

Topic II

Symmetry and Group Theory in Chemistry

II.1 Module 3: Symmetry Elements and Operations

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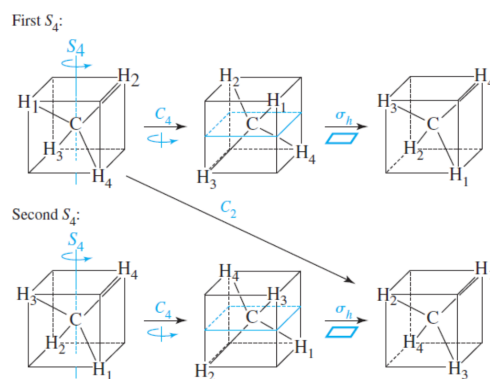
- He will upload lecture slides in advance in the future.
- An object is symmetric if one part is the same as other parts.
- The symmetry of discrete objects is described using **point symmetry**.
- **Point groups** (~ 32 for molecules) provide us with a way to indicate the symmetry unambiguously.
 - These have symmetry about a single point at the center of mass of the system.
- Extended objects (e.g., crystals) have **translational symmetry** described by **Space groups**^[1] (230 total).
- Readings: Miessler et al. (2014) Chapter 4 and https://en.wikipedia.org/wiki/Molecular_symmetry.
- **Symmetry elements**: Geometric entities about which a **symmetry operation** can be performed. In a point group, all symmetry elements must pass through the center of mass (the point).
- **Symmetry operation**: The action that produces an object identical to the initial object.

Element	Operation
Identity, E	nothing
Rotation axis, C_n	n -fold rotation
Improper rotation axis, S_n	n -fold improper rotation
Plane of symmetry, σ	Reflection
Center of symmetry, i	Inversion

- **Identity**: Does nothing to the object, but is necessary for mathematical completeness.
- **n -fold rotation**: A rotation of $360^\circ/n$ about the C_n axis ($n \in [1, \infty)$).
 - In H_2O , there is a C_2 axis, so we can perform a 2-fold (180°) rotation to get the same molecule.
 - Remember, because of quantum mechanical properties, the hydrogens are indistinguishable so when we rotate it 180° , we cannot tell it apart from the unrotated molecule.
 - Rotations are considered positive in the counterclockwise direction.

¹Not covered in this course.

- Each possible rotation operation is assigned using a superscript integer m of the form C_n^m . m is the number of sequential applications.
- The rotation $C_n^n \equiv E$ is equivalent to the identity operation (nothing is moved).
- Linear molecules have an infinite number of rotational options C_∞ because any rotation on the molecular axis will give the same arrangement.
- **Principal axis:** The highest order rotation axis.
 - By convention, the principal axis is assigned to the z -axis if we are using Cartesian coordinates.
- **Reflection:** Exchanges one half of the object with the reflection of the other half.
- **Vertical mirror plane:** A mirror plane that contains the principal axis. Denoted by σ_v .
- **Horizontal mirror plane:** A mirror plane that is perpendicular to the principal axis. Denoted by σ_h .
- **Dihedral mirror plane:** A special type of σ_v that is between sides or planes. Denoted by σ_d .
 - For example, we might have vertical mirror planes in the xz - or yz -planes. In this case, the dihedral planes would contain the lines $y = \pm x$.
- Two successive reflections are equivalent to the identity operation.
- **Inversion:** Every part of the object is reflected through the inversion center, which must be at the center of mass of the object.
 - $(x, y, z) \xrightarrow{i} (-x, -y, -z)$.
- **n -fold improper rotation:** This operation involves a rotation of $360^\circ/n$ followed by a reflection perpendicular to the axis. It is a single operation and is labeled in the same manner as “proper” rotations. Also known as S_n^m , **rotation-reflection operation**.

Figure II.1: Methane's S_4 symmetry.

- Methane has S_4 symmetry.
- Note that $S_1 \equiv \sigma_h$, $S_2 \equiv i$, and sometimes $S_{2n} \equiv C_n$. In methane, for example, $S_4^2 \equiv C_2$.
- Applied to a tetrahedron (triangular prism), is a good example.
- If n is even, we have n unique operations. There should be $C_{n/2}$.
- If n is odd, we have $2n$ unique operations. There should be C_n and σ_h .
- The absence of an S_n axis is the defining symmetry property of **chiral** molecules.
 - Formerly, we learned that chiral molecules should not have mirror planes and inversion centers.
 - Rigorously, chiral molecules must not have any improper rotation axes.

II.2 Module 4: Symmetry Point Groups

- Identifying the point groups:
 - Determine if the symmetry is special (e.g., octahedral).
 - Determine if there is a principal rotation axis.
 - Determine if there are rotation axes perpendicular to the principal axis.
 - Determine if there are mirror planes.
 - Assign point groups.
- High symmetry and low symmetry groups are the most difficult to identify.
- High symmetry:
 - Perfect tetrahedral (T_d), e.g., P_4 and CH_4 .
 - Perfect octahedral (O_h), e.g., SF_6 .
 - Perfect icosahedral (I_h), e.g., C_{60} and $B_{12}H_{12}^{2-}$.

- Low symmetry:

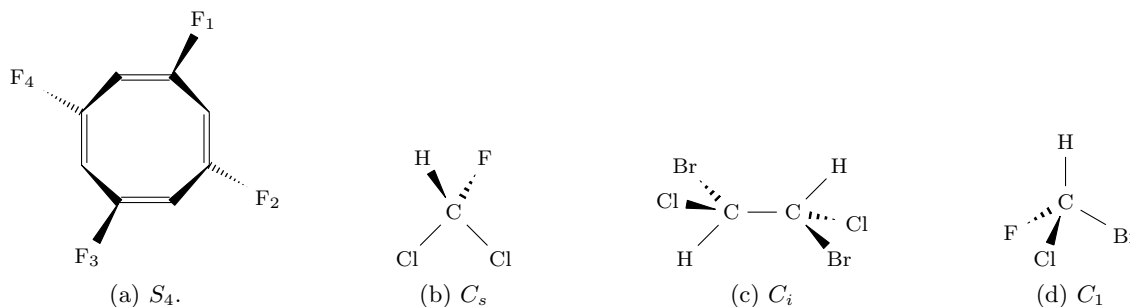


Figure II.2: Low symmetry point groups.

- Only an improper axis: S_n .
- Only a mirror plane: C_s .
- Only an inversion center: C_i .
- No symmetry: C_1 .
- C_n groups:
 - Only a C_n axis. Note that conformation is important.
- C_{nh} groups have a C_n axis and a σ_h reflection plane (such as $B(OH)_3$).
 - H_2O_2 has C_{2h} symmetry.
- All symmetry elements are listed in the top row of the corresponding characters table (Appendix C in Miessler et al. (2014)).
- C_{nv} groups have a C_n axis and a σ_v reflection plane.
 - NH_3 has C_{3v} symmetry.
 - CO has $C_{\infty v}$ symmetry since there are an infinite number of both C_n axes and σ_v mirror planes.
- D_{nh} groups: A C_n axis, n perpendicular C_2 axes, and a σ_h reflection plane.

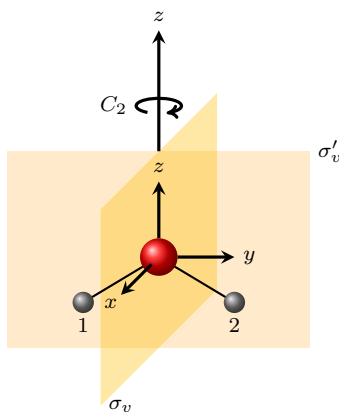
- BH_3 has D_{3h} symmetry.
- A square prism has D_{4h} symmetry.
- CO_2 has $D_{\infty h}$ symmetry.
- D_n groups: A C_n axis, n perpendicular C_2 axes, and no mirror planes.
 - A 3-bladed propeller has D_3 symmetry.
- D_{nd} groups: A C_n axis, n perpendicular C_2 axes, and a σ_d .
 - Ethane in the staggered conformation has D_{3d} symmetry.
- Local symmetry:
 - Sometimes, rigorous math analysis needs to be adjusted to physical reality.
 - If a cyclopentadienyl ring (Cp group) is bonded through the center to $\text{Mn}(\text{CO})_3$, this molecule has only C_s symmetry.
 - However, spectroscopically, there is fast rotation about the Mn–Cp bond. This means that the $\text{Mn}(\text{CO})_3$ fragment exhibits pseudo- C_{3v} symmetry while the C_5H_5 ligand exhibits pseudo- C_{5v} symmetry.
 - Often, the absolute symmetry of a molecule is very low, but the interactions are far away from the centers of interest, and do not perturb them significantly.
 - If we have platinum as a central atom bonded to two chlorines and two $\text{P}(\text{Et})_3$ groups, this molecule technically has C_1 symmetry due to the orientations of atoms within R groups (staggered), but its IR spectrum is characteristic of highly symmetric species (D_{2h}).
- Practice with identifying point groups: Otterbein University symmetry gallery ([link](#)).

II.3 Module 5: Group Theory 101

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- **Group:** A set of elements together with an operation that combines any two of its elements to form a third element satisfying four conditions called the group axioms.
- **Closure:** All binary products must be members of the group.
- **Associativity:** Associative law of multiplication must hold.
- **Identity:** A group must contain the identity operator.
- **Inverse:** Every operator must have an inverse.
- The integers with the addition operation form a group, for example.
- History:
 - Early group theory was driven by the quest for solutions of polynomial equations of degree 5 and above.
 - Early 1800s: Évariste Galois realized that the algebraic solution to a polynomial equation is related to the structure of a group of permutations associated with the roots of the polynomial (the **Galois group** of the polynomial).
 - [Link to Galois video here.](#)
 - 1920s: Group theory was applied to physics and chemistry.
 - 1931: It is often hard or even impossible to obtain a solution of the Schrödinger equation — however, a large part of qualitative results can be obtained by group theory. Almost all the rules of spectroscopy follow from the symmetry of a problem.

- We will use group theory for describing symmetry of molecules. We will use group theory to understand the bonding and spectroscopic features of molecules.
- For us, a group consists of a set of symmetry elements (and associated symmetry operations) that completely describes the symmetry of a molecule.
- **Order** (of a group): The total number of elements (i.e., symmetry operations) in the group. *Also known as **h**.*
- Rule 1: Closure.

Figure II.3: Symmetry elements for H₂O.

- H₂O is of the C_{2v} point group (refer to Figure II.3).
 - Symmetry operations: E , C_2 , $\sigma_{v(xz)}$, and $\sigma'_{v(yz)}$.
 - $\sigma_v \cdot C_2 = \sigma'_v = C_2 \cdot \sigma_v$.
 - The above property (order *does not* matter) shows that C_{2v} is an **Abelian group**.
- NH₃ is of the C_{3v} point group.
 - Symmetry operations: E , C_3^+ , C_3^- , σ_v , σ'_v , and σ''_v .
 - $\sigma''_v \cdot C_3 = \sigma_v$, but $C_3 \cdot \sigma''_v = C_3^- = C_3^2$.
 - The above property (order *does* matter) shows that C_{3v} is a **non-Abelian group**.
- Rule 2: Associativity.
 - H₂O is of the C_{2v} point group (refer to Figure II.3).

$\sigma'_v C_2 \sigma_v(1, 2)$	$= \sigma'_v C_2(2, 1)$	$\sigma'_v(C_2 \sigma_v)(1, 2)$	$= \sigma'_v E(1, 2)$	$(\sigma'_v C_2) \sigma_v(1, 2)$	$= \sigma_v \sigma_v(1, 2)$
$= \sigma'_v(1, 2)$		$= \sigma'_v(1, 2)$		$= \sigma_v(2, 1)$	
$= (1, 2)$		$= (1, 2)$		$= (1, 2)$	
- Rule 3: Identity.
- Rule 4: Inverse.
 - For a C_{2v} point group:

$$E \cdot E = E$$

$$C_2 \cdot C_2 = E$$

$$\sigma_v \cdot \sigma_v = E$$

$$\sigma'_v \cdot \sigma'_v = E$$

- Group multiplication tables.

C_{2h}	E	C_2	σ_h	i
E	E	C_2	σ_h	i
C_2	C_2	E	i	σ_h
σ_h	σ_h	i	E	C_2
i	i	σ_h	C_2	E

Table II.1: Group multiplication table for the C_{2h} point group.

- Table II.1 corresponds to the C_{2h} point group, which has E , C_2 , σ_h , and i operations.
- The operation in the top row is applied first, and the one in the left column is applied second.

- **Subgroup:** Fractional parts of groups that are groups, too.

C_{3v}	E	C_3	C_3^2	σ_v	σ'_v	σ''_v
E	E	C_3	C_3^2	σ_v	σ'_v	σ''_v
C_3	C_3	C_3^2	E	σ''_v	σ_v	σ'_v
C_3^2	C_3^2	E	C_3	σ'_v	σ''_v	σ_v
σ_v	σ_v	σ'_v	σ''_v	E	C_3	C_3^2
σ'_v	σ'_v	σ_v	σ''_v	C_3	C_3^2	E
σ''_v	σ''_v	σ''_v	σ_v	C_3^2	E	C_3

Table II.2: Group multiplication table for the C_{3v} point group.

- If $h = 6$ (as in the C_{3v} group), the subgroups can have $h = 3, 2, 1$. Why only these?^[2]
- The order 1 and 3 charts are subgroups.
- The order 2 chart is not a subgroup because C_3^2 is not an operation in the group (no closure).
- We use subgroups because they can make complex problems simpler.
 - For example, when calculating the vibrational modes of CO_2 , we use D_{2h} , a simpler finite subgroup of its actual infinite $D_{\infty h}$ point group.

II.4 Module 6: Representations

- Items of the same point group have the same vibrational modes.
- **Representation** (of a group): Any collection of quantities (or symbols) which obey the multiplication table of a group. *Also known as Γ .*
- For our purposes, these quantities are the matrices that show how certain characteristic of a molecule behave under the symmetry operations of the group.
- Operations (on a point (x, y, z) in Cartesian coordinates):
 - $E(x, y, z) = (x, y, z)$.
 - $\sigma_{xz}(x, y, z) = (x, -y, z)$.
 - $i(x, y, z) = (-x, -y, -z)$.
 - C_n : Convention is a counterclockwise rotation of the point by $\theta = \frac{2\pi}{n}$ radians.
 - S_n : Convention is a C_n followed by a σ_h .

²Lagrange's theorem.

- Matrix forms of operations:

– Identity: $E = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{bmatrix}$.

– One example of a reflection (there are two more): $\sigma_{xy} = \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & -1 \end{bmatrix}$.

– Inversion: $i = \begin{bmatrix} -1 & 0 & 0 \\ 0 & -1 & 0 \\ 0 & 0 & -1 \end{bmatrix}$.

– Rotation: Counterclockwise is $C_n(\theta) = \begin{bmatrix} \cos \theta & -\sin \theta & 0 \\ \sin \theta & \cos \theta & 0 \\ 0 & 0 & 1 \end{bmatrix}$ and clockwise is $C_n(\theta) = \begin{bmatrix} \cos \theta & \sin \theta & 0 \\ -\sin \theta & \cos \theta & 0 \\ 0 & 0 & 1 \end{bmatrix}$.

■ A derivation of this matrix can be found in the slides.

– Improper rotation: $S_n(\theta) = \sigma_h C_n(\theta)$.

- **Reducible representations** (Γ):

- A representation of a symmetry operation of a group.
- Can be expressed in terms of a representation of lower dimension.
- Can be broken down into a simpler form.
- Characters can be further diagonalized.
- Are composed of the direct sum of irreducible representations.
- Infinite possibilities.

- **Irreducible representations** (Γ_i):

- A fundamental representation of a symmetry operation of a group.
- Cannot be expressed in terms of a representation of lower dimension.
- Cannot be broken down into a simpler form.
- Characters cannot be further diagonalized.
- Small finite number dictated by point group.

- Good example of reducible/irreducible representations?

- A representation shows how certain characteristics of an object (a basis) behave under the symmetry operation of the group.

- **Conjugate elements:** Two elements X and Y for which there exists an element Z in the group such that

$$Z^{-1} \cdot X \cdot Z = Y$$

- Every element is conjugated with itself (let $Z = E$).
- If X is conjugated with Y , then Y is conjugated with X .
- If X is conjugated with Y and W , then Y and W are also conjugate.

- **Class:** A complete set of elements of a group that are conjugate to one another.

- Geometric meaning: operations in the same class can be converted into one another by changing the axis system through application of some symmetry operation of the group.

- Find the conjugates to C_3 in the C_{3v} point group (refer to Table II.2 throughout the following discussion).

- Let $X = C_3$, let Z iterate through the six symmetry elements ($E, C_3, C_3^2, \sigma_v, \sigma'_v, \sigma''_v$), and let Z^{-1} iterate through the corresponding inverses ($E, C_3^2, C_3, \sigma_v, \sigma'_v, \sigma''_v$).
- Thus, we have

$$\begin{aligned} E \cdot C_3 \cdot E &= C_3 \\ C_3^2 \cdot C_3 \cdot C_3 &= C_3 \\ C_3 \cdot C_3 \cdot C_3^2 &= C_3 \\ \sigma_v \cdot C_3 \cdot \sigma_v &= C_3^2 \\ \sigma'_v \cdot C_3 \cdot \sigma'_v &= C_3^2 \\ \sigma''_v \cdot C_3 \cdot \sigma''_v &= C_3^2 \end{aligned}$$

- It follows from the above that C_3 and C_3 are conjugates, and C_3 and C_3^2 are conjugates.
- Thus, C_3 and C_3^2 are in the same class.
- We can use a similar method to prove that σ_v, σ'_v , and σ''_v are all in the same class within the C_{3v} point group.
- Likewise E is in a class by itself.
- Thus, for the C_{3v} point group, E forms a class of order 1, C_3, C_3^2 form a class of order 2, and $\sigma_v, \sigma'_v, \sigma''_v$ form a class of order 3.
- Note, however, that *all* rotation and reflection operations are not always in the same class.
 - In the higher-order C_{5v} point group, for example, C_5 and C_5^4 are in the same class, and C_5^2 and C_5^3 are in the same class, but these two classes are distinct.

- **Similarity transformation:** The transformation

$$v^{-1} \cdot A \cdot v = A'$$

- A is a representation for some type of symmetry operation.
 - v is a similarity transform operator.
 - v^{-1} is the inverse of the similarity transform operator.
 - A' is the product.
 - A and A' are conjugates, and we say that A' is the similarity transform of A by v .
 - If A and A' are matrices of equal dimensions, A and A' are **similar matrices**, and v is the change of coordinates matrix.
- **Block-diagonal** (matrix): A matrix with nonzero values only in square blocks along the diagonal from the top left to the bottom right.

$$\left[\begin{array}{cc|cc|cc} 2 & 3 & & & & \\ 1 & 2 & & 0 & & 0 \\ \hline & & 1 & 1 & & \\ & 0 & 1 & 1 & & 0 \\ \hline & & & & 0 & 2 \end{array} \right]$$

- The above matrix is an example of a block-diagonal matrix.
- Irreducible representations are the corresponding blocks within a set of block-diagonalized matrices representing each operation in a group.

II.5 Module 7: Characters and Character Tables

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- Nocera Lecture 3 notes (on Canvas) explain how reducible and irreducible transformations are related to each other through the similarity transformations.
- For H_2O , each atom has 3 Cartesian coordinates, so our transformation matrix^[3] is 9-square.

³Some values in it are negative because of the cosine/sine definition of a rotation matrix for $\theta = 180^\circ$.

- However, we can also apply a smaller matrix to the molecule as a whole and invoke symmetry to find the position of the individual atoms.
- **Characters** (of a representation): The traces (i.e., sums of the diagonal matrix elements) of the representation matrices for each operation. *Also known as χ .*
 - The character is an invariant for each type of symmetry operation (e.g., regardless of the axis about which a C_n operation is performed, the trace of the corresponding matrix will be the same).
- Common characters:
 - C_n character: $\chi = 2 \cos \theta + 1$.
 - σ_v, σ_d character: $\chi = 1$.
 - $S_2 \equiv i$ character: $\chi = -3$.
 - S_n character: $\chi = 2 \cos \theta - 1$.
 - $\perp C_2$ axes character: $\chi = -1$.
- **Character table:** The collection of characters for a given irreducible representation, under the operations of a group. *Also known as χ table.*

Irreducible Representations	Group Symbol	Symmetry Elements				linear	quadratic
	C_{2v}	E	C_2	$\sigma_v(xz)$	$\sigma'_v(yz)$		
	A_1	1	1	1	1	z	x^2, y^2, z^2
	A_2	1	1	-1	-1	R_z	xy
	B_1	1	-1	1	-1	x, R_y	xz
	B_2	1	-1	-1	1	y, R_x	yz
Characters						Basis Functions	

Figure II.4: A character table.

- Character tables for all point groups are listed in Appendix C of Miessler et al. (2014).
- **Mulliken symbols** are used to classify irreducible representations based on degeneracy and symmetry.
 - A or B: singly degenerate (the maximum block size in the block-diagonalized irreducible transformation matrix is 1×1).
 - E: Doubly degenerate (the maximum block size in the block-diagonalized irreducible transformation matrix is 2×2).
 - T: Triply degenerate (the maximum block size in the block-diagonalized irreducible transformation matrix is 3×3).
 - A: symmetric (+) with respect to C_n .
 - B: anti-symmetric (–) with respect to C_n .
 - Subscript g: symmetric (+) with respect to i . *Etymology* short for gerade (German for symmetric).
 - Subscript u: anti-symmetric (–) with respect to i . *Etymology* short for ungerade (German for unsymmetric).
 - If the molecule has a center of inversion, we label irreducible representations with g or u .
 - Subscript 1: symmetric (+) with respect to $\perp C_2$ or σ_v .
 - Subscript 2: anti-symmetric (–) with respect to $\perp C_2$ or σ_v .
 - Superscript ': symmetric (+) under σ_h (if no i).
 - Superscript ": anti-symmetric (–) under σ_h (if no i).

- Don't mistake the operation E for the Mulliken symbol E .
- To assign Mulliken symbols, use the character table.
 - Assigning the main letter:
 - If E -character = 1 and C_n -character = 1: A .
 - If E -character = 1 and C_n -character = -1 : B .
 - If E -character = 2: E .
 - If E -character = 3: T .
 - Assigning a subscript g or u :
 - If i -character = 1: g .
 - If i -character = -1 : u .
 - This subscript can be assigned to A, B, E, T representations.
 - Assigning a superscript $'$ or $''$:
 - If σ_h -character = 1: $'$.
 - If σ_h -character = -1 : $''$.
 - This subscript can be assigned to A, B representations.
 - Assigning a subscript 1 or 2:
 - If $\perp C_2$ or σ_v -character = 1: 1.
 - If $\perp C_2$ or σ_v -character = -1 : 2.
 - This subscript can be assigned to A, B representations.
- σ , π , and δ bonds come from the Mulliken symbols!
 - Infinity tables use Greek rather than Latin letters.

D_3	E	$2C_3(z)$	$3C'_2$	linear	quadratic
A_1	1	1	1		$x^2 + y^2, z^2$
A_2	1	1	-1	z, R_z	
E	2	-1	0	$(x, y)(R_x, R_y)$	$(x^2 - y^2, xy)(xz, yz)$.

Table II.3: Character table for the D_3 point group.

- In character tables, we need to multiply each symmetry operation by the order of the class (see Table II.3).
- Basis functions show us how different functions transform under different symmetry operations.
- In the C_{2v} point group:
 - The p_x orbital has B_1 symmetry.
 - p_x transforms as B_1 .
 - p_x has the same symmetry as B_1 .
 - p_x forms a basis for the B_1 irreducible representation.
 - p_x is B_1 because (see Table 0.2 and Figure II.4) it does not change under E , it inverts under C_2 , it does not change under $\sigma_v(xz)$, and it inverts under $\sigma'_v(yz)$ (note that the C_2 axis is the z -axis, not the x -axis).
- We can apply the same procedure to other more complex functions.
 - For example, in the C_{2v} point group, we know that (with respect to orbitals):

- p_y is B_2 .
 - p_z is A_1 .
 - d_{z^2} is A_1 .
 - $d_{x^2-y^2}$ is A_1 .
 - d_{yz} is B_2 .
 - d_{xy} is A_2 .
 - d_{xz} is B_1 .
 - We can even go into the cubic functions describing the f orbitals and assign them Mulliken symbols.
- Essentially, the right hand side of a character table tells you how atomic orbitals will transform under certain symmetry operations.
 - Properties of a character table:
 1. The characters of all matrices belonging to the operations in the same class are identical in a given irreducible representation.
 - We most commonly form a **rotational class** and a **reflection class**.
 2. The number of irreducible representations in a group is equal to the number of classes of that group.
 3. There is always a totally symmetric representation for any group.
 - I.e., a representation where every character is 1.
 4. The sum of the squares of the **dimensionality** of all the irreducible representations is equal to the order of the group. Mathematically,

$$h = \sum_i [\chi_i(E)]^2$$

- For example, the dimensionalities of the D_3 point group (see Table II.3) are 1, 1, and 2, and the order is, indeed, $6 = 1^2 + 1^2 + 2^2$.
- 5. The sum of the squares of the characters multiplied by the number of operations in the class equals the order of the group. Mathematically,

$$h = \sum_{R_c} g_c [\chi_i(R_c)]^2$$

- For example, with respect to the D_3 point group (see Table II.3),

$$\begin{aligned} 6 &= (1)(1)^2 + (2)(1)^2 + (3)(1)^2 \\ &= (1)(1)^2 + (2)(1)^2 + (3)(-1)^2 \\ &= (1)(2)^2 + (2)(-1)^2 + (3)(0)^2 \end{aligned}$$

6. The sum of the products of the corresponding characters of any two different irreducible representations of the same group is zero. Mathematically,

$$\sum_{R_c} g_c \chi_i(R_c) \chi_f(R_c) = 0$$

- Basically, this means that if we treat irreducible representations as vectors in h -space, they are orthogonal.
- For example, with respect to the D_3 point group (see Table II.3),

$$\begin{aligned} 0 &= 1(1)(1) + 2(1)(1) + 3(1)(-1) \\ &= 1(1)(2) + 2(1)(-1) + 3(1)(0) \\ &= 1(1)(2) + 2(1)(-1) + 3(-1)(0) \end{aligned}$$

- **Dimensionality:** The character of the identity operation E . Also known as **dimension**.

II.6 Module 8: Using Character Tables

- A reducible representation of a group is any representation Γ of the form

$$\Gamma = \sum_i a_i \Gamma_i$$

where each Γ_i is an irreducible representation of the group and a_i is a real scalar.

- Basically, a reducible representation is any linear combination of irreducible representations.
- For example, with respect to the C_{2v} point group (see Figure II.4), $\Gamma = (7, 1, 5, 3) = 4A_1 + 2B_1 + B_2$ is a reducible representation.
- We may “factor” reducible representations by inspection, or by the...
- Decomposition/reduction formula for a reducible representation:

$$a_i = \frac{1}{h} \sum_Q N \cdot \chi(R)_Q \cdot \chi_i(R)_Q$$

- a_i is the number of times the irreducible representation appears in Γ_1 .
- h is the order of the group.
- N is the number of operations in class Q .
- $\chi(R)_Q$ is the character of the reducible representation.
- $\chi_i(R)_Q$ is the character of the irreducible representation.
- This formula cannot be applied to $D_{\infty h}$ and $C_{\infty v}$.
- Let's look at decomposing $\Gamma = (7, 1, 5, 3)$ into its component irreducible point groups using the above formula.

$$\begin{aligned} a_{A_1} &= \frac{1}{4}(1 \cdot 7 \cdot 1 + 1 \cdot 1 \cdot 1 + 1 \cdot 5 \cdot 1 + 1 \cdot 3 \cdot 1) = 4 \\ a_{A_2} &= \frac{1}{4}(1 \cdot 7 \cdot 1 + 1 \cdot 1 \cdot 1 + 1 \cdot 5 \cdot -1 + 1 \cdot 3 \cdot -1) = 0 \\ a_{B_1} &= \frac{1}{4}(1 \cdot 7 \cdot 1 + 1 \cdot 1 \cdot -1 + 1 \cdot 5 \cdot 1 + 1 \cdot 3 \cdot -1) = 2 \\ a_{B_2} &= \frac{1}{4}(1 \cdot 7 \cdot 1 + 1 \cdot 1 \cdot -1 + 1 \cdot 5 \cdot -1 + 1 \cdot 3 \cdot 1) = 1 \end{aligned}$$

- You can also find websites that will apply the formula for you.
- Basis \rightarrow reducible representation \rightarrow irreducible representations workflow:
 1. Assign a point group.
 2. Choose a basis function (bond, vibration, orbital, angle, etc.).
 3. Apply operations.
 - The following shortcuts allow us to skip matrix math in certain situations.
 - If the basis stays the same: +1.
 - If the basis is reversed: -1.
 - If it is a more complicated change: 0.
 4. Generate a reducible representation.
 5. Reduce to irreducible representation.
- We now look at an example of applying the above method to H_2O .

- H_2O is of the C_{2v} point group.
- The 9×9 identity matrix represents the identity operation on the 3 atoms in H_2O , each described by 3 Cartesian coordinates. Thus, $\chi(E) = 9$.
- The following matrix represents the C_2 symmetry operation. Thus, $\chi(C_2) = -1$.

$$\begin{array}{c} \text{O} \\ \text{H}_a \\ \text{H}_b \end{array} \begin{array}{c} \left[\begin{array}{c} x' \\ y' \\ z' \end{array} \right] \\ \left[\begin{array}{c} x' \\ y' \\ z' \end{array} \right] \\ \left[\begin{array}{c} x' \\ y' \\ z' \end{array} \right] \end{array} = \begin{bmatrix} -1 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & -1 & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 1 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & -1 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & -1 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 1 \\ 0 & 0 & 0 & -1 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & -1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 1 & 0 & 0 & 0 \end{bmatrix} \begin{array}{c} \left[\begin{array}{c} x \\ y \\ z \end{array} \right] \\ \left[\begin{array}{c} x \\ y \\ z \end{array} \right] \\ \left[\begin{array}{c} x \\ y \\ z \end{array} \right] \end{array}$$

- Note that atoms moved during the transformation do not contribute to the character of the transformation matrix.
- Since under $\sigma_v(xz)$ only O is unshifted, we need only consider its part of the transformation matrix (as follows) when looking for the character. Thus, $\chi(\sigma_v(xz)) = 1^{[4]}$.

$$\begin{bmatrix} x' \\ y' \\ z' \end{bmatrix} = \begin{bmatrix} 1 & 0 & 0 \\ 0 & -1 & 0 \\ 0 & 0 & 1 \end{bmatrix} \begin{bmatrix} x \\ y \\ z \end{bmatrix}$$

- Since under $\sigma_v(yz)$ no atom is shifted, we need to consider each (identical) part of the transformation matrix (as follows) when looking for the character. Thus, $\chi(\sigma_v(yz)) = 3$.

$$\begin{bmatrix} x' \\ y' \\ z' \end{bmatrix} = \begin{bmatrix} -1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{bmatrix} \begin{bmatrix} x \\ y \\ z \end{bmatrix}$$

- Thus, the characters of our final reducible representation is $\Gamma_{3N} = (9, -1, 1, 3)$.

■ This representation represents fully unrestricted motion of all 3 ambiguities of freedom.

- Another example: XeOF_4 .

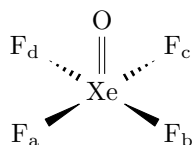


Figure II.5: XeOF_4 .

- Point group: C_{4v} .
- Basis function: F atoms.
- Let's see what happens to the fluorine atoms under the C_{4v} operations (remember that if a basis element (a fluorine atom) stays the same, it contributes +1 to the character of an operation, and if it moves, it contributes 0 to the character).

⁴Note also that $a_{11} = 1$ since the x -vector (a basis) is unchanged under $\sigma_v(xz)$, $a_{22} = -1$ since the y -vector is reversed under $\sigma_v(xz)$, and $a_{33} = 1$ since the z vector is unchanged under $\sigma_v(xz)$.

E	all unchanged	4
C_4	all move	0
C_2	all move	0
$2\sigma_v$	2 move, 2 unchanged	2
$2\sigma_d$	all move	0

Table II.4: Changes in the fluorine atoms of XeOF_4 under the C_{4v} symmetry operations.

- Thus, $\Gamma = (4, 0, 0, 2, 0)$.
- With the C_{4v} character table and the decomposition formula, we can discover that $\Gamma = A_1 + B_1 + E$.

II.7 Nocera Lecture 3

From Nocera (2008).

- 1/25:
- Corresponding blocks within block-diagonalized matrices obey the same multiplication properties as the block-diagonalized matrices, themselves.
 - One basis may not uncover all irreducible representations.
 - For example, we can use a Cartesian basis to uncover the A_1 and E irreducible representations of the D_3 point group (see Table II.3) and the basis function R_z ($E : R_z \rightarrow R_z$, $C_3 : R_z \rightarrow R_z$, $\sigma_v : R_z \rightarrow \bar{R}_z$) to uncover the A_2 irreducible representation.
 - We can also construct a character table algebraically. Continuing with the D_3 example...
 - There is one totally symmetric representation $\Gamma_i = (1, 1, 1)$.
 - $h = 6 = \sum_i [\chi_i(E)]^2$. Since each $\chi_i(E)$ can only be 1, 2, or 3, we know that $\chi_1(E) = \chi_2(E) = 1$ and $\chi_3(E) = 2$.
 - This combines with orthogonality to reveal that

$$\begin{aligned}
 0 &= \sum_{R_c} g_c \chi_1(R_c) \chi_2(R_c) \\
 &= 1 \cdot 1 \cdot 1 + 2 \cdot 1 \cdot \chi_2(C_3) + 3 \cdot 1 \cdot \chi_2(\sigma_v) \\
 &= 1 + 2\chi_2(C_3) + 3\chi_2(\sigma_v)
 \end{aligned}$$

i.e., $\Gamma_i = (1, 1, -1)$.

- To determine the last representation, we can use two rules and solve as a system of equations.

$$\begin{aligned}
 0 &= \sum_{R_c} g_c \chi_1(R_c) \chi_3(R_c) & 6 &= \sum_{R_c} g_c [\chi_3(R_c)]^2 \\
 &= 2 + 2\chi_3(C_3) + 3\chi_3(\sigma_v) & &= 4 + 2[\chi_3(C_3)]^2 + 3[\chi_3(\sigma_v)]^2
 \end{aligned}$$

From this, we can determine that $\chi_3(C_3) = -1$ and $\chi_3(\sigma_v) = 0$. Therefore, $\Gamma_i = (2, -1, 0)$.

- General algebraic procedure: Apply Property 1, then 2, then 4, then 6. Check with 5.

II.8 TA Review Session 1

- 1/22:
- We also don't need to show the equatorial electron pair in the shape picture.
 - For SeCl_4 , we need to show that the axial bonds are longer and are not straight up and down.
 - For I_3^- , it's trigonal bipyramidal EPA and linear molecular geometry — there's an extra electron pair around the central iodine atom. This makes it $D_{\infty h}$ point group.

- For SeOCl_4 , we should also make the axials longer and bent away from the oxygen atom.
 - This molecule is sp^3d hybridized, not sp^2d^2 , as it would be if the oxygen were axial.
- For $\text{IO}(\text{OH})_5$, we should show the equatorial OH's pushed away from the oxygen atom. The oxygen bond should also be a bit shorter.
- In the exam, they will specify whether we need to consider the hydrogens or not when calculating symmetry.
- For ClOF_4^- , one or the other of the lone pair/oxygen will push the equatorial fluorines a bit. You don't have to know which, just show bent.
- For XeO_2F_2 , show the lone pair pushing the axial fluorines away.
- For IF_3^{2-} , show that this is a *distorted* T.
- A tennis ball belongs to the D_{2d} point group since you have perpendicular C_2 axes punching through the seam and mirror planes at 45° angles to the C_2 axes (i.e., dihedral).
- FeF_6^{3-} also loses 4 C_2 axes.
- There are three possible isomers of IF_3O_2 (see Figure I.1), but the one with equatorial oxygens is the most stable.
 - The structure analogous to Figure I.1a has C_{2v} symmetry.
 - The structure analogous to Figure I.1b has D_{3h} symmetry.
 - The structure analogous to Figure I.1c has C_s symmetry.
- Determine the point group of NO_3^{2-} , $\text{HFC}=\text{C}=\text{CHF}$, $\text{H}_2\text{C}=\text{CF}_2$, and C_2H_6 (consider three possible conformers).
 - NO_3^{2-} is of the D_{3h} point group.
 - $\text{HFC}=\text{C}=\text{CHF}$ is of the C_2 point group.
 - $\text{H}_2\text{C}=\text{CF}_2$ is of the C_{2v} point group.
 - C_2H_6 eclipsed has D_{3h} symmetry, staggered has D_{3d} symmetry, and others have D_3 symmetry.

II.9 Module 9: Molecular Vibrations

- 1/25:
- If you have a nonlinear triatomic molecule (like H_2O), we have $3 \text{ atoms} \times 3 \text{ DOF} = 9 \text{ DOF}$.
 - This accounts for any possible perturbations of our molecules.
 - We can break this down into three types of motion:
 - Translational motion along the x -, y -, and z -axes (3 DOFs).
 - Rotational motion about the x -, y -, and z -axes (3 more DOFs).
 - The remaining degrees of freedom are vibrational.
 - Every *nonlinear* molecule has 3 translational and 3 rotational DOFs.
 - The number of vibrations is $3N - 6$, where N is the number of atoms.
 - Every *linear* molecule has 3 translational and 2 rotational DOFs.
 - The number of vibrations is $3N - 5$, where N is the number of atoms.
 - **Vibrational mode:** A perturbation of molecular structure that keeps the center of mass of the molecule in one place.

- For H_2O , there are three vibrational modes: antisymmetric stretch (one H moves away from the O as the other moves in, and then the process reverses), symmetric stretch (both H's move away from the O and then move back in), and scissoring bend (the bond angle changes).
- For molecules in general, there are these three and an additional three: wagging (in H_2O , the H's rotate out of the plane of the molecule and then back), twisting (in H_2O , both H's rotate out of the plane of the molecule, but one in one direction and one in the other), and rocking (in H_2O , both H's rotate about the O within the plane of the molecule preserving the bond angle, and then go back).
- What vibrational modes a molecule can and cannot have is governed by its point group and group theory.
- Each normal mode of vibration forms a basis for an irreducible representation of the point group of the molecule.
- Workflow to identify the vibrational properties of a molecule:
 1. Find number/symmetry of vibrational modes.
 2. Assign the symmetry of known vibrations.
 3. What does the vibration look like?
 4. Find if a vibrational mode is IR or Raman Active.
- To find vibrational modes, follow the 5-step Basis \rightarrow reducible representation \rightarrow irreducible representation workflow and then subtract translational and rotational motion.
 - For example, with H_2O , as before, we have C_{2v} point group, $3N$ basis, $\Gamma_{3N} = (9, -1, 1, 3)$, and $\Gamma_{3N} = 3A_1 + A_2 + 2B_1 + 3B_2$.
 - We now need to use the basis functions in the C_{2v} character table to determine the translational and rotational degrees of freedom. From Figure II.4, the x, y, z -translational modes come from the B_1 , B_2 , and A_1 Γ_i 's, respectively. Thus, the total translational mode, sufficient to describe all possible translations in 3 DOFs, is $\text{trans} = A_1 + B_1 + B_2$. Similarly, we can determine that the total rotational mode is $\text{rot} = A_2 + B_1 + B_2$.
 - Thus, we can determine that the total vibrational mode is $\Gamma_{3N} - \text{trans} - \text{rot} = 2A_1 + B_2$.
 - When H_2O scissors or symmetrically stretches, it maintains its C_{2v} symmetry. Thus, both of these vibrations are represented by the totally symmetric irreducible representation A_1 (hence the $2A_1$ component).
 - As to asymmetric stretch, there exist points in time where H_2O lowers its symmetry from C_{2v} to C_s , losing its C_2 and $\sigma_v(xz)$ symmetry elements but maintaining E and $\sigma_v(yz)$. This shift is encapsulated by the B_2 irreducible representation.
 - We can see all of these modes in H_2O 's infrared absorption spectrum.
 - Note that this spectrum does not tell us the energy of vibrations (we can model this with quantum mechanics), or if it is IR or Raman active (which modes will appear in the respective spectrum).
- Determine the number and symmetry types for the translations, vibrations, and rotations of PF_5 .
 - Point group: D_{3h} .
 - We could write out all 18 x, y, z -vectors, or we can use a shortcut: To construct Γ_{3N} , we can, for each symmetry element, multiply the number of atoms unmoved under the operation by the corresponding character the representation that accounts for x, y, z -movements.

$$\Gamma_{3N} = (\Gamma_{x,y,z}) \cdot (\# \text{ unmoved atoms})$$

Applying this shortcut, we have $\Gamma_{x,y,z} = E' + A_2'' = (3, 0, -1, 1, -2, 1)$. We also have $\# \text{ unmoved atoms} = (6, 3, 2, 4, 1, 4)$. Thus, $\Gamma_{3N} = (18, 0, -2, 4, -2, 4)$.

D_{3h}	E	$2C_3$	$3C_2$	σ_h	$2S_3$	$3\sigma_v$	linear	quadratic
A'_1	1	1	1	1	1	1		$x^2 + y^2, z^2$
A'_2	1	1	-1	1	1	-1	R_z	
E'	2	-1	0	2	-1	0	(x, y)	$(x^2 - y^2, xy)$
A''_1	1	1	1	-1	-1	-1		
A''_2	1	1	-1	-1	-1	1	z	
E''	2	-1	0	-2	1	0	(R_x, R_y)	(xz, yz)

Table II.5: Character table for the D_{3h} point group.

- We can now reduce it into $\Gamma_{3N} = 2A'_1 + A'_2 + 4E' + 3A''_2 + 2E''$.
 - Note that we still have our 18 degrees of freedom because each E is double degenerate, meaning that $4E'$ counts for 8 degrees of freedom and $2E''$ counts for 4 degrees of freedom.
- This combined with the fact that $\Gamma_{\text{trans}} = \Gamma_{x,y,z} = E' + A''_2$ and $\Gamma_{\text{rot}} = A'_2 + E''$ implies that $\Gamma_{\text{vibs}} = 2A'_1 + 3E' + 2A''_2 + E''$.
 - For a similar reason to the note above, we can count $12 = 3N - 6$ degrees of freedom (vibrational modes) in Γ_{vibs} .
- Although degenerate modes count for multiple DOFs, the multiple modes they count for are of the same type.
 - In an ideal world, those modes corresponding to E would be twice as intense as the others.
 - From the definition of Γ_{vibs} and the above discussion, we know that PF_5 has $2 + 3 + 2 + 1 = 8$ distinct vibrational modes.
- There are (broadly) two types of vibrations: **stretches** (ν) and **bends** (δ).
 - These two types appear in different energies in IR and Raman spectra.
- To differentiate between stretching and bending modes: Look at how the stretching modes transform. Each stretch happens along a single vector from central atom to ligand. Thus, if we let these vectors be our basis and look at how many vectors stay the same (+1) or change (-1) under each operation, we will have a reducible representation that can be decomposed. Its components will be the stretching modes and $\Gamma_{\text{vibs}} - \Gamma_{\nu}$ will equal Γ_{δ} .
 - Applied to PF_5 , we have $\Gamma_{\nu} = (5, 2, 1, 3, 0, 3) = 2A'_1 + E' + A''_2$.
 - Thus, both A'_1 representations, one E' representation, and one A''_2 representation correspond to stretching modes (or [spectroscopic] bands; note that the E' band will be twice as large). Consequently, the other two E' representations, the other A''_2 representation, and the E'' representation correspond to bending modes.

II.10 Module 10: IR and Raman Active Vibrations (part 1)

- Today, we will learn what we need to know for PSet 2. Wednesday, we will introduce a more rigorous, quantum mechanical foundation.
- Molecular vibrations can be experimentally observed by **infrared spectroscopy** or **Raman spectroscopy**.
- **Infrared spectroscopy**: A method of measuring the change in dipole moment during a vibration^[5]. Also known as **IR spectroscopy**.

⁵Refer to Labalme (2020), specifically the discussion of spectroscopy in Chapter 13.

- **Raman spectroscopy:** A method of measuring the change in the polarizability during a vibration. We impinge upon a sample of a substance with an incident light (a powerful laser). Most of the light that scatters will be a **Raleigh scatter**, but some will be a **Raman scatter** (a new wavelength).
- **Raleigh scatter:** The same wavelength is emitted as was impinged by the incident light because the laser was elastically scattered. The electrons fell the same number of energy levels that they rose after absorbing the light.
- **Raman scatter:** A new wavelength is emitted, different than the one impinged by the incident light. The electrons fell fewer (**Stokes Raman scattering**) or more (**Anti-Stokes Raman scattering**) energy levels than they rose after absorbing the light.
- **IR active** (vibration): A vibration that transforms with the same symmetry as the \vec{x} , \vec{y} , and \vec{z} vectors (see the χ tables).
- **Raman active** (vibration): A vibration having the same symmetry as the quadratic x, y, z terms, i.e., x^2, z^2, yz , etc.
- For molecules possessing inversion centers, IR and Raman activity will be mutually exclusive.
- Activity data comes from character tables.
 - All vibration representations that have a linear term are IR active, while those that have a quadratic term are Raman active.
- Continuing with the PF_5 example...
 - Table II.5 tells us that E' and A_2'' vibrations are IR active, and A_1' , E' , and E'' are Raman active.
 - Thus, since $\Gamma_{\text{vibs}} = 2A_1' + 3E' + 2A_2'' + E''$, there are $3 + 2 = 5$ IR active bonds (corresponding to $3 \cdot 2 + 2 = 8$ vibrations) and $2 + 3 + 1 = 6$ Raman active bonds (corresponding to $2 + 3 \cdot 2 + 1 \cdot 2 = 10$ vibrations).

II.11 Chapter 4: Symmetry and Group Theory

From Miessler et al. (2014).

- 1/18: • **Coincident** (axes): Two identical axes.
- For example, the C_3 rotation axis of CHCl_3 is **coincident** with the C–H bond axis.
- Snowflakes, which are often planar and have hexagonal symmetry, have a twofold (C_2), threefold (C_3), and sixfold (C_6) axis through their center and perpendicular to their plane (see Figure II.6).
- Rotations C_3^2 and C_6^5 are also symmetry operations.
- “When necessary, the C_2 axes perpendicular to the principal axis are designated with primes; a single prime (C_2') indicates that the axis passes through several atoms of the molecule, whereas a double prime (C_2'') indicates that it passes between the outer atoms” (Miessler et al., 2014, p. 77).
- 1/19: • Even though $S_2 \equiv i$ and $S_1 \equiv \sigma$, the i and σ notations are preferred because of the group theory requirement of maximizing the number of unique classes of symmetry operations associated with a molecule.
- **Point group:** A set of symmetry operations that describes a molecule’s overall symmetry.
 - Alternative steps for assigning point groups:
 1. Determine whether the molecule exhibits very low symmetry (C_1 , C_s , C_i) or high symmetry (T_d , O_h , $C_{\infty v}$, $D_{\infty h}$, I_h).

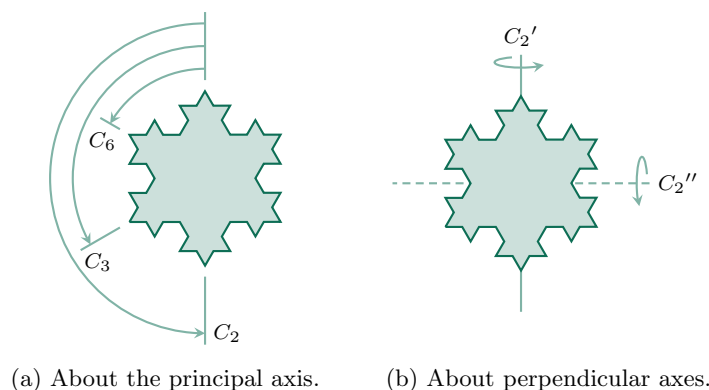


Figure II.6: Rotations of a snowflake design.

2. If not, find the highest order C_n axis for the molecule.
3. Does the molecule have any C_2 axes perpendicular to the principal C_n axis? If it does, there will be n of such C_2 axes, and the molecule is in the D set of groups. If not, it is in the C or S set.
4. Does the molecule have a mirror plane (σ_h) perpendicular to the principal C_n axis? If so, it is classified as C_{nh} or D_{nh} . If not, continue with Step 5.
5. Does the molecule have any mirror planes that contain the principal C_n axis (σ_v or σ_d)? If so, it is classified as C_{nv} or D_{nd} . If not, but it is in the D set, it is classified as D_n . If the molecule is in the C or S set, continue with Step 6.
6. Is there an S_{2n} axis collinear with the principal C_n axis? If so, it is classified as S_{2n} . If not, the molecule is classified as C_n .

• Groups of high symmetry:

- $C_{\infty v}$ (linear): These molecules are linear, with an infinite number of rotations and an infinite number of reflection planes containing the rotation axis. They do not have a center of inversion.
- $D_{\infty h}$ (linear): These molecules are linear, with an infinite number of rotations and an infinite number of reflection planes containing the rotation axis. They also have perpendicular C_2 axes, a perpendicular reflection plane, and an inversion center.
- T_d (tetrahedral): Most (but not all) molecules in this point group have the familiar tetrahedral geometry. They have four C_3 axes, three C_2 axes, three S_4 axes, and six σ_d planes. They have no C_4 axes.

■ Look for C_3 and C_2 axes.

- O_h (octahedral): These molecules include those of octahedral structure, although some other geometrical forms, such as the cube, share the same set of symmetry operations. Among their 48 symmetry operations are four C_3 rotations, three C_4 rotations, and an inversion.

■ Look for C_4 , C_3 , and C_2 axes.

- I_h (icosahedral): Icosahedral structures are best recognized by their six C_5 axes, as well as many other symmetry operations — 120 in all.

■ Look for C_5 , C_3 , and C_2 axes.

- T_h : Adds i to T_d . Example: $W[N(CH_3)_2]_6$.

1/24:

- When we have a block-diagonalized reducible representation, the irreducible representations can be obtained from the individual blocks.
 - This is because we will find that an analogous set of blocks satisfies the multiplication table for the group.

- For the C_{2v} point group, we have the following reducible representation that can be applied to each coordinate.

$$E : \begin{bmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{bmatrix} \quad C_2 : \begin{bmatrix} -1 & 0 & 0 \\ 0 & -1 & 0 \\ 0 & 0 & 1 \end{bmatrix} \quad \sigma_v(xz) : \begin{bmatrix} 1 & 0 & 0 \\ 0 & -1 & 0 \\ 0 & 0 & 1 \end{bmatrix} \quad \sigma_v(yz) : \begin{bmatrix} -1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{bmatrix}$$

- The above matrices are block diagonalized. Thus, the sets of a_{11} , a_{22} and a_{33} form three irreducible representations (and we can confirm that these obey the C_{2v} group multiplication table).

$$\begin{array}{llll} E : [1] & C_2 : [-1] & \sigma_v(xz) : [1] & \sigma_v(yz) : [-1] \\ E : [1] & C_2 : [-1] & \sigma_v(xz) : [-1] & \sigma_v(yz) : [1] \\ E : [1] & C_2 : [1] & \sigma_v(xz) : [1] & \sigma_v(yz) : [1] \end{array}$$

- Three of the C_{2v} irreducible representations can be found in this way. The fourth can be found with linear algebra and the character table properties (or by inspection).
- If one matrix in the reducible representation cannot be blocked past having a 2×2 chunk, per se, then the others must also have that 2×2 chunk (so that we have an irreducible 2×2 representation).
- “Matching the symmetry operations of a molecule with those listed in the top row of the character table will confirm any point group assignment” (Miessler et al., 2014, p. 99).
- **Symmetric:** Character of 1.
- **Antisymmetric:** Character of -1 .
- Mulliken symbol rules:

1. “Letters are assigned according to the dimension of the irreducible representation” (Miessler et al., 2014, p. 99).
 - Assign A if the dimension is 1 and the representation is symmetric to the principal rotation operation ($\chi(C_n) = 1$).
 - Assign B if the dimension is 1 and the representation is antisymmetric to the principal rotation operation ($\chi(C_n) = -1$), or if $\chi(S_{2n}) = -1$ even if $\chi(C_n) = 1$.
 - Assign E if the dimension is 2.
 - Assign T if the dimension is 3.
2. “Subscript 1 designates a representation symmetric to a C_2 rotation perpendicular to the principal axis, and subscript 2 designates a representation antisymmetric to the C_2 . If there are no perpendicular C_2 axes, 1 designates a representation symmetric to a vertical plane, and 2 designates a representation antisymmetric to a vertical plane” (Miessler et al., 2014, p. 100).
3. “Subscript g designates representations symmetric to inversion, and subscript u designates representations antisymmetric to inversion” (Miessler et al., 2014, p. 100).
4. “Single primes are symmetric to σ_h and double primes are antisymmetric to σ_h when a distinction between representations is needed (C_{3h} , C_{5h} , D_{3h} , D_{5h})” (Miessler et al., 2014, p. 100).

1/26:

- **Chiral** (molecule): A molecule that is not superimposable on its mirror image. *Also known as dissymmetric.*
 - “In general, a molecule or object is chiral if it has no symmetry operations (other than E), or if it has only proper rotation axes” (Miessler et al., 2014, p. 100).
- Raman spectroscopy makes use of higher energy radiation than IR, exciting molecules to higher electronic states that are envisioned as short-lived “virtual” states.

II.12 Columbia Group Theory

From here.

- A great introduction to the more formal mathematics underlying symmetry groups.