

Theoretical Investigations of Hydrogen Bonding Interactions of (E)-1-(1H-Benzo[d]imidazol-2-yl)-3-Phenylprop-2-en-1-one Monomers and Dimers: NBO, QTAIM and NCI Study

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Abstract: Hydrogen bonding is an essential interaction in nature and plays a crucial role in many formations of materials and biological processes, requiring a deeper understanding of its formation. Benzimidazole is an important structural unit found in a large number of natural and pharmacologically active molecules. In the present work, the electronic structures and properties and relative stabilities of a series of (E)-1-(1H-benzo[d]imidazol-2-yl)-3-phenylprop-2-en-1-one monomers and dimers have been studied by density functional theory using B3LYP 6-31+G (d, p) calculation level. The strengths of the noncovalent interactions have been analyzed in terms of the QTAIM analysis, NCI analysis and natural bond orbital approaches. It was found that the dimers are formed by double N-H...O hydrogen bond. QTAIM analysis proved the presence of intramolecular hydrogen bond in monomers and coexistence of intramolecular and intermolecular hydrogen bond in dimers. Frequency analysis shows that intermolecular N-H...O interactions are proper hydrogen bond while intramolecular C-H...N, C-H...O, C-H...H-C interactions are improper hydrogen bond. NBO and NCI analyses confirm the existence of hydrogen bonds in the studied monomers and dimers. The presence of weakly electron acceptor group on benzene ring favors the total interaction energy of dimerization.

Keywords: NBO, QTAIM, NCI, Benzimidazole, Electron Density, Hydrogen Bond

1. Introduction

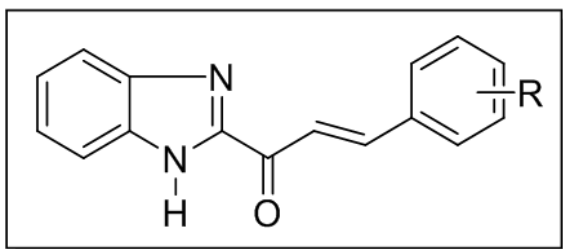
Benzimidazole (BZim) is a heterocyclic aromatic organic compound. It is an important drug and a preferred structure in medicinal chemistry [1-3]. The benzimidazole ring is one of the bioactive heterocyclic compounds that have a range of biological activities such as antivirals (anti-HIV), anticancer drugs, antibacterials, antifungals and many others [4-9]. Benzimidazole derivatives are associated with various types

of pharmacokinetics and pharmacodynamics properties. The most important benzimidazole compound in nature is N-ribosyldimethylbenzimidazole, which serves as an axial ligand of cobalt in vitamin B12 [10]. In addition, some benzimidazoles are used in coordination chemistry [11-14], optoelectronics [15].

The hydrogen bond is one of the fundamental interactions that plays a key role in many fields of chemistry, physics and biology [16-19]. It affects the stability of many important molecular structures such as water [20, 21] and DNA [22].

Knowledge of hydrogen bonding strength is essential for physical, chemical and biological applications [23]. In addition, a slight difference in molecular structure may play a role in the strength of hydrogen bonds [24, 25]. In most cases, the formation of $X-H\cdots Y$ bonds leads to a weakening of the X-H bond, accompanied by an elongation of the bond and a concomitant decrease of the vibrational stretching frequency of the X-H bond relative to the non-interactive bond, these bonds are called proper or Red shifted hydrogen bond. However, there are also many examples in which the stretching frequency of the X-H bond increases with the contraction of the bond length during the formation of the hydrogen bond, these are improper or Blue shifted hydrogen bond [26].

The electronic foundations of the improper hydrogen bond were analyzed by Alabugin *et al* [27]. The structural reorganization of the X-H bond observed for proper and improper hydrogen bonds results from the balance between hyperconjugation ($n_X \rightarrow \sigma_{X-H}^*$) which weakens the X-H bond and rehybridization which strengthens X-H bond [27]. Improper hydrogen bonds are observed when the hyperconjugative interaction is relatively low. When the hyperconjugative interaction is low and the hybrid orbital of the X atom of the X-H bond is able to undergo hybridization, rehybridization prevails, resulting in a shortening of the X-H bond and a blue shift of the X-H vibrational stretching frequency. The presence of electron-rich centers (N, O) and hydrogen atoms linked by a covalent bond to nitrogen and carbon atoms offers the possibility of different types of non-covalent bonds such as $C-H\cdots N$, $C-H\cdots O$, $N-H\cdots O$ (Figure 1).



MONOMER	R	DIMER
M1	H	$D_1=M_1\cdots M_1$
M2	4-Cl	$D_2=M_2\cdots M_2$
M3	4-F	$D_3=M_3\cdots M_3$
M4	4-NO ₂	$D_4=M_4\cdots M_4$
M5	2-Cl, 5-NO ₂	$D_5=M_5\cdots M_5$

Figure 1. Structure of M1-M5 monomers and D1-D5 dimers studied.

In this study, we applied quantum chemistry methods to determine the interaction energies and understand the nature of the different hydrogen bonds of BZims monomers and dimers. The topological properties, hyperconjugative interactions between donor and proton acceptor of different $X-H\cdots Y$ hydrogen bonds formed have been determined. The QTAIM analysis was used to assess the strength and nature of intramolecular and intermolecular hydrogen bonds. We also performed the NBO (natural bond orbital) and NCI (non-covalent interaction) analyses that suggest the existence of hydrogen bonds in the studied systems. Our calculations

will give a theoretical overview on the possibility of dimerization of BZims.

2. Materials, Methods and Parameters

2.1. Computational Methods and Materials

The monomers M1-M5 and dimers D1-D5 (Figure 1) were optimized using B3LYP [28, 29] calculation level with a 6-31+G (d, p) basis set. The optimization was performed with Gaussian 09 [30] software on Intel i7 2600 processors. In a further step, the topological analysis of the electron density was performed from the "formatted checkpoint file" obtained by the DFT calculation with AIMALL software [31]. The NBO orbital analysis method [32] (NBO: Natural orbital bond) incorporated in the Gaussian 09 software was used for NBO analyses. NCI analysis was performed with Multiwfn software [33] and NCI analysis surfaces were represented with Chemcraft software [34].

2.2. Interaction Energy

The interaction energies of the complexes were determined by the difference between the total energy of the complex and the energies of the isolated monomers, then corrected by taking account the basic set superposition error (BSSE) using the Boys-Bernardi counterpoise procedure [35] and the ZPE:

$$E_{int} = E^{Dimer} - (2 * E^{monomer}) + bsse + \Delta ZPE$$

$$\text{With } \Delta ZPE = ZPE^{Dimer} - 2 * ZPE^{monomer}$$

2.3. NBO Analysis

NBO analysis is based on an approach of transformation of the multielectronic wave functions of the molecules in a localized form that corresponds to the elements with single center (single lone pair LP) and two centered orbitals (natural bonding σ π and antibonding $\sigma^* \pi^*$ respectively). It provides an in-depth understanding of intra- and intermolecular orbital interactions in molecules between the NBOs of filled donors and the NBOs of empty acceptors [36]. For each donor NBO (i) and acceptor NBO (j), the stabilization energy associated with delocalization $i \rightarrow j$ can be estimated as follows:

$$E^{(2)} = \Delta E_{ij} = q_i \frac{F(i, j)}{\epsilon_i \epsilon_j}$$

Where q_i is the electron density in the donor's orbital, $F(i, j)$ is the non-diagonal element of the Fock matrix and ϵ_i and ϵ_j are the energies of the occupied orbitals i and vacant j . By analyzing the interactions between the different acceptor and donor NBOs as well as the resulting stabilizing energy, clear information is obtained on the origin of the stabilization of a molecule. If the stabilization energy $E^{(2)}$ associated with an interaction is high, the extent of stabilization will be greater.

2.4. QTAIM Analysis

Bader's Quantum Theory of Atoms in Molecules (QTAIM) is a useful tool for characterizing topological properties of chemical bonds [37]. The presence of chemical bonds between atoms and interatomic interactions is revealed by the presence of bond critical points (BCPs). The QTAIM method provides information on the electron density of a system that governs properties at BCPs. The QTAIM theory gives information on variations of electron density due to the formation of bonds or complexes [37]. Several parameters exist within the framework of the QTAIM theory, we have among others the electron density $\rho(r)$, its Laplacian $\nabla^2\rho(r)$, the potential local energy density $V(r)$, the local kinetic energy density $G(r)$ and the total kinetic energy density $H(r)$ at bond critical points (BCPs). In general, the parameter values of the QTAIM analysis at BCPs can describe the covalent or electrostatic nature of non-covalent interactions. According to Rozas et al [38] the interactions at the different BCPs can be classified as follows: for strong hydrogen bonds $\nabla^2\rho_{BCP} < 0, H_{BCP} < 0$; for medium hydrogen bonds $\nabla^2\rho_{BCP} > 0, H_{BCP} < 0$; for weak hydrogen bonds $\nabla^2\rho_{BCP} > 0, H_{BCP} > 0$. Also, the ratio $-\frac{G(r)}{V(r)}$ also allows a better understanding of the nature of the interaction at the different BCPs [39-41]. If $-\frac{G(r)}{V(r)} > 1$ the bond is considered non-covalent. If $0,5 < -\frac{G(r)}{V(r)} < 1$ we have a partially covalent character of the bond. If $-\frac{G(r)}{V(r)} < 0,5$ we have a closed-shell interaction. Espinosa [42] proposed a relationship between the energy of the hydrogen bond E_{HB} and the potential energy density (V_{BCP}) at BCPs: $E_{HB} = \frac{1}{2}|V_{BCP}|$.

In addition, the ellipticity ε to the different BCPs is defined as $\varepsilon = \frac{\lambda_1}{\lambda_2} - 1$; λ_1 and λ_2 are eigenvalues of the Hessian of the electron density at BCP. This quantity estimates the extent to which the electron density is deformed in one direction relative to another. Ellipticity provides a measure of π or σ character of chemical bonds. A high value of ellipticity ($\varepsilon > 0.1$) indicates a π character of the bond while a lower value reflects a σ character of the bond [37]. Ellipticity reflects the stability of the bonds [39,43].

2.5. NCI Analysis

NCI (Non covalent interaction) analysis based on the correlation between the Reduced Gradient Density (RDG) and electron density $\rho(r)$ was recently developed by Yang et al [44, 45]. The reduced gradient density is a fundamental dimensionless quantity of DFT, which is used to describe the deviation from a homogeneous distribution of electron density [44, 46]. To some degree, the NCI analysis method can be considered as an extension of the AIM analysis [45]. The non-covalent interaction analysis method provides graphical visualization of molecule regions where non-covalent interactions occur and has demonstrated its ability to distinguish hydrogen bonds, van der Waals interactions and repulsive interactions [47]. Recently, the NCI-RDG analysis detected a low interaction compared to the AIM

analysis [48]. To determine the presence of low interactions, $RDG = f(\text{sign}(\lambda_2) \times \rho(r))$ graphs were generated, where $\text{sign}(\lambda_2) \times \rho(r)$ is the product of the electronic density $\rho(r)$ by the sign of the second eigenvalue of the hessian. The sign value $\text{sign}(\lambda_2) \times \rho(r)$ is useful for predicting the nature of non-covalent interaction. For an attractive interaction $\text{sign}(\lambda_2) \times \rho(r) < 0$ (hydrogen bond) and for repulsive interactions $\text{sign}(\lambda_2) \times \rho(r) > 0$ (steric hindrance). For Van Der Waals interactions $\text{sign}(\lambda_2) \times \rho(r) = 0$.

3. Results and Discussions

3.1. Study of Monomers

The electronic properties obtained from QTAIM analysis, the geometrical parameters as well as the hyperconjugative interactions between the orbitals involved in the intramolecular hydrogen bonds of the M1-M5 monomers are presented in Tables 1 and 2. The interactions $C-H \cdots N$, $C-H \cdots H-C$ and $C-H \cdots Cl$ were revealed by the QTAIM analysis (Figure 2). From the data in Table 1, it can be seen that the distances $d(H \cdots Y)$ and $d(X \cdots Y)$ are between 2.20881-2.27482 Å and 3.03051-3.05551 Å respectively. Also, the linearity angles $\angle XHY$ are between 127.080-130.636°. These geometrical parameters indicate the presence of weak intramolecular hydrogen bond [18, 49, 50]. The $C-H \cdots N$ bond have an electron density $\rho(r)$ in the range of 0.016617-0.018498 ea_0^{-3} , a Laplacian $\nabla^2\rho(r)$ (0.052978-0.058614 ea_0^{-5}) and a positive value of the total kinetic energy density $H(r)$ at different BCPs. These observations confirm the presence of weak intramolecular hydrogen bonds in M1-M5 monomers [51-54]. In addition, for all $C-H \cdots N$ bond, the ratio $-\frac{G(r)}{V(r)} > 1$ implies that the $C-H \cdots N$ hydrogen bonds are non-covalent character [40, 41]. The interaction energies obtained for the $C-H \cdots N$ bonds are in the order of 3.3414375-3.8873625 kcal/mol, these values confirm that these hydrogen bonds are weak [42]. Nevertheless, these interactions are stable because we have a low ellipticity value at the different BCPs.

The $C-H \cdots Cl$ hydrogen bond is observed in the M4 molecule. At the Bcp of $C-H \cdots Cl$ bond we have an electron density $\rho(r) = 0,014863$ ea_0^{-3} , its Laplacian $\nabla^2\rho(r) = 0,05587$ ea_0^{-5} and a positive value of the total kinetic energy density $H(r) = 0.002198$ au. The $C-H \cdots Cl$ hydrogen bond is weak because $\rho(r) > 0$, $\nabla^2\rho(r) > 0$ and $H(r) > 0$. In addition, the ratio $-\frac{G(r)}{V(r)} > 1$ allows us to affirm that the $C-H \cdots Cl$ interaction is non-covalent in nature. Except for the M4 monomer, $C-H \cdots H-C$ interactions were observed in all monomers studied. At the Bcps of the different interactions $C-H \cdots H-C$ the electron density $\rho(r)$ varies between 0.008791-0.009004 ea_0^{-3} the Laplacian $\nabla^2\rho(r)$ is between 0.02-0.002492 au.

Table 1. Topological parameters at bonds critical points of intramolecular interactions obtained by AIM analysis of M1-M5 monomers.

	$X-H\cdots Y$	$\rho(r)_{X-H\cdots Y}$	$\nabla^2\rho(r)_{X-H\cdots Y}$	$V(r)$	$G(r)$	$H(r)$	$-\frac{G(r)}{V(r)}$	$L(r)$	E_{HB}^*	ΣE_{HB}	$\varepsilon(r)$
M1	$C-H\cdots N$	0.016346	0.052978	-0.01065	0.011946	0.001298	1.122	-0.01325	3.3414375	4.825475	0.015129
	$C-H\cdots H-C$	0.009004	0.038893	-0.00473	0.007225	0.002499	1.529	-0.00972	1.4840375		1.235422
M2	$C-H\cdots N$	0.016715	0.05402	-0.01094	0.01222	0.001284	1.117	-0.01351	3.432425	4.8788125	0.014019
	$C-H\cdots H-C$	0.008813	0.038364	-0.00461	0.007098	0.002492	1.541	-0.00959	1.4463875		1.434104
M3	$C-H\cdots N$	0.016757	0.054062	-0.01096	0.012236	0.001279	1.117	-0.01352	3.4387	4.8913625	0.014245
	$C-H\cdots H-C$	0.008791	0.039479	-0.00463	0.007249	0.00262	1.566	-0.00987	1.4526625		3.146135
M4	$C-H\cdots N$	0.018498	0.058614	-0.01239	0.013521	0.001133	1.091	-0.01465	3.8873625	6.88995	0.001775
	$C-H\cdots Cl$	0.014863	0.05587	-0.00957	0.011769	0.002198	1.230	-0.01396	3.0025875		0.042812
M5	$C-H\cdots H-C$	0.008872	0.038579	-0.00465	0.007146	0.002	1.5368	-0.00965	1.4589375	4.8662625	1.370549
	$C-H\cdots N$	0.016617	0.053709	-0.01086	0.012143	0.001	1.118	-0.01343	3.407325		0.014142

Table 2. Geometrical parameters, hyperconjugative interactions, stabilization energy $E^{(2)}$ (kcal/mol) and charge transferred CT (me).

	geometrical parameters					NBO Interaction		CT
	$X-H\cdots Y$	$d(X-H)$	$d(H\cdots Y)$	$d(X\cdots Y)$	$\angle XHY$	Transitions	$E^{(2)}$	
M1	$C-H\cdots N$	1.08660	2.27482	3.05551	127.080	$n_{N13}^{(1)} \rightarrow \sigma_{C18-H20}^*$	3.77	CT
M2	$C-H\cdots N$	1.08654	2.26372	3.04660	127.239	$n_{N13}^{(1)} \rightarrow \sigma_{C18-H20}^*$	3.92	6.87
M3	$C-H\cdots N$	1.08644	2.26282	3.04352	127.048	$n_{N13}^{(1)} \rightarrow \sigma_{C18-H20}^*$	3.97	7.144
M4	$C-H\cdots N$	1.08627	2.20881	3.03051	130.636	$n_{N13}^{(1)} \rightarrow \sigma_{C18-H20}^*$	4.91	7.144
	$C-H\cdots Cl$	1.08127	2.50603	3.21292	122.021	$n_{Cl30}^{(1)} \rightarrow \sigma_{C17-H19}^*$	2.57	9.096
M5	$C-H\cdots N$	1.08660	2.26653	3.04957	127.258	$n_{N13}^{(1)} \rightarrow \sigma_{C18-H20}^*$	3.90	5

$C-H\cdots H-C$ interactions are weak and non-covalent in nature because $\rho(r)>0$, $\nabla^2\rho(r)>0$, $H(r)>0$ and $-\frac{G(r)}{V(r)}>1$ (Table 1). QTAIM parameters $\rho(r)$, $\nabla^2\rho(r)$ and E_{int} observed at the different Bcps of the $C-H\cdots H-C$ interactions are smaller than those of the $C-H\cdots N$ and $C-H\cdots Cl$ interactions. The ellipticity values $\varepsilon(r)$ of the $C-H\cdots H-C$ interactions are large than those of the $C-H\cdots N$ and $C-H\cdots Cl$ interactions. As a result, $C-H\cdots H-C$ interactions are less stable and weak than $C-H\cdots N$ and $C-H\cdots Cl$ interactions.

NBO analysis was performed on the M1-M5 monomers and the results obtained are shown in Table 2. NBO analysis revealed the presence of $C-H\cdots N$ and $C-H\cdots Cl$ interactions. Hyperconjugative interactions $n_X \rightarrow \sigma_{X-H}^*$ with stabilization energy ranging from 2.57-4.91 kcal/mol and corresponding transferred charges ranging from 5-9.096 me were obtained confirming the presence of weak intramolecular hydrogen bonds $C-H\cdots N$ and $C-H\cdots Cl$ [27].

3.2. Dimer Studies

3.2.1. Stability, Geometrical Parameters and NBO Analysis of Complexes

D2 dimer with an interaction energy of -13.0604 kcal/mol is the most stable of the series. It is characterized by the presence of a hydrogen bond pair $N-H\cdots O$ (figure 3). The intermolecular distances $d(N\cdots O)$ and $d(H\cdots O)$ are in 2.87592 and 1.88814 Å respectively. The $\angle NHO$ angle of the $N-H\cdots O$ bond is separated by 18.849° from linearity. The variation in the length of the $N-H$ bond $\Delta d(N-H)>0$ implies an elongation its length during complexation. The vibrational stretching frequency ν_{N-H} of the monomer is 3646.9195 cm^{-1} while that of the dimer is 3435.3275 cm^{-1} .

As a result, the variations in the vibrational stretch frequencies are negative. Based on the findings, we can say that the elongation of the N-H bonds in the dimer is done in concert with a decrease in the elongation vibration frequencies of $\Delta\nu_{N-H}=-211.5919\text{ cm}^{-1}$ to the low frequencies (red shift). Thus in the D2 dimer the $N-H\cdots O$ bonds are proper hydrogen bonds [26, 55]. In Table 5 we have presented the non-covalent interactions observed in the dimers as well as the stabilization energy, the charge transferred and the variation of charges of the atoms X, H and Y involved in the interactions $X-H\cdots Y$. The NBO analysis detected the presence of the $N-H\cdots O$ hydrogen bond characterized by the transfer of the lone pair of oxygen atom to the antibonding orbital σ^* of the $N-H$ bond. Hyperconjugative interactions $n_{O47}^{(1)} \rightarrow \sigma_{N11-H12}^*$, $n_{O16}^{(1)} \rightarrow \sigma_{N42-H43}^*$ on the one hand and $n_{O47}^{(2)} \rightarrow \sigma_{N11-H12}^*$, $n_{O16}^{(2)} \rightarrow \sigma_{N42-H43}^*$ on the other hand with stabilization energies $E^{(2)}$ of 10.12 and 6.12 kcal/mol and charges transferred CT of 13.75 and 12.552 me respectively prove the presence of $N-H\cdots O$ hydrogen bonds [27].

The second stable complex in the series is the D5 dimer with an interaction energy $E_{int}=-11.94$ kcal/mol. This value is slightly lower than that of D2 dimer ($E_{int}=-13.0604$ kcal/mol). The D5 dimer is characterized by the presence of a double $N-H\cdots O$ hydrogen bond (figure 3), an intermolecular $C-H\cdots O$ hydrogen bond and intramolecular hydrogen bonds $C-H\cdots N$ and $C-H\cdots H-C$. The $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ interactions (figure 4) were revealed by the AIM analysis of the complexes (see 3.2.3 section). the intermolecular distances $d(H\cdots Y)$ and $d(X\cdots Y)$ of the $N-H\cdots O$, $C-H\cdots O$ and $C-H\cdots N$ interactions are between 1.87984-3.18209 Å and 2.87331-3.00995 Å respectively. The length $d(H\cdots H)$ of the $C_{17}-H_{19}\cdots_{17}H-C_{23}$ interaction is

2.16299 Å. The bonding angles of the intermolecular interactions $N-H\cdots O$, $C-H\cdots O$ and $C-H\cdots N$ are between 116.863 - 163.414° . The bonding angles of the $N-H\cdots O$ interactions are closer to linearity than those of the $C-H\cdots O$ and $C-H\cdots N$ interactions. Also, we observe that the intermolecular distances $d(X\cdots Y)$ and $d(H\cdots Y)$ are smaller in $N-H\cdots O$ interactions. As a result, $N-H\cdots O$ interactions are stronger than $C-H\cdots O$ and $C-H\cdots N$ interactions. The $N-H$ bonds of the $N-H\cdots O$ interactions undergo an elongation $\Delta d(N-H)$ of 10.58 and 10.18 mÅ while the $C-H$ bonds of the $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ interactions undergo a contraction in the range of 0.02-0.93 mÅ. The elongation of the $N-H$ bond coincides with a shift of vibrational stretching frequency to low frequencies (red shift). Concerning the interactions $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ a slight shift of the vibrational stretching frequency to high frequencies (blue shift) is observed (Table 4). These observations allow us to conclude that we have two types of non-covalent interactions in the D5 dimer; The $N-H\cdots O$ interaction characterized by an elongation of the $N-H$ bond ($\Delta d(N-H) > 0$) and a large red displacement of the vibrational stretching frequency is a proper hydrogen bond. The $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ interactions characterized by a

blue shift of the vibrational stretching frequencies are improper hydrogen bonds [26, 55]

The NBO analysis of D5 dimer and the NBO descriptors of hydrogen bond are recorded in Table 4. Except for the $C_{17}-H_{19}\cdots_{27}H-C_{23}$ interaction, the NBO analysis detected the presence of observed hydrogen bonds.

$n_O \rightarrow \sigma_{N-H}^*$ interactions between the lone pair of hydrogen bond acceptor and the antibonding orbital σ_{N-H}^* of hydrogen bond donor were revealed. The highest stabilization energy is obtained in the interaction $N_{42}-H_{43}\cdots O_{16}$ ($n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{42}-H_{43}}^*$) with an energy $E^{(2)}=11.06$ kcal/mol and a charge transferred $CT=12.637$ me followed by the interaction $N_{11}-H_{12}\cdots O_{47}$ ($n_{O_{47}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*$) with a stabilization energy $E^{(2)}=9.28$ kcal/mol and a charge transferred $CT=12.637$ me. For improper hydrogen bonds, the NBO analysis revealed the presence $C_{18}-H_{20}\cdots N_{13}$ ($n_{N_{13}}^{(1)} \rightarrow \sigma_{C_{18}-H_{20}}^*$, $E^{(2)}=4.37$ kcal/mol $CT=7.993$ me) and $C-H\cdots O$ ($n_{O_{47}}^{(1)} \rightarrow \sigma_{C_2-H_7}^*$, $E^{(2)}=0.05$ kcal/mol, $CT=0.07159$ me), however the stabilizing energy and charge transferred of these interactions are low compared to the $N_{11}-H_{12}\cdots O_{47}$ interaction.

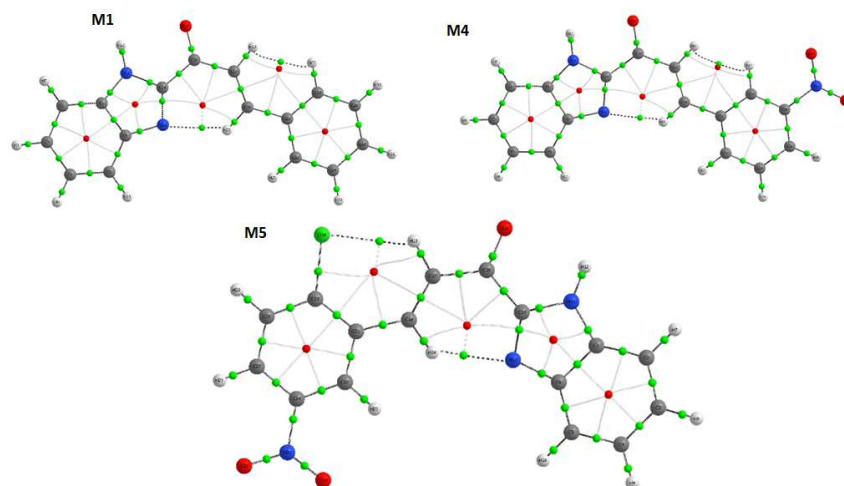


Figure 2. AIM molecular graphs of M1, M4 and M5 monomers at B3LYP 6-31+G (d, p) small green spheres are bonds critical points (Bcps).

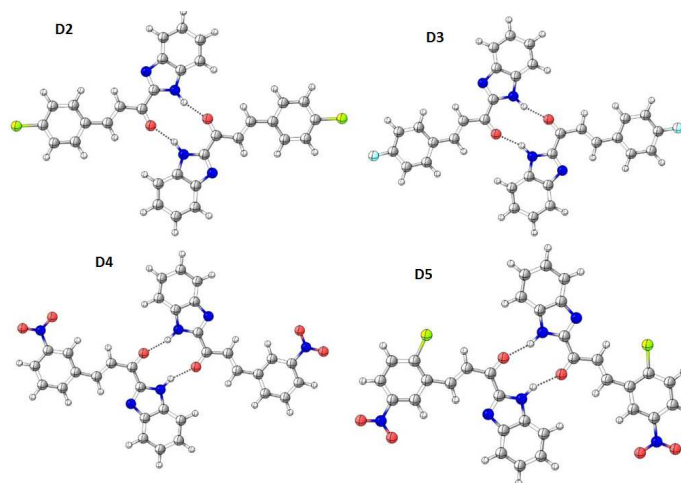


Figure 3. Optimized structures of D2-D5 dimers at B3LYP 6-31+G (d, p).

Table 3. Interaction energy $E_{int}(\text{kcal/mol})$ of **D1-D5** dimers corrected by BSSE and ZPE.

	$E_{Dimer}(\text{au})$	$E_{Monomer}(\text{au})$	$\Delta ZPE(\text{au})$	$BSSE(\text{au})$	$E_{int}(\text{au})$	E_{int}
D1	-1603.45	-801.715	0.001413	0.001237	-0.016	-10.0422
D2	-2522.64	-1261.31	0.001147	0.001287	-0.02081	-13.0604
D3	-2012.47	-1006.22	0.001394	0.001251	-0.01846	-11.581
D4	-2931.64	-1465.81	0.001296	0.001244	-0.01879	-11.7923
D5	-1801.93	-900.955	0.00127	0.001249	-0.01904	-11.9464

Table 4. Geometrical parameters of $X-H\cdots Y$, $C-H\cdots H-C$ interactions (distances en Å, angle $\angle XHY$ (deg), change in bond length Δd (mÅ) and stretching frequencies ν (cm^{-1}) and its shifts $\Delta\nu$ (cm^{-1}) for $C-H$ and $N-H$ bond.

	$X-H\cdots Y$ $C-H\cdots H-C$	$d(H\cdots Y)$ $d(H\cdots H)$	$d(X\cdots Y)$	$\angle XHY$	$\Delta d(X-H)$ $\Delta d(C-H)$	$\nu_{X-H}^{monomer}$ $\nu_{C-H}^{monomer}$	ν_{X-H}^{dimer} ν_{C-H}^{dimer}	$\Delta\nu_{X-H}$ $\Delta\nu_{C-H}$
D1	$N_{11}-H_{12}\cdots O_{47}$	1.88661	2.86706	159.887	10.19	3646.4497	3464.7658	-181.6839
	$N_{42}-H_{43}\cdots O_{16}$	1.88650	2.86694	159.883	10.19	3646.4497	3464.7658	-181.6839
	$C_{49}-H_{51}\cdots N_{44}$	2.23882	3.01365	126.449	-0.32	3196.0143	3198.2511	2.2368
	$C-H_{58}\cdots H-C$	2.16137			-0.02 0.06	3180.628	3187.9229	7.2949
	$C_{18}-H_{20}\cdots N_{13}$	2.23854	3.01343	126.453	-0.31	3196.0143	3198.2272	2.2129
	$C_{17}-H_{19}\cdots {}_{17}H-C_{23}$	2.16141			0.02 0.06	3180.628	3187.9156	7.2876
D2	$C_{33}-H_{38}\cdots O_{16}$	3.20049	3.80527	116.151	-0.77	3206.6944	3216.294	9.5996
	$C_2-H_7\cdots O_{47}$	3.2006	3.80545	116.151	-0.77	3206.6944	3216.2923	9.5979
	$N_{11}-H_{12}\cdots O_{47}$	1.88814	2.87592	161.532	11.18	3646.9194	3435.3275	-211.5919
	$N_{42}-H_{43}\cdots O_{16}$	1.88814	2.87592	161.532	11.18	3646.9194	3435.3275	-211.5919
	$N_{11}-H_{12}\cdots O_{49}$	1.91787	2.89576	159.212	10.58	3646.0336	3454.6697	-191.3639
	$N_{44}-H_{45}\cdots O_{16}$	1.88837	2.88423	164.416	9.29	3646.0336	3475.1849	-170.8487
D3	$C_2-H_7\cdots O_{49}$	3.19843	3.81206	116.777	-0.89	3198.1771	3199.25	1.0729
	$C_{18}-H_{20}\cdots N_{13}$	2.23393	3.00558	126.159	0.26	3197.4879	3194.7291	-2.7588
	$C_{17}-H_{19}\cdots {}_{27}H-C_{23}$	2.16603			-0.1 -0.07	3193.0372	3245.2407	52.2035
D4	$N_{11}-H_{12}\cdots O_{49}$	1.87241	2.86785	163.983	10.26	3647.2604	3455.0778	-192.1826
	$N_{44}-H_{45}\cdots O_{16}$	1.90681	2.88528	159.410	10.32	3647.2604	3455.0778	-192.1826
	$C_{35}-H_{40}\cdots O_{16}$	3.19900	3.80798	116.457	-0.97	3208.6194	3219.0874	10.468
	$C_{17}-H_{19}\cdots {}_{Cl}I_{30}$	2.53155	3.20379	119.509	-2.19	3258.3331	3288.7712	30.4381
	$C_{51}-H_{53}\cdots N_{46}$	2.17823	2.99290	129.873	-0.21	3185.9744	3186.8562	0.8816
	$C_{50}-H_{52}\cdots {}_{Cl}I_{63}$	2.50563	3.10349	122.398	-0.07	3258.3331	3257.3667	-0.9664
D5	$N_{11}-H_{12}\cdots O_{47}$	1.91387	2.88999	158.799	10.59	3646.7178	3451.4549	-195.2629
	$N_{42}-H_{43}\cdots O_{16}$	1.87984	2.87331	163.414	10.18	3646.7178	3451.4549	-195.2629
	$C_2-H_7\cdots O_{47}$	3.18209	3.79751	116.863	-0.93	3214.7217	3217.4894	2.7677
	$C_{18}-H_{20}\cdots N_{13}$	2.23457	3.00995	126.481	-0.31	3196.3516	3197.94	1.5884
	$C_{17}-H_{19}\cdots {}_{27}H-C_{23}$	2.16289			-0.04 -0.02	3180.4875	3180.9779	0.4904

$$\Delta\nu = \nu_{X-H}^{dimer} - \nu_{X-H}^{monomer}, \Delta d = d_{X-H}^{dimer} - d_{X-H}^{monomer}$$

Table 5. Hyperconjugative interactions, stabilization energies $E^{(2)}$ (kcal/mol) and charge transferred CT (me) as well as the variation of the charges of atoms X, Y and H with dimerization.

Dimers	Contact	Interaction	$E^{(2)}$	$\Sigma E^{(2)}$	CT	Δq_X	Δq_H	Δq_Y
D1	$N_{11}-H_{12}\cdots O_{47}$	$n_{O_{47}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	10.41		14.32	0.0539	-0.0152	0.0122
	$N_{11}-H_{12}\cdots O_{47}$	$n_{O_{47}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	5.57		11.74	0.0539	-0.0152	0.0122
	$N_{42}-H_{43}\cdots O_{16}$	$n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	10.41		14.32	0.0539	-0.0152	0.0122
	$N_{42}-H_{43}\cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	5.57		11.74	0.0539	-0.0152	0.0122
	$C_2-H_7\cdots O_{47}$	$n_{O_{47}}^{(2)} \rightarrow \sigma_{C_2-H_7}^*$	0.06	40.66	0.11834	0.03216	-0.01012	0.0539
	$C_{33}-H_{38}\cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{C_{33}-H_{38}}^*$	0.06		0.11834	0.03203	-0.01012	0.0539
	$C_{49}-H_{51}\cdots N_{44}$	$n_{N_{44}}^{(1)} \rightarrow \sigma_{C_{49}-H_{51}}^*$	4.29		7.705	-0.01551	-0.00185	0.00543
	$C_{18}-H_{20}\cdots N_{13}$	$n_{N_{13}}^{(1)} \rightarrow \sigma_{C_{18}-H_{20}}^*$	4.29		7.705	-0.01551	-0.00185	0.00543
	$N_{11}-H_{12}\cdots O_{47}$	$n_{O_{47}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	10.12		13.7468	0.06631	-0.0153	0.00029
	$N_{11}-H_{12}\cdots O_{47}$	$n_{O_{47}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	6.12	32.48	12.552	0.06631	-0.0153	0.00029
D2	$N_{42}-H_{43}\cdots O_{16}$	$n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	10.12		13.7468	0.06631	-0.0153	0.00029

Dimers	Contact	Interaction	$E^{(2)}$	$\Sigma E^{(2)}$	CT	Δq_x	Δq_H	Δq_y
D3	$N_{42} - H_{43} \cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	6.12	44.24	12.552	0.06631	-0.0153	0.00029
	$N_{44} - H_{45} \cdots O_{16}$	$n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{44}-H_{45}}^*$	13.14		17.6975	0.00076	-0.01588	0.06656
	$N_{44} - H_{45} \cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{44}-H_{45}}^*$	6.82		13.9593	0.00076	-0.01588	0.06656
	$N_{11} - H_{12} \cdots O_{49}$	$n_{O_{49}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	11.27		15.5	0.01394	-0.01758	0.08213
	$N_{11} - H_{12} \cdots O_{49}$	$n_{O_{49}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	7.96		16.11	0.01394	-0.01758	0.08213
	$C_2 - H_7 \cdots O_{49}$						
	$C_{18} - H_{20} \cdots N_{13}$	$n_{N_{13}}^{(1)} \rightarrow \sigma_{C_{18}-H_{20}}^*$	5.05		9.198	-0.00788	-0.00523	0.00023
	$N_{11} - H_{12} \cdots O_{49}$	$n_{O_{49}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	11.31		15.5	0.01552	-0.01386	0.059595
	$N_{11} - H_{12} \cdots O_{49}$	$n_{O_{49}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	5.17		10.68	0.01552	-0.01386	0.059595
	$N_{44} - H_{45} \cdots O_{16}$	$n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{44}-H_{45}}^*$	9.57		13.185	0.01581	-0.01437	0.06804
D4	$N_{44} - H_{45} \cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{44}-H_{45}}^*$	5.61	43.51	11.35	0.01581	-0.01437	0.06804
	$C_{17} - H_{19} \cdots Cl_{30}$	$n_{Cl_{30}}^{(1)} \rightarrow \sigma_{C_{17}-H_{19}}^*$	0.67		0.756	0.00392	-0.01466	0.00386
		$n_{Cl_{30}}^{(2)} \rightarrow \sigma_{C_{17}-H_{19}}^*$	2.35		4.4157			
	$C_{50} - H_{52} \cdots Cl_{63}$	$n_{Cl_{63}}^{(1)} \rightarrow \sigma_{C_{50}-H_{52}}^*$	0.73		0.834	0.00382	0.00325	0.00264
		$n_{Cl_{63}}^{(2)} \rightarrow \sigma_{C_{50}-H_{52}}^*$	2.65		5.246			
	$C_{51} - H_{53} \cdots N_{46}$	$n_{N_{46}}^{(1)} \rightarrow \sigma_{C_{51}-H_{53}}^*$	5.51		10.394	0.00455	-0.00181	-0.02136
	$C_{35} - H_{40} \cdots O_{16}$						
	$N_{11} - H_{12} \cdots O_{47}$	$n_{O_{47}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	9.28		12.637	0.01242	-0.01558	0.06314
	$N_{11} - H_{12} \cdots O_{47}$	$n_{O_{47}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*$	5.71		12.05	0.01242	-0.01558	0.06314
	$N_{42} - H_{43} \cdots O_{16}$	$n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	11.06		15.2	0.01831	-0.01482	0.05552
D5	$N_{42} - H_{43} \cdots O_{16}$	$n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{42}-H_{43}}^*$	5.10	35.57	10.58	0.01831	-0.01482	0.05552
	$C_2 - H_7 \cdots O_{47}$	$n_{O_{47}}^{(1)} \rightarrow \sigma_{C_2-H_7}^*$	0.05		0.07159	0.03285	-0.00649	0.06314
	$C_{18} - H_{20} \cdots N_{13}$	$n_{N_{13}}^{(1)} \rightarrow \sigma_{C_{18}-H_{20}}^*$	4.37		7.993	-0.01647	0.0018	0.00395

D4 Dimer is the third most stable complex in the series. It has an interaction energy $E_{int} = -11.7933 \text{ kcal/mol}$. This dimer is formed by a double $N-H \cdots O$ hydrogen bond (figure 3). In addition, the QTAIM analysis detected the presence of $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions (figure 4) (see 3.2.3 section). The intermolecular distances $d(X \cdots Y)$, $d(H \cdots Y)$ of the interactions observed in the D4 dimer range from 1.87241-3.19900 Å and 2.86785-3.80798 Å respectively. Regarding the linearity angles, they vary between 116.457-163.983°. The distances of the intermolecular interactions $N-H \cdots O$ are generally smaller than those of the intramolecular interactions $C-H \cdots Cl$ and $C-H \cdots N$ and the bonding angles $\angle NHO$ are closer to linearity. The $N-H \cdots O$ intermolecular interaction are consequently stronger than the $C-H \cdots Cl$ and $C-H \cdots N$ intramolecular interaction. With complexation, we observe an elongation of the $N-H$ bonds of the $N-H \cdots O$ interactions ($\Delta d(N-H) > 0$) while in the $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions we observe a shortening of the $C-H$ bonds ($\Delta d(N-H) < 0$). The elongation of the $N-H$ bonds is combined with a red shift of the vibrational stretching frequency $\Delta \nu_{N-H} = -192.1826 \text{ cm}^{-1}$ and the contraction of the $C-H$ bonds of the $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions is combined with a small blue shift of the vibrational stretching frequencies $\Delta \nu_{C-H}$ which is comprised between 0.8816-30.4381 cm^{-1} . It can be seen that the blue shifts of the vibrational stretching frequencies of the $C-H$ bonds are small compared to the shifts of the $N-H$ bonds. This low blue shift of the $C-H$ bonds implies that the $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions are weaker than the $N-H \cdots O$ interactions. However, the contraction of the $C-H$ bond in the $C_{50} - H_{52} \cdots Cl_{63}$ interaction is associated with a low red shift of the stretch vibration frequency

$\Delta \nu_{C-H} = -0.9664 \text{ cm}^{-1}$. Consequently, $N-H \cdots O$ interactions are proper hydrogen bonds and $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions are improper hydrogen bonds [26, 55].

The elongation of the $X-H$ bond in the $X-H \cdots Y$ interaction is generally attributed to the charge transferred of the interaction $n_Y \rightarrow \sigma_{X-H}^*$ between the lone pair of the proton acceptor and the anti-bonding orbital σ_{X-H}^* of the donor [24]. The consequence of these charges transferred are measured by the second-order stabilization energy $E^{(2)}$ obtained by the NBO analysis presented in Table 5. We observe that the largest stabilization energies $E^{(2)}$ and charge transferred are obtained for the interaction $N_{11} - H_{12} \cdots O_{49}$ ($n_{O_{49}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*$, $E^{(2)} = 11.31 \text{ kcal/mol}$, $CT = 15.52 \text{ e}$) followed by $N_{44} - H_{45} \cdots O_{16}$ interaction ($n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{44}-H_{45}}^*$, $E^{(2)} = 9.57 \text{ kcal/mol}$, $CT = 12.637 \text{ e}$). The total stabilization energy $\Sigma E^{(2)}$ in the dimer is 31.66 kcal/mol which implies a large charge transfer in the D4 dimer through $N-H \cdots O$, $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots N$ interactions.

With an interaction energy of -11.581 kcal/mol the D3 dimer is the fourth stable complex studied. It is also characterized by a double $N-H \cdots O$ interaction (figure 3) but also by the $C-H \cdots O$, $C-H \cdots Cl$ and $C-H \cdots H-C$ interactions (figure 4). revealed by the QTAIM analysis (see 3.2.3 section). The intermolecular distances of the $N-H \cdots O$ interactions ($d(H \cdots O) = 1.87241-1.90681 \text{ Å}$; $d(N \cdots O) = 3.00558-3.81206 \text{ Å}$) (Table 4) are lower than that of $C-H \cdots O$ and $C-H \cdots N$ interactions ($d(H \cdots Y) = 2.23393-3.19843 \text{ Å}$; $d(X \cdots Y) = 3.00558-3.81206 \text{ Å}$). The intermolecular distance $d(H \cdots H)$ of the $C-H \cdots H-C$ interaction is 2.16603 Å. the $\angle NHO$ bonding angles of the $N-H \cdots O$ interactions (159.410-163.983°) are closer to linearity than those of the $C-H \cdots O$ and $C-H \cdots N$

interactions (116.777-126.159°). In addition, the analysis of the bond lengths shows that the $N-H$ and $C-H$ bonds of the $N_{11}-H_{12}\cdots O_{49}$, $N_{44}-H_{45}\cdots O_{16}$ and $C_{18}-H_{20}\cdots N_{13}$ interactions undergo elongation ($\Delta d(C-H) > 0, \Delta d(N-H) > 0$) while that of the $C_2-H_7\cdots O_{49}$ interaction undergoes contraction with complexation. This variation in bond length is combined with a red shift of vibrational stretching frequency of the $N-H$ and $C-H$ bonds of the $N_{11}-H_{12}\cdots O_{49}$, $N_{44}-H_{45}\cdots O_{16}$ and $C_{18}-H_{20}\cdots N_{13}$ interactions. However, a blue shift of vibrational stretching frequency of the $C-H$ bond is observed in the $C_2-H_7\cdots O_{49}$ and $C_{17}-H_{19}\cdots {}_{17}H-C_{23}$ interactions. With these observations we can affirm that the interactions $N-H\cdots O$ and $C-H\cdots N$ are proper hydrogen bonds and the interactions $C-H\cdots O$ and $C-H\cdots H-C$ are improper hydrogen bonds [26, 55].

The NBO analysis was performed and the stabilization energy due to hyperconjugative interactions $n_Y \rightarrow \sigma_{X-H}^*$ of the $X-H\cdots Y$ interactions are recorded in Table 5. This analysis clearly shows the presence of hydrogen bonds $N_{44}-H_{45}\cdots O_{16}$ ($n_{O_{16}}^{(1)} \rightarrow \sigma_{N_{44}-H_{45}}^*, \Delta E^{(2)} = 13,14 \text{ kcal/mol}$; $n_{O_{16}}^{(2)} \rightarrow \sigma_{N_{44}-H_{45}}^*, \Delta E^{(2)} = 6,82 \text{ kcal/mol}$), $N_{11}-H_{12}\cdots O_{49}$ ($n_{O_{49}}^{(1)} \rightarrow \sigma_{N_{11}-H_{12}}^*, E^{(2)} = 11.27 \text{ kcal/mol}$; $n_{O_{49}}^{(2)} \rightarrow \sigma_{N_{11}-H_{12}}^*, \Delta E^{(2)} = 7.96 \text{ kcal/mol}$) and $C_{18}-H_{20}\cdots N_{13}$ ($n_{N_{13}}^{(1)} \rightarrow \sigma_{C_{18}-H_{20}}^*, E^{(2)} = 5.05 \text{ kcal/mol}$) [27]. These observed interactions due to the overlap between the free doublet of the atom n_Y and the antibonding orbital σ_{X-H}^* have as total stabilization energies $\Sigma E^{(2)}$ equal to 44.24 kcal/mol for the $N_{44}-H_{45}\cdots O_{16}$, $N_{11}-H_{12}\cdots O_{49}$ and $C_{18}-H_{20}\cdots N_{13}$ interactions, therefore there is a strong charge transfer in the D3 dimer which contributes to its stability.

The least stable complex is D1 dimer. It has an interaction energy $E_{int} = -10.0422 \text{ kcal/mol}$ significantly lower than that of D2-D5 dimers. It is stabilized by a $N-H\cdots O$ hydrogen bond (figure 3). and the $C-H\cdots N$, $C-H\cdots O$ and $C-H\cdots H-C$ hydrogen bond revealed by the AIM analysis (figure 4). The intermolecular distances $d(H\cdots Y)$, $d(X\cdots Y)$ are between 1.88661-3.20049 Å and 2.86694-3.80545 Å respectively. The corresponding linearity angles $\angle XHY$ vary between 116.151-159.887°. During complexation, the $N-H$ bonds of the $N-H\cdots O$ interactions lengthen ($\Delta d(N-H) > 0$) with a red shift of their vibrational stretching frequency of $\Delta \nu_{N-H} = -181.6839 \text{ cm}^{-1}$ while the $C-H$ bonds of the $C-H\cdots N$ and $C-H\cdots O$

interactions shorten with a blue shift $\Delta \nu_{C-H}$ (between $2.2129-9.5996 \text{ cm}^{-1}$) of their vibrational stretching frequencies. $N-H\cdots O$ interactions are therefore stronger than $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ interactions. Consequently, the $N-H\cdots O$ interactions are proper hydrogen bonds and the $C-H\cdots O$, $C-H\cdots N$ and $C-H\cdots H-C$ interactions are improper hydrogen bonds [26, 55]. The hyperconjugative interactions $n_X \rightarrow \sigma_{X-H}^*$ obtained by the NBO analysis allow to confirm the presence of the $N-H\cdots O$, $C-H\cdots O$ and $C-H\cdots N$ hydrogen bonds in the D1 dimer [27]. The corresponding stabilization energies vary between 0.06-10.41 kcal/mol. The total stabilization energy $\Sigma E^{(2)}$ is 40.66 kcal/mol, confirming the presence of a high charge transfer through the $N-H\cdots O$, $C-H\cdots O$ and $C-H\cdots N$ hydrogen bonds.

The formation of the $X-H\cdots Y$ hydrogen bond is related to the increase of the positive charge of the H atom and the negative charge of the X and Y atoms. The change in atomic charges resulting from the complexing of the D1-D5 dimers is recorded in Table 5. Apart from $C_{50}-H_{52}\cdots Cl_{63}$ (D4) and $C_{18}-H_{20}\cdots N_{13}$ (D5) hydrogen bonds, the atomic charges of hydrogen in the $X-H\cdots Y$ interactions of the D1-D5 dimers increase ($\Delta q_H < 0$) while that of the N, O and C atoms decrease ($\Delta q_O > 0, \Delta q_C > 0, \Delta q_N > 0$) with complexation. Nevertheless, in the intramolecular bonds of the D1 and D4 dimers, the increase in the positive charge of the hydrogen atom H is combined with an increase of the charge of the nitrogen atom. In the interactions $X-H\cdots Y$ of the D1-D5 dimers, the increase of positive charge of the hydrogen atom H and that of the negative charge of X atom observed is a signature of the hydrogen bond [27, 56]. The lone pairs of the Y atoms ($Y \equiv O, N, Cl$) involved in hyperconjugative interactions $n_Y \rightarrow \sigma_{X-H}^*$ are hybridized sp^m ($m \in 0.74 - 99.99$) with electron densities ranging from 1.90394 to 1.99144 e. These electron densities less than 2 show that these lone pairs are delocalized, particularly in the antibonding orbital σ^* , which have an electron density between 0.01378 e and 0.04825 e. the electron densities of the antibonding orbitals of the N-H, C-H bonds and the S character of the nitrogen and carbon atoms ($(N-H, (C-H)$ (Table 7) increase with complexity. this increase in the S character of nitrogen and carbon atoms indicates a rehybridization of the orbitals of the latter during the formation of the hydrogen bonds $N-H\cdots O$, $C-H\cdots O$ and $C-H\cdots N$ [27].

Table 6. Lone pairs(n_Y) and antibonding orbitals (σ_{X-H}^*) hybridization states, their respective electronic densities as well as the variation of the electronic density of the antibonding orbitals σ_{X-H}^* upon complexation.

dimers	Contact	Donor		Acceptor					
		n_Y		σ_{X-H}^*					
		Hybridization	ED	Hybridization	ED	$\Delta \sigma^*$	$ED_X(\%)$	$ED_H(\%)$	
D ₁	$N_{11}-H_{12}\cdots O_{47}$	$sp^{0.74}$	1.96219	$\sigma_{N_{11}-H_{12}}^* = 0.4892(sp^{2.01})_N - 0.8722(s)_H$	0.04462	0.02385	23.93	76.07	
	$N_{11}-H_{12}\cdots O_{47}$	$sp^{47.79}$	1.90394						
	$N_{42}-H_{43}\cdots O_{16}$	$sp^{0.74}$	1.96219	$\sigma_{N_{42}-H_{43}}^* = 0.4892(sp^{2.01})_N - 0.8722(s)_H$	0.04462	0.02385	23.93	76.07	
	$N_{42}-H_{43}\cdots O_{16}$	$sp^{47.79}$	1.90394						
	$C_{49}-H_{51}\cdots N_{44}$	$sp^{1.97}$	1.92099	$\sigma_{C_{49}-H_{51}}^* = 0.5999(sp^{1.97})_C - 0.8001(s)_H$	0.02305	0.00125	35.99	64.01	
	$C_{18}-H_{20}\cdots N_{13}$	$sp^{1.97}$	1.92099	$\sigma_{C_{18}-H_{20}}^* = 0.5999(sp^{1.97})_C - 0.8001(s)_H$	0.02305	0.00125	35.99	64.01	
	$C_2-H_7\cdots O_{47}$	$sp^{0.47}$	1.96219	$\sigma_{C_2-H_7}^* = 0.6070(sp^{2.23})_C - 0.7947(s)_H$	0.01378	0.00007	36.85	63.15	

dimers	Contact	Donor		Acceptor					
		n_Y		σ_{X-H}^*					
		Hybridization	ED	Hybridization	ED	$\Delta\sigma^*$	$ED_X(\%)$	$ED_H(\%)$	
D ₂	$N_{11} - H_{12} \cdots O_{47}$	$sp^{0.74}$	1.96227	$\sigma_{N11-H12}^* = 0.4893(sp^{2.03})_N - 0.8721(s)_H$	0.04516	0.02462	23.94	76.06	
	$N_{11} - H_{12} \cdots O_{47}$	$sp^{55.38}$	1.89373						
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{0.74}$	1.96227	$\sigma_{N42-H43}^* = 0.4893(sp^{2.02})_N - 0.8721(s)_H$	0.04516	0.02462	23.94	76.06	
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{55.38}$	1.89373						
	$N_{44} - H_{45} \cdots O_{16}$	$sp^{0.71}$	1.95863	$\sigma_{N44-H45}^* = 0.6165(sp^{2.02})_N - 0.7873(s)_H$	0.04746	0.02721	23.79	76.37	
	$N_{44} - H_{45} \cdots O_{16}$	$sp^{28.54}$	1.91131						
D ₃	$N_{11} - H_{12} \cdots O_{49}$	$sp^{0.70}$	1.96083	$\sigma_{N11-H12}^* = 0.4861(sp^{1.97})_N - 0.8739(s)_H$	0.04825	0.028	23.63	76.37	
	$N_{11} - H_{12} \cdots O_{49}$	$sp^{33.83}$	1.90013						
	$C_{18} - H_{20} \cdots N_{13}$	$sp^{1.88}$	1.92310	$\sigma_{C18-H20}^* = 0.5973(sp^{2.93})_C - 0.8020(s)_H$	0.02438	0.00205	35.68	63.32	
	$N_{11} - H_{12} \cdots O_{49}$	$sp^{0.74}$	1.96322	$\sigma_{N11-H12}^* = 0.4899(sp^{2.01})_N - 0.8718(s)_H$	0.04429	0.02398	24	76	
	$N_{11} - H_{12} \cdots O_{49}$	$sp^{49.50}$	1.89247						
	$N_{44} - H_{45} \cdots O_{16}$	$sp^{0.73}$	1.96322	$\sigma_{N44-H45}^* = 0.4897(sp^{2.01})_N - 0.8719(s)_H$	0.043397	0.023087	23.98	76.02	
D ₄	$N_{44} - H_{45} \cdots O_{16}$	$sp^{71.08}$	1.89247						
	$C_{17} - H_{19} \cdots Cl_{30}$	$sp^{0.21}$	1.99144	$\sigma_{C17-H19}^* = 0.5988(sp^{2.15})_C - 0.8009(s)_H$	0.02306	0.00132	35.86	64.14	
	$C_{17} - H_{19} \cdots Cl_{30}$	$sp^{99.99}$	1.96437						
	$C_{50} - H_{52} \cdots Cl_{63}$	$sp^{0.21}$	1.99133	$\sigma_{C50-H52}^* = 0.6062(sp^{2.33})_C - 0.7953(s)_H$	0.02208	0.00034	36.75	63.25	
	$C_{50} - H_{52} \cdots Cl_{63}$	$sp^{99.99}$	1.96472						
	$C_{51} - H_{53} \cdots N_{46}$	$sp^{1.98}$	1.91916	$\sigma_{C51-H53}^* = 0.5967(sp^{2.45})_C - 0.8025(s)_H$	0.02547	0.00373	35.60	64.40	
D ₅	$N_{11} - H_{12} \cdots O_{47}$	$sp^{0.74}$	1.96142	$\sigma_{N11-H12}^* = 0.4893(sp^{2.02})_N - 0.8721(s)_H$	0.04391	0.02325	23.94	76.06	
	$N_{11} - H_{12} \cdots O_{47}$	$sp^{62.96}$	1.89365						
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{0.74}$	1.96142	$\sigma_{N42-H43}^* = 0.4897(sp^{2.01})_N - 0.8719(s)_H$	0.04417	0.02351	23.98	76.02	
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{49.88}$	1.90488						
	$C_2 - H_7 \cdots O_{47}$	$sp^{0.74}$	1.96142	$\sigma_{C2-H7}^* = 0.6070(sp^{2.23})_C - 0.7947(s)_H$	0.01403	0.00035	37.02	62.98	
	$C_{18} - H_{20} \cdots N_{13}$	$sp^{1.97}$	1.92070	$\sigma_{C18-H20}^* = 0.5994(sp^{2.43})_C - 0.8004(s)_H$	0.02319	0.00115	35.93	64.07	

3.2.2. Hyperconjugation and Rehybridization

In $N - H \cdots O$ interactions, we have a rehybridization of the orbitals of nitrogen atoms with complexation. The orbitals of the nitrogen atoms are hybridized $sp^{2.20}$ in monomers while in dimers they are hybridized sp^n ($n \in 2.01-2.03$) (Table 7). The P character of the orbitals of the nitrogen atoms decreases (the S character increases) with complexation. This observation is also verified by the parameter $\Delta\%S(X)$ of the nitrogen atoms of the $N - H \cdots O$ interaction which is between 1.73-2.46. This presence of rehybridization during complexation allows to shorten the $N - H$ bond [27]. The presence of hyperconjugative interactions $n_O \rightarrow \sigma_{N-H}^*$ with a stabilization energy between 5.17-11.31 kcal/mol for $N - H \cdots O$ interactions increases the electron density in the bonding orbitals σ_{N-H}^* which are all polarized with respect to the hydrogen atom (Table 6), this polarization of the antibonding orbitals σ_{N-H}^* with respect to the H atom ensures a better overlap $n_O \rightarrow \sigma_{N-H}^*$ [56]. The observed variation in electron density $\Delta\sigma^*$ has the effect of weakening the N-H bonds by lengthening them. The elongation of the N-H bond ($\Delta d(N-H) > 0$) is clearly demonstrated in the geometry of the complexes where it is between 10.19-11.8 mÅ. Consequently, in the $N - H \cdots O$

interactions, the presence of the hyperconjugative interaction $n_O \rightarrow \sigma_{N-H}^*$ with high stabilization energy $E^{(2)}$ masks the hybridization effect, which leads to the elongation of N-H bonds and a red shift in their vibrational stretching frequency [24, 27].

Except for the D2 dimer, $C - H \cdots O$ interactions are observed in the different complexes studied. carbon atoms (C) are hybridized sp^m ($m \in 2.26-2.28$) in monomers and $sp^{2.23}$ in dimers. Also the variation in the percentage of the carbon atom $\Delta\%S(C)$ is between 0.29-0.51. The S character of the carbon atoms involved in $C - H \cdots O$ interactions increases, so there is a presence of rehybridization with complexation. Hyperconjugative interactions $n_O \rightarrow \sigma_{N-H}^*$ are also observed in the D1 and D5 complexes with a low stabilization energy value of 0.06 and 0.05 kcal/mol respectively. the interactions $n_O \rightarrow \sigma_{N-H}^*$ were not observed in D3 and D5 dimers. For $C - H \cdots O$ interactions, the presence of hyperconjugative interaction with a very low value of $E^{(2)}$ and its absence in certain complexes (D3 and D5) do not allow to mask the effects of rehybridization observed with complexation. As a result, the $C - H$ bonds are shortened with a blue shift of their vibrational stretching frequency [24, 55].

Table 7. Hybridization of the X atoms involved in the $X - H \cdots Y$ interaction in the monomers and dimers and the variation of the S character of the X atom ($\Delta\%S(X)$) with the complexation.

Complexes	Contact	Changes of hybridization $sp^n(X - H)$		$\Delta\%S(X)$
		Monomer	Dimer	
D1	$N_{11} - H_{12} \cdots O_{47}$	$sp^{2.20}$	$sp^{2.01}$	1.96
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{2.20}$	$sp^{2.01}$	1.96
	$C_2 - H_7 \cdots O_{47}$	$sp^{2.28}$	$sp^{2.23}$	0.51
	$C_{33} - H_{38} \cdots O_{16}$	$sp^{2.28}$	$sp^{2.23}$	0.51
	$C_{49} - H_{51} \cdots N_{44}$	$sp^{2.45}$	$sp^{2.23}$	0.15
	$C_{18} - H_{20} \cdots N_{13}$	$sp^{2.45}$	$sp^{2.23}$	0.15
	$C_{17} - H_{19} \cdots {}_{27}H - C_{23}$	$sp^{2.40}$	$sp^{2.45}$	-0.04
D2		$sp^{2.43}$	$sp^{2.40}$	-0.11
	$N_{11} - H_{12} \cdots O_{47}$	$sp^{2.20}$	$sp^{2.03}$	1.73
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{2.20}$	$sp^{2.03}$	1.73
	$N_{44} - H_{45} \cdots O_{16}$	$sp^{2.20}$	$sp^{1.98}$	2.25
D3	$N_{11} - H_{12} \cdots O_{49}$	$sp^{2.20}$	$sp^{1.98}$	2.46
	$C_{18} - H_{20} \cdots N_{13}$	$sp^{2.27}$	$sp^{2.39}$	0.35
	$C_2 - H_7 \cdots O_{49}$	$sp^{2.27}$	$sp^{2.23}$	0.38
		$sp^{2.29}$	$sp^{2.28}$	0.1
	$C_{17} - H_{19} \cdots {}_{27}H - C_{23}$	$sp^{2.41}$	$sp^{2.43}$	-0.11
D4	$N_{11} - H_{12} \cdots O_{49}$	$sp^{2.20}$	$sp^{2.01}$	1.94
	$N_{44} - H_{45} \cdots O_{16}$	$sp^{2.20}$	$sp^{2.01}$	1.9
	$C_{17} - H_{19} \cdots Cl_{30}$	$sp^{2.31}$	$sp^{2.15}$	1.56
	$C_{50} - H_{52} \cdots Cl_{63}$	$sp^{2.31}$	$sp^{2.33}$	-0.11
	$C_{51} - H_{53} \cdots N_{46}$	$sp^{2.31}$	$sp^{2.45}$	-1.2
	$C_{35} - H_{40} \cdots O_{16}$	$sp^{2.26}$	$sp^{2.23}$	0.29
	$N_{11} - H_{12} \cdots O_{47}$	$sp^{2.20}$	$sp^{2.02}$	1.9
	$N_{42} - H_{43} \cdots O_{16}$	$sp^{2.20}$	$sp^{2.01}$	1.96
D5	$C_2 - H_7 \cdots O_{47}$	$sp^{2.28}$	$sp^{2.23}$	0.42
	$C_{18} - H_{20} \cdots N_{13}$	$sp^{2.45}$	$sp^{2.43}$	0.16
		$sp^{2.43}$	$sp^{2.44}$	0.11
	$C_{17} - H_{19} \cdots {}_{27}H - C_{23}$	$sp^{2.38}$	$sp^{2.39}$	-0.03

$$\Delta\%S(X) = \%S^{dimer}(X) - \%S^{monomer}(X)$$

The $C - H \cdots N$ interactions are observed in D1, D3, D4 and D5 dimers. The hybridization states sp^m of the carbon atoms involved in the $C - H \cdots N$ interactions reveal the presence of rehybridization with the complexation. We also note the presence of hyperconjugative interactions that allows an increase of the electron density in the antibonding orbital. This electron density $\Delta\sigma^* = 0.000205$ e has the effect of lengthening the CH bond of $\Delta d(C - H) = 0.00026$ Å in the interaction $C_{18} - H_{20} \cdots N_{13}$ combined with a red shift $\Delta\nu_{C-H} = -2.7588$ cm⁻¹ of the vibrational stretching frequency of the $C - H$ bond. However, in the D1, D4 and D5 dimers, the rehybridization which has the effect of lengthening the bonds during the complexation outweighs the hyperconjugative interactions. Therefore, we have an elongation of the C-H bond with a blue shift of the vibrational stretching frequencies [24, 55]. Concerning the $C - H \cdots H - C$ interactions, the hybridization states of the carbon atoms vary little with the complexation. In addition, NBO analysis did not detect the presence of interaction between orbitals in different bonds involved in $C - H \cdots H - C$ interactions. Consequently, a blue shift of the vibrational stretching frequencies is observed.

3.2.3. AIM Analysis

In the D1-D5 dimers, the $N - H \cdots O$ interactions have an electron density $\rho(r)$ between 0.024561-0.026517 ea_0^{-3} and a positive value of the Laplacian $\nabla^2\rho(r)$ (varying between 0.082075-0.09015 ea_0^{-5}) and the total kinetic energy density $H(r)$ (varying between 0.000672-0.001158 ua). The values of $\rho(r)$, $\nabla^2\rho(r)$ and $H(r)$ indicate a decrease of electronic density in the interatomic bond path characteristic of closed-shell interactions such as hydrogen bonds [52, 54]. Also for all $N - H \cdots O$ interactions, the ratio $-\frac{G(r)}{V(r)} > 1$ highlights the electrostatic nature of these hydrogen bonds [39-41]. Ellipticity is a measure of the stability of the chemical bond. A chemical bond with a large ellipticity is potentially unstable. The $N - H \cdots O$ interactions observed in dimers are stable because they have a low ellipticity $\varepsilon(r)$ in the range of 0.02064-0.031263. The Espinoza method was used to calculate the $N - H \cdots O$ interaction energies and the results obtained are shown in Table 8. The energies of the $N - H \cdots O$ interactions vary between 5.81074-6.29393 kcal/mol. The $N - H \cdots O$ interactions are weak because the energies obtained are less than 15 kcal/mol [50].

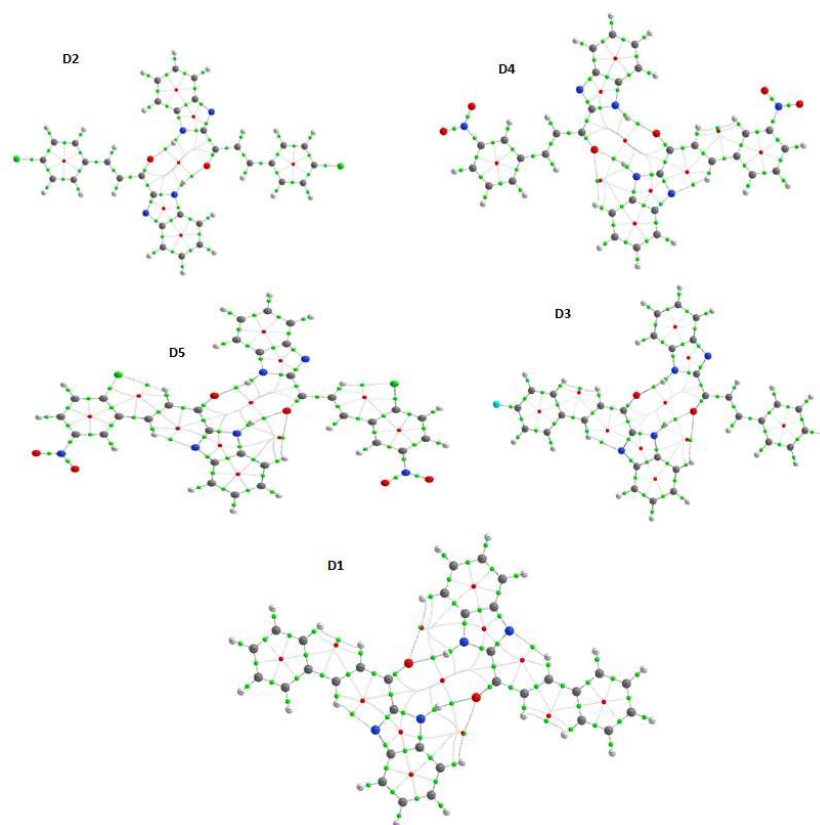


Figure 4. AIM molecular graphs of D1-D5 dimers at B3LYP 6-31+ G (d, p). Small green spheres are bond critical points (Bcps).

At the $C-H \cdots N$ interaction Bcps, the electronic density $\rho(r)$ is between 0.017788 - $0.01991 \text{ ea}_0^{-3}$, the Laplacian of the electronic density $\nabla^2\rho(r)$ is positive and varies between 0.057507 - $0.062472 \text{ ea}_0^{-5}$. In addition, the total energy density $H(r)$ between 0.001253 - 0.001265 au is positive. These features are typical of closed-shell interactions that indicate the presence of hydrogen bonding [52, 54]. In addition, all the interactions $C-H \cdots N$ have the ratio $-\frac{G(r)}{V(r)}$ greater than 1, thus highlighting the electrostatic nature of these bonds [41]. The $C-H \cdots N$ interactions are stable with an ellipticity $\varepsilon(r)$ between 0.003872 - 0.016963 . The interaction energies of $C-H \cdots N$ interactions range from 3.68662 - 4.22942 kcal/mol and are significantly lower than those of the $N-H \cdots O$ interactions.

The electron density obtained at the Bcps of the $C-H \cdots O$ interactions is lower than those of the $N-H \cdots O$ and $C-H \cdots N$ interactions. This electron density $\rho(r)$ varies between 0.00228 - $0.00231904 \text{ ea}_0^{-3}$, its Laplacian $\nabla^2\rho(r)$ is positive and varies between 0.009179 - $0.009449 \text{ ea}_0^{-5}$. Thus, a positive value of $H(r)$ is observed at the different Bcps of the $C-H \cdots O$ interactions. The values of $\rho(r)$, $\nabla^2\rho(r)$ and $H(r)$ show that $C-H \cdots O$ interactions are weak hydrogen

bonds [52, 54]. Moreover, the ratio $-\frac{G(r)}{V(r)} > 1$ for the $C-H \cdots O$ interactions highlights its electrostatic nature [40, 41]. The high value of the ellipticity $\varepsilon(r)$ observed (1.512957 - 11.458266) and the low value of the energy of the hydrogen bond E_{HB} (0.35141 - 0.36396 kcal/mol) highlight their instabilities and weaknesses with respect to the $N-H \cdots O$ and $C-H \cdots O$ hydrogen bonds.

At the Bcps of the $C-H \cdots H-C$ interactions, the electronic density $\rho(r)$ is in the range of 0.008995 - $0.009047 \text{ ea}_0^{-3}$, the corresponding Laplacian is between 0.038854 - $0.039858 \text{ ea}_0^{-5}$ with a positive energy density $H(r)$ (0.002478 - 0.002492 au). Also the interaction energy is about 1.49 kcal/mol , thus $C-H \cdots H-C$ interactions are weak hydrogen bonds [50, 54]. From more $-\frac{G(r)}{V(r)} > 1$ at the Bcps of the different $C-H \cdots H-C$ interaction, thus the $C-H \cdots H-C$ interactions are purely electrostatic [40].

The ellipticity of the $C-H \cdots H-C$ interactions is between 1.145216 - 2.031056 , so we can say that the $C-H \cdots H-C$ interactions are less stable than the $C-H \cdots N$ and $N-H \cdots O$ interactions but more stable than the $C-H \cdots O$ interaction.

Table 8. Topological analysis of the bond critical points (Bcps) of the $X-H \cdots Y$ interactions of D1-D5 dimers.

	$X-H \cdots Y$		$\rho(r)_{X-H \cdots Y}$	$\nabla^2\rho(r)_{X-H \cdots Y}$	$V(r)$	$G(r)$	$H(r)$	$L(r)$	$\varepsilon(r)$	$-\frac{G(r)}{V(r)}$	E_{HB}
D1	$C_{17}-H_{19} \cdots {}_{27}H-C_{23}$	intra	0.009047	0.038856	-0.00476	0.007235	0.002479	-0.00971	1.1452	1.520	1.49347
	$C_{18}-H_{20} \cdots N_{13}$	intra	0.017788	0.057118	-0.01175	0.013015	0.001265	-0.01428	0.0170	1.108	3.68662
	$C_{49}-H_{51} \cdots N_{44}$	intra	0.017798	0.057149	-0.01176	0.013023	0.001265	-0.01429	0.0170	1.107	3.68976
	$C-H_{58} \cdots H-C$	intra	0.009047	0.038854	-0.00476	0.007234	0.002478	-0.00971	1.1457	1.520	1.49347

$X-H\cdots Y$		$\rho(r)_{X-H\cdots Y}$	$\nabla^2\rho(r)_{X-H\cdots Y}$	$V(r)$	$G(r)$	$H(r)$	$L(r)$	$\epsilon(r)$	$-\frac{G(r)}{V(r)}$	E_{HB}
$C_2-H_7\cdots O_{47}$	inter	0.00228	0.009426	-0.00116	0.001755	0.0006	-0.00236	11.4583	1.513	0.36396
$N_{11}-H_{12}\cdots O_{47}$	inter	0.025987	0.087155	-0.01972	0.020752	0.001036	-0.02179	0.0300	1.052	6.18725
$N_{42}-H_{43}\cdots O_{16}$	inter	0.025993	0.087181	-0.01972	0.020758	0.001037	-0.0218	0.0300	1.053	6.18725
$C_{33}-H_{38}\cdots O_{16}$	inter	0.002281	0.009427	-0.00116	0.001756	0.000601	-0.00236	10.7918	1.514	0.36396
$N_{11}-H_{12}\cdots O_{47}$	inter	0.026159	0.085744	-0.01966	0.020549	0.000887	-0.02144	0.0198	1.045	6.16842
$N_{42}-H_{43}\cdots O_{16}$	inter	0.026159	0.085745	-0.01966	0.020549	0.000887	-0.02144	0.0198	1.045	6.16842
$N_{44}-H_{45}\cdots O_{16}$	inter	0.025527	0.086416	-0.01934	0.020472	0.001132	-0.0216	0.0309	1.059	6.06802
$N_{11}-H_{12}\cdots O_{49}$	inter	0.024561	0.079453	-0.01852	0.019191	0.000672	-0.01986	0.0206	1.036	5.81074
$C_2-H_7\cdots O_{49}$	inter	0.00224	0.009179	-0.00112	0.001706	0.00059	-0.0023	2.0833	1.523	0.35141
$C_{35}-H_{40}\cdots O_{16}$	Inter	0.002252	0.009236	-0.00113	0.001718	0.00059	-0.00231	2.3831	1.520	0.35454
$N_{44}-H_{45}\cdots O_{16}$	Inter	0.025138	0.082075	-0.01898	0.019751	0.000767	-0.02052	0.0212	1.041	5.95507
$N_{11}-H_{12}\cdots O_{49}$	Inter	0.026517	0.090159	-0.02006	0.021298	0.001242	-0.02254	0.0313	1.062	6.29393
$C_{17}-H_{19}\cdots Cl_{30}$	intra	0.014125	0.054501	-0.00891	0.011268	0.002357	-0.01363	0.0743	1.265	2.79556
$C_{51}-H_{53}\cdots N_{46}$	intra	0.01991	0.062472	-0.01348	0.014546	0.001071	-0.01562	0.0039	1.079	4.22942
$C_{50}-H_{52}\cdots Cl_{63}$	intra	0.014913	0.055646	-0.00961	0.011758	0.002153	-0.01391	0.0388	1.224	3.01519
$C_{18}-H_{20}\cdots N_{13}$	intra	0.017945	0.057507	-0.01187	0.013124	0.001253	-0.01438	0.0167	1.106	3.72427
$C_{17}-H_{19}\cdots {}_{27}H-C_{23}$	intra	0.008995	0.038835	-0.00473	0.007218	0.002492	-0.00971	1.2077	1.526	1.48406
$N_{42}-H_{43}\cdots O_{16}$	inter	0.026125	0.08835	-0.01977	0.02093	0.001158	-0.02209	0.0304	1.059	6.20294
$N_{11}-H_{12}\cdots O_{47}$	inter	0.024803	0.080458	-0.01872	0.019418	0.000697	-0.02012	0.0212	1.037	5.87349
$C_2-H_7\cdots O_{47}$	inter	0.002319	0.009449	-0.00115	0.001758	0.000604	-0.00236	1.5130	1.529	0.36082

4. NCI-RDG Analysis

In M1-M5 monomers and D1-D5 complexes, NCI analysis revealed the presence of hydrogen bonds, VDW interactions and repulsive interactions. The presence of the $N-H\cdots O$, $C-H\cdots N$ and $C-H\cdots O$ hydrogen bonds in the complexes is evidenced by the presence of peaks that appear at $-0,040 < \text{sign}(\lambda_2) \times \rho(r) < -0,015$ (figure 5 and 6). In figure 5 and 6, the hydrogen bonds $N-H\cdots O$, $C-H\cdots N$ and $C-H\cdots O$ are characterized by the presence of blue

isosurface between the atoms H and O, H and N. The presence of green isosurface reveals the presence of VDW interaction (figure 6). The peaks corresponding to these interactions are observed at $\text{sign}(\lambda_2) \times \rho(r) = 0$. Repulsive interactions are observed when we have the presence of a high electron density. In the $RDG = f(\text{sign}(\lambda_2) \times \rho(r))$ graphs, the peaks of repulsive interactions (steric hindrance) appear at $0,020 < \text{sign}(\lambda_2) \times \rho(r) < 0,100$. The presence of these repulsive interactions is represented by red isosurfaces in the various benzene and imidazole ring.

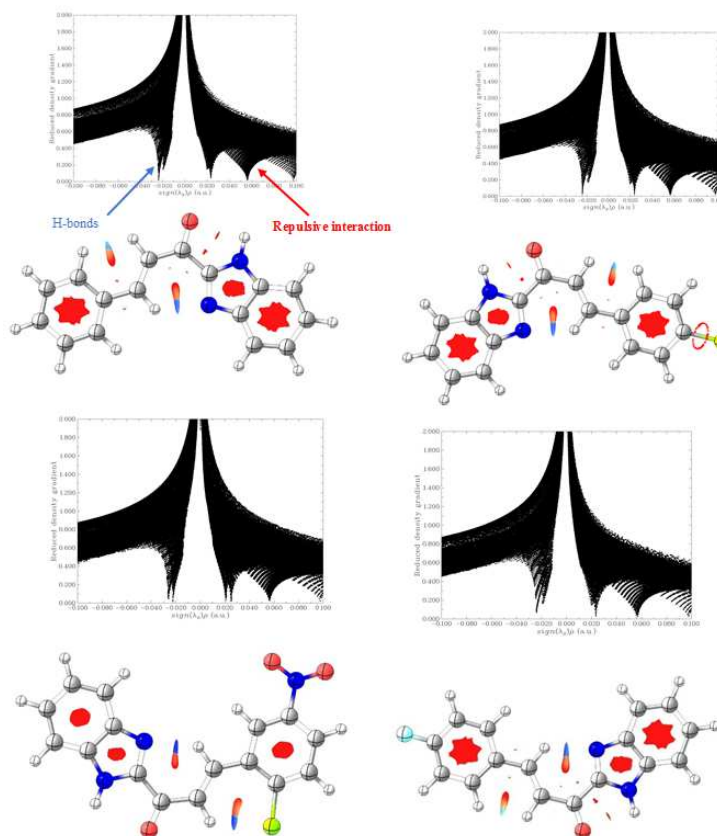


Figure 5. NCI isosurface and plot of the reduced density gradient (RDG) versus $\text{sign}(\lambda_2) \times \rho(r)$ of BZims monomer.

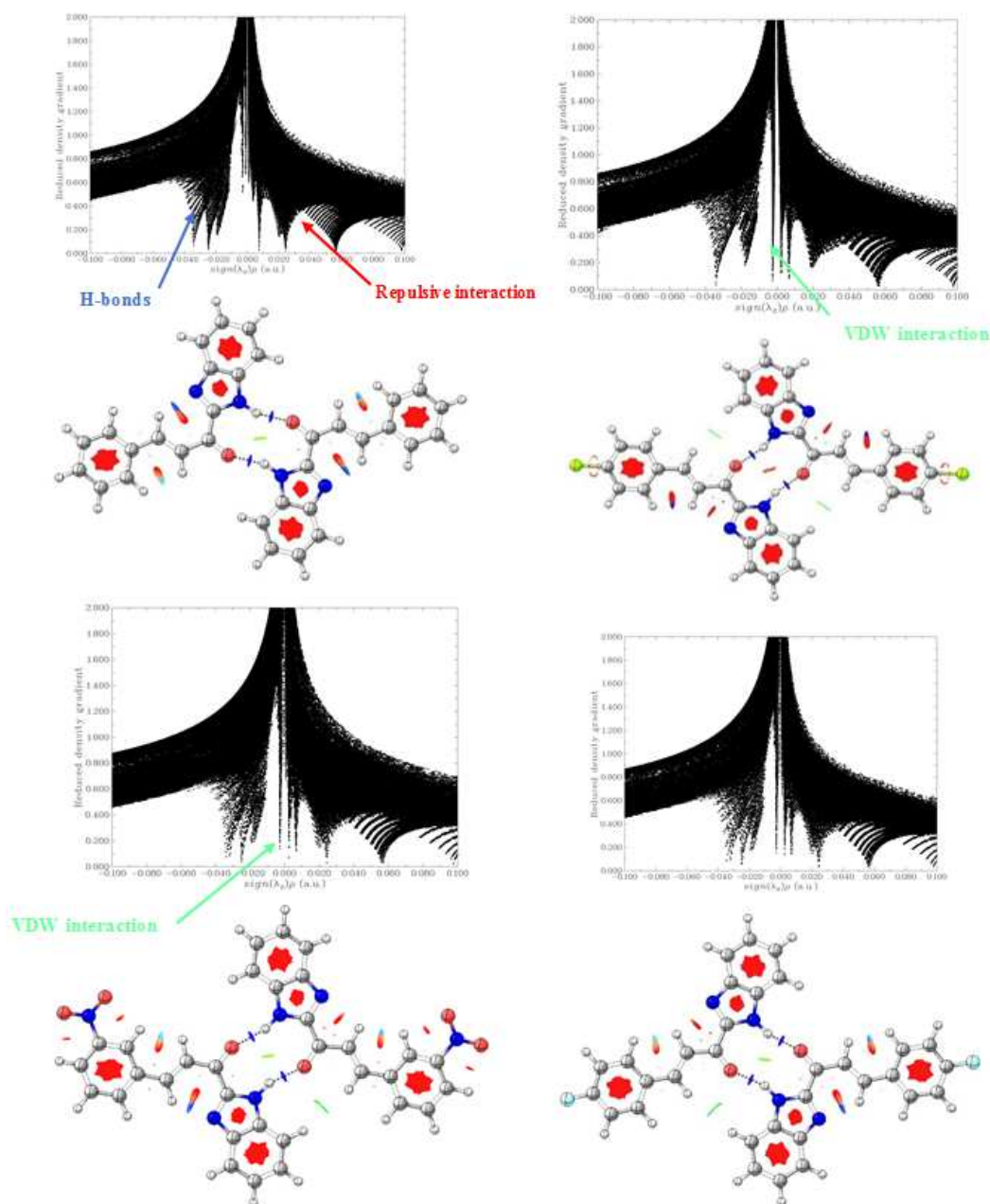


Figure 6. NCI isosurface and plot of the reduced density gradient (RDG) versus $\text{sign}(\lambda_2) \times \rho(r)$ of BZims complexes.

5. Conclusion

We carried out a detailed theoretical analysis of the monomers and dimers of BZims. Geometry optimization and harmonic frequency analysis were performed. The results of the frequency calculations show that all the monomers and dimers of the BZims are at least global structures. Our calculations clearly show that the BZims monomers and dimers have intramolecular and intermolecular hydrogen bonds. From the point of view of the interaction energy, the D2 dimer is more stable than the other forms. The stability sequence of dimers BZims is $D_2 > D_5 > D_4 > D_3 > D_1$. The presence of weakly electron acceptor group (-Cl) on benzene ring favor the total interaction energy of dimerization. Evidence for the presence of intramolecular and

intermolecular hydrogen bonding has been demonstrated by topological, geometrical, NCI and NBO analyzes. Frequency analysis results for BZims monomers and dimers indicate that the intramolecular hydrogen bond shows an increase in stretching frequencies and a blue shift in the IR spectrum, while the intermolecular hydrogen bond shows a decrease in stretching frequencies. and a red shift of the IR spectrum.

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