

Evaluation of protonation sites in two MacMillan catalysts in solution by gas phase predissociation spectroscopy and electronic structure calculations

Laís C. Tavares,^a Tatiana C. Penna,^a Mark Johnson,^{b*} and Thiago C. Correra^{a*}

^aDepartment of Fundamental Chemistry, Institute of Chemistry, University of São Paulo, Av. Prof. Lineu Prestes, 748, 05508-000, São Paulo, SP, Brazil

^bSterling Chemistry Laboratory, Yale University, P. O. Box 208107, New Haven, Connecticut 06520, United States

Email: mark.johnson@yale.edu, tcorrera@iq.usp.br

In honor of Prof. José Manuel Riveros on the occasion of his 80th anniversary

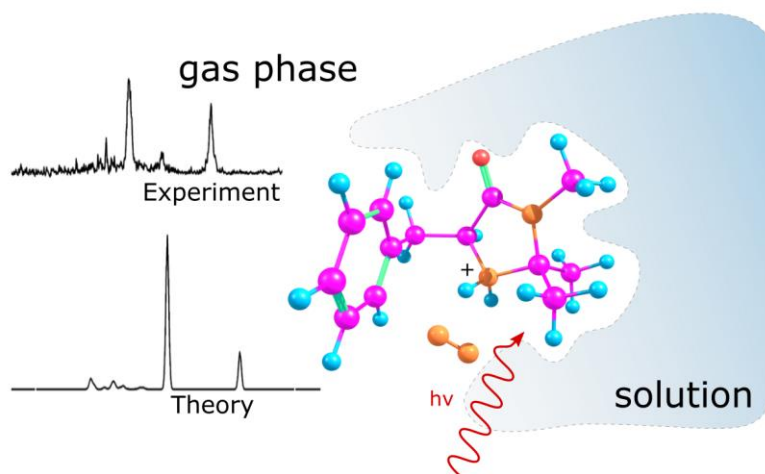
Received 01-06-2020

Accepted 05-11-2020

Published on line 05-18-2020

Abstract

The protonation sites of first and second generation MacMillan catalyst were evaluated using cryogenic ion vibrational predissociation spectroscopy. The comparison with calculated spectra showed that when isolated in the gas phase, the species are protonated at the secondary amine, as would be expected by the difference in free energy of the calculated protomers. The relative energies of the conformers were also calculated for the species in solution using the SMD model, in water, methanol and in a 1:1 (v/v) water:methanol mixture. This comparison indicates that the trends observed in the gas phase are retained in solution, and that the secondary amine is the protonation site regardless the solvent used.



Keywords: Organocatalysis, predissociation spectroscopy, MacMillan catalyst, DFT calculations.

Introduction

New challenges posed by the modern world have been pushing the boundaries of Science in general and Chemistry in particular because of its central role.¹ Advances in diverse fields ranging from nanostructured materials, renewable energy and feedstock to new drugs and treatments are a few examples where transformational change is currently underway.² In this context, Chemistry plays a fundamental role by providing insights into the pathways that drive chemical processes at the molecular level, thus enabling the understanding and development of new processes. Paramount among these processes is the catalytic activation and transformation of chemical bonds.^{3,4} It is estimated that roughly 85 to 90% of the products generated by chemical industry relies on catalysts at least in one step along the reaction sequence.⁵ Asymmetric catalysts are especially important because the absolute configuration of enantiomeric centers are essential for the pharmaceutical industries.^{6,7} About twenty years ago, a new class of organocatalysts capable of carrying out chiral synthesis opened a new era in the field.⁸⁻¹² The use of small organic molecules made it possible to obtain outstanding enantiomeric excess with high reaction yield, while having with much less impact on the environment than previous metal-based strategies due to their lower toxicity to living organisms.¹³ Moreover, these “green” organocatalysts promise to use reactants more efficiently as well as lower costs associated with purification and disposal.

Here we are concerned with the class of organocatalysts based on the imidazolidinones, also known as the MacMillan catalysts,¹⁴⁻¹⁸ with two structures illustrated in Figure 1. These compounds have been shown to be active in a series of chiral transformations with high yields and enantiomeric excess, including Friedel-Crafts, Diels Alder and Mukaiyama-Michael reactions, to name a few.¹⁹⁻²⁵ Because these reactions are usually carried out under acidic conditions, it is important to establish the intrinsic protonation sites of these catalysts in solution, as the protonation equilibrium could disturb the nature of the catalytic species.²⁶⁻²⁹ One way to monitor the structures of ionic species in solution is to extract them into a mass spectrometer using ambient ionization, where a highly refined arsenal of tools are available to identify structures. In this case, however, despite the great sensitivity and dynamic detection range of this technique, traditional mass spectrometry methods are not capable of differentiating conformers and protomers.³⁰⁻³² In the past decade, nonetheless, hybrid methods that combine mass spectrometry with cryogenic ion vibrational spectroscopy have been developed that are capable of structure determination.^{33,34} In that approach, vibrational spectra are obtained for ions that are cooled after they are extracted from solution to cryogenic temperatures. Once cold (~20K), weakly bound neutral species such as H₂ or N₂ are condensed onto the ions. These adducts or “mass tags” are then photodissociated upon resonant absorption of a single photon from a tunable IR laser. Measurement of the photodissociation yield of the tagged species as a function of the laser wavelength yields the linear vibrational spectra of the mass selected ions in the gas phase. By comparing the experimental band pattern with theoretical calculations for candidate structures of the protonated imidazolidinones, it is possible to establish their protonation sites and thus infer the speciation of these catalysts in solution.^{30,35,36}

In this work, we evaluated the protonation sites of two imidazolidinones known as the first and second generation MacMillan catalysts with structures indicated in Figure 1.

The cations of both neutral compounds corresponding to addition of a proton from solution were extracted by electrospray (ESI) from a 1:1 methanol:water mixture, which is typically used as solvent for these species. Their structures were then interrogated by analyzing the pattern of vibrational bands obtained using cryogenic predissociation spectroscopy with DFT calculations as described below. To gauge whether the gas phase results obtained correlate with the solution phase, solvation effects were modeled by the SMD method, as suggested in the literature.³⁷

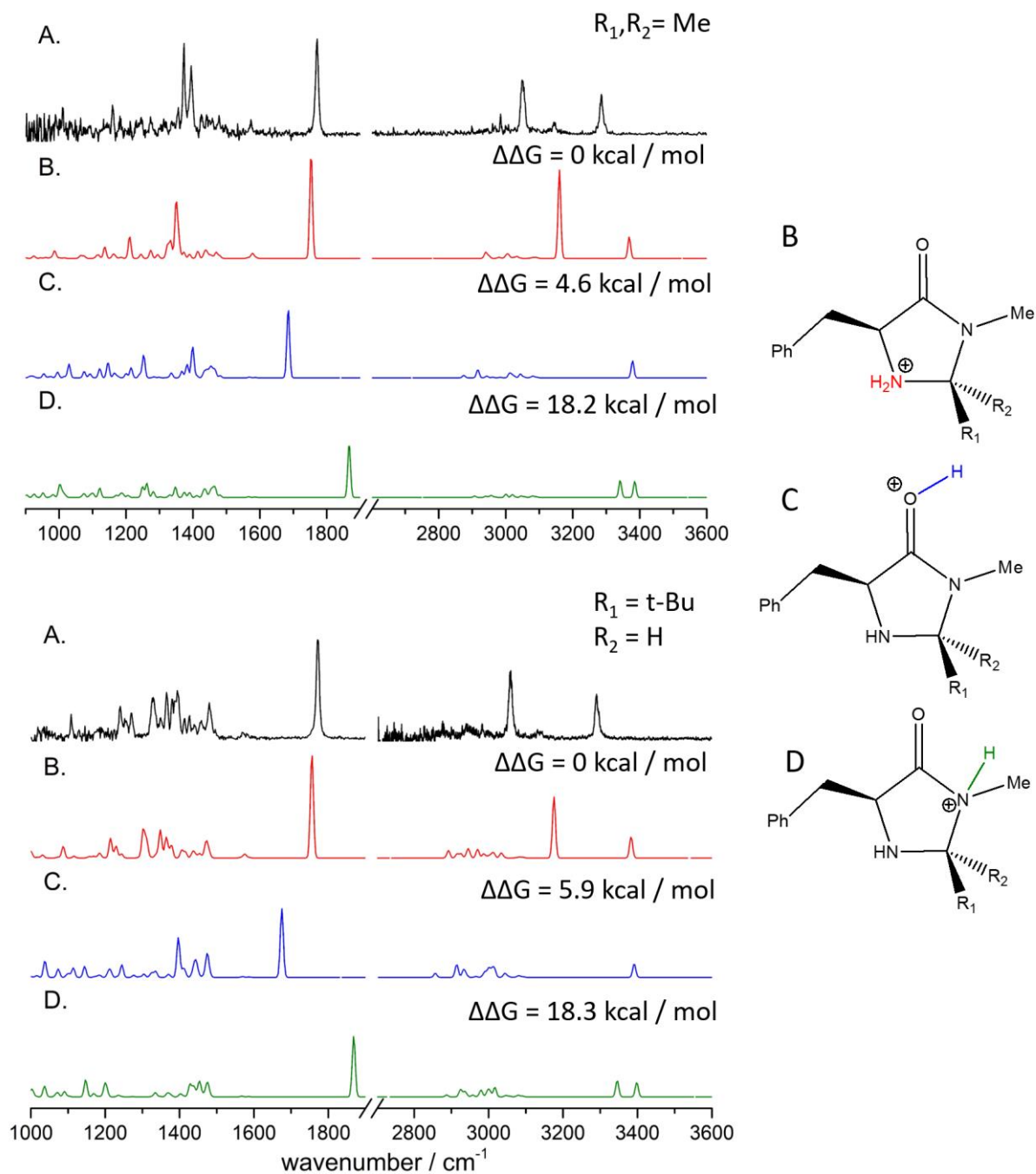


Figure 2. Top: first generation [Cat1+H]⁺ (R₁,R₂= Me) and bottom: second generation [Cat2+H]⁺ (R₁=t-But; R₂=H) MacMillan organocatalysts. A) Experimental vibrational pre-dissociation spectra of the N₂-tagged species. Theoretical vibrational spectra of the catalyst with different protonation sites at B3LYP/6-311+G(3df,2p) level of theory: (B) secondary amine; (C) amide oxygen; (D) amide nitrogen.

In summary, the theoretical analysis of the spectra strongly support the identification of the protonated secondary amine structures as the isomeric forms generated for both the Cat1 and Cat2 in an ESI ambient ion source.

Table 1. Comparison of the main observed vibrational features (Exp.) in cm^{-1} of $[\text{Cat1+H}]^+$ and $[\text{Cat2+H}]^+$ with the lowest energy protomer calculated at the B3LYP/6-311+G(3df,2p), {CAM-B3LYP/6-311+G(3df,2p)} and [M06-2X/6-311+G(3df,2p)] levels of theory

Mode assignment	$[\text{Cat1+H}]^+$		$[\text{Cat2+H}]^+$	
	Exp.	B3LYP {CAM} [M06]	Exp.	B3LYP {CAM} [M06]
NH ₂ asymmetric stretch	3286	3369 {3324} [3324]	3290	3383 {3338} [3352]
NH ₂ symmetric stretch	3051	3160 {3123} [3144]	3057	3175 {3135} [3141]
Carbonyl stretch	1770	1753 {1754} [1776]	1770	1756 {1756} [1779]
NH ₂ bending	1571	1577{1553} [1539]	1568	1575 {1551} [1542]
Multiple CH bending	1394	1350 {1362} [1373]		
	1370	1334 {1339} [1344]		
CH bending			1315-1500	1290-1490
C-N stretches			1240 - 1280	1200 - 1240

It is worth noting that, at the B3LYP/6-311+G(3df,2p) level of theory, for both $[\text{Cat1+H}]^+$ and $[\text{Cat2+H}]^+$, although the scaled low frequency bands values are in good agreement with the observed bands, the higher frequency modes are still *ca.* 100 cm^{-1} blueshifted relative to the experimental values. We note, however, that these modes involve acidic protons that are typically much more anharmonic than the skeletal modes and NH bends. This, in turn, suggests the necessity of the use of dual scale factors, as suggested for calculations carried out with more limited basis sets.³⁸

To compare the effect of different methodologies on the predicted vibrational spectra, the most stable protomer (secondary amine protonation) was also calculated in CAM-B3LYP and M06-2X functionals, as suggested in previous works of our group^{37,39} (Table 1), maintaining the same basis set. The use of these functionals led to similar improvements in the calculated spectra for both catalysts (Table 1). In the high frequency bands, the calculated bands were redshifted, reducing the observed shift for B3LYP from approximately 90 cm^{-1} to 45 cm^{-1} for the asymmetric NH₂ stretching mode. For the symmetric stretch, the improvement was a little less pronounced as the shift was reduced from approximately 115 to 80 cm^{-1} (see Figure S1 in the Supporting Information and Table 1). In the lower frequency range, the CAM-B3LYP functional led to no difference on the carbonyl stretching mode when compared to B3LYP functional. For M06-2X, however, the calculated frequency is blueshifted, but only by 6 and 9 cm^{-1} from the experimental values for $[\text{Cat1+H}]^+$ and $[\text{Cat2+H}]^+$ respectively. It is also worth noting that for $[\text{Cat1+H}]^+$, the intensity of the features observed for the bands 1394 and 1370 cm^{-1} were better represented by the CAM-B3LYP and M06-2X functionals, despite of the blueshift in the frequency values.

For both catalysts studied, the secondary amine structure is calculated to be the most stable among the candidate protomers ($\Delta\Delta G = 0 \text{ kcal/mol}$). The formation of the protonated species at the secondary amine would also be expected considering the basicity of the functional groups in this species. If the protonation at the amide would occur, for example, it would be preferable to do so at the oxygen atom ($\Delta\Delta G = 4.6$ or 5.9 kcal/mol , for $[\text{Cat1+H}]^+$ and $[\text{Cat2+H}]^+$, respectively) given that the lone pair of the amide nitrogen is less basic, with it being resonance-stabilized in this structure. For that reason, these protomers have a higher free energy ($\Delta\Delta G = 18.2$ and 18.3 kcal/mol for $[\text{Cat1+H}]^+$ and $[\text{Cat2+H}]^+$, respectively) and are therefore not expected to be observed in the gas phase.

We recognize that, despite the ample evidence for the protonation site to be located at the secondary amine, the results discussed so far are carried out in the gas phase and indirectly probe the speciation of the

cations in solution prior to extraction into the mass spectrometer by electrospray ionization (ESI). As described before in the literature, ESI ion source conditions and solvents used can promote isomerization of the species extracted, as extensively discussed for protonated *p*-aminobenzoic acid and other species.^{37,40–42}

To evaluate if the results observed in the gas phase could be transferred to solution, we carried out the continuous solvation calculation using the SMD method in water, methanol and a methanol:water 1:1 mixture at the M06-2X/cc-pVDZ level of theory, as this is the preferred functionals for SMD calculations. The relative energy obtained for each catalyst in these different condensed phases are collected in Table 2.

Table 2. Relative free energy in kcal mol⁻¹ at 298 K for the protomers of [Cat1+H⁺] and ([Cat2+H⁺] in parenthesis) in different solvents as calculated by the SMD solvation scheme at M06-2X/cc-pVDZ level of theory and gas phase results for B3LYP/6-311+(3df,2p) and M06-2X/cc-pVDZ for comparison

kcal mol ⁻¹	Gas Phase		SMD M06-2X/cc-pVDZ		
	B3LYP/ 6-311+G(3df,2p)	M06-2X/ cc-pVDZ	Methanol	Water	1:1 Mixture ^a
Secondary amine	0.0(0.0)	0.0(0.0)	0.0(0.0)	0.0(0.0)	0.0(0.0)
Amide Oxygen	4.6(5.9)	4.2(5.5)	10.5(10.6)	9.0(9.8)	9.3(10.4)
Amide Nitrogen	18.2(18.3)	15.7(17.3)	20.1(21.6)	18.6(20.4)	19.7(21.4)

a) Methanol:water 1:1 mixture in volume, 0.4:0.6 methanol:water molar fraction

These relative free energy values show that, for both catalysts and solution compositions, the same trend observed in the gas phase is predicted to be preserved in solution, with the secondary amine protonated protomer being more stable than the amide protonation at oxygen and nitrogen, respectively. In fact, when implicit solvation effects are considered, the relative free energy for the amide oxygen protonated protomer increases from approximately 5 to 10 kcal mol⁻¹ when compared to gas-phase results, while the amide protonation relative energy increases from 2 to 4 kcal mol⁻¹. This suggests that the protonated secondary amine is even more stabilized than the other protomers in solution, what is in line with what is expected for amine protonation trends from gas phase to solution.⁴³ This trend is in agreement with the pK_a values expected for the different protonation sites of imidazolidinones in water^{44,45} (Figure S2 in Supporting Information) and can be explained by the fact that the secondary amine charge is more localized than the charge in the other two protomers, which are likely more delocalized by the amide resonance. This hypothesis is supported by the NBO charge calculated at the B3LYP/6-31+G(d,p) level of theory for [Cat1+H⁺] and [Cat2+H⁺], as shown in Figure 3. As can be seen, the secondary amide shows the lowest NBO charge in comparison to the other protonation sites. Therefore, the increased charge density in the secondary amine is better solvated than the other two protomers.

These results suggest that the protonation tendency observed in the gas phase may be transferred to solution, as the relative energy values of the calculated protomers are retained.³⁷

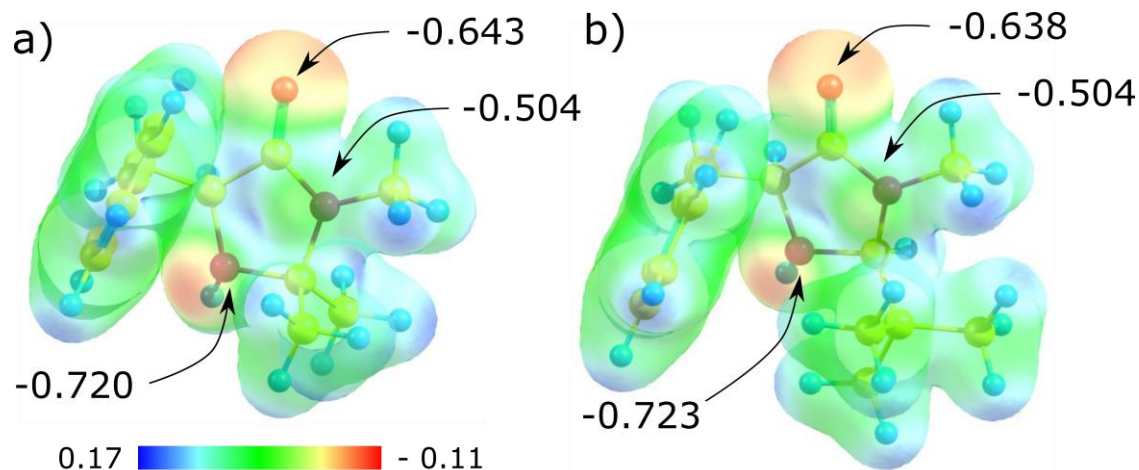


Figure 3. Representation of a) Cat1 and b) Cat2 NBOcharges at the protonation sites at the B3LYP/6-31+G(d,p) level of theory overlaid to a molecular electrostatic potential surface mapped on a 0.013458 iso-surface value. C – yellow, H – blue, N – purple and O – red.

Conclusions

The MacMillan catalysts were evaluated using Predissociation Spectroscopy and showed that the protonation site for these species occurs at the secondary amine in the gas phase, for both catalyst generations. This species is also the most stable protomer by 4.6 and 5.9 kcal mol⁻¹ for [Cat1+H⁺] and [Cat2+H⁺], respectively.

The predicted absorptions at the B3LYP/6-311+(3df,2p) level of theory accurately reproduced the bands at the lower frequency region of the spectra but underestimated the red-shifts in the NH₂ stretching bands in the highly anharmonic region above 3000 cm⁻¹. This discrepancy was less significant, however, with calculations using the CAM-B3LYP and M06-2X functionals. Solvation effects were modeled by the SMD method and showed that for both catalysts, the energy trends observed in the gas phase are conserved in solution for water, methanol and water:methanol 1:1 mixture, as expected by comparison with pK_a values in water for similar species. Therefore, we suggest that the protonation site characterized in the gas phase can be extrapolated to the situation in solution. These considerations indicate that the secondary amine is the preferential protonation site for the imidazolidinones, and the preference for this arrangement can be rationalized due to the amide resonance that reduces the amide basicity and charge density, thus stabilizing the secondary amine protonation site in the gas phase and in solution.

Experimental Section

All chemicals were purchased from Sigma Aldrich, and each of the organocatalysts Cat1 ((5S)-(-)-2,2,3-Trimethyl-5-benzyl-4-imidazolidinone monohydrochloride) and Cat2 ((2S,5S)-(-)-2-tert-Butyl-3-methyl-5-benzyl-4-imidazolidinone) were dissolved in a 1:1 methanol:acetic acid solution to a final concentration of 10⁻³ M. The ions were extracted from solution by electrospray ionization (ESI) and guided through the Yale Tandem Photofragmentation Mass Spectrometer, described in detail elsewhere.^{46,47} The ionized species pass through differential pumping stages composed of ion guides and skimmers until they reach the ion trap where they were collisionally cooled down to 60 K by a helium buffer gas containing a small fraction of nitrogen gas.

At this temperature, the ions are “tagged” with molecular nitrogen, forming an adduct that is mass selected and probed by infrared predissociation spectroscopy. An OPO/OPA parametric system (Laservision) was used to dissociate the tagged ions in a range from 1000 cm^{-1} to 3600 cm^{-1} . The photodissociation yield as a function of the laser wavelength provides de vibrational spectrum of the ion.

Experimental results were compared to electronic structure and vibrational spectrum calculations carried out at the Gaussian09 computational package.⁴⁸ All reported species were confirmed to be true minima by displaying no imaginary frequency upon vibrational analysis. DFT was used to describe the electronic structure, and different functionals, B3LYP,⁴⁹ CAM-B3LYP⁵⁰ and M06-2X⁵¹ were tested with the same basis set, 6-311+G(3df,2p). Scaling factors were used to correct the vibrational frequencies due to anharmonicity effects. The theoretical spectra were scaled by 0.967, 0.947, 0.952 at B3LYP, CAM-B3LYP and M06-2X, respectively.⁵² NBO charges were calculated at the B3LYP/6-31+G(d,p) level of theory in NBO 3.1 as implemented in Gaussian09.⁵³ MEP plots were calculated at by Gaussian G09 at this same level of theory and plotted with Chemcraft visualization program.⁵⁴ A gaussian profile of 10 FWHM was used to represent the band width in the calculated spectra reported. All optimized geometries and comparison between different methodologies can be found in the Supporting Information Material. SMD calculations were carried out for methanol and water using standard G09 parameters. Methanol:water 1:1 mixture parameters were obtained by using the experimental values for a 1:1 mixture of methanol:water ($\sim 0.4:0.6$ methanol:water molar fraction) for refraction index⁵⁵ and surface tension⁵⁶ and taking the molar fraction weighted average for the Abraham’s hydrogen bond acidity and basicity, and dielectric constant,⁵⁷ necessary for the SMD calculations. These parameters can be found in the supplementary information.

Acknowledgements

TCC would like to thank FAPESP (grants 2014/15962-5 and 2015/08539-1) and CAPES (finance code 001 and grant 23038.006960/2014-65). Individual fellowships are highly thanked by TCP to FAPESP (2017/20262-0) and LCT to PIBIC program (2015, project 598) and Santander International Mobility Fellowship (program 486/2015). M.A.J. thanks the Air Force Office of Scientific Research (AFOSR) under grant FA9550-18-1-1213, and acknowledges that the study was carried out using an instrument developed under (DURIP) FA9550-17-1-0267.

Supplementary Material

Equilibrium geometry for [Cat1+H]⁺ and [Cat2+H]⁺ with different protonation sites and predicted vibrational frequencies at B3LYP/6-311+(3df,2p) level of theory; pK_a values for imidazolidinones different protonation sites in water; comparison between diverse calculation methodologies; SMD parameters for the 1:1 methanol:water mixture.

References

1. National Research Council (U.S.). Committee on Challenges for the Chemical Sciences in the 21st Century. *Beyond the Molecular Frontier Challenges for Chemistry and Chemical Engineering*; National Academies

- Press: Washington, D.C., 2003.
- Ramsurn, H.; Gupta, R. B. *ACS Sustain. Chem. Eng.* **2013**, *1* (7), 779–797.
<https://doi.org/10.1021/sc400046y>
 - Muñoz-Muñoz, J. L.; García-Molina, M. del M.; Garcia-Molina, F.; Berna, J.; Garcia-Ruiz, P. A.; García-Moreno, M.; Rodriguez-Lopez, J. N.; Garcia-Canovas, F. J. *Mol. Catal. B Enzym.* **2013**, *91* (0), 17–24.
<https://doi.org/10.1016/j.molcatb.2013.02.003>
 - Sataraddi, S. R.; Nandibewoor, S. T. *Synth. React. Inorg. Met.-Org. Chem* **2013**, *43* (7), 809–821.
<https://doi.org/10.1080/15533174.2012.749905>
 - Chorkendorff, I.; Niemantsverdriet, J. W. *Concepts of Modern Catalysis and Kinetics*; Wiley-VCH: Weinheim, 2007.
 - Lima, V. L. E. *Quim. Nova* **1997**, *20* (6), 657–663.
<https://doi.org/10.1590/S0100-40421997000600015>
 - Federsei, H. J. *Endeavour* **1994**, *18* (4), 163–172.
[https://doi.org/10.1016/0160-9327\(95\)90525-Y](https://doi.org/10.1016/0160-9327(95)90525-Y)
 - Bertelsen, S.; Jorgensen, K. A. *Chem. Soc. Rev.* **2009**, *38* (8), 2178–2189.
<https://doi.org/10.1039/b903816g>
 - Hajos, Z. G.; Parrish, D. R. *J. Org. Chem.* **1974**, *39* (12), 1615–1621.
<https://doi.org/10.1021/jo00925a003>
 - Jensen, K. L.; Dickmeiss, G.; Jiang, H.; Albrecht, Ł.; Jørgensen, K. A. *Acc. Chem. Res.* **2011**, *45* (2), 248–264.
<https://doi.org/10.1021/ar200149w>
 - Erkkilä, A.; Majander, I.; Pihko, P. M. *Chem. Rev.* **2007**, *107* (12), 5416–5470.
<https://doi.org/10.1021/cr068388p>
 - MacMillan, D. W. C. *Nature* **2008**, *455* (7211), 304–308.
<https://doi.org/10.1038/nature07367>
 - McCort-Tranchepain, I.; Petit, M.; Dalko, P. I. *Organocatalysis*; Wiley-VCH Verlag GmbH & Co. KGaA, 2010.
<https://doi.org/10.1002/9783527628698.hgc009>
 - García-Barradas, O.; Juaristi, E. *Tetrahedron* **1995**, *51* (12), 3423–3434.
[https://doi.org/10.1016/0040-4020\(95\)00091-L](https://doi.org/10.1016/0040-4020(95)00091-L)
 - Capacci, A. G.; Malinowski, J. T.; McAlpine, N. J.; Kuhne, J.; MacMillan, D. W. C. *Nat. Chem.* **2017**, *9* (11), 1073–1077.
<https://doi.org/10.1038/nchem.2797>
 - Lee, S.; MacMillan, D. W. C. *J. Am. Chem. Soc.* **2007**, *129* (50), 15438–15439.
<https://doi.org/10.1021/ja0767480>
 - Beeson, T. D.; MacMillan, D. W. C. *J. Am. Chem. Soc.* **2005**, *127* (24), 8826–8828.
<https://doi.org/10.1021/ja051805f>
 - Austin, J. F.; Kim, S.-G.; Sinz, C. J.; Xiao, W.-J.; MacMillan, D. W. C. *Proc. Natl. Acad. Sci.* **2004**, *101* (15), 5482–5487.
<https://doi.org/10.1073/pnas.0308177101>
 - Kinsman, A. C.; Kerr, M. A. *J. Am. Chem. Soc.* **2003**, *125* (46), 14120–14125.
<https://doi.org/10.1021/ja036191y>
 - Shih, H.-W.; Vander Wal, M. N.; Grange, R. L.; MacMillan, D. W. C. *J. Am. Chem. Soc.* **2010**, *132* (39), 13600–13603.
<https://doi.org/10.1021/ja106593m>
 - Sacchetti, A.; Rossi, F.; Rossetti, A.; Pesa, R.; Mauri, E. *Chem. Pap.* **2016**, *70* (4).

<https://doi.org/10.1515/chempap-2015-0232>

22. Cho, H.; Jung, J.; Kim, J.; Park, S.; Kim, S. *Asian J. Org. Chem.* **2019**, *8* (7), 1010–1016.
<https://doi.org/10.1002/ajoc.201900220>
23. Srivastava, V. *Curr2. Organocatalysis* **2016**, *3* (3), 277–282.
<https://doi.org/10.2174/2213337202666150729230804>
24. Mountanea, O. G.; Limnios, D.; Kokotou, M. G.; Bourboula, A.; Kokotos, G. *European J. Org. Chem.* **2019**, *2019* (10), 2010–2019.
<https://doi.org/10.1002/ejoc.201801881>
25. Gramage-Doria, R.; Bellini, R.; Rintjema, J.; Reek, J. N. H. *ChemCatChem* **2013**, *5* (5), 1084–1087.
<https://doi.org/10.1002/cctc.201200541>
26. Halland, N.; Hansen, T.; Jørgensen, K. A. *Angew. Chemie Int. Ed.* **2003**, *42* (40), 4955–4957.
<https://doi.org/10.1002/anie.200352136>
27. Ahrendt, K. A.; Borths, C. J.; MacMillan, D. W. C. *J. Am. Chem. Soc.* **2000**, *122* (17), 4243–4244.
<https://doi.org/10.1021/ja000092s>
28. Burley, J. C.; Gilmour, R.; Prior, T. J.; Day, G. M. *Acta Crystallogr., Sect. C Cryst. Struct. Commun.* **2008**, *64* (1), o10–o14.
<https://doi.org/10.1107/S0108270107051396>
29. Seebach, D.; Grošelj, U.; Badine, D. M.; Schweizer, W. B.; Beck, A. K. *Helv. Chim. Acta* **2008**, *91* (11), 1999–2034.
<https://doi.org/10.1002/hlca.200890216>
30. M. Ribeiro, F. W.; Rodrigues-Oliveira, A. F.; C. Correra, T. *J. Phys. Chem. A* **2019**, *123* (38), 8179–8187.
<https://doi.org/10.1021/acs.jpca.9b05065>
31. Santos Fernandes, A.; Maître, P.; Carita Correra, T. *J. Phys. Chem. A* **2019**, *123* (5), 1022–1029.
<https://doi.org/10.1021/acs.jpca.8b09979>
32. Tripodi, G. L.; Correra, T. C.; Angolini, C. F. F.; Ferreira, B. R. V.; Maître, P.; Eberlin, M. N.; Roithová, J. *European J. Org. Chem.* **2019**, *2019* (22), 3560–3566.
<https://doi.org/10.1002/ejoc.201900171>
33. Jašíková, L.; Roithová, J. *Chem. - A Eur. J.* **2018**, *24* (14), 3374–3390.
<https://doi.org/10.1002/chem.201705692>
34. Fournier, J. A.; Johnson, C. J.; Wolke, C. T.; Weddle, G. H.; Wolk, A. B.; Johnson, M. A. *Science*. **2014**, *344* (6187).
<https://doi.org/10.1126/science.1253788>
35. Polfer, N. C. *Chem. Soc. Rev.* **2011**, *40* (5), 2211–2221.
<https://doi.org/10.1039/c0cs00171f>
36. Masson, M. A. C.; Karpfenstein, R.; de Oliveira-Silva, D.; Teuler, J.-M. M.; Archirel, P.; Maître, P.; Correra, T. *C. J. Phys. Chem. B* **2018**, *122* (43), 9860–9868.
<https://doi.org/10.1021/acs.jpcc.8b06523>
37. C. Correra, T.; S. Fernandes, A.; M. Reginato, M.; C. Ducati, L.; Berden, G.; Oomens, J. *Phys. Chem. Chem. Phys.* **2017**, *19* (35), 24330–24340.
<https://doi.org/10.1039/C7CP04617K>
38. Halls, M. D.; Velkovski, J.; Schlegel, H. B. *Theor. Chem. Acc.* **2001**, *105* (6), 413–421.
<https://doi.org/10.1007/s002140000204>
39. Rodrigues-Oliveira, A. F.; M. Ribeiro, F. W.; Cervi, G.; C. Correra, T. *ACS Omega* **2018**, *3* (8), 9075–9085.
<https://doi.org/10.1021/acsomega.8b00815>

40. Patrick, A. L.; Cismesia, A. P.; Tesler, L. F.; Polfer, N. C. *Int. J. Mass Spectrom.* **2017**, *418*, 148–155.
<https://doi.org/10.1016/j.ijms.2016.09.022>
41. Chang, T. M.; Prell, J. S.; Warrick, E. R.; Williams, E. R. *J. Am. Chem. Soc.* **2012**, *134* (38), 15805–15813.
<https://doi.org/10.1021/ja304929h>
42. Warnke, S.; Seo, J.; Boschmans, J.; Sobott, F.; Scrivens, J. H.; Bleiholder, C.; Bowers, M. T.; Gewinner, S.; Schöllkopf, W.; Pagel, K.; Von Helden, G. *J. Am. Chem. Soc.* **2015**, *137* (12), 4236–4242.
<https://doi.org/10.1021/jacs.5b01338>
43. Brauman, J. I.; Riveros, J. M.; Blair, L. K. *J. Am. Chem. Soc.* **1971**, *93* (16), 3914–3916.
<https://doi.org/10.1021/ja00745a016>
44. Cox, R. A. *Can. J. Chem.* **1998**, *76* (6), 649–656.
<https://doi.org/10.1139/v98-012>
45. Kaljurand, I.; Kütt, A.; Sooväli, L.; Rodima, T.; Mäemets, V.; Leito, I.; Koppel, I. A. *J. Org. Chem.* **2005**, *70* (3), 1019–1028.
<https://doi.org/10.1021/jo048252w>
46. Wolk, A. B.; Leavitt, C. M.; Garand, E.; Johnson, M. A. *Acc. Chem. Res.* **2014**, *47* (1), 202–210.
<https://doi.org/10.1021/ar400125a>
47. Yang, N.; Duong, C. H.; Kelleher, P. J.; Johnson, M. A. *Nat. Chem.* **2019**.
<https://doi.org/10.1038/s41557-019-0376-9>
48. Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Mennucci, B.; Petersson, G. A.; Nakatsuji, H.; Caricato, M.; Li, X.; Hratchian, H. P.; Izmaylov, A. F.; Bloino, J.; Zheng, G.; Sonnenberg, J. L.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Montgomery, J. A.; Peralta, J. E.; Ogliaro, F.; Bearpark, M.; Heyd, J. J.; Brothers, E.; Kudin, K. N.; Staroverov, V. N.; Keith, T.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Rega, N.; Millam, J. M.; Klene, M.; Knox, J. E.; Cross, J. B.; Bakken, V.; Adamo, C.; Jaramillo, J.; Gomperts, R.; Stratmann, R. E.; Yazyev, O.; Austin, A. J.; Cammi, R.; Pomelli, C.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Zakrzewski, V. G.; Voth, G. A.; Salvador, P.; Dannenberg, J. J.; Dapprich, S.; Daniels, A. D.; Farkas, Ö.; Foresman, J. B.; Ortiz, J. V.; Cioslowski, J.; Fox, D. J. *Gaussian 09, Revision D.01*; 2013.
49. Becke, A. D. *J. Chem. Phys.* **1993**, *98* (2), 1372–1377.
<https://doi.org/10.1063/1.464304>
50. Yanai, T.; Tew, D. P.; Handy, N. C. *Chem. Phys. Lett.* **2004**, *393* (1–3), 51–57.
<https://doi.org/10.1016/j.cplett.2004.06.011>
51. Zhao, Y.; Truhlar, D. *Theor. Chem. Acc.* **2008**, *120* (1–3), 215–241. <https://doi.org/10.1007/s00214-007-0310-x>
<https://doi.org/10.1007/s00214-007-0310-x>
52. Russell D. Johnson III; Johnson III, R. D. *NIST Computational Chemistry Comparison and Benchmark Database, NIST Standard Reference Database Number 101 Release 14*; 2006.
53. Glendening, E. D.; Reed, A. E.; Carpenter, J. E.; Weinhold, F. NBO Version 3.1. 1998.
54. Zhurko, G. A. *Chemcraft - Graphical Program for Visualization of Quantum Chemistry Computations*, build 536.; 2018.
55. Herráez, J. V.; Belda, R. *J. Solution Chem.* **2006**, *35* (9), 1315–1328.
<https://doi.org/10.1007/s10953-006-9059-4>
56. Vazquez, G.; Alvarez, E.; Navaza, J. M. *J. Chem. Eng. Data* **1995**, *40* (3), 611–614.
<https://doi.org/10.1021/je00019a016>

57. Marenich, A. V.; Cramer, C. J.; Truhlar, D. G. *J. Phys. Chem. B* **2009**, *113* (18), 6378–6396.

<https://doi.org/10.1021/jp810292n>

This paper is an open access article distributed under the terms of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>)