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# A new chemical compound with an unusual ratio of number of carbon and nitrogen atoms – C(N<sub>12</sub>): quantum-chemical modelling†

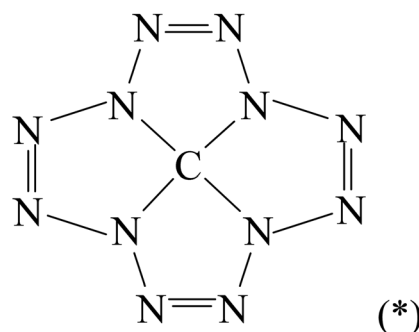
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Based on the results of a quantum chemical calculation using the DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP levels, the possibility of the existence of carbon–nitrogen compounds having the unusual carbon : nitrogen ratio 1 : 12, which is unknown for these elements at present, was shown. Data on the structural parameters are presented; it was shown that all CN<sub>4</sub> groupings are strictly planar. In addition, the bond lengths formed by nitrogen atoms and a C atom are equal to each other, and the bond angles formed by nitrogen atoms and a C atom are equal to 90.0°, or practically do not differ from this value. Thermodynamical parameters, NBO analysis data and HOMO/LUMO images for this compound are presented, too. Good agreement was found between the calculated data obtained using the above three quantum-chemical methods.

## Introduction

Carbon and nitrogen form among themselves a very significant number of chemical compounds – both neutral and cations and anions.<sup>1</sup> Among neutral compounds containing atoms of only these two elements, cyanogens C<sub>2</sub>N<sub>2</sub> (dicyan N≡C–C≡N, isocyanogen C=N–C≡N), diisocyanogen C=N–N=C), paracyanogen – a cyanogen polymer (NCCN)<sub>n</sub>, paraisocyanogen – an isocyanogen polymer, (CNCN)<sub>n</sub>,<sup>2–4</sup> percyanoalkynes, -alkenes and -alkanes,<sup>5–15</sup> dinitrilopolynes, perazidoalkynes, -alkenes and -alkanes,<sup>16,17</sup> percyanoheterocycles,<sup>18</sup> of which in total there are at least several dozen.<sup>1</sup> At the end of the 20th – beginning of the 21st centuries, in connection with the discovery of fullerenes, numerous derivatives of them were added to this list, which can be subdivided into two categories – cyanofullerenes that are a class of modified fullerenes in which cyano-groups are attached to a fullerene skeleton with the C<sub>60</sub>(CN)<sub>2n</sub> formula (*n* = 1–9), and azafullerenes that are a class of heterofullerenes in which the element substituting for carbon is nitrogen, f.e. bis[aza[60]fullerenyl] (C<sub>59</sub>N)<sub>2</sub>, diaza[60]fullerene C<sub>58</sub>N<sub>2</sub>, triaza[60]fullerene C<sub>57</sub>N<sub>3</sub> and dodecaaza[60]fullerene C<sub>48</sub>N<sub>12</sub>.<sup>19–24</sup> Tri-carbon tetranitride or simply carbon nitride C<sub>3</sub>N<sub>4</sub>, first described 100 years ago in ref. 25 and considered in more detail in a number of other works, in particular<sup>26–32</sup> and review

articles,<sup>33–36</sup> stands apart from this background. The carbon : nitrogen ratio in these compounds varies from 69 : 1 in diheterofullerene (C<sub>69</sub>N)<sub>2</sub> to 1 : 12 in tetraazidomethane C(N<sub>3</sub>)<sub>4</sub>.<sup>1–36</sup> Since a carbon atom is capable of forming four bonds with a maximum of four neighboring atoms, and a nitrogen atom – with three atoms, then for compounds of the above types, a ratio of 1 : 12 seems to be extremely possible. Nevertheless, the indicated ratio C : N = 1 : 12 can also be realized in another compound, which is an isomer of tetraazidomethane, but which, as far as is known, has not yet been considered in the literature even theoretically, namely in (dodecaazacyclododecantetraene-1,4,7,10)carbon(IV) C(N<sub>12</sub>) having the structural formula (\*)



In this connection, the given communication is devoted to the consideration of the question of the possibility of its existence and, in the case of a positive conclusion, to the determination of the parameters of its molecular structure and thermodynamic characteristics using various modern quantum-chemical methods of calculation, and namely, the density functional theory (DFT), MP2 and MP3 methods.

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## Method

To carry out the quantum-chemical calculation, in this work we used the density functional theory (DFT), which combines the standard extended split valence basis set TZVP and the hybrid functional B3PW91, described in detail in ref. 37–39 and used by us, in particular, in ref. 40–42. The use of the B3PW91/TZVP

method, in this case, is because, according to,<sup>37–39</sup> it allows one to obtain, as a rule, the most accurate (*i.e.*, close to experimental) values of the geometric parameters of molecular structures, as well as much more accurate values of thermodynamic and other physical-chemical parameters in comparison with other variants of the DFT method.

**Table 1** Bond lengths and bond angles in the CN<sub>12</sub> calculated by and DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods

Structural parameter	Calculated by		
	DFT B3PW91/TZVP	MP2/TZVP	MP3/TZVP
<b>Carbon–nitrogen bond lengths in the CN<sub>4</sub> node, pm</b>			
C1N1	149.7	151.0	148.6
C1N4	149.7	151.0	148.6
C1N7	149.7	151.0	148.6
C1N10	149.7	151.0	148.6
<b>Nitrogen–nitrogen bond lengths in macrocycle, pm</b>			
N1N2	132.1	132.4	133.0
N2N3	129.1	133.0	128.6
N3N4	132.1	132.4	133.0
N4N5	132.1	132.4	133.0
N5N6	129.1	133.0	128.6
N6N7	132.1	132.4	133.0
N7N8	132.1	132.4	133.0
N8N9	129.1	133.0	128.6
N9N10	132.1	132.4	133.0
N10N11	132.1	132.4	133.0
N11N12	129.1	133.0	128.6
N12N1	132.1	132.4	133.0
<b>Bond angles in the CN<sub>4</sub> node, deg</b>			
N1C1N4	90.0	90.2	90.3
N4C1N7	90.0	90.2	90.3
N7C1N10	90.0	90.1	90.3
N10C1N1	90.0	90.1	90.3
Bond angles sum (BAS), deg	<b>360.0</b>	<b>360.6</b>	<b>361.2</b>
<b>Non-bond angles in the CN<sub>4</sub> node, deg</b>			
N1N4N7	89.9	89.7	89.5
N4N7N10	89.9	89.7	89.5
N7N10N1	89.9	89.7	89.5
N10N1N4	89.9	89.7	89.5
Non-bond angles sum (NBAS), deg	359.6	358.8	358.0
<b>Bond angles in 5-membered cycles, deg</b>			
C1N1N2	116.7	117.1	116.8
N1N2N3	108.3	107.8	108.0
N2N3N4	108.3	107.8	108.0
N3N4C1	116.7	117.1	116.9
C1N4N5	116.7	117.1	116.8
N4N5N6	108.3	107.8	108.0
N5N6N7	108.3	107.8	108.0
N6N7C1	116.7	117.1	116.9
C1N7N8	116.7	117.1	116.8
N7N8N9	108.3	107.8	108.0
N8N9N10	108.3	107.8	108.0
N9N10C1	116.7	117.1	116.9
C1N10N11	116.7	117.1	116.8
N10N11N12	108.3	107.8	108.0
N11N12N1	108.3	107.8	108.0
N12N1C1	116.7	117.1	116.9



The calculations were carried out using the Gaussian09 program package.<sup>43</sup> As in our previous articles, in which this method of calculation was used,<sup>40–42</sup> the correspondence of the found stationary points to the energy minima in all cases was proved by calculating the second derivatives of the energy to the coordinates of the atoms; wherein, all equilibrium structures corresponding to the minimum points on the potential energy surfaces had only real (and, moreover, always positive) frequency values. Of the optimized structures for further consideration, the one with the lowest total energy was selected. Natural Bond Orbital (NBO) analysis was carried out, using NBO version 3.1, integrated with Gaussian09 program package<sup>43</sup> according to the methodology described in detail.<sup>44</sup> NBO methods are well known for excellent numerical stability and the convergence with respect to basis set expansion, sensibly proportionate to convergence of energy and other calculated wavefunction properties (unlike Mulliken analysis and related overlap-dependent methods in this case). Since the C(N<sub>12</sub>) structure maybe not trivial, a nontrivial electron density distribution in it is quite possible. As it is well known, the DFT methods when calculating structures with a nontrivial distribution of electron density in some cases can lead to an incorrect order of orbitals and even an incorrect molecular structure. To exclude such a possibility, we decided to carry out the calculation by other *ab initio* quantum chemical methods, too. However, unfortunately, it was not possible to complete the calculation with such methods with a stricter account of electronic correlation as CCSD and QCISD due to high

computational costs. That is why, in addition to the calculation by the DFT B3PW91/TZVP method, as an alternative, we used perturbation theory methods,<sup>45</sup> namely MP2 (ref. 46) and MP3 (ref. 47) in the combination with the extended split valence TZVP basis set, the each of which is noticeably less computationally intensive than the CCSD and QCISD methods. At the moment, it was also not possible to complete the calculation using the MP4 method, since in terms of computational complexity it is comparable to the CCSD method. The standard thermodynamic parameters of formation ( $\Delta H_{f,298}^0$ ,  $S_{f,298}^0$  and  $\Delta G_{f,298}^0$ ) for the C(N<sub>12</sub>) compound under examination were calculated using the G4 method described in ref. 48.

## Results and discussion

According to the data of each of the three above-mentioned methods of quantum-chemical calculation, a chemical compound of the composition C(N<sub>12</sub>) is capable of independent existence. The lengths of chemical bonds between atoms and bond angles for this chemical compound calculated by B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods, were presented in Table 1. As it can be seen from these data, all three methods give very close values of these key parameters of molecular structures, in connection with which it seems appropriate to discuss them together.

It is well known that in all those cases when a carbon atom is bonded to four other atoms through single bonds, a tetrahedral

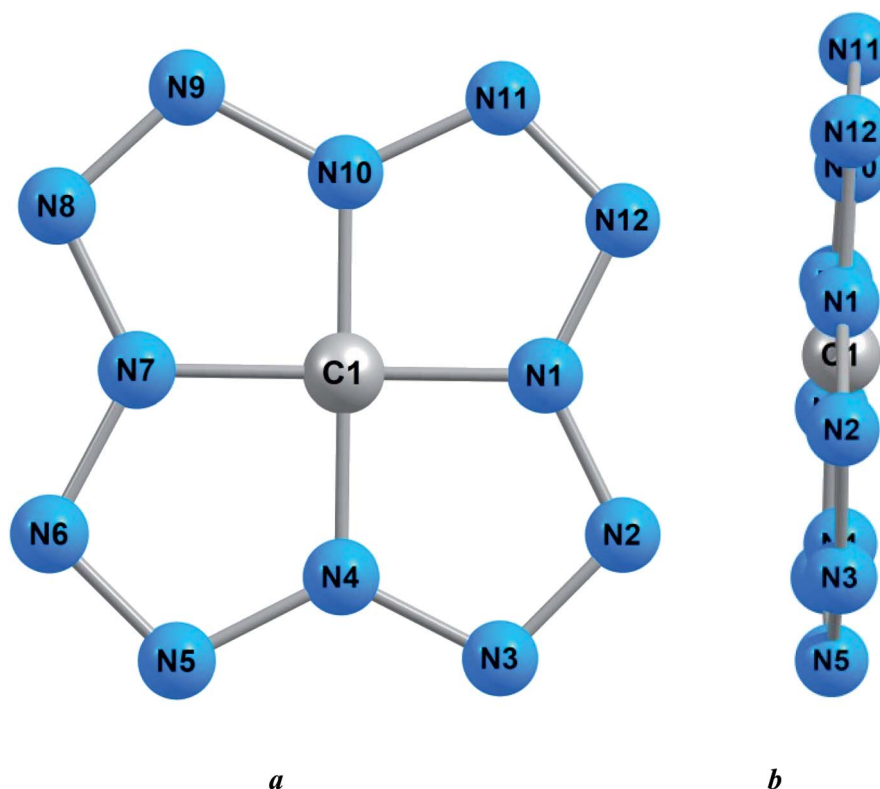


Fig. 1 Molecular structures of the C(N<sub>12</sub>) compound obtained as a result of DFT B3PW91/TZVP quantum-chemical calculation: (a) front view, (b) side view.

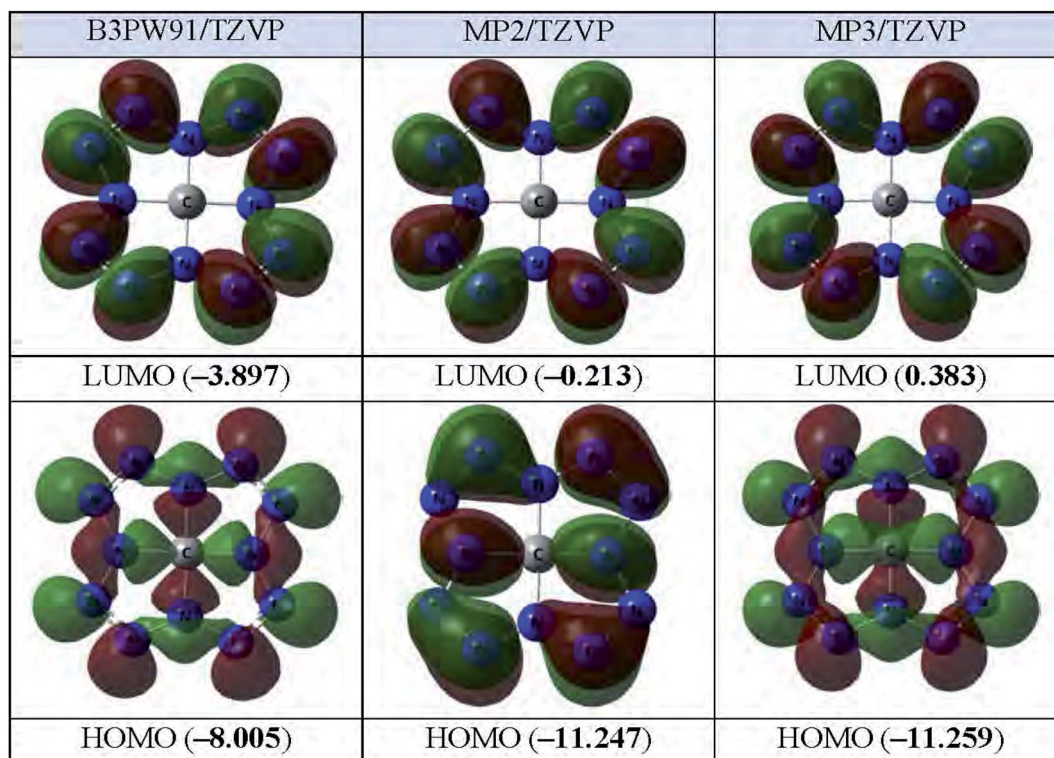


**Table 2** NBO analysis data for the  $C(N_{12})$  calculated by DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods

Calculation method	Effective charge on atom, in units of electron charge ( $\bar{e}$ )						
	C1	N1 (N4)	N7 (N10)	N2 (N6)	N3 (N5)	N8 (N12)	N9 (N11)
B3PW91/TZVP	0.2014	−0.0680	−0.0680	0.0089	0.0088	0.0088	0.0089
MP2/TZVP	0.3575	−0.1366	−0.1366	0.0237	0.0236	0.0235	0.0236
MP3/TZVP	0.4080	−0.1607	−0.1607	0.0294	0.0293	0.0293	0.0293

or close to it coordination of these same atoms relative to the carbon atom occurs. Taking this into account, one would expect that in the compound  $C(N_{12})$  with the above structural formula (\*), the nitrogen atoms bound to the C atom by single bonds should be located at the vertices of a regular tetrahedron or a polyhedron close to it. However, the calculation of the molecular structure of  $C(N_{12})$ , using each of these three quantum chemical methods, indicates that the group of four nitrogen atoms bonded to a carbon atom is *practically flat* [the sum of angles (N1N4N7), (N4N7N10), (N7N10N1) and (N10N1N4) differ from  $360^\circ$  by no more than  $2.0^\circ$ ]. What is even more remarkable, the grouping of  $CN_4$  atoms also turns out to be practically flat [the sum of angles (N1C1N4), (N4C1N7), (N7C1N10) and (N10C1N1) also differs from  $360^\circ$  by no more than  $1.2^\circ$ ] (Table 1). 5-Membered rings containing four nitrogen atoms and a carbon atom as a kind of “central atom” are also strictly planar since the sum of the internal bond angles in each

of them are the same and within the framework of each of the above calculation methods are  $540.0^\circ$ , that coincides with the sum of the interior angles in a flat pentagon ( $540^\circ$ ). Taking all this into account, it can be argued that the considered compound of carbon with nitrogen, according to the data of each of the quantum-chemical methods used by us, as a whole, has a molecular structure very close to coplanar. Each of the three quantum-chemical calculation methods used by us gives the same lengths of all four carbon–nitrogen bonds taking place in this compound, although, depending on the method, they differ slightly from each other (Table 1). As for the 12 chemical bonds formed by nitrogen atoms, they are subdivided within each of these methods into two groups containing four and eight bonds, respectively, with the same lengths within each of these groups. Wherein, for the molecular structures obtained by the DFT B3PW91/TZVP and MP3/TZVP methods, the bond lengths in the group of four bonds are shorter than the bond



**Fig. 2** The images of highest occupied (HOMO) and lowest unoccupied (LUMO) molecular orbitals in the  $C(N_{12})$  (ground state – spin singlet,  $M_S = 1$ ) according to DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods. The energies values of these molecular orbitals (in brackets) are given in eV.





**Table 3** Standard thermodynamic parameters of formation ( $\Delta H_{f,298}^0$ ,  $S_{f,298}^0$  and  $\Delta G_{f,298}^0$ ) for C(N<sub>12</sub>) calculated by B3PW91/TZVP and G4 methods

Calculation method	$\Delta H_{f,298}^0$ , kJ mol <sup>-1</sup>	$S_{f,298}^0$ , J mol <sup>-1</sup> K	$\Delta G_{f,298}^0$ , kJ mol <sup>-1</sup>
B3PW91/TZVP	1614.8	378.1	1842.2
G4	1581.5	374.3	1810.1

lengths in the group of eight bonds, while for the molecular structure obtained by the MP2/TZVP method, the opposite ratio occurs (Table 1). On the whole, the molecular structures of this compound obtained by these three methods show almost complete similarity with each other; an example of such a structure is shown in Fig. 1. The values of the dipole electric moment for this compound calculated using the each of DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP method, is practically no different from 0.00 Debye units, that, taking into account the practically coplanar structure of C(N<sub>12</sub>) obtained by each of these methods, seems to be quite expected.

Key data of NBO analysis and, namely, the values of effective charges on carbon atom and nitrogen atoms for macrocyclic compound under study obtained by DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods, are presented in Table 2.

As can be seen from these data, the values of the effective charges on individual atoms calculated by the above methods differ quite noticeably from each other, that seems quite natural because these methods take into account electron correlation in different ways. More reliable, in our opinion, are the values obtained using the DFT B3PW91/TZVP method. However, these values according to their module were considerably lesser than +4.000 $\bar{e}$  (for C1 atom) and -1.000 $\bar{e}$  (for N1, N4, N7 and N10 atoms connected with C1 atom by the chemical bonds) which would be in that case if all chemical bonds between C and N atoms would be ionic. This fact indicates that the compound under study has a very high degree of delocalization of the electron density.

The ground state of this compound is a spin singlet ( $M_s = 1$ ), and the values of the operator of the square of the proper angular momentum of the total spin of the system  $\langle S^2 \rangle$  in the framework of each of these methods are equal to 0. In addition, according to calculation data, the nearest excited state, which has a different  $M_s$  value, namely the spin triplet ( $M_s = 3$ ), is 363.7 kJ mol<sup>-1</sup> higher than the ground state (according to the G4 method). It should be noted, that testing the wave functions of the ground state for stability using the standard STABLE =

OPT procedure showed that the ground state wave function at  $M_s = 1$  is stable under the considered perturbations.

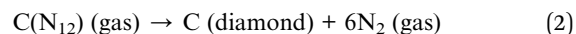
The images of the highest occupied and lowest vacant (unoccupied) molecular orbitals (HOMO and LUMO, respectively) obtained by each of these three calculation methods are presented in Fig. 2. As you can see from it, LUMO forms obtained by each of these three methods, are practically identical to each other; in this case, the electron density in the MO data is concentrated on the peripheral nitrogen atoms, while it is practically absent on the C1 carbon atom. As to HOMO, the above similarities are noted only for orbitals obtained by DFT B3PW91/TZVP and MP3/TZVP methods whereas HOMO obtained by MP2/TZVP method, has another form (Fig. 2). Despite this, the energies of these MOs obtained by the DFT B3PW91/TZVP and MP2/TZVP and MP3/TZVP methods differ quite significantly (Fig. 2).

The standard thermodynamic parameters of formation ( $\Delta H_{f,298}^0$ ,  $S_{f,298}^0$  and  $\Delta G_{f,298}^0$ ) for the chemical compound under examination were given in Table 3. As may be seen from it, all these parameters are positive, and, therefore, this compound, as it should be expected, cannot be obtained from the most thermodynamically stable simple substances formed by carbon and nitrogen (*i.e.*, graphite and molecular nitrogen N<sub>2</sub>).

It is of some interest to consider the reaction of the interaction of C(N<sub>12</sub>) with molecular oxygen, which proceeds in the gas phase according to the eqn (1)



and the reaction of decomposition of this compound according to the eqn (2)



Standard thermodynamical parameters for these reactions  $\Delta_r H_{298}^0$  and  $\Delta_r S_{298}^0$  are presented in the Table 4. As may be seen from these data, the reaction of the interaction of the C(N<sub>12</sub>) compound under consideration with molecular oxygen, on the

**Table 4** Standard enthalpies ( $\Delta_r H_{298}^0$ ) and standard entropies ( $\Delta_r S_{298}^0$ ) of the reactions (1) and (2) obtained from B3PW91/TZVP and G4 methods

Calculation method	Reaction			
	C(N <sub>12</sub> ) (gas) + O <sub>2</sub> (gas) → CO <sub>2</sub> (gas) + 6N <sub>2</sub> (gas)		C(N <sub>12</sub> ) (gas) → C (diamond) + 6N <sub>2</sub> (gas)	
	$\Delta_r H_{298}^0$ , kJ	$\Delta_r S_{298}^0$ , J K <sup>-1</sup>	$\Delta_r H_{298}^0$ , kJ	$\Delta_r S_{298}^0$ , J K <sup>-1</sup>
B3PW91/TZVP	-1806.8	788.0	-1613.0	823.7
G4	-1984.8	798.2	-1579.7	827.5



one hand, is strongly exothermic, since the  $\Delta_r H_{298}^0$  values for it are negative, and they themselves are very significant in modulus ( $\sim 2000$  kJ) within the framework of each of the calculation methods we used. On the other hand, the value of  $\Delta_r S_{298}^0$  is positive for it, and in accordance with the well-known Gibbs – Helmholtz eqn (3)

$$\Delta_r G^0(T) = \Delta_r H_{298}^0 - T\Delta_r S_{298}^0 \quad (3)$$

the  $\Delta_r G^0(T)$  value for it will be negative at any temperature, which in turn means that the process described by eqn (1) is irreversible. In this regard, it is interesting to note that a similar situation takes place for reaction (2) (Table 4), which, in principle, can be used to obtain an allotropic modification of carbon – a diamond in the framework of the isobaric process. And since the reaction (2) indicated in the article proceeds in the gas phase [in any case, the initial substance in it is  $C(N_{12})$  (gas)], then we can assume with sufficient confidence that the energy barrier to the activation of this reaction will not be too big and it will not be kinetically inhibited, which means that it can be realized within a reasonable time.

*A priori*, one would expect that the  $C(N_{12})$  compound considered in our article would be more stable than its isomeric compound, tetraazidomethane  $C(N_3)_4$ , described in ref. 16, since, as is known from organic and coordination chemistry, with the same composition of substances, as a rule, those that have a cyclic structure are more stable. Our calculation of the above parameters using the DFT B3PW91/TZVP method gave the values  $\Delta H_{f,298}^0 = 1350.5$  and  $\Delta G_{f,298}^0 = 1550.3$  kJ mol $^{-1}$ , using the G4 method, the values  $\Delta H_{f,298}^0 = 1293.0$  and  $\Delta G_{f,298}^0 = 1490.6$  kJ mol $^{-1}$ . Comparing this data with the data in Table 3, it can be found that, in the case of the first of these methods, the difference in the values of  $\Delta H_{f,298}^0$  is 265.3 kJ mol $^{-1}$ , in the case of the second one, 288.5 kJ mol $^{-1}$ , *i.e.* tetraazidomethane  $C(N_3)_4$  turns out to be a somewhat more stable compound than (dodecaazacyclododecantetraene-1,4,7,10)carbon(IV)  $C(N_{12})$ . These differences, in our opinion, are not very significant, so this circumstance in no way disproves the possibility of obtaining  $C(N_{12})$  as a result of any chemical reaction (and, possibly, even as a result of isomerization of tetraazidomethane).

## Conclusion

So, the data obtained using three different methods of quantum-chemical calculation, namely DFT B3PW91/TZVP, MP2/TZVP and MP3/TZVP methods, presented above, unambiguously predict the possibility of the existence of a new, so far unknown in chemical science, MACROTHETRACYCLIC compound of carbon with nitrogen composition  $C(N_{12})$ . Which, on the one hand, is characterized by an unusual (and so far the highest) ratio between the number of nitrogen atoms and the number of carbon atoms (12 : 1), on the other hand, it has an almost planar molecular structure. Wherein, the latter circumstance is also unusual, since it has long been well known that in those compounds where a carbon atom is bonded to four other

atoms through single chemical bonds, tetrahedral or close coordination of these atoms with respect to the carbon atom takes place. The noted results of quantum-chemical calculations, in our opinion, give all the grounds for a more careful study of such an unusual macrocyclic compound, and now it should confirm its existence experimentally since its synthesis can be important for the further development of physical chemistry and chemical technology of both these chemical elements. Playing a very important role in nature. Predicting the possibility of the existence of such an exotic chemical substance and modelling its molecular and electronic structures using modern quantum-chemical calculations, as it seems to us, can serve as a very useful tool in the decision of the problems associated with this synthesis.

## Author contributions

Conceptualization, O. V. Mikhailov; methodology, O. V. Mikhailov and D. V. Chachkov; software, D. V. Chachkov; validation, O. V. Mikhailov and D. V. Chachkov; formal analysis, O. V. Mikhailov and D. V. Chachkov; investigation, O. V. Mikhailov and D. V. Chachkov; resources, D. V. Chachkov; data curation, D. V. Chachkov; writing – original draft preparation, O. V. Mikhailov and D. V. Chachkov; writing – review & editing, O. V. Mikhailov; visualization, O. V. Mikhailov and D. V. Chachkov; supervision, O. V. Mikhailov; project administration, O. V. Mikhailov; funding acquisition, D. V. Chachkov.

## Conflicts of interest

The authors declare that they have no conflict of interest, financial or otherwise.

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