

# Data Review

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- 2 SAPT(DFT) Implementation
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  - Results
- 3 Three-Body FDDS Dispersion
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# Intermolecular Energies

► Supermolecular approach

$$E_{int} = E_{AB} - E_A - E_B$$

- Straightforward, but cannot separate different types of interactions
  - Can adopt to different electronic structure methods
  - DFT-D3 with proper functional can be both cheap and accurate
- Symmetry-Adapted Perturbation Theory
- Can give details about interactions (electrostatic, induction/polarization, exchange repulsion, London dispersion components); important in understanding their nature
  - Not as cheap as DFT-D3
  - SAPT0 is somewhat cheap, but does not include intramonomer correlation

# SAPT(DFT)

- ▶ Attempt to include intramonomer correlation in a cheap way
- ▶ Replaces HF orbitals with KS orbitals
- ▶ Needs to consider orbital response for dispersion terms
- ▶ Exchange-dispersion term needs to be estimated from scaling
- ▶ Investigate the accuracy and efficiency of SAPT(DFT)

## Three-Body Interaction

- ▶ Crucial in computing high accuracy lattice energies (5% for benzene)
- ▶ DFT-D3 does not perform well for three-body interaction
- ▶ MP2.5 scales as  $O(N^6)$ , MP2 is  $O(N^5)$  but lacks three-body dispersion
- ▶ Three-body dispersion can be implemented with SAPT(DFT) in  $O(N^5)$
- ▶ Combine MP2 with SAPT(DFT) dispersion to model three-body interaction

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## Idea of SAPT(DFT)

- ▶ SAPT energy in orders of interaction and fluctuation potentials;  $n$  denotes order in  $V$  and  $k, l$  for  $W_A, W_B$

$$H = F_A + F_B + V + W_A + W_B$$

$$E_{int} = \sum_{n=1}^{\infty} \sum_{k=0}^{\infty} \sum_{l=0}^{\infty} \left( E_{pol}^{(nkl)} + E_{exch}^{(nkl)} \right)$$

- ▶ SAPT0:  $n = 2, k = l = 0$ , no intramonomer correlation,  $O(N^5)$  cost
- ▶ Many-body SAPT:  $k + l \geq 2$ ,  $O(N^6)$  or higher cost
- ▶ SAPT(DFT): Use Kohn-Sham operator  $K_{A,B}$  instead of Fock operator  $F_{A,B}$ ,  $O(N^5)$  cost
- ▶ Primitive SAPT(DFT) works well on 1st-order terms, but not 2nd-order terms (especially dispersion). Needs orbital response for these terms



## Coupled Dispersion Energy

- Uncoupled dispersion energy in terms of frequency-dependent density susceptibility (FDDS):

$$\begin{aligned}
 E_{disp,u}^{(2)} &= -4 \sum_{ia \in A, jb \in B} \frac{|(ia|jb)|^2}{\epsilon_{ij}^{ab}} \\
 &= -\frac{1}{2\pi} \int_0^\infty d\omega \int d\mathbf{r}_A d\mathbf{r}'_A d\mathbf{r}_B d\mathbf{r}'_B \\
 &\quad \frac{1}{|\mathbf{r}_A - \mathbf{r}_B|} \frac{1}{|\mathbf{r}'_A - \mathbf{r}'_B|} \chi_0^A(\mathbf{r}_A, \mathbf{r}'_A | i\omega) \chi_0^B(\mathbf{r}_B, \mathbf{r}'_B | i\omega)
 \end{aligned}$$

- Kohn-Sham DFT constructs a fictitious system of non-interacting particles, which reproduces the density and energy of the real electronic system
- Kohn-Sham FDDS does not reflect the correct response properties of the electronic system



## Dispersion Term

- ▶ Coupled FDDS from solving TDDFT equations:

$$\chi = \chi_0 + \chi_0 \mathbf{S}^{-1} \mathbf{W} (\mathbf{S} - \chi_0 \mathbf{S}^{-1} \mathbf{W})^{-1} \chi_0$$

- ▶  $\mathbf{S}$  and  $\mathbf{W}$  corresponds to Coulomb metric and xc kernel
- ▶ Pure ALDA kernel good for pure GGA functional, but not for hybrid functional
- ▶ Exact exchange in  $v_{xc} \rightarrow$  increased  $\epsilon_{ij}^{ab} \rightarrow$  decreased  $E_{disp}^{(2)}$

$$E_{disp,u}^{(2)} = -4 \sum_{ia \in A, jb \in B} \frac{|(ia|jb)|^2}{\epsilon_{ij}^{ab}}$$

- ▶ Hybrid ALDA kernel to compensate, or localized HF (LHF) exchange to avoid increase in  $\epsilon_{ij}^{ab}$

# Coupled FDDS with hybrid kernel

$$\chi = \chi'_0 + (\chi'_0 \mathbf{S}^{-1} \mathbf{W} + \mathbf{K}') [\mathbf{S} - (\chi'_0 \mathbf{S}^{-1} \mathbf{W} + \mathbf{K}')]^{-1} \chi'_0$$

$$\mathbf{K}' = [-\xi \mathbf{K}_1(\lambda d) - \xi \mathbf{K}_2(\lambda d) + \xi^2 \mathbf{K}_{21}(\lambda)] (\mathbf{R}^T)^{-1} \mathbf{S}$$

$$[\mathbf{K}_1(\lambda d)]_{PQ} = (P|ar)\lambda_{ar}d_{ar}[(aa'|rr') + (ar'|a'r)](a'r'|\mathbf{Q}|Q)$$

$$[\mathbf{K}_2(\lambda d)]_{PQ} = (P|ar)\lambda_{ar}d_{ar}[(aa'|rr') - (ar'|a'r)](a'r'|\mathbf{Q}|Q)$$

$$\begin{aligned} [\mathbf{K}_{21}(\lambda)]_{PQ} = & (P|ar)\lambda_{ar}[(aa''|rr'') - (ar''|a''r)] \\ & [(a'a''|r'r'') - (a'r''|a''r')](a'r'|\mathbf{Q}|Q) \end{aligned}$$

$$[\mathbf{K}'_2(\lambda)]_{PQ} = (P|ar)\lambda_{ar}[(aa'|rr') - (ar'|a'r)](a'r'|\mathbf{Q})$$

$$\chi'_0 = \chi_0 - \xi \mathbf{K}_2(\lambda)$$

$$(ar|Q) = (ar|\mathbf{Q}|P)(P|\mathbf{R}|Q)$$



## Equation for $E_{disp}^{(2)}$

- ▶ Coupled Kohn-Sham (CKS) FDDS reflects correct response properties of electrons
- ▶  $O(N^5)$  scaling is limited to forming  $\mathbf{K}_1$ ,  $\mathbf{K}_2$ ,  $\mathbf{K}_{21}$  and  $\mathbf{K}'_2$ . These quantities are frequency-dependent, but could store frequency-independent intermediates from the  $O(N^5)$  contractions on disk, and the frequency-dependent contractions are only  $O(N^4)$
- ▶  $E_{disp}^{(2)}$  from coupled FDDS: (Integration is approximated by Gauss-Legendre quadrature)

$$E_{disp,r}^{(2)} = -\frac{1}{2\pi} \int_0^\infty d\omega \int d\mathbf{r}_A d\mathbf{r}'_A d\mathbf{r}_B d\mathbf{r}'_B \frac{1}{|\mathbf{r}_A - \mathbf{r}_B|} \frac{1}{|\mathbf{r}'_A - \mathbf{r}'_B|} \chi^A(\mathbf{r}_A, \mathbf{r}'_A | i\omega) \chi^B(\mathbf{r}_B, \mathbf{r}'_B | i\omega)$$



## Exchange-Dispersion Term

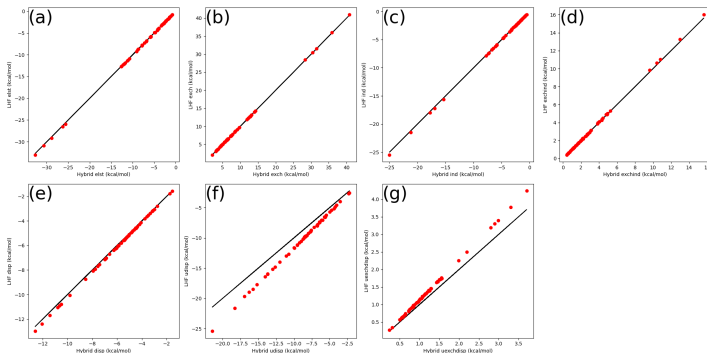
- ▶ Explicit coupled exchange-dispersion not trivial to implement; currently working on this
- ▶ Estimate from scaling uncoupled exchange-dispersion; Scale with ratio in dispersion term or with pre-fitted (with S22×5) fixed factor

$$\tilde{E}_{exch-disp,r}^{(2)} = E_{exch-disp,u}^{(2)} \cdot \frac{E_{disp,r}^{(2)}}{E_{disp,u}^{(2)}}$$

$$\tilde{E}_{exch-disp,r}^{(2)} = \alpha \cdot E_{exch-disp,u}^{(2)} (\alpha = 0.686)$$

- ▶ The value above is fitted from  $E_{exch-disp,u}^{(2)}$  with LHF orbitals
- ▶ Non-LHF orbitals have greater o-v gaps and smaller  $E_{disp,u}^{(2)}$ , needs to re-fit with non-LHF results

# LHF vs non-LHF orbitals



Hybrid vs. LHF values in kcal/mol for each term for S66 data set: (a)  $E_{elst}^{(1)}$ , (b)  $E_{exch}^{(1)}$ , (c)  $E_{ind}^{(2)}$ , (d)  $E_{exch-ind}^{(2)}$ , (e)  $E_{disp,r}^{(2)}$ , (f)  $E_{disp,u}^{(2)}$ , (g)  $E_{exch-disp,u}^{(2)}$

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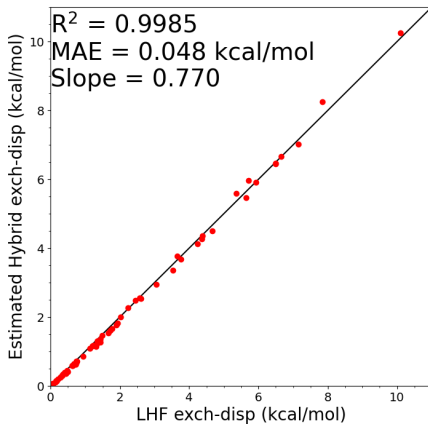
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## Exchange-Dispersion Refitting

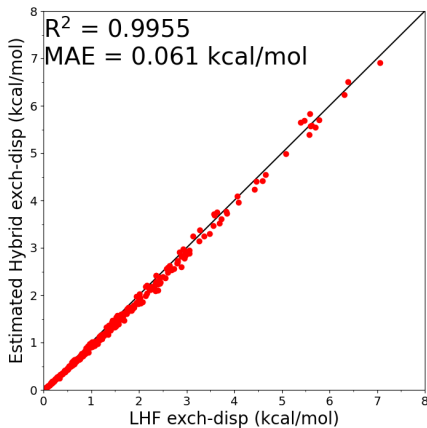
- ▶ Need to fit the uncoupled  $E_{exch-disp,r}^{(2)}$  with non-LHF orbitals on the coupled LHF orbital values (implemented in Molpro)
- ▶ Assuming coupled LHF and non-LHF orbital  $E_{exch-disp,r}^{(2)}$  from the behavior of  $E_{disp,r}^{(2)}$
- ▶ Exchange-related components depend heavily on distance between monomers, sets like S22×5 and S66×8 would be preferred
- ▶ Determine the scaling factor with S22×5, validate with S66×8



## S22×5 Fitting Results



# S66×8 Validating Results





## Termwise results

- ▶ Compared the SAPT(DFT)/aug-cc-pVTZ results of our code to SAPT(CCSD)/aug-cc-pVTZ results from Korona S2<sup>1</sup>
- ▶ Also comparing the results for S66 with SAPT2+3(CCD) $\delta$ MP2/aug-cc-pVTZ as reference. Also added SAPT0/aug-cc-pVDZ, SAPT2+/aug-cc-pVDZ and SAPT2+(3) $\delta$ MP2/aug-cc-pVTZ into comparison as side-reference.
- ▶ Errors of each system with respect to reference shown as vertical lines
- ▶ Mean absolute error (MAE) and mean unsigned relative error (MURE) listed for S2. MAE indicated by black box in the diagram
- ▶ Color scheme for S66 systems: Hydrogen-bonded (HB, red), mixed-influence (MX, green), dispersion-dominated (DD, blue)

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<sup>1</sup>T. Korona, Mol. Phys. **111**, 3705 (2013).

# Korona S2 Results

Method	MAE	MURE	Error Distribution					
			4	OB	1	0	1	UB
Electrostatics								
SAPT(DFT) hybrid	0.112	2.39						
SAPT(DFT) LHF	0.114	3.68						
SAPT0	0.520	8.61						
Exchange								
SAPT(DFT) hybrid	0.251	3.38						
SAPT(DFT) LHF	0.258	3.09						
SAPT0	1.757	12.88						
Induction								
SAPT(DFT) hybrid	0.148	2.79						
SAPT(DFT) LHF	0.192	2.97						
SAPT0	1.993	16.83						
Exchange-Induction								
SAPT(DFT) hybrid	0.144	4.03						
SAPT(DFT) LHF	0.165	4.76						
SAPT0	1.551	26.80						

# Korona S2 Results

Method	MAE	MURE	Error Distribution							
			4	OB	1	0	1	UB	4	
Dispersion										
SAPT(DFT) hybrid	0.175	3.68								
SAPT(DFT) LHF	0.141	2.77								
SAPT(DFT) non-hybrid	0.326	9.58								
SAPT0	0.811	24.86								
Exchange-Dispersion										
SAPT(DFT) hybrid	0.062	12.47								
SAPT(DFT) LHF	0.039	3.25								
SAPT0	0.265	36.11								
Total										
SAPT(DFT) hybrid	0.155	4.98								
SAPT(DFT) LHF	0.189	4.17								
SAPT(DFT) hon-hybrid	0.244	10.64								
SAPT0	1.237	19.63								

# S66 Results

Method	Total	HB	MX	DD	Error Distribution						
					4	OB	1	0	1	UB	4
Electrostatics											
SAPT(DFT) hybrid	0.374	0.556	0.177	0.311							
SAPT(DFT) LHF	0.423	0.666	0.196	0.319							
SAPT0	0.613	1.034	0.439	0.297							
SAPT2+	0.236	0.270	0.136	0.263							
SAPT2+(3) $\delta$ MP2	0.000	0.000	0.000	0.000							
Exchange											
SAPT(DFT) hybrid	0.886	1.127	0.426	0.926							
SAPT(DFT) LHF	0.886	1.121	0.431	0.928							
SAPT0	0.675	0.942	0.263	0.658							
SAPT2+	0.337	0.467	0.222	0.277							
SAPT2+(3) $\delta$ MP2	0.000	0.000	0.000	0.000							
Induction											
SAPT(DFT) hybrid	0.211	0.201	0.212	0.220							
SAPT(DFT) LHF	0.224	0.223	0.223	0.225							
SAPT0	0.241	0.200	0.261	0.271							
SAPT2+	0.327	0.384	0.250	0.318							
SAPT2+(3) $\delta$ MP2	0.152	0.179	0.121	0.145							

# S66 Results

Method	Total	HB	MX	DD	Error Distribution						
					4	OB	1	0	1	UB	4
Dispersion											
SAPT(DFT) hybrid	0.370	0.260	0.219	0.573							
SAPT(DFT) LHF	0.308	0.200	0.173	0.499							
SAPT(DFT) non-hybrid	0.635	0.581	0.419	0.822							
SAPT0	0.443	0.862	0.162	0.195							
SAPT2+	0.235	0.397	0.169	0.115							
SAPT2+(3) $\delta$ MP2	0.093	0.129	0.056	0.080							
Total											
SAPT(DFT) hybrid	0.334	0.588	0.107	0.217							
SAPT(DFT) LHF	0.234	0.382	0.046	0.199							
SAPT(DFT) non-hybrid	0.604	0.955	0.389	0.385							
SAPT0	0.990	1.197	0.692	0.965							
SAPT2+	0.230	0.235	0.138	0.280							
SAPT2+(3) $\delta$ MP2	0.105	0.056	0.082	0.169							

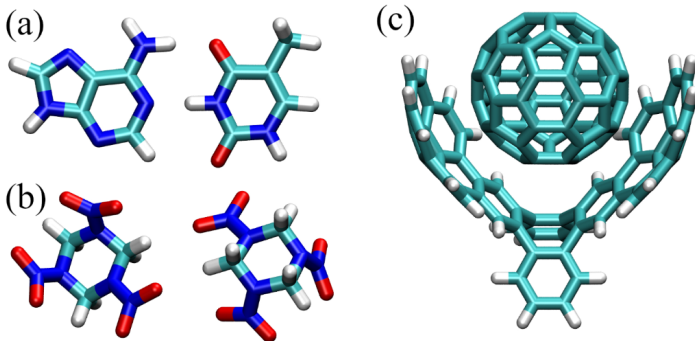


## Timing Performance

- ▶ Analyzing breakdown wall times for subroutines in SAPT(DFT) for a few systems with 500–3000 basis functions
- ▶ Using Intel Core i7-6800K processor with 6 cores for Watson-Crick adenine-thymine complex and RDX dimer
- ▶ Using Intel Core i9-10980XE processor with 18 cores for  $C_{60}$ –buckycatcher ( $N_{bf} = 3012$ ), completed entire calculation in 4.03 days
- ▶ Contribution of  $O(N^5)$  dispersion terms not dominant for smaller systems
- ▶ Cost of SCF (HF/DFT) calculations and induction terms are usually non-negligible



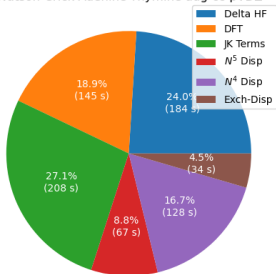
# Timing Systems



Dimer systems for timing: (a) Watson-Crick adenine-thymine complex, (b) RDX dimer, (c)  $C_{60}$ -buckycatcher complex.

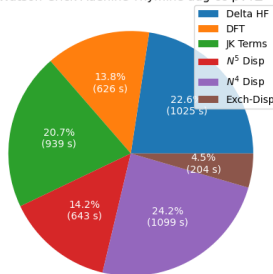
# Watson-Crick Adenine-Thymine

(a) Watson-Crick Adenine-Thymine aug-cc-pVDZ



$$N_{bf} = 536$$

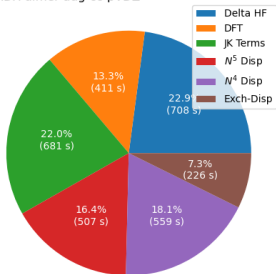
(b) Watson-Crick Adenine-Thymine aug-cc-pVTZ



$$N_{bf} = 1127$$

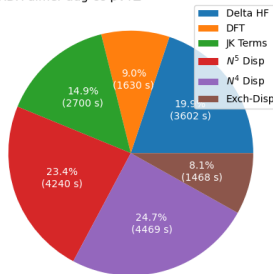
# RDX Dimer

(c) RDX dimer aug-cc-pVDZ



$$N_{bf} = 798$$

(d) RDX dimer aug-cc-pVTZ



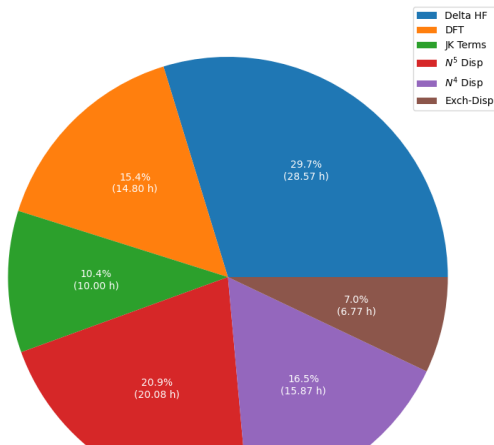
$$N_{bf} = 1656$$

## Comparison with LHF Approach

- ▶ Comparison of subroutine wall times between hybrid xc kernel approach implemented in Psi4 1.4 and LHF approach implemented in Molpro 2019.2
- ▶ Some subroutines does not exist or not included by default in the Molpro DFT-SAPT program

Subroutine	Psi4 hybrid time (h)	Molpro LHF time (h)
Delta HF	0.96	N/A
DFT	0.45	2.29
xc kernel	0.08	4.17
$O(N^5)$ objects formation	2.35	N/A
$E_{disp}^{(2)}$ time integration	0.37	3.59
$E_{exch-disp}^{(2)}$	0.41	1.99
Total	5.03	12.80

# $C_{60}$ –Buckycatcher Complex



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## 3B-69 Benchmark Set

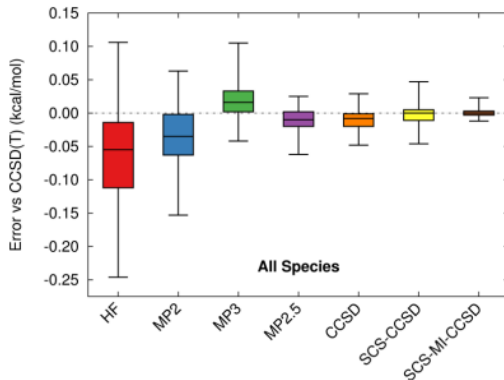
- ▶ Benchmark set for 3-body interaction energies
- ▶ 69 trimers extracted from 23 different molecular crystal structures (3 each)
- ▶ Used focal point approach to obtain CCSD(T) (and other wavefunction method) energies

$$E = E^{HF}(aQZ) + \Delta E^{MP2}(aTZ/aQZ) + \Delta E^{CCSD(T)}(aDZ)$$

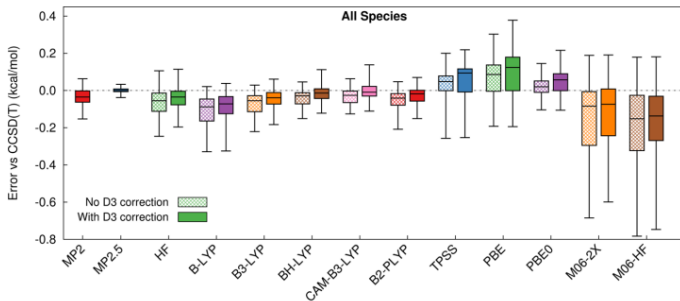
- ▶ Assessing accuracy for various wavefunction and DFT methods
- ▶ Recommended MP2.5 and SCS-MI-CCSD, both  $O(N^6)$
- ▶ Trying to assess performance of MP2 + FDDS dispersion for 3B-69 systems



# 3B-69 Wavefunctional Methods



# 3B-69 DFT Methods



## New Set from X23

- ▶ Sampling trimer geometries from X23 crystal structures
- ▶ Trying to include trimer with different intermolecular distances and alignment
- ▶ Aiming to serve as a "three-body version" of  $S22 \times 5 / S66 \times 8$
- ▶ Distance: Geometry mean of 3 pairwise closest contact distance
- ▶ Alignment: Angles of the COM triangle; mainly looking at the greatest angle

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## Three-Body Dispersion

- ▶ Three-body dispersion energy in terms of FDDS, analogous to the two-body dispersion:

$$E_{disp,r}^{(3)} = -\frac{1}{\pi} \int_0^\infty d\omega \int d\mathbf{r}_A d\mathbf{r}'_A d\mathbf{r}_B d\mathbf{r}'_B d\mathbf{r}_C d\mathbf{r}'_C \frac{1}{|\mathbf{r}_A - \mathbf{r}_B|} \frac{1}{|\mathbf{r}'_A - \mathbf{r}_C|} \frac{1}{|\mathbf{r}'_B - \mathbf{r}'_C|} \chi^A(\mathbf{r}_A, \mathbf{r}'_A | i\omega) \chi^B(\mathbf{r}_B, \mathbf{r}'_B | i\omega) \chi^C(\mathbf{r}_B, \mathbf{r}'_B | i\omega)$$

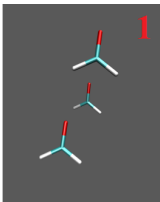
- ▶ Transform from position space into density-fitting auxiliary basis space:

$$E_{disp,r}^{(3)} = \int_0^\infty d\omega \text{Tr} (\mathbf{S}^{-1} \boldsymbol{\chi}^A \mathbf{S}^{-1} \boldsymbol{\chi}^B \mathbf{S}^{-1} \boldsymbol{\chi}^C)$$

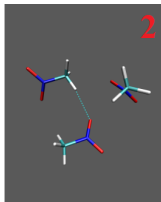
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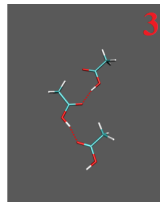
## 3B-69 Systems



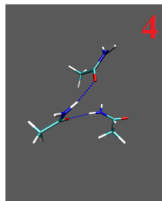
formaldehyde b



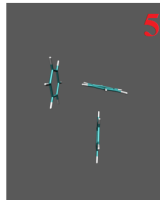
nitromethane a



acetic acid b



acetamide a



benzene c

## 3B-69 Results

- ▶ Total interaction energies in kcal/mol
- ▶ FDDS(aDZ) and FDDS(aTZ) correspond to estimated total IE with MP2 + FDDS

System	CCSD(T)	FDDS(aDZ)	FDDS(aTZ)	MP2	MP2.5
<b>1</b>	0.181	0.207	0.210	0.161	0.179
<b>2</b>	-0.122	-0.069	-0.065	-0.178	-0.143
<b>3</b>	-0.922	-0.905	-0.904	-0.937	-0.913
<b>4</b>	-0.089	-0.003	-0.003	-0.239	-0.151
<b>5</b>	-0.027	0.002	0.003	-0.061	-0.023

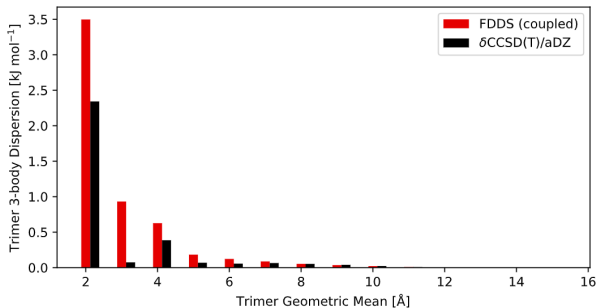


## 3B-69 Results

- ▶ Estimated 3-body dispersion energies comparison
- ▶ CCSD(T) corresponds to dispersion energy estimated by  $E^{CCSD(T)} - E^{MP2}$

System	CCSD(T)	FDDS(aDZ)	FDDS(aTZ)
<b>1</b>	0.020	0.046	0.049
<b>2</b>	0.056	0.109	0.113
<b>3</b>	0.015	0.032	0.033
<b>4</b>	0.150	0.236	0.242
<b>5</b>	0.034	0.063	0.064

# Distance Dependence



# Distance Dependence

