

Capturing non-local effects when fragmenting molecules for quantum chemical torsion scans

This manuscript ([permalink](#)) was automatically generated from [ChayaSt/fragmenter-manuscript@b95643d](#) on January 29, 2020.

Authors

- **Chaya D Stern**

 [0000-0001-6200-3993](#) ·  [ChayaSt](#) ·  [SternChaya](#)

Tri-Institutional PhD Program in Chemical Biology, Memorial Sloan Kettering Cancer Center, New York, New York 10065, USA; Computational and Systems Biology Program, Memorial Sloan Kettering Cancer Center, New York, New York 10065 USA · Funded by Grant XXXXXXXX

- **Christopher I Bayly**

 [XXXX-XXXX-XXXX-XXXX](#) ·  [johndoe](#) ·  [johndoe](#)

Department of Something, University of Whatever · Funded by Grant XXXXXXXX

- **Daniel G A Smith**

 [XXXX-XXXX-XXXX-XXXX](#) ·  [dgasmith](#) ·  [johndoe](#)

Department of Something, University of Whatever · Funded by Grant XXXXXXXX

- **Josh Fass**

 [XXXX-XXXX-XXXX-XXXX](#) ·  [maxentile](#) ·  [johndoe](#)

Department of Something, University of Whatever · Funded by Grant XXXXXXXX

- **Lee-Ping Wang**

 [XXXX-XXXX-XXXX-XXXX](#) ·  [leeping](#) ·  [johndoe](#)

Department of Something, University of Whatever · Funded by Grant XXXXXXXX

- **David L Mobley**

 [XXXX-XXXX-XXXX-XXXX](#) ·  [davidlmobley](#) ·  [johndoe](#)

Department of Something, University of Whatever · Funded by Grant XXXXXXXX

- **John D Chodera**

 [0000-0003-0542-119X](#) ·  [jchodera](#) ·  [jchodera](#)

Computational and Systems Biology Program, Memorial Sloan Kettering Cancer Center, New York, New York 10065 USA · Funded by Grant XXXXXXXX

Abstract

Accurate molecular mechanics force fields for small molecules are essential for predicting protein-ligand binding affinities in drug discovery and understanding the biophysics of biomolecular systems. The accuracy of torsion parameters is important for determining the conformational distribution of molecules, and can have a large effect on computed properties like binding affinities. Torsion parameters are usually fit to computationally costly quantum chemical (QC) torsion scans that scale poorly with molecule size. To reduce computational cost and avoid the complications of distant intramolecular interactions, molecules are generally fragmented into smaller entities to carry out QC torsion scans. Poor fragmentation schemes, however, have the potential to significantly disrupt electronic properties of the region around the torsion, leading to poor representation of the real chemical environment and the resulting torsion energy profile. Here, we show that a rapidly computable quantity, the fractional Wiberg bond order (WBO), is sensitive to the chemical environment of bonds, and can be used as a useful surrogate to assess the robustness of fragmentation schemes and identify conjugated bond sets. We use this concept to construct a validation set consisting of exhaustive fragmentations of druglike organic molecules (and their corresponding WBO distributions derived from accessible conformations) that can be used to evaluate fragmentation schemes. To illustrate the utility of the WBO in assessing fragmentation schemes that preserve the chemical environment, we propose a new fragmentation scheme that uses rapidly-computable AM1 WBOs, available essentially for free as part of an AM1-BCC partial charge assignment process, to maximize the chemical equivalency of the fragment and the substructure in the larger molecule.

1. Introduction

Molecular mechanics (MM) small molecule force fields are essential to the molecular design for chemical biology and drug discovery, as well as the use of molecular simulation to understand the behavior of biomolecular systems. However, small molecule force fields have lagged behind protein force fields given the larger chemical space these force fields must cover to provide good accuracy over the space of druglike ligands, common metabolites and other small biomolecules [1,2]. Torsion parameters are particularly problematic because historical approaches to their determination tend to produce parameters that generalize poorly [3]. This lack of generalizability has let many practitioners to aim to improve the force field accuracy by refitting torsion parameters for individual molecules in a semi automated bespoke fashion [4,5,6]. This leads to a significant barrier to setting up simulations for new projects and may not produce generalizable parameters.

In many molecular mechanics force fields (e.g., Amber [7], CHARMM [8], OPLS [9]) a low-order Fourier series, such as a cosine series, is often used to represent the contribution of torsion terms to the potential energy.

$$E_{tor} = \sum_{n=1}^N \frac{V_n}{2} [1 + \cos[n(\theta - \gamma)]]$$

where V_i is the torsion force constant which determines the amplitudes, n is the multiplicity which determines the number of minimas, and γ is the phase angle which is sometimes set to 0° or 180° to enforce symmetry around zero. In most force fields N is 4, however, some go up to 7. The torsion potential energy parameters such as amplitudes and phase angles for each Fourier term, are generally fit to the residual difference between gas phase quantum chemistry (QC) torsion energy profile and the non-torsion MM parameters [10]. The QC torsion energy profile is generated by fixing the torsion atoms and geometry minimizing all other atomic positions. Neighboring torsions can have correlated conformational preferences the low-order Fourier series does not capture[11]. 2D spline

fits, such as CMAP [12,13], have become a popular way to model non-local correlations by fitting residuals between the 2D QC torsion energy profile and the 2D MM torsion energy profile.

In order to produce a quantum chemical energy profile representing the chemical environment that the torsion of interest is to be fit to, the quantum chemical torsion profile is generally computed on smaller fragments generated by a fragmentation-and-capping process for two main reasons.

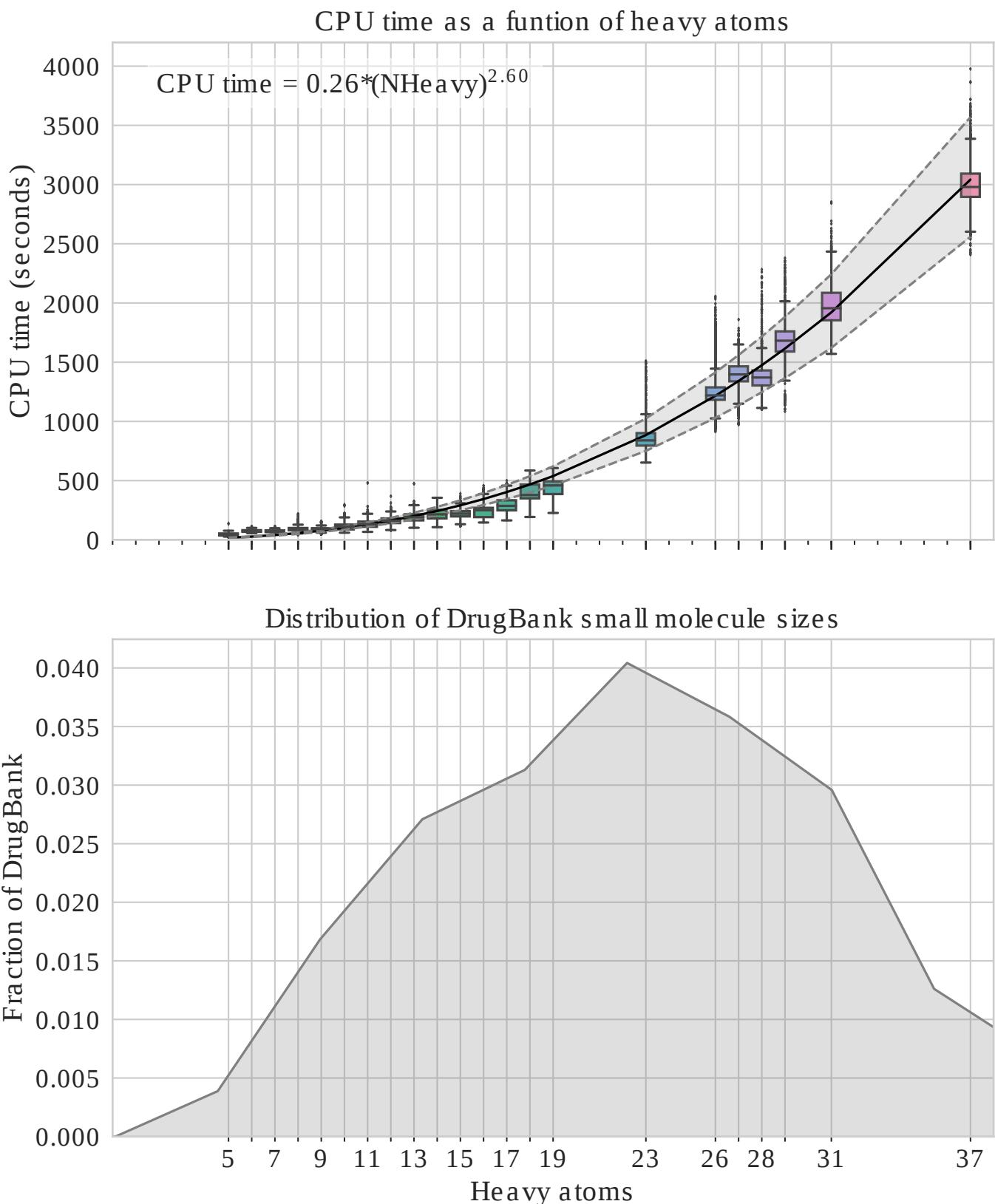


Figure 1: Fragmenting molecules is necessary to avoid high computational cost of generating QC data. [A] CPU time (wall time * nthreads) of one QC gradient evaluation at B3LYP-D3(BJ)/DZVP level of theory [14,15] vs number of heavy atoms in molecules. All computations shown here were run on an Intel(R) Xeon(R) CPU E5-2697 v4 @ 2.30GHz.

Empirically, gradient evaluations grow as $O(N^{2.6})$ where N is the number of heavy atoms. The scaling is similar on other processors shown in SI Figures 15 & 16. The black curves shows a power law fit to the data, and the grey is the 95% CI of the curve estimate. [B] Smoothed histogram of heavy atoms in small molecules from DrugBank. The average druglike molecules has 25 heavy atoms.

1. **Computational efforts scale poorly with molecule size.** Generating one dimensional QC torsion profiles are computationally expensive and become increasingly inefficient for larger molecules and/or higher dimensional QC torsion profiles. QC calculations scale poorly with the number of basis sets N , like $O(N^M)$ where formally, $M \leq 4$ for hybrid DFT. With modern implementations, hybrid DFT scales asymptotically as $N^{2.2-2.3}$ [16]. Using QC Archive data [17], we found that practically, hybrid DFT grows like $N^{2.6}$ as shown for gradient evaluations in Figure 1, A. To achieve good sampling to adequately fit the torsions, constrained geometry optimizations need to be calculated at $\leq 15^0$ intervals for a minimum of 24 constrained geometry optimizations. To avoid hysteresis in the energy profile due to orthogonal degrees of freedom [18], methods like wavefront propagation [hold for CITE torsiondrive paper] are used. This adds a factor of 2D, where D is the dimension of the QC scan, to the computational cost. We found (SI Figure 18) that on average ~60 optimizations are needed for a 1D wavefront propagated torsion drive to converge. We also found that roughly 20 gradient evaluations are needed for an optimization to converge for the molecules we optimized (SI Figure 17). Figure 1, B shows a smoothed histogram of the distribution of the number of heavy atoms in FDA approved small molecules taken from DrugBank [19]. For an average druglike molecule of 25 heavy atoms, we can estimate the cost of a 1D torsion scan to be $60 * 20 * 0.26 * 25^{2.6} \approx 1,000,000$ CPU seconds. Reducing the size of the molecule to 15 heavy atoms will reduce the cost the torsion scan by an order of magnitude ($60 * 20 * 0.26 * 15^{2.6} \approx 300,000$ CPU seconds)

2. **Intramolecular interactions complicate torsion drives and torsion parameter fitting** In larger molecules, there is a greater potential for the torsion atoms to interact with other degrees of freedom and convolute the energy profile. While this can also happen in smaller molecules such as ethylene glycol [hold to CITE torsiondrive paper] this problem is reduced when a minimal model molecule is used, albeit not completely eliminated.

While a number of algorithms for fragmenting large molecules into smaller molecular fragments have been previously proposed, few are appropriate for the purpose of generating high-quality torsion scans. Many of these algorithms fall into two categories: 1. fragmentation for synthetic accessibility [20,21,22] and 2. fragmenting molecules to achieve linear scaling for QC calculations [23,24,25,26]. Fragmentation schemes for synthetic accessibility find building blocks for combinatorial and fragment based drug design. Cleavage happens at points where it makes sense for chemical reactions and do not consider how those cuts affect the electronic properties of the fragments. For retrosynthetic applications, many cleavage points are at functional groups because those are the reactive sites of molecules. However, for our application, we especially do not want to fragment at these reactive points given how electron rich they are and how much the electronic density changes when they are altered. Fragmentation algorithms for linear scaling such as Divide-and-Conquer methods [27], effective fragment potential method [28] and systematic molecular fragmentation methods [26] require the users to manually specify where the cuts should be or which bonds not to fragment. Furthermore, besides the scheme suggested by Rai et. al. [29], none of these methods address the needs specific to fragmenting molecules for QC torsion scans. Fragments need to include all atoms involved in 1-4 interactions, since they are incorporated in the fitting procedure. We also need a systematic way to determine if remote substituents change the barrier to rotation significantly for the central bond of interest.

In this work, we use the Wiberg Bond Order (WBO) [30], which is both simple to calculate from semi-empirical QC methods and is sensitive to the chemical environment around a bond. WBOs quantify electron population overlap between bonds, or the degree of binding between atoms. Bond orders

are correlated with bond vibrational frequencies [31,32] and WBOs are used to predict trigger bonds in high energy-density material because it is correlated with the strength of the bond [33]. Wei, et. al. [34] have shown that simple rules for electron richness of aromatic systems in diaryls are good indication of torsion force constants (specifically for V_2), however, this measure was only developed for diaryls and it does not take into account substituents beyond the aromatic ring directly adjacent to the bond. Here, we develop an approach that uses the WBO to validate whether a fragmentation scheme corrupts the local chemical environment of interest, with a particular focus on fragmentation schemes suitable for QC torsion drives. Our approach uses simple heuristics to arrive at a minimal fragment for QM torsion scan that is representative of the torsion scan of the substructure in the parent molecules. For a central bond, include all atoms that are involved in the sterics of a torsion scan. Then use the WBO as a surrogate signal to determine if the fragment needs to be grown out more to restore the correct electronics around the central bonds.

The paper is organized as follows: Section 2 provides a mathematical and physical definition of the problem. Section 3 provides the motivation for using the WBO as a surrogate, evaluates how robust it is, proposes a minimal fragmentation scheme and describes a rich validation set that can be used to benchmark fragmentation schemes. Section 4 provides a discussion of the implications of this study and section 5 provides detailed methods.

2. Theory and definitions

2.1 A mathematical definition of the problem

A molecular structure can be modeled as a degree bounded graph $G = (V, E)$ where V are the nodes and E are the edges. In a molecular graph, the nodes correspond to the atoms and the bonds correspond to the edges. We define rotatable bonds as bonds that are not in rings and they are subset of edges, $e' \in E$ and G' as a set of subgraphs, where each subgraph $G' = (v', e')$:

- The number of atoms in the subgraph, $|v'|$ are $4 \leq |v'| \leq |V|$
- e' and all $e \in E$ such that e, e' share a vertex
- all v adjacent to e
- if $v \in v'$ is in a ring in G , the ring should be included in G'

The weights on the edges are given by $\delta(e'; G')$ where δ is the RMSE of the torsion potential around the central, rotatable bond in the full graph vs. a subgraph. Since $\delta(e'; G')$ is computationally expensive, we use a surrogate $\gamma(e'; G')$ which we define as the difference of the WBO on the central bond e' in G' and in the full graph G . In order to calculate the WBO, we need to cap open valences. The rule we use are defined in section 3.4.

We want to minimize $\gamma(e'; G')$, while also minimizing the cost of each subgraph. We define the cost as estimated in 1

$$c(G') \propto (|v'|)^{2.6}$$

Which leads to minimizing

$$\sum_{G'} |\gamma(e', G') - c(G')| \tag{1}$$

The search space of G' is combinatorial and its upper bound is $\binom{|V|}{4} + \binom{|V|}{5} + \dots + \binom{|V|}{|V|}$ since all subgraphs G' need to be connected and rings are not fragmented. To reduce the search space, we also define a list of functional groups that should not be fragmented in [2](#).

Given how large the search space can become, we use several heuristics as described in section 3.4.

2.2 Physical definitions

The torsion energy function (or profile) of a bond is determined by a combination of effects from conjugation, hyperconjugation, sterics and electrostatics [[34](#),[35](#),[36](#),[37](#)]. Most of these effects are non-local. For this study we define local as atoms within two bonds of the central bond of a torsion and remote as any atom beyond those two bonds.

Sterics and elecrostatics are in principle handled by non bonded terms in most force fields, so a torsion profile should represent conjugation or hyperconjugation, and only the 1-4 electrostatics. Using small fragments to generate QC torsion profiles reduces non-local electrostatics and steric interactions from convoluting the data. However, conjugation and hyperconjugation are non local properties and remote chemical changes can influence the extent of conjugation and / or hyperconjugation. In this study, we aim to mitigate the effects of remote chemical changes on conjugation and hyperconjugation by understanding how the strength of the central bond changes with remote chemical changes. Here we define conjugation and hyperconjugation and how we use these terms in this paper.

Conjugation and hyperconjugation describes the sharing of electron density across several bonds. Conjugation is formally defined as the overlap of p-orbital electrons across σ bonds [[38](#),[39](#)] such as what occurs in benzene or butadiene. Hyperconjugation is the interactions of electrons in a donating bonding orbital to an anti-bonding orbital [[40](#)]. There are several modes of hyperconjugation such as $\sigma \rightarrow \sigma^*$, $\sigma \rightarrow \pi^*$, and $\pi \rightarrow \sigma^*$. In this study, for simplicity, we use the term conjugation to refer to all modes of conjugation and hyperconjugation.

3. Results

3.1 Torsion energy barriers are sensitive to the chemical environment, which can be influenced by remote substituents

In most forcefields, torsions are defined by the quartet of atom types involved in the dihedral [[41](#),[7](#),[8](#),[9](#)]. However, the quartet of atom types do not always capture the relevant chemistry, especially when the effects are non local i.e., atoms contributing to hyperconjugation, delocalization or other non classical effects, are not part of the quartet involved in the torsion [[3](#)]. Figure [2](#), A illustrates such a case with a series of biphenyls in different protonation states. While the MM torsion profiles are all the same ([2](#) D), the QC torsion profiles are different for each protonation state ([2](#) E). The torsion energy barrier increases relative to the neutral state for the cation, anion and zwitterion, in that order. The profile changes qualitatively as well. For the neutral molecule, the lowest energy conformer is slightly out of plane, at 150° and 120° . For the zwitterion, the lowest energy conformer is at 180° . In the neutral molecule, the slightly out of plane conformer is preferred to accommodate the hydrogens. However, the central bond in the zwitterion is part of the larger conjugated system between the two aromatic rings ([2](#) B) so the planar conformer is preferred to enable conjugation.

This trend poses several problems to automatic forcefield parametrization. Most forcefields consider the central bond in the zwitterion rotatable which is reflected in the same torsion parameters for all four molecules ([2](#), D) while the QC scan clearly shows that it is not. This illustrates one of the fundamental limits of atom types in classical forcefields. At what point in this series should a new

atom type be introduced? The Open Force Field Initiative's effort on automating data driven direct chemical perception [1,42,43] addresses this problem by using SMIRKS patterns to assign parameters, and providing a framework to sample over SMIRKS space in a data driven way. In addition, this example illustrates why fragmenting molecules appropriately for QC torsion scans requires human expertise and is difficult to automate. In this case, a small change three bonds away from the torsion central bond changed the bond from a rotatable bond to a non-rotatable conjugated bond. When fragmenting molecules, we need to avoid destroying a bond's chemical environment by naively removing an important remote substituent.

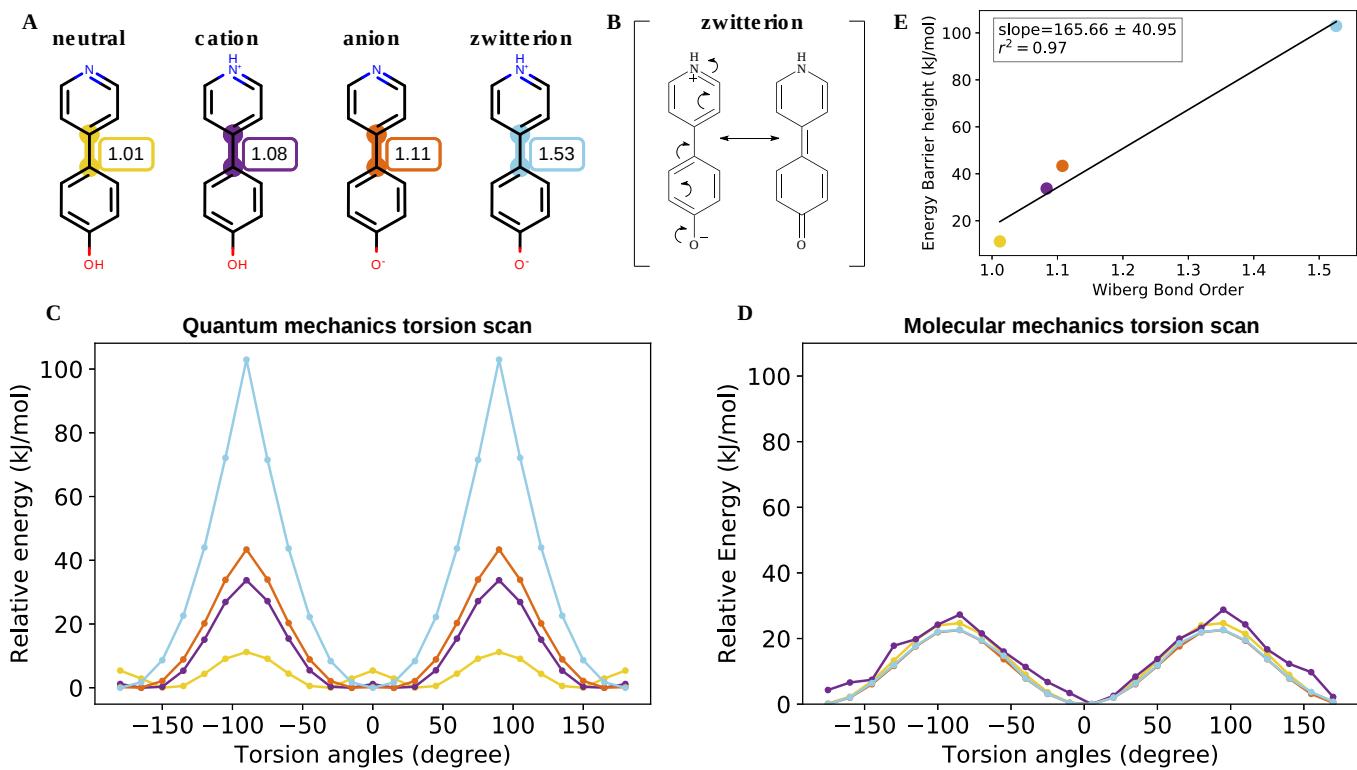


Figure 2: Torsion profiles can be sensitive to remote substituents changes in a molecule **[A]** Biphenyl protonation states and tautomers with increasing Wiberg bond order for the central bond. **[B]** The resonance structure of the biphenyl zwitterion shows that the central bond is conjugated. The Wiberg bond order and torsion scan for this bond (see **A** and **C**) are reflective of a conjugated bond. **[C]** Relative QC energy as a function of torsion angle of the central bond computed via QCArchive at B3LYP-D3(BJ) / DZVP level of theory. The colors of the QC scan corresponds to the highlighted bonds in **A**. **[D]** Same as **C** but using MM energy computed via the openff-1.0.0 force field. **[E]** Torsion barrier heights vs WBOs. The color of the data points correspond to the highlighted bonds in **A**. The QC torsion barrier height scales linearly with the WBO.

3.2 The Wiberg Bond Order quantifies the electronic population overlap between two atoms and captures bond conjugation

The Wiberg bond order (WBO) is a bond property that is calculated using orthonormalized atomic orbitals that are used as basis sets in semi-empirical QC methods[44]. Wiberg originally formulated it for the CNDO basis set [30] but it can be easily extended to other semi-empirical QC methods such as AM1 [45] and PM3 [46]. The WBO is a measure of electron density between two atoms in a bond and is given by the quadratic sum of the density matrix elements over occupied atomic orbitals on atoms A and B

$$W_{AB} = \sum_{\mu \in A} \sum_{\nu \in B} P_{\mu\nu}^2$$

The Wiberg bond order assumes atomic orbitals (AOs) are orthogonal. In ab initio calculations, however, AOs are not orthogonal but the WBO can be calculated via Löwdin normalization [47,48]

which is how it is calculated in Psi4 [49].

We calculated the WBO from AM1 calculations for the biphenyl series as shown in figure 2 A. The increase in the WBO corresponds to increasing conjugation and torsion energy barrier height of the bond. When the torsion energy barrier heights are plotted against the WBO (fig 2 E), the relationship is linear with an r^2 of 0.97.

3.3 The WBO is an inexpensive surrogate for the chemical environment around a bond

Since the WBO can be calculated from a cheap AM1 calculation, is indicative of a bond's conjugation, and is correlated with torsion energy barrier height, it is an attractive measure to use as a surrogate when automating fragmentation or interpolating torsion force constants. However, WBOs are conformational dependent [50,51] so we investigated this dependence to understand if WBOs will be a robust descriptor. In addition, we also investigated the generality of the torsion energy barrier and WBO linear relationship. In this section we will first discuss our findings and solution to the conformational dependency and then discuss how general the linear relationship is.

3.3.1 Conformation dependent variance of WBOs are higher for conjugated bonds

Since WBOs are a function of the electronic density, which is conformational dependent, WBOs change with conformation. However, not all bonds' WBOs change the same way with conformation. We found that WBOs for conjugated bonds have higher variance with respect to conformation and that bonds involved in conjugated systems have WBOs that are correlated with each other.

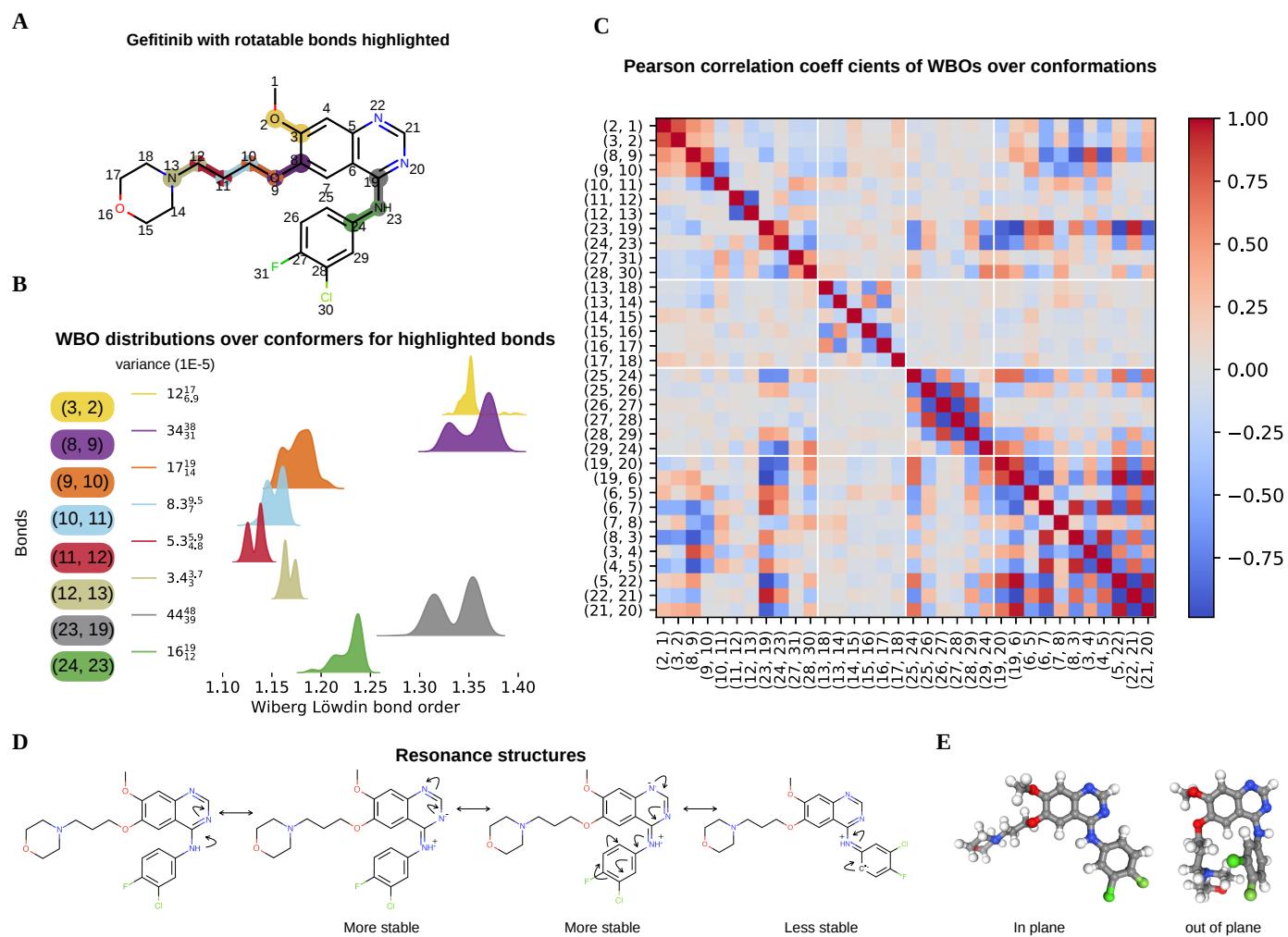


Figure 3: Variance and correlations of Wiberg bond order distributions with respect to conformations are higher for conjugated bonds [A] Gefitinib, with its rotatable bonds highlighted and numbered to correspond with figures B and C. **[B]** WBO distributions over 232 conformations of the highlighted, rotatable bonds. The colors of the distributions correspond to the colors in the highlighted bonds in A. The variance and their 95% confidence interval are shown on the left (with exponent base of 1E-5). The single non conjugated bonds (blue, (10, 11), red (11, 12), and olive (12, 13)) have lower variance than conjugated bonds (yellow (3, 2), purple (8, 9), orange (9, 10), grey (23, 19), and green(24, 24)). **[C]** Correlation plot of WBOs every bond in Gefitinib against WBOs of all other bonds over 232 conformations. The white lines indicate ring systems. Bonds in conjugated systems have higher correlations between their WBOs (see the aromatic ring systems in the two lower right diagonal squares). Both bonds (23, 19) (grey) and (24, 23) (green) have WBOs that are correlated with their neighboring ring systems, but bond (23, 19) are more correlated with the ring systems than the green bond (24, 23). **[D]** Resonance structures of Gefitinib illustrates why the grey bond (23, 19) has higher variance than the green bond (24, 23) even if both bonds are conjugated. When the double bond is on bond (23, 19), the negative charge is on a nitrogen which is the more stable form, vs the resonance structure where the double bond is on (24, 23) with the negative charge on a carbon. **[E]** The conformations of the molecule for the highest WBO and lowest in the distribution. The mode with higher WBOs has bond (23, 19) in plane with quinazoline which allows for stronger conjugation while the mode with lower WBOs has the bond out of plane so there is less electron population overlap in out of plane conformation.

To investigate how WBOs change with conformation, we used Omega [52] to generate conformers for a set of kinase inhibitors [hold for SI] and calculated the Lowdin-Wiberg bond order for each conformation from a B3LYP-D3(BJ) / DZVP [14,15,53,54] geometry optimized calculation using Psi4 [49]. Omega is a knowledge-based conformer generator that uses a modified version of MMFF94s [55] to score conformations. It has been shown to accurately reproduce experimentally observed crystallography conformers in the Platinum benchmark dataset [56]. Figure 3 illustrates the results for Gefitinib (3, A), a representative molecule. Figure 3, B shows the distribution of WBOs for all rotatable bonds color coded with the colors used to highlight the bonds in Gefitinib (3, A). Single carbon-carbon bonds, and carbon-nitrogen bonds formed by atoms numbered 10 - 13 are freely rotating. This is reflected by the tighter distribution (lower variance) of WBOs around closer to one for those bonds. The bonds involving the ether oxygens and aromatic rings (formed by atoms numbered 1-3, 8-10, 19, 23-24) exhibit higher variance. It is interesting to note the difference in the WBOs for the conjugated bonds formed by the nitrogen between the quinazoline and chloro fluro phenyl (bonds formed by atoms numbered 19, 23 and 23, 24). Both of these bonds are conjugated with their neighboring ring systems, however, While the distribution of WBOs for bond 23-19 (the grey distribution) has two modes clear modes of almost equal weights, the WBO distribution for bond 24-23 has lower variance. This is in agreement with the resonance structures shown in figure 3. The resonance strictures that have the double bond on the bond closer to the quinazoline (bond 19-23) are more stable because the negative charge is on a nitrogen. When the double bond is on the neighboring 23-24 bond, the negative charge is on an aromatic carbon which is less stable. The results are similar for other kinase inhibitors tested shown in [hold for SI]. In addition, when we inspected the conformations associated with the highest and lowest WBO in the grey distribution (fig 3, E) we found that conformations with lowest WBO on bond 19-23 had that bond out of plane while the conformation with the highest WBO value had the bond in plane which allows conjugation. We found similar results from WBOs calculated from QC torsion scans. Fig Z shows Lowdin-Wiberg bond order for each point in the QC corresponding torsion scans. The WBOs are anti-correlated with the torsion potential energy which is in line with chemical intuition. Conjugation stabilizes conformations and leads to more electronic population overlap in bonds [32]. At higher energy conformers, the aromatic rings are out of plan and cannot conjugate. Therefore the WBO is lower for those conformers. At lower energy conformations, the rings are in plane and can conjugate so the WBO is higher. We found that the trends discussed above are similar when using semi-empirical methods such as AM1 and HF3C (hold for SI)

3.3.2 Bonds in conjugated systems have highly correlated conformation-dependent WBOs

We found that certain bond orders are strongly correlated or anti-correlated with each other, indicating strong electronic coupling. As bonds in one conformation gain electron population overlap,

the coupled bonds will decrease in electron population overlap and vice versa. Figure 3 C shows the Pearson correlation coefficient for each bond WBO distribution against all other bond WBO distributions. There is a clear structure in this correlation plot. The square formed by bonds from atoms 24-29 shows that the alternating bonds in the aromatic ring (25-29) are strongly anti-correlated with each other. This trend is different in the ring formed by atoms 13-18 which is not aromatic. In this ring, bonds 13-18, 13-14, 16-15 and 16-17 (which involve electron rich atoms O and N) have Pearson correlation coefficients with absolute values higher than for the other bonds in the ring, but lower than the bonds in the aromatic ring. The bonds involved in the methoxy groups (atoms 1-3 and 8-10) are correlated with each other and also correlated to the quinazoline, albeit not as strongly. And the bonds between the chloro fluoro phenyl and quinazoline follow the same trend as their WBO distribution and resonance structures. The bond closer to the quinazoline (bond 23-19) has WBO distribution correlated with the quinazoline while the bond closer to the chloro fluoro phenyl (bond 23-24) is not as strongly coupled with the quinazoline. The trends are similar for other kinase inhibitors tested as shown in [\(hold for SI\)](#).

3.3.3 ELF10 provides a useful way to capture informative conformation-independent WBOs

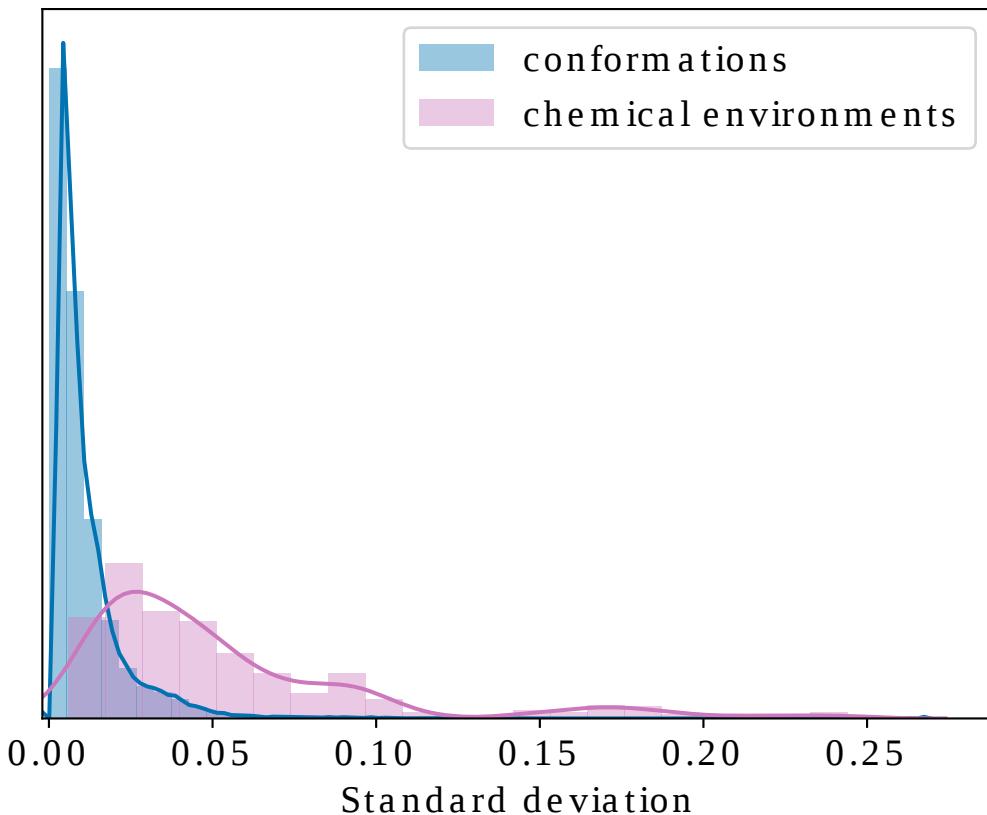


Figure 4: Standard deviations of conformation dependent WBO distributions are smaller than standard deviations of WBO distribution of the same bond in different chemical environments The distribution of standard deviations of WBO distributions over conformations is shown in blue. The distribution of standard deviations of ELF10 WBO distributions for the same bond in different chemical environments is shown in pink. The changes in WBO due to conformations are smaller than the changes in WBO due to chemical changes around the bond. The validation set (SI 10), was used to generate these distributions. The blue std distribution was calculated over 140602 WBO dependent distributions (this is the number of individual fragments in the dataset). The pink std distribution was calculated over 366 distributions of 366 bonds in different chemical environments (of the 140602 fragments)

As we have shown, the WBO is conformation dependent and this dependency can also be highly informative of the electronic coupling in a system. Figure 4 shows the distribution of standard deviations of WBO distributions with respect to conformation in blue. Most of the standard deviations fall below 0.03, which is encouragingly small. However, it can become computationally expensive to calculate the WBO for all conformations. If we want to use WBOs as a surrogate to determine if our

fragment is representative of the parent molecule in a reproducible way, we need a way to capture informative conformation-independent WBOs. Electronically Least-interacting Functional groups (ELF) conformation selection implemented in quacpac [57] resolves the issue of sensitivity of molecular mechanics electrostatic energies from QM derived charges.

Leave to Christopher Bayly to describe

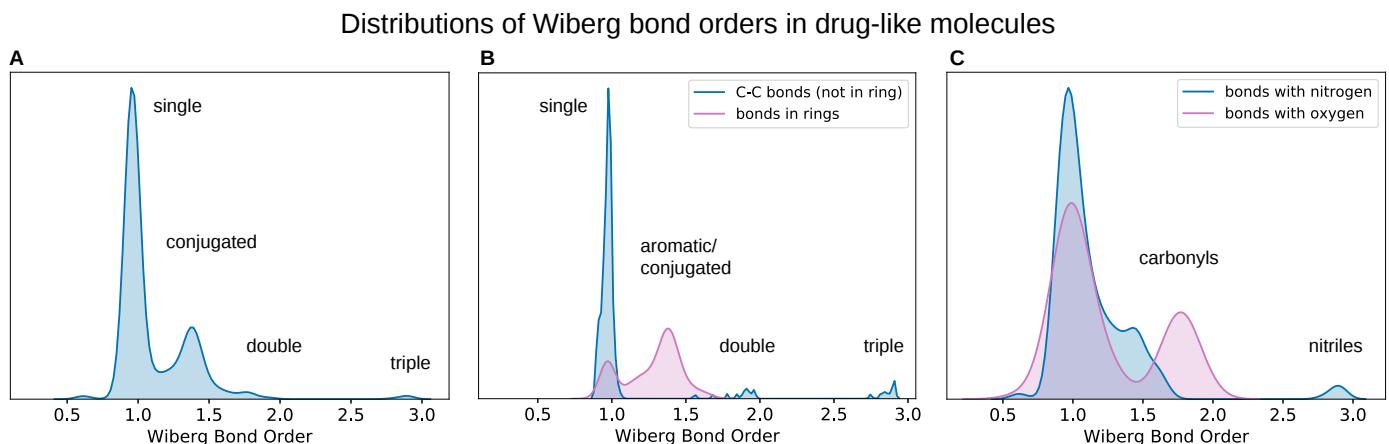


Figure 5: Distribution of WBO in drug-like molecules have peaks at expected values **[A]** The distribution of all WBOs for all bonds in molecules in set. The peak at one, close to two and close to thres correspond to single, double and triple bonds. the density between one and two correspond to aromatic and conjugated bonds. The peak at ~0.7 correspond to bonds that include sulfur and phosphorous which are longer, weaker bonds. **[B]** The blue distribution includes carbon - carbon bonds that are not in rings. The peaks at one, two and three correspond to single, double and triple bonds. The pink distribution include bonds that are in rings. The peak at one corresponds to single bonds and the density between one and 1.5 are aromatics. **[C]** The blue distribution includes bonds that have either one or two nitrogen. Many of these bonds are conjugated as demonstrated by the density around 1.5. The density at three corresponds to nitriles. The pink distribution include bonds that have oxygens. The peak at two corresponds to carbonyls.

This method can also be applied to derive WBOs that are insensitive to conformers. To check how well ELF10 estimated W_{AB} recapitulates the multiplicity of bonds, we calculated W_{AB} from ELF10 AM1 calculations for all bonds in a set of molecules shown in SI Figure 19. The distribution in figure 5 corresponds closely with bond multiplicity. The density at ~ 0.7 correspond bonds involving sulfur and phosphorous since these are weaker and longer bonds. The peak at 1.0 corresponds to C-H and C-C bonds, the peak close to 1.5 corresponds to bonds in aromatic rings, the peak close to 2.0 corresponds to double bonds, and finally the triple bonds form the last peak. Figures 5 B and D separate out different kinds of bonds to more clearly illustrate what the WBO captures. Figure 5 B shows carbon - carbon bonds not in rings (blue) and bonds in rings (pink). The carbon - carbon distribution has distinct peaks at one, two and three corresponding to single, double and triple bonds. There is also a smaller peak at 1.5 that corresponds to conjugated bonds. The pink distribution includes bonds in rings and has a peak at one and 1.5 which corresponds to aliphatic and aromatic rings, respectively. Figure 5 D shows distributions with bonds that have nitrogens (blue) and oxygens (pink). The peaks occur at the expected values; 1, 1.5 and 3 for nitrogen which corresponds to single, conjugated and triple bonds and 1 and 2 for oxygens which correspond to single and carbonyl bonds. For the rest of this section we will be focusing on the robustness and generalizability of ELF10 WBOs.

3.3.4 WBOs are a robust signal to how remote substituent changes alter a bond's torsion barrier height

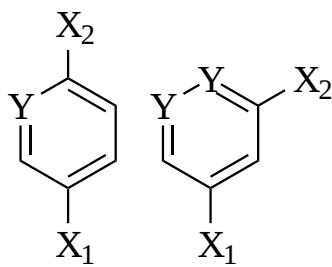
To investigate how resonance and electronic effects from remote substituents change the torsion energy of a bond, we took inspiration from the Hammett equation [58] of reactions involving benzoic

acid derivatives. The Hammett equation relates meta and para benzoic acid substituents to the acid's ionization equilibrium constants

$$\log \frac{K}{K_0} = \sigma \rho$$

Where σ is a substituent constant and ρ is a reaction constant. It aims to isolate the resonance and inductive effects of substituents from the sterics effects of a reaction. Here, we generated a combinatorial set of meta and para substituted phenyls and pyridine [6 A](#) with 25 functional groups that cover a wide range of electron donating and withdrawing groups. We then calculated the ELF10 WBO for the bond attaching the functional group to the aromatic ring for all functional groups which resulted in 128 ($25 * 5 + 3$) data points for each functional group. This allowed us to isolate the effect on a bond's WBO from remote chemical environment changes, defined as a change more than two bonds away, from other effects such as sterics and conformations. The resulting distributions are in figure [6, B](#). It is interesting to note that the trend of decreasing WBOs for more electron donating groups are anti correlates with increasing Hammett substituent constants. In [6 C](#) and [D](#), the AM1 ELF10 WBO of the bond between the functional group and benzoic acid is plotted against their Hammett para and meta substituent constants. Functional groups that are more electron donating will have more electron density on the bond attaching the functional group to the benzoic acid. The resonance and/or inductive effect destabilize the benzoate, increases its pKa, which corresponds to lower substituent constants.

A

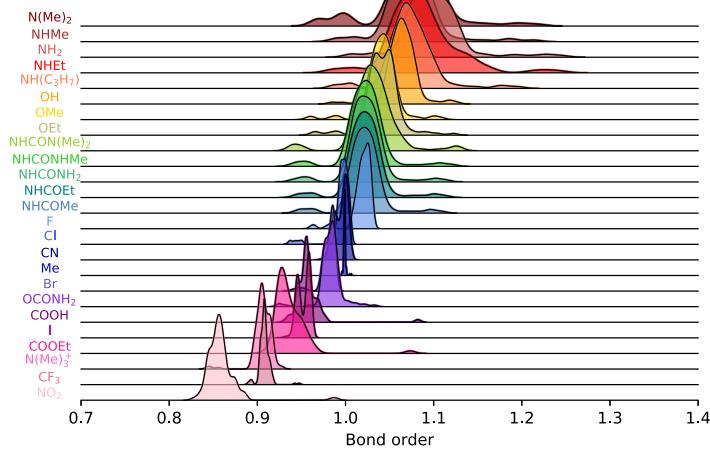


$X_1 = [N(Me)_2, NHMe, NH_2, NHEt, NH(C_3H_7), OH, OMe, OEt, NHCON(Me)_2, NHCONHMe, NHCONH_2, NHCOEt, NHCOMe, F, Cl, Br, I, CN, Me, OCONH_2, COOH, COOEt, N(Me)_3^+, NO_2]$

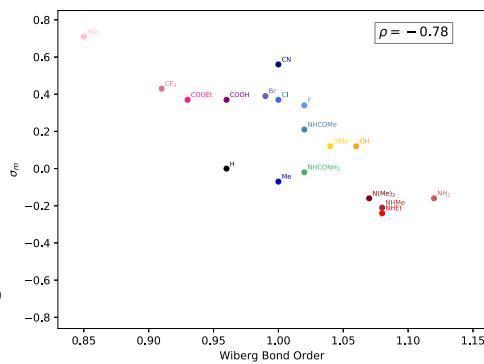
$X_2 = [X_1, H]$

$Y = [C, N]$

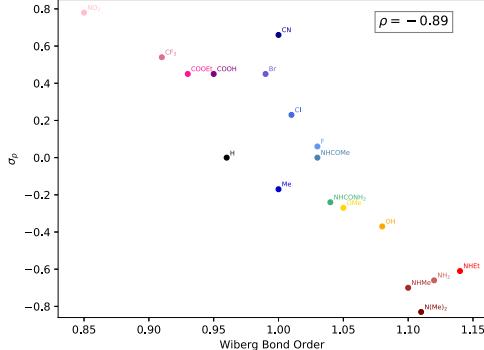
B



C



D



E

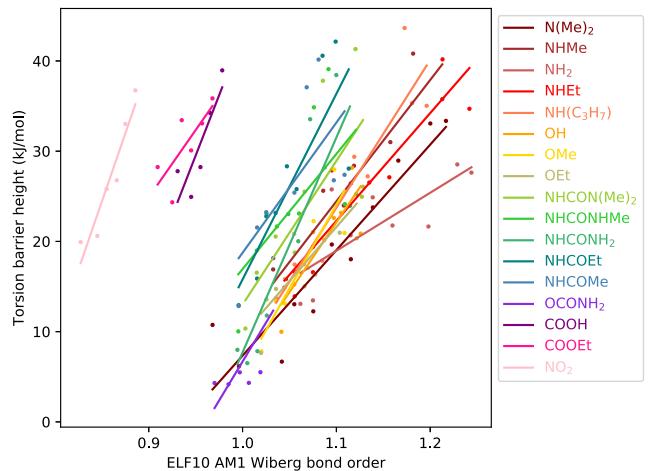


Figure 6: Change in AM1 ELF10 WBOs correlates with barrier heights in torsion profiles [A] Systems and functional groups used in the substituted phenyl set. The functional groups were chosen to span a large range of electron donating and withdrawing groups. [B] AM1 ELF10 WBO distributions for the bond between the phenyl ring and X_1 in different chemical environments [C] Hammett sigma para parameters vs AM1 ELF10 WBOs of X_1 para to carboxylic acid [D] Hammett sigma meta parameters vs AM1 ELF10 WBOs of X_1 meta to carboxylic acid [E] Selected QC torsion scan barrier heights vs ELF10 AM1 WBOs

Table 1: slope and associated statistics for torsion barrier height vs WBO for selected functional groups

functional group	slope	r^2	P value	standard error
N(Me) ₂	116.916250	0.880571	0.000019	14.352479
NHMe	134.517241	0.896103	0.000033	16.194060
NH ₂	64.266814	0.577819	0.017438	20.763035
NHEt	119.513117	0.836392	0.000552	19.978559
NH(C ₃ H ₇)	163.763962	0.871119	0.000236	23.808080
OH	154.824936	0.729133	0.003393	35.666975

functional group	slope		P value	standard error
OMe	185.309951	0.800852	0.006494	41.326214
OEt	119.662499	0.479470	0.038706	47.124984
NHCON(Me) ₂	159.312781	0.579507	0.010533	47.979491
NHCONHMe	127.645256	0.434585	0.053430	55.030363
NHCONH ₂	238.119990	0.734429	0.003158	54.120536
NHCOEt	205.802258	0.692740	0.005374	51.804680
NHCOMe	144.320372	0.457787	0.065376	64.121783
OCONH ₂	172.716947	0.508785	0.111518	84.854215
COOH	267.218639	0.739937	0.061417	91.463658
COOEt	149.012778	0.581298	0.077956	63.233457
NO ₂	302.072242	0.909174	0.003192	47.737899

To investigate how these long range effects observed in the WBOs capture changes in the bonds' torsion potential energy, we ran representative QC torsion scans for 17 of the functional groups (hold for SI). We did not run QC torsion scans for functional groups that either did not have a torsion such as halogens, were congested such as trimethyl ammonium and functional groups where the WBOs did not change by more than 0.01 for different functional groups at the meta or para position such as methyl. We chose the representative molecules for the 17 functional groups by sorting them by their WBO and selecting molecules with minimum WBO difference of 0.02. All of the resulting QC torsion scans are shown in [hold for SI]. We show a representative series of torsion scan for the nitro functional group in figure 7A. The torsion energy barrier height increase with increasing ELF10 WBO of the bond. In addition, 7B shows that the Wiberg-Lowdin bond orders are anti-correlated with the QC torsion scan which is the same result we saw for the initial bipheynl set discussed in the previous section. We also found that the linear relationship between WBOs and torsion energy barrier height shown in 2D generalizes to all functional groups tested in this set 6E. Table 1 lists the slopes and associated statistics for the fitted lines.

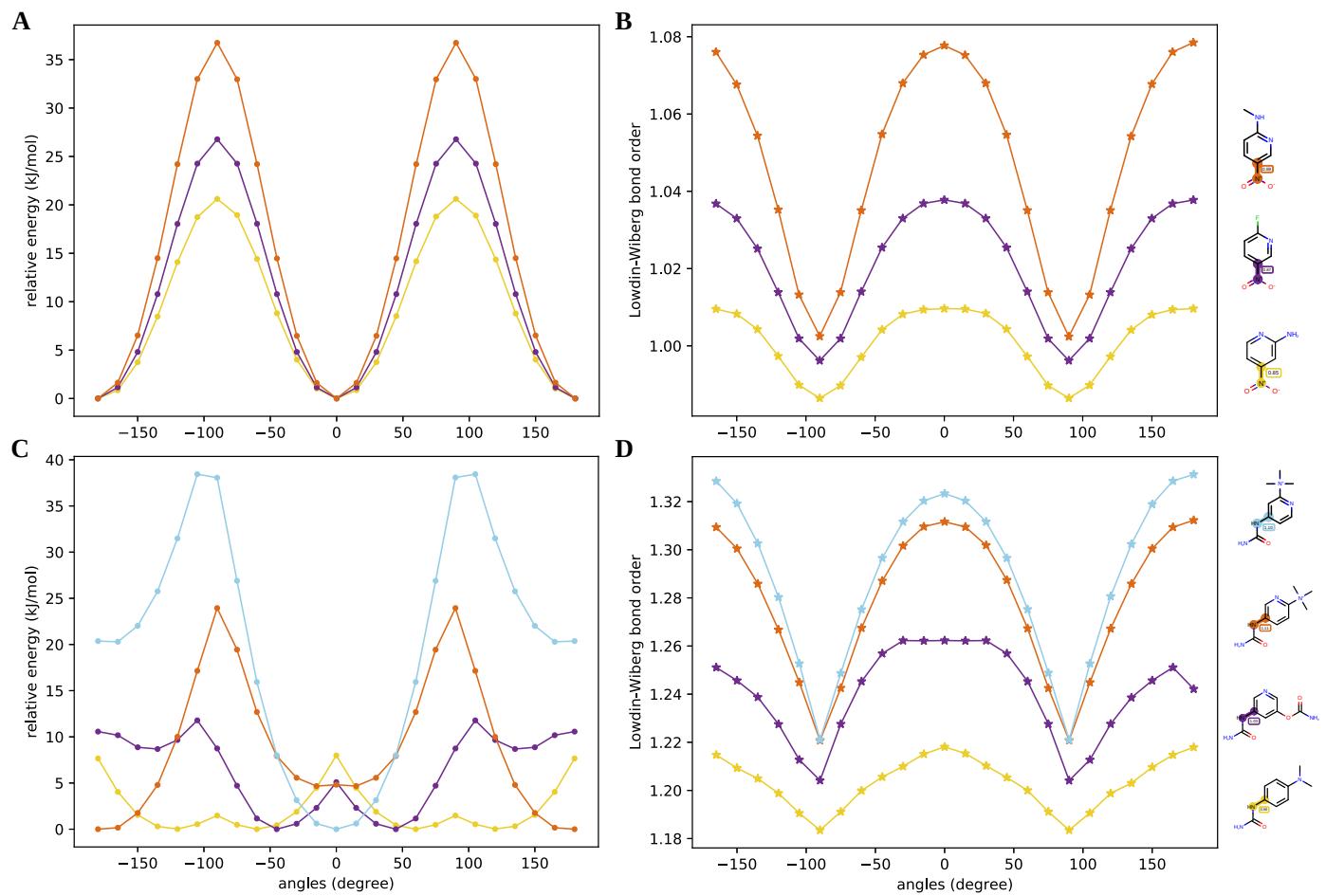


Figure 7: Löwdin-Wiberg bond orders are anti correlated with QC torsion scans [A] QC torsions for methylamino in series of different chemical environment. Barrier heights increase with increasing ELF10 AM1 WBOs [B] Löwdin-Wiberg bond orders calculated at each point in the QC torsion scan using the same level of theory. The bond orders are anti correlated with QC torsion scans [C] QC scans for urea in a series of different chemical environment. Both profiles and energy barriers change with ELF10 AM1 WBOs [D] Löwdin-Wiberg bond orders are not perfectly anti correlated to QC scans

For most functional groups, the change in WBOs correspond to changes in torsion barrier heights. [hold for SI]. However, for some functional groups, the change in WBO does not fully capture the differences in torsion scans because not only do the torsion energy barrier heights increase, but the profile changes considerably as shown in C for urea. Interestingly, the Löwdin-Wiberg bond order scans do have the same profiles D.

When we compare the standard deviations of WBO distributions with respect to conformation versus with respect to changes in chemical space (figure 4, pink distribution), we find that the changes in ELF10 WBO for remote chemical environment changes are usually bigger than the changes in WBO that arise from change in conformation. This allows us to use the difference in ELF10 WBO of parent and fragment as a good surrogate to the level of disruption of the chemical environment.

3.4 A simple fragmentation scheme can use the WBO to preserve the chemical environment around a torsion

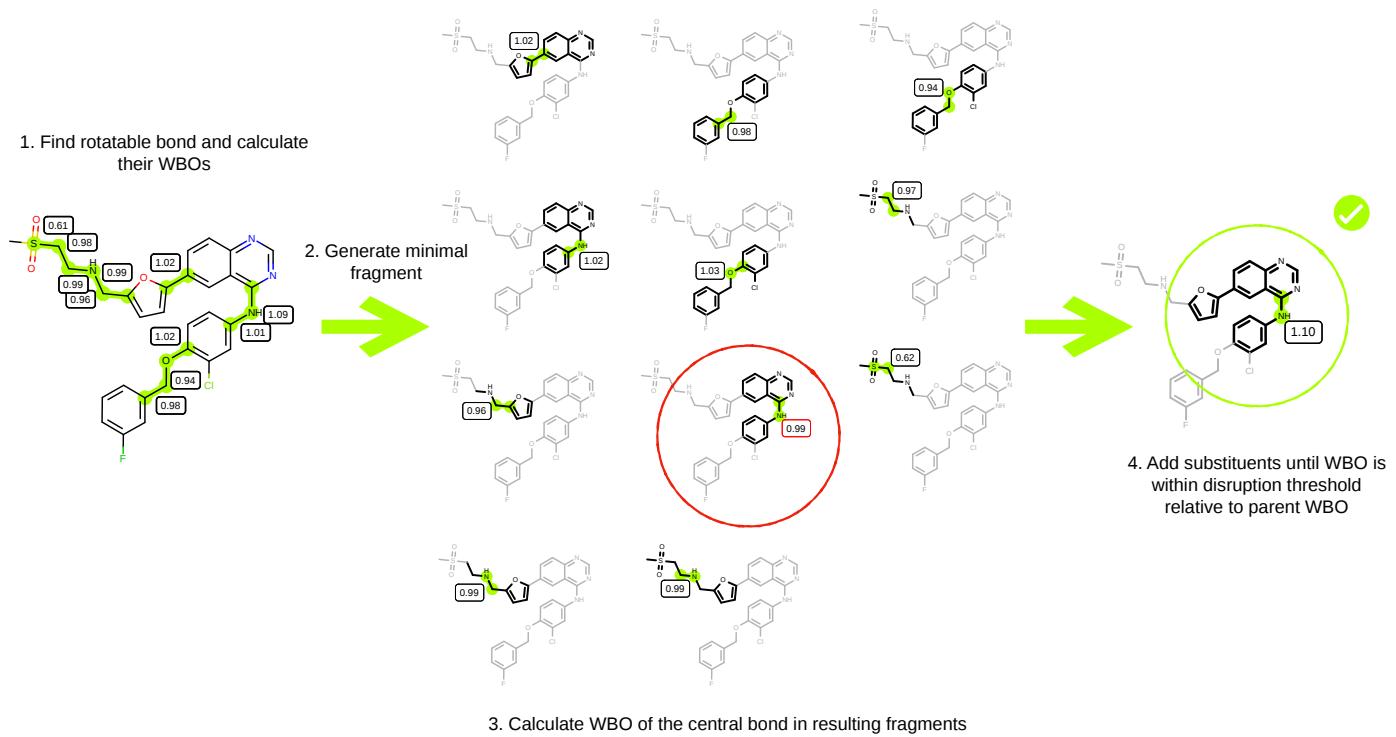


Figure 8: Illustration of fragmentation scheme using WBOs First, we find the rotatable bonds and calculate their ELF10 WBOs. Then, for each rotatable bond, we find the minimal fragment as described in the text. Then we calculate WBOs for the bonds around which the minimal fragments were generated and check if the new WBO is within the disruption threshold relative to the parent's WBO. If the absolute difference is greater than the disruption threshold, substituents are added, one at a time, and the WBO is recalculated. The fragment continues to grow until the central bond's WBO is within the disruption threshold of the WBO of the bond in the parent molecule

The WBO is a robust indicator of changes in torsion energy barrier heights for related torsions. Therefore, if a fragment's WBO changes too much from its parent WBO at the same bond, the fragmentation is probably inadequate. Using this concept, we extended the fragment-and-cap scheme (with slight changes) proposed by [29] by considering resonance via WBOs. The scheme, illustrated in figure 8 is as follows:

1. Find acyclic bond. For this step we use the SMARTS pattern `[!$(*##)&!D1]-,=; !@[$(*##)&!D1]`.
2. Keep the four atoms in the torsion quartet and all atoms bonded to those atoms (this ensures that all 1-5 atoms are included in the minimal fragment).
3. If any of the atoms are part of a ring or functional group shown in table 2, include ring and functional groups atoms to avoid ring breaking and fragmenting functional groups that contain more than one heteroatom.
4. Keep ortho substituents to the rotatable bond and the bonds directly bonded to the rotatable bond it is involved in the sterics of the torsion. This rule is slightly different than described in [29].
5. N, O and S are capped with methyl. All other open valence atoms are capped with hydrogen.
6. Calculate WBO for fragment.
7. If the fragment's WBO differs by more than a user defined threshold, continue grow out one bond at a time until the fragment's WBO is within the threshold of the parent WBO.

Table 2: Functional groups that are not fragmented. This list is not comprehensive as it only includes functional groups that were present in the validation set. Users can add their own functional groups they do not want to fragment

Chemical group	SMARTS pattern
azo	<code>[NX3] [N]=</code>
nitric oxide	<code>[N]-[O]</code>

Chemical group	SMARTS pattern
amide	[#7] [#6] (=[#8]) , [#7] [#6] (-[O-])
urea	[NX3] [CX3] (=[OX1]) [NX3]
aldehyde	[CX3H1] (=[O]) [#6]
sulfoxide	[#16X3]=[OX1] , [#16X3+] [OX1-]
sulfonyl	[#16X4] (=[OX1])=([OX1])
sulfinic acid	[#16X3] (=[OX1]) [OX2H, OX1H0-]
sulfinamide	[#16X4] (=[OX1])=([OX1]) ([NX3R0])
sulfonic acid	[#16X4] (=[OX1]) (=[OX1]) [OX2H, OX1H0-]
phosphine oxide	[PX4] (=[OX1]) ([#6]) ([#6]) ([#6])
phosphonate	P(=[OX1]) ([OX2H, OX1-]) ([OX2H, OX1-])
phosphate	[PX4] (=[OX1]) ([#8]) ([#8]) ([#8])
carboxylic acid	[CX3] (=[O]) [OX1H0-, OX2H1]
nitro	([NX3+] (=[O]) [O-]) , ([NX3] (=[O]) =O)
ester	[CX3] (=[O]) [OX2H0]

(Add discussion on the changes I made to the Pfizer scheme for our minimal fragment before we start building out and why. Add SI figure justifying it)

3.5 Fragmentation schemes can be assessed by their ability to preserve the chemical environment while minimizing fragment size

This fragmentation scheme improves upon [\[29\]](#), however, it leaves some parameters up to the user. In order to assess various thresholds and different fragmentation schemes in general, we generated a diverse set of FDA-approved drug molecules that can be a useful validation set. The goal of this set was to find molecules that are challenging to fragment. In other words, molecules that have bonds that are sensitive to remote substituent changes. To find these molecules, we first filtered DrugBank (version 5.1.3 downloaded on 2019-06-06) [\[19\]](#) with the following criteria:

1. FDA approved small molecules
2. Largest ring size has less than 14 heavy atoms
3. Smallest ring size has at least 3 heavy atoms
4. Molecule has less than 10 rotatable bonds
5. Molecule must have at least one aromatic ring
6. Molecule has only one connected component

This left us with 730 small molecules [\[hold for SI\]](#). Charged molecules exacerbates remote substituent sensitivity and many molecules are in charged states at physiological pH. To ensure that our dataset is representative of drugs at physiological pH, we used the OpenEye `EnumerateReasonableTautomers` to generate tautomers that are highly populated at pH ~7.4. This tautomer enumeration extended the set to 1289 small molecules [\[hold for SI\]](#). We then generated all possible fragments of these molecules by using a combinatorial fragmentation scheme. In this scheme, every rotatable bond is fragmented and then all possible connected fragments are generated where the smallest fragment has 4 heavy atoms and the largest fragment is the parent molecule. This scheme generated ~300,000 fragments. For each fragment, Omega was used to

generate conformers and the AM1 WBO was calculated for every bond in every conformer. This resulted in a distribution of WBOs for all bonds in all fragments. The resulting dataset is very rich where exquisitely nuanced, long distance chemical changes are detected.

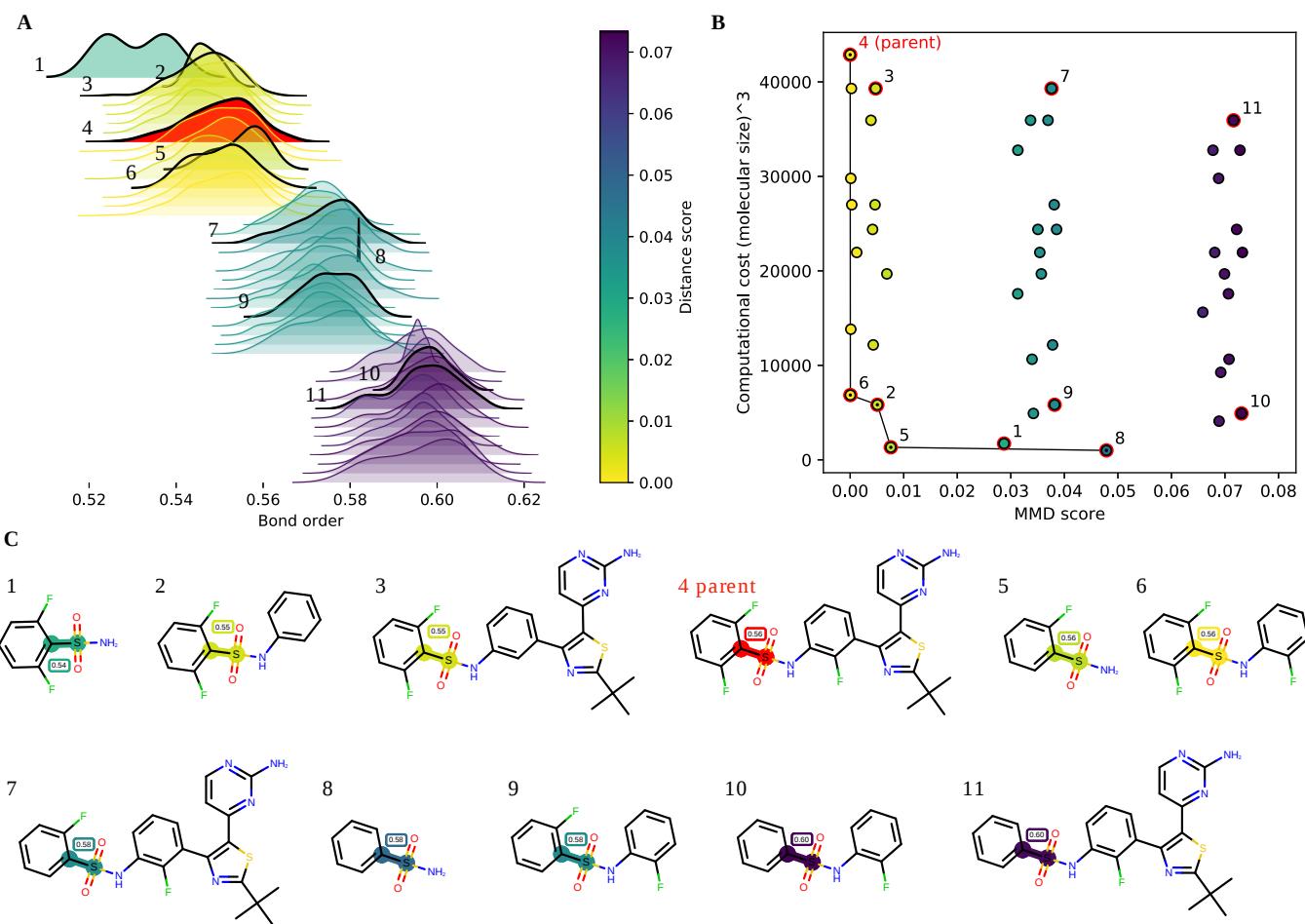


Figure 9: Changes in WBO distributions are a good indicator of remote substituent electronic effects [A] An illustrative example of the shift in the conformationally dependent WBO distributions due to crucial chemical changes such as the loss of Fluorine. The distributions are shaded with their corresponding distance score shown in the colorbar on the right. The parent molecule WBO distribution (numbered 4) is shaded red. Selected distributions are outlined and the corresponding fragments are shown in C. [B] Computational cost of fragment (heavy atoms^3) vs distance score (MMD) of the fragment indicates that it is possible to reduce the cost of torsion scans without destroying the torsion profile. The black line is the Pareto frontier, or the cheapest fragment with the best score at that size. The selected fragment should be on the Pareto front at the lower left corner. [C] Selected fragments. Bonds are highlighted with their distance score. The ELF10 WBO is shown in the boxes above the highlighted bonds.

Figure 9 shows an example of the results of exhaustive fragmentation and how this data can be used to benchmark fragmentation schemes. All rotatable bonds in the parent molecule, Dabrafenib (9C 4) were fragmented into 11 fragments (in this example, the trimethyl was not fragmented) resulting in 108 connected fragments when all connected combinations were generated. Of those 108 fragments, 44 fragments contained the bond between the sulfur in the sulfonamide and phenyl ring highlighted in fragments in 9C. When the WBOs were calculated for all Omega generated conformers for each of the 44 fragments, the resulting WBO distributions clustered into 4 distinct bins (9A). Upon closer inspection we found that the shifts of the distributions corresponded to specific remote substituent changes, in this case the loss of fluorine and the phenyl ring bonded to the nitrogen in the sulfonamide. Here, these two changes cause the distributions to shift in opposite directions. While the loss of a fluorine on the phenyl bonded to the sulfure shifts the distribution to the right, the loss of the ring bonded to the nitrogen shifts the distributions to the left illustrating that the changes are multi dimensional. Fragments 2, 3, 4, and 6 (9C) all contain two fluorine and fall in the same cluster as

the parent molecule, regardless if the rest of the molecule is included in the fragment. Fragments 7 and 9 only have one fluorine on the phenyl ring and both of their distributions are shifted to the right relative to the parent WBO distribution. Fragments 10 and 11 have no fluorine on the ring and are shifted to the right even more. Since removing the ring bonded to the nitrogen shifts the WBO distribution in the opposite direction, fragment 1, while having two fluorine, is shifted to the left of the parent distribution, fragment 5 WBO distribution overlaps with the parent WBO distribution even if it only has one fluorine, and fragment 8 is only shifted slightly to the right of the parent WBO distribution with no fluorine.

3.5.1 Scoring how well fragments preserve chemical environments using WBO distributions

Each fragment needs to be assigned a score of how well it preserves its parent chemical environment. To score each fragment, we compare the conformer dependent WBO distribution for a bond in a fragment against the WBO conformer-dependent distribution of the same bond in the parent molecule. To compare these distributions, we compute the maximum mean discrepancy [CITE] for the fragment distribution to the parent as follows:

$$MMD(P, Q) = \|\mathbb{E}_{X \sim P}[\varphi(X)] - \mathbb{E}_{Y \sim Q}[\varphi(Y)]\|_{\mathcal{H}}$$

where the feature map $\varphi : \mathcal{X} \rightarrow \mathcal{H}$ we use is squared $\varphi(x) = (x, x^2)$ and the MMD becomes:

$$MMD = \sqrt{(\mathbb{E}[X] - \mathbb{E}[Y])^2 + (\mathbb{E}[X^2] - \mathbb{E}[Y^2])^2} \quad (2)$$

where X is the parent WBO distribution and Y is the fragment WBO distribution. Including the squared mean incorporates the variance of the distribution and helps distinguish distributions both with different means and variances. It is important incorporate changes in variance given how the variance of the WBO distributions change for different chemical environments (see figure ZB and D. Change in variance corresponds to change in relative barrier heights).

In figure 9, the MMD score, which we call the distance score, is shown with the color map. The distributions in 9A are shaded with the distance score. The scores clearly differentiates the shifted distributions.

3.5.2 Good fragmentation schemes minimize both chemical environment disruption and fragment size

The goal of our fragmentation scheme is to find fragments that have a WBO distribution of the bond of interest closest the the parent while minimizing the computational cost of the fragment. We estimate the computational cost of a fragment by cubing its number of heavy atoms because DFT calculations grow by $O(n^3)$. The distance score calculated with MMD indicates how far the fragment's WBO distribution is or how much the chemical environment changed from its parent. When we plot the fragment size against this score, the points that fall on the Pareto front [CITE] are the ones where the distance score is the best for for a given fragment size or vice versa. Figure 9B shows an illustrative example of this. The fragments data points on the Pareto front have a black dot in the center. The numbers on the annotated data points correspond the the numbered fragments in 9C. Fragment 6 has is the smallest fragment with the smallest distance to the parent molecule. It has the important chemical moieties, such as all three fluorine and the ring bonded to the nitrogen. While fragments 2 and 5 are also on the Pareto front, the missing ring and fluorine increase the distance score, however, it is not clear if this difference is significant. It is interesting to note that fragment 3, which is also missing the fluorine on the ring bonded to the nitrogen, is shifted in the distance score relative to the parent by the same amount as fragment 2 from 6, even if it has all other parts of the

molecule adding credence to the fact that the small difference in the distance score does pick up on this chemical change. The trend is in the opposite direction for molecules missing a fluorine on the ring bonded to the sulfur. Fragment 9 and 10 both contain fluorine on the ring bonded to the nitrogen, but have greater distance scores than the fragments without that fluorine (data points to the bottom left of 9 and 10. Fragments not shown). Data points 7 and 11 illustrate that having larger fragments will not improve the distance score if the important remote substituents are not in the fragment. Fragment 9, while a lot smaller than fragment 7, has the same distance score because they both are missing the important fluorine. Data points 10 and 11 show the same trend for the fragments missing both fluorine. While fragments 1, 5 and 8 are all small, the loss of the ring results in larger distance scores.

In molecule 12, both the amide and ester bond are sensitive to the same circled negatively charged oxygen.

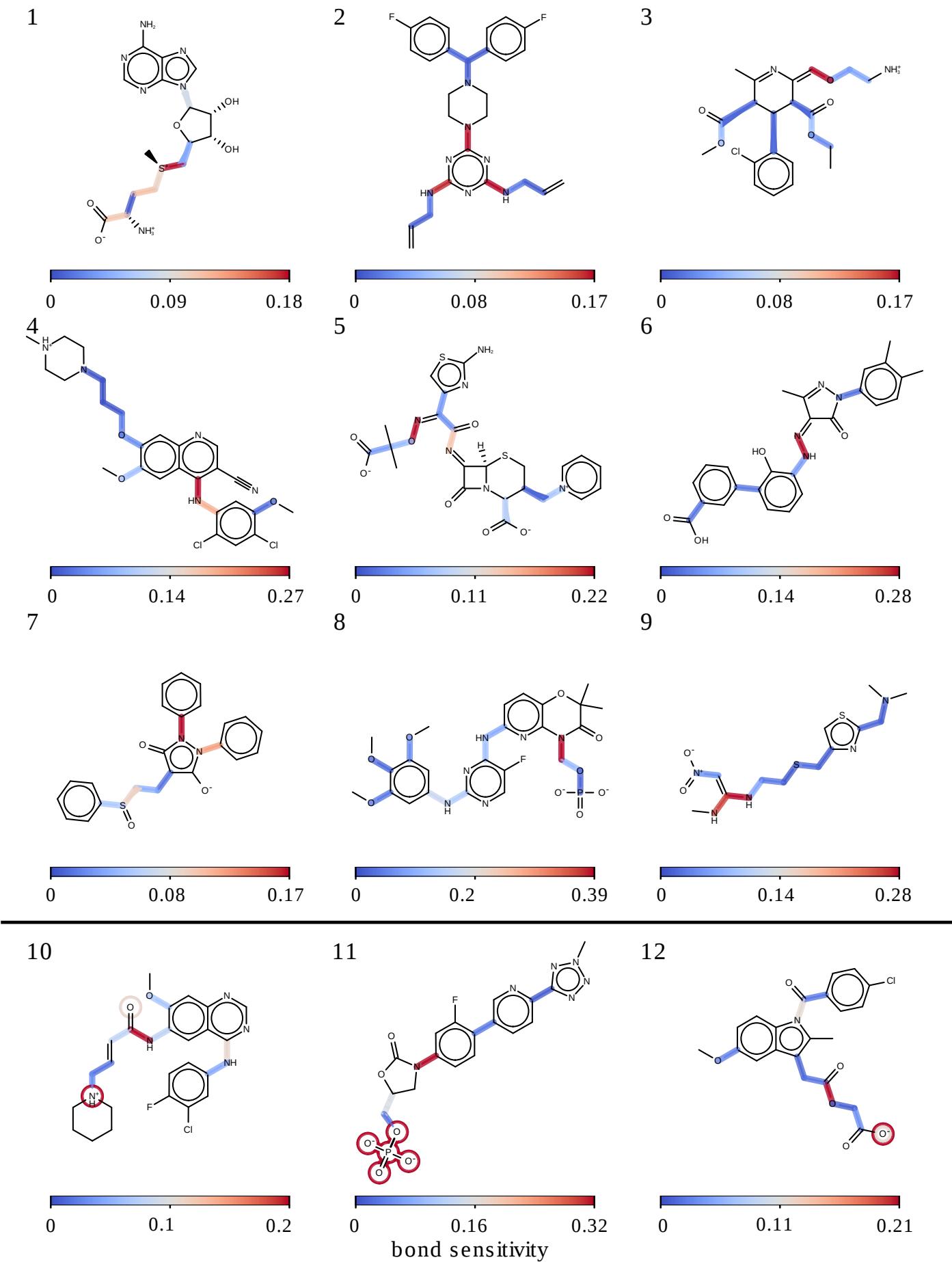


Figure 10: Some bonds are more sensitive to peripheral fragmentation and are therefore challenging to fragment

Selected molecules of the validation set. Bonds are highlighted by how sensitive they are to remote substituent changes. The bond sensitivity score is given by finding the maximum distance score of WBO distributions of the bond in fragments generated by exhaustive fragmentation, to the WBO distribution of the same bond in the parent molecule.

molecule. Fragments used to get the maximum distances include all 1-5 atoms around the highlighted bonds. Molecules 10-12 also show the atoms the bonds are sensitive to. The atoms are circled with the same color as the bond that is sensitive to it. The rest of the molecules used in the validation set are shown in SI figure [20](#)

The goal of any fragmentation scheme is to find fragments on the Pareto front that minimize both the changes in the chemical environment of the bond and fragment size. In other words, they should be on the lower left corner of the plot. To test our fragmentation scheme, we wanted to find the molecules that are challenging to fragment. To do that, we scored the WBO distributions of all resulting fragments from our exhaustive fragmentation experiment using equation [2](#) and chose 100 molecules that had bonds where fragments that included all 1-5 atoms around the central bond had the highest distance scores. Selected molecules with the bonds highlighted according to their sensitivity are shown in figure [10](#). The rest of the molecules are shown in SI figure [20](#). This set included many molecules in charged states. The sensitivity score of the bonds are given by taking the MMD of the fragment where the WBO distribution of that bond has the greatest distance relative to the WBO distribution of the bond in the parent molecule. This is a good indication of a bond's sensitivity to peripheral fragmentation because the more its WBO distribution shifts relative to the parent when fragmented, the more the electronic population overlap around that bond changes with remote chemical changes.

Not all bonds are equally sensitive to such changes. This is shown by how different the sensitivity score is for different bonds in the same molecule in figure [10](#). The general trend observed is that conjugated bonds, and bonds including atoms with lone pairs such as N, O, and S, are more sensitive to peripheral cuts. Molecules 10-12 (fig[10](#)) also show which chemical moiety the bond is sensitive to, indicated by circles around the atoms which are colored with the corresponding bond's sensitivity score. In molecule 10, the WBO distribution of the amide bond shifts significantly if the positively charged nitrogen is removed regardless if the rest of the molecule is intact (data not shown). In molecule 11, the removal of the phosphate group shifts the distribution of the red bond. In molecule 12, both the amide and ester bond are sensitive to the same negatively charged oxygen indicated by two circles around the oxygen.

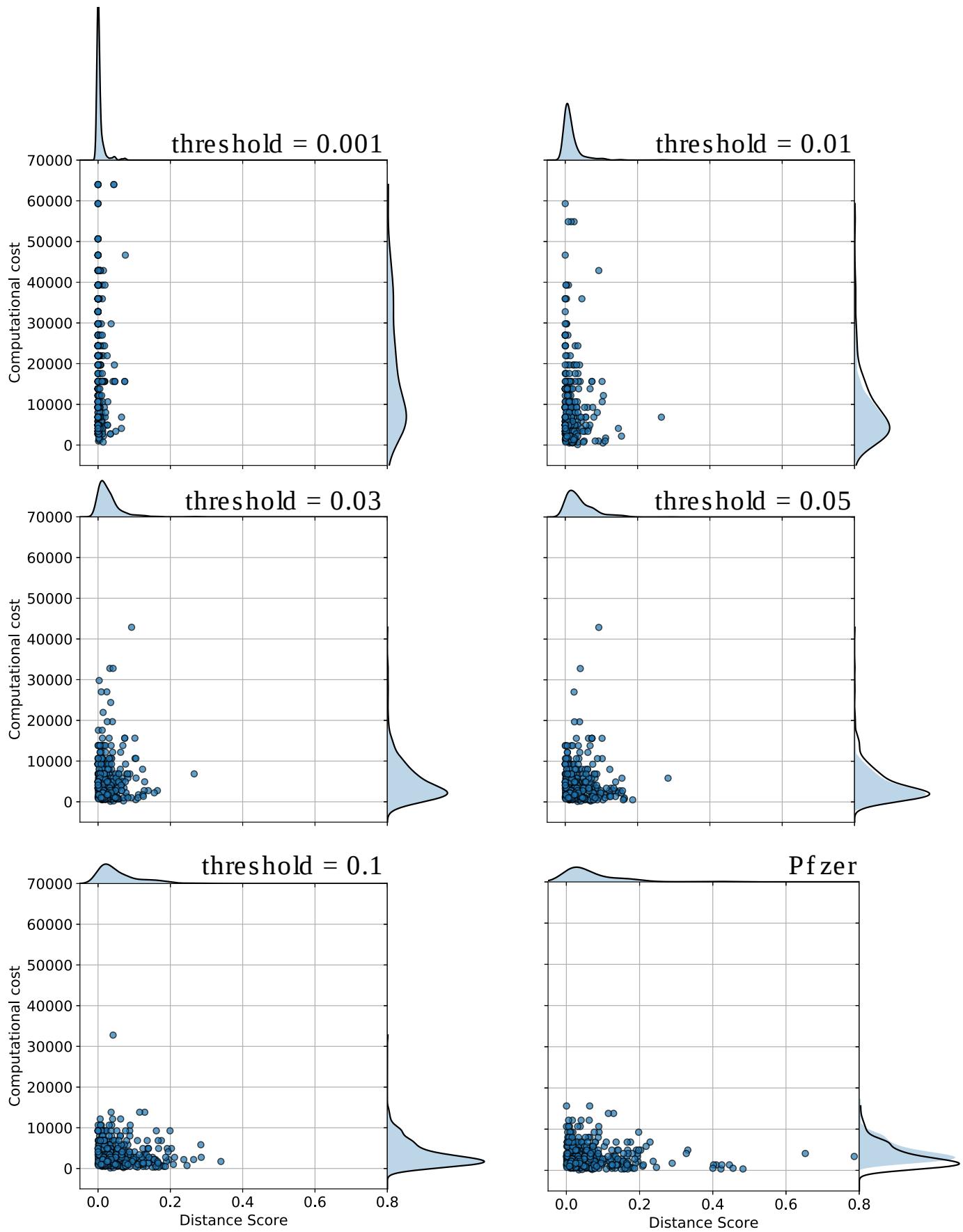


Figure 11: Assessment of fragmentation scheme illustrate an accuracy vs cost trade-off Computational cost vs distance score of the fragments for the bonds in the benchmark set shown in figure 20. Computational cost is the cubed of heavy atoms in the fragment. The threshold is the maximum amount of change allowed in the ELF10 WBO relative to the parent's ELF10 WBO during fragmentation.

We want to find the parameters for our fragmentation scheme that maximizes the number of fragments that end up in that lower left corner (illustrated in figure 9, B). To do that, we generated

fragments for the red bonds in the 100 molecules shown in figure 20 set using different disruption thresholds. For every fragment, we found the distance score of their fragments' WBO distribution and their computational cost. We then plotted all fragments from the validation set for different thresholds {figure {fig;joint_plots}}. When the threshold is low, the fragmentation scheme will generate fragments which have very good distance scores, but many of them will be too big for computational efficient QC torsion scan. On the other hand, when the disruption threshold is too low, the scheme generates fragments that are small but the distance scores are too big. For the molecules we tested, a threshold of 0.03 leads to the most fragments in the lower left quadrant (defined as cost < 10000 and score < 0.05) as shown in table 3. This threshold is similar to what we found when we looked at the distribution of standard deviations for WBO distributions with respect to conformations (4, blue). Most of them fall under 0.03. Both of these data points lead us to recommend a disruption threshold of 0.03 for our fragmentation scheme. While the current scheme does not provide a perfect solution, plots in figure 11 shows less fragments outside of the lower left region for thresholds 0.01, 0.03 and 0.05. This scheme performs better than other schemes such as the scheme Pfizer used in [29] (figure 11, lower right and table 3).

Table 3: Number of fragments in the lower left quadrant in figure 11 defined as a distance score less than 0.1 and computational cost less than 10000.

disruption threshold	fragments in lower left quadrant
0.001	153
0.005	197
0.01	229
0.03	259
0.05	231
0.07	214
0.1	200
scheme from 29	189

3.5.3 Benchmark results reveal chemical groups that induce long range effects

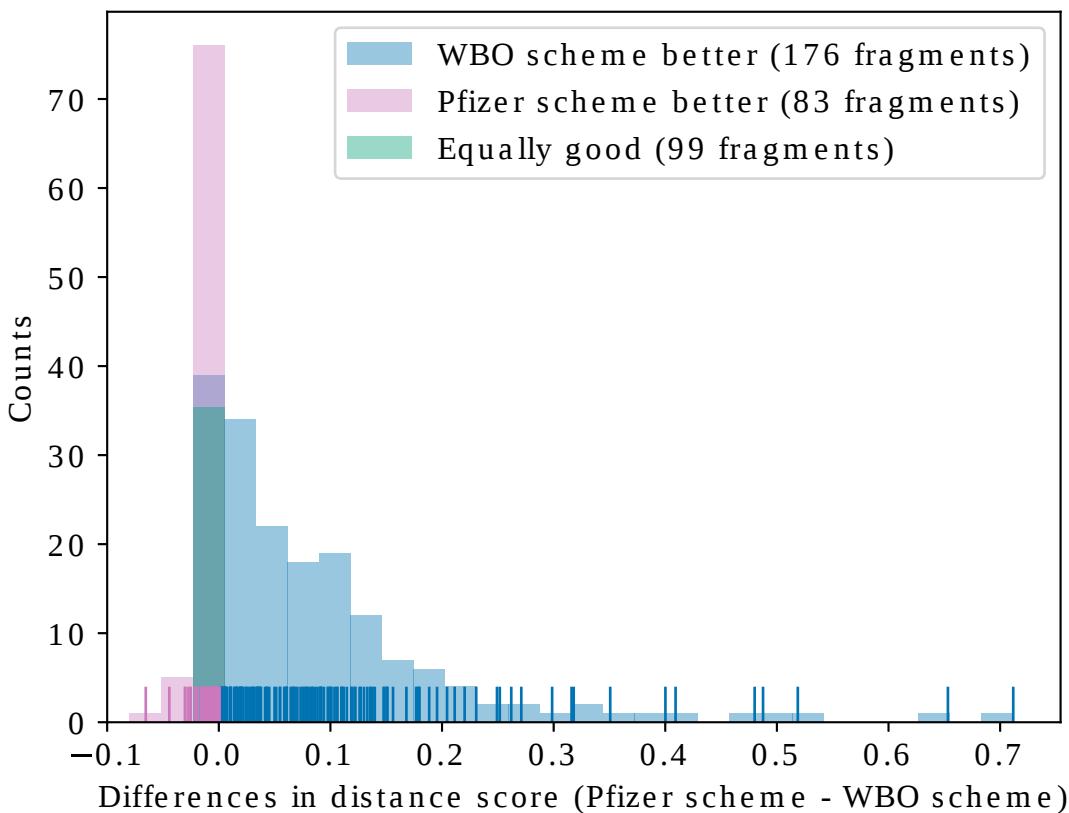
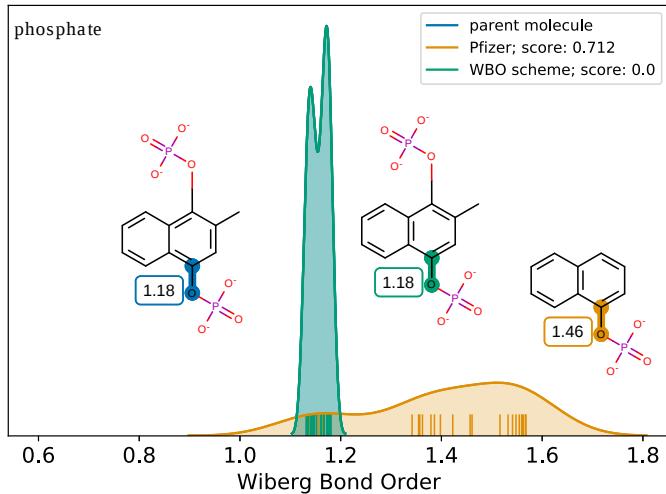
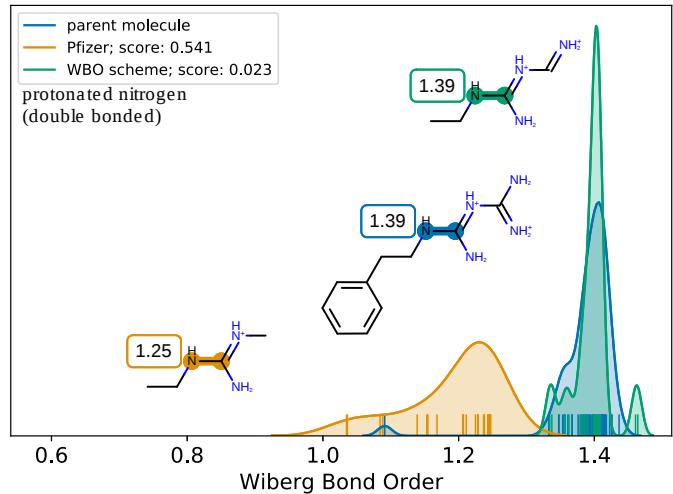
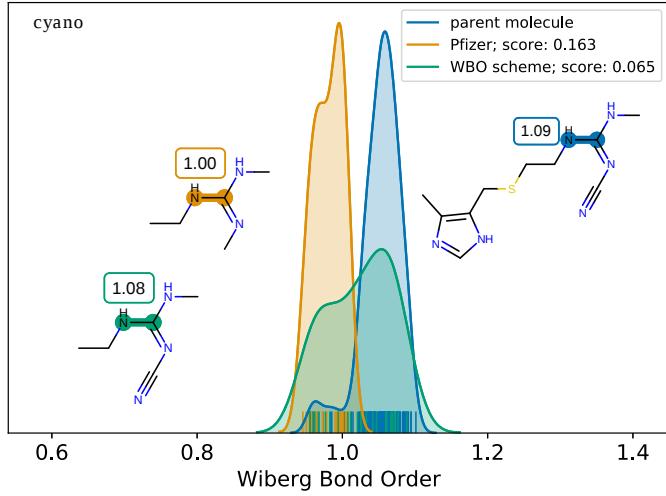
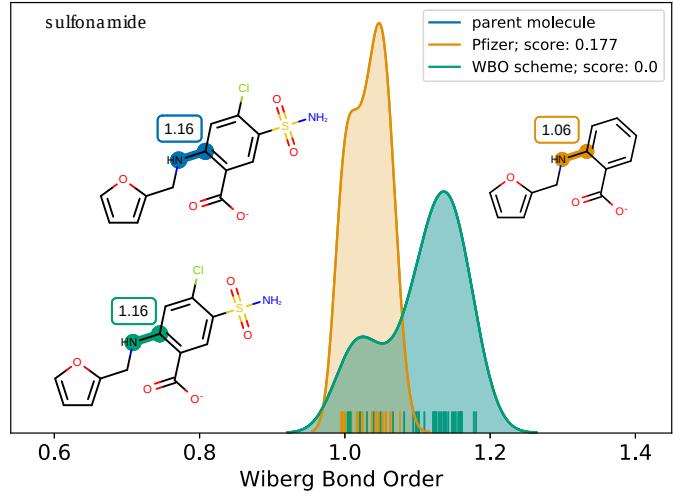
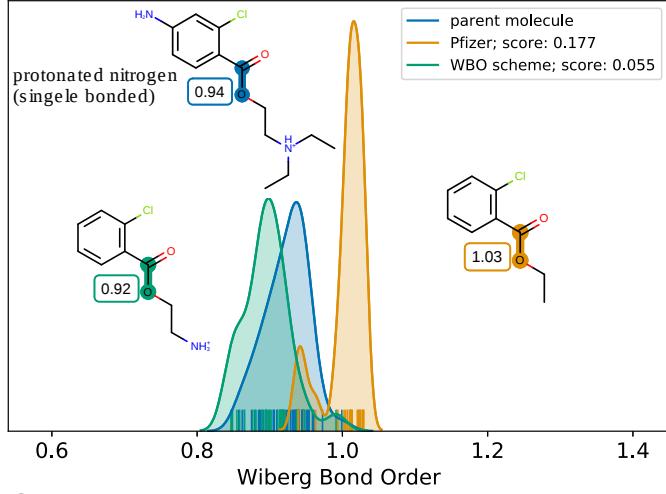
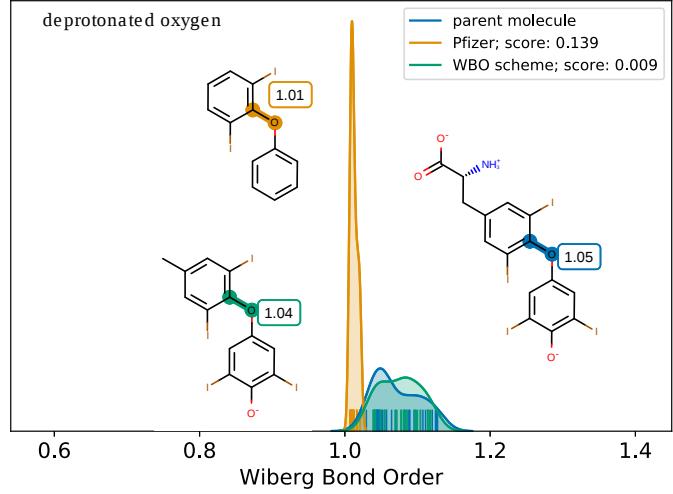
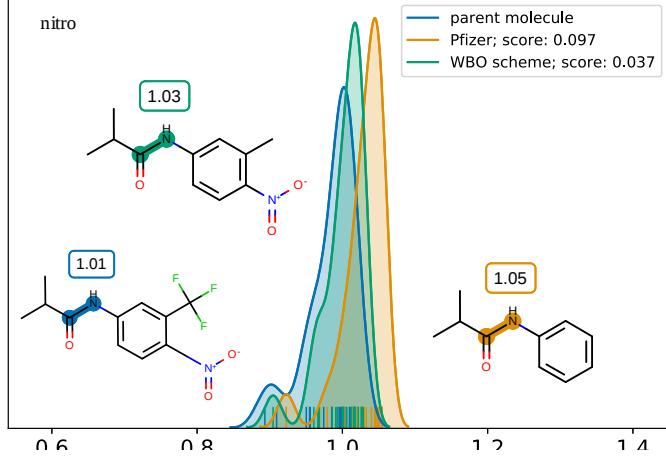
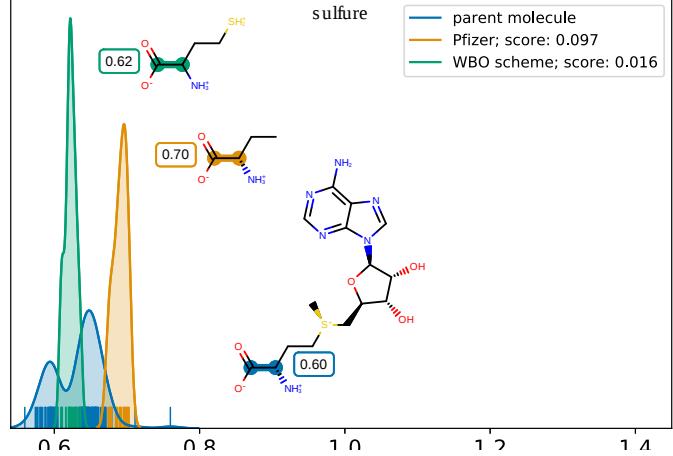


Figure 12: Using the WBO as an indicator of chemical environment disruption improves fragmentation

Distribution of differences in distance scores for fragments in the validation set (SI fig 20) generated using Pfizer's rules and our scheme using 0.03 as the disruption threshold. For many bonds, both approaches yield equally preforming fragments (shown in green). In some cases, Prizer's rules preforms better than our scheme (shown in red), however, the differences are usually very small. In most cases, using the WBO as an indicator improves the distance score (shown in blue)

In the benchmark experiment (figure 11), the distance scores measured the distance between WBO distributions generated from Omega generated conformers of the parents and fragments. Omega aims to generate low energy conformers [52] and in some cases, fragments only have one or two low energy conformers so it is not clear how accurate the distances measured are. In addition, only comparing low energy conformers do not fully capture torsion energy barriers which we also want to ensure remain accurate relative to their parent's torsion energy scan. To mitigate the above mentioned issues when validating our scheme, we also added WBOs calculated from conformers generated on a grid of torsion angles about the bonds which included higher energy conformers that are closer to conformers generated in canonical torsion scans. Furthermore, since we know that WBOs from structures in a QC torsion scan are anti correlated with the QC torsion energy scan (Z), adding these WBOs to the distributions provided a better validation of our method than only looking at the distance between omega generated WBO distributions. The differences in distances of these distributions from fragments generated with our scheme and [29] is shown in figure 12.

For many molecules, using a common sense rule based approach, such as the one used in [29] to fragmenting molecules, will yield fragments that are the same fragments generated with our scheme shown in green in figure 12, and sometimes can even preform slightly better than using the WBO as an indicator (fig 12, red). However, in many cases, especially if certain chemical groups are involved, using the WBO as an indicator significantly improves the electron population overlap about the bonds and brings them closer to their parent's chemical environment (fig 12, blue). It is important to note that when the fragment generated from both scheme are the same (green in figure 12), they are not necessarily the optimal fragment and both schemes can preform equally poorly (see SI).

A**B****C****D****E****F****G****H**

v.v

Wiberg Bond Order

Wiberg Bond Order

Figure 13: Some chemical groups induce non local effects that are captured in fragments when using the WBO as an indicator of chemical environments Wiberg bond order distributions for parent molecules (shown in blue) and fragments generated with Pfizer rules (shown in orange) and our scheme (shown in green). This figure shows eight chemical groups where the WBO distributions of the highlighted bonds change when those groups are removed. These changes are consistent across the validation set.

Upon closer inspection of the validation set, we found eight chemical groups that induce long range effects to sensitive bonds shown in figure 13. These chemical groups with representative examples are shown in figure 13. The groups are ordered by how strongly they induce long range effect, in decreasing order. The most dramatic change happens when a phosphate group is removed (figure 13, A). The variance of the WBO distribution increases which conveys an increase in relative energies of conformers in the QC torsion scans. In other molecules where phosphates are removed, the variance can decrease even if the phosphate group is ten bonds away (figure 14, F and SI). In figure 13, B, removing a protonated nitrogen that is double bonded causes the WBO distribution to shift and the variance to increase. Long range effects are seen in other molecules with similar chemical patterns up to eight bonds away (SI). Removing a nitrile group (fig 13, C) and sulfonamide group (13, D) have similar effects on the WBO distributions which is also consistent with other molecules that contain these groups up to three bonds away (SI). A protonated nitrogen and deprotonated oxygen (13 E and F) can effects bonds between 3-6 bonds away (SI). While the changes in distributions for removing a nitro group and sulfur (13, G and H) are not as big as other chemical groups, they are mostly consistent across other molecules in the validation set (SI).

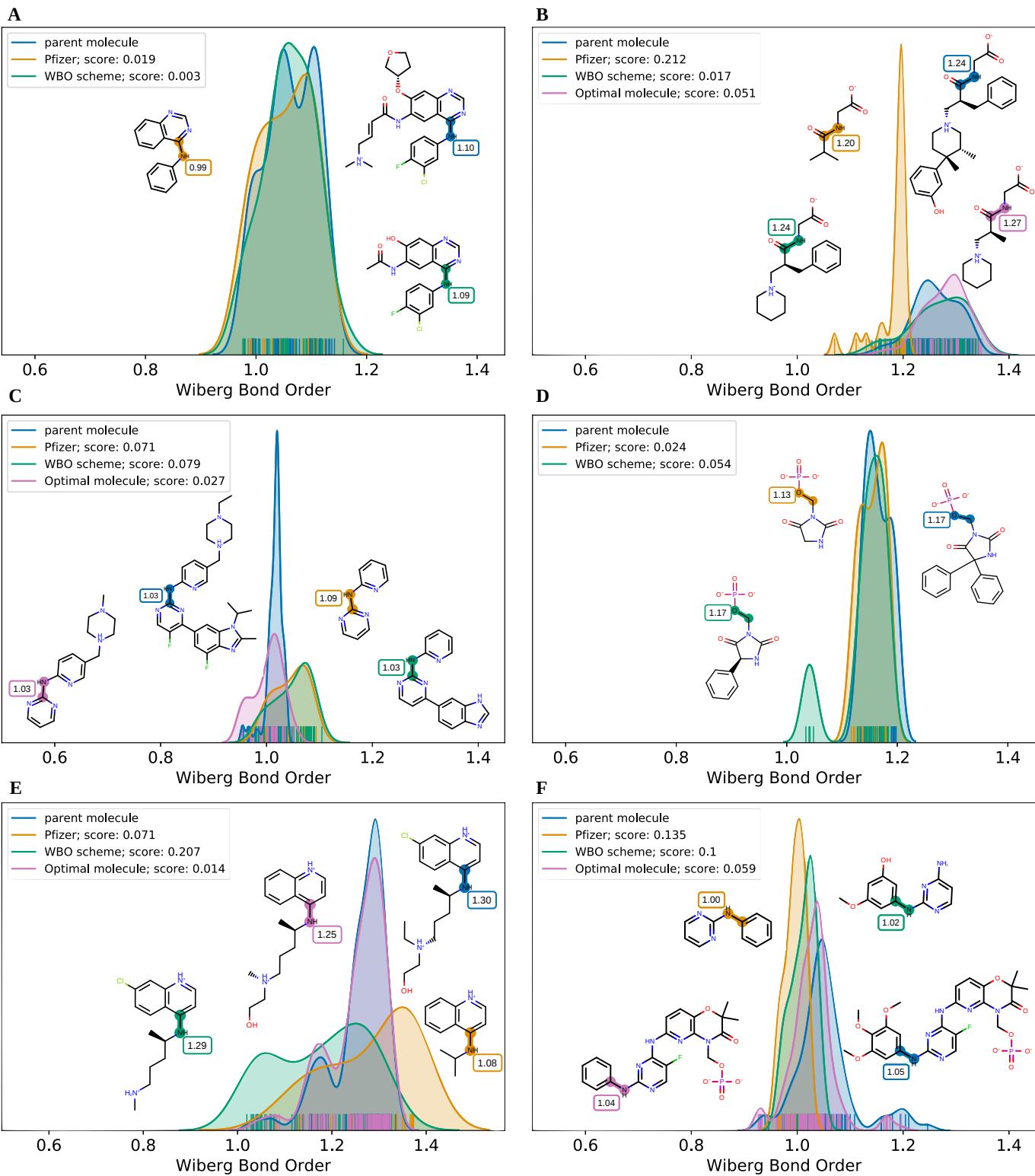


Figure 14: Using the WBO as an indicator when fragmenting can still fail to find the optimal fragment Our scheme can fail in several ways. A. A smaller fragment (shown in orange) is just as good as a larger fragment (shown in green) even if the ELF10 WBO estimate of the bond in the smaller fragment relative to its parent (shown in blue) is more than the disruption threshold. B. While our scheme finds a fragment with good overlap of the WBO distributions (shown in green), it is not the smallest fragment possible with good distributions overlap (smallest fragment with good overlap is shown in purple). C. The fragment we find is bigger than what the simple scheme finds (shown in orange) but without improving the WBO overlap (green). The optimal fragment that neither scheme generates is shown in purple. D. Our scheme finds a larger fragment that has worse WBO distribution overlap. E and F. Sometimes, almost the entire molecule is needed to achieve good WBO distribution overlap between the fragment and the parent. This is not a failure mode but inherent to the challenge of fragmenting molecules for QC calculations.

While our scheme captures long range effects that a simple rule based approach does not, it is not an optimal solution and will sometimes fail to find the most optimal fragment. By optimal we mean the

smallest fragment that retains the torsion potential of the bond in the parent molecule. Our scheme can fail in multiple ways as illustrated in figure 14 and listed below.

1. We find a fragment with good WBO distributions overlap but do not find the smallest fragment. This is shown in both 14 A and B. In A, the fragment that Pfizer scheme find is smaller and has a WBO distribution that is close to the parent's WBO distribution (MMD 0.019). In this case, the ELF10 WBO estimate of the bond in the fragment is 0.11 lower than the ELF10 WBO estimate in the parent molecule. In B, our fragment has better WBO distribution overlap with the parent WBO distribution vs using Pfizer's scheme (0.017 vs 0.212), but it is not the smallest fragment. According to the fragment highlighted in purple, the benzene ring is not required to achieve good overlap of the WBO distributions (0.051)
2. We find a fragment that is bigger than Pfizer's scheme fragment and the remote substituents do not improve the WBO distribution overlap (MMD 0.079 vs 0.071) (14 C). The better fragment is shown in purple. It is both smaller and has better overlap (MMD 0.027) than the orange and green fragment.
3. We find a fragment that is both larger and has worse overlap (0.054 vs 0.024) than what the Pfizer's scheme generates (fig 14)

While it is usually possible to find a fragment that is significantly smaller than the parent and retains remote substituent effects, the effects are sometimes more than 3-6 bonds away and a large fragment is needed to accurately represent the chemical environment of the parent molecule. Two such examples are shown in figure 14 E and F. In E, not only is the protonated nitrogen needed (shown in green), but the alcohol group is also needed to achieve good WBO distribution overlap (shown in purple). In F, the phosphate group nine bonds away from the bond of interest is needed to get the density of of the mode at 1.2 (shown in blue and purple).

4. Discussion

4.1 Combinatorial fragmentation benchmark set contains rich, chemical information that can be useful for other applications

The validation set used to benchmark our fragmentation scheme, and determine the disruption threshold to use, was specifically selected to validate a fragmentation scheme for QC torsion scan so we wanted molecules that are challenging to fragment. Therefor, the hundred molecules chosen include bonds that are sensitive to remote chemical changes and the functional groups they are sensitive to exhibited relatively large, long range effects. However, these molecules were filtered out from a larger set with a total of 1,234 molecules and ~300,000 fragments with their WBO distributions over conformations. This exhaustive fragmentation dataset, with their changing WBO distributions, provides a treasure trove of nuanced chemical data. For the purposes of this paper, we did not delve into interesting effects we observed because it is out of scope. Here we describe some of those observations, and provide some thoughts on how this kind of data can be useful for other applications.

At closer proximities that was not relevant for this study, and when functional groups are fragmented, the changes in the distributions detect varied effects. Many of these effects confirmed known considerations, such as removing a carbonyl from an amide or carboxylic group. But some were more subtle, such as changes observed vis-à-vis primary, secondary, and tertiary amines. The changes seem to pick up on subtle pKa changes for different amines [59]. In addition, the shifts in the distributions observed in figure 9 are also picking up on the effects of fluorine on sulfinamide. should I add more figures on the kind of things I saw? SI?

This kind of data can potentially be used to complement knowledge based molecular similarity applications which are usually defined locally and might not detect long range effects. In addition,

since WBO is anti-correlated with Hammett resonance parameters (fig 6 C and D), and shifts in distributions also seem to detect pKa shifts, WBOs may be useful in improving pKa predictions.

We are sharing the dataset as a benchmark set for future fragmentation schemes. It is also straight forward to generate such data for molecules that are not in the set by following the directions given in the detailed methods.

4.2 Bond orders can be used to fit force field parameters

WBO is one form of quantum mechanical electron density based bond order calculations, among many other definitions of bond orders (Pauling [60], Coulson [61], Mulliken [62] Mayer [63], Jug, [64], Politzer [65], atomic overlap matrix bond order [66], natural resonance theory bond order [67], Nalewajksi-Mrozek bond order [68], effective bond order [69], natural localized molecular orbital bond order [70], delocalization index [71], fuzzy bond order [72]) that aim to quantify the chemical bond, a key concept in chemistry, by computing the electron population overlap between atoms. These quantities attempt to provide a link between the physical understanding of molecules, a system of individual particles without explicit bonds, to the powerful, mental, graphical, models that chemists employ when thinking about molecules. Given that these quantities try to make that connection, it is not surprising that fractional bond orders captures important, chemical properties that can be useful in many applications, specifically force field parameterization. Indeed, In the MMP2 [73], MM3 [74] and MM4 [75] force fields, a Variable Electronegativity SCF derived bond order for pi-systems was used to fit bond length, bond force constants and twofold torsional force constants. Given the relationship we find in figure 6, E, we should be able to extrapolate or interpolate torsion force constants by calculating the WBO, thus avoiding running expensive QC torsion scans. The WBO comes free with an AM1-BCC [76] charging calculation at no extra cost, and is also now possible to calculate EHT electronic populations with the OFF toolkit [77]. The SMIRNOFF format provides a convenient way to add appropriate torsion terms given the data we have. For example, In figure 6, E, the lines seem to cluster into three clusters, so we can us ChemPer [43] to generate SMIRKS patterns to capture the chemistries in these clusters and interpolate the torsion force constants for the different chemical environments of those patterns. This has the potential to avoid many expensive QC torsion scans, specifically for bespoke parameter fitting for new chemical entities, and improve torsion parameters by including long range effects.

In addition, the ELF10 WBO can potentially be useful to determine if a bond is rotatable rather than relying on cheminformatics definitions. This can allow us to avoid running expensive, hard to converge, QC torsion scans for bonds that have very high torsion energy barriers. Instead,

4.3 Relative changes in WBO is not enough to capture all characteristics of QM torsion scans

The WBO is a measure of electron population overlap between atoms in a bond, so its relative changes is a good indication of conjugation and therefor a surrogate for torsion barrier heights (fig 6, E). However, as we have shown in figure 2 and [Hold for SI], relative changes in WBO for the same torsion type in different chemical environments only capture change of scale in torsion scans, not changes in the profile (relative amplitudes, periodicities and location of minima and maxima). There are several ways a torsion scan of the same torsion type in different chemical environments can change besides scale. One, symmetry around zero is lost, either with the minima and maxima shifting, or when one maxima is higher than the corresponding maxima across the symmetry line at zero degrees [Hold for SI]. In these cases, corresponding WBO scans have the same features as the QC scans. This indicates that the changes in the profiles correspond to electronic changes in the different torsions. All scans that exhibit these kind of changes have torsions types that include trivalent nitrogen which can be either in planar or pyrmidal conformations. When we measured the improper angle of the nitrogen involved in the torsion scan, along the scan, we found that the scans that were

shifted relative to most other scans in the series, had improper angles that were pyrmidal throughout the scan. The other molecules in the series all had improper angles that became planar at lower energies to allow conjugation [Hold for SI]. A trivalent nitrogen will be pyramidal if the lone pair is not conjugated, and the scans that have improper angles along the entire QC scan all fall on the lower end of WBOs within the series which indicates that the lone pairs are not conjugated with the phenyl ring. Another interesting observation is that in most cases where the nitrogen is pyramidal throughout the scan, its chirality stays the same, however, for the scans where the relative barrier heights are not symmetric, the chirality of the nitrogen flips during the scan [Hold for SI].

Another way QC torsion scan profiles can change besides scale, are when the the relative heights of the minimas or maximas are different or new minima or maxima are observed. Urea in figure Z, C and in [hold for SI] is an extreme example of this kind of change, but other series with bulkier X~₁~ [hold for SI] exhibit similar changes. In these cases, the correspond WBO scans do not have these features, but have similar profiles to other WBO scans in the series. This indicates that the observed changes in the profiles do not implicate changes in conjugation. Furthermore, when their corresponding improper angles (when relevant) were measured, we found that all of them become planar at low energy which imply that all molecules in the series are conjugated along the phenyl ring and X₁. Here, steric interactions cause the changes in QC profiles. In all scans where one minima relative to the other minima in the scan is higher, and this is not the case for other scans in the series, a bulky group is at the meta position, while the other scans have X₂ at para positions. In addition, for other scans where the profiles do not change as much and X₁ is not as bulky, but the barrier heights are out of order of increasing ELF10 WBO (a lower ELF10 WBO has a higher torsion energy barrier than another molecule with a higher ELF10 WBO in the series), X₂ is at the meta position.

The torsion parameters in forcefields are supposed to model both conjugation, a quantum chemical phenomenon that is not well modeled in classical forcefields, and corrections for 1-4 nonbonded interactions. To increase transferability of torsion parameters, torsion parameters should not include non bonded interactions beyond the 1-4 atoms. However, in general, it is difficult to separate the contributions of sterics and conjugation in a QC torsion scan. Here, it seems like the WBO scans along torsion scans, and relative ELF10 WBOs can provide a way to separate these contributions. If a torsion profile changes relative to another torsion profile of the same torsion type, and their WBO scans along the torsion profile only change in scale, or if their relative barrier heights are not in the same order as their ELF10 WBO estimate, the changes are probably coming from nonbonded interactions beyond the 1-4 atoms. However, if changes in QC torsion scans of the same torsion types are accompanied with the same profile changes in the corresponding WBO scan, then the profile change is inherent to the electron population overlap of the bond and might need a different torsion type.

4.4 Using Bond orders when fragmenting molecules captures long range effects that simple rules do not

Relative changes in bond orders between the fragments and their parent molecules, are a good indication of disruption of electron density around a central bond. QC torsion scans capture both steric and conjugation effects so torsion force field parameters should capture both short range non bonded corrections and conjugation. However, simple fragmentation rules assumes that including 1-5 atoms around the central bond (including rings and functional groups) is enough to capture conjugation effects in addition to sterics. While this assumption holds true for many molecules, it is not sufficient for sensitive bonds and functional groups that have long range effects, as we have shown. Therefor, using the WBO to detect changes to a bond's chemical environment, which is sensitive to nonlocal effects, ameliorates some deficiencies in rule based fragmentation schemes. But there is a trade off and the solution is not always optimal. While WBO calculations are cheap relative to QC torsion scans, they still require several semi-empirical QM calculations, the ELF10 estimate is sometimes not an adequate estimate, and heuristics used to add substituents to minimize electron density disruption do not always find the optimal path.

One way to speed up conjugation perception, is using RDKit's conjugation detector which relies on rules. However, these rules are binary; a bond is either considered fully conjugated or not. However, Conjugation is a continuous property and the extent of conjugation determines relative barrier heights. Boolean values do not allow us to detect such relative changes to a bond's chemical environment that will effect torsion barrier height. A better approach is to extend the concept of H-TEQ (hyperconjugation for torsional energy quantification) developed in [35,78] to include not just the 1-4 atoms in bonds adjacent to conjugated systems, or aromatic rings in biaryls, but other atoms in the molecule to obtain V_1 and V_2 estimates, and use those values instead of WBOs to determine the disruption of electron population overlap around the bond.

However, both of the above solutions, while reducing the cost of extent of conjugation detection, will still rely on needing to find the optimal path to grow our the fragment. A data-driven approach, which can find the optimal fragment is the ideal solution. The OFF QC datasets on QCArchive [17] all include Wiberg Löwdin bond orders for all conformations. Given the sheer number of data available on QCArchive, and the long range information WBOs captures, it will be possible to train an ML model to find which parts of a molecule are electronically coupled and need to be conserved when fragmenting to reproduce the parent molecule chemical environment. This data-driven approach will reduce the cost of fragmenting and also potentially find the most optimal fragment that minimizes both electronic density disruption and fragment size. It can also provide data needed to determine how many torsion SMIRKS types are needed for maximum transferability and chemical coverage.

5. Detailed method

5.1 QCArchive automates QC data generation and archives the resulting data

5.1.1 Submitting computations to QCArchive ensures reproducibility

[Hold for Daniel to write up. Please describe how QCArchive runs torsion scans and how to submit and retrieve data]

5.1.2 Details on QC and MM torsion scans.

All QC calculations were computed at B3LYP-D3(BJ) / DZVP. This level of theory was chosen based on benchmark conducted by the Open Force Field consortium for fitting Parsley [79]

Molecular mechanics torsion scans were run on QCArchive using OpenMM [80] for the energy and gradient evaluations. The molecules were parameterized with the OpenFF parsley Force Field (v1.0.0) [81]

5.2 Calculating Bond orders

5.2.1 AM1 WBO and AM1 ELF10 WBO

To calculate AM1 ELF10 WBO, we used OpenEye's QUACPAC toolkit [82] (OpenEye version 2019.Apr.2). The ELF10 WBO comes along free after an AM1-BCC charge fitting procedure. For ELF10 WBOs generated in this paper, we used the `get_charges` function in the `chemi.py` module in `fragmenter` versions v0.0.3 and v0.0.4. To calculate AM1 WBO for individual conformers, we used the `OEAssignPartialCharges` with the `OECharges_AM1BCCSym` option from the QUACPAC toolkit for each conformer generated with Omega [83] (OpenEye versio 2019.Apr.2) which is called for the `get_charges` function. For AM1 WBOs calculated to verify the results from the validation set, we generated conformers using the `generate_grid_conformer` function in the `chemi.py` module in `fragmenter` version v0.0.4.+25.gbb12030

5.2.2 Wiberg Bond Orders calculated with larger basis sets

[Leave to Daniel to describe how Wiberg bond orders are calculated in QCArchive (it only calculates Wiberg-Löwdin so provide that formula and why the Löwdin normalization is needed)]

5.3 Datasets

5.3.1 Kinase inhibitor dataset

The kinase inhibitor dataset consists of 43 FDA approved kinase inhibitors (smi files in the SI) with their Omega generated conformers (OpenEye veriso 2019.Apr.2, `generate_conformers` function in the `chemi.py` module in `fragmenter` version v0.0.4). AM1 WBOs were calculated as described above, for all conformers of all 43 kinase inhibitors. B3LYP-D3(BJ) / DZVP Wiberg-Löwdin bond orders were calculated for 9 kinase inhibitors and Omega generated conformers after a B3LYP-D3P(BJ) / DZVP geometry optimization. The DFT results are available on QCArchive as an `OptimizationDataset` named `Kinase Inhibitors: WBO Distributions`.

The variance of the WBO distributions were calculated using the numpy [84] `var` function version 1.16.2 and their confidence intervals were calculated using arch `IIDBootstrap` function [85] version 4.8.1. To calculate the correlation matrix, we calculated the Pearson correlation coefficient with the numpy [84] `corrcoef` function version 1.16.2. Scripts and data used to generate and analyze this dataset are in github.com/choderalab/fragmenter_data/manuscript-figures/kinase_inhibitors_wbos

5.3.2 Substituted phenyl dataset

The substituted phenyl dataset consists of 3,458 substituted phenyl molecules where the substituents chosen to span a large range of electron donating and withdrawing groups. We arrived at 3,200 molecules by attaching 26 different functional groups to 5 scaffolds (Figure 6, A) at the X_1 position, and then attach the 26 functional group (and H) at the X_2 position for a total of 133 molecules per functional group ($26 * 5 + 3$ (for molecules with H at the X_2 position)). The AM1 ELF10 WBOs were calculated as described above. We selected molecules for QC torsion scans as follows: 1. From the 26 functional groups, we only selected molecules from 18 functional groups, skipping X_1 s that either did not have a rotatable bond (fluorine, chlorine, iodine, bromine, nitrile, oxygen), were too congested (trifluoromethyl, trimethylammonium) or where the WBOs on the bonds attaching X_1 to the phenyl ring did not change much with different chemical group at the X_2 position (methyl). 2. For the 18 functional groups, we chose molecules that were evenly spaced along the WBO range of that functional group, up to 15 molecules. While all the skipped functional groups for X_1 were allowed to be at X_2 , we did not include the negative oxygen at X_2 because OpenFF have not yet benchmarked the level of theory to use for anions. 3. After selection, we had 140 molecules that we submitted to QCArchive for both QC and MM torsion scan. The dataset is available on QCArchive as a `TorsionDriveDataset` named `OpenFF Substituted Phenyl Set 1`. This dataset also includes the biphenyl torsion scans shown in Figure 2

There is another substituted phenyl set on QCArchive whose results are not shown in this paper because it was run with a different level of theory as the default OpenFF level of theory, included anions which we did not yet decide how to handle and did not have good coverage of WBO ranges.

5.3.3 Exhaustive fragmentation dataset

The exhaustive fragmentation dataset was generated by filtering DrugBank version (version 5.1.3 downloaded on 2019-06-06) [19] with criteria described in section 4 and repeated here for clarity: 1.

FDA approved 2. Ring sized between 3 and 14 heavy atoms 3. Rotatable bonds between 4 and 10 4. At least one aromatic ring 5. Only 1 connected component

This left us with 730 molecules. To expand states, we used `OEGetReasonableTautomers` from QUACPAC (OpenEye version 2019.Apr.2) in the `states.py` module in `fragmenter` (version v0.0.2+175.g6fbbf32 for this original set, but versions v0.0.3 and v0.0.4 will generate the same results with the same options). We set `pKaNorm` to True so that the ionization state of each tautomer is assigned to a predominant state at pH ~ 7.4. This generated 1289 molecules.

We then used the `CombinatorialFragmenter` from `fragmenter` version v0.0.2+179.g0e7e9e3 (versions v0.0.3 and v0.0.4 will generate the same fragments with the same options) to generate all possible fragments for each molecules. We set the option `functional_groups` to False so that all functional groups besides rings will also get fragmented so we can use the data to explore which functional groups should not be fragmented. We used the default settings for all other options (`min_rotor` is 1 and `min_heavy_atoms` is 4 so that the smallest fragments have at least one torsion. `max_rotors` is the number of rotors in the parent molecules so that the largest fragment generated is one less rotor than the parent molecule). This generated ~300,000 fragments.

We then used Omega (OpenEye version 2019.Apr.2) to generate conformers for each fragment and calculated each conformer's WBOs as described above. All scripts used to generate this dataset are in github.com/choderalab/fragmenter_data/combinatorial_fragmentation. The dataset is available as zip files on (hold for link to where we will host the data).

5.4 Fragmenting molecules

The `fragmenter` package provides several fragmentation schemes with various options. Below we discuss different modes of fragmentation and their options.

5.4.1 Exhaustive fragmentation generates all possible fragments of a parent molecule.

This functionality is provided by the `CombinatorialFragmenter` class in the `fragment.py` module. To use this class, the user needs to provide an openeye molecule. `fragmenter` provides a list of functional groups SMARTS in a yaml file located in `fragmenter/data/fgroup_smarts_combs.yaml` that it will not fragment by default. This list is given in table 4. The list is different than the default list used on the `WBOFragmenter` because here the carbon bonded to the functional groups are also tagged. To allow all functional groups to be fragmented, the user can set the parameter `functional_groups = False`. This option will fragment all bonds besides bond in rings. The user can also provide their own dictionary of functional group SMARTS patterns that they wish to avoid fragmenting.

Table 4: Default functional groups that the `CombinatorialFragmenter` will not fragment. This list is not comprehensive and is different than the list used for the `WBOFragmenter`

Chemical group	SMARTS pattern
amide	<code>[NX3R0] [CX3] (=[OX1])</code>
sulfone	<code>[#16X4] (=[OX1])=([OX1])</code>
phosphine_oxide	<code>[PX4] (=[OX1])([CX4])([CX4])</code>
phosphon	<code>[PX4] (=[OX1])([OX2])([OX2])</code>
trimethyl	<code>[CX4!H] ([CX4H3])([CX4H3])([CX4H3])</code>
tri_halide	<code>[#6]((([F,Cl,I,Br])[F,Cl,I,Br])[F,Cl,I,Br])</code>

Chemical group	SMARTS pattern
carboxylic_acid	[CX3] (=O) [OX2H1]
ester	[CX3] (=O) [OX2H0]
dimethyl	[CX4H1] ([CX4H3]) ([CX4H3])
carbonyl	[CX3R0]=[OX1]
alkyne	[CX2]#[CX2]
nitrile	[NX1]#[CX2]

The user can also set the minimum and maximum number of rotatable bonds, and minimum heavy atoms in a fragment.

5.4.2 Generate minimal fragments

The `PfizerFragmenter` implements the scheme developed at Pfizer and described in [29]. It uses the same list of functional groups as the `WBOFragmenter` uses. The user can also provide their own SMARTS patterns of functional groups not to fragment.

5.4.3 Using the WBO as a surrogate for changes in chemical environment.

The `WBOFragmenter` implements the FBO scheme described in this paper. The functional groups that are not fragmented are given in table 2. Users can add more SMARTS patterns if needed.

When the WBO on a central bond in a minimal fragment has been disrupted more than the threshold, remote atoms need to be added onto the fragment. However, there are multiple ways to grow out a fragment and enumerating all possible ways to find the best one can become too computationally expensive. Therefore, we need to use heuristics to decide where to add the next atoms. The two heuristics available in `fragmenter` are:

- 1. Shortest path length** Here, the bond with the shortest path to the central bond is added next. The rationale for this heuristic is that atoms closer to the central bond will have a greater influence to the bond's chemical environment. If more than one connected bond has the shortest path to the central bond, the bond with the higher WBO is added next.
- 2. Greatest WBO** Here, the bonds connected to the fragment that have the greatest WBO is added next. The rationale for this heuristic is that bonds with higher WBO are more likely to be involved in extended conjugation that can influence the central bond.

Both of these heuristics will sometimes give different results [\[Hold for SI figure\]](#). We found that for the benchmark set we tested, the shortest path heuristic preformed better, or found more optimal fragments when compared to using the greatest WBO heuristic [\[Hold for SI\]](#).

Currently, `fragmenter` depends on OpenEye to provides three modes of fragmentation described here. In the future, `fragmenter` will be incorporated into the `openforcefield` toolkit and will have the option to use RDKit [86], an open source cheminformatic libraries. However, given that RDKit supports EHT instead of AM1, the fragments might be different.

Conclusion

We have shown that the ELF10 WBO estimate is a simple, yet informative quantity about the extent of binding between two connecting atoms, thus descriptive of a bond's chemical environment, its level of conjugation, and resistance to rotation. We can use the change in WBO of a bond to quantify the amount of disruption of its chemical environment due to remote chemical substituent changes, specifically for bonds that are sensitive to peripheral chemical changes such as bonds in or adjacent to conjugated systems, or bonds involving atoms that have lone pairs.

We used this concept to extend a rule-based fragmentation scheme to improve the resulting fragments, by adding remote substituents until the change in WBO is lower than a user defined threshold. We generated a validation set using exhaustive fragmentation to benchmark fragmentation schemes and found that a threshold of 0.03 will find the most fragments that minimize both fragment size and distance to the parent's conformation distribution about the bond. We found eight chemical groups that have long-range effects on sensitive bonds and their inclusion is necessary to recapitulate a parent's chemical environment even if they are 3-6 bonds away from the central bond.

References

1. Toward Learned Chemical Perception of Force Field Typing Rules

Camila Zanette, Caitlin C. Bannan, Christopher I. Bayly, Josh Fass, Michael K. Gilson, Michael R. Shirts, John D. Chodera, David L. Mobley

Journal of Chemical Theory and Computation (2018-12-04) <https://doi.org/gft4hf>

DOI: [10.1021/acs.jctc.8b00821](https://doi.org/10.1021/acs.jctc.8b00821) · PMID: [30512951](https://pubmed.ncbi.nlm.nih.gov/30512951/) · PMCID: [PMC6467725](https://pubmed.ncbi.nlm.nih.gov/PMC6467725/)

2. OPLS3: A Force Field Providing Broad Coverage of Drug-like Small Molecules and Proteins

Edward Harder, Wolfgang Damm, Jon Maple, Chuanjie Wu, Mark Reboul, Jin Yu Xiang, Lingle Wang, Dmitry Lupyan, Markus K. Dahlgren, Jennifer L. Knight, ... Richard A. Friesner

Journal of Chemical Theory and Computation (2015-12) <https://doi.org/f76wpm>

DOI: [10.1021/acs.jctc.5b00864](https://doi.org/10.1021/acs.jctc.5b00864) · PMID: [26584231](https://pubmed.ncbi.nlm.nih.gov/26584231/)

3. Accuracy evaluation and addition of improved dihedral parameters for the MMFF94s

Joel Wahl, Joel Freyss, Modest von Korff, Thomas Sander

Journal of Cheminformatics (2019-08-07) <https://doi.org/gf6rz2>

DOI: [10.1186/s13321-019-0371-6](https://doi.org/10.1186/s13321-019-0371-6) · PMID: [31392432](https://pubmed.ncbi.nlm.nih.gov/31392432/) · PMCID: [PMC6686419](https://pubmed.ncbi.nlm.nih.gov/PMC6686419/)

4. Paramfit: Automated optimization of force field parameters for molecular dynamics simulations

Robin M. Betz, Ross C. Walker

Journal of Computational Chemistry (2014-11-21) <https://doi.org/f6svdh>

DOI: [10.1002/jcc.23775](https://doi.org/10.1002/jcc.23775) · PMID: [25413259](https://pubmed.ncbi.nlm.nih.gov/25413259/)

5. Rapid parameterization of small molecules using the force field toolkit

Christopher G. Mayne, Jan Saam, Klaus Schulten, Emad Tajkhorshid, James C. Gumbart

Journal of Computational Chemistry (2013-09-02) <https://doi.org/f5ggrj>

DOI: [10.1002/jcc.23422](https://doi.org/10.1002/jcc.23422) · PMID: [24000174](https://pubmed.ncbi.nlm.nih.gov/24000174/) · PMCID: [PMC3874408](https://pubmed.ncbi.nlm.nih.gov/PMC3874408/)

6. Fitting of Dihedral Terms in Classical Force Fields as an Analytic Linear Least-Squares Problem

Chad W. Hopkins, Adrian E. Roitberg

Journal of Chemical Information and Modeling (2014-07-09) <https://doi.org/f6cffs>

DOI: [10.1021/ci500112w](https://doi.org/10.1021/ci500112w) · PMID: [24960267](https://pubmed.ncbi.nlm.nih.gov/24960267/)

7. A Second Generation Force Field for the Simulation of Proteins, Nucleic Acids, and Organic Molecules

Wendy D. Cornell, Piotr Cieplak, Christopher I. Bayly, Ian R. Gould, Kenneth M. Merz, David M. Ferguson, David C. Spellmeyer, Thomas Fox, James W. Caldwell, Peter A. Kollman

Journal of the American Chemical Society (1995-05) <https://doi.org/dbzh27>

DOI: [10.1021/ja00124a002](https://doi.org/10.1021/ja00124a002)

8. CHARMM: A program for macromolecular energy, minimization, and dynamics calculations

Bernard R. Brooks, Robert E. Brucolieri, Barry D. Olafson, David J. States, S. Swaminathan, Martin Karplus

Journal of Computational Chemistry (1983) <https://doi.org/bqh7f2>

DOI: [10.1002/jcc.540040211](https://doi.org/10.1002/jcc.540040211)

9. Development and Testing of the OPLS All-Atom Force Field on Conformational Energetics and Properties of Organic Liquids

William L. Jorgensen, David S. Maxwell, Julian Tirado-Rives

Journal of the American Chemical Society (1996-01) <https://doi.org/fvftxj>
DOI: [10.1021/ja9621760](https://doi.org/10.1021/ja9621760)

10. Automated conformational energy fitting for force-field development

Olgun Guvench, Alexander D. MacKerell Jr.

Journal of Molecular Modeling (2008-05-06) <https://doi.org/bzphqw>

DOI: [10.1007/s00894-008-0305-0](https://doi.org/10.1007/s00894-008-0305-0) · PMID: [18458967](https://pubmed.ncbi.nlm.nih.gov/18458967/) · PMCID: [PMC2864003](https://pubmed.ncbi.nlm.nih.gov/PMC2864003/)

11. Machine learning of correlated dihedral potentials for atomistic molecular force fields

Pascal Friederich, Manuel Konrad, Timo Strunk, Wolfgang Wenzel

Scientific Reports (2018-02-07) <https://doi.org/gczmpn>

DOI: [10.1038/s41598-018-21070-0](https://doi.org/10.1038/s41598-018-21070-0) · PMID: [29416116](https://pubmed.ncbi.nlm.nih.gov/29416116/) · PMCID: [PMC5803249](https://pubmed.ncbi.nlm.nih.gov/PMC5803249/)

12. Empirical force fields for biological macromolecules: Overview and issues

Alexander D. Mackerell

Journal of Computational Chemistry (2004) <https://doi.org/dbhsbb>

DOI: [10.1002/jcc.20082](https://doi.org/10.1002/jcc.20082) · PMID: [15264253](https://pubmed.ncbi.nlm.nih.gov/15264253/)

13. ff19SB: Amino-Acid Specific Protein Backbone Parameters Trained Against Quantum Mechanics Energy Surfaces in Solution

Chuan Tian, Koushik Kasavajhala, Kellon Belfon, Lauren Raguette, He Huang, Angela Migues, John Bickel, Yuzhang Wang, Jorge Pincay, Qin Wu, Carlos Simmerling

American Chemical Society (ACS) (2019-06-17) <https://doi.org/gf6rz8>

DOI: [10.26434/chemrxiv.8279681](https://doi.org/10.26434/chemrxiv.8279681)

14. A consistent and accurate ab initio parametrization of density functional dispersion correction (DFT-D) for the 94 elements H-Pu

Stefan Grimme, Jens Antony, Stephan Ehrlich, Helge Krieg

The Journal of Chemical Physics (2010-04-21) <https://doi.org/bnt82x>

DOI: [10.1063/1.3382344](https://doi.org/10.1063/1.3382344) · PMID: [20423165](https://pubmed.ncbi.nlm.nih.gov/20423165/)

15. Optimization of Gaussian-type basis sets for local spin density functional calculations. Part I. Boron through neon, optimization technique and validation

Nathalie Godbout, Dennis R. Salahub, Jan Andzelm, Erich Wimmer

Canadian Journal of Chemistry (1992-02) <https://doi.org/c78qjn>

DOI: [10.1139/v92-079](https://doi.org/10.1139/v92-079)

16. A quantitative study of the scaling properties of the Hartree–Fock method

Douglas L. Strout, Gustavo E. Scuseria

The Journal of Chemical Physics (1995-06) <https://doi.org/b3323h>

DOI: [10.1063/1.468836](https://doi.org/10.1063/1.468836)

17. The MolSSI QC Archive <https://qcarchive.molssi.org/>

18. Building a More Predictive Protein Force Field: A Systematic and Reproducible Route to AMBER-FB15

Lee-Ping Wang, Keri A. McKiernan, Joseph Gomes, Kyle A. Beauchamp, Teresa Head-Gordon, Julia E. Rice, William C. Swope, Todd J. Martínez, Vijay S. Pande

The Journal of Physical Chemistry B (2017-04-06) <https://doi.org/f92nv5>

DOI: [10.1021/acs.jpcb.7b02320](https://doi.org/10.1021/acs.jpcb.7b02320) · PMID: [28306259](https://pubmed.ncbi.nlm.nih.gov/28306259/)

19. DrugBank 5.0: a major update to the DrugBank database for 2018

David S Wishart, Yannick D Feunang, An C Guo, Elvis J Lo, Ana Marcu, Jason R Grant, Tanvir Sajed,

Daniel Johnson, Carin Li, Zinat Sayeeda, ... Michael Wilson
Nucleic Acids Research (2017-11-08) <https://doi.org/gcwtzk>
DOI: [10.1093/nar/gkx1032](https://doi.org/10.1093/nar/gkx1032) · PMID: [29126136](https://pubmed.ncbi.nlm.nih.gov/29126136/) · PMCID: [PMC5753335](https://pubmed.ncbi.nlm.nih.gov/PMC5753335/)

20. RECAPRetrosynthetic Combinatorial Analysis Procedure: A Powerful New Technique for Identifying Privileged Molecular Fragments with Useful Applications in Combinatorial Chemistry

Xiao Qing Lewell, Duncan B. Judd, Stephen P. Watson, Michael M. Hann
Journal of Chemical Information and Computer Sciences (1998-04-11) <https://doi.org/d4z4pf>
DOI: [10.1021/ci970429i](https://doi.org/10.1021/ci970429i) · PMID: [9611787](https://pubmed.ncbi.nlm.nih.gov/9611787/)

21. Break Down in Order To Build Up: Decomposing Small Molecules for Fragment-Based Drug Design with eMolFrag

Tairan Liu, Misagh Naderi, Chris Alvin, Supratik Mukhopadhyay, Michal Brylinski
Journal of Chemical Information and Modeling (2017-04-04) <https://doi.org/f9x9bg>
DOI: [10.1021/acs.jcim.6b00596](https://doi.org/10.1021/acs.jcim.6b00596) · PMID: [28346786](https://pubmed.ncbi.nlm.nih.gov/28346786/) · PMCID: [PMC5433162](https://pubmed.ncbi.nlm.nih.gov/PMC5433162/)

22. The Properties of Known Drugs. 1. Molecular Frameworks

Guy W. Bemis, Mark A. Murcko
Journal of Medicinal Chemistry (1996-01) <https://doi.org/fshj3p>
DOI: [10.1021/jm9602928](https://doi.org/10.1021/jm9602928) · PMID: [8709122](https://pubmed.ncbi.nlm.nih.gov/8709122/)

23. pyEFP: Automatic decomposition of the complex molecular systems into rigid polarizable fragments

Alexey V. Odinokov, Nikita O. Dubinets, Alexander A. Bagaturyants
Journal of Computational Chemistry (2017-12-26) <https://doi.org/gcq4qs>
DOI: [10.1002/jcc.25149](https://doi.org/10.1002/jcc.25149) · PMID: [29280158](https://pubmed.ncbi.nlm.nih.gov/29280158/)

24. Approximate ab initio energies by systematic molecular fragmentation

Vitali Deev, Michael A. Collins
The Journal of Chemical Physics (2005-04-15) <https://doi.org/ch4zhg>
DOI: [10.1063/1.1879792](https://doi.org/10.1063/1.1879792) · PMID: [15945620](https://pubmed.ncbi.nlm.nih.gov/15945620/)

25. Fragmentation Methods: A Route to Accurate Calculations on Large Systems

Mark S. Gordon, Dmitri G. Fedorov, Spencer R. Pruitt, Lyudmila V. Slipchenko
Chemical Reviews (2011-08-26) <https://doi.org/b8tc8n>
DOI: [10.1021/cr200093j](https://doi.org/10.1021/cr200093j) · PMID: [21866983](https://pubmed.ncbi.nlm.nih.gov/21866983/)

26. Systematic fragmentation of large molecules by annihilation

Michael A. Collins
Physical Chemistry Chemical Physics (2012) <https://doi.org/gf6v2d>
DOI: [10.1039/c2cp23832b](https://doi.org/10.1039/c2cp23832b) · PMID: [22373545](https://pubmed.ncbi.nlm.nih.gov/22373545/)

27. Linear-scaling semiempirical quantum calculations for macromolecules

Tai-Sung Lee, Darrin M. York, Weitao Yang
The Journal of Chemical Physics (1996-08-15) <https://doi.org/bdtpqw>
DOI: [10.1063/1.472136](https://doi.org/10.1063/1.472136)

28. Flexible effective fragment QM/MM method: Validation through the challenging tests

A. V. Nemukhin, B. L. Grigorenko, I. A. Topol, S. K. Burt
Journal of Computational Chemistry (2003-07-11) <https://doi.org/dpwk5b>
DOI: [10.1002/jcc.10309](https://doi.org/10.1002/jcc.10309) · PMID: [12868106](https://pubmed.ncbi.nlm.nih.gov/12868106/)

29. Comprehensive Assessment of Torsional Strain in Crystal Structures of Small Molecules and Protein-Ligand Complexes using ab Initio Calculations

Brajesh K. Rai, Vishnu Sresht, Qingyi Yang, Ray Unwalla, Meihua Tu, Alan M. Mathiowetz, Gregory A. Bakken

Journal of Chemical Information and Modeling (2019-10) <https://doi.org/ggfvzc>

DOI: [10.1021/acs.jcim.9b00373](https://doi.org/acs.jcim.9b00373) · PMID: [31573196](#)

30. Application of the pople-santry-segal CNDO method to the cyclopropylcarbinyl and cyclobutyl cation and to bicyclobutane

K. B. Wiberg

Tetrahedron (1968-01) <https://doi.org/fvwkkh>

DOI: [10.1016/0040-4020\(68\)88057-3](https://doi.org/10.1016/0040-4020(68)88057-3)

31. Bond Order Analysis Based on the Laplacian of Electron Density in Fuzzy Overlap Space

Tian Lu, Feiwu Chen

The Journal of Physical Chemistry A (2013-04-02) <https://doi.org/f4t9v3>

DOI: [10.1021/jp4010345](https://doi.org/10.1021/jp4010345) · PMID: [23514314](#)

32. Resonance interactions in acyclic systems

K. B. Wiberg, C. M. Breneman, K. E. Laidig, R. E. Rosenberg

Pure and Applied Chemistry (1989-01-01) <https://doi.org/c6rd43>

DOI: [10.1351/pac198961040635](https://doi.org/10.1351/pac198961040635)

33. Predicting Trigger Bonds in Explosive Materials through Wiberg Bond Index Analysis

Lenora K. Harper, Ashley L. Shoaf, Craig A. Bayse

ChemPhysChem (2015-11-06) <https://doi.org/f3jt5h>

DOI: [10.1002/cphc.201500773](https://doi.org/10.1002/cphc.201500773) · PMID: [26458868](#)

34. Torsional Energy Barriers of Biaryls Could Be Predicted by Electron Richness/Deficiency of Aromatic Rings; Advancement of Molecular Mechanics toward Atom-Type Independence

Wanlei Wei, Candide Champion, Zhaomin Liu, Stephen J. Barigye, Paul Labute, Nicolas Moitessier

Journal of Chemical Information and Modeling (2019-08-20) <https://doi.org/ggh677>

DOI: [10.1021/acs.jcim.9b00585](https://doi.org/acs.jcim.9b00585) · PMID: [31430147](#)

35. Atom Types Independent Molecular Mechanics Method for Predicting the Conformational Energy of Small Molecules

Zhaomin Liu, Stephen J. Barigye, Moeed Shahamat, Paul Labute, Nicolas Moitessier

Journal of Chemical Information and Modeling (2018-01-05) <https://doi.org/gcxm7s>

DOI: [10.1021/acs.jcim.7b00645](https://doi.org/10.1021/acs.jcim.7b00645) · PMID: [29253333](#)

36. Hyperconjugation not steric repulsion leads to the staggered structure of ethane

Vojislava Popristic, Lionel Goodman

Nature (2001-05) <https://doi.org/dff6j7>

DOI: [10.1038/35079036](https://doi.org/10.1038/35079036) · PMID: [11385566](#)

37. Gauche Effect in 1,2-Difluoroethane. Hyperconjugation, Bent Bonds, Steric Repulsion

Lionel Goodman, Hongbing Gu, Vojislava Popristic

The Journal of Physical Chemistry A (2005-02) <https://doi.org/dtg782>

DOI: [10.1021/jp046290d](https://doi.org/10.1021/jp046290d) · PMID: [16833433](#)

38. Zur Kenntnis der ungesättigten Verbindungen. Theorie der ungesättigten und aromatischen Verbindungen

Johannes Thiele

39. IUPAC - conjugated system (C01267)<https://goldbook.iupac.org/terms/view/C01267>

40. Hyperconjugation*

Robert S. Mulliken, Carol A. Rieke, Weldon G. Brown

Journal of the American Chemical Society (1941-01) <https://doi.org/brj5s3>

DOI: [10.1021/ja01846a008](https://doi.org/10.1021/ja01846a008)

41. A new force field for molecular mechanical simulation of nucleic acids and proteins

Scott J. Weiner, Peter A. Kollman, David A. Case, U. Chandra Singh, Caterina Ghio, Giuliano Alagona, Salvatore Profeta, Paul Weiner

Journal of the American Chemical Society (1984-02) <https://doi.org/dcvwwx>

DOI: [10.1021/ja00315a051](https://doi.org/10.1021/ja00315a051)

42. Escaping Atom Types in Force Fields Using Direct Chemical Perception

David L. Mobley, Caitlin C. Bannan, Andrea Rizzi, Christopher I. Bayly, John D. Chodera, Victoria T. Lim, Nathan M. Lim, Kyle A. Beauchamp, David R. Slochower, Michael R. Shirts, ... Peter K. Eastman

Journal of Chemical Theory and Computation (2018-10-11) <https://doi.org/gffnf3>

DOI: [10.1021/acs.jctc.8b00640](https://doi.org/acs.jctc.8b00640) · PMID: [30351006](#) · PMCID: [PMC6245550](#)

43. ChemPer: An Open Source Tool for Automatically Generating SMIRKS Patterns

Caitlin C. Bannan, David Mobley

American Chemical Society (ACS) (2019-06-21) <https://doi.org/gf66hw>

DOI: [10.26434/chemrxiv.8304578.v1](https://doi.org/10.26434/chemrxiv.8304578.v1)

44. Bond order and valence indices: A personal account

I. Mayer

Journal of Computational Chemistry (2006) <https://doi.org/ddssrj>

DOI: [10.1002/jcc.20494](https://doi.org/10.1002/jcc.20494) · PMID: [17066501](#)

45. Development and use of quantum mechanical molecular models. 76. AM1: a new general purpose quantum mechanical molecular model

Michael J. S. Dewar, Eve G. Zoebisch, Eamonn F. Healy, James J. P. Stewart

Journal of the American Chemical Society (1985-06) <https://doi.org/fd8bwp>

DOI: [10.1021/ja00299a024](https://doi.org/10.1021/ja00299a024)

46. Optimization of parameters for semiempirical methods I. Method

James J. P. Stewart

Journal of Computational Chemistry (1989-03) <https://doi.org/cg5pck>

DOI: [10.1002/jcc.540100208](https://doi.org/10.1002/jcc.540100208)

47. On the Non-Orthogonality Problem Connected with the Use of Atomic Wave Functions in the Theory of Molecules and Crystals

Per-Olov Löwdin

The Journal of Chemical Physics (1950-03) <https://doi.org/dj2c35>

DOI: [10.1063/1.1747632](https://doi.org/10.1063/1.1747632)

48. On the quantum theory of valence and bonding from the ab initio standpoint

Mario A. Natiello, Jorge A. Medrano

Chemical Physics Letters (1984-03) <https://doi.org/bdfk5f>

DOI: [10.1016/0009-2614\(84\)85645-6](https://doi.org/10.1016/0009-2614(84)85645-6)

49. Psi4 1.1: An Open-Source Electronic Structure Program Emphasizing Automation, Advanced Libraries, and Interoperability

Robert M. Parrish, Lori A. Burns, Daniel G. A. Smith, Andrew C. Simmonett, A. Eugene DePrince III, Edward G. Hohenstein, Uğur Bozkaya, Alexander Yu. Sokolov, Roberto Di Remigio, Ryan M. Richard, ... C. David Sherrill

Journal of Chemical Theory and Computation (2017-06-06) <https://doi.org/gcz64j>

DOI: [10.1021/acs.jctc.7b00174](https://doi.org/acs.jctc.7b00174) · PMID: [28489372](#)

50. Can we treat ab initio atomic charges and bond orders as conformation-independent electronic structure descriptors?

T. Yu. Nikolaienko, L. A. Bulavin, D. M. Hovorun

RSC Advances (2016) <https://doi.org/gf66tp>

DOI: [10.1039/c6ra17055b](https://doi.org/10.1039/c6ra17055b)

51. Resonance interactions in acyclic systems. 3. Formamide internal rotation revisited. Charge and energy redistribution along the C-N bond rotational pathway

Kenneth B. Wiberg, Curt M. Breneman

Journal of the American Chemical Society (1992-01) <https://doi.org/c85zcc>

DOI: [10.1021/ja00029a005](https://doi.org/10.1021/ja00029a005)

52. Conformer Generation with OMEGA: Algorithm and Validation Using High Quality Structures from the Protein Databank and Cambridge Structural Database

Paul C. D. Hawkins, A. Geoffrey Skillman, Gregory L. Warren, Benjamin A. Ellingson, Matthew T. Stahl

Journal of Chemical Information and Modeling (2010-03-17) <https://doi.org/d4rb6g>

DOI: [10.1021/ci100031x](https://doi.org/ci100031x) · PMID: [20235588](#) · PMCID: [PMC2859685](#)

53. Density-functional thermochemistry. III. The role of exact exchange

Axel D. Becke

The Journal of Chemical Physics (1993-04) <https://doi.org/d2fq9j>

DOI: [10.1063/1.464913](https://doi.org/10.1063/1.464913)

54. Effect of the damping function in dispersion corrected density functional theory

Stefan Grimme, Stephan Ehrlich, Lars Goerigk

Journal of Computational Chemistry (2011-03-01) <https://doi.org/b3ztst>

DOI: [10.1002/jcc.21759](https://doi.org/10.1002/jcc.21759) · PMID: [21370243](#)

55. MMFF VI. MMFF94s option for energy minimization studies

Thomas A. Halgren

Journal of Computational Chemistry (1999-05) <https://doi.org;brxdg7>

DOI: [10.1002/\(sici\)1096-987x\(199905\)20:7<720::aid-jcc7>3.0.co;2-x](https://doi.org/10.1002/(sici)1096-987x(199905)20:7<720::aid-jcc7>3.0.co;2-x)

56. Benchmarking Commercial Conformer Ensemble Generators

Nils-Ole Friedrich, Christina de Bruyn Kops, Florian Flachsenberg, Kai Sommer, Matthias Rarey, Johannes Kirchmair

Journal of Chemical Information and Modeling (2017-10-18) <https://doi.org/gb4v2v>

DOI: [10.1021/acs.jcim.7b00505](https://doi.org/acs.jcim.7b00505) · PMID: [28967749](#)

57. QUACPAC 2.0.2.2 — Applications,

v2019.Nov.2 <https://docs.eyesopen.com/applications/quacpac/index.html>

58. The Effect of Structure upon the Reactions of Organic Compounds. Benzene Derivatives

Louis P. Hammett

Journal of the American Chemical Society (1937-01) <https://doi.org/dz8d4r>
DOI: [10.1021/ja01280a022](https://doi.org/10.1021/ja01280a022)

59. Predicting and Tuning Physicochemical Properties in Lead Optimization: Amine Basicities
Martin Morgenthaler, Eliane Schweizer, Anja Hoffmann-Röder, Fausta Benini, Rainer E. Martin, Georg Jaeschke, Björn Wagner, Holger Fischer, Stefanie Bendels, Daniel Zimmerli, ... Klaus Müller
ChemMedChem (2007-08-13) <https://doi.org/cw6dkr>
DOI: [10.1002/cmdc.200700059](https://doi.org/10.1002/cmdc.200700059) · PMID: [17530727](#)

60. The Dependence of Interatomic Distance on Single Bond-Double Bond Resonance1
Linus Pauling, L. O. Brockway, J. Y. Beach
Journal of the American Chemical Society (1935-12) <https://doi.org/fsdwx>
DOI: [10.1021/ja01315a105](https://doi.org/10.1021/ja01315a105)

61. The electronic structure of some polyenes and aromatic molecules. VII. Bonds of fractional order by the molecular orbital method
Proceedings of the Royal Society of London. Series A. Mathematical and Physical Sciences (1939-02-07) <https://doi.org/bnqx76>
DOI: [10.1098/rspa.1939.0006](https://doi.org/10.1098/rspa.1939.0006)

62. Electronic Population Analysis on LCAO-MO Molecular Wave Functions. I
R. S. Mulliken
The Journal of Chemical Physics (1955-10) <https://doi.org/d24mtt>
DOI: [10.1063/1.1740588](https://doi.org/10.1063/1.1740588)

63. Charge, bond order and valence in the AB initio SCF theory
I. Mayer
Chemical Physics Letters (1983-05) <https://doi.org/b5m8r5>
DOI: [10.1016/0009-2614\(83\)80005-0](https://doi.org/10.1016/0009-2614(83)80005-0)

64. A maximum bond order principle
Karl Jug
Journal of the American Chemical Society (1977-11) <https://doi.org/dvfv63>
DOI: [10.1021/ja00466a009](https://doi.org/10.1021/ja00466a009)

65. Bond Orders of Homonuclear Diatomic Molecules
Peter Politzer
The Journal of Chemical Physics (1969-03-15) <https://doi.org/d28qgt>
DOI: [10.1063/1.1671457](https://doi.org/10.1063/1.1671457)

66. Covalent bond orders in the topological theory of atoms in molecules
Jerzy Cioslowski, Stacey T. Mixon
Journal of the American Chemical Society (1991-05) <https://doi.org/fd6w4x>
DOI: [10.1021/ja00011a014](https://doi.org/10.1021/ja00011a014)

67. Natural resonance theory: II. Natural bond order and valency
E. D. Glendening, F. Weinhold
Journal of Computational Chemistry (1998-04-30) <https://doi.org/bcrs8w>
DOI: [10.1002/\(sici\)1096-987x\(19980430\)19:6<610::aid-jcc4>3.0.co;2-u](https://doi.org/10.1002/(sici)1096-987x(19980430)19:6<610::aid-jcc4>3.0.co;2-u)

68. Modified valence indices from the two-particle density matrix
Roman F. Nalewajski, Janusz Mrozek

International Journal of Quantum Chemistry (1994-08-05) <https://doi.org/d7xjk6>
DOI: [10.1002/qua.560510403](https://doi.org/10.1002/qua.560510403)

69. Reaching the Maximum Multiplicity of the Covalent Chemical Bond

Björn O. Roos, Antonio C. Borin, Laura Gagliardi

Angewandte Chemie International Edition (2007-02-19) <https://doi.org/c5fht5>

DOI: [10.1002/anie.200603600](https://doi.org/10.1002/anie.200603600) · PMID: [17225237](#)

70. Chemical bonding in hypervalent molecules. The dominance of ionic bonding and negative hyperconjugation over d-orbital participation

Alan E. Reed, Paul v. R. Schleyer

Journal of the American Chemical Society (1990-02) <https://doi.org/bkgqqb>

DOI: [10.1021/ja00160a022](https://doi.org/10.1021/ja00160a022)

71. Spatial localization of the electronic pair and number distributions in molecules

R. F. W. Bader, M. E. Stephens

Journal of the American Chemical Society (1975-12) <https://doi.org/bt73sh>

DOI: [10.1021/ja00859a001](https://doi.org/10.1021/ja00859a001)

72. Overlap populations, bond orders and valences for “fuzzy” atoms

I Mayer, P Salvador

Chemical Physics Letters (2004-01) <https://doi.org/bgrrjh>

DOI: [10.1016/j.cplett.2003.11.048](https://doi.org/10.1016/j.cplett.2003.11.048)

73. The MMP2 calculational method

Joseph T. Sprague, Julia C. Tai, Young Yuh, Norman L. Allinger

Journal of Computational Chemistry (1987-07) <https://doi.org/dpjxnn>

DOI: [10.1002/jcc.540080504](https://doi.org/10.1002/jcc.540080504)

74. Molecular mechanics (MM3) calculations on conjugated hydrocarbons

Norman L. Allinger, Fanbing Li, Liqun Yan, Julia C. Tai

Journal of Computational Chemistry (1990-08) <https://doi.org/fs344c>

DOI: [10.1002/jcc.540110709](https://doi.org/10.1002/jcc.540110709)

75. Molecular mechanics (MM4) vibrational frequency calculations for alkenes and conjugated hydrocarbons

Neysa Nevins, Norman L. Allinger

Journal of Computational Chemistry (1996-04) <https://doi.org/ftx79r>

DOI: [10.1002/\(sici\)1096-987x\(199604\)17:5/6<730::aid-jcc9>3.0.co;2-v](https://doi.org/10.1002/(sici)1096-987x(199604)17:5/6<730::aid-jcc9>3.0.co;2-v)

76. Fast, efficient generation of high-quality atomic charges. AM1-BCC model: II. Parameterization and validation

Araz Jakalian, David B. Jack, Christopher I. Bayly

Journal of Computational Chemistry (2002-10-18) <https://doi.org/cktk6g>

DOI: [10.1002/jcc.10128](https://doi.org/10.1002/jcc.10128) · PMID: [12395429](#)

77. openforcefield/openforcefield

Open Force Field Initiative

(2020-01-29) <https://github.com/openforcefield/openforcefield>

78. Atom Type Independent Modeling of the Conformational Energy of Benzylic, Allylic, and Other Bonds Adjacent to Conjugated Systems

Candide Champion, Stephen J. Barigye, Wanlei Wei, Zhaomin Liu, Paul Labute, Nicolas Moitessier

79. The Open Force Field 1.0 small molecule force field, our first optimized force field (codename “Parsley”)

David Mobley Chodera Yudong Qiu, Simon Boothroyd, Lee-Ping Wang, and John (2019-10-10) [//openforcefield.org/news/introducing-openforcefield-1.0/](https://openforcefield.org/news/introducing-openforcefield-1.0/)

80. OpenMM 7: Rapid development of high performance algorithms for molecular dynamics

Peter Eastman, Jason Swails, John D. Chodera, Robert T. McGibbon, Yutong Zhao, Kyle A. Beauchamp, Lee-Ping Wang, Andrew C. Simmonett, Matthew P. Harrigan, Chaya D. Stern, ... Vijay S. Pande

PLOS Computational Biology (2017-07-26) <https://doi.org/gbppkv>

DOI: [10.1371/journal.pcbi.1005659](https://doi.org/journal.pcbi.1005659) · PMID: [28746339](https://pubmed.ncbi.nlm.nih.gov/28746339/) · PMCID: [PMC5549999](https://pubmed.ncbi.nlm.nih.gov/PMC5549999/)

81. openforcefield/openforcefields: Version 1.0.0 “Parsley”

Jeff Wagner

Zenodo (2019-10-12) <https://doi.org/ggjgjk>

DOI: [10.5281/zenodo.3483227](https://doi.org/10.5281/zenodo.3483227)

82. QUACPAC | Tautomer / Protomer Enumeration & Charge Assignment | Database Preparation

OpenEye Scientific Software

<https://www.eyesopen.com/quacpac>

83. OMEGA | Multi-Conformer Structure Databases | Database Preparation | OpenEye Scientific

OpenEye Scientific Software

<https://www.eyesopen.com/omega>

84. The NumPy Array: A Structure for Efficient Numerical Computation

Stéfan van der Walt, S Chris Colbert, Gaël Varoquaux

Computing in Science & Engineering (2011-03) <https://doi.org/d8k4p9>

DOI: [10.1109/mcse.2011.37](https://doi.org/10.1109/mcse.2011.37)

85. bashtage/arch: Release 4.8.1

Kevin Sheppard, Stanislav Khrapov, Gábor Lipták, Rob Capellini, Esvhd, Hugle, JPN, Xavier RENE-CORAIL, Michael E. Rose, Jbrockmendel

Zenodo (2019-03-28) <https://doi.org/ggjfwh>

DOI: [10.5281/zenodo.2613877](https://doi.org/10.5281/zenodo.2613877)

86. RDKit <http://www.rdkit.org/>

Supporting Information

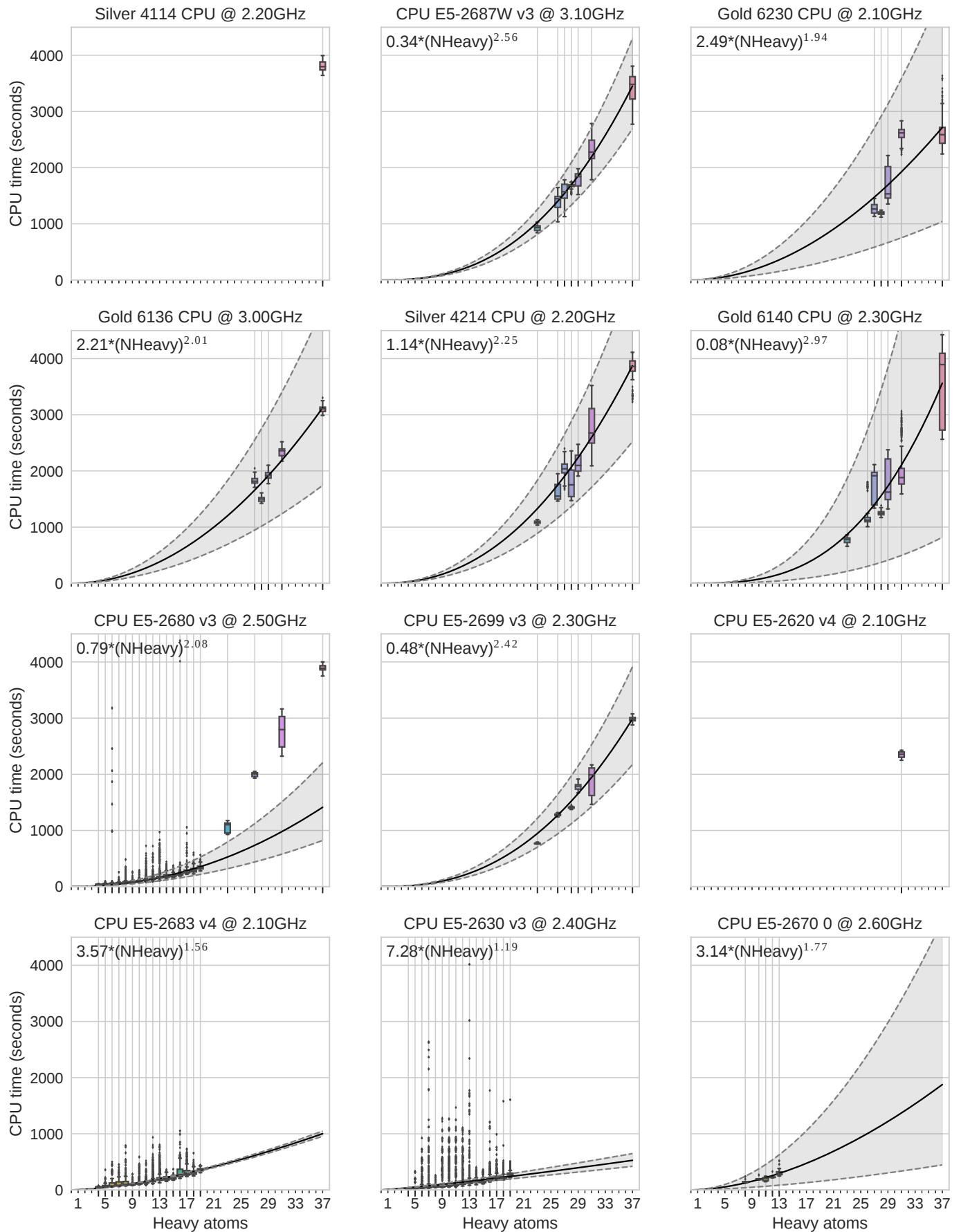


Figure 15: QC gradient evaluations scale similarly on various processor CPU time (wall time * nthreads) for one gradient evaluation vs. heavy atoms in molecules. All CPUs shown in this figure are Intel(R) Xeon(R)

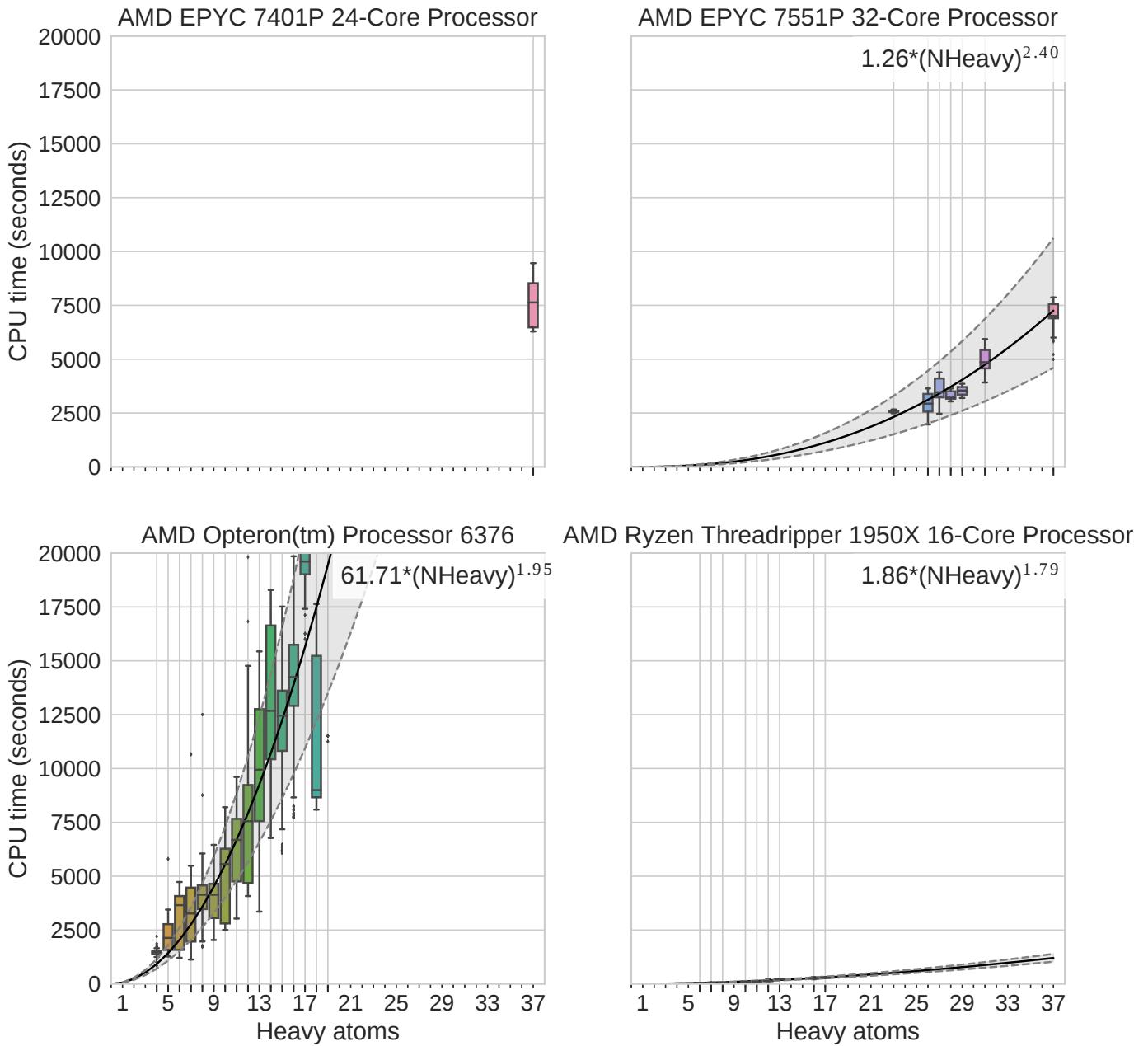


Figure 16: QC gradient evaluations scale similarly on various processor Same as Figure 15 but on AMD processors

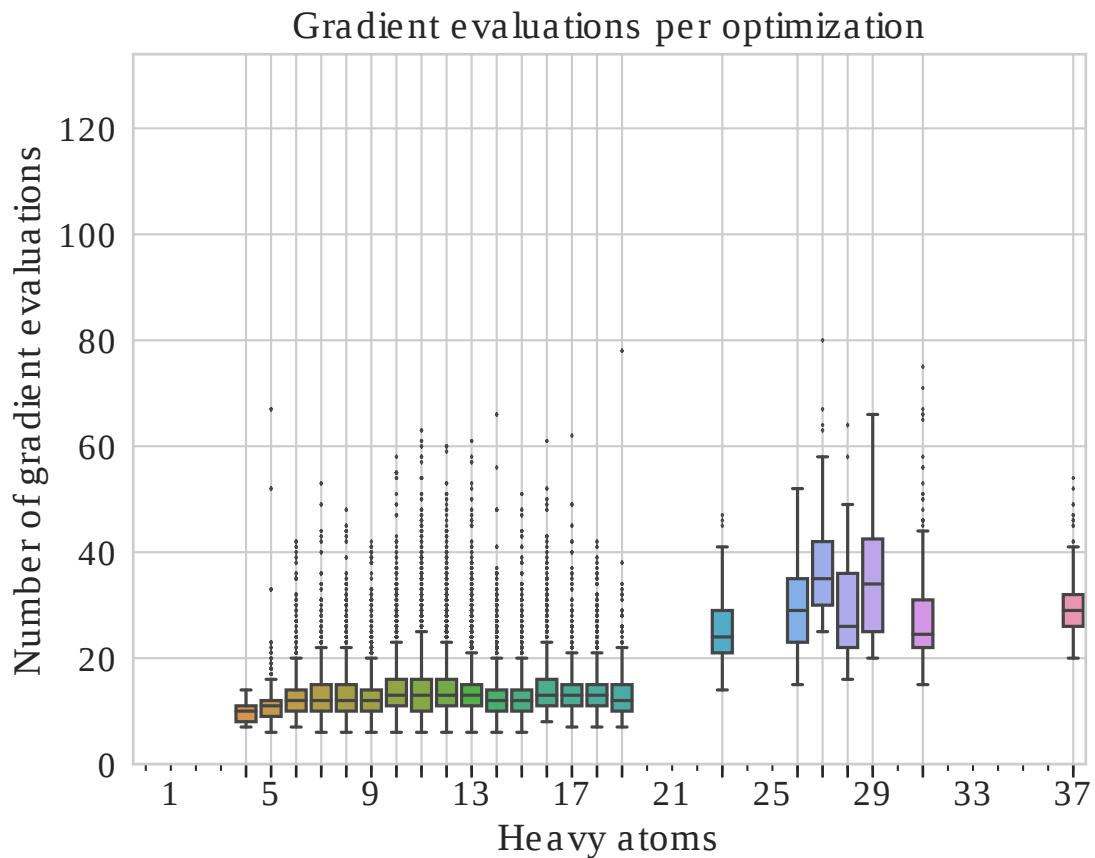


Figure 17: Distributions of number of gradient evaluations per optimizations for different size molecules The number of gradient evaluations per optimization depends on many factors such as initialization and tolerance, but there is also a slight dependency on molecular size as shown in this figure

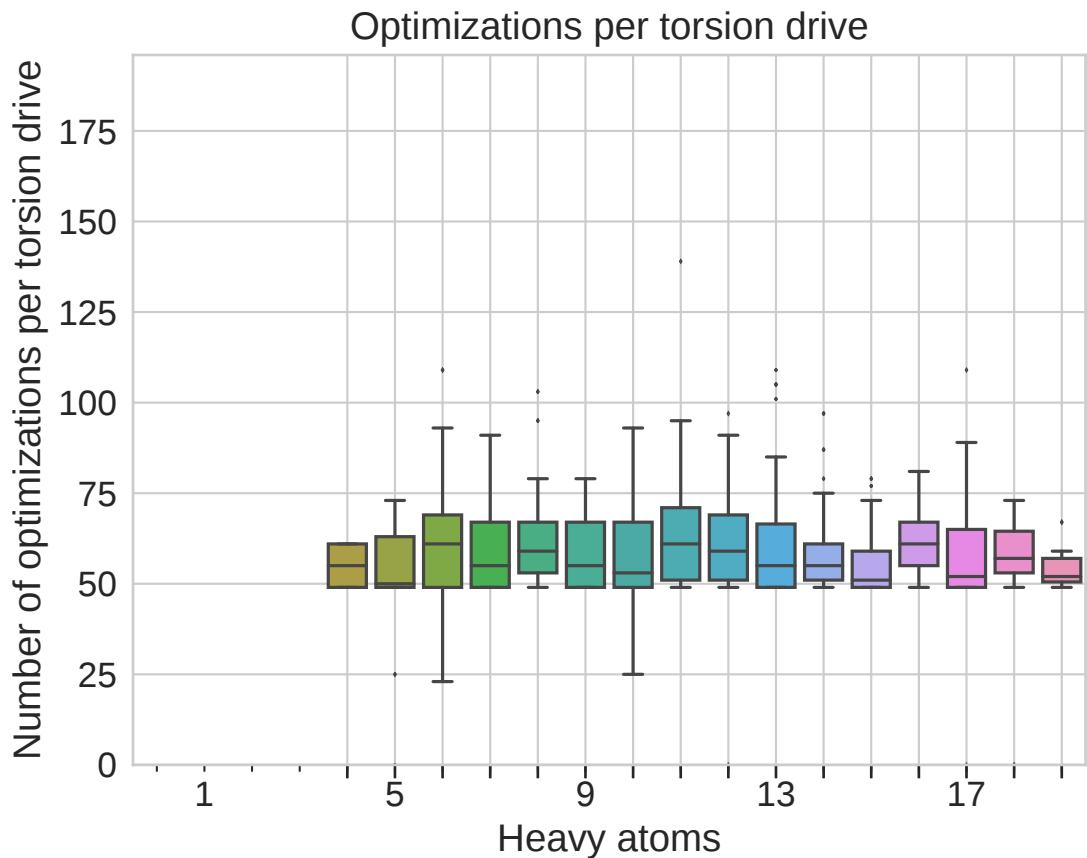


Figure 18: Distribution of optimizations per torsion drive This figures shows the distributions of optimizations per torsion drive when using wavefront propagation

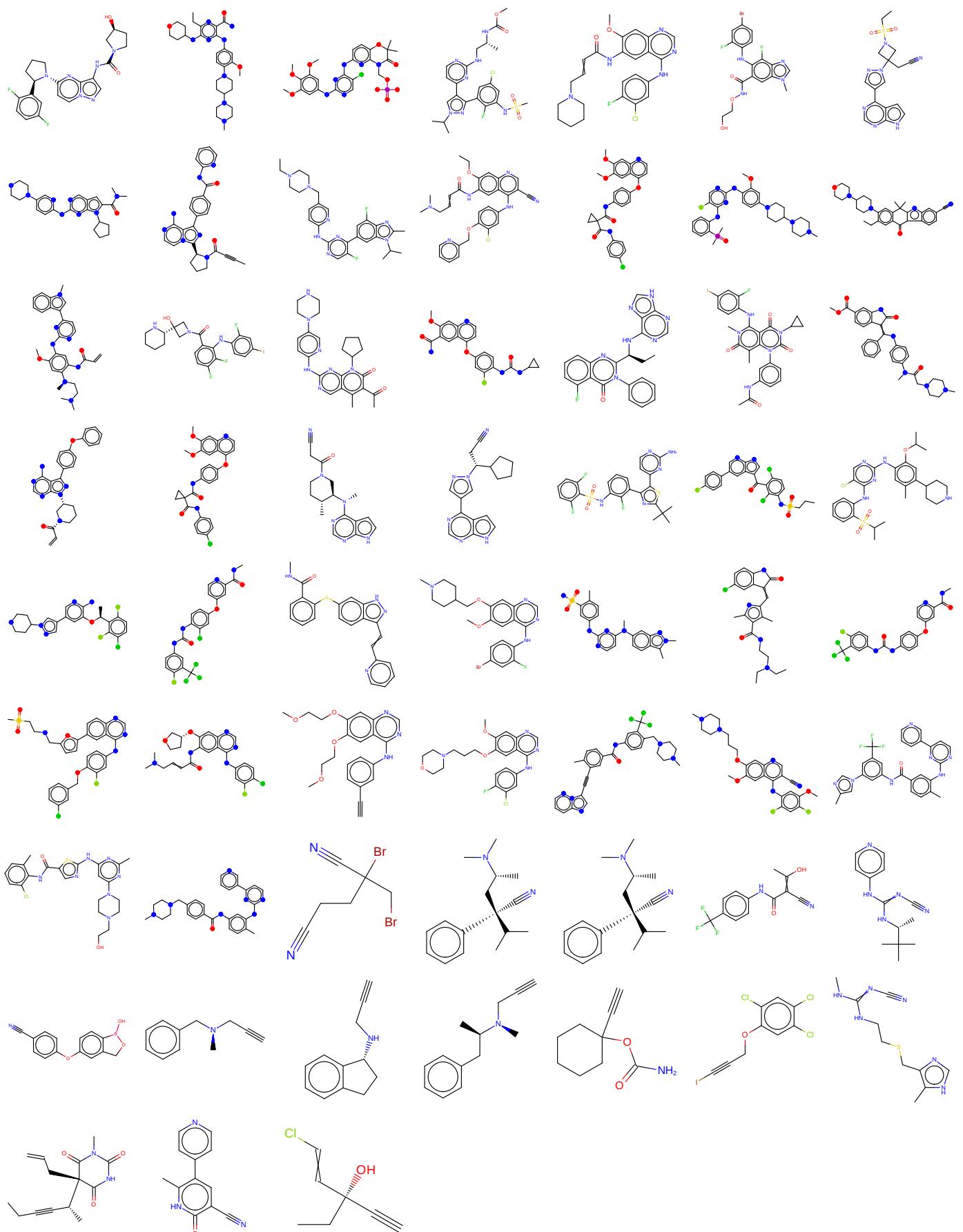


Figure 19: Druglike molecules used to calculate ELF10 AM1 WBOs This set of molecules were selected to cover bonds of different types and multiplicities

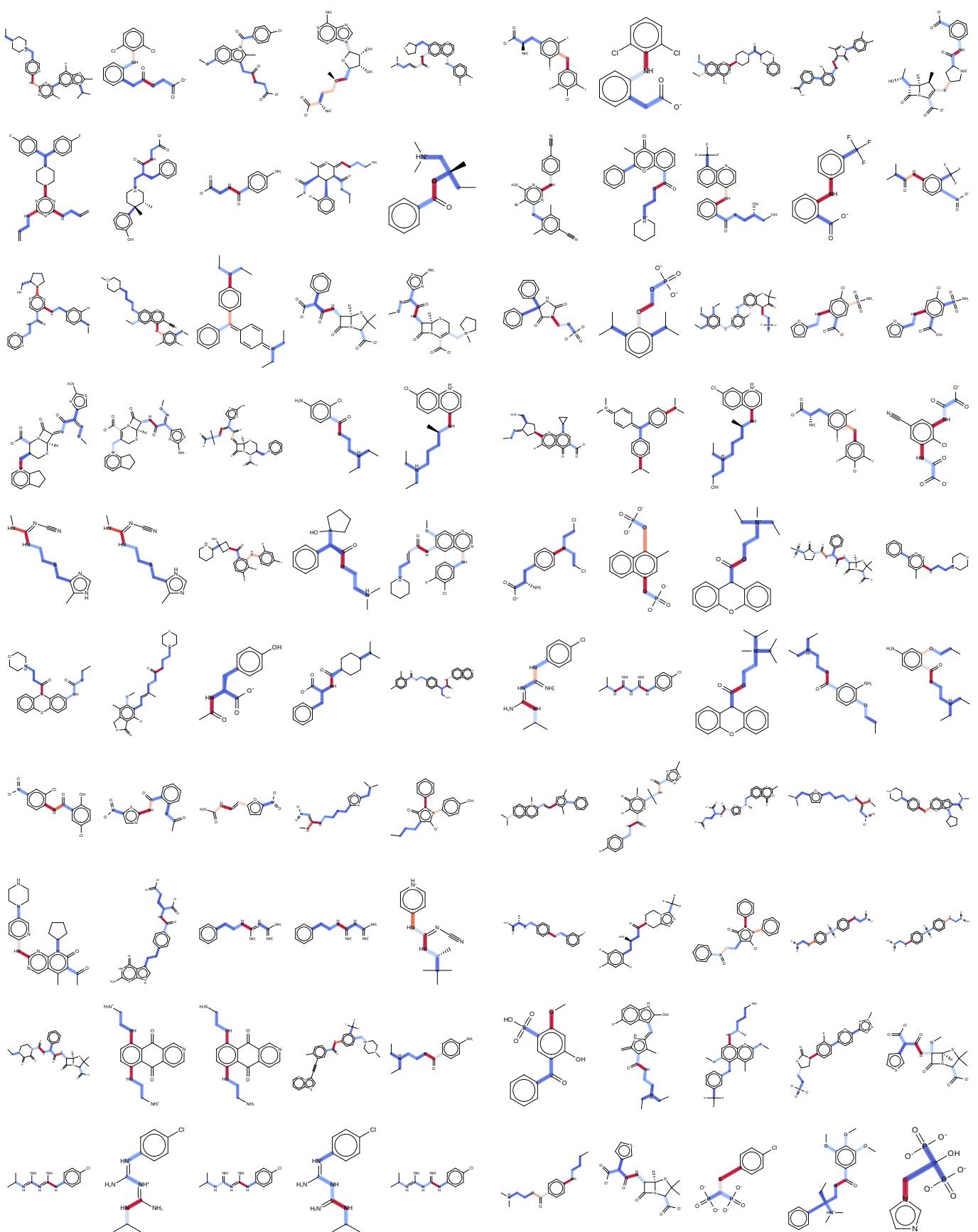


Figure 20: Validation set All molecules used in the validation set of fragmentation schemes. The bonds are highlighted by how sensitive they are to remote fragmentation. The redder bonds are more sensitive while the WBO distributions around the blue bonds do not change much with remote fragmentation.