Exploration of the Accessible Chemical Space of Acyclic Alkanes

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Saturated acyclic alkanes show steric strain if they are highly branched and, in extreme cases, fall apart rapidly at room temperature. Consequently, attempts to count the number of isomeric forms for a given molecular formula that neglect this physical consideration will inevitably overestimate the size of the available chemical space. Here we derive iterative equations to enumerate the number of isomers (both structural and optical are considered separately) for the alkane series that take into account the inherent instability of certain carbon skeletons. These function by filtering out certain substructures from the graph representation of the molecule which have been found to be thermodynamically unstable. We use these relations to report new estimates of the size of physically accessible chemical space for acyclic alkanes and show that for large molecules there are more isomers that are structurally disallowed than there are allowed.

1. INTRODUCTION

The familiar concepts of isomerism were developed in the 1860s,¹ when it was realized that there were different possible structural formulas for the same atomic composition. This realization provided crucial evidence for the validity of structural formulas and the existence of structured molecules. Since then various methods have been devised to calculate the precise number of possible isomers (both structural and stereochemical) for a given molecular formula.

The problem of the number of isomeric hydrocarbons of the methane series has concerned mathematicians and chemists since 1874. It was in this year that Cayley first formulated iterative equations to formally enumerate the number of isomers of the alkanes.² This has continued to be a topic of interest, particularly with Henze and Blair making several enumerations for alkanes and homologous sequences of alkane derivatives using recursion algorithms in 1931.³ Shortly afterward, motivated by this problem of chemical isomerism, Pólya⁴,⁵ introduced a powerful enumeration algorithm that used molecular symmetry, weighting factors, and generating functions.

More recent applications of isomer enumeration have been seen in the field of combinatorial chemistry. Various estimates have been proposed for the size of chemical space of small organic molecules, ranging from 10¹⁵–10²⁰⁰ compounds.⁶ Enumeration techniques are used to chart the size and composition of the small molecule chemical universe, from which the structural diversity of a combinatorial library can be quantified.⁷ Davies and Freyd have also considered the physical reality of the astronomical numbers involved in isomer enumeration—they calculated that C₁₆₇H₃₃₆ is the smallest alkane for which the number of stereoisomers exceeds the classical figure for the number of particles in the universe (~10⁸⁰).⁸

In many of these approaches to isomer enumeration it is assumed that all conceivable molecules can exist. However, alkanes may be very strained, even without cyclic structures, as multiple branching pushes side chains close together. Neighboring quaternary centers are likely to lead to highly strained structures.⁹ Klein¹⁰,¹¹ has considered self-avoidingly embedding alkanes onto a tetrahedral (diamond) lattice, thus simulating the sp³ bond angles of unsaturated hydrocarbons. The smallest acyclic alkane isomers which cannot fit onto the lattice are shown in Figure 1.

Any unsaturated hydrocarbon which cannot be fit on a diamond lattice is clearly very strained. However, tetra-tert-
established by Henze and Blair and Davies and Freyd to isomerism. We formally develop the mathematical relations that cannot be made. Never be made and suggested that C16 is the smallest alkane unstable. On this basis we made the prediction that C17 could that these two molecules are both thermodynamically dissociation of one molecule into two, strongly suggesting hermic (Figure 3). However, entropic effects will favor the C16 and C17 structures into radicals to be slightly endot- calculations suggest that the homolytic dissociation of both certain structures.

Analysis of the strain energy from molecular mechanics calculations was used to calculate the activation energy of the dissociation of highly branched alkanes into radicals. These results, summarized in Figure 2, suggest that tri-tert-butyl-isopropylmethane (C16H34) will spontaneously undergo thermolysis at −16 °C, while tetra-tert-butylmethane (C17H36) will undergo thermolysis at −145 °C. Additional DFT calculations suggest that the homolytic dissociation of both C16 and C17 structures into radicals is embedded in a solid argon or nitrogen matrix. However, in previous studies and in this paper we concern ourselves with molecules that can be made and stored at room temperature.

In this paper we return to the classic example of alkane isomerism. We formally develop the mathematical relations established by Henze and Blair and Davies and Freyd to enumerate the number of isomeric hydrocarbons of the methane series, while excluding any isomers that contain the unstable highly branched C16 and C17 substructures. Thus our isomer count (both structural and stereochemical isomers are considered here) represents only those molecules which are realizable/ synthesizable at room temperature. We have implemented these new mathematical relations using the Java programming language, and we have enumerated the numbers of isomers for the alkanes up to C300H600. We then analyze the effects that these physical constraints have on the size of accessible chemical space for the saturated hydrocarbons, by comparison with a simple isomer counting scheme that takes no account of the inherent instability of certain structures.

2. ENUMERATION METHODOLOGY

In the standard approach to isomer enumeration each isomer is represented as a molecular graph. The graphs are

Figure 4. The iterative generation of alkyl radicals (rooted trees). Here three smaller alkyl radicals (containing a, b, and c carbon atoms) are joined to create a new radical of size a+b+c+1. Radicals can also be zero-order, consisting of a single H-atom.

Studies within our group on acyclic alkane conformations led us to the question of the simplest acyclic alkane that cannot be made. Certainly, extraordinary molecules have never been synthesized and theoretical studies have shown it must be extremely strained.

"hydrogen deleted" in the sense that only carbon atoms and C—C bonds are explicitly represented. Thus structural isomers of the alkane CnH2n+2 can be uniquely represented by n vertices, each of which is connected to one, two, or three edges (signifying a primary, secondary, or tertiary carbon atom, respectively). An enumeration is first made of alkyl radicals or graph-theoretically speaking "rooted trees". These trees can be iteratively generated from three smaller rooted trees (i.e., alkyl radicals) and so on as shown below in Figure 4.

The number of structural isomers R(n) for a given rooted tree of size n can be calculated from the iterative relations:

\[ R(0) = 1 \]
\[ R(1) = 1 \]
\[ R(n+1) = \sum_{a+b+c=n} R(a)R(b)R(c) + \sum_{2a+b+c=n} \left( \frac{R(a+1)}{2} \right) R(c) + \sum_{3a+b+c=n} \left( \frac{R(a+2)}{3} \right) \]  

The three terms correspond to the cases of no pairs, one pair, and three of a kind with regard to the size of smaller radicals a, b, and c. The binomial coefficient

\[ \binom{m + k - 1}{k} = \frac{m!}{k!(m-k)!} \]

is equal to the number of ways of choosing k things, each of one of m types with repeated types allowed.

The various rooted trees thus obtained can be joined together to form alkanes as shown in Figure 5. Any graph representation with n vertices has a unique "centroid", defined as either an edge that when removed yields two rooted trees with n/2 vertices or as a vertex that when removed yields a set of trees each with fewer than n/2 vertices. A centroid edge is only possible for graphs with an even number of vertices. Every structural isomer can be constructed in this fashion, so the number of structural isomers T(n) for the alkane CnH2n+2 can be calculated from the iterative relation
The first term counts the number of isomers with a centroid edge. The remaining terms count the number of isomers with a centroid vertex and correspond to the cases of no pairs, one pair, two pairs, three of a kind, and four of a kind with regards to rooted tree size.

The enumeration method outlined thus far counts the number of structural isomers for the alkane series. It can be further modified to count all possible stereoisomers for the alkane series, utilizing the notion of a stereotree. A stereotree is formed from smaller rooted trees as above; however, the ordering of rooted trees is important: two such orderings are equivalent if one is obtainable from the other by an even permutation. The formulas are modified as follows:

\[
T(n) = \sum_{2a=n} \left( \binom{R(a) + 1}{2} + \sum_{a+b+c+d=n-1}^{} R(a)R(b)R(c)R(d) + \sum_{2a+b+c=d=n-1}^{} \binom{R(a)+1}{2}R(c)R(d) + \sum_{c+d=n/2}^{} \binom{R(a)+1}{2}R(c) + \sum_{4a=n-1}^{} \frac{R(a)+3}{4} \right) (3)
\]

is the number of ways of choosing a sequence of \( k \) things each of \( m \) types with repeated types allowed, where two such sequences are equivalent if one is obtainable from the other by an even permutation.

\[
T(n) = \sum_{2a=n} \left( \binom{R(a) + 1}{2} + \sum_{a+b+c+d=n-1}^{} R(a)R(b)R(c)R(d) + \sum_{2a+b+c=d=n-1}^{} \binom{R(a)+1}{2}R(c)R(d) + \sum_{c+d=n/2}^{} \binom{R(a)+1}{2}R(c) + \sum_{4a=n-1}^{} \frac{R(a)+3}{4} \right) (3)
\]

The number of isomers for the alkane series up to \( C_{2000}H_{4002} \) was calculated using these relations, and the results are summarized below in Table 1. (For larger numbers all digits are given in the Supporting Information). The asymptotic behavior for alkane enumerations approaching the many atom limit has been described in the work of Pólya, Otter, and more recently by Davies and Bytautas and Klein. It has shown that the behavior of \( R(n) \) and \( T(n) \) as \( n \) increases can be described by asymptotic forms, the simplest of which is of the form \( AB^n/n^2.21 \) In the limit of very large values of \( n \), the growth factor (i.e., \( R(n+1)/R(n) \)) is given by \( B \) in the above relationship. Applying the asymptotic analysis techniques of Otter to our data, we obtained values for \( A \) and \( B \). Rather than a least-squares regression, we used the logarithmic relationship derived in eq 7.

\[
\ln(T(n)) \times n^{2.5} = \ln(A) + n\ln(B)
\]
which can, in theory, be synthesized. We treat the two cases (C16 disallowed and C17 disallowed) separately to quantitatively study the effect that a more or less stringent condition has on the isomer count.

(i) C17 Elimination. An isomer is eliminated if the C17 (tetra-tert-butylmethane) substructure occurs at least once somewhere in the entire structure. In the enumeration methodology outlined above the occurrence of this substructure could arise (a) in the formation of the rooted trees or (b) as the rooted trees come together at the centroid edge or vertex, or a combination of both (Figure 6).

In addition to the previously defined alkyl radical generating function \( R(n) \), we define the terms \( T(n) \) to enumerate those radicals tertiary at their root (i.e. with three nonzero branches) and \( T(t) \) to enumerate those radicals tertiary at the root, with each adjacent site being quaternary. These values are given by the following relations:

\[
0 \leq n \leq 3, T(n) = 0
\]

\[
Tr(n + 1) = \sum_{0 < a < b < c, a + b + c = n} R(a)R(b)R(c) + \sum_{0 < a < b < c, 2a + c = n} R(a)R(b)R(c) + \sum_{0 < a < b < c, 2a + c = n} R(a)R(b)R(c)
\]

\[
0 \leq n \leq 12, T(n) = 0
\]

\[
Tt(n + 1) = \sum_{3 < a < b < c, a + b + c = n} Tt(a)Tr(b)Tr(c) + \sum_{3 < a < b < c, 2a + c = n} Tt(a)Tr(b)Tr(c) + \sum_{3 < a < b < c, 2a + c = n} Tt(a)Tr(b)Tr(c)
\]

Thus to avoid the scenario in Figure 6a trees which are allowed, herein enumerated as \( R'(n) \), must not connect a \( Tt \) radical with two nonzero radicals at any point in their iterative formation. The first term of the iteration (eq 9) represents the case for which three nonzero radicals of different size are joined together. In this case, none of these radicals must be a \( Tt \) branch so these are subtracted from the available number of radicals \( a, b, \) and \( c \). The second term has a zero radical (i.e., a hydrogen atom) so there are no restrictions. The next three terms represent the case of a pair (two radicals the same size) in which there are no, one, and two hydrogen radicals, respectively. The final term considers the scenario of three nonzero radicals of the same size.

\[
0 \leq n \leq 15, R'(n) = R(n)
\]

\[
R'(n + 1) = \sum_{a + b + c = n} (R'(a) - T'(a))R'(b) - T'(b)(R'(c) - T'(c)) + \sum_{b + c = n} R'(b)R'(c) + \sum_{0 < a < b < c, 2a + c = n} R'(a)R'(b)R'(c) + \sum_{0 < a < b < c, 2a + c = n} R'(a)R'(b)R'(c) + \sum_{2a = n} (R'(a) - T'(a) + 1)(R'(c) - T'(c)) + \sum_{2a = n} (R'(a) + 1) + R'(n) + \sum_{3a = n} (R'(a) - T'(a) + 2)
\]
where $T_t(n)$ and $T_t'(n)$ are the allowed versions of $T(n)$ and $T(n)$ respectively, constructed in an entirely analogous fashion but using allowed radicals only in their construction:

\[
T_t'(n + 1) = \sum_{0 \leq a < b < c \leq n} R'(a)R'(b)R'(c) + \sum_{0 \leq a < b < c \leq n} \left( \frac{R'(a) + 1}{2} \right) R'(c) + \sum_{3a = n} \left( \frac{R'(a) + 2}{3} \right)
\]

\[
T_t'(n + 1) = \sum_{3 \leq a < b < c \leq n} T_t'(a)T_t'(b)T_t'(c) + \sum_{3 \leq a < b < c \leq n} \left( \frac{T_t'(a) + 1}{2} \right) T_t'(c) + \sum_{3a = n} \left( \frac{T_t'(a) + 2}{3} \right)
\]

Turning to the formation of disallowed structures as branched trees are brought together (Figure 6b), we then proceed to enumerate alkane structures adapting the relationship in eq 3 to include only those isomers that are disallowed at the centroid, $N'(n)$. The first two terms describe a centroid edge, which connects at least one $T_t$ rooted tree with a tree which is tertiary or more branched (see Figure 6b(i)). All subsequent terms describe a centroid vertex. We account for the two disallowed scenarios in Figure 6b(ii) by summation of all centroids with four tertiary branched radicals and all centroids with at least one $T_t$ branched radical and three nonzero radicals. Care is taken to avoid double counting, so for the case of four tertiary radicals, we exclude the existence of $T_t$ radicals so the number of possible radicals for each branch is $T_t'(a)\cdot T_t'(a)$. The second term is obtained by subtracting all isomers which contain no $T_t$ radicals from all possible isomers to leave only those structures which contain at least one $T_t$ radical at the centroid. This process is repeated for the different scenarios which correspond to the cases of no pairs, one pair, two pairs, three of a kind, and four of a kind with regards to rooted tree size.

\[
N'(n) = \sum_{2a = n} \left( \frac{T_t'(a) + 1}{2} \right) + T_t'(a)(T_t'(a) - T_t'(a)) + \sum_{a < b < c < d \leq n} \left( T_t'(a) - T_t'(a))(T_t'(b) - T_t'(b))(T_t'(c) - T_t'(c))(T_t'(d) - T_t'(d)) \right)
\]

\[
+ \sum_{a < b < c < d \leq n} \left( R'(a)R'(b)R'(c)R'(d) - (R'(a) - T_t'(a))(R'(b) - T_t'(b))(R'(c) - T_t'(c))(R'(d) - T_t'(d)) \right)
\]

\[
+ \sum_{c<d=n+1} \left( \frac{T_t'(a) - T_t'(a) + 1}{2} \right) (T_t'(c) - T_t'(c))(T_t'(d) - T_t'(d)) \right)
\]

\[
+ \sum_{2a + c = n+1} \left( \frac{T_t'(a) - T_t'(a) + 1}{2} \right) (T_t'(c) - T_t'(c)) \right)
\]

\[
+ \sum_{2a + c = n+1} \left( \frac{T_t'(a) - T_t'(a) + 1}{2} \right) (T_t'(c) - T_t'(c)) \right)
\]

\[
+ \sum_{3a = n+1} \left( \frac{T_t'(a) - T_t'(a) + 3}{4} \right) + (T_t'(a) - T_t'(a) + 3) \right)
\]

(13)

Subtracting $N'(n)$ from the number of isomers with all possible centroids (similarly enumerated using only allowed rooted trees $R'(n)$) then yields the number of structural isomers which do not contain the C17 substructure anywhere in the molecule. An analogous procedure was followed to obtain values for the number of allowed stereoisomers (full details of the enumeration are given in the Supporting Information). The results are shown in Table 2.

For C20H42, 11 structural isomers are disallowed in our enumeration method. The strain energy of these disallowed isomers was evaluated by molecular mechanics. Our earlier studies suggest that a strain energy of about 300 kJ mol$^{-1}$ is sufficient to make the molecule unstable at room temperature.\textsuperscript{14} In all cases the strain energy of these isomers is greater than that calculated for the isolated C17 molecule (416.05 kJ mol$^{-1}$), and so these isomers are also likely to be unstable. By comparison, the strain energy of two isomers not containing the C17 substructure (but with some steric crowding) is significantly less than that of both the C17 molecule and the C16 molecule (335 kJ mol$^{-1}$). We conclude that our counting scheme is correct to include these less-strained structures.

This problem of steric overcrowding severely limits the number of possible isomers that could exist as $n$ becomes large. Since the number of “virtual” isomers (i.e., all those that can be drawn on paper without physical consideration, as calculated
will approach infinity as \(b\) approach zero even though the number of possible isomers
the ratio of possible to impossible isomers will asymptotically increase without any substructure filtering) increases exponentially, 

\[
\text{Table 3. Numbers of Alkane Isomers Which Are Allowed, i.e., Do Not Contain the Unstable C16 Substructure Anywhere in the Molecule, Also Expressed as a Fraction of the Total Possible Number Neglecting Physical Considerations}
\]

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<th>(n)</th>
<th>(N_{\text{allowed}})</th>
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<th>(\frac{N_{\text{allowed}}}{N_{\text{disallowed}}})</th>
<th>(N_{\text{allowed}})</th>
<th>(\frac{N_{\text{allowed}}}{N_{\text{possible}}})</th>
<th>(\frac{N_{\text{allowed}}}{N_{\text{disallowed}}})</th>
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(ii) C16 Elimination. An isomer is eliminated if the C16 (tri-tert-butyl-isopropyl)methane substructure occurs at least once somewhere in the entire structure. The recursion relations in this case are significantly more complex than for C17 since the undesirable substructure, being smaller, can arise in the carbon skeleton in many more ways. As before, the undesirable substructure could arise (a) in the formation of the rooted trees or (b) as the rooted trees come together at the centroid edge or vertex, or a combination of both, shown in Figure 11.

In addition to previously defined \(R(n)\), \(T(n)\), and \(T(n)\), here we need to define the term \(S(n)\) to enumerate secondary radicals and the term \(T(n)\) which is used to enumerate those radicals which are tertiary at their root and where each branch also possesses a tertiary root or one where one branch is secondary and the other two are tertiary (i.e., \(T(n)\) is a subgroup of \(T(n)\)). These values are obtained from the following recursion relations:

\[
S(n + 1) = \sum_{b+c=n}^{R(b)R(c)} + \sum_{2a=n} \left( \frac{R(a) + 1}{2} \right) \quad (14)
\]

\[
0 \leq n \leq 12, T(n) = 0
\]
are more ways in which it can arise in the carbon skeleton. Structural isomers which do not contain the C16 substructure can arise from the connection of a edge (Figure 10b(i)), the disallowed C16 substructure can arise in one of three ways: (i) A to avoid overlap with the previous case. (iii) Three tertiary radicals that always form the disallowed C16 skeleton upon connection with all possible centroids (similarly enumerated using only allowed radicals, \(R'(n)\)), these substructures are excluded to avoid overlap with the previous case. (iii) Three tertiary radicals that will not lead to an undesirable alkane structure.

We then proceed to enumerate alkane structures adapting the relationship in eq 3 to include only those isomers that are disallowed at the centroid, \(N'(n)\), which can occur in any of five ways shown in Figure 10b. In the case of a centroid edge (Figure 10b(i)), the disallowed C16 substructure can arise from the connection of a Ts and a Tr radical or of a TT and an S radical. In the case of a centroid vertex, the C16 substructure can arise in one of three ways: (i) A Ts radical and three nonzero radicals are brought together. (ii) A TT radical and two nonzero radicals are brought together. One of the four branches around the centroid must be zero to avoid overlap with the previous case. (iii) Three tertiary and one secondary or four tertiary radicals are brought together. These must not be Ts radicals to avoid overlap with the first case.

These structures are enumerated for the different scenarios which correspond to a pair of no pairs, one pair, two pairs, three of a kind, and four of a kind with regards to rooted tree size. Subtracting this total from the number of isomers with all possible centroids (similarly enumerated using only allowed rooted trees \(R'(n)\)) then yields the number of structural isomers which do not contain the C16 substructure anywhere in the molecule. The results are shown in Table 3. For full details of the recursion relation refer to the Supporting information.

Since the C16 substructure is smaller than the C17, there are more ways in which it can arise in the carbon skeleton.

\[
Ts(n + 1) = \sum_{a+b+c=n}^{3a<b<c} Tr(a)Tr(b)Tr(c)Tr(d) + \sum_{a+b+c=n}^{3a<b<c} (S + a)Tr(a)Tr(b)S(c) + \sum_{a+b+c=n}^{2a+b+c=n} \left( \frac{Ts(a) + 1}{2} \right) S(c) + S(a) + Ts(a) + Ts(b) + Ts(c) + \sum_{a+b+c=n}^{3a=n} \left( \frac{Tr(a) + 2}{3} \right) + \frac{Tr(a) + 1}{2} S(a) \right) + \sum_{2a+b+c=n}^{3a=n} \left( \frac{R'(a) - Ts'(a) + 1}{2} \right) \left( R'(c) - Ts'(c) \right) + \sum_{a+b+c=n}^{3a=n} \left( R'(a) - Tr(a) + 1 \right) + R'(n) + \sum_{a+b+c=n}^{3a=n} \left( R'(a) - Ts'(a) + 2 \right) \right) (16)
\]

Unsurprisingly the proportion of disallowed structures increases more rapidly, and by the time C\(_{2000}H_{4002}\) is reached the ratio of allowed to disallowed structures is 2.4:1. The decline in the proportion of allowed isomers is shown in Figure 12.

4. RESULTS AND DISCUSSION

Assuming all isomer counts were of the form \(AB^n/n^{2.5}\) we constructed logarithmic plots using data in the range \(n = 100\) to \(n = 2000\) as detailed earlier in the text. These plots yielded values of \(A\) and \(B\) for our “allowed” isomer counts (i.e., those that filter out the C16 and C17 undesired substructures), and in every case the linear correlation coefficient was greater than 0.999 so the behavior is described well by the above form. For the numbers of allowed isomers smaller values of \(B\) are obtained than for the total number of “paper” isomers—this is because their growth is slowed by the increasing proportion of disallowed isomers. Hence disallowing the C16 substructure leads to a greater reduction in the growth factor as the proportion of disallowed isomers is larger. From the values of \(A\) and \(B\) thus obtained (Figure 13), we can easily calculate and make predictions of the ratio of allowed:total number of “virtual” (i.e., unfiltered) isomers or equally the ratio of allowed: disallowed isomers.

\[
\frac{T_{\text{allowed}}}{T_{\text{paper}}} = \frac{A_d}{A_p} \left( \frac{B_d}{B_p} \right)^n
\]

(17)

\[
\frac{T_{\text{allowed}}}{T_{\text{paper}}} = \left( \frac{1}{\frac{A_d}{A_p} \left( \frac{B_d}{B_p} \right)^n} - 1 \right)
\]

(18)

Since the growth factor of the total number is greater than for the allowed isomer counts (\(B_d > B_p\)), the denominator of the ratio of allowed to disallowed isomers tends to infinity.
as \( n \) increases, and therefore this ratio tends asymptotically to zero. That is to say, it becomes increasingly difficult to make alkanes as they get larger, since the number of unfavorable structures grows faster than does the number of favorable structures. This argument that the number of crowding-constrained structures becomes an ever smaller fraction of the total theoretical count has been made qualitatively by Klein,\(^{10}\) which as is shown in Figure 14, ratios calculated by fitting to asymptotic relations are in excellent agreement with the explicitly calculated values. Hence we can use these relations to extrapolate beyond the upper limit of our data to predict that the first alkane for which there are equal numbers of allowed and disallowed structural isomers is \( \text{C}_{3783} \text{H}_{7568} \) if \( \text{C}_{16} \) is disallowed or \( \text{C}_{33477} \text{H}_{66956} \) if \( \text{C}_{17} \) is disallowed.

5. CONCLUSION

If an acyclic alkane structure with a molecular weight above 53 kD is randomly generated, it is more likely than not that the substance cannot exist. Our enumeration strategy overestimates the odds of the successful construction of such a structure, because we consider only a \( \text{C}_{16} \) forbidden unit. There are many other units that will also be forbidden which do not include this substructure. For example, if the methyl groups on propane are substituted with isopropyl groups, and this process is repeated several times on the new structure, a different impossible structure will result (Figure 15). However, the smallest unstable structure is likely to be rather larger in these series. \( \text{C}_{31} \text{H}_{64} \) in Figure 15 is very likely to be stable on the basis of its low strain energy, although \( \text{C}_{63} \text{H}_{128} \) is unlikely to exist. Figure 14 shows that the number of structures excluded by disallowing the \( \text{C}_{16} \) unit is much greater than the number excluded by disallowing the \( \text{C}_{17} \) unit. While we cannot enumerate all impossible substructures, they must all be larger than \( \text{C}_{17} \) and so will exclude a smaller fraction of the possible isomers.
suggest that C_{3783}H_{7568} is the smallest alkane for which a reported the smallest alkane that cannot be made. We now C_{2000}H_{4002} the odds of successfully generating an isomer diminish as the number of carbon atoms increase. For the smallest such alkane may be smaller still. The odds randomly chosen isomer has a less than even chance of C_{1000}H_{2002} and C_{2000}H_{4002} including all digits and the full that of being struck by lightning. 25 It might be suggested this paper, along with the Java source code, is available at http://www-jmg.ch.cam.ac.uk/data/isomercount/

These calculations demonstrate that chemical space is restricted for larger molecules. In our earlier work, 15 we reported the smallest alkane that cannot be made. We now suggest that C_{733}H_{568} is the smallest alkane for which a randomly chosen isomer has a less than even chance of actually existing, because of the need to exclude C_{16} substructures. If all impossible substructures were considered, the smallest such alkane may be smaller still. The odds diminish as the number of carbon atoms increase. For C_{3000}H_{4002} the odds of successfully generating an isomer are 30:1 against. For C_{3000}H_{10002} the odds are less than that of being struck by lightning. 25 It might be suggested that these are moving toward molecules that are so big they cannot be made.

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Supporting Information Available: Isomer counts for C_{1000}H_{2002} and C_{2000}H_{4002} including all digits and the full algorithm used to enumerate isomers filtering the C_{16} substructure. This material is available free of charge via the Internet at http://pubs.acs.org.

REFERENCES AND NOTES

Figure 15. Isomers for which the successive addition of isopropyl groups leads to impossible structures. Strain energies of more than 300 kJ mol^{-1} are likely to be unstable.

An applet which performs the enumerations outlined in this paper, along with the Java source code, is available at http://www-jmg.ch.cam.ac.uk/data/isomercount/


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