

# Exercise Session 6 IESM Fall 2023-2024

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#### Last Week's Annoucement

As mentioned last week, during your work on exercises for this course you have the opportunity to help Zhenyu Cai, a PhD student in the CHILI lab who is developing new teaching tools \* More details about the project: https://drive.google.com/file/d/ 1GNR9NSpvh3s0nmvWXwMBONjNC3mAAW53/view?usp=sharing \* Consent form (if you agree to participate): https://forms.gle/hc6K8f5yRUA5T9Lz9 \* Your decision to participate will have no influences on your grades in any way. We will remain unaware of your consent status. \* If you previously said "no" to cotnirbuting but wish to change your answer, let him know so he can re-launch your permissions \* Thanks for considering!



# Exercise 6 - DFT vs (Post) HF Methods





## Exercise 6 - DFT vs (Post) HF Methods: Theory

- (Post) HF methods are wavefunction-based (we need to find the wavefunction)
- DFT shifts the focus: we need to find the ground-state charge density
- Why? For N electrons, wavefunction is a complex function of 3N variables, but the ground-state charge density is a function of 3 variables
- The universal functional not known, but proven to exist
- Everything that is unknown is contained in  $E_{XC}$

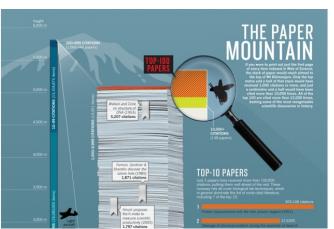
$$E(\rho) = T_0(\rho) + J(\rho) + \int v_{\text{ext}} \rho + E_{XC}$$

Each functional will treat the XC part differently



## Exercise 6 - DFT (continued)

DFT is the workhorse of electronic structure methods:



 In the top 100 most cited papers (ever!!) in the scientific community, 12 are on DFT



#### Comments on DFT

- Kohn-Sham formulation: ficticious molecular orbitals (non-interacting)
- If the exact XC functional is known ground state energies, charge densities and HOMO (Koopman's theorem) are known
- Usually fast and widely available
- What can DFT do?
  - Atomic and cell geometries (fixed V,P)
  - Formation energy
  - Properties related to ground state



#### Comments on DFT - downsides

- DFT also has some downsides we will see this in practice
  - Difficulties with capturing systems dispersion
  - Band gap problem LUMO cannot be associated with KS orbitals (derivative discontinuity, deviation from piecewise linearity)
  - No magic solutions :( we must make approximations and test our decisions

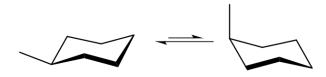


#### Comments on orbitals

- Orbitals are spatial wave functions, essentially probability amplitudes
- In practice, our calculated orbitals are mathematical formulations which approximate our true system
- Different calculations of orbitals (KS orbitals, canoncial HF orbitals, Dyson orbitals) can disagree qualitatively
- Be careful with overinterpreting orbitals



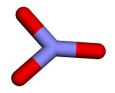
## Exercise 6.1 - Methylcyclohexane A-value



- You will perform calculations with HF and MP2 and different DFT functionals, add results to collaborative spreadsheet (linked also on Moodle)
- Points of comparison:
  - $\psi$  or  $\rho$  based?
  - how accurate (w.r.t. experimental reference)?
  - computational time
- DFT is a world on its own depending on the functional chosen you can go from cheap, very off calculations to expensive and more reliable ones



## Exercise 6.2 - Geometric properties: NO<sub>3</sub> radical



- Calculate N-O bond lengths and O-N-O bond angles
  - Experiments:  $D_3^h$ , N-O 1.24 Å and O-N-O 120°
- Compare results (HF, MP2 vs DFT)
- You will visualize the KS orbitals what can they tell us?

Exercise Session 6

- Changes in the electronic structure between different species
- General size, shape of expected one-electron orbitals (hopefully)



## Exercise 6 - Tips

- DFT will be further explored during lectures and the next exercises
- Here we used as reference papers that can be useful for further understanding DFT1, DFT2, orbitals
- Calculations for Exercise 6.1 will be done in a collaborative way to speed up the exercise, add your results to collaborative spreadsheet (linked also on Moodle)