

Redox-Neutral α -Oxygenation of Amines: Reaction Development and Elucidation of the Mechanism

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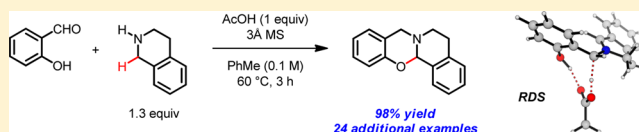
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Supporting Information

ABSTRACT: Cyclic secondary amines and 2-hydroxybenzaldehydes or related ketones react to furnish benzo[*e*][1,3]-oxazine structures in generally good yields. This overall redox-neutral amine α -C–H functionalization features a combined reductive N-alkylation/oxidative α -functionalization and is catalyzed by acetic acid. In contrast to previous reports, no external oxidants or metal catalysts are required. Reactions performed under modified conditions lead to an apparent reductive amination and the formation of *o*-hydroxybenzylamines in a process that involves the oxidation of a second equivalent of amine. A detailed computational study employing density functional theory compares different mechanistic pathways and is used to explain the observed experimental findings. Furthermore, these results also reveal the origin of the catalytic efficiency of acetic acid in these transformations.



INTRODUCTION

As drug discovery programs have come to rely on high-throughput screenings of diverse chemical libraries, the ability to rapidly construct complex, heterocyclic small molecules from simple starting materials is of great importance.¹ The N,O-acetal moiety can be found in a diverse set of natural products² and useful synthetic intermediates.³ Benzoxazines in particular have been studied as nonsteroidal progesterone receptor agonists,⁴ as antibacterial agents⁵ and as non-nucleoside reverse transcriptase inhibitors for the treatment of human immunodeficiency virus (HIV)⁶ as well as for a wide array of other applications.⁷ For example, benzo[*e*][1,3]oxazines such as PD 102 807 (**2a**) have been identified as potent, selective inhibitors of the m4 muscarinic receptor, which have made such compounds important leads in Parkinson's disease research.⁸

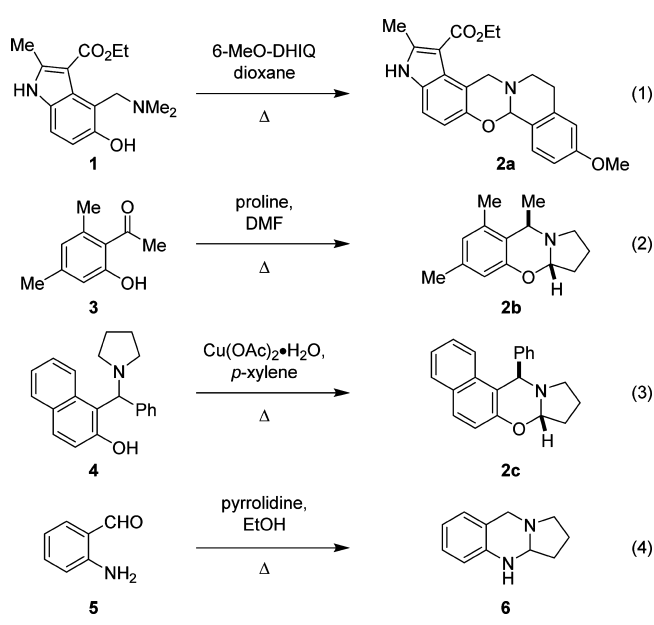
A number of methods for the synthesis of benzo[*e*][1,3]-oxazines have been reported (Scheme 1).^{3i,l,o,q,v,9} One early approach to polycyclic benzoxazines such as N,O-acetal **2a** involves the addition of a 3,4-dihydroisoquinoline (DHIQ) to a phenolic Mannich base (e.g., **1**), proceeding via an *o*-quinone methide intermediate and generally resulting in low to moderate yields (eq 1).^{8,9g,o–q,10} An intriguing and unanticipated entry to the N,O-acetal motif was reported by Cohen et al. in 1979 (eq 2).^{9h} Proline was found to react with 2-hydroxyacetophenones (e.g., **3**) via a decarboxylative process to yield products such as **2b**. Unfortunately, this method exhibited a rather narrow substrate scope. The presence of a methyl group in the ortho-position of the ketone was reported to be

crucial; replacement with a hydrogen substituent led to the recovery of **3** and pyrrolidine (from the decarboxylation of proline). The use of pipercolic acid (piperidine-2-carboxylic acid) in place of proline resulted in the formation of only trace amounts of the corresponding product. Recently, during the preparation of this paper, Maycock and co-workers^{9r} reported an oxidative, copper(II) acetate-catalyzed synthesis of benzoxazines such as **2c** from *o*-aminomethylnaphthols (e.g., **4**) and -phenols (eq 3). This report was closely followed by an independent publication by Jana and co-workers^{9t} in which the essentially identical transformation was described with superstoichiometric amounts of Ag₂O as the oxidant. All previous methods for benzo[*e*][1,3]oxazine synthesis involve either a prefunctionalized amine moiety (an amino acid or imine), an external oxidant, and/or a metal catalyst.¹¹ In 2008, one of our groups¹² reported the synthesis of amins such as **6** from *o*-aminobenzaldehydes (e.g., **5**) and unactivated secondary amines such as pyrrolidine (eq 4). These reactions feature a combined reductive N-alkylation/oxidative α -amination and function most efficiently in alcoholic solvents in the absence of any additives.¹³ The overall redox-neutral nature of this reaction distinguishes it from oxidative approaches to the C–H functionalization of amines, which continue to dominate most of the research efforts conducted in this area.¹⁴ The Seidel group has worked extensively on developing redox-neutral

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Scheme 1. Selected Previous Approaches to N,O-Acetals and Aminals

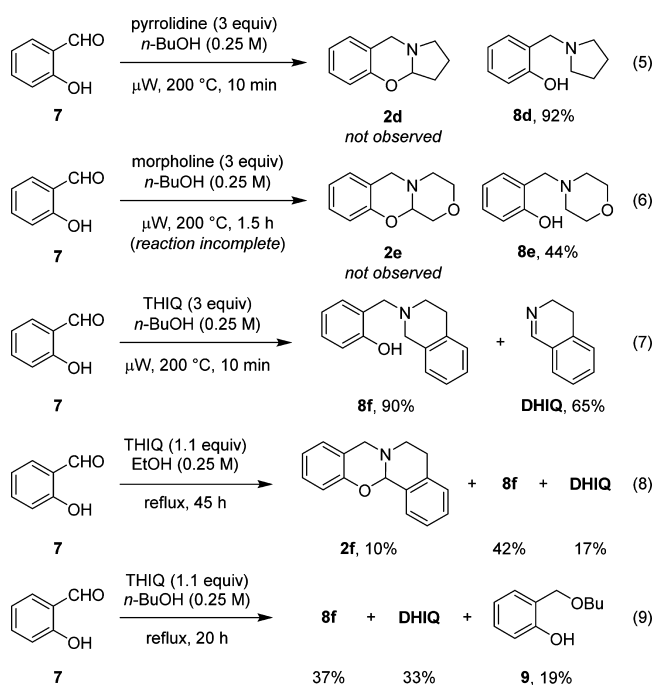


methods for the α -functionalization of amines,¹⁵ many of which involve iminium isomerization through azomethine ylide intermediates.^{16,17} Our two groups recently published a joint computational and experimental study of the amination reaction (eq 4) that revealed some interesting mechanistic features.^{12c} Simple iminium ions do not appear to play a role in this reaction, and the rate-determining step most likely involves a 1,6-proton transfer event. On the basis of the ability of *o*-aminobenzaldehydes to undergo these condensations with amines, we decided to explore the analogous reaction with salicylaldehydes in an effort to gain access to the N,O-acetal functionality in a facile, redox-neutral fashion. Here we report the successful development of this α -oxygenation, the scope of the reaction, and a detailed computational study of the mechanism.

EXPERIMENTAL RESULTS AND DISCUSSION

Evaluation of Various Reaction Conditions. To facilitate reaction development, we began our investigation using microwave conditions that had proven successful in the analogous aminal formation (Scheme 2).^{12d} Surprisingly, a reaction of pyrrolidine with salicylaldehyde (7) in *n*-butanol solvent (optimized conditions for aminal formation) did not lead to desired N,O-acetal product 2d. Instead, 2-hydroxybenzylamine 8d, the apparent product of a reductive amination, was isolated in 92% yield (eq 5). Similar observations were made in the corresponding reactions of morpholine (eq 6) and 1,2,3,4-tetrahydroisoquinoline (THIQ) (eq 7). In the case of THIQ, 3,4-dihydroisoquinoline (DHIQ) was isolated as a second product in 65% yield. This indicates that THIQ functions as the reductant in the formation of 8f. In order to avoid the formation of undesired product 8f, milder conditions were employed and the amount of THIQ was reduced. Heating a 1:1.1 mixture of 7 and THIQ under reflux in ethanol did, indeed, lead to isolation of N,O-acetal 2f, albeit in only 10% yield alongside a substantial amount of 8f and DHIQ (eq 8). When an otherwise identical reaction was performed under reflux in *n*-butanol (eq 9), a trace amount of

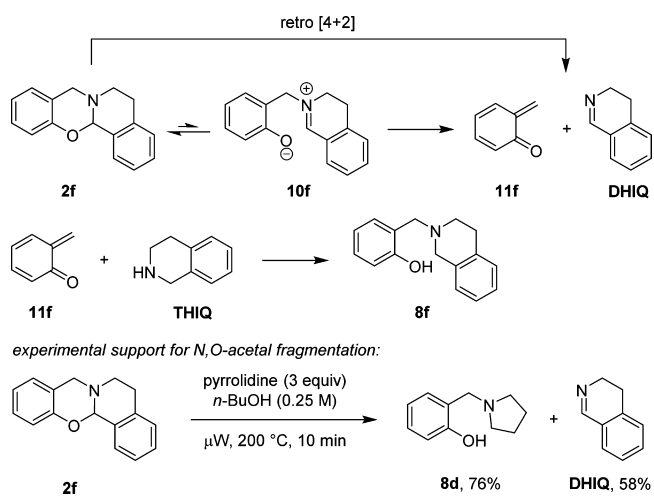
Scheme 2. Initial Studies on N,O-Acetal Formation



N,O-acetal was observed alongside 8f (37%) and DHIQ (33%). In addition, *n*-butyl ether 9 was isolated in 19% yield.

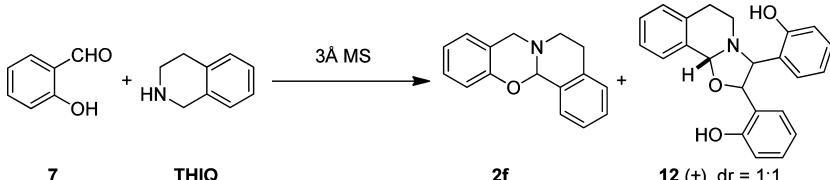
There are a number of different mechanistic scenarios that could account for the formation of reduced product 8f (Scheme 3). First, 8f could be formed from the desired product 2f.

Scheme 3. Potential Reaction Pathways and Experimental Support



Fragmentation of 2f, either via a retro [4 + 2] reaction or a stepwise pathway via zwitterion 10f, would result in the formation of *o*-quinone methide 11f and DHIQ. Reaction of the highly reactive 11f with THIQ would be expected to readily form 8f.¹⁸ Alternatively, the formation of 8f and DHIQ could be explained by reduction of 10f (or the regioisomeric zwitterion from the condensation of 7 and THIQ) via intermolecular hydride transfer from THIQ with concurrent oxidation of the latter to DHIQ (not shown). The formation of 9 (eq 9) is consistent with the intermediacy of 11f but not with the hydride transfer pathway. To obtain further insights into

Table 1. Optimization of N,O-Acetal-Forming Reaction between Salicylaldehyde and THIQ



entry	7, mmol	THIQ, mmol	additive (equiv)	solvent (M)	T, °C	t, h	yield of 2f, %	yield of 12, %
1	2	1	—	PhMe (0.25)	rt	48 ^a	35	36
2	2	1	PhCO ₂ H (0.2)	PhMe (0.25)	rt	18	38	48
3	1.1	1	PhCO ₂ H (0.2)	PhMe (0.25)	rt	24	54	25
4	1.1	1	2-EHA ^b (0.2)	PhMe (0.25)	rt	24	62	21
5	1	1.1	2-EHA (0.2)	PhMe (0.25)	rt	48 ^a	38	trace
6	1	1.1	2-EHA (0.2)	DMF (0.25)	rt	48 ^a	23	trace
7	1	1.1	2-EHA (0.2)	EtOH (0.25)	rt	48 ^a	22	31
8	1	1.1	2-EHA (0.2)	MeCN (0.25)	rt	48 ^a	62	32
9	1	1.1	2-EHA (1.3)	PhMe (0.25)	rt	48 ^a	75	trace
10	1	1.3	2-EHA (1.3)	PhMe (0.25)	rt	48	94	trace
11	1	1.3	2-EHA (1.3)	PhMe (0.25)	60	1.5	95	trace
12	1	1.3	2-EHA (1.3)	PhMe (0.1)	60	3	97	—
13 ^c	1	1.3	2-EHA (1.3)	PhMe (0.1)	60	6 ^a	64	—
14	1	1.3	AcOH (1.3)	PhMe (0.1)	60	3	98	—
15	1	1.3	AcOH (0.2)	PhMe (0.1)	60	3	59	—
16	1	1.3	AcOH (1.0)	PhMe (0.1)	60	3	98	—

^aReaction incomplete. ^b2-EHA = 2-ethylhexanoic acid. ^cNo molecular sieves.

the course of the reaction, benzoxazine **2f** was subjected to high temperatures in the presence of an excess of pyrrolidine (Scheme 3). In the event, *o*-hydroxybenzyl pyrrolidine **8d** and DHIQ were isolated in good yields, providing additional support for the fragmentation pathway.

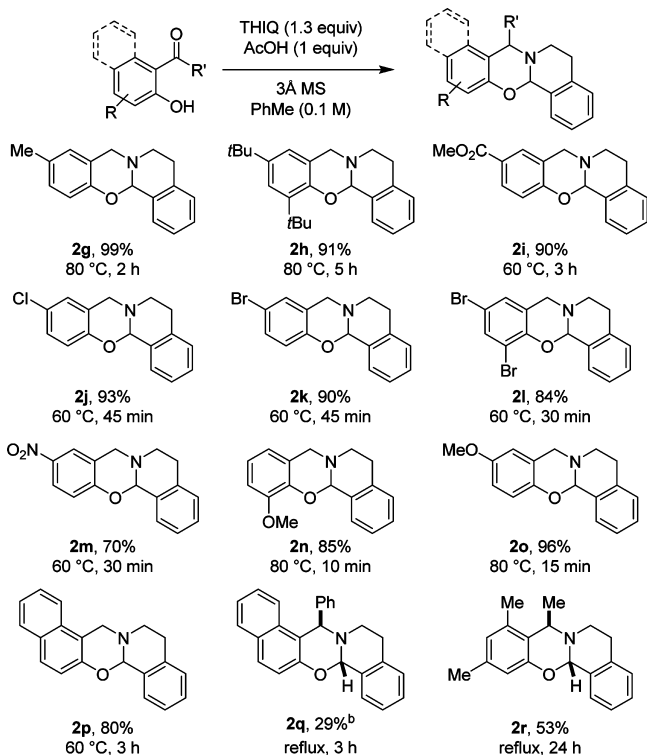
Due to the formation of undesired side products at higher temperatures and with nucleophilic solvents, we decided to evaluate the reaction under milder conditions (Table 1).

Toluene, a solvent that had previously been shown to be optimal for other redox-isomerization reactions,^{15h-j,l-o} was selected as the reaction medium. To further facilitate product formation, molecular sieves were added to sequester the water released during the condensation. Remarkably, the desired reaction was found to proceed at room temperature in the absence of any acid additives to provide N,O-acetal **2f** in 35% isolated yield (entry 1). In addition, the apparent [3 + 2] product **12** was formed in 36% yield, consistent with the intermediacy of an azomethine ylide.^{12c,19} The relative stereochemistry of **12** was not confirmed unambiguously, but literature precedent suggests that the phenolic groups should be trans-configured.¹⁹ The 1:1 diastereomeric ratio is the result of conformational instability of the N,O-acetal (see Supporting Information). Addition of catalytic amounts of benzoic acid dramatically accelerated the rate of the reaction while increasing the yield of both **2f** and **12** (entry 2). Reduction of the amount of salicylaldehyde (**7**) led to a more favorable product ratio with partial suppression of the [3 + 2] product **12** (entry 3). 2-Ethylhexanoic acid (2-EHA) performed slightly better than benzoic acid (entry 4), and acetic acid was later found to be still better (entry 14). Several protic and aprotic solvents were evaluated as potential alternatives to toluene, but all either resulted in lower yields (entries 6 and 7) or promoted the [3 + 2] reaction (entry 8). The formation of product **12** was completely suppressed with a reduction in solvent concentration (entry 12). Elevating the temperature to 60 °C and

using acetic acid in stoichiometric amounts led to the best result and allowed for the isolation of **2f** in 98% yield following a reaction time of 3 h (entry 16).

Substrate Scope. Optimized conditions were employed to evaluate the scope of the α -oxygenation with a number of different salicylaldehydes and related *o*-hydroxy ketones (Scheme 4). Salicylaldehydes with simple alkyl groups appended to the ring provided the corresponding products in good yields but required a higher temperature of 80 °C to achieve reasonable reaction rates (**2g** and **2h**). Both electron-withdrawing and electron-donating groups were tolerated, although more electron-deficient salicylaldehydes such as 3,5-dibromo- (**2l**) and 5-nitrosalicylaldehyde (**2m**) provided products with slightly decreased yields. *o*-Hydroxyketones required higher temperatures and afforded N,O-acetal products in relatively low yields but as single diastereomers (**2q** and **2r**). While ketones with some steric demand in the 6-position did yield the desired products, neither 2-hydroxyacetophenone nor 2-hydroxybenzophenone underwent the formation of N,O-acetals with THIQ under a variety of conditions, an observation that is in line with Cohen's findings on the related decarboxylative process.^{9h}

The scope of the reaction with regard to other amines was evaluated next (Scheme 5). Not surprisingly, cyclic secondary amines with benzylic protons in α -position to the ring nitrogen proved to be the most reactive substrates. Tetrahydroisoquinolines with methoxy groups appended to the aryl ring, upon reaction with **7**, resulted in the formation of products in excellent yields (**2s** and **2t**). A THIQ derivative with a phenyl group at the 1-position required more forcing conditions in order to form the corresponding N,O-acetal **2u**. Nevertheless, this highly substituted product was obtained in 72% yield. N,O-Acetal products could also be obtained with pyrrolidine, piperidine, and azepane. However, acyclic amines such as methylbenzyl amine failed to undergo the title reaction.

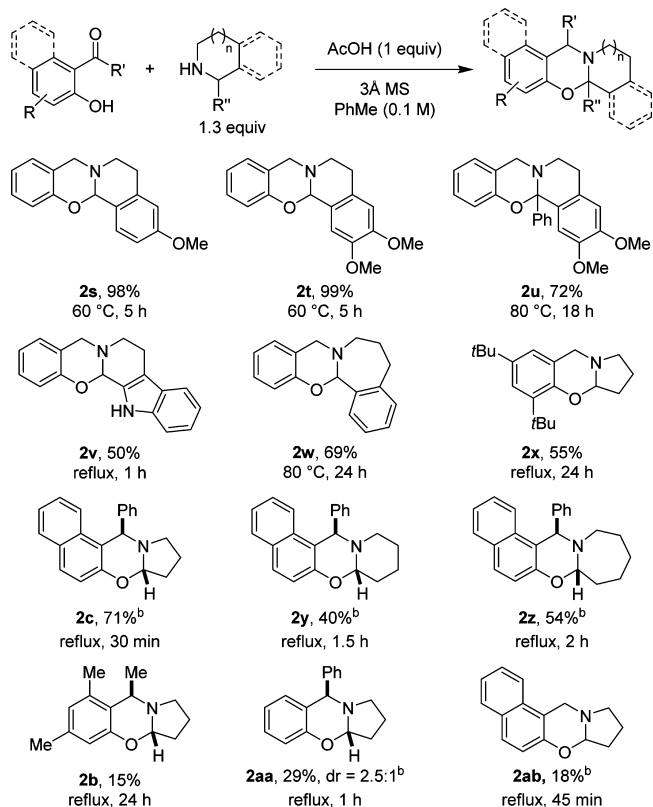
Scheme 4. Variation of Salicylaldehyde Moiety^a

^aAll reactions were performed on a 1 mmol scale. ^bFor this reaction, 1 mmol of amine, 2 equiv of ketone, and xylenes (0.1 M) were used.

Interestingly, attempted reactions with pyrrolidine and parent salicylaldehyde (**7**) did not yield N,O-acetal **2d** under a variety of conditions. Despite different experimental and computational attempts, this result could so far not be rationalized. Pyrrolidine underwent reaction with 3,5-di-*t*-butylsalicylaldehyde to form the corresponding N,O-acetal **2x** in 55% yield. Formation of product **2c** from pyrrolidine and 2-hydroxy-1-benzoylnaphthalene proceeded in 71% yield but required more forcing conditions (reflux in xylenes). Maycock and co-workers^{9r} did not observe benzoxazine **2c** as a product with the same starting materials in an experiment conducted at 130 °C in xylenes. This failure to obtain product **2c** is likely due to the fact that neither acidic additives nor molecular sieves were employed. While products **2c**, **2y**, **2z**, and **2b** were all isolated as single diastereomers, product **2aa**, which is different in that it lacks a substituent in the 6-position of the aryl ring, was formed as a 2.5:1 mixture of diastereomers.

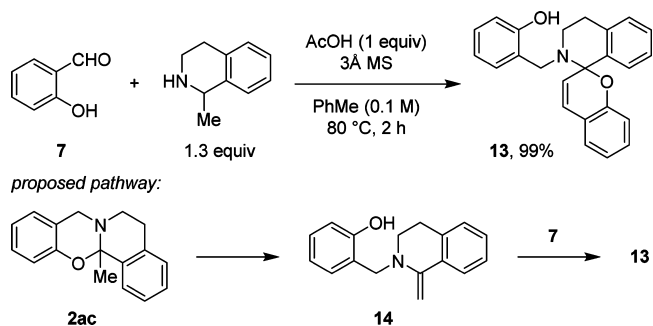
Interestingly, when 1-methyl THIQ was subjected to the reaction conditions, the desired N,O-acetal **2ac** was not obtained. Instead, the spirobenzopyran **13** was obtained in essentially quantitative yield (Scheme 6). In this case, N,O-acetal **2ac** or the corresponding zwitterion (not shown) could undergo transformation into or exist in equilibrium with enamine **14**. The latter can engage a second molecule of salicylaldehyde to give **13**. Reactions of structurally related enamines with salicylaldehydes have been previously reported.²⁰

To demonstrate that the redox-neutral synthesis of N,O-acetals is amenable to scale-up, the reaction of salicylaldehyde (**7**) and THIQ was performed on a 1-g scale in benzene as the solvent (Scheme 7). In the absence of molecular sieves, heating of the reaction mixture under reflux for a period of 24 h

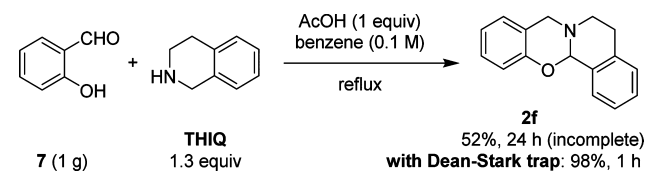
Scheme 5. Variation of Secondary Amine^a

^aAll reactions were performed on a 1 mmol scale. ^bFor these reactions, 1 mmol of amine, 2 equiv of aldehyde or ketone, and xylenes (0.1 M) were used.

Scheme 6. Reaction of Salicylaldehyde with 1-Methyltetrahydroisoquinoline

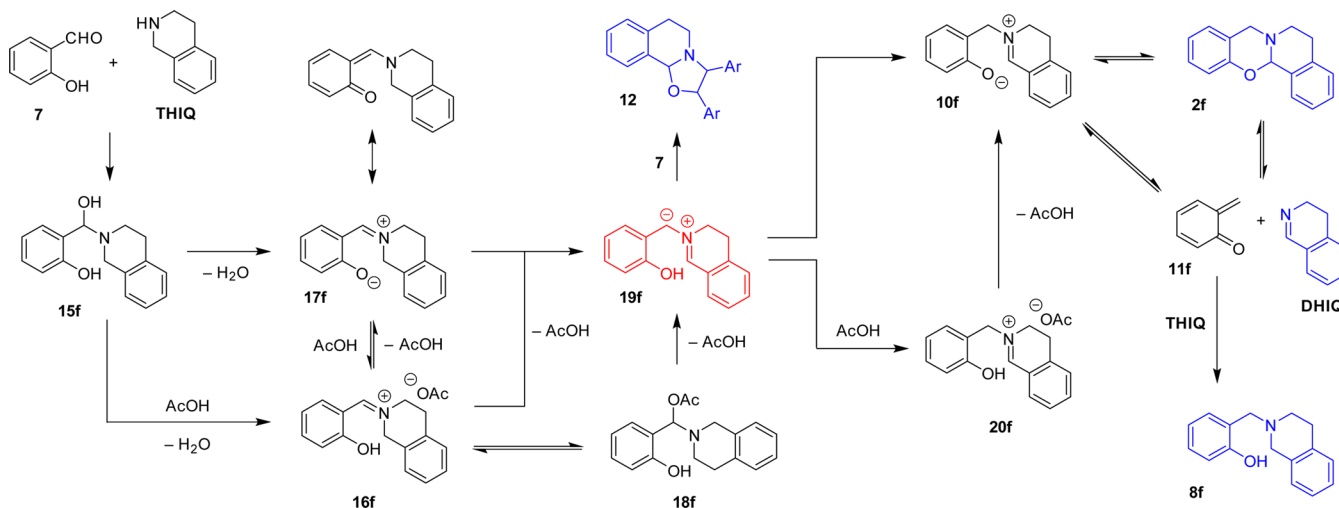


Scheme 7. Effect of Water on Condensation Reaction



resulted in an incomplete reaction and furnished the expected N,O-acetal **2f** in only 52% yield. Remarkably, an otherwise identical reaction performed in the presence of a Dean-Stark apparatus (for water removal) was completed after only 1 h and provided **2f** in nearly quantitative yield (98%, Scheme 7). These experiments nicely illustrate not only the ease with which this

Scheme 8. Possible Mechanistic Pathways for Formation of N,O-Acetal 2f and Benzylamine 8f



reaction can be performed under optimized conditions but also the importance of removing water from the reaction mixture.

Conceivable Mechanistic Pathways. A network of interrelated pathways presented itself when we considered possible mechanisms that would account for the formation of all observed products from the reaction of salicylaldehyde (7) and THIQ (Scheme 8). Based on the isolation of apparent [3 + 2] product **12**, it appears highly likely that the overall transformation involves the intermediacy of azomethine ylide **19f**. The reaction most likely initiates by addition of THIQ to salicylaldehyde (7) to form N,O-acetal **15f**, a step that may be facilitated by the presence of acetic acid. Subsequent elimination of water could occur either with the assistance of acetic acid, yielding iminium **16f**, or in a concerted intramolecular fashion to give zwitterion/quinoidal species **17f**. Due to the presence of acetic acid, **16f** and **17f** may exist in equilibrium. Azomethine ylide **19f** could be formed from **17f** via a 1,6-proton transfer; an analogous step was established in the formation of the corresponding aminals.^{12c} Another pathway to **19f** would involve deprotonation of **16f**. Alternatively, **18f**, which could exist in equilibrium with **16f**, could suffer concerted loss of acetic acid to generate azomethine ylide **19f**, consistent with a proposal by Yu and co-workers²¹ for a related process. Azomethine ylide **19f** would then progress to zwitterion **10f** either by a stepwise protonation/deprotonation pathway via **20f** or by direct proton transfer. Ring closure finally leads to N,O-acetal product **2f**. N,O-Acetal **2f** can undergo further transformation to “reduced” product **8f** via the addition of THIQ to *o*-quinone methide **11f**, formed in a formal retro-[4 + 2] reaction that also generates DHIQ. It should be noted that the retro-[4 + 2] step may occur in a stepwise manner via zwitterion **10f**. Facile ring-opening of benzoxazines and the potential existence of an equilibrium between **2f** and **10f** is supported by an observation about the appearance of benzoxazine **2f**. While **2f** is a white solid in pure form, solutions of **2f** turn bright yellow in the presence of an acid (e.g., acetic acid or silica gel), suggesting the formation of a new species.

Another observation consistent with the existence of zwitterions in equilibrium with benzoxazines is that benzoxazine products with an electron-deficient phenolic ring (i.e., **2i**, **2l**, and **2m**) exhibit broadened peaks in their ¹H NMR spectra (see Supporting Information). Presumably, the electron-

withdrawing groups stabilize the phenoxide, allowing the N,O-acetal to rapidly equilibrate with zwitterionic form **10l**. This process is suppressed or slowed down at lower temperatures, as illustrated by a series of ¹H NMR spectra that were recorded at temperatures between 20 and -60 °C (Figure 1).

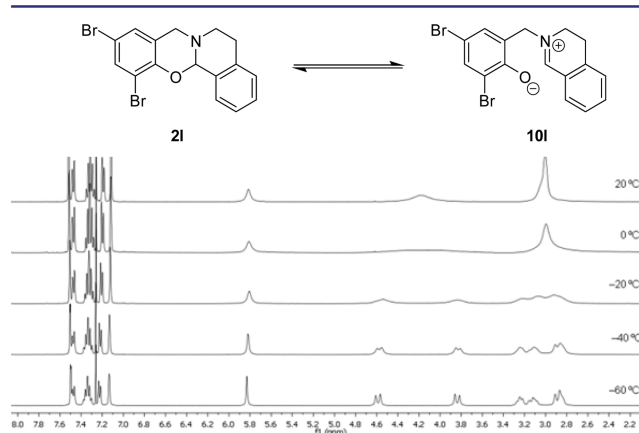
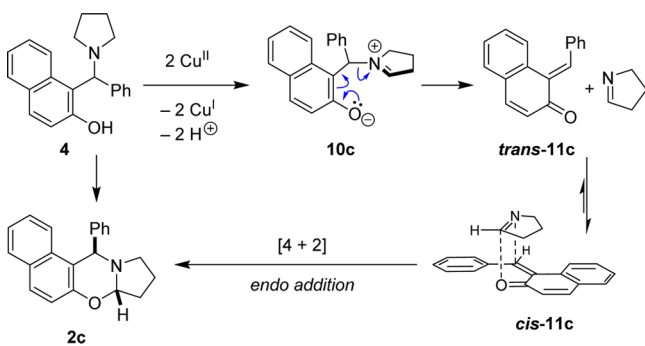


Figure 1. Temperature-dependent ¹H NMR spectra of **2l** in CDCl₃ (400 MHz).

With regard to the above-mentioned oxidative N,O-acetal syntheses reported by Maycock and co-workers^{9r} (eq 3) and Jana and co-workers,^{9t} different mechanisms were proposed by the two groups. Jana and co-workers proposed an initial oxidation of **4** at the benzylic position and deprotonation of the resulting iminium ion, followed by a pathway that is based on our previously established mechanism for the corresponding aminal formation, namely, 1,6-proton abstraction to generate an azomethine ylide and subsequent proton transfer and ring closure (not shown). Interestingly, Maycock’s mechanistic proposal is radically different (Scheme 9). It involves oxidation of amine **4** at the seemingly less activated endocyclic position rather than the benzylic position to give intermediate **10c**. While **10c** may undergo direct ring closure to product **2c**, the observed diastereoselectivity was rationalized via a different pathway. According to Maycock and co-workers, intermediate **10c** undergoes fragmentation to *o*-quinone methides *trans*-**11c** and *cis*-**11c** and 1-pyrroline. The isomer *cis*-**11c** is proposed to

Scheme 9. Proposed Mechanism of Maycock and Co-workers^{9r} for Oxidative N,O-Acetal Formation


engage in an endo-[4 + 2] cycloaddition with 1-pyrroline to selectively form **2c** in the observed relative configuration. An alternative and more likely explanation that accounts for the essentially exclusive formation of **2c** over its other diastereomer is based on equilibration between the two possible diastereomers via zwitterion **10c**. In fact, the equilibration of diastereomers of closely related benzoxazines was studied in detail by Fülöp, Kleinpeter, and co-workers,^{9q} who concluded that the diastereomer corresponding to **2c** represents the thermodynamically more stable product.

COMPUTATIONAL RESULTS AND DISCUSSION

In order to investigate the underlying mechanism and to identify the most important pathways for the formation of benzoxazines under these conditions, we have employed density functional theory calculations [M06-2X-D3/def2-

TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM]. We will first discuss the mechanisms for the uncatalyzed and acetic-acid-catalyzed reaction using the model system salicylaldehyde (**7**) and THIQ before we analyze the influence of substituents at the carbonyl and variation of the amine as well as potential side reactions.

Uncatalyzed Reaction in Toluene. Although the uncatalyzed reaction between the aldehyde and the amine results in low yields of the corresponding benzoxazines (Table 1), this background reaction is important for the acid-catalyzed reaction as well. Therefore, we first carefully analyzed the mechanism for the prototypic reaction between salicylaldehyde (**7**) and tetrahydroisoquinoline (THIQ) in toluene solution in the absence of any catalyst (cf. Scheme 8). The calculated free energy profile is depicted in Figure 2 (black lines), and selected calculated structures are discussed in Figures 3–5.

In the first step of this transformation, the aldehyde **7** and THIQ form the hemiaminal **15f** in a slightly endergonic reaction ($\Delta G = +1.9$ kcal·mol⁻¹). Next, water is eliminated from the hemiaminal, yielding the zwitterionic intermediate **17f**. This reaction could occur either in a concerted mechanism ($\Delta G^\ddagger = +22.5$ kcal·mol⁻¹, via *cis*-TS1, Figure 3) or in a stepwise reaction through an iminium ion. In line with previous investigations of the synthesis of tetrahydroquinazolines,^{12c} the putative iminium ion obtained from the elimination of a hydroxy group was located 38 kcal·mol⁻¹ above *cis*-TS1 and is not shown in Figure 2. As a consequence, the concerted elimination is also preferred over the stepwise elimination of hydroxide and subsequent deprotonation in these transformations. In principle, both eliminations to the *cis* and the *trans* zwitterions **17f** are possible (Figure 3). Our calculations

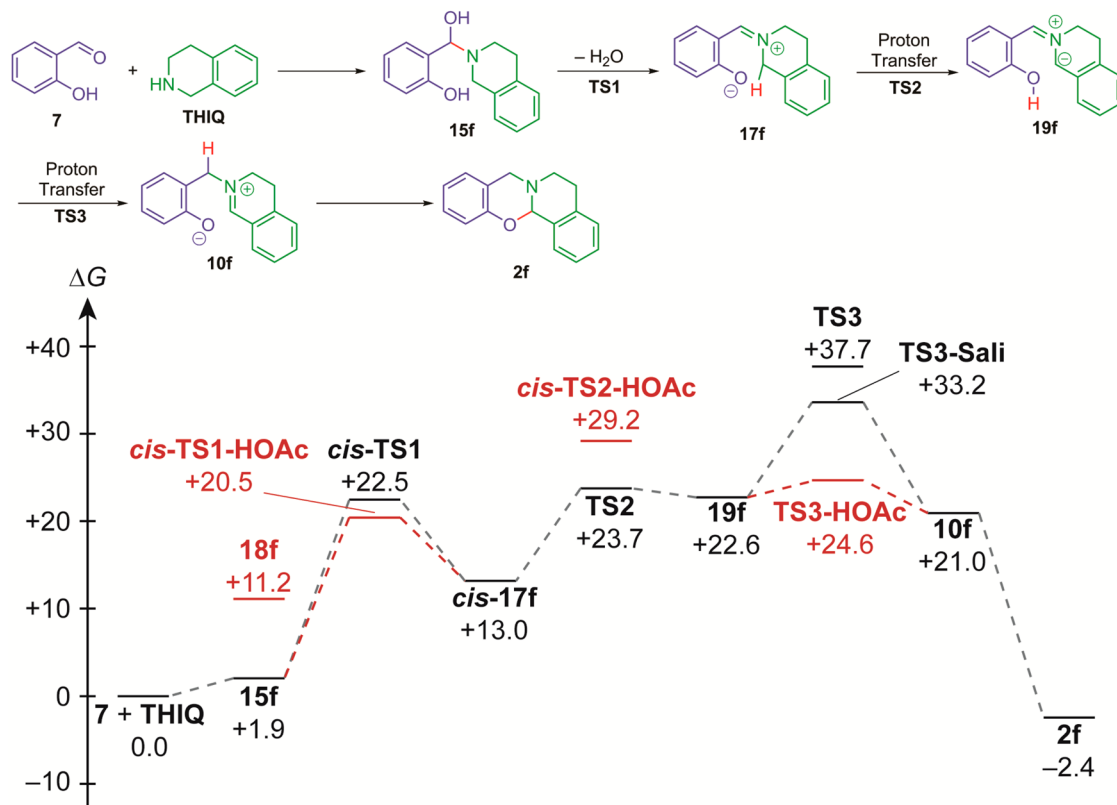


Figure 2. Free energy profile [in kcal·mol⁻¹, M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM] for uncatalyzed (black) and acetic-acid-catalyzed (red) transformation of **7** and THIQ to benzoxazine **2f** in toluene.

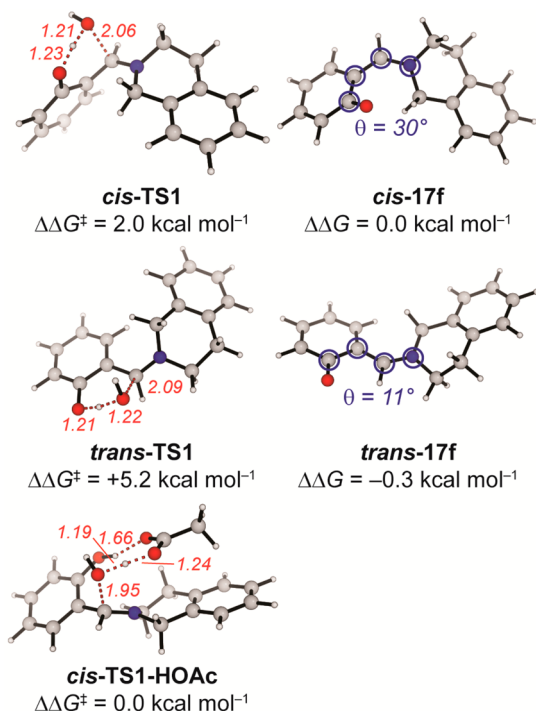


Figure 3. Calculated structures [M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM], relative free energies (in kcal·mol⁻¹), selected bond lengths (in Å), and dihedrals for different conformers of transition state TS1 and zwitterion 17f.

predict *cis*-TS1 to be significantly favored over *trans*-TS1 ($\Delta\Delta G^\ddagger = 3.2 \text{ kcal}\cdot\text{mol}^{-1}$), while the product *cis*-17f is essentially isoenergetic to its isomer *trans*-17f. The slight thermodynamic preference for the *trans* conformer can be rationalized by the greater planarity of the exocyclic π -system (as reflected by the dihedral angle θ in Figure 3). Analysis of the charge distribution [e.g. natural bond orbital (NBO) or ChelpG] in 17f as well as of smaller model systems revealed that the zwitterionic and neutral resonance structures should be equally important (see Supporting Information for more details). The next step of the mechanism requires abstraction of one of the α -hydrogens of the heterocycle, which, in an intramolecular reaction, is possible only from the *cis* conformation of 17f. However, previous calculations on the corresponding aminobenzaldehyde-derived intermediates have shown that *cis*- and *trans*-17f can be directly interconverted with small barriers.^{12c}

A subsequent proton transfer via TS2 leads to the azomethine ylide 19f in another endergonic transformation (Figure 4). The endergonicity of this step is also reflected in the short O–H bond length of the late transition state TS2. This reaction could proceed via either a 1,6-hydride shift or a 1,6-proton transfer, and our charge calculations (NBO or ChelpG) indicate that the latter is more likely due to a significant positive charge on the transferred hydrogen atom.

The azomethine ylide 19f then undergoes another, rate-limiting proton transfer yielding the zwitterion 10f. This transformation can either occur in an intramolecular reaction (TS3, Figure 5) or in a salicylaldehyde-mediated reaction (TS3-Sali, Figure 5). The entropic penalty ($-T\Delta S$) for the intermolecular reaction through TS3-Sali involving a second molecule of salicylaldehyde 7 is compensated by the very favorable activation enthalpy rendering TS3-Sali the preferred

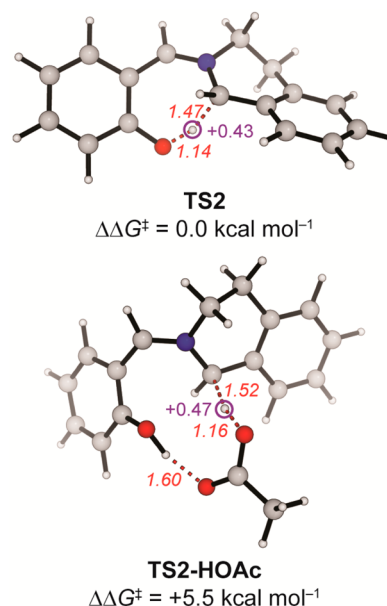


Figure 4. Calculated structures [M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM], relative free energies (in kcal·mol⁻¹), selected bond lengths (in Å), and selected NBO charges for transition state TS2.

pathway. The oxazinane 2f is subsequently obtained by a barrierless cyclization of the zwitterionic intermediate 10f in an overall exergonic reaction ($\Delta G = -2.4 \text{ kcal}\cdot\text{mol}^{-1}$).

As an alternative pathway, azomethine ylide 19f may be trapped via the reaction with excess salicylaldehyde 7, affording the [3 + 2] adduct 12 (Scheme 8, Figure 6). As the relative stereochemistry of the experimentally isolated adduct has not been determined, we have investigated all four possible stereoisomers and their corresponding transition states. Our calculations indicate that *RRS*-12 is the most stable stereoisomer of the four [$\Delta\Delta G(\text{RRR}-12) = +2.2$, $\Delta\Delta G(\text{RSR}-12) = +2.9$, and $\Delta\Delta G(\text{RRS}-12) = +3.6 \text{ kcal}\cdot\text{mol}^{-1}$; see Supporting Information for more details] and only *RRS*-12 is formed in an exergonic reaction ($\Delta G = -1.5 \text{ kcal}\cdot\text{mol}^{-1}$, Figure 6). The lowest-energy transition states were calculated to be *RRR*-TS4 ($\Delta G^\ddagger = 32.6 \text{ kcal}\cdot\text{mol}^{-1}$) and *RRS*-TS4 ($\Delta G^\ddagger = 33.0 \text{ kcal}\cdot\text{mol}^{-1}$). As the formation of *RRR*-12 is endergonic, the computational data predict the formation of *RRS*-12 to be the preferred [3 + 2] pathway (Figure 6). Both pathways leading to the benzoxazine 2f and the alternate product *RRS*-12 are similar in both activation and reaction free energies. While formation of the [3 + 2] adduct has a slightly smaller activation free energy (kinetic preference), the benzoxazine is preferred thermodynamically. This also explains why the [3 + 2] cycloaddition is facilitated when the aldehyde is present in large concentrations and used in excess over the amine (e.g., entries 1 and 2 in Table 1).

Acetic-Acid-Catalyzed Reaction in Toluene. As the rate-limiting step for the uncatalyzed reaction was calculated to be rather high and significant accelerations could be observed in the experiments with acetic acid as a catalyst, we subsequently investigated how acetic acid can catalyze the synthesis of benzoxazinanes (Figures 2–4).

Previous calculations on similar redox isomerizations by Yu and co-workers,²¹ employing MP2/6-31+G(d)//B3LYP/6-31+G(d)/CPCM, have already highlighted the crucial role of

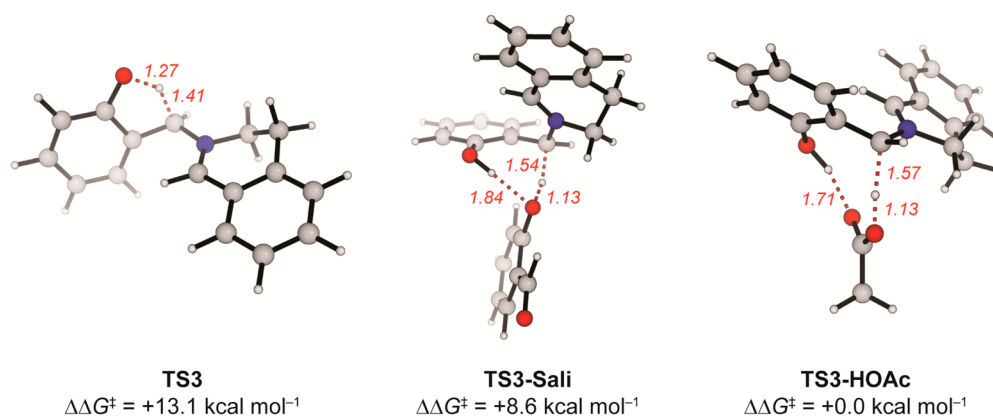


Figure 5. Calculated structures [M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM], relative free energies (in kcal·mol⁻¹), and selected bond lengths (in Å) for transition state TS3.

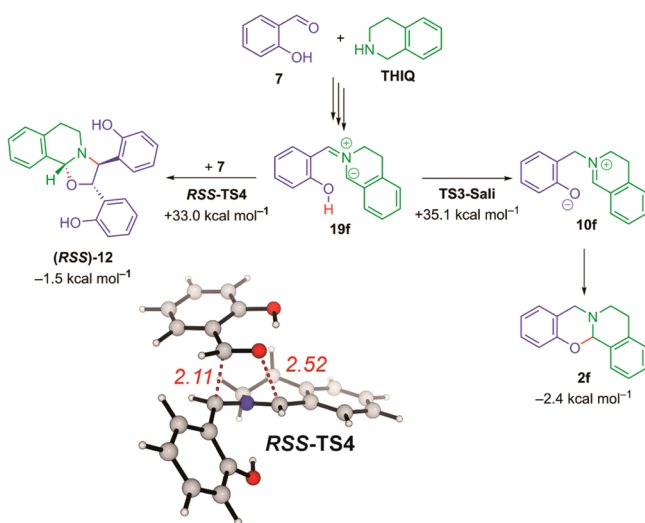


Figure 6. Calculated activation and reaction free energies of different pathways involving the azomethine ylide **19f** [M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM] and transition state RSS-TS4 with selected bond lengths (in Å).

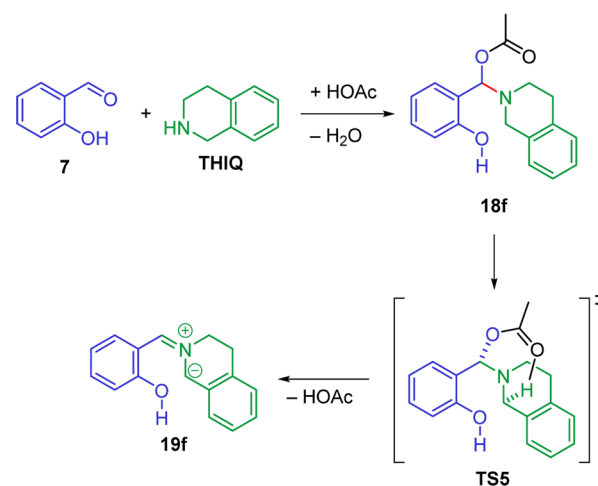
acetic acid in these transformations and are possibly important for the transformations under investigation.

In a first step, we analyzed whether the acetylated hemiaminal **18f** could eliminate acetic acid with formation of the azomethine ylide **19f** as proposed by Yu and co-workers (Scheme 10).²¹ However, according to our calculations, the formation of hemiaminal **18f** is significantly endergonic ($\Delta G = +11.2$ kcal·mol⁻¹) and the transition state for the elimination of acetic acid TS5 ($\Delta G^\ddagger = +35.3$ kcal·mol⁻¹) was found to be comparable in energy to the uncatalyzed reaction [$\Delta G^\ddagger(\text{TS3-Sali}) = +33.2$ kcal·mol⁻¹]. Based on these results, this mode of activation by acetic acid does not explain the rate acceleration and has to be rejected for these transformations.

Next, we investigated whether acetic acid can act as a proton shuttle within the transition states TS1–3 and thereby lower the activation energy of each step. The activation free energies for the acetic-acid-catalyzed reactions are summarized in Figure 2 and the optimized structures are depicted in Figures 3–5.

The dehydration of the hemiaminal **15f** is slightly facilitated by acetic acid acting as proton shuttle (*cis*-TS1-HOAc versus *cis*-TS1, Figure 3), while the proton transfer yielding the azomethine ylide (TS2 and TS2-HOAc, Figure 4) is actually

Scheme 10. Potential Acceleration of Acetic Acid (cf. Ref 21)



destabilized by acetic acid. As the barrier for the uncatalyzed reaction is already very small (with respect to **19f**), the additional entropy penalty ($-T\Delta S$) cannot be compensated by the more favorable enthalpy. As a consequence, the intramolecular proton transfer is preferred over the intermolecular process for this step. In contrast, a large stabilization has been calculated for the rate-limiting proton transfer TS3 in TS3-HOAc (Figure 5), indicating a substantial stabilization of the transition state. In summary, this large difference in free energy for TS3 ($\Delta\Delta G^\ddagger = 8.6$ kcal·mol⁻¹) is also the origin of the favorable acetic acid catalysis. This role of acetic acid in N,O-acetal formation has a parallel in the corresponding synthesis of aminals. In the latter case, the solvent, ethanol, has been shown to serve as the proton shuttle.^{12c}

Comparison of Selected Carbonyl-Amine Combinations. To better understand the observed reactivities, we next analyzed selected carbonyl-amine combinations including electron-rich and -poor carbonyls and two different amines.

Independent of the combinations of carbonyl and amine, the overall reaction free energies are all found within a relatively small range ($-5.4 < \Delta G < +2.4$ kcal·mol⁻¹, Table 2), indicating that the substituents on the carbonyl and the choice of amine are less important for the thermodynamics of the overall reaction. The fact that some intermediates (e.g., for **19ad** or **19d**) are higher in energy than the corresponding acetic-acid-catalyzed transition states indicate that acetic acid can

Table 2. Calculated Free Energies for Different Carbonyl-Amine Combinations in Toluene^a

15	+1.9	+6.4	+5.6	-0.7	+2.1	-0.2	+1.8
18	+11.2	+20.9	+19.0	+9.1	+2.3	+0.7	+13.6
cis-TS1	+22.5	+27.9	+24.8	+19.5	+24.0	+18.4	+21.6
cis-TS1-HOAc	+20.5	+25.9	+25.4	+20.2	+20.6	+15.4	+18.0
cis-17	+13.0	+20.4	+16.6	+9.5	+15.9	+7.5	+9.8
TS2	+23.7	+30.3	+26.3	+21.4	+23.7	--- ^b	--- ^b
TS2-HOAc	+29.2	+24.8	+28.2	+24.5	+29.6	--- ^b	--- ^b
19	+22.6	+29.3	+22.8	+20.7	+22.3	+26.1	+31.7
TS3	+37.7	+45.5	+38.9	+32.7	+39.3	+36.7	+38.1
TS3-Sali	+33.2	+44.9	+37.7	+30.4	+32.9	+32.4	+41.0
TS3-HOAc	+24.6	+31.9	+31.5	+21.6	+23.7	+22.9	+25.8
10	+21.0	+28.0	+22.5	+13.2	+24.9	+22.1	+24.5
2	-2.1	+2.4	-3.1	-4.5	-2.6	-5.4	-3.1

^aEnergies are given in kcal·mol⁻¹, M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM. ^bAll attempts to locate a transition state for these reactions failed, and potential energy surface scans indicate a barrierless reaction.

coordinate to the intermediates, which leads to a further stabilization. While the combination of hydroxyacetophenone and THIQ resulted in an endergonic reaction (\rightarrow **2ad**) and no detectable product formation, the dimethyl analogue yields the corresponding benzoxazine **2r** in an exergonic reaction in line with an isolated yield of 53% (Scheme 4). Employing truncated model systems, we could show that this difference can be attributed to a relief of 1,3-strain present in the reactant **3** but not in the corresponding oxazine (see Supporting Information for more details).

A pronounced substituent effect is observed, however, for the zwitterionic intermediates **10**, as already indicated by the broad NMR peaks (Figure 1) for the dibromo compound **2l**. The electron-withdrawing bromo substituents in **10l** result in a stabilization of 7.8 kcal·mol⁻¹ compared to **10f**, while the electron-donating methoxy group leads to a destabilization in **10o** ($\Delta\Delta G = +3.9$ kcal·mol⁻¹). The additional benzene ring of tetrahydroisoquinoline compared to pyrrolidine only translates to a small difference in free energy (cf. **10f** and **10d** in Table 2) indicating that the interaction with the negatively charged alcoholate is more important for the stability than an interaction with the iminium substructure.

For very bulky substrates (e.g., leading to **2r** or **2x**), the additional steric interactions in the transition states **cis-TS1-HOAc** and **TS3-Sali** completely compensate any stabilization and render them higher in energy than **cis-TS1** and **TS3**, respectively.

Investigation of Side Reactions. Among the possible pathways for side reactions, we first analyzed the feasibility of an intermolecular reduction of the intermediate zwitterion **10f**

by excess THIQ (Figure 7). The high activation free energy for the hydride transfer ($\Delta G^\ddagger = 38.4$ kcal·mol⁻¹) renders this

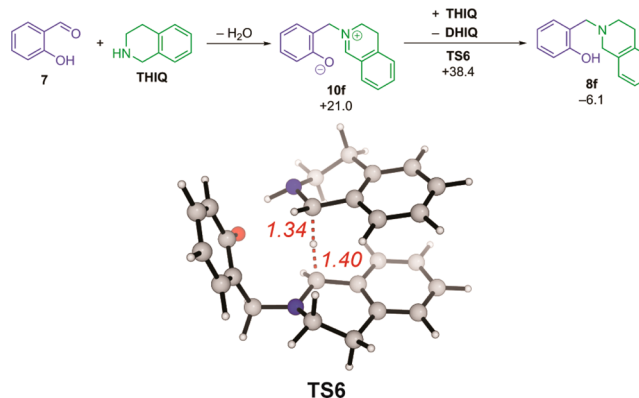
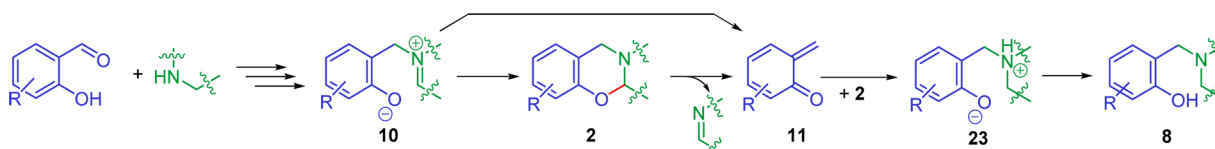


Figure 7. Calculated transition state **TS6** with selected bond lengths (in Å) and activation and reaction free energies (in kcal·mol⁻¹) for an intermolecular reduction of the intermediate zwitterion **10f** by THIQ [M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM].

pathway unlikely under the reaction conditions employed (Table 1). However, a large excess of the amine would favor this reaction and could slightly reduce the activation free energy.

Another possibility, which is also in line with the experimental isolation of dihydroquinoline (DHIQ), is a retro-Diels–Alder reaction followed by a nucleophilic attack

Table 3. Calculated Free Energies for Intermediates of Reductive Isomerization for Different Carbonyl-Amine Combinations in Toluene^a


	2f	2ad	2r	2l	2o	2d	2x
10	+21.0	+28.0	+22.5	+13.2	+24.9	+22.1	+24.5
2	-2.1	+2.4	-3.1	-4.5	-2.6	-5.4	-3.1
11	+22.8	+25.4	+21.0	+21.3	+19.5	+23.8	+20.8
23	+24.4	---	---	+17.3	+23.5	+19.5	+21.4
8	-6.1	+1.2	-3.9	-8.8	-5.8	-6.8	-4.8

^aEnergies are given in kcal·mol⁻¹, M06-2X-D3/def2-TZVPP/IEFPCM//TPSS-D2/6-31+G(d,p)/IEFPCM.

of the amine on the formed quinone methide. The energies of all intermediates for these transformations are summarized in Table 3. In all cases under investigation, the alternate product **8** is thermodynamically more stable than the corresponding cyclic N,O-acetal **2** ($0.8 < \Delta\Delta G < 4.3$ kcal·mol⁻¹) and the intermediates of the putative retro-[4 + 2] reaction, **11** and the corresponding imine, are 20–25 kcal·mol⁻¹ higher in energy. Table 3 further shows that the reduced amines dihydroquinoline and pyrrolidine are comparable in stability ($\Delta\Delta G = 1.0$ kcal·mol⁻¹). However, we were not able to locate any transition states for any of these transformations. Therefore, we investigated the potential energy landscape around the hemiaminals **2** by performing two-dimensional relaxed potential energy surface scans at the TPSS-D2/6-31G(d)/IEFPCM level of theory (Figure 8). Regardless of the

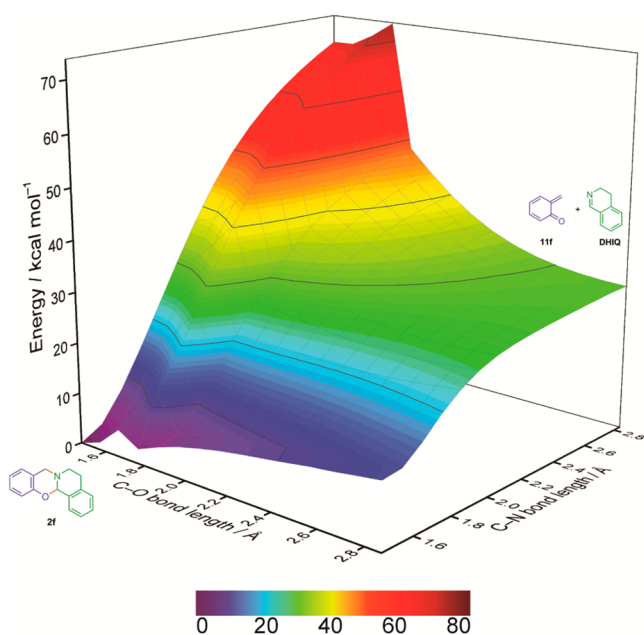


Figure 8. Calculated potential energy surface scan for the putative retro-hetero-Diels–Alder reaction involving **2f** [in kcal·mol⁻¹, TPSS-D2/6-31G(d)/IEFPCM].

proposed mechanism (e.g., stepwise versus concerted cycloaddition), these scans result in a barrierless combination of the quinone methide **11f** and the imine DHIQ, yielding the experimentally observed N,O-acetal **2f**. From these results, we have to conclude that both a putative retro-[4 + 2] reaction and the subsequent nucleophilic attack would proceed without significant barriers. These results are in agreement with previous kinetic studies by the groups of Freccero, Kresge, Richard, Rokita, Mayr, and others.²²

As a consequence, none of the pathways considered can account for the unusual reactivity of pyrrolidine and salicylaldehyde, and a different reason has to be responsible for the experimental observations.

COMPUTATIONAL DETAILS

The conformational space of all intermediates for the benzoxazine synthesis was explored by use of the OPLS-2005²³ force field and a modified Monte Carlo search routine implemented in MacroModel 9.9.²⁴ An energy cutoff of 20 kcal·mol⁻¹ was used for the conformational analysis, and structures with heavy-atom root-mean-square deviation (RMSD) less than 1–2 Å after the initial force field optimization were assumed to be the same conformer. The remaining structures were subsequently optimized by employing the meta-GGA functional TPSS²⁵ with Grimme's dispersion-correction D2,²⁶ and the double- ζ basis set 6-31+G(d,p). Solvation by toluene was taken into account by using the integral equation formalism polarizable continuum model (IEFPCM) for all calculations (optimizations, frequencies, and single points).²⁷ It has recently been shown that the use of a polarizable continuum model does not have a large impact on the calculated frequencies but is necessary for the location of transition states in some cases.²⁸ Vibrational analysis verified that each structure was a minimum or a transition state. Following the intrinsic reaction coordinates (IRC) confirmed that all transition states connected the corresponding reactants and products on the potential energy surface. Two-dimensional potential energy surface scans were performed at the TPSS-D2/6-31G(d)/IEFPCM level of theory. Thermal corrections were calculated from unscaled harmonic vibrational frequencies at the same level of theory for a standard state of 1 mol·L⁻¹ and 298.15 K. Entropic contributions to the reported free energies were calculated from partition functions evaluated with quasiharmonic approximation of Truhlar and co-workers.²⁸ This method uses the same approximations as the usual harmonic oscillator except that all vibrational frequencies lower than 100 cm⁻¹ are set equal to 100 cm⁻¹ to correct for the breakdown of the harmonic oscillator approximation for low frequencies. Electronic energies were subsequently obtained

from single-point calculations of the TPSS-D2 geometries employing the meta-hybrid M06-2X functional,²⁹ the large triple- ζ def2-TZVPP basis set,³⁰ IEFPCM for toluene, and Grimme's dispersion-correction D3 (zero-damping),³¹ a level expected to give accurate energies.³² An ultrafine grid corresponding to 99 radial shells and 590 angular points was used throughout this study for numerical integration of the density.³³ All density functional theory (DFT) calculations were performed with Gaussian 09,³⁴ and the additional D3 corrections for single-point calculations were carried out with Grimme's DFT-D3 program.³¹

CONCLUSIONS

We have developed a mild and highly efficient synthesis of benzoxazines from the direct condensation of salicylaldehydes and secondary amines. This redox-neutral process can be used to rapidly create a wide range of polycyclic N,O-acetals. In addition, a reductive amination of salicylaldehydes in which excess amine serves as reductant was discovered. The mechanism of the α -oxygenation was elucidated by DFT calculations that correlate well with experimental results. Further studies on this and related reactions are ongoing.

ASSOCIATED CONTENT

Supporting Information

Experimental procedures, characterization data, Cartesian coordinates, energies of all reported structures, and details of computational methods. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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