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INTERDISCIPLINARY APPLICATIONS OF PAULING’S METALLIC ORBITAL AND UNSYNCHRONIZED RESONANCE TO PROBLEMS OF MODERN PHYSICAL CHEMISTRY: CONDUCTIVITY, MAGNETISM, MOLECULAR STABILITY, SUPERCONDUCTIVITY, CATALYSIS, PHOTOCONDUCTIVITY AND CHEMICAL REACTIONS

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\textbf{ABSTRACT:} The resonating valence bond theory, emphasizing unsynchronized resonance and metallic orbitals, introduced by Pauling 50 years ago, has become an easily visualized tool for describing some electron transfer processes. It has yielded simple, qualitative and useful descriptions of a wide variety of phenomena including the stability and geometry of molecules, conductivity, photoconductivity, superconductivity, magnetism, surface catalyzed reactions, chemical carcinogenesis as well as the stability of the metallic state.
I. Introduction

Linus Pauling’s resonating valence bond (VB) theory, hereafter denoted RVB, yields simple consistent explanations of several molecular phenomena, and synchronized resonance has become very popular, appearing in most chemistry textbooks. Although less familiar, unsynchronized resonance of covalent bonds through a metal has also proved to be a very useful concept for understanding electrical conductivity and other metallic properties.

General VB theory admits not only various spin-coupling schemes that label regions of a molecule or of a crystal where chemical bonds are formed, but also changes of orbital configuration arising from transfer of electrons between orbitals such that some atoms become formally negatively charged and others positively charged. The RVB theory considers that two electrons with opposite spins and occupying orbitals on adjacent atoms form a covalent bond, which may have some ionic character, and that these covalent bonds resonate among alternative positions in a metal.

Consider Pauling’s original lithium example. Much more important to the stabilization of a Li crystal than synchronized resonant structures (Fig. 1) are unsynchronized resonant structures such as shown in Fig. 2 in which one bond resonates independently from one position to another. Such unsynchronized resonance requires, however, that the atom receiving a bond (Li+ or Li) have an orbital available for receiving the electron. It is the possession of this orbital, the metallic orbital, in addition to the orbitals required for occupancy by unshared electron pairs and
bonding electrons, by all or many of the atoms in the condensed phase that makes possible
unsynchronized resonance of covalent bonds and gives rise to metallic properties. By a
succession of shifts by single bonds, a negative charge is seen to move in the appropriate
direction from atom to atom in a crystal\textsuperscript{6}.

Fifty years ago Pauling identified the metallic orbital in describing the nature of transition
metals\textsuperscript{1}. In 1987 he employed the concept of metallic orbital and the same statistical arguments
to explain superconductivity in some copper oxide compounds\textsuperscript{4}. The present authors recent VB
and molecular orbital (MO) calculations showed that the metallic orbital and unsynchronized
RVB structures yield useful descriptions of a wide variety of phenomena\textsuperscript{5-22}. This paper presents
a concise review of RVB theory with new applications based on both ab initio and semiempirical
calculations. In Sec. II we describe conductivity in metals and organic polymers,
photoconductivity in nickel dihalides, superconductivity in YBa\textsubscript{2}Cu\textsubscript{3}O\textsubscript{7-\delta}, magnetism of iron,
dissociation of diatomic molecules on 3d-transition metal surfaces, O\textsubscript{4} formation, and chemical
carcinogenesis—all in the RVB formalism\textsuperscript{5-22}. Computational details are given in Sec. III.
Conclusions follow in Sec. IV.

\section*{II.1 Generality of RVB Theory}

Pauling’s idea was based on empirical arguments when he demonstrated the value of
unsynchronized resonant structures as a tool for understanding the properties of metals.
However, we can also show that RVB theory works using ab initio calculations. In the lithium
case, consider, for example, a Li\textsubscript{4} cluster with D\textsubscript{2h} symmetry (Figs. 1 and 2). Labeling the
orbitals of atom 1 as 1 (normal orbital, occupied in common Kekule covalent or ionic structures)
and 1\textsuperscript{'} (metallic orbital, occupied only in Pauling structures) and so on, the structures can be
represented as ordered sequences of numbers (labels of the occupied orbitals). A Kekule covalent structure, for example, with bonds 1-3 and 2-4 would then be represented as:

$$1324 = A[\phi_1(r_1) \phi_2(r_2) \phi_3(r_3) \phi_4(r_4)] \theta$$  \[1\]

A typical Pauling structure, say 133´4, would then be written

$$133´4 = A[\phi_1(r_1) \phi_3(r_3) \phi_3(r_3) \phi_4(r_4)] \theta$$  \[2\]

in which the 1-3 bond involves the normal orbitals of atoms 1 and 3 but the 3´-4 bond involves the metallic orbital of atom 3 and the normal orbital of atom 4. Here \(\theta\) denotes a \(S = 0\) spin wavefunction, (cf, ref. 5). RVB calculations of the Li\(_4\) D\(_{2h}\) cluster show that the binding energy per atom increases from 1.21 eV to 2.18 eV as, in addition to the synchronized resonant structures, i.e., Kekule structures (Fig. 1), unsynchronized resonant structures, called Pauling structures (Fig. 2), are introduced.

In addition to predictions of the behavior of general metallic clusters, these RVB calculations yield a finding that links conformation, electronic structure, and stability. The finding may be stated: As Pauling structures enable each atom to make a covalent bond with all its neighbors, the system will seek a conformation that makes the bonds as equal in length as possible, leading to more metal-metal pathways and to increase of the coordination number of the atoms. This finding applies particularly well to even \(n\) systems, for which all the electrons can be paired in covalent links, and immediately predicts high stability for \(n = 6\) in a plane and \(n = 8\) in three dimensions. It has been shown that among the planar alkali clusters, \(n = 6\) should be a magic number\(^5\). Our first ab initio VB calculations on lithium clusters with explicit use of unsynchronized resonance structures confirmed the importance of the metallic orbital and the

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5. Reference number for magic number discussion.
covalent character of the metal-metal bond. Pauling structures for Li$_4$ substantially stabilize the ground state electron structure, and the pivotal switch of their bonds does provide a VB representation for electron conduction.

Harcourt and co-workers have done some important work using the VB formalism. These authors provide an alternative representation which involves one-electron transfer into antibonding MOs. This formulation, which can now be somewhat modified, might also be considered. For the Li($^+$) Li–Li$^-$–Li* Pauling structure, the (truly) divalent Li($^-$) uses orthogonal s and p atomic orbitals (AOs) to form the two bonds. Either Heitler-London type wavefunctions (as in Ref. 5) or (essentially) orthogonal localized MOs can then be used to formulate the wavefunctions for electron-pair bonds.

We have also recast the RVB theory in a nonorthogonal ab initio VB formalism and applied it to the study of small anionic lithium clusters. The Pauling structures have large weight compared to the Kekule structures, which emphasizes the importance of RVB theory in the VB description of these systems. The preference of small anionic clusters for linear geometries and the influence of the extra electron have a clear connection to Pauling structures.

RVB theory can also describe conductivity in some organic polymers. Polyacetylene, (CH)$_x$, is a particularly important example in which the degenerate ground state should lead to important soliton excitations in which the $\pi$ band is half-filled, implying the possibility of metallic conductivity. In conjugated polymers, carbon sp$^2$ hybridization leads formally to one unpaired electron per carbon atom. Consequently, the electronic structure is determined by chain symmetry (i.e., the number and kind of atoms in the repeat unit) with the result that such polymers can be expected to exhibit semiconducting and possibly metallic properties. A chain
of atoms (or more generally, monomers made up of groups of atoms) can be dimerized in two distinct patterns that have the same energy. Labeling the atoms (or monomers 1,2,3, . . .), one could have the bonding structures 1=2–3=4–5=6 or 1–2=3–4=5–6 denoted as A and B phases. Clearly, if all monomers are equivalent and the chain is infinite in extent, the energies of these bonding structures are the same. In the language of quantum field theory, the vacuum state is two-fold degenerate. The domain wall is a nonlinear shape-preserving excitation which propagates freely called a soliton. A moving soliton converts A-phase material into B-phase material (and vice versa). The undoped trans-(CH)$_x$ resembles a semiconductor with a band gap but through doping the electronic properties can be controlled\textsuperscript{30}. At dilute doping levels, the intrinsic resistivity of the material dominates but heavy doping can lead to a regime of high conductivity in which the amount of unsynchronized resonance becomes so great as to make the conducting state more stable than the insulating state. The electrical conductivity in trans-(CH)$_x$ can be rationalized in terms of unsynchronized resonating structures (Fig. 3.) In Figure 3, the apparently pentavalent carbon carbon atoms in the second and third VB structures arise only if two non-orthogonal localized MOs are used to accommodate the four $\pi$ electrons. The types of unsynchronized resonance used for the two cases differ, see for example refs. 24-26.

In the presence of an applied electromagnetic field, electrons tend to move from atom to atom. By a succession of bond shifts, negative charge moves toward the anode. In a similar way positive charges (electron holes) move toward the cathode. The probability of resonance is small for a chain of atoms with alternating single and double bonds - there are only two structures. As the sequences are interrupted, for example, by oxidation (or reduction) of a carbon center, the probability of unsynchronized resonance is increased. The mobile charge carriers introduced in the $\pi$-electron system are responsible for the electrical conductivity of conducting polymers. It
must be noted, however, that charge transport is limited by microscopic defects and sample imperfections.

We now consider RVB in describing insulating materials. Mott-Hubbard theory predicts that the band gap in insulating transition metal compounds involves transitions of the type d_i^6d_j^{10} \rightarrow d_i^{6-1}d_j^{11}, where i and j label transition metal sites. We note, however, that for these materials, a unified theoretical description of the observed band structure has not been achieved. The 3d orbital interaction is at the center of the controversy. Other authors have interpreted the optical absorption gap in terms of a p \rightarrow d transition, an interpretation which involves only one transition metal atom. In fact, photoconductivity experiments in nickel dihalide compounds (NiCl_2, NiBr_2, and NiI_2) indicate an optical gap corresponding to p \rightarrow d charge-transfer transitions. By applying RVB theory to describe the photoconductivity in nickel dihalide compounds, it is possible to unify these interpretations. Based on the band structure and on the nature of the chemical bond of insulating nickel difluoride constructed using the s, p, d Multiple Scattering Xα (MS-Xα) spin-polarized orbitals of an octahedral (NiF_6)^4^- cluster, unsynchronized resonance of the covalent bonds can be depicted as follows:

$$M-X \rightarrow M^+ + X^-$$ [3]

where X represents F, Cl, Br, and I. As required for unsynchronized resonance of covalent bonds, the atom receiving a bond must have an orbital available for electron reception. MS-Xα calculations indicated that this orbital can be identified as the unoccupied metallic 3d:e_g spin minority molecular orbital (the 4e_g molecular orbital in the NiF_6^-4 cluster has large Ni:3d character). The p \rightarrow d charge transition creates the M^+ state Ni(3d^9) as well as the hole in the p band. Therefore, the structure required for unsynchronized bond resonance (M^+ X – M^-) is
attained when the photohole is transferred from the anion p band to the 3d state of the nearest-neighbor nickel ion, creating the M$^+$ state Ni(3d$^7$). The process can be described as:

$$3d_i^8 3d_j^8 \rightarrow 3d_i^9 L 3d_j^8 \rightarrow 3d_i^9 3d_j^7$$  \[4\]

where L is a ligand hole. The creation of the M$^+$ and M$^-$ charge polarized states is a charge-transfer process in which one electron moves from the ligand to the metal and a hole moves from the ligand to another nickel in the opposite direction. The photohole excited state 3d$^9$ 2p$^5$ 3d$^8$ decays to the 3d$^9$ 2p$^6$ 3d$^7$ state. The global process is equivalent to the shift of a covalent bond from one nickel to another in the crystal, making RVB equivalent to Mott-Hubbard theory in the sense that two metal atoms are involved in the electron transfer. There is a resonance from 2Ni$^{+2}$ to Ni$^+$ and Ni$^{+3}$. A direct consequence of this interpretation is that the magnitude of the band gap is determined by

p $\rightarrow\square$ d charge-transfer energy, i.e., there is a dependence of the band gap on the electronegativity of the ligand and it is possible to reduce the conductivity gap by an appropriate choice of ligand. The difference between the insulator NiO and the conductor NiS, for example, is that in the insulator the electrons are localized in the 2p band and require energy for the first step of the unsynchronized resonant-valence-bond mechanism, while for metallic NiS the first step is facilitated by itinerant electrons of the hybridized Ni(3d) and S(3p) band, with no charge transport gap$^6$.

It has been noted that (NiF$_6$)$_{4^-}$ with a $^3A_{2g}$ ground-state is a hypoligated transition metal complex$^{27}$. A VB description for this complex involves resonance between 15VB structures, each of which involves four (polar) Ni-F electron-pair bonds (each as ($\sigma$NiF$^2$) ($\sigma^*$NiF$^0$)), using a simple MO formulation of the wavefunction for the electron-pair) and two Ni-F Pauling three-
electron bonds (each as \((\sigma_{\text{NiF}^2}) (\sigma_{\text{NiF}^1})\)). The two (parallel-spin) odd-electrons occupy the two antibonding MOs. With this type of VB description, perhaps the photoconductivity could also be formulated via the transfer of an electron from an antibonding \(\sigma^*\text{NiF} \) MO located around an adjacent metal center\(^{27}\).

II.2. Superconductivity

In BCS superconductivity theory, the interaction of electrons and phonons causes a clustering of electrons that move in phase with a phonon when the energy of this interaction is greater than the phonon energy\(^4\). In RVB theory the electronic structure of the metal is described in successive steps involving integrals of decreasing magnitude. At first, two electrons with opposite spins and occupying orbitals on adjacent atoms form a covalent bond, which may have some ionic character. Second, these covalent bonds resonate among alternative positions in the metal. RVB theory is in many ways equivalent to the conventional band theory of metals, but it has the advantage that it far more easily permits discussion of the dependence of metal properties on the identity of the constituent metal atoms. Pauling\(^4\) suggested that for the superconductor La (\(T_c = 6\)K) the rate of \(\text{M}^- \text{M}^0 \rightarrow \text{M}^0 \text{M}^- \) (electrons moving to the right on the crest of the phonon), is greater than that of the \(\text{M}^+ \text{M}^0 \rightarrow \text{M}^0 \text{M}^+ \) (holes moving to the right on the trough of the phonon) because the exchange integral is smaller for the motion of the hole as a result of the smaller size of \(\text{M}^+ \) relative to \(\text{M}^- \). This characteristic causes holes to fall behind electrons in following the phonons.

The mechanism of superconductivity of copper oxide compounds can be described by the theory of unsynchronized resonance of covalent bonds. Although electron pairs can move along a string of atoms in synchronism, if some O atoms are missing, so that vacancies interrupt the
sequence, then there would be the possibility of unsynchronized resonance. If the segments are short enough, the amount of unsynchronized resonance can become so great as to make the conducting state more stable than the insulating state, as is the case for metals. Suitable doping may be required and the interactions with La (Ba, Sr) are also essential for high-temperature superconductivity. The critical temperature \( T_c \) increases as scattering of electrons by phonons decreases, i.e., as the strength of the electron-phonon interaction decreases.

In summary, several structural features, including electron transfer between atoms of different electronegativity, oxygen deficiency, and unsynchronized resonance of valence bonds, as well as tight binding of atoms, cooperate to confer metallic properties and high-temperature superconductivity on copper oxide superconductors. The superconducting transition temperature decreases considerably when group VIII, IB and IIB metal ions are substituted for Cu ions. We have, in collaboration with other authors\(^7-^9\), investigated the substitution of CuO by NiS in these superconductors and reported x-ray diffraction, resistivity, and ac magnetic susceptibility findings for a set of samples with nominal composition \( \text{YBa}_2\{(\text{Cu}_3\text{NiS})_x\}\text{O}_{4-\delta} \). We found a small increase in \( T_c \) and a decrease of the room temperature resistivity in a certain substitution of CuO by NiS. These results seem to agree with RVB theory which predicts increasing \( T_c \) as unsynchronized resonance increases, which is the case when semiconducting CuO rings are substituted for conducting NiS rings. Our fluorescence x-ray emission spectra show that sulphur atoms enter the lattice and form oxyanion groups\(^9\).

**II.3. Magnetism in Iron**

A theory of superconductivity should explain the Meissner effect, which shows that a superconductor is a perfect diamagnetic system; magnetic flux is excluded from all but a thin
penetrable region near the surface. A complete quantitative treatment of the magnetic behavior of transition metals does not exist. Magnetism in Fe, Co, and Ni is problematic because 3d electrons are neither completely localized onto single centers nor entirely delocalized. Both the localized Heisenberg picture and the itinerant-electron theory developed by Stoner, implemented in connection with band theory, have been invoked to explain different aspects of the observed magnetic properties of Fe, Co, and Ni\textsuperscript{10}. Further, molecular field theory, either from the view of localized electrons or itinerant electrons, has been unable to satisfactorily explain the observed magneton number. Over the past decade a large number of experimental and theoretical studies have been performed on small metallic clusters, including bimetallic systems, which are of great interest in surface physics and chemistry, nucleation, heterogeneous catalysis, alloy formation and magnetism.

For the magnetic 3d transition metals, Fe, Co, and Ni, systematic studies by various authors show serious discrepancies with experiment. A long-standing goal is better comprehension of the origin of magnetism in these metals. For iron, in particular, chemical properties cannot be divorced from magnetic properties, since ferromagnetism leads to shifts on the order of an electron volt or more in energy levels due to spin polarization. Pauling also made important contributions to the phenomenon of magnetism in transition metals and proposed an interesting correlation between the magnetic properties of transition metals and the bonding of 4s and 3d electrons, metallic orbitals, and resonance of covalent bonds. The present authors\textsuperscript{10} reported ab initio Hartree-Fock calculations on the quintet spin state of Fe\(_2\) in a study of the electronic, magnetic, and chemical bonding properties as well as the hyperfine interactions. Good agreement was found with band-theoretical and experimental parameters. The calculations support Pauling’s model of occupation and bonding of valence orbitals, resonance of covalent
bonds, and conduction-band metallic orbitals. The calculated magneton number was found to be in good agreement with experiment.

II.4. O$_4$ Molecule

Various experimental and theoretical studies of O$_4$ lead to different descriptions of its stability and geometry (see Ref. 18 and references contained therein.) The present authors carried out ab initio MCSCF (multi-configuration self consistent field) calculations to clarify discrepancies regarding the structure and stability of this molecule. The study indicated a stable O$_4$ molecule. The system is found to be more stable for D$_{2h}$ symmetry than for D$_{4h}$ symmetry, in agreement with experimental studies of the O$_2$ dimer$^{18}$. 

Further support for D$_{2h}$ symmetry for O$_4$ is provided by Pauling's RVB theory. An easy visualization of O$_4$ formation is as a two step charge transfer process involving two O$_2$ molecules (see Fig. 4).

The first step of the mechanism is O$_2$ $\rightarrow$ O$_2$ electron transfer with corresponding bond transfer from one O atom of one O$_2$ molecule (which acquires a positive charge) to an O atom (which acquires a negative charge) of the other O$_2$ molecule. RVB theory requires that the atom receiving a bond must have an orbital available for electron reception. In the case of O$_2^-$, this orbital is identified with the $2\pi^*$ antibonding molecular orbital. In this process, the first electron transfer creates the O$_2^+ -$ O$_2^-$ bond. The O — O bond of the O$_2^-$ unit is weakened due to the occupation of the $2\pi^*$ antibonding orbital while the O — O bond of the O$_2^+$ unit is stronger due to the loss of an electron from an antibonding MO. The second step of the mechanism is O$_2^-$ $\rightarrow$ O$_2^+$ electron back-donation. This second interaction is similar to the first one in the sense that for both interactions the orbital receiving the electron is a $2\pi^*$ antibonding
MO. In this RVB mechanism the electron completes a cycle and forms new bonds. The resulting O$_4$ molecule must have D$_{2h}$ symmetry since the bonds between the two O$_2$ units, O$_2$—O$_2$, are different from the O—O bonds in each O$_2$ unit. This prediction of D$_{2h}$ symmetry by RVB theory is in agreement with experiment and with MCSCF calculations.$^{18}$

More recent work$^{28-29}$ yields additional details of O$_4$ formation. The O$_2$ structures have S = 0 spin excited states and S = 1 spin ground states and the O$_4$ VB structure does not retain double-bond character for the O$_2$ moieties. The (Linnett-type) VB structures 6 and 7 of ref. 28 with two Pauling three-electron bonds ($\pi_x$)$^2$(\$\pi^*_x$)$^1$ and ($\pi_y$)$^2$(\$\pi^*_y$)$^1$ for the six $\pi$ electrons, and parallel spins for the two antibonding electrons, are the appropriate VB structures for the O$_2$ ground state. The primary dimerization$^{28}$ process involves the spin pairings of the antibonding unpaired electrons to generate an increased-valence structure for D$_{2h}$ O$_4$ with fractional intermolecular bonds. This spin pairing process alone, however, does not stabilize the dimer relative to the two separated O$_2$ monomers. Although the VB calculation which are reported in ref. 29 are not necessarily definitive, they do indicate that dispersive interactions, with some help from charge-transfer, are primarily responsible for the small stabilization of the dimer. Attention has been focused on the importance of dispersive interactions by others; see, for example, refs. 2 and 3 of ref. 29.

II.5. Dissociation of Diatomic Molecules on Transition Metal Surfaces

Heterogeneous catalysis serves today as the basis for most petroleum and chemical technologies and has substantial importance for the life sciences as well as for environmental protection. It
has been intensively investigated with the tools of physics, quantum chemistry, and surface
science. A particularly interesting and important area of catalysis is the interaction of diatomic
molecules (CO, N₂, NO, H₂, and O₂) with transition metal surfaces which can lead to breaking
and making chemical bonds and trigger important surface catalyzed reactions. In particular, the
CO interaction with transition metals has been described in terms of charge transfer from the
metal to CO (back donation) and from CO to metal (forward donation) charge transfer, i.e., the
Blyholder model. This model does not consider, however, the existence of tilted states or the
unusually low CO stretching frequencies observed for CO on some transition metal surfaces.
The present authors have published a number of papers in which ab initio molecular orbital
calculations and the unsynchronized resonating-valence-bond mechanism formed the basis for a
modification of the Blyholder model that explains tilted states, dissociation of diatomic
molecules and low stretching frequencies observed on some transition metal surfaces. The RVB
mechanism of CO dissociation is schematically represented in Fig. 5 where A = C, B = O.

The first electron-transfer step leads to bond formation between the surface, M(s), which
acquires a positive charge (M⁺). It is the possession of the lowest unoccupied 2π* molecular
orbital, as required by Pauling's theory, that makes possible the interaction of CO as a Lewis acid
with the metal surface. The primary interaction in the CO adsorption process on a transition
metal surface is the so-called metal → (2π*) CO back-donation. The metal attacks the carbon
atom where the unoccupied 2π* molecular orbital is mainly localized. The charge transfer
process leads to a build up of negative charge mainly on carbon, although a significant portion
also delocalizes onto oxygen. In the next step, the C-O bond is transferred to the metal
surface. Then CO dissociation occurs when the electron returns to the surface, restoring
electrical neutrality as it must in a catalytic process. The electron traverses a closed loop while
dissociating the CO molecule. Note that the condition for the dissociation is determined by the amount of charge-transfer in the first step, which is generally larger in the tilted configuration (and the consequent C-O weakening.) If there is no metal $\rightarrow$ CO charge transfer, then molecular adsorption will be observed.

Experiments have established that CO adsorption on the first-row transition metal surfaces, Sc to Fe, leads to dissociation, whereas for Co to Cu, CO remains adsorbed on the surface. Ab initio charge transfer calculations for the monocarbonyl complexes are in agreement with these experimental results. It was found that each metal from Sc to Fe is positively charged in the monocarbonyl M-CO system consistent with the unsynchronized resonating-valence bond mechanism of CO dissociation. For Co, Ni, and Cu, however, no net charge transfer was found. Furthermore, higher metal $\rightarrow$ CO charge transfer is obtained for the inclined molecule ($\sim 45^\circ$ relative to the surface.) The tilted state was identified as the precursor state for CO dissociation. The observed preference for charge transfer in the bent structure is consistent with the present model, since a bent and charged adsorbed molecule is predicted as the intermediate state in the second step of the unsynchronized RVB mechanism. Our model was also extended to N$_2$ (A, B = N) interaction with transition metal surfaces as well as the diatomic molecules O$_2$, H$_2$, and NO.

In general, in the tilted configuration, large charge transfer weakens A-B bonds and lowers stretching frequencies$^{11-17, 21}$. We note, however, that more accurate VB calculations would be helpful in order to shed more light on how the CO bond and other diatomic-molecule bonds are fully broken. For discussions regarding the types of unsynchronized resonances which could be relevant for the phenomena described in Fig. 5, see refs. 24-26.

II.6. Chemical Carcinogenesis
Over fifty years ago Hammet suggested the correlation of biological response with electronic parameters\textsuperscript{33}. Since the pioneering work of Miller\textsuperscript{34}, it has been recognized that chemical carcinogens are, or are converted by metabolism into, electrophilic reactants that exert their biological effects by covalent interaction with cellular macromolecules (DNA, RNA and proteins).\textsuperscript{35-38} In our previous work\textsuperscript{19}, large electron affinities (EAs) were calculated for several families of carcinogens, confirming experiments and other computational studies that show that these species react as electrophiles.\textsuperscript{35-38} The present analysis of the DNA-carcinogen interaction implies that the ultimate chemical metabolite becomes covalently bound to liver DNA as a consequence of HOMO (DNA) \textarrow{\rightarrow} LUMO (carcinogen) electron donation. This electron transfer, which leads to DNA-carcinogen bond formation, can be described in the framework of RVB theory (see Fig. 6). The benefit of introducing Pauling’s theory is that it permits the rationalization of the electron transfer process in terms of the molecular structure of the isolated molecules. Using the frontier orbital approximation, the DNA-carcinogen interaction is treated as an electron transfer from the HOMO of DNA to the LUMO of the carcinogen. This unsynchronized resonance requires that the system receiving a bond have an orbital available for electron reception. It is the existence of such an orbital, the LUMO of the carcinogen with the appropriate energy that makes possible DNA \textarrow{\rightarrow} carcinogen electron donation and subsequently DNA-carcinogen bond formation. Chemical carcinogens can be divided into two main groups: those that act directly on DNA, and those that act indirectly. In the majority of compounds that can induce cancer, the ultimate carcinogen is a metabolite of the original compound.

Nitrosamines, for example, are not carcinogens per se; they must be metabolized to be converted into ultimate carcinogens.\textsuperscript{38-40} Nitrosoureas, unlike nitrosamines, are direct-acting carcinogens.
For dimethylnitrosamine (DMN), a considerable body of evidence indicates that damage to cellular DNA and proteins is caused by diazomethyl and dazomethane metabolites\textsuperscript{39}.

Notwithstanding the considerable research on carcinogens, many features of their formation and metabolism are still not fully understood. There is evidence that different carcinogen-derived alkylating agents react at specific guanine sites\textsuperscript{40}. Oxygen alkylation is believed to be a critical mutagenic event, whereas nitrogen alkylation may be innocuous. The DNA to carcinogen electron transfer mechanism\textsuperscript{20} combined with MO calculations show that carcinogens are converted by metabolism into strong electrophilic reactants. In the framework of RVB theory, this electron transfer leads to the formation of a DNA-carcinogen covalent chemical bond. We have calculated the electron affinity (EA) of known carcinogens including some common species such as water, ethanol, and glucose, which are found in animal plasma. The ultimate carcinogens have the highest EAs compared to the original compound or to the other metabolites. On the other hand, ionization potential (IP) calculations of the 35 tautomers of the DNA bases indicate that guanine has the lowest IP, being the preferential base for carcinogen electrophilic attack. An analysis of HOMO contour maps shows that DNA alkylation occurs at specific nitrogen and oxygen sites, in agreement with experiment\textsuperscript{35-41}.

To obtain an approximate view of the nucleophilic sites of DNA, we considered the HOMO energy of the DNA/RNA base units. The tautomeric forms of these bases were also calculated. Among the DNA bases, guanine has the highest HOMO energy and can be considered the preferential site for carcinogen attack. The major HOMO components of the guanine base are localized at specific nitrogen and oxygen sites indicating that these atoms are the most appropriate centers for interaction with the carcinogen. Experimental evidence leads to the same conclusion\textsuperscript{34-36}. Pauling and Cameron\textsuperscript{42} pointed out that vitamin C has anticancer action.
According to the present model one can consider that the protector agent competes with the substrate (DNA, proteins) in donating an electron to the carcinogen. We found that the ascorbate HOMO energy (-3.19 eV) lies well above the guanine HOMO energy (-7.91 eV), indicating that the ascorbate ion is energetically favored for the interaction with the carcinogen relative to the DNA base. It is known that ascorbic acid metabolizes (almost 100%) to ascorbate anion in the human body. In general, anions have high HOMO energies and hence they can react as protectors. It has been reported that the increase of degenerative diseases has been correlated to the reduction of anions in air\textsuperscript{43}. 

In the absence of more sophisticated calculations we suggest that the carcinogen-DNA interaction be described as an electron transfer from the HOMO of the substrate to the LUMO of the carcinogen. It is found that this electron transfer occurs preferentially from the guanine base. Molecular orbital calculations indicate that chemical carcinogens have electrophilic character, in agreement with experimental results. Carcinogenesis involves many complex factors, however, the overall picture emerging from the present simple analysis corresponds to the interpretation of the observed action of chemical carcinogens.

### III. METHODS

Various forms of ab initio VB theory are in current use. We used the approach in which structures of arbitrary type are admitted and a unique set of nonorthogonal orbitals, expressed as linear combinations of a global set of basis functions, are fully optimized\textsuperscript{5}. The basis function used for the Li\textsubscript{n} cluster calculations consist, for each atom, of a 7-term gaussian contraction for the lithium 1s core plus a valence basis having 3 single s type gaussians with exponents 0.020, 0.047, 1.051, and 2 p type gaussians (p\textsubscript{x}, p\textsubscript{y}) with exponent 0.1135. The MS-X\textalpha{} method was
employed to describe the electronic structure$^{10}$ of nickel halides clusters of the type [NiX$_6$]$^{4-}$. The interatomic distances were taken from experiment. The $\alpha$ value is 0.72 for all the muffin-tin regions of the cluster. The self consistent field (SCF) ab initio HF as well as the semiempirical AM1 (Austin Method 1)$^{44}$ MO calculations were performed for selected systems studied here. Different gaussian basis sets were employed. Diffuse functions and Hartree-Fock gradient techniques were used to obtain a better description of the wavefunction in the valence region and consequently of the anion properties that depend on a reasonable description of this region. For the systems $\text{M}_n\text{AB}$ ($\text{M} = \text{Sc to Cu}, \text{AB} = \text{CO}, \text{N}_2, \text{NO}, \text{O}_2,$ and $n = 1-5$) an effective core potential (ECP) with the Hay and Wadt basis set were used$^{45}$. Electron correlation is included at the MP2 level (second order Moller-Plesset correlation energy correction). Geometry optimizations were obtained at different levels of calculation. The EAs of the carcinogens were calculated using Koopmans theorem ($\text{EA} = -\text{E}_{\text{LUMO}}$) and using total energy differences ($\Delta\text{SCF adiabatic}$): $\text{EA} = E_{\text{(neutral)}} - E_{\text{(anion)}}$. The bond distances for the ground state configurations of $\text{O}_4$ and $\text{O}_2$ were fully optimized using the MCSCF method. The computer codes used were MOPAC 93$^{46}$, GAUSSIAN 94$^{47}$ and GAMESS$^{48}$.

IV. CONCLUSIONS

After almost 200 years since its introduction, “the concept of chemical bond is still one of the most valuable concepts in chemistry,” but the quantum mechanical concept of resonance of covalent bonds, introduced by Pauling, has turned out to be an illuminating concept in the modern theory of the chemical bond. Although, one-electron transfers rather than electron-bond
pair bond transfers may provide a more important guide to electronic processes in chemistry and
the Pauling RVB formulation, A-B C-D ↔ A(+)B-C(-)-D, may not often be the appropriate way
to represent electronic reorganization theory, the easily visualized RVB model has proven to be a
very useful guide to simple qualitative descriptions of electron transfer processes in chemistry.
In addition to the numerous applications discussed in this work, further investigations could
reveal an even wider variety of phenomena that can be explained by the RVB theory.

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FIGURE CAPTIONS

Fig. 1  Synchronized resonance Kekule structures of Li$_4$

Fig. 2  Unsynchronized resonance Pauling structures of Li$_4$

Fig. 3  Unsynchronized resonating structures in trans-(CH)$_x$

Fig. 4  Resonating valence band theory for the D$_{2h}$ structure of O$_4$

Fig. 5  Resonating valence band model for diatomic molecular dissociation on transition metal surfaces

Fig. 6  Unsynchronized resonance description of the DNA → carcinogen electron donation and bond formation
Li Li Li — Li
|   | ↔ |    |
Li Li Li — Li
Fig. 1

\[ \text{Li}^- \text{Li} \]

\[ \text{Li} \quad \text{Li}^+ \]
Fig. 3

\[ \text{Diagram showing a reaction process.} \]
Fig. 4
Fig. 6