Continuous

Nuclear Magnetic Resonance

Physics 122 Lab



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Textbooks

NUCLEAR MAGNETIC RESONANCE

Introduction:

For the purposes of this experiment, the nuclei of a collection of atoms may be regarded in lowest approximation as an assembly of non-interacting magnetic moments in a magnetic field. The Hamiltonian for a single nuclear magnetic moment in a magnetic field is

$$H = g\mu_N \overline{I} \bullet \overline{H} \tag{1}$$

where \overline{I} is the angular momentum of the nucleus; \overline{H} , the magnetic field present at the nucleus; g, the nuclear Land'e g-factor, and μ_N , the nuclear magneton.

$$\mu_N = \frac{e\hbar}{2mc} \tag{2}$$

This equation is in cgs units, as are all the subsequent equations in the writeup of this experiment. Note that when people refer to the "magnetic moment in nuclear magnetons", they mean the product, gI.

Assuming a magnetic field present along the z-axis of magnitude H_z , the expectation values of H are

$$Energy = \langle H \rangle = g\mu_N m_I H_z \tag{3}$$

The energy levels are then of the following form.



For the case I=1/2. The transition indicated by the arrow has the frequency

$$v_0=g\mu_NH_Z/h$$

(4)

for all values of I since the selection rule is $m_I=1$. For the proton this transition is at 4.2576 kHz / gauss. Note that this energy separation can also be described using angular frequency $\omega_0 = 2\pi v_0$.

(**A**)

Transitions from one m_I level to another are induced by an rf magnetic field at frequency v (angular frequency ω) oriented along an axis perpendicular to z, say along the x-axis. The perturbation Hamiltonian is

$$\overline{H}' = g\mu_N \overline{I}\overline{H}_{rf}$$
(5)

$$H_{rf} = H_{rf} \cos \omega t \hat{x} \tag{6}$$

 \hat{x} being a unit vector along the x axis.

Solution for the transition probability predicted by (5) is given by

$$P = \frac{\left(\omega_{o}H_{rf}/H_{Z}\right)^{2}}{\left(\omega_{o}-\omega\right)^{2} + \left(\omega_{o}H_{rf}/H_{Z}\right)^{2}}\sin^{2}\sqrt{\left(\omega_{o}-\omega\right)^{2} + \left(\omega_{o}H_{rf}/H_{Z}\right)^{2}}\frac{t}{2}$$
(7)

The sin^2 factor accounts for the fact that, once the nuclei have made the transition, they are in a position to again interact with the rf magnetic field and return to their original energy levels; and then of course the nuclei can again interact and repeat the process, however, by a variety of interactions, so that their oscillations get out of phase with one another. The time required for this to occur is the "transverse relaxation time", T₂. After a time T₂ the *sin*² factor becomes its average, which is ½. Equation (7) then becomes,

$$P = \frac{1}{2} \frac{\left(\omega_0 H_{rf} / H_z\right)^2}{\left(\omega_0 - \omega\right)^2 + \left(\omega_0 H_{rf} / H_z\right)^2}$$
(8)

(ref: N.F. Ramsey, Molecular Beams, p. 146-150).

Equation (7) does not take T_2 into account. When that is done, the sin² oscillations damp out exponentially with time constant T_2 . These damped oscillations are often referred to as the "wiggles".

When transitions occur, power can be emitted or absorbed, depending upon the relative populations of the two levels. But at any temperature T, the population ratio is given by the Boltzman factor:

$$\frac{N(\frac{1}{2})}{N(-\frac{1}{2})} = e^{-hv_0/kT} \equiv 1 - \frac{hv_0}{kT}$$
(9)

A very slight absorption is possible at room temperature and several thousand gauss.

When resonance occurs, the thermal equilibrium is disturbed and the populations are no longer given by (9). Then when the nuclei are out of resonance, they relax to thermal equilibrium in a time T_1 , which is called the "longitudinal relaxation time". One must not attempt to repeat the experiment in a time less than T_1 , otherwise one will not see a signal. This is important in the experiment, since the static magnetic field H_z is modulated at frequency v_{mod} ; during each cycle of the modulation, the experiment is repeated. Thus it is important that

$$\frac{1}{v_{\rm mod}} >> T_l$$

General Experiment

It is the purpose of this experiment to observe the phenomenon of nuclear magnetic resonance (or "NMR") and to gain some familiarity with the instrumentation used for its observation.

The NMR is observed by measuring the corresponding change in quality- factor, or "Q", of a coil containing a sample. This coil together with a capacitor constitute the tank circuit of a radio frequency oscillator which operates at very low oscillation levels and hence in the extreme non-linear portion of its characteristic. Small changes in Q of the tank circuit result in relatively large changes in oscillation level. It is these changes in oscillation level, after demodulation, that we observe as the NMR signal.

The sample to be tested is placed between the poles of a Varian DC magnet capable of up to a 9 kiloGauss field. On the inside of the large dc magnet poles are placed coils that are driven at an audio frequency to provide a slowly sweeping AC field at a small fraction of the DC field strength. As the combination of the DC and varying AC field sweep through the NMR range, a signal is produced on the oscilloscope screen. The block diagram below shows the basic system components. A large DC power supply capable of 1500V at up to 1.5 amps supplies regulated power to the large magnet. An audio oscillator provides a continuous audio sine wave to a power amplifier that provides AC power to the relatively small coils between the large magnet poles, and to a phase shifter to synchronize a horizontal sweep signal for the oscilloscopes horizontal deflection amplifier. The amplitude modulated signal from the RF oscillator is fed to a frequency counter to provide an accurate frequency measurement and to an AM detector to provide a demodulated signal for the oscilloscopes vertical deflection amplifier.





Function of components:

a. RF Oscillator -

The oscillator in use is a conventional marginal oscillator. The basic circuit is that of Proffitt and Gardiner - "Instructional NMR Instrument' Journ. Chem. Educ. 4.1 152 (1966)

This is a standard rf oscillator at the frequency ν which furnishes a linearly polarized rf magnetic field perpendicular to H₀ and of magnitude H_{rf} cos ω t Note that there are now three magnetic fields present: the large DC field H₀, plus a time dependent modulation H_{mod}cos ω _{mod}t both along \hat{z} , as well as the rf field perpendicular to these two.

The oscillator can be modeled as a series RLC circuit, where L is the inductance of the rf coil around the sample and C is the capacitance of the tuning capacitors in the oscillator. R is a series resistance present to account for all power losses in the RLC circuit including the effect of the nuclear magnetic resonance. As any beginning physics text shows, the impedance of the circuit at resonance is equal to R and the "quality factor", Q, (defined as the energy stored in the circuit divided by the energy lost per cycle) is $\omega L/R$, where ω is 2π times the oscillation frequency. As a consequence, at resonance, the impedance of the circuit is proportional to L/Q. (see for example, ED. M Purcell, Electricity and Magnetism, 1969, p. 278.)

When connected as a oscillator, the voltage level of the oscillations is proportional to the impedance of the circuit, and so to L/Q. But Q represents effects of a number of losses:

$$\frac{1}{Q} = \frac{1}{Q_0} + \frac{1}{Q_{dielectric}} + \frac{1}{Q_{NMR}}$$
(11)

Here Q_0 describes the losses in the coil itself, $Q_{dielectric}$ describes the losses in the dielectric sample holder and Q_{NMR} the losses in the sample itself. In order for the NMR to be observable, Q_0 and $Q_{dielectric}$ must be very large, so that their reciprocals will be negligible.

When NMR occurs, a macroscopic magnetic moment, M, appears in the sample, proportional to H_{rf} .

$$M_{x} = 2H_{rf} \left(\chi' \cos \omega t + \chi'' \sin \omega t \right)$$

The quantities χ' and χ'' are called the "rf susceptibilities". One may show that χ'' is proportional to $1/Q_{NMR}$:.

$$\frac{1}{Q_{NMR}} = \frac{4\pi\chi''}{1 + 4\pi\chi'} = 4\pi\chi''$$
(12)

As the field is swept through resonance at the modulation frequency v_{mod} , the oscillation level at the rf oscillator per half cycle is like this:



The envelope traces out the resonance absorption curve of the sample determined by χ ". Since the abscissa is H = H₀ + H_{rf}cos ω t, the envelope is χ " (H).

As can be seen in the oscilloscope photos below, the actual waveforms look significantly different from the drawing above.



The small change in the magnetic field causes only a small change in the coils value and a correspondingly small change in the actual signal amplitude. The change is significant, however, and is enough to detect and display. The photos show the 100 Hz magnetic sweep on the lower trace and the rf signal on the upper traces. The left photo shows the rf signal at over 1.0 volt p/p amplitude with only tiny bumps in the signal discernable. Offsetting the trace and zooming in on the top shows a usable signal of about 5 mV in the right photo.

NMR Oscillator- Functions of Controls:



b) <u>Detector</u> - This is just a rectifier with a time constant long compared to l/v_o . and short compared to $1/v_{mod}$. An especially simple form might be

Oscillation Frequency Adjust

100 pf



Oscillation level

During the negative half cycle the diode is non-conducting. It conducts during the positive half cycle of the input voltage to keep the capacitor C charged. We require

$$\omega >> \frac{1}{RC} >> \omega_{\rm mod} \tag{13}$$

The output at the detector, over one half cycle of the modulating frequency, is,



(assuming that the conditions of equation (13) are satisfied). This is the vertical signal to the oscilloscope.

Oscilloscope – We can use two ways of generating the horizontal drive for observation

of the signal.

A) Horizontal drive synchronous with modulation.



Here only the amplifier in the Time Base is used to get the sweep signal to the horizontal deflection plates using, Display mode of Amplifier, and Triggering mode of External Source. During one full cycle of the AC portion of the magnetic field, the CRT beam will be swept across the CRT face in one direction and then back across in the other direction. Since the waveform peaks will usually not align this way, a phase shifter circuit is used between the audio sweep signal and scope amplifier to allow adjustment of the opposite direction sweep timing to align the peaks.



The **Phase Shifter** box has both a level control and a phase control. The level control is adjusted to obtain a full horizontal scan on the CRT and the phase control is adjusted to align the opposite scan waveforms.

INTENSITY

RN



The result of non-aligned sweeps is shown on the left, aligned on the right and an actual correctly aligned scope waveform below.



B) Internal time base triggered by either modulation or the input signal.



The horizontal sweep can be generated by triggering the time base internally (Display Mode-Time Base, Triggering Source-Internal) using the modulation signal from the Vertical Amplifier. This technique is helpful if an accurately linear scale is needed on the scope screen, such as measuring timing of the wiggles.

1/vmod

DC Magnet and Supply

The Varian Regulated DC Magnet and Power Supply are shown below. This supply can be adjusted to hold the large magnet at a very stable gauss level from near zero to about 9 kiloGauss.



Frequency Counter



A Hewlett Packard 5328A Frequency Counter monitors the frequency of the RF oscillator to determine the actual frequency of an observed NMR signal.

Gauss Meter



The AlphaLabs DC Magnetometer uses a Hall effect probe that is placed between the two magnet poles to measure the DC magnetic flux.

Audio Signal Generator



The Tektronix SG 502 Oscillator plugs into a Tektronix TM Power Supply Mainframe and feeds audio sine waves to the audio power amplifier.

Audio Power Amplifier



The power amplifier driving the AC magnet coils is an Altec Lansing stereo amplifier modified for single channel use with input and monitor connections on the front panel.

Basic Procedure

Set up the experimental components according to the block diagram on page 5. Have your instructor check the set-up then try to find a proton resonance. Pure distilled water will not yield an easily observable signal, you will need some paramagnetic ion relaxer. If $CuSO_4$ is used the solution should be a pale blue.

After obtaining a signal, vary the RF oscillation level, audio power level, probe position in the magnet and the paramagnetic ion concentration to obtain the best signal to noise ratio. Now do a field calibration over the range available to you.

Samples containing protons in H_2O , glycerol, polyethylene and paraffin as well as several other nuclei are provided. All can be observed using this apparatus. Their gyromagnetic ratios and isotopic abundances are listed below (1Tesla =10 kiloGauss).

Isotope	NMR Freq MHz/Tesla	Natural Abundance	Spin
¹ H	42.5759	99.9844%	1/2
⁷ Li	16.546	92.57%	3/2
¹¹ B	13.660	81.17%	3/2
¹⁹ F	40.0541	100%	1/2
²⁷ AI	11.094	100%	5/2
⁵¹ V	11.193	100%	7/2

The nuclei below are more difficult to observe; signal to noise ratio is < 5.

Isotope	NMR Freq MHz/Tesla	Natural Abundance	Spin
²⁰³ TI	24.332	29.5%	1/2
²⁰⁵ TI	24.570	70.5%	1/2
³¹ P	17.235	100%	1/2
¹¹⁷ Sn	15.186	7.61%	1/2
¹¹⁹ Sn	15.869	8.58%	1/2
⁸⁷ Rb	13.932	27.2%	3/2

Use care in handling the fluorine, boron, vanadium, phosphorous, and thallium samples; the triflouracetic and phosphoric acids are highly corrosive and thallium is <u>extremely</u> poisonous.

 Using the Hall effect gauss meter, measure the g factor for the proton in H₂O. This is best done by plotting the nuclear resonance frequency as a function of magnetic field over a substantial range, say 5 MHz - 30 MHz. The least-squares slope of the Line will be the g factor. You will be limited to an accuracy of perhaps 0.1% by the accuracy of the gauss meter. However, the magnet power supply is stable to at least one part in 10⁵; thus it should be possible to measure ratios of NMR frequencies for different elements to this precision in a constant magnetic field, by tuning the oscillator frequency to the different resonances.

- 2. Using the known g factor of the proton (2 x 2.79270) measure the nuclear resonance frequencies for several of the other isotopes listed, and calculate their g factors. Compare with published values. (See CRC Handbook of Chemistry and Physics and note that the g factor times the spin is the "nuclear moment".)
- 3. (a) With the scope horizontal sweep set up for synchronous drive (vert amp rather than time base), measure the width of the proton resonance in water or glycerol in gauss. In this configuration, the horizontal deflection of the oscilloscope is proportional to the magnetic field. The easiest way to calibrate the x-axis is to measure the static field (or better, the NNM frequency) at those points where the nuclear magnetic resonance disappears from the oscilloscope screen. For liquids, the intrinsic NMR line width is very narrow, and virtually all of the observed line width is due to nonuniformities in the applied magnetic field over different parts of the sample.
 - (b) An alternate method of determining the line width is to measure T₂, the transverse relaxation time or "ring down" time characterizing the "wiggles" which follow rapid transit through the nuclear resonance. The nuclear absorption is described by

$$\chi''(\omega) = \frac{Const.}{1 + (\omega_0 - \omega)^2 T_2^2}, \qquad \omega_0 = \gamma H \qquad \gamma = \frac{g\mu_N}{\hbar}$$

The envelope of the "wiggles" is given by the Fourier Transform of $\chi^{"}(\omega)$

$$\chi''(t) = \int_{-\infty}^{\infty} e^{i\omega t} \chi''(\omega) d\omega \propto e^{-t/T_2}$$

The full width at half maximum of $\chi''(\omega)$ is 2/T₂ or, in gauss, $\Delta H = \frac{2}{\gamma T_2}$

Find a sample which displays wiggles when the field sweep is adjusted for fast passage (how fast?), through the resonance, and with the scope set up for internal time base, trigger the oscilloscope internally from the field sweep waveform (vertical channel) using its own internal time base. Then plot the log of the wiggle. amplitude versus time to obtain T_2 . If possible, compare with the line width H obtained in (a).

4. The fluorine resonance may also be observed in teflon, a solid white plastic (CF₂). Likewise the proton resonance can be seen in solid polyethylene and paraffm. NMR line widths in solids are typically much larger than in liquids ("dipolar broadening"). The dipolar broadening is averaged out in liquids by the tumbling motion of the atoms, which is much more rapid than the nuclear precession rate. See if you can observe this effect in the solid samples provided.

Running the NMR Experiment

<u>The Marginal oscillator:</u> (3 MHz < v_0 < 30 MHz)

(1) RF probes:

You have the choice of a **high-frequency** (larger wire, fewer turns) probe capable of providing RF oscillation frequencies higher than 10 MHz, and a **low-frequency** (smaller wire, more turns) probe capable of providing RF oscillation frequencies less than 12 MHz. Since the maximum DC magnetic field is limited to 8500 gauss, you need both probes to measure NMR signals from ¹H and ¹⁹F (the **high-frequency** probe) and from ⁷Li, ¹¹B, and ³¹P (the **low-frequency** probe).





- (2) After attaching the selected RF probe to the oscillator, Insert the desired sample into the probe and position the assembly so that the sample is at the center of the magnet.
- (3) Turn on the rack instruments with the switch on the power strip. This will also supply 12.6 volts to the RF Oscillator tube filaments.
- (4) Turn on the power supply for oscillator plate voltage;
- (5) Set the sensitivity for the "Base Level" to "Hi"
- (6) Set the dial for the "Base Level" to 6 o'clock;



(7) Set the capacitor (frequency) dial on the side of the oscillator box for the desirable frequency v.

The frequency, v. (in unit of MHz) is read with the HP 5328A Frequency Counter set up as shown below.



FUNCTION – FREQ A FREQ RESOLUTION – 10 KHz SAMPLE RATE - mid range DELAY - OFF (fully counter clockwise) LEVEL A - PRESET (fully counter clockwise) SLOPE - + ATTEN - 1 AC DC - AC COM - SEP

DC magnetic field (50 <Bd, < 8500 gauss)

Use a piece of masking tape to hold the gauss meter probe on the side of the RF probe,



or a magnet pole, so that the magnetic field \mathbf{B}_{dc} , at the sample can be read as the magnet power is turned on and adjusted. The gauss meter is battery powered and must be turned on separately from the instrument rack power. Set the gauss meter to the 19,999 position. Note: residual magnetism in the magnets iron poles will cause a reading on the gauss meter with magnet power still off.

-Gauss Meter Probe

The DC magnet power supply is a serious piece of vacuum tube electronics as can be seen



in the picture below. Care must be taken in its operation to avoid damage to the system. The tubes will not regulate the magnet current properly if the filaments are not at operating temperature, so they must be allowed to warm up prior to applying high voltage, and the *high voltage must be turned* off before turning the filament supplies off at shut down to avoid destroying tubes or other components. (yes, I have seen this happen)

Magnet Supply



To turn on the magnet:

- 1. Turn on the water. (The magnet is water cooled)
- 2. Check to make sure the autotransformer Voltage Control, sometimes referred to by the trade names variac or powerstat, is in the off position, fully counter clockwise.

3. Turn on the Filament power switch (green light at left will turn on), wait at least one minute for the vacuum tube filaments to warm up. (*This power supply uses vacuum tube technology and the tube filaments must be warm before the tubes will operate correctly. In high power circuitry like this the tubes will not control properly when the filaments are not at operating temperature, but they can still conduct power and cause damage.*)

4. With the Voltage Control in the off position (counter clockwise), turn on the High Voltage switch (Red Light at right will turn on). Turn up the Voltage Control (clockwise increases power) until the left hand, Regulator Range meter reads in the darkest band.



This indicates the correct operating voltage being dropped across the series pass regulator tubes. To the left of the darkest band you will lose magnet current regulation. To the right of the darkest band is another dark band in which regulation will still occur, but this is not an optimum regulating point. To the far right beyond the dark bands the pass tubes will be operating at too high a power level and might be damaged. *IN NO CASE SHOULD YOU LET THIS METER GO OFF SCALE ON THE RIGHT*

- 5. Adjust the coarse and fine controls to obtain the desired magnet gauss value, watch the voltage meter and adjust the autotransformer as needed to see that it stays in the heaviest black band region
- 6. Adjust the "Voltage Control" slowly up until the indicator of the "Regulation Range Meter" moves from the far left to the first black band on. the left side of the meter; (Note: *this is, the range of -regulation that you should operate the power* supply)
- 7. Adjust the "Coarse Current" first and the "Fine Current" next to set the magnetic field B_{dc} , to the desired value as determined by the Table on page 12 for $g\mu_N/h$ and the following relation

$$v_0 = \left(g\mu_N/h\right)B_{dc}$$

You must continue to adjust the "Voltage Control" such that the regulation range indicator remains in the first black band on the left.

AC magnetic field (B_{ac}): (0 < B_{ac} < 25 Gauss)

 Use a combination of the sine wave generator (Tektronix SG502) and audio power amplifier to drive the pair of AC coils to produce an oscillating magnetic field B_{ac} as follows;



2) Set the signal generator with Frequency-. 40 to 270 Hz

Amplitude: 0.5 volts RMS



- (3) Connect the output of the signal generator to the Audio In connector of the Power Amplifier.
- (4) Connect the output at the rear of the audio power amplifier to the AC Magnet coils and the front panel, monitor output, to the input of the phase- shifter.
- (5) Set the Audio Power control at 4 on the power amplifier so that the output voltage is roughly 3.5 volts(Note: *this produces an oscillating magnetic field B_{ac} with peak-to-peak amplitude of about 14 gauss);*
- (6) Connect the NMR voltage signal (the output of the marginal oscillator) to the oscilloscope mainframe (Tektronix 7603) vertical amplifier (AM-6565). Set "VOLTS/DIV" to 1.



(7) Connect the output of the phase shifter to the input of the scopes Time Base.

(8) Note- the phase-shifter is used to add an adjustable phase to the sinusoidal wave from sweeping the AC magnet, so that the output of the phase-shifter is in synchronization with the oscillating magnetic field B_{ac} . You can only adjust the phase when you have already found the NMR signal from the sample (see the Measurement Procedure below); (9) Set the switch on the phase-shifter to "1 kHz";

(10) Adjust "Level" on the phase-shifter until the signal just fills the oscilloscope screen along the x-axis (total of 10 divisions).

Measurement procedures:

(1) Set the RF Oscillator to the desired frequency and Fine tune B_{dc}, by adjusting the "Fine Current" on the DC magnet supply until you observe the NMR signal as shown in the scope waveform on page 11.

(Note- since the screen only displays the signal over a range of 25 gauss or less, you should not dial the "fine Current" too fast or you will miss the signal)

(2) Synchronization of the phase-shifter output and B_{ac}

Once a signal is observed, you will typically see two peaks on the screen that move in opposite directions as you change B_{dc} this indicates that the phase-shifter output is out of sync with the alternating magnetic field B_{ac} ;

Adjust the phase dial until the two peaks overlap on the screen,

Now the phase-shifter output along the x-axis is exactly proportional to B_{ac} ;

(3) Calibration of the x-axis as the magnetic field axis

Since the x-axis is proportional to \mathbf{B}_{ac} , you can determine the proportionality constant ηB (in unit of gauss/division) by adjusting B_{dc} , until the two peaks move together to the far left and then to the far right. The proportionality constant is then obtained by dividing the net change $\Delta \mathbf{B}_{dc}$ by 10 (the total number of divisions along x-axis)

$$\eta_B = \Delta B_{dc} / 10$$

Using this method, you will find that \mathbf{B}_{dc} , ~ 14 gauss when the audio amplifier output is at about 3.5 volts, or 4 on the Audio Power control knob.

(4) Calibration of the x-axis as the frequency axis"

You may convert the x-axis to the frequency axis with a proportionality constant

 $\eta_{\rm RF}\cdots u\sin g\cdots \eta_{\rm RF}(g\mu_{\rm N}/h)\eta_{\rm B}$

<u>Determination of the relation</u> $V_0 = (g\mu_N / h)B_{dc}$ and $g\mu_N / h$ for ¹H, ¹⁹F, ⁷Li ^{II}B, ³¹P

- (1) Adjust B_{dc} to move the two NMR signal peaks to the middle of the screen and record B_{dc} and v_0
- (2) Repeat the procedure for a series of v_0 , and B_{dc} ;

Measure the Full-Width-at-Half-Maximum (FWHM) of the NMR signal peaks for all five **nuclei** on the screen (in unit of division) and use

 $\eta_{RF} = (g\mu_N/h)\eta_B$ to express the result in terms of frequency.

Turning off the system:

- (1) First turn down "Voltage Control" (the heavy variac) on the DC magnet power supply to zero;
- (2) Then turn off "High Voltage" on the DC magnet power supply;
- (3) Turn off "Filament" on the DC magnet power supply,
- (4) Turn off the power strip on the floor supplying power to the instrument rack
- (5) Turn off the computer;
- (6) Turn off the cooling water to the DC magnet.

Computer-aided measurement of NMR signals with a LabVIEW program:

This lab is equipped with a computer-aided data acquisition system based on the LabVIEW-6i program (National Instruments) on a PC. The program, 'NMR_2002.vi", is placed on the C Drive in the folder "Physicsl22Lab_folder". Such a program enables one to perform

(i) **signal averaging** so that weak NMR signals from ¹⁹F, ⁷Li, ^{II}BI ³¹P or even other nuclei can be detected with much better signal-to-noise ratio;

(ii) **digital signal processing** so that the segments of the NMR signal when the magnetic field is ramped up or down can be displayed and analyzed separately;

- (iii) **save the data** in **text files** so that the results can be analyzed and plotted for laboratory reports.
- (1) Connect the phase-shifter output to "ACH0" of BNC-2090;
- (2) Connect the NMR signal to "ACH2" (ACH! Does not work) of BNC-2090;
- (3) Set the frequency of the function generator to f_{audio} ,= 250Hz
- (4) Open up the program "NMR-2002.vi" on the C Drive in the folder "Physicsl22Lab_folder";
- (5) Set "Sampling Frequency (f_s)":,to "50000"
- (6) Set "Number of data points per waveform (N_s)" to $f_s/_{faudio}$ = 200;
- (7) Set "Number of waveforms to be averaged";
- (8) If you only want to see the data with no intention of saving them, do not press "Save the Data,?" button;

If you do want to save the data, press "Save the data ?";

When the data acquisition is completed, a window will pop up for you to enter the file name and to select where you want the file to go. The data file can be open with MS Excel or KaleidaGraph, both are available on the computer. You may also send the data out through Ethernet from this computer.

(9) To analyze the waveform when the magnetic field is ramped up or down, you can slice the required data from the data file using Excel or KaleidaGraph.

A quick sample of a Proton NMR run in Excel is shown on the next page.



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