the factor
\[ \frac{\omega + \omega_0 + k |V_A|^2}{\omega - \omega_0} \]
\[ \frac{k |V_A|^2}{2 \omega^2} \]
has an extreme value.

Then we obtain a possible maximum width of the potential well
\[ \Delta x \approx \frac{1}{8} \left( \frac{L_A}{r} \right)^2 \frac{a^2}{r} \].
If \( \Delta x < a \equiv r |V_A|/\Delta x \), that is, \( \Delta x < a \equiv a \beta / \Delta x \), the unstable localized solution cannot exist. Then we have a stability criterion
\[ L_A / r < 2(\alpha / \alpha)^{1/2} \]. \hspace{1cm} (A.39)

Mikhailovskaya and Mikhailovskii also obtained another stability criterion
\[ L_A / r < (r / a)^{1/2} \beta ^{-1/6} \]. \hspace{1cm} (A.40)

Beyond the turning point \( x = x_A \), the localized solution is expressed by use of the Airy function \[ \psi(x) \approx \left( \frac{a^2}{U'(x_A)} \right)^{1/6} (x_A - x)^{-1/6} \]
\[ \times \exp \left\{ - \frac{2}{3} \left( \frac{U'(x_A)}{a^2} \right)^{1/3} (x_A - x)^{2/3} \right\} \],
where \( U'(x_A) \) is roughly given by \( \Delta x / r x_A \), that is, \[ U'(x_A) \approx \frac{1}{8} \frac{L_A a}{r^3 \beta} \].

Then the characteristic length of exponential decay which we denote by \( \delta x \) is given by
\[ \delta x \sim 2(a r^3 \sqrt{\beta / L_A})^{1/4} \].

Finally, we can derive the stability criterion Eq. (A.40) from the condition \( \delta x > x_A - x_e \) where \( x_e \) is defined by \( \omega_0 \equiv |V| / \omega_0 \).

Galeev also obtained almost the same criterion as Eq. (A.40). He mentioned that if the transparency condition of a hill between \( x_A \) and \( x_e \) is satisfied, that is, if
\[ \int_U(x) \, dx < 1 \], \hspace{1cm} (A.42)
the region of existence of the localized solution expands into the ion \( \text{Landau} \) damping region. The transparency condition Eq. (A.42) is essentially the same as \( \delta x > x_A - x_e \). However, his statement is not right because the ion \( \text{Landau} \) damping region does not lie in a region \( x < x_e \), but in a region \( x \approx x_i \),
where \( x_i \) is defined \( k |v| = \omega_0 \). Eq. (A.40) means that the localized region of the perturbation extends into the region of the electron \( \text{Landau} \) damping.

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A General Formulation for the Evaluation of Two-center Moment Integral – An Extension

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(Z. Naturforschg. 22 a, 1701—1704 [1967]; received 27 June 1967)

In two recent studies \(^1\) \(^2\) we presented a general formulation for the evaluation of two-center moment integrals as well as the rotational properties of s, p, and d atomic orbitals needed in the quantum mechanical calculations of one-electron properties. In the present investigation, we extend the scope of our previous formulation by presenting a series of tabulations useful both to inorganic and organic chemists.

(a) One- and Two-center Transition Moment Integrals

Our most recent investigation of the intensities in the spectrum of XeF\(4\) \(^3\) has revealed certain useful internal relationships existant in one- and two-center transition moments. These are conveniently reported in Tables 1 and 2. It should be noticed that, in the case of the two-center integrals,
the bra functions are defined in a right-handed coordinate system, while the ket functions are defined in a left-handed coordinate system. Obviously, if both functions are defined in right-handed coordinate systems, \( M_2 \) as defined in our original paper \(^1\) becomes

\[
M_2' = (-1)^{1+m} M_2,
\]

while if both are defined in left-handed coordinate systems, it becomes

\[
M_2'' = (-1)^{r+m'} M_2.
\]

(b) Rotational Transformations of \( f \) Orbitals for the Evaluation of Two-center Integrals

In a recent note we commented on the rotational transformations of \( s, p, \) and \( d \) orbitals necessary for the evaluation of two-center integrals \(^2\). In the present note we extend our previous formulation to the \( f \) orbitals. It should be noted that although the \( f \) orbitals do not play as important a role in bonding as do the \( d \) orbitals, they do, nonetheless, account for the subtle aspects of the chemistry of the elements with atomic numbers 51 to 70 and 83 to 102. The \( 5f \) orbitals, for instance, are believed to be responsible for the stability of \( \text{MO}_2^{2+} \) and \( \text{MO}_2^{+} \) ions \(^4\) of some of the actinides \((M = \text{U}, \text{Np} \text{ or Pu})\), while the \( 4f \) orbitals, it has been suggested \(^5-8\), even contribute to the stability of the compounds of a few elements lighter than the lanthanides.

Thus, we define \( \xi, \eta, \alpha, \beta, \) and \( \gamma \) precisely in the same manner as in our previous note \(^2\), and if for ease of writing we set

\[
\begin{align*}
\delta_{+} & = \delta_{+} (x^a - 3 y^c), \\
\delta_{B} & = \delta_{B} (x^a - 3 y^c), \\
\delta_{C} & = \delta_{C} (5 x^a - x^c), \\
\delta_{D} & = \delta_{D} (5 x^a - 3 r^c), \\
\delta_{E} & = \delta_{E} (5 x^a - 3 r^c), \\
\delta_{F} & = \delta_{F} (y^a - y^c), \\
\delta_{G} & = \delta_{G} (y^a - y^c), \\
\end{align*}
\]

then for the \( f \) orbitals \(^9, 10\) we have

\[
\begin{align*}
\delta_{ac} & = \delta_{ac} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right) \\
\delta_{Bo} & = \delta_{Bo} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right) \\
\delta_{Co} & = \delta_{Co} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right) \\
\delta_{Do} & = \delta_{Do} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right) \\
\delta_{Fo} & = \delta_{Fo} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right) \\
\delta_{Go} & = \delta_{Go} \left( 3 \alpha + 3 \gamma \right) - \eta^6 \cos \left( 3 \alpha - 3 \gamma \right)
\end{align*}
\]

\(^5\) Z. Z. Hugus, J. Am. Chem. Soc. 74, 1076 [1952].
\[ f_{z(x^2-y^2)} = \left\{ -V/6(f^2 \eta \cos(3 \alpha + 2 \gamma) + \xi \eta^2 \cos(3 \alpha - 2 \gamma)) \right\} f_{A_0}^2 + \left\{ (f^2 - 5 \xi \eta) \sin(2 \alpha + 2 \gamma) \right\} f_{B_0} + \left\{ (f^2 - 5 \xi \eta) \sin(2 \alpha - 2 \gamma) \right\} f_{E_0} + \left\{ (f^2 - 5 \xi \eta \cos(2 \alpha + 2 \gamma) \right\} f_{D_0} + \left\{ (f^2 - 5 \xi \eta \cos(2 \alpha - 2 \gamma) \right\} f_{E_0} \]

\[ f_{x^2} = \left\{ -V/8(f^4 \eta^2 \cos(3 \alpha + 2 \gamma) + f^4 \eta^2 \cos(3 \alpha - 2 \gamma)) \right\} f_{A_0} + \left\{ -V/8(f^4 \eta^2 \cos(3 \alpha + 2 \gamma) + f^4 \eta^2 \cos(3 \alpha - 2 \gamma)) \right\} f_{B_0} + \left\{ f^4 \eta^2 \cos(2 \alpha + 2 \gamma) \right\} f_{C_0} + \left\{ f^4 \eta^2 \cos(2 \alpha - 2 \gamma) \right\} f_{D_0} + \left\{ f^4 \eta^2 \cos(2 \alpha + 2 \gamma) \right\} f_{E_0} + \left\{ f^4 \eta^2 \cos(2 \alpha - 2 \gamma) \right\} f_{F_0} \]

\[ f_{y^2} = \left\{ -V/8(f^4 \eta^2 \cos(3 \alpha + 2 \gamma) + f^4 \eta^2 \cos(3 \alpha - 2 \gamma)) \right\} f_{A_0} + \left\{ -V/8(f^4 \eta^2 \cos(3 \alpha + 2 \gamma) + f^4 \eta^2 \cos(3 \alpha - 2 \gamma)) \right\} f_{B_0} + \left\{ f^4 \eta^2 \cos(2 \alpha + 2 \gamma) \right\} f_{C_0} + \left\{ f^4 \eta^2 \cos(2 \alpha - 2 \gamma) \right\} f_{D_0} + \left\{ f^4 \eta^2 \cos(2 \alpha + 2 \gamma) \right\} f_{E_0} + \left\{ f^4 \eta^2 \cos(2 \alpha - 2 \gamma) \right\} f_{F_0} \]

\[ f_{x^2-y^2} = \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{A_0} + \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{B_0} + \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{C_0} + \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{D_0} + \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{E_0} + \left\{ f^2 \eta \cos(3 \alpha + 2 \gamma) - f^2 \eta \cos(3 \alpha - 2 \gamma) \right\} f_{F_0} \]

\[(c) \text{ Carbon-Carbon Overlaps and Transition Moment Integrals} \]

In the simple LCAO-MO Hückel formalism, orbitals defined on different centers are assumed to be orthonormal, while, on the other hand, in higher order approximations overlaps between non-adjacent carbon π orbitals are indeed included. It can be shown that the overlap \( S_{ij} \) for a pair of homonuclear Slater 2p orbital \( \xi \) and \( \eta \) is given by

\[ S_{ij} = e^{-\rho^2}(1 + (\gamma/2) + (\gamma/10) + (\gamma/120)), \]

where \( \rho \) is the effective nuclear charge (for carbon \( Z = 3.25 \)) and \( R \) is the internuclear distance. \( \gamma = Z R / a_0 \)

nuclear distance in Angstroms (Å). The broken curve of Fig. 1 gives the variation of \( S_{C-C} \) with the C—C distance ranging from the distance of C≡C in acetylene (1.20 Å) to the one of C—C in ethane (1.54 Å). The solid curves, on the other hand, give the overlaps (\( \gamma = 0 \)) as well as the transition moment integrals (\( \gamma = 1 \)) for a pair of carbon SCF atomic functions given by Clementi. The radial function used for carbon is the ground state's (\(^3\)P) and is given by

\[
R(2p) = 0.5777 \varphi_2(1.4209) + 0.23561 \varphi_2(2.5873) + 0.24762 \varphi_2(0.9554) + 0.01090 \varphi_2(6.3438)
\]

Fig. 1. \( \langle \pi' | Z | \pi \rangle \) values of carbon as a function of inter-nuclear distance in Å.

---

![Flowchart](image)

**Fig. 2.** (see Appendix, p. 1704).
where \( \psi_n(\xi) = N_n \frac{r^{-1}}{\pi} e^{-\xi r}, \) \( N_n \) being the normalization constant. Table 3, on the other hand, gives the values of these integrals as a function of their internuclear distance.

### Appendix

In a recent study\(^1\) we reported a general formulation for the evaluation of two-center moment integrals, and presented therein a computer program for the coding of the said formulation. Unfortunately, however, for some unexplainable reason, the part that dealt with the coding of the one-center integrals, the right-hand side of the flow chart of Fig. 2, seems to be totally erroneous. Consequently, we would like herewith to present the correct version in Fig. 2 (see p. 1703).

<table>
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<th>Distance (Å)</th>
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<th>( \gamma = 1 )</th>
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<td>0.3486</td>
</tr>
<tr>
<td>1.8</td>
<td>0.1897</td>
<td>0.3228</td>
</tr>
<tr>
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<td>0.1656</td>
<td>0.2975</td>
</tr>
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<tr>
<td>3.3</td>
<td>0.0138</td>
<td>0.0472</td>
</tr>
</tbody>
</table>

\(^{12}\) E. Clementi, IBM J. Res. Develop. 9, 2 [1965].

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### Wie genau kann man heute in der EPR-Spektroskopie g-Faktoren messen?

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In the past few years theoretical aspects of the \( g \)-factors of aromatic hydrocarbon radicals have been developed. Since the \( g \)-factor variation of these radicals is only of the order of \( 10^{-4} \) precise measurements of these quantities are clearly called for. Absolute \( g \)-factor measurements are performed according to the equation \( g = C \omega / \omega_p \). A survey of the different determinations of the conversion factor \( C = 2 \gamma_p m/c/e \) is given and a value of \( C = 3.0419980 \times 10^{-3} \pm 0.5 \) ppm is proposed for a well defined proton NMR probe. The most important effects are discussed which may shift the measured EPR and NMR frequencies \( \omega \) and \( \omega_p \).

In der paramagnetischen Elektronenresonanz(EPR)-Spektroskopie der Übergangselemente wird bereits seit geraumer Zeit der \( g \)-Faktor der Elektronenhülle theoretisch und experimentell ausführlich untersucht\(^1\). Im Gegensatz zu den Ionen der Übergangsmetalle liegen bei den aromatischen Kohlenwasserstoff-Radikalen die Variationen des \( g \)-Faktors nur in der Größenordnung von \( 10^{-4} \) (100 ppm). Zur Untersuchung von Gesetzmäßigkeiten ist also eine beträchtliche Meßgenauigkeit erforderlich. Da es nunmehr – zumindest im Prinzip – möglich geworden ist, die \( g \)-Faktoren selbst sehr komplizierter organischer Radikale theoretisch vorherzusagen\(^2\), sind auch vom experimentellen Standpunkt aus diese Größen inter-
