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Castet, Frédéric; Gillet, Adrien; Bureš, Filip; Plaquet, Aurélie; Rodriguez, Vincent; Champagne, Benoît

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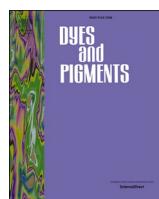
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## Second-order nonlinear optical properties of Λ-shaped pyrazine derivatives

Frédéric Castet<sup>a,\*</sup>, Adrien Gillet<sup>a</sup>, Filip Bureš<sup>b</sup>, Aurélie Plaquet<sup>c</sup>, Vincent Rodriguez<sup>a, \*\*</sup>, Benoît Champagne<sup>c, \*\*\*</sup>

<sup>a</sup> Institut des Sciences Moléculaires (ISM, UMR CNRS 5255), Université de Bordeaux, 351 Cours de La Libération, 33405, Talence, France

<sup>b</sup> Institute of Organic Chemistry and Technology, Faculty of Chemical Technology, University of Pardubice, Studentská 573, Pardubice, 532 10, Czech Republic

<sup>c</sup> Unité de Chimie Physique Théorique et Structurale, Chemistry Department, Namur Institute of Structured Matter, University of Namur, Rue de Bruxelles, 61, B-5000, Namur, Belgium

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### ABSTRACT

The linear and nonlinear optical (NLO) properties of a series of Λ-shaped derivatives containing a 4,5-dicyanopyrazine acceptor unit, *N,N*-dimethylamino donor groups and systematically enlarged π-conjugated linkers are investigated by means of UV/Visible and Hyper-Rayleigh scattering spectroscopies. Density functional theory calculations are also carried out to rationalize the magnitude and symmetry of the NLO responses. The results demonstrate that these compounds possess two low-lying excited electronic states close to each other in energy, which are accessible through one-photon optical transitions respectively polarized perpendicular and parallel to the two-fold molecular axis. These two states contribute additively to the first hyperpolarizability, which exhibits a dominant dipolar character. We also show that the NLO responses significantly deviate from Kleinman's index permutation symmetry.

### 1. Introduction

The design of organic chromophores having the ability to deliver large second order nonlinear optical (NLO) responses has been a continuous source of research for the past thirty years, in connection with a broad field of applications ranging from smart materials (including optical telecommunications, data storage, information and signal processing) [1,2] to biology (including *in vivo* bioimaging and therapy) [3–6]. Most of systems reported to date are (one-dimensional) dipolar compounds, which comprise a π-conjugated linker connected at both ends to electron-donating (D) and electron-withdrawing (A) chemical substituents. Varying the nature of the D/A groups and the length of the π-conjugated bridge by means of chemical design remains the most common strategy for tuning the magnitude of the first hyperpolarizability ( $\beta$ ) [7–15].

However, if 1D compounds present the advantage to enable electric-field poling in chromophore-polymer composite materials [1], they are limited by the fact that any increase of the first hyperpolarizability is most often accompanied by a redshift of the absorbance maxima, which translates into a loss of transparency. An alternative to circumvent the

nonlinearity/transparency tradeoff is to exploit the dimensionality of the NLO chromophores. In that respect, two-dimensional octupolar chromophores that exhibit three-fold symmetry, with crystal violet as prototypical compound, have proven to be of major interest for electro-optics, since their potentially high  $\beta$  values are associated with a zero dipole moment and with increased transparency [16–18]. The enhanced nonlinear response of these compounds originates from the existence of two degenerate low-lying excited states accessible by one-photon absorption, which both contribute to the first hyperpolarizability.

NLO chromophores exhibiting  $C_{2v}$  symmetry with a rigid Λ-shaped geometry also enable the possibility of modulating the magnitudes and dimensionality of the second order NLO responses, and offer an interesting compromise combining the advantages of 1D and 2D systems [19–23]. As exemplified by 3,5-dinitro-aniline [24] or (dicyanomethylene)pyran derivatives [25], NLO compounds with  $C_{2v}$  symmetry possess two low-lying excited electronic states close to each other in energy, the first transition ( $S_1$ ) being polarized perpendicular to the symmetry ( $z$ ) axis (*i.e.* along  $x$ ), whereas the second ( $S_2$ ) is polarized parallel to  $z$ . As a consequence, the  $\beta$  tensor contains two main

\* Corresponding author.

\*\* Corresponding author.

\*\*\* Corresponding author.

E-mail address: [frederic.castet@u-bordeaux.fr](mailto:frederic.castet@u-bordeaux.fr) (F. Castet).

components,  $\beta_{zzz}$ , which in a Sum-Over-States (SOS) picture depends only on  $S_2$ , and  $\beta_{zxx}$ , which depends both on  $S_1$  and  $S_2$ . Important aspects related to the large  $\beta_{zxx}$  components lie in the fact that harmonic lights polarized perpendicular to the transition dipole are not reabsorbed, which reduces losses in the second-harmonic intensity, and that fundamental and harmonic lights are propagating with different polarization, which offers the possibility of phase-matching. Theoretical investigations further demonstrated that specific ratios between the off-diagonal ( $\beta_{zxx}$ ) and diagonal ( $\beta_{zzz}$ ) tensor components can be achieved by finely tuning the D/A strengths and the angle between the two molecular branches [26,27]. In addition, it has been evidenced that the presence of the low-lying excited state with transition dipole perpendicular to the symmetry axis was responsible for significant deviation from Kleinman's index permutation symmetry [24,28].

Herein, we report a detailed investigation of the linear and nonlinear optical properties of a series of  $\Lambda$ -shaped pyrazine derivatives [29], which involve systematically enlarged  $\pi$ -conjugated segments with 1, 4-phenylene rings as well as acetylene units (Fig. 1). Our study combines Hyper-Rayleigh Scattering (HRS) measurements to Density Functional Theory (DFT) calculations, and focuses on the impact of the nature and length of the  $\pi$ -conjugated system on the magnitude and symmetry of the second-order NLO response, in relation with the structural symmetry of the compounds. In particular, we introduce a new experimental approach to evaluate the deviations from Kleinman's symmetry, which are interpreted in the light of theoretical calculations.

## 2. Experimental and theoretical details

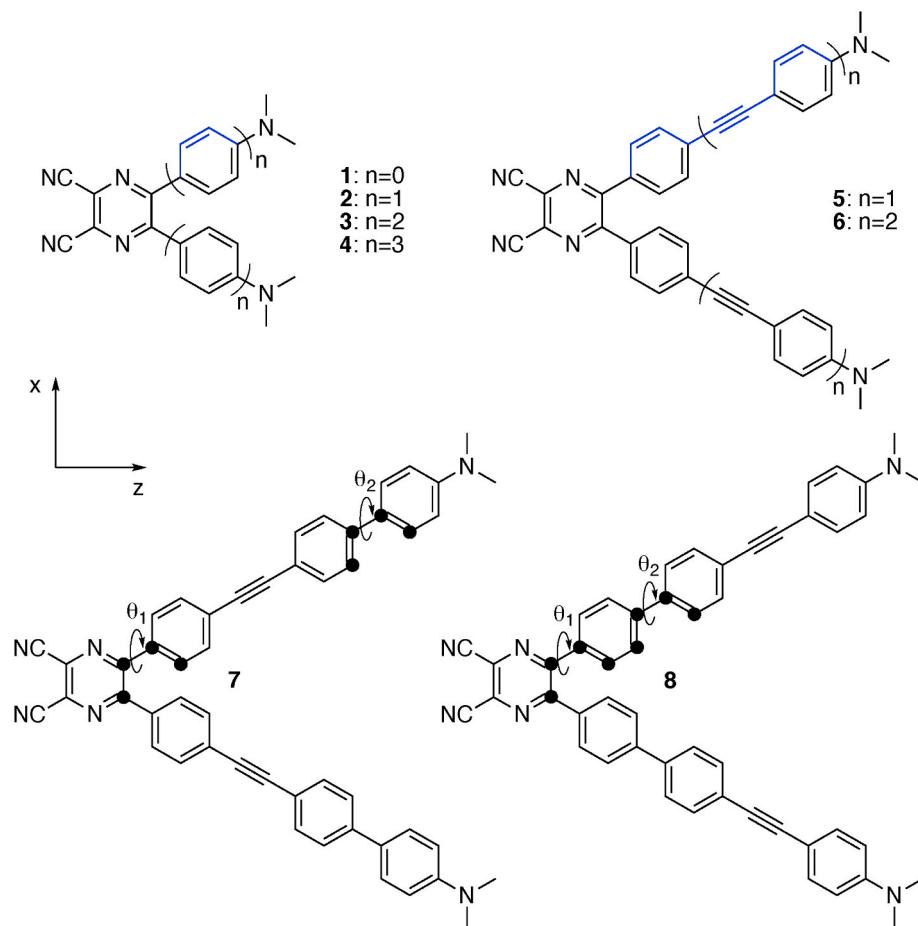
### 2.1. Hyper-Rayleigh Scattering measurements

UV/vis absorption spectra were recorded on a PerkinElmer Lambda 650 spectrophotometer in dichloromethane (DCM). HRS experiments were performed on diluted solutions with concentrations ranging from  $10^{-4}$  to  $10^{-5}$  mol/L in DCM, checking the linear dependence of absorbance. The first hyperpolarizability was determined from the intensity of the incoherent scattered light at the second harmonic frequency of an IR diode pumped picosecond laser Nd:YVO<sub>4</sub> ( $\lambda = 1064$  nm, pulse duration 65 ps, repetition rate 2 kHz). The scattered light was collected at 90° and focused into a Jobin-Yvon Horiba spectrograph, according to an experimental setup reported previously [30].

The HRS signal of incoherent and uncorrelated molecular scatterers in dilute solution is related to the molecular first hyperpolarizability as follows:

$$I^{2\omega} \propto C_{chr} \langle \beta_{HRS}^2 \rangle (I^\omega)^2 \times 10^{-\varepsilon_{2\omega}\ell} C_{chr} \quad (1)$$

where  $I^\omega$  is the intensity of the incident light beam of frequency  $\omega$  and  $C_{chr}$  is the concentration of the NLO chromophores. The last term (where  $\varepsilon_{2\omega}$  is the molar extinction coefficient at frequency  $2\omega$  and  $\ell$  the optical path length) accounts for the absorption losses at the second harmonic wavelength.  $\langle \beta_{HRS}^2 \rangle$  is the orientational average of the first hyperpolarizability tensor over all possible molecular orientations. Since for a non-polarized/natural incident light both horizontal ( $H \parallel X$ ) and vertical ( $V \parallel Z$ ) polarizations have equal weights,  $\beta_{HRS}$  can be expressed as the



**Fig. 1.** Pyrazine derivatives investigated in this work. The Cartesian frame used in NLO calculations is shown on the left. The dihedral angles around single bonds ( $\theta_{1-3}$ ) are defined between dotted carbon atoms starting from the pyrazine core, as exemplified on compounds **7** and **8**. The conjugated segment considered to compute the BLA is shown in blue in compounds **1–6**.

sum of two contributions:

$$\beta_{HRS} = \beta_{HRS}(-2\omega; \omega, \omega) = \sqrt{\langle \beta_{HRS}^2 \rangle} = \sqrt{\langle \beta_{ZZZ}^2 \rangle + \langle \beta_{ZXX}^2 \rangle} \quad (2)$$

The  $\langle \beta_{ZZZ}^2 \rangle$  and  $\langle \beta_{ZXX}^2 \rangle$  invariants can be written in terms of molecular  $\beta$  tensor components as follows:

$$\begin{aligned} \langle \beta_{ZZZ}^2 \rangle &= \frac{1}{7} \sum_{\zeta} \beta_{\zeta\zeta\zeta}^2 + \frac{4}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\eta}^2 + \frac{2}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\zeta} \beta_{\zeta\eta\eta} \\ &+ \frac{4}{35} \sum_{\zeta \neq \eta} \beta_{\eta\zeta\zeta} \beta_{\zeta\zeta\eta} + \frac{4}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\zeta} \beta_{\eta\eta\zeta} + \frac{1}{35} \sum_{\zeta \neq \eta} \beta_{\eta\zeta\zeta}^2 \\ &+ \frac{4}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\zeta\eta} \beta_{\eta\xi\xi} + \frac{1}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\eta\zeta\zeta} \beta_{\eta\xi\xi} \\ &+ \frac{4}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\zeta\eta} \beta_{\xi\xi\eta} \\ &+ \frac{2}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\eta\xi}^2 + \frac{4}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\eta\xi} \beta_{\eta\xi\xi} \end{aligned} \quad (3)$$

$$\begin{aligned} \langle \beta_{ZXX}^2 \rangle &= \frac{1}{35} \sum_{\zeta} \beta_{\zeta\zeta\zeta}^2 + \frac{4}{105} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\zeta} \beta_{\zeta\eta\eta} - \frac{2}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\zeta} \beta_{\eta\eta\zeta} \\ &+ \frac{8}{105} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\eta}^2 + \frac{3}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\eta}^2 - \frac{2}{35} \sum_{\zeta \neq \eta} \beta_{\zeta\zeta\eta} \beta_{\eta\zeta\zeta} \\ &+ \frac{1}{35} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\eta\eta} \beta_{\zeta\xi\xi} - \frac{2}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\zeta\zeta} \beta_{\eta\eta\xi} \\ &- \frac{2}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\zeta\eta} \beta_{\eta\xi\xi} \\ &+ \frac{2}{35} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\eta\xi}^2 - \frac{2}{105} \sum_{\zeta \neq \eta \neq \xi} \beta_{\zeta\eta\xi} \beta_{\eta\xi\xi} \end{aligned} \quad (4)$$

Experimentally,  $\langle \beta_{ZZZ}^2 \rangle$  and  $\langle \beta_{ZXX}^2 \rangle$  can be obtained independently from measurements performed in the VV (vertically polarized incident and scattered lights) and HV (horizontally polarized incident light and vertically polarized scattered light) configurations, respectively. Their ratio, referred to as the depolarization ratio (DR), depends on the symmetry of the NLO chromophores and ranges between 1.5 for octupolar compounds and 9 for pure dipolar molecules [31]:

$$DR = \frac{\langle \beta_{ZZZ}^2 \rangle}{\langle \beta_{ZXX}^2 \rangle} \quad (5)$$

However, more accurate determination of  $\beta_{HRS}$  and DR are obtained by considering the full range of polarization angles  $\Psi$  of the incident light. Considering a phase retardation of  $\pi/2$  in accordance with the experimental setup (which combines a rotating half-wave plate and a fixed quarter-wave plate, see Ref. 30 for details), the intensity of the harmonic light collected perpendicular to the incident beam and vertically polarized implies a quartic dependence in  $\cos\Psi$ , following the expression established by Bersohn [32]:

$$\begin{aligned} I_{\Psi\Psi}^{2\omega} \propto & \langle \beta_{ZXX}^2 \rangle \cos^4\Psi + \langle \beta_{ZZZ}^2 \rangle \sin^4\Psi \\ & + \langle (\beta_{ZXZ} + \beta_{ZZX})^2 - 2\beta_{ZZZ}\beta_{ZXX} \rangle \cos^2\Psi \sin^2\Psi \end{aligned} \quad (6)$$

In number of previously reported works [12,24,25,28,30], NLO measurements of rod-like and  $\Lambda$ -shaped compounds were analyzed assuming a planar  $C_{2v}$  molecular symmetry (where the molecules lay in a mean  $xz$  plane with  $z$  the twofold symmetry axis), as well as time reversibility of the second-order optical process. The latter approximation, known as Kleinman's symmetry, is strictly fulfilled only in the static limit and assumes the invariance of the  $\beta$  tensor components with respect to permutation of their Cartesian indices, such that  $\beta_{ijk} = \beta_{kji}$ . Assuming both planar  $C_{2v}$  and Kleinman symmetries implies that  $\beta_{zyy} = \beta_{yyz} = 0$ ,  $\beta_{zzz} \neq 0$  and  $\beta_{zxx} = \beta_{xzx} \neq 0$  in equations (3) and (4). The intensity of the harmonic light given in equation (6) can then be expressed as a function of the two non-zero independent  $\beta$  components:

$$\begin{aligned} I_{\Psi\Psi}^{2\omega}(C_{2v}) \propto & \frac{1}{105} \beta_{zzz}^2 ((15 + 18R + 27R^2) - (24 + 68R + 4R^2) \cos^2\Psi \\ & + (12 + 48R - 12R^2) \cos^4\Psi) \end{aligned} \quad (7)$$

where  $R = \beta_{zxx}/\beta_{zzz}$ . Accordingly, the expressions of  $\beta_{HRS}$  and DR can be simplified into:

$$\beta_{HRS}(C_{2v}) = |\beta_{zzz}| \left( \frac{6}{35} + \frac{16}{105} R + \frac{38}{105} R^2 \right)^{1/2} \quad (8)$$

$$DR(C_{2v}) = \frac{\frac{1}{35} + \frac{6}{35} R + \frac{9}{35} R^2}{\frac{1}{35} - \frac{2}{105} R + \frac{11}{105} R^2} \quad (9)$$

In the work reported herein, we rely on spherical harmonics in addition to Cartesian coordinates. This formalism, in which the first hyperpolarizability is decomposed in terms of dipolar ( $\beta_{J=1}$ ) and octupolar ( $\beta_{J=3}$ ) tensorial components, allows us to analyze the HRS data without imposing any structural symmetry constraint on the  $\beta$  tensor. Assuming Kleinman's conditions, the three HRS invariants of equation (6),  $\langle \beta_{ZZZ}^2 \rangle$ ,  $\langle \beta_{ZXX}^2 \rangle$  and  $\langle (\beta_{ZXZ} + \beta_{ZZX})^2 - 2\beta_{ZZZ}\beta_{ZXX} \rangle$ , can be rewritten as:

$$\langle \beta_{ZZZ}^2 \rangle = |\beta_{J=1}|^2 \left( \frac{9}{45} + \frac{6}{105} \rho^2 \right) \quad (10)$$

$$\langle \beta_{ZXX}^2 \rangle = |\beta_{J=1}|^2 \left( \frac{1}{45} + \frac{4}{105} \rho^2 \right) \quad (11)$$

$$\langle (\beta_{ZXZ} + \beta_{ZZX})^2 - 2\beta_{ZZZ}\beta_{ZXX} \rangle = |\beta_{J=1}|^2 \left( -\frac{2}{45} + \frac{22}{105} \rho^2 \right) \quad (12)$$

where the nonlinear anisotropy parameter  $\rho = |\beta_{J=3}|/|\beta_{J=1}|$  denotes the relative magnitude of the octupolar and dipolar contributions to the first hyperpolarizability. The total HRS intensity and the depolarization ratio are expressed as:

$$\beta_{HRS} = |\beta_{J=1}| \sqrt{\frac{2}{3} \left( \frac{1}{3} + \frac{1}{7} \rho^2 \right)} \quad (13)$$

$$DR = \frac{9 \left( 1 + \frac{2}{7} \rho^2 \right)}{1 + \frac{12}{7} \rho^2} \quad (14)$$

and equation (6) becomes:

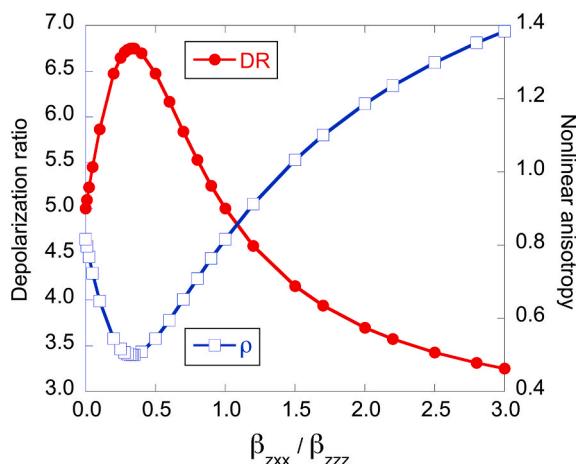
$$I_{\Psi\Psi}^{2\omega} \propto |\beta_{J=1}|^2 \left( \frac{1}{45} + \frac{4}{105} \rho^2 \right) [\cos^4\Psi + \Delta \sin^4\Psi + \Delta' \cos^2\Psi \sin^2\Psi] \quad (15)$$

According to Kleinman symmetry,  $\Delta = DR$  and  $\Delta' = 7 - DR$ . Since the  $\rho$  and DR parameters are linked through equation (14), the values of the two parameters can be determined from the polarization curves by using a fitting procedure implying  $\rho$  as the only adjustable parameter. Assuming in addition that molecular structures have a  $C_{2v}$  symmetry, the  $\rho$  and DR parameters evolve as a function of  $R = \beta_{zxx}/\beta_{zzz}$  as shown in Fig. 2.

Finally, to address the applicability of the time-reversibility assumption, it is further possible to evaluate the deviation from the Kleinman symmetry by introducing an additional parameter in equation (15), in such a way that  $\Delta = DR_{NK}$  and  $\Delta' = 7 - DR_{NK} + \delta_{NK}$ . In this case, the "non-Kleinman" (NK) depolarization ratio,  $DR_{NK}$ , and the deviation from Kleinman symmetry,  $\delta_{NK}$ , are optimized according to a two-parameter fitting procedure.

## 2.2. Quantum chemical calculations

Molecular structures were optimized at the density functional theory (DFT) level using the hybrid meta-GGA M06-2X exchange-correlation functional (XCF) including 54% of Hartree-Fock (HF) exchange, in association with the 6-311G(d) basis set. Real values for all vibrational



**Fig. 2.** Evolution of the depolarization and anisotropy ratios as a function of  $R = \beta_{zxx}/\beta_{zzz}$  when assuming both  $C_2v$  and Kleinman symmetries (equations (9) and (14)).

frequencies confirmed that the geometries correspond to minima on the potential energy surface. Linear and nonlinear optical properties (in the static limit and using an incident wavelength of 1064 nm) were determined using time-dependent (TD) DFT with the same functional and the larger 6-311+G(d) basis set containing additional diffuse functions. The M06-2X XCF was demonstrated to perform well for calculating the second-order NLO properties of a large variety of organic chromophores [33–37]. Solvent effects were taken into account in both geometry optimizations and calculations of the optical responses by using the integral equation formalism of the polarizable continuum model (IEF-PCM) [38], with dichloromethane as solvent. All calculations were performed using the Gaussian09 program package [39].

### 3. Results and discussion

#### 3.1. Structural and ground-state electronic properties

All molecules exhibit a  $C_2$  symmetry. The bond length alternation (BLA) along the conjugated chain connecting the donor and acceptor units (i.e. between the most distant carbon atoms of the first and last phenyl units, see Fig. 1), as well as the torsional angles around single bonds (between the pyrazine and phenyl rings and between consecutive phenyl units when relevant), are listed in Table 1 for all compounds. The energy of the frontier MOs and the ground-state dipole moments are also provided, together with the net charge  $\delta$  of the *N,N*-dimethylamino donor and dicyano-pyrazine acceptor moieties (calculated by summing the Mulliken atomic populations). For a conjugated chain containing  $N$  carbon atoms, the BLA was calculated as:

$$BLA = \frac{1}{N-2} \sum_{i=1}^{N-2} (-1)^{i+1} (d_{i+1,i+2} - d_{i,i+1}) \quad (16)$$

where  $d_{ij}$  is the interatomic distance between carbons  $i$  and  $j$ . A positive value of the BLA indicates a dominant aromatic character of the conjugated linker, while  $BLA < 0$  reveals a dominant quinoid (charge transfer) character. A value equal to zero in 1–4 indicates that the aromatic and quinoid resonance forms equivalently contribute to the ground-state electronic structure according to a 2-state resonance picture.

Increasing the number of phenyl rings in compounds 1–4 concomitantly destabilizes the HOMO and stabilizes the LUMO, inducing a lowering of the energy gap. From compound 1 to 2, the enlargement of the conjugation length strongly increases the dipole moment. Further addition of phenyl units in compounds 3 and 4 slightly lowers the dipole moment, due to a decrease in the efficiency of the charge transfer between the *N,N*-dimethylamino and dicyano-pyrazine units. This lowering of the quinoid character from 2 to 4 is evidenced by the decrease of the difference in the net charges of the D and A units, as well as by the increase of the BLA values from  $-0.026$  to  $+0.024$  Å.

Replacing the central phenyl ring in compound 4 by an acetylenic linker (compound 5) gives rise to a larger donor-to-acceptor charge transfer and strongly enhances the molecular dipole, while introducing a second phenylacetylene unit (from compound 5 to 6) has a weak impact. Finally, comparing the results calculated for compounds 7 and 8 shows that the position of the acetylenic bridge impacts the ground-state intramolecular charge transfer (ICT), a larger dipole being obtained with the triple bond closer to the *N,N*-dimethylamino donor units.

#### 3.2. Linear optical properties

Due to the  $\Lambda$ -symmetry of the molecules, the two lowest-energy transitions,  $S_0 \rightarrow S_1$  and  $S_0 \rightarrow S_2$ , are optically allowed. The transition energies, wavelengths, oscillator strengths, as well as the main electronic excitations contributing to these two optical transitions, are gathered in Table 2. The experimental and calculated UV/Vis spectra are provided in Figs. S1 and S6. In all compounds, the  $S_0 \rightarrow S_2$  transition is 0.1–0.2 eV higher in energy than  $S_0 \rightarrow S_1$ , and associated to a larger oscillator strength. As already observed for  $\Lambda$ -shape compounds such as 3,5-dinitroaniline [24] or 1,3-diamino-4,6-dinitrobenzene [19] derivatives, the transition dipole moment associated to the  $S_0 \rightarrow S_1$  excitation ( $\vec{\mu}_{01}$ ) is parallel to the transverse  $x$ -axis, while the one corresponding to  $S_0 \rightarrow S_2$  ( $\vec{\mu}_{02}$ ) is aligned along the  $C_2$  symmetry axis (see Figs. S7 and S8). As discussed below, these two perpendicular charge-transfer excitations are responsible for the 2D NLO character of the chromophores.

Although the computed transition wavelengths globally underestimate the experimental absorption maximum, DFT calculations using the vertical approximation reproduce well the relative trends. Within the 1–4 series, the transition wavelength increases from 1 to 2, and then decreases when introducing additional 1,4-phenylene units in

**Table 1**

Bond length alternation (BLA, Å), torsional angles ( $\theta_{1-3}$ , deg.), HOMO and LUMO energies (eV), HOMO-LUMO gap ( $E_g$ , eV), ground-state dipole moment ( $\mu_g$ , D), and sum of Mulliken charges within the dicyano-pyrazine and *N,N*-dimethylamino units ( $\delta_A$  and  $\delta_D$ , |e|), calculated at the IEF-PCM/M06-2X/6-311+G(d) level in dichloromethane.

Compound	BLA	$\theta_1, \theta_2, \theta_3$	HOMO	LUMO	$E_g$	$\mu_g$	$\delta_A$	$\delta_D$
1	/	/	-7.39	-1.40	5.98	13.81	-1.46	1.46
2	-0.026	36	-6.58	-1.89	4.70	17.84	-0.92	1.05
3	0.014	39, 34	-6.57	-2.14	4.43	17.23	-1.00	0.92
4	0.024	40, 38, 35	-6.56	-2.20	4.36	16.68	-0.85	0.82
5	0.049	40	-6.50	-2.27	4.23	19.26	-0.95	1.03
6	0.060	41	-6.46	-2.37	4.10	19.50	-0.87	0.97
7	0.045	41, 35	-6.55	-2.34	4.21	17.93	-0.85	0.81
8	0.045	40, 36	-6.47	-2.24	4.23	19.22	-0.76	0.98

**Table 2**

Experimental maximum absorption wavelengths ( $\lambda_{exp}$ , nm), and wavelengths ( $\lambda_{ge}$ , nm), energies ( $\Delta E_{ge}$ , eV), squared transition dipoles ( $\mu_{ge}^2$ , a.u.<sup>2</sup>) oscillator strengths ( $f_{ge}$ ), photoinduced charge displacements ( $\Delta q$ , |e|), charge transfer distances ( $\Delta r$ , Å), dipole moment variations ( $\Delta \mu_{ge}$ , D), and weights of the main electronic excitations ( $\geq 10\%$ ), calculated at the IEF-PCM/M06-2X/6-311+G(d) level in dichloromethane for the two lowest-energy optical transitions.

Compound	$\lambda_{exp}$ [29]	$\lambda_{ge}$	$\Delta E_{ge}$	$\mu_{ge}^2$	$f_{ge}$	$\Delta q$	$\Delta r$	$\Delta \mu_{ge}$	Electronic excitations
1:S <sub>0</sub> → S <sub>1</sub>	355	323	3.84	3.92	0.37	0.55	2.34	6.21	98% H→L
1:S <sub>0</sub> → S <sub>2</sub>		317	3.91	4.41	0.42	0.47	0.71	1.60	98% H→L+1
2:S <sub>0</sub> → S <sub>1</sub>	471	421	2.94	7.31	0.53	0.77	3.29	12.14	92% H→L
2:S <sub>0</sub> → S <sub>2</sub>		398	3.11	7.57	0.58	0.76	2.95	10.84	94% H→L+1
3:S <sub>0</sub> → S <sub>1</sub>	425	399	3.11	9.07	0.69	0.87	4.64	19.49	77% H→L + 10% H-2→L
3:S <sub>0</sub> → S <sub>2</sub>		376	3.30	15.37	1.24	0.86	4.72	19.35	53% H→L+1 + 29% H-1→L + 10% H-2→L+1
4:S <sub>0</sub> → S <sub>1</sub>	402	371	3.35	10.29	0.84	0.85	5.68	23.19	57% H→L + 26% H-2→L
4:S <sub>0</sub> → S <sub>2</sub>		355	3.49	16.75	1.43	0.89	6.10	26.09	43% H-1→L + 26% H→L+1 + 15% H-2→L+1
5:S <sub>0</sub> → S <sub>1</sub>	433	416	2.98	13.93	1.02	0.84	5.13	20.78	72% H→L + 10% H-2→L
5:S <sub>0</sub> → S <sub>2</sub>		392	3.17	23.90	1.85	0.84	5.35	21.60	45% H-1→L + 34% H→L+1
6:S <sub>0</sub> → S <sub>1</sub>	402	400	3.10	22.93	1.74	0.73	6.14	21.46	43% H→L + 21% H-2→L
6:S <sub>0</sub> → S <sub>2</sub>		387	3.20	39.56	3.10	0.74	6.50	22.92	36% H-1→L + 17% H→L+1 + 14% H→L+3 + 13% H-1→L+1
7:S <sub>0</sub> → S <sub>1</sub>	404	391	3.17	16.50	1.28	0.77	5.76	21.34	49% H→L + 27% H-2→L
7:S <sub>0</sub> → S <sub>2</sub>		373	3.28	28.06	2.28	0.80	6.36	24.24	39% H-1→L + 21% H→L+1 + 11% H-2→L+1
8:S <sub>0</sub> → S <sub>1</sub>	400	381	3.26	14.13	1.13	0.80	6.39	24.61	52% H→L + 19% H-2→L
8:S <sub>0</sub> → S <sub>2</sub>		368	3.37	29.74	2.45	0.82	6.74	26.46	43% H-1→L + 19% H→L+1 + 11% H-1→L+2 + 11% H→L+3

compounds **3** and **4**. The energy differences between the S<sub>1</sub> and S<sub>2</sub> excited states for compounds **1** to **3** are, in order, equal to 0.07, 0.17 and 0.19 eV. Interpreted using a Valence-Bond or Essential-State model, this increase can be ascribed to the strengthening of the interaction between the two branches of the molecules. In compound **4**, the splitting between the two first excited states reduces to 0.14 eV, suggesting a lowering in the strength of the interaction. Reversely, the transition probabilities toward the S<sub>1</sub> and S<sub>2</sub> states regularly increases from **1** to **4**, according to the increase of the transition dipole moments. This increase is particularly marked for transition dipoles associated to the S<sub>0</sub> → S<sub>2</sub> excitation, which are aligned parallel to the charge transfer axis of the molecules.

Compared to compound **5**, the addition of a second phenylacetylene unit in compound **6** blue-shifts the first (second) absorption band by 0.12 (0.03) eV, and reduces the splitting between S<sub>1</sub> and S<sub>2</sub> from 0.19 to 0.10 eV. The strengths of the S<sub>0</sub> → S<sub>1</sub> and S<sub>0</sub> → S<sub>2</sub> transitions are also strongly increased. A slight blue-shift of the two absorption bands is also observed from **7** to **8**, showing that the position of the triple bond within the conjugated branches impact the vertical transition energies, but without any change in the S<sub>1</sub>-S<sub>2</sub> gap. Note that the S<sub>0</sub> → S<sub>1</sub> transition probability decreases while that of the S<sub>0</sub> → S<sub>2</sub> transition slightly increases going from **7** to **8**.

As expected for push-pull conjugated dyes of C<sub>2</sub> symmetry, the S<sub>0</sub> → S<sub>1</sub> absorption band is dominated by a  $\pi \rightarrow \pi^*$  electronic transition between the highest occupied molecular orbital (HOMO, H) and lowest unoccupied molecular orbital (LUMO, L), while the S<sub>0</sub> → S<sub>2</sub> is dominated by H-1→L and H→L+1 electronic excitations. As illustrated in Fig. 3, the HOMOs are delocalized over the donor moiety of the pyrazine derivatives, whereas the LUMOs are spread over the acceptor one, revealing a significant intramolecular charge transfer upon light excitation. However, the evolution observed in the transition energies do not always follow the evolution of the HOMO-LUMO gap, since other electronic excitations do contribute to the S<sub>0</sub>→S<sub>1</sub> and S<sub>0</sub>→S<sub>2</sub> absorption bands.

To gain better insight on the ICT occurring upon the excitation process, Fig. 3 also illustrates the difference between the total electron densities of the ground and excited state:

$$\Delta\rho = \rho_{S_1} - \rho_{S_0} = \Delta\rho^+ + \Delta\rho^- \quad (17)$$

where  $\Delta\rho^+ > 0$  and  $\Delta\rho^- < 0$  correspond to increasing and decreasing areas of the electron density, respectively. In line with previous studies on rod-like push-pull compounds [40–42], the photo-induced charge

displacement (reported in Table 2) is evaluated as:

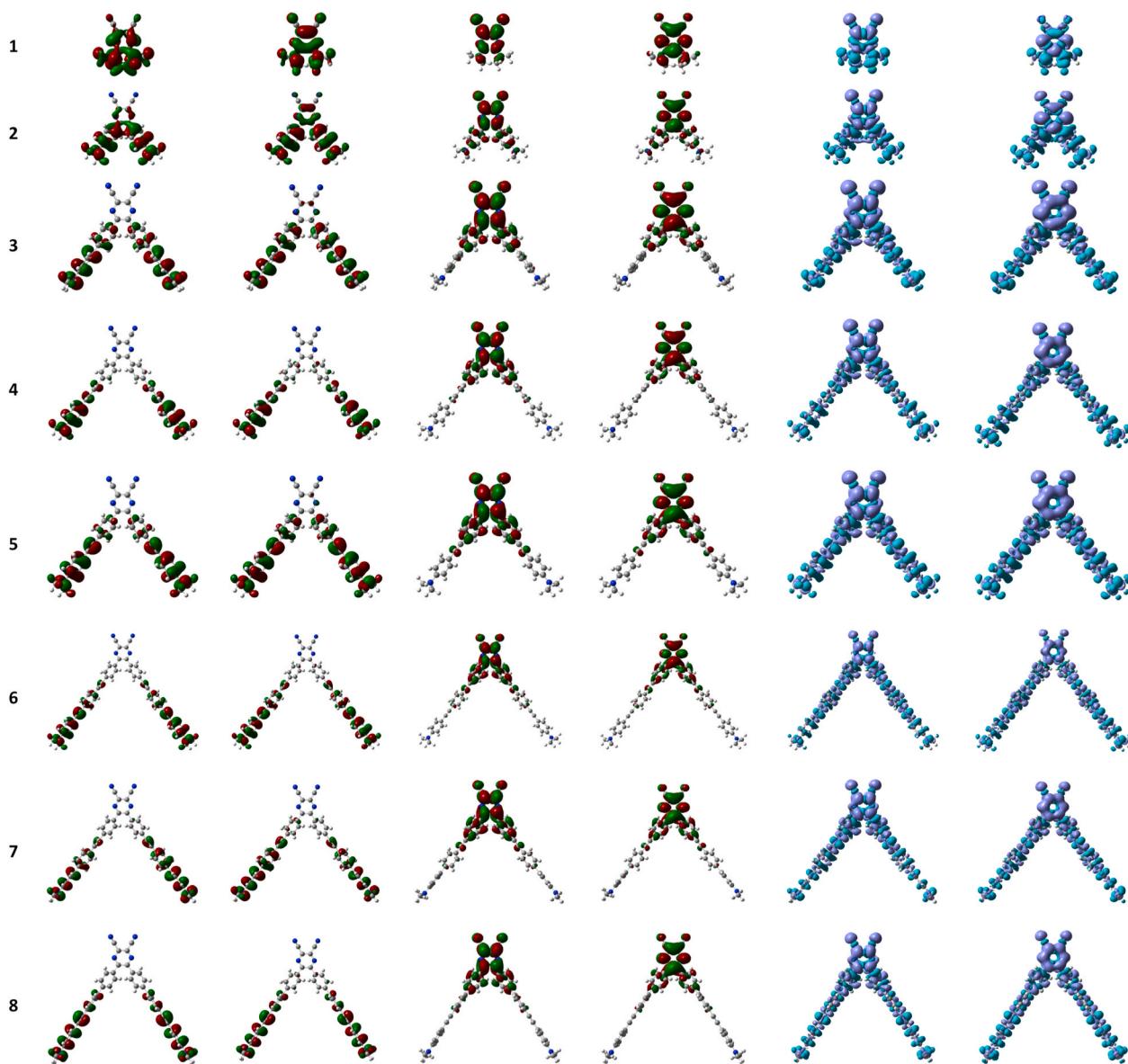
$$\Delta q = \int \Delta\rho^+(\mathbf{r}) d\mathbf{r} = - \int \Delta\rho^-(\mathbf{r}) d\mathbf{r} \quad (18)$$

While the charge transfer distance  $\Delta r$  is assimilated to the distance between the centroids of the  $\Delta\rho^+$  and  $\Delta\rho^-$  volumes. The change in dipole moment upon light excitation can then be expressed as  $\Delta\mu_{ge} = \Delta q \times \Delta r$ . As reported in Table 2, the  $\Delta\mu_{ge}$  values increase with the length of the conjugated linker for both the S<sub>1</sub> and S<sub>2</sub> states. This enhancement is particularly marked in the **1**–**4** series of compounds, and mainly originates from the regular increase of the charge transfer distances  $\Delta r$ , while  $\Delta q$  values saturate from compound **3** to **4**. A similar enhancement of  $\Delta\mu_{ge}$  values, although of weaker amplitude, is observed upon addition of a phenylacetylene unit from **5** to **6**. We also note that the photo-induced charge displacement is slightly larger in compound **8** than in compound **7** for both the S<sub>1</sub> and S<sub>2</sub> states.

### 3.3. Nonlinear optical properties

The results calculated at the IEF-PCM/M06-2X/6-311+G(d) level, for an incident wavelength  $\lambda$  of 1064 nm and in the static limit ( $\lambda = \infty$ ), are collected in Table 3. The independent non-negligible components of the static first hyperpolarizability tensor are gathered in Table S1. Before analyzing the data, it is interesting to assess if the present family of compounds could be treated as having a planar C<sub>2v</sub> symmetry, as assumed in earlier works for other Λ-shaped chromophores [24,25,28]. The contribution of the out-of-plane components to the total HRS hyperpolarizability can be evaluated by comparing the full  $\beta_{HRS}$  value to the one evaluated by using equation (8). As reported in Table 3, the  $\beta_{HRS}(C_{2v})$  values differ from the full  $\beta_{HRS}$  values by about –20% for all compounds, except **1** and **8** for which the differences reach –27% and –14%, respectively. These results evidence that assuming a planar C<sub>2v</sub> symmetry would not be fully justified for the compounds investigated here.

The data collected in Table S1 also show that the ratio between the main transverse and longitudinal components,  $R = \beta_{xxx}/\beta_{zzz}$ , decreases as increasing the length of the conjugated linker, both in the **1**–**4** molecular series and from **5** to **6**. The R value also decreases significantly from compound **7** to **8**, as the acetylenic bridge is shifted from position 2 to position 3 with respect to the pyrazine unit. It is also informative to rely on the Sum-Over-States (SOS) formalism to address the individual contributions of the low-lying excited states S<sub>1</sub> and S<sub>2</sub> to the two main



**Fig. 3.** From left to right: HOMO-1, HOMO, LUMO and LUMO+1 of the eight pyrazine derivatives, and electron density differences associated to the  $S_0 \rightarrow S_1$  and  $S_0 \rightarrow S_2$  transitions. Dark (light) blue lobes in density maps are associated with positive (negative)  $\Delta\rho$  values.

**Table 3**

Static and dynamic HRS data calculated at the IEF-PCM/M06-2X/6-311+G(d) level in dichloromethane: first hyperpolarizability ( $\beta_{HRS}$ ), anisotropy ratio ( $\rho$ ) and depolarization ratio ( $DR$ ).  $\beta_{HRS}(C_{2v})$  values are calculated using equation (8), with deviations with respect to the full  $\beta_{HRS}$  values given in parentheses. All first hyperpolarizability values are reported in a.u. according the T convention [43]. 1 a.u. of  $\beta = 3.6213 \times 10^{-42} \text{ m}^4 \text{V}^{-1} = 3.2064 \times 10^{-53} \text{ C}^3 \text{m}^3 \text{J}^{-2} = 8.639 \times 10^{-33} \text{ esu}$ .

Compound	Calc. ( $\lambda = \infty$ )				Calc. ( $\lambda = 1064 \text{ nm}$ )		
	$\beta_{HRS}$	$\rho$	$DR$	$\beta_{HRS}(C_{2v})$	$\beta_{HRS}$	$\rho$	$DR$
1	1386	0.79	5.11	1010 (-27%)	1386	0.67	5.75
2	10685	0.68	5.69	8366 (-22%)	18291	0.52	6.64
3	15492	0.65	5.85	12489 (-19%)	30603	0.58	6.24
4	13286	0.65	5.85	10701 (-19%)	25744	0.61	6.07
5	25678	0.66	5.81	20403 (-21%)	59990	0.61	6.09
6	30752	0.63	5.97	24987 (-19%)	83428	0.57	6.33
7	21955	0.65	5.87	17682 (-19%)	53742	0.58	6.27
8	21181	0.58	6.28	18322 (-14%)	48496	0.50	6.74

first hyperpolarizability components. Since the  $S_0 \rightarrow S_2$  excitation is associated to a transition dipole moment ( $\vec{\mu}_{02}$ ) parallel to the  $C_2$  axis, and since the dipole moment difference ( $\Delta \vec{\mu}_{02}$ ) is oriented along the same axis due to the molecular symmetry, only the  $S_2$  state contributes to the  $\beta_{zzz}$  component according to the following expression:

$$\beta_{zzz}^{S_2} = 6 \frac{\mu_{01}^2 \Delta \mu_{02}}{\Delta E_{02}^2} \quad (19)$$

Similarly, the  $S_1$  state contributes to the  $\beta_{zxx}$  component since  $\vec{\mu}_{01}$  and  $\Delta \vec{\mu}_{01}$  are respectively oriented perpendicular and parallel to the symmetry axis:

$$\beta_{zxx}^{S_1} = 2 \frac{\mu_{01}^2 \Delta \mu_{01}}{\Delta E_{01}^2} \quad (20)$$

Note that the  $S_2$  state also contributes to  $\beta_{zxx}$  through a cross-term (not considered here) implying the dipolar coupling with  $S_1$ :

$$\beta_{zxx}^{S_1/S_2} = 4 \frac{\mu_{01} \mu_{12} \mu_{02}}{\Delta E_{01} \Delta E_{02}} \quad (21)$$

where  $\mu_{12} = \langle 1 | \hat{\mu}_x | 2 \rangle$  and  $\hat{\mu}_x$  is the dipole moment operator along  $x$ . Although the values of  $\beta_{zxx}^{S_1}$  and  $\beta_{zzz}^{S_2}$  (calculated using spectroscopic data of Table 2) correlate with the full  $\beta_{zxx}$  and  $\beta_{zzz}$  components (see Table S1 and Fig. S9), the two sets of values differ significantly, which evidence that the two main components of the  $\beta$  tensor cannot be reduced to the sole contributions of the low-lying  $S_1$  and  $S_2$  states.

We continue the discussion by comparing the theoretical results with experimental measurements. The HRS data of the eight investigated pyrazine derivatives, as measured in DCM solutions, are reported in Table 4. 3D representations of the scattered light as a function of the incident power and chromophore concentration (equation (1)), as well as polarization scans of  $I_{\Psi\Psi}^{2\omega}$  at constant incident power (equation (15)) are reported in Figs. S2–S5. Note that NLO measurements for some compounds are subject to significant errors due to two-photon induced fluorescence effects overlapping with the HRS signal. Fig. 4 compares the experimental and theoretical results, and illustrates the evolution of  $\beta_{HRS}$  and DR values within three subgroups of compounds presenting progressive structural variations, {1–4}, {2,5,6} and {7–8}.

Within the {1–4} series, elongating the conjugation path from 1 to 3 leads to a significant enhancement of  $\beta_{HRS}$ , while further addition of a phenylene linker in compound 4 is detrimental to the HRS response. Compared to experiments, both static and dynamic TDDFT calculations underestimate the HRS responses of the compounds, but reproduce well the trend. This non monotonic evolution of  $\beta_{HRS}$  results from a combination of different effects, which can be interpreted with the help of the SOS expressions (equations (19) and (20)) and the computational results collected in Table 2. From 1 to 2, the increase of  $\beta_{HRS}$  originates from the decrease of the excitation energies towards the  $S_1$  and  $S_2$  states,

combined with the increase of the dipole moment differences ( $\Delta \mu_{ge}$ ) and transition dipole moments ( $\mu_{ge}$ ). From 2 to 3,  $\beta_{HRS}$  increases despite the blueshift of the low-lying absorption bands, because of the large enhancement of  $\mu_{ge}$  and  $\Delta \mu_{ge}$  values. Finally, the lowering of  $\beta_{HRS}$  from 3 to 4, well predicted by TDDFT calculations, cannot be rationalized by assuming that only the  $S_1$  and  $S_2$  states contribute to the first hyperpolarizability. Indeed, equations (19) and (20) predict an increase of the longitudinal and transverse  $\beta$  components, revealing significant (negative) contributions of higher energy excited states or possibly of the  $\beta_{zxx}^{S_1/S_2}$  coupling term (equation (21)).

It is also worth noticing that the evolution of  $\beta_{HRS}$  from 1 to 4 also follows that of the molar extinction coefficient ( $\epsilon_\lambda$ ) measured at the harmonic wavelength ( $\lambda = 532$  nm), which increases from 0 to 3000 and  $3500 \text{ L mol}^{-1} \text{ cm}^{-1}$  from 1 to 3, and then decreases to  $750 \text{ L mol}^{-1} \text{ cm}^{-1}$  for 4. Although absorption losses at the second harmonic wavelength are accounted for through a corrective term in equation (1), the slight overlap of the absorption band edge and the scattered light suggests that frequency dispersion effects also contribute to (and reinforce) the non monotonic evolution of  $\beta_{HRS}$  with the number of 1,4-phenylene rings in the linker. This is confirmed when comparing the evolution with the molecular size of the computed static and dynamic HRS hyperpolarizabilities (Fig. 4, top left panel).

Experimental anisotropy and depolarization ratios of chromophores 1–4, respectively smaller than 1 and close to 5, indicate that the dipolar component of  $\beta$  is larger than the octupolar one. TDDFT calculations also predict a dominant dipolar character of the second-order NLO responses, although the calculated DR values are systematically larger than the measured ones. This dipolar character is further illustrated in Fig. 5, which provides a unit-sphere representation (USR) of the calculated static first hyperpolarizability [44,45] (see SI for details). As previously evidenced for a  $C_2$ -symmetric push-pull bifluorene derivative [46], the effective dipoles are oriented along the charge-transfer axes of the two molecular branches, which gives a helmet-like shape to the projection of the  $\beta$  tensor onto the pyrazine plane.

Importantly, the  $DR_{NK}$  values estimated for compounds 3 and 4 are much larger than the ones obtained assuming Kleinman symmetry, and reach values larger than 5, in closer agreement with the calculations. This result evidences the limit of applicability of the Kleinman approximation when the frequency of the incident optical field approaches the two-photon resonance. Such deviations from index permutation symmetry have been previously revealed by means of polarization dependent electric field-induced second harmonic generation and electrooptical absorption measurements for other  $\Lambda$ -shaped derivatives, and were attributed to the presence of the low-lying perpendicular transitions [24]. They are further illustrated here by the calculated dynamic values of the  $\beta_{zxx}$  and  $\beta_{xxz}$  components (Table S2), which significantly differ from each other.

We now consider the {2,5,6} series. Comparing compounds 2 and 5,

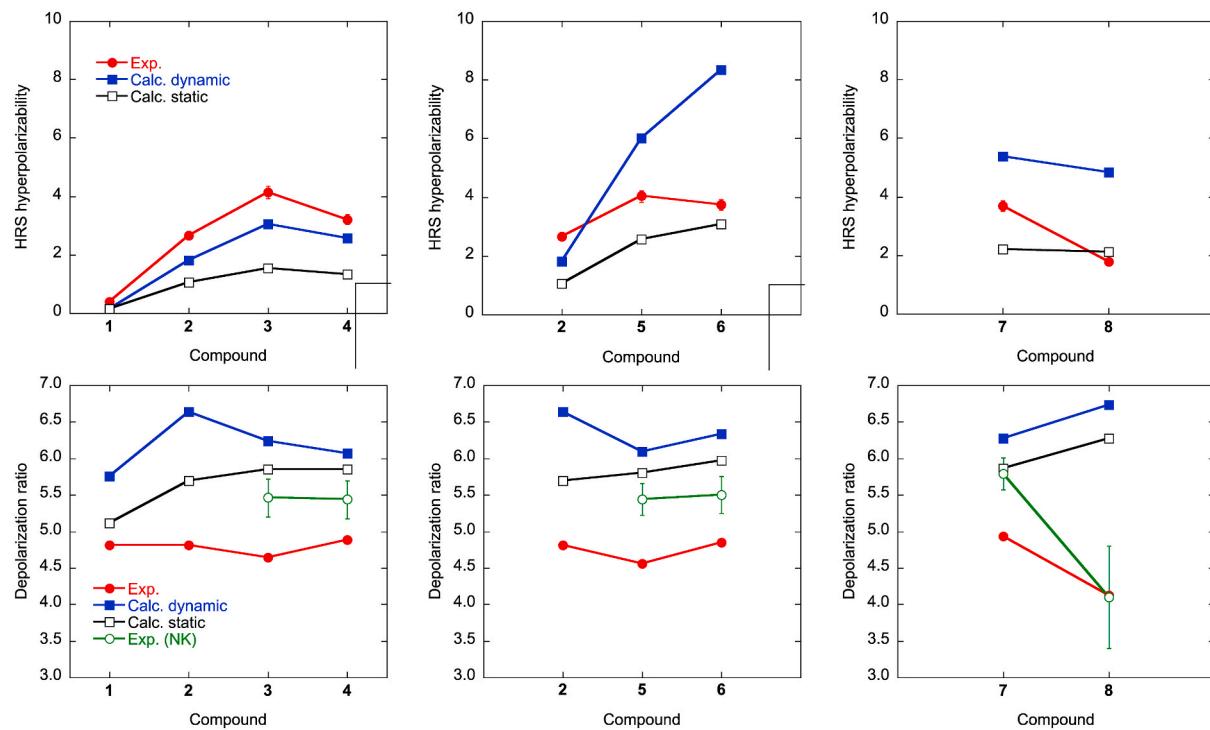
**Table 4**

HRS data measured in dichloromethane: first hyperpolarizability ( $\beta_{HRS}$ ), anisotropy ratio ( $\rho$ ), depolarization ratios estimated with (DR) and without ( $DR_{NK}$ ) assuming Kleinman symmetry, and deviation from Kleinman ( $\delta_{NK}$ , see equation (15)).  $\Delta_{DR} = (DR_{NK} - DR)/DR$ . First hyperpolarizability values are reported in a.u. according the T convention [43]. 1 a.u. of  $\beta = 3.6213 \times 10^{-42} \text{ m}^4 \text{ V}^{-1} = 3.2064 \times 10^{-53} \text{ C}^3 \text{ m}^3 \text{ J}^{-2} = 8.639 \times 10^{-33} \text{ esu}$ . Experimental errors are deduced from the fits (see SI).

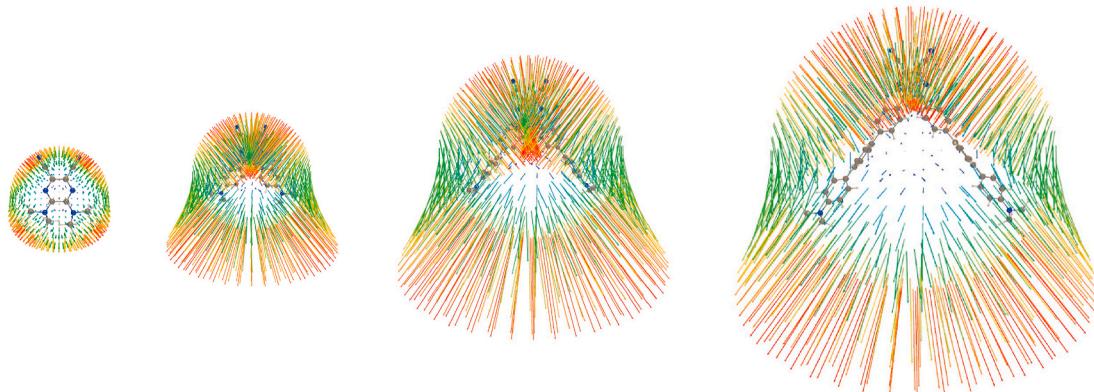
Compound	$\beta_{HRS}$	$\rho$	DR	$DR_{NK}$	$\delta_{NK}$	$\Delta_{DR}$
1	$4060 \pm 210$	0.85 <sup>a</sup>	4.81	–	–	–
2	$26700 \pm 1350$	$0.86 \pm 0.11^b$	4.81	–	–	–
3	$41000 \pm 2050$	$0.90 \pm 0.02$	4.64	$5.46 \pm 0.26$	$1.83 \pm 0.53$	0.18
4	$32140 \pm 1600$	$0.84 \pm 0.02$	4.89	$5.44 \pm 0.26$	$1.19 \pm 0.50$	0.11
5	$40400 \pm 2050$	$0.92 \pm 0.02$	4.56	$5.66 \pm 0.22$	$2.57 \pm 0.47$	0.24
6	$37530 \pm 1900$	$0.85 \pm 0.02$	4.85	$5.50 \pm 0.25$	$1.45 \pm 0.49$	0.13
7	$36980 \pm 1850$	$0.83 \pm 0.02$	4.94	$5.79 \pm 0.22$	$1.81 \pm 0.42$	0.17
8	$17710 \pm 890$	$1.04 \pm 0.10^b$	4.13	$4.10 \pm 0.70$	$0.09 \pm 1.63$	-0.01

<sup>a</sup> Conjectured owing to the very weak HRS signal.

<sup>b</sup> Large errors are due to two-photon fluorescence.



**Fig. 4.** Evolution of the HRS hyperpolarizability (in  $10^4$  a.u., top) and depolarization ratio (bottom) in the series of compounds 1–4 (left), 2,5,6 (middle) and 7–8 (right).



**Fig. 5.** Unit-sphere representation of the static first hyperpolarizability calculated at the TDDFT/M06–2X/6–311+G(d) for compounds 1 to 4 (from left to right). Arrows represent the magnitude of the second-order contribution to the induced dipole when an external field is applied normally to the surface of a virtual sphere centered on the center of mass of the molecules. See SI and Fig. S10 for details on theUSR and color scales.

the addition of a phenylacetylene unit enhances  $\beta_{HRS}$  by a factor of 2.7, which is also reproduced by the calculations that predict an enhancement ratio of 3.3 and 2.4 of dynamic and static responses, respectively. This strong enhancement originates from the large increase of the transition dipole moments and of the dipole moment variations associated to these two electronic transitions (see Table 2). Adding a second phenylacetylene unit (from compound 5 to 6) gives rise to a lowering (~7%) of  $\beta_{HRS}$ , which can be ascribed to the blueshift of the main absorption band. Although TDDFT calculations reproduce well the energy upshift of the  $S_0 \rightarrow S_1$  and  $S_0 \rightarrow S_2$  transitions, the lowering of  $\beta_{HRS}$  is not reproduced, because of the large increase of the strength of the transition towards the two excited states. Similarly to the 1–4 series, the magnitude of  $\beta_{HRS}$  correlates with the evolution of the molar extinction coefficient at the harmonic wavelength ( $\epsilon_{532} \sim 0, 4000$  and  $500 \text{ L mol}^{-1} \text{ cm}^{-1}$  for compounds 2, 5 and 6, respectively). The computed dynamic DR values are overestimated compared to experiments, but in global

agreement with the experimental trend, showing a minimum for compound 5. Here also, the non-Kleinman DR values estimated for compounds 5 and 6 are much closer from the computed ones.

Finally, we compare the NLO responses of compounds 7 and 8, which incorporate three phenyl rings and one acetylenic bridge between the pyrazine unit and the terminal *N,N*-dimethylamino donor groups. The measured HRS hyperpolarizability is larger by 55% when the acetylenic linker is located in position 2 with respect to the pyrazine, which is consistent with a decrease of  $\epsilon_{532}$  values (from  $\sim 500$  to  $\sim 0 \text{ L mol}^{-1} \text{ cm}^{-1}$ ). Although of much smaller amplitude (~10%), the lowering of  $\beta_{HRS}$  from 7 to 8 is reproduced by dynamic TDDFT calculations, and can be ascribed to the increase of the transition energies as well as to the decrease of the dipole of the  $S_0 \rightarrow S_1$  transition. On the other hand, TDDFT calculations fail to reproduce the decrease of the DR from compound 7 to 8. The unexpectedly low DR and  $DR_{NK}$  values measured for 8 might find origin in experimental errors due to large two-photon

fluorescence effects.

#### 4. Conclusions

The second-order NLO responses of push-pull  $\Lambda$ -shaped pyrazine derivatives have been studied by means of HRS experiments and DFT calculations. This combination of techniques allowed to gain deep insight into the evolution of the first hyperpolarizabilities when varying the nature and length of the  $\pi$ -conjugated bridge linking the dicyanopyrazine acceptor unit to the *N,N*-dimethylamino donor groups. It is evidenced that the NLO responses of this series of compounds have a dominant dipolar character, and result from the additive contribution of two low-lying perpendicular electronic absorption bands. The largest structures exhibit large NLO responses, which points out the potential interest of these systems for optoelectronic applications. Results also clearly demonstrate that the second-order polarizability is not Kleinman-symmetric, and could not be finely interpreted assuming a planar  $C_{2v}$  symmetry for the molecular structures. More generally, this work illustrates the complementarity of experimental measurements and quantum chemical calculations for rationalizing the effects of structural symmetry, chemical modifications and two-photon resonance on the NLO properties of  $\pi$ -conjugated dyes.

#### CRediT authorship contribution statement

**Frédéric Castet:** Investigation, Conceptualization, Methodology, Formal analysis, Writing - original draft, Writing - review & editing. **Adrien Gillet:** Investigation. **Filip Bureš:** Resources, Writing - review & editing. **Aurélie Plaquet:** Investigation. **Vincent Rodriguez:** Conceptualization, Methodology, Formal analysis, Writing - original draft, Writing - review & editing. **Benoît Champagne:** Conceptualization, Methodology, Formal analysis, Writing - original draft, Writing - review & editing.

#### Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.dyepig.2020.108850>.

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