

Effect of substituents on the excited-state dynamics of the modified DNA bases 2,4-diaminopyrimidine and 2,6-diaminopurine†

Zsolt Gengeliczki,^a Michael P. Callahan,^c Nathan Svadlenak,^c
Csaba István Pongor,^b Bálint Sztáray,^{bd} Leo Meerts,^g Dana Nachtigallova,^{*e}
Pavel Hobza,^e Mario Barbatti,^f Hans Lischka^{*ef} and Mattanah S. de Vries^{*c}

Received 1st September 2009, Accepted 9th February 2010

First published as an Advance Article on the web 8th April 2010

DOI: 10.1039/b917852j

To explore the excited state dynamics of pyrimidine derivatives, we performed a combined experimental and theoretical study. We present resonant two-photon ionization (R2PI) and IR-UV double resonance spectra of 2,4-diaminopyrimidine and 2,6-diaminopurine seeded in a supersonic jet by laser desorption. For 2,4-diaminopyrimidine ($S_0 \rightarrow S_1$ 34 459 cm^{-1}), we observed only the diamino tautomer with an excited state lifetime bracketed between experimental limits of 10 ps and 1 ns. For 2,6-diaminopurine, we observed two tautomers, the 9H- ($S_0 \rightarrow S_1$ 34 881 cm^{-1}) and 7H- ($S_0 \rightarrow S_1$ 32 215 cm^{-1}) diamino forms, with excited state lifetimes of 6.3 ± 0.4 ns and 8.7 ± 0.8 ns, respectively. We investigated the nature of the excited state of 2,4-diaminopyrimidine by means of multi-reference *ab initio* methods. The calculations of stationary points in the ground and excited states, minima on the S_0/S_1 crossing seam and connecting reaction paths show that several paths with negligible barriers exist, allowing ultrafast radiationless deactivation if excited at energies slightly higher than the band origin. The sub-nanosecond lifetime found experimentally is in good agreement with this finding.

Introduction

Gas phase laser spectroscopy provides the means to study the intrinsic properties of biologically relevant molecules in isolation. Such studies on RNA and DNA bases have revealed unique photophysical properties that are sensitive to subtle structural differences. In many cases, the biologically most relevant tautomeric form has a sub picosecond excited state lifetime, while other *tautomeric forms of the same compound* are much longer lived.^{1,2} We have even found that the Watson–Crick structure, adopted by the guanine-cytosine

(GC) base pair in DNA, appears to have a much shorter excited state lifetime than other *structures of the same base pair*.³ Theoretical models explain these short lifetimes by a rapid internal conversion in which the excited state (S_1) is coupled to the ground state (S_0) *via* pathways with no or a very small barrier leading to conical intersections.⁴ For the longer lived structures, small differences in relative energies cause the existence of barriers that lead to discrete spectra and lifetimes that can be two orders of magnitude longer. This rapid internal conversion pathway provides selected isomers with significantly enhanced photochemical stability, absent in the other longer lived structures. It is conceivable that these differences between excited state lifetimes of different bases and base-pair structures could have played a significant role in prebiotic chemistry.³

The five naturally occurring nucleic acid bases^{5–9} exhibit an ultrafast excited state relaxation in the gas phase. For DNA bases it has been shown^{1,10–24} that the energetically lowest conical intersections responsible for fast deactivation to the ground state are characterized by ring puckering modes. The most favorable ring puckering conical intersections result from the change of the HC6C5R (R = H, CH₃) dihedral angle in uracil and thymine and the HN1C2H dihedral angle in adenine. This model also explains why by contrast 2-aminopurine, in which the C2 position is substituted by the amino group, has a long excited state lifetime and strong fluorescence.^{25–27} Alternatively, the puckering at the C6 atom was also suggested to be responsible for adenine relaxation.²⁸

Recently, surface-hopping dynamics studies of 4-aminopyrimidine (4-APy)^{29–31} and 9H-adenine²⁴ showed that while the former relaxes into the ground state *via* different conical

^a Department of Chemistry, Stanford University, Stanford, CA 94305-5080, USA. E-mail: dana.nachtigallova@uochb.cas.cz, hans.lischka@univie.ac.at, devries@chem.ucsb.edu; Fax: (+1) 805-893-4120; Tel: (+1) 805-893-5921

^b Institute of Chemistry, Eötvös Loránd University Budapest, 1/A Pázmány P. stny., Hungary 1117

^c Department of Chemistry and Biochemistry, University of California, Santa Barbara, CA-93106-9510, USA

^d Department of Chemistry, University of the Pacific, Stockton, CA-95211, USA

^e The Institute of Organic Chemistry and Biochemistry, Flemingovo nám., 2, 166 10 Praha 6, Czech Republic

^f Institute for Theoretical Chemistry, University of Vienna, Waehringerstrasse 17, A1090 Vienna, Austria

^g Molecular and Biophysics Group, Institute for Molecules and Materials, Radboud University, 6500 GL Nijmegen, The Netherlands

† Electronic supplementary information (ESI) available: The calculated vertical ionization potentials for all the 2,4-diaminopyrimidine tautomers (Fig. S1 and Table S1). The vertical excitation energies calculated at the RICC2 method with various basis sets (Table S2). The interpolation curves between the S1min_C2 and S1min_C6 minima towards various MXS structures calculated at the CASSCF and MR-CISD(17)+Q methods (Fig. S2). This material is available free of charge on the Internet. See DOI: 10.1039/b917852j

intersections formed by puckering at the C6, N1, and C2 atoms, the latter relaxes exclusively through conical intersections formed by puckering at the C2 atom. This demonstrates the role of hindering imposed by the imidazole ring of adenine. Fig. 1 shows the structures and numbering of these two compounds as well as the 2-amino substituted analogues studied in the present paper, 2,4-diaminopyrimidine (2,4-DAPy) and 2,6-diaminopurine (2,6-DAPu). Fig. 1b and c show the lowest energy tautomers for the latter two compounds. One may expect that a structural modification of the pyrimidine ring in each of these molecules affects locations and accessibility of conical intersections. In particular, amino-substitution in the C2 position should affect conical intersections associated with puckering at that site, while addition of an imidazole ring to form purine bases should affect conical intersections associated with ring deformation at the C5 and C6 sites. Thus, in 2,4-DAPy puckering at the C2 position is affected, and in 2,6-DAPu puckering at both positions is modified. Therefore, it is especially useful to compare the photochemistry of these compounds.

In addition to these theoretical aspects there exists a further motivation for studying 2,4-DAPy and 2,6-DAPu bases. Joyce *et al.* have proposed that alternative nucleobases, which may form base pairs with geometries similar to Watson–Crick structures, may have played a role in constructing the first genetic code on the early Earth.³² Xanthine and 2,4-DAPy can form an unnatural base pair that fits the Watson–Crick geometry and can be incorporated into RNA and DNA by polymerases.^{33–36} Another base pair mimicking the Watson–Crick geometry is formed by 2,6-diaminopurine and uracil. For each of these molecules, there exists a plausible, prebiotic synthetic route.^{37–41}

Purines and pyrimidines can exist in a variety of different tautomeric forms, which can exhibit drastically different photophysical behavior.^{42–45} Therefore, investigating the effects of modification of the hetero-aromatic ring, in particular a possible immobilization of certain parts of this ring by its substitution at locations affecting the accessibility of conical intersections can refine our understanding of the intrinsic photostability of these alternative nucleobases.^{9,29–31}

In the series of prebiotic and alternate RNA/DNA bases, we previously studied the nucleobase xanthine.⁴⁶ In this paper we report the one-color R2PI spectra of 2,4-diaminopyrimidine (2,4-DAPy) and 2,6-diaminopurine (2,6-DAPu) as well as their IR-UV double resonance spectra. Because pyrimidine derivatives have relatively high ionization energies unavailable by two-photon ionization,⁴⁷ we measured the ionization energy of 2,4-DAPy by photoelectron spectroscopy to rule out three-photon resonant ionization schemes. To explain the nature and dynamics of the excited state of 2,4-DAPy, we characterized the stationary points in its ground and excited states and its S_1/S_0 conical intersections (minima on the crossing seam, MXS) as well as pathways to internal conversion by means of CASSCF, MR-CISD, and CASPT2 methods.

Experimental and theoretical methods

2,4-Diaminopyrimidine and 2,6-diaminopurine were purchased from Sigma-Aldrich and used without further purification.

The experimental setups have been described in detail elsewhere and only brief descriptions will be given here.^{48,49}

Laser spectroscopy

The desorption laser, a Nd:YAG operating at 1064 nm, is attenuated to 1 mJ cm^{-2} and focused to a spot approximately 0.5 mm diameter within 2 mm in front of the nozzle orifice. We translate the sample in order to expose fresh sample to successive laser shots. The nozzle consists of a pulsed valve with a nozzle diameter of 1 mm and a backing pressure of 6 atm of argon drive gas. The neutral molecules are skimmed and then ionized with a frequency doubled dye laser. We detect the ions in a reflectron time-of-flight mass spectrometer.

We obtain resonant two-photon ionization (R2PI) spectra by monitoring mass selected peaks while tuning the one-color, two-photon ionization wavelength. We measure UV-UV double resonance spectra with two laser pulses separated in time by 200 ns. Ionization laser intensities are approximately 3 mJ pulse^{-1} and are strongly attenuated to avoid saturation. The first pulse serves as a “burn” pulse, which removes the ground state population and causes depletion in the ion signal of the second “probe” pulse, provided both lasers are tuned to a resonance of the same tautomer. IR-UV double resonance spectra are obtained in an analogous way with the burn laser operating in the near-IR region. IR frequencies are produced in an OPO setup (LaserVision) pumped by a Nd:YAG laser operating at its fundamental frequency. For this work, we operated within the range of $3200\text{--}3800 \text{ cm}^{-1}$, which encompasses NH and OH modes. Typical IR intensities in the burn region are 12 mJ pulse^{-1} and the bandwidth is 3 cm^{-1} . Excited state lifetimes are measured in a two-color, two-photon ionization experiment. The pump and probe photons are separated by a time delay of 100 ps to 100 ns, and the ion signal is recorded as a function of the time delay. The lifetime of the excited state is given as the time constant of the exponential decay of the ion signal.

Photoelectron spectroscopy

He-I photoelectron spectra were recorded on a custom-built ATOMKI ESA-32 instrument.⁴⁹ The spectrometer is equipped with a Leybold-Heraeus UVS 10/35 high-intensity helium discharge photon source. The sample was introduced into the ionization chamber *via* solid inlet probe at an elevated temperature of 260 °C. The spectra were calibrated against Ar $^2P_{3/2}$ and $^2P_{1/2}$ peaks. The maximum error in the ionization energies is estimated to be less than 0.04 eV, and the energy resolution of the He-I spectra was better than 40 meV (FWHM) as determined from Ar 3p ionizations. In the extraction of the vertical ionization potentials, Shirley background and pseudo-Voigt peaks were fitted on the experimental spectra. The adiabatic ionization potential can be estimated as the onset of the ionization and has a somewhat higher uncertainty than the vertical ionization potentials.

Quantum chemical calculations

Identifying the tautomers. To assign the IR-UV double resonance spectra, we computed the structures of the possible tautomers using density functional theory (DFT) and *ab initio*

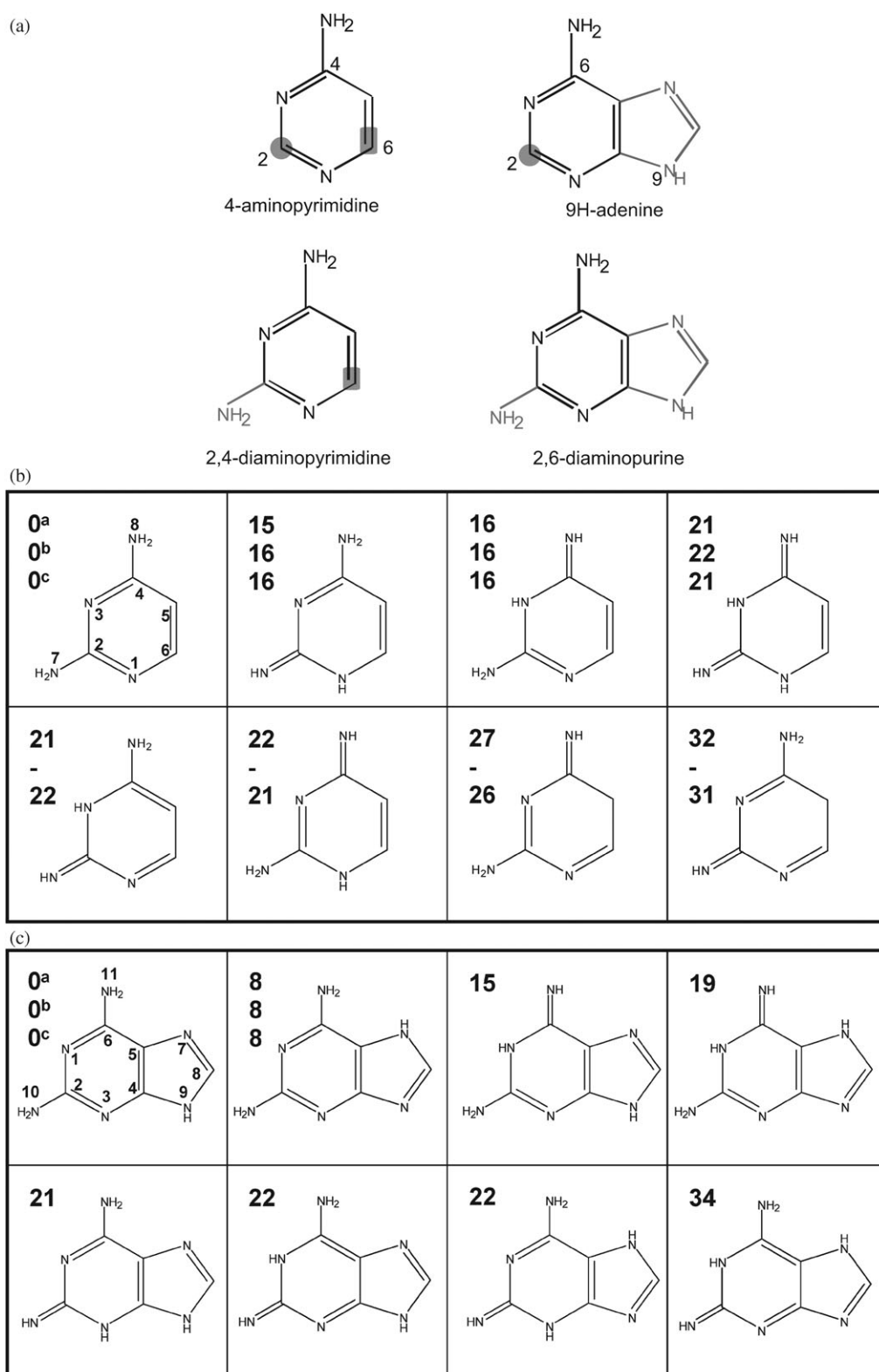


Fig. 1 (a) Structures of the four related compounds discussed in the text. Calculations predict conical intersections for 4-Apyr due to ring deformations at positions C2 and C6, C5, and N1. Amino substitution in the C2 position affects the conical intersection associated with that position, while addition of the 5 membered ring for the purines affects the conical intersections associated with ring deformation in the other positions. (b) The tautomers and numbering scheme of 2,4-diaminopyrimidine. Relative energies obtained at the (a) B3LYP/6-311+G(2d,p) (b) MP2(FC)/6-311+G(2d,p) and (c) G3 levels are in kcal mol⁻¹. (c) The tautomers and numbering scheme of 2,6-diaminopurine. Relative energies obtained at the (a) B3LYP/6-311+G(2d,p) (b) MP2(FC)/6-311+G(2d,p) and (c) G3 levels are in kcal mol⁻¹.

methods. We combined Becke's three-parameter hybrid functional⁵⁰ with the Lee–Yang–Parr exchange correlation functional^{51,52} (B3LYP) and the 6-311 + G(2d,p) basis set.^{53,54} We used the same basis set in frozen core second order Møller–Plesset perturbation (MP2) calculations.^{55–60} We verified the equilibrium structures by the absence of imaginary vibrational frequencies. The calculated vibrational frequencies were scaled by 0.9618⁶¹ (DFT) or 0.9496⁶² (MP2) to account for electronic structure method deficiency and anharmonicity. To establish the relative energies of the tautomers, we also applied the Gaussian-3 (G3) composite method.⁶³

Ionization energies. To assign the photoelectron spectrum, we calculated ionization energies for 2,4-diaminopyrimidine, using the B3LYP functional and 6-311 + G(2d,p) basis set. We calculated the adiabatic ionization potential as the energy difference between the equilibrium geometries of the ionic state and the ground state neutral molecule, estimating the first vertical ionization energy as the difference between the energies of the ionic state and the neutral molecule at the equilibrium geometry of the latter. Then, at the equilibrium geometry of the ion, we performed time-dependent DFT (TD-DFT) calculations to compute the vertical excitation energies of the ion.^{64–66} Vertical ionization energies are then the sum of the adiabatic ionization energies and vertical excitation energies of the ion. We also performed outer valence Green's function (OVGF) calculations with the 6-311 + G(2d,p) basis set at the equilibrium geometries of the neutral tautomers obtained in the DFT calculations.^{67–74} All calculations were carried out using the Gaussian 03 Rev. C. 02. quantum code package.⁷⁵

Excited state investigations of 2,4-diaminopyrimidine. We performed the calculations using the complete active space self-consistent field (CASCCF) and multi-reference interaction (MR-CI) methods for 2,4-DAPy. In the MR-CI approach, single and double excitations from the CI reference space are included (MR-CISD) and generalized interacting space restrictions are adopted.⁷⁶ We constructed the orbital space for CASCCF wavefunctions using 14 electrons in 10 orbitals, *i.e.* composed of eight π orbitals and two lone pairs located on the ring nitrogen atoms. We used a state-averaging procedure using three states (SA-3) at the CASCCF level throughout the calculations. Based on the CAS (14,10), we constructed the MR-CISD reference space by moving orbitals with natural occupation larger than 0.9 and smaller than 0.1 to the doubly occupied and virtual spaces, respectively, resulting in a MR-CI reference space composed of six electrons in five orbitals (MR-CISD(6,5)). All single and double excitations from this reference space were allowed; all core orbitals were frozen. To reduce the computational cost of the calculations, up to a total number of 17 doubly occupied orbitals were frozen in MRCI calculations of the reaction paths. We tested this procedure against full calculations for selected typical examples, taking into account the size-consistency effects by means of Pople's correction method, indicated by +Q.⁷⁷ We used the 6-31G** basis set throughout the calculations.^{78,79} For comparison and to verify the reliability of the applied basis set we also applied the complete active space self-consistent-field second-order perturbation theory (CASPT2) method^{80,81} with the same

reference CAS space and resolution-of-identity coupled cluster to the second-order (RICC2)^{82,83} method.

We determined the minima on the ground and excited S_1 surfaces and on the seam of conical intersections (MXS) by the CASSCF(14,10) method. We constructed reaction paths between structures of the S_1 minima and the MXSs using the method of linear interpolation of internal coordinates (LIIC). The CASSCF and MR-CISD energies of relevant points of the reaction paths were plotted as a function of mass-weighted distances between each point of the path and the S_1 minimum. We also determined the structures of the transition states for selected reaction paths using the CASSCF(14,10) method. We confirmed the character of the stationary point by Hessian calculations within a selected space of internal coordinates relevant for the ring puckering modes.

We simulated the absorption spectrum of 2,4-DAPy employing the RICC2 and CASSCF methods, using the Gaussian broadening method described by Barbatti *et al.*⁸⁴ We used the same active space and number of states in the average procedure as specified above. We generated five hundreds points by a Wigner distribution in the ground vibrational state of the ground electronic state, computed based on a Wigner distribution for the ground vibrational and electronic state, taking each nuclear degree of freedom within the harmonic approximation. The Wigner distribution composed of 500 different geometries was projected onto the excited state by multiplying it by the Einstein coefficient B computed for each point. We used a phenomenological broadening of 0.05 eV.

We performed the optimization of conical intersections using the analytic gradient and non-adiabatic coupling vectors^{85–89} available in the COLUMBUS program system.^{90–92} We performed the CASPT2 calculations with the MOLCAS program package^{93–95} and RICC2 computations were performed using the Turbomole program system.⁹⁶ The absorption spectrum was simulated using the NEWTON-X program package.^{84,97}

Results and discussion

2,4-Diaminopyrimidine R2PI and IR-UV double resonance spectroscopy

Fig. 2 shows the R2PI spectrum of 2,4-diaminopyrimidine in the wavelength range of 34 350–35 100 cm^{-1} . In a one-color experiment, we might not be able to observe the $S_1 \leftarrow S_0$ origin transition if the ionization potential of the molecule is too high. Therefore, we determined the ionization potential by photoelectron spectroscopy, as detailed below. For 2,4-diaminopyrimidine we found a relatively low vertical ionization potential of 8.30 ± 0.05 eV and an adiabatic ionization potential of 7.86 ± 0.05 eV, which corresponds to 33 471 cm^{-1} and 31 698 cm^{-1} photon wave numbers, respectively, in terms of the one-color experiments. We, therefore, tentatively assign the red-most peak at 34 459 cm^{-1} as the $S_1 \leftarrow S_0$ origin transition. If this assignment is correct, the intensity distribution in the spectrum suggests significant geometric deformation of the excited state relative to the ground state.

The R2PI spectrum consists of a single tautomer, based on IR-UV double resonance experiments at different UV probe wavelengths, as detailed below. Fig. 3 shows the IR spectrum,

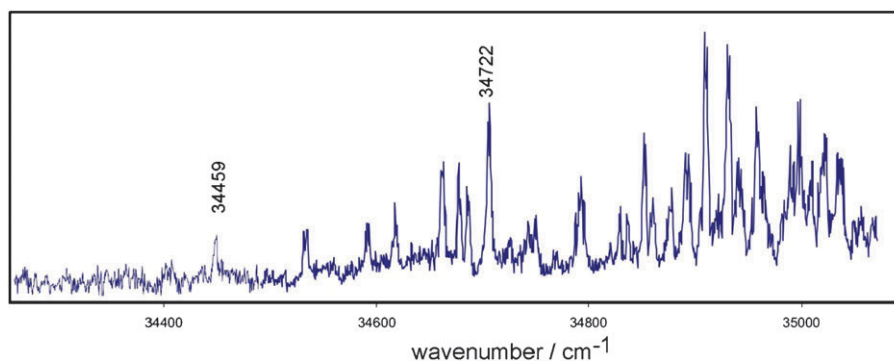


Fig. 2 The R2PI spectrum of 2,4-diaminopyrimidine. The $S_1 \leftarrow S_0(\pi\pi^*)$ origin is at 34459 cm^{-1} . Data were not corrected for variations in laser intensities.

obtained at a probe frequency of 34516 cm^{-1} . The spectrum shows four distinct peaks in the frequency range of $3400\text{--}3600\text{ cm}^{-1}$. By fitting the experimental data with a sum of Lorentzian curves, we extracted vibrational frequencies of 3452 , 3464 , 3563 , and 3578 cm^{-1} . This indicates the presence of four free N–H stretching modes in the molecule and suggests that we observe the diamino form (see Fig. 1b). We computed the relative energies of all tautomers at the B3LYP/6-311+G(2d,p), MP2(FC)/6-311+G(2d,p) and G3 levels. At every level the di-amino form was the lowest energy tautomer with the second most stable tautomer at a relative energy 15 kcal/mol higher than the diamino form. We calculated the vibrational frequencies for all tautomers at the DFT and MP2 levels. The di-amino tautomer provides an excellent match between the calculated and the experimental frequencies. Table 1 summarizes the vibrational frequencies.

To determine that *only* the diamino tautomer is present in the R2PI spectrum, we set the IR frequency to 3578 cm^{-1} and scanned the R2PI spectrum again. The IR band at 3578 cm^{-1} is the most diagnostic band for the diamino tautomer

compared with the theoretically calculated IR bands for the other tautomers. The R2PI spectrum obtained while hole burning at this IR frequency showed a complete depletion in the ion signal for the entire spectral range when the IR was on. This result implies that all peaks in the R2PI spectrum are due to the diamino tautomer.

Photoelectron spectroscopy

Fig. 4 shows the photoelectron spectrum of 2,4-diaminopyrimidine. The adiabatic ionization potential, derived from the onset, is $7.86 \pm 0.05\text{ eV}$. We derived the vertical ionization potentials by fitting the sum of pseudo-Voigt curves on the observed bands. Based on the intensities and the widths of the bands, we find vertical ionization potentials corresponding to six ionic states. An additional experimentally unresolved broad band starts at an onset of 13 eV . It is implausible that precise ionization energies can be extracted from the spectrum above 13 eV , and we only note that the first maximum of this range is at 13.51 eV .

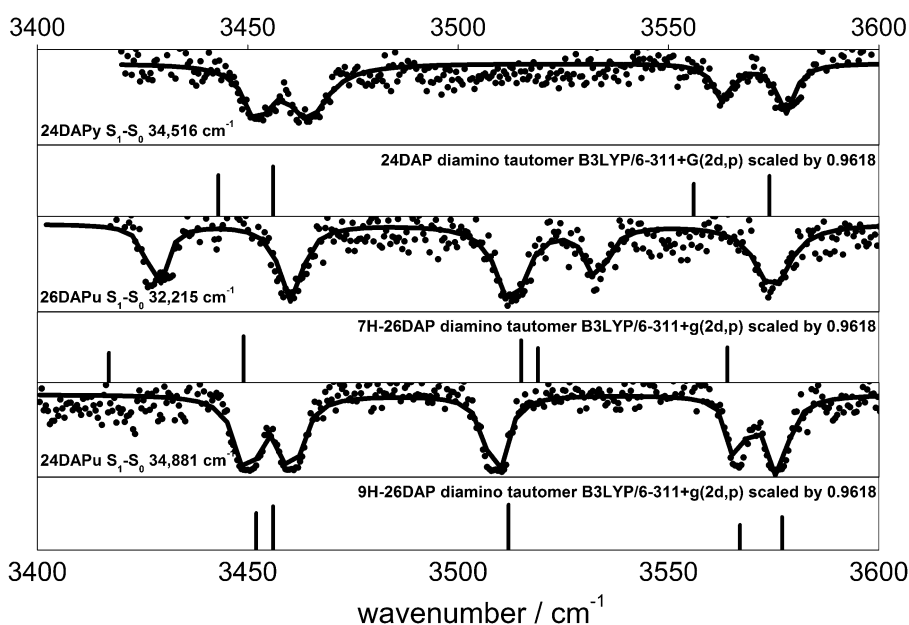


Fig. 3 The IR-UV double resonance spectra of 2,4-diaminopyrimidine and 2,6-diaminopurine in the N–H stretch IR wavelength range. The stick spectra represent the best matching calculated vibrational frequencies at the B3LYP/6-311+G(2d,p) level. A scaling factor of 0.9618 was applied.

Table 1 Calculated vibrational frequencies for tautomers of 2,4-diaminopyrimidine in the N–H region. All frequencies have been scaled and given in cm^{-1} . MP2 scaling factor: 0.9496; B3LYP scaling factor: 0.9618. For numbering scheme, see Fig. 1. For comparison, the experimental vibrational frequencies are 3452, 3464, 3563, and 3578 cm^{-1}

Tautomer	MP2 6-311+G(2d,p) ω/cm^{-1}	B3LYP 6-311+G(2d,p) ω/cm^{-1}	Vib. Mode
24DAP diamino tautomer	3532	3574	N7H as
	3518	3556	N8H as
	3410	3456	N7H s
	3400	3443	N8H s
1H, 7 imino	3527	3564	N8H as
	3440	3483	N1H
	3403	3446	N8H s
	3351	3386	N7H
	3496	3534	N7H as
3H, 8 imino	3403	3445	N3H
	3387	3431	N7H s
	3346	3380	N8H
	3467	3507	N1H
	3422	3469	N3H
1H,3H 7,8 imino	3359	3392	N10H
	3344	3378	N11H
		3535	N8H as
		3454	N3H
		3435	N8H s
1H, 8 imino		3387	N7H
		3508	N7H as
		3492	N1H
		3411	N7H s
		3328	N8H
5H, 8 imino		3593	N7H as
		3465	N7H s
		3316	N8H
		2986	CH
		3559	N8H as
5H, 7 imino		3436	N8H s
		3345	N7H
		2972	CH

Considering the relative energies of the tautomers of 2,4-diaminopyrimidine, it is very unlikely that we observe more than one tautomer in the gas phase, even at the applied inlet temperature of $260 \text{ }^\circ\text{C}$. However, we performed restricted OVG calculations at the 6-311+G(2d,p) level for all the possible isomers, employing single point calculations on the DFT equilibrium geometries. The outer valence Green's function method is a useful tool in the assignment of photoelectron spectra of organic molecules. The vertical ionization potentials are usually predicted within 0.3 eV .⁹⁸ We obtained the best match with the experimental values for the most stable

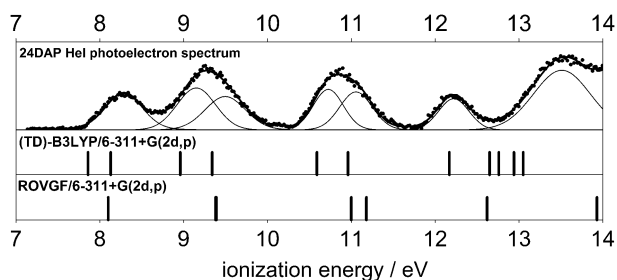


Fig. 4 He-I photoelectron spectrum of 2,4-diaminopyrimidine. pseudo-Voigt shaped peaks were fitted on the observed bands to extract the vertical ionization potentials. The adiabatic ionization potential was taken as the ionization onset at 7.86 eV . The adiabatic and vertical ionization energies were also computed at the (TD)-B3LYP/6-311+G(2d,p) and ROVGF/6-311+G(2d,p) levels and represented by the stick spectra. See text for details.

tautomer, the diamino form. Fig. 4 lists the calculated ionization potentials. Photoelectron spectroscopy can also be regarded as a tool for probing the ionic states. To obtain adiabatic and vertical ionization energies for the diamino form, we carried out TD-B3LYP/6-311+G(2d,p) calculations at the equilibrium geometry of the ion. According to both methods, the first peak of the spectrum can be assigned to the ionization of the HOMO, that is, the removal of an electron from the π system.

Table 2 lists the adiabatic and the lowest vertical ionization energies. The predicted vertical ionization energies of all the possible tautomers are compared to the experimental spectrum in the Supporting Information (Fig. S1 and Table S1, ESI).[†]

2,6-Diaminopurine R2PI and IR-UV double resonance spectroscopy

Previously, we reported a resonant two-photon ionization spectrum of 2,6-diaminopurine between 32000 cm^{-1} and 34000 cm^{-1} without identifying the observed tautomer. In the present study, we extended the R2PI spectrum up to 35400 cm^{-1} , as shown in Fig. 5. Two groups of peaks can be distinguished, based on hole burning, with possible $S_1 \leftarrow S_0$ origins at 32215 cm^{-1} and 34881 cm^{-1} , respectively. We carried out IR-UV double resonance experiments on these two possible origins. The resulting IR spectra, shown in Fig. 3, are not identical, indicating the presence of at least two different tautomers in the gas phase. Both spectra exhibit five

Table 2 Experimental excitation and ionization energies (in eV) of 2,4-diaminopyrimidine and 2,6-diaminopurine

2,4-diaminopyrimidine		
Transition		$h\nu/\text{eV}$
$S_1 \leftarrow S_0$		4.27
$D_0 \leftarrow S_0$	IE_{ad}	7.86
	IE_{vert}	8.30
7H-2,6-diaminopurine		
$S_1 \leftarrow S_0$		3.99
9H-2,6-diaminopurine		
$S_1 \leftarrow S_0$		4.32

IR bands. With the UV probe set at 34881 cm^{-1} IR frequencies appear at 3450 , 3460 , 3509 , 3566 , and 3576 cm^{-1} . These frequencies match very well with the calculated IR frequencies of 2,6-diaminopurine. We assign the peaks at 3566 and 3576 cm^{-1} to the antisymmetric combinations of the N–H stretches in the amino groups, the peaks at 3450 and 3460 cm^{-1} to the symmetric combinations of the same stretches, and the peak at 3509 cm^{-1} to either the N9–H or the N7–H stretching mode. A hydrogen atom on the N9 position seems to be more plausible, because it might have a smaller effect on the N–H stretching modes of the amino groups, as discussed in the next paragraph.

With the UV probe at 32215 cm^{-1} , the five IR bands appear at 3428 , 3460 , 3513 , 3533 , and 3575 cm^{-1} . When compared to the former IR spectrum, significant red shifts of two peaks are apparent. The peak at 3450 cm^{-1} is shifted to 3428 cm^{-1} , and the peak at 3566 cm^{-1} is shifted to 3533 cm^{-1} . If the assignment of the first IR spectrum is correct, this shift can be explained by placing the hydrogen atom for this tautomer on the N7 position instead of the N9 position. That would perturb the N–H stretching modes in the N11 amino group. DFT (B3LYP/6-311+G(2d,p)) and *ab initio* (MP2(FC)/6-311+G(2d,p)) calculations of the vibrational frequencies of the various tautomers support this qualitative assignment. Table 3 shows the comparison of the calculated vibrational frequencies with the experimental values. The two diamino forms provide the best match. Furthermore, when calculating

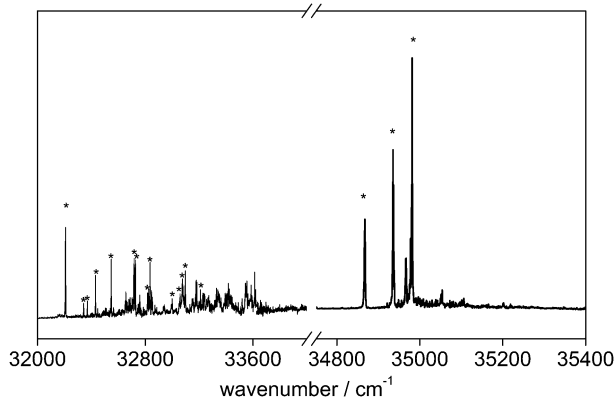


Fig. 5 The R2PI spectra of the 7H and 9H tautomers of 2,6-diaminopurine. The $S_1 \leftarrow S_0(\pi\pi^*)$ origins are at 32215 cm^{-1} and 34881 cm^{-1} , respectively. Asterisks indicate UV wavelengths that were checked for ion signal depletion at diagnostic IR bands.

the relative energies of the tautomers the N9H di-amino tautomer is the most stable form, followed by the N7H di-amino tautomer, at every level of calculation.

Excited state lifetimes

To obtain a measure of excited state lifetimes we performed two color pump–probe experiments, in which we resonantly excited the molecule to the S_1 state, followed by ionization out of the excited state with a 266 nm photon from a second laser, with a variable delay between the two pulses. Fig. 6 shows the results for 2,6-DAPu. Fitting the decay curves with a single exponential decay, we obtained excited state lifetimes of $6.3 \pm 0.4\text{ ns}$ for the N9H tautomer and $8.7 \pm 0.8\text{ ns}$ for the N7H tautomer. Using laser pulses of about 5 ns pulse width imposes a lower limit of the order of a few nanoseconds on these lifetime measurements.

In the case of 2,4-DAPy, the lifetime is shorter than this experimental limit so we were unable to perform two-color experiments and our attempts to carry out UV-UV double resonance experiments also failed. In this case we can use the peak width in the R2PI spectrum to obtain a rough estimate of the lower limit of the excited state lifetime. Fig. 7 shows the peak at 34720 cm^{-1} together with simulations of the rotational envelope with different Lorentzian linewidths. The simulated lineshapes assume a dominantly b-type transition and a rotational temperature of 20 K. These parameters are consistent with our simulations of the peaks in the 2,6-DAPur spectrum, for which the width is constrained by the experimental resolution only and not by lifetime broadening. The results, as shown in Fig. 7, suggest an upper limit to the linewidth of 0.5 cm^{-1} corresponding to a lower limit of the excited state lifetime of 10 ps. Therefore, we can bracket the experimental window of our lifetime measurements for 2,4-DAPy roughly between 10^{-11} and 10^{-9} s . This is clearly shorter than that of 2,6-DAPur, but it is longer than the lifetimes of 1.8 or 2.4 ps and 5.2 or 6.4 ps, reported for uracil and thymine, respectively, with excitation at 267 nm.⁹⁹ We also note that the UV spectra of uracil and thymine are broad while that of 2,4-DAPy is sharp.⁵

This lifetime is clearly shorter than that of 2,6-DAPu, but consistent with that of other pyrimidine bases excited at the band origin.^{100,101} Not surprisingly, this 10 ps–1 ns time range spans values much longer than the typical 1–6 ps found for pyrimidine bases excited close to the first band maximum, well above any barriers towards conical intersections.^{9,99} The sub-nanosecond lifetime of band-origin excited 2,4-DAPy is a strong indication that this molecule deactivates by means of internal conversion just like 4-APy does. In order to examine how the internal conversion takes place, we carried out extensive analysis of the excited-state surfaces of this molecule, which will be discussed in the next section.

Excited-state analysis of 2,4-diaminopyrimidine

Stationary points in the ground and excited states. Table 4 shows the vertical excitation energies calculated at the CASSCF, MR-CISD, and CASPT2 levels. At the CASSCF level, the first excited state is of $\pi\pi^*$ character while the second excited state is due to excitation from the lone pairs located on

Table 3 Calculated vibrational frequencies for tautomers of 2,6-diaminopurine in the N–H region. All frequencies have been scaled and given in cm^{-1} . MP2 scaling factor: 0.9496; B3LYP scaling factor: 0.9618. For numbering scheme see Fig. 1. For comparison, the experimental vibrational frequencies are 3450, 3460, 3509, 3566, and 3576 cm^{-1} (UV probe 34881 cm^{-1}), and 3428, 3460, 3513, 3533, and 3575 cm^{-1} (UV probe 32215 cm^{-1})

Tautomer	MP2 6-311+G(2d,p) ω/cm^{-1}	B3LYP 6-311+G(2d,p) ω/cm^{-1}	Vib. Mode
9H-26DAP diamino tautomer	3505	3577	N11H as
	3498	3567	N10H as
	3438	3512	N9H
	3380	3456	N11H s
	3379	3452	N10H s
7H-26DAP diamino tautomer	3493	3564	N10H as
	3454	3519	N11H as + N7H
	3356	3515	N7H + N11H s
	3375	3449	N10H s
	3346	3417	N11H s
1,9H, 11 imino		3525	N11H as
		3507	N9H
		3453	N1H
		3426	N10H s
		3377	N11H
1,7H, 11 imino		3508	N7H
		3500	N10H as
		3465	N1H
		3406	N10H s
		3351	N11H
3,9H, 10 imino		3586	N11H as
		3512	N9H
		3486	N3H
		3460	N11H s
		3391	N10H
1,9H, 10 imino		3543	N11H as
		3511	N9H
		3460	N1H
		3437	N11H s
		3392	N10H
3,7H, 10 imino		3522	N11H as + N7H
		3520	N7H + N11H as
		3484	N3H
		3417	N11H s
		3390	N10H
1,7H, 10 imino		3516	N11H as
		3485	N7H
		3460	N1H
		3422	N11H s
		3392	N10H

the nitrogen atoms of the pyrimidine ring ($n\pi^*$ transition). The calculated energy gap between these states is 0.34 eV. We obtained the same ordering of states at the CASPT2 and RICC2 levels, with a smaller energy gap amounting to 0.2 and 0.1 eV, respectively. Increasing the flexibility of the basis set does not change the ordering of the states and the energy gap between them (see Table 4 and Supplementary Material for CASPT2 and RICC2 results, respectively). At the MR-CISD+Q level the ordering is reversed with an energy gap of 0.3 eV.

The experimentally measured excitation energy at the band origin is 4.27 eV. To estimate the energy of the band maxima we have performed a simulation of the absorption spectra (see Fig. 8 for the absorption spectrum simulated with RICC2 method). From the figure we calculated the energy difference between the band maximum and band origin as 0.63 eV (the same result was obtained employing the CASSCF method). Adding this shift to the experimental value of the band origin results in a band maximum at 4.91 eV, in a good agreement with the calculated vertical excitation energies.

We found two minima on the S_1 surface (Fig. 9a) using CASSCF optimization, with energies of 4.75 and 5.08 eV above the ground state minimum (Table 4). Both structures show a distortion from the planarity of the ring which is more pronounced in the latter case. For both minima, the CASSCF wavefunctions show strong coupling between the $n\pi^*$ and $\pi\pi^*$ configurations with almost equal weight. Unlike the case of 4-aminopyrimidine,^{29,30} we did not succeed in locating a third minimum which is planar and of $\pi\pi^*$ character. Any attempt to find this minimum ended up in conformation S1min_C6.

The energy ordering of S1min_C6 and S1min_C2 remains the same at the MR-CISD(8)+Q level and at the CASPT2 level with an energy gap of about 0.2 and 0.3 eV, respectively. The energy ordering is reversed when 17 orbitals are frozen in the CI procedure, with the energetic changes being, however, quite small. The latter calculations place the S1min_C2 below the S1min_C6 by about 0.2 eV. The difference between the results with 8 and 17 orbitals frozen is however not significant for the discussion of reaction paths calculated by means of linear interpolation curves and, thus, the less accurate but

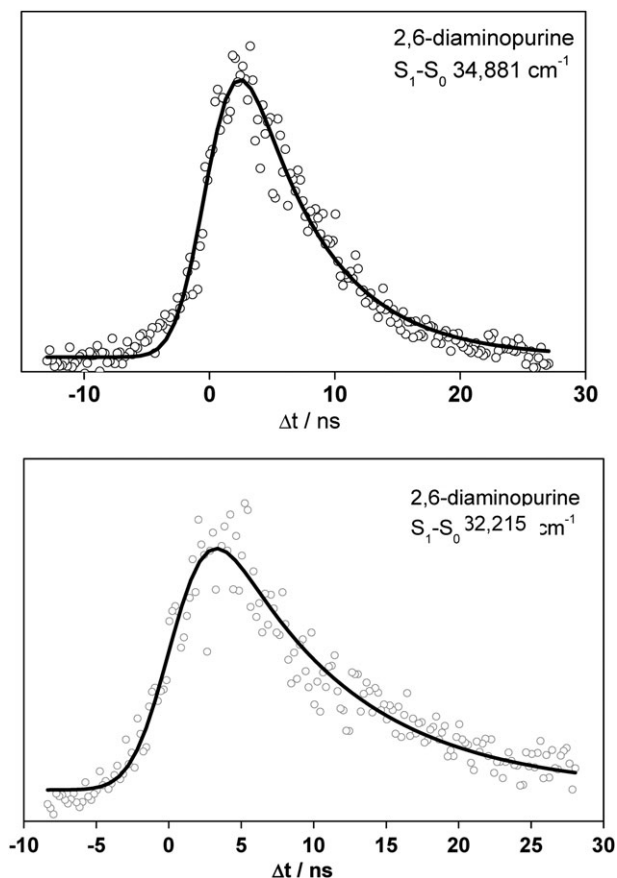


Fig. 6 Exponential decay of the ion signal of 2,6-diaminopurine in the two-color R2PI experiment. The extracted lifetime of the excited state is 8.7 ± 0.8 ns for the 7H tautomer and 6.3 ± 0.3 ns for the 9H tautomer.

more computationally feasible approach with 17 orbitals frozen can be used.

There is a substantial energetic relaxation of the first excited states when moving from the Franck–Condon region to the S1min_C6 and S1min_C2 optimized structures. This result is consistent with the experimentally observed low intensity of the 0–0 transition. Although the amount of relaxation energy from 2^1A into the S1min_C2 structure, calculated with different methods, is not as uniform as in the case of relaxation into the S1min_C6 structure (see Table 4), results for all methods used show that both minima located on the S_1 surface are energetically accessible from the Franck–Condon region.

Minima on the crossing seam. Table 4 lists the CASSCF and MR-CISD energies and the character of five different minima located on the S_0/S_1 crossing seam optimized at the CASSCF level; Fig. 9b shows the corresponding structures. All structures display strong out-of-plane deformations with the twist around either the CN or CC bonds. We adopted the Cremer–Pople classification scheme in performing the conformational analysis of these structures.^{102,103} Using this approach, we identified four types of structure deformations, in particular screw-boat (S), boat (B), half-chair (H) and envelope (E) conformations. The 3H_4 , 1S_2 , 1S_6 and B_{14} MXSs are energetically grouped closely together at the CASSCF level within 0.4 eV; all MXSs shown are energetically accessible starting from the Franck–Condon region. The structures and the energetics of all MXS points are very similar to those found by Barbatti *et al.*^{29,30} for 4-aminopyrimidine although the results of conformational analysis are not the same (see Table 4). This situation indicates that there is only a small (if any) energetic effect of the additional amino group on the puckering of the pyrimidine ring and subsequent formation of conical intersection.

In all cases there are two major configurations which contribute to the CASSCF wavefunction ($\pi\pi^*$ and $n\pi^*$) (see Table 4). Although it is difficult to distinguish lone-pair and π orbitals in such distorted structures, the main character of the singly occupied orbitals is indicated in the Table 4. The second configuration contributing to the wavefunction in all cases is the closed-shell configuration (CS). In agreement with the results found for 4-aminopyrimidine, the character of the singly occupied orbital was identified as π orbital.

We performed MR-CISD calculations with 8 and 17 orbitals frozen in the CI procedure. The MR-CI and CASPT2 procedures resulted in relatively large splittings, usually around 0.8 eV, of the S_0 and S_1 surfaces at the MXS points determined at the CASSCF level. We estimated the energies and MXS structures at the MR-CISD level from an extrapolation of the LIIC curves (these results are obtained from the MR-CISD(17)+Q calculations, see below). Table 4 lists the CASSCF and these estimated MR-CISD(17)+Q results. When the dynamic correlation is included, there is a change in the ordering of various MXS structures and the 1S_2 structure becomes the most stable one. However, all structures are placed in a relatively small energy range, with the difference between the most and least stable structure 0.8 eV, in good agreement with the results of 4-aminopyrimidine.

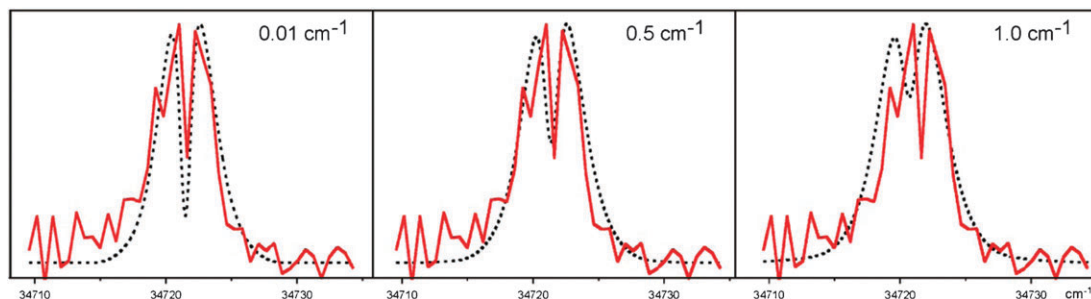


Fig. 7 Detail of the 34720 cm^{-1} peak in the 2,4-DAPy REMPI spectrum, compared with simulations of the rotational envelope at three different Lorentzian linewidths.

Table 4 Energies (in eV) of stationary points and conical intersections for 2,4-diaminopyrimidine relative to the ground state minimum structure calculated at the CASSCF(14,10), MR-CISD(6,5)+Q and CASPT2 levels and comparison with 4-aminopyrimidine. The 6-31G** basis set was used. The experimental excitation energy (band maximum) was estimated to be 4.91 eV^a

		CASSCF	MR-CISD(17)+Q ^b	CASPT2 ^c	4AP ^d MR-QDPT2
2 ¹ A($\pi\pi^*$)	ΔE	5.189	5.420 (5.406)	5.024 (4.938)	4.79 (5.34)
	f^e	0.065		0.063	
3 ¹ A($n\pi^*$)	ΔE	5.528	5.044 (5.142)	5.204 (5.110)	4.71 (5.88)
	f^e	0.013		0.012	
S1min_C6	$\pi\pi^* + n\pi^*$	4.754	4.743 (4.708)	4.573 (4.528)	4.22 (4.93)
S1min_C2	$\pi\pi^* + n\pi^*$	5.075	4.632 (4.917)	4.831 (4.822)	4.33 (5.02)
MXS(³ H ₄)	CS + $\pi\pi^*$	4.594	4.2 ^f	4.3 (4.4) ^f	4.62 (4.60) ^g
MXS(¹ S ₂)	CS + $\pi\pi^*$	4.645	3.9 ^f	4.2 (4.2) ^f	4.36 (4.49) ^h
MXS(¹ S ₆)	CS + $\pi\pi^*$	4.759	4.5 ^f	4.5 (4.5) ^f	4.71 (4.79)
MXS(B _{1,4})	CS + $\pi\pi^*$	4.949	4.1 ^f	4.3 (4.3) ^f	(4.97)
MXS(⁶ E)	CS + $\pi\pi^*$	5.328	4.7 ^f		(5.42)

^a For estimation of the experimental band maximum see text. ^b The results with eight frozen orbitals are given in parentheses. ^c The results obtained with the 6-311G(2d,p) basis set are given in parentheses. ^d MR-QDPT2 energies (CASSCF energies are given in parentheses) calculated for the relevant structures of 4-aminopyrimidine. ^e Oscillator strength. ^f Estimated from an extrapolation of the LIIC curves. ^g Assigned as ³S₄ for 4-APy. ^h Assigned as E₂ for 4-APy.

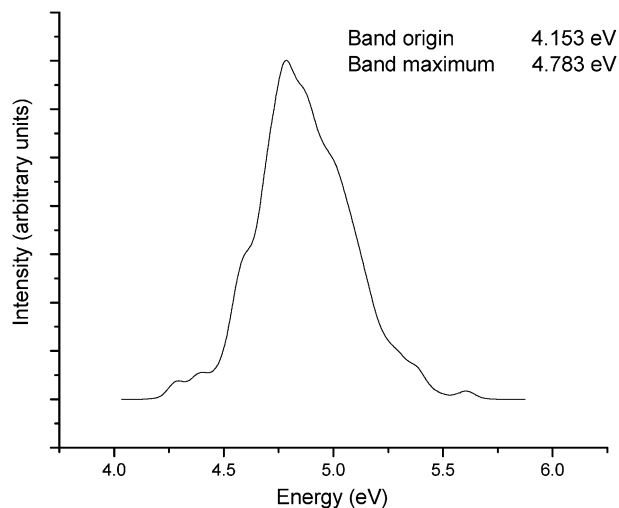


Fig. 8 Absorption spectrum of 2,4-DAPy calculated at the RICC2 method using the SVP basis set.

The CASPT2 energies, also estimated from the LIIC curves, agree reasonably well with the MR-CI results. Thus, the inclusion of the dynamic correlation effects does not change the parameters for prediction of the dynamics of the molecular systems studied.

Interpolation curves. We calculated LIIC curves for the reaction paths between the S1min_C2 and S1min_C6 minima and the four lowest MXS structures at the CASSCF and MR-CISD levels. The LIIC curve for the reaction path S1min_C6 to ¹S₆ was calculated using the CASPT2 method as well. (See Fig. 10 for the reaction path between S1min_C6 and ¹S₆. Other reaction paths are reported in the ESI).[†] To make the studies of reaction paths computationally more feasible we performed MR-CISD calculations with 17 frozen orbitals. A comparison of the character of the interpolation curves calculated at the MR-CISD(8)+Q and MR-CISD(17)+Q levels for the reaction path between S1min_C6 and ¹S₆ (see Fig. 10 and ESI)[†] shows

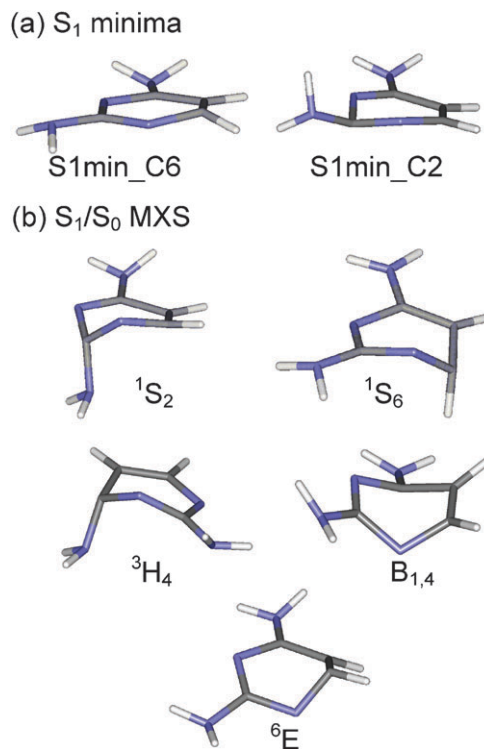


Fig. 9 (a) Structures of the two minima found on the S₁ potential energy surface optimized at the CASSCF(14,10)/6-31G** level. (b) Structures of the five different minima located on the S₀/S₁ crossing seam optimized at the CASSCF(14,10)/6-31G** level.

that the characters of the interpolation curves are very similar justifying the use of the latter approach. The barriers for the reaction paths estimated from the highest point of the interpolation curves are presented in Table 5. These values are similar to those found by Zechmann and Barbatti for 4-APy.¹⁰⁴ Since this estimation provides only an upper bound for the reaction barrier, we have optimized the transition states along the selected reaction paths, particularly the paths from

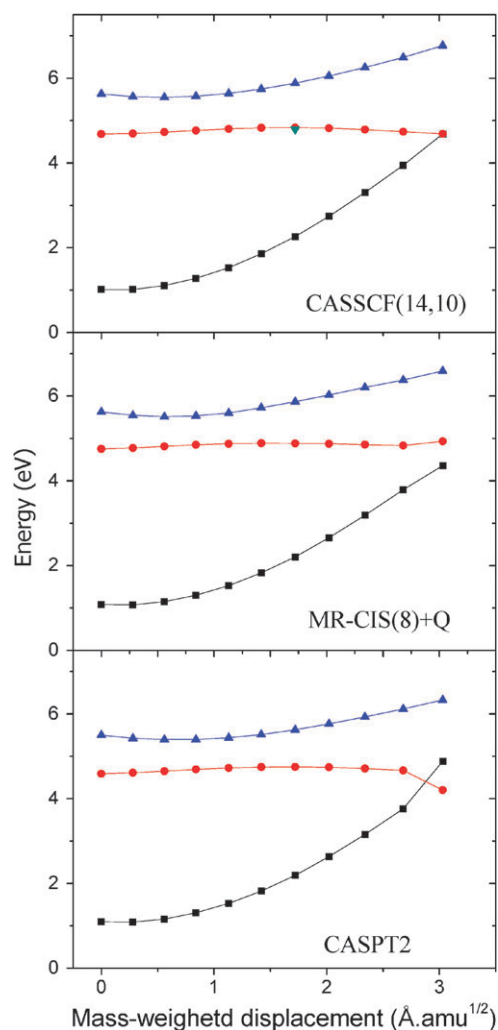


Fig. 10 Reaction path between S1min_C6 and 1S_6 calculated using CASSCF(14,10), MR-CIS(8)+Q, and CASPT2 methods. Other reaction paths are reported in the ESI.†

S1min_C6 towards 1S_6 and 3H_4 MXS structures. The energies of those structures are indicated in the graphs (see ESI)† and in Table 5.

Relatively small barriers exist for the reaction paths between the S1min_C6 and the 1S_6 and $B_{1,4}$ and 3H_4 MXSs. The only sizeable barrier we found was for the reaction paths toward the 1S_2 structure. This finding reflects some immobilization of the ring by substitution of the hydrogen atom by an amino group at the C2 atom. The barriers connecting the S1min_C2 with MXSs are generally larger with values of approximately

0.8 eV. Starting from the highest points of the interpolation curves we determined the true saddle points for the reaction paths towards the 1S_6 and 3H_4 structures. At the CASSCF and MR-CISD(17)+Q levels these calculations reduced the reaction barriers to 0.17 and 0.12 eV at for 3H_4 , and to 0.04 and 0.02 eV for 1S_6 , respectively, indicating that both conical intersections should be easily accessible leading to a subsequent ultrafast deactivation to the ground state.

Comparing the results for 2,4-DAPy and those previously reported for 4-APy shows that the additional amino-group on the pyrimidine ring does not significantly influence the energetics of the stationary points and conical intersections, as well as the characters of interpolation curves. The only exception is the 1S_2 structure and the interpolation curve towards it, as a result of an immobilization of the pyrimidine ring at the C2 position. Since 4-APy usually does not access the pathway with C2 atom deformation, one would not expect significant differences in the dynamics of these two systems, provided comparable initial conditions are used. Therefore, in analogy to the previously described surface hopping dynamics for 4-APy, the present reaction pathways for 2,4-DAPy indicate that ultrafast internal conversion to the ground state should be expected provided that sufficient initial kinetic energy is available to surpass the small energy barriers to the 1S_6 and 3H_4 conical intersections.

In the current experiments, the pump energy is in the range of 34 000 to 35 000 cm^{-1} which corresponds to 4.2–4.3 eV. Neglecting any vibrational corrections, this energy range is close to or even below the calculated energies of the saddle points (using the CASPT2 energies of MXSs and energy barriers of the relevant saddle points collected in Table 4). Therefore, the pump energy used in the experiment is not sufficient to overcome the barrier to reach the conical intersections, producing the much longer lifetime experimentally observed for 2,4-DAPy.

When the relevant coordinates involve ring deformations, lifetimes can be significantly different for specific structures. In particular, if deformation involving C5 and C6 leads to internal conversion in 2,4-DAPy then that pathway would be significantly altered for 2,6-DAPy, in which the puckering at C6 is strongly hindered. This observation explains why the lifetime for 9H-adenine is longer than that calculated for 4-APy and also explains the very long lifetime (6–8 ns) for 2,6-DAPy, in which both the C2 position and the C5=C6 positions are modified. In this latter case the subtle difference between the N7H and N9H tautomer lifetimes requires further analysis. This effect of immobilization of the C5=C6 twist is analogous to the findings of Zgierski *et al.*, which showed that

Table 5 Barriers (in eV) located on the interpolation curves

	CASSCF	CISD(17)+Q
S1min_C6 to 1S_6	0.145 Saddle point: 0.040	0.082 Saddle point: 0.024
S1min_C6 to $B_{1,4}$	0.419	0.153
S1min_C6 to 3H_4	0.407 Saddle point: 0.174	0.251 Saddle point: 0.118
S1min_C6 to 1S_2	1.041	1.042
S1min_C2 to 1S_2	0.614	0.976
S1min_C2 to $B_{1,4}$	0.686	0.974
S1min_C2 to 1S_6	0.788	0.830
S1min_C2 to 3H_4	0.870	0.857

in solution 5,6-trimethylenecytosine and 5,6-trimethylenuracil do not exhibit subpicosecond excited-state lifetimes characteristic of the naturally occurring pyrimidine bases.²¹

Conclusions

We recorded resonant two-photon ionization spectra of 2,4-diaminopyrimidine and 2,6-diaminopurine in the frequency range of 32 000 to 36 000 cm⁻¹. IR-UV double resonance experiments supported by DFT and *ab initio* calculations suggest that we observed the most stable tautomers of both compounds. We also recorded and assigned the photoelectron spectrum of 2,4-diaminopyrimidine, placing the adiabatic ionization potential at 7.86 ± 0.05 eV, and the first vertical ionization potential at 8.30 ± 0.04 eV. The S₁ ← S₀ (ππ*) origin of 2,4-diaminopyrimidine, as measured with one-color, two-photon resonant ionization, is 34 459 cm⁻¹. The excited state lifetime measured at this pump wavelength is shorter than the time resolution (approx. 1 ns) of the instrument and longer than 10 ps, based on the sharp and resolved peaks in the R2PI spectrum.

According to the IR-UV double resonance experiments, both the N7H and N9H tautomers of 2,6-diaminopurine are easily observable in the cold jet. The S₁ ← S₀(ππ*) origin is at 32 215 cm⁻¹ for the N7H tautomer and 34 881 cm⁻¹ for the N9H tautomer. The measured excited state lifetime is 8.7 ± 0.8 ns for the former and 6.3 ± 0.4 ns for the latter.

This long lifetime is consistent with reduction of ring deformations in both the C2 position and the C5=C6 twist, affecting all conical intersections that lead to fast excited state dynamics in 4-APy.

To understand the photodynamics of 2,4-diaminopyrimidine, we carried out quantum chemical modeling. Based on the results of the relative energies of stationary points on the S₁ surface and its crossings with the ground state at MXS points, together with the nature of the interpolation curves we can expect ultrafast dynamics in the excited state of 2,4-diaminopyrimidine which is predicted to be comparable to that of 4-aminopyrimidine. The additional NH₂ group in the C2 position blocks the path to only one of the conical intersections predicted for the analogous case of 4-aminopyrimidine, namely the one associated with ring deformation at C2. However, there are still other alternatives for almost barrierless access to conical intersections, associated with deformations at the C5, C6, and N1 positions. The actually observed lifetimes will depend on the chosen excitation energy which needs to be large enough to surmount the existing small energy barriers if ultrafast processes are to occur.

The present experiments show a significant reduction of the lifetime of 2,4-DAPy as compared to the nanosecond lifetimes determined for 2,6-DAPu, indicating a larger internal conversion rate for the former molecule. However, the current experiments involve a very narrow excitation window very close to the band origin, and therefore exhibit the longest possible lifetime. The pump energies in the range of 34 000 to 35 000 cm⁻¹ are close to or possibly just below the computed energy barriers, which explains the still rather large lifetime of 2,4-DAPy compared to that of other pyrimidine bases excited at the center of the absorption band.

Acknowledgements

This study is based upon work supported by National Science Foundation (CHE-0911564), and Hungarian National Science Fund (OTKA T60669, F61153). Zsolt Gengeliczki gratefully acknowledges the generous support of the Rosztoczy Foundation. This work was supported by grants from the Ministry of Education of the Czech Republic (Center for Biomolecules and Complex Molecular Systems, LC512). It was part of research project Z40550506 and by the Austrian Science Fund within the framework of the Special Research Program F16 (Advanced Light Sources) and Project P18411-N19. Support from the Preaemia Academiae, Academy of Sciences of the Czech Republic, awarded to PH in 2007, is gratefully acknowledged.

References

- 1 C. M. Marian, *J. Phys. Chem. A (USA)*, 2007, **111**, 1545–1553.
- 2 K. Seefeld, R. Brause, T. Haber and K. Kleinermanns, *J. Phys. Chem. A (USA)*, 2007, **111**, 6217–6221.
- 3 A. Abo-Riziq, L. Grace, E. Nir, M. Kabelac, P. Hobza and M. S. de Vries, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 20–23.
- 4 A. L. Sobolewski, W. Domcke and C. Hättig, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 17903–17906.
- 5 B. B. Brady, L. A. Peteanu and D. H. Levy, *Chem. Phys. Lett.*, 1988, **147**, 538–543.
- 6 D. C. Luhrs, J. Viallon and I. Fischer, *Phys. Chem. Chem. Phys.*, 2001, **3**, 1827–1831.
- 7 H. Kang, B. Jung and S. K. Kim, *J. Chem. Phys.*, 2003, **118**, 6717–6719.
- 8 C. Z. Bisgaard, H. Satzger, S. Ullrich and A. Stolow, *ChemPhysChem*, 2009, **10**, 101–110.
- 9 C. Canuel, M. Mons, F. Piuze, B. Tardivel, I. Dimicoli and M. Elhanine, *J. Chem. Phys.*, 2005, **122**, 7.
- 10 H. R. Hudock, B. G. Levine, A. L. Thompson, H. Satzger, D. Townsend, N. Gador, S. Ullrich, A. Stolow and T. J. Martinez, *J. Phys. Chem. A (USA)*, 2007, **111**, 8500–8508.
- 11 T. Climent, R. Gonzalez-Luque, M. Merchan and L. Serrano-Andres, *Chem. Phys. Lett.*, 2007, **441**, 327–331.
- 12 L. Serrano-Andres, M. Merchan and A. C. Borin, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 8691–8696.
- 13 S. Perun, A. L. Sobolewski and W. Domcke, *J. Am. Chem. Soc.*, 2005, **127**, 6257–6265.
- 14 H. Chen and S. H. Li, *J. Phys. Chem. A (USA)*, 2005, **109**, 8443–8446.
- 15 C. M. Marian, *J. Chem. Phys.*, 2005, **122**, 10.
- 16 N. Ismail, L. Blancafort, M. Olivucci, B. Kohler and M. A. Robb, *J. Am. Chem. Soc.*, 2002, **124**, 6818–6819.
- 17 L. Blancafort, B. Cohen, P. M. Hare, B. Kohler and M. A. Robb, *J. Phys. Chem. A (USA)*, 2005, **109**, 4431–4436.
- 18 L. Blancafort and M. A. Robb, *J. Phys. Chem. A (USA)*, 2004, **108**, 10609–10614.
- 19 L. Blancafort, *J. Am. Chem. Soc.*, 2006, **128**, 210–219.
- 20 S. Matsika, *J. Phys. Chem. A (USA)*, 2004, **108**, 7584–7590.
- 21 M. Z. Zgierski, S. Patchkovskii, T. Fujiwara and E. C. Lim, *J. Phys. Chem. A (USA)*, 2005, **109**, 9384–9387.
- 22 K. A. Kistler and S. Matsika, *J. Chem. Phys.*, 2008, **128**, 215102.
- 23 K. A. Kistler and S. Matsika, *J. Phys. Chem. A (USA)*, 2007, **111**, 2650–2661.
- 24 M. Barbatti and H. Lischka, *J. Am. Chem. Soc.*, 2008, **130**, 6831–6839.
- 25 A. Broo, *J. Phys. Chem.*, 1998, **A102**, 526–531.
- 26 E. Nir, K. Kleinermanns, L. Grace and M. S. de Vries, *J. Phys. Chem. A (USA)*, 2001, **105**, 5106–5110.
- 27 S. Perun, A. L. Sobolewski and W. Domcke, *Mol. Phys.*, 2006, **104**, 1113–1121.
- 28 E. Fabiano and W. Thiel, *J. Phys. Chem. A (USA)*, 2008, **112**, 6859–6863.
- 29 M. Barbatti and H. Lischka, *J. Phys. Chem. A (USA)*, 2007, **111**, 2852–2858.

- 30 M. Barbatti, M. Ruckebauer, J. J. Szymczak, A. J. A. Aquino and H. Lischka, *Phys. Chem. Chem. Phys.*, 2008, **10**, 482–494.
- 31 M. Barbatti, B. Sellner, A. J. A. Aquino and A. Lischka, *Nonadiabatic Excited-State Dynamics of Aromatic Heterocycles: Toward the Time-Resolved Simulation of Nucleobasis*, Springer Science, Business Media B.V., 2008.
- 32 G. F. Joyce, A. W. Schwartz, S. L. Miller and L. E. Orgel, *Proc. Natl. Acad. Sci. U. S. A.*, 1987, **84**, 4398–4402.
- 33 M. J. Lutz, J. Horlacher and S. A. Benner, *Bioorg. Med. Chem. Lett.*, 1998, **8**, 1149–1152.
- 34 M. J. Lutz, H. A. Held, M. Hottiger, U. Hubscher and S. A. Benner, *Nucleic Acids Res.*, 1996, **24**, 1308–1313.
- 35 J. A. Piccirilli, T. Krauch, S. E. Moroney and S. A. Benner, *Nature*, 1990, **343**, 33–37.
- 36 S. Yamazaki, A. L. Sobolewski and W. Domcke, *Phys. Chem. Chem. Phys.*, 2009, **11**, 10165–10174.
- 37 H. J. Cleaves, K. E. Nelson and S. L. Miller, *Naturwissenschaften*, 2006, **93**, 228–231.
- 38 J. P. Ferris, O. S. Zamek, A. M. Altbuch and H. Freiman, *J. Mol. Evol.*, 1974, **3**, 301–309.
- 39 S. Miyakawa, H. J. Cleaves and S. L. Miller, *Origins Life Evol. Biosphere*, 2002, **32**, 209–218.
- 40 M. P. Robertson, M. Levy and S. L. Miller, *J. Mol. Evol.*, 1996, **43**, 543.
- 41 R. Saladino, C. Crestini, V. Neri, J. R. Brucato, L. Colangeli, F. Ciciriello, E. Di Mauro and G. Costanzo, *ChemBioChem*, 2005, **6**, 1368–1374.
- 42 E. Nir, M. Muller, L. I. Grace and M. S. de Vries, *Chem. Phys. Lett.*, 2002, **355**, 59–64.
- 43 E. Nir, C. Janzen, P. Imhof, K. Kleinermanns and M. S. de Vries, *J. Chem. Phys.*, 2001, **115**, 4604–4611.
- 44 E. Nir, I. Hunig, K. Kleinermanns and M. S. de Vries, *Phys. Chem. Chem. Phys.*, 2003, **5**, 4780–4785.
- 45 K. A. Seefeld, C. Plutzer, D. Lowenich, T. Haber, R. Linder, K. Kleinermanns, J. Tatchen and C. M. Marian, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3021–3026.
- 46 M. P. Callahan, B. Crews, A. Abo-Riziq, L. Grace, M. S. de Vries, Z. Gengeliczki, T. M. Holmes and G. A. Hill, *Phys. Chem. Chem. Phys.*, 2007, **9**, 4587–4591.
- 47 R. Tembreull, C. H. Sin, H. M. Pang and D. M. Lubman, *Anal. Chem.*, 1985, **57**, 2911–2917.
- 48 G. Meijer, M. S. de Vries, H. E. Hunziker and H. R. Wendt, *Appl. Phys. B: Photophys. Laser Chem.*, 1990, **51**, 395–403.
- 49 B. Csakvari, A. Nagy, L. Zanathy and L. Szepes, *Magy. Kem. Foly.*, 1992, **98**, 415–419.
- 50 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 51 C. T. Lee, W. T. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter*, 1988, **37**, 785–789.
- 52 B. Miehlich, A. Savin, H. Stoll and H. Preuss, *Chem. Phys. Lett.*, 1989, **157**, 200–206.
- 53 A. D. Mclean and G. S. Chandler, *J. Chem. Phys.*, 1980, **72**, 5639–5648.
- 54 R. Krishnan, J. S. Binkley, R. Seeger and J. A. Pople, *J. Chem. Phys.*, 1980, **72**, 650–654.
- 55 C. Moller and M. S. Plesset, *Phys. Rev.*, 1934, **46**, 618–622.
- 56 J. A. Pople, M. Headgordon and K. Raghavachari, *J. Chem. Phys.*, 1989, **90**, 4635–4636.
- 57 M. J. Frisch, M. Headgordon and J. A. Pople, *Chem. Phys. Lett.*, 1990, **166**, 275–280.
- 58 M. J. Frisch, M. Headgordon and J. A. Pople, *Chem. Phys. Lett.*, 1990, **166**, 281–289.
- 59 M. Headgordon and T. Headgordon, *Chem. Phys. Lett.*, 1994, **220**, 122–128.
- 60 S. Saebo and J. Almlöf, *Chem. Phys. Lett.*, 1989, **154**, 83–89.
- 61 M. P. Andersson and P. Uvdal, *J. Phys. Chem. A (USA)*, 2005, **109**, 2937–2941.
- 62 A. P. Scott and L. Radom, *J. Phys. Chem.*, 1996, **100**, 16502–16513.
- 63 L. A. Curtiss, K. Raghavachari, P. C. Redfern, V. Rassolov and J. A. Pople, *J. Chem. Phys.*, 1998, **109**, 7764–7776.
- 64 R. E. Stratmann, G. E. Scuseria and M. J. Frisch, *J. Chem. Phys.*, 1998, **109**, 8218–8224.
- 65 R. Bauernschmitt and R. Ahlrichs, *Chem. Phys. Lett.*, 1996, **256**, 454–464.
- 66 M. E. Casida, C. Jamorski, K. C. Casida and D. R. Salahub, *J. Chem. Phys.*, 1998, **108**, 4439–4449.
- 67 J. V. Ortiz, *J. Chem. Phys.*, 1988, **89**, 6353–6356.
- 68 L. S. Cederbaum, *J. Phys. B: At. Mol. Phys.*, 1975, **8**, 290–303.
- 69 W. von Niessen, J. Schirmer and L. S. Cederbaum, *Comput. Phys. Rep.*, 1984, **1**, 57.
- 70 V. G. Zakrzewski and W. Vonniessen, *J. Comput. Chem.*, 1993, **14**, 13–18.
- 71 V. G. Zakrzewski and J. V. Ortiz, *Int. J. Quantum Chem.*, 1995, **53**, 583–590.
- 72 J. V. Ortiz, *Int. J. Quantum Chem.*, 1988, **22**, 431.
- 73 J. V. Ortiz, *Int. J. Quant. Chem. Symp.*, 1989, **23**, 321.
- 74 J. V. Ortiz, V. G. zakrzewski and O. Dolgounircheva, in *Conceptual Perspectives in Quantum Chemistry*, ed. J. L. Calais and E. S. Kryachko, Kluwer Academic, 1997, p. 465.
- 75 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, J. Montgomery, J. A., T. Vreven, K. N. Kudin, J. C. Burant, J. M. Millam, S. S. Iyengar, J. Tomasi, V. Barone, B. Mennucci, M. Cossi, G. Scalmani, N. Rega, G. A. Petersson, H. Nakatsuji, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, M. Klene, X. Li, J. E. Knox, H. P. Hratchian, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, P. Y. Ayala, K. Morokuma, G. A. Voth, P. Salvador, J. J. Dannenberg, V. G. Zakrzewski, S. Dapprich, A. D. Daniels, M. C. Strain, O. Farkas, D. K. Malick, A. D. Rabuck, K. Raghavachari, J. B. Foresman, J. V. Ortiz, Q. Cui, A. G. Baboul, S. Clifford, J. Cioslowski, B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi, R. L. Martin, D. J. Fox, T. Keith, M. A. Al-Laham, C. Y. Peng, A. Nanayakkara, M. Challacombe, P. M. W. Gill, B. Johnson, W. Chen, M. W. Wong, C. Gonzalez and J. A. Pople, Gaussian, Inc., Wallingford, CT, 2004.
- 76 A. Bunge, *J. Chem. Phys.*, 1970, **53**, 20.
- 77 J. A. Pople, R. Seeger and R. Krishnan, *Int. J. Quantum Chem.*, 1977, 149–163.
- 78 W. J. Hehre, R. Ditchfie and J. A. Pople, *J. Chem. Phys.*, 1972, **56**, 2257.
- 79 J. S. Binkley, J. A. Pople and W. J. Hehre, *J. Am. Chem. Soc.*, 1980, **102**, 939–947.
- 80 K. Andersson, P. A. Malmqvist and B. O. Roos, *J. Chem. Phys.*, 1992, **96**, 1218–1226.
- 81 K. Andersson, P. A. Malmqvist, B. O. Roos, A. J. Sadlej and K. Wolinski, *J. Phys. Chem.*, 1990, **94**, 5483–5488.
- 82 A. Kohn and C. Hattig, *J. Chem. Phys.*, 2003, **119**, 5021–5036.
- 83 C. Hattig, *J. Chem. Phys.*, 2003, **118**, 7751–7761.
- 84 M. Barbatti, G. Granucci, M. Persico, M. Ruckebauer, M. Vazdar, M. Eckert-Maksic and H. Lischka, *J. Photochem. Photobiol., A*, 2007, **190**, 228–240.
- 85 M. Dallos, H. Lischka, R. Shepard, D. R. Yarkony and P. G. Szalay, *J. Chem. Phys.*, 2004, **120**, 7330–7339.
- 86 H. Lischka, M. Dallos, P. G. Szalay, D. R. Yarkony and R. Shepard, *J. Chem. Phys.*, 2004, **120**, 7322–7329.
- 87 H. Lischka, M. Dallos and R. Shepard, *Mol. Phys.*, 2002, **100**, 1647–1658.
- 88 R. Shepard, H. Lischka, P. G. Szalay, T. Kovar and M. Ernzerhof, *J. Chem. Phys.*, 1992, **96**, 2085–2098.
- 89 R. Shepard, in *Modern Electronic Structure Theory*, ed. D. R. Yarkony, World Scientific, Singapore, 1995, vol. 1, p. 345.
- 90 H. Lischka, R. Shepard, R. M. Pitzer, I. Shavitt, M. Dallos, T. Muller, P. G. Szalay, M. Seth, G. S. Kedziora, S. Yabushita and Z. Y. Zhang, *Phys. Chem. Chem. Phys.*, 2001, **3**, 664–673.
- 91 H. Lischka, R. Shepard, F. B. Brown and I. Shavitt, *Int. J. Quantum Chem.*, 1981, 91–100.
- 92 H. Lischka, R. Shepard, I. Shavitt, R. M. Pitzer, M. Dallos, T. Muller, P. G. Szalay, F. B. Brown, R. Ahlrichs, H. J. Boehm, A. Chang, D. C. Comeau, R. Gdanitz, H. Dachsels, C. Ehrhardt, M. Ernzerhof, P. Hoescht, S. Irlle, G. Kedziora, T. Kovar, V. Parasuk, M. J. M. Pepper, P. Sharf, H. Shiffer, M. Schindler, M. Schuler, M. Seth, E. A. Stahlberg, J.-G. Zhao, S. Yabushita, Z. Zhang, M. Barbatti, S. Matsika, M. Schuurmann, D. R. Yarkony, S. R. Brozell, E. V. Beck and

-
- J.-P. Blaudeau, *Columbus, an ab initio electronic structure program* release 5.9.1, www.univie.ac.at/columbus, 2006.
- 93 G. Karlstrom, R. Lindh, P. A. Malmqvist, B. O. Roos, U. Ryde, V. Veryazov, P. O. Widmark, M. Cossi, B. Schimmelpfennig, P. Neogrady and L. Seijo, *Comput. Mater. Sci.*, 2003, **28**, 222–239.
- 94 P. A. Malmqvist, A. Rendell and B. O. Roos, *J. Phys. Chem.*, 1990, **94**, 5477–5482.
- 95 B. O. Roos and P. R. Taylor, *Chem. Phys.*, 1980, **48**, 157–173.
- 96 R. Ahlrichs, M. Bar, M. Haser, H. Horn and C. Kolmel, *Chem. Phys. Lett.*, 1989, **162**, 165–169.
- 97 M. Barbatti, G. Granucci, M. Ruckebauer, J. Pittner, M. Persico and H. Lischka, *NEWTON-X: a package for Newtonian dynamics close to the crossing seam*, www.univie.ac.at/newtonx, 2007.
- 98 I. P. Csonka, U. Szepes and A. Modelli, *J. Mass Spectrom.*, 2004, **39**, 1456–1466.
- 99 H. Kang, K. T. Lee, B. Jung, Y. J. Ko and S. K. Kim, *J. Am. Chem. Soc.*, 2002, **124**, 12958–12959.
- 100 Y. G. He, C. Y. Wu and W. Kong, *J. Phys. Chem. A (USA)*, 2003, **107**, 5145–5148.
- 101 Y. G. He, C. Y. Wu and W. Kong, *J. Phys. Chem. A (USA)*, 2004, **108**, 943–949.
- 102 D. Cremer, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1984, **40**, 498–500.
- 103 D. Cremer and J. A. Pople, *J. Am. Chem. Soc.*, 1975, **97**, 1354–1358.
- 104 G. Zechmann and M. Barbatti, *Int. J. Quantum Chem.*, 2008, **108**, 1266–1276.