

**Mechanisms of the Aqueous Solvolysis of the Ring-Opening of the Lactone Ring of Goniiodomin A**

**ABSTRACT**

The phycotoxin Goniiodomin A is a Dinoflagellate of the Alexandrium genus. It is a macrocyclic lactone that contains six oxygen heterocycles with a sixth ring being a hemiketal attached to the carboxylate oxygen that forms the lactone ring. In is found widely in sea water and is known to break the lactone connection slowly in water. While lactones are known to undergo hydrolysis to produce a carboxylic acid and an alcohol, the structure of the lactone is such that it is very likely that nucleophilic substitutions play the key role in the loss of the lactone via the carboxylate acting as a leaving group. The cleavage of the lactone is sensitive to the pH of the water as one changes the pH from ~6 to 8 the rate of cleavage dramatically increases. Treatment of kinetic data previously published indicates that there are likely two competing first-order reactions, an  $S_N1$  and an intramolecular  $S_N2$  reaction.

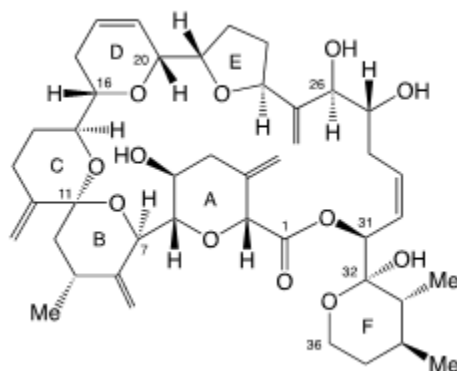
**Keywords :** marine phytotoxin, mechanism of lactone hydrolysis

**1. INTRODUCTION**

The phycotoxin Goniiodomin A (GDA,**1**) is a Dinoflagellate of the Alexandrium genus. GDA is produced by four species of the genus - *hiranoi*, *A. pseudogonyaulax*, *A. monilatum* and *A. taylorii*. [1] [2] [3] [4] While Morakami and Takeda determined the structure of GDA [1] [5] and its structure was recently confirmed by X-ray crystallography. [6] GDA is a polyketide macrolide containing six oxygen heterocycles, five macrocyclic rings with two of the five constituting spiro ketals. The sixth ring contains a hemiketal, which will be the focus in this manuscript.

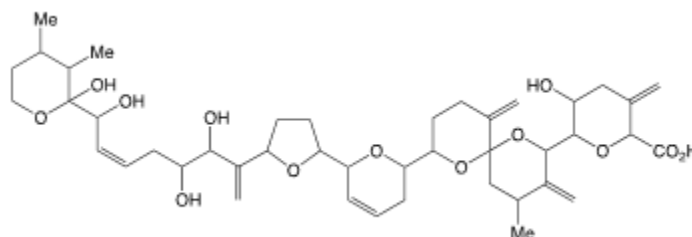
In 2020 Onofrio reported that GDA degrades in pure water, presumably at pH 7 and significantly faster in sea water where the pH is approximately 8. [7] Further studies by Hintze [8] found that the rate of

degradation of **1** in deionized water (~pH 6.5), MilliQ-Q water (pH 7.2) and in culture medium (pH 8.2) became



GDA (**1**)

significantly faster with increase of the pH. The product of the degradation is known to be a carboxylic acid such as the seco acid of GDA (**2**). At first blush one might expect the reaction to proceed with



GDA-SA (**2**)

simply hydrolysis of the lactone. However, an alternative mechanism would be that of solvolysis with the leaving group being the carboxylate anion with production of a carbocation at C<sub>31</sub>. While the carboxylate is one of the worst known leaving groups in nucleophilic substitutions, there are two aspects the structure of **1**, which might make the solvolytic mechanism the preferred one. The first is that the forming carbocation is allylic since the **the** developing positive charge on C<sub>31</sub> is allylic. The second one is the OH group at C<sub>32</sub> might interact as a neighboring group in which case an epoxide would be formed or possibly the epoxide might simply exist on the pathway of a concerted reaction. In the latter case it would not form the C<sub>31</sub> carbocation **would not be formed** but rather the C<sub>32</sub> carbocation, which would be stabilized by the ether oxygen lone pair. The solvolytic degradation of **1** was confirmed by Harris with H<sub>2</sub><sup>18</sup>O. [9]

The purpose of this study is to treat computationally a model system to determine the solvolytic mechanism. The rather dramatic change in rate vs pH found by Hintze[8] suggests that there might be two competing mechanisms

## 2. KINETICS OF THE SOVOLYTIC REACTIONS

As mentioned above, Hintze studied the degradation of **1** at three different pHs. Included in his thesis were the plots of his kinetic data (concentration vs. time in days). It was found that these kinetic data all arose from first order reactions – straight lines were found when plotting the natural log of the concentration at time  $t$  divided by the initial concentration ( $t=0$ ). Our kinetic treatment of these data are presented in Fig. 1.

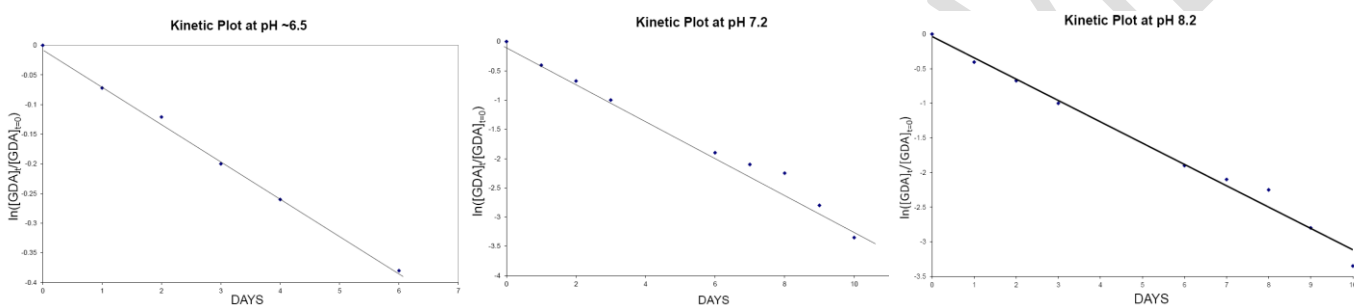
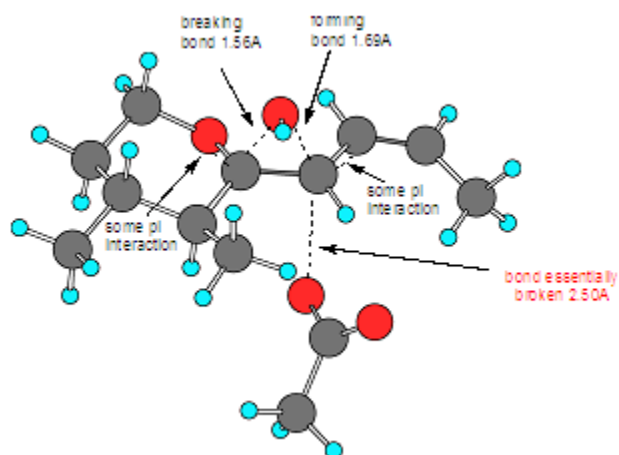


Fig. 1. The kinetically treated data of Hintze at three different pH

The  $t_{1/2}$  half-lives of these three kinetic treatments are 3.4, 2.0 and 1.2 days, respectively. The dramatic change in  $t_{1/2}$  lives suggests that there might be two competing first order reactions occurring, one being dominant at pH 6 and the other at pH 8. What is different at these two pHs? Given that it is likely that both are nucleophilic substitutions, it is possible that the hydroxide ion might be involved in the major reaction occurring at pH 8, since the hydroxide concentration is two orders of magnitude higher at pH 8 than at pH 6. At this point it was possible that the dominant reaction at pH 6 is  $S_N1$  and at pH 8  $S_N2$ . The calculations described below were done to confirm this proposal.

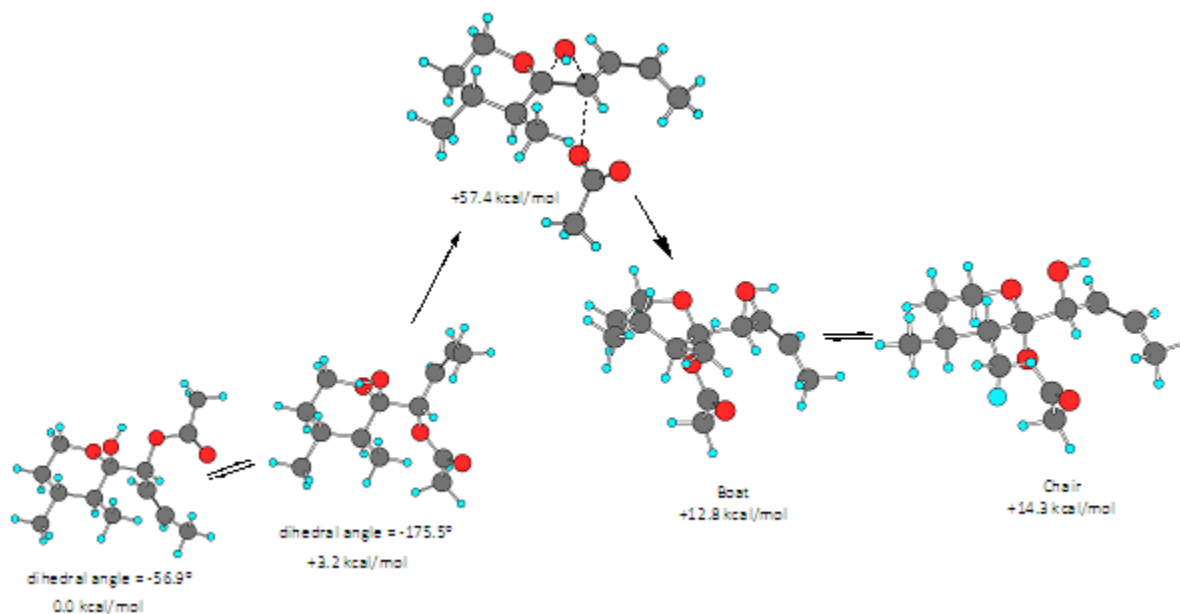
## 3. DFT CALCULATIONS.

The calculations [10] described below were done to explain the proposal based on the kinetic results. At pH 6 most likely an  $S_N1$  reaction occurs with a unimolecular ionization with the carboxylate acting as the leaving group. Using the model system mentioned above a transition structure (4) was located, which suggested a concerted rearrangement with the forming carbocation (Fig. 2). As seen the ionization is assisted by the



**Fig. 2.** The concerted, but asynchronous, transition structure for the  $S_N1$  reaction dominant at pH 6

double bond attached to  $C_{31}$  as well as migration of the OH group from  $C_{32}$  to  $C_{31}$  and is further stabilized by interaction of the hemiacetal oxygen attached to  $C_{32}$ , which is crucial in stabilizing the final rearranged carbocation. Intrinsic Reaction Coordinate (IRC) calculations produced the reactant and product as shown in Fig. 3. In the case of the reactant the IRC gave a conformer of the that in the x-ray structure. [6]



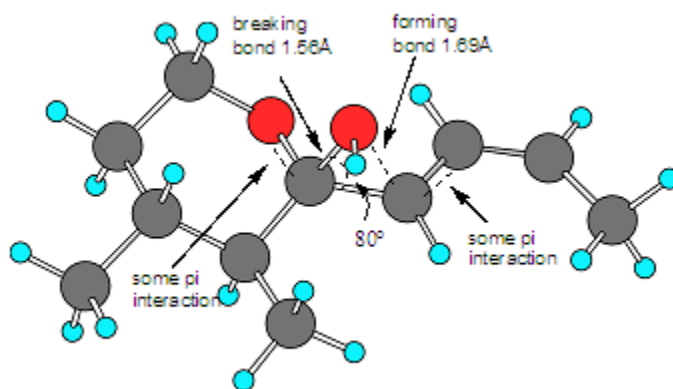
**Fig. 3** The computed concerted reaction pathway for the  $S_N1$  reaction dominant at pH 6

The IRC reactant conformer has a dihedral angle of the two carbons connecting the acyclic chain to the carboxylate carbon of the lactone of  $-175.5$  and its conformer's dihedral angle is  $-56.9^\circ$ . The x-ray conformer

has a dihedral angle that would prevent OH group participation in the concerted reaction. The IRC gave a product with the six-membered ring in a boat conformation. Interestingly the chair conformer of the product was found to be higher in energy than the boat conformer.

While the DFT calculations were carried out in a water medium,  $S_N1$  reactions are not properly described by these calculations [11], since it is known that it first forms a contact ion pair, which in turn forms a solvent separated ion pair. With this type of calculation, these ion pairs are not taken into account and the leaving group has only one choice, to collapse to the newly formed carbocation produced by the concerted reaction. The activation energy is also very large, since while the calculation is modeled with a surrounding solvent of water, no water molecules are present in the vicinity of the ionization of the C-O bond to “assist” in the ionization, such as the formation of the solvent separated ion pair. Consequently the computed activation energy is likely much higher than the experimental one.

However, it is possible to obtain a better estimate of the overall energetics of carbocation reactions by carrying out calculations on the carbocation which is formed. There is good evidence for this in recent computations of carbocation rearrangements in terpene chemistry. [12] Beginning with the carbocation that is formed by loss of the carboxylate group in our model system a transition structure (Fig. 5) was located very similar to that located above (Fig. 2).



**Fig. 4. The  $S_N1$  transition structure (with an overall positive charge) located for the model system carbocation**

Perhaps more dramatic is the difference in the overall energetics of the carbocation reaction (Fig. 5) than

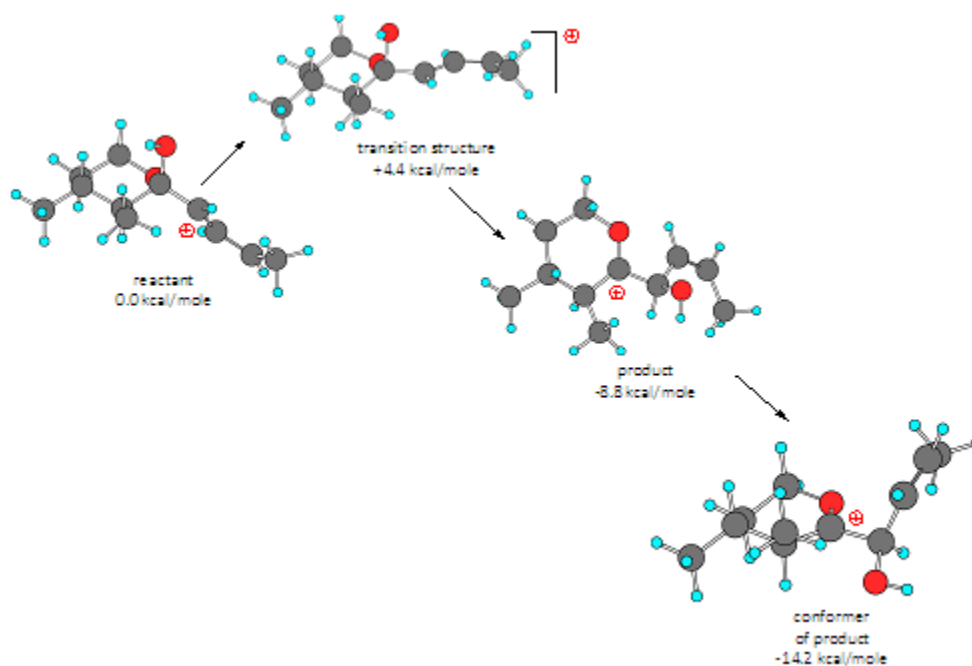


Fig. 5. The overall DFT pathway of the carbocation in the  $S_N1$  reaction dominant at pH 6

that in Fig. 3. Note the very low activation energy (+4.0 kcal/mol) and the overall energy of reaction between the reactant and the IRC product of -14.5 kcal/mole. It is apparent that major driving force for this concerted rearrangement is the ability of the ether oxygen in the oxane ring to stabilize the carbocation on C<sub>32</sub>. This is not unexpected given that the methoxy group has a large negative Hammett  $\sigma_{para}$  constant.

Turning to the dominant reaction at pH 8, a DFT study was conducted with the addition of a hydroxide ion. A transition structure was located (Fig. 6) for the removal of the hydrogen on the OH group on C<sub>32</sub>.

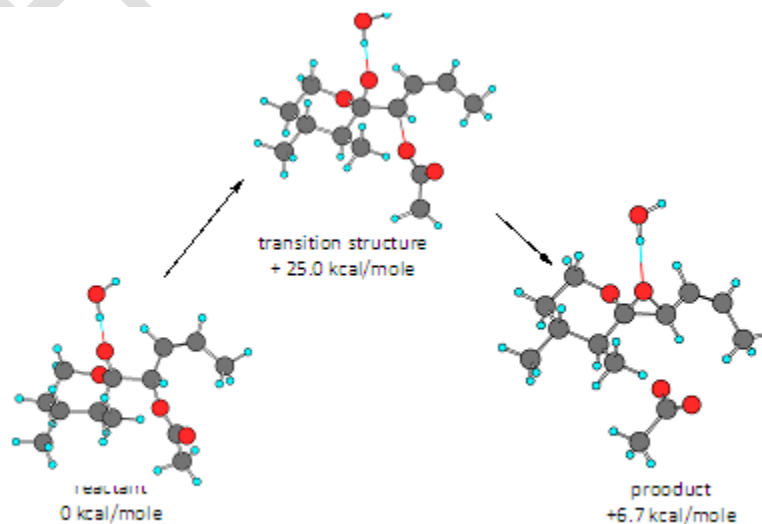


Fig. 6. The reaction pathway which includes a hydroxide ion of the solvolysis of the ester at pH 8 or under basic conditions

Again, the overall reaction studied in the modeled water solution suffers from no individual water molecules present in the vicinity of the structures along the reaction pathway. For example, the carboxylate leaving group would be hydrated with water molecules, which would of course likely change the overall energetics of the reaction. The pathway was also studied beginning with the anion of the reactant (Fig. 7). It is seen that

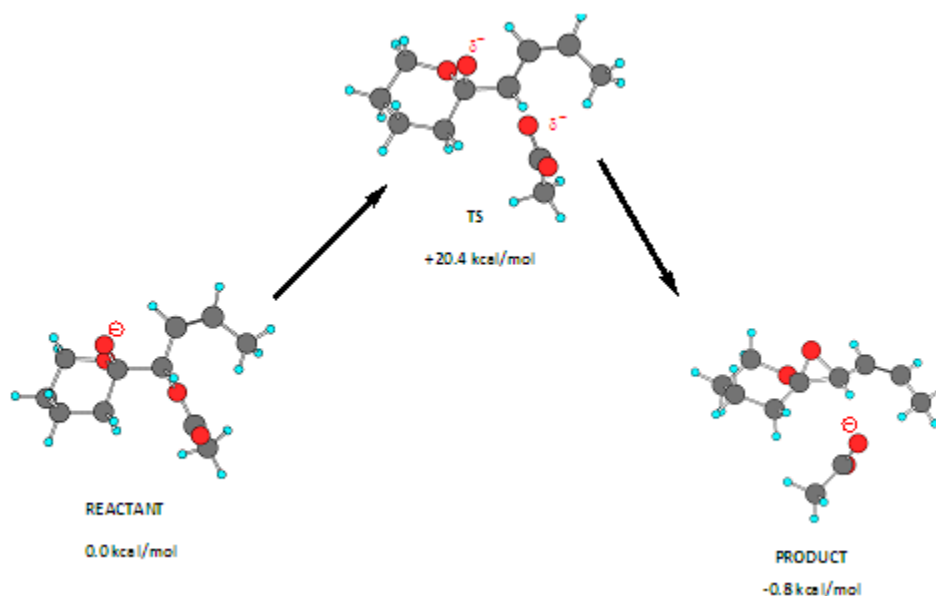


Fig. 7. The  $S_N2$  reaction at pH 8 beginning with the proton removed from the oxygen at  $C_{32}$

this reaction is indeed an intramolecular  $S_N2$  reaction. In this case the attacking nucleus is the negatively charged oxygen on  $C_{32}$ . While the overall reaction energy is now negative, it is rather small at  $-0.8 \text{ kcal/mol}$ . Again, these energetics are different from what one would expect experimentally because of lack of water molecules interacting with the structures along the reaction pathway.

The question at this point is: Why is the dominant reaction at pH8 a first order reaction rather than second order, since two reactants are involved prior to rate determining step? Note that one of the reactants is the hydroxide ion, which has a much higher concentration than the reactant. Hence this is simply a pseudo first order reaction, since the *change* in the concentration of the substrate is far more than that of the hydroxide.

Whether or not the epoxide is stable or not under the basic reaction conditions is an interesting question. It has two pathways that ring opening of the epoxide can occur as shown in Fig. 8, with the two products being diastereomers and the product giving rise to the *R-R* one favored because of steric hindrance at the carbon undergoing the nucleophilic attack. Interestingly the  $S_N1$  reaction would also give rise to a  $C_{31}S$ - $C_{32}S$  glycol. However under the aqueous conditions the  $C_{32}$  carbon might very well be racemized by mutarotation of the hemiketal.

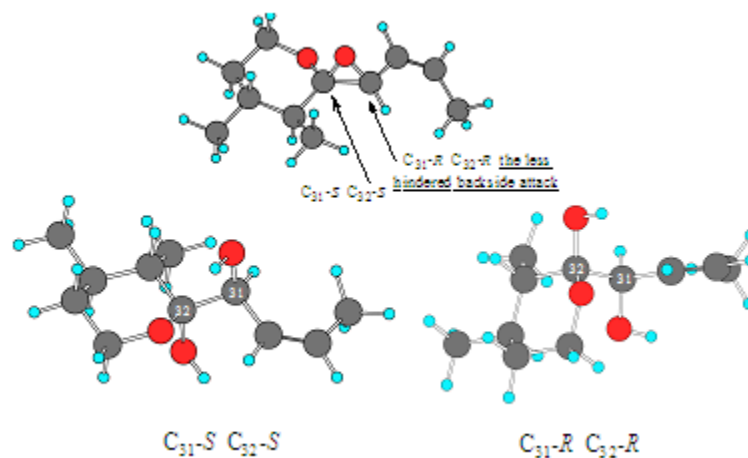


Fig. 8. Potential products of base catalyzed opening of the epoxide

#### 4. CONCLUSIONS

The DFT results suggest that indeed there might be competing first-order nucleophilic reactions. At the lower pH the  $S_N1$  dominates and at higher pH and intramolecular  $S_N2$  reaction is predicted to dominate.

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