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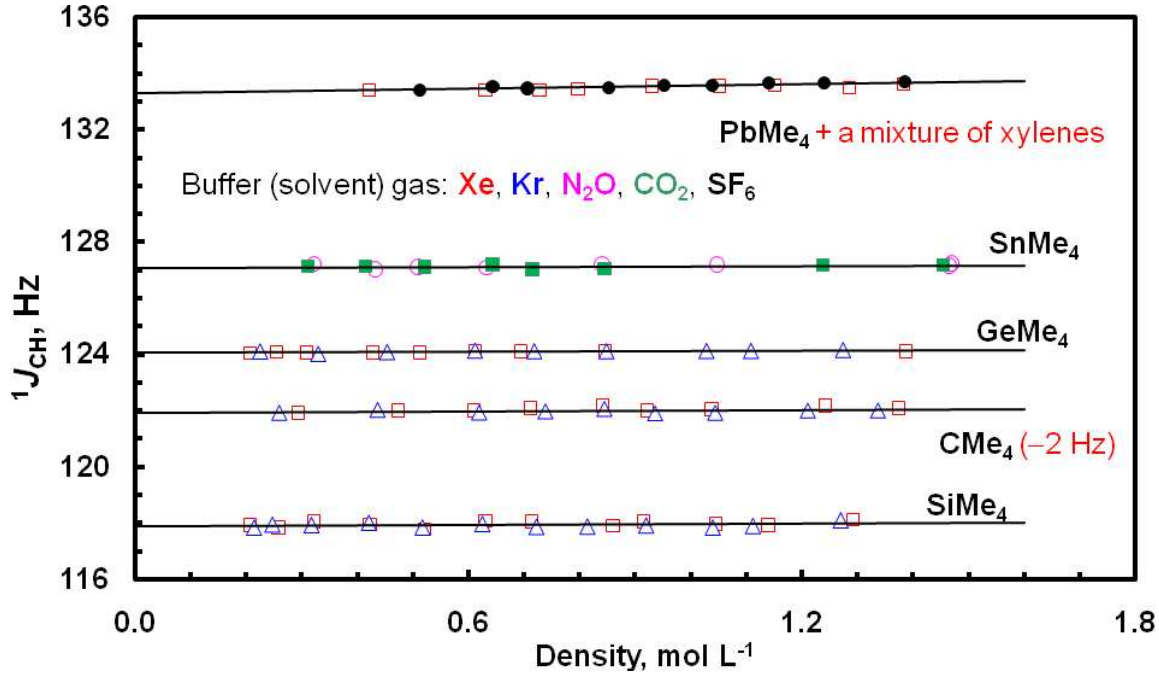
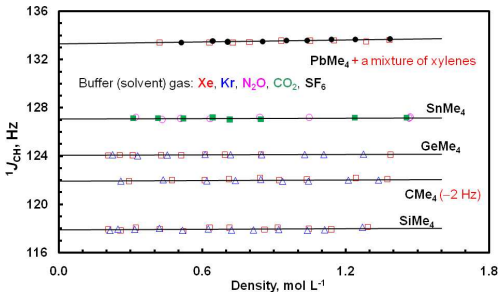
A graphical and textual abstract

The gas-phase $^{1}J_{0,CH}$s in 'isolated' molecules of EMe_4 were determined and discussed in terms of their geometric/electronic properties obtained from the DFT calculations.



Small version (40 x 80 mm) – the EXCEL file



$^1J_{CH}$ couplings in Group14/IVA tetramethyls from the gas-phase NMR and DFT structural study: a search for the best computational protocol†

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Abstract: Four tetramethyl compounds EMe_4 (E = C, Si, Ge, and Pb) were studied by high-resolution NMR spectroscopy in gaseous and liquid states at 300 K. Extrapolation of experimental vapor-phase C–H *J*-couplings to a zero-pressure limit permitted determining the $^1J_{0,CH}$s in methyl groups of their nearly isolated molecules. Theoretical predictions of the latter NMR parameters were also performed in a locally dense basis sets/pseudopotential (Sn, Pb) approach, by applying a few DFT methods pre-selected in calculations of other gas-phase molecular properties of all these species and $SnMe_4$ (bond lengths, C–H stretching IR vibrations). A very good agreement theory *vs.* experiment was achieved with some computational protocols for the all five systems. The trends observed in their geometry and associated coupling constants ($^1J_{CH}$s, $^2J_{HH}$s) are discussed and rationalized in terms of the substituent-induced rehybridization of the methyl group (treated as a ligand) carbon, by using Bent's rule and newly proposed, theoretically derived values of the Mulliken electronegativity (χ) of related atoms and groups. All these χ data for the Group-14/IVA entities were under a lot of controversy for a very long time. As a result, the recommended χ values are semi-experimentally confirmed for the first time and only a small correction is suggested for χ(Ge) and $\chi(GeMe_3)$.

Keywords: tetramethyl compounds, carbon family, tin and lead organoderivatives, gas-phase NMR spectra, molecular properties, carbon-proton coupling constants, pseudopotentials, local dense basis sets, Bent's rule, Mulliken electronegativity, tetrahedral distortion

† Electronic supplementary information (ESI) available: Tables S1-S9 and Figs S1-S8 (11 pages, PDF). See DOI: 10.1039/

‡ Physical image *vs.* molecular structure relation, Part 18. For Part 17, see ref. 79.

1 Introduction

High-resolution nuclear magnetic resonance (NMR) spectroscopy is undoubtedly the most valuable technique for determining the structure and motion of molecules in all states of aggregation. Among various NMR parameters, magnetic shieldings σ (or related chemical shifts, δ) and scalar spin-spin coupling constants [hereafter referred to as *J*-couplings or J_{XY}s] are the most important physical observables. Nowadays, first-principles calculations of the latter NMR data are available and published for many objects.[1-4] But their performance is not as straightforward as predicting the σ data. As viewed from the classical (nonrelativistic) Ramsey theory of J_{XY}s,[5] there are four interactions between nuclear spins in molecules that can contribute to an indirect *J*-coupling, *i.e.,* the Fermi contact (FC), spin-dipole (SD), paramagnetic spin-orbit (PSO), and diamagnetic spin-orbit (DSO) term. Although, in most cases, the FC contribution dominates the isotropic *J* value, none of these terms can be *a priori* excluded in any reliable J_{XY} calculations.[1-4] Moreover, more advanced relativistic approach is needed for *J*-couplings embracing heavier elements of the 5th and higher rows of the periodic table.[4,6] Fortunately, the impact of such atoms (*e.g.,* beyond Kr) on J_{XY}s that do not directly involve these nuclei is small unless a heavy nucleus is located directly in the coupling path.[3,4] Currently, density-functional theory (DFT) computations seem to be the most promising in *J* evaluations for small and medium-sized molecules, such as, *e.g.,* various organo-derivatives of tin[7] and lead. It is obvious that all theoretical calculations may be verified only when accurate experimental data are accessible. The most valuable slices of such data come from the gas-phase studies, because results of this kind are not obscured by bulk effects arising from the surrounding medium and molecular effects coming from specific interactions.

In this paper we present experimental results from the vapor-phase measurements of $^1J_{CH}$ couplings for a complete set of tetramethyl derivatives of the main Group-14/IVA elements, namely, 2,2-dimethylpropane (neopentane, CMe_4), tetramethylsilane (TMS, $SiMe_4$), tetramethylgermane ($GeMe_4$), tetramethyltin ($SnMe_4$), and tetramethyllead ($PbMe_4$). Among all these isostructural systems EMe_4, where the central element E = C to Pb, only the Sn derivative was studied in such a way very recently.[8] These species were analyzed by means of 500 MHz ^{1}H NMR spectra taken for their gaseous mixtures with krypton (Kr), xenon (Xe), sulfur hexafluoride (SF_6), nitrous oxide (N_2O), and carbon dioxide (CO_2) used as buffer gases. Indirect $^1J_{CH}$ couplings, evaluated from the ^{13}C satellite lines of related methyl group signals, were then extrapolated to zero-pressure limit.[9] Such a procedure leads to determining the

$^1J_{0,CH}$s practically free from solvent effects. The latter experimental data were subsequently used in verification of related NMR J_{CH}^{theor}s predicted with various locally dense basis sets (LDBS)[10-13]/effective core pseudopotential (ECP, for Sn and Pb) approaches, by using a few pre-selected DFT-level treatments. All these two-step protocols were *a priori* tested in computations of other gas-phase properties of these objects EMe_4, such as bond lengths and C–H stretching infrared (IR) vibrations. A general methodology of quite accurate calculations of the geometries and spectroscopic properties of these organoelement compounds was established in this way. Related $^1J_{CH}$s were also measured for neat (pure liquid) samples. A consideration of such NMR data brings up the magnitude of the gas-to-solution effects on the one-bond $^{13}C–^1H$ *J*-coupling. Finally, the impact of the change in a central E atom on J_{XY}s in the attached methyl groups (treated as ligands) is discussed in terms of Bent's rule, new theoretical values of the Mulliken electronegativity of atoms (χ_a) or functional groups (χ_g), and distortion of a tetrahedral symmetry around the methyl carbons in all analyzed molecules. These latter efforts were also targeted a semi-empirical verification of newly determined χ values mentioned above, being under considerable controversy for a very long time.

Four crucial and inseparable issues were addressed in this work: (*i*) the highly reliable representation of a molecular geometry of all compounds EMe_4, (*ii*) adequate calculations of $^1J_{CH}$s and $^2J_{HH}$s operative in these isostructural model systems, (*iii*) the interpretation of trends in the J_{XY} values observed for the species EMe_4 in terms of changes in the molecular shape and nett charge on their methyl carbons (reflected by the recent Mulliken electronegativities of attached groups), and (*iv*) the experimental NMR data-based validation of these electronegativity data for the all five Group 14/IVA elements and related EMe_3 substituents. As far as we know, such a comprehensive internally consistent structure- and NMR-oriented investigation of the title systems has not yet been published.

2 Experimental details

2,2-Dimethylpropane (neopentane, Merck-Schuchardt, 98%) from a lecture bottle with a liquified gas, tetramethylsilane (TMS, Aldrich, ≥99.9%), tetramethylgermane (Aldrich, 98%) and tetramethyllead (Aldrich, 65 wt.% solution in xylene) from a glass container, were used without further purification. As reported in detail elsewhere,[6,8,9b,14-16] gaseous samples were obtained by condensation of the studied compound and pure commercially available solvent gas (Kr, Xe, or SF_6) transferred from the calibration part of our vacuum line. The solute gas,

EMe_4, was applied in small concentration (~0.005 mol/L, pressure ~100 Torr) and was mixed with different quantities of a gaseous solvent (from ~0.2 to ~1.5 mol/L).

One-dimensional ^{1}H NMR spectra were recorded at 300 K on a Varian INOVA FT-NMR spectrometer operating at the 500.6075 MHz proton Larmor frequency. The FID acquisition time was set to 2 s and the spectral width from 400 to 1000 Hz was applied. $^1J_{CH}$ couplings were directly evaluated in Hz as differences between the outside ^{13}C satellites of methyl group signals. Liquid TMS was always used as an external NMR chemical-shift reference standard.

3 Computational details

All electronic structure 'gas-phase' calculations, *i*.e., for isolated vibration-free objects at 0 K in vacuum, were performed applying the Gaussian 09 suite of programs,[17] by starting with the tetrahedral models of EMe_4 systems. The hybrid B3LYP functional as implemented in the Gaussian code was employed,[18] because its use has been found suitable for a large variety of previous computations, including the Sn and Pb containing species.[7,19-21] This exchange-correlation density functional has been one of the more successful tools for predicting different magnetic properties of organic systems.[1,18,22-25] The threshold and convergence parameters were tightened in all geometry optimizations (carried out by taking advantage of the T_d symmetry) and frequency calculations by specifying 'Int=UltraFine', 'SCF=VeryTight', and 'Opt=VeryTight' for numerical integration grids and convergence criteria in Gaussian;[16] the former two keywords were also used in subsequent single-point NMR predictions. Moreover, the five pure *d*-functions have been mainly applied (5D option), see also Table S1 (ESI†). Corresponding vibrational wavenumbers, ωs, were computed analytically at the same theory level in a rigid rotor-harmonic oscillator approximation. These data were also applied to verify that all structures, localized as stationary points on pertinent Borne-Oppenheimer energy hypersurfaces, represent real global minima (NImag = 0). Finally, a gauge-including atomic orbital (GIAO) formalism[26] was employed to ensure the gauge independence of related calculated J_{XY}s. Several pre-selected functional/basis set combinations were used as DFT protocols at two stages. A locally dense basis sets (LDBS)] approach,[10-13] was applied in the vast majority of calculations. In addition, simulations of the impact of nonpolar molecules of CCl_4 and C_6H_6 on the geometry and NMR response properties of species EMe_4 were performed in an improved equilibrium solvation protocol[27] of the polarizable continuum model of solvation (IEF-PCM).[28]

Thus, in the first step of a ground-state-geometry optimization, a centrally situated E atom was best represented by an economical def2-TZVPP[29] (hereafter referred to as **A**) or def2-

TZVPPD[30] (**B**) basis set of triple-ζ valence quality, while the small double-ζ Gaussian-type orbitals (GTOs), that is, 6-31G(*d*,*p*) (**C**), 6-31+G(*d*,*p*) (**D**) or 6-31+G(2*df*,*p*) (**E**) bases,[17] were applied for the C and H atoms in methyl groups. The usage of basis sets **C** and **D** for analogous organometallics was found as a fully sufficient approach.[19,20] The aug-cc-pVTZ (C,[31] Si,[32] Ge[33]) and aug-cc-pVTZ-PP (Sn and Pb)[34] (**F**) basis sets were also *a priori* tested for the E atom, but with rather worse results. In these **A**, **B**, and **F** bases an all-electron description concerns C, Si, and Ge – however, the innermost orbitals of Sn and Pb are represented by small-core quasi-relativistic energy potentials known as effective core pseudopotentials (ECPs).[7] In turn, five other basis sets were exploited for C and H in the second computational step, instead of the Pople style bases mentioned above, namely, IGLO-II (**G**), IGLO-III (**H**), aug-cc-pVTZ-J (**I**), pcJ-2 (**J**), and pcJ-3 (**K**). The former two Huzinaga-type atomic basis sets[35] modified by Kutzelnigg and coworkers[36,37], also known as HII and HIII bases,[38-41] have been widely used for calculations of NMR properties;[36-45] *e.g.,* the compact basis set **G** of polarized triple-ζ valence quality consists of an [9*s*5*p*1*d*]/(5*s*4*p*1*d*) C and [5*s*1*p*](3*s*1*p*) H functions in the [primitive]/(contracted GTOs)] notation.[41] These latter aug-cc-pVTZ-J[46] and pcJ-*n* bases[23] were specifically developed for *J* calculations. As for the E atoms, the following basis sets of quadruple-ζ valence quality were used for the J_{CH} prediction: def2-QZVPP[29] (**L**), def2-QZVPPD[30] (**M**), (aug-)cc-pVQZ (C,[31] Si,[32] Ge[33]) and (aug-)cc-pVQZ-PP (Sn and Pb)[34] [**N** and **O** (augmented)].

In addition, a few runs were conducted in order to verify the adopted methodology. In one series, the B3LYP functional was applied, at both computational steps, in conjunction with the all-electron basis sets **G** available for C-Sn. Moreover, owing to some reported[22] disadvantages of B3LYP in $^1J_{CH}$ calculations, two other density functionals were used, *i.e.,* PBE[47] and PBE0,[48] in a PBE/**I**(*C,H*),**L**(*E*)//PBE0/**E**(*C,H*),**A**(*E*) approach[49] similar to that employed for hydrocarbons.[22]

On the whole, 15 different basis sets **A**-**O** were used in conjunction with related DFT functionals in the two computational steps; for their application in 14 different B3LYP/basis sets combinations tested to obtain the best results, see Table S1 (ESI†). All atomic orbitals inaccessible within Gaussian[17] were downloaded from the EMSL base (ver. 1.2.2)[50] and applied in their standard forms with optimized general contractions. The only exceptions were **G** orbitals for the Ge and Sn atoms, created by O. Malkina and V. G. Malkin, taken from an earlier release of the StoBe package.[51] Such Sn basis has been used very recently.[52] Only results from two most suitable procedures [*i.e.,* B3LYP/**G**(*C,H*),**L**(*E*)//B3LYP/**C**(*C,H*),**A**(*E*)

and, especially, B3LYP/**G**(*C,H*),**M**(*E*)// B3LYP/**C**(*C,H*),**B**(*E*) (henceforth called protocols I and II, respectively)] are discussed in depth, for the sake of brevity. An employment of a basis set **J** instead of **G** in both above protocols (referred to as approaches I-pcJ-2 and II-pcJ-2, respectively) leads to similar or slightly worse results. Generally, the use of IGLO-II (**G**) in the second step was found the most reliable and efficient protocol of the DFT NMR calculations presented here. The relative computer time required for a single-point *in vacuo* B3LYP-prediction of *J*-couplings in $Ge(Me)_4$ with the use of basis sets **G**, **H**, **J**, and **K** was determined as 1.0, 1.9, 4.7, and 34.4, respectively.

4 Results and discussion

In the gaseous phase, nuclear spin–spin coupling constants are modified by pairwise interactions between the coexisting molecules and by additional multiple interactions. At a constant temperature, the appropriate equation for a measured one-bond heteronuclear X,Y *J*-coupling value in a binary mixture of two gases, *i.e.,* a solute gas (A) and buffer gas (B), is expressed as

$$J_{XY} = J_{0,XY} + J_{AA,XY}\cdot\rho_A + J_{AB,XY}\cdot\rho_B + \ldots. \qquad (1)$$

where $J_{0,XY}$ is the *J*-coupling between the nuclei X and Y at a zero-density limit, whereas couplings $J_{AA,XY}\cdot\rho_A$ and $J_{AB,XY}\cdot\rho_B$ are solely due to intermolecular effects coming from binary collisions of the A–A and A–B molecules, respectively. Usually, the density of a solute gas A (ρ_A) is kept sufficiently low and so the above relation, eqn (1), is simplified to a linear equation of the form (Eq. 2)

$$J_{XY} = J_{0,XY} + J_{AB,XY}\cdot\rho_B \qquad (2)$$

valid in a middle range of pressures,[9a] as the term $J_{AA,XY}\rho_A$ can be safely ignored if only micrograms of a solute A are present in the gas-phase solution. Usually, this approximation is additionally verified when at least two different gaseous solvents (buffers) are carefully used in the same apparatus and $J_{0,XY}$s obtained by an extrapolation remain the same within an experimental error. It is usually assumed that an averaged value of the parameter $J_{0,XY}$ derived in this way is very close to its *in vacuo* value for an isolated solute molecule.

In this work, one-bond J_{CH} couplings between constituent atoms of the methyl groups (treated as ligands surrounding the centrally situated E atoms) in species EMe_4 were measured with good accuracy. Also their determination was made with very high precision, because methyl group ^{1}H NMR signals of all these systems were flanked by sharp lines of more or less

symmetrically disposed ^{13}C satellites separated by J_{CH}s. The $J_{0,CH}$ values obtained from the extrapolation of such experimental data points to a zero density of the buffer gas are summarized in Table 1, where intermolecular effects on such couplings are also listed. As can easily be seen in Fig. 1, the J_{CH}s are affected by different buffer gases in different ways. The greatest concentration dependence of $^{13}C–^{1}H$ *J*-couplings was found for $PbMe_4$; $J_{AB,CH} > 200$ Hz·mL mol^{-1}. Related J_{CH} values measured for the neat samples of all tetramethyls are also given in Table 1, together with related literature data for liquids or solutions.[53-55] Inspection of this table shows a small difference $\Delta J_{CH} = J_{CH}(\text{neat}) – J_{0,CH,av}(\text{gas})$ slowly growing from CMe_4 to $PbMe_4$, which may be recognized as a modulation of the $J_{0,CH}$ coupling – a solvent independent characteristic of the solute molecule – induced by a surrounding medium. Indeed, there is good, $R^2 = 0.9948$, expected in view of ref. 24, quasi-linear correlation $\Delta J_{CH} = f\,[2(\varepsilon–1)/(2\varepsilon+1)]$ between the *J* data and values of the relative permittivity (formerly called dielectric constant, ε) of three normally liquid tetramethyls,[56] accounted *via* the reaction field function (Fig. S1, ESI†); unfortunately, the value of ε for $PbMe_4$ is not available in the literature. The above observation is also in line with the finding that the solvent induced change in the $^{1}J_{CH}$ coupling in a given solvent depends on the *J* value itself.[24]

The foregoing vapor-phase $J_{0,CH}$s and related liquid/solution data were subsequently compared with the high-quality computational results on J_{CH}s in all five species EMe_4 obtained for the first time. It is well-known that the Fermi-contact (FC) contribution to this type of *J*-couplings is a dominating term.[1] Furthermore, this contribution is very sensitive to geometry and an applied calculational protocol (basis-set size and inclusion of the electron-correlation, in particular), while the SD, PSO, and DSO terms are less sensitive to these factors.[1,38] Hence, special efforts were made in choosing a suitable DFT treatment. The use of experimental geometries often employed in *J* calculations on small molecules[39,57] was discarded at the beginning, because our aim was to elaborate an accurate and cheap (from the viewpoint of calculation time) methodology available for other similar organic/organometallic species. Accordingly, only a few pre-selected computational approaches were used here.

4.1 Selection of two-step computational protocol

In the first stage, reliable equilibrium ground-state geometries of tetramethyls of all members of the periodic table Group 14/IVA were modeled, while in the second estimations of their NMR spectroscopic parameters were made at a higher theory level, on the optimized geometries. The same DFT-level approach had to always be used for all objects, so that such

computed J_{XY}s are comparable and mutually consistent. The presence of heavy atoms in $SnMe_4$ and $PbMe_4$ forced the use of relativistic effective core pseudopotentials (ECPs) to describe their innermost orbitals. Recent results[21] on several diatomics demonstrate that the B3LYP functional combined with a mixture of LANL08d[58] (as an ECP) for Sn and Pb, and of a triple-ζ 6-311G(*d*,*p*) basis set for H and C atoms provides a sensible compromise between accuracy and computational cost. But an application of def2-TZVP[29] instead of LANL08d was capable of predicting some molecular properties of these species slightly more accurately.[21] As a consequence, only a few sufficiently flexible basis sets were tested and used in this work, for which the ECP approximation is fully consistent with an associated all-electron treatment available for the whole carbon group elements, *i.e.,* atoms C through Pb.

It is well known, that NMR parameters depend critically on the electron density in the nuclear regions which may not be accurately modeled by the ECPs.[1] However, we were mainly interested in the reliable evaluation of J_{XY}s within the methyl groups around E atoms. Hence, our approach was fully compatible with the ECP-based studies on NMR properties of organic ligands linked to the metal atoms.[59] Thus, all J_{CH}s in species EMe_4 were calculated with Gaussian 09 using the GIAO formalism coupled with B3LYP, as this DFT protocol was found suitable for many similar computations performed previously.[1,18,24]

Generally, a locally dense basis sets (LDBS) method[10-13] was applied. However, in the geometry optimization step, this idea was employed in a manner opposite to that of the originally formulated one by Chesnut *et al.*[10] *i.e.,* the E atom was described by the large (locally dense) basis set to reproduce the E–C bond distances as well as possible. Indeed, some problems with the choice of basis sets were reported for predicting the E–H bond lengths in Group-14/IVA tetrahydrides.[60] The LDBS approach was successfully used in the calculations of many molecular properties including NMR chemical shifts[10] and J_{XY}s.[12,13] Thus, fully balanced basis functions of the same level were employed here for C and H atoms of methyl groups, that is, double-zeta split-valence Pople's bases and more extended Huzinaga-Kutzelnigg-type IGLO-II (or IGLO-III) atomic orbitals in our first and second step, respectively. The former compact basis set was widely adapted for calculating the NMR response of many different tin-containing species.[38,42,52] It is also known that subsequent use of IGLO sets at the geometries B3LYP-optimized in conjunction with double-ζ Pople's bases lead to adequate prediction of the experimental J_{CH} couplings.[41,43-45]

Let us consider our results on some interatomic distances in five molecules of EMe_4. All such geometric data obtained by a pre-selected DFT protocol II are collected in Table S2

(ESI†); for the plot, related statistics and Cartesian coordinates of all these species, see Figs S2 and S8-S10 (ESI†), respectively. Inspection of Table S2 reveals an excellent agreement (R^2 = 0.9998) with the experimental r_g bond lengths from an electron diffraction analysis of vapor-phase samples,[61-65] while perfect consistency (R^2 = 0.9999) was found for the carefully examined CMe_4.[61] But such r_g data[65] for $PbMe_4$ seem likely to be underestimated, especially for the Pb···H distance. Obviously, there is some dissimilarity in the physical meaning of these two quantities, *i.e.,* thermally averaged r_gs measured for vibrating species at room temperature *vs.* equilibrium r_es computed for motionless objects at 0 K. But their ratio is approximately constant over a wide distance range as these geometric parameters are related by the equation $r_e \cong r_g - 3au^2/2$,[66] where a and u is an associated Morse anharmonicity constant and a root-mean-square vibration amplitude for a related atom pair, respectively. A very good agreement (R^2 = 0.9993) was also achieved between our C–E distances and those B3LYP-optimized within a relativistic elimination of small components (RESC) approach[67] (ESI†, Table S2). Surprisingly, the greatest difference between such r_e data is found for $SiMe_4$. Some shortening of the C–E bond lengths was, however, found in a relativistic DFT study on less sterically crowded molecules $H–EMe_3$ possessing the post-*d* main-group central atom Ge-Pb.[68]

Very good conformity was also found (R^2 = 0.9994 and 0.9998 for the protocols I and II, respectively) in a comparison of *in vacuo* computed antisymmetric C–H IR stretchings, $\omega_{as}(CH_3)$, and their vapor-phase experimental counterparts, $\nu_{as}(CH_3)$, determined by the same researchers,[69] see Table S3 and Fig. S3 (ESI†). Unexpectedly, two such values were stated[69] for CMe_4 but only one compatible with the other independently reported[70] values of this fundamental. Present calculations supported these one-value results. So, the problem of a relation between two types of molecular parameters recurs again that is commonly overcome with a linear scaling theoretical data, by using the equation $\nu = \lambda \cdot \omega$[71,72] or $\nu = a + b \cdot \omega$.[72] A better agreement obtained in the approach II also justifies a modification of the basis set **A** to **B**.[30] The aforementioned findings confirmed our belief that the computed T_d symmetric geometries of title tetramethyls are sufficiently accurate, especially those found with protocols I and II. Hence, all subsequent predictions of their NMR spectroscopic properties based on such molecular structures should also be reliable.

4.2 DFT calculation results on $^1J_{CH}$s

At this point, we present our gas- and solution-phase computational results on total values of J_{CH}s in five molecules EMe_4 obtained with the approaches I and II, including their

partitioning into all four Ramsey contributions with a dominant FC term participation as expected;[1-4,6,25] see Table 2 and S4 (as its extended version, ESI†). The *J* values denoted as CCl_4 or C_6H_6 are from the implicit solvent IEF-PCM[27,28] simulations mimicking solvation of solutes EMe_4 by molecules of carbon tetrachloride or benzene. A typical approach taking into account the scaling theoretical results according to the measured *J* data was used for assessing the $J_{XY}^{theor} = f(J_{XY}^{exp})$ correlations, as in our previous works[44,45,73] and in line with a recent benchmarking study.[25] Generally, a fairly good agreement with an experiment was found (R^2 = 0.9847, 0.9879, 0.9845, 0.9840, 0.9892, and 0.9930 for the approaches I, II, I-pcJ-2, II-pcJ-2, II-C_6H_6, and II-CCl_4, respectively). The only one outlying data point for CMe_4 was at first glance a considerable surprise in consideration of a relative simplicity of J_{CH}-coupling predictions for hydrocarbons (*vide infra*). The best linear four-point plot $J_{CH}^{theor} = f(J_{0,CH})$ for gaseous samples is shown in Fig. 2. In either case, *i.e.,* using IGLO-II *vs.* IGLO-III or pcJ-2 *vs.* pcJ-3, the better conformity in the second computational step was found with the basis set of smaller size. This suggests a fortuitous cancellation of calculational errors on the geometry and NMR spectroscopic properties occurring with the use of smaller basis sets.[3] On the other hand, a little difference (1.1 Hz, protocol II) between the gas-phase experimental data for CMe_4 and its predicted $^1J_{CH}$ value is considerably smaller than *e.g.,* the 6-7 Hz discrepancy obtained at a similar theory level for some strained carbon atoms.[41] An omission of the CMe_4 point gives perfect relationships with R^2 of 0.9999, 0.9999, 0.9997, 0.9997, 0.9995, and 0.9999, respectively. In this case, a slope of 1.036 and an intercept of –5.34 Hz [or only –0.34 Hz, after the +5 Hz arbitral correction for a zero-point vibrational (ZPV) contribution to the C–H ^1J-couplings[22,25,40]] was found for the best approach II, close to the ideal values of unity and zero. Hence, one can infer that $J_{0,CH}$s [or $J_{CH}(CCl_4)$s] and pertinent J_{CH}^{theor}s are perfectly correlated for the four remaining tetramethyls. The use of the recommended PBE/**I**(*C,H*),**L**(*E*)//PBE0/**E**(*C,H*),**A**(*E*) protocol[22] afforded R^2 of 0.9702 and 0.9998 after exclusion of CMe_4. Even worse results were found in the all-electron B3LYP/**G**//B3LYP/**G** approach available for the E atoms from C to Sn; R^2 of only 0.9456 and 0.9965 (without CMe_4). In view of the latter finding, the employment of the LDBS approach is fully justified.

It is worth mentioning that our PCM calculations for CCl_4 solutions failed to reproduce the observed trends, because they predict a slight decrease of J_{CH}s *vs.* $J_{0,CH}$s, from -0.00_9 Hz (Si) to -0.09_5 Hz (Pb) in the I-CCl_4 data and from -0.01_2 Hz (Si) to -0.08_8 Hz (Pb) in II-CCl_4 [Table S4 (ESI†)], in contradiction to a small increase accessed experimentally (Table 1); the protocol II again turned out to be better tool, however. A lot more remarkable is the

'abnormal' position of the CMe_4 data point on the plot $J_{CH}^{theor} = f(J_{0,CH})$, see Fig. 2. This observation, apparently not fulfilling a 'periodic correlation', was made many years ago for liquid/solution samples of all five tetramethyls,[53-55] but was only tentatively rationalized by (a) the effectiveness of an E–C orbital overlap[53a] or (b) the C–H bond rehybridization.[54] The usage of the latter factor as an only criterion was criticized for molecules with the possible impact of electronegativity.[74] The most probable explanation of this irregularity will be given afterwards (*vide infra*).

The next crucial data shown in Table 2 relates to spin-dipole (SD) contributions to the total values of J_{CH}^{theor}s in five systems EMe_4. In this instance, our data fulfilled the expected 'periodic correlation', at first sight. Thus, the increase of this generally small term was found as one goes down the periodic table from CMe_4 (0.18 Hz) to $PbMe_4$ (0.44 Hz), regardless of the computation level used. Such a finding can be explained by a gradual change in the central E atom that modulates interactions of a nuclear magnetic moment with an electron spin within the methyl groups in question. An increase of the SD term with the electronegativity is known for the directly coupled nuclei,[38,57] including $^1J_{EH}$s in tetrahydrides EH_4 (E = C to Sn).[38] To gain further information, various correlations were considered, by using the newly proposed values of atomic Mulliken electronegativities (χ_as) and group electronegativities (χ_gs) of the EMe_3 substituents attached to the methyl carbons in species EMe_4 *versus* the SD term magnitudes. All these χ_a and χ_g values (Table 3) were theoretically determined by Giju *et al.* [68] with the inclusion of relativistic effects. In the cited work, an issue of the controversial electronegativity of the Group 14/IVA elements and, especially, related EMe_3 substituents is discussed in details, but without experimental evidences for the correctness of these novel χ data. The NMR results considered in the present paper afforded such a possibility, for the first time (see also below). Thus, for SD term values mentioned above the best least squares regression line ($R^2 = 0.997$) was really found with the recent χ_gs, but again after omission of the CMe_4 outlier. The plot with this bounding point is shown in Fig. 3. Obviously, any linear correlation embracing the SD terms of five species EMe_4 does not exist.

4.3 Discussion of observed trends

As seen in Tables S2 and S3 as well as Figs S2 and S3 (ESI†), some 'periodic tendencies' in the geometry and infrared properties of EMe_4 are clearly shown. Indeed, related interatomic distances and, especially, C–H stretching IR frequencies change monotonically. It should be highlighted that differences in the computed tetrahedral-like geometry around the methyl

carbon atoms in these species agree with Bent's model of isovalent hybridization.[75-78] According to this concept, in a molecule with inequivalent R groups, smaller bond angles are formed between the electronegative R groups since a central atom tends to direct the sp^n-hybridized orbitals of greater *p* character towards its more electronegative R substituents. Based on the foregoing χ_g data of the groups attached to the methyl carbons (Table 3), it is clear that the fractional *s* character of a carbon 2*s* orbital forming the C–H bond in species EMe_4 follows the order Si < Ge < Sn < Pb, exactly the same as is observed from the $^1J_{CH}$ data. Thus, an 'abnormal large' *J*-value found for CMe_4 as an outlier (*vide supra*) is rationalized now to some degree, because such C-H couplings depend mainly on the FC interactions occurring for *s* electrons.[54,74]

In order to discuss the gas-phase $J_{0,CH}$-couplings in terms of the χ_as and χ_g values,[68] two series of these data were considered for the SD terms as previously; the point due to CMe_4 was found always to be a strongly outlying one. In this case, a better four-point regression line plot was found for χ_as; $R^2 = 0.963$ (Fig 4) *vs.* $R^2 = 0.950$ for χ_gs (Fig. S4, ESI†). It was also of interest to see how strong other linear relationships expected for compounds EMe_4 are, *i.e.,* $^1J_{0,CH} = f(^2J_{HH})$, $\angle HCH = f(^2J_{HH})$, $\chi_a = f(^2J_{HH})$, and $\chi_g = f(^2J_{HH})$. The $^2J_{HH}$s = 6.51 × $^2J_{DH}$s (provided that there is no primary isotope effect on the *J* values) for species $E(CH_2D)_4$ in CCl_4 solution[55] and our PCM computed HCH bond angles were used accordingly (Table 3). Corresponding plots reflecting mutually consistent correlations (R^2 = 0.994, 0.996, 0.973, and 0.957, respectively), again after omission of CMe_4, are shown in Figs S5-S8 (ESI†). Thus, the 2° opening of the HCH angle on going from Si to Pb causes an increase in the $^2J_{HH}$ value (from −14.05 to −10.94 Hz)[55] and an increase in the *s* character of the C–H carbon sp^n-hybridized bonding orbitals, as expected. All the above data dealing with Mulliken electronegativities, based on a combination of experimental and DFT results, seem to prove the legitimacy of newly reported χ_a and χ_g values.[68] As far as we know, such χ data for this type of Group-14/IVA entities are confirmed for the first time. The analysis of the five plots [Figs 3, 4, S4, S7, and S8 (ESI†)] suggests, however, the need for a decrease of χ_a to ~4.54 eV and increase of χ_g to ~3.88 eV for Ge and -$GeMe_3$, respectively, because there is $0.989 \leq R^2 \leq 0.999$ for the above correlations with the use of these corrected χ values.

Finally, let us come back to some geometric parameters predicted for the molecules EMe_4 (Table 3). It is worth noting that all changes in their ECH and HCH bond angles also follow the trends anticipated from Bent's rule;[75-78] the influence of simulated 'solvation' by molecules of CCl_4 or C_6H_6 on the geometry is negligible. Indeed, such angles found for

CMe_4, being close in magnitude to those for $GeMe_4$, reproduce the relation $J_{CH}(C) \leq J_{CH}(Ge)$ very well and so excellently explain the 'abnormal' position of the CMe_4 data point in Fig. 2. Furthermore, the C–H distance in this object is in between those found for $SiMe_4$ (the longest length, the greatest *p* character) and $GeMe_4$. It seems that the computed geometry around all five methyl carbons in species EMe_4 is mainly dependent on the C–E distance, which forces large modification of the methyl group shape. Indeed, its strongly 'stretched' geometry, most distorted from the regular tetrahedron ($\angle$ECH = 111.46°, $\angle$HCH = 107.41°) was found for $SiMe_4$ with the shortest C–E bond length; Table 3 and S2 (ESI†). In contrast, a fully relaxed, practically idealized tetrahedral coordination was computed for the methyl carbons in $PbMe_4$ possessing the longest C–E distance. Hence, the greatest $J_{0,CH}$ coupling in this compound is attributable in part to a relatively small difference in χ_gs of its carbon substituents (H, $PbMe_3$) and to the greatest *s* character of the C–H bond among all tetramethyls under study.

In view of the foregoing facts, the 'abnormal' couplings $^1J_{CH}$ and $^2J_{HH}$ in CMe_4, discussed here *via* related outlying data points, are attributable to the shortest C–E distance and greatest electronegativity of its E atom (or -EMe_3 group) which characterize this molecule. On the contrary, there is a monotonic increase in the C–E distance for all four remaining tetramethyls (E = Si to Pb), associated with a diminished internal steric congestion around the E atom, and an increase in pertinent values of χ_a and χ_g on going down a Group 14/IVA. As a result, their $^1J_{CH}$ couplings follow the observed Si < Ge < Sn < Pb sequence. The case of the system CMe_4 is significantly different in this respect and that is the most probable origin of the aforesaid 'anomalies' in its molecular NMR response properties.

5 Conclusions

This investigation affords the experimental one-bond C–H *J*-couplings in tetramethyl species EMe_4 (E = C to Pb) in the vapor and liquid states. Such gas-phase $J_{0,CH}$s, obtained as *J* values extrapolated to a zero-pressure limit, are only slightly different from the J_{CH}s measured for liquids. The former data show linear dependences on density of the gaseous solvents used. The present study also demonstrates that current DFT protocols allow for a successful prediction of the experimental NMR *J*-couplings and other molecular properties as well as for their interpretation in terms of the electronic and geometric structure. From the computational results on title compounds it is concluded that: (1) Reliable geometries of all these molecules can be obtained applying the B3LYP functional in the LDBS approach that consists of the use of a def2-TZVPPD basis set for a central E atom (ECPs for Sn and Pb) and 6-31G(*d*,*p*) for the

ligand (methyl group) atoms; this procedure can be also proposed for the other structurally similar systems. (2) Subsequent LDBS calculations of J_{CH}s, carried out with an IGLO-II basis sets for the C and H atoms and def2-QZVPPD for the E atoms (protocol II), afforded *J* values being in good agreement with the experiment. (3) The use of pcJ-2 instead of IGLO-II gives a similar or worse result, but at the expense of computation time. (4) All four Ramsey contributions to J_{CH}s in compounds EMe_4 were computed with the Fermi-contact term being dominant participant. (5) Strong linear correlations were shown between the SD terms, $J_{0,CH}$s, and J_{HH}s *vs.* recently reported Mulliken electronegativities (χs) of the E atoms and EMe_3 groups, excluding the CMe_4 case. (6) These χ values were positively verified, for the first time – however, a small correction of χ_a (Ge) and χ_g for -$GeMe_3$ is proposed. (7) Predicted alterations in the C–E distance and bond angles around the methyl carbons in species EMe_4 in conjunction with the χ_gs of their substituents and Bent's rule permitted an explanation of the trends observed in experimental $^1J_{CH}$s and $^2J_{HH}$s in terms of gradual changes in the *s* character of carbon sp^n-hybridized orbitals forming the C–H bonds. (8) 'Abnormal' molecular NMR properties of CMe_4 discussed above appear to result from the shortest C–E bond length and the greatest electronegativity of its E atom and EMe_3 group among all tetramethyls under study.

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Tables

Table 1. Experimental NMR $^1J_{CH}$ data for gaseous mixtures, liquids, and solutions of the species EMe_4

Solute gas (A)	Solvent gas (B)	$J_{0,CH}$ [Hz]	$J_{AB,CH}$[a] [Hz mL mol^{-1}]	$J_{0,CH,av}$[b] [Hz]	J_{CH}(neat)[c] [Hz]	ΔJ_{CH}[c,d] [Hz]	J_{CH} for neat,[e] solution in C_6H_6[f] or CCl_4,[g] [Hz]
CMe_4	Kr	123.95(5)	23(59)	123.93(4)	124.06(1)[h]	0.13(5)	124.5(1),[e] 124.3,[f] 124.0(2)[g]
	Xe	123.92(6)	163(63)				
$SiMe_4$	Kr	117.88(5)	52(67)	117.88(4)	118.15(2)	0.27(6)	118.2(1),[e] 118.2,[f] 117.8(2)[g]
	Xe	117.88(7)	99(83)				
$GeMe_4$	Kr	124.05(2)	76(30)	124.05(1)	124.40(1)	0.35(2)	124.0(4),[e] 124.4,[f] 124.6(2)[g]
	Xe	124.05(1)	53(21)				
$SnMe_4$[i]	CO_2	127.12(8)	32(10)	127.13(5)	127.62(2)	0.49(7)	127.2(1),[e] 127.7,[f] 127.8(2)[g]
	N_2O	127.13(7)	64(10)				
$PbMe_4$	SF_6	133.28(4)	314(42)	133.29(3)	134.07(2)[j]	0.78(5)[j]	134.0(2),[e] 134.2,[f] 134.3(2)[g]
	Xe	133.31(5)	205(52)				

[a] Density dependence of J(CH); see Eq. 2. [b] Weighted averages. [c] This work. [d] Differences [$J(CH)_{neat} - J_0(CH)_{av}$].[e] Ref. [53b]. [f] Ref. [54]. [g] Ref. [55]. [h] Measured for a liquid at vapor pressure of ~2 atm. [i] Ref. [8]. [j] For 65 wt.% solution in xylene.

Table 2. $^1J_{CH}$ couplings for the species EMe_4 computed[a] in vacuum and in CCl_4 solution, with their decomposition into all four Ramsey terms, [Hz]

Compd	Medium	FC	SD	PSO	DSO	Total J^{theor}
CMe_4	gas	119.70	0.18	0.92	0.73	121.52
	CCl_4	119.67	0.18	0.92	0.73	121.50
$SiMe_4$	gas	114.25	0.29	1.46	0.68	116.68
	CCl_4	114.24	0.29	1.46	0.68	116.67
$GeMe_4$	gas	120.70	0.33	1.13	0.98	123.13
	CCl_4	120.67	0.33	1.13	0.98	123.10
$SnMe_4$	gas	124.15	0.39	1.21	0.63	126.38
	CCl_4	124.08	0.39	1.22	0.63	126.31
$PbMe_4$	gas	130.51	0.44	1.10	0.57	132.62
	CCl_4	130.41	0.44	1.11	0.57	132.53

[a] Protocol II, and II-CCl_4 was applied, respectively.

Table 3. Selected computed[a] or experimental structural, spectroscopic[b] and Mulliken electronegativity data[c] for all species EMe_4 and their constituent units

Compd	∠ECH (gas), [°]	∠HCH (gas), [°]	∠HCH (CCl_4), [°]	$^2J_{HH}^{theor}$ (CCl_4), [Hz]	$^2J_{HH}^{exp}$ (CCl_4), [Hz] [b]	χ_a for the E atom, [eV] [c,d,e]	χ_g for the EMe_3 group, [eV] [c,e]
CMe_4	111.07	107.83	107.85	-11.93	-12.56	6.73	4.15
SiMe_4	111.46	107.41	107.46	-13.47	-14.05	4.96	3.80
GeMe_4	110.86	108.05	108.09	-12.36	-12.96	4.71 (~4.54)	3.85 (~3.88)
SnMe_4	110.51	108.41	108.47	-12.12	-12.37	4.31	3.95
PbMe_4	109.48	109.46	109.52	-10.68	-10.94	3.85	4.04

[a] Protocol II, and II-CCl_4 was applied, respectively. [b] Ref. [55]. [c] Ref. [68]. [d] χ_a for the H atom = 7.26 eV, ref. [68].
[e] The proposed corrected value is in parenthesis, see the text.

Figures

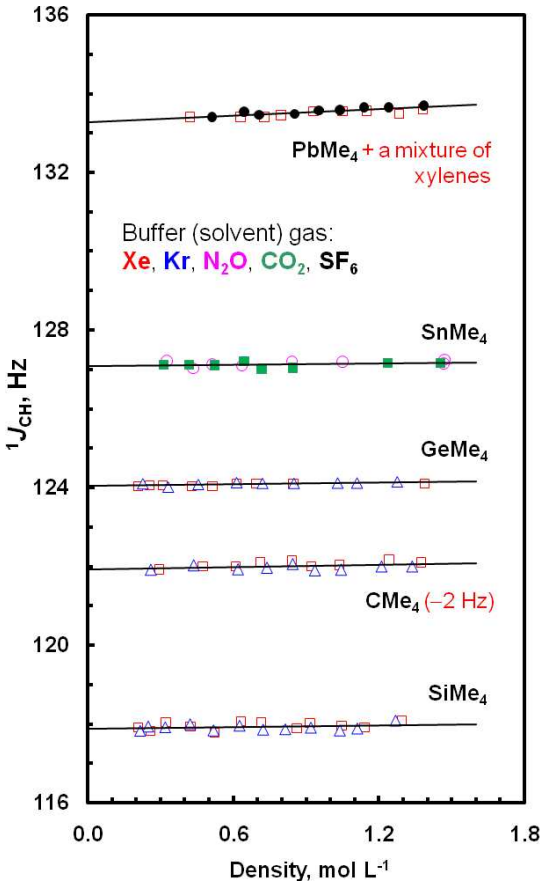


Fig. 1. The dependence of the $^1J_{CH}$ data in compounds EMe_4 on the density of solvent gases at 300 K (□-Xe, Δ-Kr, ○-N_2O, ■-CO_2, ●-SF_6). The data points for CMe_4 are shown with the (–2 Hz) vertical shift, for clarify.

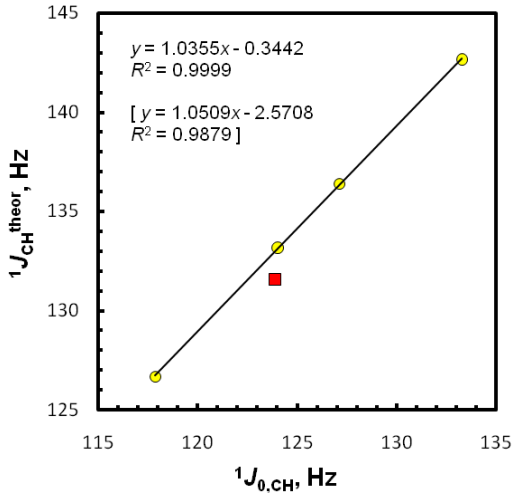


Fig. 2. Correlation between (+5 Hz corrected) $^1J_{CH}$ computed (protocol II) and extrapolated $^1J_{0,CH}$ values. Statistics for the five-point plot with the CMe_4 data (■) are in the bracket.

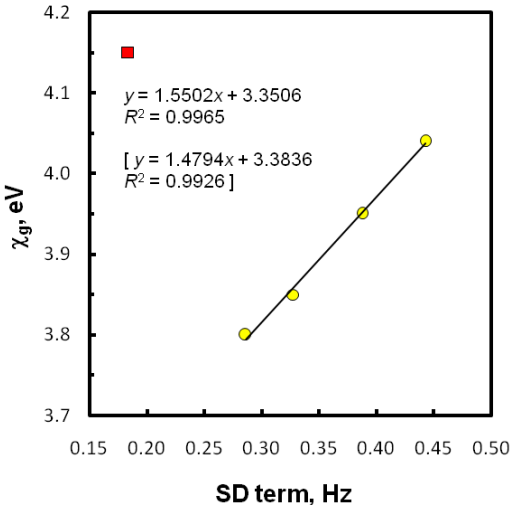


Fig. 3. The group electronegativity *vs*. SD term (protocol II) values; the CMe_4 point (■) was omitted. Statistics with the χ_g corrected for $GeMe_4$ (3.88 eV) are in the bracket.

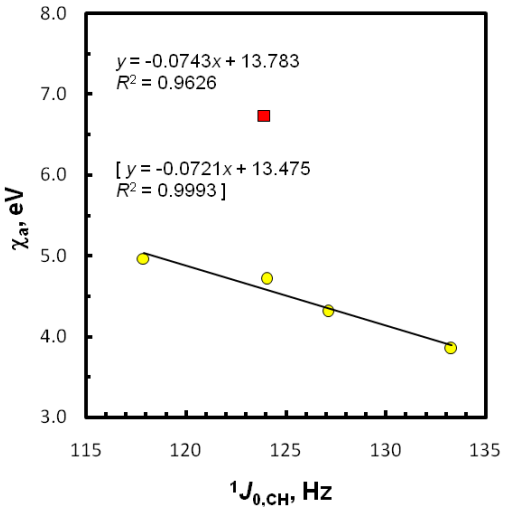
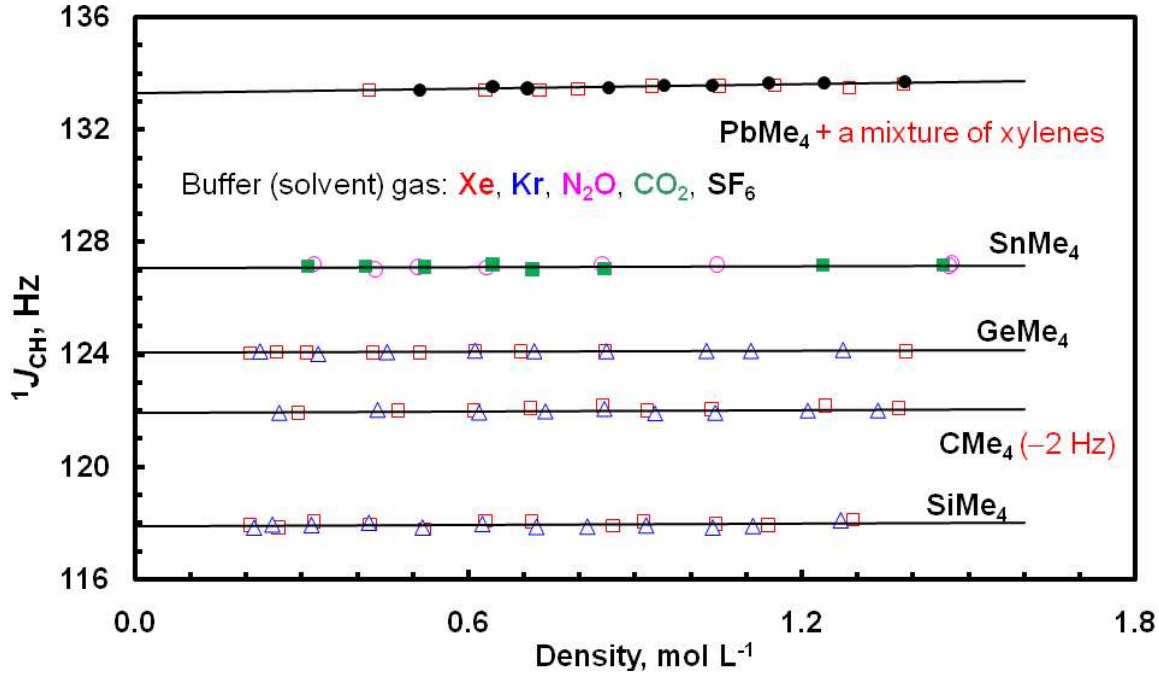


Fig. 4. Correlation between the atom electronegativity and $^1J_{0,CH}$ data; the CMe_4 point (■) was omitted. Statistics with the χ_a corrected for Ge (4.54 eV) are in the bracket.

A graphical and textual abstract

The gas-phase $^{1}J_{0,CH}$s in 'isolated' molecules of EMe_4 were determined and discussed in terms of their geometric/electronic properties obtained from the DFT calculations.



Small version (40 x 80 mm) – the EXCEL file