Evidence for the existence of two stable sites for cobalt impurity atoms in aluminum

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AN ABSTRACT OF THE THESIS OF Arun Venkatachar for the
Master of Science in Physics presented August 17, 1971.

Title: Evidence for the Existence of Two Stable Sites for
Cobalt Impurity Atoms in Aluminum.

APPROVED BY THE MEMBERS OF THE THESIS COMMITTEE:

Donald G. Howard, Chairman

Bruce W. Brown

Rudi H. Nussbaum

Raymond W. Sommerfeldt

Using Fe$^{57}$ Mössbauer spectroscopy, two alternative sites
occupied by cobalt (10^{-4} at. %) impurity atoms in aluminum have
been isolated. The substitutional site A is the stable position
after annealing the sample above 840 K, followed by a rapid
quench. The impurity atoms in site A are characterized by a
single line Mossbauer spectrum (indicative of a cubic environment), a room-temperature $f = 0.502$ (r. m. s. displ. 0.071 Å) and an I. S. (rel. to Fe) = $-0.421$ mm/sec. For anneals below 770 K the cobalt atoms migrate to site B, which is characterized by a well resolved quadrupole doublet (indicative of a non-cubic environment), a large change in the value of the room-temperature $f = 0.766$ (r. m. s. displ. 0.071 Å), I. S. = $-0.150$ mm/sec (increased s-electron density at the nucleus). All linewidths are approximately 0.21 mm/sec, indicating high uniformity of impurity sites. The site distribution [A] / [B] varies from about 5 % to 95 % for anneals between 770 K and 830 K.
EVIDENCE FOR THE EXISTENCE OF TWO STABLE SITES
FOR COBALT IMPURITY ATOMS IN ALUMINUM

by

ARUN VENKATACHAR

A thesis submitted in partial fulfillment of
the requirements for the degree of
MASTER OF SCIENCE
in
PHYSICS

Portland State University
August 1971
TO THE OFFICE OF GRADUATE STUDIES:

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August 17, 1971
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I. INTRODUCTION

A crystal can act very much like a rigid body insofar as at least a fraction of low energy gamma rays are emitted or absorbed by a radioactive atom (embedded in the crystal) in a recoil-free fashion; that is, essentially without any energy loss, since the crystal as a whole takes up the recoil momentum. This phenomenon is called the Mössbauer effect and the probability for a recoiless emission (or absorption) is commonly referred to in the literature as the Lamb-Mössbauer or Debye-Waller factor $f$.

A number of investigators (1-6) have utilised the Mössbauer effect to study dilute alloys of Fe$^{57}$ (and its parent Co$^{57}$) in aluminum. There exists a general lack of agreement in the results of their investigations. The main purpose of this thesis was to re-examine the Co-Al system using not only Mössbauer line shapes (as used by previous workers) but also precision $f$-measurements. We have found that a combination of these two techniques can yield more information than just using line shape data.

As a result of our investigations, we have found that at very low concentrations ($10^{-6}$ at. % Co), cobalt atoms occupy two distinct sites in the host lattice. Co atoms in site A are characterised by a cubic environment and a low room-temperature $f$ ($= 0.502$). On the other hand, site B is characterised by a non-cubic environment and a much higher
room-temperature $f (= 0.766)$. In addition, using $f$-measurements we have followed the transformation of Co atoms from one site to the other, both as a function of time as well as temperature. These measurements have been useful in determining the nature of site B.

This thesis is divided into three sections: the first section contains the theory, where a brief analysis of the Debye-Waller factor is given. The influence of internal fields on line shapes has also been discussed in this section. In the experimental section, the emphasis is on the alloy preparation and the annealing procedure used. The third and final section contains the presentation and interpretation of data.
II. REVIEW OF LITERATURE

The earliest paper pertaining to the use of the Mössbauer effect in studying dilute alloys of iron in Aluminum is that due to Bara et al (1). The source used by Bara and his co-workers was a foil of Al on which was deposited Co$^{57}$ in the form of an aqueous solution of Co$^{57}$Cl$_2$. The sample was then diffused at 873 K in an argon atmosphere for 100 minutes. A room temperature line shape measurement on the source revealed that the spectrum consisted of a single line pattern. After further diffusion at the same temperature, the room temperature spectrum changed with the single line growing weaker and the gradual appearance of a six line, magnetically split pattern. The authors interpreted the six line pattern as the result of long paramagnetic spin relaxation times associated with an extremely dilute solution of magnetic iron impurity in a non-magnetic host lattice. Next, they studied the temperature variation of the hyperfine spectrum and found that at 573 K the six line pattern completely disappeared and only a doublet remained. They considered this to be a quadrupole doublet and attributed its formation to a structural change in the sample.

Sprouse and Kalvius (2) used a Mössbauer source prepared by introducing coulomb excited Fe$^{57}$ nuclei into an Al foil by the technique of recoil implantation. The Mössbauer spectrum measured at room temperature consisted of a broad single line pattern. The
broadening was attributed to an unresolved quadrupole coupling in the crystal.

Besides the above mentioned source experiments, work has also been done with absorbers made of dilute alloys of iron in aluminum. Bush et al (3) used a 0.05 wt % alloy (made by melting 99.999 % pure Al and Fe containing 68 % Fe$^{57}$) for their experiments. The Mössbauer spectrum taken after annealing the sample at 913 K for a period of 65 hours was found to be a single line pattern. Next, they annealed the sample for 128 hours at 593 K and found that the Mössbauer spectrum had changed to a quadrupole doublet. They attributed the single line to be characteristic of a substitutional solid solution of Fe in Al and the doublet pattern to be the result of iron precipitating out of the solid solution. In addition, they showed that contrary to phase diagram studies the precipitate did not correspond to FeAl$_3$ but was indeed a true precipitate of iron. They have suggested a distorted f.c.c. structure for the precipitate.

Janot and Lelay (4) working with dilute alloys (0.006-0.24 wt % of Fe in Al) as absorbers have been able to isolate two precipitate phases in addition to the solid solution phase mentioned above. Both these precipitates are characterised by quadrupole effects visible in their Mössbauer spectra. They have found a 'plate like' precipitate is formed by cold-working the sample prior to annealing it at 593 K. The 'rod like' precipitate was formed by heat treating the sample at 723 K for 13 hours.

Nasu et al (5) measured the shift of the solid solution
line as a function of temperature, and from this they calculated how much of the total shift relative to metallic iron is due to thermal shift and how much is due to the isomer shift.

Recently, Preston and Gerlach (6) have measured the Mössbauer spectra of dilute alloys of Fe in Al with the iron concentration ranging between 0.005 and 1.2 %. The spectrum analysis reveals the presence of two phases— a solid solution phase similar to that obtained by previous workers, and a multiple line component which they interpret to be the result of Fe$^{57}$ in Fe$_4$Al$_{13}$. They have found that these two phases can be easily distinguished from each other by the difference in their behaviour when either the concentration or temperature is varied.
TABLE I
COMPARISON OF PARAMETERS DERIVED FROM LINE SHAPE MEASUREMENTS REVIEWED IN THIS SECTION

<table>
<thead>
<tr>
<th>Component</th>
<th>Isomer Shift $\delta$ (mm/sec)</th>
<th>Quad. Splitting $\Delta E$ (mm/sec)</th>
<th>Line Width $\Gamma$ (mm/sec)</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Single Line</td>
<td>- 0.310</td>
<td>-</td>
<td>0.800</td>
<td>a, 1</td>
</tr>
<tr>
<td></td>
<td>- 0.440</td>
<td>-</td>
<td>0.660</td>
<td>b, 2</td>
</tr>
<tr>
<td></td>
<td>- 0.423</td>
<td>-</td>
<td>0.250</td>
<td>c, 3</td>
</tr>
<tr>
<td></td>
<td>- 0.424</td>
<td>-</td>
<td>0.280</td>
<td>d, 4</td>
</tr>
<tr>
<td></td>
<td>- 0.430</td>
<td>-</td>
<td>0.250</td>
<td>e, 5</td>
</tr>
<tr>
<td></td>
<td>- 0.422</td>
<td>-</td>
<td>0.242</td>
<td>f, 6</td>
</tr>
<tr>
<td>Quadrupole Split</td>
<td>- 0.740</td>
<td>0.320</td>
<td>-</td>
<td>a, 1</td>
</tr>
<tr>
<td>Line</td>
<td>- 0.140</td>
<td>0.270</td>
<td>0.260</td>
<td>c, 3</td>
</tr>
<tr>
<td>Ppt. Phase I</td>
<td>- 0.100</td>
<td>0.200</td>
<td>0.280</td>
<td>d, 4</td>
</tr>
<tr>
<td>Ppt. Phase II</td>
<td>- 0.200</td>
<td>0.300</td>
<td>0.350</td>
<td>d, 4</td>
</tr>
<tr>
<td>** $Fe_4A1_13$</td>
<td>- 0.010</td>
<td>0.382</td>
<td>0.265</td>
<td>f, 6</td>
</tr>
<tr>
<td></td>
<td>- 0.202</td>
<td></td>
<td>0.265</td>
<td></td>
</tr>
<tr>
<td></td>
<td>- 0.392</td>
<td></td>
<td>0.265</td>
<td></td>
</tr>
</tbody>
</table>

* All Isomer shifts quoted in table are with respect to natural iron.
** This spectra was fitted by using three lines of equal width and line positions given in column 2 of table.

b) Sprouse et al
c) Bush et al Scripta Met. 1, 75 (1967)
f) Preston and Gerlach
Debye-Waller Factor

Consider the emission of a gamma ray from a radioactive nucleus embedded in a crystal. Due to the strength and short range nature of the nuclear forces an assumption (7) can be made that the nuclear decay is not influenced by the state of the lattice and the lattice conditions in turn do not depend on the nuclear state. Under these conditions, we can write the matrix element describing the transition in which a gamma ray of momentum $\hbar \mathbf{k}$ is emitted by the nucleus whose center of mass coordinate is $\mathbf{X}$, while the lattice goes from an initial state $L_i$ to some final state $L_f$, as a produce of two matrix elements (7).

1) \[ M_L = \langle L_i | e^{i\mathbf{K} \cdot \mathbf{X}} | L_f \rangle <\!N> \]

The transition probability $P(L_i \rightarrow L_f)$ is proportional to the square of the above matrix element and has been shown by Lipkin (8) to have the form:

2) \[ P(L_i \rightarrow L_f) = |\langle L_i | e^{i\mathbf{K} \cdot \mathbf{X}} | L_f \rangle|^2 \]

The Mössbauer effect arises if during the nuclear transition there is no simultaneous lattice transition. For such a transition $L_i \rightarrow L_f$, and the gamma ray is emitted recoilessly. The probability for a Mössbauer transition (or the Debye-Waller factor), $f$, will be:
For a crystal with harmonic forces the Debye-Waller factor $f$ takes a simple form (7):

$$ f = \exp \left( \frac{4\pi \langle x^2 \rangle}{\lambda^2} \right) $$

where $\langle x^2 \rangle$ is the time averaged mean squared displacement of the impurity atom along the direction of gamma ray emission, and $\lambda$ is the wave length of the gamma quantum. Equation (4) describes $f$ to a high degree of accuracy, including the case where the atom is an impurity vibrating in localized mode (9). For harmonically bound crystals the mean squared displacement is related to the phonon frequency spectrum $G_h(\omega)$ in a mono-atomic crystal (15), while for an impurity atom $G_\text{i}(\omega)$ stands for the "dynamic response function", (15) which describes the coupling of the impurity modes to the unperturbed normal modes of the host. In that case $G_\text{i}(\omega)$ also depends on the impurity-host mass ratio and it may also include localized vibrational modes (15).

5) $$ \langle x^2 \rangle = \frac{\hbar}{m} \int G(\omega) \frac{\coth \left( \frac{\hbar \omega}{2kT} \right) G(\omega)}{\omega} \, d\omega $$

where $G(\omega)$ is normalized: $$ \int G(\omega) \, d\omega = 1 $$

At high temperatures ($\hbar \omega/kT < 2\pi$), $\coth (\hbar \omega/kT)$ in equation (5) can be expanded:

6) $$ \langle x^2 \rangle = \frac{kT}{m} \int \left[ 1 + \frac{1}{12} \left( \frac{\hbar \omega}{kT} \right)^2 - \frac{1}{720} \left( \frac{\hbar \omega}{kT} \right)^4 + \ldots \right] \frac{G(\omega)}{\omega^2} \, d\omega $$

Using the moments of the frequency spectrum defined as $$ \bar{n}_n = \int G(\omega) \omega^n \, d\omega $$
we can write the high temperature limit as follows:

\[ <x^2> = \frac{k}{m} \left[ \omega^{-2} T + \frac{1}{12} \left( \frac{\hbar}{k} \right)^2 T^{-1} - \frac{1}{720} \left( \frac{\hbar}{k} \right)^4 \omega^2 T^{-3} + \ldots \right] \]

Each term in the above expansion can be described in terms of an effective Debye frequency spectrum such that its nth moment \( \omega_D(n) \) equals the real moment \( \omega^n \). The \( \omega_D(n) \) defines weighted Debye temperatures \( \Theta_D(n) \) according to:

\[ \Theta_D(n) = \frac{\hbar}{k} \omega_D(n) = \frac{n^3}{3} \frac{\omega^n}{\omega_D(n)} \]

for \( n \geq -3 \)

From equations (7) and (8) we find that at high temperatures and for harmonic forces:

\[ -\ln f_h = \frac{6RT}{k} \left( \frac{1}{\Theta_D(-2)} \right)^2 \left[ 1 + \left( \frac{\Theta_D(2)}{6T} \right)^2 + \ldots \right] \]

The first term contributes more than 95% for \( T/\Theta_D < 0.72 \). We therefore see that high temperature measurements yield the lattice parameter \( \Theta_D(-2) \).

So far, we have assumed that the forces present in the crystal are harmonic, while in fact all crystals have some inherent anharmonicity. Marradudin and Flinn (10) have shown that as a result of high temperature anharmonicity, we have the expression:

\[ \ln f = \ln f_h \left[ 1 + \varepsilon(-2) T \right] \]

where \( f_h \) is the harmonic Mössbauer fraction and \( \varepsilon(-2) \) is the anharmonicity parameter at high temperatures.
The gamma ray emitted is not monochromatic, but has an intensity distribution which is Lorentzian in character and is given by the expression:

$$I(E) = \frac{\Gamma^2/4}{(E-E_0)^2 + \Gamma^2/4}$$

$E_0$ is the energy of the nuclear transition; $\Gamma$ is the natural width of the emission band and is equal to the full width at half maximum of the above Lorentzian line shape. From Heisenberg's uncertainty principle, $\Gamma$ can be related to $\tau$, the mean life time of the decaying nucleus, by the relationship:

$$\Gamma = \frac{\hbar}{\tau}$$

Wertheim (12) has shown that for thin absorbers, the experimentally determined Mössbauer spectrum which is a result of the folding of the absorption and emission lines, is also Lorentzian in shape with a line width equal to the sum of the line widths of the absorption and emission lines. If both these lines have natural line widths, the Mössbauer line width is just twice the line width of the gamma ray. Natural line widths are rare in Mössbauer research, as various factors such as source and absorber thickness, velocity resolution of apparatus, internal fields, etc., broaden and change the
shape of the line.

Due to the narrow line width of the nuclear transition involved in the Mössbauer effect, resonant absorption is extremely sensitive to energy of the gamma radiation. For this reason small interactions between the nucleus and orbital electrons, which cannot be observed directly by other methods, manifest themselves very markedly in the Mössbauer effect. It is thus the influence of the electronic environment on the nuclear transition, both in emission and absorption that determines the hyperfine structure of the Mössbauer spectrum.

The main interactions that reveal themselves in the Mössbauer effect are:

a) Nuclear Isomer shift
b) Nuclear quadrupole coupling
c) Magnetic hyperfine interactions

The nuclear isomer shift is the nuclear analogue of two hyperfine interactions observed in the electronic spectrum: the isotope shift and the isomeric shift. These effects are changes in wavelength of spectral terms of different isotopes and in the isomeric states of one and the same isotope. The nuclear isomer shift, which manifests itself as a shift from zero velocity of the centroid of the Mössbauer spectrum, is due to the electrostatic interaction between the charge distribution of the nucleus and those electrons which have a non-vanishing probability in the region occupied by the nucleus. An expression (12) can be derived for this interaction between the nuclear charge and the electronic charge:
where $e |\psi(o)|^2$ is the electronic charge density, $Z$ the atomic number for the impurity atom $<R^2>_{gr}$ & $<R^2>_{ex}$ the mean squared radii for the ground and excited states respectively (see figure 1). The experimental quantity measured in the Mössbauer effect is the difference (see equation 12) between a source characterized by a given electron density $|\psi_s(o)|^2$ and an absorber characterized by a different electron density $|\psi_a(o)|^2$. Thus:

$$\Delta E = Ze^2 \left[ <R^2>_{ex} - <R^2>_{gr} \right] |\psi_s(o)|^2 - |\psi_a(o)|^2$$

The nuclear isomer shift is always measured as a relative quantity, and is expressed as a product of a nuclear factor and an electronic one.

The nuclear quadrupole coupling results in an interaction of the nuclear quadrupole moment $Q$ with the gradient of the electric field due to the other charges present in the crystal. The contribution to field gradient comes from three sources:

a) Charges external to the impurity atom; the ionic charges of the lattice may produce a field gradient at a given atomic site.

b) Electrons belonging to the atom; example: in transition ions the inner d or f shells are not completely filled and may cause an electric field gradient at the nucleus.

c) Polarization of the inner shells of the atom; the filled inner shells have spherical symmetry and
normandy give no contribution to the field gradient. However, both external charges outside the atom and its own unpaired electrons can polarize these inner shells, which then will contribute to the field gradient at the nucleus.

For the 14.4 kev transition in Fe\textsuperscript{57} the ground and exited states are characterized by spins 3/2 and 1/2. In the presence of an electric field gradient the excited level is split into two sub-levels (see figure 2). The Mössbauer spectrum will consist of two components corresponding to the transitions: $\pm 3/2 \rightarrow 1/2$ and $\pm 1/2 \rightarrow 1/2$. If there is an electric field of cubic symmetry acting on the emitting nucleus there will be no field gradient and only one line in the Mössbauer spectrum corresponding to the transition $3/2 \rightarrow 1/2$. On the other hand, a non-zero electric field gradient indicates a departure from cubic symmetry and the spectra will consist of two components.

The magnetic hyperfine interactions arise due to the effective magnetic field acting on the nucleus. The main contributions to the field are:

a) The Fermi contact interaction: direct coupling between the nucleus and unpaired s-electron density.

b) The contribution of the orbital magnetic moment.

c) Dipole interaction with the electronic spin.

Wertheim (12) has shown that in the case of the 14.4 kev transition in Fe\textsuperscript{57}, six different transitions (see figure 3) are possible if the
Iron nucleus is acted upon by a magnetic field. Therefore, in the natural Iron absorber (commonly used for line shape measurements) the spectrum is split into six completely resolved lines of natural width.
Figure 1. Isomer shift

Figure 2. Quadrupole splitting in $^{57}$Fe

Figure 3. Magnetic hyperfine splitting
IV. EXPERIMENTAL

Source Preparation

A piece of Aluminum was cut with a jeweler's saw from a single crystal of 99.999% purity obtained from Semi-Metals Inc. This piece was rolled into a foil of thickness 0.35 mm. A small rectangular (8 mm x 6 mm) piece was cut from this foil and the surface degreased with tri-chloroethylene. The foil was then dropped into a test tube containing a 1 N solution of NaOH so as to dissolve as much of the oxide (Al₂O₃) layer as possible. The foil was then washed with distilled water to remove all traces of the alkali. After allowing the surface to dry a small amount of beef-albumin was deposited on the Al surface. The albumin serves as a wetting agent. Next approximately 1.5 milli-curies of activity in the form of CoCl₂ in 0.1 N HCl was pipetted onto the metal surface. The activity was measured by placing the foil under a NaI scintillation counter and determining the count rate of the 122.0 kev gamma ray, then comparing this count rate with that for a Cu source with known activity concentration.

The foil was then placed on a small quartz plate and gently introduced into a quartz tube closed at one end. The tube was then inserted into a furnace heated to about 898 K and the source diffused in an atmosphere of hydrogen for 3 days. The hydrogen atmosphere serves two main purposes: reduces CoCl₂ to metallic Cobalt and
secondly prevents the Al surface from getting oxidized. A line
shape was taken after this initial heat treatment and the results
of this measurement indicated that the sample required additional
diffusion. The source was put back in the furnace at the same
temperature and the diffusion process continued for another week.
For the second diffusion we placed the system under vacuum (10^{-4}\text{mm}
of Hg). The high vacuum served as an added precaution against
oxidation of the sample. The diffusion temperature and time were
chosen after a careful study of the solubility of Co in Al (26).
After the completion of this second diffusion, the sample was
removed and scrubbed with pumice soap and then etched with 1 N \text{NaOH}
solution. This removes all traces of activity left on the surface.
The count rate from the foil was measured again and it was found
that there was about 0.8 milli-curies of activity associated with
the source (Al II).

Annealing Procedure

The annealing of the Al source was carried out in a Lindberg
high temperature tube furnace. Silicon Carbide heating elements are
used to heat the furnace. These elements cross the heating chamber
at right angles with closer spacing at the chamber ends then at the
center. This arrangement of the elements gives a uniform heat
distribution in the chamber. The furnace is capable of operating in
the temperature range 773 K to 1773 K. The sample is placed in the
above mentioned quartz tube and introduced in the furnace. The tube
is 69 cm. in length and 1.8 cm. in diameter. The annealing was
carried out in an atmosphere of hydrogen. The temperature of the source was measured by means of a chromel-alumel thermo couple placed in the heating chamber of the furnace. The tube was so positioned that the sample lay as close to the thermo couple as possible. The temperature fluctuations in the furnace were in the order of \( \pm 2 \) K. The quenching of the source was carried out by plunging the tube into a large beaker containing cold water.

**Line Shape Measurements**

Mössbauer line shape measurements involve the determination of the counting rate of the gamma radiation emitted by a source and passing through an absorber as a function of the instantaneous Doppler velocity of the source. This is achieved by using the speaker drive described by Nees (12). The absorber used in measuring line shapes with Al II was natural Iron - a 7.6 mg/cm² foil of Iron containing a natural abundance (2%) of Fe\(^{57}\). Due to the nuclear Zeeman effect mentioned in the theory section of this thesis, the Mössbauer spectrum will consist of six well resolved lines. Of these six lines, only the inner two closest to zero velocity have been used by us.

**Debye-Waller Factor Measurements**

The Debye-Waller factor \( f \) was measured both at room temperatures and elevated temperatures using the black, wide absorber technique (1). The experimental value of \( f \) has to be corrected for:

a) Re-radiation from the black absorber
b) Blackness of the black absorber.

c) Finite amount of resonant absorption at $I_\infty$.

Besides these corrections, one has to make an additional correction for the Compton scattering from the Beryllium window in the furnace used for high temperature $f$ measurements. These corrections as well as the experimental set up used has been described in detail by Housley (14).

Due to the low Atomic number of $\text{At}$ there are no host x-rays of energy comparable to the 14.4 kev gamma ray. An energy spectrum was taken for the sample, using an $\text{Xe} - \text{CH}_4$ filled proportional counter, an FET preamplifier and a 512 multi-channel pulse height analyzer. The spectrum (see Fig. 4) shows two well resolved peaks corresponding to the 14.4 kev gamma ray and 6.3 kev $K_\alpha$ x-ray of iron. Thus, no additional corrections have to be made for the presence of the host x-rays. Using the corrections discussed in this sub-section, we have found the statistical accuracy for $f$-measurements to be better than 0.5%.
Figure 4. Energy spectrum of Fe$^{57}$ in Aluminum
V. DATA PRESENTATION AND INTERPRETATION

The variation of the Debye-Waller factor $f$ with temperature has been experimentally determined in the range $293 \, \text{K} - 545 \, \text{K}$. The results of these measurements are shown in figure 5. The expected temperature dependence of $(-\ln f)$ on the basis of the Debye model has also been calculated. The results of these calculations are the solid lines shown in figure 5. The method used to fit the parameters $\Theta_D(-2)$ and $c(-2)$ has been described in detail by Nussbaum et al. (15)

Examining the curve marked A in figure 5, one notices the excellent agreement between the experimentally determined value of $f$ and its theoretical counterpart in the temperature range $293 \, \text{K} - 500 \, \text{K}$. Above $491 \, \text{K}$ the measured value of $f$ no longer follows the predicted line but is greater than that predicted by the Debye model. From equation 4 we see that $-\ln f = <x^2>$ where, $<x^2>$ is the mean squared displacement of the Fe$^{57}$ impurity atoms in the Aluminum matrix. An increase in $f$ therefore, corresponds to a decrease in the value of $<x^2>$. Since, the mean squared displacement depends on the local environment surrounding the impurity atom any changes in its value indicate a subsequent change in the environment. Moreover, at $545 \, \text{K}$, the value of $f$ is not constant but is found to increase with time (see figure 6). Based on these observations, we propose that the increase in the value of $f$ is caused by the diffusion of the Co$^{57}$
Figure 5. Temperature dependence of the Debye-Waller factor for cobalt impurity atoms in sites A and B respectively.
Figure 6. Time dependence of the Debye-Waller factor for Co$^{57}$ at 545 K.
impurity atoms from one lattice site to another, non-equivalent site which is characterized by a much higher value of the Debye-Waller factor. These two sites will be referred to in this thesis as sites A and B respectively. A detailed analysis of sites A and B is given below:

I. SITE A

Annealing the sample near its melting temperature for 8-10 days results in the creation of site A. The temperature range used for creating this state lies between 840 K and 930 K. There is no direct evidence, such as x-ray diffraction, etc., to indicate the type of lattice sites that the \( ^{57} \)Co atoms could occupy in the host matrix. The atomic radius of cobalt (\( R_{Co} = 1.25 \, \text{Å} \)) is about 14% less in magnitude than the radius of aluminum (\( R_{Al} = 1.43 \, \text{Å} \)). On the basis of size ratios, Haag's rule (18) would predict a substitutional solid solution for the Co-Al system. If cobalt atoms do occupy substitutional sites in the Al matrix, the Mössbauer spectrum should consist of a sharp, well defined single line pattern. This is so because of the following reasons:

a) There is no magnetic ordering in the phase, so there will be no broadening of the line due to any magnetic effects.

b) The amount of cobalt atoms is very small (approximately \( 10^{-4} \) at. % in the case of Al II) so assuming a perfectly random distribution for the \( ^{57} \)Co atoms, there is very little likelihood that a given cobalt atom will
Figure 7. Mössbauer spectrum for Co$^{57}$ atoms occupying site A.
have a second cobalt atom near it to affect its Mössbauer spectrum.

c) Since the substitutional Co atoms are not quite the right size, their presence undoubtedly causes a local distortion of the host lattice. The symmetry about the Co atom will however remain essentially cubic and so the substitution of the cobalt atom merely introduces cubically symmetric distortions. Under these circumstances, the Co atom experiences no electric field gradients to broaden its spectrum.

The Mössbauer spectrum for Co$^{57}$ atoms in site A, shown in figure (7) was taken after the sample (Al II) was subjected to 10 days of annealing at 898 K. The spectrum consists of a slightly broadened single line pattern. The measured line shape parameters are listed in table (II). We have been unable to give a suitable explanation for the line broadening mentioned above as another sample (Al III) was prepared (under identical conditions to that used for Al II) in which the site A spectrum consisted of a single line having natural width. On the basis of these line shape measurements on Al II and Al III, we conclude that site A is substitutional in character.

The next step in the analysis of the data is to obtain values for the lattice parameters $\theta_D$(-2) and $\epsilon$(-2) for Co atoms in the substitutional site. It must be noted that since beyond 500 K a significant number of Co atoms have diffused from the substitutional
### TABLE II
LINE SHAPE AND LATTICE DYNAMICAL PARAMETERS FOR IRON ATOMS OCCUPYING SITE A

<table>
<thead>
<tr>
<th>ISOMER SHIFT (mm/sec)</th>
<th>LINE WIDTH (mm/sec)</th>
<th>DEBYE TEMP (Fe)</th>
<th>DEBYE TEMP (Al)</th>
<th>ANHARM. PARAMETER ( \epsilon(-2) ) (K)(^{-1} )</th>
<th>FORCE CONSTANT RATIO ( \alpha' / \alpha )</th>
</tr>
</thead>
<tbody>
<tr>
<td>-0.421</td>
<td>0.230</td>
<td>246.1 K</td>
<td>396.0 K</td>
<td>~0.0</td>
<td>~0.80</td>
</tr>
</tbody>
</table>

* Relative to natural iron
** Value taken from reference (22)
site to site B, as is evident from the anomalous increase in the value of $f$ (see figure 5). Therefore, the data above 500 K cannot be used to determine $\Theta_D(-2)$ and $\varepsilon(-2)$. Table (II) lists the values obtained for these lattice parameters.

II. SITE B

Annealing the sample for 8 - 10 days between 545 K and 780 K results in the formation of Co$^{57}$ atoms occupying site B. Site B can be readily distinguished from the substitutional site A on the basis of the following characteristics:

a) Non-cubic symmetry - Co atoms in site B have a non-cubic environment. This is clearly revealed by the Mössbauer spectrum taken after the sample was annealed for 9 days at 706 K. The spectrum is shown in figure 8. The line shape is characterized by a well defined quadrupole doublet with the two peaks being equal in intensity, thereby indicating the existence of an electric field gradient at the site of the Co impurity. The line shape parameters have been listed in Table III.

b) Increase in $s$-electron density - the isomer shift relative to natural Iron for the impurity atoms in site B is less in magnitude than that in site A. This corresponds to an increase in $s$-electron density for Fe$^{57}$ atoms in site B. Using the 'Walker Chart' (see Appendix A), the electronic configuration for
Figure 8. Mössbauer spectrum for Co$^{57}$ atoms occupying site B.
Iron atoms in site B can be written as $3d^{7.4s^{0.95}}$. The corresponding electronic configuration for site A impurity atoms can be written either as $3d^{7}4s^{0.72}$ or $3d^{7.15}4s^{0.85}$. The importance of this data is not, however, yet fully understood, mostly because the theory of metals and alloys has not been sufficiently developed. However, isomer shifts may prove to be a useful tool for investigating the environments surrounding the impurity atoms.

c) Tight binding - f-measurements on At II (with Co$^{57}$ atoms in site B) reveal that the mean squared displacement is much smaller compared to that for site A, under identical temperature conditions. This is clearly seen in curve B (figure 5) which represents the temperature dependence of f with 90% of the impurity atoms in site B (insufficient annealing at 545 K resulted in this partial transformation of cobalt atoms from site A to site B). We have corrected for the incomplete transformation and the results of this calculation is the curve marked B in figure (5). Analyzing the data shown in figure (5) we find the Debye temperature $\theta_D(-2)$ to be much higher than that for site A. The various lattice parameters derived from f-measurements have been listed in Table III.

To fully understand the nature of the crystal environment which produces
### TABLE III
LINE SHAPE AND LATTICE DYNAMICAL PARAMETERS FOR IRON ATOMS OCCUPYING SITE B

*(Relative to natural iron)*

<table>
<thead>
<tr>
<th><em>ISOMER SHIFT</em> ($\delta$) (mm/sec)</th>
<th>LINE WIDTH ($\Gamma$) (mm/sec)</th>
<th>QUAD. SPLITTING ($\Delta E$) (mm/sec)</th>
<th>DEBYE TEMP. ($\Theta_D$) (K)</th>
<th>$\epsilon(-2)$ ($\text{K}^{-1}$)</th>
<th>FORCE CONSTANT RATIO $\frac{a'}{a}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$-0.150$</td>
<td>$0.210$</td>
<td>$0.279$</td>
<td>$\sim 350.0$ K</td>
<td>$\sim 0$</td>
<td>$\sim 1.60$</td>
</tr>
</tbody>
</table>

* *
site B, a number of quenching experiments were performed on Al II. By this procedure it is possible to 'freeze-in' a high temperature state, and yet examine the Mössbauer effect at room temperature where the Debye-Waller factor, and hence the total fraction of recoilless gamma rays emitted is large. We started these measurements with the impurity atoms all in site B (after a long anneal at 545 K), and then re-annealed the sample at a number of temperatures lying between 545 K and 898 K long enough to establish a new equilibrium configuration. After a rapid quench (assuming that no changes occurred in the impurity distribution) we determined the room temperature value of the Debye-Waller factor. The times required to reach equilibrium at each annealing temperature was approximately of the order of 8 to 10 days. The results of these f-measurements are shown in figure (9), where we have plotted K(T) as a function of the annealing temperature T. K(T) is the fraction of site B impurity atoms and has been defined as follows:

\[ K(T) = \frac{f_{\text{exp}} - f_A}{f_B - f_A} \]

where \( f_{\text{exp}} \) is the room temperature f measured after quenching Al II from a temperature T and \( f_A \) and \( f_B \) are the f values (at room temperature) for sites A and B respectively. For \( K(T) = 1 \), we have all the Co\( ^{57} \) atoms occupying site B in the host matrix and for \( K(T) = 0 \), all the atoms are in site A. For the case, \( 0 < K(T) < 1 \), which results when Al II is quenched from temperatures between 760 K and 825 K, the measured line shape is a mixture of a single line pattern and a quadrupole doublet (see figure 10)
Figure 9. Temperature dependence of $K(T)$. 
Figure 10. Mössbauer spectrum taken after annealing Al II at 793 K for 3 days.
characteristic of the co-existence of the two sites. We can therefore conclude that the two impurity sites are independent and that there exists a temperature dependent equilibrium of their concentrations. Based on these observations we propose the following two models for site B:

MODEL I. The impurity atoms could be paired off in the form of dumbbells (split <100> interstitials), as in the case of Co$^{57}$ in gold (21, 17). Since this model requires two impurities to be in the nearest neighbor positions before such a pair can be formed, the dependence on diffusion enters the picture quite naturally. If the atoms are assumed to be hard spheres with the diameter given by the nearest neighbor spacing of the metal (see Fig. 11), it is possible to determine the size of the hole created by removing one of the Al atoms. This hole is too small to accommodate two cobalt atoms. More room can be created for the pair by forming these dumbbells in dislocations or by distributing a vacancy among the Al atoms surrounding the pair. The Mössbauer spectrum for site B (Figure 8) does not reveal any excessive line broadening, indicating the absence of Co$^{57}$ atoms in dislocations. This favors the view that a formation of a dumbbell requires the presence of vacancies amongst the Al atoms close to the pair. This distribution of vacancies coupled with the slow diffusion of Co in Al could explain the long times involved in creating site B.

This model gives a satisfactory explanation for both the quadrupole splitting and high f's observed for site B. In this configuration the impurity atoms occupy positions which lack cubic symmetry, thereby creating a non-zero electric field gradient (at the
Figure 11. Position of the impurity dumbbell in the Al lattice. The symmetry axis of the dumbbell can lie along any of the three [100] directions.
site of the Co atom). In addition the crowding effect experienced by the impurity atom (in the dumbbell configuration) will result in a mean squared displacement much smaller in magnitude than that for the impurity atom in a substitutional site. There is one major drawback to this model: it fails to predict the slow times involved in breaking up dumbbells. So far, all attempts to modify the theory in order to account for the slow break up time have met with little success.

MODEL II. The impurity atoms could be a part of an intermetallic phase that precipitates out of the solid solution. From the Co-Al phase diagram, one expects Co to precipitate out as the intermetallic compound CoAl₃ (26). CoAl₃ has a monoclinic structure (3) with 100 atoms per unit cell and five different types of Co atom environment. Since each type of Co atom environment should have a characteristic Mössbauer spectrum, one would expect a complicated line shape for this phase (6). The Mössbauer line shape for site B is however indicative of only one type of Co atom environment, therefore ruling out the possibility that the precipitate formed is CoAl₃. However, Co may very well be precipitating out in the form of some other intermetallic compound.

An alternate view is to consider the possibility of the formation of small clusters of Co within the host matrix. The screened potential around a Co atom in Al, is given by the expression (25):
$V(R) = \frac{Z e^2 \cos(2k_F R - \delta)}{k_F R}$

$V(R)$ is the electrostatic potential at a distance $R$ from the impurity atom and $k_F$ the Fermi wave number for aluminum. This potential has been plotted in figure (12) and shows the existence of an attractive force between Co atoms at nearest neighbor distances. Co atoms therefore have a tendency to cluster and possibly form Guinier-Preston zones (see Appendix B).

The formation of a new phase (whether an inter-metallic ppt. or Co clusters) involves the process of nucleation (24). The latter in turn, depends on the following three steps (25):

a) Assembly of the proper kind of atoms by diffusion.

b) Structural change into one or more intermediate structures.

c) Formation of the new phase.

Due to the low solubility of Co in aluminum, step (a) is very slow. The slowness of step (a) coupled with steps (b and c), which are also time dependent (24), can explain the long times needed to form site B. Similarly, the break up of these clusters (or precipitates) is slow.

Model II, on the other hand cannot give a satisfactory explanation for the high $f$'s and observed line shape splitting as we have no direct information regarding the exact structure of this new phase.

* From the classical nucleation theory, the break up is analogous to the slow evaporation of atoms from the surface of a liquid drop.
Figure 12. Electrostatic potential per unit positive charge about an Co atom in an Al matrix.
VI. SUMMARY AND CONCLUSIONS

Using Mössbauer spectroscopy, two distinct lattice sites occupied by Co (10^{-4} at. %) in Aluminum have been isolated and tentatively identified. The substitutional site A is the stable position after annealing a dilute alloy above 840 K for a period of 8 - 10 days. Co atoms in site A are characterized by a single line Mössbauer spectrum and a room temperature $f = 0.502$. Annealing below 770 K for 8 - 10 days results in the migration of the impurity atoms to site B. This site is characterized by a well defined quadrupole doublet (indicative of a non-cubic environment) and a much higher value for the Debye-Waller factor ($f = 0.766$). The site distribution A / B varies from 95% to 5% for anneals between 770 K and 825 K. All linewidths are approximately 0.21 mm/sec, indicating a high uniformity of impurity sites in the host matrix.

Two possible models for site B have been proposed in this thesis. Model I, considers the impurity atoms to be in the form of dumbbells (split <100> interstitials). In such a configuration, two Co^{57} atoms are crowded into the same substitutional site. The impurity atom therefore has a non-cubic environment (resulting in the observed splitting of the spectrum) and the crowding effect causes a decrease in the mean squared displacement of the Co atom (or increase in $f$). In addition, the formation of a dumbbell is directly related to the diffusion of Co atoms in Al. As the diffusion of Co below 770 K is
very slow, the formation of site B is a slow process (as borne out by observations). However, the model fails to account for the slow break-up rate of site B.

In model II, the Co impurity atoms are assumed to be in the form of either an intermetallic precipitate or Guinier-Preston zones. Such a model is capable of explaining (on the basis of the classical nucleation theory) the slow rate of formation and break-up of site B. On the other hand, due to lack of information regarding the structure of these precipitates (or zones) it is uncertain whether they can explain the observed line splitting and high f's.

As yet, we are unable to choose between the two models. A number of experiments are in progress which might shed more light on the precise nature of site B. Recently, we have prepared a more concentrated alloy of cobalt (10⁻² at. % Co) in Al. Preliminary measurements made on this alloy reveal the presence of only one type of impurity site, namely site B at all temperatures below 832 K. This is at variance with the absorber experiments conducted by Bush et al (3) on an alloy of iron (5·10⁻² at. % Fe) in Al, which clearly shows the existence of two impurity sites (identical to site A and site B). This difference in the results of the source and absorber measurements indicate that, although one studies the nuclear physics of iron in both type of measurements (source and absorber), the metallurgy involved may be quite different. Source measurements involve the metallurgy of the Co-Al system, whereas absorber work is concerned with the Fe-Al system. Therefore, we feel that a portion of the inconsistency in the results of previous workers (1-6) could be explained by the difference between source and absorber measurements mentioned above.
### TABLE IV

**SUMMARY OF DATA FOR SITES A AND B OCCUPIED BY CO\(^{57}\) IN Al**

<table>
<thead>
<tr>
<th>LATTICE SITE</th>
<th>ISOMER SHIFT (\delta) (mm/sec)</th>
<th>QUADRUPOLE SPLITTING (\Delta E) (mm/sec)</th>
<th>LINE WIDTH (\Gamma) (mm/sec)</th>
<th>DEBYE TEMP (\Theta_D) (K)</th>
<th>REL. FORCE CONST. (\alpha/\alpha)</th>
<th>MOSSBAUER FRACTION (f)</th>
<th>MEAN SQUARE DISPLACEMENT (&lt;x^2&gt;) Å(^2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>-0.421</td>
<td>-</td>
<td>0.230</td>
<td>246.0 K</td>
<td>~0.80</td>
<td>0.502 ±0.001</td>
<td>0.013</td>
</tr>
<tr>
<td>B</td>
<td>-0.150</td>
<td>0.279</td>
<td>0.210</td>
<td>350.0 K</td>
<td>~1.60</td>
<td>0.766 ±0.001</td>
<td>0.005</td>
</tr>
</tbody>
</table>

* Isomer shifts quoted in table are with respect to natural iron.*


25. G. M. Hood, Phil. Mag. 21, 305 (1970)

APPENDIX A

WALKER CHART

Isomer shifts have been measured in various host materials and a large amount of information has been obtained for metals and alloys. Walker et al (20) reported the information of the Fe$^{57}$ isomeric shift in various solids relative to stainless steel in the form of the "Walker Chart". In this chart the total s-electron density is plotted as a function of the percentage of 4s character for various 3d electron configurations. In order to interpret the observed isomeric shift according to the "Walker Chart", it is necessary to convert from the value relative to iron to that relative to stainless steel. The value of Fe$^{57}$ isomeric shift in iron relative to stainless steel is known to be about 0.09 mm/sec. Therefore, the value of the isomeric shift in A into relative to stainless steel will be given by:

18) \[ I.S. \text{ (St.)} = I.S. \text{ (Fe)} + 0.09 \]

where I.S. (Fe) is the measured isomer shift relative to natural iron. Once the isomer shift (rel. to S.S.) is known, the corresponding electronic configuration can be directly read off from the chart.
APPENDIX B

GUINIER-PRESTON ZONES

The ideal Guinier-Preston zone is a region of the matrix with high solute concentration, but in which all the atoms are situated on the sites occupied in the random solid solution. A zone has a different composition and may have differing interatomic distances from those of the matrix, but its structure is continuous with that of the matrix. These clusters are generally small (consisting of a few atoms) and may be either plate shaped or spherical depending on the extent of the change produced in the lattice parameter (of Al) by the addition of the solute atom.

Nearly ideal zones with no deformation of the surrounding lattice are found in alloys of aluminum with silver or zinc. More usually, the size difference of the solute and solvent atoms will produce some distortions, and the atoms in and around the zone will be displaced from their positions in the random solid solution. It is characteristic of a cluster or zone, as distinct from a coherent precipitate, that the atom sites do not approximate to a new regular arrangement under these circumstances; the displacements may be regarded as large elastic strains and vary from the center to the outside of the zone.

Direct images of Guinier-Preston zones has been obtained using electron microscopy and x-ray diffraction. The diffraction technique
has one major drawback, that is, it requires a minimum concentration of the solute atoms (normally 2%). On the other hand, electron microscopy can detect these zones in alloys having solute concentration as low as 0.05%.