

***1--Nonlinear dynamics in Optics***  
***2-Chaos in optics and***  
***application to Cognitive processes***

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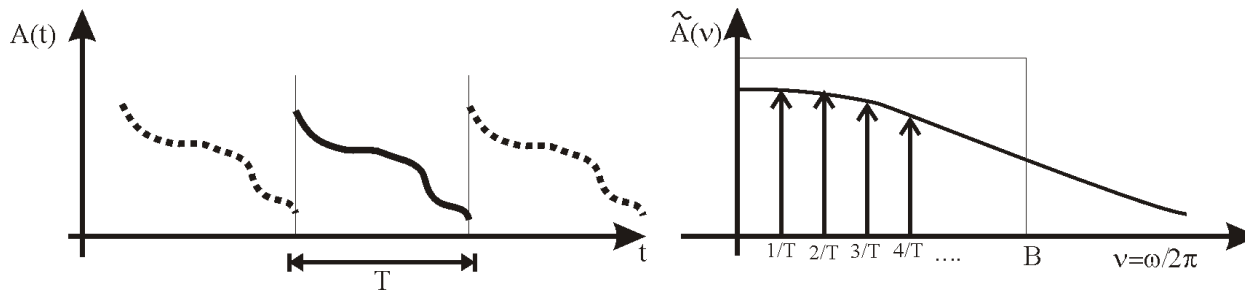
# ***Nonlinear dynamics in Optics***

## ***0 - HOW TO ORGANIZE OBSERVATIONS***

## Degrees of freedom of a signal (Shannon)

Two real n° for each Fourier component, namely, **modulus and phase**

$$A(\omega) = |A(\omega)|e^{i\varphi(\omega)}.$$



A measuring device has a band **B** and is active for a time **T**

Beyond **T**, assume  $A(t)$  repeats left and right, thus use a **series expansion**: keep only harmonics  $A(v)$ ,

spaced by  $1/T$ , and only those within **B**, for a total **B·T**. Each one yields two reals, thus the n° **N** of degrees of freedom is (**Shannon**):

$$N = 2BT$$

Apply to TV screen. Let us visualize a chessboard of white-black squares.

Visual permanence time  **$t_p = 1/15$  s**; to have motion feeling, need 2 squares within  $t_p$ , hence  **$T = 1/30$  s**. TV channel band  **$B = 5$  MHz**. Thus

$$N = 2 \cdot 5 \cdot 10^6 \cdot \frac{1}{30} \cong 0,3 \cdot 10^6$$

N° horizontal lines  **$N^{1/2} = 600$**

## Space case :optical resolution (pixel)

Image formation through a lens (Abbe, 1880) .

Shine an object with a plane wave; information elements will scatter the single  $k$  direction to many waves of equal color but different directions  $k$ . The amplitude  $\tilde{A}(\vec{k})$  results from summing the different features  $A(r)$  of the 2D domain  $\vec{r}(x, y)$  with different phases (**Fourier transform**)

$$A(k) = \int A(r) e^{ik \cdot r} dr .$$

Let's go through a lens. Each plane wave converges to a point of the focal plane (= Fourier plane).

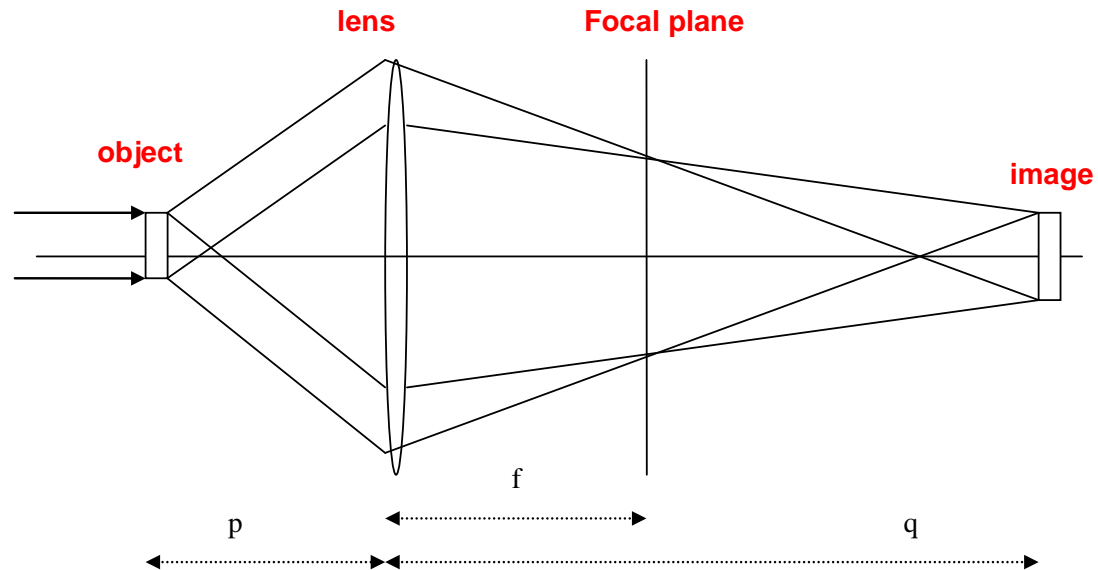
However a detector senses NOT the *amplitude but the energy* of each wave (phase lost).

We do not get the signal on the focal plane, but let each point act as a source of a spherical wave. The spherical waves interfere with their mutual phases; in particular they recombine on the image plane yielding an image similar (besides a *magnification*) to the object. Object and image planes are related by *Gauss formula*

$$\frac{1}{p} + \frac{1}{q} = \frac{1}{f}$$

$p$ ,  $q$  and  $f$  being the distances of object, image and focus from the lens. Thus the object is an inverse Fourier transform

$$A(r) = \mathfrak{F}^{-1}[A(k)] = \int A(k) e^{-ik \cdot r} dk$$



- Image formation (**Abbe**) : each scattered wave is focused on a point on the focus; points on  **$f$**  are sources of spherical waves which sum with the right phases only on the image

- **Image formation**

$$\underbrace{A(r)}_{\text{object}} \Rightarrow \Rightarrow \underbrace{A(k)}_{\text{focal plane}} \Rightarrow \Rightarrow \underbrace{A(r)}_{\text{image}}$$

- this occurs only in the visible where transparent media **refract**, i.e. bend the rays (lens).
- At X rays no lenses. Going in the far field collect all the square Fourier transform

$$|A(k)|^2$$

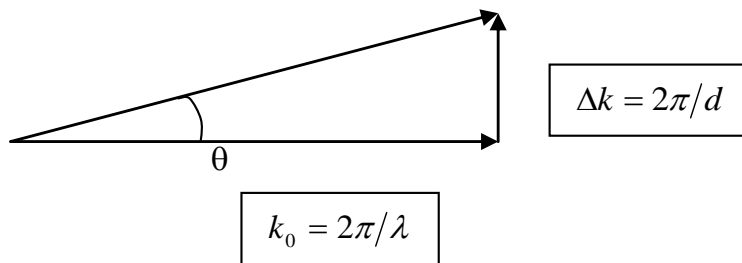
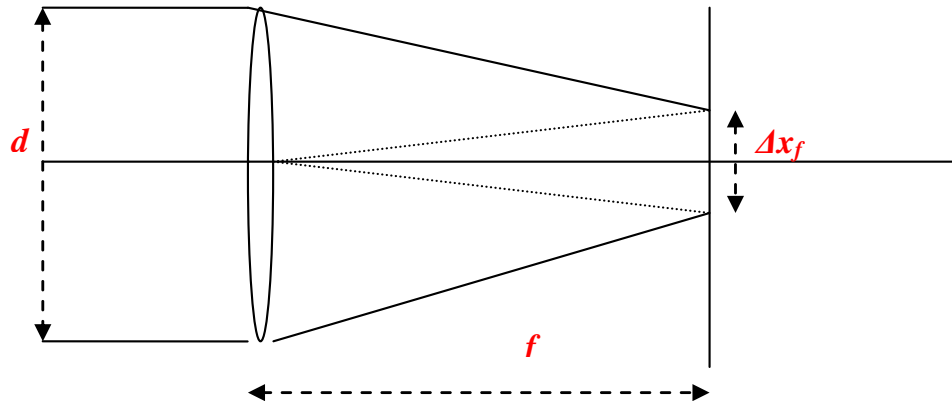
- **Problem of phase:** *how to reconstruct the signal in crystallography*

## PSF(point spread function) and space Shannon (pixels) .

Due to limited  $d$ , large angles are lost .

A plane wave in Fourier is not integrated between  $\pm\infty$  but between  $\pm d/2$  ; as a rect function angles are spread in a cone

$$\theta = \Delta k / k = \frac{2\pi/d}{2\pi/\lambda} = \frac{\lambda}{d} .$$



- Have a spot

$$\Delta x_f = f \cdot \theta = \lambda f / d$$

- Thus the focal image of a plane wave is  $\Delta x_f$  called **PSF** (point spread function):

- The Fourier band is not  $\Delta k = \infty$  yielding  $\Delta x_f = 0$ , ma  $\Delta k = 1/\Delta x_f = d/\lambda f$

The number of distinct spots (**pixel** = picture elements) with an object of length  $L$  è  $L/\Delta x_f$ .

Multiply by 2 (amplitude+phase) and get the n°  $N$  of degrees of freedom of a 1-dimensional image.  
(**Shannon**)

$$N = 2L \cdot \Delta k = 2L \frac{d}{f\lambda}$$

For 2D have  $N^2$

For a slide of side  $d=3cm$  and focus  $f=10cm$ , in green light ( $\lambda=0,5$  micrometer)

$N=10^4$  thus  **$N^2=10^8$  pixel=100 megapixel.**

In microscopes  **$d=f$**  ( in jargon: numerical **numerical aperture =1**)



# ***Nonlinear dynamics in Optics***

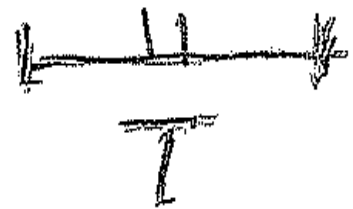
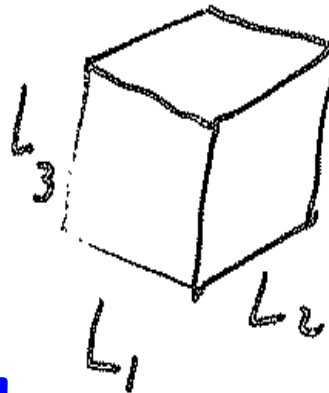
## ***1-LASER BASICS***

## The e.m. field and its quantization

$$\vec{E}(x, y, z, t) = \sum_{\vec{k}} \vec{E}(\vec{k}, t) e^{i(k_1 x + k_2 y + k_3 z) - i\omega_k t}$$

$$\omega_k = c k$$

$$k_i = n_i \frac{2\pi}{L_i} \quad (i = 1, 2, 3)$$



$$\delta^3 k = \frac{(2\pi)^3}{V}$$

$$\omega_{1'} = n_{1'} \frac{2\pi}{L}$$

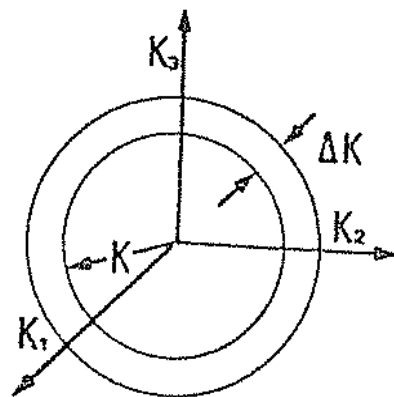


Fig. 1

**Rayleigh**

$$M = 2 \cdot \frac{4\pi k^2 \Delta k}{\int^3 k} = \boxed{8\pi \frac{\nu^2}{c^3} \Delta \nu V}$$

$$= 8\pi \frac{V}{\lambda^3} \cdot \left( \frac{\Delta \nu}{\nu} \right)$$

**Planck 1900**

$$\frac{dW}{d\nu} = \frac{dM}{d\nu} \cdot h\nu \cdot \bar{n}_1(\nu) = \boxed{\frac{8\pi \nu^2}{c^3} \cdot V \cdot h\nu \frac{1}{e^{h\nu/kT} - 1}}$$

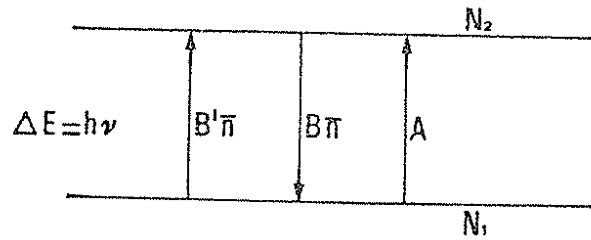


Fig. 2

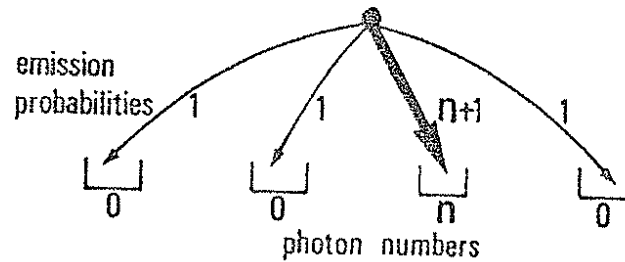


Fig. 3

**Einstein 1915**

$$\frac{N_2}{N_1} = e^{-\Delta E / kT} \quad \text{where } k \text{ is Boltzmann constant}$$

$$N_1 B \bar{n} = N_2 B \bar{n} + N_2 \cdot A$$

$$\bar{n} = \frac{A/B}{\frac{B'}{B} \frac{N_1}{N_2} - 1} \equiv \bar{n}_i(\nu) \cdot \Delta \nu$$

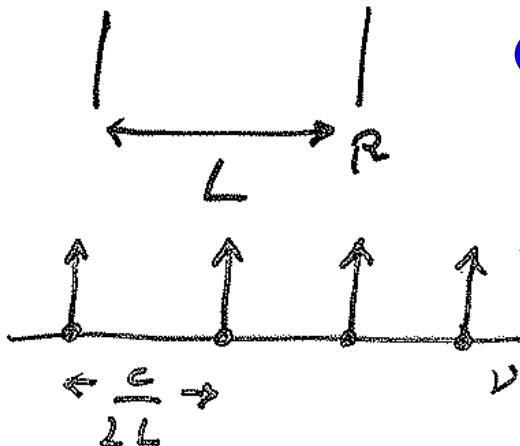
$$B' = B, \quad \frac{A}{B} = \frac{8\pi \nu^2}{c^3} V \Delta \nu = M$$

# L A S E R

$$n > 1$$



Fabry-Perot cavity



$$t_c = \frac{L}{c} \frac{1}{1-R}$$

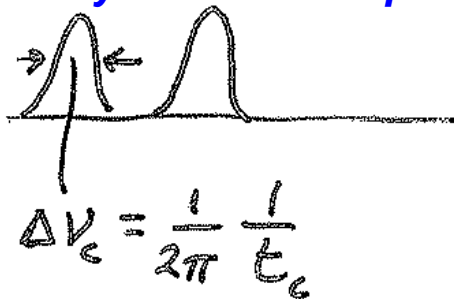
$R < 1$

$$m \cdot \frac{\lambda}{2} = L$$

$$\Delta \nu_{i,i+1} = \frac{c}{2L}$$

FSR

Fabry-Perot ideal spectrum



[Free spectral range]

$$Q_{\text{finesse}} = \frac{\Delta \nu_{i,i+1}}{\Delta \nu_c} = \frac{\pi}{1-R}$$

Fabry-Perot real spectrum

Rate

$$\dot{n} = B N \cdot n - n/t_c + B N_2$$

$$\dot{N} = -B N \cdot n + P - A N_2$$

Threshold

$$N \geq \frac{1}{B \cdot t_c}$$

Pump

$$P \Rightarrow B N n > \frac{n}{t_c} > \frac{M}{t_c}$$

**Concept of cross section**

$$B n \equiv \sigma \phi = \sigma \cdot c \frac{n}{V}$$

$1/s$        $cm^2 / cm^2 \cdot s$

$$\boxed{\sigma = \frac{B V}{c}}$$

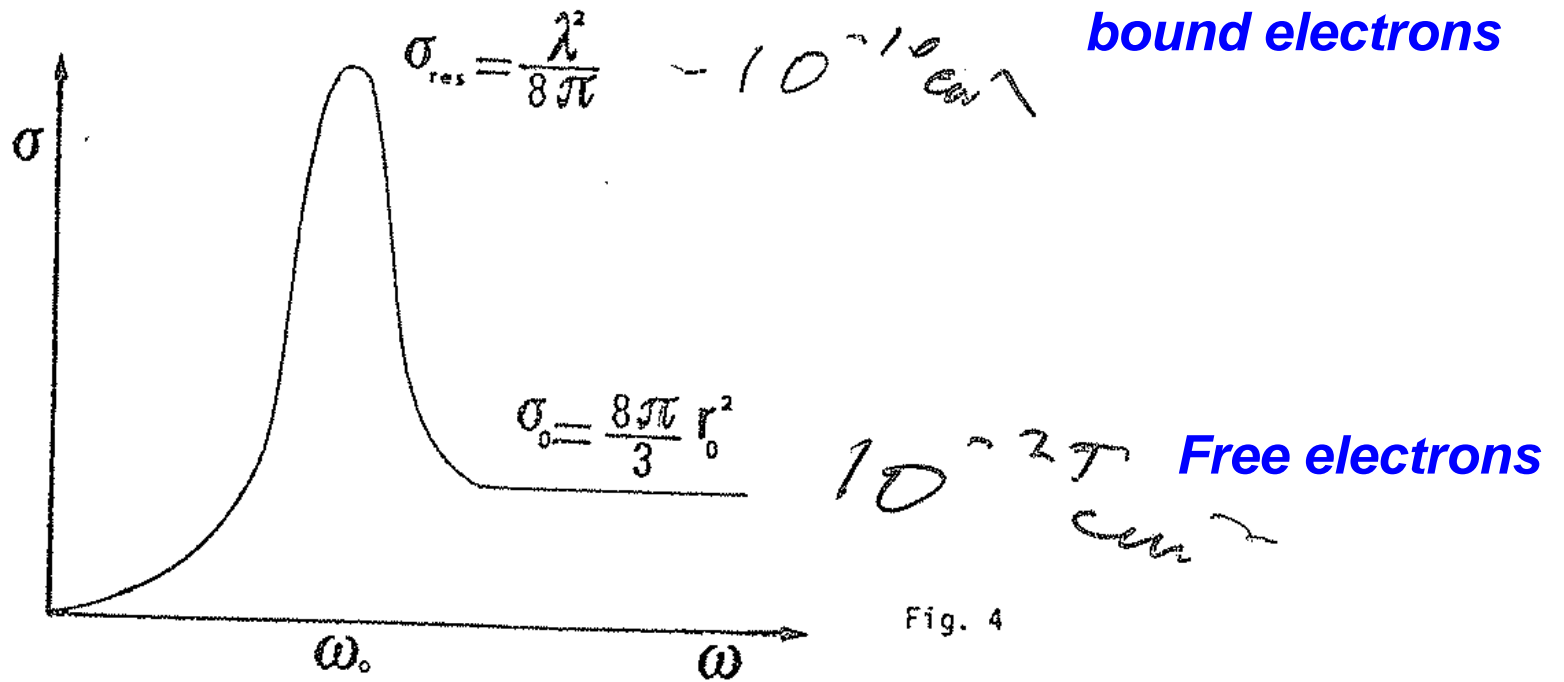
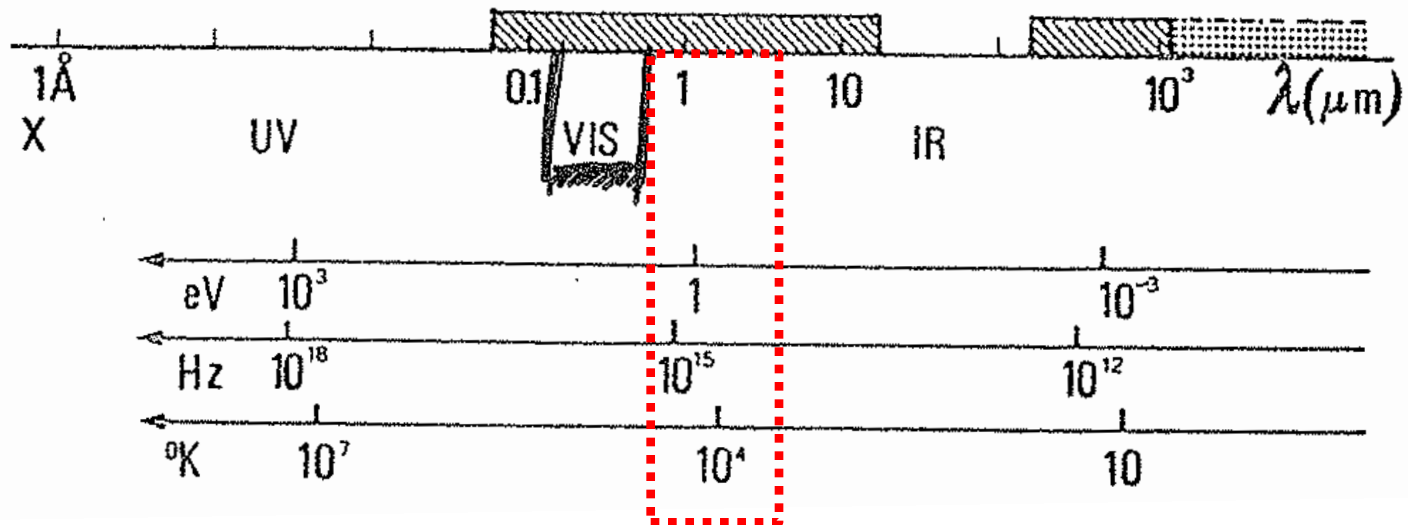
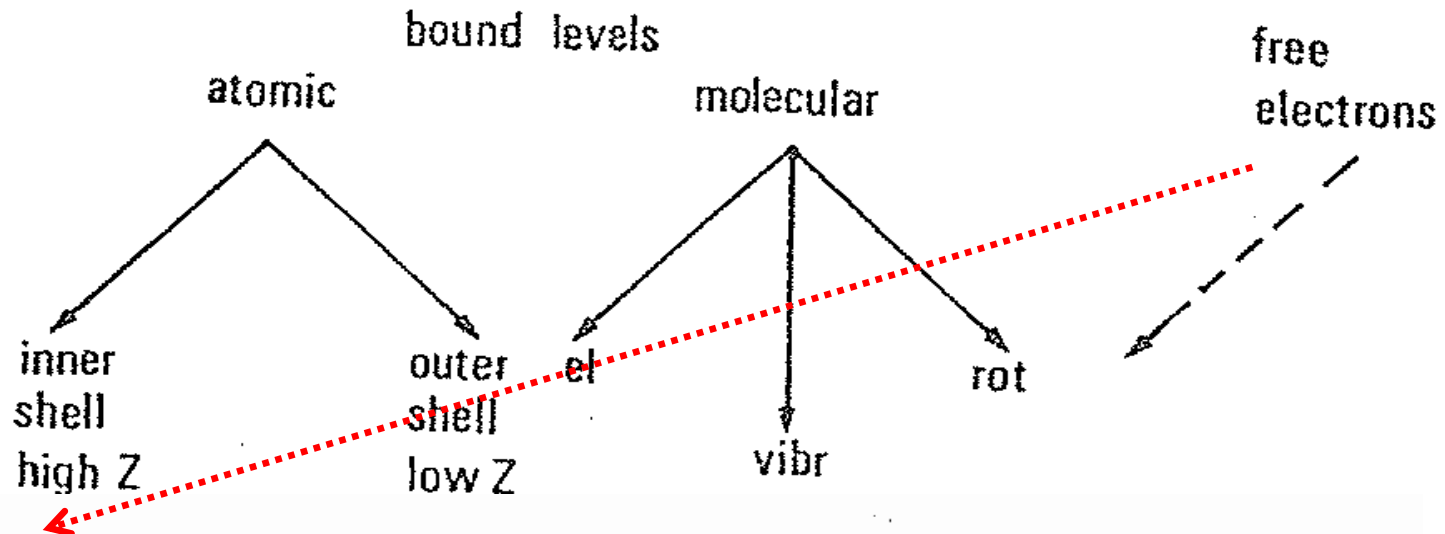
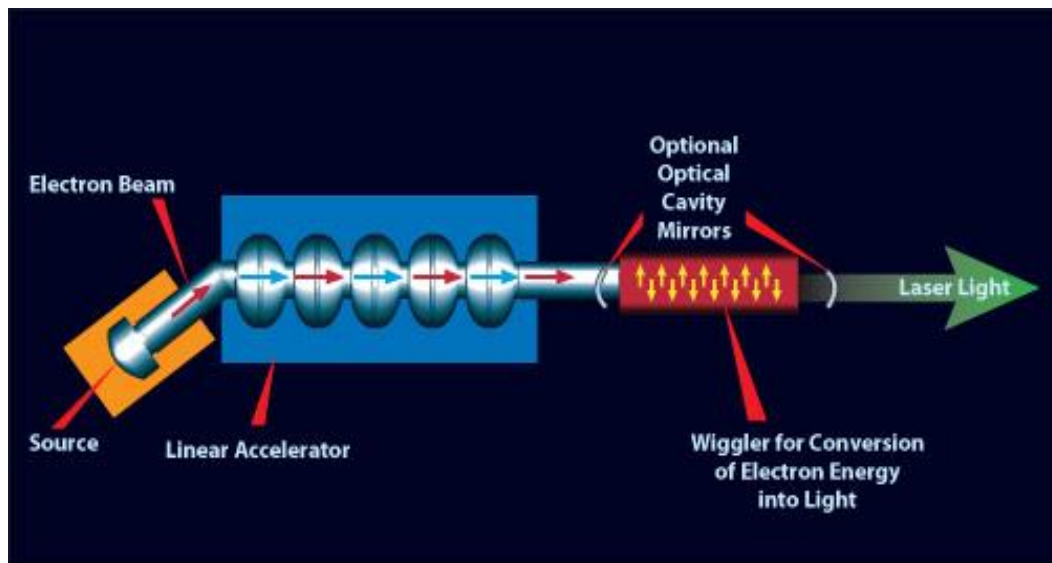


Fig. 4

# LASERS



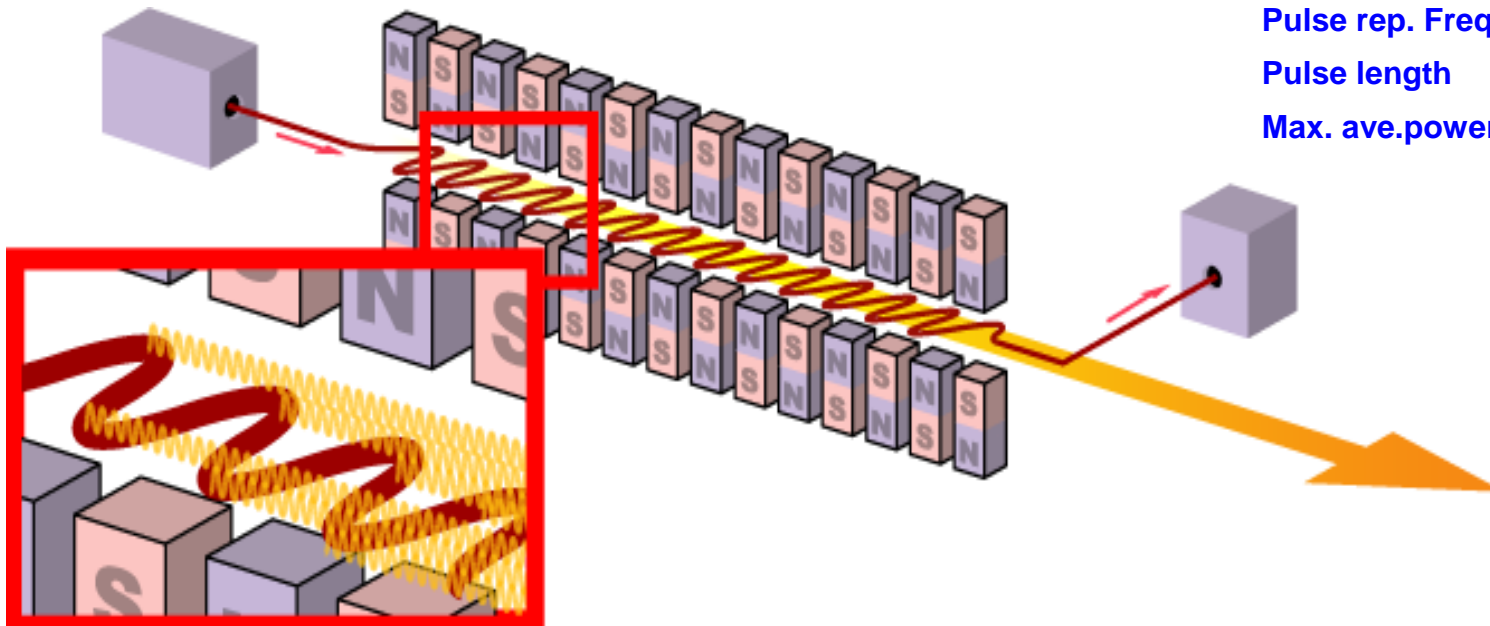




Wavelength range (IR) 1-14 $\mu$ m  
 Energy/pulse 120  $\mu$ J  
 Pulse rep. Freq. Up to 75 MHz  
 Pulse length 500-1700 fs  
 Max.ave. power

>10 kW

Wavelength range (UV/VIS) 250-1000 nm  
 Energy/pulse 20  $\mu$ J  
 Pulse rep. Freq. Up to 75 MHz  
 Pulse length 300-1700 fs  
 Max. ave.power >1 kW



- electron energy  $E$ ,
- period of the undulator magnet  $\lambda_u$
- its magnetic field  $B$ ,

$$\lambda = \lambda_u (1 + K^2) / 2\gamma^2 \quad [\text{double Doppler}]$$

$\lambda_u$  = undulator period (e.g. = 1cm)

$K = eB\lambda_u / (2\pi mc) = \text{pitch parameter}$  (e.g.=2)

$\gamma = E / (mc^2) = \text{relativistic factor}$  (e.g. =  $10^3$  for  $E=0.5$  GeV)

$N$  = number of undulator periods

drawback of FELs : setups large and expensive, used only at few large facilities. Most ambitious project currently pursued in Hamburg : goal hard X-ray output , wavelengths down to 0.085 nm and pulse duration below 100 fs. So far, wavelengths down to 6.5 nm achieved

# *Linac Coherent Light Source (LCLS) at SLAC- Stanford.*

## *Electron Beam*

*Electron energy, GeV*      *14.3*    ( $\gamma=2.8 \times 10^4$ )

*Peak current, kA*              *3.4*

*Pulse duration, fs*            *230*

## *Undulator*

*Period, cm*                      *3*

*Field, T*                          *1.32*

*K*                                    *3.7*

*Gap, mm*                        *6*

*Total length, m*               *100*

## *Radiation*

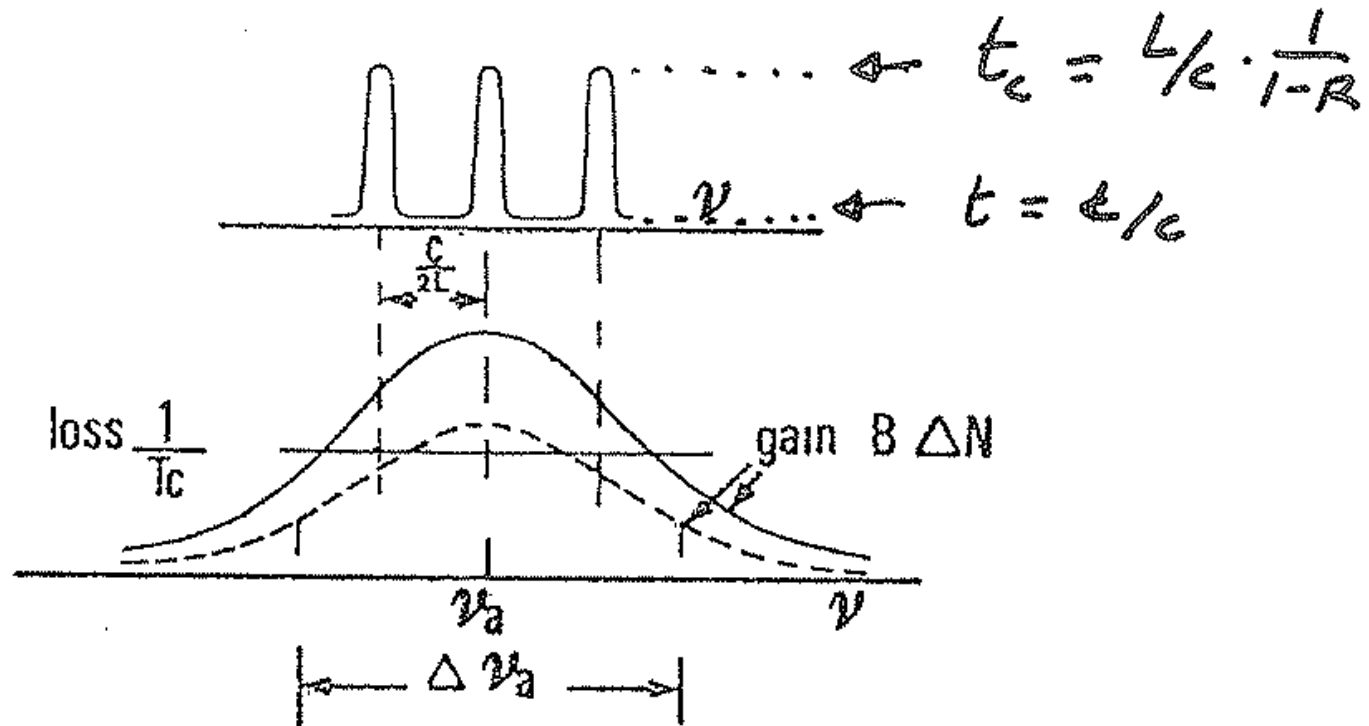
*Wavelength, nm*               *0.15*

*Bunches/sec*                   *120*

*Average Brightness*           *$4 \times 10^{22}$*

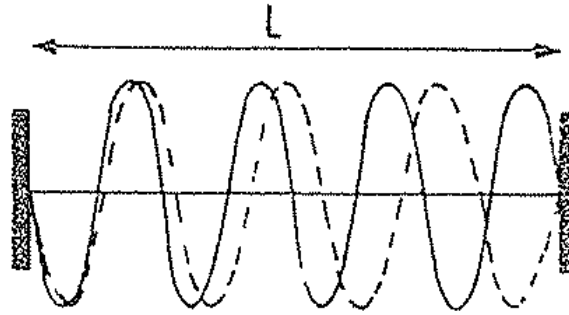
*Peak power, GW*                *$10^{10}$*

**Threshold:  $\text{Gain} = BN > 1/t_c$**



**As gain increases, from 1 to 3 lasing modes**

*Two adjacent modes  
have different wavelength*



*Since pop. depletion (saturation) depends on local intensity  
different modes exploit different regions of the laser material  
(space inhomogeneity)*

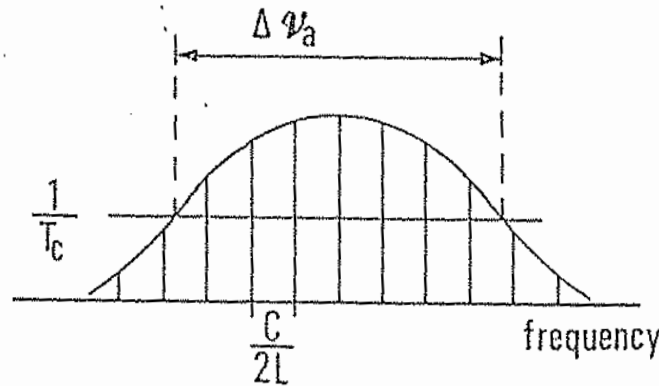


*space inhomog.*

MODE LOCKING  $\rightarrow$  SHORT PULSES

***N # of modes above threshold = material width/mode separation***

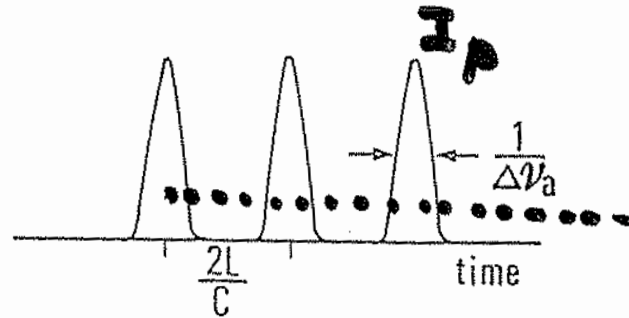
$$N = \frac{\Delta \nu_a}{\frac{c}{2L}}$$



a)

***If modes coherent in phase, then Fourier spectrum of repetitive pulses***

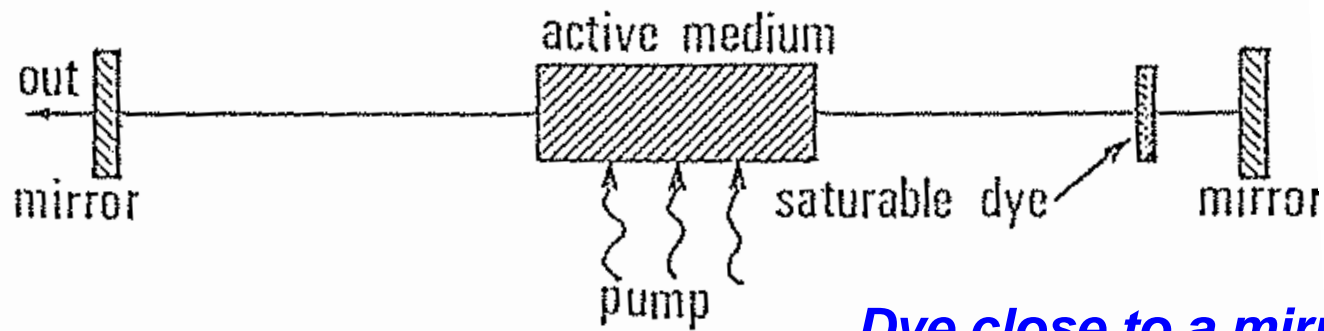
$$\Delta t = \frac{1}{N} \text{ sep.}$$



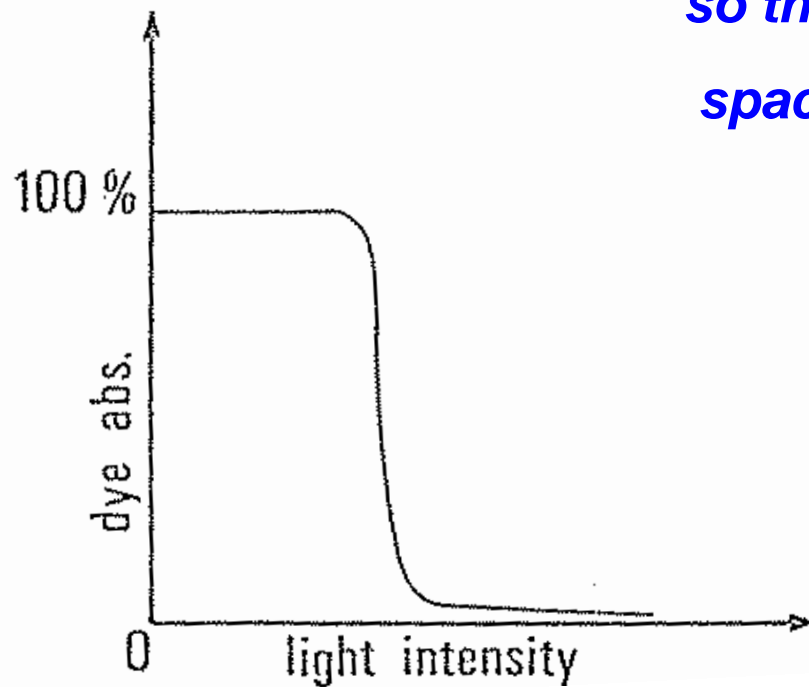
$$I_p = N^2 I,$$

$$\langle I \rangle = N I,$$

***Pulse duration = (pulse separation)/N***

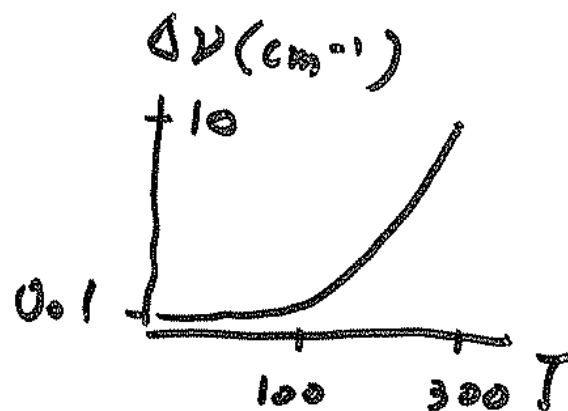
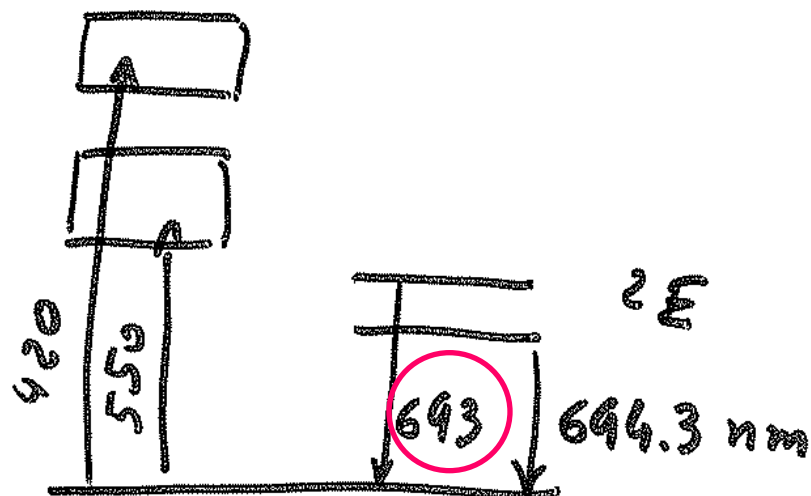


*Dye close to a mirror,  
so that all modes have same  
space phase*



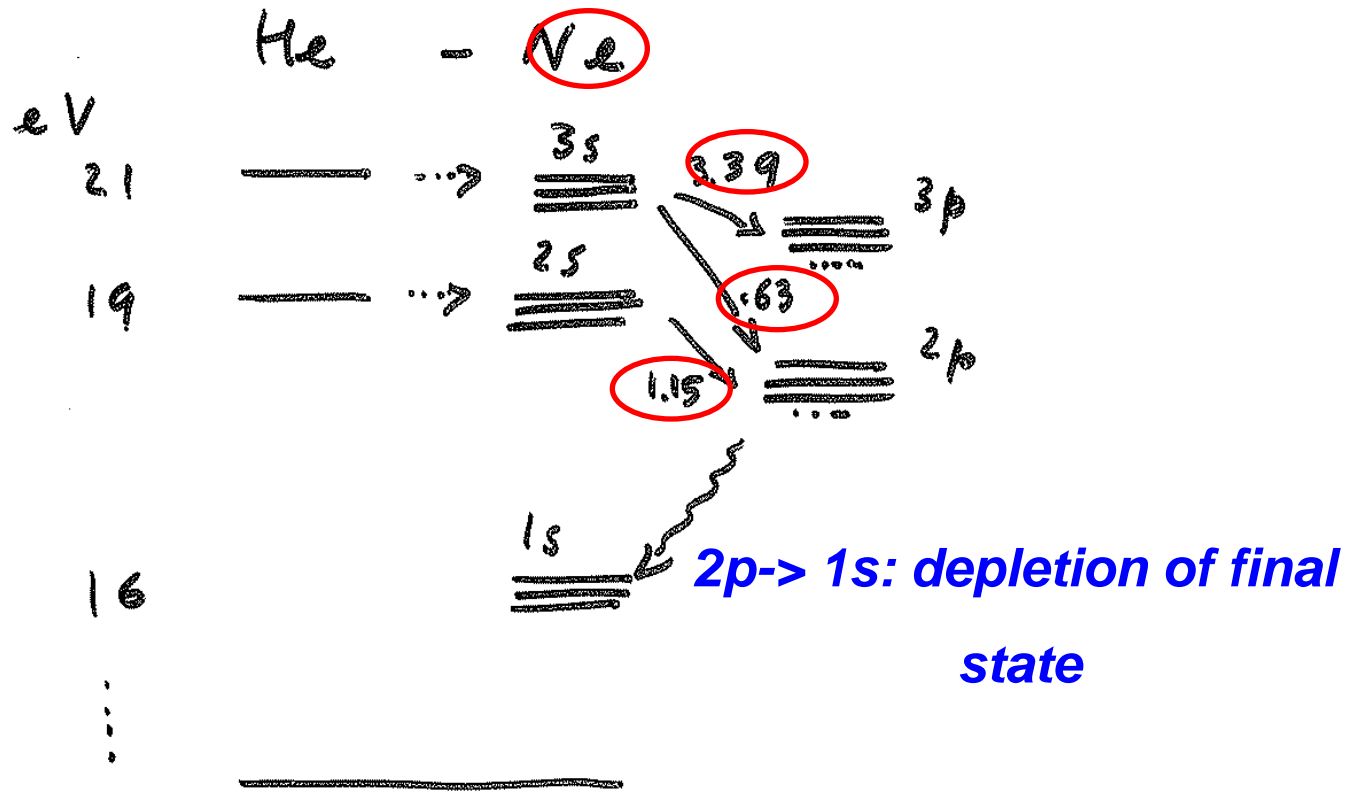
Ruby  $\text{Cr}^{3+} : \text{Al}_2\text{O}_3$

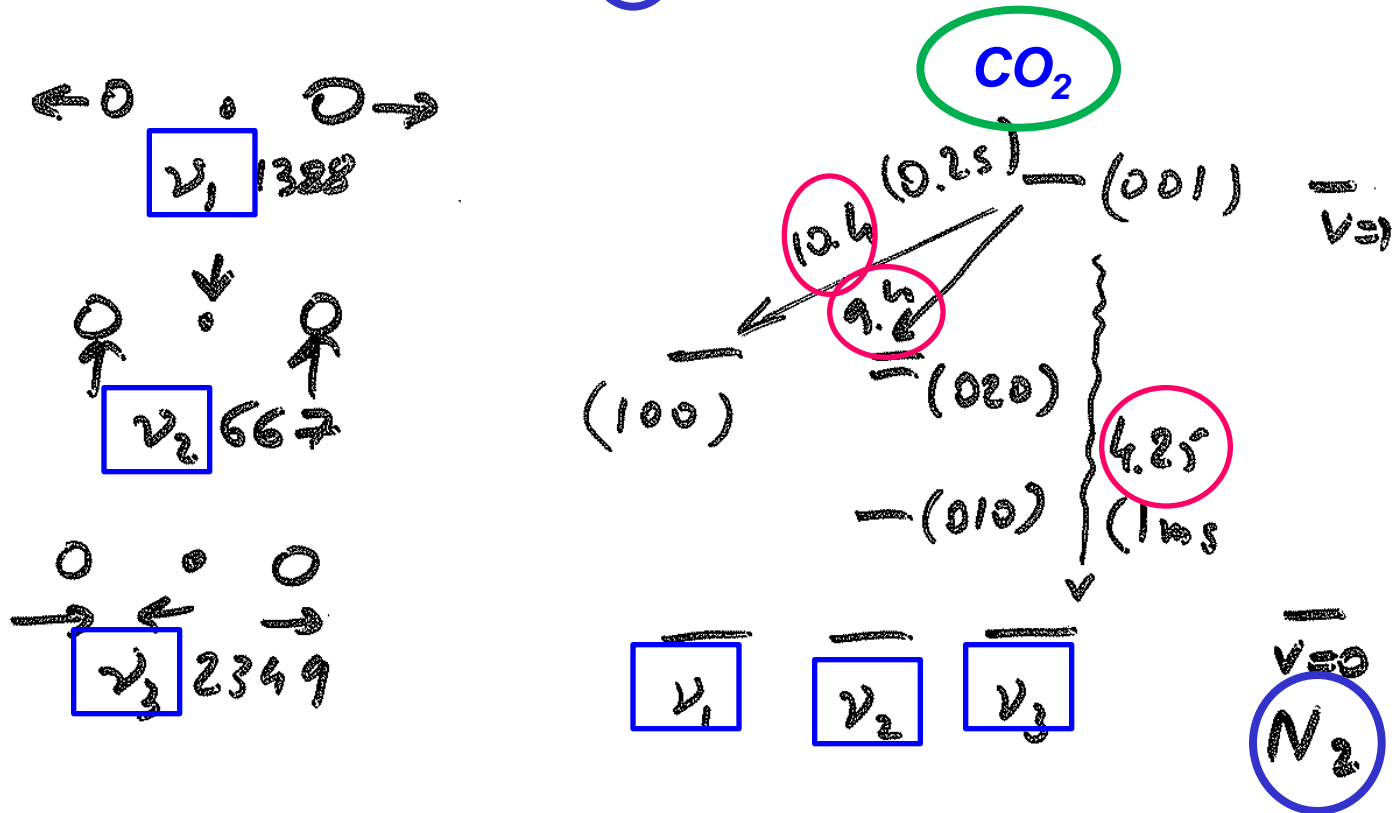
• • x





 Radiative transitions



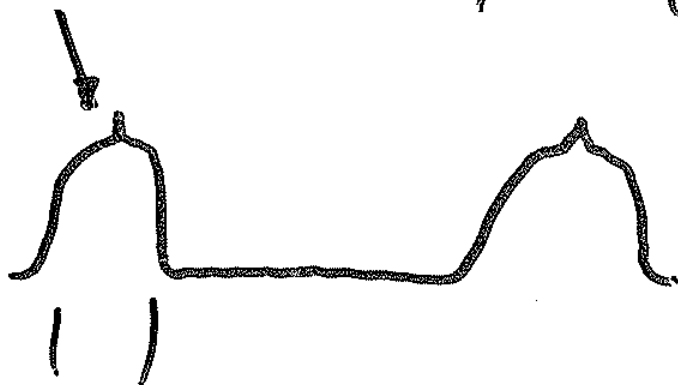


$\nu_{1-3}$  are three vibrational modes : transition frequencies ( $\text{cm}^{-1}$ )  
and  $\bigcirc$  corresponding wavelengths (micron)

$$\tau_{\text{coll vib}}^{-1} \sim 10^2 \text{ s}^{-1}/\text{torr}$$

$$\tau_{\text{coll rot}}^{-1} \sim 10^3 \text{ s}^{-1}/\text{torr}$$

$$T_2^{-1} = 7 \text{ MHz/torr (Lamb dip)}$$



$$E_{\text{rot}} = \frac{\hbar^2}{2I} J(J+1)$$

$$\nu = \Delta E \sim J+1$$

$$\Delta \nu \sim \text{cm}^{-1}$$

$$\Delta \nu_D \sim 50 \text{ MHz}$$

$$\Delta \nu_{\text{rot}} = 50 \text{ GHz}$$

***Dipole moments and linewidths  
of laser materials***

**Let us** evaluate the dipole source due to a radiating atom.

It will be the expectation value of the dipole operator over the atomic state  $|\psi\rangle$ , i.e.,

$$\langle d \rangle = \mathbf{e} \langle \mathbf{r} \rangle = \mathbf{e} \langle \psi | \mathbf{r} | \psi \rangle$$



$$\psi = |s\rangle$$

$$\langle \mathbf{r} \rangle = 0$$



$$\psi = |p\rangle$$

$$\langle \mathbf{r} \rangle = 0$$



$$\psi = a|s\rangle + b|p\rangle$$

$$\langle \mathbf{r} \rangle \neq 0$$

$$\langle \mathbf{r} \rangle \equiv \int \vec{\mathbf{r}} |\psi(\mathbf{r})|^2 d^3\mathbf{r}$$

Fig. 1.8

Atomic wavefunctions for pure  $s$  and  $p$  states and for a linear combination with amplitudes  $a$  and  $b$

When interacting with a constant E field, an atom is driven back and forth in a reversible way as shown in fig. 1.9 at a rate

$$\Omega = \mu E / \hbar \quad (\text{CmVm}^{-1} / \text{Js} = \text{s}^{-1}) \quad (1.21)$$

where  $\mu = e \langle s | r | p \rangle$  is the transition matrix element (taken for simplicity as a real number) and  $\Omega$  is called the Rabi frequency. The wavefunctions at different times are sketched in fig. 1.9 and any time the instantaneous wavefunction  $\psi(t)$  is given by

$$|\psi(t)\rangle = a(t)|s\rangle + b(t)|p\rangle, \quad (1.22)$$

where  $a(t)$  and  $b(t)$  have sinusoidal variations as shown in the figure. Correspondingly the induced dipole

$$\langle d \rangle = \mu (a^* b + ab^*) \quad (1.23)$$

*[because of the quantum evolution of a and b due to the field E]*

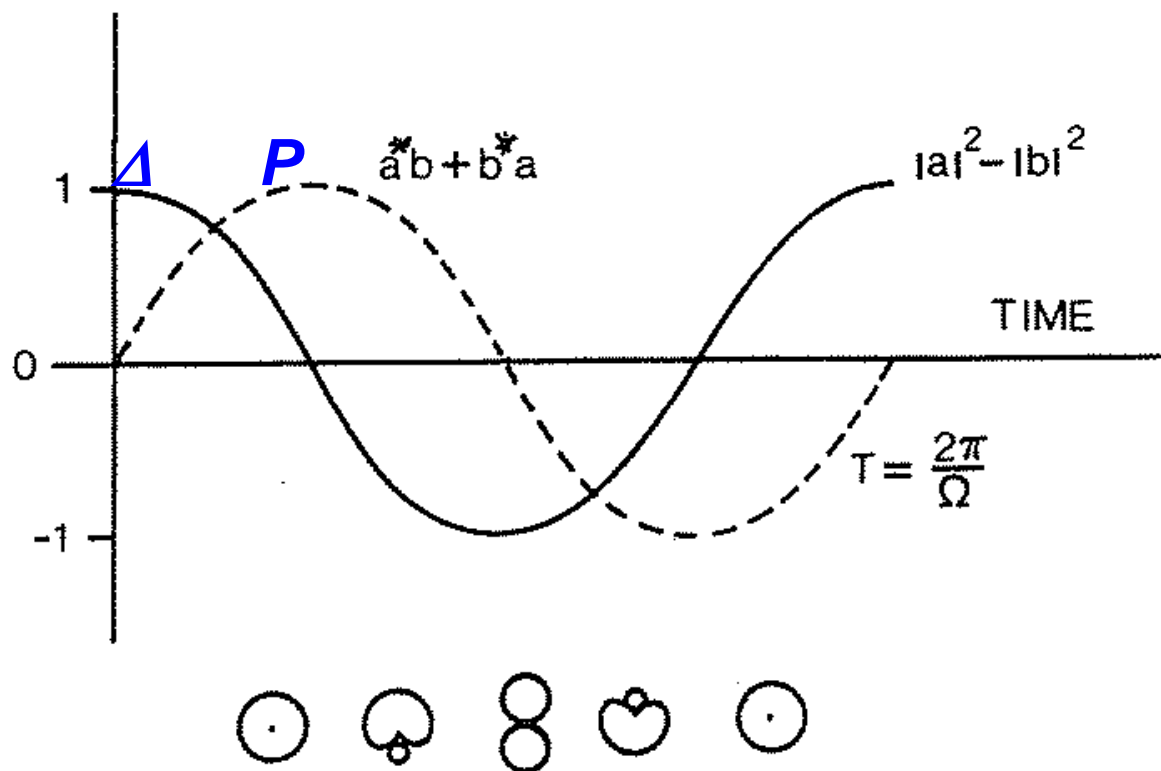


Fig. 1.9

Population difference (solid line) induced dipole (dashed line) and atomic orbitals (ff. below) for different time during a Rabi period

goes as the dashed line in fig. 1.9. As shown in eq. (1.23) it depends on the phase of the wavefunction, whereas the population inversion  $|b|^2 - |a|^2$  is phase independent. Both  $|a|$  and  $|b|$  return to equilibrium values  $|a| = 1$ ,  $b = 0$  by spontaneous emission processes

. On a faster scale, phase destroying processes, as collisions, interrupt the coherent Rabi precession as shown in fig. 1.10. The average interruption time is called  $T_2$ , and its reciprocal is the homogeneous linewidth

$$1/T_2 = \Delta\nu_a \geq \Delta\nu_{sp}$$

Since  $\langle d \rangle / \mu = \Omega T_2$ , then the average dipole is  $\langle d \rangle = \frac{\mu^2 E T}{h}^2 = \frac{\mu^2 E}{h \Delta\nu}$

and the polarization ( $\rho$  being the atomic density)

$$P = \rho \langle d \rangle = \frac{\rho \mu^2}{h \Delta\nu} E \quad (1.24)$$



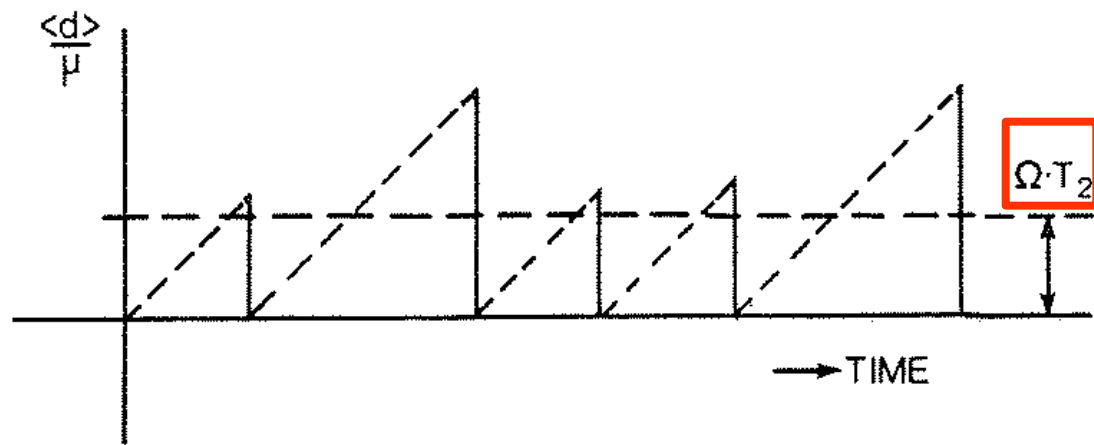


Fig. 1.10

*Interruptions due to decay processes, giving a non zero polarization*

## *The 3 so-called Maxwell-Bloch equations ruling field-atom interaction*

$$\left\{ \begin{array}{l} \Delta = \cos \Omega t \\ P = \sin \Omega t \end{array} \right. \quad \Omega = \mu E / \hbar$$

$$[\Delta^2 + P^2 = 1 (\text{circle})]$$

$$\left\{ \begin{array}{l} dP/dt = \Omega \Delta \end{array} \right. \quad (2)$$

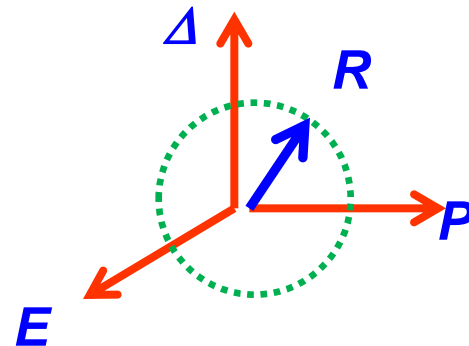
$$\left\{ \begin{array}{l} d\Delta/dt = -\Omega P \end{array} \right. \quad (3)$$

*On the other hand, from Maxwell, E evolution linear in P:*

$$dE/dt = P \quad (1)$$

*Thus, interaction atom-field as the motion of a vector R on a 3-D space (precession of R around E*

*at angular rate  $\Omega = \mu E / \hbar$  )*



## **Lasers of Class A, B and C: how time scales rule the dynamics**

**Couple e.m. field in cavity with  $N$  atoms.**

**Keeping only the first mode  $E$  which goes unstable,  $E$  has as**

**linear** source a polarization  $P$ .  $P$  depends on both  $E$  and population inversion  $\Delta$  in a **nonlinear way** (terms in  $\bigcirc$ ) as shown by Maxwell-Bloch eqs.

**Furthermore, we add dissipative terms to account for:**

**i) Losses of  $E$ ,  $P$  and  $\Delta$ , at rates  $k$ ,  $\gamma_{\perp}$ ,  $\gamma_{\parallel}$  respectively**

**ii) incoherent supply of energy (pump)**

$$\dot{E} = -kE + gP,$$

$$\dot{P} = -\gamma_{\perp}P + gE\Delta,$$

$$\dot{\Delta} = -\gamma_{\parallel}(\Delta - \Delta_0) - 4gPE.$$

Here,  $k$ ,  $\gamma_{\perp}$ ,  $\gamma_{\parallel}$  are the loss rates for **field**, **polarization** and **population**;  $g$  is a coupling constant and  $\Delta_0$  is the population inversion that would be established by the pump, in absence of coupling with  $E$ .

While the terms in  $\left( \right)$  are due to field-atom coupling, the terms  $k$ ,  $\gamma_{\perp}$ ,  $\gamma_{\parallel}$  represent dissipation, that is, coupling with the environment (precisely, the field escapes from the cavity; and polarization  $P$  and population  $\Delta$  decay at different rates

The following classification has been introduced ([Arecchi, #111](#))

*Class A* (e.g., He–Ne, Ar, Kr, dye):  $\gamma_{\perp} \simeq \gamma_{\parallel} \gg k$

The two last equations (1) can be solved at equilibrium (adiabatic elimination procedure) and one single nonlinear field equation describes the laser.  $N = 1$

*Class B* (e.g., ruby, Nd, CO<sub>2</sub>):  $\gamma_{\perp} \gg k \lesssim \gamma_{\parallel}$

Only polarization is adiabatically eliminated [middle eq. (1)] and the dynamics is ruled by two rate equations for field and population.  $N = 2$  allows also for self-excited oscillations.

*Class C* (e.g., FIR lasers):  $\gamma_{\parallel} \approx \gamma_{\perp} \simeq k$

The complete set of eqs. (1) has to be used, i.e.

## **Class A: single eq.**

$$dE/dt = f(E)$$

where

$$f(E) = -\alpha E$$

*below*

$$f(E) = +\alpha E - \beta |E|^2 E$$

*above*

} *threshold*

## **Class B: two coupled eqs:**

$$\left\{ \begin{array}{l} dE/dt = f(E, \Delta) \\ d\Delta/dt = g(E, \Delta) \end{array} \right.$$

or, calling  $I = E^2$  the intensity

$$\left\{ \begin{array}{l} dI/dt = F(I, \Delta) \\ d\Delta/dt = G(I, \Delta) \end{array} \right.$$

*that in fact are the rate eqs. for  $n = I$  (photon  $n^\circ$   $n$  prop. to intensity  $I$ ) and  $N = \Delta$ , given beforehand*

Table 2 summarizes the relevant parameters for the most common lasers

**[remember  $T_{cavity}=10^{-6}$  sec]**

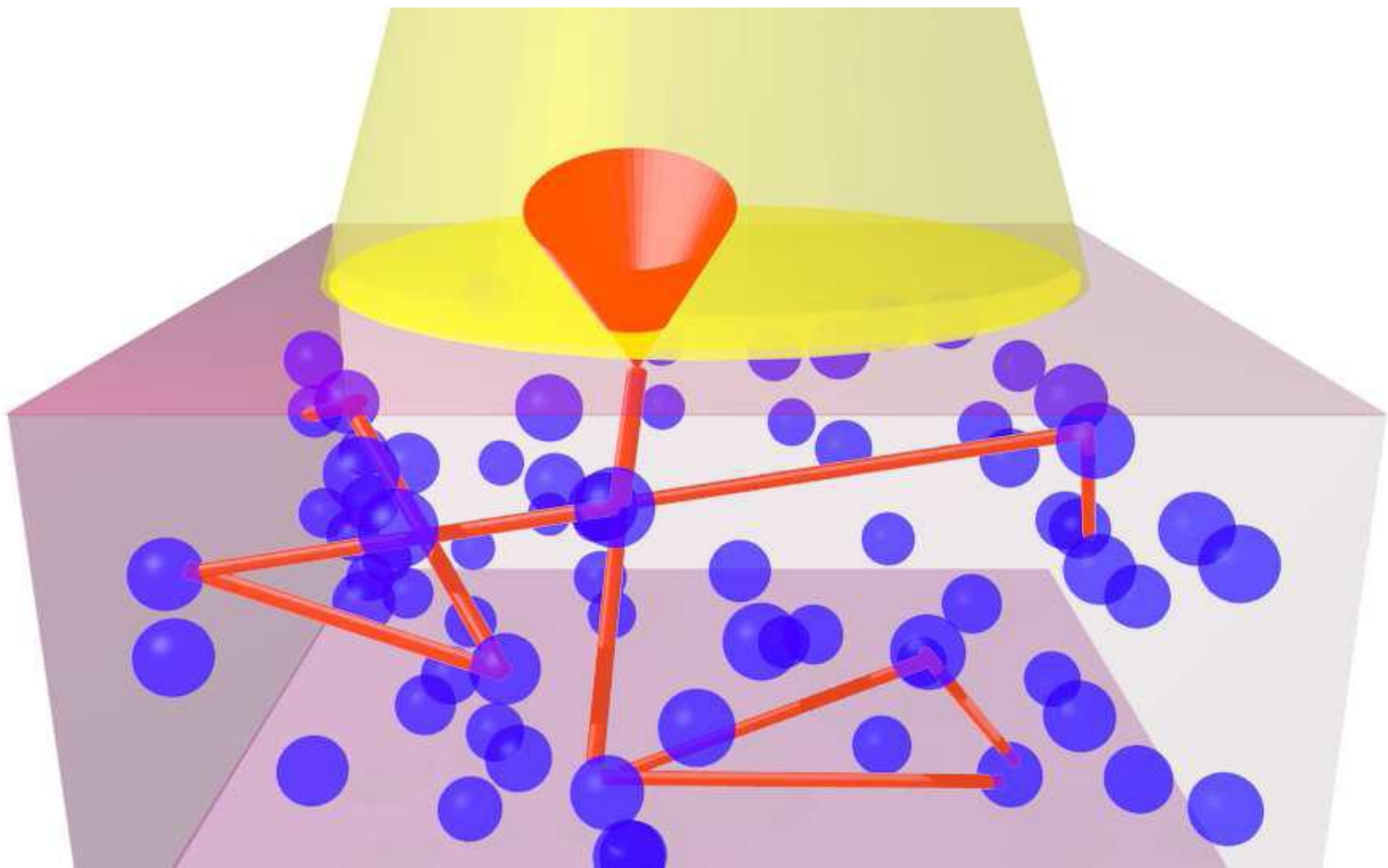
	Ruby Nd	Dye	He-Ne	Ar <sup>+</sup>	CO <sub>2</sub>	Semi- conduct.
$\tau_{sp}$ (sec)	$10^{-3}$	$10^{-8}$	$10^{-8}$	$10^{-8}$	$0.2^{(o)}$	$10^{-9}$
$\tau_2$ (sec)	$10^{-12}$	$10^{-13}$	$10^{-8}$	$10^{-8}$	$(4.4 \times 10^7 / \text{torr})^{-1}$	$10^{-14}$
$\tau_2^*$ (sec)	$10^{-10}$	$10^{-13}$	$10^{-10}$	$\frac{1}{3} 10^{-10}$	$(3.14 \times 10^8)^{-1}$	--
$\sigma (\text{cm}^2)$	$3 \times 10^{-20}$	$3 \times 10^{-16}$	$10^{-12}$	$10^{-12}$	$10^{-15}$	$10^{-15}$
$\rho (\text{cm}^{-3})$	$10^{19}$	$10^{15}$	$10^9$	$10^{10}$	$10^{14}$	$10^{18}$
$\alpha (\text{cm}^{-1})$	0.3	3	$10^{-3}$	$10^{-2}$	0.1	50
$l (\text{cm})$	10	0.1	$10^{3(*)}$	$10^{2(*)}$	10	$3 \times 10^{-2}$

**[Laser for  
 $\exp(\alpha l) > 1$  ]**

Table 2

(o) The inversion decay time due to collisions is  $0.4 \times 10^{-2}$  s

(\*) High reflectivity mirrors reduce the cavity length to 10 - 100 cm

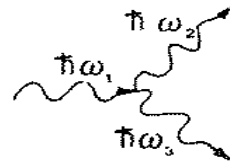


**The RANDOM LASER-** ZnO scatterers at random locations (blue) are optically pumped from above (wide yellow beam). The pumping yields an inversion of the atomic occupation number causing stimulated emission (orange light paths). The emitted intensity multiply scatters and concentrates . At the laser threshold the system experiences a phase transition and coherent intensity escapes the system through its surfaces (orange cone). The scatterers radius is 600nm,  $\lambda = 723\text{nm}$  .

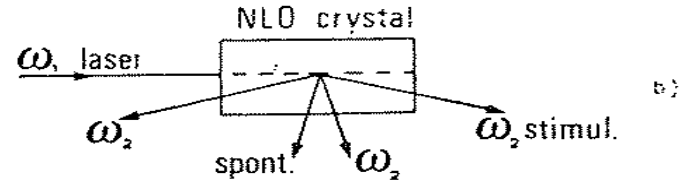


## ***2- NONLINEAR OPTICS***

# Spontaneous vs Stimulated



a)



b)

Spont. ( $\hbar\omega = E_n$ )  $\omega_1 = \omega_2 + \omega_3$

Stimul. ( $\hbar k = p$ )  $\vec{k}_1 = \vec{k}_2 + \vec{k}_3$

$P_i = \epsilon_0 \chi_{ij}^{(1)} E_j$  **Linear polarization**

$P_i = \epsilon_0 \chi_{ijk}^{(2)} E_j E_k$  **quadratic polarization**  
(3 quanta involved)

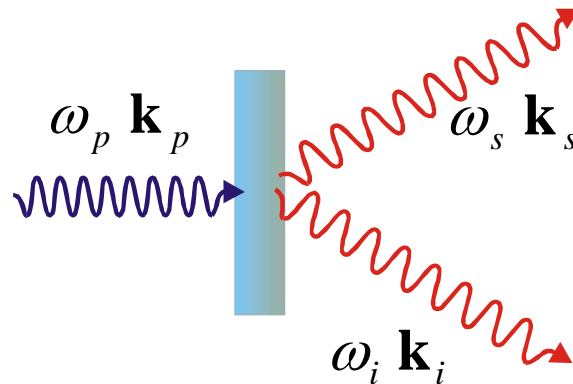
### 3- quanta NLO processes

Nature of the quanta		Name of the process
2	3	
light	molecular vibrations	Raman
light	optical phonons in solids	Raman
light	acoustical phonons in solids	Brillouin
light	sound waves in liquids	Brillouin
light	light	parametric conversion (sum or difference of frequency, second harmonic generation, etc.)

***Parametric down conversion-***

***Entangled photons***

**Spontaneous  
Parametric down-  
conversion (SPDC)**

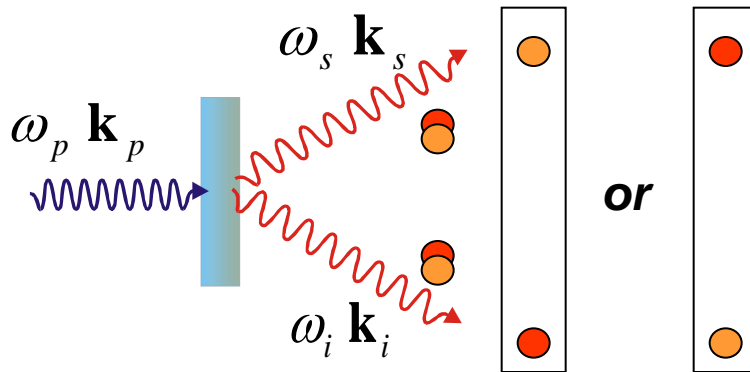


$$\omega_p = \omega_i + \omega_s$$

$$\mathbf{k}_p = \mathbf{k}_i + \mathbf{k}_s$$

**Energy and  
momentum  
conservation**

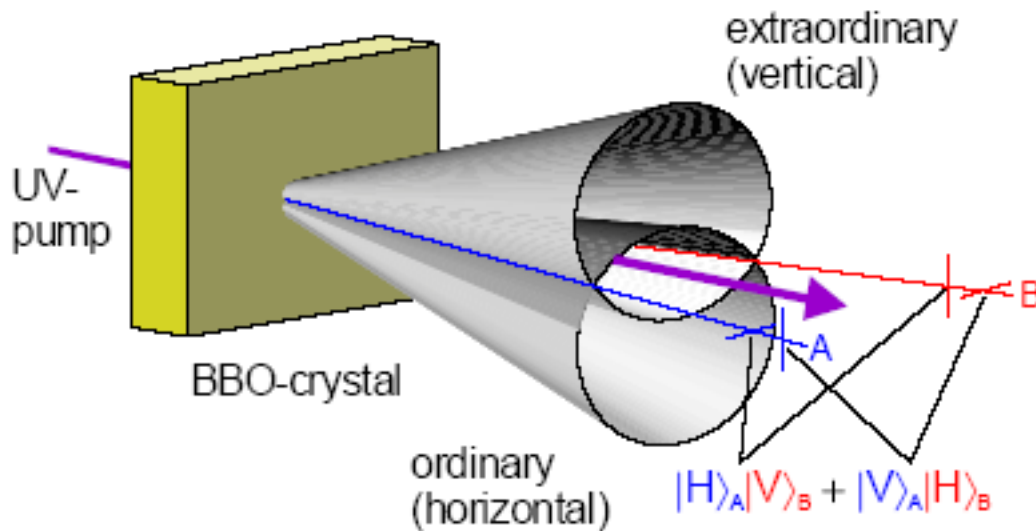
➡  $|\psi\rangle = (|\bullet\rangle_s |\bullet\rangle_i + |\bullet\rangle_s |\bullet\rangle_i)$  **SPDC Entangled state**



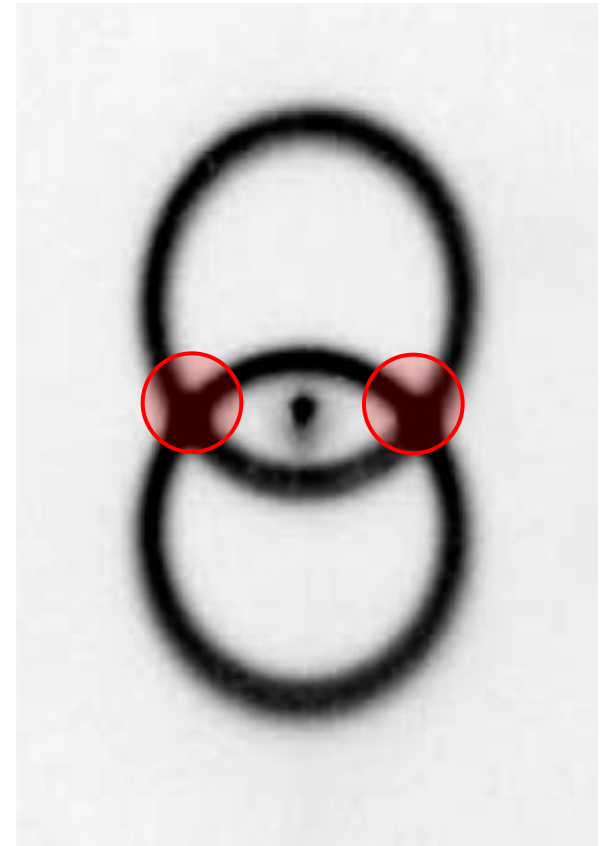
**The properties of a single  
photon are not defined  
individually  
but are completely correlated  
to those of the other**

# Polarization Entanglement

- In type II crystals:



$$|\psi\rangle = (|H\rangle_1|V\rangle_2 + |V\rangle_1|H\rangle_2)$$



4 photon (self actions):

- " self focusing
- " defocusing
- " phase modul. (broadening)
- " amplit. " (steepening)



$$P_i = \chi_{ijkl}^{(3)} E_j^* E_k E_l$$

$$n = n_0 + n_2 |E|^2$$

***3- COHERENCE,  
PHOTON STATISTICS,  
LASER PHASE TRANSITION***



## - Young interferometer

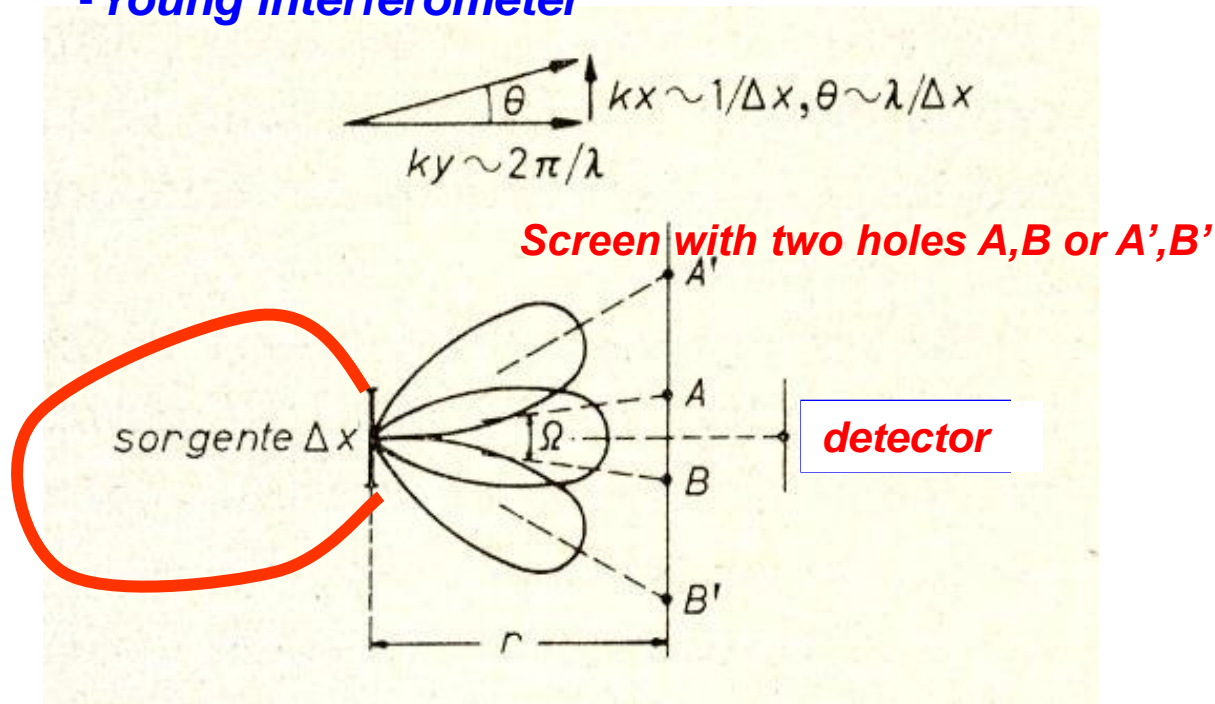


Photo-current proportional to square modulus of field

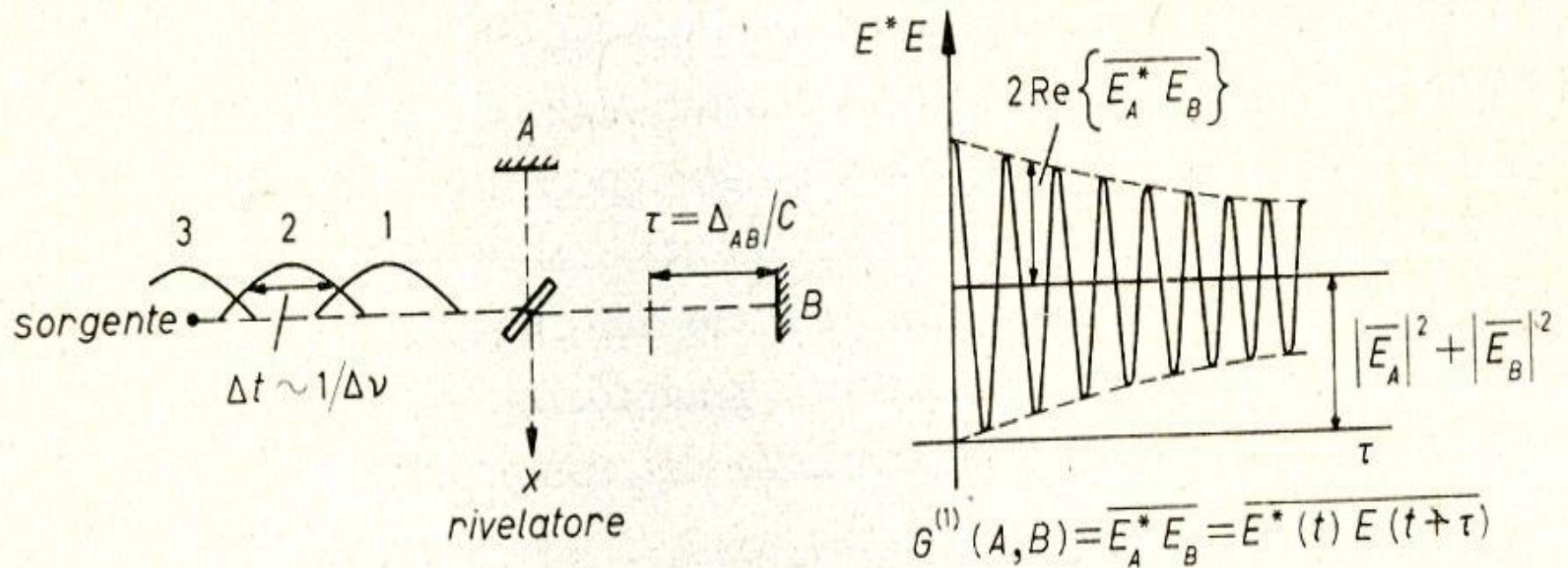
$$\langle |E_1 + E_2|^2 \rangle \longrightarrow I_1 = |E_1|^2 \quad I_2 = |E_2|^2 \quad \langle E^*_1 E_2 \rangle$$

1-st order correlation function  $G^{(1)}(1,2) = \langle E^*_1 E_2 \rangle$

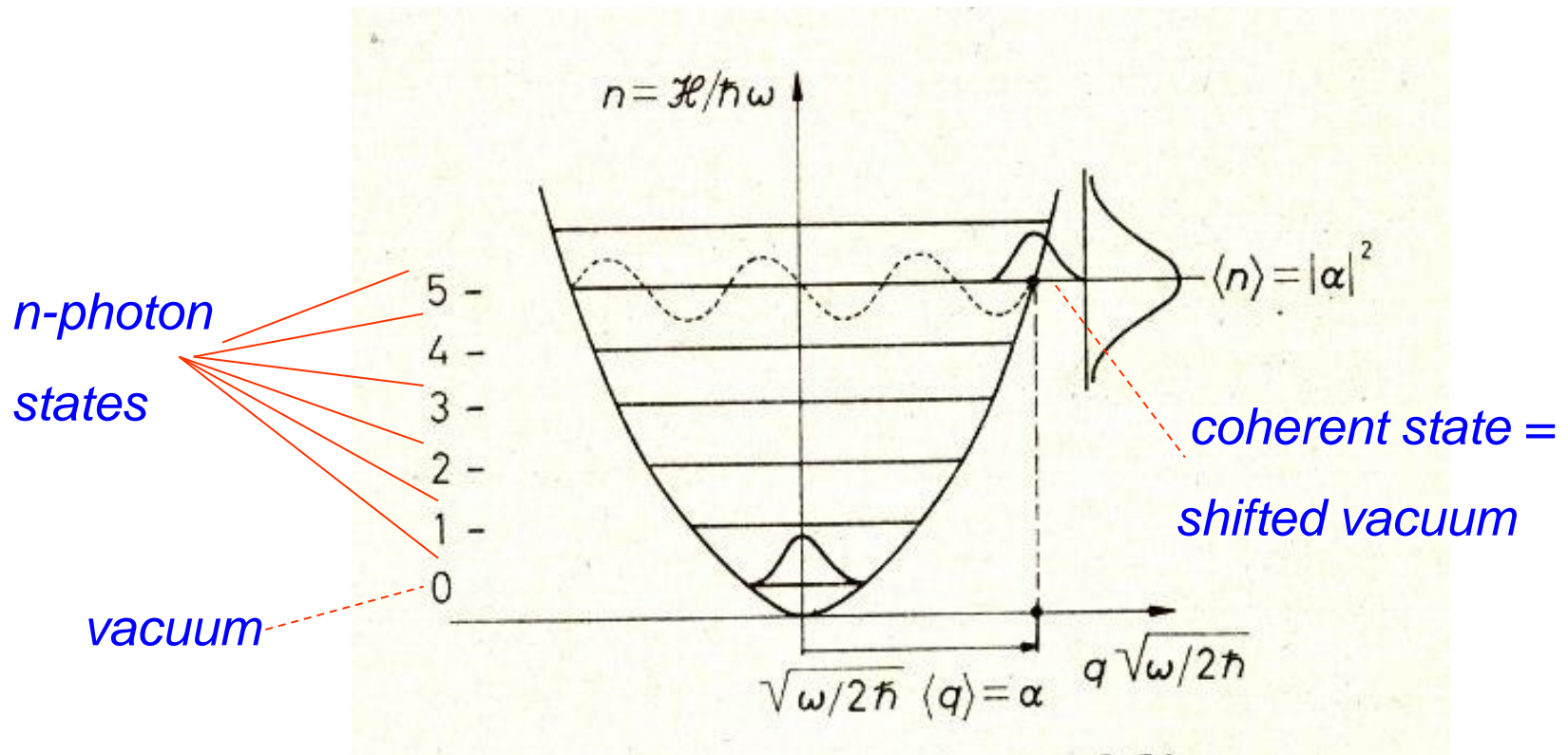
Coherence area

$$S_{AB} = \frac{\lambda^2 \cdot r^2}{(\Delta x)^2}$$

## Michelson Interferometer



## Harmonic Oscillator: energy $H$ versus coordinate $q$

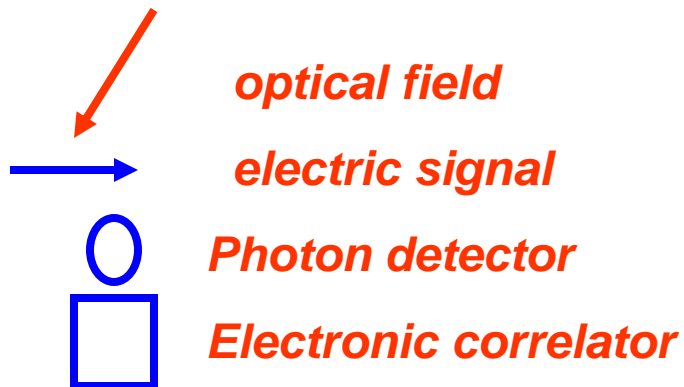
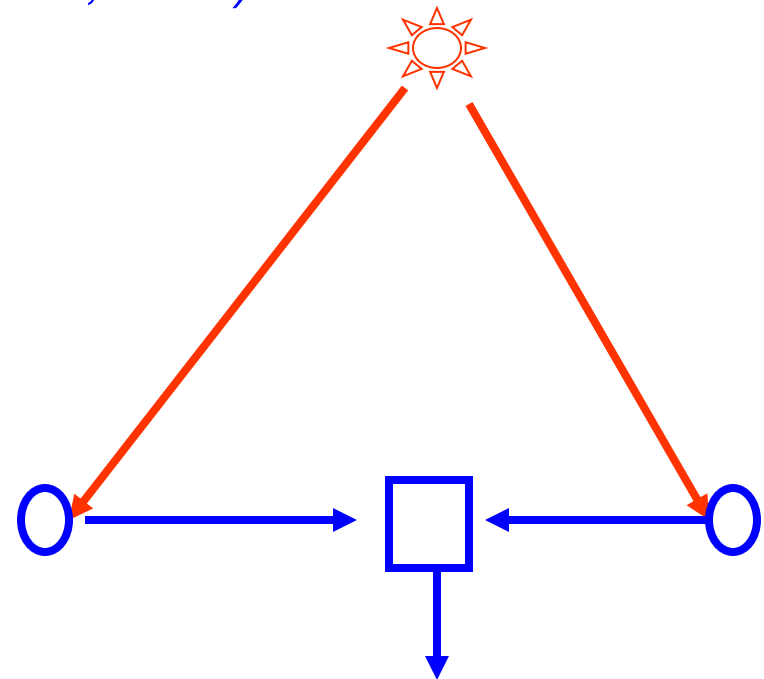
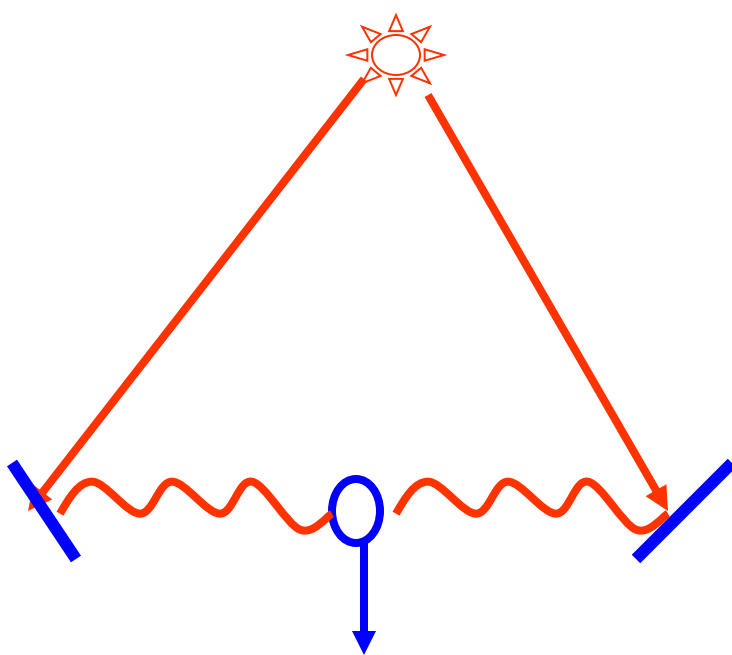


$$\Delta q \Delta p = (n + 1/2)\hbar$$

$$\Delta q \Delta p = 1/2\hbar$$

$$G^{(n)} = [G^{(1)}]^n,$$

*Stellar Interferometers: **field** (Michelson, 1925)  
and **intensity** (Hanbury-Brown & Twiss, 1956)*



## Hanbury-Brown & Twiss: intensity correlation

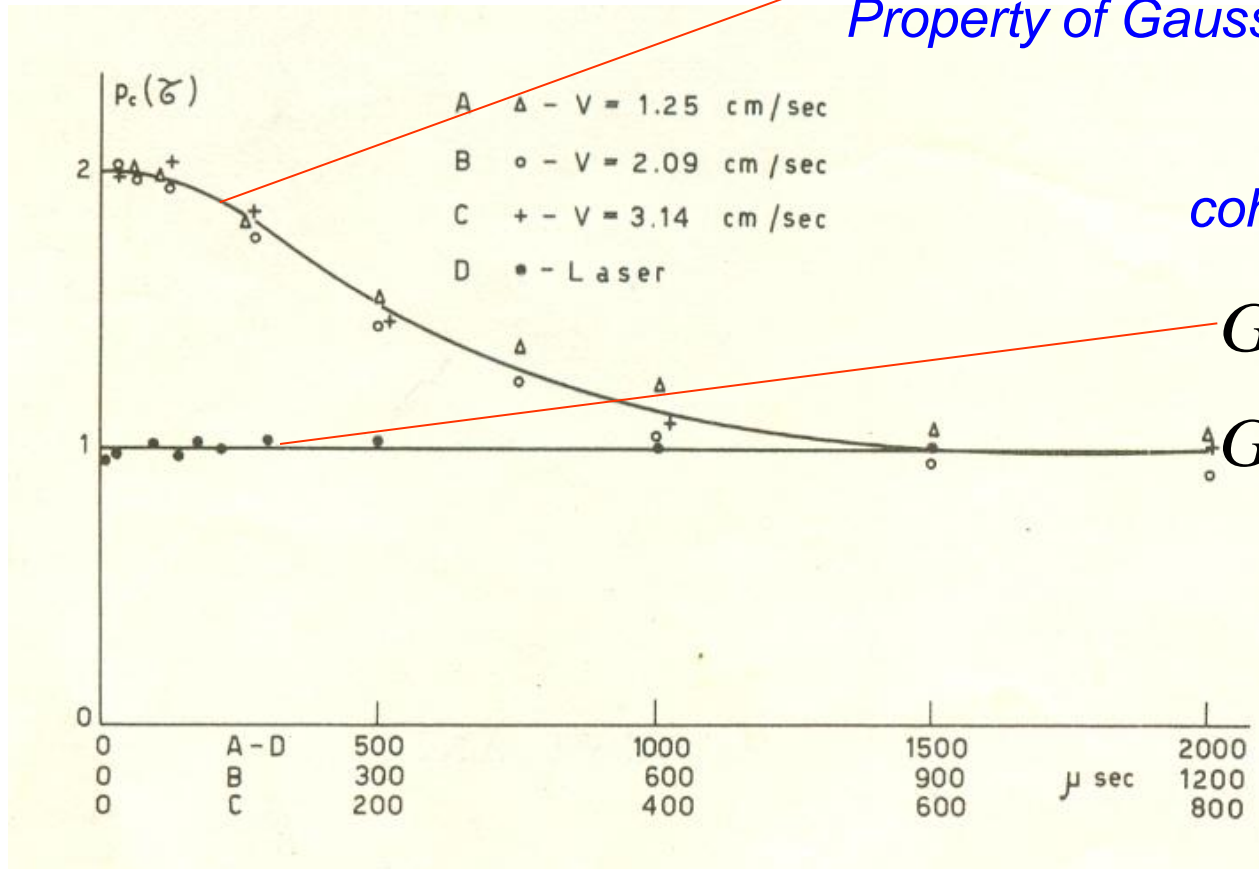
$$G^{(2)}(1,2) = \langle |E_1|^2 |E_2|^2 \rangle = I_1 I_2 + |G^{(1)}|^2$$

*Property of Gaussian processes*

*coherent state*

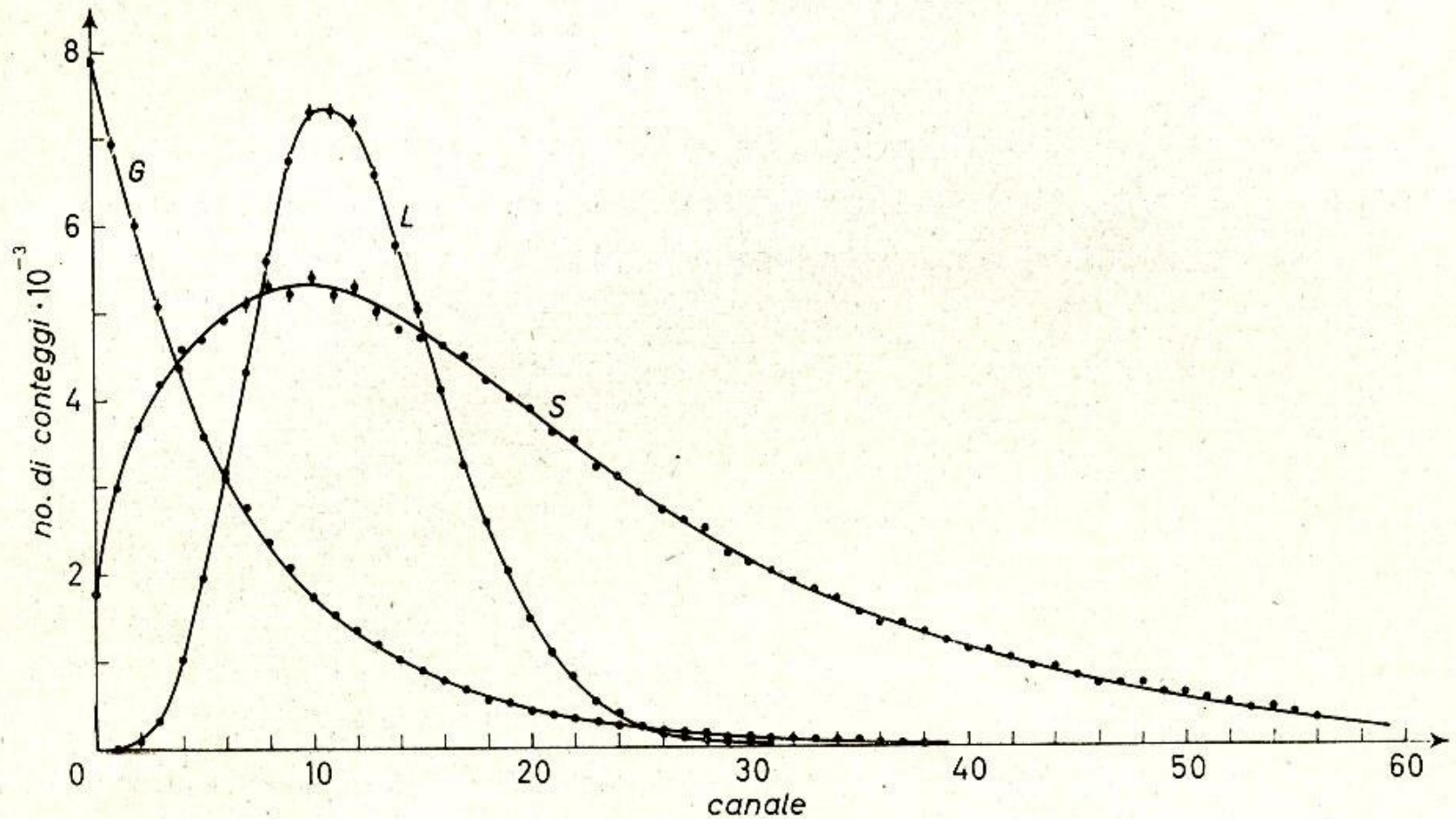
$$G^{(2)} = [G^{(1)}]^2,$$

$$G^{(n)} = [G^{(1)}]^n,$$



(Arecchi, Gatti, Sona-1966)

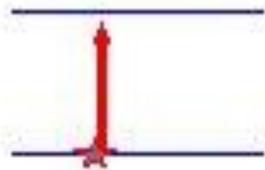
## Statistics of photocounts(Arecchi, 1965)



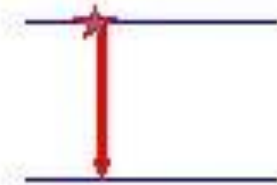
***L= laser; G= Gaussian light; S= superposition of L and G***

$P$  = polarization; energy  $W = -P \cdot E$

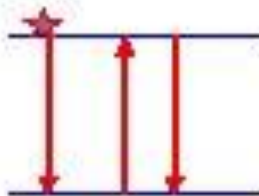
1



$$P = -\alpha E$$



$$P = \alpha E$$



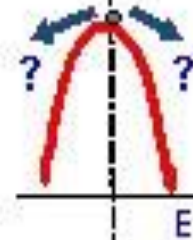
$$P = \alpha E - \beta E^3$$

$W(E)$

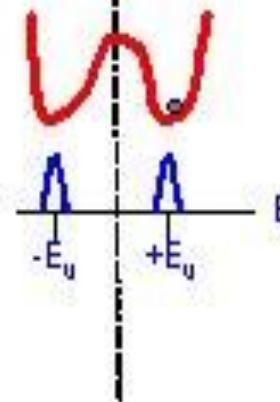
$p(E)$



**Stationary field  
probability**



$$p(E) = \exp(-W)$$





*Energy curves*

*$W(E)$  [red]*

*and*

*field probabilities*

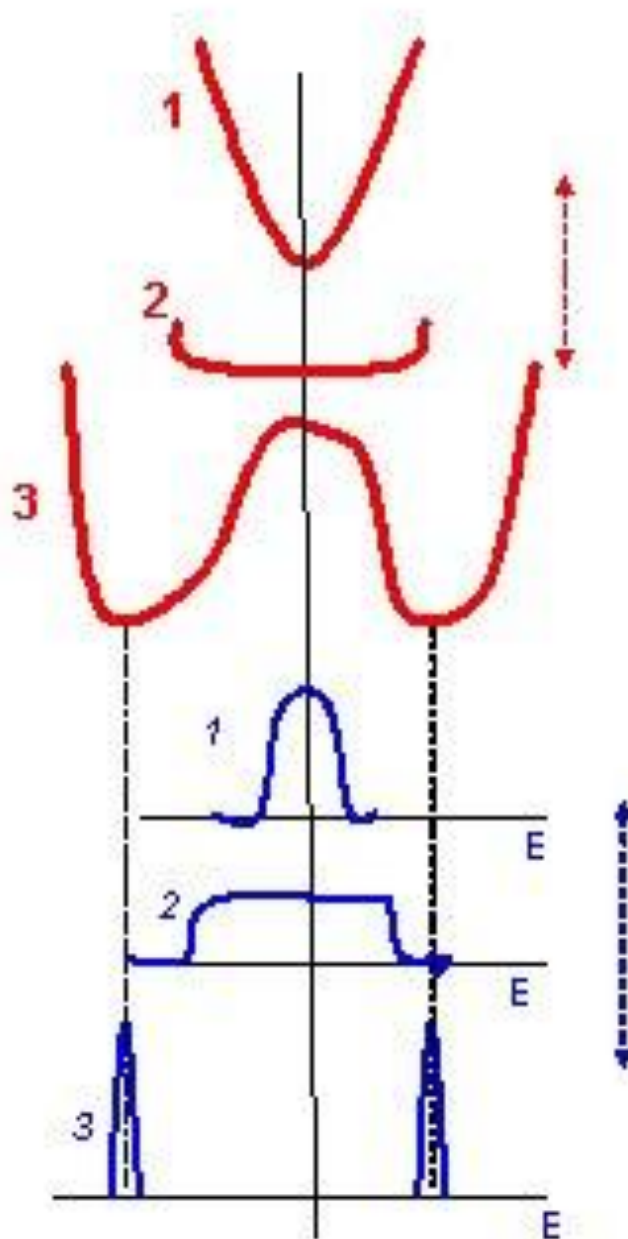
*$p(E)$  [blue]*

*1-below*

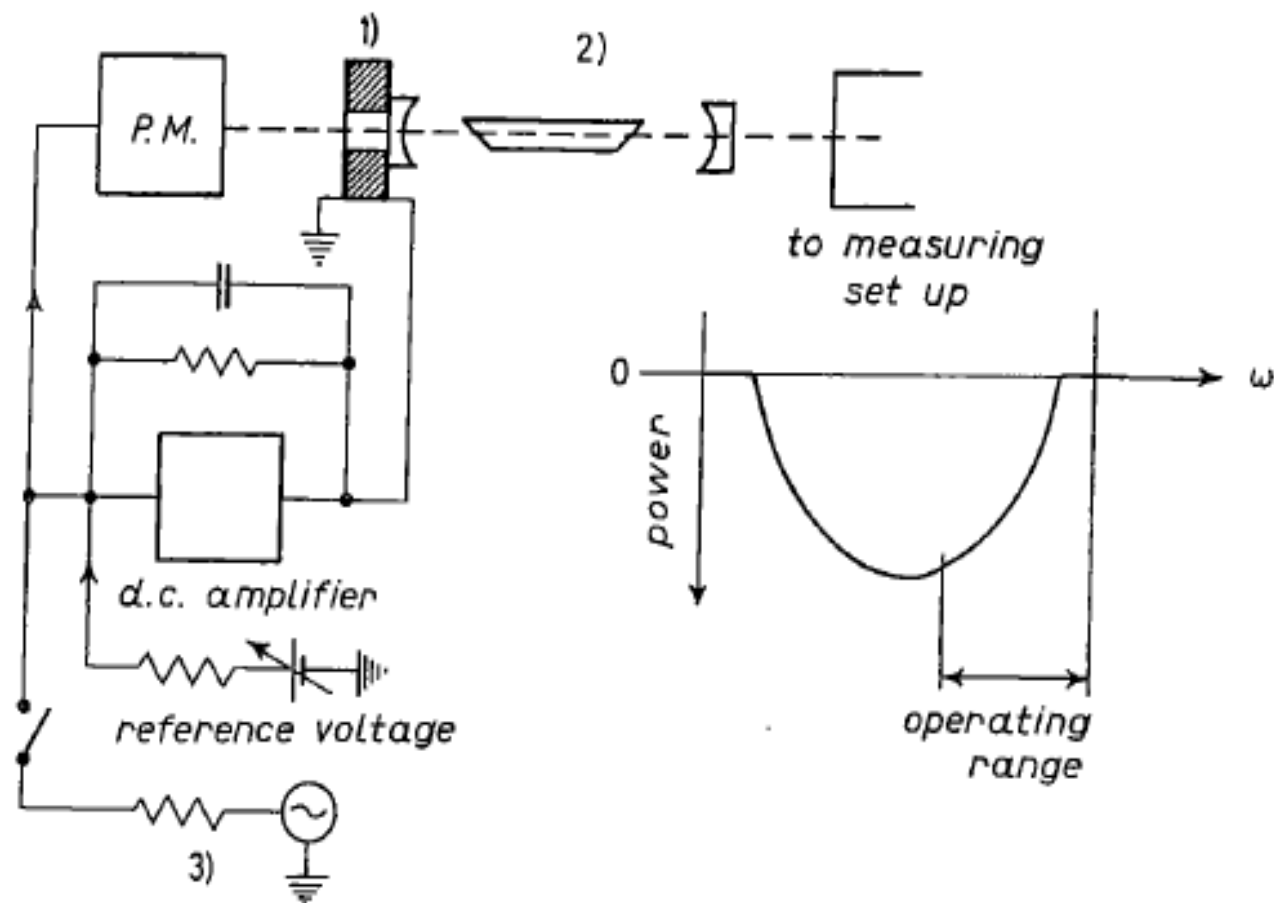
*2-at*

*3-above*

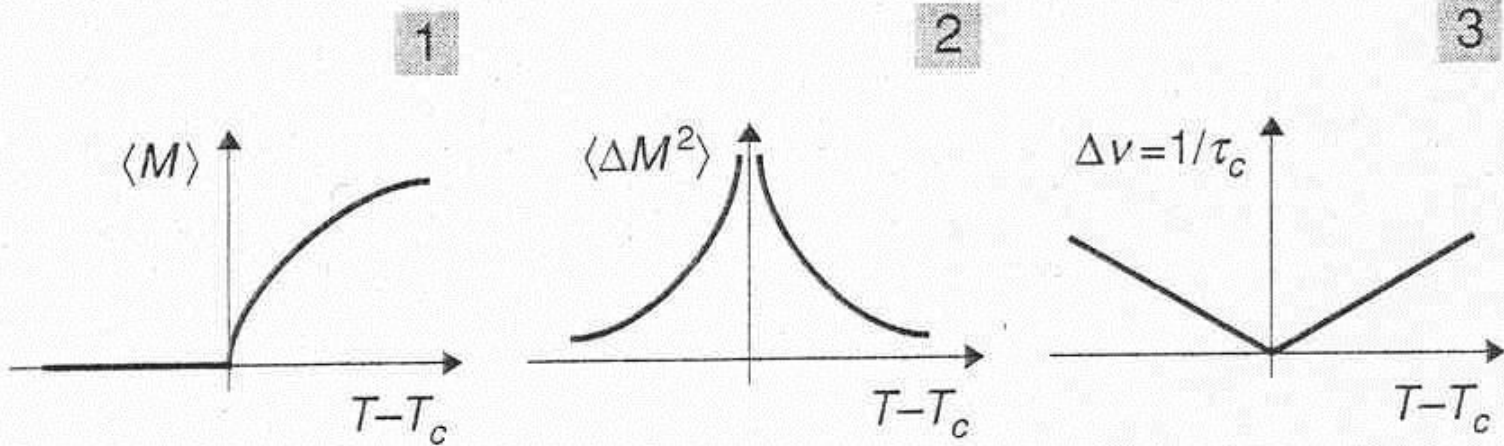
*threshold*







- Experimental set-up used for performing measurements of Gaussian noise superposed on a laser beam around threshold. 1) Piezoceramic disc; 2) laser length 20 cm, inner diameter 1 mm; 3) optional a.c. drive to control the operating range.

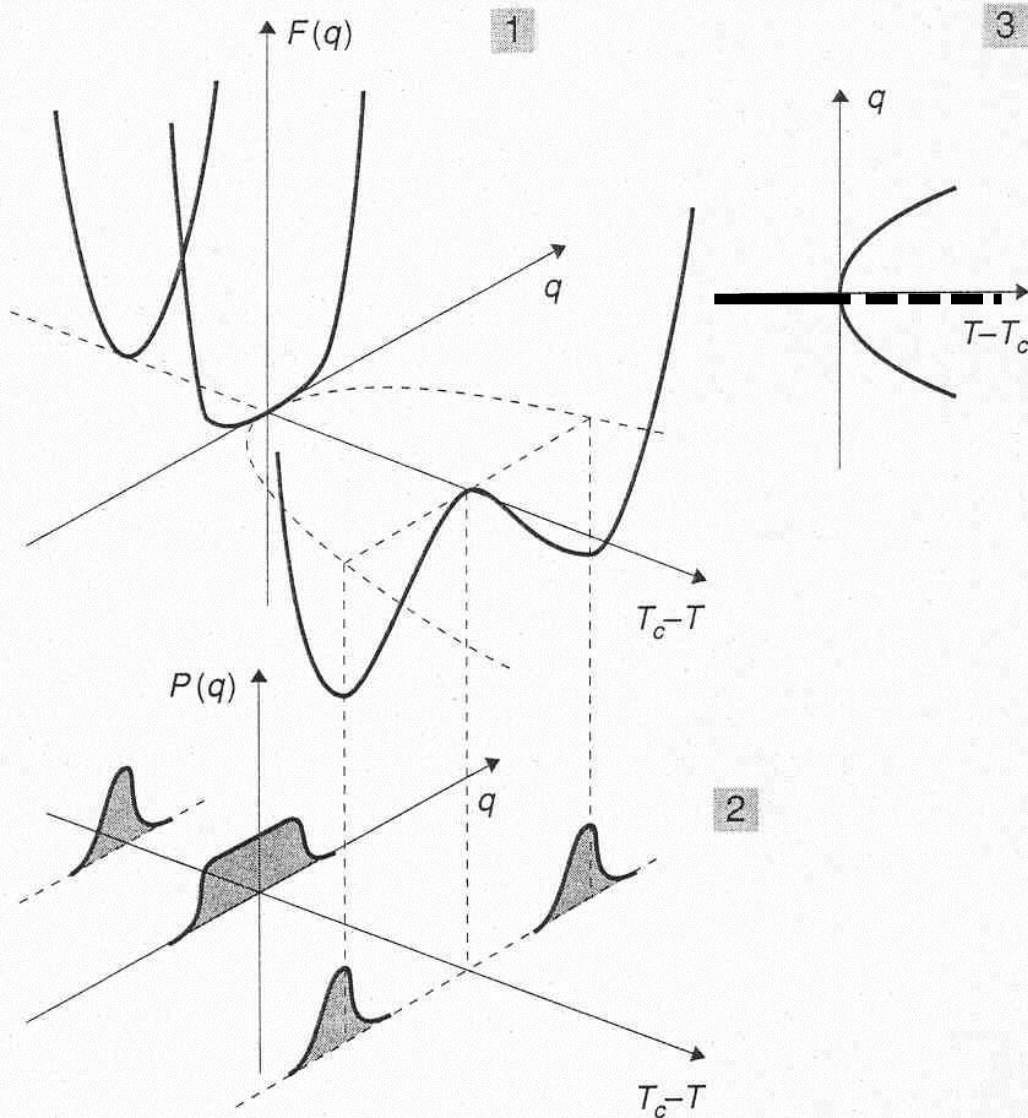


## **2nd order phase transition in magnetic material :**

$\langle M \rangle$  = order parameter (total magnetization),

$\langle \Delta M^2 \rangle$  = fluctuations (diverge at  $T_c$ )

$\Delta \nu = 1/\tau_c$  = linewidth of fluctuations (zero at  $T_c$  = critical slow down)



# Landau model

$F$  = free energy,

$q$  = order parameter,

$T$  = temperature,

$T_c$  = critical temperature.

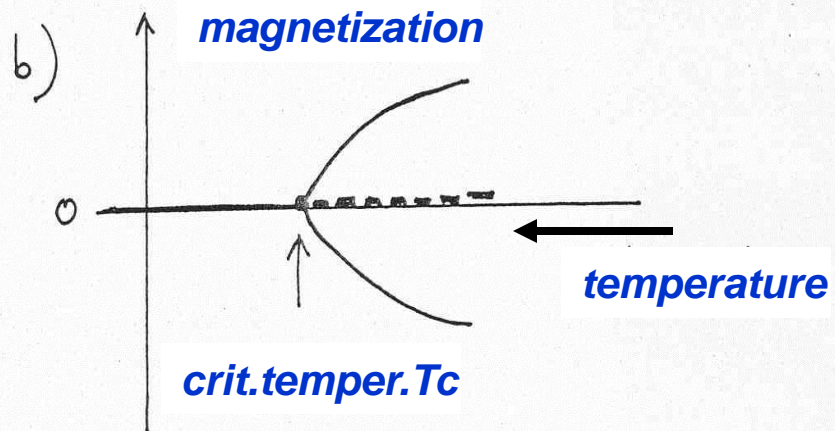
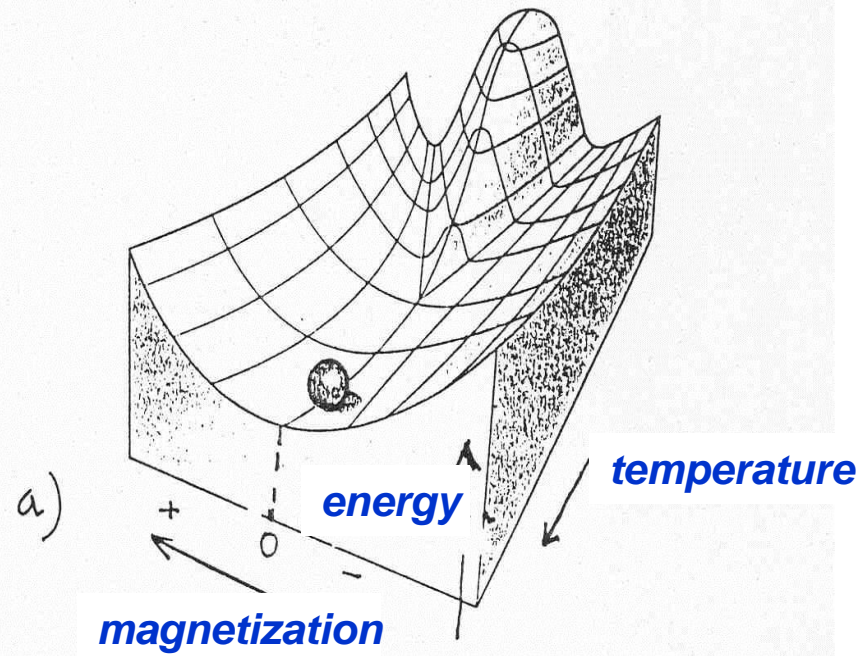
$P(q)$  = equilibrium

probability distribution  
of  $q$ .

$$P(q) = N \exp \left[ - F(q) / k_B T \right]$$

$$F = F_0(T) + \alpha q^2 + \beta q^4$$

$$\alpha = a(T - T_c) \dots \text{with} \dots a > 0$$



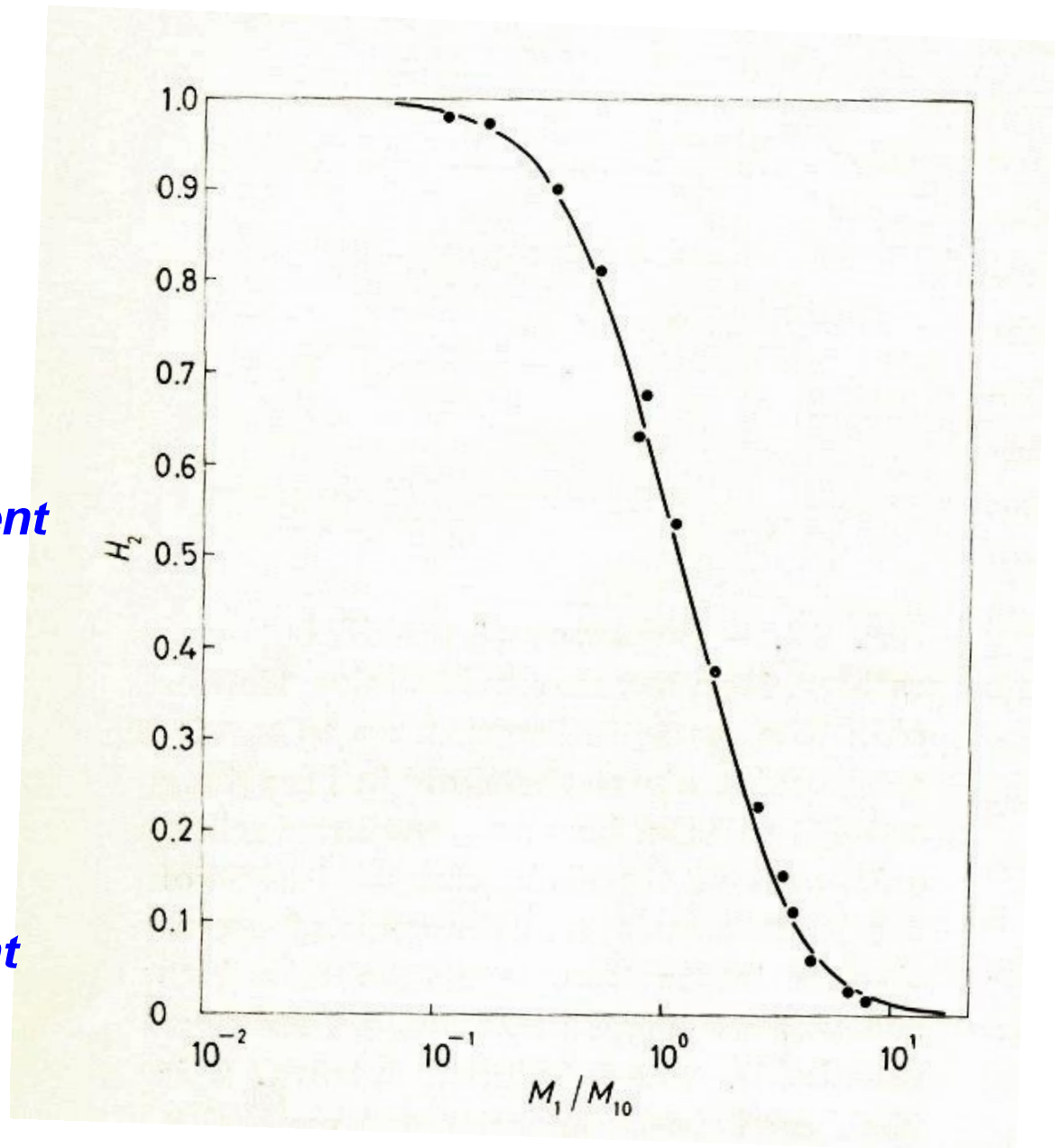


$H_2$  is a  
combination of first &  
second moment of  
photon distribution  
versus the first moment  
 $M_1$  normalized to its  
threshold value  $M_{10}$

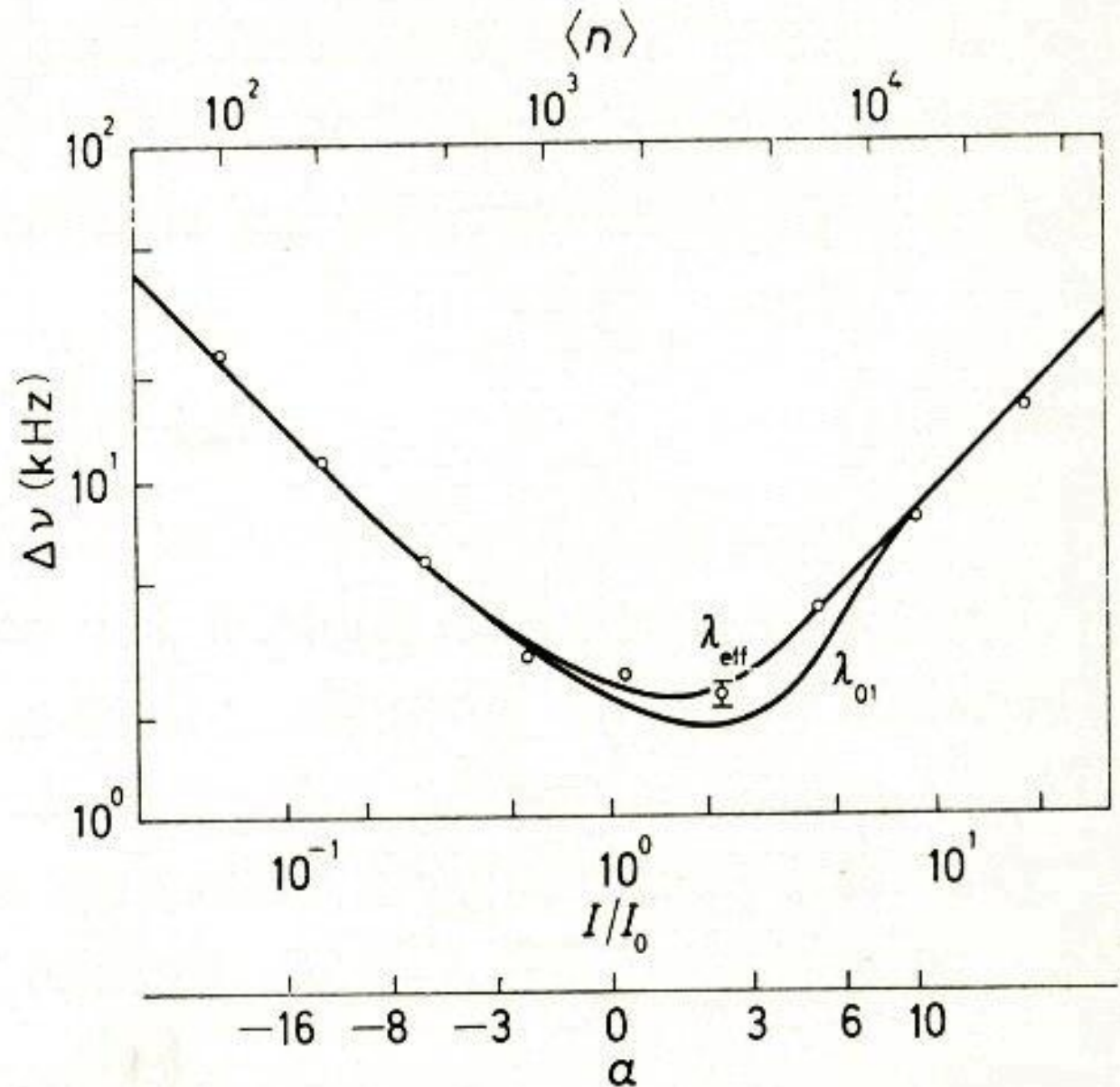
$H_2=1$  for Gaussian

$H_2=0$  for coherent light

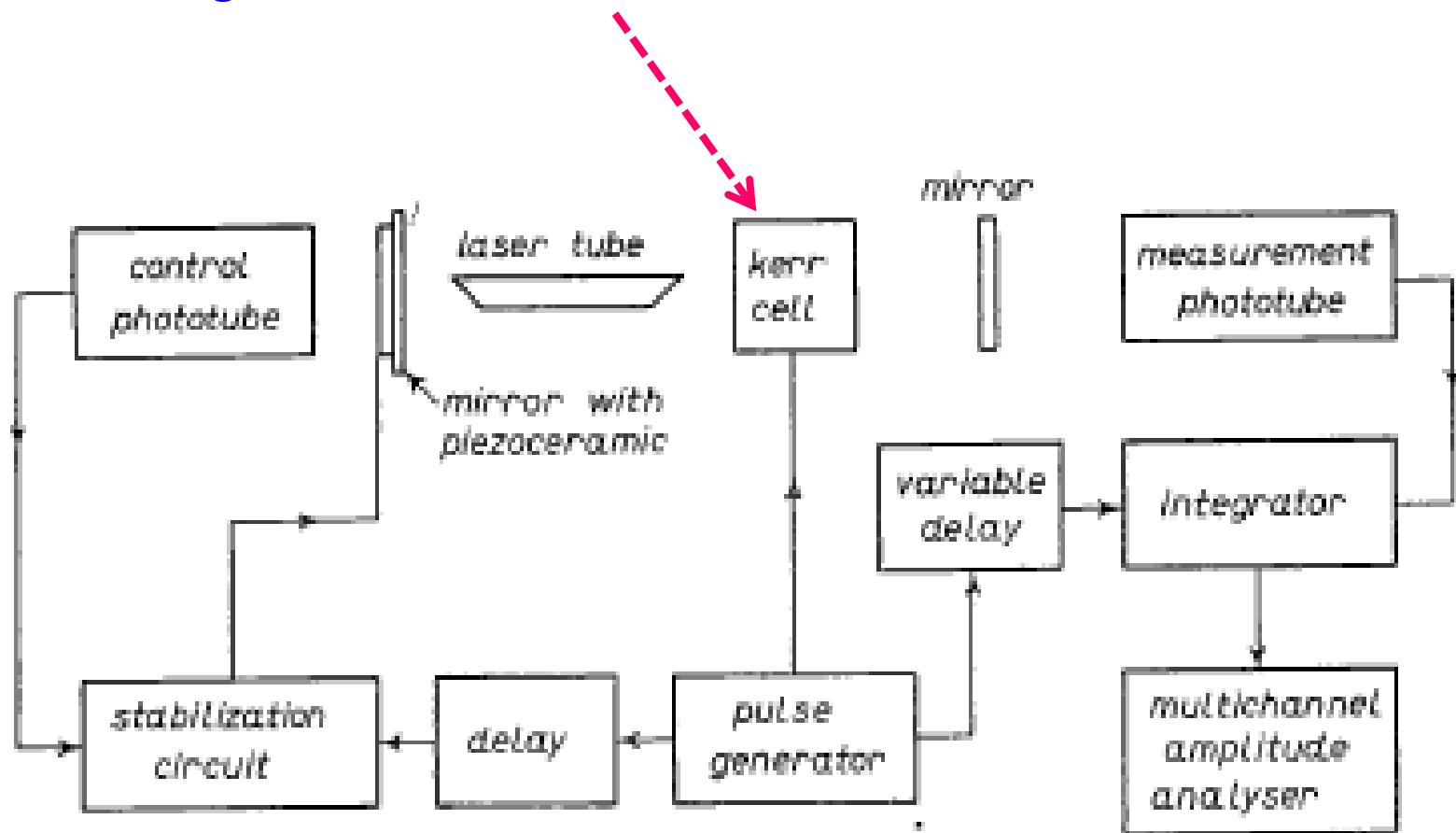
(Arecchi 1966)



Frequency width  
of laser light  
fluctuations  
versus  
laser intensity  
normalized to  
threshold value  
  
(Arecchi 1967)

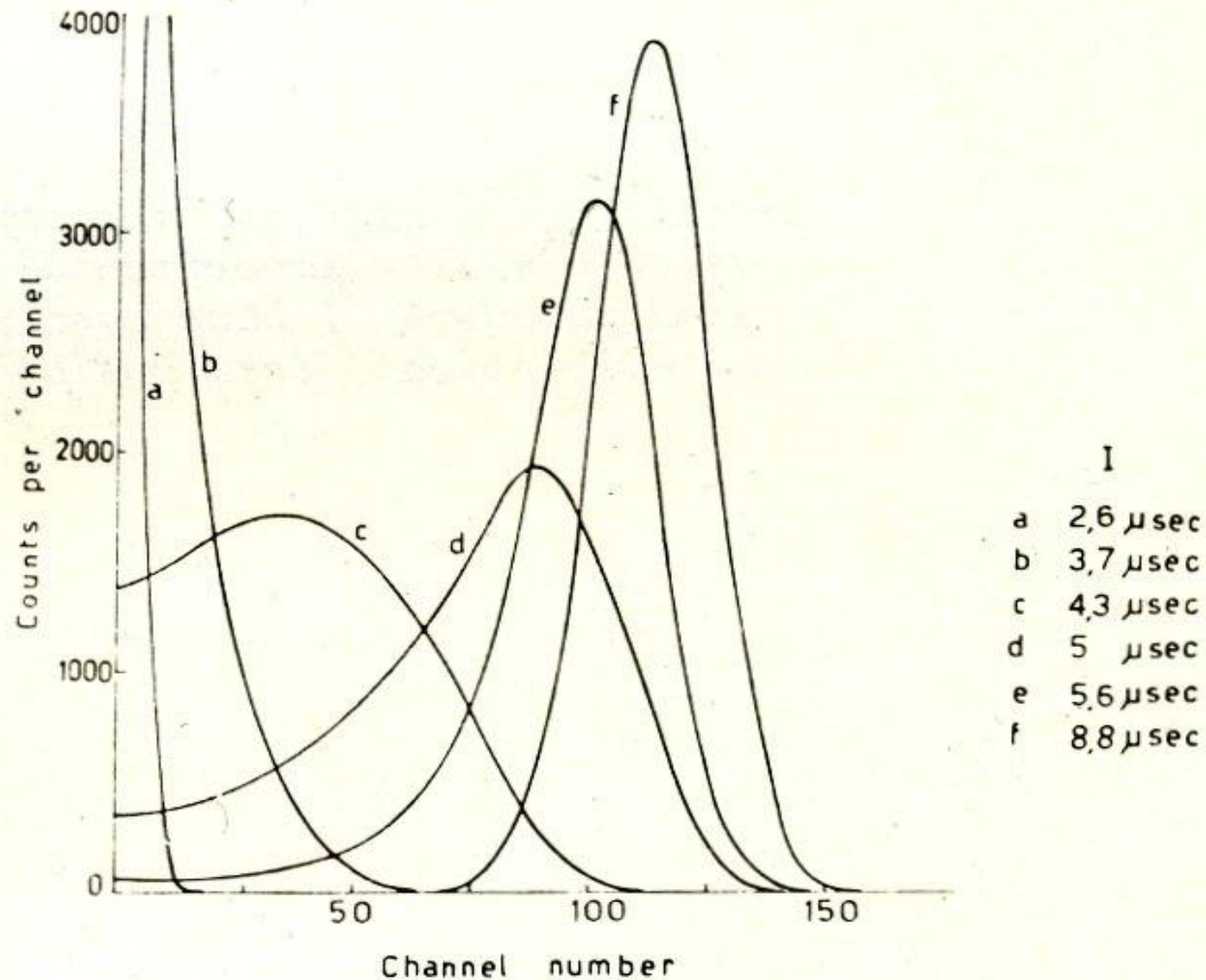


*Laser transient statistics: high gain; losses switched from high to low via a Kerr shutter*



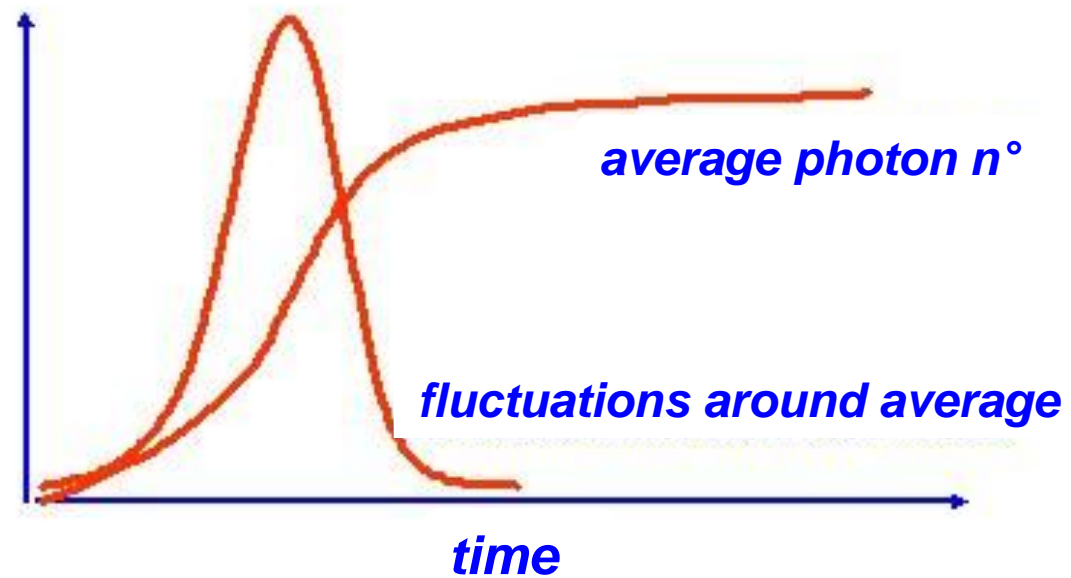
– Experimental set-up for the transient experiment.

**TRANSIENT STATISTICS:** Photon statistics sampled over 50 nanosec windows at different times after  $t=0$  (sudden onset of mirrors)





## *Transient photon statistics*



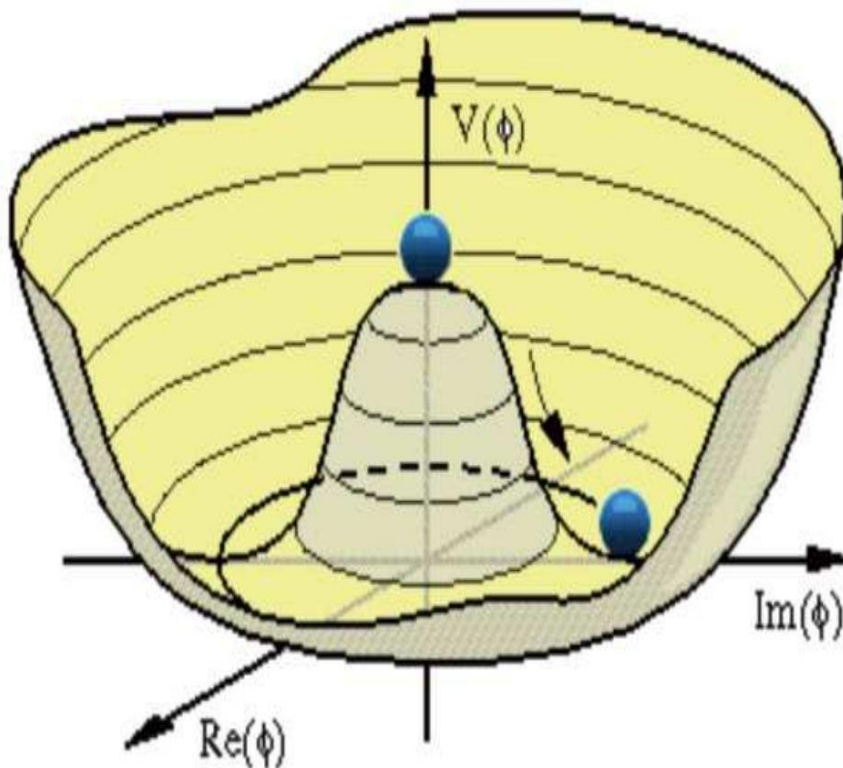
*( Arecchi et al.1967)*

# Higgs boson as laser with symmetry breaking in phase

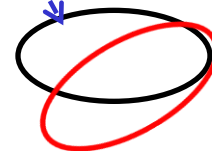
Lowering temperature, parabola inverts and ball falls, but it is confined in a potential “mexican hat”; no tangential constraints = **massless Goldstone boson**

As a laser above threshold, it has definite intensity (radius of circle) but phase undefined (angular position  $\Phi$  along circumference);

However from outside we can impose phase  $\Phi_0$  = minimum energy = **MASS**



$\text{Re } \Phi = x$  ,  $\text{Im } \Phi = y$



*Fields arising by symmetry breaking after Big-bang are massless ; in the Higgs field , tilting the circumference provides a mass (different from one particle to the other , as red and green) .*

