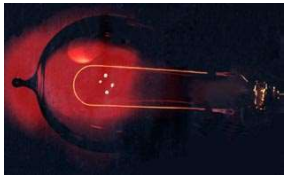
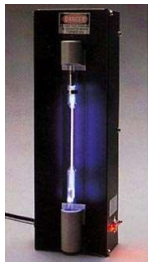
Generation of light – Light sources



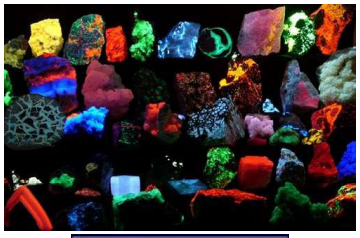


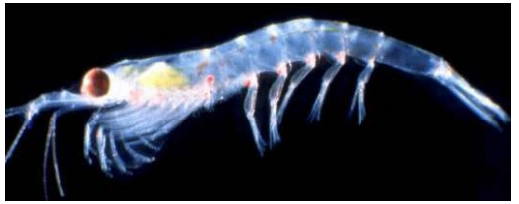
Luminescence

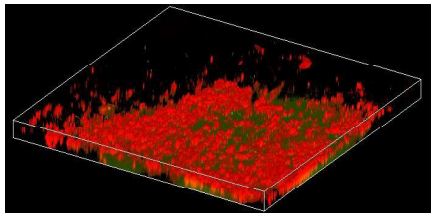
Laser

Black-body radiation

Luminescence







Repetition

- Types of energy states in atoms and molecules are independent (not coupled)
- Energy states are non-continuous, but discrete
- Transition between states involves packets (quanta) of energy

$$E_{total} = E_e + E_v + E_r$$

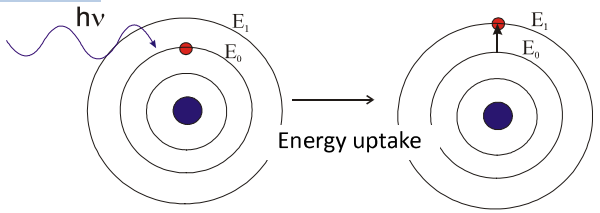
Scales of transition energies between different states are different:

$$E_e > E_v > E_r$$

Consider a **single atom**

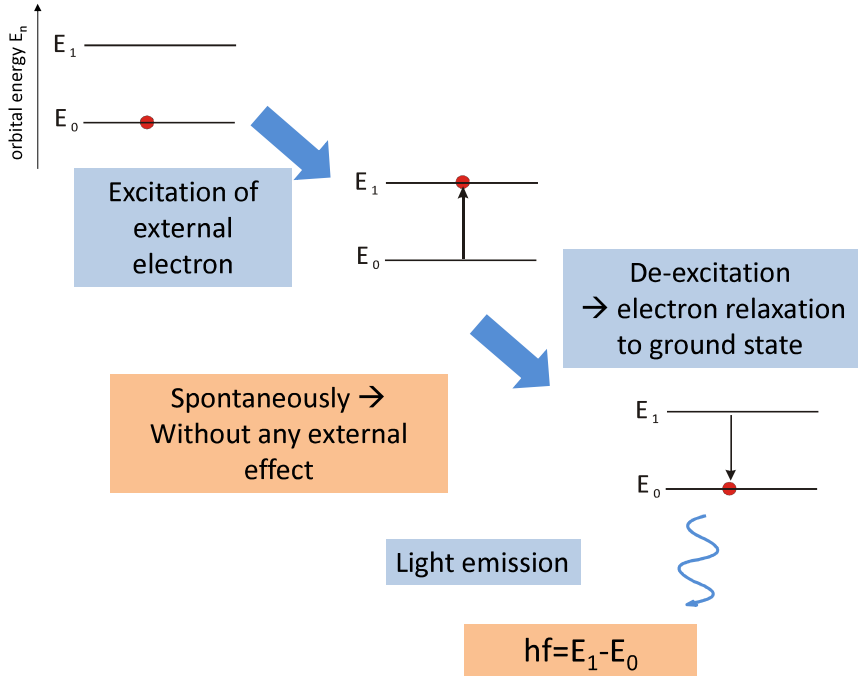
- Energy states are discrete
- Electrons occupy the lowest possible energy state (ground state)
- Pauli exclusion principle: no two identical fermions (particles with half-integer spin) may occupy the same quantum state simultaneously

Excitation energy

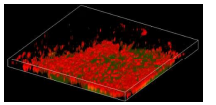


Ground state

Excited state



Excitation modes



- absorption of radiation (UV/VIS) : *photoluminescence*

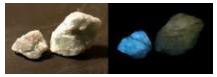
- chemical reaction: *chemo/bio-luminescence*

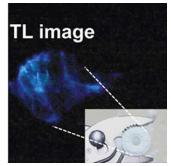
- Injection of charges: *electroluminescence*

- friction → mechanical deformation: *triboluminescence*

-thermally activated ion recombination: *thermoluminescence*

- Sound waves: *sonoluminescence*





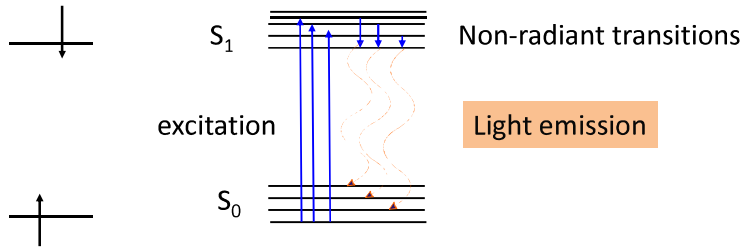
Luminescence:

spontaneous light photon emission by electrons when they return from their excited state to their original (ground) state of lower energy

$hf=E_1-E_0$

The emitted photon energy is characteristic for the electronic orbitals, thus for the atom/molecule.

The energy of the electronic orbitals in molecules is perturbed by the discrete states of molecular vibrations



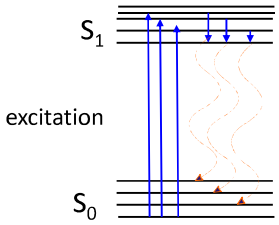
First excited singlet state

the spin states of the electrons

sum of all e^- spin quantum numbers=0

Fluorescence

De-excitation by photon emission between singlet states



Fluorescence

De-excitation by photon emission between singlet states

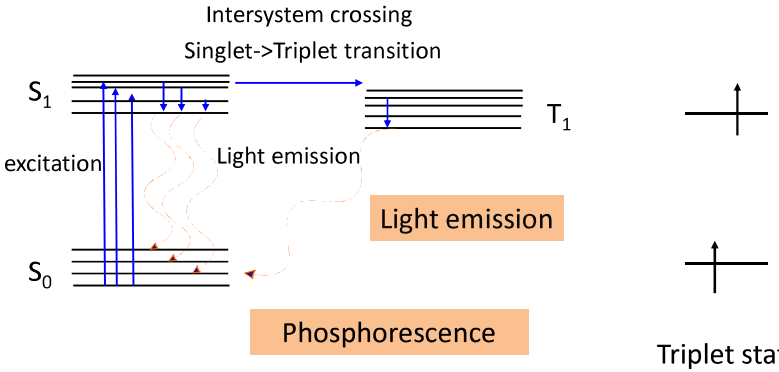
Kasha's rule:

fluorescence originates always from the vibrational state of lowest energy within the lowest electronic excited state.

$E_{excitation} \geq E_{fluorescence}$

$\lambda_{excitation} \leq \lambda_{fluorescence}$

Stokes-shift



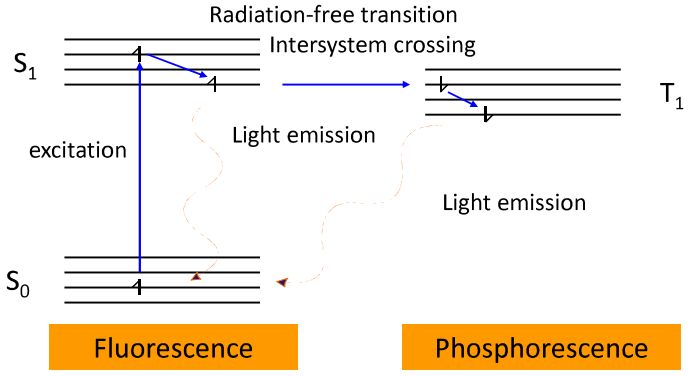
Triplet state

spontaneous light photon emission from metastable excited state

sum of all e^- spin quantum numbers=1

Metastable state

Emitted photon energies



Stokes-shift

$E_{excitation} \geq E_{fluorescence} > E_{phosphorescence}$

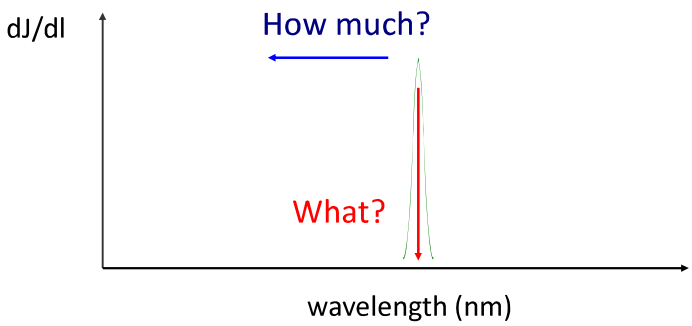
$\lambda_{excitation} \leq \lambda_{fluorescence} < \lambda_{phosphorescence}$

Characteristics of emitted light

Wavelength distribution of emitted light

Emission spectrum

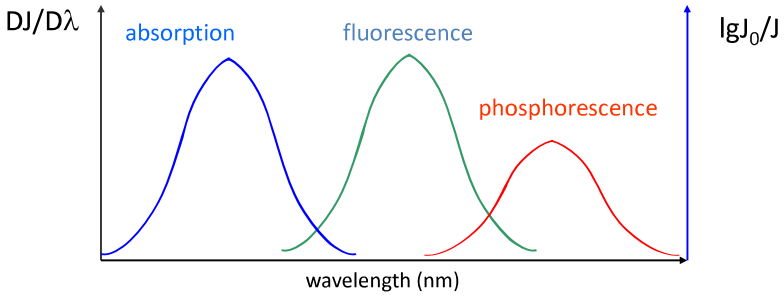
Atoms line spectrum



Wavelength distribution of emitted light

Emission spectrum

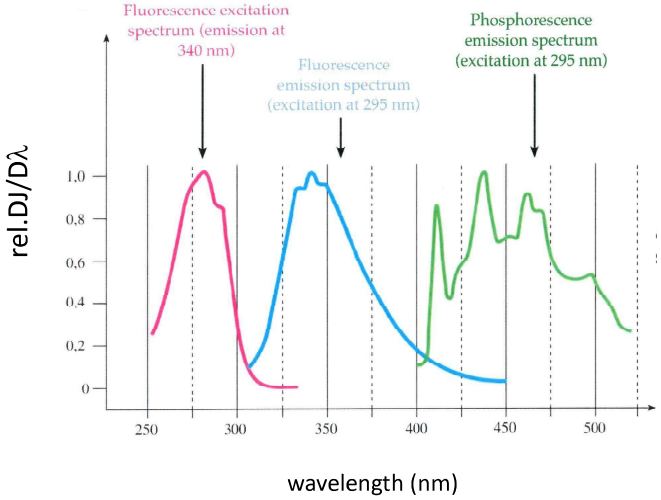
molecules: band spectrum



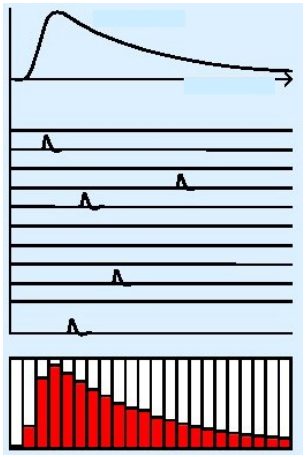
$\lambda_{excitation} \leq \lambda_{fluorescence} < \lambda_{phosphorescence}$

Stokes shift

E.g.: Corresponding spectra of triptophane



Excited-state lifetime

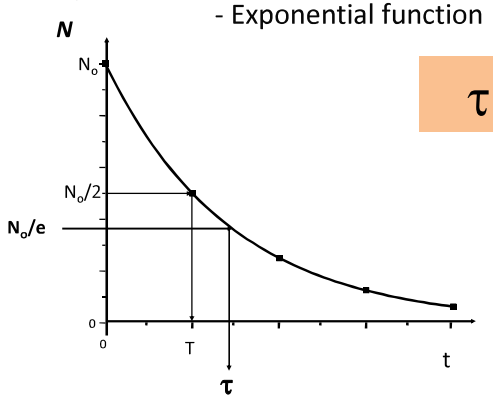


Single photon counting

Measuring of time gap between excitation and photon emission.

Statistical analysis of large number of measurements.

Number of excited electrons $\longrightarrow N = N_0 e^{-\frac{t}{\tau}}$ (time after excitation)



- Exponential function

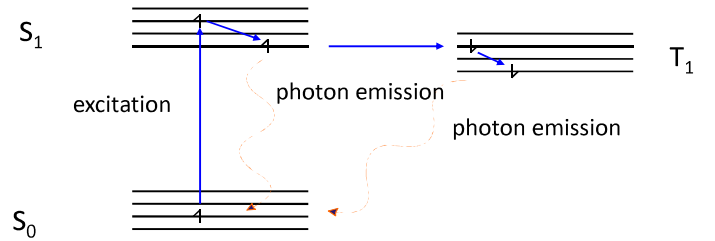
τ: life time

the time during which the number of excited electrons decreases to its e^{th}.

Typical excited-state lifetimes

Lifetime

the time during which the number of excited electrons decreases to its e^{th}.



Fluorescence	Phosphorescence
short	long
$10^{-9} – 10^{-7}$ s	$10^{-3} – 10^{2}$ s

Is excitation always followed by photon emission?

- Excited state decay can be caused by mechanisms other than photon emission and are therefore often called "non-radiative rates„.
- These can include: chemical reaction, dynamic collisional quenching, near-field dipole-dipole interaction, internal conversion and intersystem crossing.

Is excitation always followed by photon emission?

Quantum yield

$A \xrightarrow{h\nu} B$

Reciprocal of the number of absorbed photons for one photon emission

Fluorescence quantum yield (Q_F)

$$Q_F = \frac{\text{number.of.photons.emitted}}{\text{numbe.of.photons.absorbed}}$$

$$Q_F \leq 1$$

Types of luminescence:

- fluorescence
- phosphorescence

They can be characterized by:

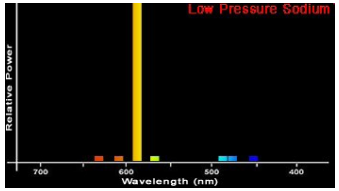
- emission spectrum:
  - Types
  - position of peaks
  - amplitude
- lifetime
- quantum yield

Application fields of luminescence

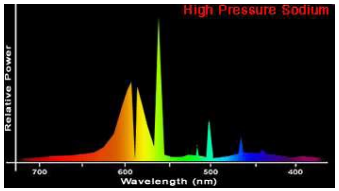
- Light sources (lightning, fertilization, sunbeds, photomedicine...)
- concentration determination (flame photometer)
- luminescence spectroscopy
- luminescence microscopy
- dosimetry (see later)
- archeology
- architecture
- safety control … many more

Luminescent light sources

Metal vapor lamps

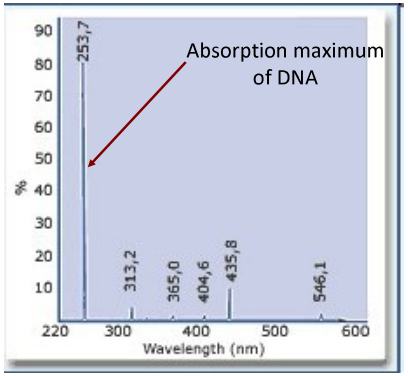


Low-pressure Na-vapor lamp



High-pressure Na-vapor lamp

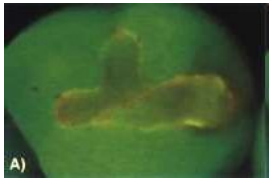
Low-pressure Hg-vapor lamp



Emission spectrum

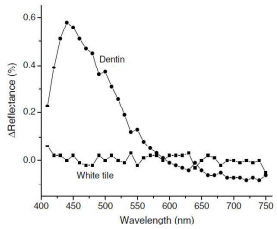
„germicid lamp"

Application in dental medicine

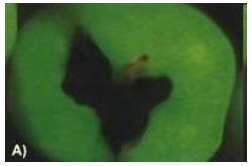


Red fluorescence indicates the activity of identifies cariogenic bacteria

Auto-fluorescence of teeth. When teeth are illuminated with high intensity blue light they will start to emit light in the green part of the spectrum.



Lee, Journal of Biomedical Optics 20(4), 040901 (April 2015)



amalgam restoration

0 – 14	No special measures.
15 – 20	Usual prophylactic measures.
21 – 30	More intensive prophylaxis or restoration: indication is dependent on: *Caries activity. *Caries risk. * Recall interval, etc.
from 30	Restoration and more intensive prophylaxis.

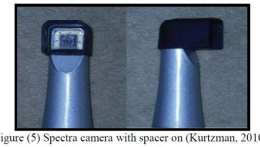
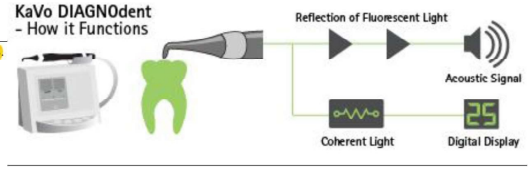
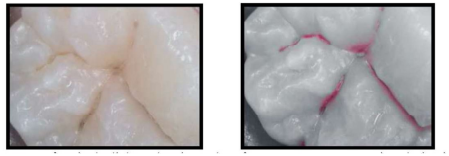


Figure (5) Spectra camera with spacer on (Kurtzman, 2010).

Table 2: Interpretation of Spectra data (Kurtzman, 2010).

Displayed Color	GREEN → BLUE → RED → ORANGE → YELLOW
Displayed Number	1 → 5
Depth of Involvement	Sound Enamel / Initial Enamel Caries / Deep Enamel Caries / Initial Dentin Caries / Deep Dentin Caries





SOPROCARE. (A) Carious lesion invisible in DAYLIGHT mode. (B) Carious lesion visible in CARIO mode

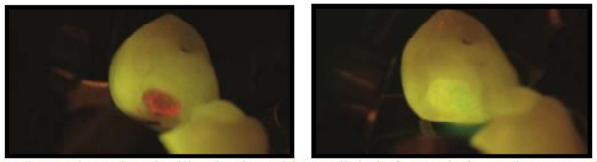
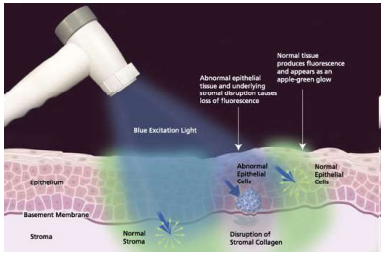


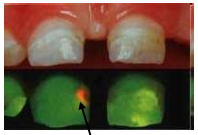
Figure (8) Photos showed cavity illumination with Facelight before and after caries excavation (21).

Mualla, Journal of Dental and Medical Sciences. 15:3, 2016, PP 65-75

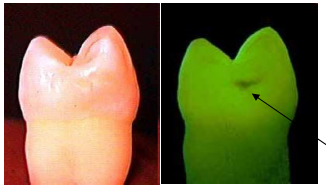


Healthy and malignant tissues different fluorescent properties

Teeth
native and fluroescent
images



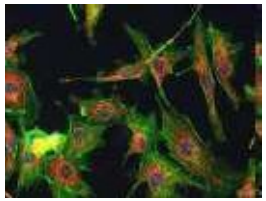
Active caries



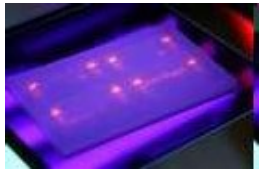
caries

Tooth
native and fluroescent
image

Luminescent microscopy



Laboratory application in many ways



And more...

Damjanovich, Fidy, Szöllősi: Medical Biophysics

II. 2.2
- 2.2.4
- 2.2.6

VI.3.3
- 3.3.1
- 3.3.2 pp. 411-413
- 3.3.3

Question of the week

How can you explain that a typical life time of fluorescence is shorter than that of the phosphorescence?