

# Relaxation and Dynamic Nuclear Polarization Transfer DNP

Nuclear Overhauser Effect

Rotating Frame Nuclear Overhauser Effect

- The NMR signal intensity is proportional to  $\gamma^{5/2} = (\gamma \cdot \gamma^{3/2})$
- $\gamma$  concerns **excitation**, determining the **population** difference
- $\gamma^{3/2}$  concerns **detection**, determining the intensity of the nuclear magnetic moment. Furthermore, the detector sensitivity is proportional to frequency (directly proportional to  $\gamma$ ), but the noise that increases with  $\gamma^{1/2}$  must be accounted for.

- An artificial increase of intensity can be obtained by means of polarization transfer from a more sensitive nucleus
- NOE and ROE: **transfer through space**. The “insensitive” nucleus must relax mainly by the dipolar coupling with proton

## Wiener Process



A continuous-time stochastic process  $W(t)$  for  $t \geq 0$  with  $W(0)=0$  and such that the increment  $W(t)-W(s)$  is Gaussian with mean 0 and variance  $t-s$  for any  $0 \leq s < t$ , and increments for nonoverlapping time intervals are independent. **Brownian motion** (i.e., **random walk with random step sizes**) is the most common example of a Wiener process.

Weisstein, Eric W. "Wiener Process." From MathWorld--A Wolfram Web Resource.  
<http://mathworld.wolfram.com/WienerProcess.html>

# $T_1$ for a $I = \frac{1}{2}$ nucleus

—————  $N_{\beta 0}$   $N_{\alpha 0}$  e  $N_{\beta 0}$  are the Boltzmann equilibrium populations

—————  $N_{\alpha 0}$

$\Delta_0 = N_{\alpha 0} - N_{\beta 0}$  population difference at equilibrium

The populations of the two levels can be varied by employing a suited r.f.

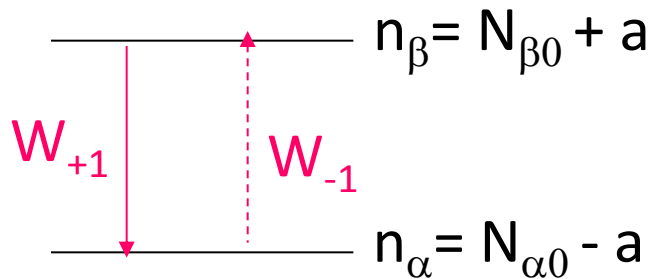
—————  $n_{\beta} = N_{\beta 0} + a$

$n_{\alpha}$  and  $n_{\beta}$  out of equilibrium populations

—————  $n_{\alpha} = N_{\alpha 0} - a$

$\Delta = n_{\alpha} - n_{\beta}$  population difference out of equilibrium

- After the perturbation the systems is reverting to equilibrium
- Considering the systems under a microscopic view:



$$\frac{W_{+1}}{W_{-1}} = \frac{N_{\alpha 0}}{N_{\beta 0}}$$

in order to attain the Boltzmann distributon

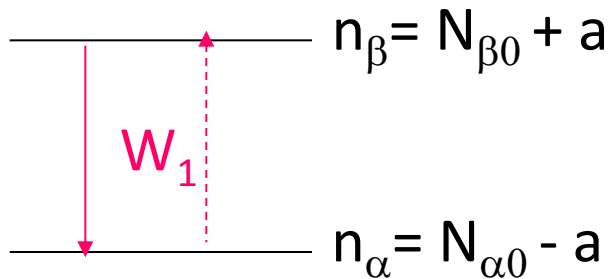
Since the populations' ratio is very close to 1, usually for simplicity sake, it is considered:

$$W_{-1} \approx W_{+1}$$

$W$ : transition probability per unit time and per nucleus

the subscripts +1 e -1 indicate that the transition are single quantum

Therefore the deviation  $a$  is required to account for that at equilibrium the Boltzman distribution is attained



We write the *gain-loss equations*. The first is the gain term.

$$\frac{dn_\beta}{dt} = W_1(-a) - W_1 a = -2aW_1$$

$$\frac{dn_\alpha}{dt} = W_1 a - W_1(-a) = 2aW_1$$

$$\frac{d(n_\alpha - n_\beta)}{dt} = 4aW_1$$

The population difference  $n_\alpha - n_\beta$  is proportional to  $M_z$

$$M_z(t) = N_0(n_\alpha \mu_{z\alpha} + n_\beta \mu_{z\beta}) = \frac{1}{2} N_0 \gamma \hbar (n_\alpha - n_\beta)$$

$$M_0 = \frac{1}{2} N_0 \gamma \hbar (N_{\alpha 0} - N_{\beta 0}) \quad \text{therefore}$$

$$M_z(t) - M_0 = \frac{1}{2} N_0 \gamma \hbar [(n_\alpha - n_\beta) - (N_{\alpha 0} - N_{\beta 0})] = \frac{1}{2} N_0 \gamma \hbar (-2a)$$

recalling

$$\frac{dM_z(t)}{dt} = -\frac{M_z(t) - M_0}{T_1}$$

$$2W_1 = \frac{1}{T_1}$$

# Relaxation Mechanisms

## Yoga realizado por un maestro santiaguense



Esta versión de postura demanda de una fuerza extraordinaria en el cuello, hombros, y glúteos, además de una concentración extrema.

En la provincia de Santiago se realiza esta práctica casi desde que se nace y sin supervisión.



The motion of the spins is decoupled from that of the molecules :



they are like the needles of compasses on ships in the storm



**Bounty Rounds Cape Horn**

<http://cleangreengems.com/bounty/studycenter.htm>

molecular collisions do not change spin orientation

# Spontaneous Emission

- Nel ritorno delle popolazioni di spin alla distribuzione di equilibrio di Boltzmann
- l'emissione spontanea non riveste alcun ruolo perché le frequenze di risonanza degli spin sia elettronici che nucleari sono troppo basse
- essa è proporzionale a  $\nu^3$  ed è trascurabile per frequenze  $< 1$  THz.

# Emissione stimolata

- Sono necessari dei campi magnetici fluttuanti alla frequenza di Larmor
- Nel caso dei solidi le vibrazioni del reticolo (i cosiddetti fononi) mediano lo scambio di energia tra sistema di spin e reticolo. Di qui l'origine del nome spin-reticolo per il rilassamento longitudinale
- In soluzione i campi magnetici fluttuanti sono forniti dalle fluttuazioni rispetto al c.m. della parte anisotropa dei tensori delle varie interazioni magnetiche ed elettriche dovute al moto molecolare.

Il rilassamento sarà più veloce:

- quanto più grande sono i **campi magnetici fluttuanti**
- in EPR sono molto più grandi che in NMR anche in virtù del momento magnetico molto più intenso
- quanto più intense le **componenti di moto** di frequenza adatta nel moto molecolare
- le frequenze in gioco in EPR sono almeno 100 volte maggiori che in NMR
- i  $T_1$  in EPR sono molto più piccoli che in NMR

# Meccanismo dipolare

- Per i nuclei di  $^1\text{H}$  e per nuclei di atomi direttamente legati all'idrogeno il meccanismo di rilassamento più importante è quello dipolare
- Nel caso di moto veloce la velocità di rilassamento di un  $^{13}\text{C}$  vicino ad un  $^1\text{H}$  è:

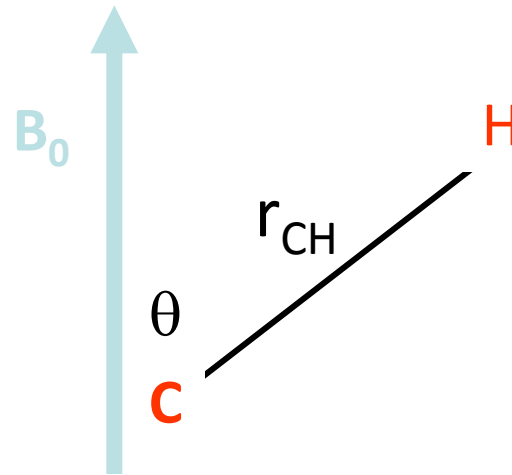
L'importanza del meccanismo dipolare si ricava da misure di NOE

molto rilevante in presenza di e<sup>-</sup> spaiati

$$\frac{1}{T_{1dip}} = \left( \frac{\mu_0}{4\pi} \right)^2 \hbar^2 \gamma_C^2 \gamma_H^2 \frac{1}{r_{C-H}^6} \tau_C$$

# Nuclear Overhauser Effect in un sistema eteronucleare $^1\text{H}^{13}\text{C}$

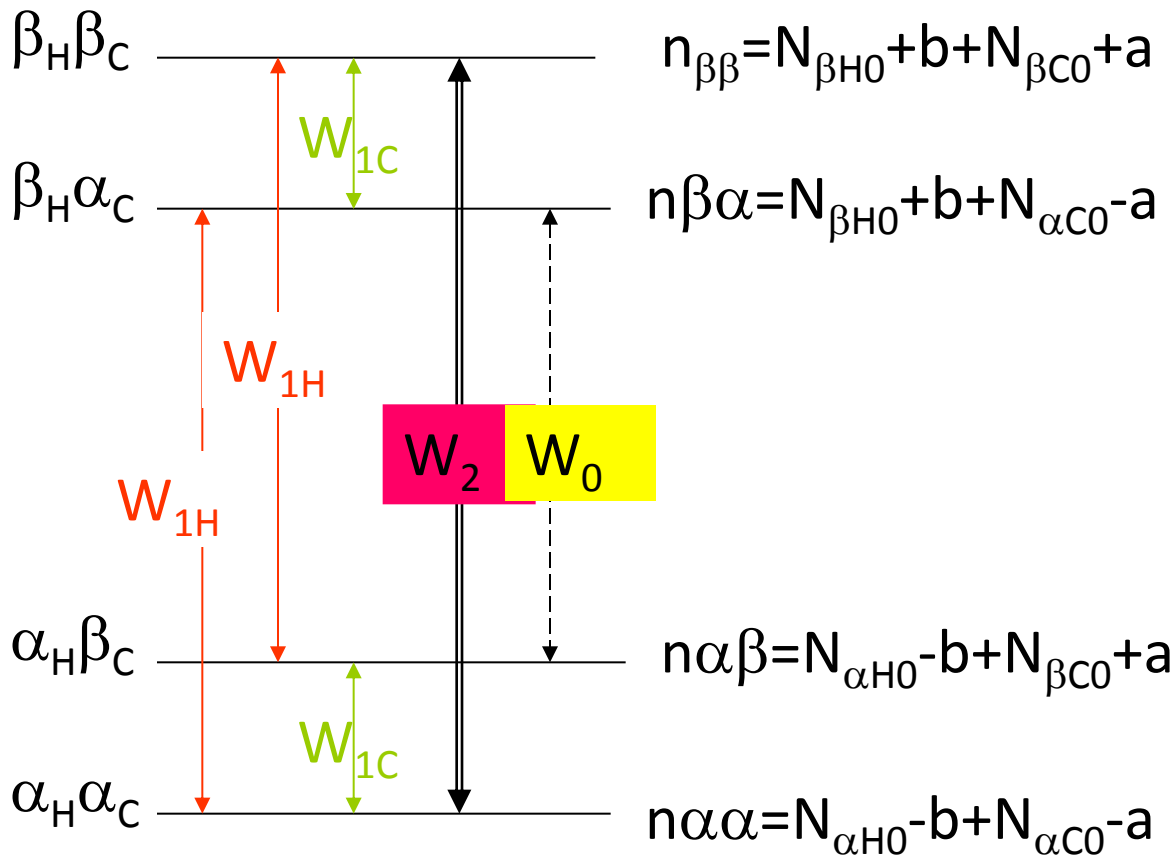
- I nuclei sono vicini nello spazio e si considera esclusivamente l'interazione dipolare attraverso lo spazio.



Il dipolo magnetico del H genera un campo magnetico sul C e viceversa. A causa dei moti molecolari  $\theta$  (e  $r$  nel caso in cui la distanza C-H non sia fissa) variano nel tempo.

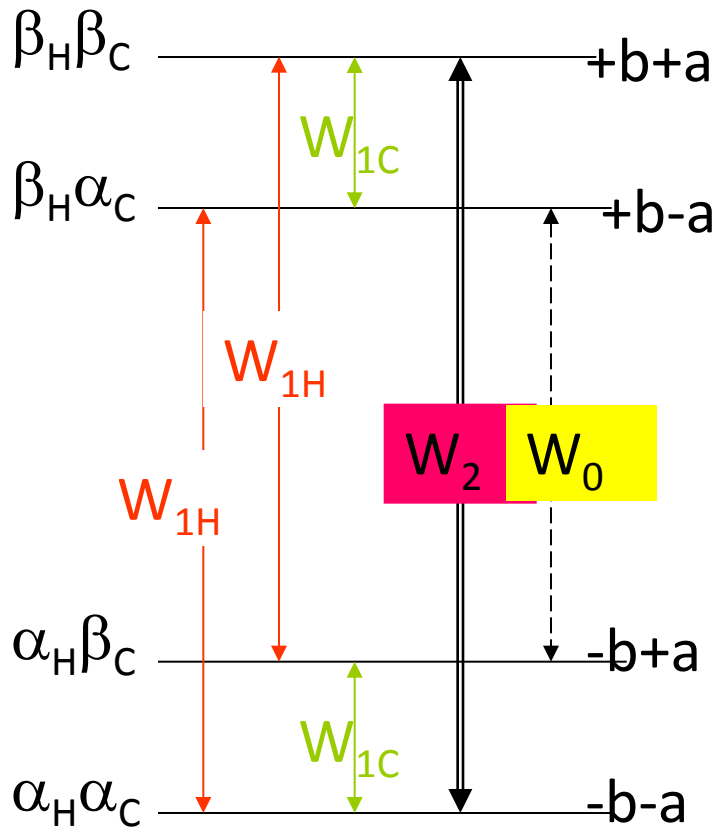
I campi magnetici fluttuanti nel sistema d'assi del laboratorio inducono il mutuo rilassamento dei due nuclei.

# Energy Levels and Populations for the System CH



In the presence of dipolar coupling *cross-relaxation* occurs

# Variation on Time of Populations



$$\frac{dn_{\alpha\beta}}{dt} = W_{1H}b + W_{1c}(-a) + W_0(b-a) - W_{1H}(-b) - W_{1C}a - W_0(-b+a) = 2bW_{1H} - 2aW_{1C} + 2(b-a)W_0$$

$$\frac{dn_{\beta\beta}}{dt} = W_{1H}(-b) + W_{1c}(-a) + W_2(-b-a) - W_{1H}b - W_{1C}a - W_2(a+b) = -2bW_{1H} - 2aW_{1C} - 2(a+b)W_2$$

$$\frac{dn_{\beta\alpha}}{dt} = W_{1H}(-b) + W_{1C}a + W_0(-b+a) - W_{1H}b - W_{1C}(-a) - W_0(b-a)$$



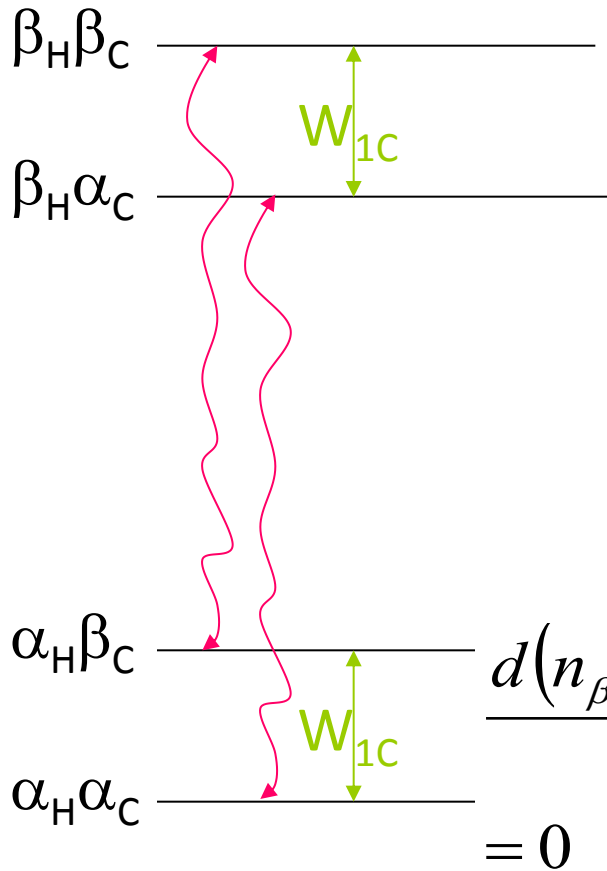
$$\frac{dn_{\beta\beta}}{dt} = -2bW_{1H} - 2aW_{1C} - 2(a+b)W_2$$

$$\frac{dn_{\beta\alpha}}{dt} = -2W_{1H}b + 2W_{1C}a - 2W_0(b-a)$$

for one  $^{13}\text{C}$  transition:

$$\begin{aligned} \frac{d(n_{\beta\alpha} - n_{\beta\beta})}{dt} &= 4aW_{1C} - 2W_0(b-a) + 2(a+b)W_2 \\ &= 2a(2W_{1C} + W_2 + W_0) + 2b(W_2 - W_0) \end{aligned}$$

# Steady-state NOE



The experiment implies the continuous irradiation of proton transitions

$$\Delta H = 0$$

At the steady state:

$$\frac{d(n_{\beta\alpha} - n_{\beta\beta})}{dt} = 2a(2W_{1C} + W_2 + W_0) + 2b(W_2 - W_0) = 0$$

concerns one  $^{13}\text{C}$  transition

$$2a(2W_{1C} + W_2 + W_0) + 2b(W_2 - W_0) = 0$$

$$\frac{2a}{2b} = -\frac{W_2 - W_0}{2W_{1C} + W_2 + W_0} = \frac{\Delta C - \Delta C_0}{\Delta H - \Delta H_0}$$

$\Delta C$  is the  $^{13}\text{C}$  population difference

$\Delta C_0$  is the equilibrium  $^{13}\text{C}$  population difference

analogously for  $^1\text{H}$

During the steady state experiment  $\Delta H = 0$

$$\frac{\Delta C - \Delta C_0}{-\Delta H_0} = -\frac{W_2 - W_0}{2W_{1C} + W_2 + W_0}$$

$$\frac{\Delta C - \Delta C_0}{\Delta H_0} = \frac{W_2 - W_0}{2W_{1C} + W_2 + W_0}$$

$$\frac{\Delta H_0}{\Delta C_0} = \frac{\gamma_H}{\gamma_C}$$

Where:

$S_0$  the intensity of the C signal for the unperturbed system

$S_s$  that in the presence of steady state NOE

$$\frac{S_s - S_0}{S_0 \frac{\gamma_H}{\gamma_C}} = \frac{W_2 - W_0}{2W_{1C} + W_2 + W_0}$$

$$\frac{S_s - S_0}{S_0} = \frac{\gamma_H}{\gamma_C} \frac{W_2 - W_0}{2W_{1C} + W_2 + W_0} = \eta$$

$\eta$ : Nuclear Overhauser enhancement factor

$$\frac{S_s}{S_0} = 1 + \eta$$

$S_s/S_0$  is the ratio of the intensities of the carbon signal in the presence of steady state NOE and in the absence of proton irradiation

In the case of exclusive **dipolar relaxation** and **extreme narrowing** hold:  $W_2 : W_{1C} : W_0$  12: 3 : 2

which leads to:

$$1 + \eta = 1 + \frac{1}{2} \frac{\gamma_H}{\gamma_C}$$

Since  $\gamma_H/\gamma_C \approx 4$ , the C signals with NOE have intensity  $1 + \eta = 1 + 2$

**NB** the effect magnitude **does not depend** on the number of close protons

# Internuclear Distances by means of NOE

- When the dipolar interaction between the two nuclei is only one of the mechanisms that cause relaxation the observed NOE is not full
- The other transition probabilities,  $W_{1C}^*$ , acting as leakage pathways, decrease the NOE

$$\eta = \frac{\gamma_H}{\gamma_C} \frac{W_2 - W_0}{2W_{1C} + W_2 + W_0 + 2W_{1C}^*}$$

In the homonuclear case  
for overwhelming  $W^*$

$$\frac{1}{\eta} = 2 + \frac{2W^*}{W_2 - W_0}$$

In the case of extreme narrowing:

$$W_1 \propto \frac{3\tau_c}{r^6} \quad W_0 \propto \frac{2\tau_c}{r^6} \quad W_2 \propto \frac{12\tau_c}{r^6}$$

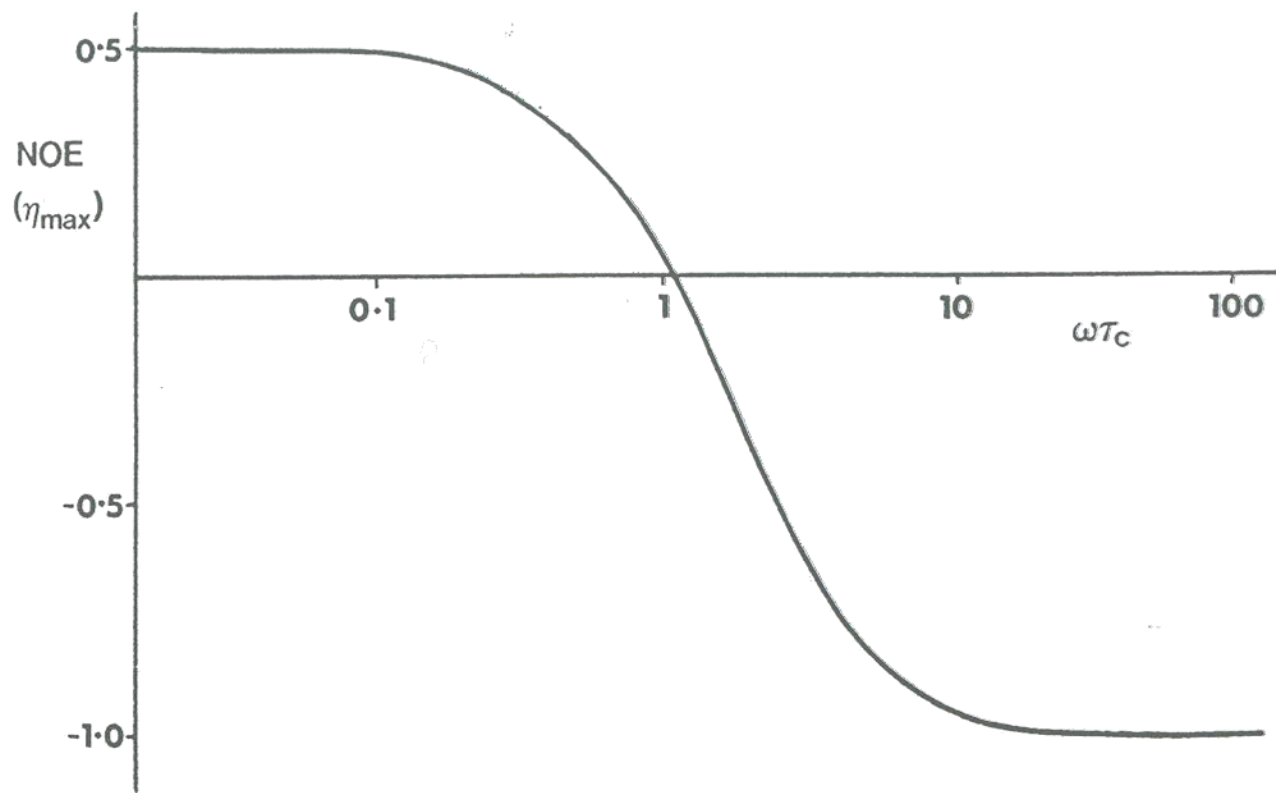
$$e \quad W_2 - W_0 \propto \frac{10\tau_c}{r^6}$$

$$\frac{1}{\eta} \propto 2 + 2\rho^* \frac{r^6}{\tau_c}$$

For analogous molecules  
it can be expected that  
 $\rho^*/\tau_c$  are the same, so  
that the measured **NOEs**  
are proportional to  $r^{-6}$ .

In this way information on  
distances is obtained

# Steady State NOE $\eta$ : Dependence on the Motional Regime

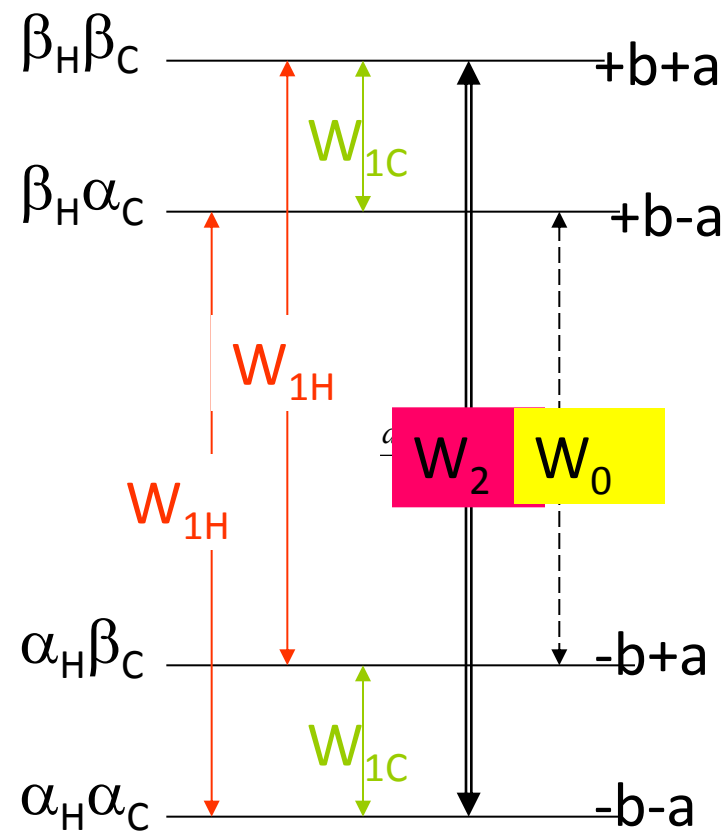


**FIGURE 2.6**

Dependence of maximum homonuclear NOE enhancement on  $\omega\tau_c$ . Note the log scale of  $\omega\tau_c$ .



# TOE (Transient NOE)



Starting condition:  $\pi$  pulse on  $^1\text{H}$

$$\Delta H_{180} = -\Delta H_0 \quad 2b = 2\Delta H_0$$

$$\Delta C = \Delta C_0 \quad 2a = 0$$

since  $\Delta H_0$  and  $\Delta C_0$  are constants

$$\frac{d(\Delta H - \Delta H_0)}{dt} = \frac{d\Delta H}{dt}$$

$$\frac{d(\Delta C - \Delta C_0)}{dt} = \frac{d\Delta C}{dt}$$

$$\frac{dn_{\beta\beta}}{dt} = -2bW_{1H} - 2aW_{1C} - 2(a+b)W_2$$

$$2a = -(\Delta C - \Delta C_0)$$

$$\frac{dn_{\beta\alpha}}{dt} = -2W_{1H}b + 2W_{1C}a - 2W_0(b-a)$$

$$2b = -(\Delta H - \Delta H_0)$$

$$\frac{dn_{\alpha\beta}}{dt} = 2W_{1H}b - 2W_{1C}a + 2W_0(b-a)$$

$$\frac{d\Delta C}{dt} = \frac{d(n_{\beta\alpha} - n_{\beta\beta})}{dt} = 2a(2W_{1C} + W_2 + W_0) + 2b(W_2 - W_0)$$

$$\frac{d\Delta H}{dt} = \frac{d(n_{\alpha\beta} - n_{\beta\beta})}{dt} = 2b(2W_{1H} + W_2 + W_0) + 2a(W_2 - W_0)$$

$$\frac{d(\Delta C - \Delta C_0)}{dt} = -(\Delta C - \Delta C_0)(2W_{1C} + W_2 + W_0) - (\Delta H - \Delta H_0)(W_2 - W_0)$$

$$\frac{d(\Delta H - \Delta H_0)}{dt} = -(\Delta H - \Delta H_0)(2W_{1H} + W_2 + W_0) - (\Delta C - \Delta C_0)(W_2 - W_0)$$

Performing the following substitutions:

$$x = (\Delta H - \Delta H_0) \quad y = (\Delta C - \Delta C_0)$$

$$R_H = 2W_{1H} + W_2 + W_0$$

$$R_C = 2W_{1C} + W_2 + W_0$$

$$\sigma_{HC} = W_2 - W_0$$

one recognizes two coupled first order differential equations

## Solomon Equations

$$\frac{dx}{dt} = -R_H x - \sigma_{HC} y$$

$$\frac{dy}{dt} = -R_C y - \sigma_{HC} x$$

# Solution for the homonuclear case

The solution is simple for  $R_H = R_C$  (homonuclear case)

$$y(\tau) = 1/2 \exp(-R\tau) [\exp(\sigma\tau) - \exp(-\sigma\tau)] = \exp(-R\tau) \sinh(\sigma\tau)$$

- for  $\tau > 0$   $y(\tau)$  is the product of an increasing function and one decreasing

- it must possess a maximum

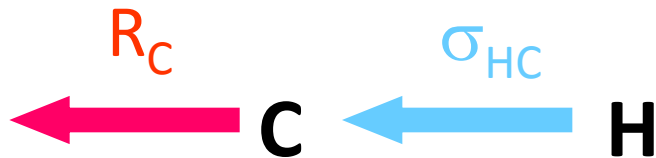
$$\tau_{Max} = \frac{1}{2\sigma} \ln \frac{R + \sigma}{R - \sigma}$$

The value of the maximum increment for the homonuclear transient NOE is::

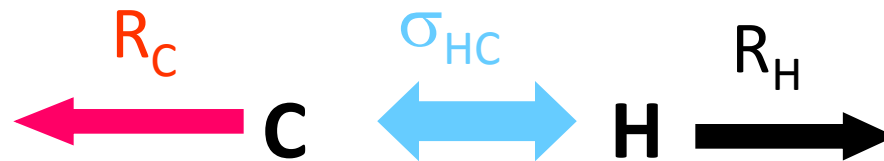
$$\eta_{Max}^{transiente} = \left( \frac{R + \sigma}{R - \sigma} \right)^{-\left[ \frac{R - \sigma}{2\sigma} \right]} - \left( \frac{R + \sigma}{R - \sigma} \right)^{-\left[ \frac{R + \sigma}{2\sigma} \right]}$$

# Steady State NOE and TOE: Comparison

**steady state NOE: asymmetric**

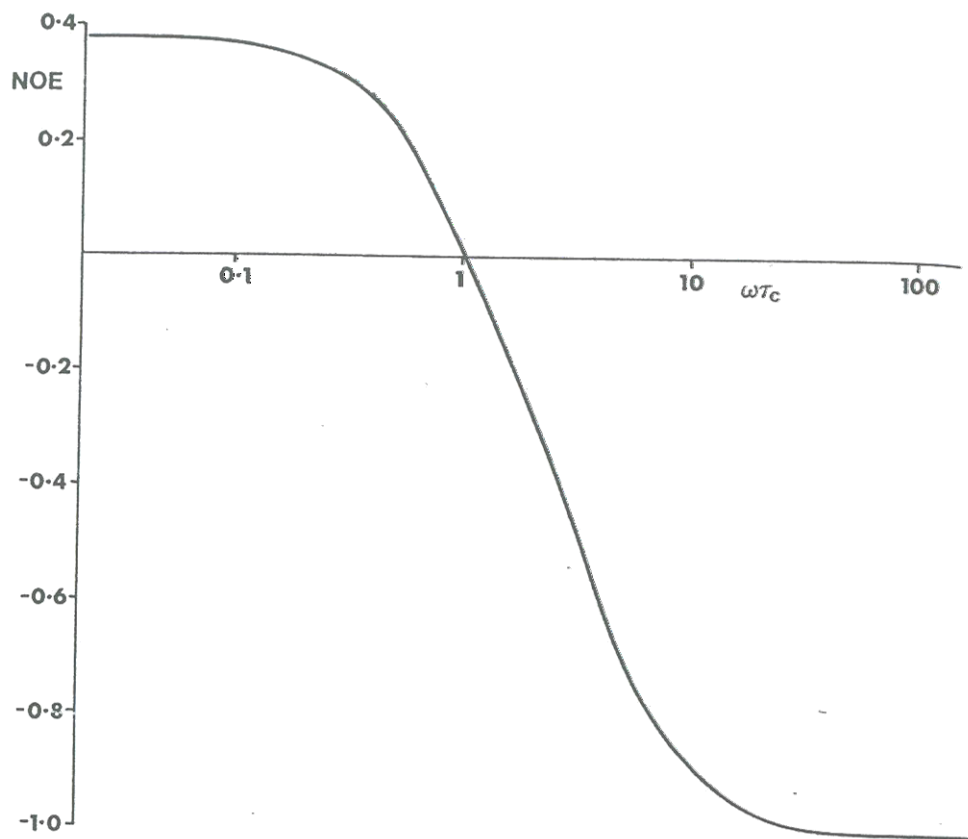


**transient NOE: symmetric**



R refers to self-relaxation,  $\sigma$  to mutual relaxation

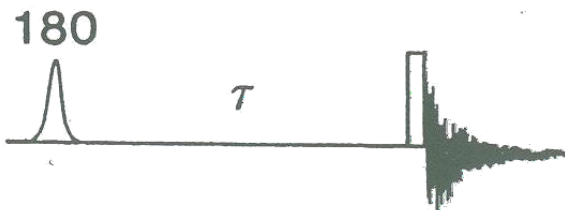
# Dependence of TOE $\eta$ on the Motional Regime



**FIGURE 4.16**

Plot of  $\eta_{\max}^{\text{transient}}$  (the maximum enhancement attainable in a two-spin homonuclear system, using a transient NOE experiment) against  $\omega\tau_c$ . Note the similarity to the curve for steady-state enhancements vs.  $\omega\tau_c$  (Figure 2.6).

selective



# Rotating Frame Nuclear Overhauser Effect

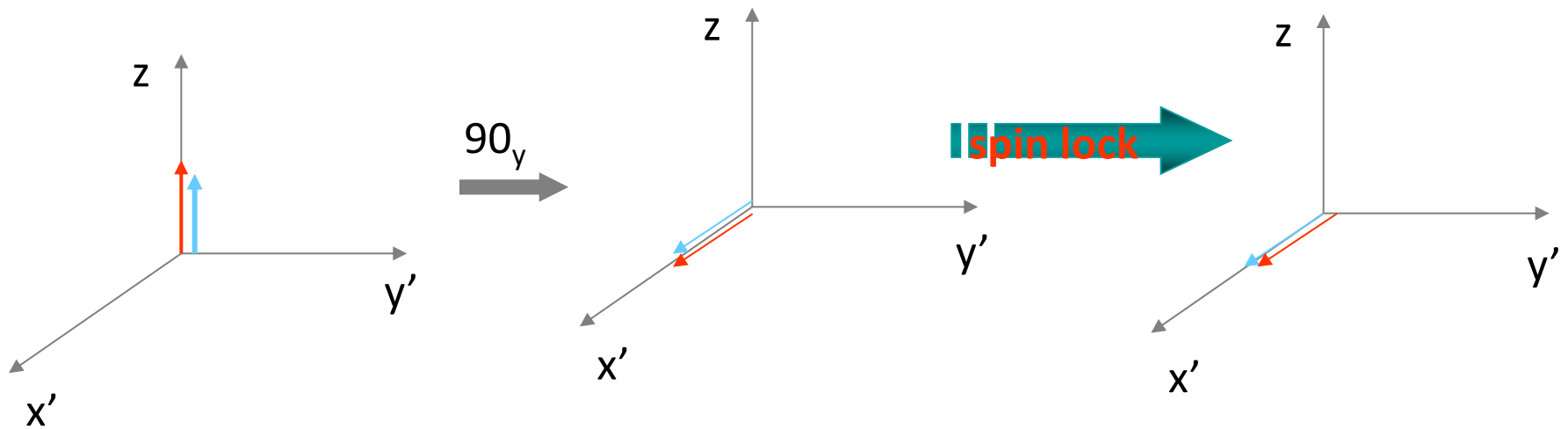
~~transverse cross-relaxation~~

There is no polarization transfer between spin I and spin S  
transverse magnetizations

It happens exclusively when  $\omega_I = \omega_S$ , i.e. degenerate nuclei, or  
when the two nuclei **look degenerate**, that is they have the same  
Lamor frequency

The former case is not interesting because it is impossible to detect the magnetization transfer between two degenerate spins

We can pretend that the two nuclei have the same Larmor frequency by the **spin lock**

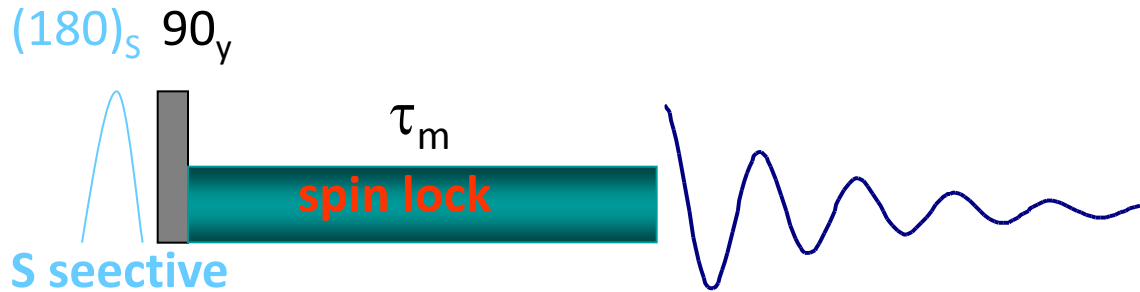


they can cross talk because they are close



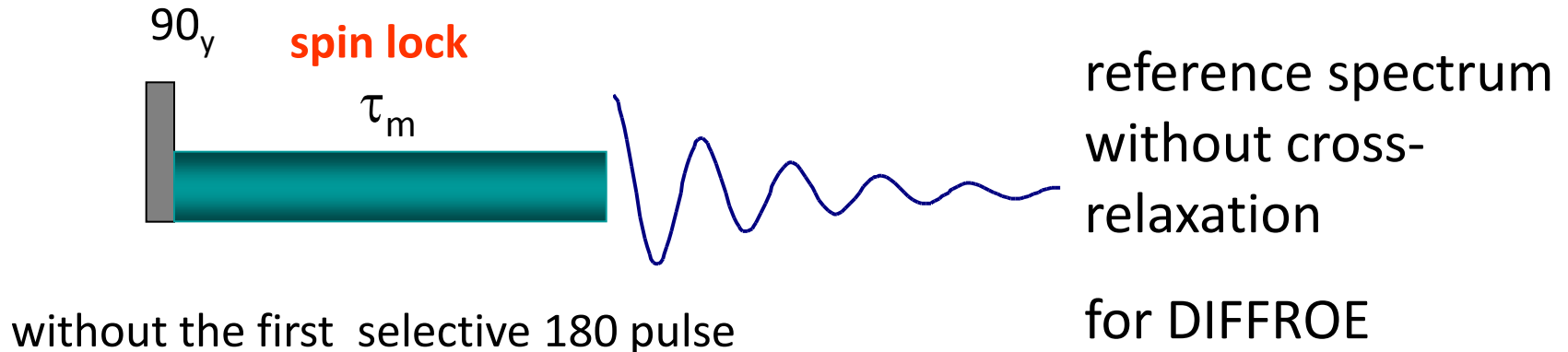
- ✓ The simplest spin lock consists of a r. f. magnetic field on  $x'$
- ✓ The field must be strong enough so that the effective field is on the  $x'$  axis for both spins, which have different off-sets
- ✓ In the rotating frame the transverse magnetizations of both spins appear static
- ✓ the magnetization is called **spin locked**
- ✓ since the two vectors,  $M_I$  and  $M_S$ , do not diverge during the application of the spin lock, they look to possess the same Larmor frequency
- ✓ under these conditions **transverse cross-relaxation** can take place
- ✓ the result can be detected by recording the usual FID after the removal of the spin lock

# 1D ROE Experiment



La combinazione dell'impulso selettivo di  $\pi$  e di quello (non selettivo) di 90 genera uno stato in cui  $M_x$  è lungo  $x'$  mentre  $M_y$  è lungo  $-x'$

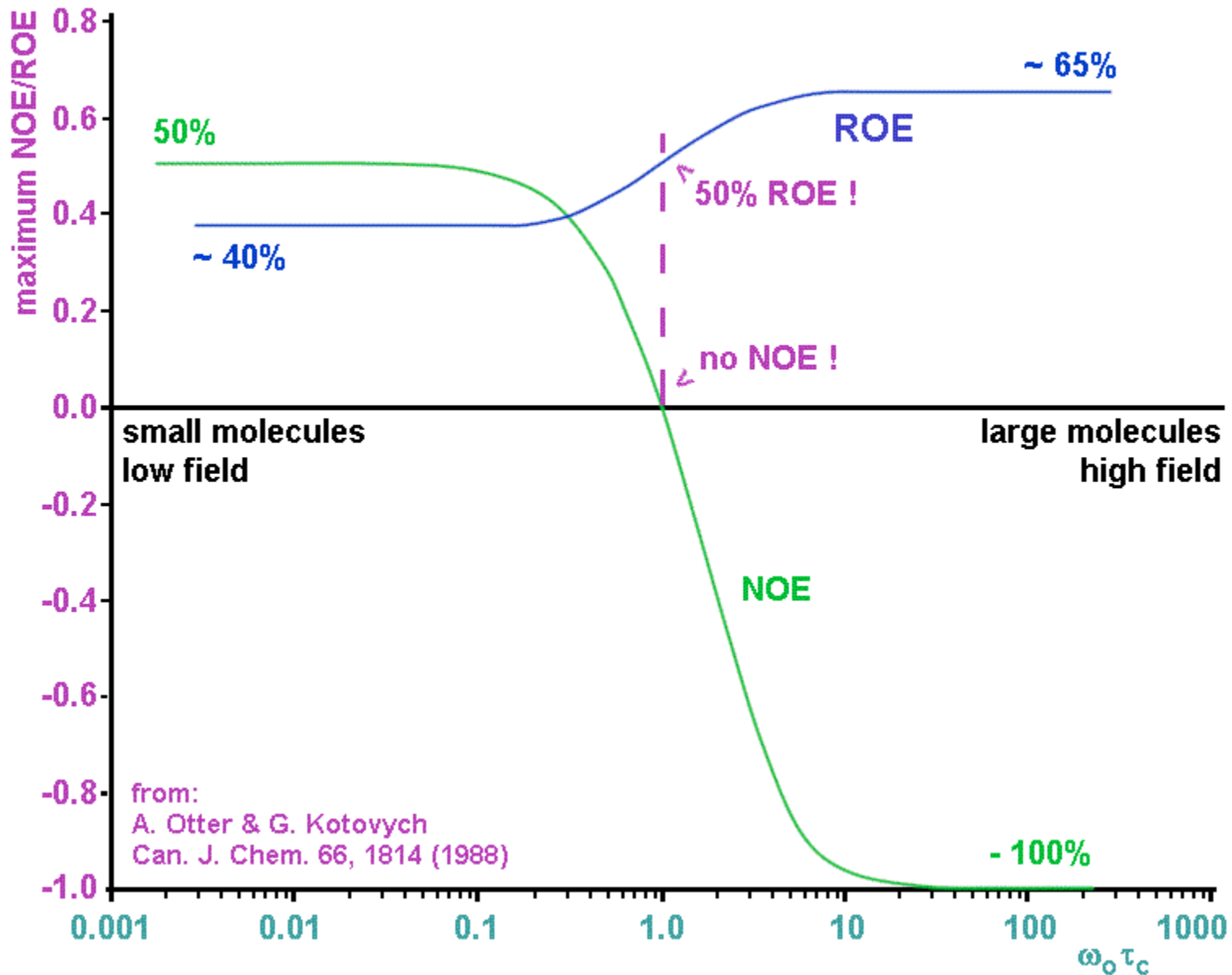
Durante lo spin-lock può avvenire la cross-relaxation trasversale



# Dependence on the molecular tumbling frequency

- The NOE is positive for motions faster than the Larmor frequency
- is negative for slow motions
- it is zero for intermediate motional regimes
- The TOE displays an analogous dependence as NOE, its maximum value is slightly lower
- The ROE is always positive and it does not become zero because of the motional regime !!!

# $\eta$ Comparison for NOE and ROE



In the case of **off-resonance ROE**, that is the effective B is not lying on the transverse plane, there is a combination of ROE (for the transverse components) and of NOE per le componenti longitudinali

- The signal intensity is proportional to  $\gamma \cdot \gamma^{3/2}$
- By means of NOE an increase of intensity is obtained by an artificial increase of the Boltzmann factor (the population difference)
- It is needed that the weak nucleus is relaxed mainly by the dipolar coupling with protons.  $^{15}\text{N}$  and  $^{29}\text{Si}$  have  $\gamma < 0$  and their increments are  $< 0$  (-5 per  $^{15}\text{N}$ )

# Dynamic Nuclear Polarization Electron-Nucleus in Soluzione

- The term DNP is properly used to address the polarization transfer from e<sup>-</sup> to nucleus
- It can be obtained both in solid and in liquid state
- The Overhauser Effect is the mechanism for the DNP in solution