

On the electronic structure of methyl butyrate and methyl valerate^{*,**}

Małgorzata A. Śmialek^{1,a}, Denis Dufflot², Nykola C. Jones³, Søren Vrønning Hoffmann³, Lucia Zuin⁴, Michael Macdonald⁴, Nigel J. Mason⁵, and Paulo Limão-Vieira⁶

¹ Department of Control and Power Engineering, Faculty of Ocean Engineering and Ship Technology, Gdańsk University of Technology, Gabriela Narutowicza 11/12, Gdańsk 80-233, Poland

² Univ. Lille, CNRS, UMR 8523 – PhLAM – Physique des Lasers Atomes et Molcules, Lille 59000, France

³ ISA, Department of Physics and Astronomy, Aarhus University, Ny Munkegade, Building 1520, Aarhus C 8000, Denmark

⁴ Canadian Light Source Inc., 44 Innovation Boulevard, Saskatoon S7N 2V3, Canada

⁵ School of Physical Sciences, Ingram Building, University of Kent, Canterbury CT2 7NH, UK

⁶ Atomic and Molecular Collisions Laboratory, CEFITEC, Department of Physics, Universidade NOVA de Lisboa, Caparica 2829-516, Portugal

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Abstract. We present novel results of the analysis of the electronic structure of two aliphatic esters: methyl butyrate and methyl valerate. High-resolution photoabsorption spectra were collected and analyzed over the energy range 4.0–10.8 eV and showed for both the molecules not only a clear band of the HOMO to LUMO transition, but also vibronic structure associated with the first Rydberg-valence transition. Photoelectron spectra recorded from 9 to over 28 eV revealed many ionization states with the first adiabatic ionization energies found to be 9.977 eV and 9.959 eV for methyl butyrate and methyl valerate, respectively. Ab initio calculations have been performed in order to help assign the photoabsorption and photoelectron features. Photolysis life times in the atmosphere were calculated revealing that photolysis is not competitive over hydroxyl radical scavenging in the process of removal of these esters from the atmosphere.

1 Introduction

The knowledge of the electronic structure and properties of esters is of common interest in many disparate areas and therefore has been studied over many decades, however the data still needs to be updated either because they are outdated, were recorded with poor resolution, or are simply absent. Esters occur both naturally, being secreted from plants, and as a by-product of industrial processes. These compounds are widely used in the food and flavoring industry due to their distinct aroma [1,2], but they also are of potential interest to astrochemists, who find the smallest members of this group in the interstellar medium, which is of interest for their possible role in the origins of life [3]. Both molecules of interest can serve as insecticides [4]. Methyl valerate in its pure form is used as plasticizer [5] and shows potential for use in the production of biofuels [6]. Due to their low vapour pressure that makes them easy to handle, both esters are produced on a massive scale and thus are likely to be released into

the atmosphere. Since it was shown that the main products of photolysis of such esters are carbon monoxide and dioxide, it is crucial to gain an insight into their structure, properties and photoabsorption cross sections [7].

Previously we have reported our findings on methyl formate [8], ethyl formate [9], isobutyl formate [10], ethyl acetate [11], isobutyl acetate [12] and some acetates and propionates [13]. Here we present our findings on a further two methyl esters: butyrate and valerate. The high-resolution VUV photoabsorption and photoelectron spectra presented here were measured for the first time for these compounds. The measured photoabsorption cross-sections have been used to calculate the photolysis lifetimes of these compounds in the upper atmosphere (20–50 km) of the Earth. The experimental findings are supported by theoretical calculations that allowed the determination of the ionization energies in the photoelectron spectra of both compounds and revealed predominantly mixed valence-Rydberg character of the transitions resolved in the photoabsorption spectra.

2 Experimental

2.1 The samples

The liquid samples used in both the VUV photoabsorption measurements (PA) and the photoelectron spectroscopy

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^a e-mail: smialek@pg.edu.pl

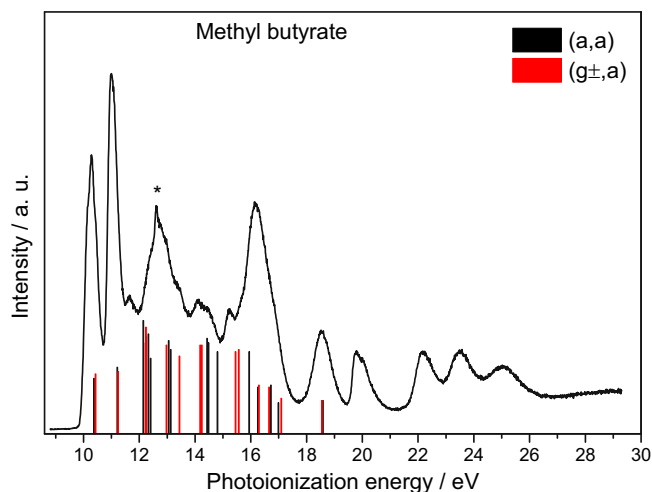


Fig. 1. Photoelectron spectrum of methyl butyrate, $C_5H_{10}O_2$, in the 9.0–29.0 eV region, collected at 80 eV initial photon energy; * denotes 2B_1 H_2O contamination; the stick spectrum represents the computed ionization energies at the M06-2X/aug-cc-pVTZ geometry with the P3+ method for the most abundant conformers: (a, a), black, and (g±, a), red.

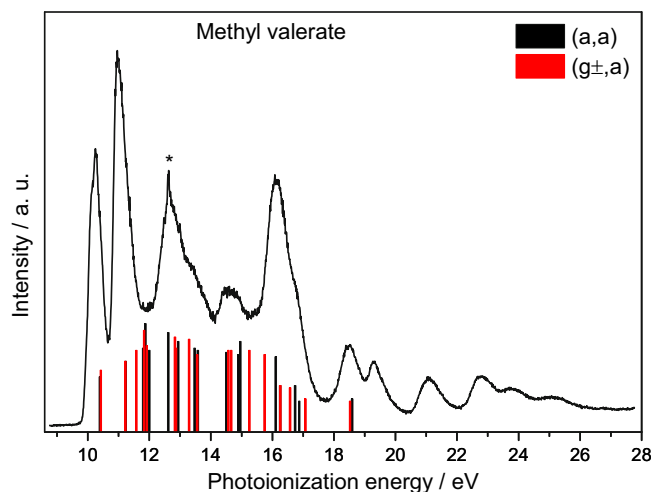


Fig. 2. Photoelectron spectrum of methyl valerate, $C_6H_{12}O_2$, in the 9.0–28.0 eV region, collected at 80 eV initial photon energy; * denotes 2B_1 H_2O contamination; the stick spectrum represents the computed ionization energies at the M06-2X/aug-cc-pVTZ geometry with the P3+ method for the most abundant conformers: (a, a), black, and (g±, a), red.

(PES) experiments were purchased from Sigma-Aldrich with a stated purity of 99%. The samples were degassed by repeated freeze-pump-thaw cycles with no further purification.

2.2 Photoelectron measurements

The photoelectron spectra of methyl butyrate, Figure 1, and methyl valerate, Figure 2, were measured at the VLS-PGM beamline [14] at the Canadian Light Source facility in Saskatoon, Canada, using a Double Toroidal Coincidence Spectrometer, that was designed and used for analysis of noble gases and small, diatomic molecules [15], but it has recently been demonstrated that it may also be used for measurements of more complex systems [16]. In our experiments we have used the data collected by the 180° toroidal detector, set at a pass energy of 4 eV. The measured resolution from the nitrogen calibration spectra was 60 meV. The photoexcitation energy of the photoelectron spectrum presented here was 80 eV, recorded with the entrance and exit slits of the VLS-PGM beamline set at $50 \mu\text{m}$.

Spectra presented here were calibrated against the $X^2\Sigma_g^+$, $\nu' = 0$ and $A^2\Pi_u$, $\nu' = 0$ peaks of N_2^+ , rounded to three decimal places [17,18].

The stars marked on the spectra in the figures denote the 2B_1 ionic state from some water contamination could also be used to check the accuracy of the calibration of the energy scale.

2.3 Photoabsorption measurements

The high-resolution VUV photoabsorption spectra of methyl butyrate, Figure 3, and methyl valerate, Figure 4, were measured at the UV1 beam line of the ASTRID synchrotron light source at Aarhus University, Denmark. The

experimental apparatus has been described in detail previously [19].

The sample pressure is measured using a capacitance manometer (Baratron). To ensure that the data were free of any saturation effects [20,21], the cross sections were measured over the pressure range 0.07–1.27 mbar with typical attenuations below 50%. A background scan is recorded with the cell evacuated. Absolute photoabsorption cross sections are then obtained using the Beer–Lambert attenuation law

$$I_t = I_0 \times \exp(-n\sigma x), \quad (1)$$

where I_t is the radiation intensity transmitted through the gas sample, I_0 is that through the evacuated cell, n is the molecular number density of the sample gas, σ is the absolute photoabsorption cross section, and x is the absorption path length (15.5 cm) [22]. A small amount of water contamination (<15%) was observed in the spectra recorded for these samples. The water contribution was subtracted in the present spectra. Due to this, the accuracy of the cross-section is estimated to be around $\pm 15\%$.

3 Computational methods

All the calculations have been performed using the Gaussian 16 rev B.01 code [23]. The geometry of the conformers was optimised at the MP2 level, as well as with density functional theory (DFT), using the ultra-fine grid size. As discussed in detail in the Supplementary Information (SI), several methods were tested on the relative energies of the conformers and compared with previous works [24,25]. The M06-2X [26] was thus employed. The optimized geometry of the conformers was obtained using the tight criteria and unscaled zero-point vibrational energies were calculated within the

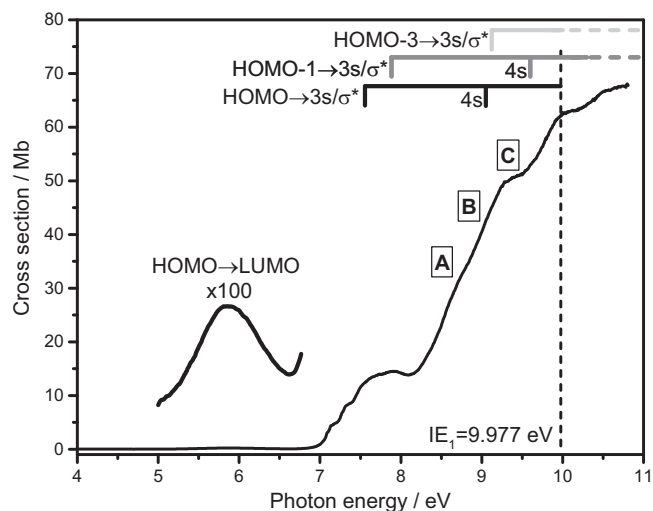


Fig. 3. Photoabsorption spectrum of methyl butyrate, $C_5H_{10}O_2$, in the 4.0–10.8 eV region; close-up shows the first valence HOMO → LUMO transition. See text for further details.

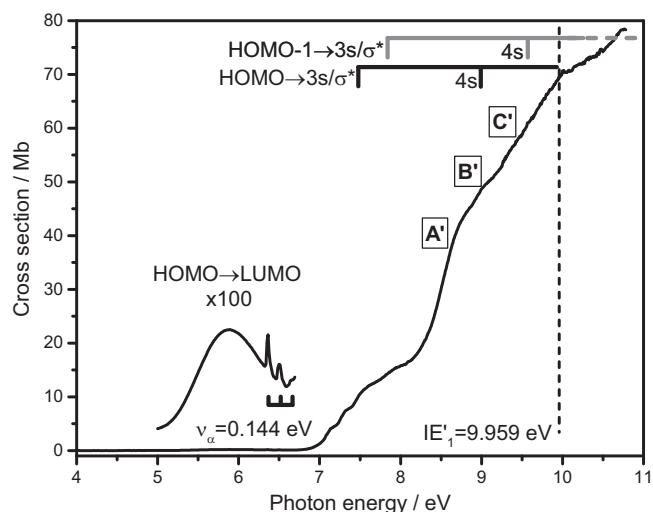


Fig. 4. Photoabsorption spectrum of methyl valerate, $C_6H_{12}O_2$, in the 4.0–10.8 eV region; close-up shows the first valence HOMO → LUMO transition. See text for further details.

harmonic approximation. The basis set used for these calculations is Dunning's aug-cc-pVTZ [27,28]. For the calculation of higher Rydberg states, this basis set was supplemented with a set of $(8s8p3d)$ diffuse functions taken from Kaufmann et al. [22,29] centred on the carbon atom close to the middle of the molecule. The oscillator strengths were evaluated with the length gauge. The assignment of the transitions was performed by visual inspection of the Natural Transition Orbitals (NTO's) [30] using the Chemcraft software [31]. Finally, the ionization energies were obtained at the OVGF, P3 and renormalized P3+ levels (see [32,33] for recent reviews). Detailed

information on computational methods and structure on both molecules can be found in the SI.

4 Structure of methyl butyrate and methyl valerate

According to previous results, there are eight possible conformers for methyl valerate [24] and four for methyl butyrate [25]. Nonetheless, the analysis of the rovibrational spectroscopic data shows that for both of these esters, only the (a, a) , with C_S symmetry, and $(g\pm, a)$, with no symmetry, conformers are present in almost the same quantities. Therefore, only these two conformers were investigated.

In Figures 5 and 6 the structures together with the highest occupied molecular orbital, HOMO, HOMO-1, HOMO-2 localization on both the conformers and LUMO for the (a, a) one, are shown for methyl butyrate and methyl valerate. The bond lengths can be found in the SI. The orbitals in square brackets denote the core ones, the remaining are valence orbitals. For the (a, a) conformer of methyl butyrate the electron configuration is $[1a'^2 2a'^2 3a'^2 4a'^2 5a'^2 6a'^2 7a'^2] 8a'^2 9a'^2 10a'^2 11a'^2 12a'^2 13a'^2 14a'^2 15a'^2 1a''^2 16a''^2 2a''^2 17a''^2 18a''^2 3a''^2 19a''^2 20a''^2 21a''^2 4a''^2 5a''^2 22a''^2$, whereas for $(g\pm, a)$ one the electronic configuration is simply $[1a^2 2a^2 3a^2 4a^2 5a^2 6a^2 7a^2] 8a^2 9a^2 10a^2 11a^2 12a^2 13a^2 14a^2 15a^2 16a^2 17a^2 18a^2 19a^2 20a^2 21a^2 22a^2 23a^2 24a^2 25a^2 26a^2 27a^2 28a^2$.

Similarly, for methyl valerate the configuration of the (a, a) conformer is $[1a'^2 2a'^2 3a'^2 4a'^2 5a'^2 6a'^2 7a'^2] 8a'^2 9a'^2 10a'^2 11a'^2 12a'^2 13a'^2 14a'^2 15a'^2 16a'^2 17a'^2 1a''^2 18a''^2 2a''^2 19a''^2 3a''^2 20a''^2 21a''^2 4a''^2 22a''^2 23a''^2 5a''^2 24a''^2 6a''^2 7a''^2 25a''^2$ and for the $(g\pm, a)$ one: $[1a^2 2a^2 3a^2 4a^2 5a^2 6a^2 7a^2] 8a^2 9a^2 10a^2 11a^2 12a^2 13a^2 14a^2 15a^2 16a^2 17a^2 18a^2 19a^2 20a^2 21a^2 22a^2 23a^2 24a^2 25a^2 26a^2 27a^2 28a^2 29a^2 30a^2$.

In all cases the HOMO, as in previously analyzed ester molecules, in the neutral ground state is localized predominantly on the terminal oxygen in-plane lone pair. In the case of methyl butyrate the HOMO is $(22a')^2$ for the (a, a) conformer and $(28a)^2$ for the $(g\pm, a)$ one, whereas for methyl valerate the HOMO is $(25a')^2$ for the (a, a) conformer and $(30a)^2$ for $(g\pm, a)$.

Since the distinct presence of both conformers contribute equally to the spectra, in this work we will refer to the states as the HOMO, HOMO-1, etc., meaning e.g. for HOMO of methyl butyrate both $(22a')^2$ for (a, a) conformer and $(28a)^2$ for the $(g\pm, a)$ one.

5 Results and discussion

5.1 Photoelectron spectra

The photoelectron spectra of methyl butyrate and methyl valerate are shown in Figures 1 and 2. Based on the calculated values for both conformers, which are shown for methyl butyrate in Table 1 and methyl valerate in Table 2, it was possible to identify most of the ionization energies of

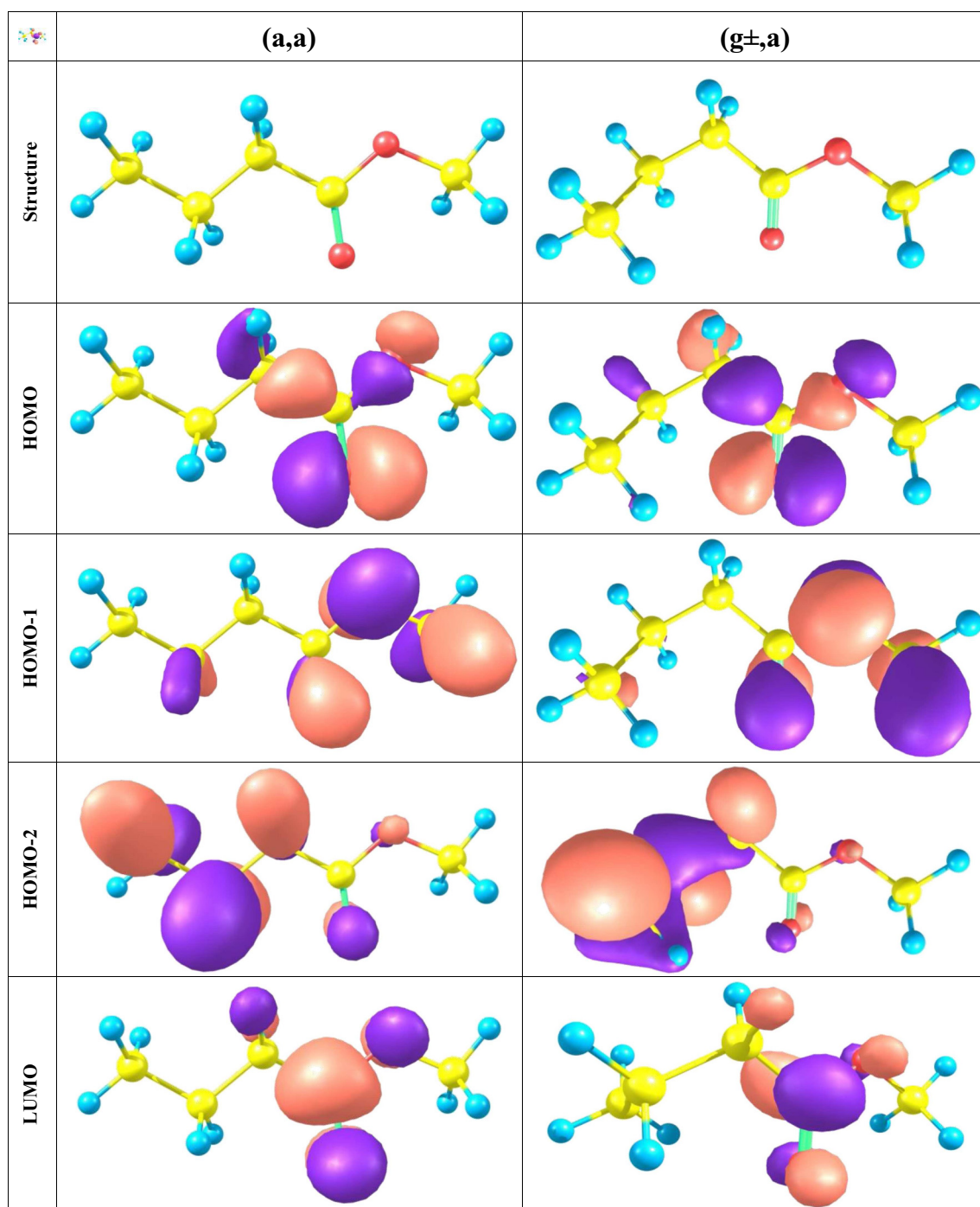


Fig. 5. The structure and HOMO, HOMO-1, HOMO-2, LUMO localization on both the (a, a) and $(g\pm, a)$ conformers of methyl butyrate, $C_5H_{10}O_2$.

both molecules. The values obtained based on the experimental spectra are also shown in the tables for comparison with the results of the calculations. The best accuracy was obtained for energies obtained from calculations using the P3+ method.

In the case of methyl butyrate it was not possible to resolve unambiguously the ionization energies of the third, fourth and fifth orbitals, since they appeared too

close together. For methyl valerate it was the position of additional 2 orbitals for the (a, a) conformer and one for $(g\pm, a)$ that were impossible to determine from the experimental spectrum.

The analysis of the first ionic state of both esters allowed not only the determination of the value of the first adiabatic and vertical ionization energies of the compounds, but also revealed a vibronic progression, similar to



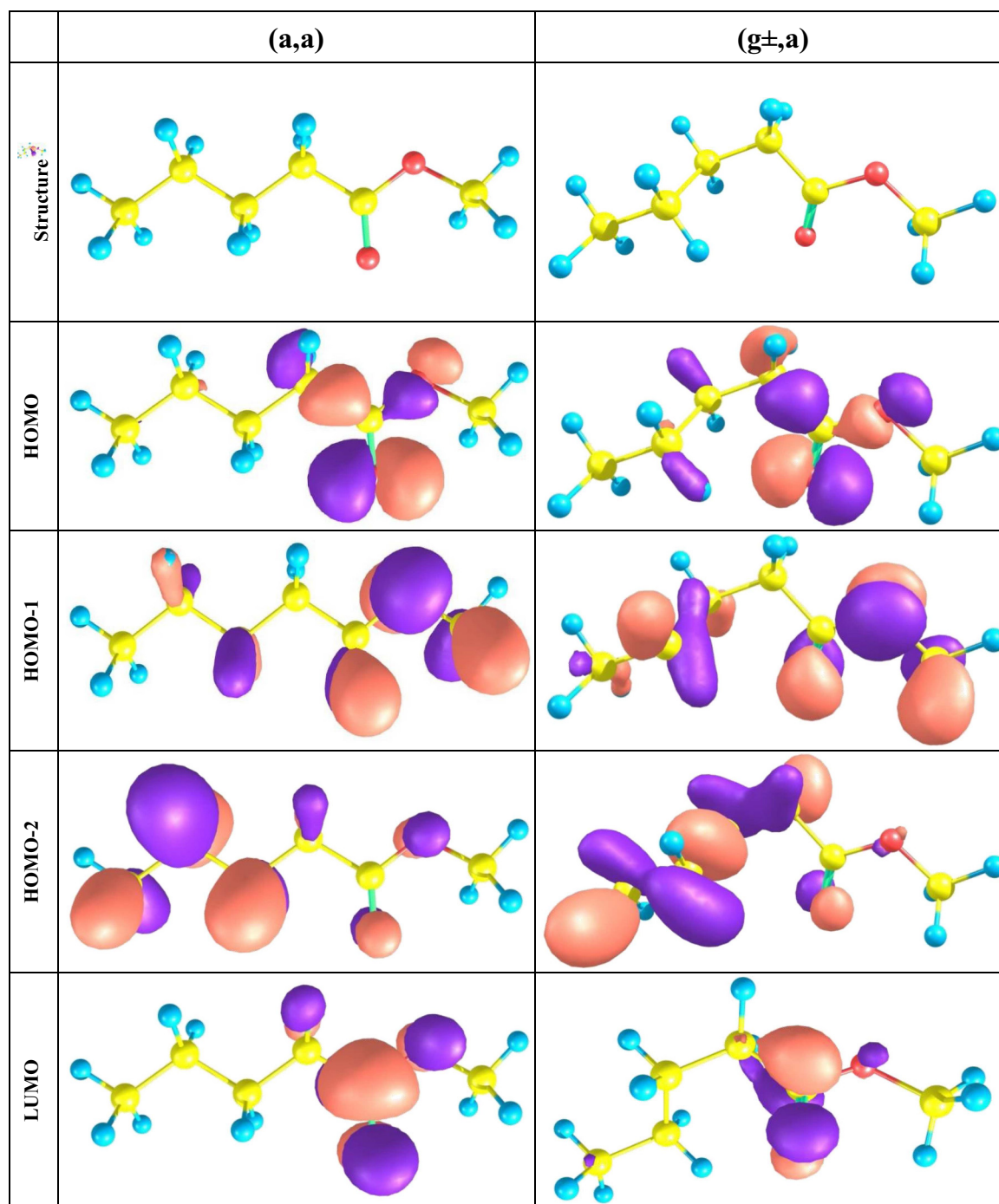


Fig. 6. The structure and HOMO, HOMO-1, HOMO-2, LUMO localization on both the (a, a) and $(g\pm, a)$ conformers of methyl valerate, $C_6H_{12}O_2$.

one seen for previous esters. This vibrational structure is depicted in Figure 7, top for methyl butyrate and bottom for methyl valerate. The obtained values are summarized in Table 3. With an average spacing of $\nu'_A = 0.153$ eV for methyl butyrate and $\nu'_{A'} = 0.151$ eV for methyl valerate, the vibration corresponds to a combination of C–O stretch combined with C=O stretch, as found previously for other esters.

5.2 Valence states, transitions and Rydberg series in photoabsorption spectra

According to the calculations presented in Tables 4 and 5, the first absorption bands that are centered at 5.877 eV for methyl butyrate and 5.883 eV for methyl valerate (Figs. 3 and 4) correspond to the first HOMO \rightarrow LUMO valence transition.



Table 1. Calculated vertical ionization energies and pseudo-intensities (pole strengths, PS) of methyl butyrate for (a, a) and $(g\pm, a)$ symmetries at the M06-2X/aug-cc-pVTZ geometry calculated with propagator methods, compared with experimental values, all in eV.

Configuration	Calculated/eV						Experimental/eV	
	Koopmans	OVGF	PS	P3	PS	P3+		
(a, a)								
$22a'^{-1}$	12.115	10.744	0.895	10.595	0.891	10.387	0.885	10.281
$5a''^{-1}$	12.541	11.460	0.899	11.359	0.894	11.224	0.890	11.002
$4a''^{-1}$	13.045	12.285	0.915	12.200	0.912	12.160	0.911	–
$21a'^{-1}$	13.309	12.379	0.911	12.402	0.907	12.338	0.905	–
$20a'^{-1}$	13.670	12.491	0.902	12.543	0.897	12.420	0.894	–
$19a'^{-1}$	14.409	13.334	0.909	13.193	0.906	13.071	0.902	13.166
$4a''^{-1}$	14.155	13.274	0.905	13.202	0.900	13.138	0.898	13.166
$3a''^{-1}$	15.434	14.571	0.909	14.488	0.904	14.446	0.903	14.471
$18a'^{-1}$	15.662	14.624	0.907	14.556	0.902	14.495	0.901	14.471
$17a'^{-1}$	16.186	15.025	0.905	14.933	0.900	14.815	0.897	14.708
$2a''^{-1}$	17.334	16.026	0.907	16.014	0.899	15.953	0.897	15.692
$16a'^{-1}$	18.321	16.518	0.893	16.510	0.890	16.268	0.881	16.202
$1a''^{-1}$	18.564	16.879	0.893	16.882	0.887	16.735	0.882	16.884
$15a'^{-1}$	19.088	17.170	0.886	17.194	0.881	17.006	0.874	17.325
$14a'^{-1}$	20.873	18.857	0.886	18.801	0.882	18.603	0.875	18.551
$(g\pm, a)$								
$28a'^{-1}$	12.007	10.750	0.897	10.612	0.893	10.414	0.887	10.281
$27a'^{-1}$	12.557	11.470	0.898	11.364	0.893	11.222	0.888	11.002
$25a'^{-1}$	13.203	12.225	0.908	12.284	0.903	12.206	0.901	–
$26a'^{-1}$	13.136	12.301	0.913	12.272	0.909	12.219	0.908	–
$24a'^{-1}$	13.355	12.399	0.910	12.308	0.907	12.228	0.904	–
$23a'^{-1}$	14.337	13.225	0.908	13.091	0.905	12.956	0.900	12.973
$22a'^{-1}$	14.525	13.578	0.902	13.512	0.897	13.427	0.895	13.487
$21a'^{-1}$	15.314	14.268	0.907	14.241	0.902	14.170	0.900	14.110
$20a'^{-1}$	15.400	14.386	0.907	14.294	0.902	14.227	0.900	14.110
$19a'^{-1}$	16.771	15.497	0.908	15.514	0.900	15.435	0.897	15.233
$18a'^{-1}$	16.892	15.618	0.908	15.626	0.900	15.553	0.898	15.233
$17a'^{-1}$	18.289	16.506	0.894	16.512	0.890	16.277	0.882	16.202
$16a'^{-1}$	18.514	16.795	0.893	16.800	0.886	16.641	0.881	18.884
$15a'^{-1}$	19.130	17.255	0.888	17.256	0.883	17.079	0.876	17.325
$14a'^{-1}$	20.829	18.797	0.885	18.745	0.882	18.534	0.875	18.551

The spectra of both molecules present distinct similarities in shape and energy position of detected transitions. Calculations reveal the highly mixed character of the transitions within the investigated range, thus there are no distinct features present, apart from the second band that was assigned to HOMO \rightarrow $3s/\sigma^*$ and HOMO-1 \rightarrow $3s/\sigma^*$ transitions, localized on the CH bond in both compounds. The mixed character of the transitions together with the complexity of the molecules and continuous overlap of the states at higher energies made it difficult to perform any unambiguous assignments. In the case of methyl butyrate, apart from a few members of some Rydberg series, a shoulder feature, marked A, was assigned to the HOMO-1 \rightarrow LUMO transition for the (a, a) conformer together with a mixed transition HOMO-1 \rightarrow LUMO + HOMO \rightarrow $3d$ for the $(g\pm, a)$ conformer, centered at 8.683 eV. A band labeled B at 9.124 eV was identified either as HOMO-3 \rightarrow $3s/\sigma^*$ for (a, a) or a mixed transition from HOMO-4 in $(g\pm, a)$. The structure centered at 9.311 eV is most likely either another

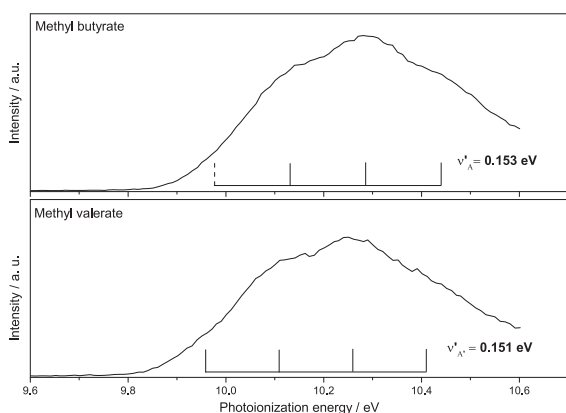
mixed transition from HOMO-4 orbital, but in (a, a) , or a HOMO \rightarrow $4p$ one in $(g\pm, a)$.

Similarly, for methyl valerate, there are 3 bands A' , B' , C' marked in the spectrum. The first, at 8.562 eV, was also assigned to a combination of HOMO-1 \rightarrow LUMO for the (a, a) conformer together with a mixed transition HOMO-1 \rightarrow LUMO + HOMO \rightarrow $3d$ for $(g\pm, a)$ conformer. The B' band was tentatively assigned at 8.744 eV and corresponds to HOMO-1 \rightarrow $3p\pi$ + HOMO \rightarrow $3d\sigma$ for the (a, a) conformer mixed with HOMO-1 \rightarrow + HOMO-3 \rightarrow $3s\sigma/\sigma^*$ for $(g\pm, a)$ one. The last band C' at about 9.107 eV was assigned based on the calculations to either a mixed transition from HOMO-2 in (a, a) conformer or also a mixed one from HOMO-3 in $(g\pm, a)$ conformer.

The HOMO \rightarrow $3s/\sigma^*$ transition in the case of both methyl butyrate and methyl valerate is associated with rather complex vibronic structure, marked for both esters in Figure 8. For both esters we identified three vibrational modes ν_a , ν_b , ν_c (labeled with prime for methyl valerate) with an average energy spacing of 180, 68 and 48 meV,

Table 2. Calculated vertical ionization energies and pseudo-intensities (pole strengths PS) of methyl valerate for (a, a) and $(g\pm, a)$ symmetries at the M06-2X/aug-cc-pVTZ geometry calculated with propagator methods, compared with experimental values, all in eV.

Configuration	Calculated/eV						Experimental/eV	
	Koopmans	OVGF	PS	P3	PS	P3+		PS
(a, a)								
$25a'^{-1}$	12.076	10.712	0.895	10.594	0.891	10.394	0.885	10.240
$7a''^{-1}$	12.516	11.456	0.901	11.354	0.895	11.228	0.892	11.000
$24a'^{-1}$	12.943	11.916	0.905	11.900	0.901	11.803	0.898	–
$6a''^{-1}$	12.827	12.039	0.914	11.926	0.910	11.876	0.909	–
$5a''^{-1}$	13.088	12.026	0.905	12.092	0.900	11.999	0.897	–
$23a'^{-1}$	13.596	12.735	0.910	12.662	0.906	12.616	0.905	–
$22a'^{-1}$	14.308	13.201	0.909	13.069	0.905	12.942	0.901	–
$21a'^{-1}$	14.660	13.609	0.905	13.551	0.901	13.471	0.898	13.457
$4a''^{-1}$	14.597	13.751	0.904	13.637	0.899	13.577	0.897	13.575
$20a'^{-1}$	15.771	14.630	0.903	14.594	0.899	14.500	0.896	14.520
$19a'^{-1}$	16.276	15.102	0.903	14.994	0.899	14.894	0.895	14.811
$3a''^{-1}$	16.087	15.003	0.911	15.002	0.902	14.961	0.901	14.811
$2a''^{-1}$	17.584	16.209	0.904	16.176	0.896	16.108	0.894	16.152
$18a'^{-1}$	18.304	16.493	0.893	16.491	0.889	16.253	0.881	16.152
$1a''^{-1}$	18.572	16.886	0.893	16.882	0.886	16.740	0.881	18.84
$17a'^{-1}$	18.989	17.054	0.886	17.063	0.881	16.874	0.874	18.84
$16a'^{-1}$	20.880	18.848	0.886	18.788	0.882	18.590	0.875	18.501
$(g\pm, a)$								
$32a'^{-1}$	12.041	10.746	0.897	10.611	0.894	10.424	0.888	10.240
$31a'^{-1}$	12.497	11.445	0.901	11.347	0.895	11.223	0.892	11.000
$30a'^{-1}$	12.631	11.639	0.905	11.673	0.900	11.578	0.897	11.489
$29a'^{-1}$	12.855	11.989	0.912	11.896	0.908	11.831	0.906	–
$28a'^{-1}$	13.037	12.009	0.906	12.009	0.902	11.916	0.899	–
$27a'^{-1}$	13.924	12.989	0.909	12.895	0.905	12.833	0.903	–
$26a'^{-1}$	14.146	13.053	0.906	12.979	0.902	12.870	0.898	–
$25a'^{-1}$	14.497	13.530	0.908	13.378	0.904	13.293	0.902	13.305
$24a'^{-1}$	14.682	13.697	0.902	13.638	0.897	13.556	0.895	13.575
$23a'^{-1}$	15.799	14.717	0.904	14.647	0.900	14.567	0.897	14.520
$22a'^{-1}$	15.978	14.853	0.904	14.745	0.900	14.662	0.897	14.520
$21a'^{-1}$	16.523	15.277	0.907	15.314	0.898	15.254	0.897	15.402
$20a'^{-1}$	17.190	15.857	0.906	15.828	0.898	15.748	0.895	15.768
$19a'^{-1}$	18.307	16.503	0.893	16.501	0.889	16.263	0.881	16.152
$18a'^{-1}$	18.476	16.736	0.892	16.734	0.885	16.574	0.880	18.84
$17a'^{-1}$	19.147	17.256	0.887	17.250	0.882	17.076	0.875	17.335
$16a'^{-1}$	20.829	18.789	0.885	18.728	0.881	18.524	0.874	18.501

**Fig. 7.** First band of photoelectron spectrum of methyl butyrate, $C_5H_{10}O_2$ (top) and methyl valerate, $C_6H_{12}O_2$ (bottom), in the 9.6–10.7 eV region, collected at 80 eV initial photon energy with the most intense vibrational modes ν'_A and $\nu'_{A'}$ resolved, respectively.**Table 3.** Experimental values of adiabatic and vertical ionization energies of methyl butyrate and methyl valerate together with the vibrational mode ν'_A that can be distinguished in the first ionization band, all in eV.

	Methyl butyrate	Methyl valerate
Adiabatic	9.977	9.959
Vertical	10.286	10.259
ν'_A	0.153	0.151

respectively. The first mode, ν_a , is present in all PA spectra of esters and corresponds to the most distinct vibration, resulting from the combination of C–O and C=O stretch. The ν_b mode is most likely to be O=C–O deformation combined with a C–O stretch. A similar structure with an energy spacing of c.a. 90 meV was seen in the spectrum of ethyl formate [9]. The mode of average spacing of 48 meV, ν_c , is ascribed to an OCC out-of-plane bend,

Table 4. Calculated electronic spectrum of methyl butyrate for (a, a) and $(g\pm, a)$ symmetries, compared with experimental values.

State	E/eV	f_L	Main character	Mixed	Exp./eV
(a, a)					
$^1A''$	5.816	0.0005	$22a' \rightarrow \text{LUMO}$		5.877
$^1A'$	7.452	0.0150	$22a' \rightarrow 3s/\sigma^*(\text{CH})$		7.557
$^1A''$	7.987	0.0119	$5a'' \rightarrow 3s/\sigma^*(\text{CH})$		7.882
$^1A''$	8.089	0.0001	$22a' \rightarrow 3p\pi$		
$^1A'$	8.146	0.0102	$22a' \rightarrow 3p\sigma$		
$^1A'$	8.222	0.0212	$22a' \rightarrow 3p\sigma$		
$^1A'$	8.332	0.0626	$5a'' \rightarrow \text{LUMO}$		8.683
$^1A''$	8.481	<0.0001	$4a'' \rightarrow 3s/\sigma^*(\text{CH})$		
$^1A'$	8.549	0.0068	$22a' \rightarrow 3d\sigma$		
$^1A''$	8.653	0.0118	$5a'' \rightarrow 3p\sigma$		
$^1A'$	8.741	0.0096	$5a'' \rightarrow 3p\pi$		
$^1A'$	8.793	0.0065	$22a' \rightarrow 3d\sigma$		
$^1A''$	8.799	0.0016		HOMO-4 \rightarrow LUMO + HOMO $\rightarrow 3d\pi$	
$^1A''$	8.810	<0.0001	$5a'' \rightarrow 3p\sigma$		
$^1A''$	8.828	0.0003		HOMO-4 \rightarrow LUMO + HOMO $\rightarrow 3d\pi$	
$^1A'$	8.849	0.0114	$21a' \rightarrow 3s/\sigma^*(\text{CH})$		9.124
$^1A'$	8.886	0.0049	$22a' \rightarrow 3d\sigma$		
$^1A''$	8.964	0.0001	$22a' \rightarrow 3d\pi$		
$^1A''$	9.087	0.0012	$5a'' \rightarrow 3d\sigma$		
$^1A'$	9.091	0.0209	$22a' \rightarrow 4s\sigma$		9.05
$^1A'$	9.105	0.0055		HOMO-4 $\rightarrow 3s/\sigma^*(\text{CH})$ + HOMO $\rightarrow 4p\sigma$	
$^1A''$	9.147	0.0192	$4a'' \rightarrow 3p\pi$		
$^1A'$	9.149	0.0737		HOMO-4 $\rightarrow 3s/\sigma^*(\text{CH})$ + HOMO-2 $\rightarrow 4p\pi$	9.311
$^1A''$	9.161	<0.0001	$22a' \rightarrow 4p\pi$		
$^1A'$	9.180	0.0116	$22a' \rightarrow 4p\sigma$		
$(g\pm, a)$					
1A	5.808	0.0007	$28a \rightarrow \text{LUMO}$		5.877
1A	7.497	0.0118	$28a \rightarrow 3s/\sigma^*(\text{CH})$		7.557
1A	8.025	0.0141	$27a \rightarrow 3s/\sigma^*(\text{CH})$		7.882
1A	8.069	0.0031	$28a \rightarrow 3p$		
1A	8.132	0.0040	$28a \rightarrow 3p$		
1A	8.225	0.0229	$28a \rightarrow 3p$		
1A	8.312	0.0505		HOMO-1 \rightarrow LUMO + HOMO $\rightarrow 3d$	8.683
1A	8.532	0.0372	$28a \rightarrow 3d$		
1A	8.636	0.0083	$27a \rightarrow 3p$		
1A	8.695	0.0033		HOMO-2 $\rightarrow 3s/\sigma^*(\text{CH})$ + HOMO-1 $\rightarrow 3p$	
1A	8.742	0.0031		HOMO-4 \rightarrow LUMO + HOMO $\rightarrow 3d$	
1A	8.764	0.0095		HOMO-1 $\rightarrow 3p$ + HOMO $\rightarrow 3d$	
1A	8.775	0.0040		HOMO $\rightarrow 3d$ + HOMO-1 $\rightarrow 3p$	
1A	8.800	0.0079	$28a \rightarrow 3d$		
1A	8.809	0.0107	$25a \rightarrow 3s/\sigma^*(\text{CH})$		
1A	8.841	0.0216		HOMO-4 $\rightarrow 3s/\sigma^*(\text{CH})$ + HOMO-1 $\rightarrow 3p$	9.124
1A	8.899	0.0016	$28a \rightarrow 3d$		
1A	8.902	0.0026	$28a \rightarrow 3d$		
1A	8.948	0.0007	$28a \rightarrow 4s$		
1A	9.044	0.0034	$28a \rightarrow 4p$		
1A	9.110	0.0103	$27a \rightarrow 3d$		
1A	9.133	0.0439	$26a \rightarrow 3p$		9.311
1A	9.148	0.0040	$28a \rightarrow 4p$		
1A	9.171	0.0220		HOMO $\rightarrow 4p$ + HOMO-3 $\rightarrow 3p$	
1A	9.183	0.0078	$28a \rightarrow 4p$		

Table 5. Calculated electronic spectrum of methyl valerate for (a, a) and $(g\pm, a)$ symmetries, compared with experimental values.

State	E/eV	f_L	Main character	Mixed	Exp./eV
(a, a)					
$^1A''$	5.817	0.0005	$25a' \rightarrow$ LUMO		5.883
$^1A'$	7.449	0.0127	$25a' \rightarrow 3s\sigma/\sigma^*(CH)$		7.478
$^1A''$	7.983	0.0111	$7a'' \rightarrow 3s\sigma/\sigma^*(CH)$		7.852
$^1A''$	8.063	<0.0001	$25a' \rightarrow 3p\pi$		
$^1A'$	8.127	0.0017	$25a' \rightarrow 3p\sigma$		
$^1A'$	8.207	0.0080	$25a' \rightarrow 3p\sigma$		
$^1A''$	8.322	0.0002	$6a'' \rightarrow 3s\sigma/\sigma^*(CH)$		
$^1A'$	8.327	0.0676	$7a'' \rightarrow$ LUMO		8.562
$^1A'$	8.465	0.0392	$7a'' \rightarrow 3p\sigma$		
$^1A'$	8.608	0.0019		HOMO $\rightarrow 3d\sigma$ + HOMO-3 $\rightarrow 3p\sigma$	
$^1A''$	8.631	0.0068	$7a'' \rightarrow 3p\sigma$		
$^1A'$	8.714	0.0199		HOMO-1 $\rightarrow 3p\pi$ + HOMO $\rightarrow 3d\sigma$	8.744
$^1A''$	8.728	0.0012		HOMO $\rightarrow 3p\pi$ + HOMO-3 \rightarrow LUMO	
$^1A'$	8.728	0.0045		HOMO $\rightarrow 3d\sigma$ + HOMO-1 $\rightarrow 3p\pi$	
$^1A'$	8.777	0.0045	$5a'' \rightarrow 3s\sigma/\sigma^*(CH)$		
$^1A''$	8.784	0.0002		HOMO-3 \rightarrow LUMO + HOMO $\rightarrow 3d\pi$	
$^1A''$	8.797	0.0014		HOMO-1 $\rightarrow 3d\sigma$ + HOMO-3 \rightarrow LUMO	
$^1A'$	8.842	0.0049	$25a' \rightarrow 3d\sigma$		
$^1A''$	8.876	0.0044	$5a'' \rightarrow 3s\sigma/\sigma^*(CH)$		
$^1A'$	8.926	0.0605		HOMO-2 $\rightarrow 3p\pi$ + HOMO-1 $\rightarrow 3p\pi$	9.012
$^1A''$	8.939	0.0002	$25a' \rightarrow 3d\pi$		
$^1A''$	8.966	0.0570		HOMO-2 $\rightarrow 3p\sigma$ + HOMO-1 $\rightarrow 3d\sigma$	9.107
$^1A'$	9.019	0.0085		HOMO-3 $\rightarrow 3d\sigma$ + HOMO-4 $\rightarrow 3p\sigma$	
$^1A''$	9.074	0.0093	$24a' \rightarrow 3p\pi$		
$^1A'$	9.102	0.0078		HOMO $\rightarrow 3d\sigma$ + HOMO-3 $\rightarrow 3p\sigma$	
$(g\pm, a)$					
1A	5.812	0.0007	$30a \rightarrow$ LUMO		5.883
1A	7.506	0.0120	$30a \rightarrow 3s\sigma/\sigma^*(CH)$		7.574
1A	8.016	0.0122	$29a \rightarrow 3s\sigma/\sigma^*(CH)$		7.962
1A	8.05	0.0024	$30a \rightarrow 3p$		
1A	8.104	0.0020	$30a \rightarrow 3p$		
1A	8.188	0.0141	$30a \rightarrow 3p$		
1A	8.304	0.0643		HOMO-1 \rightarrow LUMO + HOMO $\rightarrow 3d$	8.562
1A	8.399	0.0018	$28a \rightarrow 3s\sigma/\sigma^*(CH)$		
1A	8.422	0.0231		HOMO-3 $\rightarrow 3s\sigma/\sigma^*(CH)$ + HOMO $\rightarrow 3d$	
1A	8.526	0.0231		HOMO $\rightarrow 3d$ + HOMO-3 $\rightarrow 3s\sigma/\sigma^*(CH)$	
1A	8.648	0.0076	$29a \rightarrow 3p$		
1A	8.663	0.0066	$29a \rightarrow 3p$		
1A	8.709	0.0040	$30a \rightarrow 3d$		
1A	8.720	0.0180		HOMO-1 $\rightarrow 3p$ + HOMO-3 $\rightarrow 3s\sigma/\sigma^*(CH)$	8.744
1A	8.758	0.0038		HOMO-5 \rightarrow LUMO + HOMO-4 $\rightarrow 3s\sigma/\sigma^*(CH)$	
1A	8.776	0.0069		HOMO-1 $\rightarrow 3p$ + HOMO-3 $\rightarrow 3s\sigma/\sigma^*(CH)$	
1A	8.814	0.0198		HOMO $\rightarrow 3d$ + HOMO-2 \rightarrow LUMO	
1A	8.846	0.0020	$30a \rightarrow 3d$		
1A	8.877	0.0015	$30a \rightarrow 3d$		
1A	8.883	0.0022		HOMO $\rightarrow 3d$ + HOMO-1 $\rightarrow 3d$ + HOMO-3 $\rightarrow 3s\sigma/\sigma^*(CH)$	
1A	8.911	0.0106		HOMO-3 $\rightarrow 3p$ + HOMO-1 $\rightarrow 3d$	
1A	8.915	0.0123		HOMO-2 $\rightarrow 3p$ + HOMO $\rightarrow 3d$	
1A	8.947	0.0575		HOMO-3 $\rightarrow 3p$ + HOMO-1 $\rightarrow 3d$	9.012
1A	8.991	0.0110	$28a \rightarrow 3p$		
1A	9.032	0.0248		HOMO $\rightarrow 4s$ + HOMO-3 $\rightarrow 3p$ + HOMO-1 $\rightarrow 3d$	9.107

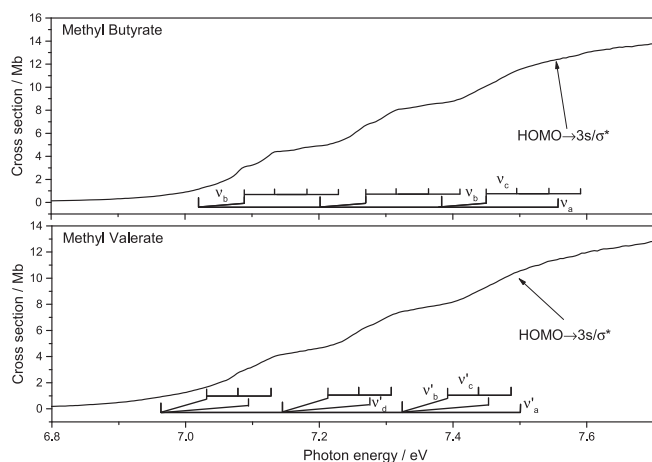


Fig. 8. First band of photoabsorption spectrum of methyl butyrate, $C_5H_{10}O_2$ (top) and methyl valerate, $C_6H_{12}O_2$ (bottom), in the 6.8–7.8 eV range with the most intense vibrational modes ν_a , ν_b , ν_c and $\nu_{a'}$, $\nu_{b'}$, $\nu_{c'}$ and $\nu_{d'}$ resolved, respectively.

seen in the PA spectrum of isobutyl acetate [10]. For methyl valerate we also resolved a ν'_d mode of 130 meV and assigned this to skeletal C–C stretching, which was also observed in methyl and ethyl formates [8,9].

In the photoabsorption spectrum of methyl valerate, a series of three peaks, evenly spaced with $\nu_\alpha = 144$ meV appear at about 6.5 eV (Fig. 4). Such feature has not been observed previously for other ester molecules [8–12]. This series may be an additional vibrational mode, such as a series of C–C skeletal stretching, already seen here as ν_d associated with the ν_a mode.

Over the whole range of the photoabsorption spectrum investigated, only the first two members of the Rydberg series converging to the two highest ionic states were resolved, mainly due to state overlap as a result of the presence of the two conformers for each compound. The other transitions were assigned based purely on the results from the computer calculations and the value of the oscillator strength, since it was not possible to unambiguously determine the value of the ionization energy from HOMO-2, HOMO-3 and HOMO-4 for both esters. The resolved states are marked in Figure 3 for methyl butyrate and Figure 4.

The values obtained for Rydberg series converging to the first two ionization energies are summarized in Table 6 with the value of the quantum defect calculated using the well-known Rydberg formula. As for previously investigated esters, the ns series values of δ are between 0.7 and 0.9 for members of all transition families with the quantum defect values being higher for the transitions converging to HOMO-1 ionization band than those converging to the HOMO.

5.3 Atmospheric photolysis

There is a very little information on the possible interaction of free radicals with methyl butyrate and methyl valerate. There is also no information in the literature

on the photolysis rate and thus the lifetimes of these esters in the upper atmosphere. Therefore, the present absolute cross sections can be used in combination with solar actinic flux [34] measurements from the literature to estimate the photolysis rate of both methyl esters in the atmosphere from an altitude close to the ground, to the stratopause at 50 km. Details of the calculation programme were published previously [35] and the quantum yield for dissociation following absorption in that programme is assumed to be unity. The reciprocal of the photolysis rate at a given altitude corresponds to the local photolysis lifetime.

Photolysis lifetimes of less than 1 sunlit day were calculated at altitudes above 30 km for methyl butyrate, which means that the molecules can be broken up quite efficiently by VUV absorption above this altitude. Also at ground level the lifetimes are shorter than one day. UV photolysis is therefore expected to play a significant role in the tropospheric and stratospheric removal of methyl butyrate.

For both methyl butyrate and methyl valerate the stratospheric lifetime is long, about 6 days at 30 km. The lifetimes at 10 km are extremely long for these molecules, reaching thousands of years. Therefore, photolysis will not be the main sink mechanism for the removal of the molecules from the atmosphere. This is similar to previously analyzed esters, ethyl and methyl formate, for which UV photolysis was also not expected to play a significant role, but in contrast to isobutyl formate, where photolysis may be a major removal mechanism.

The only relevant investigations found for these esters presented kinetic studies of hydroxyl radical reactions [36]. The lifetimes obtained there with typical tropospheric OH concentrations of 10^6 molecules per cm^3 yielded 3.5 days for methyl butyrate and 2.4 days for methyl valerate. This would certainly suggest that the reaction with OH radical would be a prevalent sink mechanism for both methyl butyrate and methyl valerate.

6 Conclusions

For the first time the complete electronic spectra of methyl butyrate and methyl valerate are presented here together with absolute photoabsorption cross sections from 4.5 to 10.8 eV. The structures that can be observed in the spectrum can be assigned to both valence and Rydberg transitions, based on ab initio calculations of vertical excitation energies and oscillator strengths of these molecules. Fine structure, that was resolved both in the photoelectron and photoabsorption spectra, has been assigned to vibrational series involving, predominantly, excitations of c.a. 180 meV, attributed to C=O and C–O stretching. Other vibrations resolved in the photoabsorption spectrum of both molecules correspond to further stretches and deformations that were also seen in the previously analyzed molecules. The theoretical calculations presented here are in good agreement with experimental data. The photoabsorption cross sections were used to calculate the photolysis lifetimes of methyl butyrate and methyl valerate for the Earth's troposphere and stratosphere. From this and

Table 6. Energies (eV), quantum defects and assignments of the ns Rydberg series converging to the first two ionic electronic ground state of methyl butyrate and methyl valerate.

Vertical transition energy/eV	Quantum defect, δ	Assignment
Methyl butyrate		
	$IE_1 = 10.286$ eV	
7.56	0.77	$3s\sigma^*$
9.05	0.87	$4s\sigma$
	$IE_2 = 11.002$ eV	
7.88	0.91	$3s\sigma^*$
9.60	0.88	$4s\sigma$
Methyl valerate		
	$IE_1 = 10.259$ eV	
7.48	0.79	$3s\sigma^*$
8.99	0.73	$4s\sigma$
	$IE_2 = 11.000$ eV	
7.85	0.92	$3s\sigma^*$
9.57	0.91	$4s\sigma$

comparing our results with those obtained from kinetic studies of reaction with hydroxyl radicals, we conclude that photolysis is not a leading removal mechanism for either methyl butyrate or methyl valerate.

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Author contribution statement

MAS performed the PA and PES measurements, overall data analysis and wrote the manuscript. DD performed

computations and analysis of their outcomes. SVH and NCJ performed PA measurements, whereas LZ and MMD – PES measurements. PLV performed calculations for the photolysis section. NJM provided the samples for both synchrotron runs. All authors commented on the manuscript. All the authors have read and approved the final manuscript.

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