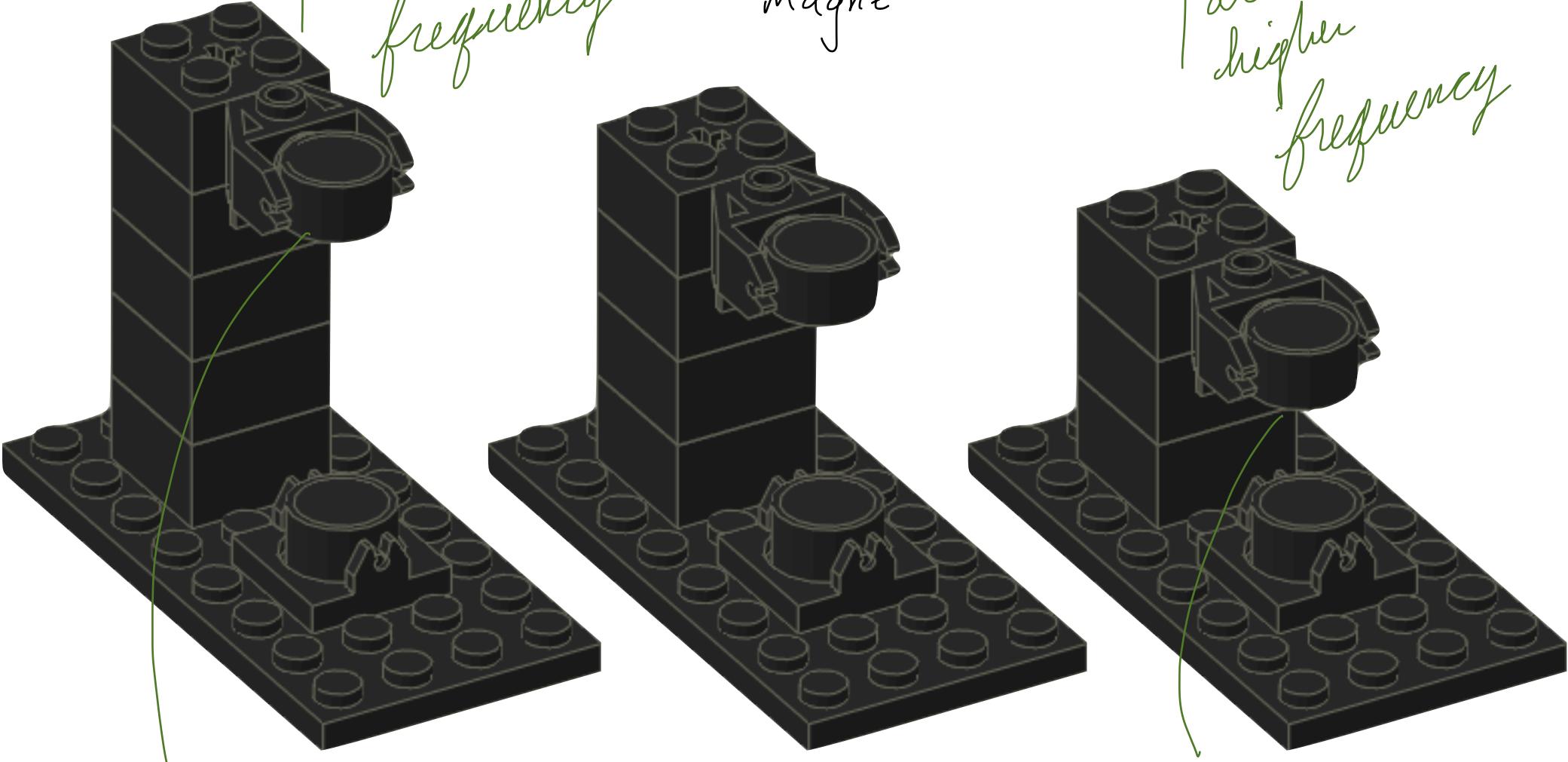


Magnets placed in a magnetic field align with the magnetic field.

When perturbed the magnets will "resonate".

The frequency of the resonance depends on the strength of the magnetic fields



weaker field

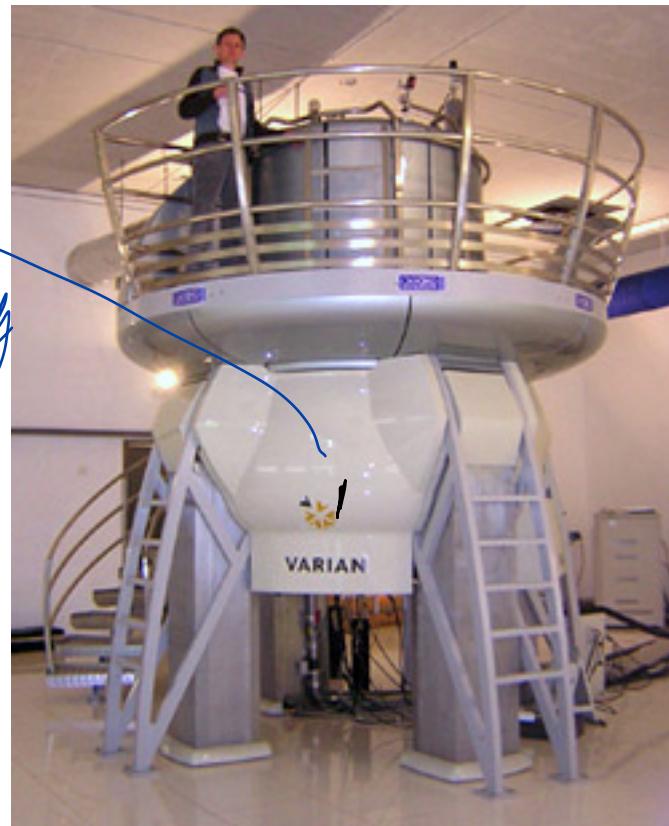
$^3H + ^2H$  not  
magnets

spin  $\frac{1}{2}$

stronger field

$^1H$  is a  
magnet

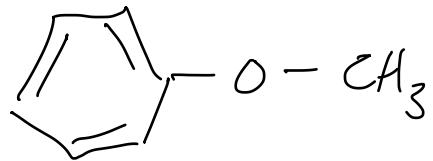
a superconducting  
wire with  $e^-$   
going round  
and round



- 1.4 T magnet will  
cause  $H$  to align +  
resonate at  
 $\sim 60,000,000$  Hz

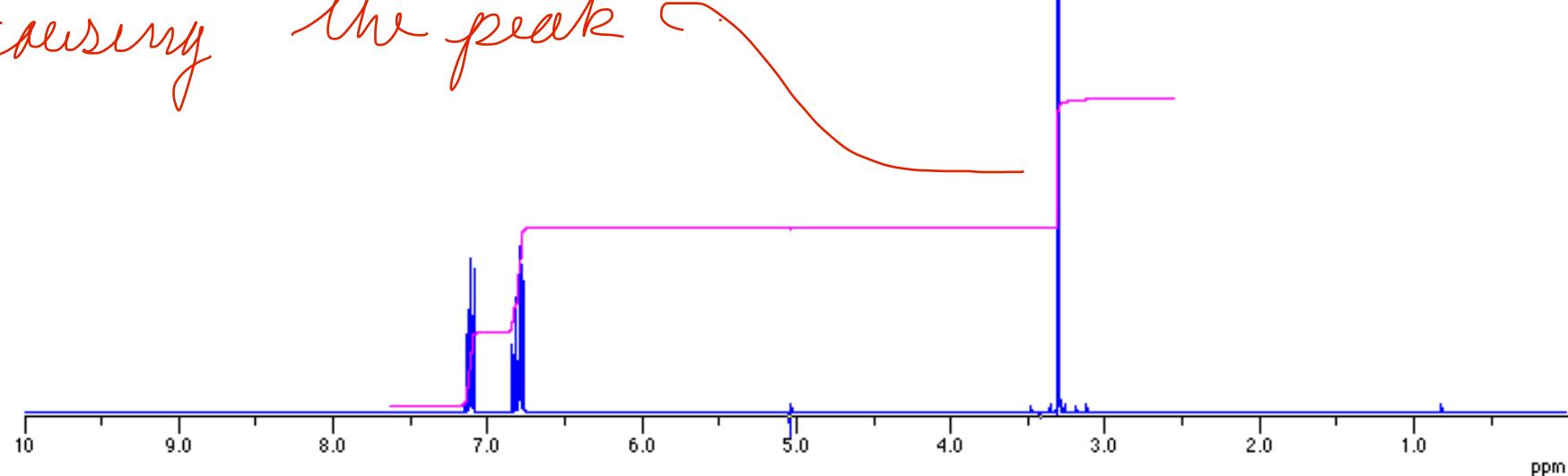
900 MHz, (21.2 T) NMR Magnet at HWB-NMR, Birmingham, UK  
[https://en.wikipedia.org/wiki/Nuclear\\_magnetic\\_resonance#/media/File:HWB-NMR\\_-\\_900MHz\\_-\\_21.2\\_Tesla.jpg](https://en.wikipedia.org/wiki/Nuclear_magnetic_resonance#/media/File:HWB-NMR_-_900MHz_-_21.2_Tesla.jpg)

sample experiences a uniform  
magnetic field



The # of peaks is related to the number of different H's in our molecule.

relative #'s of <sup>1</sup>H atoms causing the peak



The chemical environment

The # of H atoms on neighboring C, N, or O atoms

magnetic field varies based on position



[www.radiologyinfo.org](http://www.radiologyinfo.org)

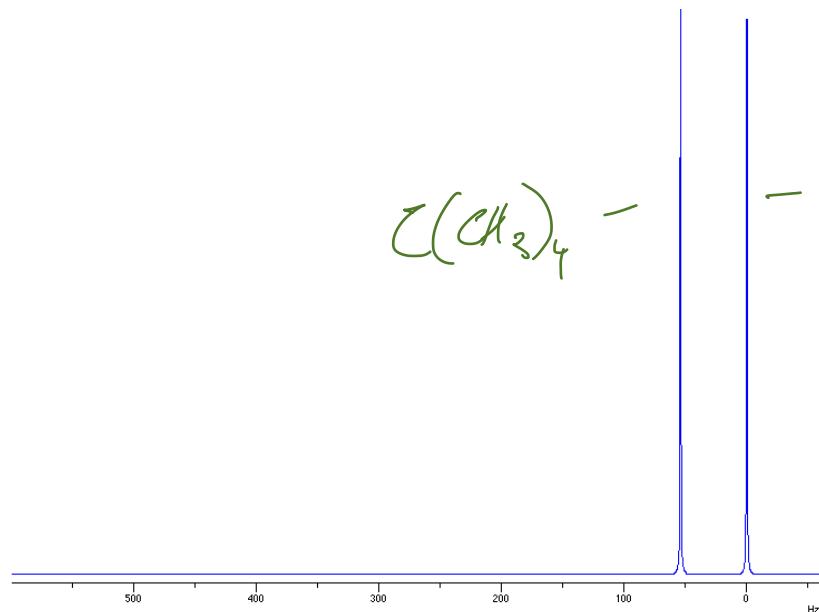


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Magnetic Resonance Imaging  $H_2O$

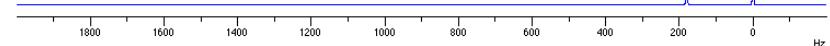
Maps the varying concentrations of water molecules in tissues to create an image.

Resonance frequencies depend on magnetic strength

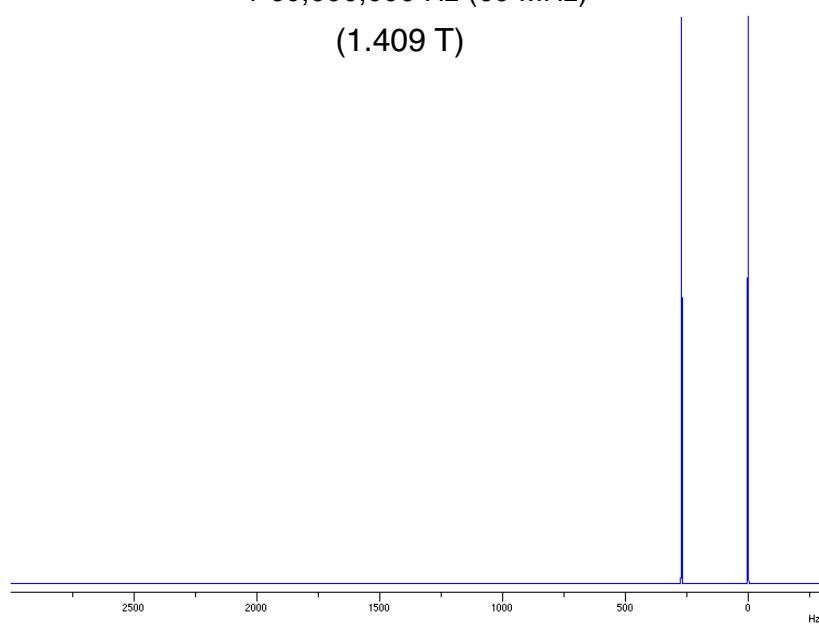


+ 60,000,000 Hz (60 MHz)  
(1.409 T)

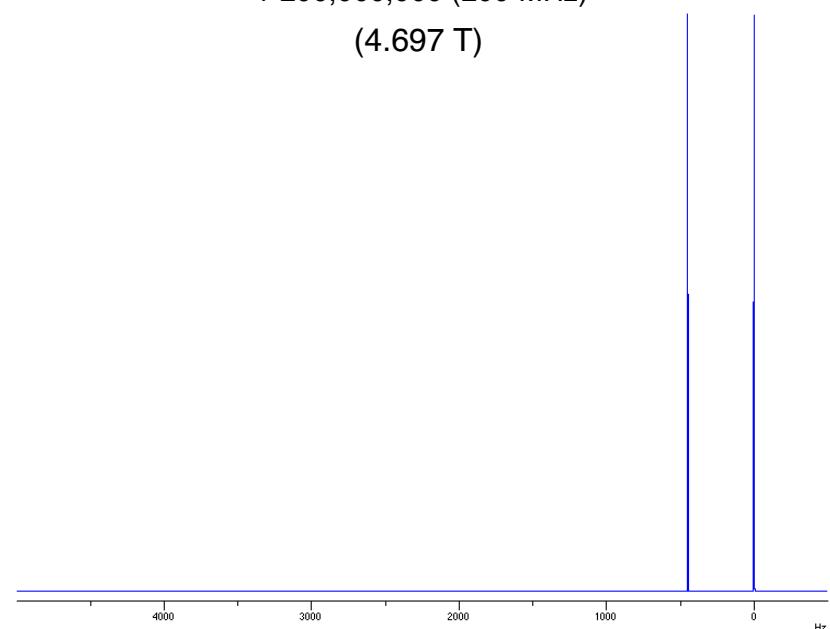
tetramethylsilane  
— TMS



+ 200,000,000 (200 MHz)  
(4.697 T)



+300,000,000 Hz (300 MHz)  
(7.046 T)



+ 500,000,000 Hz (500 MHz)  
(11.743 T)

$$\delta \text{ ppm} = \frac{\nu(\text{peak}) \text{ Hz} - \nu(\text{TMS}) \text{ Hz}}{\nu(\text{TMS}) \text{ MHz}}$$

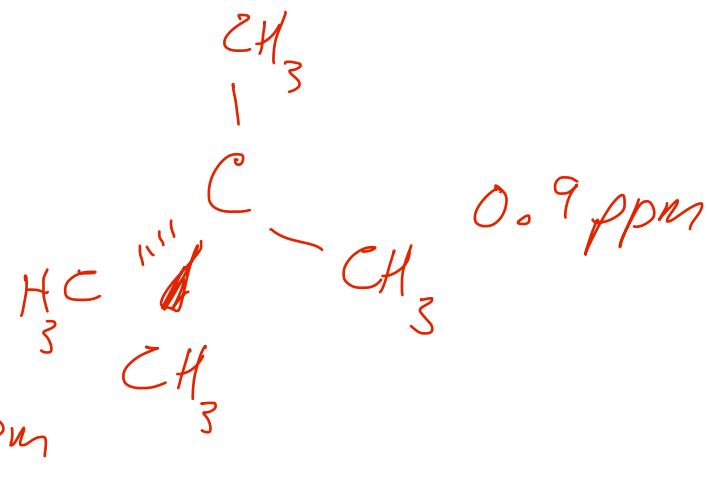
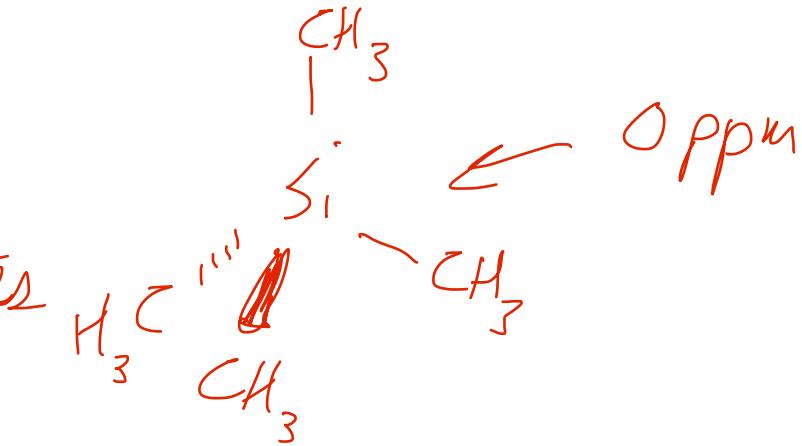
reference molecule

TMS = tetramethyl silane

1.409 T magnet TMS resonates at 60,000,000 Hz

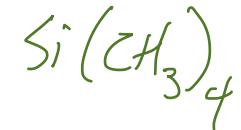
neopentane resonates at 60,000,054 Hz

$$\frac{(60,000,054 - 60,000,000) \text{ Hz}}{60 \text{ MHz}} = 0.9 \text{ ppm}$$





TMS



$$\delta \text{ ppm} = \frac{\nu(\text{peak}) \text{ Hz} - \nu(\text{TMS}) \text{ Hz}}{\nu(\text{TMS}) \text{ MHz}}$$

the  $CH_3$ 's →

here are

"deshield" as

shielded  
by  $Si^-$   
density

compared to

TMS... see a

stronger applied

field, and resonate  
at higher frequency

donated  
from the  
 $Si^-$

high  
freq

low  
freq

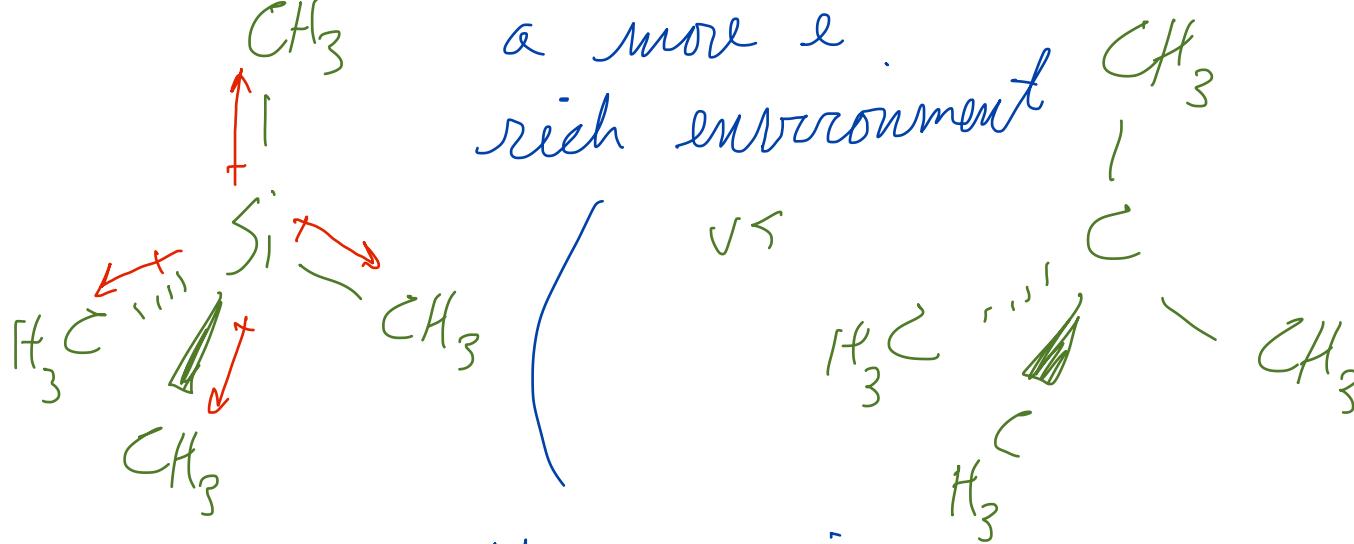
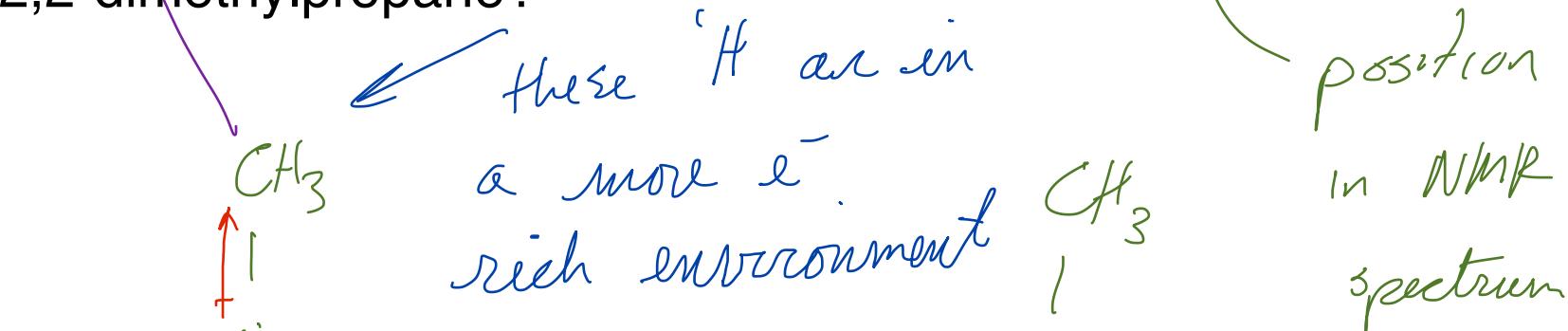
ppm

for historical reasons its "backwards"

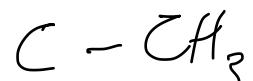
shielding causes these H's to "see" a weaker field  
thus resonate at a lower freq

What gives rise to differences in **chemical shift**?

Why do the H's of tetramethylsilane resonate at a different frequency than 2,2-dimethylpropane?



these moving  
e<sup>-</sup>'s are creating a  
magnetic field which shields  
the H's from the applied field



TMS



$^1\text{H}$ 's resonating over here would be "deshielded" by electro-negative atoms pulling the  $\text{e}^-$ 's away most organic  $\text{H}$ 's are over here

$^1\text{H}$ 's resonating over here would be strongly shielded by  $\text{e}^-$  density donated from neighboring atoms metal - H compounds

