

Optical spectroscopy in materials science 12.

Ellipsometry Emission spectroscopy

Kamarás Katalin

MTA Wigner FK

kamaras.katalin@wigner.mta.hu

Emission part by Hajnalka Tóháti, Wigner RCP



Ellipsometry

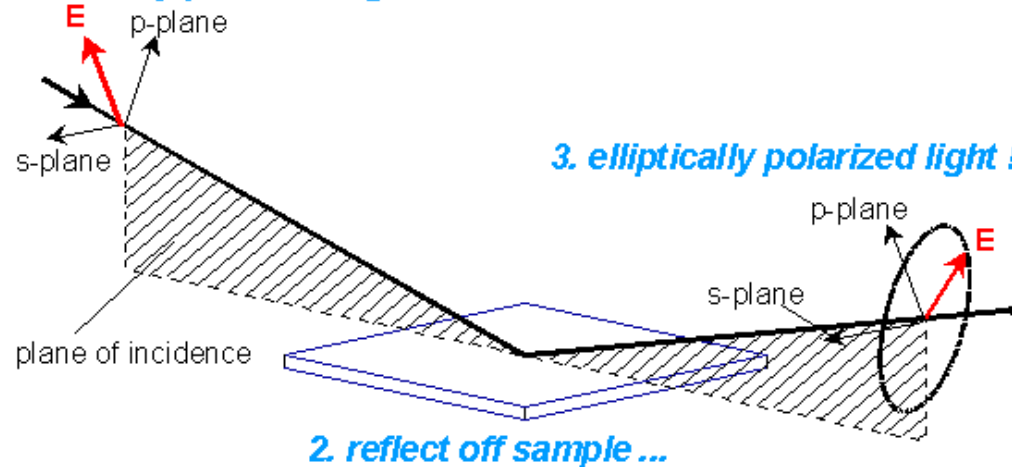
Fried Miklós
 Lohner Tivadar MTA EK MFA
 Petrik Péter
 SOPRA → Semilab

Snell's law:
 (Snellius – Descartes-törvény:
 Snellius - Gesetz:)

$$n_a \sin \varphi_a = n_b \sin \varphi_b$$

From the boundary conditions of Maxwell's equations:
 the tangential components of E and H are continuous at the interface

1. linearly polarized light ...



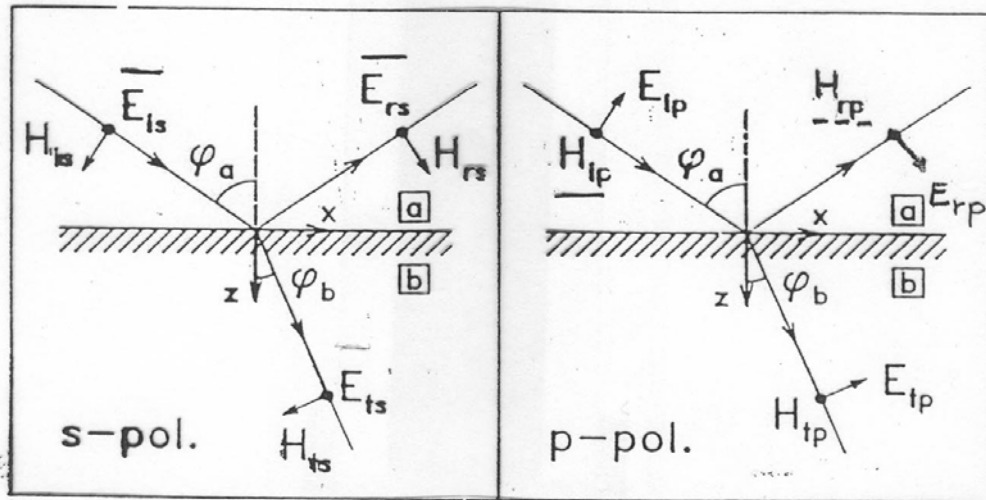
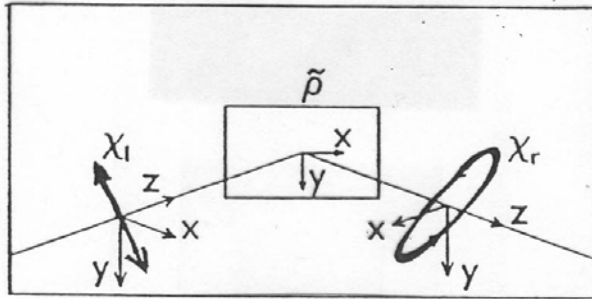
making use of

$$\mathbf{k} \times \mathbf{E} = \frac{\omega}{c} \mathbf{H}$$

$$|\mathbf{k}| |\mathbf{E}| = \frac{\omega}{c} |\mathbf{H}|$$

$$H = \frac{kc}{\omega} E = nE$$

Fresnel's equations



$$E_{rs} + E_{is} = E_{ts}$$

$$n_a \cos \varphi_a (E_{rs} - E_{is}) = -n_b \cos \varphi_b E_{ts}$$

$$n_a (E_{ip} - E_{rp}) = n_b E_{tp}$$

$$\cos \varphi_a (E_{rp} + E_{ip}) = \cos \varphi_b E_{tp}$$

Fresnel coefficients:

$$r_p = \frac{E_{rp}}{E_{ip}} \quad r_s = \frac{E_{rs}}{E_{is}} \quad t_p = \frac{E_{tp}}{E_{ip}} \quad t_s = \frac{E_{ts}}{E_{is}}$$

$$r_p = |r_p| e^{i\theta_{rp}} \quad r_s = |r_s| e^{i\theta_{rs}}$$

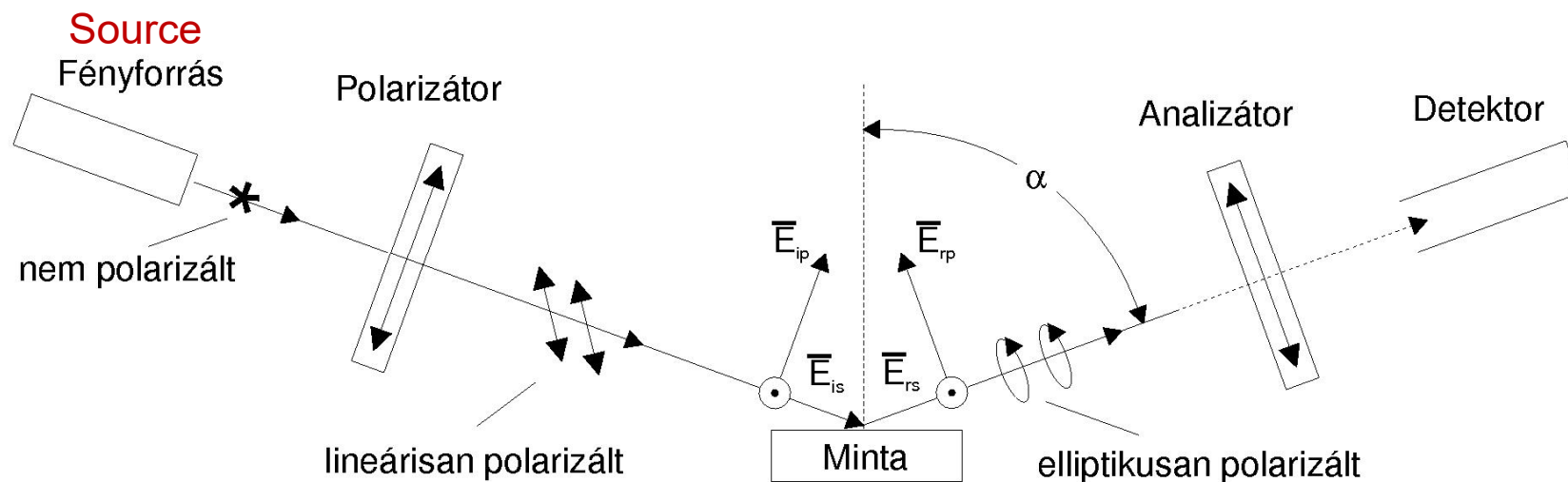
$$R = r^* r = |r|^2$$

$$r_p, r_s, t_p, t_s = f(n_a, n_b, \varphi_a, \varphi_b)$$

Measured quantities

Incident light: linear polarization

Reflected light: elliptical polarization



$$\rho = \frac{r_p}{r_s} = \tan \Psi e^{i\Delta}$$

Sample

Ellipsometric angles:

$$\tan \Psi = \left| \frac{r_p}{r_s} \right|$$

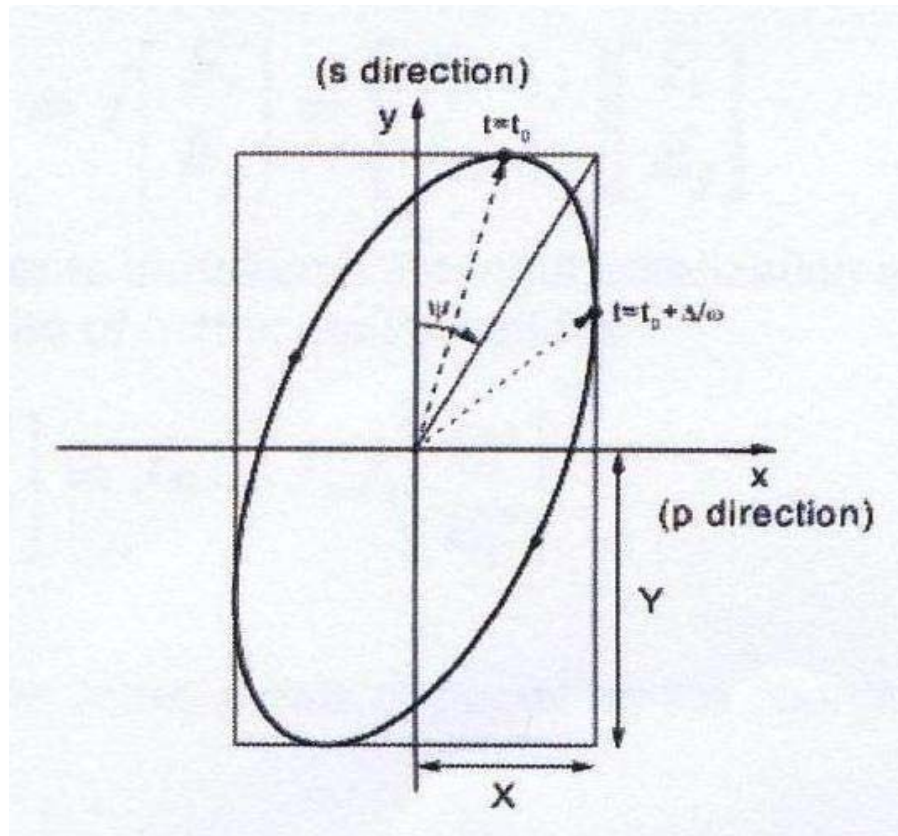
$$\Delta = \theta_{rp} - \theta_{rs}$$

Spectroscopic ellipsometry:

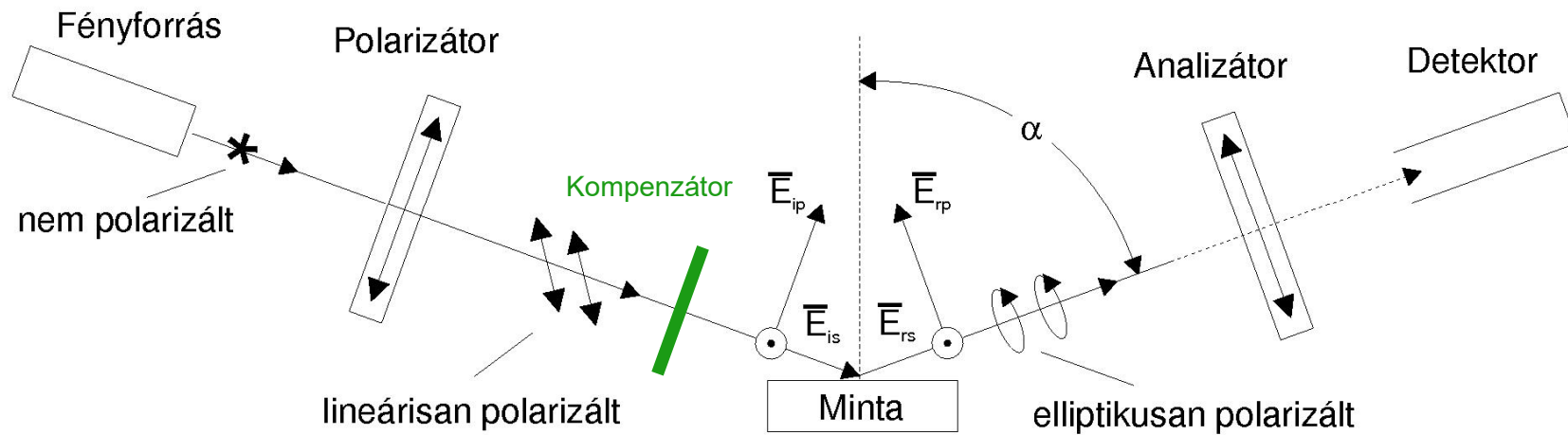
$$\Psi(\omega), \Delta(\omega)$$

Ellipsometric angles

Forrás: Tamáska István, 2009



Measurement: rotating analyzer setup



Polarizer: fixed angle (P)
Analyzer rotating: A(t)

$$I_{\text{det}}(A) = 1 + \frac{|\rho|^2 \cos^2 P - \sin^2 P}{|\rho|^2 \cos^2 P + \sin^2 P} \cos 2A + \frac{\text{Re}(\rho) \sin 2P}{|\rho|^2 \cos^2 P + \sin^2 P} \sin 2A$$

$$I_{\text{det}}(A_i) \sim 1 + \alpha(\rho, P) \cos 2A_i + \beta(\rho, P) \sin 2A_i$$

$$\tan \Psi = \sqrt{\frac{1 + \alpha}{1 - \alpha}} \tan P \quad \cos \Delta = \frac{\beta}{\sqrt{1 - \alpha^2}}$$

Evaluation: Isotropic, infinite two-phase model

$$n_b^2 = \varepsilon_r(\text{sample}) = n_a^2 \left(\sin^2 \varphi_a + \frac{(1-\rho)^2}{(1+\rho)^2} \sin^2 \varphi_a \tan^2 \varphi_a \right)$$

$$\rho = \rho' + i\rho''$$

If $n_a = 1$:

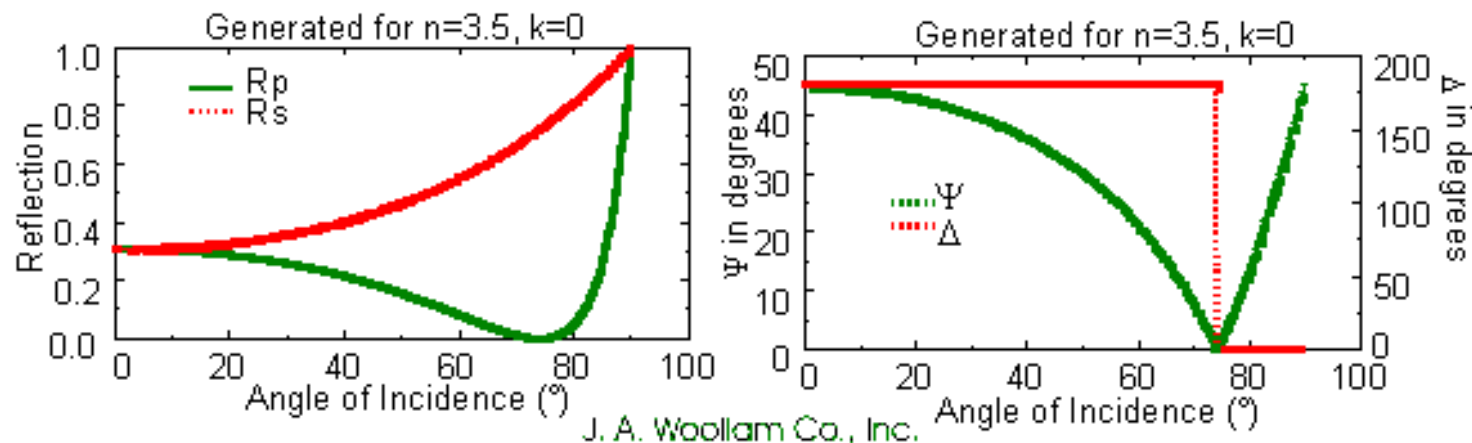
$$\varepsilon_r' = \sin^2 \varphi_a + \frac{(1-\rho^2)^2 - 4(\rho'')^2}{1+\rho^2 + 2(\rho')^2} \sin^2 \varphi_a \tan^2 \varphi_a$$

$$\varepsilon_r'' = -\frac{4\rho''(1-\rho^2)}{1+\rho^2 + 2(\rho')^2} \sin^2 \varphi_a \tan^2 \varphi_a$$

$$\varepsilon_r'' > 0 \Rightarrow \rho'' = \tan \Psi \sin \Delta < 0 \Rightarrow -\pi < \Delta < 0$$

convention! (measured quantity: $\cos \Delta$)

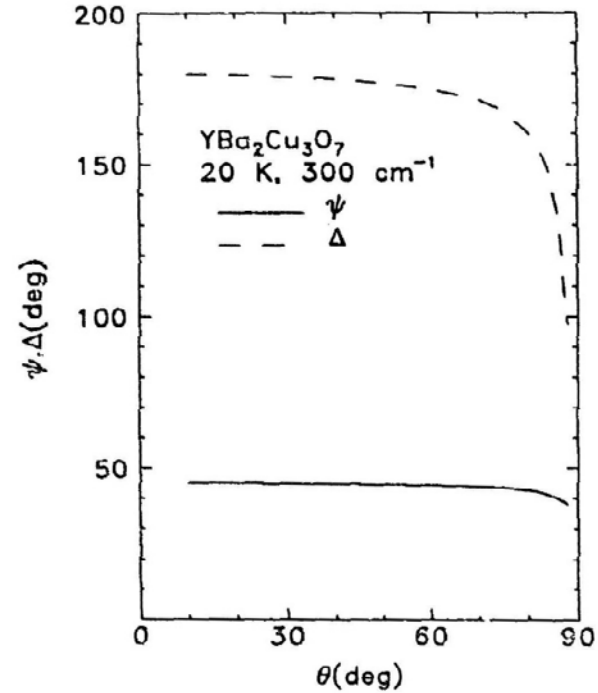
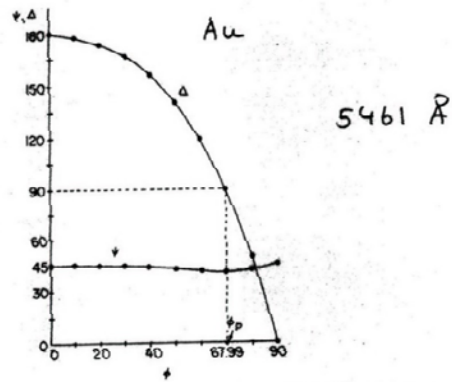
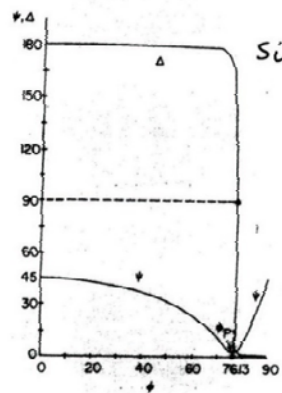
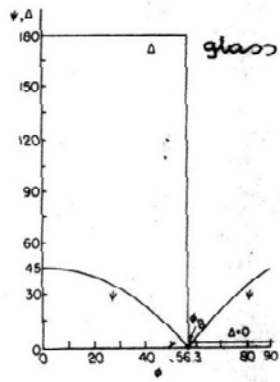
Accuracy of measurement



Brewster angle: $r_p=0$ $\Psi = 0, \Delta = \frac{\pi}{2}$
 $\tan \varphi_a(B) = n_b/n_a$

cos Δ is the measured quantity – optimal range is where it is the most sensitive to Δ
 however, sensitivity to angle of incidence is also large there
 sensitivity of Ψ to angle is small around minimum

Dependence on angle of incidence



R.M.A. Azzam, N.M. Bashara: Ellipsometry and Polarized Light.
North-Holland, Amsterdam, 1977

$\text{YBa}_2\text{Cu}_3\text{O}_7$ 20 K

K.Kamarás, D.van der Marel, C.C.Homes, T.Timusk:
Physica C **235**, 1085 (1994)



Advantages - disadvantages

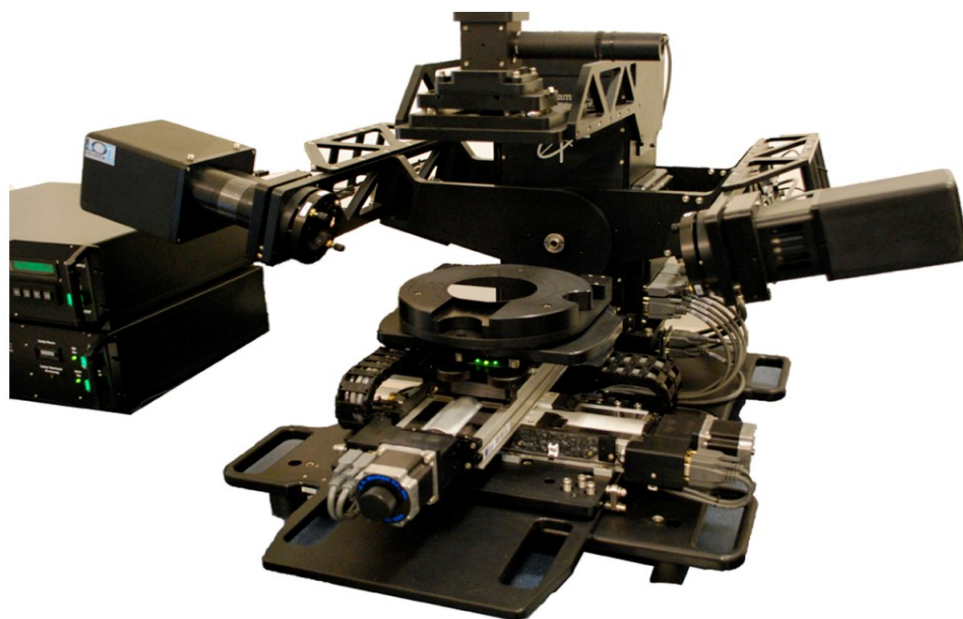
Advantages:

- direct determination of complex dielectric function (with appropriate model)
- no reference needed
- scattered light, small surface discontinuities cause small errors
- non-destructive
- remote sensing possible (visible range)

Disadvantages:

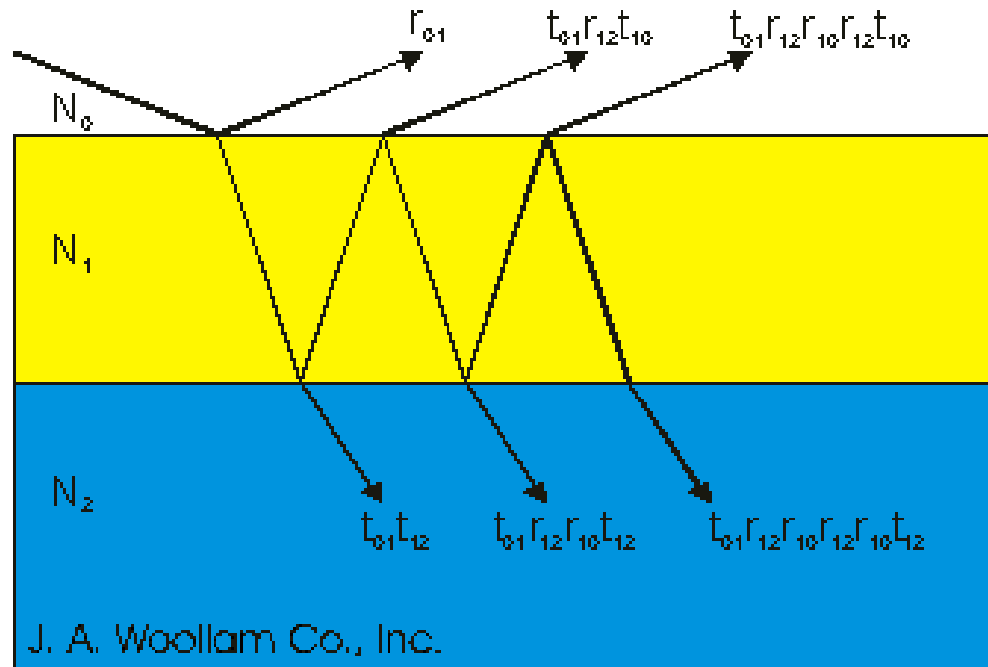
- large angle of incidence – large light spot, large sample area required
- evaluation complicated
- many parameters of the sample have to be known beforehand.

Modern ellipsometer



Woollam M2000DI – (MFA)
Rotating compensator
spectroscopic ellipsometer
range: **190-1700 nm**
minimum focus spot **0.15 mm**

Evaluation: multilayer systems

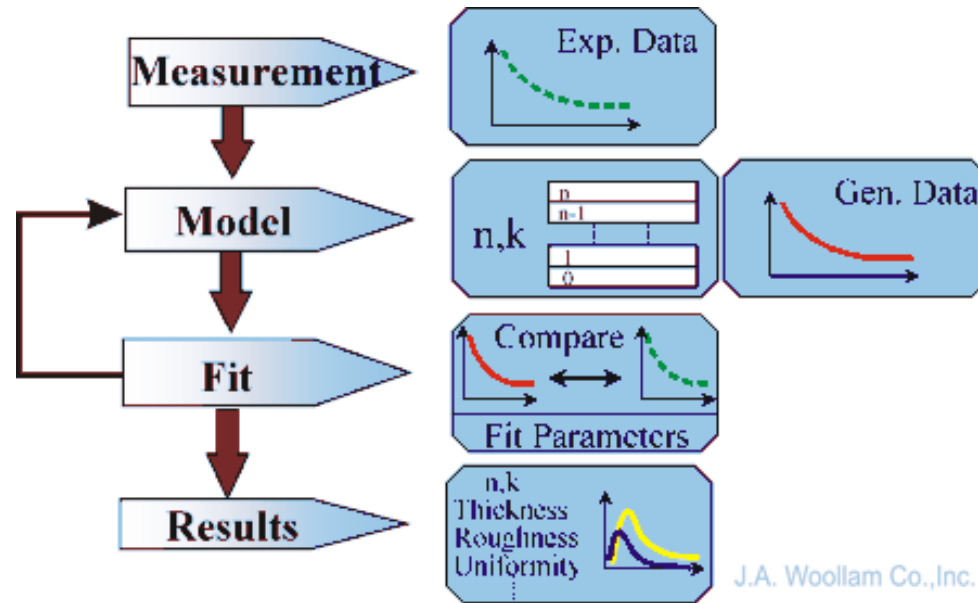


$$\rho(\varphi_a, \varepsilon_b, d_b, \varepsilon_c, d_c \dots)$$

knowing (n-2) parameters,
any two unknown quantities
can be determined
(e.g. thickness)

www.jawoollam.com

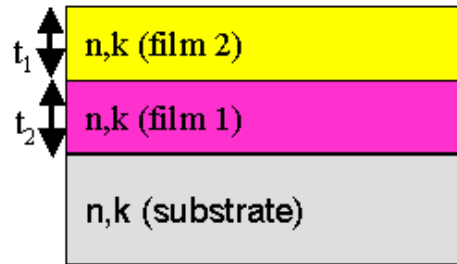
Fitting procedures



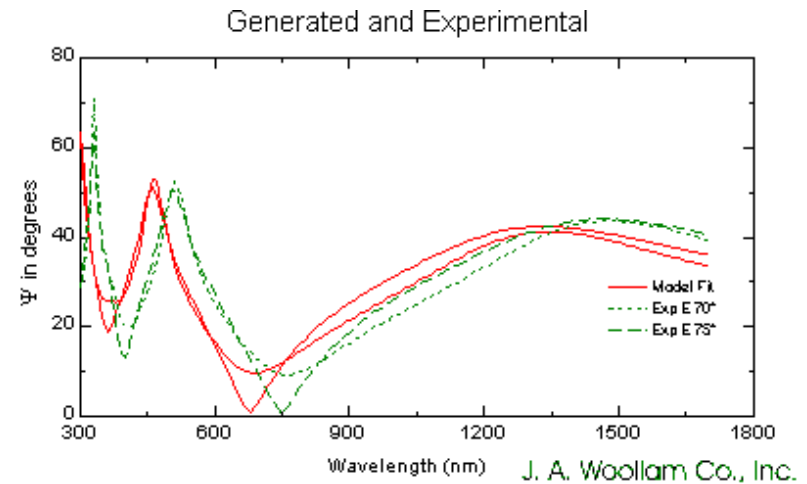
If $n(\omega)$ is known, more than two parameters can be fitted

more angles of incidence – more information

model



2	amorphous si	500 Å
1	si02	1000 Å
0	si	1 mm



Sensitivity of ellipsometry

$\Psi, \Delta = 0.01 - 0.02^\circ \rightarrow 0.01\text{nm}$ sensitivity on layer thickness

d (nm)	Δ	Ψ
0	179.257	10.448
0.1	178.957	10.448
0.2	178.657	10.449
0.3	178.356	10.450
0.4	178.056	10.451
0.5	177.756	10.453
1	176.257	10.462

Precise calibration (e.g. angle of incidence) is crucial!

Technology: process monitoring
process control

Source: Tamáska István, 2009

Application

- quick determination of dielectric function
- thickness measurement, technology control
- investigation of distribution in layered systems
(comparison with model calculations)
- ideal for semiconductors, multilayers
- small sensitivity in case of transparent and strongly absorbing samples

*“Kramers-Kronig transformation is arbitrary –
ellipsometry gives directly the dielectric function”*

What’s wrong with this sentence?



Combination of ellipsometry and Kramers-Kronig analysis

K. Kamarás, K.-L.Barth, F.Keilmann, R.Henn, M.Reedyk, C.Thomsen,
 M.Cardona, J.Kircher, P.L.Richards, J.-L.Stehlé:
 J. Appl. Phys. **78**, 1235 (1995)

SrTiO₃

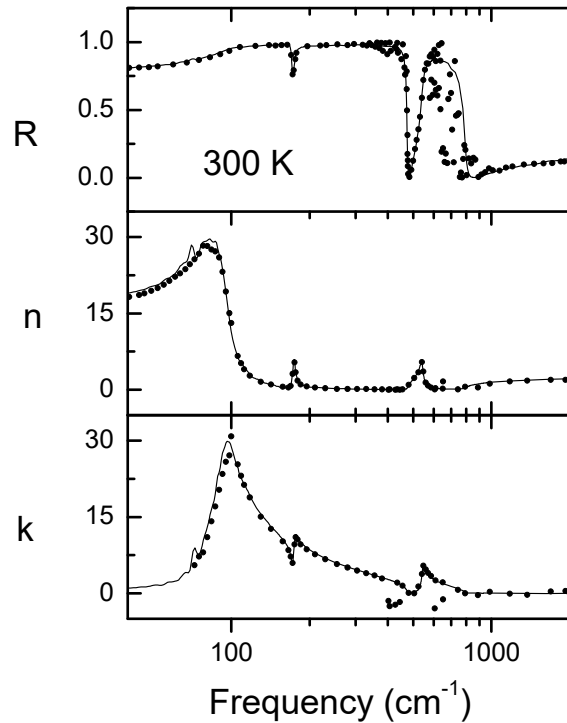


Fig. 1. Kamaras et al.

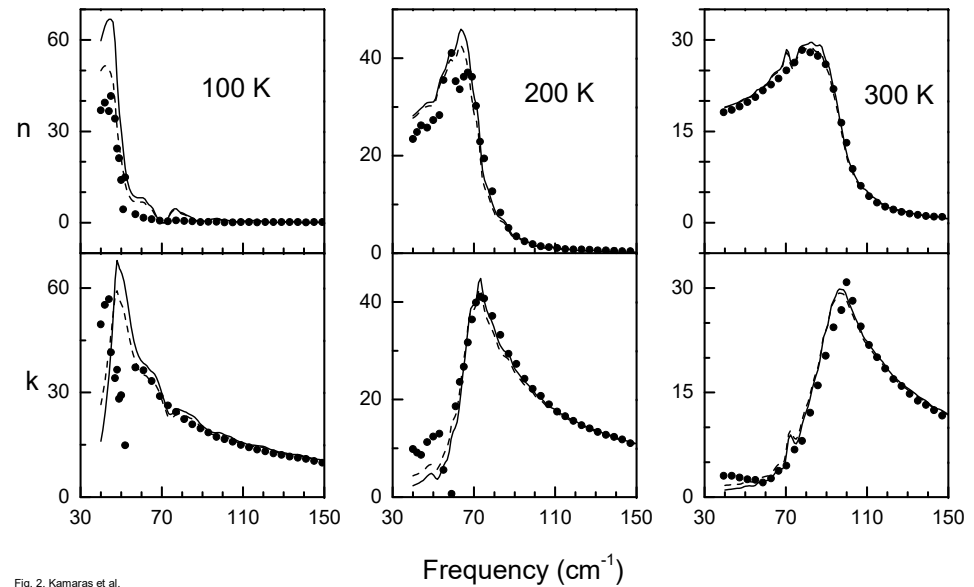


Fig. 2. Kamaras et al.

scaling of normal-incidence reflectance to ellipsometry
 low-frequency extrapolations depend on slope of curve



Take-home message

- Basics of ellipsometry: illumination of sample with linearly polarized light under finite angle; analyzing polarization state of reflected (elliptically polarized) light
- Measured quantity: ratio of Fresnel coefficients $\rho = \frac{r_p}{r_s} = \tan \Psi e^{i\Delta}$
- Ellipsometric angles ψ , Δ depend on sample dielectric function and angle of incidence
- Pseudodielectric function (isotropic, infinite, two-phase model)
- Multilayer systems: any 2 parameters can be determined when the others are known (mostly thickness of known materials)
- Modeling, process control, remote sensing

Összefoglalás

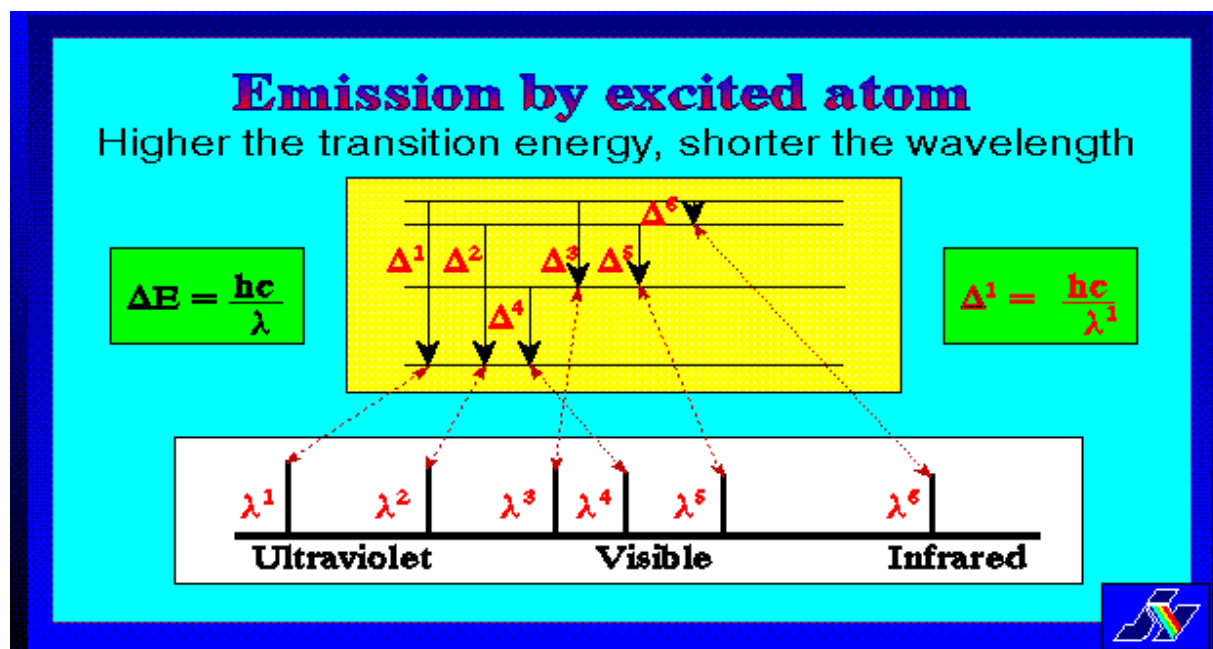
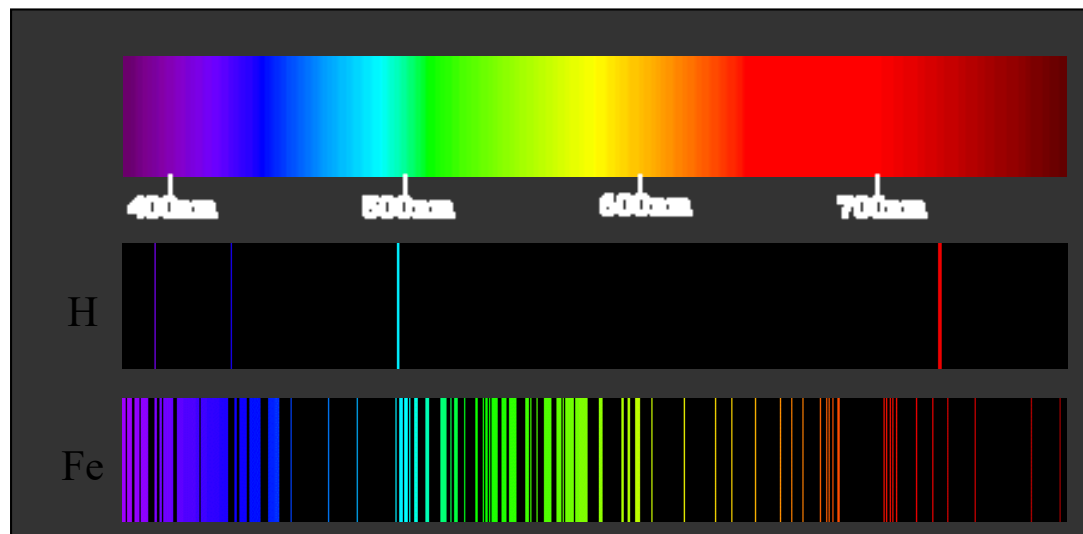
- Ellipszometria alapjai: minta megvilágítása lineárisan polarizált fénnel véges beesési szöggel ; a visszavert (elliptikusan polarizált) fény analizálása
- Mért mennyiség: Fresnel-együtthatók aránya $\rho = \frac{r_p}{r_s} = \tan \Psi e^{i\Delta}$
- A ψ , Δ ellipszometrikus szögek a minta dielektromos függvényétől és a beesési szögtől függenek
- Pszeudodielektromos függvény (izotrop, végtelen, kétfázisú modell)
- Többrétegű rendszerek: bármely 2 paraméter meghatározható, ha a többi ismert (legtöbbször ismert anyagokból álló rétegek vastagsága)
- Modellezés, folyamatirányítás, távoli érzékelés

Emission spectroscopy

- **Atomic emission spectroscopy (AES)**
 - Flame test
 - Flame emission photometry
 - Atomic absorption spectrophotometry
 - Inductively coupled plasma

- Molecular spectroscopy
 - IR emission spectroscopy
 - Luminescence

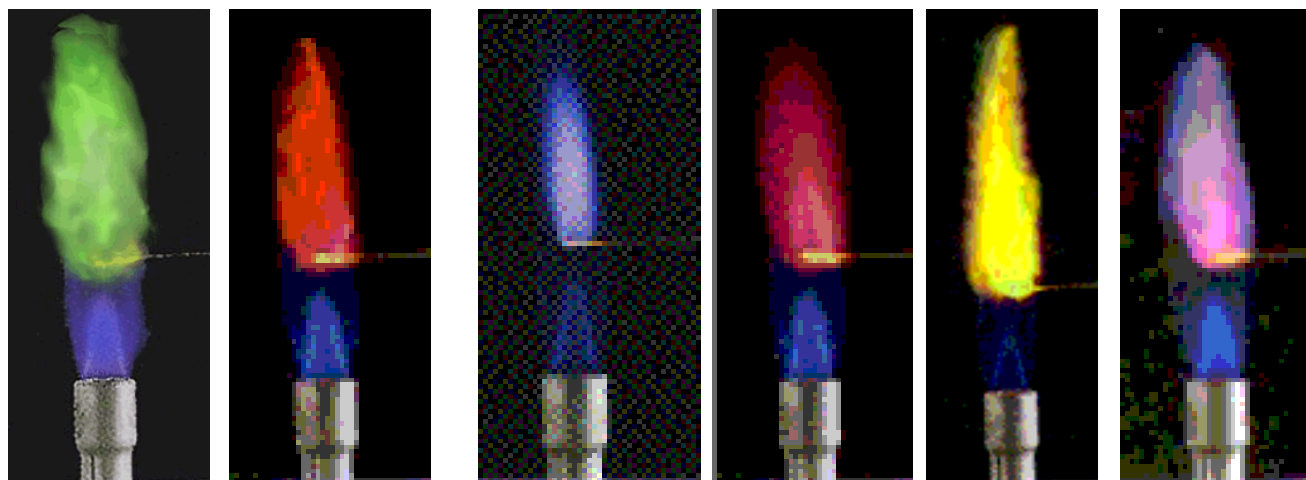
Emission spectroscopy



Atomic emission spectroscopy - Flame test

Walt Wolland, Bellevue Community College

<http://www.800mainstreet.com/s/s.html>



Ba

Ca

K

Li

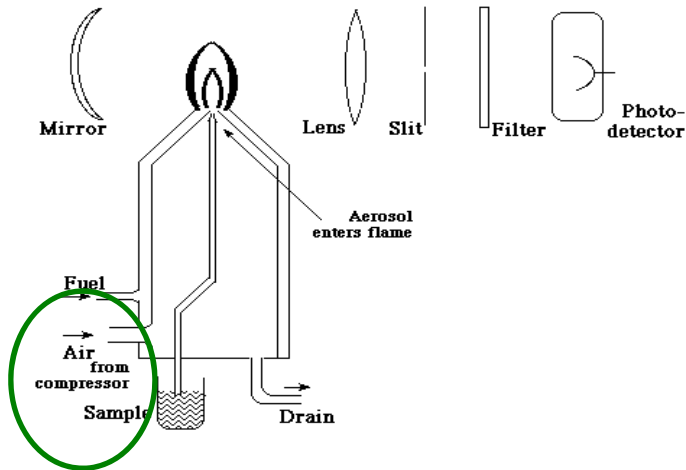
Na

Rb

Quantitative methods: flame photometry, atomic absorption spectroscopy (AAS)

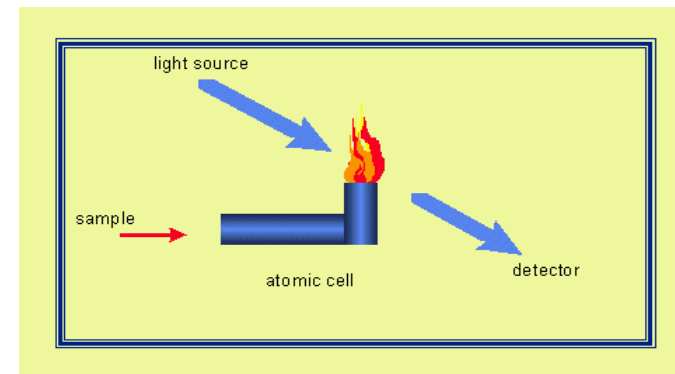
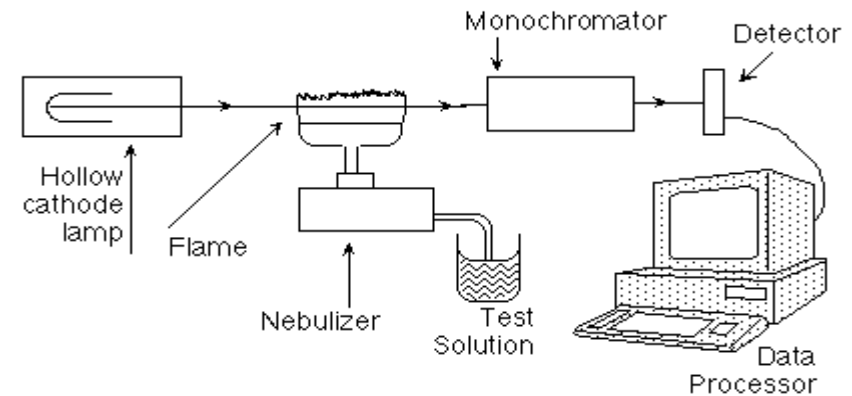
Atomic emission spectroscopy

Flame Emission Photometry (FEP)



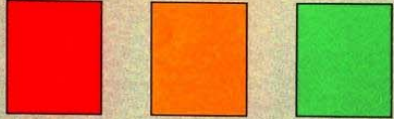
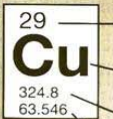
- **Excitation:** thermal energy of the flame
- **Flame:** - most frequent: acetylene and air
- vaporization → homogeneous atomic cloud
- excitation, but not ionization, of atoms
- **Vaporizer** → sample solution
- **Temperature:** 2000 – 3000 °C
- **Requirement:** constant composition, temperature and structure of the flame

Atomic Absorption Spectroscopy (AAS)

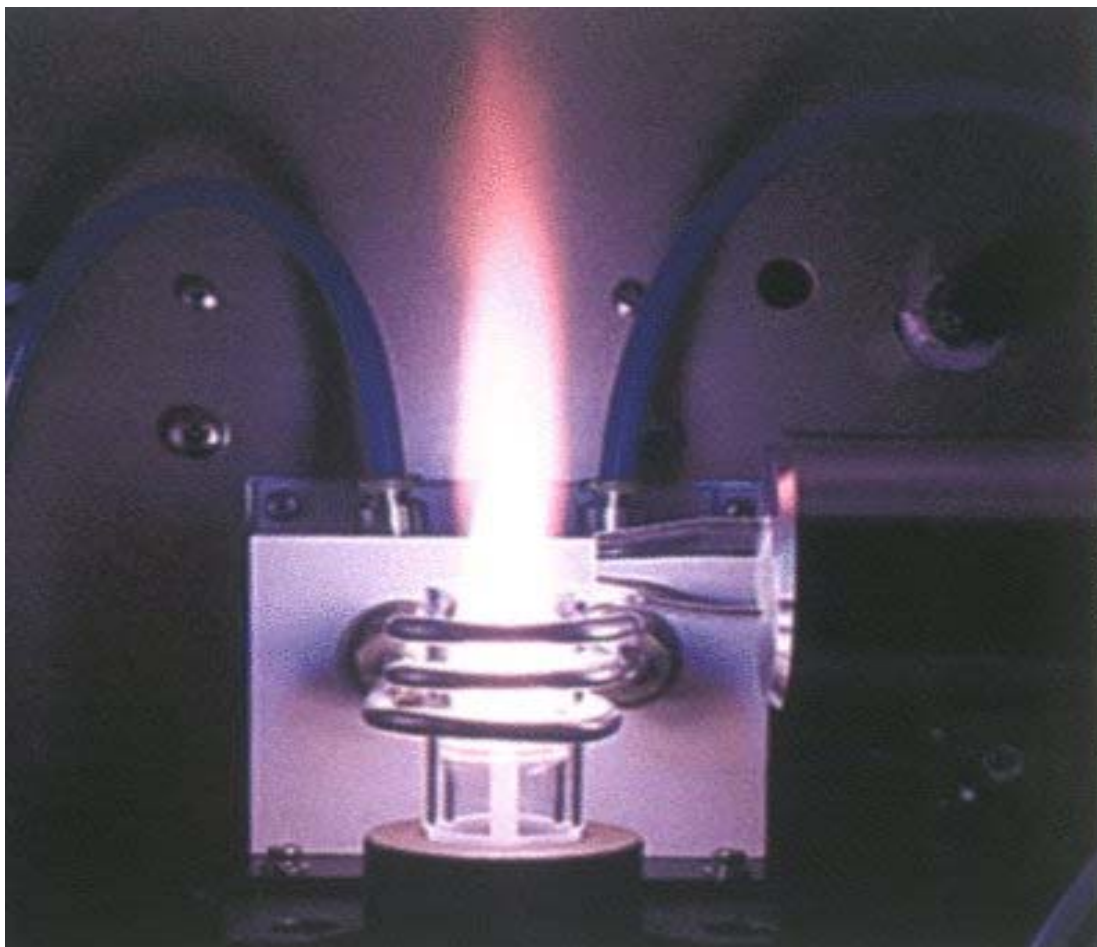


<http://www.cce.vt.edu/ewr/environmental/teach/smprimer/aa/aa.html>
<http://www.resonancepub.com/atomicsspec.htm>

Detectable elements

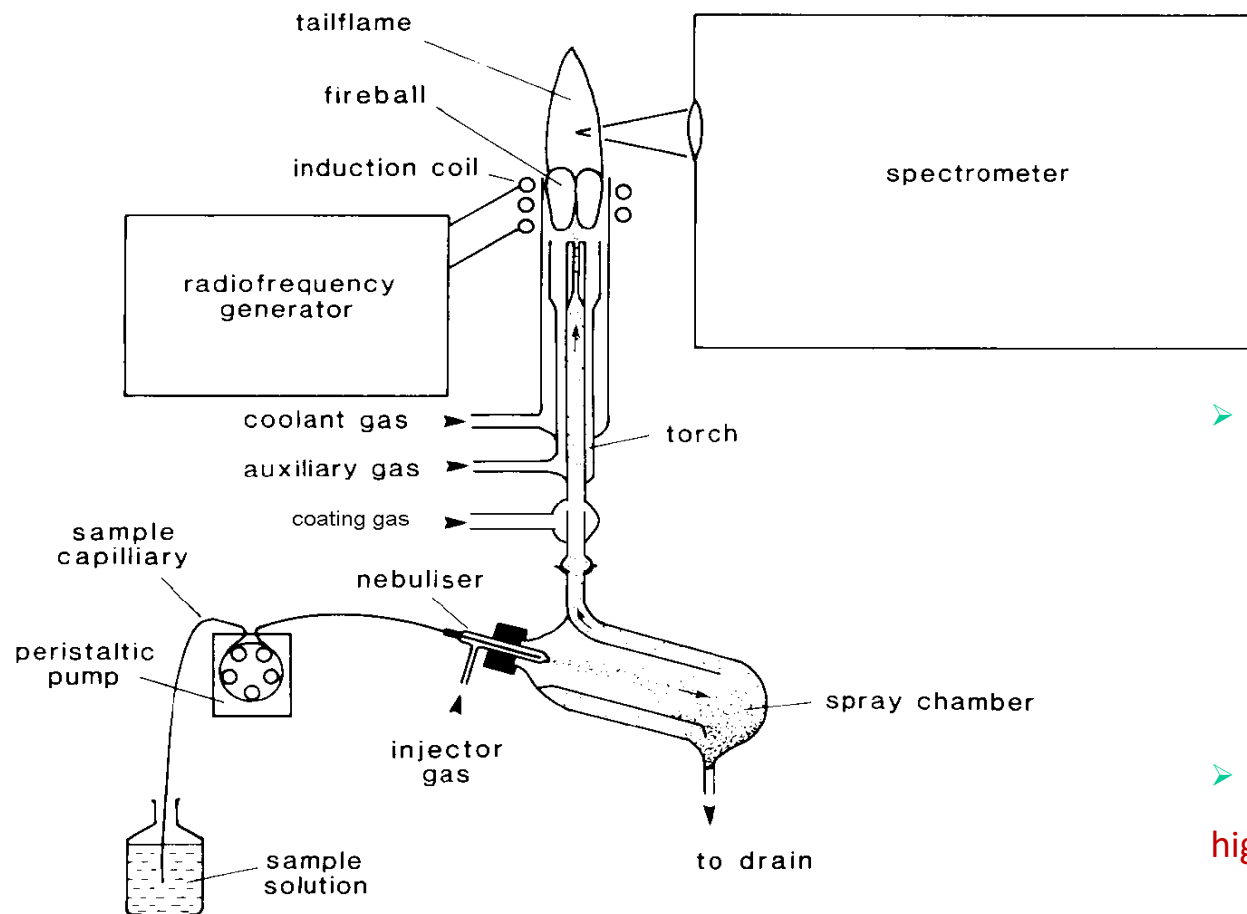
3 Li 670.8 6.941	4 Be 234.9 9.01218	 <p>Fuel-oxidant mixtures Two colours indicate use of either of two mixtures</p>										5 B 249.8 10.81	6 C 12.011	7 N 14.0067	8 O 15.999
11 Na 589.0 22.9898	12 Mg 285.2 24.305	 <p>ATOMIC NUMBER SYMBOL WAVELENGTH ATOMIC WEIGHT</p>										13 Al 309.3 26.9815	14 Si 251.6 28.086	15 P 213.6 30.9738	16 S 32.06
19 K 766.5 39.102	20 Ca 422.7 40.08	21 Sc 391.2 44.9559	22 Ti 368.4 47.88	23 V 318.5 50.9414	24 Cr 357.9 51.996	25 Mn 279.5 54.938	26 Fe 248.3 55.847	27 Co 240.7 58.9332	28 Ni 232.0 58.71	29 Cu 324.8 63.546	30 Zn 213.9 65.37	31 Ga 287.4 69.72	32 Ge 265.2 72.59	33 As 193.7 74.9216	34 Se 196.0 78.96
37 Rb 780.0 85.4678	38 Sr 460.7 87.62	39 Y 410.2 88.9058	40 Zr 360.1 91.22	41 Nb 334.4 92.9064	42 Mo 313.3 95.94	43 Tc 98.9062	44 Ru 349.9 101.07	45 Rh 343.5 102.9055	46 Pd 247.6 106.4	47 Ag 328.1 107.868	48 Cd 228.8 112.40	49 In 303.9 114.82	50 Sn 224.6 118.69	51 Sb 206.8 121.75	52 Te 214.3 127.60
55 Cs 852.1 132.9055	56 Ba 553.6 137.34	57 La 550.1 138.9055	72 Hf 307.3 178.49	73 Ta 271.5 180.9479	74 W 265.1 183.85	75 Re 346.1 186.2	76 Os 280.9 190.2	77 Ir 208.9 192.22	78 Pt 265.9 195.09	79 Au 242.8 196.9665	80 Hg 253.7 200.59	81 Tl 276.8 204.37	82 Pb 217.0/283.3 207.2	83 Bi 223.1 208.9806	84 Po 209
87 Fr 223	88 Ra 226.0254	89 Ac 227													
Lanthanides		* 58 Ce 140.12	59 Pr 495.1 140.9077	60 Nd 492.5 144.24	61 Pm 145	62 Sm 429.7 150.4	63 Eu 459.4 151.96	64 Gd 407.9 157.25	65 Tb 432.7 158.9254	66 Dy 421.2 162.50	67 Ho 410.4 164.9303	68 Er 400.8 167.26	69 Tm 371.8 168.9342	70 Yb 398.8 173.04	71 Lu 336.0 174.967
Actinides		** 90 Th 232.0381	91 Pa 231.0359	92 U 358.5 238.029	93 Np 237.0482	94 Pu 244	95 Am 243	96 Cm 247	97 Bk 247	98 Cf 251	99 Es 254	100 Fm 257	101 Md 258	102 No 259	103 Lr 260

Inductively coupled plasma (ICP)



 JOBIN YVON - SPEX
Groupe HORIBA

ICP Kitchen area

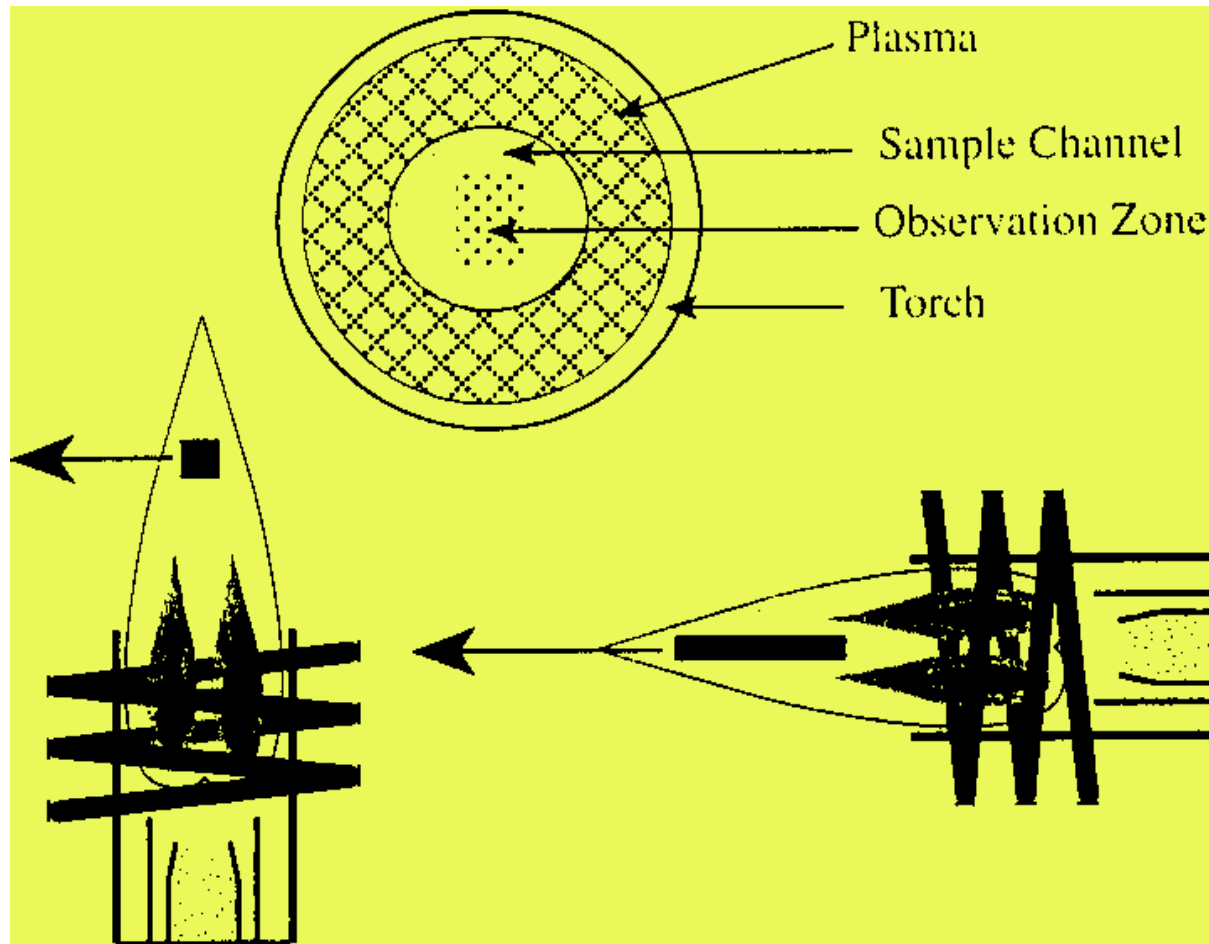


- Ar gas excited at radio frequency
- Temperature up to 8000° C
- Low concentration

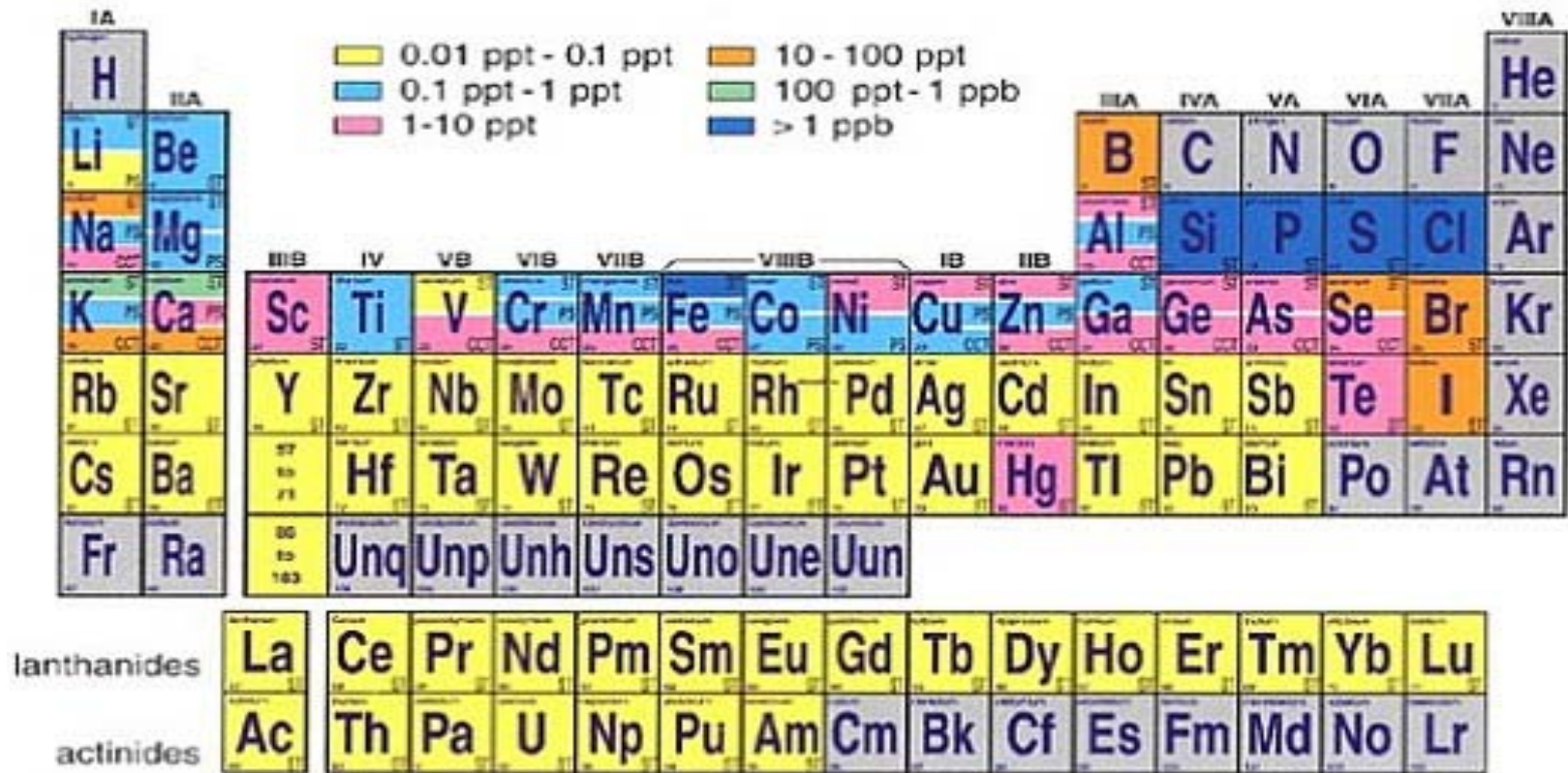
- Plasma torch: 3 concentric quartz tubes, streaming Ar
 - ❖ Outside → cooling
 - ❖ Central → plasma
 - ❖ Inside → carrier gas for sample
- Plasma state: electric excitation → high temperature fireball

Observation in the upper part of the fireball: low concentration, more lines

ICP plasma



ICP-AES – Detection limits

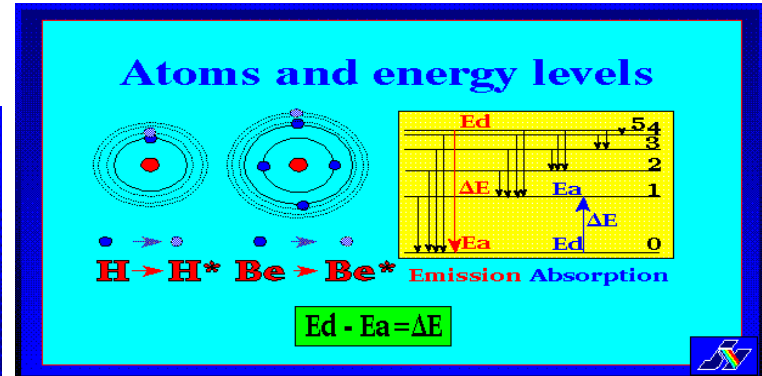
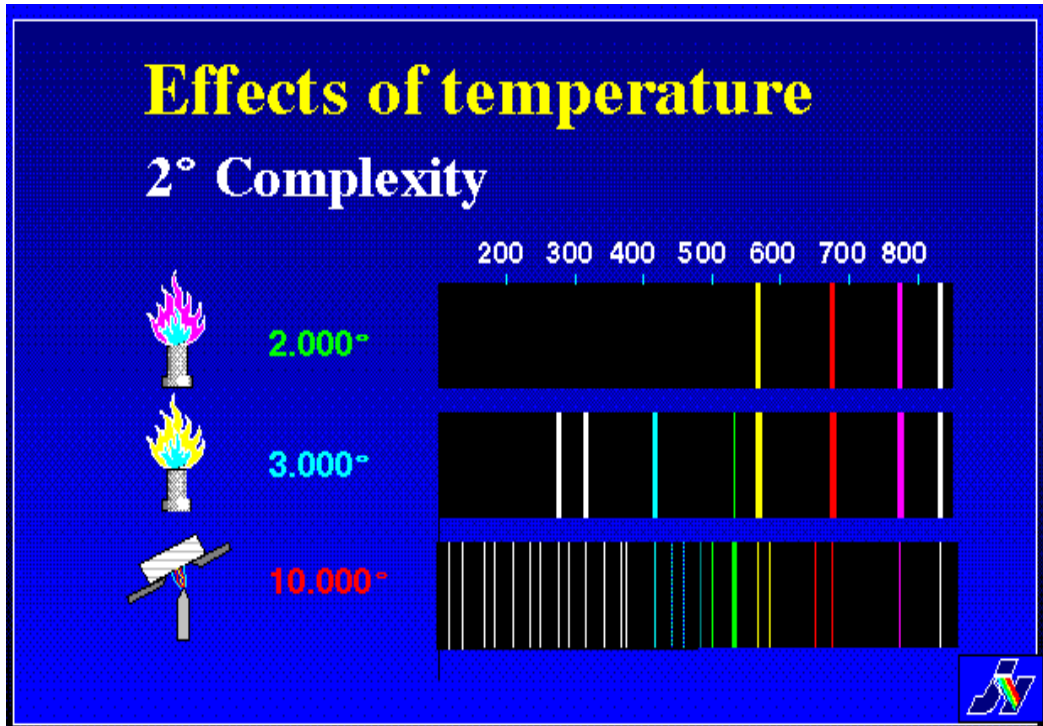


3 sigma detection limits in de-ionized water

ST = Standard PS = PlasmaScreen CCT = Collision Cell



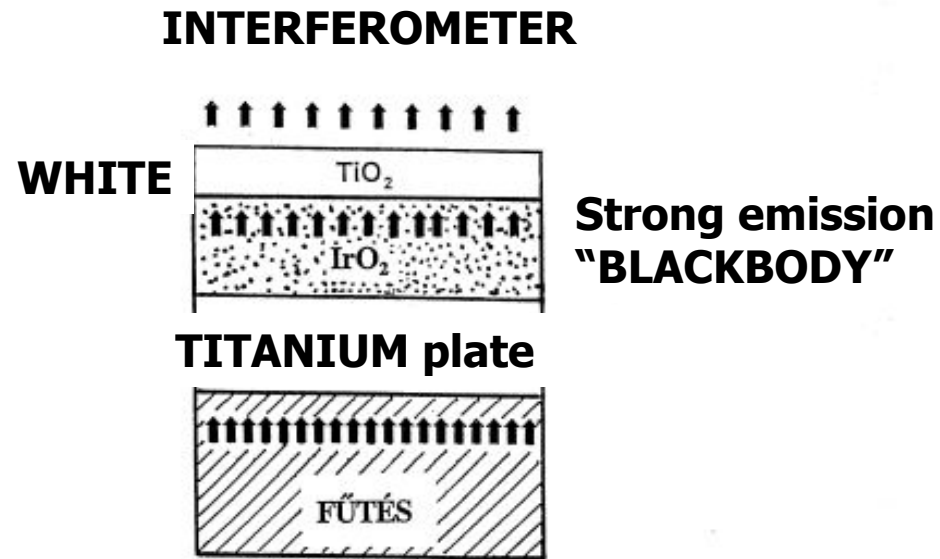
Emission spectra: effect of temperature



Infrared emission spectroscopy

Temperature: 100 – 200 °C

Keresztury Gábor, Mink János,
Kristóf János
MTA Kémiai Kutatóközpont,
Veszprémi Egyetem

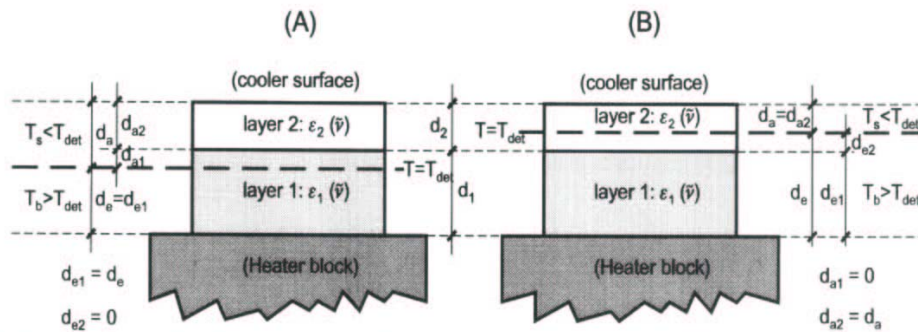


The model of layer structure

Self-absorption

The spectral shape depends on:

- effective thickness of the layers (d_1, d_2)
- thickness of emitting, absorbing layers (d_e, d_a)



G. Keresztury, J. Mink, J. Kristóf:
Anal. Chem. **67**, 3782 (1995)

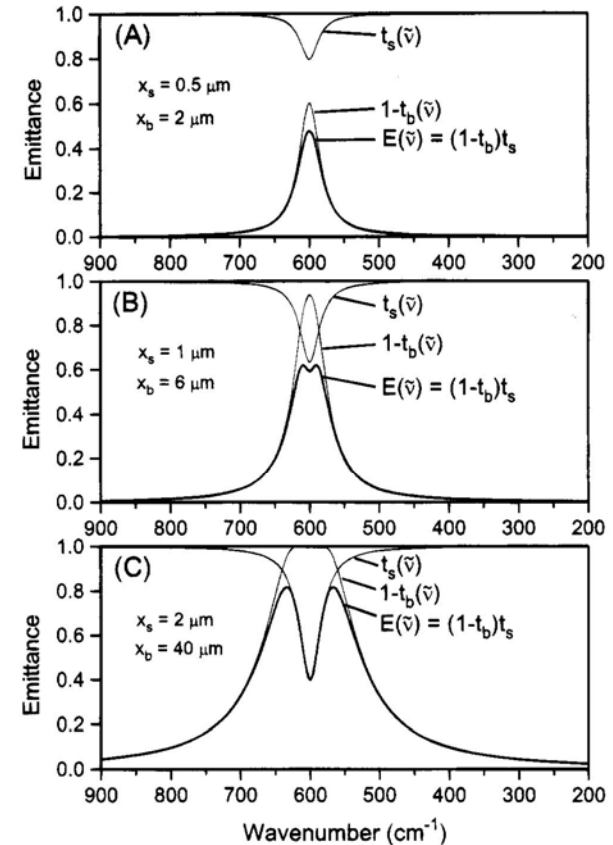


Figure 2. Simulation of the effect of self-absorption using eq 12 for different effective thicknesses of emitting bulk (x_s) and absorbing surface layer (x_b). (Lorentzian band shapes with $\tilde{\nu}_0 = 600$, $l_0 = 2000$, and $w = 30 \text{ cm}^{-1}$.)

t_b – transmittance of the bulk

t_s – transmittance of the surface

E - emittance

Molecular spectroscopy – Luminescence

Light emission by excited molecules



Fluorescence of different sized CdSe quantum dots

Joseph R. Lakowicz – Principles of fluorescence spectroscopy, 3rd edition

Types of luminescence

(a) Excitation Mode	Luminescence Type
absorption of radiation (UV/VIS)	photoluminescence
chemical reaction	chemiluminescence, bioluminescence
thermally activated ion recombination	thermoluminescence
injection of charge	electroluminescence
high energy particles or radiation	radioluminescence
friction	triboluminescence
sound waves	sonoluminescence

(b) Excited State (Assuming Singlet State)	Luminescence Type
first excited singlet state	fluorescence, delayed fluorescence
lowest triplet state	phosphorescence

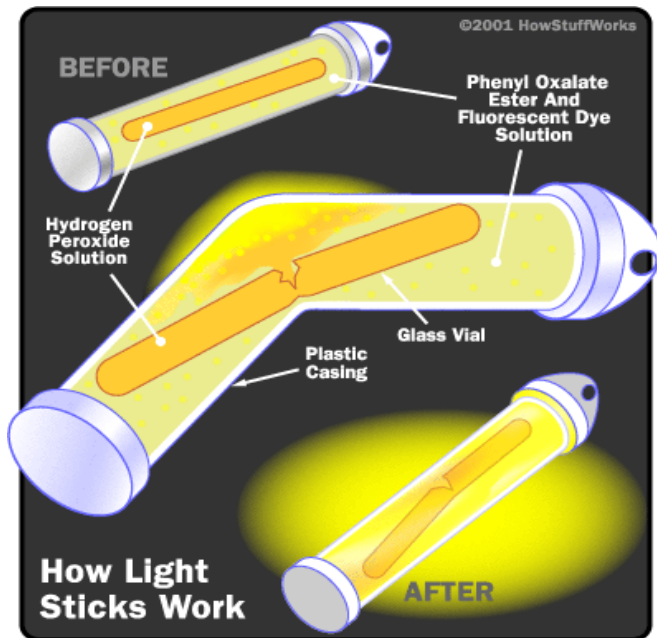
Chemiluminescence/bioluminescence

Chemiluminescence – is the emission of light as the result of a chemical reaction



Bioluminescence – one type of chemiluminescence; the light is produced and emitted by a living organism

e.g. firefly, deep-sea fish, jellyfish, squids, bacteria, planktons, mushrooms



Luminescence

Fluorescence

- Emission: From excited singlet state
- Transition: Allowed
- Emission rate: Fast: 10^8 s^{-1}
- Average lifetime: 1 – 10 ns

➤ Example:



Phosphorescence

- From excited triplet state
- „Forbidden”
- Slow: $10^3 - 10^0 \text{ s}^{-1}$
- ms – s



0 sec

1 sec

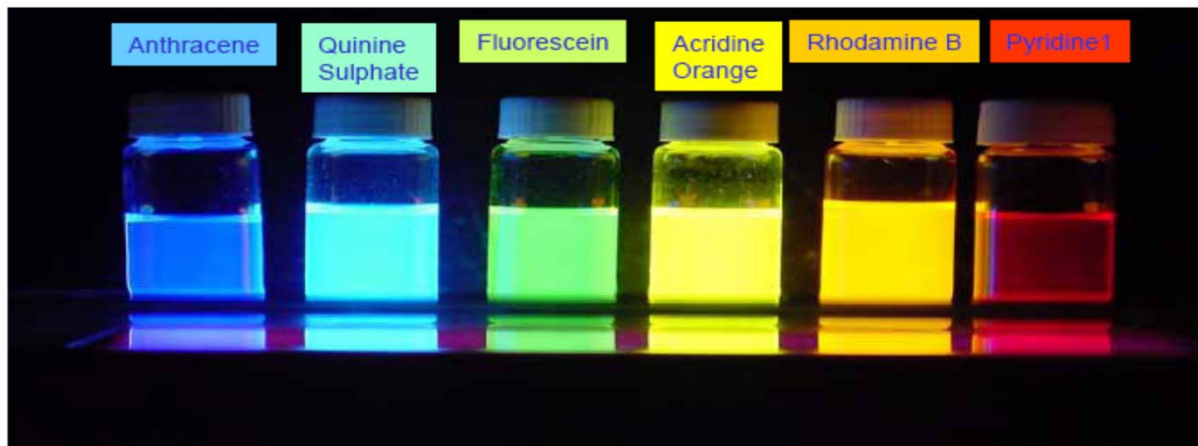
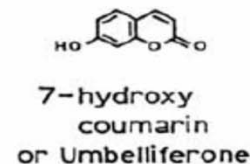
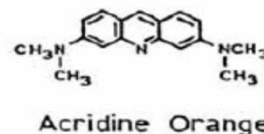
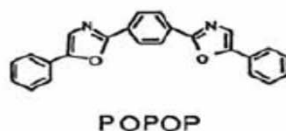
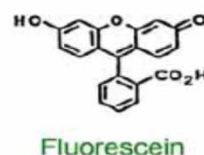
640 sec



Molecular spectroscopy

Fluorescence – Typical fluorophores

- Typically aromatic molecules
- Usually no fluorescence in condensed state



Molecular spectroscopy

Fluorescence – the beginning



Sir John Fredrich William Herschel

1792 – 1871

On a case of superficial colour presented by a homogeneous liquid internally colourless. By Sir John Frederick William Herschel, Philosophical Translation of the Royal Society of London (1845) 135:143–145. Received January 28, 1845 — Read February 13, 1845.

"The sulphate of quinine is well known to be of extremely sparing solubility in water. It is however easily and copiously soluble in tartaric acid. Equal weights of the sulphate and of crystallised tartaric acid, rubbed up together with addition of a very little water, dissolve entirely and immediately. It is this solution, largely diluted, which exhibits the optical phenomenon in question. Though perfectly transparent and colourless when held between the eye and the light, or a white object, it yet exhibits in certain aspects, and under certain incidences of the light, an extremely vivid and beautiful celestial blue colour, which, from the circumstances of its occurrence, would seem to originate in those strata which the light first penetrates in entering the liquid, and which, if not strictly superficial, at least exert their peculiar power of analysing the incident rays and dispersing those which compose the tint in question, only through a very small depth within the medium.

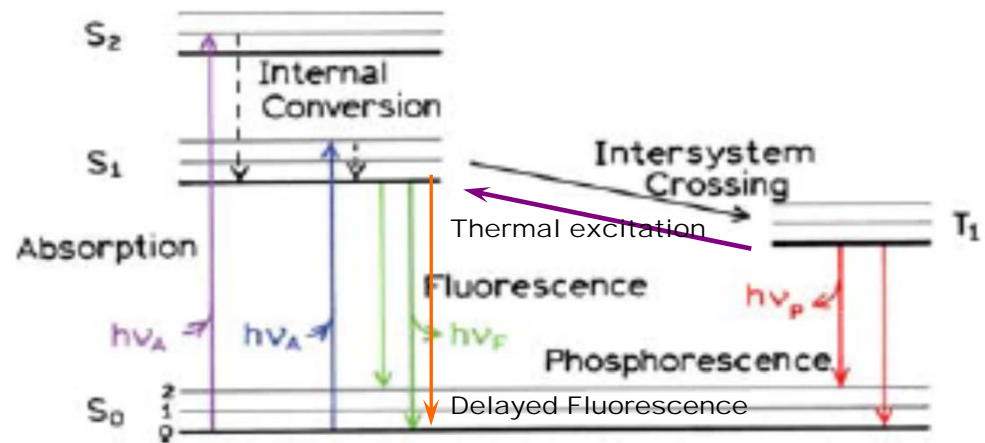
Fluorescence of quinine is the most widely used example up to now

Luminescence – Jablonski diagram



Professor Alexander Jablonski

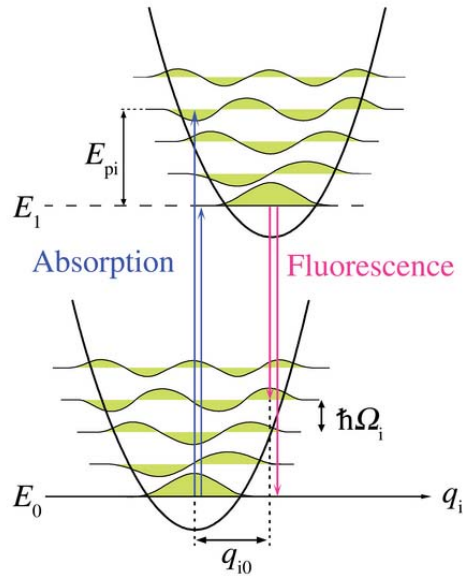
1898 – 1980



One form of a Jablonski diagram.

Luminescence in molecules

<http://www.shsu.edu/~chemistry/chemiluminescence/JABLONSKI.html>



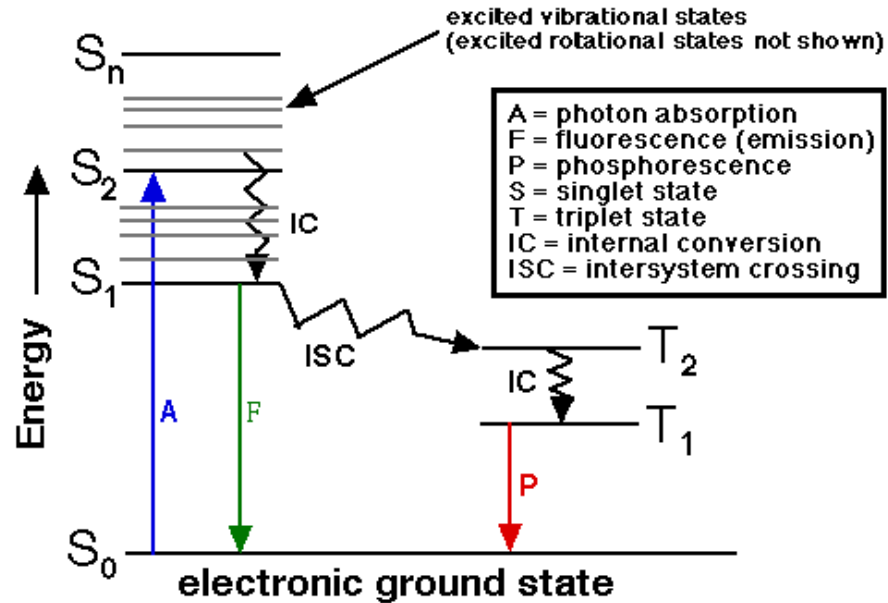
Franck-Condon principle
 R: configuration coordinate
 absorption (vertical)

↓
 relaxation

↓
 emission (vertical)

↓
 relaxation

http://en.wikipedia.org/wiki/Franck-Condon_principle



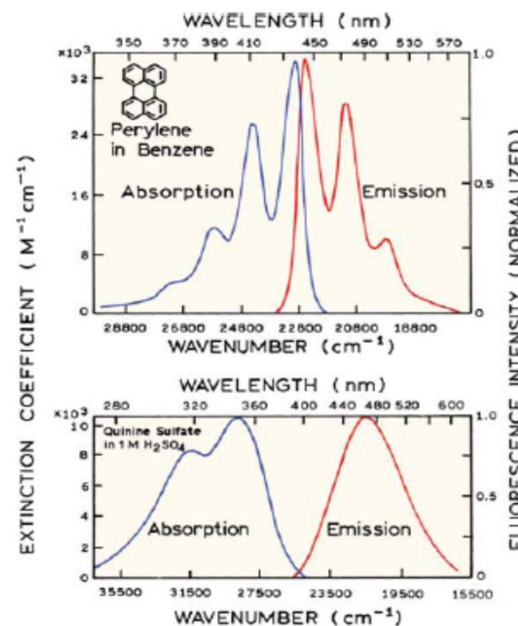
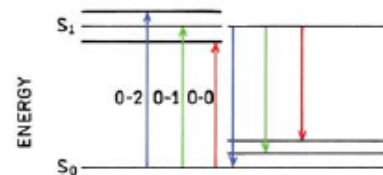
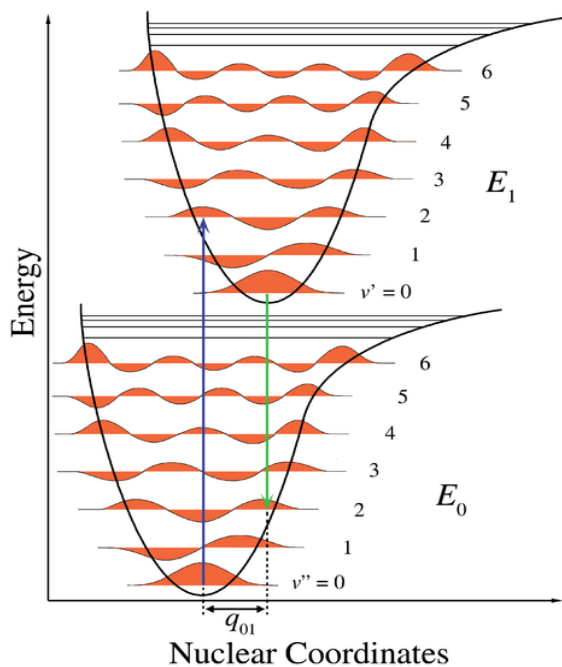
Jablonski diagram

- Intersystem crossing:** singlet– triplet
- Internal conversion:** into vibrationally excited state of higher singlet
- Fluorescence:** singlet - singlet
- Phosphorescence:** singlet – triplet (delayed)

Fluorescence basics

Mirror image

Franck-Condon principle



During an electronic transition a change from one vibrational energy level to another will be more likely to happen if the two vibrational wavefunctions overlap more significantly

Electronic excitation does not greatly alter the nuclear geometry

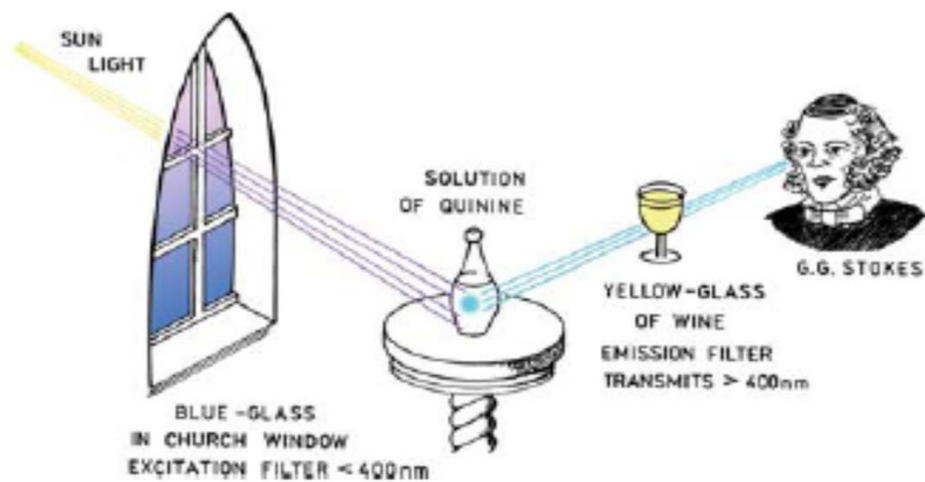
Molecular spectroscopy

Fluorescence – Stokes shift



Sir George Gabriel Stokes

1819 – 1903



Experimental schematic for detection of the Stokes shift.

Visual observation of Stokes shift

Molecular spectroscopy

Fluorescence – Lifetime (τ) and quantum yield (Q)

- These are the two most important characteristics

$$\text{quantum yield} = Q = \frac{\text{number of emitted photons}}{\text{number of absorbed photons}} = \frac{\Gamma}{\Gamma + k_{nr}}$$

where Γ – emission rate of fluorophores

k_{nr} – number of non-radiative transitions to ground state S_0

if there is Stokes shift, $Q < 1$

Lifetime: average time between excitation and emission

$$\tau = \frac{1}{\Gamma + k_{nr}}$$

if $k_{nr} = 0$, *intrinsic* lifetime

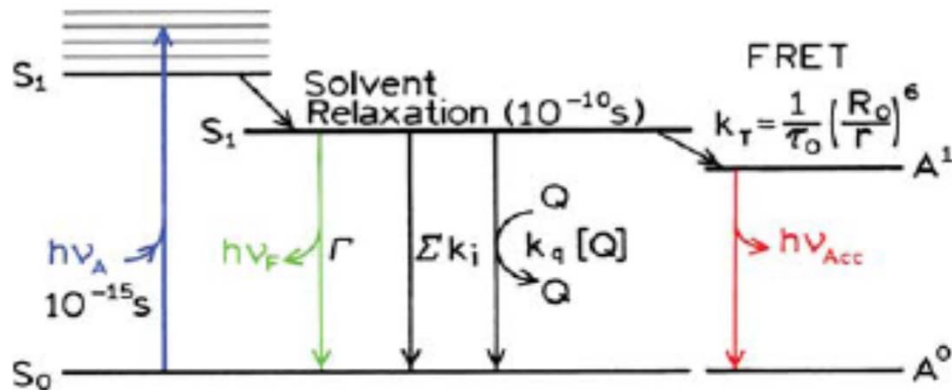
Molecular spectroscopy

Fluorescence – Quenching

Quenching – intensity of fluorescence decreases

Reasons:

- collision with other molecules (quenchers)
- formation of non-fluorescent complexes
- resonance energy transfer (RET)



Jablonski diagram with collisional quenching and fluorescence resonance energy transfer (FRET). The term Σk_i is used to represent non-radiative paths to the ground state aside from quenching and FRET.

Molecular spectroscopy

Fluorescence – Resonance energy transfer

- emission spectrum of donor overlaps with absorption spectrum of acceptor
- no intermediate photon
- dipole-dipole interaction between donor and acceptor

By Alex M Mooney - Own work, CC BY-SA 3.0,
<https://commons.wikimedia.org/w/index.php?curid=23197114>

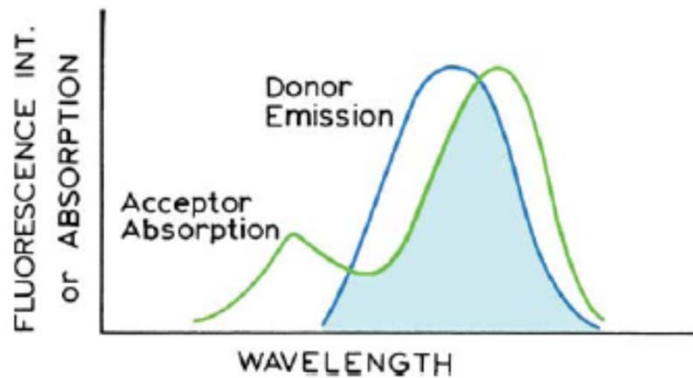
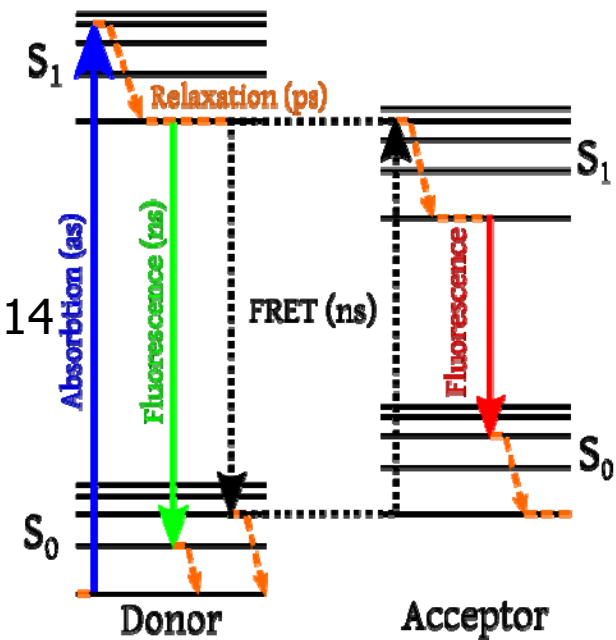


Figure 1.16. Spectral overlap for fluorescence resonance energy transfer (RET).



Molecular spectroscopy

Fluorescence – Eu-based fluorophores

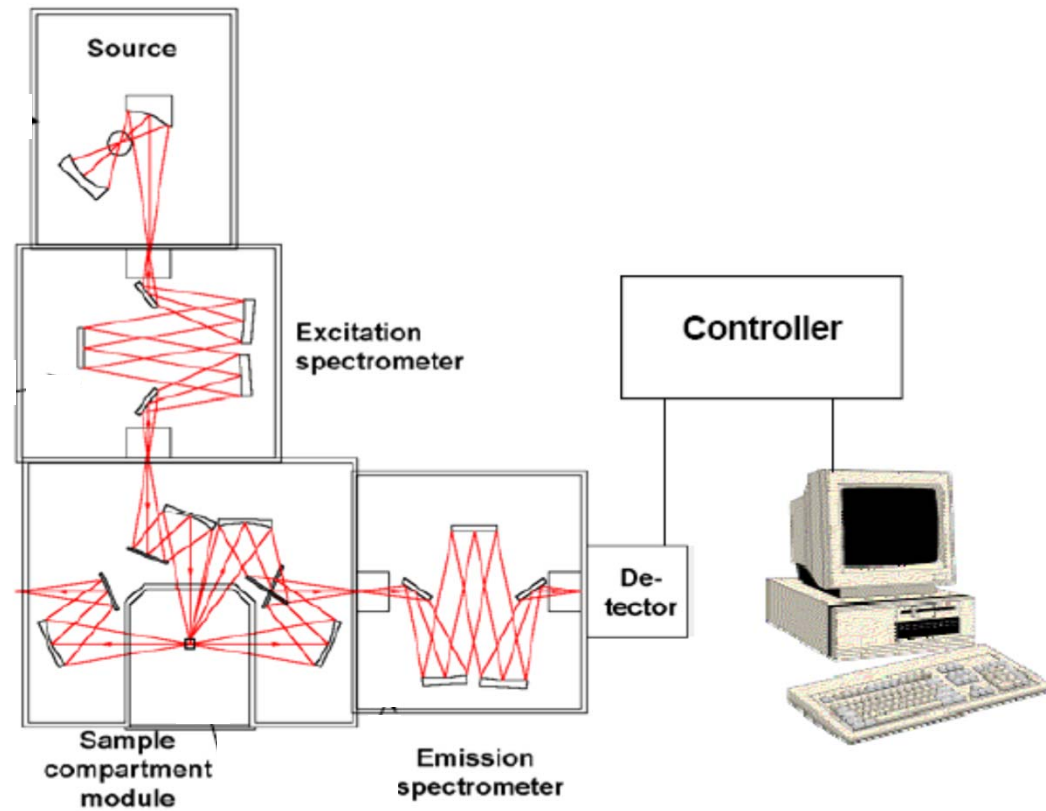


White light illumination

UV light (365 nm)



Spectrofluorimeter – Fluorolog 3



Take-home message

- Atomic emission spectroscopy: flame test, flame emission photometry, atomic absorption spectroscopy, inductive coupled plasma
- Infrared emission spectroscopy: vibrational levels, self-absorption
- Molecular spectroscopy: types of luminescence
- Jablonski diagram: absorption, fluorescence, phosphorescence, internal conversion, intersystem crossing
- Quantum yield and lifetime
- Resonance energy transfer

Összefoglalás

- Atomi emissziós spektroszkópia: lángfestés, lángfotometria, atomabszorpciós spektroszkópia
- Infravörös emissziós spektroszkópia: rezgési szintek, önabszorpció
- Molekulaspektroszkópia: lumineszcencia típusok
- Jablonski-diagram: abszorpció, fluoreszcencia, foszforeszcencia, belső konverzió, intersystem crossing
- Kvantumhatásfok és élettartam
- Rezonáns energiatranszfer