Chemical Science



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View Article Online
View Journal | View Issue



Cite this: Chem. Sci., 2018, 9, 4505

Copper-catalyzed asymmetric hydrogenation of 2-substituted ketones via dynamic kinetic resolution†

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A new class of tunable heterophosphole dimeric ligands have been designed and synthesized. These ligands have enabled the first examples of Cu-catalyzed hydrogenation of 2-substituted-1-tetralones and related heteroaryl ketones *via* dynamic kinetic resolution, simultaneously creating two contiguous stereogenic centers with up to >99:1 dr and 98:2 er. The ligand-Cu complexes were isolated and characterized by single crystal X-ray, and DFT calculations revealed a novel heteroligated dimeric copper hydride transition state

Received 26th January 2018 Accepted 19th April 2018

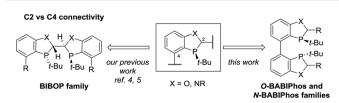
DOI: 10.1039/c8sc00434j

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Transition metals such as Pd, Pt, Rh, Ru and Ir have been widely utilized for a large variety of catalytic transformations.1 However, the long-term sustainability of such processes has become an increasing concern due to the high cost, limited abundance and observed toxicity of noble metals.2 To overcome these issues, in recent years, there has been a major endeavor in the field to explore new synthetic methods utilizing nonprecious metal catalysis.3 To this end, we have recently reported several novel methodologies employing Fe, Ni, Co and Cu-centered catalysts.4 For example, our laboratories have developed the first Cu-catalyzed asymmetric propargylation reaction of aldehydes^{4a} whereas the key to obtaining high enantioselectivities was the use of our bidentate MeO-BIBOP ligand to control the stereochemical course of the reaction. MeO-BIBOP belongs to a family of P-chiral benzooxaphosphole (BOP) ligands which have demonstrated superior abilities to catalyze a range of asymmetric transformations.5 The modular design of the ligand core allows for high tunability of steric and electronic properties of the ligand to afford the desired reactivities and stereoselectivities (Scheme 1). Encouraged by our success with the BIBOP-Cu system, we aspired to design new bidentate benzooxaphosphole ligands to promote other nonprecious metal catalyzed fundamental reactions such as asymmetric hydrogenation of ketones. Herein we report the discovery of a new class of heterophosphole dimeric ligands (BABIPhos) and their application in the first copper-catalyzed asymmetric hydrogenation of 2-substituted-1-tetralones and related heteroaryl ketones via dynamic kinetic resolution.

The Cu-catalyzed asymmetric reduction of ketones has been described in the literature using both silanes and hydrogen.6 The silane-based methods often require cryogenic temperatures and generate stoichiometric amounts of silane byproducts rendering them unattractive for scaleup.7 On the other hand, greener alternatives based on catalytic hydrogenation are more challenging and significantly less developed. Shimizu and coworkers reported the first asymmetric Cu-catalyzed ketone hydrogenation using BDPP as ligand obtaining good selectivities with ortho-substituted aryl and heteroaryl ketones.8 Beller's group described a more general system using monodentate binaphthophosphepine ligands that provided selectivities up to 77% ee.9 More recently, Hatcher and co-workers reported the use of modified BoPhoz ligands for ketone hydrogenation to afford optically enriched alcohol products with 75:25 to 98:2 er.10 As part of our long-standing interest in non-precious metal catalysis and engineering innovative catalytic systems, we

[†] Electronic supplementary information (ESI) available. CCDC 1816601 and 1816602. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8sc00434j



Scheme 1 Dimeric benzoheterophosphole ligands.

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sought to develop more efficient methods for Cu-catalyzed asymmetric ketone hydrogenation.

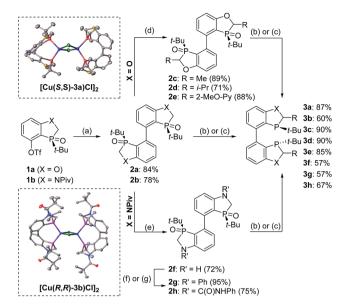
2-Aryl-1-tetralone derivatives were chosen as initial substrates for the study because the resulting chiral alcohols were required for one of our ongoing programs. Furthermore the relatively low pK_a of the keto-benzylic position offers an opportunity for dynamic kinetic resolution (DKR), creating two adjacent stereocenters in one single operation. It should be noted that no examples of Cu-catalyzed asymmetric reduction of ketones via DKR have been previously reported.11

The asymmetric hydrogenation of racemic 2-phenyl-1tetralone 4a was first evaluated using protocols reported by Shimizu,8 Beller9 and Hatcher10 (Table 1). Low conversions were observed under Shimizu's BDPP and Beller's CatASium KPh conditions. Hatcher's modified BoPhoz ligand provided 70% conversion after 24 h at 400 psi with moderate stereoselectivity of 88:12 dr and 83:17 er for the major diastereoisomer. Evaluation of chiral BIBOP ligand provided full conversion exclusively to the cis-diastereomer with 77:23 er, while the reaction with MeO-BIBOP gave less gratifying results. Phenyl and cyclohexyl BIBOP analogs exhibited no reactivity in this reaction. Despite the observed modest stereoselectivities in the initial screening, these results clearly showed that the anticipated DKR has occurred under the reaction conditions.

To further improve the reactivity and selectivity of the reaction, we decided to synthesize the alternative BOP dimer via the C4-coupling which might provide new ligands with different properties (Scheme 1).12 Our quest for this type of biaryl ligands began by exploring the reductive homocoupling of the corresponding benzooxaphosphole oxide triflate 1a (Scheme 2). After

Table 1 Cu-Catalyzed Asymmetric Hydrogenation of 2-Phenyl-1tetralone^a

100%, >99:1 dr, 91:9 ert



Scheme 2 Synthesis of O-BABIPhos and N-BABIPhos. Conditions: (a) 2a (1 equiv.), Pd₂(dba)₃ (2.5 mol%), BI-DIME (5 mol%, 2a) or DTBPF (5 mol%, 2b), Et₄NI (2 equiv.), Zn dust (3 equiv.), DMAc, 140 °C; (b) $(Me_2HSi)_2O$ (3 equiv.), $Ti(Oi-Pr)_4$ (2.3 equiv.), THF, 65 °C; (c) Et_3SiH (3 equiv.), Et_3N (3 equiv.), toluene, 110 °C; (d) LDA (2.5 equiv.); then RX (2.5 equiv.; 2c: Mel, 2d: i-Prl, 2e: 2-methoxy-6-(phenylsulfonyl)pyridine), THF, -78 °C to r.t.; (e) NaOH, MeOH/H₂O, 50 °C; (f) **2g**: PhBr, Pd₂(dba)₃ (5 mol%), RuPhos (20 mol%), NaOt-Bu (2.4 equiv.), THF, reflux; (g) 2h: PhNHCO (10 equiv.), i-Pr2NEt (3.6 equiv.), THF, reflux. BI-DIME = 3-(tert-butyl)-4-(2,6-dimethoxyphenyl)-2,3-dihydrobenzo[d][1,3]oxaphosphole, DTBPF = 1,1'-bis(di-tert-butylphosphino)ferrocene, LDA = lithium diisopropylamide, RuPhos = 2-dicyclohexylphosphino-2',6'-diisopropoxybiphenyl.

extensive investigation of the Pd-13 and Ni-catalyzed14 protocols for dimerizing aryl halides and triflates,15 it was found that under Pd-catalysis in the presence of BI-DIME, aryl triflate 1a was converted to the corresponding biaryl 2a in 84% yield. Additionally, the oxaphosphole ring of 2a was alkylated to form C2, C2'-substituted oxide analogs 2c-e. The phosphine oxides were subsequently reduced to the corresponding phosphines **3a,c-e** (O-BABIPhos) giving access to a new class of bidentate ligands. This route was demonstrated on several hundred grams scale.

With these ligands in hand, the ketone hydrogenation was carried out, and we were pleased to find that the reaction with O-BABIPhos ligand 3a provided high reactivities and selectivities up to 91:9 er of single diastereoisomer 5a (Table 1). However, modification of the R group in O-BABIPhos did not further improve the enantioselectivity. To obtain more selective ligands, a new class of dimeric azaphosphole ligands, N-BABI-Phos were designed and prepared to further modulate the electronic properties of the ligand by introducing various functionalities on the nitrogen atom (Scheme 2). Dimerization of the nitrogen-containing triflate 1b proved to be much more challenging and a wide-scope ligand screen led to the identification of DTBPF as the best ligand that gave good yield of biaryl product 2b. Further modification at nitrogen gave quick access to N-BABIPhos analogs 3f-h.

^a Conversions were determined by comparison of relative HPLC integration of alcohol to ketone at 220 nm, diastereo- and enantiomeric ratios (dr and er) were determined by chiral HPLC. b t-AmOH as solvent.

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We were delighted to find that the Cu-catalyzed hydrogenation of 2-phenyl-1-tetralone in *t*-AmOH using *N*-BABIPhos **3b** as the ligand proceeded in complete diastereoselectivity and 97 : 3 er (Table 2, entry 1). Reactions with other azaphospholes bearing different *N*-substitutions gave overall less satisfactory results (entries 2–4). *t*-AmOH proved to be the solvent of choice for this transformation. Lower reactivity was observed when performing the reactions in *t*-BuOH as solvent while i-PrOH provided good reactivity but only 92 : 8 er (entries 5–6)¹⁶. The nature of ancillary phosphine played a key role in the reactivity of our new catalytic system. Thus, no reaction was observed in the absence of ancillary phosphine or using electron-deficient triarylphosphines (entries 7–8). The source of this phenomenon was further elucidated *via* DFT calculations (*vide infra*).

Both the Cu–O-BABIPhos and Cu–N-BABIPhos complexes were obtained in quantitative yields by treatment of CuCl with a stoichiometric amount of respective ligand in dichloromethane at reflux temperature for 2 h. The molecular structures of the complexes were unambiguously characterized by single crystal X-ray crystallography¹⁷ revealing a chloride-bridged dicopper complex such as $[Cu(\mu\text{-Cl}) \ (\kappa_2\text{-}N\text{-BABIPhos})]_2$ with a distorted tetrahedral geometry at both copper(i) centers, as observed for similar Cu complexes in the literature (Scheme 2).¹⁸

The scope of the reaction was next examined (Table 3). A variety of racemic 2-aryl-1-tetralones were smoothly reduced under our standard conditions with excellent enantioselectivities (96:4 to 98:2 er, products 5a-h). Reactions with heterocycle-containing substrates were equally effective. Pyridine, furan, thiophene and pyrazole-fused cyclohexanones underwent the Cu-catalyzed hydrogenation in high stereoselectivities (96:4 to 97:3 er, products 5i-l). Furthermore, the hydrogenative DKR of 2-alkyl-1-tetralones 4m and 4n proved to be highly enantioselective with good diastereoselectivities. Substituted indanone substrate 40 was reduced under the

Table 2 Optimization of reaction conditions^a

Entry	Ligand	PAr ₃	%5 a ^b	dr^c	er ^c
1	3 b	$P(3,5-xy y)_3$	100	>99:1	97:3
2	3f 3g	$P(3,5-xylyl)_3$ $P(3,5-xylyl)_3$	50 100	>99 : 1 >99 : 1	96:4 86:14
4	3h	$P(3,5-xylyl)_3$	50	>99:1	94:6
5 ^d	3b	$P(3,5-xylyl)_3$	66	>99:1	96:8
6^e	3 b	$P(3,5-xylyl)_3$	100	>99:1	92:8
7	3b	_	0	_	_
8	3 b	$P(3,5-CF_3-Ph)_3$	1	_	_

 $[^]a$ Conditions: H_2 (450 psi), $CuCl_2$ (5 mol%), N-BABIPhos (5 mol%), PAr_3 (5 mol%), ER_4 (25 mol%), ER_4 (25 mol%), ER_4 (25 mol%), ER_4 (25 mol%), ER_4 (26 mol%), ER_4 (27 mol%), ER_4 (28 mol%), ER_4 (29 mol%), ER_4 (29 mol%), ER_4 (5 mol%), ER_4 (6 mol%), ER_4 (6 mol%), ER_4 (7 mol%), ER_4 (7 mol%), ER_4 (8 mol%), ER_4 (9 mol%), ER_4 (9 mol%), ER_4 (10 mol%),

Table 3 Scope of Cu-catalyzed asymmetric hydrogenation via DKR^a

reaction conditions, albeit with a more modest er (76:24). Finally, hydrogenation of a 1-cyclohexyl-2-tetralone (4p) was also successful furnishing the desired product in synthetically useful enantioselectivity (84:16).

The resulting optically enriched tetralols are valuable building blocks for synthesis of bioactive molecules. For example, compound 5q (>99:1 dr, 97:3 er from asymmetric hydrogenation of 4q, Scheme 3) can be readily deoxygenated to 2-aryltetralin 6 (97:3 er) which is a key precursor to pharmaceutically important compounds for treatment of arrhythmias by inhibiting the Na⁺/Ca²⁺ exchange mechanism.¹⁹ The only known asymmetric route to such 2-aryltetralins required the use of Pd-catalyzed arylation of 1-tetralone employing an equimolar amount of highly toxic tributyltin methoxide.²⁰

Scheme 3 Synthesis of 7-methoxy-2-phenyl tetralin. Conditions: (a) H_2 (700 psi), $CuCl_2$ (5 mol%), **3b** (5 mol%), $P(3,5-xylyl)_3$ (5 mol%), KOt-Bu (0.25 equiv.), t-AmOH, 20 °C, 24 h; (b) Et_3SiH (8 equiv.), TFA (4 equiv.), CH_2Cl_2 , 0 °C.

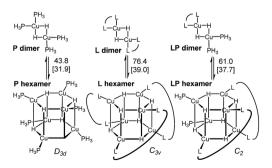
^a Isolated yields; diastereomeric and enantiomeric ratios (dr and er) were determined by chiral HPLC. ^b i-PrOH as solvent.

Scheme 4 Generic catalytic cycle for the Cu-catalyzed ketone reduction.

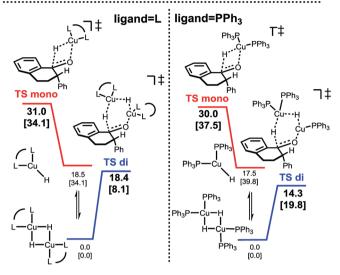
To understand the observed mixed ligand effect and to facilitate further development of the efficient catalysts, we initiated DFT study of the mechanism of this transformation. The reaction is believed to proceed *via in situ* formation of copper hydrides, where multiple species could be acting as the hydride source (Scheme 4).²¹

Copper hydrides are known to exist in oligomeric form, with the exact nature of the oligomer strongly depending on the identity of the ligand and the reaction conditions22. Thus, for the LCuH species, where L is a phosphine, there is evidence for existence of the corresponding dimers23, trimers4, pentamers55, hexamers26 and octamers27. For biaryl bidentate ligands, analogous to our ligand of interest, only a SEGPHOS-based system²⁸ has been examined29; NMR and MS experiments did not provide any conclusive results about the exact nature of this LCuH species, but suggested the presence of oligomers. Kinetic studies also pointed to the oligomeric character of the copper hydride. Previously published computational studies support the relative instability of the monomer³⁰. Given the well-known propensity of the LCuH species to form less reactive oligomers, the greater reactivity observed here when two different ligands are employed [bidentate L and monodentate P(3,5-xylyl)₃] may arise from the formation of different aggregates. Specifically, we theorize that the homoaggregates [L alone or P(3,5-xylyl)₃ alone] are less reactive than mixed aggregates incorporating the two different ligands.

DFT analysis of some model higher aggregates demonstrated that this is not the case, and the heteroligated aggregates average the stabilities of the homoligated systems (Scheme 5). This finding led us to propose the involvement of transition states incorporating two Cu atoms, where different ligands can



Scheme 5 Energy of the model dimers relative to the corresponding hexameric CuH aggregates. Ligands are truncated (see ESI† for details).³¹



Scheme 6 Relative energetics of monomeric (red) and dimeric (blue) hydride transfer pathways for bidentate (left) and monodentate (right) ligands.³¹

be coordinated to each Cu, to explain the mixed ligand effect. To the best of our knowledge, such transition states incorporating dimer Cu species had not been previously calculated for this type of reaction.

In analyzing hydride transfer transition states for the reduction of substituted tetralone, transition states incorporating two copper centers were lower than those incorporating one copper center for both monodentate and bidentate ligands (Scheme 6).

Following discovery of this lower energy reaction pathway, we analyzed the relative energetics of the most stable homo- and hetero-ligated dimeric transition states (Scheme 7).

These results indicate that the heteroligated dimeric transition state is lower in energy than the monoligated dimeric transition states, providing more than 1000-fold acceleration of the hydride transfer. Interestingly, the stabilization of the

Scheme 7 Relative energetics of the homoligated (leftmost and center) and mixed dimeric hydride transfer transition states.

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electron-rich L₁
enhances
H nucleophilicity

H electron-poor L₂
enhances C=O
electron-pilicity

Scheme 8 Favorable electronics for the dimeric hydride transfer transition state.

heteroligated dimeric transition state is not a purely entropic phenomenon. That is, P TS and LP TS differ in energy, even though both have the same molecularity. We hypothesize that the mixed ligand environment better supports the charge distribution required for the effective hydride transfer (Scheme 8). Specifically, one Cu center engages in Lewis acid activation of the carbonyl, for which an electron-deficient L2 would be superior. And, the second copper center delivers nucleophilic hydride, which is assisted by an electron-rich L1.

In summary, a new class of heterophosphole dimeric ligands (*O*-BABIPhos and *N*-BABIPhos) have been designed and synthesized. These ligands are highly modular and tunable and have enabled the first example of Cu-catalyzed hydrogenation of 2-substituted-1-tetralones and related heteroaryl ketones *via* dynamic kinetic resolution, simultaneously establishing two contiguous stereogenic centers with up to >99:1 dr and 98:2 er. The ligand–Cu complexes were isolated and characterized by single crystal X-ray, and DFT calculations revealed a novel heteroligated dimeric copper hydride transition state. Application of this new series of oxa- and azaphosphole bisphosphine ligands to other important catalytic reactions is currently under investigation and will be disclosed in due course.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

We thank Scott Pennino and Keith McKellop of Boehringer Ingelheim Pharmaceuticals for HRMS analysis. M. C. K. thanks the NIH (GM087605 M. C. K.) and Boehringer Ingelheim Pharmaceuticals for financial support. Computational support was provided by XSEDE (TG-CHE120052).

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