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TU DRESDEN

ADVANCED PRACTICAL COURSE

LAB REPORT

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# Nuclear Magnetic Resonance

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# 1 Introduction

## 1.1 Motivation

*Nuclear Magnetic Resonance* is a physical phenomenon that can be observed while placing an ensemble of nuclei into a static magnetic field and stimulate it with a high-frequent alternating field. A necessary condition for this effect is that the atoms of the sample have a *nuclear spin* different from zero. It is the central concept that is used for *NMR-Spectroscopy*, a standard methodology for the investigation of the structure and interaction of complex molecules and solid state bodies by measuring local magnetic fields, and the *magnetic resonance tomography* which is an imaging technique used in clinical diagnostics for describing the morphologic and physiologic build-up of tissues and organs. For all of those applications some important parameters of particular physical compensation-processes, the so called *relaxation times*  $T_1$  and  $T_2$  need to be quantified. In the following experiment exactly those material-characteristic observables are determined for an ensemble of  $^{57}\text{Fe}$ -nuclei. But at first some basic knowledge.

## 1.2 Nuclear Zeeman-Effect

Every quantum mechanic angular momentum - especially every spin - is correlated with a magnetic moment  $\mu$ . The proportionality factor is called the *gyromagnetic ratio*  $\gamma$ . So the intrinsic magnetic momentum matching to the nuclear spin  $\vec{I}$  considered in the experiment is given by:

$$\vec{\mu} = \gamma \vec{I} \quad (1)$$

$$\gamma = g_I \frac{\mu_N}{\hbar} \stackrel{^{57}\text{Fe}}{=} 0.8661 \cdot 10^7 \text{ T}^{-1} \text{ s}^{-1} \quad (2)$$

Where  $\mu_N$  is the *nuclear magneton* and  $g_I$  is the Landé-factor which both are core-specific parameters. If those magnetic moments are placed in a static magnetic field  $\vec{B} = (0, 0, B)^T$  then the Hamiltonian of the system and its Eigenvalues to the Eigenstates of the spin-operator  $|I, m_I\rangle$  are given by:

$$\mathcal{H} = -\vec{\mu} \vec{B} \stackrel{(1)}{=} -\gamma \mathcal{I}_z B \quad (3)$$

$$\langle \mathcal{H} \rangle = \langle I, m_I | \mathcal{H} | I, m_I \rangle = -\gamma B \langle I, m_I | \mathcal{I}_z | I, m_I \rangle = -\gamma B \hbar m \equiv E_m \quad (4)$$

Where the Eigenvalues of the Spin-operator for given spin-quantumnumber  $I$  and magnetic quantum number  $m_I = -I, \dots, I$  were used. So the outer magnetic field annuls the  $2I + 1$ -fold degeneration of the energystates. The nuclear spin-quantum-number of  $^{57}\text{Fe}$  is  $I = 1/2$  so there are two additional energystates with a energydifference:

$$\Delta E = \hbar \gamma B = \hbar \omega_L \quad (5)$$

As equation (5) suggests, there may occur optical transitions between the terms which lead to an emission of photons with the angular frequency  $\omega_L$ . One finds that this frequency is equivalent to the *Larmor-Frequency* that describes the precession of a magnetic moment caused by the torsional moment  $\vec{M} = \vec{\mu} \times \vec{B}$  in a magnetic field. The classical description of this process leads to the same result as quantum mechanics do. If we consider the ensemble of  $N$  nuclei as canonical, then the number of spins in the state  $|s, m\rangle$  in the thermodynamic equilibrium is given by the Boltzman-statistics:

$$N(s, m) = N \cdot \frac{e^{-\frac{E_m}{k_B T}}}{Z} = N \cdot \frac{e^{-\frac{\hbar \gamma B m}{k_B T}}}{Z} \quad (6)$$

Where  $Z = \text{const.}$  is the canonical partition function and  $T$  is the absolute temperature of the environment. This implies that the spins prefer to be polarised not uniformly but in the direction

of the B-field so this leads to a mean magnetic moment  $\langle \vec{\mu} \rangle \neq 0$ . This leads to an observable macroscopic magnetisation in the volume  $V$ :

$$\vec{M} = \frac{d\vec{\mu}}{dV} \cong \frac{N}{V} \langle \vec{\mu} \rangle \neq 0 \quad (7)$$

These changes in magnetisation are used to induce voltages that can be measured.

## 2 Experimental procedure

### 2.1 Preparation of a high frequency resonant circuit

First one has to prepare a copper coil with a diameter big enough to hold an iron powder assay. After the coil is wrapped one has to sold it onto the contacts of a stick, which provides a mechanism to tune the measured frequency to find the resonance frequency. While solding one has to be careful to make sure, that one do not take to much tin, because the resistances of tin and copper are different. In the same way the soldered point should connect the contact and the coil directly, otherwise this could influence the results while tuning to the resonance frequency.

### 2.2

### 2.3

## 3 Data Analysis

## 4 Discussion and conclusions

## References

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