), 5.79 – 5.70 (m, 1H), 5.43 – 5.40 (m, 1H), 5.35 (dd, *J* = 10.8, 1.2 Hz, 1H), 5.19 (dd, *J* = 16.8, 1.2 Hz, 1H), 5.16 (dd, *J* = 17.4, 1.8 Hz, 1H), 5.12 (dd, *J* = 10.2, 1.2 Hz, 1H), 3.98 (dd, *J* = 10.2, 6.0 Hz, 1H), 3.96 – 3.91 (m, 2H), 3.82 (dd, *J* = 10.2, 4.2 Hz, 1H), 1.43 (s,

18H); ¹³C NMR (100 MHz, CDCl₃) δ 153.3, 151.9, 150.6, 150.3, 144.7, 133.7, 132.8, 128.8, 119.5,
117.9, 83.7, 72.4, 70.6, 57.2, 27.9; HRMS (ESI-TOF): exact mass calcd for C₂₂H₃₁N₅NaO₅
[M+Na]⁺ requires m/z 468.2217, found m/z 468.2221.

(S)-9-(3,6-dihydro-2H-pyran-3-yl)-9H-purin-6-diBocamine (3gd)

Colorless oil; 73.4 mg, 88% yield, 94% ee; $[\alpha]_D^{23} = 53.55$ (c = 0.600, CHCl₃); **HPLC** CHIRALCEL IE, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 22.153 min (minor), 23.503 min (major); ¹**H** NMR (600 MHz, CDCl₃) δ 8.84 (s, 1H), 8.26 (s, 1H), 6.31 (ddt, *J* = 10.2, 3.6, 1.8 Hz, 1H), 6.05 – 6.02 (m, 1H), 5.21 – 5.18 (m, 1H), 4.38 (dddd, *J* = 17.2, 3.6, 2.4, 1.2 Hz, 1H), 4.27 (dq, *J* = 17.2, 2.4 Hz, 1H), 4.04 (dt, *J* = 12.0, 1.8 Hz, 1H), 4.01 (dd, *J* = 12.0, 3.0 Hz, 1H), 1.45 (s, 18H); ¹³C NMR (150 MHz, CDCl₃) δ 153.0, 152.0, 150.7, 150.4, 144.4, 133.5, 129.1, 121.4, 83.8, 68.5, 65.4, 47.4, 27.9; **HRMS** (ESI-TOF): exact mass calcd for C₂₀H₂₈N₅O₅ [M+H]⁺ requires m/z 418.2085, found m/z 418.2081.

(S)-9-(3,6-dihydro-2H-pyran-3-yl)-9H-purin-6-amine (3ge)



White solid; m.p. 192.6-194.4 °C; 41.2 mg, 95% yield, 95% ee; $[\alpha]_D^{20} = 92.36$ (c = 0.550, CH₃OH); **HPLC** CHIRALCEL IF, *n*-hexane/2-propanol = 50/50, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 24.745 min (minor), 32.330 min (major); ¹**H NMR** (400 MHz, Methanol-*d*₄) δ 8.22 (s, 1H), 8.14 (s, 1H), 6.34 (ddt, *J* = 10.0, 3.2, 1.6 Hz, 1H), 6.06 (ddt, *J* = 10.0, 4.8, 2.0 Hz, 1H), 5.09 (s, 1H), 4.39 – 4.33 (m, 1H), 4.28 – 4.22 (m, 1H), 4.00 (t, *J* = 3.2 Hz, 2H); ¹³**C NMR** (150 MHz, CDCl₃) δ 157.3, 153.7, 150.1, 141.7, 134.6, 122.1, 120.1, 69.2, 66.4, 48.9; **HRMS** (ESI-TOF): exact mass calcd for C₁₀H₁₂N₅O [M+H]⁺ requires m/z 218.1036, found m/z 218.1039.

(3S,4R,5R)-5-(6-diBocamino-9H-purin-9-yl)tetrahydro-2H-pyran-3,4-diol (3gf)



White solid; m.p. 124.6-126.3 °C; 79.3 mg, 88% yield, 93% ee, >20:1 dr; $[\alpha]_D^{20} = 37.62$ (c = 0.505,

CHCl₃); **HPLC** CHIRALCEL OD, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, λ = 256 nm, retention time: 5.005 min (minor), 8.857 min (major); ¹**H NMR** (600 MHz, CDCl₃) δ 8.77 (s, 1H), 8.05 (s, 1H), 4.78 (td, *J* = 9.6, 4.8 Hz, 1H), 4.46 (dt, *J* = 9.6, 4.2 Hz, 1H), 4.11 (dd, *J* = 11.4, 4.8 Hz, 1H), 4.08 – 3.99 (m, 3H), 3.97 (s, 1H), 3.71 (dd, *J* = 12.6, 1.8 Hz, 1H), 3.52 (s, 1H), 1.46 (s, 18H); ¹³**C NMR** (150 MHz, CDCl₃) δ 153.6, 151.9, 150.9, 150.4, 145.2, 129.2, 84.5, 70.5, 69.4, 68.4, 67.4, 56.2, 28.0; **HRMS** (ESI-TOF): exact mass calcd for C₂₀H₃₀N₅O₇ [M+H]⁺ requires m/z 452.2140, found m/z 452.2137.

(3S,4R,5R)-5-(6-amino-9H-purin-9-yl)tetrahydro-2H-pyran-3,4-diol (3gg)



White solid; m.p. 265.5-267.2 °C; 46.0 mg, 92% yield; ¹H NMR (600 MHz, Methanol- d_4) δ 8.18 (s, 2H), 4.78 (td, J = 10.2, 4.8 Hz, 1H), 4.46 (dd, J = 10.2, 3.0 Hz, 1H), 4.06 (dd, J = 11.4, 4.8 Hz, 1H), 4.02 – 3.92 (m, 3H), 3.74 (dd, J = 12.0, 1.2 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 157.3, 153.5, 151.2, 142.4, 120.4, 72.0, 70.8, 70.1, 68.8, 56.4; HRMS (ESI-TOF): exact mass calcd for C₁₀H₁₄N₅O₃ [M+H]⁺ requires m/z 252.1091, found m/z 252.1093.

(S)-2-(6-diBocamino-9H-purin-9-yl)but-3-en-1-yl 4-methylbenzenesulfonate (3gh)



White solid; m.p. 52.2-54.7 °C; 104.0 mg, 93% yield; ¹H NMR (400 MHz, CDCl₃) δ 8.62 (s, 1H), 8.03 (s, 1H), 7.51 (d, J = 8.0 Hz, 2H), 7.22 (d, J = 8.0 Hz, 2H), 6.18 (ddd, J = 17.2, 10.4, 6.4 Hz, 1H), 5.44 (d, J = 10.4 Hz, 1H), 5.37 (q, J = 6.4 Hz, 1H), 5.31 (d, J = 17.2 Hz, 1H), 4.66 (dd, J = 10.8, 7.2 Hz, 1H), 4.43 (dd, J = 10.8, 4.0 Hz, 1H), 2.40 (s, 3H), 1.47 (s, 18H); ¹³C NMR (100 MHz, CDCl₃) δ 152.8, 151.8, 150.7, 150.5, 145.7, 144.0, 131.6, 130.1, 128.9, 127.8, 121.8, 84.0, 68.3, 57.4, 27.9, 21.7; HRMS (ESI-TOF): exact mass calcd for C₂₆H₃₄N₅O₇S [M+H]⁺ requires m/z 560.2173, found m/z 560.2172.

(2R,3R)-2-(6-diBocamino-9H-purin-9-yl)-3,4-dihydroxybutyl 4-methylbenzenesulfonate (3gi)



White solid; m.p. 128.4-129.9 °C; 109.2 mg, 92% yield, 96% ee, >20:1 dr; $[\alpha]_D^{23} = 2.17$ (c = 0.400, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 70/30, flow rate = 1.0 mL/min, $\lambda = 256$ nm, retention time: 9.662 min (major), 15.002 min (minor); ¹H NMR (600 MHz, CDCl₃) δ 8.65 (s, 1H), 8.10 (s, 1H), 7.54 (d, *J* = 8.4 Hz, 2H), 7.24 (d, *J* = 7.8 Hz, 2H), 5.06 (dt, *J* = 9.0, 4.2 Hz, 1H), 4.58 (dd, *J* = 11.4, 9.0 Hz, 1H), 4.52 (dd, *J* = 10.8, 4.8 Hz, 1H), 4.20 (td, *J* = 6.0, 4.2 Hz, 1H), 3.41 (dd, *J* = 11.2, 6.0 Hz, 1H), 3.20 (dd, *J* = 11.2, 6.6 Hz, 1H), 2.40 (s, 3H), 1.47 (s, 18H); ¹³C NMR (150 MHz, CDCl₃) δ 153.1, 151.6, 150.6, 145.7, 145.7, 131.7, 130.2, 128.5, 127.8, 84.4, 70.1, 67.7, 62.9, 57.0, 27.9, 21.7; **HRMS** (ESI-TOF): exact mass calcd for C₂₆H₃₆N₅O₉S [M+H]⁺ requires m/z 594.2228, found m/z 594.2226.

(3R,4R)-4-(6-diBocamino-9H-purin-9-yl)tetrahydrofuran-3-ol (3gj)



Colorless oil; 69.6 mg, 85% yield, 94% ee, 20:1 dr; $[\alpha]_D^{23} = -5.50$ (c = 0.800, CHCl₃); **HPLC** CHIRALCEL ID, *n*-hexane/2-propanol = 70/30, flow rate = 0.8 mL/min, λ = 256 nm, retention time: 12.100 min (major), 16.352 min (minor); ¹**H NMR** (600 MHz, CDCl₃) δ 8.87 (s, 1H), 8.12 (s, 1H), 5.02 (dt, *J* = 5.4, 3.0 Hz, 1H), 4.61 (dt, *J* = 6.0, 3.0 Hz, 1H), 4.44 (dd, *J* = 10.2, 5.4 Hz, 1H), 4.32 – 4.28 (m, 2H), 3.89 (dd, *J* = 10.2, 3.6 Hz, 2H), 1.47 (s, 18H); ¹³**C NMR** (150 MHz, CDCl₃) δ 153.3, 152.2, 150.8, 150.7, 142.8, 128.9, 84.2, 76.6, 74.1, 70.0, 63.0, 28.0; **HRMS** (ESI-TOF): exact mass calcd for C₁₉H₂₇N₅NaO₆ [M+Na]⁺ requires m/z 444.1854, found m/z 444.1857.

7. Copies of NMR spectra for the products







S27















S34












- 500

- 0 - -500











S41







S43

80

70 60

210 200 190 180 170 160 150 140 130 120 110 100 90 f1 (ppm) -20000

0

-10

50 40 30 20 10









S46

















- 800 - 600 - 400 - 200 - 0 - -200

S50











S54







~ 157.31 ~ 153.53 ~ 150.69 ~ 141.97 ~ 134.68 120.05 119.34 .63.95 .61.18 .61.18 .49.43 .49.28 .49.00 .48.86 .48.72 .48.57 NH_2 650 600 но 550 3gb 500 450 400 350 300 250 200 150 100 50 - 0 -50 20 210 200 190 180 170 160 150 140 130 120 110 100 90 f1 (ppm) 80 70 60 50 40 30 10 0 -10 20



210 200 190 180 170 160 150 140 130 120 110 100 90 80 70 60 50 40 30 20 10 0 -10 f1 (ppm)















- 157.28 ~ 153.48 ~ 151.17 - 142.37 1000 - 120.40 71.99 70.82 68.85 56.37 49.43 49.43 49.43 49.14 49.14 48.72 48.72 NH₂ 900 ОН 800 OH Ċ 3gg 700 600 500 400 300 200 100 - 0 20 210 200 190 180 170 160 150 140 130 120 110 100 90 80 70 60 50 40 30 20 10 f1 (ppm) 0 -10



S62









8. Copies of HPLC spectra for racemic and chiral products





Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	16.395	234.905	460.178	98.54	98.19
2	17.837	3.480	8.492	1.46	1.81
Total:		238.385	468.670	100.00	100.00



你刀印木								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	13.112	60.805	132.222	50.20	58.49			
2	17.760	60.323	93.827	49.80	41.51			
Total:		121.128	226.050	100.00	100.00			



你刀垣禾								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	13.320	7.561	17.654	2.48	3.96			
2	17.232	297.434	427.810	97.52	96.04			
Total:		304.995	445.464	100.00	100.00			





积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	21.165	835.005	1257.164	97.77	97.64
2	23.288	19.036	30.342	2.23	2.36
Total:		854.042	1287.507	100.00	100.00



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	13.952	65.324	146.815	49.81	52.56
2	15.745	65.823	132.537	50.19	47.44
Total:		131.147	279.352	100.00	100.00



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	13.873	1471.461	2707.963	96.07	95.37			
2	15.707	60.210	131.486	3.93	4.63			
Total:		1531.671	2839.449	100.00	100.00			



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	10.800	12.853	42.513	49.76	51.32
2	11.783	12.976	40.328	50.24	48.68
Total:		25.829	82.841	100.00	100.00



Total:		214.157	692.852	100.00	100.00
2	11.790	6.884	22.991	3.21	3.32
1	10.732	207.273	669.861	96.79	96.68
Peak	Retention Time min	Area mAU*min	Height mAU	Area %	Height %
积分约	吉果				



Total:		135.459	408.564	100.00	100.00
2	12.773	68.170	192.383	50.33	47.09
1	11.652	67.289	216.181	49.67	52.91
	min	mAU*min	mAU	%	%
r cak	Relention Time	Alea	пеідпі	Alea	Height



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.492	921.928	2602.760	96.97	96.82
2	12.808	28.784	85.352	3.03	3.18
Total:		950.711	2688.111	100.00	100.00



小刀 汨木								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	10.953	69.934	96.693	50.03	51.96			
2	13.017	69.849	89.405	49.97	48.04			
Total:		139.783	186.099	100.00	100.00			



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	11.008	32.086	46.581	3.04	3.84			
2	12.845	1022.821	1167.253	96.96	96.16			
Total:		1054.907	1213.834	100.00	100.00			




100.00

100.00

106.611

积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	23.547	16.879	20.581	2.04	2.61			
2	25.702	810.764	769.282	97.96	97.39			
Total:		827.644	789.863	100.00	100.00			



Peak	Retention Time min	Area mAU*min	Height mAU	Area %	Height %
1	14.323	87.924	164.606	49.71	55.56
2	16.192	88.963	131.657	50.29	44.44
Total:		176.887	296.263	100.00	100.00



∠ Total:	16.353	300.441	532.092	1.51	1.44			
0	10.050	4 500	7.050	4.54	1.14			
1	14,182	295,911	524,439	98.49	98.56			
	min	mAU*min	mAU	%	%			
Peak	Retention Time	Area	Height	Area	Height			
积分结果								



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	6.902	270.838	993.512	49.63	56.25
2	8.345	274.899	772.874	50.37	43.75
Total:		545.737	1766.386	100.00	100.00



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	6.885	863.523	2931.162	97.17	98.16			
2	8.505	25.136	55.071	2.83	1.84			
Total:		888.659	2986.233	100.00	100.00			



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	12.310	141.474	248.413	50.11	54.71			
2	14.330	140.836	205.656	49.89	45.29			
Total:		282.310	454.069	100.00	100.00			



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	12.292	14.956	28.214	2.48	3.23			
2	14.260	587.604	846.342	97.52	96.77			
Total:		602.560	874.556	100.00	100.00			



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	20.483	309.501	185.687	49.26	44.24
2	24.770	318.784	234.034	50.74	55.76
Total:		628.285	419.722	100.00	100.00



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	18.970	1739.579	1110.341	98.25	97.84
2	25.622	30.905	24.504	1.75	2.16
Total:		1770.484	1134.844	100.00	100.00



Total:		199.976	424.463	100.00	100.00
2	14.777	99.812	201.482	49.91	47.47
1	13.025	100.164	222.981	50.09	52.53
	min	mAU*min	mAU	%	%
Peak	Retention Time	Area	Height	Area	Height



积分结果							
Peak	Retention Time	Area	Height	Area	Height		
	min	mAU*min	mAU	%	%		
1	13.050	1.595	4.038	4.25	5.12		
2	14.757	35.927	74.819	95.75	94.88		
Total:		37.523	78.857	100.00	100.00		



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.550	150.275	207.446	50.53	57.07
2	13.470	147.137	156.064	49.47	42.93
Total:		297.411	363.510	100.00	100.00



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.653	23.414	36.713	5.25	7.72
2	13.247	422.238	438.630	94.75	92.28
Total:		445.653	475.343	100.00	100.00



100.00

100.00



1515.795

积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	11.888	1.652	4.125	5.60	7.58			
2	14.988	27.865	50.296	94.40	92.42			
Total:		29.517	54.421	100.00	100.00			





积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	19.610	14.455	27.845	3.69	4.38
2	21.112	377.396	607.486	96.31	95.62
Total:		391.851	635.331	100.00	100.00



Total:		239.857	231.116	100.00	100.00
2	14.640	120.246	104.513	50.13	45.22
1	11.457	119.610	126.604	49.87	54.78
	min	mAU*min	mAU	%	%
Peak	Retention Time	Area	Height	Area	Height



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.307	2243.472	2336.370	97.06	96.95
2	14.690	67.921	73.500	2.94	3.05
Total:		2311.393	2409.871	100.00	100.00



Total:		201.816	309.749	100.00	100.00
2	23.000	100.428	145.289	49.76	46.91
1	21.122	101.388	164.460	50.24	53.09
	min	mAU*min	mAU	%	%
Peak	Retention Time	Area	Height	Area	Height



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	21.060	33.257	59.903	5.24	6.88
2	22.550	601.410	811.266	94.76	93.12
Total:		634.667	871.169	100.00	100.00



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	9.702	415.150	1386.466	50.10	52.65
2	10.767	413.562	1246.943	49.90	47.35
Total:		828.713	2633.409	100.00	100.00



积分组	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	9.748	49.121	177.300	6.09	7.03
2	10.738	757.358	2345.581	93.91	92.97
Total:		806.478	2522.881	100.00	100.00



Total:		204.964	514.366	100.00	100.00
2	13.132	102.208	270.734	49.87	52.63
1	12.035	102.756	243.631	50.13	47.37
	min	mAU*min	mAU	%	%
Peak	Retention Time	Area	Height	Area	Height



积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	12.065	40.407	104.673	8.21	8.40			
2	13.032	452.012	1141.062	91.79	91.60			
Total:		492.419	1245.735	100.00	100.00			





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100.00

254.135

积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	22.160	63.600	107.763	12.43	14.96
2	23.753	448.263	612.519	87.57	85.04
Total:		511.864	720.282	100.00	100.00



Total:	190.182	203.803	100.00	100.00
350 350 250 250 0 250 150 300 250 150 0	$\begin{array}{c} 130.102\\ \hline \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\$	1 - 17.402	2	
-50]	10.0	15.0 [min]	20.0	25.0 28.0

积分结果								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	17.402	199.742	216.948	98.19	98.06			
2	20.738	3.687	4.283	1.81	1.94			
Total:		203.429	221.231	100.00	100.00			



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	5.773	166.552	705.659	50.09	52.22
2	6.612	165.939	645.688	49.91	47.78
Total:		332.491	1351.346	100.00	100.00



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	5.817	1.401	6.913	2.24	2.85
2	6.797	61.103	235.760	97.76	97.15
Total:		62.505	242.673	100.00	100.00



Total:		724.045	1412.335	100.00	100.00
2	18.412	360.507	667.987	49.79	47.30
1	16.930	363.538	744.348	50.21	52.70
	min	mAU*min	mAU	%	%
- Gar	Retention Time	Alea	Height	Alea	neight



枳分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	16.857	3.138	6.826	4.18	4.91
2	18.233	71.924	132.077	95.82	95.09
Total:		75.063	138.903	100.00	100.00



Total:		207.545	807.051	100.00	100.00
2	9.612	103.413	366.023	49.83	45.35
1	8.117	104.133	441.028	50.17	54.65
	min	mAU*min	mAU	%	%
Peak	Retention Time	Area	Height	Area	Height



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	8.133	20.010	85.060	10.63	12.65
2	9.602	168.248	587.338	89.37	87.35
Total:		188.257	672.398	100.00	100.00





100.00

100.00

136.317

积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	8.798	239.091	877.729	93.85	94.40
2	10.423	15.679	52.025	6.15	5.60
Total:		254.770	929.754	100.00	100.00





积分约	吉果			1	
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	7.980	1.604	4.534	0.73	1.19
2	9.965	216.949	376.140	99.27	98.81
Total:		218.553	380.674	100.00	100.00



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	10.977	140.576	209.451	49.93	55.59
2	13.203	140.986	167.341	50.07	44.41
Total:		281.562	376.792	100.00	100.00



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.000	4.210	7.193	2.47	3.58
2	13.123	166.365	193.951	97.53	96.42
Total:		170.575	201.145	100.00	100.00



Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.187	42.734	128.863	49.86	52.87
2	12.457	42.977	114.857	50.14	47.13
Total:		85.711	243.720	100.00	100.00



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	11.372	14.796	46.683	4.13	5.08
2	12.470	343.328	872.736	95.87	94.92
Total:		358.124	919.419	100.00	100.00



Peak	Retention Time	Area	Height	Area	Height		
	min	mAU*min	mAU	%	%		
1	17.010	164.110	244.684	48.98	54.43		
2	19.405	170.958	204.879	51.02	45.57		
Total:		335.069	449.562	100.00	100.00		



积分约	告果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	17.220	10.882	15.626	3.21	3.81
2	19.225	328.255	394.209	96.79	96.19
Total:		339.137	409.836	100.00	100.00



积分约	吉果	_		-	
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	21.907	81.845	125.358	49.25	52.92
2	23.648	84.331	111.540	50.75	47.08
Total:		166.176	236.897	100.00	100.00



积分约	吉果			_	
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	22.153	8.345	13.556	2.92	3.82
2	23.503	277.015	341.377	97.08	96.18
Total:		285.360	354.934	100.00	100.00





积分组	吉果	-			
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	24.745	46.606	27.319	2.50	2.95
2	32.330	1816.801	899.994	97.50	97.05
Total:		1863.407	927.314	100.00	100.00



伏万き	ā 衆				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	4.885	150.833	395.726	49.60	67.90
2	8.750	153.260	187.059	50.40	32.10
Total:		304.093	582.784	100.00	100.00



积分约	积分结果								
Peak	Retention Time	Area	Height	Area	Height				
	min	mAU*min	mAU	%	%				
1	5.005	16.426	44.164	3.42	7.37				
2	8.857	463.372	555.129	96.58	92.63				
Total:		479.798	599.293	100.00	100.00				



积分约	吉果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	9.798	25.753	55.234	26.87	38.25
2	10.883	25.653	48.983	26.77	33.92
3	15.092	22.223	28.155	23.19	19.50
4	26.563	22.210	12.036	23.17	8.33
Total:		95.839	144.408	100.00	100.00



权分约	若 果				
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	9.662	148.706	334.856	97.89	98.55
2	15.002	3.211	4.930	2.11	1.45
Total:		151.917	339.786	100.00	100.00



秋分站朱								
Peak	Retention Time	Area	Height	Area	Height			
	min	mAU*min	mAU	%	%			
1	12.185	119.459	269.271	53.00	62.87			
2	16.160	105.945	159.019	47.00	37.13			
Total:		225.403	428.290	100.00	100.00			



积分结果					
Peak	Retention Time	Area	Height	Area	Height
	min	mAU*min	mAU	%	%
1	12.100	237.663	499.750	96.80	97.34
2	16.352	7.853	13.670	3.20	2.66
Total:		245.516	513.420	100.00	100.00