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INFRARED SENSITIVITY OF DIATOMIC GASES

FOR

THE ANHARMONIC VIBRATING ROTATOR MODEL

W. Malkmus
A. Thomson

May 5, 1961

This work was conducted under Contract
No. AF19(604)-8584, ARPA Order No. 116,
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CONVAIR SAN DIEGO
Convair Division of General Dynamics Corporation
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INFRARED EMISSIVITY OF DIATOMIC CASES FOR THE ANHARMONIC VIBRATING ROTATOR MODEL

W. Malkmus
A. Thomson

ABSTRACT

In order to compute the emissivity of a diatomic gas, a simplified model of a diatomic molecule is assumed: an anharmonic oscillator with the first approximation to the vibration-rotation interaction. For a given band, the frequency of emitted radiation is expressed as a quadratic function of the quantum number \( m \), which is solved to express \( m \) as a function of \( \omega \). This expression is substituted for \( m \) in the equations used for computing the average line intensity and average line spacing. By applying the random Elsasser model to the fundamental and superposed higher order bands, closed-form solutions are obtained for the emissivity as a function of \( \omega \) for certain limiting cases. Experimental data, where available, are used for the line widths and total absorption of the bands. The harmonic oscillator approximation is used to estimate strengths of higher order bands for which experimental data are not available. This analysis is applied to HCl, HF, CO, and NO for temperatures ranging from 300 to 7000°K.

I. INTRODUCTION

A first approximation to the infrared spectral emissivity of a diatomic gas can be made by assuming a harmonic oscillator model for the molecule.\(^6,9\) This has the advantage of simplicity, but the very nearly symmetric intensity distribution which it yields does not closely match the asymmetric shape which is found experimentally or calculated by a line-by-line method.\(^1\)
To improve the approximation, the model is assumed to be an anharmonic oscillator with the first approximation to the vibration-rotation interaction. This will improve the approximation to the actual spectral emissivity without resort to the more detailed approach of considering the emission from each individual spectral line.

II. DEVELOPMENT

The energy levels of an anharmonic oscillator, in the first approximation to the vibration-rotation interaction, are given by

\[ E(v, J) = \omega_v \left( v + \frac{1}{2} \right) - \omega_e x_e \left( v + \frac{1}{2} \right)^2 + \omega_e y_e \left( v + \frac{1}{2} \right)^3 + \omega_e z_e \left( v + \frac{1}{2} \right)^4 + B_e J(J + 1) - \alpha_e \left( v + \frac{1}{2} \right) J(J + 1). \]  

(1)

(Refer to the treatment of Stull and Plass\(^1\) for notation and basic equations.) The frequency of a transition for \( |\Delta v| = 1 \) is then given by

\[ \omega = E(v + 1, J') - E(v, J) \]

\[ = \omega_v + B_e [J'(J' + 1) - J(J + 1)] \]

\[ - \alpha_e \left( (v + \frac{3}{2}) J'(J' + 1) - (v + \frac{1}{2}) J(J + 1) \right), \]  

(2)

where

\[ \omega_v = \omega_v - 2(v+1) \omega_e x_e + [3(v+1)^2 + \frac{1}{4}] \omega_e y_e + [4(v+1)^3 + (v+1)] \omega_e z_e. \]  

(3)

In terms of \( m = J + 1 \) for the R-branch, \( m = -J \) for the P-branch,

\[ \omega = \omega_v + 2B_e m - \alpha_e [m(m + 1) + 2(v + \frac{1}{2}) m]. \]  

(4)

This equation expresses \( \omega \) as a quadratic function of \( m \). On solving for \( m \), this becomes

\[ m = \frac{B_e - \alpha_e (v + 1) + \sqrt{[B_e - \alpha_e (v + 1)]^2 - \alpha_e (\omega - \omega_v)}}{\alpha_e}. \]  

(5)
Note that Equation (4) implies a maximum value for \( \omega \) of
\[
\omega_{\text{max}} = \omega_v + \frac{[B_e - \alpha_e (v + 1)]^2}{\alpha_e}
\]
for
\[
m = \frac{B_e - \alpha_e (v + 1)}{\alpha_e}.
\]

Since \([B_e - \alpha_e (v + 1)]/\alpha_e > 0\) for all molecules under consideration, this model will always have a band head in the R-branch. Hence, the upper sign before the radical in Equation (5) refers to the main portion of the band and the lower sign to the returning R-branch. (This convention for the \(\tau\) or \(\pm\) sign will be used throughout this analysis.)

The average line spacing in the region of a given \( m \) is
\[
d = 2 |B_e - \alpha_e (v + 1 + m)|.
\]
This becomes [expressed in terms of \( \omega \) by use of Equation (5)],
\[
d(\omega) = 2 \sqrt{[B_e - \alpha_e (v + 1)]^2 - \alpha_e (\omega - \omega_v)}
\]
for the entire band, including the returning R-branch.

The energy term \( E_v(j) \) [Equation (6), Stull and Plass\(^1\)] in terms of the transition frequency \( \omega \), is given by
\[
E_v(j) = \frac{B_e - (v + 1) \alpha_e}{\alpha_e^2} \left\{ 2 \left[ B_e - (v + 1) \alpha_e \right] \left[ B_e - (v + 1) \alpha_e \right] \right. \\
\left. \tau \sqrt{\left[ B_e - (v + 1) \alpha_e \right]^2 - \alpha_e (\omega - \omega_v)} \right] \\
\left. - \left( 1 + \frac{\gamma \alpha_e}{B_e - (v + 1) \alpha_e} \right) \alpha_e (\omega - \omega_v) \right\}.
\]

Several approximations are used in the analysis. The factor \( \omega_v' \), defined by Equation (5)\(^1\) is approximated by
\[
\omega_v' = \omega_v \left[ 1 - \exp \left( - \frac{\hbar \omega_v}{kT} \right) \right].
\]
The Herman and Wallis factor $F_{v,j}^{v',j'}$ is approximated by

$$F = 1 + c \left[ \frac{B_e - \alpha_e (v + 1) \mp \sqrt{[B_e - \alpha_e (v + 1)]^2 - \alpha_e (\omega - \omega_v)}}{\alpha_e} \right]$$

(12)

where

$$c = \frac{-8 B_e \theta}{\omega_e}$$

(13)

($\theta$ is the ratio of the zero\textsuperscript{th} to the first order term in the expansion of the electric dipole moment). The partition functions $G_R(T)$ and $G(T)$ are approximated by

$$G_R(T) = \frac{kT}{B_e h c}$$

(14)

and

$$G(T) = \frac{kT}{B_e h c} \frac{\exp \left[ -E(0,0) \frac{h c}{kT} \right]}{1 - \exp \left( -\frac{h c \omega_v}{kT} \right)}.$$  

(15)

Stull and Plass\textsuperscript{1} give the following equation for the integrated absorption of a vibration-rotation line:

$$S_{v,j}^{v',j'} = \frac{\alpha_{\nu}^{v'} \omega_{\nu} \omega_{\nu}^{v'}}{G_R(v,T) \omega_{\nu}^{v'}} \exp \left[ -E_v(J) \frac{h c}{kT} \right] F_{v,j}^{v',j'} \times$$

$$\left[ J \delta_{j-1,j'} + (J + 1) \delta_{j+1,j'} \right] \left[ 1 - \exp \left( -\frac{\omega_{\nu}^{v'} \omega_{\nu} \omega_{\nu}^{v'}}{kT} \right) \right].$$

(16)

Since

$$J \delta_{j-1,j'} + (J + 1) \delta_{j+1,j'} = |m|,$$

8
and by use of the preceding approximations, this becomes

\[ S_{v+1}(\tau)(\omega) = \frac{\alpha_{v+1}^e B e^{\frac{\hbar c}{\nu_{v+1}}} \omega}{\nu_{v+1}^e k T} \times \]

\[ \exp \left[ -1.439 \frac{[B_e - \alpha_e(v+1)]}{\alpha_e^2 T} \left\{ \frac{2[B_e - \alpha_e(v+1)]}{2[B_e - \alpha_e(v+1)]} \right\} \right] \times \]

\[ \frac{\nu_{v+1}^e}{\nu_{v+1}^e} \left[ \frac{B_e - \alpha_e(v+1)}{\nu_{v+1}^e} \right] - \left( 1 + \frac{\alpha_e}{B_e - \alpha_e(v+1)} \right) \alpha_e(\omega - \nu_v) \right] \times \]

\[ \left( 1 + \frac{\alpha_e}{B_e - \alpha_e(v+1)} \right) \alpha_e(\omega - \nu_v) \right) \right] \times \]

\[ \left( 1 + \frac{\alpha_e}{B_e - \alpha_e(v+1)} \right) \alpha_e(\omega - \nu_v) \right) \right) \right] \times \]

(17)

Experimental data are used, where available, for the total band absorption \( \alpha_{v+1}^e \). A convenient approximation is made to Stull and Plass' Equation (8); namely,

\[ \frac{\alpha_{v+1}^e(T)}{\alpha_{v+1}^e(T_0)} = \frac{T_0}{T} \frac{\omega_0^e(T)}{\omega_0^e(T_0)} \frac{\omega_{v+1}^e(T)}{\omega_{v+1}^e(T_0)} \exp \left[ \left( E(v) - E(0, 0) \right) \frac{hc}{kT} \left( \frac{1}{T_0} - \frac{1}{T} \right) \right] \]

(18)

so that \( \alpha_{v+1}^e \) can be computed for any temperature if it is known for one temperature.

If data are not available for higher-order bands, by assuming the harmonic oscillator approximation and using Stull and Plass' Equations (4) and (9), one obtains

\[ \frac{\alpha_{v+1}^e(T)}{\alpha_{v+1}^e(T)} = \frac{\nu_v^e + \frac{1}{2}}{\nu_v^e + 1} \exp \left[ - \left( E(v^e) - E(v) \right) \frac{hc}{kT} \right] \]

(19)

In the weak line approximation, the emissivity is given by

\[ \epsilon = 1 - \exp \left[ - \frac{\hbar c}{p S(\omega)} \sqrt{g(\omega)} \right] \]

(20)

where

\[ S(\omega) = \sum_v \left[ S_v^{+1}(-)(\omega) + S_v^{+1}(+)(\omega) \right] \]

(21)
An additional approximation is made in the strong line approximation:

$$\Pi_1 \left[ 1 - \text{erf} \left( x_1 \right) \right] = \exp \left( 2\pi^{-\frac{1}{2}} \sum_1 x_1 \right).$$

In general, this approximation causes a change in the emissivity of less than 5%. (It gives the same result as for the statistical model.)

In the strong line approximation,

$$\epsilon_\omega = 1 - \exp \left[ -2\alpha_0 \frac{1}{4} (p^2 + \frac{\sigma}{\omega}) \sum_1 \omega \right],$$

where

$$\sum_1 \omega = \sum_v \left[ \left( S_v^{+1} - \omega \right)^2 + \left( S_v^{-1} + \omega \right)^2 \right],$$

and $\alpha_0$ is the average half-width of the line at 1-atm pressure.

A detailed discussion of the range of validity of the various approximations is given by Plass. For example, Plass Figure 2 indicates that for pure CO, when $\epsilon_\omega \approx 0.6$, the weak line approximation is accurate within 10% when $\beta > 3$. (This requires the pressure to be above some minimum value which is largest ($p \approx 50$ atm) when $T \approx 1200^\circ K$.) The strong line approximation is accurate within 10% when $0 < 1$. (For $T \approx 1200^\circ K$ this implies a maximum $p \approx 17$ atm.) A lower limit to the validity of the strong line approximation is imposed by the presence of Doppler broadening, which has been neglected. The Doppler and Lorentz widths are comparable for $p \approx 1/2$ atm at $T \approx 1200^\circ K$.

For pure HF, the strong line approximation may be used for pressures up to a certain value (a maximum $p \approx 35$ atm for $T \approx 3000^\circ K$) and above a minimum determined by the Doppler broadening. (The Doppler and Lorentz widths are comparable for $p \approx 0.4$ atm and $T \approx 3000^\circ K$.)
III. DISCUSSION

Results of this analysis are presented (Figures 1 to 32) for HCl, HF, CO, and NO for temperatures ranging from 300 to 7000°K. The presence of isotopes other than Cl$^{35}$ and Cl$^{37}$ was ignored. Emission from as many as 20 bands was considered.

Data of Benedict, et al.,$^3$ for HCl were used to make a direct comparison with the more elaborate calculations of Stull and Plass.$^1$ Their value of 130 ± 7 cm$^{-2}$ atm$^{-1}$ for the fundamental band strength is probably too low (cf. Babrov, et al.,$^4$ 150 ± 5 cm$^{-2}$ atm$^{-1}$; Weber and Penner,$^5$ 150 cm$^{-2}$ atm$^{-1}$).

Benedict and Flyler's$^6$ value of 260 cm$^{-2}$ atm$^{-1}$ was used for the CO fundamental band strength (cf. Weber and Penner,$^5$ 237 cm$^{-2}$ atm$^{-1}$). Kuipers' data$^7$ were used for HF, and Weber and Penner's$^5$ for NO.

Average line widths were computed as described by Stull and Plass.$^1$ It should be noted that the supplementary computation made by these authors for emission from line wings in the frequency range beyond a band head is omitted here.

From 20 to 50 points were computed for each curve. Each set of four curves represents a charge of between 0.01 and 0.02 hour's time on an IBM 704 digital computer.

A comparison is made (Figures 33 and 34) with other published computations of diatomic gas emissivities. Figure 33 compares Equation (20) to the results of one particular computation by Stull and Plass$^1$ (for HCl at 2400°K in the weak line approximation for p$l$ = 100 atm-cm) in which they considered the emission from approximately 2900 individual spectral lines. Penner and Gray$^8$ recently published a comparative computation based on a simple harmonic oscillator model, which is also shown in Figure 33.

The curve representing Equation (20) follows quite closely the curve computed by Stull and Plass,$^1$ which is presumably the most accurate available theoretical computation. The agreement would be slightly closer
Figure 1 Spectral Emissivity of CO AT T = 300°C

(Wave Number, cm⁻¹ vs. Wave Number, cm⁻¹)
FIGURE 2 SPECTRAL EMISSIVITY OF CO AT T = 600°K
(WEAK LINE APPROXIMATION)
FIGURE 3
SPECTRAL EMISSIVITY OF CO AT T = 1200°K
(WEAK LINE APPROXIMATION)
Figure 4: Spectral Emissivity of CO at T = 1800 K
FIGURE 5
SPECTRAL EMISSIVITY OF CO AT T = 2400°K
(WEAK LINE APPROXIMATION)
Figure 6: Spectral Emissivity of CO at T = 3000°K (Weak Line Approximation)
FIGURE 7  SPECTRAL EMISSIVITY OF CO AT T = 5000 K

WEAK LINE APPROXIMATION

WAVE NUMBER, cm⁻¹

PI = 10² ATM - CM

1

10
FIGURE 9
SPECTRAL EMISSIVITY OF NO AT T = 300°K
(WEAK LINE APPROXIMATION)

WAVE NUMBER, CM\(^{-1}\)

\(p = 10^3\) ATM-CM

\(10^2\)
FIGURE 10 SPECTRAL EMISSIVITY OF NO AT T = 600°K
(WEAK LINE APPROXIMATION)
FIGURE 12  SPECTRAL EMISSIVITY OF NO AT T = 1800°K
(WEAK LINE APPROXIMATION)
FIGURE 15
SPECTRAL EMISSIVITY OF NO AT T = 5000 K
(WEAK LINE APPROXIMATION)
Figure 16 Spectral emissivity of NO at T = 7000°K (Weak line approximation)
Figure 17 Spectral Emissivity of HCl at T = 300K

Wave Number, cm⁻¹

P(2) = 10^6 atm²·cm⁻¹

N = 10^2
FIGURE 20 SPECTRAL EMISSIVITY OF HCl AT T = 1800°K
(STRONG LINE APPROXIMATION)
(Strong Line Approximation)

Figure 21: Spectral Emissivity of HCl at T = 2400K

Wave Number, cm⁻¹

$P_{21} = 10^6$ atm cm⁻¹
FIGURE 22 SPECTRAL EMISSIVITY OF HCl AT T = 3000°K
(STRONG LINE APPROXIMATION)
FIGURE 23
SPECTRAL EMISSIVITY OF HC4 AT T = 5000°K
(STRONG LINE APPROXIMATION)

WAVE NUMBER, CM⁻¹

P₂ = 10⁴ ATM²-CM⁻²
FIGURE 24 SPECTRAL EMISSIVITY OF HCA AT T = 7000°K
(STRONG LINE APPROXIMATION)
Figure 25: Spectral Emissivity of HF at T = 300 K.

Graph showing wave number (cm⁻¹) on the x-axis and spectral emissivity on the y-axis. The graph includes multiple curves indicating different pressures, with labels p_e = 10⁶ atm cm⁻³.
Figure 26 SPECTRAL EMISSIVITY OF HF AT T=600°K (STRONG LINE APPROXIMATION)
Figure 27: Spectral Emissivity of HF at T = 1200 K.
FIGURE 28  SPECTRAL EMISSIVITY OF HF AT T = 1800°K
(STRONG LINE APPROXIMATION)
(Strong Line Approximation)

Figure 3: Spectral Emittance of HF at T = 5000K.

Wave Number, cm⁻¹

\[ p_2 = 10^4 \text{ ATM} \cdot \text{cm} \]

\[ 10^2 \]
FIGURE 32  SPECTRAL EMISSIVITY OF HF AT T=7000°K
(STRONG LINE APPROXIMATION)
Figure 33: Comparison of published data of spectral emissivity
Figure 34: Comparison with other published data of spectral emissivity computed for NO at $T=3000^\circ K$ in the weak line approximation.
if the previously mentioned correction for line wing emission beyond the band head had been included in this analysis. Presumably the tail drawn by Stull and Plass \(^1\) in the region near 3300 cm\(^{-1}\) was computed for one particular combination of path length and pressure, perhaps for \(p = 100\) atm and \(l = 1\) cm.

Figure 34 shows the results of a computation by Breene\(^{10}\) (for NO in air at 3000°K, density ratio = 10, thickness = 5 cm). Penner, Sulzmann, and Ludwig's\(^8\) results for the simple harmonic oscillator model are also shown. It should be noted that for comparison all three of these computations utilized Breene's\(^{10}\) multiplicative factor of 1.88 for the average path length (i.e., for a 5-cm-thick layer, a value of \(l = 9.4\) cm was used).

A glance at Figure 34 shows that Equation (20) is not much more successful than the simple harmonic oscillator approximation in matching Breene's\(^{10}\) computation. The correction for line wing emission would add a tail to the curve representing Equation (20) and extend it somewhat farther past 2000 cm\(^{-1}\). (Since the line widths are of the order of 2 cm\(^{-1}\), this addition would not be very significant.) The highly asymmetric peaks in emissivity indicated by Equation (20) seem more reasonable than the nearly symmetric peaks shown by Breene.\(^{10}\) (Note, in comparison, that for HCl, the inclusion of the Herman and Wallis\(^{11}\) \(F\)-factor has caused a considerable augmentation of emissivity at lower frequencies and diminution at higher frequencies which makes the peaks nearly equal. This factor is negligible for the more symmetric NO molecule so that considerably more asymmetry would be expected on this basis.)

Emissivities have been computed (but not presented here) for HCl and HF in the weak line approximation and for CO and NO in the strong line approximation for the temperatures considered here. Also for several intermediate temperatures, emissivities have been computed for all four molecules in both approximations. It is planned to extend the computations to other molecules and also to consider Doppler broadening as well as Lorentz broadening.
REFERENCES
