

pling theory.

*Work supported by Defense Atomic Support Agency, under NWER Subtask HC010, Contract No. DASA-01-70-C-0032.

¹R. E. Olson, "Absorbing-Sphere Model for Calculating Thermal Energy Ion-Ion Recombination Reaction Rates" (to be published); R. E. Olson, J. R. Peterson, and J. Mosely, *J. Chem. Phys.* **53**, 3391 (1970).

²J. R. Peterson, W. Aberth, and J. Mosely, to be published; J. Mosely, W. Aberth, and J. R. Peterson, "Ion-Ion Mutual Neutralization Cross Sections for O_2^+ + NO_3^- and NO^+ + NO_3^- ," in Proceedings of the 23rd Annual Gaseous Electronics Conference, Hartford, Conn., October 1970 (to be published).

³M. E. Gardner, *Phys. Rev.* **53**, 75 (1938).

⁴J. Sayers, *Proc. Roy. Soc., Ser. A* **169**, 83 (1938).

⁵B. H. Mahan and J. C. Person, *J. Chem. Phys.* **40**, 392 (1964).

⁶T. S. Carlton and B. H. Mahan, *J. Chem. Phys.* **40**, 3683 (1964).

⁷H. Y. Yeung, *Proc. Phys. Soc., London* **71**, 341 (1958).

⁸S. McGowan, *Can. J. Phys.* **45**, 439 (1967).

⁹W. H. Aberth and J. R. Peterson, *Phys. Rev. A* **1**, 158 (1970).

¹⁰R. D. Rundel, K. L. Aitken, and M. F. A. Harrison, *J. Phys. B: Proc. Phys. Soc., London* **2**, 954 (1969).

¹¹W. Aberth, J. R. Peterson, D. C. Lorents, and C. J. Cook, *Phys. Rev. Lett.* **20**, 979 (1968); J. Mosely, W. Aberth, and J. R. Peterson, *Phys. Rev. Lett.* **24**,

435 (1970).

¹²H. G. Ebert, J. Booz, and R. Koeppe, *Z. Phys.* **181**, 187 (1964).

¹³M. N. Hirsh, P. N. Eisner, and J. A. Slevin, *Rev. Sci. Instrum.* **39**, 1547 (1968).

¹⁴M. N. Hirsh, P. N. Eisner, and J. A. Slevin, *Phys. Rev.* **178**, 175 (1969).

¹⁵M. N. Hirsh, E. Poss, and P. N. Eisner, *Phys. Rev. A* **1**, 1615 (1970).

¹⁶This relation assumes that recombination is the only volume loss process and that diffusion controls the ion densities in the immediate vicinity of the chamber walls, and hence controls the mass-spectrometer currents. This model, described in a forthcoming paper, agrees closely with rigorous calculations [E. P. Gray and D. E. Kerr, *Ann. Phys. (New York)* **17**, 276 (1962)] for the case of a single-positive, single-negative-ion plasma under these experimental conditions.

¹⁷Although the NO^+ -electron recombination coefficient [C. S. Weller and M. A. Biondi, *Phys. Rev.* **172**, 198 (1968)] is about the same as the NO^+ -ion recombination coefficient, electrons can be neglected because their density during irradiation, about $7 \times 10^6 \text{ cm}^{-3}$, is two decades below that of the ions, and in the afterglow it decreases more rapidly than the ions (see Ref. 14).

¹⁸Negative-ion reactions with neutral NO , O_3 , and NO_2 have been suggested [N. G. Adams *et al.*, *J. Chem. Phys.* **52**, 3133 (1970)] which could couple the NO_2^- and NO_3^- densities. We see no gross effects attributable to these reactions.

¹⁹R. E. LeVier and L. M. Branscomb, *J. Geophys. Res.* **73**, 27 (1968).

Ground-State Energy of a Charged-Boson Gas

Deok Kyo Lee

Department of Physics and Astronomy, University of Kansas, Lawrence, Kansas 66044

(Received 1 March 1971)

The ground-state energy of a charged-boson gas at high density is re-examined and it is found that Brueckner's exact result for the second-order energy is reproduced precisely by Feenberg's variation-perturbation method.

In recent years the charged-boson gas has been an interesting subject in the theoretical study of many-body boson problems. The ground-state energy of the system at high density was first evaluated exactly to second order by Brueckner,¹ who carried out complete summations of one- and two-ring diagrams using the Bogoliubov theory; his exact result is²

$$E/N = -0.8031/r_s^{3/4} + 0.0280 + O(r_s^{3/4}), \quad (1)$$

which agrees *numerically* with the variational result obtained by Lee and Feenberg³ in the Bijl-Dingle-Jastrow (BDJ) wave-function space. Because of this numerical agreement considerable energy has been expended in attempts to prove that two results agree analytically. The purpose of this report is to clarify this point by showing that the variational energy is higher in second order than the exact energy by an amount exactly equal to the energy shift generated by the three-phonon vertex.⁴

In order to evaluate the difference between the exact and variational energies in second order, here we introduce

$$t = \frac{1}{\sqrt{2}} \left[\frac{\hbar^2}{4\pi m e^2 \rho} \right]^{1/4} k, \quad \bar{t}'' = \bar{t} - \bar{t}', \quad \omega = [1 + t^4]^{1/2}, \quad S = t^2/\omega, \quad \{f\} = f + f' + f'' = f(t) + f(t') + f(t''). \quad (2)$$

Then, Brueckner's formula for the exact second-order energy term can be expressed as

$$\mathcal{E}^{(2)}[\text{exact}] = \mathcal{E}^{(2)}(\text{1 ring}) + \mathcal{E}_x^{(1)} + \mathcal{E}_{Y\lambda}^{(1)}, \quad (3)$$

where

$$\mathcal{E}^{(2)}(\text{1 ring}) = \frac{1}{4\pi^4} \int \frac{d\vec{t}d\vec{t}'}{SS'} (\omega + t^2)(1-S)^2(1-S')^2, \quad (4)$$

$$\mathcal{E}_x^{(1)} = \frac{1}{8\pi^4} \int \frac{d\vec{t}d\vec{t}'}{SS'(t'')^2} (1-S)(1-S')(1+SS'), \quad (5)$$

$$\mathcal{E}_{Y\lambda}^{(1)} = -\frac{1}{24\pi^4} \int \frac{d\vec{t}d\vec{t}'}{SS'S''\{\omega\}} \left[\left\{ \frac{1}{\omega} \right\} - SS'S'' \left\{ \frac{1}{t'^2} \right\} \right]^2. \quad (6)$$

Also, Lee and Feenberg's expression for the variational second-order energy is, from Eq. (57) of Ref. 3,

$$\mathcal{E}^{(2)}[\text{variation}] = (2\pi^4)^{-1} \int d\vec{t}d\vec{t}'(1-S)(1-S')(1-S'')^2. \quad (7)$$

Re-expressing the integrals of Eqs. (4)-(7) so that integrands are linear in the function ω and symmetric in the variables t , t' , and t'' , we find after some lengthy algebra that the energy difference in second order becomes

$$\mathcal{E}^{(2)}[\text{exact}] - \mathcal{E}^{(2)}[\text{variation}] = \Delta_1 + \Delta_2, \quad (8)$$

with

$$\begin{aligned} \Delta_1 &= -\frac{1}{24\pi^4} \int \frac{d\vec{t}d\vec{t}'}{SS'S''\{\omega\}} [SS'S''(2\{\omega\} - \{S\omega\} - \{S\}\{S\omega\} + 2\{S^2\omega\})^2 \\ &= -\frac{1}{24\pi^4} \int \frac{d\vec{t}d\vec{t}'}{SS'S''\{\omega\}} (1-S)^2(1-S')^2(1-S'')^2 \left[2\left\{ \frac{S^2\omega}{1-S} \right\} - \left\{ \frac{S}{1-S} \right\} \{S\omega\} \right]^2, \end{aligned} \quad (9)$$

$$\begin{aligned} \Delta_2 &= \frac{1}{12\pi^4} \int d\vec{t}d\vec{t}' \left[\left(4 - \frac{1}{S} \right) + 2\{S\} + SS'S'' \right] \{\omega\} + \left(\{S\} \left\{ \frac{1}{S} \right\} - 3 \right) \{S\omega\} + \left\{ \frac{\omega}{S} \right\} + \{S^3\omega\} \\ &\quad + \left(4 + \frac{1}{SS'S''} - 2 \left\{ \frac{1}{S} \right\} - \{S\} \right) \{S^2\omega\}. \end{aligned} \quad (10)$$

It is not difficult to show that the expression for Δ_1 is exactly the same as the second-order perturbation correction [Eq. (6) of Ref. 4] derived for liquid He⁴ by Davison and Feenberg,⁴ who realized that three-phonon components, which are absent in the BDJ-type wave function, contribute to the energy in second order. On the other hand, the integral of Eq. (10) can be reduced to

$$\Delta_2 = \frac{1}{2\pi^4} \int d\vec{t}d\vec{t}' (\vec{t} \cdot \vec{t}') \frac{(1-S)^2(1-S')^2}{SS'} = 0. \quad (11)$$

Thus, we arrive at the significant conclusion that Brueckner's exact result¹ for the ground-state energy is reproduced precisely by Feenberg's variation-perturbation method.^{3,4} The details of our analysis are extremely complicated, and will be reported elsewhere together with the numerical value of the perturbation correction Δ_1 .

¹K. A. Brueckner, Phys. Rev. **156**, 204 (1967).

²In Eq. (1), the energy and the mean particle separation r_s are measured in units of Rydberg ($me^4/2\hbar^2$) and the Bohr radius (\hbar^2/me^2), respectively.

³D. K. Lee and E. Feenberg, Phys. Rev. **137**, A731 (1965).

⁴T. B. Davison and E. Feenberg, Ann. Phys. (New York) **53**, 559 (1969).