II. ITINERANT ELECTRON FERROMAGNET MODEL

It is the purpose of this section to present an elementary theory, unifying several existing theories, of a single band, itinerant electron FM. In particular, we shall develop a theory, appropriate for 3-d electrons, for the Curie temperature, T_c , and its pressure derivative, $\partial T_c/\partial P$; and we shall show how estimates of the effective exchange, I, times the density of states at Fermi level, $N(\varepsilon_F)$, can be made from the measurements of $\partial T_c/\partial P$. The theory presented here follows quite closely the earlier work of Wohlfarth, Edwards and Wohlfarth, Shiga, 3 and Wohlfarth and Bartel, but includes details which have not been discussed in these earlier works.

For our model we assume that the exchange splitting is given by nI ζ where I is the effective intra-atomic exchange (accounting for the electron correlations) between the itinerant electrons, n is the number of d-electrons per atom, and ζ is the relative magnetization per electron arising from single-particle excitations. In the Stoner theory, the exchange splitting is $2k_B\theta' f$ where $k_B\theta'$ is the molecular field approximation interaction; thus $k_B\theta' = 1/2$ nI. The single particle excitations are described by the Stoner equations $\theta' = 1/2$ nI.

$$1/2 n(1 \pm \zeta) = \int_0^\infty f(\varepsilon, \eta^{\pm}) N(\varepsilon) d\varepsilon , \qquad (1)$$

where

$$f(\varepsilon,\eta) = \left\{ \exp \left[(\varepsilon - \eta)/k_B T \right] + 1 \right\}^{-1} ,$$

and

$$\eta^{\pm} = \mu \pm 1/2 \; \text{nIr} \, \pm \, \mu_B^{\, H} \qquad . \label{eq:eta_BH}$$

Here μ is paramagnetic chemical potential, H is the applied magnetic field, and $N(\varepsilon)$ is the density of states. In the limit as T \rightarrow T such that $\zeta \rightarrow 0$ and

letting H = 0, we can expand the Fermi function exponentials in the expression for ζ , Eq. (1), and obtain the well known result

$$I \int_{0}^{\infty} N(\varepsilon) \left| \frac{\partial f}{\partial \varepsilon} \right|_{T=T_{c}} d\varepsilon = 1 .$$
(2)

Within the framework of the model, Eq. (2) can be solved for T_c if $N(\varepsilon)$ is known. Even if we don't know $N(\varepsilon)$ we can solve Eq. (2) by use of the Sommerfeld expansion. To terms quadratic in T_c we obtain L_c

$$T_c^2 = T_F^2(\overline{I} - 1) / \overline{I} , \qquad (3)$$

where

$$\overline{I} = IN(\varepsilon_{\overline{F}})$$
 (4)

Here $N(\varepsilon_F)$ is the density of states per atom per spin at the paramagnetic Fermi level, and T_F is the effective degeneracy temperature defined by 9,10

$$T_{F}^{2} = \left\{ \frac{\pi^{2}}{6} \quad k_{B}^{2} \left[\left(\frac{N'(\varepsilon_{F})}{N(\varepsilon_{F})} \right)^{2} - \frac{N''(\varepsilon_{F})}{N(\varepsilon_{F})} \right]^{-1} \right\} , \qquad (5)$$

where $N^{(m)}(\varepsilon_F)$ is the m-th derivative with respect to energy evaluated at ε_F . The expression for T_c^2 , Eq. (3), is identical to what one would obtain from the singularity in the exchange enhanced susceptibility where the F-integral of Lang and Ehrenreich¹⁵ is expanded by means of a Sommerfeld expansion. In order for the system to be FM, we have from Eq. (3) the Stoner criterion, $\overline{I} \ge 1$.

The expression for T_c^2 , Eq. (3), is general to the extent that we have not specified the origin or nature of I and we have not restricted $N(\varepsilon_F)$. To find