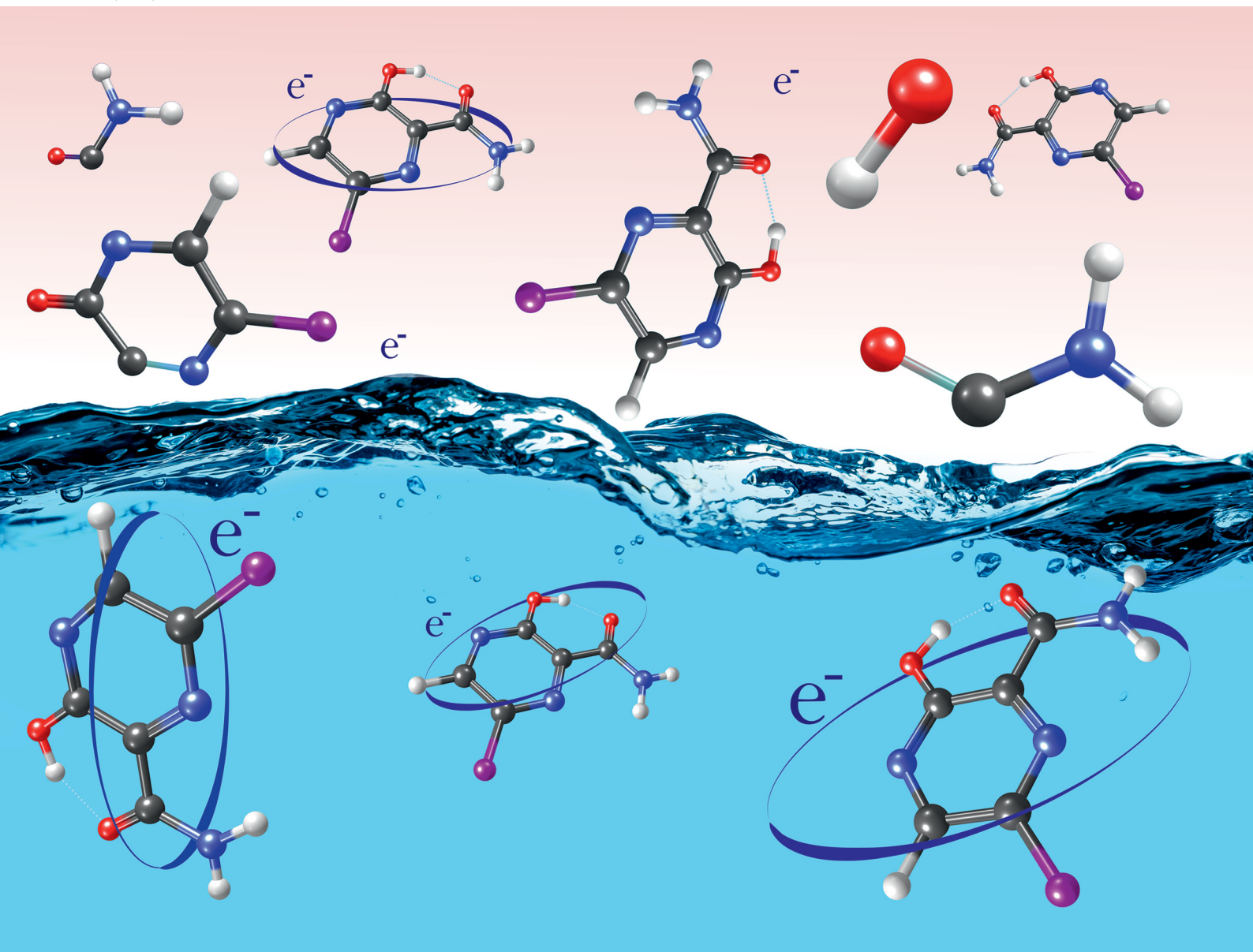


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Electron attachment to isolated and microhydrated favipiravir†

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Electron attachment and its equivalent in complex environments, single-electron reduction, are important in many biological processes. Here, we experimentally study the electron attachment to favipiravir, a well-known antiviral agent. Electron attachment spectroscopy is used to explore the energetics of associative (AEA) and dissociative (DEA) electron attachment to isolated favipiravir. AEA dominates the interaction and the yields of the fragment anions after DEA are an order of magnitude lower than that of the parent anion. DEA primary proceeds *via* decomposition of the CONH₂ functional group, which is supported by reaction threshold calculations using *ab initio* methods. Mass spectrometry of small favipiravir–water clusters demonstrates that a lot of energy is transferred to the solvent upon electron attachment. The energy gained upon electron attachment, and the high stability of the parent anion were previously suggested as important properties for the action of several electron-affinic radiosensitizers. If any of these mechanisms cause synergism in chemo-radiation therapy, favipiravir could be repurposed as a radiosensitizer.

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1 Introduction

The present study is motivated by the fact that many biological processes based on reduction can be related to electron attachment (EA). This relation was explored by the Modelli and Pshenichnyuk groups for a wide range of biological processes, from metabolic pathways to the functionality of the olfactory system.^{1–7} The electron reduction properties can also influence the transport of drugs through biological membranes and influence their target binding properties.^{8,9} Recently, the RNA

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† Electronic supplementary information (ESI) available: Low *m/z* part of the spectra depicted in Fig. 6, panels (c) and (d). Comparison of the reaction energies calculated using B3LYP and M062x functionals, Cartesian coordinates of favipiravir conformers and its DEA anion and neutral fragments at B3LYP and M062x levels of theory. See DOI: 10.1039/d1cp02686k



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inhibitor efficiency of favipiravir tautomers was related to the energy difference between its highest occupied (HOMO) and lowest unoccupied (LUMO) molecular orbitals.¹⁰ Unoccupied orbitals can become singly occupied upon electron attachment and appear as so-called shape resonances in the electron scattering spectrum. A present study of the electron attachment to favipiravir may therefore provide an important key to better understand its activity.

Another important motivation is based on the relation of low-energy electrons to the synergism observed in the concomitant chemo-radiation therapy of cancer.^{11,12} Two facts motivated a range of studies on the relation of low-energy electrons to the synergistic action of radiation with a range of chemotherapeutic and radio-sensitizing drugs.^{13,14} The first fact is a large amount of available secondary low-energy electrons in the irradiated tissue.^{15,16} The second fact is that despite the different possible modes of action,¹⁷ most of the known small molecule radiosensitizers and their important functional groups have high electron affinities.¹⁸ Processes such as the formation of reactive anions and radical species *via* dissociative electron attachment,^{19–22} enhanced linear energy transfer,²³ transport properties⁸ due to associative electron attachment, or DNA sensitization^{24–30} have been proposed as possible sources of synergism observed in concomitant chemo-radiation therapy. Several of the processes were then used to suggest novel radiosensitizers.^{31–34} However, in many cases, the suggested molecules with “ideal” electron-attachment properties fail to exhibit the synergism³⁵ or are biologically incompatible.³⁶ Since drug repurposing³⁷ is becoming still more important and antivirals are often chosen as cancer chemo or even chemo-radio therapeutics,^{38–43} we explore favipiravir as a promising candidate for an electron affinic radiosensitizer.

Favipiravir (Fig. 1) has already been established as a broad-range antiviral^{44–47} with good biocompatibility⁴⁸ and known pharmacokinetics.^{49,50} Here, we explore its behavior in the reactions with low-energy electrons using electron attachment

spectroscopy of the isolated and microhydrated molecule and perform basic computational modeling to support our experimental data.

2 Methods

2.1 Experimental

Favipiravir was purchased from Santiago Labs with stated purity of 99% and its electron-induced chemistry was explored on two experimental setups. The isolated molecule was studied using the TEM-QMS setup⁵¹ and hydration effects were studied using the CLUSTER Beam (CLUB) apparatus in the M. Fárník group.⁵²

The TEM-QMS apparatus is an electron attachment spectrometer combining a trochoidal electron monochromator (TEM) with a quadrupole mass spectrometer (QMS) analyser originally built in the M. Allan group.⁵³ With the TEM,⁵⁴ electrons thermo-emitted from an iridium-yttrium cathode, are selected according to their kinetic energy, narrowing the initial electron energy distribution function. In the present experiment, the electron-energy resolution was around 150 meV as estimated from the FWHM of the 0 eV peak in the anion yield for AEA to sulphur hexafluoride.⁵⁵ The electrons collide with sample molecules in the reaction chamber and the formed ions are extracted towards the QMS. Two modes of operation are possible, with ion yield measurement as a function of incident electron energy at a constant m/z of QMS or ion yield measurement as a function of m/z at constant incident electron energy. Favipiravir was introduced using the direct insertion probe.⁵⁶ Sample powder was loaded into a glass bulb and placed at the end of the probe, which was inserted into a resistively heated copper cylinder with the constant temperature kept at 340 K. The sublimed molecules effused into the reaction zone through a 1 cm long capillary. For calibration of the energy axis, we used SF₆ and CO₂ gases, which were introduced through the same inlet. The 4.3 eV resonance of O[−] from CO₂⁵⁷ was used to calibrate the electron-energy axis and 0 eV resonance of SF₆[−] from SF₆ to determine the energy threshold for electron transmission through the TEM. In the present case, the threshold is around 0.1 eV and constant electron current is reached around 0.25 eV as demonstrated on SF₆[−] signal in Fig. 2. Anion yields lower than this value presented in this work, particularly the yield of parent anion, are therefore underestimated.

On the CLUB setup, we only used the neutral cluster source and reflectron time-of-flight mass spectrometer. Experimental details for the negative and positive mass spectra measurements can be found in ref. 58 and 59, respectively. The molecular target was prepared by co-expansion of He buffer gas and favipiravir vapors through a conical 90 μm nozzle into the vacuum. Clusters of microhydrated molecules were prepared by an approach developed in our laboratory based on the addition of a small amount of water into the buffer gas through the Nafion membrane.⁶⁰ During all experiments, the sample was sublimed at a temperature of approximately 85 °C. To test



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research is focused on environmental effects on reaction dynamics and elementary processes in radiation interaction with living tissue, studied experimentally on model systems ranging from isolated molecules and clusters to self-assembled nanostructures.

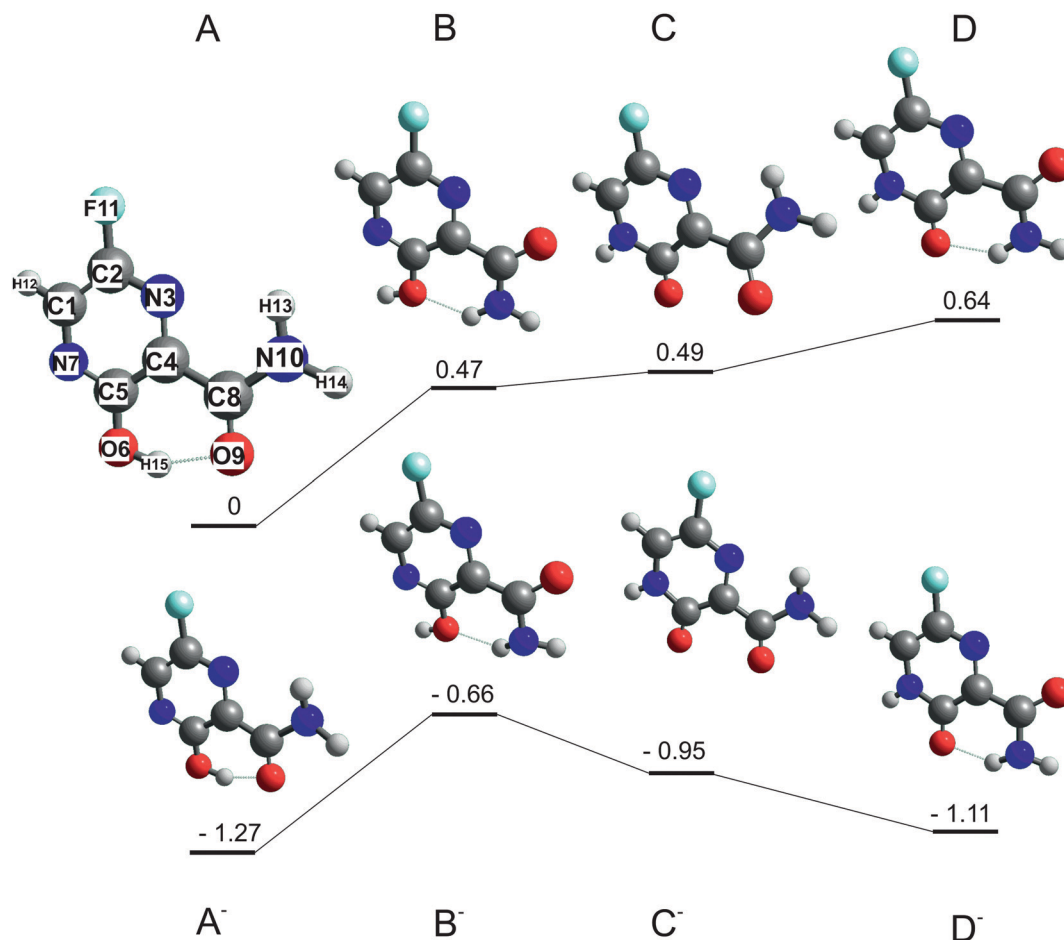


Fig. 1 Optimized structures of stable conformers A, B and keto tautomers B, C of favipiravir and their respective anions. Energies in eV relative to the most stable neutral conformer A calculated at the B3LYP/aug-cc-pVTZ level of theory.

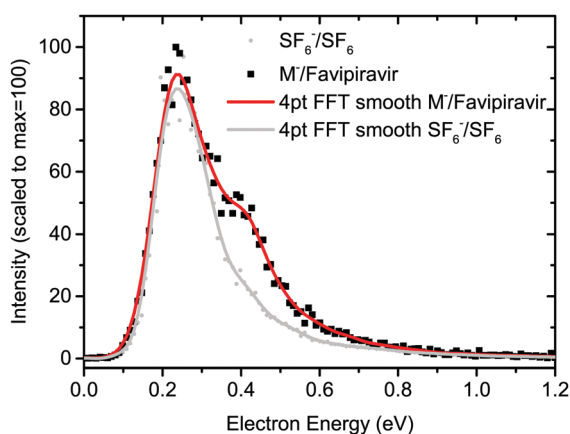


Fig. 2 Electron energy-dependent ion yields for AEA to favipiravir and sulfur hexafluoride. The drop of the anion yield below ~ 0.25 eV is caused by decrease of the incident electron current below this value. Anion yields at energies lower than ~ 0.25 eV presented in this work are, therefore, underestimated.

the thermal stability of the molecule, we heated the sample up to 118°C . Even at this elevated temperature, we did not observe any new product ions due to thermal decomposition in the ionization MS (see ESI[†]).

2.2 *Ab initio* calculations

Two favipiravir conformers, differing in the orientation of the CONH_2 functional group, its keto form, and their respective anions, were pre-optimized at the B3LYP/6-31(d) level of theory and then optimized at the B3LYP/aug-cc-pVTZ level of theory. The structures of neutrals and respective energies of neutrals and anions optimized at the B3LYP/aug-cc-pVTZ level of theory are in Fig. 1. The order of the conformers was checked by energy calculation using G3MP2⁶¹ method and method error by re-optimizing the B3LYP⁶² structures using M062x⁶³ functional and the same aug-cc-pVTZ basis set. Adiabatic electron affinities were calculated as a difference of the neutral and anion energies in their optimized geometry and they are listed together with other parameters in Table 1.

Then, energetic thresholds for individual reaction channels of DEA to favipiravir were calculated at the B3LYP/aug-cc-pVTZ theory to support the experimental results. Threshold energies were obtained using the formula:

$$E_{\text{th}} = E_{\text{M}_a^-} + E_{\text{M}_b} - E_{\text{M}} \quad (1)$$

where $E_{\text{M}_a^-}$ and E_{M_b} are the energies of the anion and neutral fragments respectively (if there are multiple neutral fragments,

Table 1 Relative energies and dipole moments of optimized ground state neutral and anion conformers of favipiravir computed at the B3LYP/aug-cc-pVTZ, G3MP2 and M062x/aug-cc-pVTZ level of theory. All energies are in eV relative to the conformer A, which is the lowest energy neutral conformer at all levels of theory. All dipole moments are in D

Conformer	Energy B3LYP	Energy G3MP2	Energy M062x	Dipole moment B3LYP
A	0	0	0	3.2
B	0.47	0.44	0.79	5.7
C	0.49	0.57	0.34	6
D	0.64	0.72	1.09	5.3
A ⁻	-1.27	-1.14	-0.68	4.8
B ⁻	-0.66	-0.54	-0.3	7.8
C ⁻	-0.95	-0.91	-0.9	9.2
D ⁻	-1.11	-0.73	-0.5	9.1

then E_{M_b} is the sum of all neutral fragment energies), and E_M is the energy of the neutral parent molecule. These energies correspond to the sum of electronic and zero-point correction energies in the Gaussian output file. Again the method error was checked by recalculating the values at M062x/aug-cc-pVTZ level (see ESI†).

Finally, virtual orbital energies of favipiravir were calculated at the HF/6-31G* level of theory, using the structures optimised at the B3LYP/aug-cc-pVTZ level of theory. These energies were then scaled to obtain vertical attachment energies according to the empirical formula of Aflatooni *et al.*⁶⁴:

$$VAE = [\varepsilon_{VO} - 2.5553]/1.3749 \quad (2)$$

where VAE is the vertical attachment energy and ε_{VO} is the virtual orbital energy.

All calculations were performed in Gaussian 16⁶⁵ and the results were analyzed and visualized using Chemcraft.⁶⁶

3 Results and discussion

3.1 Isolated molecule

Structures of the most stable isomers and keto tautomers of favipiravir are shown in Fig. 1. Conformer A is the most stable conformer of neutral favipiravir, independent of the computational method (Table 1), due to the aromatic nature of the ring and being stabilized by O(9)–H(15) hydrogen bond (atom numbers are shown on the structure of conformer A of Fig. 1). The energy of the conformer is (depending on the method) ~ 0.5 eV below the other studied structures and therefore will dominate the gas phase distribution of the sublimed favipiravir. The recent work of Antonov⁶⁷ also demonstrated the stability of this conformer in various solvents. We will therefore relate all energies in the following discussion to this most stable conformer A. The energy difference due to different conformers of the neutral precursor molecule may be easily estimated from Table 1.

The calculated adiabatic electron affinity of the molecule is 1.3 eV (Fig. 1 and Table 1). Therefore, favipiravir can form a stable valence-bound anion, which is also close in geometry to the neutral molecule. However, in its anionic state, the energy gap between the enol A conformer and keto D conformer is

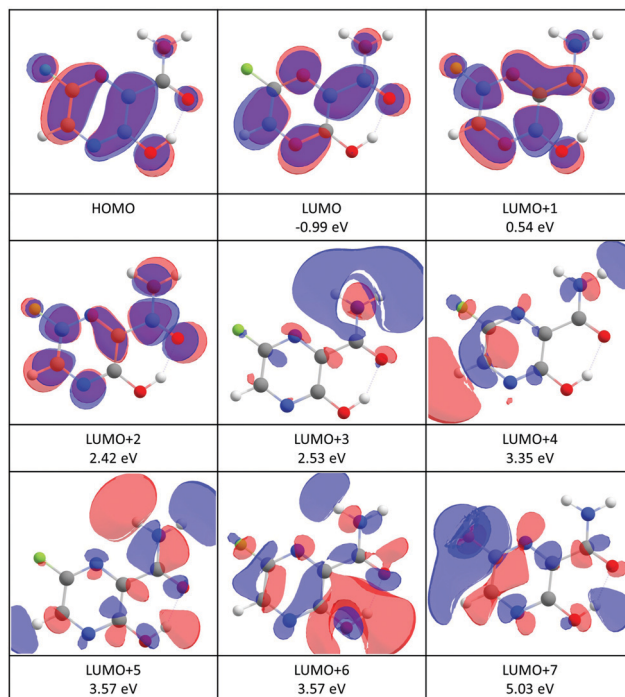


Fig. 3 Contour plots of the highest occupied (HOMO) and low-lying virtual orbitals of favipiravir. The orbitals were calculated at the HF/6-31G(d) level of theory. The numbers in eV are vertical attachment energies according to the formula (2).

much lower than in the case of the neutral molecule and therefore both forms may be present. This may influence the dissociation dynamics, as will be discussed later.

An estimated value of the vertical electron affinity at the B3LYP/aug-cc-pVTZ level of theory is 0.3 eV. The value may vary due to the limitations of the DFT approach,⁶⁸ however, its positive character is clear also from our estimation of the negative value of the LUMO vertical attachment energy (Fig. 3).

Favipiravir has a large dipole moment. The most stable conformer A has the lowest dipole moment (3.3 D) of all explored conformers, above the commonly set limits for the formation of dipole supported states of the anion (ref. 69 and references cited therein).

The incident electron energy dependent ion yields for the most stable anionic products of electron attachment to isolated favipiravir are shown in Fig. 4 and relative intensities of the bands, obtained as the area of Gaussian peaks fitted to the data, are listed in Table 2. The yields peak at energies of around 0, 1, 2, 3, 4, 5, 7, and 9 eV. The 0 eV peak is observed only for the parent anion. The 2, 3, and 5 eV peaks can be assigned to shape resonances according to estimated vertical attachment energies of LUMO+2, 4, and 7 (see Fig. 3). The yields peaking at 1 or 4 eV can also be due to the attachment to virtual orbitals, however, their low energy onset may be shifted due to the reaction endothermicity, which will be discussed later. The high energy peaks in the spectrum can be formed by core excited Feshbach resonances or core excited shape resonances. Unfortunately, the photo-electron spectrum of favipiravir, which would allow

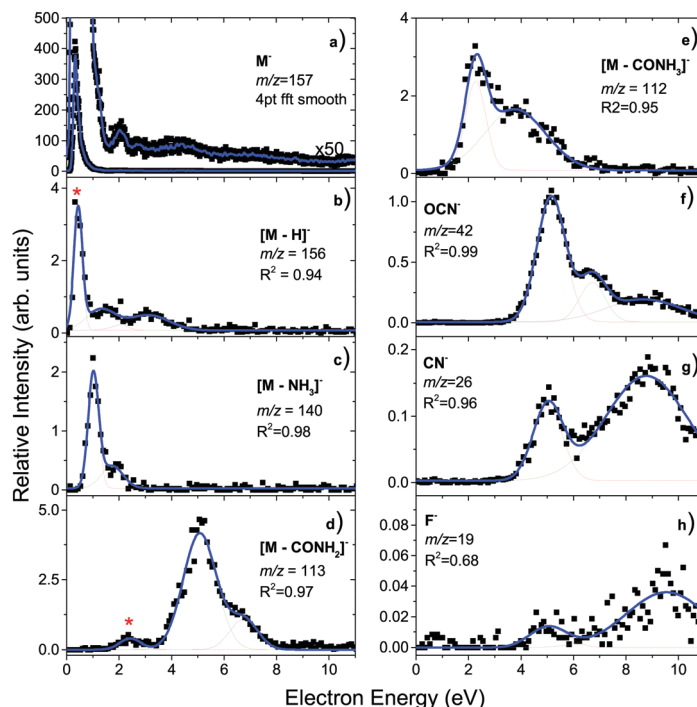


Fig. 4 Electron-energy dependent ion yields for the main products of EA to favipiravir. The contributions of ion signal from neighboring m/z due to low selectivity of QMS are marked by *.

us to better identify the Feshbach resonances^{70–72} has not yet been measured.

Positions of the Feshbach resonances can be, however, estimated on the basis of known resonances for components of the molecule,^{56,73} such as pyrazine or small amides. From the known electron energy loss spectrum of pyrazine,⁷⁴ which matches the central favipiravir ring structure, we expect that Feshbach resonances in favipiravir can occur as low as at 4 eV. Further excited states of pyrazine are available at 5 eV, 6.5 eV, and 7.5 eV. For the amide functional group, excited states may be expected at higher energies of 7 eV and 9 eV, based on electron energy loss and DEA data of small amides.^{75,76} However, for amides also an alternative process of core excited shape resonances was suggested,⁷⁷ but has yet to be confirmed.⁷⁸

The product anion yield is dominated by the parent favipiravir anion M^- . Several mechanisms for the formation of the anion at low energies are possible. As we discussed in the previous paragraphs, the attachment is exothermic and therefore attachment at 0 eV can be expected. The high dipole moment and similar geometries of the neutral and anion allow for dipole-supported vibrational Feshbach resonances, which are known DEA to various molecules.^{79–81} The electron-energy dependent ion yield curve of the M^- anion is shown in detail in Fig. 2. First, we can not see the onset at expected 0 eV but at ~ 0.1 eV due to the electron transmission function of the monochromator, which can be well demonstrated on the comparison with SF_6^- ion yield with the known 0 eV resonance and similar onset. Second, there is a small bump at energies ~ 0.4 eV, which may be an indication of vibrational structure. However, the vibrational mode can not be unambiguously

assigned because the spectrum is shifted by the electron transmission function of the monochromator. In bare pyrazine (the central cyclic structure of favipiravir), the structures in electron scattering were assigned to breathing modes of the ring.⁸² In the present case, the energy is high, if we exclude a possibility that more structures occur in the part of the ion yield to which we are blind, the modes involved will be rather O–H or C–H stretches. However, the bump in the spectrum may also be simply an experimental artefact or indication of attachment *via* the low-lying π shape resonance. Vertical attachment to LUMO+1 may occur for electrons with energies ~ 0.5 eV (Fig. 3) according to the scaling of Aflatooni *et al.*⁶⁴

A clear structure around 2 eV in the spectrum of M^- , in the panel a) of Fig. 4 can be also assigned to a shape resonance of LUMO+2.

We can see that for several bands in the spectrum of favipiravir, there are multiple possible explanations of the undergoing attachment mechanism. Their unambiguous identification requires further experiments. We will not speculate here and will instead focus on the exit channel of the DEA, which was directly studied in the present experiment.

The main DEA channels of favipiravir are listed in Table 2, together with their calculated threshold energies and information about the relative intensities of the main peaks in the experimentally measured ion yields (Fig. 4). As already mentioned, AEA dominates the interaction. In Table 2, where the intensity of the near 0 eV peak of M^- is set to 100, the intensity of the second most intense peak in the yield of EA products to favipiravir corresponds to the loss of the $CONH_2$ functional group at energies of incident electrons ~ 5 eV, which has a

Table 2 Main EA channels. Reaction thresholds obtained from calculations using B3LYP functional with the aug-cc-pVTZ basis set. Peak positions and relative intensities of anions from electron-energy dependent ion yields. All energies are presented in units of eV. Relative intensities in arbitrary units with maximum set to 100. The error of the relative intensity is influenced by the settings of the ion optics of the mass spectrometer and may reach 20%. The error of the peak position is on the level of the current TEM resolution ~ 150 meV. Structures of fragment conformers labelled A, B, C... are available in the ESI

m/z	Anion	Neutral	$M + e^- \rightarrow$ products calculated threshold	Experimental peak energy (relative intensity)		
157	$C_5H_4FN_3O_2^-$			$\sim 0(100);$	2.2(0.06)	
156	$C_5H_3FN_3O_2^-$			1.8(0.2);	3.2(0.12)	
	$[M-H]^-$ A	H(15)	0.64			
	$[M-H]^-$ B	H(13)	1.03			
	$[M-H]^-$ C	H(15)	1.03			
	$[M-H]^-$ D	H(12)	2.05			
	$[M-H]^-$ E	H(13)	2.22			
	$[M-H]^-$ F	H(12)	2.50			
140	$C_5HFN_2O_2^-$			1.1(1.4);	1.9(0.26)	
	$[M-NH_3]^-$	NH ₃	0.17			
	$[M-OH]^-$ A	OH	2.96			
	$[M-OH]^-$ B	OH	3.17			
	$[M-OH]^-$ C	OH	4.92			
	$[M-OH]^-$ D	OH	5.03			
113	$C_4H_2FN_2O^-$			5.0(6.0);	6.8(1.1)	8.5(0.67)
	$[M-CONH_2]^-$ A	CO + NH ₂	3.24			
	$[M-CONH_2]^-$ B	CO + NH ₂	3.46			
	$[M-CONH_2]^-$ C	CO + NH ₂	3.62			
	$[M-CONH_2]^-$ A	CONH ₂	4.39			
	$[M-CONH_2]^-$ B	CONH ₂	4.61			
	$[M-CONH_2]^-$ C	CONH ₂	4.77			
112	$C_4HFN_2O^-$			2.3(1.8);	4.3(3.3)	
	$[M-CONH_3]^-$ A	CONH ₃	0.86			
	$[M-CONH_3]^-$ A	CO + NH ₃	1.31			
	$[M-CONH_3]^-$ B	CONH ₃	1.96			
	$[M-CONH_3]^-$ B	CO + NH ₃	2.41			
	$[M-CONH_3]^-$ C	CONH ₃	2.35			
	$[M-CONH_3]^-$ C	CO + NH ₃	2.8			
	$[M-CONH_3]^-$ D	CONH ₃	2.4			
	$[M-CONH_3]^-$ D	CO + NH ₃	2.85			
42	OCN ⁻	—	—	5.0(1.4);	6.6(0.4);	
	OCN ⁻	$C_4H_4FN_2O$	1.93			
	OCN ⁻	$H_2 + C_4H_2FN_2O$ A	4.1			
	OCN ⁻	$H_2 + C_4H_2FN_2O$ B	4.87			
	OCN ⁻	$H_2 + C_4H_2FN_2O$ C	4.95			
26	CN ⁻	—	—	5.0(0.17);	8.7(0.55)	
	CN ⁻	$H_2O + C_4H_2FN_2O$ A	1.85			
	CN ⁻	$C_4H_4FN_2O_2$	1.97			
	CN ⁻	$HF + C_4H_3N_2O_2$	2.61			
	CN ⁻	$H_2O + C_4H_2FN_2O$ B	2.62			
	CN ⁻	$H_2O + C_4H_2FN_2O$ C	2.7			
19	F ⁻	—	—	5.0(0.02);	9.4(0.1)	
	F ⁻	(M-F) A	1.72			
	F ⁻	(M-F) B	2.15			
	F ⁻	(M-F) C	2.15			
	F ⁻	(M-F) D	2.34			

relative intensity of 6. It is important to mention that the relative intensity ratios in the present experiment may be influenced by the ion optics and quadrupole mass filter transmission settings, which are mass-dependent. To minimize these effects we tuned the ion optics for maximum transmission at $m/z = 79$, Br⁻ anion, which is present as a background in the TEM-QMS setup and is approximately in the center of the m/z fragment distribution. However, the relative intensities may still vary by up to $\sim 20\%$.

$[M-CONH_2]^-$ can be formed by the simple breaking of the C(4)–C(8) bond. The electron energy-dependent spectrum for its formation is shown on panel d) of Fig. 4. The calculated threshold energy for this process is 4 eV (Table 2). This means that there is sufficient energy in the system for this process to occur at incident electron energies above 4 eV, while it is energetically inaccessible at incident electron energies below 4 eV. The onset of the most intense peak in the spectrum is around 3.5 eV, below the calculated reaction threshold for

simple C(4)–C(8) bond cleavage, and is therefore energetically inaccessible. Only when we consider further dissociation of the CONH₂ fragment into more stable CO and NH₂ molecules, the calculated threshold energy shifts below the experimentally observed threshold. Additionally, there is a peak with a maximum at around 2 eV, which is an experimental artefact, caused by the low selectivity of the QMS. The low energy peak is a contribution from neighboring intense fragment ion, [M–CONH₃][–], as can be read from its spectra discussed in the next paragraph.

The second most intense fragment ion is [M–CONH₃][–], whose spectrum is shown in Fig. 4, panel e). In certain configurations of the anion, the H(15) hydrogen of the OH group can be easily transferred to the amino group. Then, the formation of CO and NH₃ neutral fragments together with the C₄HFNO[–] anion is possible at energies as low as 0.86 eV, explaining the first peak in the ion yield. The second peak in the spectrum can be due to anions formed after H(12) removal, which requires more than 2 eV.

The spectrum of the third most intense fragment, OCN[–] is shown in panel f) of Fig. 4. Due to its high electron affinity (EA), OCN[–] is a common ion formed by DEA to many biologically relevant molecules in the gas phase,^{83–90} as well as clusters⁵⁸ and molecules deposited on surfaces.^{91,92} The high electron affinity of OCN (∼3.6 eV)⁹³ allows for very complex rearrangement and fragmentation reactions to be induced by the attachment of low-energy electrons.^{83,86} In the present case, the most straightforward pathway for OCN[–] extraction is the cleavage of C(4)–C(8) bond followed by the elimination of H₂ from the resulting CONH₂[–] anion. As we have seen already for the [M–CONH₂][–] channel the C(4)–C(8) bond is strong and therefore the reaction energies for such mechanism are above 4 eV. Within experimental and computational errors, the mechanisms can still explain all the peaks observed in the OCN[–] spectrum. However, we also found a more energetically favorable channel, which can be opened by hydrogen transfer to C(4) carbon allowing for N(7)C(5)O(6) anion extraction at energies below 2 eV together with a C₄H₄FN₂O neutral co-fragment.

Another intense fragmentation channel results in an ion corresponding to the neutral loss of 17 Da from the parent ion (Fig. 4, panel c)), which could correspond to either OH or NH₃ loss. The calculated reaction thresholds allow us to exclude the OH loss channel, as it occurs only at energies above 2.9 eV, higher than both observed peaks in the spectrum of this anion. On the other hand, the elimination of NH₃ has a threshold of only 0.17 eV, which is well below the onset of the observed signal, meaning that it is accessible.

The [M–H][–] ion yield is dominated by a near 0 eV peak, which is due to the low selectivity of QMS and overlap with the neighboring signal of the M[–] anion, which can not be fully separated in the present experiment. Energetically, this channel is only accessible above 0.64 eV. H(15) or H(13) hydrogen loss may occur at energies of the first real peak in the spectrum with a maximum at ∼1.8 eV and cleavage of H(12) is possible at the higher energies of the second peak with a maximum at ∼3.2 eV.

Another common biomolecular fragment ion with high electron affinity, CN[–] can be formed from the decomposition of the CONH₂ functional group or the ring of favipiravir. The energetic thresholds for both mechanisms are similar, at 1.85 eV and 1.97 eV respectively. In the first mechanism, a stable water molecule is formed as a neutral co-product. In the second mechanism, the C(1)–N(7) atoms are easiest to extract after hydrogen migration from C(1) to C(2). We also found several other possible reaction pathways that are possible for the observed CN[–] formation, with an onset around 4 eV (Fig. 4, panel g). In some cases, the reaction energy is lowered by closing the ring after the extraction of CN[–] anion forming the C₃N ring. Such rearrangement, however, may proceed over a reaction barrier, which can not be identified using the simple calculations present here.

Finally, with a relative intensity of only ∼0.1% that of the parent ion signal, we observe F[–] (Fig. 4, panel h)). The signal levels for this ion are near the detection limits of the setup, there appear to be three peaks in the spectrum, at ∼1 eV, ∼5 eV, and ∼9.5 eV. The calculated threshold energy for simple cleavage of the C(2)–F(11) bond is 2.15 eV, which explains the two higher energy resonances, but not the low energy resonance. This unassigned low-energy signal is possibly an experimental artefact, and is most likely caused by background ions in the experiment.

3.2 Hydration

To explore environmental effects, we measured EA to favipiravir in clusters with attached water molecules, prepared in a molecular beam using the CLUB experimental setup. Negative ion spectra of the molecule under dry and hydrated conditions taken in the energy range 0.6 to 8.6 eV is shown in Fig. 5. We can see that for the dry conditions, the intensity of the DEA channels is already reduced with respect to parent anion intensity. Using the CLUB apparatus, we measured reasonable signal only for the *m/z* = 112, 113 fragment anions. Let's compare the relative intensities of these anions to the parent anion in the two experiments. Dividing the integral intensity of the fragment *m/z* = 112 + *m/z* = 113 ion signal by that of *m/z* = 157 parent ion signal from the energy dependent ion yields in Fig. 4, we obtain the ratio of fragment to parent ion signal in the QMS-TEM setup of ∼1/10. Dividing the integral intensity of the the fragment *m/z* = 112 + *m/z* = 113 peak to that of *m/z* = 157 parent peak in the cumulative MS in Fig. 5, we obtain the ratio of fragment to parent ion signal in CLUB experiment at dry conditions to be 1/33, which is much lower. Additionally, the ratio 1/33 is only the highest estimate, since the electron current in CLUB experiment significantly drops in the near 0 eV region of the parent ion resonance, resulting in an apparent reduction of the parent ion signal.

The main reasons for suppression of fragmentation in the CLUB experiment under dry conditions is the detection time of only tens of microseconds, in comparison to the TEM-QMS experiment, where it is hundreds of microseconds. The much shorter detection timescale of the CLUB setup means that there is less time for fragmentation or autodetachment to occur.

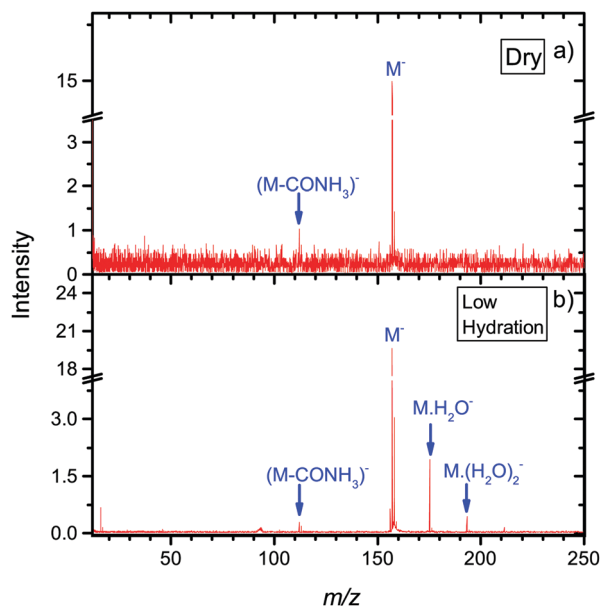


Fig. 5 Mass spectra from the CLUB experiment of anions formed after electron attachment to isolated or microhydrated neutral favipiravir in molecular beams. Negative ion MS are prepared by summing spectra taken with 0.2 eV step in the 0.6 eV to 8.6 eV range.

These effects of the detection time scale were studied in detail by Asfandiarov and co-workers.^{94,95} Another parameter influencing the fragmentation is the temperature of the precursor molecules, which is lower in the molecular beam of the CLUB than in the effusive beam of the TEM-QMS.⁹⁶

Further stabilization of parent anion with respect to both DEA and autodetachment can be induced by hydration. Under hydrated conditions the ratio of $m/z = 112, 113$ to parent anion signal further decreases to $\sim 1/90$.

Under hydrated conditions, anions are stabilized by energy transfer to the solvent. The total energy available for transfer to surrounding water molecules and their subsequent evaporation (TET) can be written as a sum of adiabatic electron affinity (AEA) and energy of the incoming electron (E_e):

$$\text{TET} = E_e + \text{AEA} \quad (3)$$

For halouracils, energy of several eV was estimated to be transferred to the solvent after electron attachment.²³ We also postulated that such energy transfer to the solvent can increase linear energy transfer (LET) value after interaction of the ionizing radiation with an environment containing electron affinic molecules and can explain their radiosensitizing action. The largest effect was previously observed for bromouracil, for which the number of evaporated water molecules after electron attachment was similar to that after electron ionization. In Fig. 6 we can see that the situation is also very similar for favipiravir. The figure shows a comparison of electron impact ionization MS of favipiravir at 70 eV and negative ion MS after electron attachment at 1.4 eV for two different hydration conditions. We can see that the observed numbers of water molecules, n , m , and p , attached to anions $M(\text{H}_2\text{O})_n^-$ and cations $M(\text{H}_2\text{O})_m^+$ and $M(\text{H}_2\text{O})_p^+$ are similar. Despite the well-known fragmentation efficiency of electron impact ionization,⁹⁷ the “soft” EA leads to a

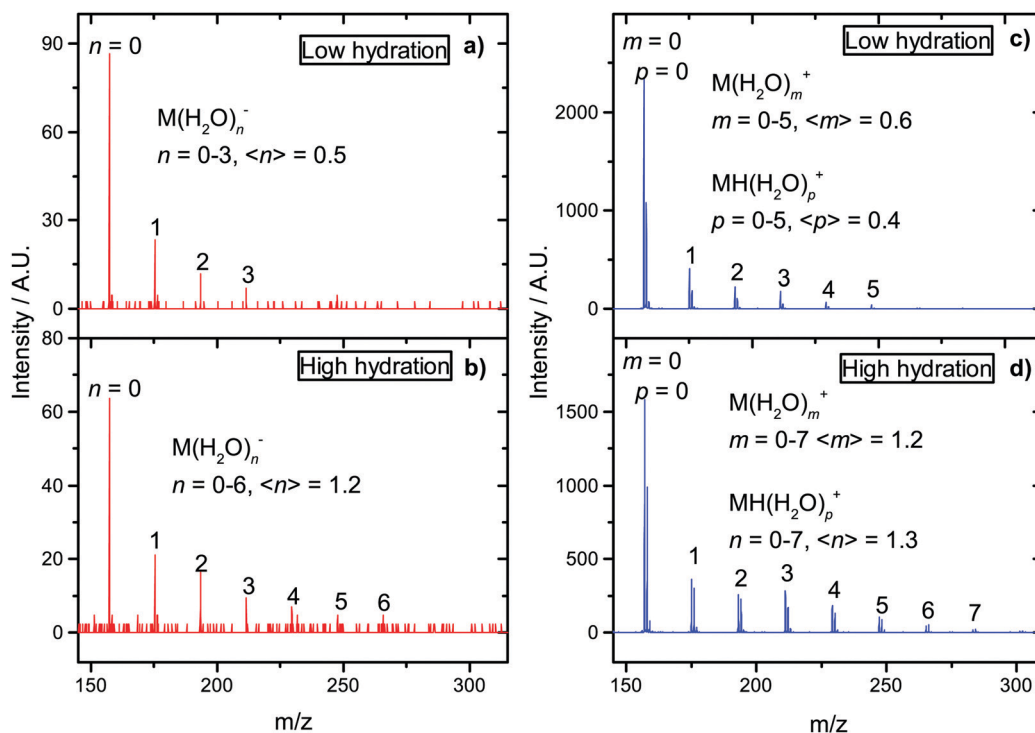


Fig. 6 Mass spectra from the CLUB experiment for hydrated favipiravir in molecular beams. Negative ion MS (a) and (b) at the electron-energy 1.4 eV and positive ion MS (c) and (d) at the electron-energy 70 eV. The values in brackets represents peak intensity weighted average.

similar fragmentation, in the form of loss of neutral water molecules from the clusters. This is caused by the high electron affinity of favipiravir, which results in water evaporation from cluster after electron attachment. Therefore, if our hypothesis that energy transfer to the solvent after EA enhances LET and causes radiosensitization is correct, favipiravir should demonstrate radiosensitizing effects comparable to halogenated uracils.

4 Conclusions

We present an experimental study of electron attachment to favipiravir as an interesting hetero-aromatic, important biochemical, and promising radiosensitizing molecule. Electron attachment to favipiravir leads primarily to the formation of the parent anion with a long lifetime. Such stability may be important for the transport and radiosensitizing properties of the molecule⁸ but may also allow for multiple electron reduction.⁹⁸ The threshold DFT calculations helped us to interpret the fragmentation pattern of the molecule with main fragmentation reactions occurring on the CONH₂ group. The estimation of vertical attachment energies demonstrates several possible virtual states available for attachment *via* shape resonances that may be related to favipiravir's antiviral action.¹⁰ The mass spectrometry of small clusters of the form Fav(H₂O)_n, reveals that a large amount of energy is transferred to the solvent after electron attachment. This energy may contribute to favipiravir's reactivity in its reduced form but also support our hypothesis that favipiravir may be repurposed as a radiosensitizer.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 S. A. Pshenichnyuk and A. Modelli, *J. Chem. Phys.*, 2014, **140**, 034313.
- 2 S. A. Pshenichnyuk and A. Modelli, in *ETS and DEAS Studies of the Reduction of Xenobiotics in Mitochondrial Intermembrane Space*, ed. V. Weissig and M. Edeas, Springer New York, New York, NY, 2015, pp. 285–305.
- 3 S. A. Pshenichnyuk and A. S. Komolov, *J. Phys. Chem. Lett.*, 2015, **6**, 1104–1110.
- 4 S. A. Pshenichnyuk, A. Modelli, E. F. Lazneva and A. S. Komolov, *J. Phys. Chem. A*, 2016, **120**, 2667–2676.
- 5 S. A. Pshenichnyuk, R. G. Rakhmeyer, N. L. Asfandiarov, A. S. Komolov, A. Modelli and D. Jones, *J. Phys. Chem. Lett.*, 2018, **9**, 2320–2325.
- 6 S. A. Pshenichnyuk, A. Modelli and A. S. Komolov, *Int. Rev. Phys. Chem.*, 2018, **37**, 125–170.
- 7 S. A. Pshenichnyuk and A. Modelli, in *Electron Attachment to Isolated Molecules as a Probe to Understand Mitochondrial Reductive Processes*, ed. V. Weissig and M. Edeas, Springer US, New York, NY, 2021, pp. 101–124.
- 8 R. Meißner, J. Kočíšek, L. Feketeová, J. Fedor, M. Fárník, P. Limão-Vieira, E. Illenberger and S. Denifl, *Nat. Commun.*, 2019, **10**, 2388.
- 9 C. W. Fong, Toxicology of platinum anticancer drugs: oxidative stress and antioxidant effect of stable free radical Pt-nitroxides, Eigenenergy, Adelaide, Australia research report, 2019.
- 10 H. M. Yasir and F. H. Hanoon, IOP Conference Series: Materials Science and Engineering, 2020, vol. 928, p. 072066.
- 11 R. Schürmann, S. Vogel, K. Ebel and I. Bald, *Chem. – Eur. J.*, 2018, **24**, 10271–10279.
- 12 Y. Dong, Y. Wang, P. Zhuang, X. Fu, Y. Zheng and L. Sanche, *J. Phys. Chem. B*, 2020, **124**, 3315–3325.
- 13 *Advances In Atomic, Molecular, and Optical Physics*, ed. E. Arimondo, C. C. Lin and S. F. Yelin, Academic Press, 2017, vol. 66, pp. 545–657.
- 14 J. D. Gorfinkiel and S. Ptasinska, *J. Phys. B: At., Mol. Opt. Phys.*, 2017, **50**, 182001.
- 15 S. M. Pimblott and J. A. LaVerne, *Radiat. Phys. Chem.*, 2007, **76**, 1244–1247.
- 16 E. Alizadeh and L. Sanche, *Chem. Rev.*, 2012, **112**, 5578–5602.
- 17 C. D. Willey, E. S.-H. Yang and J. A. Bonner, in *Clinical Radiation Oncology*, ed. L. L. Gunderson and J. E. Tepper, Elsevier, Philadelphia, 4th edn, 2016, pp. 63–79.e4.
- 18 P. Wardman, *Clin. Oncol.*, 2007, **19**, 397–417.
- 19 L. Chomicz, M. Zdrowowicz, F. Kasprzykowski, J. Rak, A. Buonaugurio, Y. Wang and K. H. Bowen, *J. Phys. Chem. Lett.*, 2013, **4**, 2853–2857.
- 20 J. Kopyra, C. Koenig-Lehmann, I. Bald and E. Illenberger, *Angew. Chem., Int. Ed.*, 2009, **48**, 7904–7907.
- 21 J. Rak, L. Chomicz, J. Wiczak, K. Westphal, M. Zdrowowicz, P. Wityk, M. Żyndul, S. Makurat and Ł. Golon, *J. Phys. Chem. B*, 2015, **119**, 8227–8238.
- 22 C. R. Wang, J. Nguyen and Q. B. Lu, *J. Am. Chem. Soc.*, 2009, **131**, 11320–11322.
- 23 J. Poštulka, P. Slaviček, J. Fedor, M. Fárník and J. Kočíšek, *J. Phys. Chem. B*, 2017, **121**, 8965–8974.
- 24 M. Rezaee, D. J. Hunting and L. Sanche, *Int. J. Radiat. Oncol., Biol., Phys.*, 2013, **87**, 847–853.
- 25 D. Reimitz, M. Davidková, O. Mestek, J. Pinkas and J. Kočíšek, *Radiat. Phys. Chem.*, 2017, **141**, 229–234.
- 26 Y. Dong, L. Zhou, Q. Tian, Y. Zheng and L. Sanche, *J. Phys. Chem. C*, 2017, **121**, 17505–17513.
- 27 C. Wagner and H.-A. Wagenknecht, *Chem. – Eur. J.*, 2005, **11**, 1871–1876.
- 28 F. Xiao, X. Luo, X. Fu and Y. Zheng, *J. Phys. Chem. B*, 2013, **117**, 4893–4900.

- 29 J. Rackwitz, M. L. Ranković, A. R. Milosavljević and I. Bald, *Eur. Phys. J. D*, 2017, **71**, 32.
- 30 J. Rackwitz, J. Kopyra, I. Dabkowska, K. Ebel, M. L. Ranković, A. R. Milosavljević and I. Bald, *Angew. Chem., Int. Ed.*, 2016, **55**, 10248–10252.
- 31 L. Chomicz, M. Zdrowowicz, F. Kasprzykowski, J. Rak, A. Buonaugurio, Y. Wang and K. H. Bowen, *J. Phys. Chem. Lett.*, 2013, **4**, 2853–2857.
- 32 M. Zdrowowicz, L. Chomicz, M. Żyndul, P. Wityk, J. Rak, T. J. Wiegand, C. G. Hanson, A. Adhikary and M. D. Sevilla, *Phys. Chem. Chem. Phys.*, 2015, **17**, 16907–16916.
- 33 J. Ameixa, E. Arthur-Baidoo, R. MeiÅŸner, S. Makurat, W. Kozak, K. Butowska, F. Ferreira da Silva, J. Rak and S. Denifl, *J. Chem. Phys.*, 2018, **149**, 164307.
- 34 R. MeiÅŸner, S. Makurat, W. Kozak, P. Limão-Vieira, J. Rak and S. Denifl, *J. Phys. Chem. B*, 2019, **123**, 1274–1282.
- 35 P. Spisz, M. Zdrowowicz, W. Kozak, L. Chomicz-Mańka, K. Falkiewicz, S. Makurat, A. Sikorski, D. Wyrzykowski, J. Rak, E. Arthur-Baidoo, P. Ziegler, M. S. Rodrigues Costa and S. Denifl, *J. Phys. Chem. B*, 2020, **124**, 5600–5613.
- 36 T. F. M. Luxford, S. A. Pshenichnyuk, N. L. Asfandiarov, T. Perečko, M. Falk and J. Kočišek, *Int. J. Mol. Sci.*, 2020, **21**, 8173.
- 37 S. Pushpakom, F. Iorio, P. A. Eysers, K. J. Escott, S. Hopper, A. Wells, A. Doig, T. Williams, J. Latimer, C. McNamee, A. Norris, P. Sanseau, D. Cavalla and M. Pirmohamed, *Nat. Rev. Drug Discovery*, 2019, **18**, 41–58.
- 38 S. H. Kim, J. H. Kim, A. Kolozsvary, S. L. Brown and S. O. Freytag, *J. Neuro-Oncol.*, 1997, **33**, 189–194.
- 39 L. Sleire, H. E. Førde, I. A. Netland, L. Leiss, B. S. Skeie and P. Øyvind Enger, *Pharmacol. Res.*, 2017, **124**, 74–91.
- 40 X. Zhang, F. Wang, C. Zhang, S. Wu, X. Zheng, T. Gong, R. Ding, K. Chen and D. Bai, *Inorg. Chem. Commun.*, 2018, **94**, 92–97.
- 41 Z. Zhang, L. Zhou, N. Xie, E. C. Nice, T. Zhang, Y. Cui and C. Huang, *Signal Transduction Targeted Ther.*, 2020, **5**, 113.
- 42 D. Bai, Y. Tian, K. Chen, X. Zhang, F. Wang, Y. Cheng, X. Zheng, K. Xiao and X. Dong, *Dyes Pigm.*, 2020, **182**, 108635.
- 43 S. Huq, J. Kedda, T. Zhao, R. Serra, A. Ding, M. Morales, J. Ehresman, H. Brem, G. L. Gallia, D. M. Sciubba and B. Tyler, *Neurosurgery*, 2020, **67**, 309.
- 44 M. Van Herp, H. Declerck and T. Decroo, *Lancet*, 2015, **385**, 2350.
- 45 K. Rosenke, H. Feldmann, J. Westover, P. W. Hanley, C. Martellaro, F. Feldmann, G. Saturday, J. Lovaglio, D. Scott, Y. Furuta, T. Komeno, B. Gowen and D. Safronetz, *Emerging Infectious Disease J.*, 2018, **24**, 1696.
- 46 Z. F. Udawadia, P. Singh, H. Barkate, S. Patil, S. Rangwala, A. Pendse, J. Kadam, W. Wu, C. F. Caracta and M. Tandon, *Int. J. Infect. Dis.*, 2021, **103**, 62–71.
- 47 K. Shiraki and T. Daikoku, *Pharmacol. Ther.*, 2020, **209**, 107512.
- 48 Y. Furuta, K. Takahashi, Y. Fukuda, M. Kuno, T. Kamiyama, K. Kozaki, N. Nomura, H. Egawa, S. Minami, Y. Watanabe, H. Narita and K. Shiraki, *Antimicrob. Agents Chemother.*, 2002, **46**, 977–981.
- 49 Y.-X. Du and X.-P. Chen, *Clin. Pharm. Ther.*, 2020, **108**, 242–247.
- 50 M. G. Ison and M. H. Scheetz, *EBioMedicine*, 2021, **63**, 103204.
- 51 J. Langer, M. Zawadzki, M. Fárník, J. Pinkas, J. Fedor and J. Kočišek, *Eur. Phys. J. D*, 2018, **72**, 112.
- 52 M. Fárník and J. Lengyel, *Mass Spectrom. Rev.*, 2018, **37**, 630–651.
- 53 M. Stepanović, Y. Pariat and M. Allan, *J. Chem. Phys.*, 1999, **110**, 11376–11382.
- 54 A. Stamatovic and G. J. Schulz, *Rev. Sci. Instrum.*, 1970, **41**, 423–427.
- 55 L. G. Christophorou and J. K. Olthoff, *J. Phys. Chem. Ref. Data*, 2000, **29**, 267–330.
- 56 M. Zawadzki, T. F. M. Luxford and J. Kočišek, *J. Phys. Chem. A*, 2020, **124**, 9427–9435.
- 57 R. Dressler and M. Allan, *Chem. Phys.*, 1985, **92**, 449–455.
- 58 J. Kočišek, K. Grygoryeva, J. Lengyel, M. Fárník and J. Fedor, *Eur. Phys. J. D*, 2016, **70**, 98.
- 59 J. Kočišek, J. Lengyel and M. Fárník, *J. Chem. Phys.*, 2013, **138**, 124306.
- 60 J. Kočišek, A. Pysanenko, M. Fárník and J. Fedor, *J. Phys. Chem. Lett.*, 2016, **7**, 3401–3405.
- 61 L. A. Curtiss, K. Raghavachari and J. A. Pople, *J. Chem. Phys.*, 1993, **98**, 1293–1298.
- 62 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 63 Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215–241.
- 64 K. Aflatoon, G. A. Gallup and P. D. Burrow, *J. Phys. Chem. A*, 1998, **102**, 6205–6207.
- 65 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian ~16 Revision C.01*, Gaussian Inc., Wallingford CT, 2016.
- 66 G. Zhurko, *Chemcraft – Graphical Program for Visualization of Quantum Chemistry Computations*, available online: <https://chemcraftprog.com>.
- 67 L. Antonov, *Theor. Chem. Acc.*, 2020, **139**, 145.
- 68 J. Simons, *Annu. Rev. Phys. Chem.*, 2011, **62**, 107–128.
- 69 H. Hao, J. Shee, S. Upadhyay, C. Ataca, K. D. Jordan and B. M. Rubenstein, *J. Phys. Chem. Lett.*, 2018, **9**, 6185–6190.

- 70 L. Sanche and G. J. Schulz, *Phys. Rev. A*, 1972, **5**, 1672–1683.
- 71 D. Spence, *J. Chem. Phys.*, 1977, **66**, 669–674.
- 72 M. Zawadzki, M. Ranković, J. Kočišek and J. Fedor, *Phys. Chem. Chem. Phys.*, 2018, **20**, 6838–6844.
- 73 V. S. Prabhudesai, A. H. Kelkar, D. Nandi and E. Krishnakumar, *Phys. Rev. Lett.*, 2005, **95**, 143202.
- 74 C. Fridh, L. Åsbrink, B. Jonsson and E. Lindholm, *Int. J. Mass Spectrom. Ion Phys.*, 1972, **8**, 101–118.
- 75 M. V. Muftakhov and A. I. Fokin, *Rapid Commun. Mass Spectrom.*, 1997, **11**, 1923–1925.
- 76 J. Gingell, N. Mason, H. Zhao, I. Walker and M. Siggel, *Chem. Phys.*, 1997, **220**, 191–205.
- 77 Z. Li, M. Ryszka, M. M. Dawley, I. Carmichael, K. B. Bravaya and S. Ptasińska, *Phys. Rev. Lett.*, 2019, **122**, 073002.
- 78 J. Fedor, *Phys. Rev. Lett.*, 2020, **124**, 199301.
- 79 R. Abouaf and H. Dunet, *Eur. Phys. J. D*, 2005, **35**, 405–410.
- 80 P. D. Burrow, G. A. Gallup, A. M. Scheer, S. Denifl, S. Ptasińska, T. Märk and P. Scheier, *J. Chem. Phys.*, 2006, **124**, 124310.
- 81 M. Allan, M. Lacko, P. Papp, Š. Matejčík, M. Zlatar, I. I. Fabrikant, J. Kočišek and J. Fedor, *Phys. Chem. Chem. Phys.*, 2018, **20**, 11692–11701.
- 82 I. Nenner and G. J. Schulz, *J. Chem. Phys.*, 1975, **62**, 1747–1758.
- 83 C. Koenig-Lehmann, J. Kopyra, I. Dabkowska, J. Kočišek and E. Illenberger, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6954–6961.
- 84 T. Hamann, A. Edtbauer, F. Ferreira da Silva, S. Denifl, P. Scheier and P. Swiderek, *Phys. Chem. Chem. Phys.*, 2011, **13**, 12305–12313.
- 85 A. Keller, J. Kopyra, K. V. Gothelf and I. Bald, *New J. Phys.*, 2013, **15**, 083045.
- 86 F. Ferreira da Silva, C. Matias, D. Almeida, G. García, O. Ingólfsson, H. D. Flosadóttir, B. Ómarsson, S. Ptasińska, B. Puschnigg, P. Scheier, P. Limão-Vieira and S. Denifl, *J. Am. Soc. Mass Spectrom.*, 2013, **24**, 1787–1797.
- 87 M. M. Dawley and S. Ptasińska, *Int. J. Mass Spectrom.*, 2014, **365–366**, 143–151.
- 88 M. M. Dawley, K. Tanzer, I. Carmichael, S. Denifl and S. Ptasińska, *J. Chem. Phys.*, 2015, **142**, 215101.
- 89 M. Muftakhov, P. Shchukin and R. Khatymov, *Radiat. Phys. Chem.*, 2021, **184**, 109464.
- 90 M. V. Muftakhov and P. V. Shchukin, *Rapid Commun. Mass Spectrom.*, 2019, **33**, 482–490.
- 91 M. M. Dawley, C. Pirim and T. M. Orlando, *J. Phys. Chem. A*, 2014, **118**, 1228–1236.
- 92 H. Abdoul-Carime, P. Cloutier and L. Sanche, *Radiat. Res.*, 2001, **155**, 625–633.
- 93 S. E. Bradforth, E. H. Kim, D. W. Arnold and D. M. Neumark, *J. Chem. Phys.*, 1993, **98**, 800–810.
- 94 N. L. Asfandiarov, S. A. Pshenichnyuk, R. G. Rakhmeyer, R. F. Tuktarov, N. L. Zaitsev, A. S. Vorobev, J. Kočišek, J. Fedor and A. Modelli, *J. Chem. Phys.*, 2019, **150**, 114304.
- 95 N. L. Asfandiarov, M. V. Muftakhov, S. A. Pshenichnyuk, P. Papp, M. Danko, M. Lacko, J. Blaško, Š. Matejčík and A. Modelli, *J. Chem. Phys.*, 2017, **147**, 234302.
- 96 H. Pauly, *Atomic and Molecular Beam Methods*, vol. **1**, 1998.
- 97 J. Lengyel, A. Pysanenko, V. Poterya, J. Kočišek and M. Fárnik, *Chem. Phys. Lett.*, 2014, **612**, 256–261.
- 98 R. Schürmann, T. F. M. Luxford, I. S. Vinklársek, J. Kočišek, M. Zawadzki and I. Bald, *J. Chem. Phys.*, 2020, **153**, 104303.