

An accurate description of the ground and excited states of CH

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With the high accuracy afforded by the sextuple correlation consistent basis set of Dunning, we have calculated energy levels, dissociation energies, equilibrium distances, and other spectroscopic constants for eleven valence and four Rydberg states of the CH radical. Comparisons with experimental and previous theoretical results are made for each state that has been treated. An understanding of their binding is attempted by means of simple valence bond–Lewis diagrams.
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I. INTRODUCTION

The CH radical was first detected in the laboratory as early as 1918,^{1,2} and has been the subject of numerous spectroscopic investigations in the optical, infrared, far infrared, and microwave regions seeking to establish various spectroscopic constants, binding energies, equilibrium distances, dipole moments, lifetimes of excited states (lately rotationally resolved), hyperfine parameters, photodissociation and photoionization processes, as well as dissociative recombination mechanisms.^{3–92} Its presence (along with other hydrides) in extraterrestrial regions^{93–117} and in flames^{118–139} has been a strong reason for the lasting interest in this radical, which interest recently has increased due to the presence of CH in reactions on surfaces of metal catalysts such as Ru and Pd.^{140,141} In a series of articles, Herzberg and co-workers^{16,19,26,30} have obtained spectra of CH and CD by flash photolysis of normal and deuterated diazomethane (as their precursor), and from the analysis of these spectra, they have obtained spectroscopic constants for the ground and excited states of CH and CD as high as about 65 000 cm⁻¹ (~8 eV). Some of these constants have been refined by other workers, but the data collection in Ref. 30 (almost duplicated in Ref. 48) still seems to be the main source of such constants. Up until 1985 the spectroscopy of states above 50 000 cm⁻¹ was not well known except for the data provided by Herzberg's work.³⁰ Since then, using the photodissociation of organic precursors of CH, resonant multiphoton ionization spectra with mass and photoelectron analysis have been obtained and have yielded new states and previously unobserved bands, and some inconsistencies of previous assignments have been resolved.^{56,59,64,66,136} Yet, very few constants for these states have been obtained so far from the analysis of these new spectra. The existence of the spectroscopically elusive $a^4\Sigma^-$ state which lies just above the ground state has been detected in the gas phase by laser photoelectron spectrometry of CH⁻ (Refs. 142, 143) and later by laser magnetic resonance.¹⁴⁴ Along with the spectroscopic constants of CH, similar data for its CD isotope have

also been obtained^{8,9,11,21,29–32,48,145–147} as well as constants for its ¹³CH isotope^{148–151} and the hyperfine parameters of X²Π.¹⁴⁹ Spectra of CH in a low temperature matrix have recently been obtained.^{152,153}

Theoretically, first-row hydrides were the main targets of early molecular computations beyond H₂ and H₂⁺. One reason for this was the relative abundance of early spectroscopic data for these molecules (and their cations) thus furnishing equilibrium separations, spectroscopic constants, and ground state symmetries. Another reason was the fact that only few basis functions were needed for H, thus allowing sufficient functions to be placed on the other nucleus for a satisfactory representation of the orbitals. Also, the correlation energy was slightly easier to estimate, since upon separation H has no correlation energy; in addition, there is little difference between the inner shells of the hydride and of the corresponding first-row atom. Some of the early treatments were qualitative^{154–156} and some employed Slater-type molecular orbitals for valence electrons without self consistency considerations.^{157,158} However, Hartree–Fock–Roothaan type calculations were employed soon, some with limited CI, and some with further semiempirical correlation corrections and exponent optimization in the expansions of the Slater-type functions. The very first simple calculation of CH apparently was that of Niira and Oohata¹⁵⁷ in 1952 while two years later Higuchi¹⁵⁹ did the first CI calculation. Over the years, the progressively developed sophisticated methods were also applied to the first and second-row hydrides; semiempirical methods estimating spectroscopic constants by combining experimental and theoretical results were also used. The CH radical has been a part of many of these calculations.^{160–234} A reasonably detailed bibliography of calculations of first-row hydrides up to 1966 was given by Cade and Huo,¹⁶⁶ while Meyer and Rosmus¹⁸⁶ give references to such calculations from 1966 to 1975.

The volume of theoretical work naturally deals with much the same topics as the experimental papers. Since we are only sketching previous work on CH (our references are certainly not exhaustive), we will touch only briefly on pre-

vious theoretical work most relevant to our present purposes. Liu and Verhaegen¹⁷⁰ employed *ab initio* LCAO-MO-SCF calculations with semiempirical correlation corrections and produced limited potential-energy curves (PEC) for seven states and obtained acceptable spectroscopic constants for six of them. The most extensive totally *ab initio* work on CH was that of Lie *et al.*¹⁷⁵ and Hinze *et al.*^{188,189} who employed a CI method with Slater orbitals as basis functions. They calculated very accurate PECs for the first five states of CH as well as their properties, spectroscopic constants, transition probabilities, lifetimes, and hyperfine parameters. Meyer and Rosmus¹⁸⁶ investigated the ground state of CH and other hydrides by PNO-CI and CEPA methods with Gaussian basis sets with an eye towards investigating the reliability of these methods. Their results, however, are somewhat inferior to those of Lie *et al.* Sun and Freed¹⁹⁶ have used quasidenerate MBPT with Slater orbitals as basis functions and have obtained a large number of CH potential curves and spectroscopic constants for eight states. However, they have not given absolute energy values; the calculations do not converge beyond 5 bohr and the derived constants of the five lowest states do not compare favorably with experimental data. Van Dishoeck²⁰⁵ has investigated the photodissociation processes of CH by a multireference CI method with Gaussian basis sets (MRDCI) and has generated potential curves for states up to about 9 eV from the ground state. However, the minimum of the $X^2\Pi$ ground state is about 16 mH higher than the latest value (*vide infra*) probably because of the limited size of the basis set and the somewhat large threshold of 10 μ H, which generated about 5000 configuration functions (CF). Notice that with the method used, the computed properties do not correspond to the extrapolated (near full-CI) energies but to the wavefunction corresponding to the 5000 CFs.²²⁷ Also, no equilibrium distances (except for the $X^2\Pi$ state) or other spectroscopic constants were given, since this was not the purpose of the work, and the binding (dissociation) energies for some states were small compared to the experimental values (almost half as small for the B state). Grev and Schaefer²²³ using the CCSD(T) method with four basis sets have calculated the ground state and atomization energies, with and without core correlation, and the heats of formation of CH and other species. Peterson *et al.*²²⁴ and Peterson and Dunning²³³ have done benchmark calculations for CH and other species (testing various basis sets and methods) and have obtained accurate energies and spectroscopic constants, but only for the ground state of each species. Hettema and Yarkony²²⁸ employing the full Breit-Pauli spin-orbit Hamiltonian with MRCI functions have studied the spin-forbidden radiative decay of the $CH(a^4\Sigma^-)$ state and have found its lifetime to be between 12 and 8 s depending on the vibrational state. They have also generated potential curves for the X , a , A , and C states and have computed their dipole moments. Martin,²³⁴ using the CCSD(T) method with correlation consistent basis sets, has carried out a very accurate calculation for the ground (only) state of CH and some other hydrides, and has obtained reliable spectroscopic constants and dissociation energies, but he has not given absolute energy values.

Although the ground state of CH has been well charac-

terized, there is still a lack of accurate data for the excited states, especially the ones above 50 000 cm^{-1} . With spectroscopic experiments appearing in the literature involving highly excited states it is obvious that accurate values of constants for these states are needed to corroborate the experimental findings and to help experimentalists to correctly assign new bands and lines.

Thus, the purpose of the present work is to produce accurate PECs and accurate spectroscopic constants, and to investigate the binding modes for as many states as possible. We generate curves for all the states arising from the $2S$ state of H and the $3P$, $1D$, $1S$, $5S$, the doublet states of $3P$, $1P$, and one state stemming from the $3D$ of C. We also compute their equilibrium distances, binding energies, and vibrational and rotational interaction constants as well as their dipole moments. For easy reference, Table I shows the quantities of interest in the present work which have been computed by some of the previous authors.

II. BASIS SETS AND COMPUTATIONAL APPROACH

The correlation consistent cc-pVnZ basis sets of Dunning and coworkers²³⁵ have been employed throughout the present work. For the H atom the $n=4$ basis set was employed in all cases. For the C atom the $n=6$ (sextuple) basis was selected but with the functions i of $l=6$ angular momentum removed. For the CH Rydberg states arising from the $3P$ and $1P(2p^13s^1)$ states of C we have used the corresponding augmented basis set but without the diffuse functions of h symmetry. Thus, our largest basis set expansion reads (17s11p6d5f4g2h/6s3p2d1f) generally contracted to [8s7p6d5f4g2h/4s3p2d1f] containing 182 Gaussian functions.

Starting with a CASSCF calculation, we have distributed the five valence electrons in ten active orbitals ($2s+2p+3d$ of C and $1s$ of H) for the valence states, and in eleven active orbitals (adding a $3s$ diffuse function on C) for the Rydberg states. This generated about 800 and about 1400 CFs, respectively, depending on the molecular symmetry.

All CASSCF vectors were optimized under C_{2v} symmetry and equivalence restrictions, thus acquiring axial symmetry. Dynamical valence correlation was obtained by single and double excitations out of the CAS (i.e., CASSCF+1+2=MRCI) using the internal contraction scheme as implemented in the MOLPRO 96.4 package.²³⁶ Of course, at the CI level, calculated states conform to the irreducible representations of the C_{2v} point group, therefore do not possess pure axial symmetry. In particular states of Σ^+ , Σ^- , Π^\pm and Δ^\pm symmetries are calculated as A_1 , A_2 , B_1 , and A_1 (or A_2), respectively. The uncontracted MRCI space ranges from 1 500 000 to 4 000 000 CFs while the internally contracted space ranges from 300 000 to 1 500 000 CFs, depending on the molecular symmetry. For a stand alone C atom, spherical symmetry was implemented by performing state averaged CASSCF calculations before the MRCI ones.

The energy loss for the $CH(X^2\Pi)$ state due to internal contraction has been estimated by Peterson *et al.*²²⁴ to be about 1 mh at the MRCI/cc-pVQZ level. Because of the large size of the basis sets used here, no correction for the basis set superposition error was deemed necessary. Size ex-

TABLE I. Comparison of the best energies and spectroscopic parameters as calculated by various authors and as determined experimentally. In all CI calculations the C(1s) electrons were kept frozen unless indicated otherwise. (Energies up to four decimal places.)

State	-Energy (hartree)	T_e (eV)	D_e , [D_0] (eV)	r_e , [r_0] (Å)	μ (D)	ω_e (cm^{-1})	$\omega_e \chi_e$ (cm^{-1})	α_e (cm^{-1})	\bar{D}_e (10^{-4}) (cm^{-1})	Reference
$X^2\Pi$...	0.0	2.98	1.09	1.97	3100				159 (1954) ^a
	38.464		2.97		1.54					160 (1958) ^b
	38.2796		2.47	1.104		3053	55.5	0.4712		166 (1967) ^c
	38.2794			1.124 ^d	1.57					168 (1968) ^d
	38.479		3.46	1.106		3108				170 (1970) ^e
	38.4104		3.51	1.118	1.450	2886.1	82.0	0.589	14.4 ^f	175 (1973) ^g
	38.4083		3.47	1.122	1.43	2841.7	64.4	0.532		186 (1975) ^h
	...	0.0		1.133		2519.1	19.17	0.737		196 (1981) ⁱ
	38.3852		[3.32]	1.124		2828.3	62.52	0.520		198 (1983) ^j
	38.4069	0.0	3.45	1.1199						205 (1987) ^k
	38.4195		3.609	1.104 ^l						223 (1992) ^l
	38.4702		3.585	1.104 ^l						223 (1992) ^l
	38.4186		3.59	1.1201		2850.2	64.0	0.5374		224 (1993) ^m
	38.4138 ⁿ	0.0		1.1224		2851.9	66.6			228 (1994) ⁿ
	38.4207		3.613	1.1202						233 (1997) ^o
	38.4720		3.608	1.1184						233 (1997) ^o
	...		3.631 ^p	1.11958		2857.88	63.839 ^p			234 (1998) ^p
	...			1.11808		2861.72	64.554 ^q			234 (1998) ^q
	38.4217		3.615	1.1204	1.4057	2851.0	62.15	0.542	14.85	present work
	...	0.0	3.640	1.119786 _g	1.46±0.06	2860.75	64.44	0.5365	14.7	Exp. 48, 86
$a^4\Sigma^-$...	0.09		1.08	1.07	3500				159 (1954) ^a
	38.449									150 (1958) ^b
	38.2344			1.1026 ^d	0.89					168 (1968) ^d
	38.445			1.093		3359				170 (1970) ^e
	38.3865		2.84	1.086	0.663	3145.7	71.8	0.553	14.3 ^f	175 (1973) ^g
	...	0.481		1.114		2947.3	83.35	0.467		196 (1981) ⁱ
	...	0.716		1.0912		3117.3	80.6			228 (1994) ⁿ
	38.3942	0.747	2.863	1.0892	0.6531	3090.9	102.17	0.723	15.19	present work
	...	0.742	[2.723]	[1.0977]						Exp. 143, 144
	$A^2\Delta$...	2.75		1.10	1.41	3300			
38.364										150 (1958) ^b
38.1803				1.1026 ^d	0.91					168 (1968) ^d
38.371				1.090		3111				170 (1970) ^e
38.3031			1.90	1.102	0.904	2970.3	98.5	0.697	15.2 ^f	175 (1973) ^g
...		3.074		1.123		2688.8	80.17	0.812		196 (1981) ⁱ
38.2794			[1.45]	1.111		2863.6	96.15	0.696		198 (1983) ^j
...		3.00 ^k	1.88							205 (1987) ^k
...		2.922		1.1073		2926.9	103.8			228 (1994) ⁿ
38.3151		2.901	1.975	1.1056	0.8434	2911.1	91.99	0.675	15.42	present work
...	2.870	2.010	1.1031	0.77±0.07	2914.10	81.40	0.6354	15.4	Exp. 30, 86, 74	
$B^2\Sigma^-$...	3.07		1.13	1.89	3100				159 (1954) ^a
	38.1583			1.1861 ^d	1.54					168 (1968) ^d
	38.358			1.124		2543				170 (1970) ^e
	38.2908		0.23	1.173	1.389	2141.7	223.2	1.933	22.6 ^f	175 (1973) ^g
	...	3.180		1.270		2081.1	385.7	1.579		196 (1981) ⁱ
	...	3.24 ^k	0.29							205 (1987) ^k
	38.3026	3.241	0.372	1.1768	1.3285	2167.1	173.72	1.11	19.38	present work
	...	3.231	0.409	1.1640		2246.42	225.7	1.4823	16.3	Exp. 30, 89
$C^2\Sigma^+$...	3.65		1.11	1.41	3200				159 (1954) ^a
	38.1283			1.1132 ^d	0.94					168 (1968) ^d
	38.333			1.097		3085				170 (1970) ^e
	38.2627		0.78	1.111	0.955	2887.5	106.8	0.771	15.5 ^f	175 (1973) ^g
	...	4.095		1.129		2558.9	88.83	0.794		196 (1981) ⁱ
	...	4.02 ^k	0.86							205 (1987) ^k
	...	3.9904		1.1179		2853.3	133.0			228 (1994) ⁿ
	38.2758	3.983	0.910	1.1164	0.9055	2837.3	87.76	0.429	14.91	present work
	...	3.943	0.940	1.1143		2840.2	125.96	0.7185	15.55 ^f	Exp. 30, 48

TABLE I. (Continued.)

State	-Energy (hartree)	T_e (eV)	D_e , [D_0] (eV)	r_e , [r_0] (Å)	μ (D)	ω_e (cm^{-1})	$\omega_e X_e$ (cm^{-1})	α_e (cm^{-1})	\bar{D}_e (10^{-4}) (cm^{-1})	Reference
$E^2\Pi$...	7.09		1.14	1.19	3100				159 (1954) ^a
	38.210			1.167		2643				170 (1970) ^e
	...	7.558		1.137		3284.9	905.2	0.261		196 (1981) ⁱ
	...	7.43 ^k								205 (1987) ^k
	38.1513	7.358		1.1437	0.8334	2743.0	57.63	0.512	14.24	present work
	...	(7.488) ^f		(1.15)						Exp. 48

^aMinimal STF basis set with small CI; dipole moments from SCF calculations.

^bMinimal STF basis set with small CI plus semiempirical correlation corrections; all at 2.124 bohr.

^cSCF with optimized STF exponents; Basis set: C ($5s4p2d1f$); H ($3s1p$); potential curve (1.6 to 3.5 bohr).

^dSame as (c) but without potential curve; all at the indicated experimental r 's ($A^2\Delta$ state's r used in a $4\Sigma^-$ state).

^eLCAO-MO-SCF with a semiempirical correlation estimate; basis set as in (c).

^fThis quantity corresponds to $D_{v=0}$ (10^{-4}) cm^{-1} .

^gMedium size CI with NO's and an optimized STF basis set (C/H: $6s4p2d2f4s3p2d$); potential curves.

^hCEPA with the GTO basis set: C: $4s4p2d1f$; H: $4s2p1d$.

ⁱQuasidegenerate MBPT with the STF basis set (C/H): ($4s3p1d/2s1p$).

^jLarge CASSCF with the basis set: C: $6s6p3d$; H: $4s3p$; active space: $5\sigma 3\pi$ and 1δ (propert.) or 2δ (energies).

^kMRDCI ($10\mu\text{H}$ threshold) with customized basis sets. T_e 's are vertical excitations from r_e of the ground state.

^lCCSD(T) with customized basis set; r_e is the equilibrium distance at the SCF level; 1st line: frozen core; 2nd line: all electrons correlated.

^mCASSCF+1+2 with the correlation consistent basis set cc-pV5Z.

ⁿSA-CASSCF/CI with the cc-pVTZ basis set; the energy value corresponds to $r=2.116$ bohr.

^oCCSD(T) with the basis sets (C/H): cc-pV(SZ/QZ), 1st line: frozen core, and cc-pCV(QZ/QZ), 2nd line all electrons correlated.

^pCCSD(T) with the cc-pV6Z basis set. Extrapolated to infinite-basis limit gives $D_e=3.643$ eV and $D_0=3.468$ eV; $\omega_e y_e=0.305$, $\omega_e z_e=-0.004$ cm^{-1} .

^qCCSD(T) "best estimate" with cc-pVnZ basis sets including core correlation; $\omega_e y_e=0.300$, $\omega_e z_e=-0.005$ cm^{-1} .

^rThe T_0 value is 7.313 eV; the T_e estimate is uncertain.

tensivity errors are also small; at the MRCI level our largest such error is about 0.4 mh for the $H^2\Pi$ Rydberg state and 0.08 mh for the valence states, as obtained by subtracting the CI fragments from the MRCI supermolecule.

Excited states of $^2\Pi$ and $^2\Sigma^+$ symmetry have been obtained via the state average (SA) methodology employing $w(1,1,1,1)$ weighting vectors. Energy losses due to the SA approach are not significant; for instance, for the states $X^2\Pi$ and $C^2\Sigma^+$ equilibrium energy differences with and without the SA amount to 0.2 and 1 mh, respectively.

Spectroscopic constants for the four isotopic species ^{12}CH , ^{12}CD , ^{13}CH , and ^{13}CD were extracted by fitting 20 to 30 equidistant energy points (0.02 bohr apart) around the equilibrium geometry, and then applying a standard Dunham analysis.²³⁷

III. RESULTS AND DISCUSSION

To estimate the sufficiency of our basis set, we computed the energy levels of the C atom in a spherically averaged manner as previously indicated. It was found that the SCF energy of the 3P ground state was $-37.688\,612$ h, just $7\,\mu\text{h}$ above the numerical result.²³⁸ Table II lists the absolute MRCI energies and energy gaps (ΔE) of the $^3P(2s^22p^2)$, $^1D(2s^22p^2)$, $^1S(2s^22p^2)$, $^5S(2s^12p^3)$, $^3P(2s^22p^13s^1)$, $^1P(2s^22p^13s^1)$, and $^3D(2s^12p^3)$ terms calculated as indicated in the previous section. Nearly all energy gaps are in excellent agreement with the experiment, showing that the chosen basis set and correlation treatment are adequate for all the computed CH states. The two small deviations from this agreement are the $^1P(2s^22p^13s^1)\leftarrow^3P(2s^22p^2)$ and $^3D(2s^12p^3)\leftarrow^3P(2s^22p^2)$ energy splittings, which are overestimated by 0.060 and 0.148 eV, respectively.

Table III displays total energies (E), binding energies (D_e), equilibrium bond lengths (r_e), dipole moments (μ), and energy gaps (T_e) of all the calculated, bond CH states. Two more states have been calculated, the $b^4\Pi$ and $d^6\Sigma^-$, but they are repulsive and thus not listed in this table. Tables IV–VII list spectroscopic constants for four isotopic species ($^{12}\text{C-H}$, $^{12}\text{C-D}$, $^{13}\text{C-H}$, $^{13}\text{C-D}$). PECs for all computed states are shown in Fig. 1.

In what follows we discuss the important characteristics of every state. An effort is made to analyze the chemical binding, using simple valence bond-Lewis (vbL) pictures.

A. The ground $X^2\Pi$ state

We can envisage the formation of the $X^2\Pi$ state as the result of $\text{C}(^3P; M=\pm 1) + \text{H}(^2S)$ reaction. The bonding can be represented by the following vbL diagram:

TABLE II. Total energies E (hartree) of the 3P , 1D , 1S , 5S , 3P , 1P , and 3D carbon states and corresponding energy splittings ΔE (eV) with respect to the ground 3P state at the MRCI level of theory. Experimental values in parentheses.

State	$-E$	ΔE^a
$^3P(2s^22p^2)$	37.788 854	0.0 (0.0)
$^1D(2s^22p^2)$	37.742 195	1.270 (1.260)
$^1S(2s^22p^2)$	37.689 894	2.693 (2.680)
$^5S(2s2p^3)$	37.635 375	4.176 (4.179)
$^3P(2s^22p^13s^1)$	37.512 859	7.510 (7.545)
$^1P(2s^22p^13s^1)$	37.504 361	7.741 (7.681)
$^3D(2s2p^3)$	37.491 569	8.090 (7.942)

^aExperimental values averaged over M_J , Ref. 239.

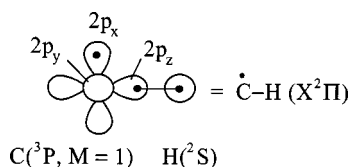
TABLE III. Total energies E (hartree), dissociation energies D_e (kcal/mol), bond distances r_e (Å), dipole moments μ (D), and energy gaps T_e (kcal/mol) of the calculated states of the C–H system.

State	Method ^d	$-E$	D_e ^b	r_e	μ	T_e
$X^2\Pi$	CASSCF	38.365 769	71.51	1.1291	1.3835	
	MRCI	38.421 680	83.37	1.1204	1.4057	0.0
	MRCI+Q	38.422 8	83.68	1.1204		
	Exp. ^c		83.94	1.119786	1.46±0.06	0.0
$a^4\Sigma^-$	CASSCF	38.348 569	59.03	1.0834	0.6292	
	MRCI	38.394 231	66.03	1.0892	0.6531	17.22
	MRCI+Q	38.395 0	66.23	1.0898		
	Exp. ^d		62.80±0.23	1.0977	...	17.11±0.18
$A^2\Delta$	CASSCF	38.252 503	31.15	1.1236	0.7634	
	MRCI	38.315 088	45.54	1.1056	0.8434	66.89
	MRCI+Q	38.316 5	46.05	1.1052		
	Exp. ^e		46.35	1.1031	0.77±0.07	66.19
$B^2\Sigma^-$	CASSCF	38.251 060	-2.84	1.2353	1.3854	
	MRCI	38.302 579	8.59	1.1468	1.3285	74.74
	MRCI+Q	38.303 6	8.94	1.1748		
	Exp. ^f		9.43	1.1640	...	74.51
$C^2\Sigma^+$	CASSCF	38.207 095	1.22	1.1300	0.7762	
	MRCI	38.275 824	20.98	1.1164	0.9055	91.85
	MRCI+Q	38.277 8	21.80	1.1134		
	Exp. ^g		21.68	1.1143	...	90.93
$D^2\Sigma^+$	CASSCF					
	MRCI	38.204 769	9.35	1.6635	1.4203	136.11
	MRCI+Q	38.208 7	9.26	1.6547		
	Exp.	
$c^4\Sigma^-$	CASSCF	38.134 148	12.79	1.8586	0.1479	
	MRCI	38.170 874	22.31	1.7866	0.2630	157.38
	MRCI+Q	38.171 5	22.60	1.7839		
	Exp.	
$E^2\Pi$	CASSCF	38.087 085		1.1663	0.5834	
	MRCI	38.151 279		1.1437	0.8334	169.68
	MRCI+Q	38.153 2		1.1426		
	Exp. ^h			1.15	...	172.68
$F^2\Pi$	CASSCF	38.071 897	58.08	1.1696	4.0236	
	MRCI	38.132 337	75.22	1.3751	4.4172	181.57
	MRCI+Q	38.134 6	75.76	1.3751		
	Exp. ^h		...	1.20	...	187.62
$G^2\Sigma^+$	CASSCF					
	MRCI	38.123 104	69.30	1.1482	6.1699	187.36
	MRCI+Q	38.125 1	69.94	1.1508		
	Exp. ^h		...	1.221	...	188.55
		(local minimum)				
	CASSCF					
	MRCI	38.064 757	32.68	2.6323	8.7422	223.97
	MRCI+Q	38.071 1	36.02	2.6642		
	Exp.	
$H^2\Pi$	CASSCF	38.040 009	43.75	1.4185	2.1774	
	MRCI	38.101 144	61.03	1.3762	2.0212	201.14
	MRCI+Q	38.105 2	62.41	1.3651		
	Exp.	
$I^2\Sigma^+$	CASSCF					
	MRCI	38.096 382	57.85	1.2639	0.1610	204.15
	MRCI+Q	38.099 4	59.12	1.2591		
	Exp.	
$J^2\Delta$	CASSCF					
	MRCI	38.069 471	48.92	1.6661	0.3515	221.05
	MRCI+Q	38.073 2	46.18	1.6612		
	Exp.	

^aMRCI+Q refers to the multireference Davidson correction.^bAll D_e values are with respect to adiabatic products.^cThe D_e, r_e, μ values are from Refs. 48, 86, and 25, respectively.^dThe data reported, Ref. 143, correspond to $D_0, r_0,$ and T_0 values; see text.^eThe r_e, T_e values are from Ref. 86, and μ is from Ref. 74.^fReferences 89 and 240. For the experimental D_e see text.^g D_e from Ref. 30, the rest of experimental findings from Ref. 48.^hReference 48.

TABLE IV. Harmonic frequencies ω_e , anharmonicities $\omega_e\chi_e$, rotational vibrational couplings α_e , and centrifugal distortions \bar{D}_e in cm^{-1} of the $^{12}\text{C-H}$ system in different states at the MRCI level. Experimental values in parentheses.

State	ω_e	$\omega_e\chi_e$	α_e	$\bar{D}_e(10^{-4})$
$X^2\Pi$	2851.0 (2860.75) ^a	62.15 (64.44)	0.542 (0.5365)	14.85 (14.7)
$a^4\Sigma^-$	3090.9 ...	102.17 ...	0.723 ...	15.19 ...
$A^2\Delta$	2911.1 (2914.10) ^a	91.99 (81.40)	0.675 (0.6354)	15.42 (15.4)
$B^2\Sigma^-$	2167.1 (2246.42) ^b	173.72 (225.7) ^c	1.11 (1.4823)	19.38 (16.3)
$C^2\Sigma^+$	2837.3 (2840.2) ^d	87.76 (125.96)	0.429 (0.7185)	14.91 (15.55)
$D^2\Sigma^+$	1542.8 ...	164.54 ...	0.327 ...	4.57 ...
$c^4\Sigma^-$	1390.3 ...	36.66 ...	0.059 ...	3.80 ...
$E^2\Pi$	2743.0 ...	57.63 ...	0.512 ...	14.24 ...
$G^2\Sigma^{+e}$	2475.1 808.2	147.27 18.52	0.946 0.060	16.80 1.10
$I^2\Sigma^+$	2892.2 ...	106.07 ...	0.043 ...	7.05 ...
$J^2\Delta$	1743.6 ...	41.63 ...	-0.076 ...	3.69 ...

^aReference 86.^bReference 89.^cReference 21.^dReference 48.^eThe first entry corresponds to the global minimum, the second to the local minimum; see text and Fig. 1.

The CASSCF equilibrium Mulliken populations are (C/H)

$$2s^{1.81}2p_z^{1.27}2p_x^{0.98}2p_y^{0.06}3d^{0.06}/1s^{0.79}2p_z^{0.01}2p_y^{0.02},$$

supporting the picture above. At infinity, the corresponding C atomic distributions are

$$2s^{1.93}2p_z^{1.0}2p_x^{1.0}2p_y^{0.05}3d^{0.03},$$

indicating a near degeneracy or GVB $2s-2p_y$ correlation. In total $0.2e^-$ are transferred from H to C. Therefore, upon completion of the C+H interaction, $\sim[0.12(2s_C) + 0.18(1s_H)]$ electrons are promoted to the $2p_z$ orbital of C.

As it can be seen from Tables I and III, our total MRCI energy of $-38.421\,680$ hartree is the lowest valence correlated energy reported so far in the literature. We also report a $D_e = 83.37$ kcal/mol at the MRCI level which becomes 83.68 kcal/mol when the multireference Davidson correction for unlinked clusters is taken into account. Further, if we add to this value a core correlation correction of about 0.13 kcal/mol (*vide infra*), we obtain a final $D_e = 83.81$ kcal/mol, 0.13 kcal/mol lower than the experimental value. Our value of r_e is 1.1204 Å; assuming a 0.002 Å decrease due to core correlation effects^{233,234} our final r_e value is 1.1184 Å. Also, our

TABLE V. Harmonic frequencies ω_e , anharmonicities $\omega_e\chi_e$, rotational vibrational couplings α_e , and centrifugal distortions \bar{D}_e in cm^{-1} of the $^{12}\text{C-D}$ system in different states at the MRCI level. Experimental values in parentheses.^a

State	ω_e	$\omega_e\chi_e$	α_e	$\bar{D}_e(10^{-4})$
$X^2\Pi$	2093.3 (2100.35) ^b	33.64 (34.16)	0.213 (0.212)	4.31 (4.32)
$a^4\Sigma^-$	2269.4 ...	44.09 ...	0.272 ...	4.38 ...
$A^2\Delta$	2137.4 (2203.3)	49.39 (78.50)	0.266 (0.260)	4.48 (4.5)
$B^2\Sigma^-$	1591.1 (1652.5)	94.14 (123.8)	0.437 (0.341)	5.62 (6.36)
$C^2\Sigma^+$	2083.2 (2081.3)	50.43 (66.79)	0.174 (0.283)	4.36 (4.5)
$D^2\Sigma^+$	1132.8 ...	95.93 ...	0.136 ...	1.31 ...
$c^4\Sigma^-$	1020.8 ...	19.81 ...	0.023 ...	1.10 ...
$E^2\Pi$	2014.0 (2025)	30.30 ...	0.199 ...	4.13 (4.0)
$G^2\Sigma^{+c}$	1817.3 593.4	78.35 9.80	0.377 0.024	4.91 0.319
$I^2\Sigma^+$	2123.5 ...	57.83 ...	0.019 ...	2.04 ...
$J^2\Delta$	1280.2 ...	22.30 ...	-0.030 ...	1.07 ...

^aReference 48 (except for $X^2\Pi$).^bReference 147.^cThe first entry corresponds to the global minimum, the second to the local, see text and Fig. 1.

dipole moment $\mu = 1.406$ D. Lie *et al.*¹⁷⁵ in their remarkable 1973 work on CH report an energy of $-38.410\,44$ hartree, a $D_e = 80.94$ kcal/mol, a $r_e = 2.113$ bohr = 1.1182 Å, and a $\mu = 1.45$ D. Peterson and Dunning²³³ give a total (valence-correlated) energy of $-38.420\,700$ hartree, a $D_e = 83.33$ kcal/mol and a $r_e = 1.1202$ Å at the CCSD(T)/cc-pV(5Z/QZ) level. When they take into account core correlation effects [at the CCSD(T)/cc-pCV(QZ/QZ) level] D_e is improved by 0.13 kcal/mol, giving their best value of 83.46 kcal/mol, and their r_e decreases by 0.0018 Å. A similar de-

TABLE VI. Harmonic frequencies ω_e , anharmonicities $\omega_e\chi_e$, rotational vibrational couplings α_e , and centrifugal distortions \bar{D}_e in cm^{-1} of the $^{13}\text{C-H}$ system in different states at the MRCI level.

State	ω_e	$\omega_e\chi_e$	α_e	$\bar{D}_e(10^{-4})$
$X^2\Pi$	2842.5	61.78	0.538	14.67
$a^4\Sigma^-$	3081.6	101.30	0.716	15.01
$A^2\Delta$	2902.4	91.43	0.669	15.24
$B^2\Sigma^-$	2160.6	172.69	1.10	19.15
$C^2\Sigma^+$	2828.8	87.31	0.425	14.73
$D^2\Sigma^+$	1568.4	152.94	0.341	4.38
$c^4\Sigma^-$	1386.1	36.45	0.058	3.75
$E^2\Pi$	2734.8	57.27	0.507	14.07
$G^2\Sigma^{+a}$	2467.7 805.8	146.37 18.40	0.937 0.060	16.60 1.08
$I^2\Sigma^+$	2883.5	105.45	0.043	6.97
$J^2\Delta$	1738.4	41.38	-0.075	3.64

^aThe first entry corresponds to the global minimum, the second to the local; see text and Fig. 1.

TABLE VII. Harmonic frequencies ω_e , anharmonicities $\omega_e\chi_e$, rotational vibrational couplings α_e , and centrifugal distortions \bar{D}_e in cm^{-1} of the $^{13}\text{C}-\text{D}$ system in different states at the MRCI level.

State	ω_e	$\omega_e\chi_e$	α_e	$\bar{D}_e(10^{-4})$
$X^2\Pi$	2081.7	33.27	0.210	4.22
$a^4\Sigma^-$	2256.8	43.46	0.268	4.29
$A^2\Delta$	2125.5	48.84	0.261	4.38
$B^2\Sigma^-$	1582.3	93.11	0.430	5.49
$C^2\Sigma^+$	2071.6	49.91	0.171	4.27
$D^2\Sigma^+$	1126.5	94.96	0.134	1.28
$c^4\Sigma^-$	1015.1	19.59	0.023	1.08
$E^2\Pi$	2002.8	29.95	0.196	4.04
$G^2\Sigma^+$	1807.2	77.47	0.371	4.80
	590.1	9.69	0.024	0.312
$I^2\Sigma^+$	2111.7	57.20	0.019	1.20
$J^2\Delta$	1273.1	22.05	-0.030	1.05

^aThe first entry corresponds to the global minimum, the second to the local; see text and Fig. 1.

crease of 0.00192 \AA is found by Martin²³⁴ upon comparing the CCSD(T) results between the cc-pCV5Z and the aug-cc-pV5Z uncontracted levels. The corresponding experimental values are $D_e = 83.94 \text{ kcal/mol}$, $r_e = 1.119786(8) \text{ \AA}$,⁸⁶ practically equal to $r_e = 1.1199 \text{ \AA}$ given by Huber and Herzberg.⁴⁸ Morino *et al.*,¹⁴⁷ by estimating B_e more accurately, have obtained an experimental $r_e = 1.118056(29) \text{ \AA}$, which is in excellent agreement with our core correlation corrected results. Finally, notice that the experimental value of the dipole moment is $1.46 \pm 0.06 \text{ D}$.²⁵

B. The $a^4\Sigma^-$ and $c^4\Sigma^-$ states

Figure 1 shows that the $a^4\Sigma^-$ state traces its origin to the ground state fragments, while the $c^4\Sigma^-$ state correlates

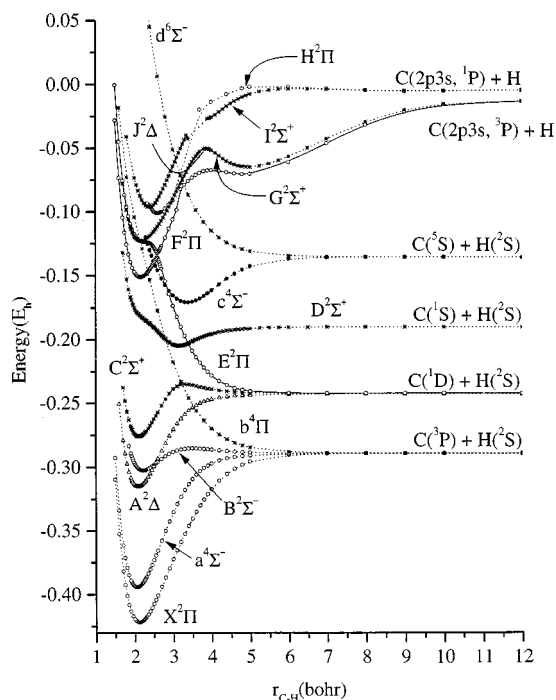


FIG. 1. Potential energy curves of the lowest 11 valence and 4 Rydberg states of CH. All energies have been shifted by $+38.00$ hartree.

to the $^5S(2s^12p^3)$ carbon state. Despite their large energy separation ($T_e = 140.2 \text{ kcal/mol}$) our findings indicate that these two quartets interact substantially. In other words, the 5S carbon state plays a prominent role in the bonding mechanism of the $a^4\Sigma^-$ state. The CASSCF atomic equilibrium and infinite separation Mulliken distributions are (C/H)

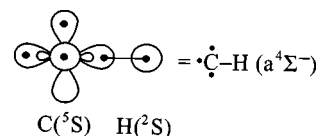
$$r_e: 2s^{1.42}2p_z^{0.81}2p_x^{0.98}2p_y^{0.98}3d^{0.03}/1s^{0.72}2p^{0.03},$$

$$r_\infty: 2s^{1.95}2p_z^{0.04}2p_x^{1.0}2p_y^{1.0}3d^{0.01}/1s^{1.0}.$$

We see that upon interaction $0.5 e^-$ from the $2s_C$ and $0.3 e^-$ from the $1s_H$ have been promoted to the $2p_z$ orbital of C. We claim that the *in situ* C atom finds itself to a considerable degree in the 5S excited state, with a corresponding CAS leading configuration

$$|a^4\Sigma^- \rangle = 0.99 | 1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^1 1\pi_y^1 \rangle,$$

and the following vBL picture of the molecule:



The $a^4\Sigma^-$ state was first observed in 1970 by Feldmann¹⁴² using laser photoelectron spectroscopy of CH^- . Kasdan *et al.*¹⁴³ in 1975 obtained a $T_0 = 0.742 \pm 0.008 \text{ eV} = 17.11 \pm 0.18 \text{ kcal/mol}$, which combined with the experimental D_0 of the $X^2\Pi$ state (3.465 eV)⁴⁸ gives $D_0 = 2.723 \pm 0.01 \text{ eV} = 62.80 \pm 0.23 \text{ kcal/mol}$ for the $a^4\Sigma^-$ state. Our MRCI $T_0 (= T_e + \Delta\omega_e/2 - \Delta\omega_e\chi_e/4)$ and $D_0 (= D_e - \omega_e/2 + \omega_e\chi_e/4)$ are 17.53 and 61.65 kcal/mol , respectively (Tables III and IV). The discrepancy of about 1 kcal/mol in D_0 between experiment and theory can be attributed to corresponding differences of D_e of $X^2\Pi$ (0.57 kcal/mol) and of T_0 (0.42 kcal/mol), which amount to 0.99 kcal/mol . Nelis *et al.*¹⁴⁴ employing a laser magnetic resonance method reported an experimental (vibrationally averaged) bond distance $r_0 = 1.09767 \text{ \AA}$ in fair agreement with our $r_e = 1.0892 \text{ \AA}$. Notice that this is the shortest bond distance of all examined states. Corresponding theoretical values by Lie *et al.*¹⁷⁵ are: $T_0 = 15.43 \text{ kcal/mol}$, $D_0 = 60.88 \text{ kcal/mol}$, and $r_e = 1.0864 \text{ \AA}$.

The $c^4\Sigma^-$ state has never been observed experimentally. The only theoretical investigation we are aware of is the SCF calculation of Liu and Verhaegen¹⁷⁰ at $r = 2.9$ bohr ($= 1.535 \text{ \AA}$) with semiempirical correlation corrections. They gave a $T_e (c^4\Sigma^- \leftarrow X^2\Pi) = 200 \text{ kcal/mol}$.

The PEC of this state is illustrated in Fig. 1, and from Table III we see that $T_e (c^4\Sigma^- \leftarrow X^2\Pi) = 157.38 \text{ kcal/mol}$, $D_e = 22.31 \text{ kcal/mol}$, with respect to $\text{C}(^5S) + \text{H}(^2S)$, and $r_e = 1.7866 \text{ \AA}$. The leading CASSCF configurations are

$$|c^4\Sigma^- \rangle = -0.55 | 1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^1 1\pi_y^1 \rangle$$

$$+ 0.51 | 1\sigma^2 2\sigma^1 3\sigma^2 1\pi_x^1 1\pi_y^1 \rangle$$

$$- 0.40 | 1\sigma^2 2\sigma^1 3\sigma^1 4\sigma^1 1\pi_x^1 1\pi_y^1 \rangle$$

$$- 0.26 | 1\sigma^2 2\sigma^2 4\sigma^1 1\pi_x^1 1\pi_y^1 \rangle,$$

with CAS equilibrium Mulliken populations (C/H)

$$2s^{1.17}2p_z^{0.83}2p_x^{1.0}2p_y^{1.0}/1s^{0.98}.$$

The -0.55 component above corresponds to the main CF of the $a^4\Sigma^-$ state, while the 0.51 and -0.40 components correspond to the asymptotes of $a^4\Sigma^-(^3P+^2S)$ and $c^4\Sigma^-(^5S+^2S)$, respectively.

C. The $A^2\Delta$ state

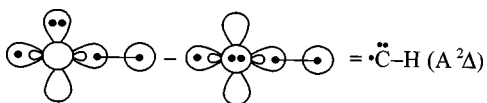
This state traces its ancestry to $C(^1D)+H(^2S)$, as shown in Fig. 1. Its CASSCF wave-function at equilibrium and its Mulliken distributions at r_e and r_∞ are as follows:

$$|A^2\Delta\rangle \approx 0.69\{|1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^2\rangle - |1\sigma^2 2\sigma^2 3\sigma^1 1\pi_y^2\rangle\},$$

$$r_e: 2s^{1.55}2p_z^{0.73}2p_x^{0.97}2p_y^{0.97}3d^{0.05}/1s^{0.67}2p^{0.05},$$

$$r_\infty: 2s^{1.93}2p_z^{0.05}2p_x^{1.0}2p_y^{1.0}3d^{0.03}/1s^{1.0}.$$

It is obvious that $0.38e^-$ from $2s_C$ and $0.28e^-$ from $1s_H$ are transferred to the $2p_{z,C}$ orbital ($0.38+0.28+0.05=0.71$). We observe that the *in situ* C atom is in a s^1p^3 valence excited state, which by symmetry cannot be other than the $^3D(2s^1 2p^3)$ state, lying 7.942 eV above the ground 3P state (Table II). It is remarkable that the interaction of $C(^1D)+H(^2S)$ to form an $A^2\Delta$ state has as a result the excitation of C to the 3D state, 6.68 eV above the 1D state. From Tables I and III we see that our T_e ($A^2\Delta \leftarrow X^2\Pi$) = 66.89 kcal/mol, $r_e = 1.1056$ Å, $D_e = 45.54$ kcal/mol, and our dipole moment $\mu = 0.8434$ D are in excellent agreement with the experimental values.^{48,83,86,74} However, considering the strong involvement ($\sim 70\%$) of the 3D state in the σ -bond formation, the intrinsic bond strength of this state is 153 kcal/mol (6.68×0.70 eV + 45.5 kcal/mol). The values of Lie *et al.*¹⁷⁵ are also in good agreement with the experiment ($r_e = 1.1023$ Å, $T_e = 67.36$ kcal/mol, and $D_e = 43.82$ kcal/mol). A vBL picture conforming to the discussion above is



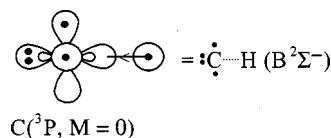
D. The $B^2\Sigma^-$ state

This weakly bound state is depicted in Fig. 1 and correlates to the ground $C(^3P)+H(^2S)$. The leading CASSCF configurations at equilibrium and the corresponding Mulliken populations are

$$|B^2\Sigma^-\rangle \approx 0.79|1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^1 1\pi_y^1\rangle - 0.40\{|1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^1 1\pi_y^1\rangle + |1\sigma^2 2\sigma^2 3\sigma^2 1\pi_x^1 1\pi_y^1\rangle\},$$

$$2s^{1.74}2p_z^{0.55}2p_x^{0.98}2p_y^{0.98}3d^{0.03}/1s^{0.68}2p^{0.03}.$$

The following vBL diagram suggests that the two atoms are held together by a half σ -bond.



At 3.5 bohr our MRCI calculations reveal an energy barrier of 2.32 kcal/mol, first observed experimentally at about 4 bohr by Herzberg and Johns.³⁰ We assume that this is due to the participation of the $^3D(2s^1 2p^3)$ excited state of C, as evinced by the population analysis. As Table I shows, Herzberg and Johns³⁰ and Kepa *et al.*⁸⁹ report an $r_e = 1.1640$ Å while in the Huber and Herzberg compilation⁴⁸ $r_e = 1.1975$ Å. In Table III we report a $r_e = 1.1768$ Å as well as $T_e = 74.74$ kcal/mol and $D_e = 8.59$ kcal/mol which compare well with the respective experimental values of 74.51 and 9.43 kcal/mol [Tables I and III, using $D_e(X) - T_e(B)$]. The corresponding theoretical values of Lie *et al.*¹⁷⁵ are $r_e = 1.1727$ Å, $T_e = 75.09$ kcal/mol, and $D_e = 5.30$ kcal/mol.

E. The $b^4\Pi$ state

As it is shown in Fig. 1, this is the first repulsive state correlating to the ground state fragments. Although not apparent from its plot, this state possesses a calculated van der Waals minimum of 4.5 cm⁻¹ at about 9.0 bohr.

F. The $C^2\Sigma^+$ state

The Hartree-Fock $^1D(M=0)$ wavefunction of C is

$$|^1D\rangle = (1/6)^{1/2}\{2|2s^2 2p_z^2\rangle - |2s^2 2p_x^2\rangle - |2s^2 2p_y^2\rangle\}.$$

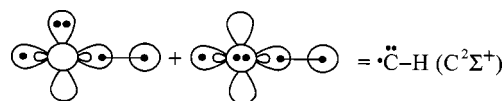
As $H(^2S)$ approaches $C(^1D)$ from infinity the interaction is repulsive at first, leading eventually to an energy barrier of 4.66 kcal/mol at about 3.3 bohr (Fig. 1). Lie *et al.*¹⁷⁵ have calculated its height to be 6.6 kcal/mol at about 3.3 bohr, and it has also been observed experimentally.³⁰ As the interatomic distance approaches equilibrium, the strong interaction with the $D^2\Sigma^+$ state induces a decrease in the p_z electronic density accompanied by a simultaneous increase in the densities of p_x and p_y . At equilibrium, the leading CAS configurations and the atomic Mulliken CAS distributions at r_e and r_∞ are

$$|C^2\Sigma^+\rangle \approx 0.69\{|1\sigma^2 2\sigma^2 3\sigma^1 1\pi_x^2\rangle + |1\sigma^2 2\sigma^2 3\sigma^1 1\pi_y^2\rangle\},$$

$$r_e: 2s^{1.54}2p_z^{0.72}2p_x^{0.98}2p_y^{0.98}3d^{0.02}/1s^{0.67}2p^{0.07},$$

$$r_\infty: 2s^{1.93}2p_z^{1.47}2p_x^{0.28}2p_y^{0.28}3d^{0.03}/1s^{1.0}.$$

The distributions and the CAS wavefunctions above suggest the following bonding diagram:



According to Tables I and III the experimental values^{30,48} of $D_e = 21.68$ kcal/mol (with respect to the adiabatic products), $r_e = 1.1143$ Å and $T_e = 90.93$ kcal/mol are in good agreement with our MRCI results of 20.98 kcal/mol,

1.1164 Å, and 91.85 kcal/mol, respectively. The corresponding values of Lie *et al.*¹⁷⁵ are 18.0 kcal/mol, 1.1113 Å, and 92.72 kcal/mol. Notice that this is the last state that Lie *et al.* have calculated.

G. The $D^2\Sigma^+$ state

The $D^2\Sigma^+$ PEC is illustrated in Fig. 1 and correlates to the $C(^1S)+H(^2S)$ fragments. Its rather ‘‘irregular’’ shape is due to a strong interaction with the $C^2\Sigma^+$ and $G^2\Sigma^+$ states. The $C(^1S)$ state is described by

$$|1S\rangle = (1/3)^{1/2} \{ |2s^2 2p_z^2\rangle + |2s^2 2p_x^2\rangle + |2s^2 2p_y^2\rangle \}.$$

The leading CAS configurations at equilibrium are

$$|D^2\Sigma^+\rangle \approx 0.83 |1\sigma^2 2\sigma^2 3\sigma^2 4\sigma^1\rangle - 0.29 |1\sigma^2 2\sigma^2 3\sigma^1 (1\pi_x^2 + 1\pi_y^2)\rangle.$$

This is essentially the same as the asymptotic CAS wavefunction of the $C^2\Sigma^+$ state, namely:

$$|C^2\Sigma^+\rangle \approx 0.78 |2s^2 2p_z^2\rangle |1s^1\rangle - 0.40 |2s^2 (2p_x^2 + 2p_y^2)\rangle |1s^1\rangle.$$

As the system moves past the equilibrium point, an intense valence–Rydberg mixing takes place at 2.8 bohr with the $G^2\Sigma^+$ state, which correlates to the carbon Rydberg $^3P(2p^1 3s^1)$ state. Although the $D^2\Sigma^+ \leftarrow X^2\Pi$ transition is symmetry allowed, it has not been observed spectroscopically, probably due to unfavorable Franck–Condon factors.

As shown in Table III, at the MRCI level of theory we have obtained $r_e = 1.6635$ Å, $D_e = 9.35$ kcal/mol, and $T_e = 136.11$ kcal/mol.

H. The $d^6\Sigma^-$ state

This is a purely repulsive state (Fig. 1) originating from the $^5S(2s^1 2p^3)$ valence state of C, as is the $c^4\Sigma^-$ state (*vide supra*). It presents a van der Waals attraction of 3.15 cm^{-1} at about 9 bohr.

I. The $E^2\Pi$, $F^2\Pi$, and $H^2\Pi$ states

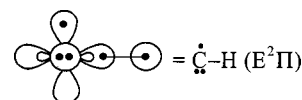
The E , F , and $H^2\Pi$ states correlate to the $^1D(2s^2 2p^2)$, 3P and $^1P(2s^2 2p^1 3s^1)$ Rydberg states of the C atom, respectively. These carbon states span an energy range of 6.42 eV (Table II). The three excited $^2\Pi$ states exhibit avoided crossings at different interatomic distances as illustrated in Fig. 1. The avoided crossing between the E and $F^2\Pi$ states occurs at 2.58 bohr giving the E state a volcanic type appearance with a local minimum at 2.16 bohr. As the H atom approaches from infinity, it is confronted by the 1D electron distribution of C given by $(1/2)^{1/2} \{ |2s^2 2p_x^1 2\bar{p}_z^1\rangle - |2s^2 2\bar{p}_x^1 2p_z^1\rangle \}$. This is an improper distribution for attractive interaction due to the singlet coupling of the p electrons, as opposed to the triplet coupling which leads to the ground state. At the avoided crossing the E state exchanges electron distributions with the descending F state, thus creating the local minimum mentioned above. The generated energy barrier of the E state (the lower of the split levels at the avoided crossing) is 9.69 kcal/mol from the local mini-

um, which has a $r_e = 1.1437$ Å and a $T_e = 169.68$ kcal/mol. The corresponding experimental parameters⁴⁸ are $r_e = 1.15$ Å, and $T_e = 172.7$ kcal/mol. Notice that the state labeled as $D^2\Pi$ in the experimental work^{30,48} is the $E^2\Pi$ state in our work due to the intervening unobserved $D^2\Sigma^+$ state.

The leading CAS configurations and the corresponding Mulliken distributions at the local minimum are

$$|E^2\Pi\rangle \approx 0.92 |1\sigma^2 2\sigma^2 1\pi_x^1 1\pi_y^2\rangle - 0.15 |1\sigma^2 3\sigma^2 1\pi_x^1 1\pi_y^2\rangle - 2s^{1.47} 2p_z^{0.14} 2p_x^{0.98} 2p_y^{1.83} 3d^{0.04} / 1s^{0.47} 2p^{0.06},$$

corresponding to the following vBL bonding diagram:



Upon interaction a total of $0.5e^-$ are transferred from the H to the C atom.

The local minimum of $E^2\Pi$ mirrors the $^3D(2s^1 2p^3, M = \pm 1)$ valence carbon state, 8.0 eV above its ground 3P state, as evidenced from the 3D configuration:

$$^3D(2s^1 2p^3, M = \pm 1) = (1/2)^{1/2} \{ |2s^1 2p_x^1 2p_y^2\rangle - |2s^1 2p_x^1 2p_z^2\rangle \}.$$

The removal of the $2p_z^2$ component, detrimental to attractive interaction, necessitates the involvement of the following valence state of C, not shown in Fig. 1 and lying 1.39 eV above the 3D state:

$$^2P(2s^1 2p^3, M = \pm 1) = (1/2)^{1/2} \{ |2s^1 2p_x^1 2p_y^2\rangle + |2s^1 2p_x^1 2p_z^2\rangle \}.$$

This means that the local minimum corresponds to an intrinsic bond strength (with respect to the 3D state) of 100 kcal/mol.

The $F^2\Pi$ state has a local minimum around 4.8 bohr with an energy barrier of 1.9 kcal/mol and a global wedge-like minimum at $r_x = 2.6$ bohr = 1.375 Å, which is the point of avoided crossing with the $E^2\Pi$ state (*vide supra*). At the MRCI level this ‘‘minimum’’ (the higher of the split levels) is 75.22 kcal/mol, and it lies 181.57 kcal/mol above the ground state (‘‘ T_e ’’). Both minima are with respect to the asymptotic $C(^3P)+H(^2S)$ level. The corresponding experimental values,⁴⁸ also given in Tables I and III are: $r_e = 1.20$ Å and $T_e = 187.6$ kcal/mol (in Ref. 48 this state is designated as $E^2\Pi$).

A state of $^2\Pi$ symmetry correlating to the 3D term of C lends its character to the $H^2\Pi$ state at around 3.2 bohr, which in turn passes its character to the $F^2\Pi$ state via an avoided crossing at about 3.1 bohr (Fig. 1). As a result, the minimum of the $H^2\Pi$ state can be considered as the continuation of the $F^2\Pi$ state. These observations are corroborated by the main CAS configurations at the global ‘‘minimum’’ of the F state:

$$\begin{aligned}
 |F^2\Pi\rangle \approx & 0.72|1\sigma^2 2\sigma^2 3\sigma^1 4\sigma^1 1\bar{\pi}_x^1\rangle \\
 & -0.30|1\sigma^2 2\sigma^2 3\sigma^1 4\bar{\sigma}^1 1\pi_x^1\rangle \\
 & -0.46|1\sigma^2 3\sigma^2 1\pi_x^1 1\pi_y^2\rangle.
 \end{aligned}$$

The 0.72 and -0.30 components express the repulsive part of the $E^2\Pi$ state, while the -0.46 component reflects the attractive part of the $F^2\Pi$ state. The shoulder of this state at about 2.4 bohr, just to the left of the avoided crossing, is caused by an interaction with the $H^2\Pi$ state. There are no spectroscopic data for this last state. For technical reasons (convergence problems at the CI level) we were unable to calculate a full $H^2\Pi$ curve (Fig. 1). Our MRCI results for this state (Table III) are: $D_e = 61.0$ kcal/mol (with respect to the asymptotic products), $r_e = 1.3762$ Å and $T_e = 201.1$ kcal/mol.

J. The $G^2\Sigma^+$ state

The PEC of this state is depicted in Fig. 1 and correlates adiabatically to the $^3P(2p^1 3s^1)$ Rydberg state of C. It has two minima, a local one at about 5.0 bohr with $D_e = 32.7$ kcal/mol, and a global one at $r_e = 1.1482$ Å with $D_e = 69.30$ kcal/mol. The barrier between them is 9.1 kcal/mol measured from the local minimum and sustains at least three vibrational modes (Table IV). It is a result of an avoided crossing with the $I^2\Sigma^+$ state (*vide infra*); this is supported by the leading CASSCF configurations at equilibrium:

$$\begin{aligned}
 |G^2\Sigma^+\rangle \approx & -0.78|1\sigma^2 2\sigma^2 3\sigma^2 5\sigma^1\rangle \\
 & -0.44|1\sigma^2 2\sigma^2 3\sigma^2 4\sigma^1\rangle \\
 & +0.35|1\sigma^2 2\sigma^2 3\sigma^2 6\sigma^1\rangle.
 \end{aligned}$$

The -0.44 component reflects the $D^2\Sigma^+$ state (valence–Rydberg mixing) while the 0.35 component corresponds to the $I^2\Sigma^+$ state.

The experimental values⁴⁸ are $r_e = 1.221$ Å and $T_e = 188.55$ kcal/mol, while our MRCI results are $r_e = 1.1482$ Å and $T_e = 187.36$ kcal/mol (Table III). Notice that in Ref. 30, the G state is referred to as F state.

K. The $I^2\Sigma^+$ state

This state is illustrated in Fig. 1 and correlates to the $^1P(2p^1 3s^1)$ Rydberg carbon state. Due to technical reasons (convergence problems at the CI level) a part of the PEC from 3.35 to 3.85 bohr has not been completed. The leading CAS configurations at equilibrium are:

$$\begin{aligned}
 |I^2\Sigma^+\rangle \approx & 0.81|1\sigma^2 2\sigma^2 3\sigma^2 6\sigma^1\rangle \\
 & +0.41|1\sigma^2 2\sigma^2 3\sigma^2 4\sigma^1\rangle \\
 & +0.19|1\sigma^2 2\sigma^2 3\sigma^2 5\sigma^1\rangle.
 \end{aligned}$$

The 0.41 component represents the valence–Rydberg mixing with the $D^2\Sigma^+$ state and the 0.19 component reflects the mixing with the $G^2\Sigma^+$ state.

Our MRCI results, listed in Table III, are $D_e = 57.85$ kcal/mol, $r_e = 1.2639$ Å, and $T_e = 204.15$ kcal/mol. There are no experimental results in the literature for this state.

L. The $J^2\Delta$ state

Part of the $J^2\Delta$ state around equilibrium is depicted in Fig. 1; it stems from the 3D valence state of C. No experimental values exist in the literature. Our MRCI results, listed in Table III, are: $D_e = 48.9$ kcal/mol, $r_e = 1.666$ Å, and $T_e = 221$ kcal/mol.

Although there are no experimental results for the $H^2\Pi$, $I^2\Sigma^+$, and $J^2\Delta$ states, an unidentified state of either $^2\Pi$, $^2\Sigma^+$, or $^2\Delta$ symmetry has been reported^{30,48} with $T_e = 213$ kcal/mol.

IV. SYNOPSIS

Using large, correlation consistent basis sets and MRCI (CASSCF+1+2) methods, we have computed PECs for 15 states of the CH radical, spanning an energy range of 9.6 eV. Our results can be considered in quantitative agreement with existing experimental findings. In particular, for the $X^2\Pi$ state (after correction for core correlation effects), our D_e value is smaller than the experimental one by 0.43 kcal/mol ($=150$ cm⁻¹) and our r_e value smaller by 0.0014 Å compared to Ref. 86 and larger by 0.00034 compared to Ref. 147. It is interesting that the bond lengths of all bound states reported in Table III are improved upon applying a uniform 0.002 Å contraction due to core-valence correlation effects.^{233,234}

In general, 0.2–0.5 e^- are transferred from H to C upon molecule formation. Dipole moments range from 0.16 D ($I^2\Sigma^+$) to a remarkably high value of 6.17 D ($G^2\Sigma^+$). Finally, the multireference Davidson correction allows us to claim that for the first ten states, valence correlation has been extracted to within 2 mhartree.

Note added in proof. While in press X. Li and Y.-P. Lee [J. Chem. Phys. **111**, 4942 (1999)] reported a $T_0(D^2\Pi \leftarrow X^2\Pi) = 58\,980.592(53)$ cm⁻¹, in excellent agreement with our $T_0 = T_e + \Delta\omega_e/2 = 169.68$ kcal/mol + (-54.0 cm⁻¹) = 59 292 cm⁻¹. Please note that the $D^2\Pi$ state of Li and Lee is referred to as $E^2\Pi$ in the present paper (see text).

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