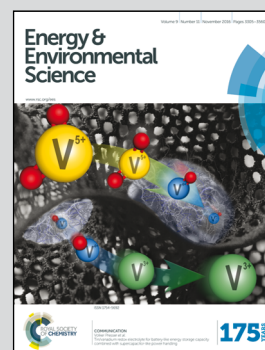


Showcasing the research on thermal decomposition of $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite by Dr Emilio J. Juarez-Perez, Mr Zafer Hawash, Dr Sonia R. Raga, Dr Luis K. Ono, and Prof. Yabing Qi at the Energy Materials and Surface Sciences Unit (EMSS) in the Okinawa Institute of Science and Technology Graduate University (OIST).

Thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite into NH_3 and CH_3I gases observed by coupled thermogravimetry–mass spectrometry analysis

Thermal gravimetric and differential thermal analysis (TG-DTA) coupled with quadrupole mass spectrometry (MS) instrumentation and first principles calculations were employed to elucidate the chemical nature of released gases during the thermal decomposition of $\text{CH}_3\text{NH}_3\text{PbI}_3$. Contrarily to the common wisdom that $\text{CH}_3\text{NH}_3\text{PbI}_3$ is decomposed into CH_3NH_2 and HI, the major gases released during decomposition are CH_3I and NH_3 .

As featured in:



See Yabing Qi et al.,
Energy Environ. Sci., 2016, 9, 3406.



Cite this: *Energy Environ. Sci.*,
2016, 9, 3406

Received 13th July 2016,
Accepted 1st September 2016

DOI: 10.1039/c6ee02016j

www.rsc.org/ees

Thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite into NH_3 and CH_3I gases observed by coupled thermogravimetry–mass spectrometry analysis†

Emilio J. Juarez-Perez, Zafer Hawash, Sonia R. Raga, Luis K. Ono and Yabing Qi*

Thermal gravimetric and differential thermal analysis (TG-DTA) coupled with quadrupole mass spectrometry (MS) and first principles calculations were employed to elucidate the chemical nature of released gases during the thermal decomposition of $\text{CH}_3\text{NH}_3\text{PbI}_3$. In contrast to the common wisdom that $\text{CH}_3\text{NH}_3\text{PbI}_3$ is decomposed into CH_3NH_2 and HI, the major gases were methyl iodide (CH_3I) and ammonia (NH_3). We anticipate that our findings will provide new insights into further formulations of the perovskite active material and device design that can prevent methylammonium decomposition and thus increase the long-term stability of perovskite-based optoelectronic devices.

Introduction

Photovoltaic technologies based on hybrid organometal halide perovskites have already achieved outstanding efficiencies.¹ The main challenge now is to overcome their limited lifetime under operating conditions.² The degradation mechanisms of perovskites have been a topic under intense investigation.^{3–6} The fundamental understanding of physicochemical processes that trigger degradation is needed to further engineer perovskite solar cells for enhanced stability. Generally speaking, extrinsic factors (*e.g.*, light, temperature, humidity, and O_2) can contribute to the degradation processes. Different degradation pathways have been proposed and the topic is still under debate. But it is clear that $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite loses its excellent light harvesting properties over time because it evolves into PbI_2 after a nominal loss of $\text{CH}_3\text{NH}_3\text{I}$.⁷ The degradation from $\text{CH}_3\text{NH}_3\text{PbI}_3$ to PbI_2 is most likely accompanied by a release of gases *via* simple sublimation or assisted chemical reaction.

Energy Materials and Surface Sciences Unit (EMSS), Okinawa Institute of Science and Technology Graduate University (OIST), 1919-1 Tancha, Kunigami-gun, Onna-son, Okinawa 904-0495, Japan. E-mail: Yabing.Qi@OIST.jp

† Electronic supplementary information (ESI) available: Details for synthesis of $\text{CH}_3\text{NH}_3\text{PbI}_3$ single crystals, TG/DTA-MS measurements, and DFT calculations, supplementary figures and table. See DOI: 10.1039/c6ee02016j

Broader context

In the past few years, we have all witnessed the unprecedented zeal for the rapid growth of the organometal halide perovskite solar cell field. The tremendous efforts devoted to device fabrication and optimization have led to PCEs exceeding 20%, which gives perovskite solar cells a competitive advantage over many other well-known solar technologies. On the other hand, some challenges remain with the lifetime as the key. At present, even the most long-lasting perovskite solar cells can only retain stable PCEs up to a few thousands of hours, which is way behind, for example, silicon cells. To further improve perovskite solar cells, we must overcome/mitigate the degradation issue. For this reason, a thorough detailed understanding of perovskite solar cell degradation processes becomes an imperative first step.

Therefore, the determination of the composition of these released gases during controlled thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ is expected to pinpoint the degradation pathways. A suitable method to identify the mass loss directly from perovskite degradation is the thermal gravimetric and differential thermal analysis (TG-DTA). Using TG-DTA, we can detect the endothermic/exothermic phase transitions and also quantify the percentage of mass loss during the heating of the material. However, this technique does not characterize the chemical nature of the released gases. For that reason, we connected a mass spectrometer (MS) to the gas exhaust line from the TG-DTA equipment in order to identify unambiguously the molecular mass of the released gases. Typically, the TG trace for $\text{CH}_3\text{NH}_3\text{PbI}_3$ shows two consecutive mass loss steps, with the former ascribed to the nominal loss of methylammonium iodide $\text{CH}_3\text{NH}_3\text{I}$ (*ca.* ~25%, 300–420 °C) and the latter to the inorganic PbI_2 component (~75% above 420 °C).^{8–16} A careful comparison of the TG trace during the first mass loss step in $\text{CH}_3\text{NH}_3\text{PbI}_3$ and the TG trace of the $\text{CH}_3\text{NH}_3\text{I}$ salt sublimation shows that the former is at least composed of two intermediate mass loss steps evolving at different rates; see the portion of TG-DTA traces indicated by blue arrows in Fig. 1A and B.

Different interpretations can be found in the literature regarding the chemical nature of these released gases during



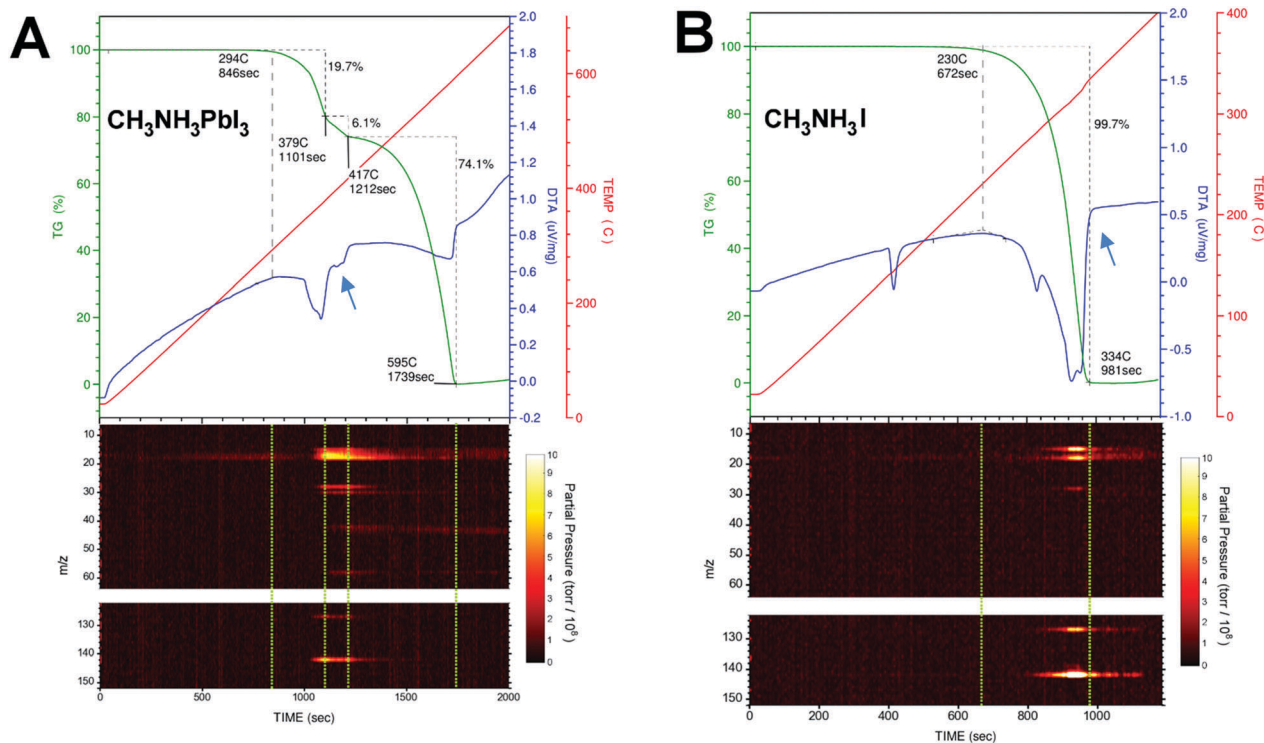


Fig. 1 TG-DTA traces (upper panel, dark green and blue color, respectively) and the m/z peaks registered simultaneously during the thermal degradation (heating rate of $20\text{ }^{\circ}\text{C min}^{-1}$) of (A) $\text{CH}_3\text{NH}_3\text{PbI}_3$ and (B) $\text{CH}_3\text{NH}_3\text{I}$. Blue arrows indicate an endothermic process present in the DTA trace in $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite, but absent in the $\text{CH}_3\text{NH}_3\text{I}$ salt. Dashed green lines in the m/z vs. time contour plots are guides to the eye delimiting the mass loss steps in the TG/DTA graph.

thermal degradation. Dualeh *et al.*¹¹ and Nenon *et al.*¹³ proposed that CH_3NH_2 and HI are the gases released sequentially¹¹ during the first mass loss step of $\text{CH}_3\text{NH}_3\text{PbI}_3$ decomposition. On the other hand, Williams *et al.*¹⁵ found that the gases released are CH_3I and NH_3 . In the study by Williams *et al.*, the determination of CH_3I and NH_3 was clearly demonstrated for $\text{CH}_3\text{NH}_3\text{I}$, but it is ambiguous for the perovskite sample because of the limitation of the methodology employed. They performed the simultaneous thermal analysis-Fourier Transform infrared spectroscopy (STA-FTIR) measurements on samples of perovskite precursor solution. Therefore, the reported infrared spectra of the gases released during the perovskite formation/degradation processes are convoluted with the thermal degradation of the solvent, and the assignment of CH_3I and NH_3 is ambiguous in this case. The assignment of degradation gas species can get complicated by the use of perovskite solutions instead of solid samples (*e.g.*, small perovskite crystals used in this work). For example, a close inspection of Fig. 4a in the report by Nenon *et al.*¹³ reveals that there is no 31 m/z peak for methylamine in the “Degraded” zone. Peak 31 m/z only appears during perovskite formation and drying steps. Also, their MS scan reached only up to 100 m/z leaving out of sight the 142 m/z CH_3I signature peak.

It is noteworthy that the presence of CH_3I molecules in the early stage of formation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ using vacuum deposition was proposed by Liu *et al.* in their X-ray photoelectron spectroscopy (XPS) study.¹⁷ Also, a recent study by Xu *et al.* provided

further evidence using soft X-ray spectroscopy and density functional theory calculations.¹⁸

Results and discussion

Simultaneous thermogravimetry (TG) and differential thermal analysis (DTA) coupled with mass spectrometry (TG-DTA/MS) measurements were carried out under an inert atmosphere (He) to identify the chemical compositions of the released gases during the thermal decomposition process of $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite monocrystals (volume $\sim 1\text{ mm}^3$, weight of $\sim 7\text{--}15\text{ mg}$) and $\text{CH}_3\text{NH}_3\text{I}$ polycrystalline powder. The gases generated during the TG-DTA measurements are collected using a quadrupole mass filter, which is employed primarily for residual gas analysis (RGA). In this work, the starting and end points of the mass loss steps are delimited using the slope changes caused by thermal events in the DTA trace, which is recorded simultaneously during the TG measurements. Fig. 1A and B show the TG-DTA traces and the mass-to-charge ratio (m/z) peaks recorded simultaneously during the thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ and $\text{CH}_3\text{NH}_3\text{I}$. For simplicity, only the relevant peaks in the m/z vs. time contour plots in Fig. 1 are shown for discussion. The complete MS results can be found in Fig. S1 (ESI[†]). Table 1 summarizes the m/z peaks observed during the thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ and $\text{CH}_3\text{NH}_3\text{I}$ together with their assignments. Dashed green lines in the m/z vs. time contour plots are guide to the eye, delimiting the mass loss steps in the TG-DTA graph.



Table 1 Assignments of parent and fragmentation peaks in the TG-DTA/MS spectra

<i>m/z</i>	Detected ion	Parent/fragment molecule
15	CH ₃ ⁺	CH ₃ I fragment
16	NH ₂ ⁺	NH ₃ fragment
17	NH ₃ ⁺	NH ₃ parent peak
18	NH ₄ ⁺	NH ₃ + H ⁺ proton transfer
18	H ₂ O ⁺	H ₂ O parent peak ²¹
28	N ₂ ⁺	N ₂ parent peak
30	CH ₄ N ⁺	(CH ₃) ₃ N fragment
42	C ₂ H ₄ N ⁺	(CH ₃) ₃ N fragment
44	C ₂ H ₆ N ⁺	(CH ₃) ₃ N fragment
58	C ₃ H ₈ N ⁺	(CH ₃) ₃ N Hydrogen dissociation
59	(CH ₃) ₃ N ⁺	(CH ₃) ₃ N parent peak
127	I ⁺	CH ₃ I fragment
141	CH ₂ I ⁺	CH ₃ I Hydrogen dissociation
142	CH ₃ I ⁺	CH ₃ I parent peak

Contrary to common wisdom, CH₃NH₂ (methylamine, $T_{bp} = -6$ °C, $m/z = 31$) and HI (hydrogen iodide, $T_{bp} = -34$ °C, $m/z = 128$) gases were not detected during the thermal degradation of CH₃NH₃PbI₃ or CH₃NH₃I. Although the $m/z = 30$ and 127 fragment peaks were observed, their corresponding parent molecules of $m/z = 31$ and 128 amu were not detected; see Fig. 1, Fig. S2 and S3 (ESI[†]). When performing measurements with MS systems care should be taken to consider the fragmentation of molecules during the ionization process (electron bombardment) within the MS system. As the control experiment, commercially available CH₃NH₂ and HI gases were fed directly into our MS system and $m/z = 31$ and 128 were indeed detected (Fig. S4a and b, ESI[†]) in addition to the $m/z = 30$ and 127 fragment peaks. In the thermal degradation process of CH₃NH₃PbI₃ and CH₃NH₃I, because the parent molecular peaks corresponding to $m/z = 31$ and 128 peaks were missing, the amounts of CH₃NH₂ and HI gases can be concluded to be negligible. Instead, the m/z peaks observed in our TG-DTA/MS measurements can be unambiguously assigned to NH₃ (ammonia gas, $T_{bp} = -33$ °C, $m/z = 17$) and CH₃I (methyl iodide, $T_{bp} = 42$ °C, $m/z = 142$) gases; see Fig. 1, Fig. S2, S3 (ESI[†]) and Table 1. It is noteworthy to mention that Nenon *et al.*¹³ performed MS in the range from $m/z = 1$ to 100, so they could not observe the m/z peaks at 127, 141, and 142, the latter two of which correspond to CH₃I.

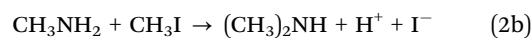
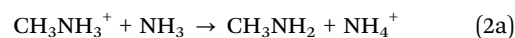
Based on the observations above, it is proposed that the first mass loss step during the thermal degradation of CH₃NH₃PbI₃ and CH₃NH₃I under an inert atmosphere proceeds as:



Therefore, the decomposition reactions of the methylammonium ions (CH₃NH₃⁺) assisted by iodide proceed through the so-called reverse Menshutkin reaction (a bimolecular nucleophilic substitution S_N2 simultaneously breaking a C–N bond and forming a C–I bond) as it is well described for quaternary ammonium salts.^{19,20} But, in contrast to the quaternary ammonium salts, the decomposition product of methylammonium *via* reverse Menshutkin reaction forms ammonia instead of trimethylamine.

The high concentrations of NH₄⁺ ($m/z = 18$), N₂ ($m/z = 28$) and H₂, ($m/z = 2$), which were also detected by our MS system, indicate that NH₃ undergoes further decomposition and rearrangement reactions (Fig. S2 and S3, ESI[†]). Interestingly, during the CH₃NH₃PbI₃ thermal degradation, unidentified traces can be detected only at the end of the first mass loss release (Fig. 1a and Fig. S2, ESI[†]). The m/z fragments at 30, 42, 44, 58, and 59 were tentatively identified as the fragmentation pattern of trimethylamine (CH₃)₃N ($T_{bp} = 3$ °C, $m/z = 59$, see Fig. S5, ESI[†]), which corroborates with the proposed pathway (eqn (2c)). On the other hand, we did not observe intermediates such as CH₃NH₂ (eqn (2a)) and (CH₃)₂NH (eqn (2b)), possibly due to the detection limit or the fact that the kinetics of the chain reaction is too fast to follow and we could only observe the final products (*e.g.*, trimethylamine).

Therefore, the proposed mechanism for the formation of (CH₃)₃N is as follows:



Here we propose that the previously formed NH₃ can react with the remaining methylammonium ions trapped in the PbI₃[−] octahedral network *via* a simple acid–base reaction (eqn (2a)), and the excess of CH₃I reacts with the as-formed primary amines (eqn (2b)) and secondary amines (eqn (2c)) *via* the conventional Menshutkin reaction until the formation of the tertiary amine (CH₃)₃N that ends the chain reaction set. We hypothesize that the (2a) reaction needs to proceed with trapped methylammonium ions inside the bulk perovskite, which would explain why these trace amounts of (CH₃)₃N are not observed during the thermal degradation of CH₃NH₃I.

Photovoltaic devices usually operate under light irradiation and, most likely, with moisture and oxygen invasion. Therefore, the chemical stability of perovskite under light irradiation and ambient conditions was studied using a modified TG-DTA/MS setup; see Fig. S9a (ESI[†]). Overall, the main difference observed when using an oxygen or water rich carrier gas during the TG-DTA/MS measurements was a different TG trace during the high temperature interval time. In all experiments performed with such oxidizing agents in the carrier gas, the PbI₂ is partially converted to lead oxide (Pb_xO_y) ashes,²² which remain intact after the measurement is finished; see Fig. S9, panels b, c, e and f (ESI[†]). On the other hand, the common feature is that the gases released during the thermal decomposition of CH₃NH₃PbI₃ above 300 °C are still ammonia and methyl iodide. Specifically, the 142 m/z signature peak for methyl iodide release is clearly observed for all cases. The detection of the CH₃I release at low temperature using our TG-DTA/MS setup equipment is non-trivial because such a low release rate of CH₃I will be below the detection limit if using the same measurement conditions as before. Therefore, we have performed a specifically designed measurement to obtain the experimental results confirming that the proposed decomposition pattern also takes place at



low temperatures such as 80 °C; see Section S3 in the ESI† for full details.

First-principles density functional theory (DFT) calculations using second order perturbation correction (Double-Hybrid Functional, see the ESI† files for full details) were carried out to gain further insight into the thermochemistry of the two chemical reaction pathways suggested for the perovskite degradation, represented by chemical eqn (3a) and (3b):



ΔG (Gibbs free enthalpy) evaluation in the temperature range of 25–600 °C reveals that both reactions are thermodynamically spontaneous and reaction (3a) shows ~ 13 –14 kcal mol⁻¹ in ΔG lower than reaction (3b). Therefore, both degradation processes are favorable thermodynamically, but it does not explain why eqn (3a) is kinetically favored compared to (3b). A description of the reaction mechanism is therefore needed, in which the molecular structures of reactants, intermediates and products are correlated with their energy reaction pathways. In this case, the proposed thermodynamic cycle describing the degradation mechanism consists of 3 processes: (1) reactants infinitely separated are brought together to one specific interacting state; (2) chemical transformation of these reactants to molecular products by means of a transition state (TS); and (3) infinite distance separation of the products. Fig. 2 depicts this hypothetical thermodynamic cycle for the ((3a), blue) and ((3b), red) reaction pathways. Based on our calculations, the interacting disposition of reactants as depicted in Fig. 2, reaction (3b) (state 6), is favored

by ~ 30 kcal mol⁻¹ compared to reaction (3a) (state 2). However, a TS (state 3) that connects reactants (state 2) and products (state 4) can only be obtained for the reaction (3a) pathway (three short movie files showing the relaxed scan calculations along the reaction coordinate for both chemical reactions and the imaginary frequency oscillation for the TS state 3 are given in the ESI†). The activation energy (enthalpy difference between state 3 and state 2) is only 0.4–0.2 kcal mol⁻¹ in the temperature range studied. On the other hand, there are no stationary ground state geometrical dispositions for HI and CH₃NH₂ molecules where such molecules do not react evolving to state 6. Consequently, a TS for the reaction (3b) pathway cannot be found. Therefore, the absolute lowest energy state for the molecular gas system is defined as CH₃I and NH₃ molecular products infinitely separated (state 5). State 6, however, remains as a local minimum in enthalpy energy but in the reactant side of the (3b) chemical path. Only this state 6 could produce directly the degradation products (HI and CH₃NH₂) (state 7) going up the hill in enthalpy energy. In summary, these first-principles calculations support the experimental results, *i.e.* the molecular ionic pair (CH₃NH₃⁺ and I⁻) decomposes to energetically favored thermal degradation products (CH₃I and NH₃) in strong contrast to the previously suggested decomposition pathway to CH₃NH₂ and HI. However, it is not possible to completely rule out that under certain experimental conditions different from what was used in this experimental/theoretical study, the favored pathway would be decomposition to CH₃NH₂ and HI. Also, these first-principles DFT calculations are carried out using state 1 as the initial state of the system and studying its evolution until obtaining the degradation products (state 5 or 7).

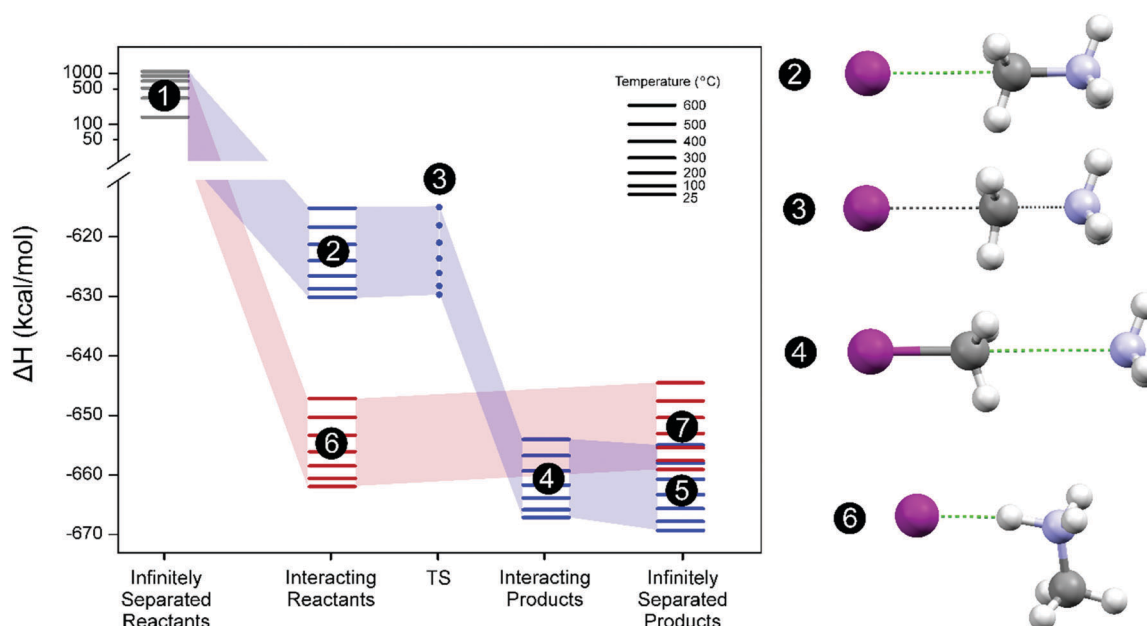


Fig. 2 Left panel: Thermodynamic cycle depicting the decomposition process of CH₃NH₃⁺ assisted by I⁻ for the chemical reaction pathways producing CH₃I and NH₃ (eqn (3a), blue) and CH₃NH₂ and HI (eqn (3b), red). Enthalpy energy profiles with reference to the infinitely separated reactant system (CH₃NH₃⁺ + I⁻) at 25 °C, $\Delta H^\circ = 0$. Colored lines mean optimized ground state system and dot transition state (TS). Right panel: Optimized molecular geometries for states 2–4 and 6. White color sphere: H; purple: I; grey: C; cyan: N. Inset: Temperature range for the calculated thermochemical variables of the state.



Deliberately, we have left out of this theoretical study state 0 of the system (the solid crystal phase for $\text{CH}_3\text{NH}_3\text{I}$ or $\text{CH}_3\text{NH}_3\text{PbI}_3$) because it is out of the scope and it is unnecessary to explain the evolution of the molecular gas system. However, we have included in the ESI† an estimation of such crystallization enthalpy from state 0 to state 1.

Conclusions

In summary, by employing coupled TG-DTA and MS measurements, we demonstrate experimentally that the thermal degradation of $\text{CH}_3\text{NH}_3\text{PbI}_3$ and $\text{CH}_3\text{NH}_3\text{I}$ releases NH_3 and CH_3I as degradation products. Also, first-principles DFT calculations support the experimental results and show that only such degradation gas release can be allowed by a thermo- and kinetically favored chemical path of reaction in the temperature range of 20–600 °C and under atmospheric pressure. This result is in sharp contrast with the common wisdom. Our findings may serve as the first cornerstone in the way to obtaining long-term stabilized perovskite based photovoltaic devices.

Acknowledgements

This work was supported by funding from the Energy Materials and Surface Sciences Unit of the Okinawa Institute of Science and Technology Graduate University and JSPS KAKENHI Grant Number 15K17925. We thank Steven D. Aird, the Technical Editor at the Okinawa Institute of Science and Technology Graduate University, for valuable suggestions in revising the manuscript.

Notes and references

- 1 N. R. E. L. (NREL), Best Research-Cell Efficiencies, http://www.nrel.gov/ncpv/images/efficiency_chart.jpg, Accessed 16 Mar, 2016.
- 2 X. Li, M. Tschumi, H. Han, S. S. Babkair, R. A. Alzubaydi, A. A. Ansari, S. S. Habib, M. K. Nazeeruddin, S. M. Zakeeruddin and M. Gratzel, *Energy Technol.*, 2015, **3**, 551–555.
- 3 N. Aristidou, I. Sanchez-Molina, T. Chotchuangchutchaval, M. Brown, L. Martinez, T. Rath and S. A. Haque, *Angew. Chem., Int. Ed.*, 2015, **54**, 8208–8212.
- 4 D. Bryant, N. Aristidou, S. Pont, I. Sanchez-Molina, T. Chotchuangchutchaval, S. Wheeler, J. R. Durrant and S. A. Haque, *Energy Environ. Sci.*, 2016, **9**, 1655–1660.
- 5 A. D. Sheikh, A. Bera, M. A. Haque, R. B. Rakhi, S. D. Gobbo, H. N. Alshareef and T. Wu, *Sol. Energy Mater. Sol. Cells*, 2015, **137**, 6–14.
- 6 R. K. Misra, S. Aharon, B. Li, D. Mogilyansky, I. Visoly-Fisher, L. Etgar and E. A. Katz, *J. Phys. Chem. Lett.*, 2015, **6**, 326–330.
- 7 T. A. Berhe, W.-N. Su, C.-H. Chen, C.-J. Pan, J.-H. Cheng, H.-M. Chen, M.-C. Tsai, L.-Y. Chen, A. A. Dubale and B.-J. Hwang, *Energy Environ. Sci.*, 2016, **9**, 323–356.
- 8 T. Baikie, Y. Fang, J. M. Kadro, M. Schreyer, F. Wei, S. G. Mhaisalkar, M. Graetzel and T. J. White, *J. Mater. Chem. A*, 2013, **1**, 5628–5641.
- 9 Y. Dang, Y. Liu, Y. Sun, D. Yuan, X. Liu, W. Lu, G. Liu, H. Xia and X. Tao, *CrystEngComm*, 2015, **17**, 665–670.
- 10 L. Dimesso, M. Dimamay, M. Hamburger and W. Jaegermann, *Chem. Mater.*, 2014, **26**, 6762–6770.
- 11 A. Dualeh, P. Gao, S. I. Seok, M. K. Nazeeruddin and M. Gratzel, *Chem. Mater.*, 2014, **26**, 6160–6164.
- 12 J.-H. Im, C.-R. Lee, J.-W. Lee, S.-W. Park and N.-G. Park, *Nanoscale*, 2011, **3**, 4088–4093.
- 13 D. P. Nenon, J. A. Christians, L. M. Wheeler, J. L. Blackburn, E. M. Sanehira, B. Dou, M. L. Olsen, K. Zhu, J. J. Berry and J. M. Luther, *Energy Environ. Sci.*, 2016, **9**, 2072–2082.
- 14 B. Wang, K. Young Wong, X. Xiao and T. Chen, *Sci. Rep.*, 2015, **5**, 10557.
- 15 A. E. Williams, P. J. Holliman, M. J. Carnie, M. L. Davies, D. A. Worsley and T. M. Watson, *J. Mater. Chem. A*, 2014, **2**, 19338–19346.
- 16 W. Zhang, M. Saliba, D. T. Moore, S. K. Pathak, M. T. Horantner, T. Stergiopoulos, S. D. Stranks, G. E. Eperon, J. A. Alexander-Webber and A. Abate, et al., *Nat. Commun.*, 2015, **6**, 6142.
- 17 L. Liu, J. A. McLeod, R. Wang, P. Shen and S. Duhm, *Appl. Phys. Lett.*, 2015, **107**, 061904.
- 18 W. Xu, L. Liu, L. Yang, P. Shen, B. Sun and J. A. McLeod, *Nano Lett.*, 2016, **16**, 4720–4725.
- 19 I. Yushina, B. Rudakov, I. Krivtsov and E. Bartashevich, *J. Therm. Anal. Calorim.*, 2014, **118**, 425–429.
- 20 M. Sawicka, P. Storoniak, P. Skurski, J. Błażejowski and J. Rak, *Chem. Phys.*, 2006, **324**, 425–437.
- 21 C. Muller, T. Glaser, M. Plogmeyer, M. Sendner, S. Doring, A. A. Bakulin, C. Brzuska, R. Scheer, M. S. Pshenichnikov and W. Kowalsky, et al., *Chem. Mater.*, 2015, **27**, 7835–7841.
- 22 A. L. Roe, K. F. Hayes, C. Chisholm-Brause, G. E. Brown Jr, G. A. Parks, K. O. Hodgson and J. O. Leckie, *Langmuir*, 1991, **7**, 367–373.

