About the nature of the wave function and its dimensionality:  
the case of quantum chemistry

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Abstract: The problem of the $3N$ dimensions of the wave function is of particular interest in the philosophy of physics. In this work, we will recall the main positions about the nature and dimensionality of the wave function and we will introduce a new perspective, coming from quantum chemistry. For this, we will bring to light the formal operations that underlie the independent electron approximation. On this basis, we will point out how quantum chemistry can offer new arguments that contribute to the debate about the ontology of wave function.

1. Introduction

The wave function is a central element in quantum mechanics, since it represents the state of the system and participates in its dynamics through its evolution according to the Schrödinger equation. However, even today, almost a hundred years after the advent of quantum mechanics, the meaning of the wave function remains a matter of debate. In this context, the problem of the $3N$ dimensions of the wave function is of particular interest in the philosophy of physics. In fact, the debates around the issue have an important impact on the way in which we conceive the world around us. This is clearly manifested by the intense discussions that have taken place in recent years (see Monton 2006, Ney and Albert 2013).

From his early work, Schrödinger tried to endow the wave function with a physical meaning, first as a kind of vibration in the atom (Schrödinger 1926a) and later as a tool for obtaining the electron density (Schrödinger 1926b). With these proposals, Schrödinger intended to develop an ontology of the wave function in a space of three dimensions, in agreement with the world in which we live. However, some years later he was disappointed with those ideas.
In recent years, different views regarding the dimensionality of wave function have been proposed. On the one hand, it is possible to propose an ontology in which the wave function is the most relevant element of quantum theory, so that its mathematical nature is directly related to “reality” (e.g. Albert 2013). In spite of the fact we only perceive three dimensions, the authors who advocate for this position, commonly called “wave function realism”, consider that the real physical space has actually $3N$ dimensions. On the other hand, the wave function can be conceived as a mere mathematical artifact belonging to the formalism of quantum mechanics. The defenders of this view usually postulate a “primitive ontology” that inhabits a real space with only three dimensions (e.g. Monton 2013, Allori 2013). Finally, there is a third position that tries to reconcile the two previous ones. It proposes an ontological picture in which both the space of $3N$ dimensions and that of three dimensions coexist (e.g. Monton 2006). This discussion is still relevant not only in the philosophy of physics, but also in physics.

In this work we will introduce a new perspective, coming from quantum chemistry, an area of study arising from the direct interaction between quantum mechanics and molecular chemistry. In the field of quantum chemistry, the question about the $3N$ dimensions of the wave function has not been discussed as deeply as in the context of quantum mechanics. In fact, even the most theoretical objects of chemists exist in real space; hence, it is natural to try to turn the wave function into a three-dimensional entity. In the context of quantum chemistry, it is common to use the so-called orbital approximation, which allows chemists to write the total wave function of a system as a product of mono-electron wave functions (Atkins and de Paula 2006). Under this approximation, the wave function of an electron depends only on the variables of this electron; therefore, it evolves in the space of three dimensions (Lowe and Peterson 2006). We will consider how quantum chemists use this approximation, which can be conceived as a particular case of the so-called independent electron approximation. On this basis, we will show that it is possible to formalize the quantum chemists’ strategies as the result of the application of two mathematical operations: first, a projection in Hilbert space and, then, a change of variables. This formalization will allow us to go beyond the approximation itself by
propose a new version of the idea of a primitive ontology, now from the perspective of quantum chemistry.

For these purposes, we have structured the work in the following way. In Section 2 we will introduce a brief historical review of Schrödinger’s proposals on this subject. Then, in Section 3, we will recall the main positions about the nature and dimensionality of the wave function. We will continue with the detailed description of the orbital approximation, as a particular case of the independent electron approximation in Section 4. In Section 5, we will bring to light the formal operations that underlie the independent electron approximation. Finally, in Section 6, we will point out how quantum chemistry can offer new arguments that contribute to the debate about the ontology of wave function.

2. The Schrödinger’s wave function

In 1926 Schrödinger published a series of papers in which he introduced his theory of wave mechanics. He postulated the now well-known Schrödinger equation,

\[ i\hbar \frac{\partial}{\partial t} \Psi = \left[ -\frac{\hbar^2}{2m} \nabla^2 + V \right] \Psi \]  

where \( \Psi(\vec{r}_1, \vec{r}_2, ..., \vec{r}_N) \) represents the wave function, which depends on the spatial coordinates \( \vec{r}_i = (x_i, y_i, z_i) \) of all the particles of the system. Then, the wave function depends on \( 3N \) dimensions, where \( N \) is the number of particles in the system.

The dimensionality of the wave function was a problem that Schrödinger did not ignore, and he addressed it from the very beginning. Very early in his work, he wanted to give a physical meaning to the wave function by associating it with a vibration process in the atom (Schrödinger 1926a). However, he later pointed out that it is only in the case of a single electron that the interpretation as a vibration in real space of three dimensions can be sustained (Schrödinger 1926b). When the system has two or more particles, the wave function inhabits a space of six or more dimensions, so it cannot be conceived as a wave in the common three-dimensional space.

In the last of his articles of 1926, Schrödinger emphasizes the idea of dispensing with the wave function in the direct interpretation of physical phenomenon, relegating its use to obtain what he called the density of electricity (Schrödinger 1926c), a quantity that he
supposes does have a direct physical meaning and is defined in the space of three dimensions. The process by which he obtains this quantity is described in a letter to Lorentz:

\[ \psi \ast \psi \text{ (just as } \psi \text{ itself) is a function of } 3N \text{ variables or, as I want to say, of } N \text{ three dimensional spaces, } \vec{r}_1, \vec{r}_2, ..., \vec{r}_N. \text{ Now first let } r_1 \text{ be identified with the real space and integrate } \psi \ast \psi \text{ over } \vec{r}_2, ..., \vec{r}_N; \text{ second, identify } r_2 \text{ with the real space and integrate over } \vec{r}_1, \vec{r}_3, ..., \vec{r}_N; \text{ and so on. The } N \text{ individual results are to be added after they have been multiplied by certain constants which characterize the particles (their charges, according to the former theory). I consider the result to be the electric charge density in real space. (Schrödinger in Przibram 1967, page 56, emphasis of us)}

In this way Schrödinger offers a wave-function ontology based on electric charge density. However, later Schrödinger set aside this approach, and by 1935 expressed his regret in the following terms: “I am long past the stage where I thought that one can consider } \psi \text{-function as somehow a direct description of reality” (quoted in Fine 1996: 82).}

The aim of developing an ontology for the wave function in three dimensions, which motivated Schrödinger in the 1920’s, reappeared in the present-day discussions about the nature of the wave function

3. Positions about the nature of wave function and its dimensionality

The problem about the nature of the wave function and its dimensionality has been addressed in the light of questions such as: does the wave function have a real physical meaning or is it merely a mathematical artifact? If it has physical meaning, what does it represent? And, in that case, does the real physical space have \(3N\) dimensions or only the three that we perceive in our daily life?

In the recent philosophical literature, at least three positions can be identified regarding the dimensionality of the wave function: one that conceives the wave function as the basic ontological item in a \(3N\)-dimensional physical space; an opposite view that retains the three-dimensional physical space and deprives the wave function of a basic ontological role; and an intermediate position that admits the coexistence of both spaces.
Some authors, such as David Albert, defend an ontology where the wave function in $3N$ dimensions represents the physical reality. The problem of this position, commonly called “wave function realism”, is that the real ontology turns out to be radically different from the way in which the world around us is perceived. According to Albert, the appearances in the three-dimensional space are a dynamical result: ultimately, what it means to be an object in the real world “is to occupy a certain location in the causal map of the world”, and the form in which geometric appearances are constructed is due to a question of *dynamics* (Albert 2013: 54).

Peter Lewis (2013), in turn, although conceiving the wave function as the central element of quantum mechanics, considers that its dependence on $3N$ variables is only the most convenient way to represent the correlation between quantum particles. Lewis admits that the common use of the notion of *spatial* corresponds to a three-dimensional space. Nevertheless, he does not rule out the possibility that, in a future paradigm, the use of ‘*spatial*’ will change in favor of a notion corresponding to a $3N$-dimensional world.

The stances that relegate the wave function to a secondary role stand at the other end of the spectrum. This is the case of Bradley Monton (2013), who claims that the wave function can be defined by the properties of the system. From this perspective, the information about the system contained in the wave function is “carried” by the properties in the real space of three dimensions; hence, the wave function itself is not indispensable.

In the same end of the spectrum stand those positions that look for a *primitive ontology* as the reference of quantum mechanics. According to this view, any physical theory must try to account for the world in which we live, a world that is constituted by entities in a three-dimensional space. These entities are the foundation of the real world, and the theory must describe them and account for their time evolution. Then, the theory must include primitive variables associated with this ontology, and non-primitive variables that describe the way in which the former change over time (Allori 2013). In the context of this position, the wave function turns out to be a *non-primitive variable*, which describes the dynamics of particles, whose representative magnitudes are the *primitive variables*.

The third intermediate position is briefly outlined by Morton (2006), who introduces two variants. In the first one, two independent spaces coexist, one of $3N$ dimensions where
the wave function evolves, and the three-dimensional space where the $N$ particles evolve. What remains to be explained is the type of connection that links these two spaces: the problem is that, although there is a lawful relationship, in principle there is no causal link between them. In the second variant, the two spaces are hypersurfaces of a space of $(3N+3)$ dimensions, which might be the stage for a kind of causal connection, however it is still unclear what kind of connection could link the two spaces.

A common feature of all the arguments in the debate is that they rely on very abstract elements: the discussions are confined to the quantum mechanics as a formal theory. But if science is a practice, the way in which quantum mechanics is used should also be taken into account. But quantum mechanics is not used in the same way in different disciplines. In particular, it is interesting to consider how quantum chemists integrate quantum mechanics in the core of their theoretical practice in order to open new ontological questions.

4. A different approach from quantum chemistry

Schrödinger original view about the *electric charge density* was rapidly dismissed in the physicists’ community and, for this reason, it was never taken seriously into account in the philosophy of physics. But this is not the case in quantum chemistry, where it reappears under the name of ‘*electronic density*’ as the central element of the discipline.

The issue of the dimensionality of the wave function is not an object of debate in the field of quantum chemistry. Although both physicists and chemists deal with phenomena that occur in a three-dimensional space, physicists are used to appeal to very abstract entities that go beyond the features of our perceived physical world. By contrast, even the most abstract entities studied by chemistry inhabit the three-dimensional space. This particularity is inherited by quantum chemistry: molecules, its main object of study, are three-dimensional objects, and the links between their components are relationships in three-dimensional space. Therefore, it is no surprising to find that a basic theoretical move in quantum chemistry is to turn the wave equation into a three-dimensional entity.

In this theoretical background, the usual story proceeds as follows. Given the impossibility to apply quantum mechanics in an exact way to the study of chemical
systems, alternative approximate methods have been developed. The most popular of them is the so-called \textit{orbital approximation}. This is not a very precise approximation since it ignores electronic interactions. Nevertheless, it is a powerful tool for the study of molecular structure. As Peter Atkins and Julio de Paula point out in their famous textbook:

The wave function of a many-electron atom is a very complicated function of the coordinates of all the electrons, and we should write it \( \psi(\vec{r}_1, \vec{r}_2, \ldots) \), where \( \vec{r}_i \) is the vector from the nucleus to electron \( i \). However, in the \textit{orbital approximation} we suppose that a reasonable first approximation to this exact wavefunction is obtained by thinking of each electron as occupying its “own” orbital, and write

\[
\psi(\vec{r}_1, \vec{r}_2, \ldots) = \psi(\vec{r}_1)\psi(\vec{r}_2) \ldots
\]

We can think of the individual orbitals as resembling the hydrogenic orbitals, but corresponding to nuclear charges modified by the presence of all the others electrons in the atom. This description is only approximate, but it is a useful model for discussing the chemical properties of atoms, and is the starting point for more sophisticated descriptions of atomic structure. (Atkins and de Paula 2006: 336).

This approach is what justifies the fact that phrases like “the 2p orbital of a fluorine atom is compact” are common in the chemists’ discourse. According to quantum mechanics, the 2p orbital is an eigenfunction of the Hamiltonian of an atom with only one electron. Since a fluorine atom has 9 electrons, the 2p orbital is not an eigenfunction of its Hamiltonian. However, under the orbital approximation chemists can talk about the 2p orbital of a fluor atom and to make inferences with it.

The orbital approximation is very imprecise because it does not take into account the interaction between the electrons in the atoms. However, the procedure can be refined in the so-called \textit{independent electron approximation}, by taking into account the interactions but at the same time retaining of an orbital for each electron. Consider a chemical system of \( N \) electrons that interact with each other, and that are influenced by the Coulomb potential of the positive charges of nuclei. The Hamiltonian of that system is given by the following equation (in atomic units),

\[
H = -\frac{1}{2} \sum_{i=1}^{N} \nabla_i^2 - \sum_{i=1}^{N} V(\vec{r}_i) + \sum_{i=1}^{N-1} \sum_{j=i+1}^{N} \frac{1}{\vec{r}_{ij}}
\]
where the \( r_i \) are the coordinates of each electron of the system, the first term corresponds to the kinetic energy, the second term includes the traditional coulombic potentials \( V(r_i) \) due to the nucleus-electron attraction, and the third term represents the electric repulsion between each pair of different electrons, where the distance between them is \( r_{ij} = |r_i - r_j| \).

The difficulty to solve this equation lies in the third term, since it couples the electron motions. A way to solve this problem is to consider that each electron moves in an average potential due to the rest of electrons, in such a way that its motion is decoupled from the others; in this way, the Hamiltonian turns out to be a function of a single electron and the Schrödinger equation can be solved using independent wave functions \( \psi_i(r_i) \) for each electron. Precisely, the Hamiltonian takes the form

\[
H' = -\frac{1}{2} \sum_{i=1}^{N} \nabla_i^2 - \sum_{i=1}^{N} V(r_i) + \sum_{i=1}^{N} u(r_i)
\]  

(3)

Now, the new problem is to determine the form of the potential \( u(r_i) \) for each electron, a problem not trivial at all, since it involves fundamental information about the interaction between all the remaining electrons inside a chemical system. The solution to this difficulty has been the subject of intense research in the field of quantum chemistry. Just to mention two of the most important approaches: the Hartree-Fock method and the density functional theory (see Gill 1998). In the former, the wave functions \( \psi_i \) are used to obtain the potential \( u(r_i) \), while in the latter the electron densities \( \rho(r) \) is appealed to.

The simplest form to solve the problem is to consider that electrons do not interact with each other, so the term corresponding to the interelectronic interaction is neglected and the equation (3) becomes,

\[
H_{AO} = -\frac{1}{2} \sum_{i=1}^{N} \nabla_i^2 - \sum_{i=1}^{N} V(r_i) = \sum_{i=1}^{N} h(r_i)
\]  

(4)

where \( h(r_i) \) represents a hydrogen-like Hamiltonian. In this way, the orbital approximation is recovered. Since each mono-electron Hamiltonian \( h(r_i) \) corresponds to an hydrogenic ion, its eigenfunctions are the same as those of the Hamiltonian of the hydrogen atom, i.e., 1s, 2s, 2p, … etc., and are called atomic orbitals. It is interesting to stress that these atomic
orbitals are one of the foundations for the construction of the electronic configurations of the atoms according to the periodic table.

As it is quite clear, either when interelectronic interactions are considered or when they are neglected, the wave function of the total system, which depends on the coordinates of all the electrons, is approximated by combinations of monoelectron wave functions, each one of them is a function of the coordinates of a single electron. This shows that, long before arriving at the problem of interpretation the $3N$ dimensions of the wave function, quantum chemists face a calculation problem. Then, they intervene the theory, modify it: the wave function of $3N$ dimensions is transformed into $N$ wave functions of three dimensions in real space, one for each electron. As we will see, this strategy, which originally arises as a computational need, becomes the conceptual basis to conceive the nature of quantum-chemical systems.

5. Dimensional marginalization

In order to understand how the wave function is conceived by quantum chemistry, but now from the viewpoint of physics, the strategy described in the previous section will be mathematically formalized.

The wave function $\psi(\vec{r}_1, \vec{r}_2, \ldots, \vec{r}_N)$ of a molecule is defined in a space of $3N$ dimensions, since it is a function of the variables $\vec{r}_1, \vec{r}_2, \ldots, \vec{r}_N$. As usual in these cases we assume that the wave function is separable (the results can be generalized to the more general case):

$$\psi(\vec{r}_1, \vec{r}_2, \ldots, \vec{r}_N) = \psi(\vec{r}_1)\psi(\vec{r}_2) \ldots \psi(\vec{r}_N)$$  \hspace{1cm} (5)

However, in the independent electron approximation, this function is somehow projected onto a three dimensional space. Indeed, according to the standard procedure, it is said that the function $\psi(\vec{r}_i)$ corresponds to the orbital occupied by electron $i$, and this orbital is represented in the physical space of three dimensions. This means that the label that identifies the electron $i$ is transferred from the variable to the function

$$\psi(\vec{r}_1) \rightarrow \psi_1(\vec{r})$$  \hspace{1cm} (6)
Therefore, a change of variables is introduced, which makes possible to represent the different orbitals in the same physical space.

From a mathematical viewpoint, this procedure can be formalized as the result of the successive application of a projector $\hat{P}_i$, followed by an operator $\hat{C}$ that changes the variables to mono-electron wave functions:

a) In the first step, a projector $\hat{P}_i$ is defined as an operator that selects the mono-electron wave function that corresponds to electron $i$ and eliminates the rest. This can be done by means of the following mathematical operation:

$$\hat{P}_i\psi(...)=\int ... \int \psi(...) \psi^*(\vec{r}_1)d\vec{r}_1 ... \psi^*(\vec{r}_{i-1})d\vec{r}_{i-1} \psi^*(\vec{r}_{i+1})d\vec{r}_{i+1} ... \psi^*(\vec{r}_N)d\vec{r}_N$$

(7)

where $\psi(...): \psi(\vec{r}_1, ... , \vec{r}_{i-1}, \vec{r}_i, \vec{r}_{i+1}, ... , \vec{r}_N)$. Then, the application of $\hat{P}_i$ to $\psi(\vec{r}_1, \vec{r}_2, ... , \vec{r}_N)$ gives the desired result:

$$\hat{P}_i\psi(\vec{r}_1, \vec{r}_2, ... , \vec{r}_N) = \psi(\vec{r}_i)$$

(8)

b) In the second step, the operator $\hat{C}$ can be defined as:

$$\hat{C}\psi(\vec{r}_1) = \psi_1(\vec{r})$$

$$\hat{C}\psi(\vec{r}_2) = \psi_2(\vec{r})$$

$$\vdots$$

$$\hat{C}\psi(\vec{r}_N) = \psi_N(\vec{r})$$

(9)

Thus, the successive application of $\hat{P}_i$ and $\hat{C}$ describe the implicit operation in the independent electron approximation:

$$\hat{C}\hat{P}_i\psi(\vec{r}_1, \vec{r}_2, ... , \vec{r}_N) = \psi_i(\vec{r})$$

(10)

Since this operation extracts a mono-electron wave function from the total wave function of $3N$ dimensions and projects it onto a three dimensional space, it can be called dimensional marginalization.

6. Toward an ontology of quantum chemistry
As explained in the previous sections, although according to quantum mechanics the wave function of a molecule cannot be expressed in terms of just three variables, in the field of quantum chemistry endowing each electron with its own wave function is a standard procedure that lies at the very core of the discipline and, as a consequence, has more implications than a mere approximation.

When trying to solve the Schrödinger equation applied to chemical systems, quantum chemists use the orbital approximation for simple cases, and the independent electron approximation for more complex situations. By means of these strategies, the energy levels in chemical systems can be computed; besides this, they make possible to describe the electronic structure in systems of relative complexity.

According to their names, these strategies used by quantum chemists are mere approximations. As such, they should be only formal tools designed to obtain approximate solutions of an equation that cannot be solved with complete precision. As Norton (2012: 207) says, “approximations merely describe a target system inexacty”; they do not even carry the novel semantic import carried by idealizations. In Frigg and Hartmann’s words, “approximation is a purely formal matter” that is introduced in a mathematical context (2017: 8). As a mere tool, the effectiveness of an approximation lies in the adequacy of the results that it allows to obtain. If the practical or formal obstacles to obtain the exact solution were overcome, the approximation could be removed. To the extent that approximations are in principle eliminable, they carry no conceptual import: they do not discover new properties of the target nor supply new elements for explanations.

However, the above characterizations do not apply to the independent electron approximation used in quantum chemistry. The idea that each electron can be described by its three-dimensional wave function is at the core of the discipline and shapes the quantum-chemical picture of the molecule: a structure given by the geometrical disposition of the nuclei, and electrons that can be conceived as individuals that “occupy” the orbitals, identified by the wave functions of the hydrogen atom (1s, 2s, etc.). The orbital approximation is an approximation since it ignores the interaction between the electrons of this quantum-chemical target. But even when the interactions are reintroduced, the global wave function of 3N-dimensions is not recovered: the target is still described in terms of
monoelectronic wave functions. In this case, quantum chemists use to talk about, say, how the 1s orbital was modified with the new correction, but the idea of each electron with its orbital is not set aside. In the context of this picture, the Hartree-Fock method and the density functional theory are only approximations to compute the potential that each electron “sees” in the presence of the rest of the electrons of the molecule. But even if the computational obstacles could be removed and the exact inter-electronic interactions could be computed, this would not remove the assumption that the behavior of each electron is described by its own wave function, now modified accordingly.

This quantum chemical picture also plays an essential role in the explanations of the processes that occur in atoms and molecules. For example, in order to explain that the SF$_6$ compound exists while the OF$_6$ does not, although both oxygen and sulfur have the same external electronic structure (two s-electrons and four p-electrons), it is usually argued that the difference is that the sulfur is in the third period while the oxygen is in the second period of the periodic table. Let us see the argument. Sulfur has two electrons in a 3s orbital and four in two 3p orbitals; therefore, it has a free 3p orbital, but also has five free 3d orbitals, and they allow it to extend its valence shell to be bonded with many fluorine atoms. In the case of oxygen, the external electronic structure consists in two electrons in a 2s orbital and four in two 2p orbitals. But the difference is that there are no 2d orbitals, so oxygen only has one free orbital (one 2p-orbital). Then it can only be linked to 2 fluorine atoms and the OF$_6$ does not exist while the OF$_2$ does. Explanations of this kind do not make any sense from the physical point of view, because according to quantum mechanics there are not mono-electronic wave functions, but only multi-electronic wave functions in a space of $3N$ dimensions. However, even if considered approximate, they offer a clear understanding of many molecular phenomena: the chemistry of molecules could not exist without them. And the scientific status of these studies cannot be denied in the light of their empirical success not only in describing and predicting phenomena, but mainly in creating new substances.

The above considerations point to the fact that, actually, the so-called ‘independent electron approximation’ is not a mere approximation, but is at the basis of a conceptual framework specific of quantum chemistry. In this framework, the notion of orbital as
monoelectronic wave function plays a central role, since it is indispensable in the
descriptions of the electronic structure in atoms and molecules and the explanation and
production of molecular phenomena. This point to the fact that, in the light of the empirical
success of quantum chemistry, it is not easy to conceive the description in terms of
monoelectronic wave functions as a mere fiction that could be left aside in favor of a more
precise description of the molecular realm. On the contrary, it should be accepted that there
is a quantum-chemical ontology, that is, the particular reference of quantum chemistry, that
is essentially different than the ontology of quantum mechanics (whatever it be) (see

Now, the question is: what is the relationship between the quantum-chemical
ontology and that based on pure quantum mechanics? According to a traditional
reductionist view, in spite of its scientific efficiency, the quantum-chemical ontology is an
appearance arising from a coarse description of the quantum world. However, the
reductionist position faces several difficulties, in general derived from the fact that
incompatible assumptions, coming from structural chemistry and quantum mechanics,
coexist in quantum chemistry and constitute the theoretical body of the new discipline
(Lombardi 2014, Fortin, Lombardi, and Martínez González 2017, 1018). If the
antireductionist stance is adopted, then the quantum-chemical ontology can be conceived as
the primitive ontology of quantum chemistry, different from the primitive ontology
corresponding to the quantum realm (see Allori 2013, Esfeld, Lazarovici, Hubert, and Dürr
2013, egg and Esfeld 2015). But the full discussion of the issue of the intertheory relation
in this case is beyond the limits of the present article, and will be addressed in a future
work.

References


