



HAL
open science

Ni⁺ reactions with aminoacetonitrile, a potential prebiological precursor of glycine

Al Mokhtar Lamsabhi, Otilia M6, Manuel Y6ñez, Jean-Claude Guillemin, Violette Haldys, Jeanine Tortajada, Jean-Yves Salpin

► **To cite this version:**

Al Mokhtar Lamsabhi, Otilia M6, Manuel Y6ñez, Jean-Claude Guillemin, Violette Haldys, et al.. Ni⁺ reactions with aminoacetonitrile, a potential prebiological precursor of glycine. *Journal of Mass Spectrometry*, 2008, 43, pp.317-326. 10.1002/jms.1313 . hal-00259424

HAL Id: hal-00259424

<https://hal.science/hal-00259424>

Submitted on 28 Feb 2008

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Ni⁺ reactions with aminoacetonitrile, a potential pre-biological precursor of glycine

Al Mokhtar Lamsabhi*, Otilia Mó, Manuel Yáñez*

Departamento de Química C-9, Facultad de Ciencias, Universidad Autónoma de Madrid, Cantoblanco, 28049-Madrid, Spain.

Jean-Claude Guillemin*

Sciences Chimiques de Rennes – Ecole Nationale Supérieure de Chimie de Rennes –CNRS – 35700, Rennes France

Violette Haldys, Jeanine Tortajada, Jean-Yves Salpin*

Laboratoire d'Analyse et Modélisation pour la Biologie et l'Environnement – Université d'Evry Val d'Essonne – CNRS – Bâtiment Maupertuis, Boulevard François Mitterrand, 91025 Evry, France

Corresponding author: Jean-Yves Salpin

Tel: 33 1 69 47 76 47 Fax: 33 1 69 47 76 55

e-mail : jean-yves.salpin@univ-evry.fr

Number of pages (including Table and Figures) : 27

Abstract

The gas-phase reactions between Ni⁺(²D_{5/2}) and aminoacetonitrile, a molecule of pre-biological interest as possible precursor of glycine, have been investigated by means of mass spectrometry techniques. The MIKE spectrum reveals that the adduct ions [NC-CH₂-NH₂, Ni⁺] spontaneously decompose by losing HCN, H₂, and H₂CNH, the loss of hydrogen cyanide being clearly dominant. The structures and bonding characteristics of the aminoacetonitrile-Ni⁺ complexes as well as the different stationary points of the corresponding potential energy surface (PES) have been theoretically studied by DFT calculations carried out at B3LYP/6-311G(d,p) level. A cyclic intermediate, in which Ni⁺ is bisligated to the cyano and the amino group, plays an important role in the unimolecular reactivity of these ions, because it is the precursor for the observed losses of HCN and H₂CNH. In all mechanisms associated with the loss of H₂, the metal acts as hydrogen carrier favoring the formation of the H₂ molecule. The estimated bond dissociation energy of

aminoacetonitrile- Ni^+ complexes (291 kJ/mol) is larger than those measured for other nitrogen bases such as pyridine or pyrimidine and only slightly smaller than that of adenine.

Introduction:

The reactions between transition metal cations and organic or inorganic molecules have attracted a great deal of attention in the last two decades, because these processes are involved in a significant number of relevant processes in chemistry and biochemistry¹⁻⁴. Transition metals are generally present in the biological media as solvated ions or complexed by different kinds of peptides and proteins. They can also interact with other biomolecules such as nucleic acids including different effects that can vary from the stabilization of the helix to transcription failures.⁵⁻¹¹ Metal cations may play also an important role in astrochemistry. In this case the interactions may take place strictly in the gas-phase or in the surface of dust particles or meteorites, but normally involve small molecules. Among them, aminoacetonitrile presents a particular interest as a precursor of glycine in astrochemical media. As a matter of fact glycine may be formed in meteorites¹² through the interaction of ammonia, formaldehyde and hydrogen cyanide that produce aminoacetonitrile, which is then finally hydrolyzed to yield glycine. This process can be, however, perturbed by the interaction with metal cations, such as Ni^+ , which as shown in the present study destroy aminoacetonitrile by the loss of HCN. Also, some photochemical studies suggest that glycine can be formed in ices in the interstellar medium or cometary bodies¹³⁻¹⁶, as well. The spectroscopic characterization of this compound by means of infrared spectroscopy in Ar matrices was carried out recently.¹⁷ Several theoretical studies at different levels of accuracy on aminoacetonitrile,¹⁷⁻¹⁹ have been also reported in the literature. However, very little is known about the intrinsic reactivity of this compound and only very recently its gas-phase basicity has been measured by means of ion cyclotron resonance (ICR) techniques.²⁰ We aim here at characterizing the reactivity of this interesting species with respect

to Ni^+ , a prototype of open-shell transition metal cation with a $3d^9$ ($^2D_{5/2}$) electronic ground state. The interplay between mass spectrometry techniques and density functional theory (DFT) calculations would allow to gain some understanding on the behavior and bonding of aminoacetonitrile- Ni^+ that, in principle, may be useful to rationalize the behavior of more complicated systems which present similar basic sites.

Experimental

Synthesis Aminoacetonitrile has been prepared as recently reported.²⁰

Mass spectrometry:

All experiments were carried out using a VG Analytical ZAB-HSQ hybrid mass spectrometer of BEqQ geometry which has been described in detail previously.²¹ Complexes were generated by the CI-FAB method.²²⁻²⁷ The CI-FAB source was constructed from VG Analytical EI/CI and FAB ion source parts with the same modifications described by Freas et al.²² The conventional FAB probe tip has been replaced by a nickel foil of high purity. “Naked” metal ions were generated by bombardment with fast xenon atoms (Xe gas 7-8 keV kinetic energy, 1-2 mA of emission current in the FAB gun). The organic samples were introduced via a probe in a non-heated source. We can assume that due to the relatively high pressure in that source (10^2 - 10^3 Pa), efficient collisional cooling of the generated ions takes place. Therefore we will consider that excited states of the Ni^+ ions which could be formed in these experimental conditions are not likely to participate in the observed reactivity as already postulated by Hornung et al.²⁶ The ion beam of the Ni^+ adduct complexes formed with aminoacetonitrile were mass-selected (using an acceleration voltage of 8 kV) with the magnetic analyser B. The ionic products of unimolecular fragmentations, occurring in the second field-free (2nd FFR) region following the magnet, were analyzed by means of Mass-analyzed Ion Kinetic Energy MIKE^{28,29} by scanning the electric sector E.

The CAD (Collision Activated Dissociation) experiments were carried out in the same fashion but introducing Argon in the cell as the collision gas. The pressure of argon in the collision cell was adjusted so that the main beam signal was reduced by approximately 30%. The spectra were recorded at a resolving power (R) of ~ 1000

Computational details:

All quantum chemistry calculations presented in this paper have been carried out with the B3LYP hybrid DFT method^{30,31} as implemented in the Gaussian03 series of programs.³² The geometries of the different species under consideration were optimized using the all-electron basis (14s9p5d/9s5p3d) of Wachters³³ and Hay³⁴ augmented by a set of *f* functions for Ni and the 6-311G** basis set for remaining atoms of the system. The same basis set expansion and method were used to calculate the harmonic vibrational frequencies, in order to classify the stationary points of the potential energy surface (PES) as local minima or transition states, and to evaluate the corresponding zero-point energies (ZPE), which were scaled by the empirical factor 0.9806.³⁵ In all the cases, the $\langle S^2 \rangle$ expectation value showed that the spin contamination of the unrestricted wave function was always very small.

The bonding characteristics, as well as the electron density redistributions triggered by Ni⁺ association were analyzed by means of the atoms in molecules (AIM) theory.³⁶ For this purpose we have located the relevant bond critical points and evaluated the charge density at each of them. To perform the AIM analysis we have used the AIMPAC series of programs.³⁷ Also a second order perturbation method in the framework of the natural bond orbital (NBO) approach³⁸ was used to evaluate the interactions between orbitals of the base and orbitals of the metal, involved in the dative bonds from the former to the latter and possible back donations from the latter to the former. Since all complexes are open-shell systems, the NBO analysis has to be carried out for both the α - and β -sets of MOs. However, for the sake of simplicity, hereafter we will provide only the information corresponding to orbital

interactions within the β -subset, because this is the subset that contains the $3d$ as well as the $4s$ unoccupied orbitals. The α natural orbital set exhibits a similar behavior although in this case only the empty $4s$ orbital is within the subset.

Results and discussion:

Mass spectra

The $^{58}\text{Ni}^+$ ions react with neutral aminoacetonitrile to produce $^{58}\text{Ni-NCCH}_2\text{NH}_2^+$ adduct ions at m/z 114. The unimolecular decomposition of the $^{58}\text{Ni-NCCH}_2\text{NH}_2^+$ complex have been investigated by means of MIKE analysis to obtain information related to the structure and reactivity of this complex. The MIKE spectrum is shown in Figure 1. The $^{58}\text{Ni-NCCH}_2\text{NH}_2^+$ ion undergoes fragmentation according to several dissociation pathways. The main fragmentation corresponds to loss of $[\text{H,C,N}]$ to produce $^{58}\text{Ni-C,H}_3\text{N}^+$ ion at m/z 87, the base peak of MIKE spectrum. This reactivity differs from that observed with alkanenitriles, for which loss of 27 daltons is not observed. As a matter of fact, previous studies have demonstrated that the unimolecular reactivity of Ni^+ /alkanenitrile complexes is characterized by loss of the intact ligand, and beginning with *n*-propyl cyanide, losses of H_2 and alkenes.³⁹⁻
⁴² A dramatic increase of the two latter processes with the chain length has been noted. On the other hand, elimination of hydrogen cyanide has been reported for α - and β -unsaturated alkenenitriles.⁴³ A second peak associated with loss of H_2 is observed at m/z 112. Such a dehydrogenation process has been already observed for both alkanenitriles³⁹⁻⁴¹ and primary amines.⁴⁴⁻⁴⁶ but its intensity is presently rather weak compared to what has been reported for primary amines. Another significant difference between the unimolecular reactivity of Ni^+ /aminoacetonitrile and Ni^+ /amine systems is the absence of elimination of nickel hydride NiH in the former case. To complete this survey, another two small peaks are also observed at m/z 113 and m/z 85. They correspond to the elimination of H^\cdot and $[\text{C,H}_3\text{N}]$, respectively.

Ions at m/z 85 have been already observed in significant intensity on metastable spectra Ni^+ /alkenenitriles adducts⁴³, or more recently on the CID spectrum of electrospray-generated $[\text{Ni}(\text{NC}-\text{CH}_2-\text{N}_3), -\text{N}_2]^+$ complexes.⁴⁷ Finally, it is worth noting that bare Ni^+ cation is not observed under metastable conditions while it is the predominant process for short-length alkylnitriles.⁴¹

Under CAD conditions one can note in the resulting CAD spectrum displayed in Figure 2, that the intensity of the peak corresponding to the loss of $[\text{H,C,N}]$ increases showing that this decomposing channel is favored when energy is provided. Formation of m/z 85 and 87 ions could correspond to elimination of methanimine and hydrogen cyanide, respectively. Since the ability of transition metal ions to be dicoordinated is well known, the competitive losses of neutral HCN and CH_2NH might be probably arise from Ni^+ -bound heterodimers such as $[\text{HCN}-\text{Ni}^+-\text{NH}=\text{CH}_2]$, which may undergo competitive dissociations leading to the two fragment ions at m/z 85 and m/z 87.

We can also observe in this spectrum the presence of two minor ions at m/z 58 and 60. The first one corresponds to bare $^{58}\text{Ni}^+$ generated by the elimination of the intact ligand. But, this process occurs at a very minor extent compared to monofunctional molecules such as nitriles and primary amines. The second one might be H_2Ni^+ . However, formation of this particular ion has never been observed in previous studies, neither with amines nor with nitriles. This ion at m/z 60 could also correspond, at least to some extent, to $^{60}\text{Ni}^+$. As a matter of fact, the observation of dehydrogenation in the metastable spectrum of the $[\text{Ni}-\text{NCCH}_2\text{NH}_2]^+$ complex strongly suggests that the ^{58}Ni -complex generated in the FAB source may interfere with the corresponding dehydrogenation product of the ^{60}Ni complex, namely $[\text{Ni}-\text{NCCH}_2\text{NH}_2, -\text{H}_2]^+$, which in turn could give rise to bare $^{60}\text{Ni}^+$ ion. Additional experiments such as MIKE and CAD spectrum of both m/z 112 and $[\text{Ni}-\text{NCCH}_2\text{NH}_2]^+$

complex (m/z 116) could have raised the uncertainty about the structure of the m/z 60 ion, but could not be performed because of failure of our FAB instrument. However, complementary electrospray experiments performed by Dr. D. Schröder confirmed that NiH_2^+ species is indeed generated. These spectra were recorded using a VG BIO-Q tandem QHQ mass spectrometer (Q stands for quadrupole and H for hexapole), which has been described elsewhere.⁴⁸ 1 mg of NiI_2 was dissolved in 1 ml distilled water, then 50 μl of a 5% solution of freshly made $\text{H}_2\text{NCH}_2\text{CN}$ in distilled water was added and the solution was measured, thus allowing ions of the type $(\text{H}_2\text{NCH}_2\text{CN})\text{NiI}^+$ to be generated. By adopting harsh ionization conditions, in-source loss of atomic iodine is observed, thereby allowing reduction of Ni(II) to Ni(I)⁴⁹ and formation of $[\text{Ni-NCCH}_2\text{NH}_2]^+$ complex. MS/MS spectra of this ion with either argon or xenon as collision gas exhibit a m/z 60 at elevated collision energy, undoubtedly attributed to NiH_2^+ , as hydrogen loss(es) is not observed with the QHQ instrument. Note that in sector experiments, hydrogen losses are often much preferred in detection.⁵⁰ Finally, one may assume that Ni^+ -bound heterodimer such as $[\text{H}_2\text{-Ni}^+\text{-NH=CH.CN}]$ could also be the precursor of both m/z 112 and m/z 60 ions.

In order to assess the mechanisms behind the aforementioned experimental findings we have carried out a detailed study of the $[\text{aminoacetonitrile-Ni}^+]$ potential energy surface by means of DFT calculations.

Coordination of Nickel⁺

The optimized geometries of the different conformers of $\text{Ni}^+\text{-NCCH}_2\text{NH}_2$ and a selection of some structural parameters are shown in Figure 3. Total energies and Ni^+ binding energies of all the structures (minima, transition states and fragments) considered in this study are summarized in Table S1 of the supporting information.

Previous studies about the reactivity of transition-metal ions (and notably Ni⁺) with nitriles have suggested the co-existence of both "end-on" and "side-on" coordination modes.^{40,41,51} The first one corresponds to the interaction of the metallic center with the lone pair of the nitrogen of the cyano group in a linear M⁺·N≡C arrangement (for monofunctional ligands) while the second one involves interaction with the π-system. Accordingly, two possible types of coordination have been considered: π-type interactions with the C≡N group which yields structures **a** and **d** (side-on), and σ-type interactions with the nitrogen lone pair of either the cyano or the amino groups, which leads to structures **b** and **c**, respectively (end-on). The most stable conformer has been found to be the complex side-on **a**, which is stabilized through the interaction of the metal cation with the nitrogen lone pair of the amino group and also through a π-type interaction with the C≡N group. As a matter of fact, a second order NBO analysis shows both interactions to be rather strong, even though the former is stronger than the latter. As illustrated in Table 1, for complex **a** there are two empty *sd*-type hybrid orbitals on the metal which participate in the interaction with aminoacetonitrile. The first of these interactions corresponds to a dative bond from one of the C-N π-bonding orbitals toward the second empty hybrid of Ni, with a 53 % participation of the 4*s* orbital. The second one is another dative bond from the lone pair of the amine group nitrogen into the first empty hybrid on Ni, with a 39% 4*s* character. Both dative interactions are followed by a backdonation from 3*d* occupied orbitals of Ni towards π_{CN}* antibonding orbital. The charge transfer from the π_{CN} bonding and the population of the π_{CN}* antibonding orbital is reflected in the lengthening of the bond (by 0.014 Å), as well as in a decrease of the charge density at the corresponding bond critical point (see Figure 4). The involvement of the amino lone-pair in the interaction with the metal cation results also in a significant lengthening (0.048 Å) of the C-NH₂ bond and in a concomitant decrease of the charge density at the corresponding bond critical point. It is worth noting that, coherently with the NBO picture, the molecular

graph of complex **a** shows two bond paths with origin in Ni^+ , connecting the metal to the amino nitrogen and to the CN group, as well as the presence of a ring critical point.

The second less stable complex **b**, which lies 18 kJ/mol above the global minimum **a**, corresponds to the end-on interaction of Ni^+ with the cyano nitrogen lone pair. The NBO analysis shows the existence of a very strong interaction between this lone pair and a sd hybrid on Ni with a large s character (77%), followed by a backdonation from an occupied d orbital of Ni towards a π_{CN}^* antibonding orbital, which results in a slight lengthening of the CN bond (see Figure 3). The less stable complex **d** is that in which Ni^+ only interacts with the CN group. The difference in energy with the complex **a** (70 kJ mol^{-1}) gives a qualitative estimate of the extra-stabilization provided by the amino group. The NBO analysis indicates that both π - and σ -interactions take place (see Table 1) although the former are clearly dominant. As in the other complexes a backdonation from occupied $3d$ orbitals of the metal cation into the π_{CN}^* antibonding orbital also occurs. These interactions and the subsequent polarization of the C-C and the C-NH₂ bonds result in a lengthening of both the CN and the CC bonds by 0.03 Å and 0.037 Å, respectively and in a shortening of the C-NH₂ linkage by 0.032 Å. The attachment of Ni^+ to the amino group leads to structure **c** which is 33 kJ/mol less stable than the global minimum. In this case, only a dative bond from the amino nitrogen lone pair toward an empty sd hybrid on Ni, with a 53% of s character is found. As expected this involves a lengthening of the C-NH₂ bond by 0.048 Å and a concomitant shortening of the C-C bond by 0.018 Å.

Reactivity of the $[\text{Ni-NCCH}_2\text{NH}_2]^+$ adducts

As mentioned above, the MIKE and MIKE-CAD spectra of $[\text{}^{58}\text{Ni-NCCH}_2\text{NH}_2]^+$ shows different fragmentation processes, that we may attribute to loss of H₂, HCN or CH₂NH. The

loss of HCN and H₂ being the dominant ones, we will concentrate our attention in these two processes in particular.

As we shall discuss shortly, the different mechanisms associated with the two dominant unimolecular processes may have its origin not only in the global minimum but also in the other less stable aminoacetonitrile-Ni⁺ complexes. This is consistent with previous experimental studies dealing with the reactivity towards nitriles, which suggested that a single type of interaction could not account for all the fragmentation processes observed.^{40,43} However, as shown in Figure 5 all of them are connected through rather small activation barriers, but more importantly, well below the entrance channel. The most stable structure **a** may evolve to complex **b** by a simple rotation of Ni in the plane molecule, through a barrier of 54 kJ/mol. A rotation of the metal cation out of the plane of the molecule would connect **a** and **d**. The evolution from **a** to **c**, through an energy barrier of 45 kJ/mol, implies an internal rotation of the -NH₂Ni group.

HCN-Loss

The observed loss of HCN dominates both the MIKE and CAD spectra and requires unavoidably a hydrogen shift towards the carbon atom of the CN group. This is possible from either the global minimum **a** or from the local minimum **b**. Both processes correspond to H-transfer from the methylene group toward the carbon atom of the CN group leading to the cyclic minimum **2** (see Figure 6), where nickel is bisligated to the two nitrogen atoms. However, the process with origin in the global minimum **a** should be discarded because it involves an activation barrier which is not only above the entrance channel (See Figure 6), but also much higher than the barrier associated with the **a** → **b** isomerization.

The most favorable process from the cyclic minimum **2** is an α -C-C bond insertion of the metal yielding minimum **3**, which may eventually dissociate into HCN + [NiCH=NH₂]⁺ or alternatively into [NiNCH]⁺ + HC-NH₂, the former process being much more favorable from

the thermodynamic point of view than the latter, which is in agreement with the experimental finding that the dominant loss corresponds to HCN and not to $[C,H_3,N]$. One may also consider the possibility of inverting the order of the processes, by inserting first the metal into the C-C bond and having the hydrogen shift as a second step. The global minimum **a** is a good starting point for such a mechanism due to the bridging position of the metal between the two basic sites of the aminoacetonitrile. In fact the insertion leads to the local minimum **1** through the transition state **TSa_1**. It can be noticed that the insertion is accompanied by an internal rotation of the $C\equiv N$ group, because as soon as the CC bond cleaves, the interaction of the nitrogen atom of the CN fragment with the metal cation is very strong, reflecting its high intrinsic basicity. From minimum **1**, two hydrogen shifts can be envisaged. The most favorable one, involving the **TS1_3** transition state, involves a methylene hydrogen and leads also to structure **3**, already found in the first mechanism discussed above with origin in the cyclic minimum **2**. The second possible hydrogen shift involves a amino hydrogen and leads to a very stable local minimum **4**, which lies 172 kJ mol^{-1} below the most stable aminoacetonitrile- Ni^+ adduct, **a**. As shown in Figure 6, minimum **4** could also be formed by a hydrogen shift from **3**. Similarly to minimum **3**, structure **4** would eventually dissociate either by losing HCN or $[C,H_3,N]$. There is a difference however between both mechanisms as far as the loss of HCN is concerned. When dissociation originates in minimum **4**, in the accompanying ion product, Ni^+ is attached to the imino nitrogen of $H_2C=NH$, while when the dissociation originates in minimum **3** it is attached to the carbon atom of $HC-NH_2$. In summary, the observed loss of HCN can be associated with two different mechanisms with origin in adducts **a** and **b**, respectively, that lead to the same precursor **3** and which involved rather similar activation barriers. The alternative mechanisms for the loss of HCN have as precursor structure **4**, but the activation barriers to reach this minimum are much higher than

those to yield **3**, and therefore we may conclude that the majority of the $[\text{Ni}, \text{H}_3, \text{C}, \text{N}]^+$ product ions will be in the form of $[\text{NiCH}=\text{NH}_2]^+$ complexes.

It is worth noting also that the loss of $[\text{C}, \text{H}_3, \text{N}]$ in the form of $\text{CH}=\text{NH}_2$ is slightly endothermic, while the loss of $\text{CH}_2=\text{NH}$ is highly exothermic. Hence, very likely the observed $[\text{C}, \text{H}_3, \text{N}]$ loss comes mostly from the fragmentation of structure **4**. It is also important to emphasize that according to our theoretical survey the $[\text{H}, \text{C}, \text{N}, \text{Ni}]^+$ cation has always a HCNNi^+ connectivity. Actually an ion with an H-Ni-CN structure lies much higher in energy.

Some other mechanisms with origin in the local minimum **2** lead also to the loss of HCN (or the loss of H_2), but they involve larger activation barriers than those discussed above (see Figure 1S of the supporting information).

Some mechanisms leading to the loss of HNC instead of HCN have also been investigated. However, all hydrogen transfers either from the amino group or from the CH_2 toward the cyano group, involved activation barriers that were well above the entrance channel (51 and 101 kJ mol^{-1} , respectively). Since these mechanisms are not likely to take place they have not been included in Figure 6.

H₂-Loss

Many possibilities may be considered for the loss of H_2 . In our survey we have analyzed a large number of processes, but for the sake of conciseness we are going to summarize the most favorable ones. As we shall show along this section, starting from complexes **c** or **a**, the exit channels are much lower in energy than the one mentioned above (see Figure 7). All these mechanisms have in common that the nickel atom acts as hydrogen carrier favoring the formation of the H_2 molecule. In fact, starting from **c**, two pathways are opened, a hydrogen transfer from the amino group to nickel (through **TSc₁₀**) followed by a second H-transfer from the methylene group (through **TS10-12**), or a process where these two

steps are inverted, i.e., the first step is a H-transfer from the methylene group (through **TSc_8**) toward Ni, and the second one a H-transfer from the amino group (**TS8_12**). In both cases the same local minimum **12** is reached. As illustrated in Figure 7, the last possibility is clearly more favorable. Even though **TS8_12** is slightly above the entrance channel by 26 kJ mol⁻¹, this process can be observed because under the experimental conditions the local minima may have enough internal energy to overpass barriers slightly higher than the entrance channel. On the other hand, since the process corresponds to a H-transfer the existence of tunneling effects should not be discarded. Both situations would be consistent with the fact that the H₂ loss is observed in much less proportion than the loss of HCN. Finally, note that we also considered a multi-center transition state (MCTS) for the **9** → **13** interconversion, since MCTS usually provide rather low-lying pathways.^{52,53} However, it turned out that optimization of such a MCTS failed, because one of the hydrogen atom moves during the optimization process and the initial structure finally collapses back to the intermediate **9**.

The most stable adduct **a** may be also a precursor for the loss of H₂, but as shown in Figure 7, the mechanism through the intermediates **9**, **11** to reach **13**, involves a quite high activation barrier associated with the **9** → **11** isomerization, and therefore should not compete with the mechanism with origin in adduct **c** and preliminary **a** → **c** interconversion.

As a final remark, it is also worth noting that the estimated bond dissociation energy of aminoacetonitrile-Ni⁺ complexes (291 kJ/mol) is larger than those measured for other nitrogen bases such as pyridine (256 ± 15)⁹ or pyrimidine (244 ± 9)⁵⁴ and only slightly smaller than that of adenine (297 ± 10)⁵⁵, in which the metal ion also forms a chelate structure bridging between the N7 position of the imidazolic ring and the amino group.

Conclusions

The MIKE spectra of $[\text{NC-CH}_2\text{-NH}_2, \text{Ni}^+]$ ions show that they spontaneously decompose by losing HCN, H_2 , and $[\text{H}_3\text{C,N}]$, the loss of hydrogen cyanide being clearly dominant. A survey of the corresponding potential energy surface, shows that the direct insertion of the metal cation into the C-C bond of the most stable adduct or the insertion through the cyclic intermediate **1**, in which Ni^+ is bisligated to the cyano and the amino group are the most favourable mechanisms leading to the observed losses of HCN and $[\text{H}_3\text{C,N}]$. The loss of $\text{CH}=\text{NH}_2$ is predicted to be endothermic, while the loss of $\text{CH}_2=\text{NH}$ is clearly exothermic, so that we can safely conclude that the observed loss of $[\text{H}_3\text{C,N}]$ corresponds mostly to $\text{CH}_2=\text{NH}$. Our theoretical calculations also indicate that in the $[\text{H,C,N,Ni}]^+$ accompanying ion, the metal is attached to the N atom of the HCN molecule. In all the mechanisms associated with the loss of H_2 , the metal acts as hydrogen carrier favoring the formation of the H_2 molecule. The topology of the PES also shows that the m/z 87 peak observed in both the MIKE and the CAD spectra corresponds exclusively to the loss of HCN, because the mechanisms associated with the loss of HNC involve activation barriers much higher in energy than the entrance channel.

Acknowledgement: This work has been partially supported by the DGI Project No. CTQ2006-08558/BQU and by the Project MADRISOLAR, Ref.: S-0505/PPQ/0225 of the Comunidad Autónoma de Madrid. Authors would like to warmly acknowledge Professor Detlef Schröder for kindly performing complementary experiments. A.M.L gratefully acknowledges a Juan de la Cierva contract from the Ministerio de Educación y Ciencia of Spain. We also acknowledge a generous allocation of computer time at the Centro de Computación Científica de la Universidad Autónoma de Madrid. J.-C. G. thanks the GDR CNRS “Exobiologie” and the Centre National d'Etudes Spatiales for financial support.

Supporting information Total, zero-point energies (ZPE) and relative energies of the various structures considered are provided as supporting information, together with potential energy surfaces associated with alternate mechanisms leading to the ions observed experimentally.

Bibliography

1. Spiro TG. In *Metal Ions In Biology*, (eds). Wiley: New York, 1980;
2. Sigel A, Sigel H. In *Metal Ions in Biological Systems: Nickel and its Role in Biology*, *Metal Ions in Biological Systems: Nickel and its Role in Biology*, (eds). Marcel Decker: New York, 1988;
3. Fontijn A. In *Gas-Phase Metal Reactions*, (eds). North Holland: Amsterdam, 1992;
4. Sigel A, Sigel H. In *Metal Ions Biological Systems: Intercations of Metal Ions with Nucleotides, Nucleic Acids and their constituents*, *Metal Ions Biological Systems: Intercations of Metal Ions with Nucleotides, Nucleic Acids and their constituents*, (eds). Marcel Decker: New York, 1996;
5. Anwander EHS, Probst MM, Rode BM. The Influence of Li^+ , Na^+ , Mg^{2+} , Ca^{2+} , and Zn^{2+} Ions on the Hydrogen-Bonds of the Watson-Crick Base-Pairs. *Biopolymers* 1990; **29**: 757-769.
6. Cerda BA, Wesdemiotis C. Li^+ , Na^+ , and K^+ binding to the DNA and RNA nucleobases. Bond energies and attachment sites from the dissociation of metal ion-bound heterodimers. *J. Am. Chem. Soc.* 1996; **118**: 11884-11892.
7. Burda JV, Sponer J, Hobza P. Ab Initio study of the interaction of guanine and adenine with various mono- and bivalent metal cations (Li^+ , Na^+ , K^+ , Rb^+ , Cs^+ ; Cu^+ , Ag^+ , Au^+ ; Mg^{2+} , Ca^{2+} , Sr^{2+} , Ba^{2+} ; Zn^{2+} , Cd^{2+} , and Hg^{2+}). *J. Phys. Chem.* 1996; **100**: 7250-7255.
8. Gadre SR, Pundlik SS, Limaye AC, Rendell AP. Electrostatic investigation of metal cation binding to DNA bases and base pairs. *Chemical Communications* 1998; 573-574.
9. Rodgers MT, Stanley JR, Amunugama R. Periodic trends in the binding of metal ions to pyridine studied by threshold collision-induced dissociation and density functional theory. *J. Am. Chem. Soc.* 2000; **122**: 10969-10978.
10. Russo N, Toscano M, Grand A. Bond energies and attachments sites of sodium and potassium cations to ONA and RNA nucleic acid bases in the gas phase. *J. Am. Chem. Soc.* 2001; **123**: 10272-10279.
11. Csaszar K, Spackova N, Stefl R, Sponer J, Leontis NB. Molecular dynamics of the frame-shifting pseudoknot from beet western yellows virus: The role of non- Watson-

- Crick base-pairing, ordered hydration, cation binding and base mutations on stability and unfolding. *J. Mol. Biol.* 2001; **313**: 1073-1091.
12. Peltzer ET, Bada JL, Schlesinger G, Miller SL. The chemical conditions on the parent body of the Murchison meteorite: some conclusions based on amno, hydroxy and dicarboxylic acids. *Adv. Space Res.* 1984; **4**: 69-74.
 13. Sandford SA, Bernstein MP, Dworkin JP, Cooper GW, Allamandola LJ. The production of amino acids in interstellar ices - Implications for meteoritic organics. *Meteor. & Planet Sci.* 2002; **37**: A125-A125.
 14. Bernstein MP, Dworkin JP, Sandford SA, Cooper GW, Allamandola LJ. Racemic amino acids from the ultraviolet photolysis of interstellar ice analogues. *Nature* 2002; **416**: 401-403.
 15. Muñoz-Caro GM, Meierhenrich UJ, Schutte WA, Barbier B, Segovia AA, Rosenbauer H, Thiemann WHP, Brack A, Greenberg JM. Amino acids from ultraviolet irradiation of interstellar ice analogues. *Nature* 2002; **416**: 403-406.
 16. Meierhenrich UJ, Muñoz-Caro GM, Schutte WA, Barbier B, Segovia AA, Back A, Rosenbauer H, Thiemann WHP. Amino acids from ultraviolet irradiation of interstellar ice analogues. *Geochim. Cosmochim. Acta* 2002; **66**: A505-A505.
 17. Bernstein MP, Bauschlicher CW, Sandford SA. The infrared spectrum of matrix isolated aminoacetonitrile, a precursor to the amino acid glycine. *Adv. Space Res.* 2004; **33**: 40-43.
 18. Basiuk VA, Kobayashi Y. DFT study of HCN and N C-C N reactions with hydrogen species. *Int. J. Quantum Chem.* 2004; **99**: 92-101.
 19. Chaban GM. Anharmonic vibrational spectroscopy of nitriles and their complexes with water. *Journal of Physical Chemistry A* 2004; **108**: 4551-4556.
 20. Bouchoux G, Guillemin J-C, Lemahieu N, McMahon TB. Protonation thermochemistry of aminoacetonitrile. *Rapid Commun. Mass Spectrom.* 2006; **20**: 1187-1191.
 21. Harrison AG, Mercer RS, Reiner EJ, Young AB, Boyd RK, March RE, Porter CJ. A Hybrid Beqq Mass-Spectrometer for Studies in Gaseous Ion Chemistry. *Int. J. Mass Spectrom. Ion Processes* 1986; **74**: 13-31.
 22. Freas RB, Ross MM, Campana JE. Chemical Ionization Fast-Atom Bombardment Mass-Spectrometry - Ion Molecule Reactions. *J. Am. Chem. Soc.* 1985; **107**: 6195-6201.
 23. Freas RB, Campana JE. Reactions of Sputtered Copper Cluster Ions. *J. Am. Chem. Soc.* 1985; **107**: 6202-6204.
 24. Mestdagh H, Morin N, Rolando C. Gas-Phase Ion Chemistry - a Comparative-Study of Reaction of 1st Row Transition-Metal Cations with 2-Methyl Propane. *Tetrahedron Lett.* 1986; **27**: 33-36.

25. Drewello T, Eckart K, Lebrilla CB, Schwarz H. Fast-Atom-Bombardment as a Convenient Means to Generate Metal-Ion Complexes in the Gas-Phase. *Int. J. Mass Spectrom. Ion Processes* 1987; **76**: 13-15.
26. Hornung G, Schroder D, Schwarz H. Diastereoselective Gas-Phase Carbon-Carbon Bond Activation Mediated by Bare Co^+ Cations. *J. Am. Chem. Soc.* 1995; **117**: 8192-8196.
27. Chamot-Rooke J, Tortajada J, Morizur J-P. Studies of the reactions of ortho-, meta- and para-methylanisoles with transition metal ions M^+ ($\text{M} = \text{Cr}, \text{Fe}, \text{Co}, \text{Ni}, \text{Cu}$) in the gas phase. *European Journal of Mass Spectrometry* 1995; **1**: 471-478.
28. Cooks RG, Beynon JH, Caprioli RM, Lester GR. In *Metastable Ions*, (eds). Elsevier: New York, 1973;
29. Cooks RG. In *Collision Spectroscopy*, (eds). Plenum Press: New York, 1978;
30. Becke AD. A new mixing of Hartree-Fock and local-density-functional theories. *J. Chem. Phys.* 1993; **98**: 1372-1377.
31. Lee C, Yang W, Parr RG. Development of the Colle-Salvetti correlation-energy formula into a functional of the electron density. *Physical Review B* 1988; **37**: 785-789.
32. Frisch MJ, Trucks GW, Schlegel HB, Scuseria GE, Robb MA, Cheeseman JR, Zakrzewski VG, J. A. Montgomery J, Vreven T, Kudin KN, Burant JC, Millam JM, Iyengar SS, Tomasi J, Barone V, Mennucci B, Cossi M, Scalmani G, Rega N, Petersson GA, Nakatsuji H, Hada M, Ehara M, Toyota K, Fukuda R, Hasegawa J, Ishida M, Nakajima T, Honda Y, Kitao O, Adamo C, Jaramillo J, Gomperts R, Stratmann RE, Yazyev O, Austin J, Cammi R, Pomelli C, Ochterski J, Ayala PY, Morokuma K, Voth GA, Salvador P, Dannenberg JJ, Zakrzewski VG, Dapprich S, Daniels AD, Strain MC, Farkas O, Malick DK, Rabuck AD, Raghavachari K, Foresman JB, Ortiz JV, Cui Q, Baboul AG, Clifford S, Cioslowski J, Stefanov BB, Liu G, Liashenko A, Piskorz P, Komaromi I, Martin RL, Fox DJ, Keith T, Al-Laham MA, Peng CY, Nanayakkara A, Challacombe M, Gill PMW, Johnson B, Chen W, Wong MW, Gonzalez C, Pople JA. *Gaussian03, Revision C.02*. Gaussian, Inc. Wallingford CT. **2003**.
33. Wachters AJH. Gaussian Basis Set for Molecular Wavefunctions Containing Third-Row Atoms. *J. Chem. Phys.* 1970; **52**: 1033-1036.
34. Hay PJ. Gaussian basis sets for molecular calculations. The representation of 3d orbitals in transition-metal atoms. *J. Chem. Phys.* 1977; **66**: 4377-4384.
35. Scott AP, Radom L. Harmonic Vibrational Frequencies: An Evaluation of Hartree-Fock, Moeller-Plesset, Quadratic Configuration Interaction, Density Functional Theory, and Semiempirical Scale Factors. *J. Phys. Chem.* 1996; **100**: 16502-16513.
36. Bader RFW. In *Atoms in Molecules. A Quantum Theory*, (eds). Clarendon Press: Oxford, 1990;
37. Cheeseman J, Bader RFW. *AIMPAC*, **2000**.

38. Reed AE, Curtiss LA, Weinhold F. Intermolecular interactions from a natural bond orbital, donor-acceptor viewpoint. *Chemical Reviews* 1988; **88**: 899-926.
39. Lebrilla CB, Drewello T, Schwarz H. The Reaction Of Linear Nitriles With Transition-Metal Ions Fe⁺, Co⁺, Ni⁺ In The Gas-Phase - Does The Back-Bonding Ability Of The Metal-Ions Govern The Geometry Of The Rcn/M⁺ Complexes And What Is The Effect On The Oxidative Addition Of Internal Ch Bonds. *Int. J. Mass Spectrom. Ion Processes* 1987; **79**: 287-294.
40. Chen LZ, Miller JM. Ion Molecule Reactions Of Transition-Metal Ions With Nitriles In The Gas-Phase - Competitive Formation Of Both End-On And Side-On Coordination. *Canadian Journal Of Chemistry-Revue Canadienne De Chimie* 1991; **69**: 2002-2007.
41. Chen LZ, Miller JM. Ion-Molecule Reaction Of Alkanenitriles And Transition-Metal Ions In The Gas-Phase - A Study On Fragmentation Mechanism Of The Adducts. *J. Am. Soc. Mass Spectrom.* 1991; **2**: 120-124.
42. Eller K, Schwarz H. Organometallic chemistry in the gas phase. *Chem. Rev.* 1991; **91**: 1121-1177.
43. Chen LZ, Miller JM. Formation And Fragmentation Of Gas-Phase Ion-Molecule Complexes Of Transition-Metal Ions With Organic-Molecules Containing 2 Functional-Groups. *J. Am. Soc. Mass Spectrom.* 1992; **3**: 451-459.
44. Tolbert MA, Beauchamp JL. Homolytic and heterolytic bond dissociation energies of the second row group 8, 9, and 10 diatomic transition-metal hydrides: correlation with electronic structure. *J. Phys. Chem.* 1986; **90**: 5015-5022.
45. Karrass S, Pruesse T, Eller K, Schwarz H. Revised and modified mechanisms for the reactions of bare transition metal ions M⁺ (M = Fe, Co, Ni) with n-C₃H₇X (X = NH₂, OH) in the gas phase. *J. Am. Chem. Soc.* 1989; **111**: 9018-9023.
46. Chen LZ, Miller JM. The Chemistry Of Transition-Metal Ion Interactions With Cyclohexylamine In The Gas-Phase. *Rapid Commun. Mass Spectrom.* 1992; **6**: 492-497.
47. Couto N, Duarte MF, Fernandez MT, Rodrigues P, Barros MT, Costa ML, Cabral BJC. Complexation of transition metals by 3-azidopropionitrile. An electrospray ionization mass spectrometry study. *J. Am. Soc. Mass Spectrom.* 2007; **18**: 453-465.
48. Schröder D, Weiske T, Schwarz H. Dissociation behavior of Cu(urea)(⁺) complexes generated by electrospray ionization. *Int. J. Mass Spectrom.* 2002; **219**: 729-738.
49. Schlangen M, Schröder D, Schwarz H. Pronounced ligand effects and the role of formal oxidation states in the nickel-mediated thermal activation of methane. *Angew. Chem., Int. Ed. Engl.* 2007; **46**: 1641-1644.
50. Fiedler A, Schröder D, Schwarz H, Tjelta BL, Armentrout PB. "Bare" iron methoxide cation: A simple model to probe the mechanism of beta-hydrogen transfer in organometallic compounds. *J. Am. Chem. Soc.* 1996; **118**: 5047-5055.

51. Prusse T, Drewello T, Lebrilla CB, Schwarz H. Chain-Length-Dependent Ligand Effects on the Direction of Fe⁺-Mediated Activation and Cleavage of CC Bonds in Alkenyl Nitriles in the Gas-Phase. *J. Am. Chem. Soc.* 1989; **111**: 2857-2861.
52. Noll RJ, Yi SS, Weisshaar JC. Bimolecular Ni⁺(²D_{5/2}) + C₃H₈ reaction dynamics in real time. *J. Phys. Chem. A* 1998; **102**: 386-394.
53. Blomberg M, Yi SS, Noll RJ, Weisshaar JC. Gas-phase Ni⁺(²D_{5/2})+n-C₄H₁₀ reaction dynamics in real time: Experiment and statistical modeling based on density functional theory. *J. Phys. Chem. A* 1999; **103**: 7254-7267.
54. Amunugama R, Rodgers MT. Periodic trends in the binding of metal ions to pyrimidine studied by threshold collision-induced dissociation and density functional theory. *Journal of Physical Chemistry A* 2001; **105**: 9883-9892.
55. Rodgers MT, Armentrout PB. Influence of d orbital occupation on the binding of metal ions to adenine. *J. Am. Chem. Soc.* 2002; **124**: 2678-2691.

Table 1. NBO analysis of aminoacetonitrile-Ni⁺ adducts showing the energies^a (kJ mol⁻¹) associated with dative bonds from π_{CN} bonding orbitals of the cyano group and from the nitrogen atoms lone pairs toward the empty orbitals of the metal. The last two rows report the percentage of the *s* and *d* character of the Ni⁺ empty hybrid orbital. This analysis corresponds to π spin-orbitals.

Complex	a	b	c	d
$\pi_{\text{C}\equiv\text{N}} \rightarrow \text{LP}^* (1) (\text{Ni})$	-	-	-	54.1
$\pi_{\text{C}\equiv\text{N}} \rightarrow \text{LP}^* (2) (\text{Ni})$	61.9	-	-	18.1
$n_{\text{N}(\equiv\text{C})} \rightarrow \text{LP}^* (1) (\text{Ni})$	-	125.4	-	23.8
$n_{\text{N}(\equiv\text{C})} \rightarrow \text{LP}^* (2) (\text{Ni})$	-	-	-	-
$n_{\text{N}} \rightarrow \text{LP}^* (1) (\text{Ni})$	87.5	-	97.0	-
$n_{\text{N}} \rightarrow \text{LP}^* (2) (\text{Ni})$	-	-	-	-
$nd_{\text{Ni}} \rightarrow \pi^*_{\text{C}\equiv\text{N}}$	69.0	23.0	-	62.2
LP* (1) Ni	39% <i>s</i> , 61% <i>d</i>	77% <i>s</i> , 23% <i>d</i>	53% <i>s</i> , 47% <i>d</i>	93% <i>s</i> , 7% <i>d</i>
LP* (2) Ni	53 % <i>s</i> , 47% <i>d</i>	97% <i>s</i> , 3% <i>d</i>	42 % <i>s</i> , 58 % <i>d</i>	1% <i>s</i> , 99% <i>d</i>

^a Only interaction energies larger than 8 kJ mol⁻¹ are included

Figure Captions

1. MIKE spectrum of $[\text{}^{58}\text{Ni-NCCH}_2\text{NH}_2]^+$ ions. Mass to charge ratios of the fragment ions are deduced from the ratio of their kinetic energy to that of the precursor ion (m/z 114).
2. CAD spectrum of $[\text{}^{58}\text{Ni-NCCH}_2\text{NH}_2]^+$ ions. Mass to charge ratios of the fragment ions are deduced from the ratio of their kinetic energy to that of the precursor ion (m/z 114).
3. Optimized geometries of the aminoacetonitrile- Ni^+ adducts. Bond lengths are in Å and bond angles in degrees.
4. Molecular graphs of the aminoacetonitrile- Ni^+ adducts. Red dots represent bond critical points and yellow dots ring critical points. Electron densities are in a.u.
5. Isomerization barriers between the most stable aminoacetonitrile- Ni^+ adduct **a** and the other aminoacetonitrile- Ni^+ adducts. Relative energies are in kJ mol^{-1} .
6. Energy profile associated with the different mechanisms that lead to HCN and H_2CNH losses in aminoacetonitrile- Ni^+ gas-phase reactions. Energies relative to the most stable aminoacetonitrile- Ni^+ adduct **a** are in kJ mol^{-1} .
7. Energy profile associated with the loss of H_2 in aminoacetonitrile- Ni^+ gas-phase reactions. Energies relative to the most stable aminoacetonitrile- Ni^+ adduct **a** are in kJ mol^{-1} .

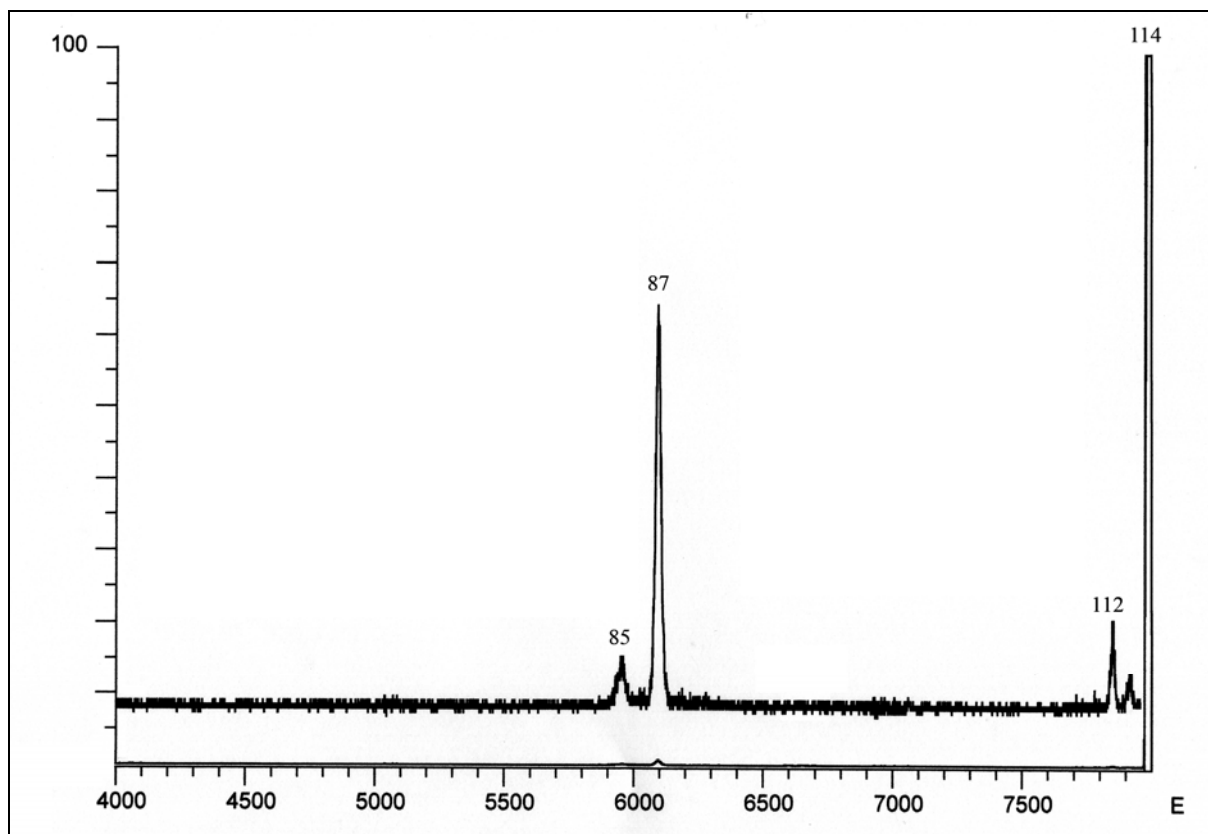


Figure 1

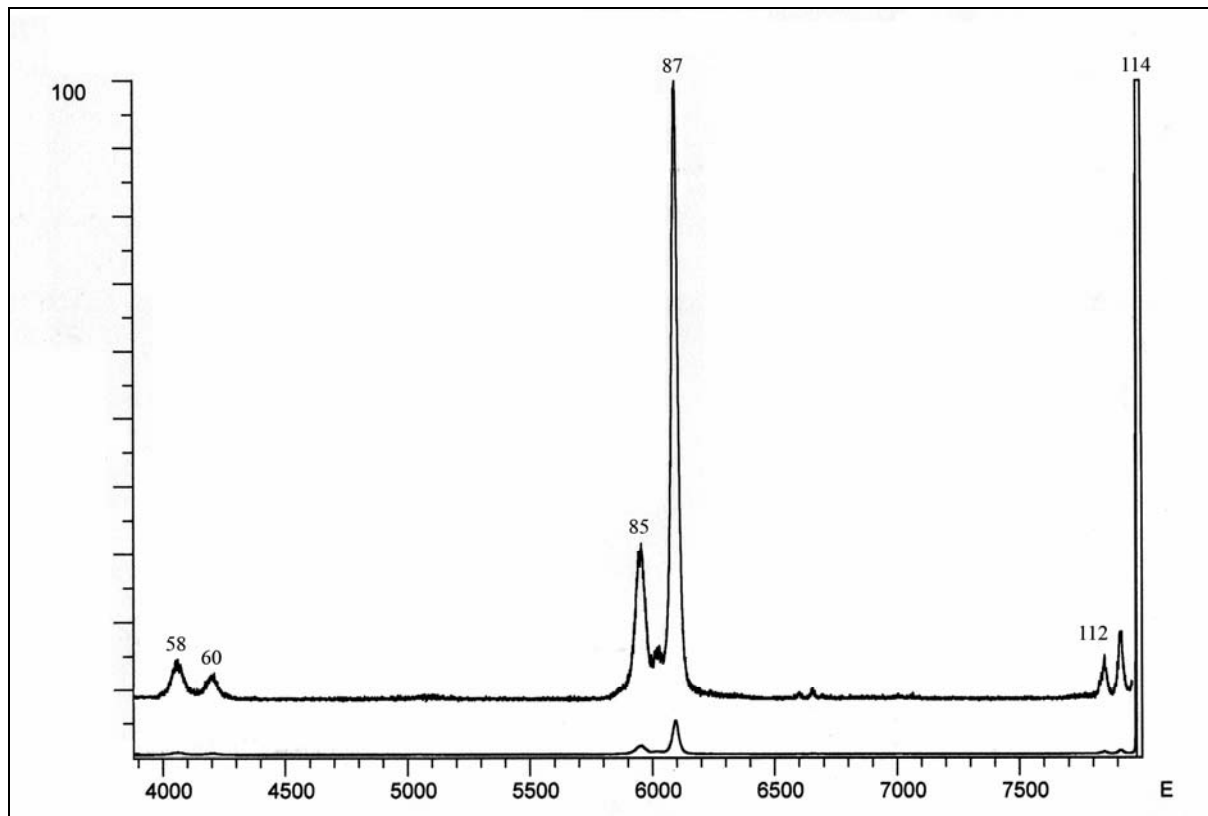


Figure 2

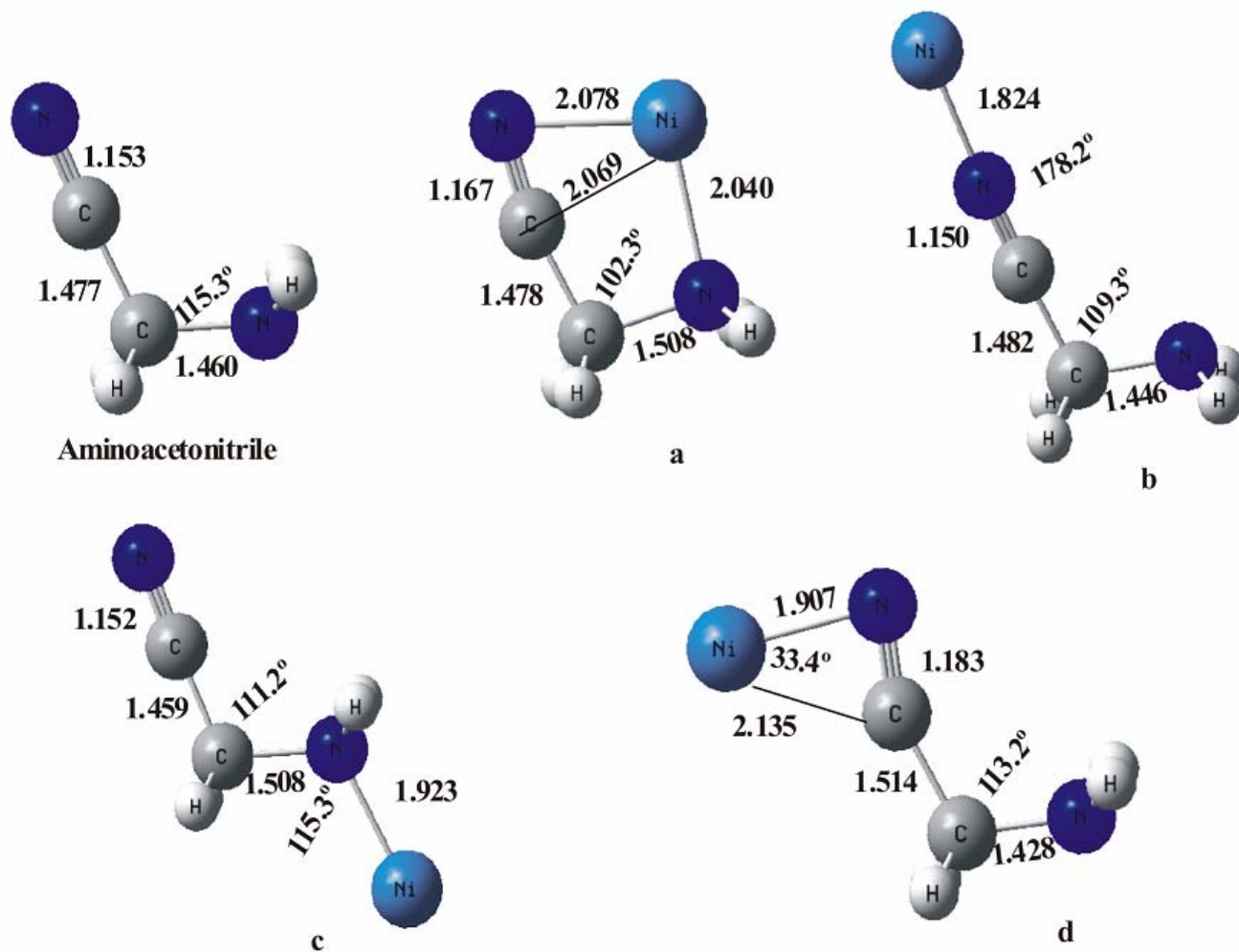


Figure 3

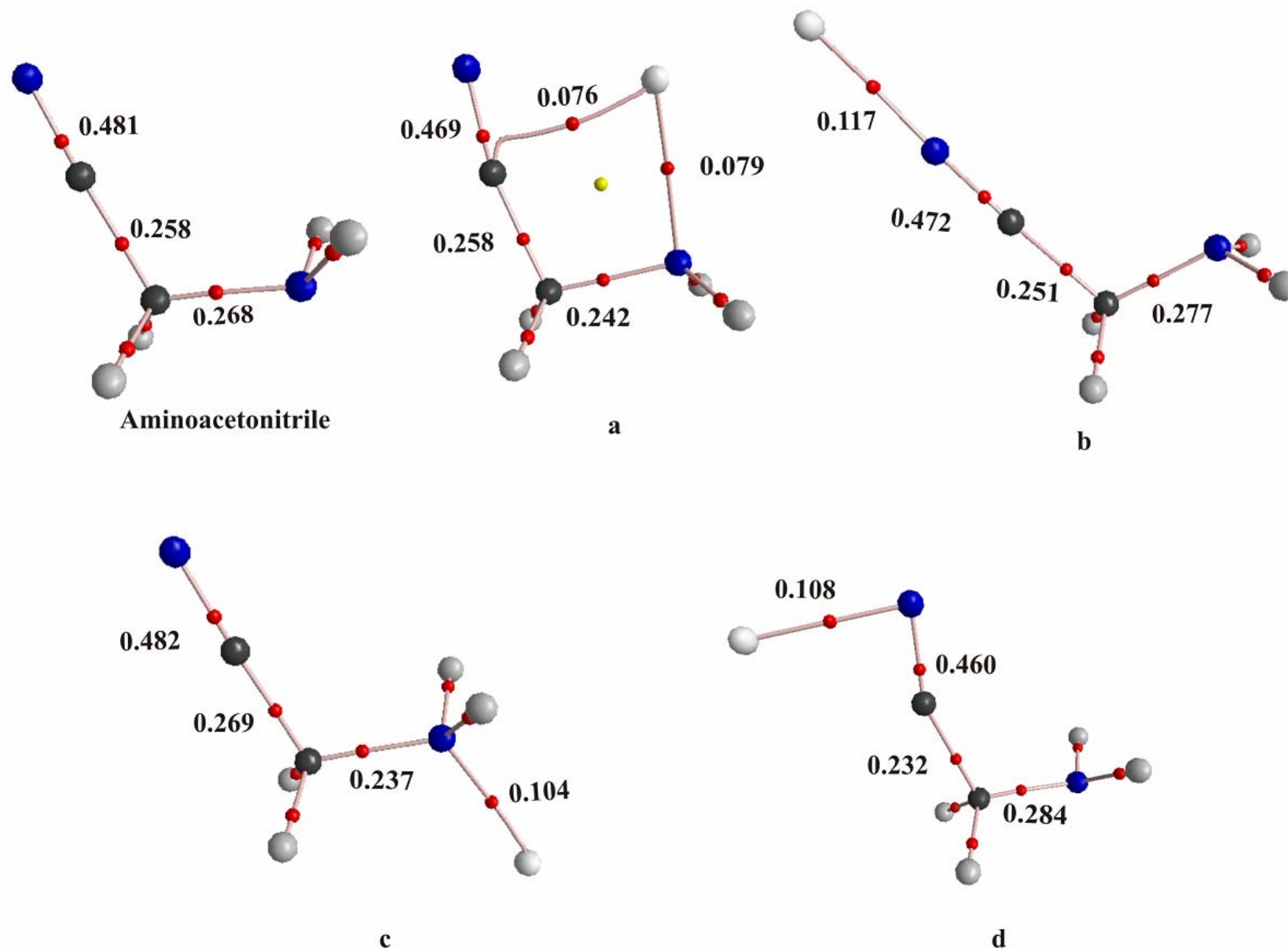


Figure 4

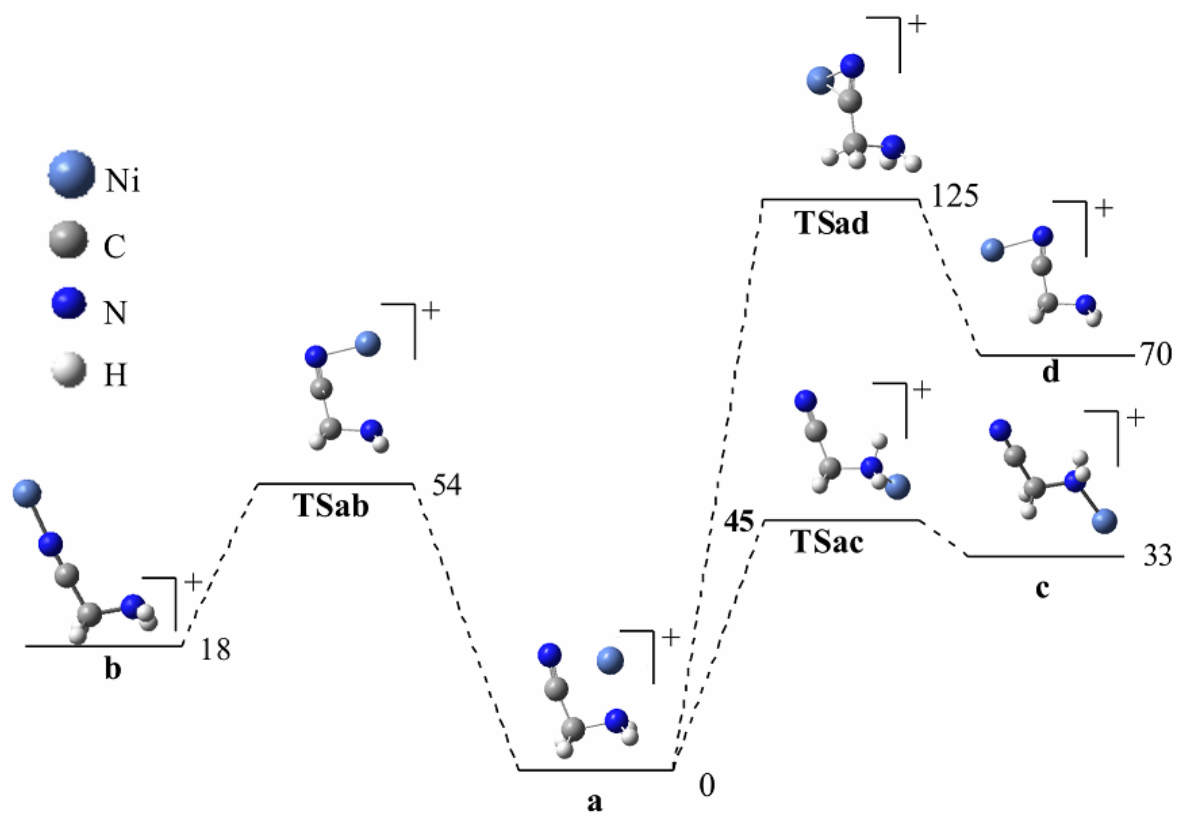


Figure 5

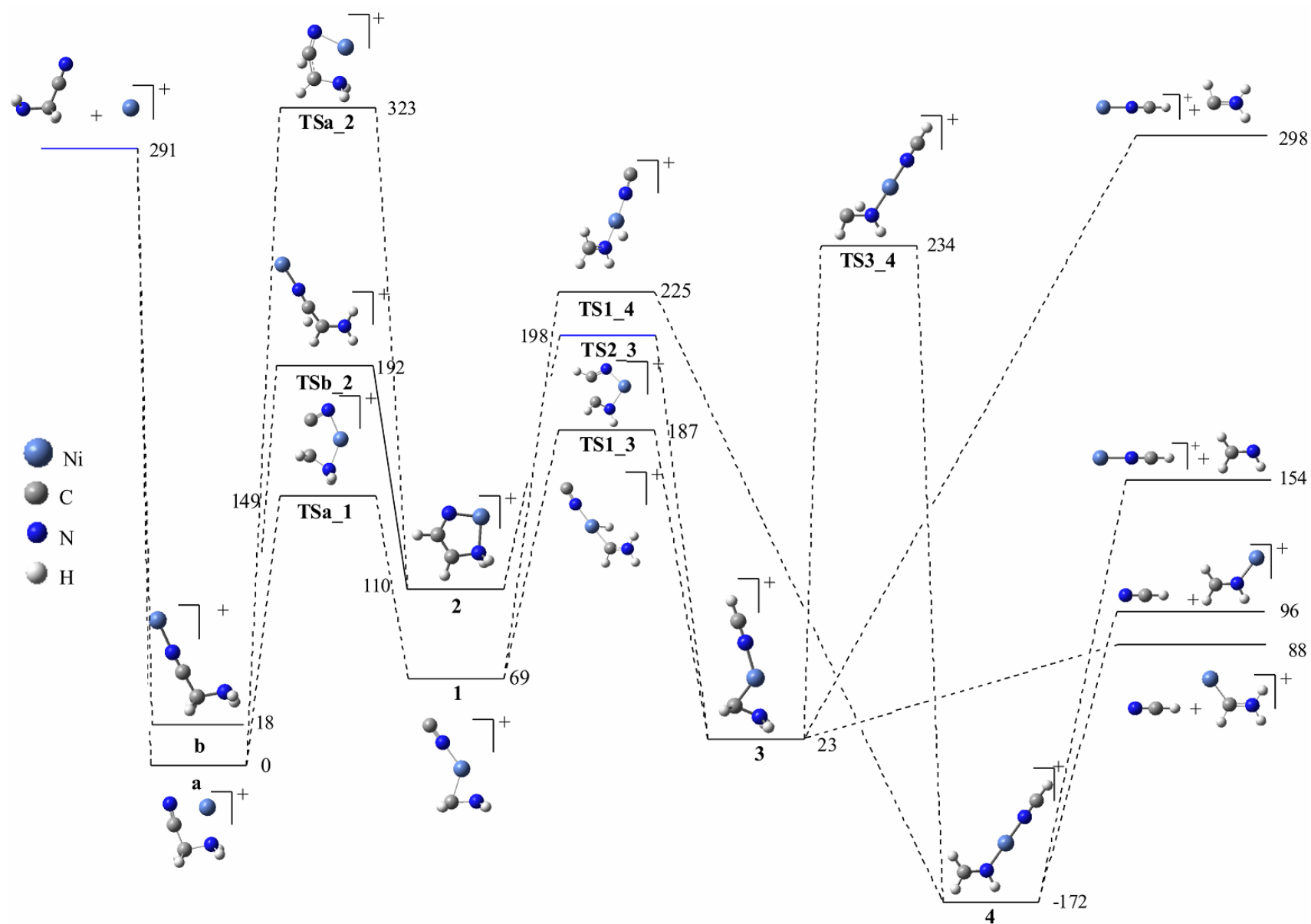


Figure 6

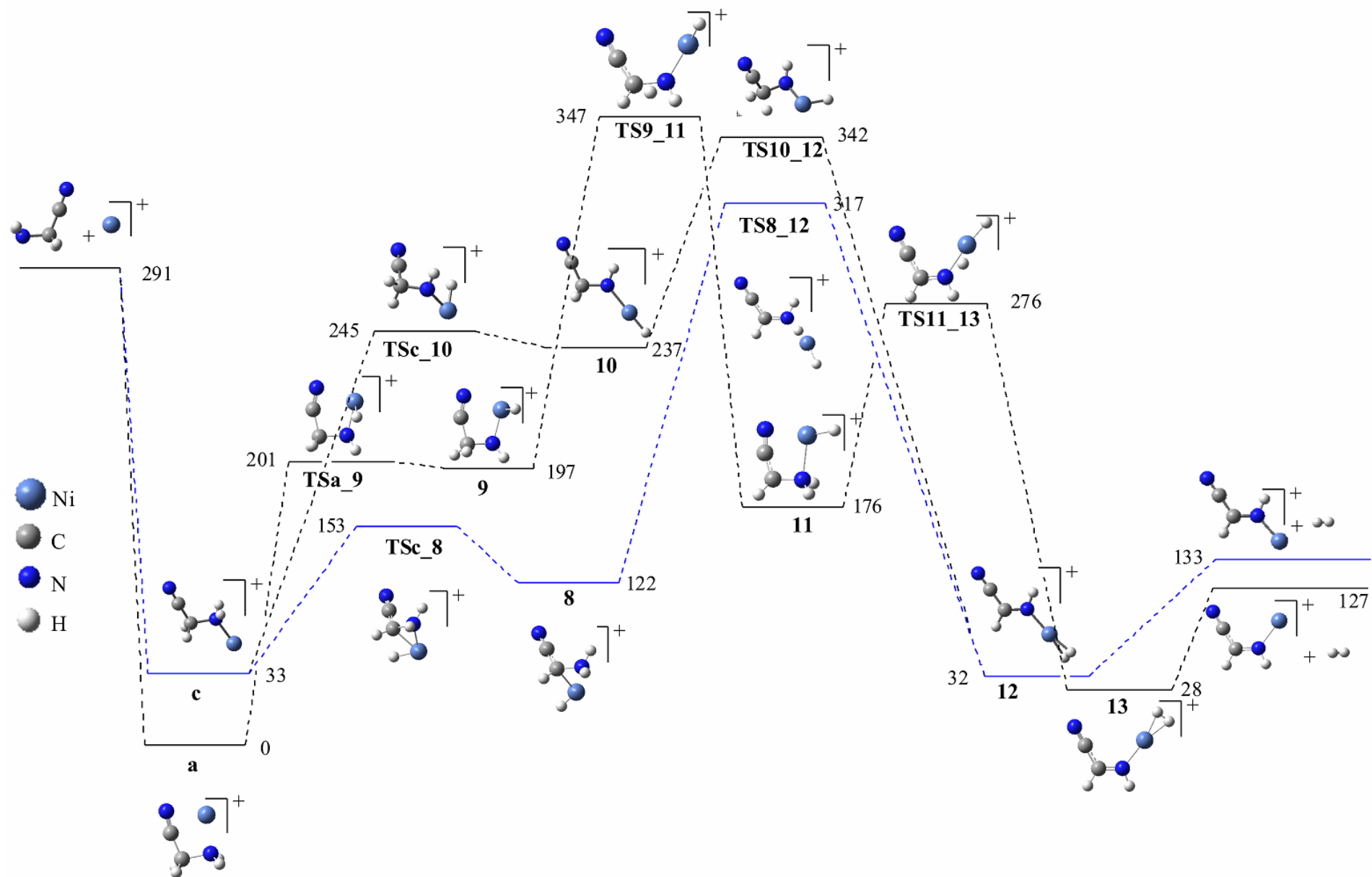


Figure 7