

# Quantitative Spectroscopy

## Line Formation and Stellar Abundances

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Moletai, 2008-08-14

# Who is this guy?

Born in 1972, raised in Marburg, Germany

MSc in Astrophysics (1996, U of London)

Diploma in Physics (1998, U of Heidelberg)

PhD in Astrophysics (2002, U of München)

Postdoc at MPE in Garching

since 2003 research fellow in Uppsala (German and Swedish funding)

since 2008 lecturer at Uppsala University

Research interests: stars from B to K, esp. at low(est) metallicity, chemical evolution of the Galaxy, quantitative spectroscopy, atomic diffusion, Gaia (coordinator for the computation of synthetic observables)



# What will be covered

- I. **Theoretical background** (introductory)  
observables, radiative transfer, opacities and line formation  
model atmosphere output  
how lines depend on  $T_{\text{eff}}$ ,  $\log g$ ,  $\log \varepsilon(X)$  etc.
- II. **Methods of stellar-parameter and chemical-abundance determination**  
fundamental stellar parameters  
photometry (in a nutshell)  
spectroscopy (a practical selection)
- III. **Exercise** (tomorrow afternoon)

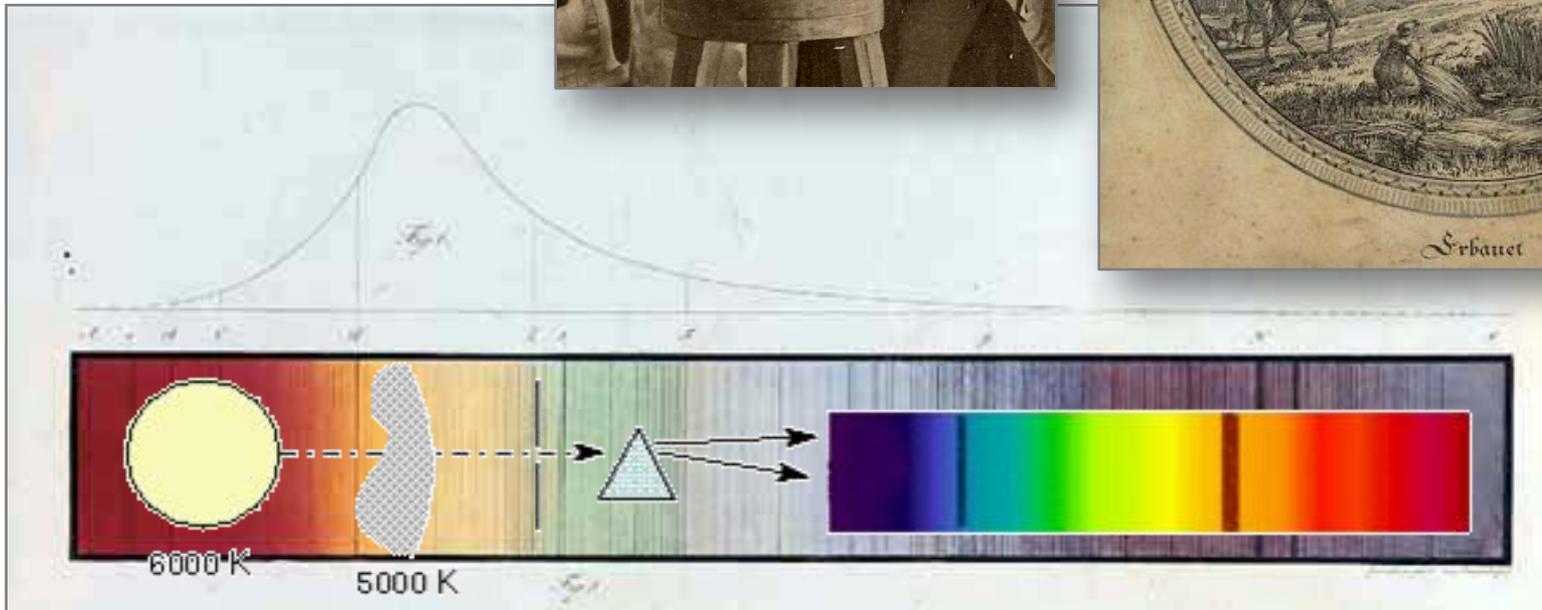
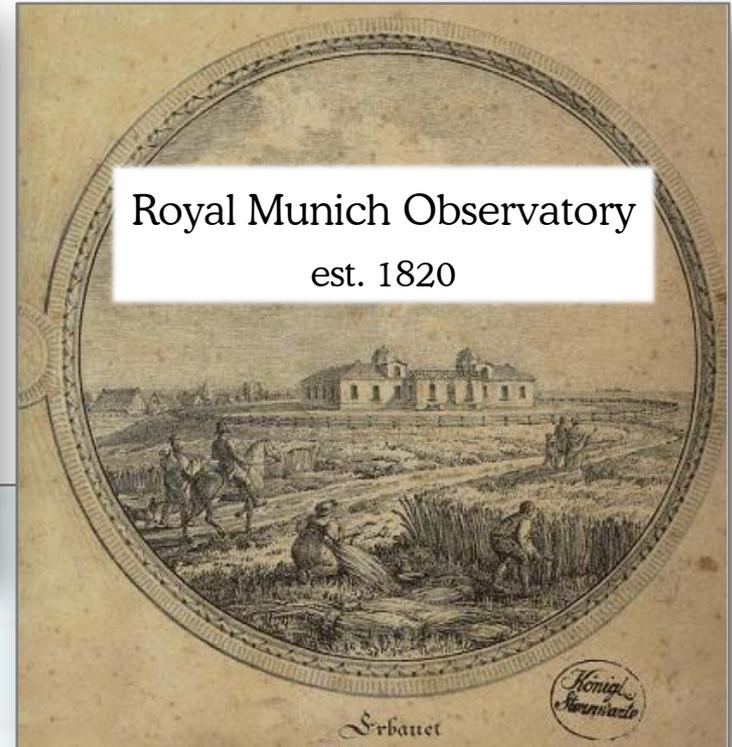


# Stellar spectroscopy in the early days

Joseph von Fraunhofer  
(1787–1826)



Royal Munich Observatory  
est. 1820



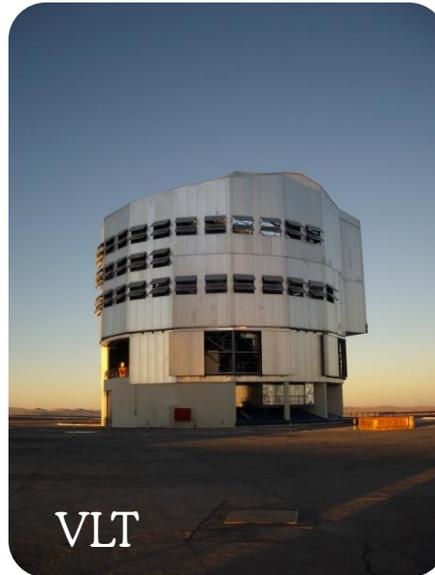
# Stellar spectroscopy today

observation vs. theory

telescope

spectrograph

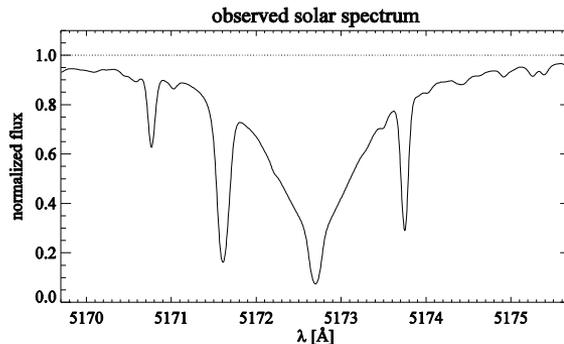
CCD



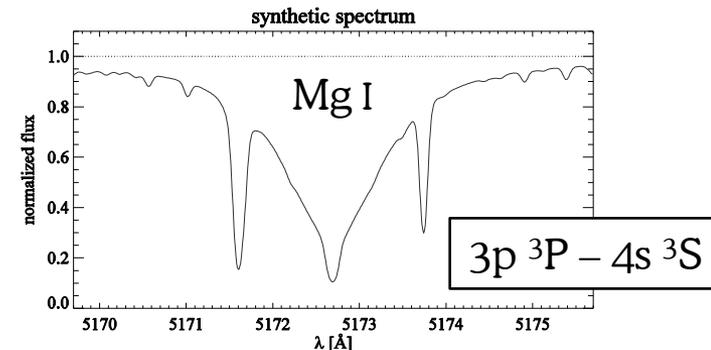
concepts

approximations

numerical model



comparison  
to constrain  
thermodynamic variables  
and abundances



NOT

FIES

E2V 42-40

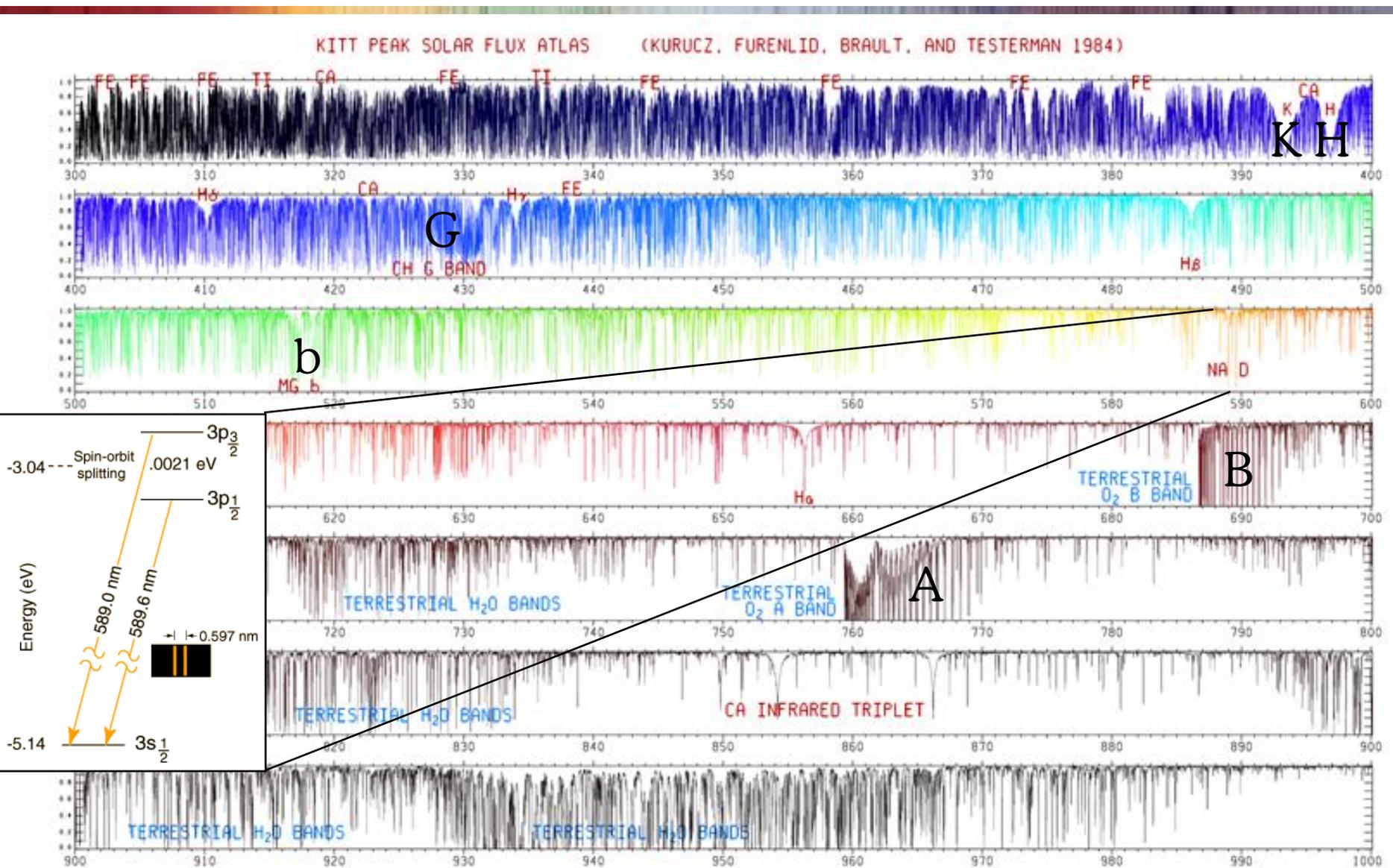
$$\frac{dI_\nu}{d\tau_\nu} = -I_\nu + S_\nu$$

LTE

$$\frac{d}{dx} \mathcal{F} = 0$$

MLT

# The Solar spectrum



# What are stellar parameters?

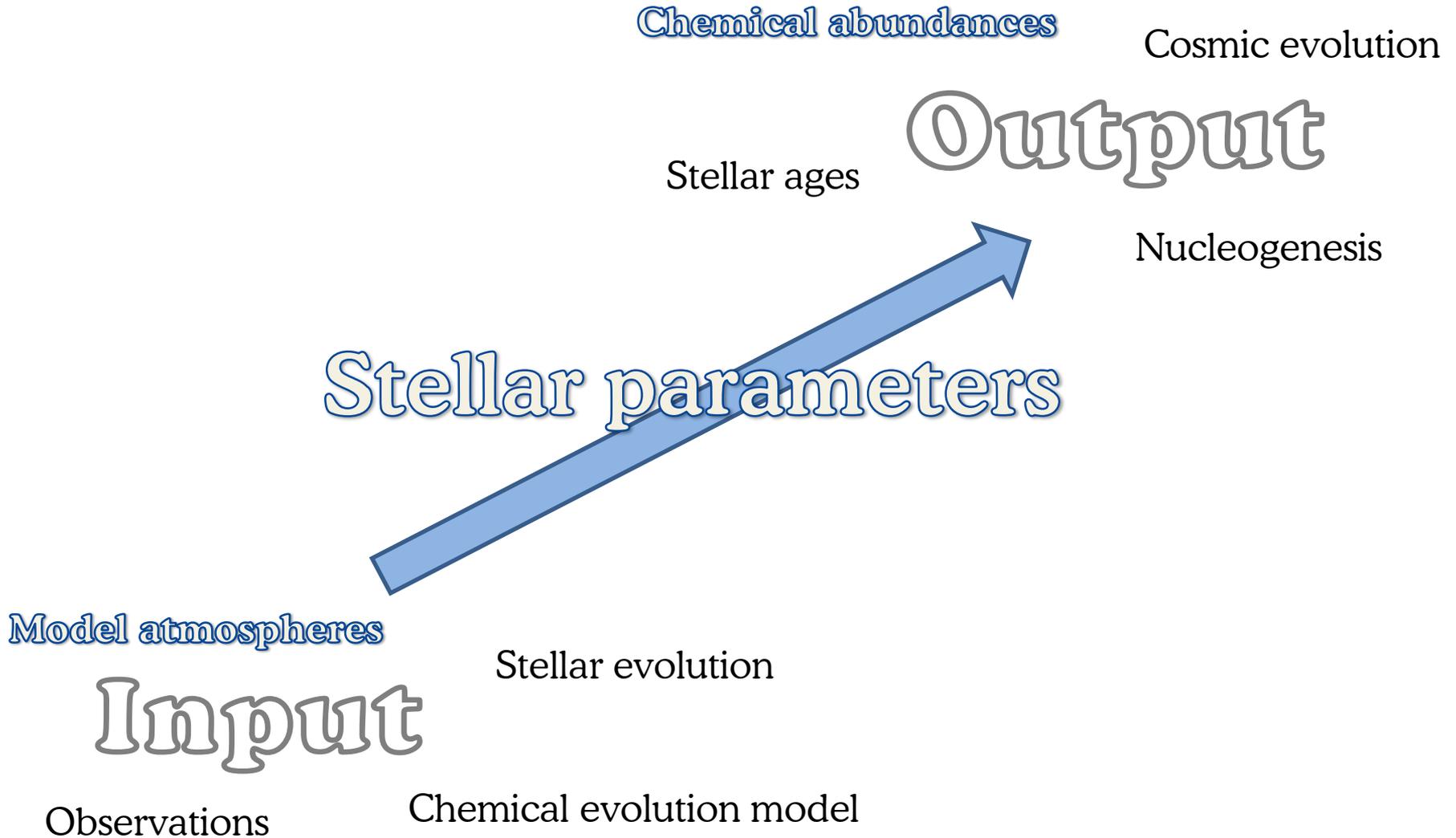
There are different ways of looking at what defines a stars:

stellar-structure view  $M, \mathcal{L}, X, Y, Z, R, v_{\text{rot}}, t, \dots$

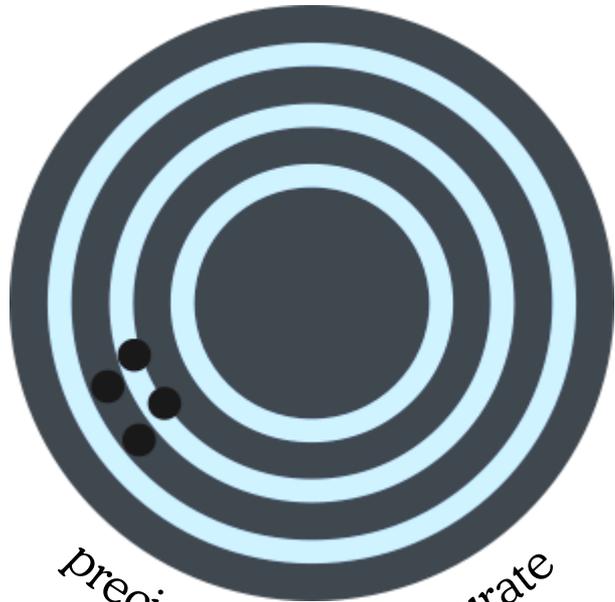
stellar-atmosphere view  $\mathcal{F}_v, T_{\text{eff}}, \log g, [X_i/H], v_{\text{rot}} \sin i, \dots$   
 $\log (G M / R^2)$

While the prior is (often) more fundamental, the latter is more directly related to observations (photospheres!) and generally speaking more applicable. In this lecture, I will follow the latter view.

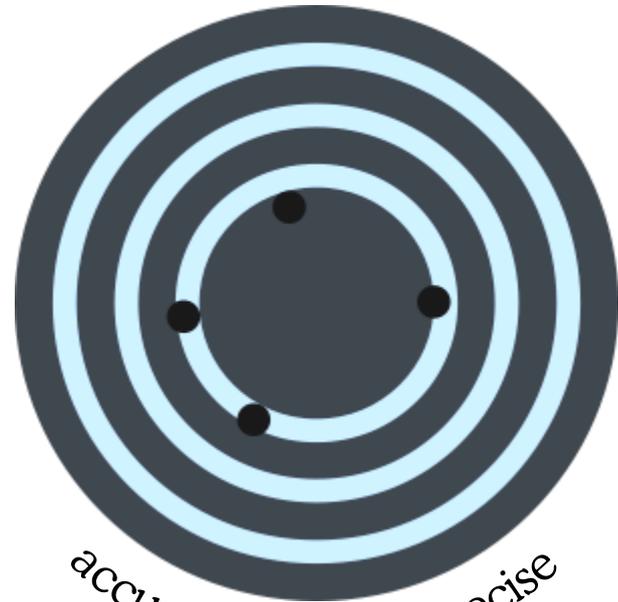
# Linking input to output



# Precision *vs.* accuracy



*precise, but not accurate*



*accurate, but not precise*

NB: Some projects may require high precision *and* accuracy, while for others it will suffice to reach some level of precision.

# Stellar atmosphere: a definition

descriptive: the layers of a star from which we receive photons = the layers we can see

physical:  $0 \leq \tau_\nu \leq 10$

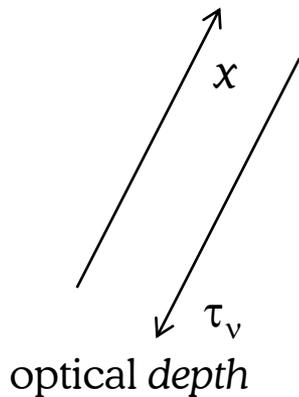
where  $\tau_\nu = (-) \int_0^L \kappa_\nu \rho dx$  is the **optical height**,

$x$  measures the geometrical path [cm],

$\rho$  is the mass density [ $\text{g cm}^{-3}$ ],

$\kappa_\nu$  is the mass absorption coefficient [ $\text{cm}^2 \text{g}^{-1}$ ] and

$L$  is the path length (see Gray, ch. 5, p. 113)



simple extinction law:  $\mathcal{I}(\nu) = \mathcal{I}_0(\nu) \exp(-\tau_\nu)$

# Stellar atmospheres: typical figures

## The Sun

$$M = 2 \times 10^{33} \text{ g} = M_{\odot}$$

$$R = 7 \times 10^{10} \text{ cm} = R_{\odot}$$

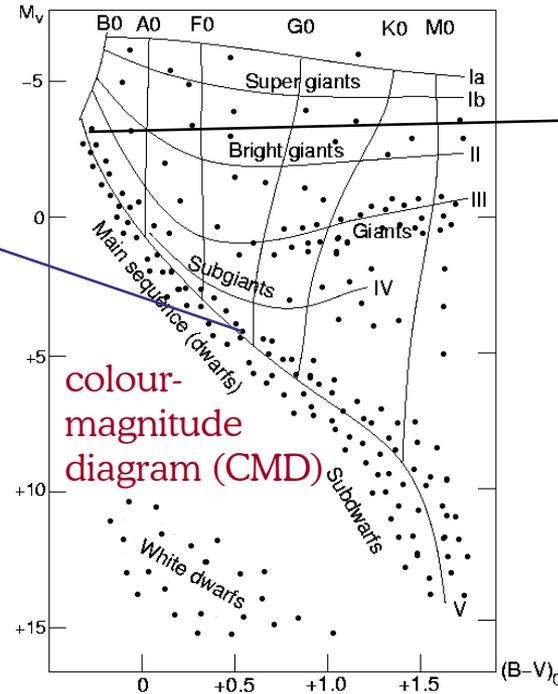
$$\mathcal{L} = 4 \times 10^{33} \text{ erg/s} = L_{\odot}$$

photosphere:

$$\Delta R \approx 200 \text{ km} < 10^{-3} R_{\odot}$$

$$n \approx 10^{15} \text{ cm}^{-3}$$

$$T \approx 6000 \text{ K}$$



an O star

$$M \sim 50 M_{\odot}$$

$$R \sim 20 R_{\odot}$$

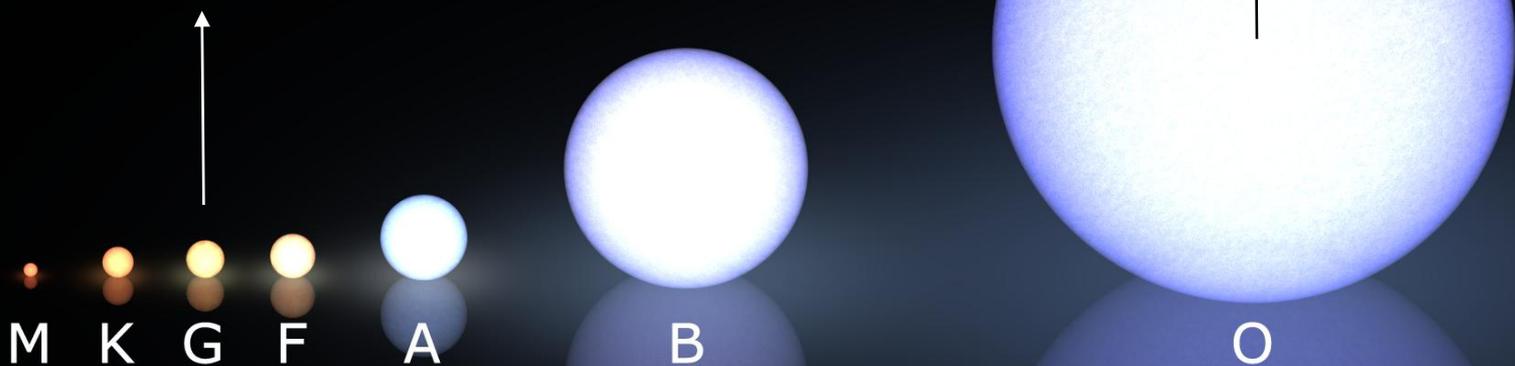
$$\mathcal{L} \sim 10^6 L_{\odot} (\propto M^3)$$

photosphere:

$$\Delta R \approx 0.1 R_{\odot}$$

$$n \approx 10^{14} \text{ cm}^{-3}$$

$$T \approx 40\,000 \text{ K}$$



# Abundance nomenclature

**Mass fractions:** let  $X$ ,  $Y$ ,  $Z$  denote the mass-weighted abundances of H, He and all other elements (“metals”), respectively, normalized to unity ( $X + Y + Z = 1$ ).

example:  $X = 0.739$ ,  $Y = 0.249$ ,  $Z = 0.012$  for the Sun

**The 12 scale:**  $\log \varepsilon(X) = \log (n_X / n_H) + 12$  ( $\log \varepsilon(H) \equiv 12$ )

example:  $\log \varepsilon(O)_{\odot} \approx 8.7$  dex, i.e., oxygen, the most abundant metal, is 2000 times less abundant than H in the Sun (the exact value is currently hotly debated!)

**Square-bracket scale:**  $[X/H] = \log (n_X / n_H)_{\star} - \log (n_X / n_H)_{\odot}$

example:  $[\text{Fe}/\text{H}]_{\text{HE0107-5240}} = -5.3$  dex, i.e., this star has an iron abundance a factor of 200 000 below the Sun (Christlieb *et al.* 2002)

# Intensity and flux

The Sun is one of the few stars whose surface we can resolve  $\Leftrightarrow$  measure the so-called specific intensity

$$\mathcal{I}_\nu = dE_\nu / \cos \vartheta \, dA \, d\Omega \, dt \, d\nu \quad [\text{J} / \text{m}^2 \text{ rad s Hz}]$$

Usually, we measure stellar fluxes

$$\mathcal{F}_\nu = \int dE_\nu / dA \, dt \, d\nu \quad [\text{J} / \text{m}^2 \text{ s Hz}]$$

Clearly, the **flux**  $\mathcal{F}_\nu = \int \mathcal{I}_\nu \cos \vartheta \, d\Omega$  and it **measures** the **anisotropy of the radiation field**.

Example: the Solar flux above the Earth's atmosphere

$$\mathcal{F}(\odot) = 1.36 \text{ kW} / \text{m}^2$$

# Flux constancy and luminosity

Stellar atmospheres are much too cool and tenuous to fuse nuclei  
⇒ the energy coming from the stellar core is merely transported through the atmosphere, either by radiation or convection.

$$\mathcal{F}(x) = \text{energy} / \text{unit area} / \text{unit time} \quad [\text{J m}^{-2} \text{s}^{-1}] = [\text{W m}^{-2}]$$

$$d \mathcal{F}(x) / dx = 0 \quad (\text{generally: } \nabla \mathcal{F} = 0)$$

The spectrum of  $\mathcal{F}$  (i.e.  $\mathcal{F}_\nu$ ) will change with  $r$ , but not the integral value.

If  $\mathcal{F}_{\text{rad}} \gg \mathcal{F}_{\text{conv}}$ , then one speaks of **radiative equilibrium**.

(Karl Schwarzschild 1873–1916)

The total energy output of a star is called its **luminosity**

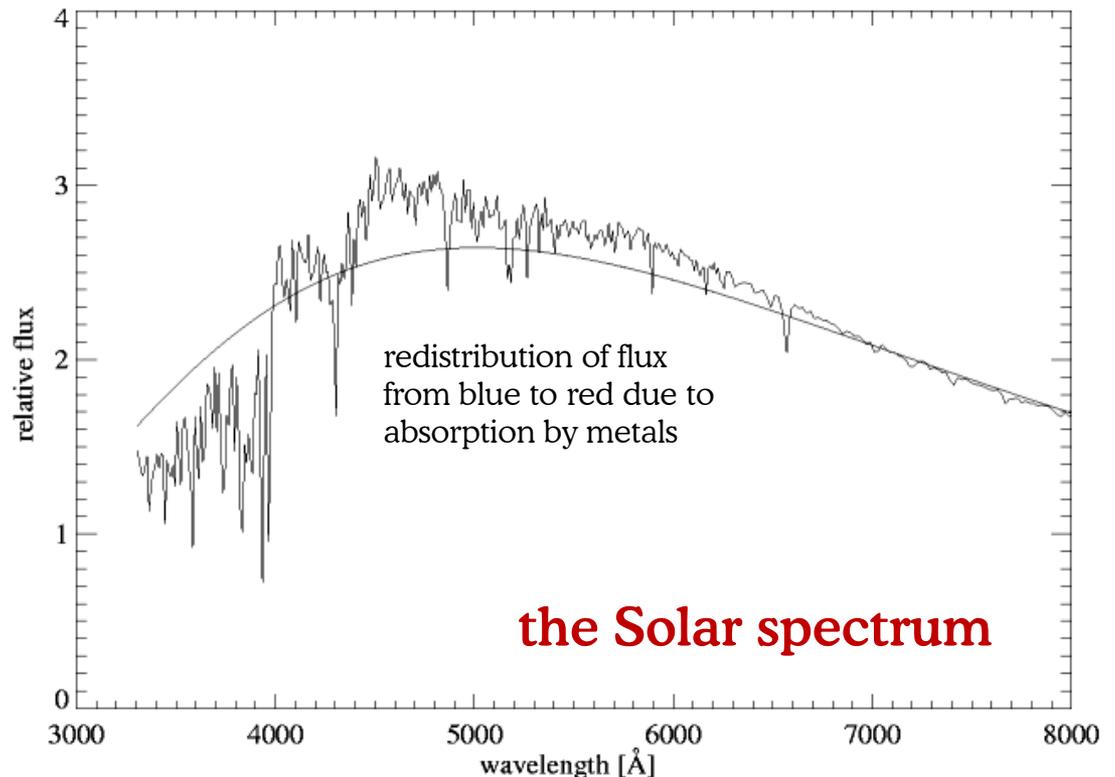
$$\mathcal{L} = 4\pi R^2 \mathcal{F}(R)$$

# Stellar spectra

Luckily, **stars** (and other celestial bodies) are not in thermodynamic equilibrium (TE) and **do not shine like blackbodies**.

(Astronomy would be the dullest of all sciences!)

In contrast to  $\mathcal{B}_\lambda$ ,  $\mathcal{I}_\lambda$  depends on plasma properties and the viewing angle. **One cannot use TE to describe starlight.**



# TE statistics

Particle velocities are assumed to be Maxwellian:

$$\frac{n(v)}{n_{\text{tot}}} dv = \left( \frac{m}{2\pi kT} \right)^{\frac{3}{2}} e^{-\frac{mv^2}{2kT}} dv$$

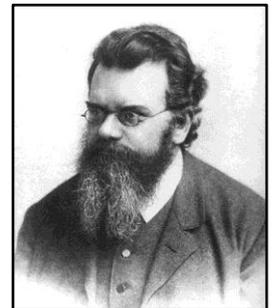
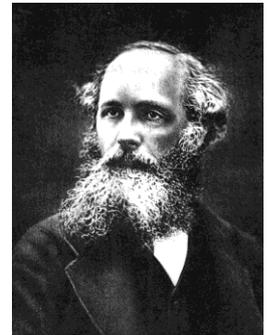
Excitation follows the Boltzmann distribution:

$$\frac{n_u}{n_{\text{tot}}} = \frac{g_u}{u(T)} e^{-\frac{\chi_u}{kT}}$$

Ionization can be computed via the Saha equation:

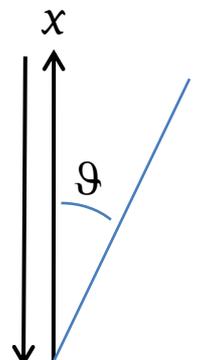
$$\frac{n_{\text{II}}}{n_{\text{I}}} P_e = \frac{(2\pi m_e)^{3/2} kT^{5/2}}{h^3} \frac{2u_{\text{II}}(T)}{u_{\text{I}}(T)} e^{-\frac{I}{kT}}$$

In **local thermodynamic equilibrium** (LTE), these are applied *locally*.



# The basics of radiative transfer

When the stellar photons interact with the stellar-atmosphere matter, photons can be absorbed and re-emitted. This is the basic message of the **radiative transfer equation**.



$$d\mathcal{I}_\nu = -\kappa_\nu \rho \mathcal{I}_\nu dx + j_\nu \rho dx$$

or  $j_\nu$ : emission coefficient

$$\cos\vartheta \, d\mathcal{I}_\nu / d\tau_\nu = +\mathcal{I}_\nu - \mathcal{S}_\nu$$

with

$$\mathcal{S}_\nu = j_\nu / \kappa_\nu$$

the **source function**

In LTE,  $\mathcal{S}_\nu = \mathcal{B}_\nu$  the **Planck function**

$\mathcal{B}_\nu$  has a number of wonderful properties: it does not depend on material properties (only  $T$ ) and increases monotonically with increasing  $T$  for all  $\nu$ .

The integral  $\int \mathcal{B} \cos\vartheta \, d\Omega$  yields  $\sigma T^4$  (Stefan-Boltzmann law).

Similarly,  $T_{\text{eff}}$  is defined:  $\mathcal{F}_{\text{Bol}}(\text{Earth}) = \left(\frac{\theta}{2}\right)^2 \sigma T_{\text{eff}}^4$   $\theta$ : angular diameter  
 bolometric flux above  
 Earth's atmosphere

# Opacities

## Continuous opacity

Caused by *bf* or *ff* transitions

In the optical and near-IR of cool stars,  $H^-$  ( $I = 0.75$  eV) dominates:

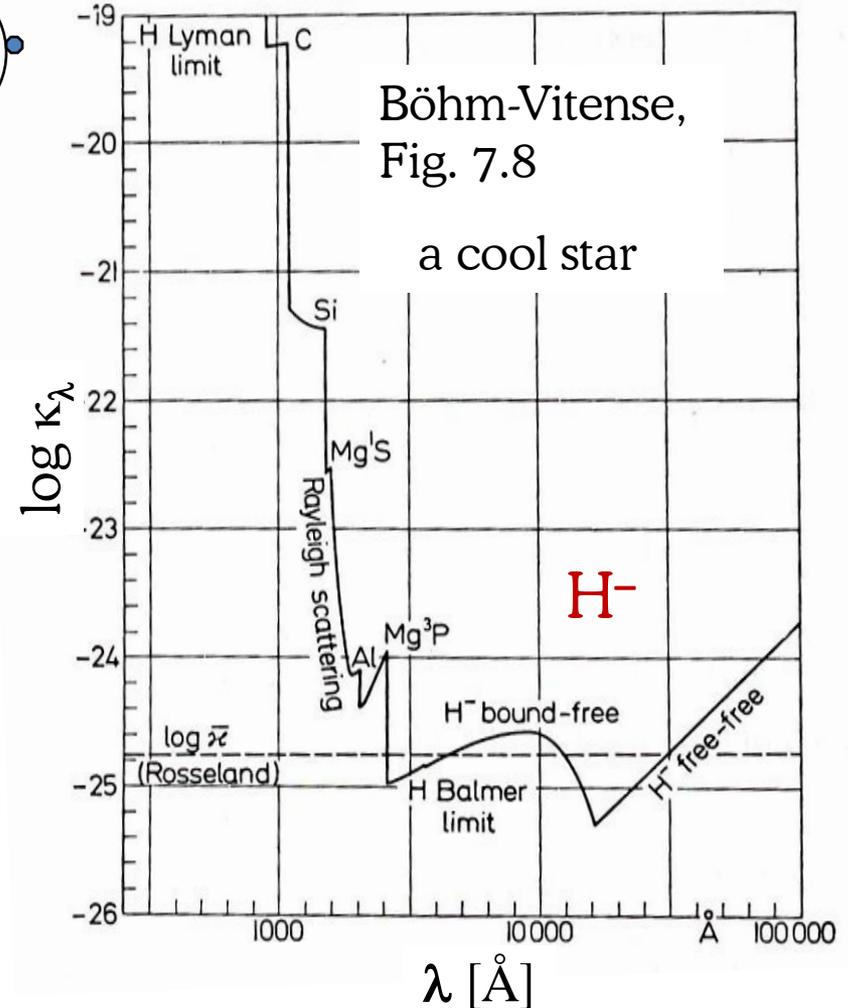
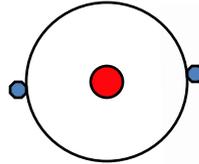
$$\kappa_{\nu}(H^-_{bf}) = \text{const. } T^{-5/2} P_e \exp(0.75/kT)$$

NB: There is only 1  $H^-$  per  $10^8$  H atoms in the Solar photosphere.

## Line opacity (*all the lines you see!*)

Caused by *bb* transitions

Need to know  $\log gf$ , damping and assume an abundance



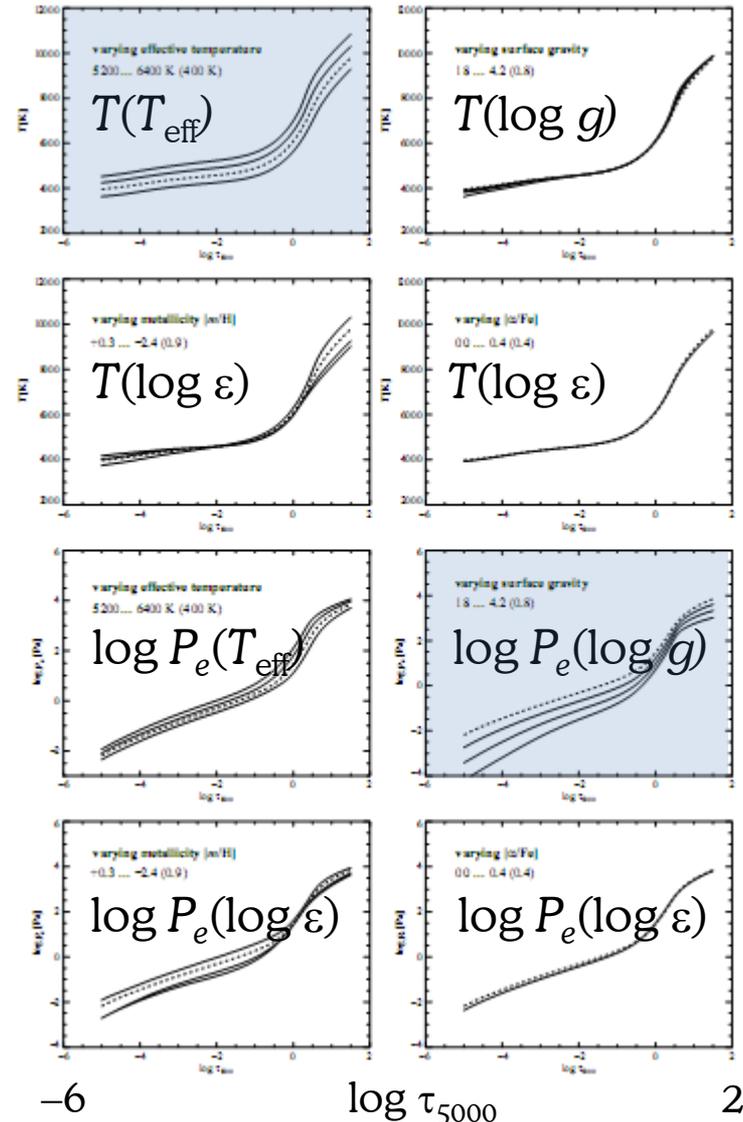
# Model atmosphere output

A 1D model atmosphere is a tabulation of various quantities as a function of (optical) depth:

- $T$  (temperature)
- $P_g$  (gas pressure)
- $P_e$  (electron pressure)
- $\mathcal{F}_v$  (esp. *surface flux*) etc.

as computed under certain input assumptions:

- $T_{\text{eff}}$  (effective temperature)
- $\log g$  (surface gravity)
- $\log \varepsilon(X_i)$  (chemical composition)
- hydrostatic equilibrium
- LTE (local thermodynamic equilibrium)
- MLT (mixing-length theory) and
- a statistical representation of opacities (either via opacity distribution functions, ODF, or opacity sampling, OS).



# How spectral lines originate

The formation of absorption lines can be qualitatively understood by studying how  $\mathcal{S}_\nu$  changes with depth.

$$W_\lambda \propto d \ln \mathcal{S}_\nu / d \tau_\nu$$

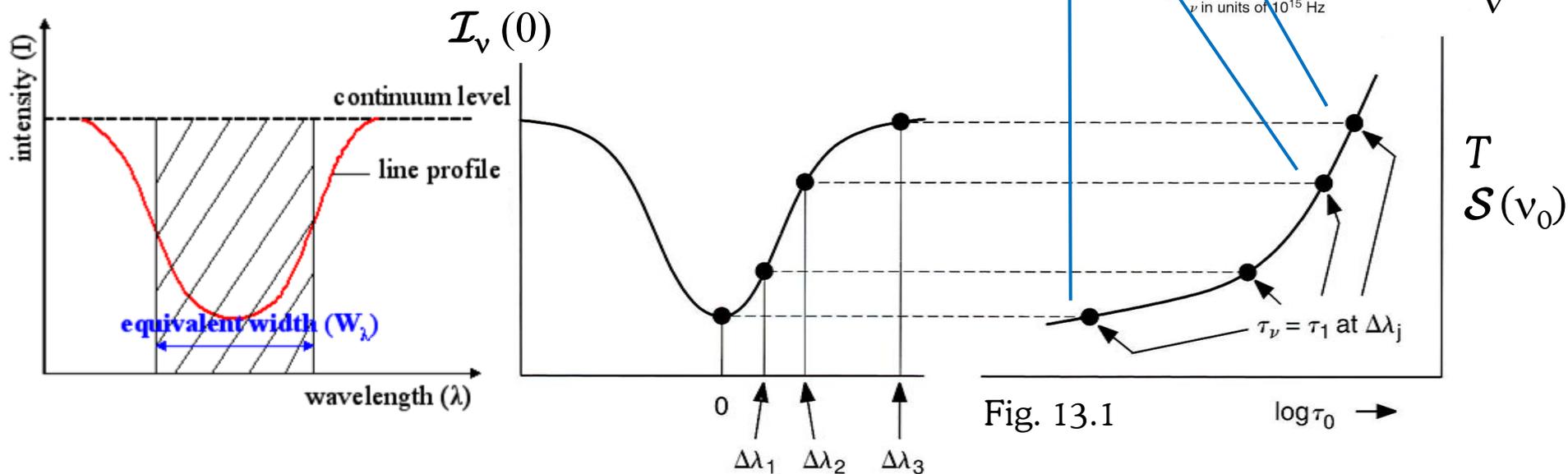


Fig. 13.1

# Spectral lines as a function of abundance

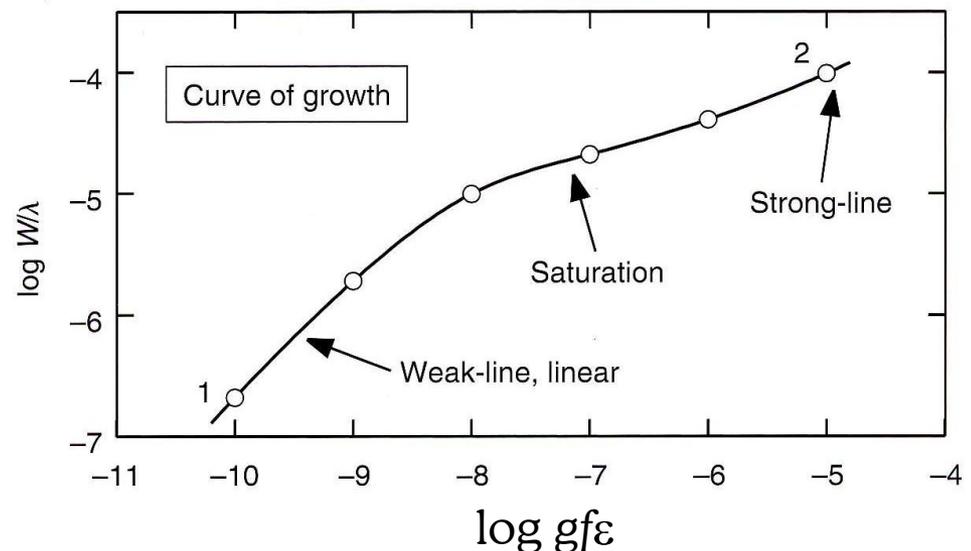
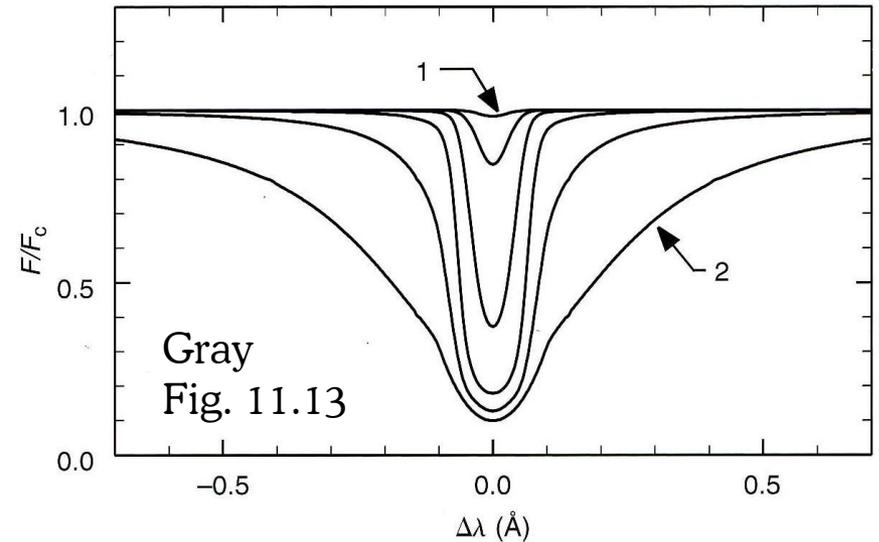
Starting from low  $\log \varepsilon$  (low  $\log gf$ ), the line strength is directly proportional to  $\log gf\varepsilon$ :

$$W_\lambda \propto gf n_X$$

When the line centre becomes optically thick, the line begins to saturate. The dependence on abundance lessens. Only when damping wings develop, the line can grow again in a more rapid fashion:

$$W_\lambda \propto \text{sqrt}(gf n_X)$$

Weak lines are thus best suited to derive the elemental composition of a star, given that they are well-observed (blending!)



# Broadening of spectral lines

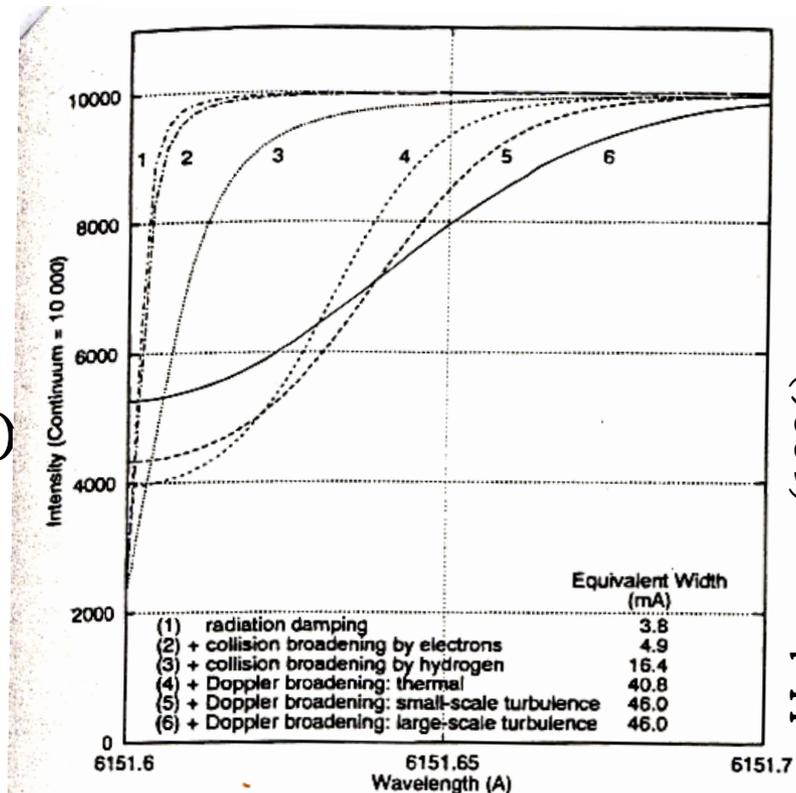
There are numerous broadening mechanisms which influence the strength and apparent shape of spectral lines:

microscopic

1. natural broadening  
(reflecting  $\Delta E \Delta t \geq h/2\pi$ )
2. thermal broadening
3. microturbulence  $\xi_{\text{micro}}$   
(treated like extra thermal br.)
- (4. isotopic shift, *hfs*, Zeeman effect)
5. collisions (H:  $\gamma_6$ ,  $\log C_6$ ;  $e^-$ :  $\gamma_4$ )  
(important for strong lines)

macro

6. macroturbulence  $\Xi_{\text{rt}}$
7. rotation
- (8. instrumental broadening)



Holweger (1996)

Fig. 3. Synthetic (half-)profiles of Fe I 6151.6 Å (Mult. 62, E.P. = 2.2eV) showing the cumulative effect of various broadening mechanisms.

# Microturbulence and damping

If lines of intermediate or high strength return too high abundances, then the microturbulence or the damping constants are (both) underestimated (the  $gf$  values can also be systematically off).

**Use an element with lines of all strengths to determine  $\xi$ .** In most cases, this will be an iron-group elements.

Hydrodynamic (“3D”) models are presently in an adolescent phase and will hopefully do away with the need for micro/macro-turbulence.

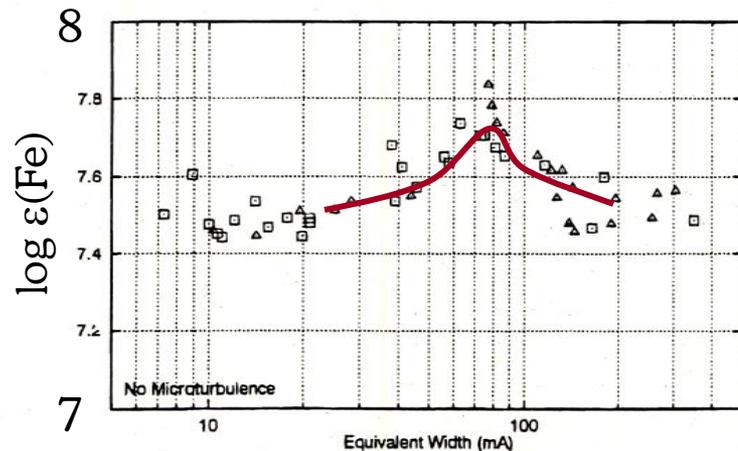


Fig. 7. Same as Fig. 6, but neglecting microturbulence.

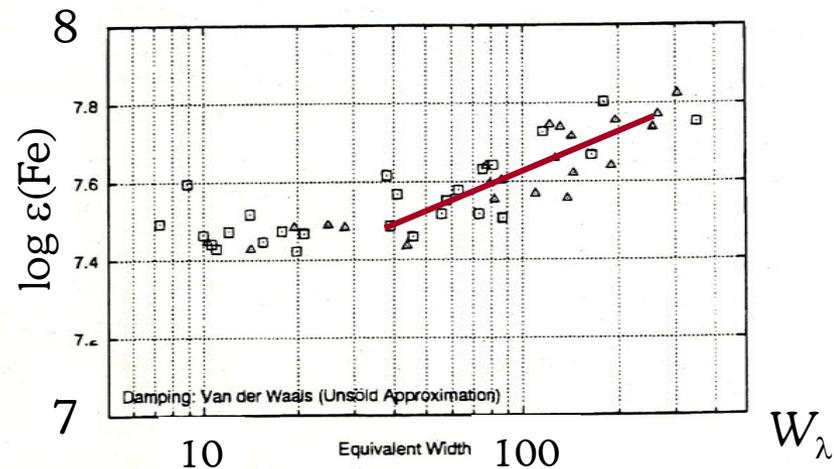
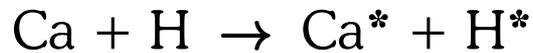


Fig. 5. Iron abundances derived from individual solar Fe I lines and Hanover  $gf$ -values. The two samples shown are from [4] (squares) and [18] (triangles). The deviation of the stronger lines indicates that the adopted damping constants are too small.

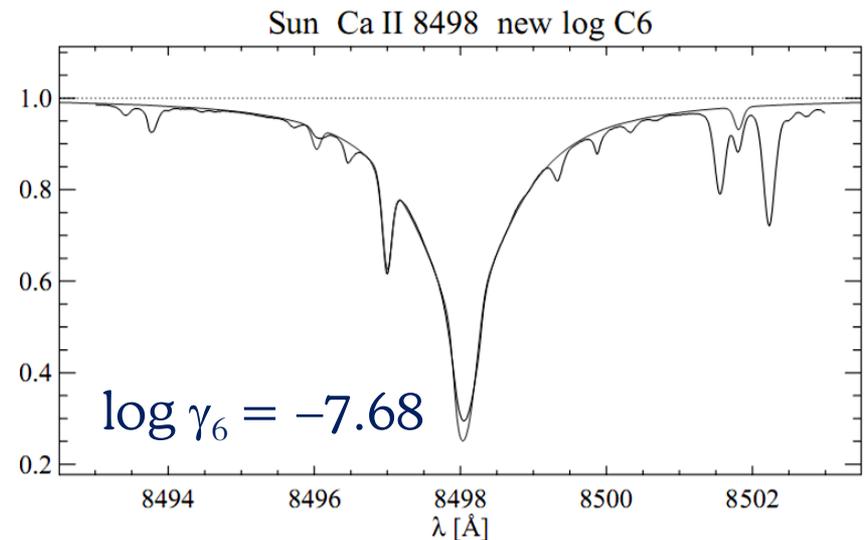
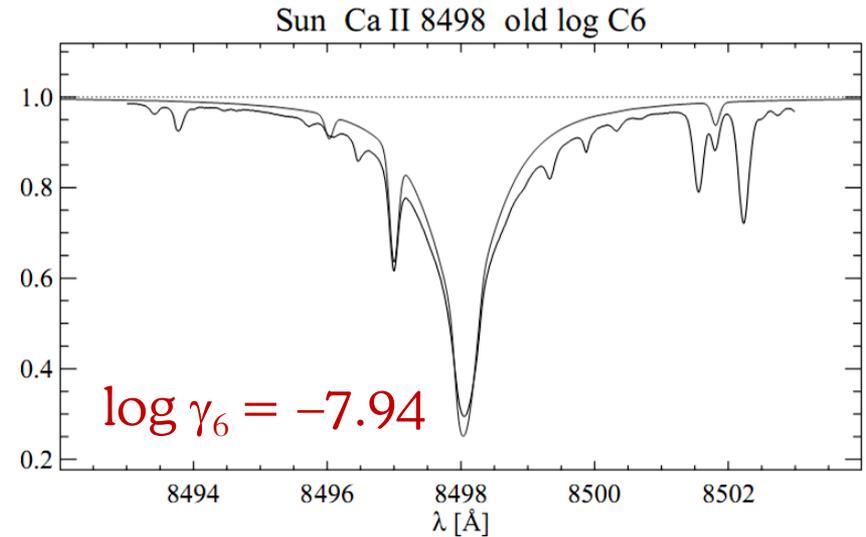
# Broadening of spectral lines: an example

The Ca II triplet lines are broadened by elastic collisions with hydrogen:



Detuning  $\Delta\nu = C_n / R^n$ : here  $C_6$

Progress in the QM description of this interaction has led to a better understand of the profiles of these (and many other) lines (Anstee & O'Mara 1991, 1995).



$F/F_c$

# Spectral lines as a function of $T_{\text{eff}}$

The **strength of a weak line** is **proportional to** the ratio of line to continuous absorption coefficients,  $l_{\nu} / \kappa_{\nu}$ . Evaluation of this ratio can tell us about the  $T_{\text{eff}}$  sensitivity of spectral lines:

$$R = l_{\nu} / \kappa_{\nu} = \text{const. } T^{5/2} / P_e \exp-(\chi + 0.75)/kT$$

for a neutral line of an element that is mostly ionized.

$$\text{Fractional change with } T: 1/R \, dR/dT = (\chi + 0.75 - I)/kT^2$$

$\Rightarrow$  depending  $\chi$  on **neutral lines decrease with  $T_{\text{eff}}$**  by between 10 and 30% per 100 K (typically 0.07 dex per 100 K). Lines of different  $\chi$  can be used to constrain  $T_{\text{eff}}$  (**excitation equilibrium** condition).

For **ionized lines** of mainly ionized elements, one finds low sensitivities to  $T_{\text{eff}}$ , except those **with a large  $\chi$** . These **become stronger with  $T_{\text{eff}}$**  by up to 20% per 100 K.

# Spectral lines as a function of $\log g$

The  $T_{\text{eff}}$  sensitivity of spectral lines may be surpassed by sensitivities with respect to other stellar parameters.

## Sensitivity to $\log g$ in cool stars?

Case 1: (weak) neutral line of an element that is mainly ionized

$W_\lambda$  is proportional to the ratio of line to continuous absorption coefficients,  $l_\nu / \kappa_\nu$ .

$$n_{r+1} / n_r = \Phi(T) / P_e \quad \Leftrightarrow \quad n_r \approx \text{const. } P_e$$

$\Rightarrow l_\nu / \kappa_\nu \neq f(P_e)$       neutral lines do not depend on  $\log g$

Case 2: ionized line of an element that is mainly ionized

**(universal)**  $\log g$  sensitivity via the continuous opacity of  $\text{H}^-$

**NB:** for strong lines, a damping-related  $\log g$  sensitivity comes into play.

# LTE vs. NLTE

Occupation, excitation & ionization are assumed to be local properties  
⇒ **Saha-Boltzmann statistics**

Assuming the  $T$ - $P$ - $\tau$  relation to be known, all you need to calculate a line strength is

- (a) the level energies and statistical weights involved
- (b) the transition probability
- (c) broadening mechanisms (microturbulence, van-der-Waals damping)

*Photons carry non-local information*

Occupation, excitation & ionization depend on the microphysics (radiation field, collisions etc.)

**One needs to know (and master!) a whole lot of atomic physics.**

One also needs to solve the involved numerical problem of radiative transfer plus **rate equations**:

$$n_i \sum_{j \neq i} (R_{ij} + C_{ij}) = \sum_{j \neq i} n_j (R_{ji} + C_{ji})$$

**While LTE may be an acceptable approximation for a cool-star photosphere on the whole, it can be very wrong for specific lines.**

# Fundamental stellar parameters

$T_{\text{eff}}$ : via  $\mathcal{F}_{\text{Bol}}$  and  $\theta$  (see IRFM below).

To get  $\theta$ , one uses interferometry and model-atmosphere theory (limb darkening!).

$\log g$ : Newton's law, needs  $M$  and  $R$ .

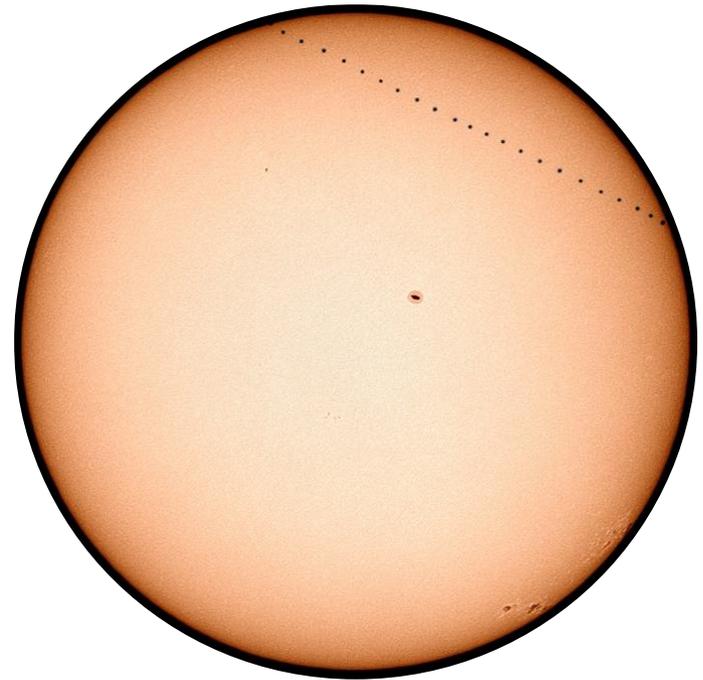
So usually one needs  $\pi$  (parallax) and  $\theta$ . Gaia is the key  $\pi$  mission (launch 2012).

$M$  needs to be inferred from stellar evolution.

Exception: eclipsing binaries.

$[m/H]$ : via meteorites (only for the Sun), which lack important (volatile) elements like CNO and noble gases.

In principle, asteroseismology can provide compositions of other stars.



# Photometry: pros vs. cons

Photometry is

- ✓ an efficient way of determining stellar parameters,
- ✓ can probe very deep,
- ✓ freely available (surveys!),
- ✓ comparatively cheap to obtain.

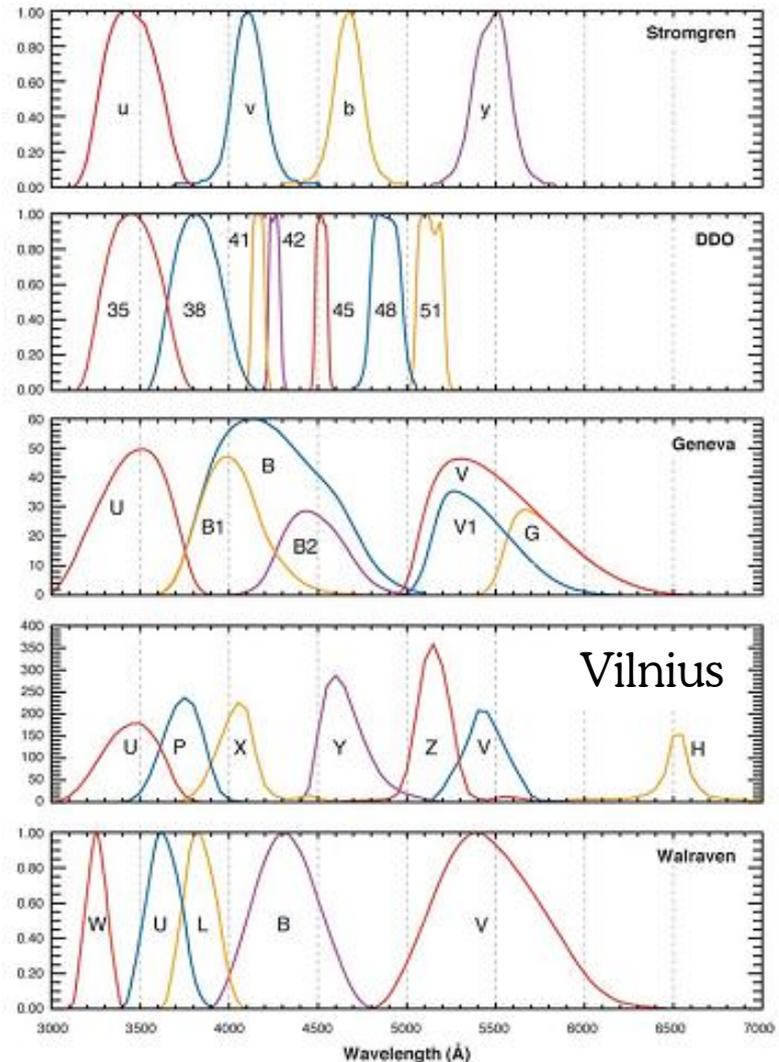
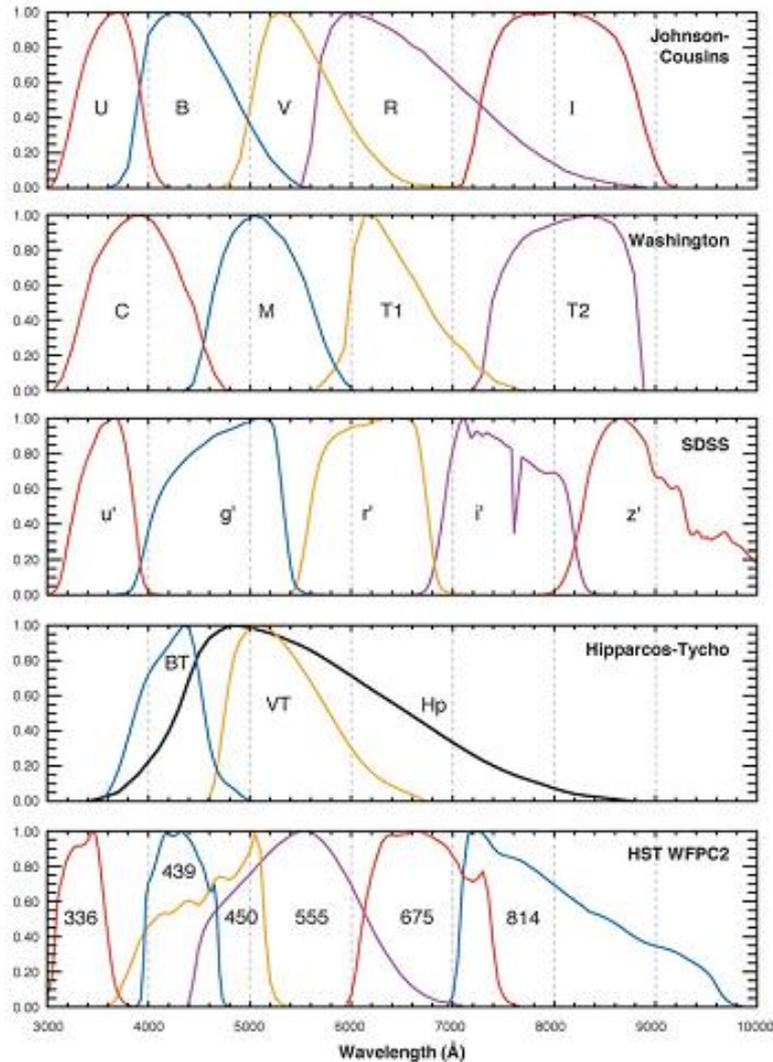
However, photometry is

- ❑ limited in which parameters can be derived,
- ❑ subject to extra parameters (reddening!)
- ❑ subject to parameters that cannot be determined well ( $\xi$ ,  $[\alpha/\text{Fe}]$ ).



(c) F. Bresolin

# Photometric standard systems

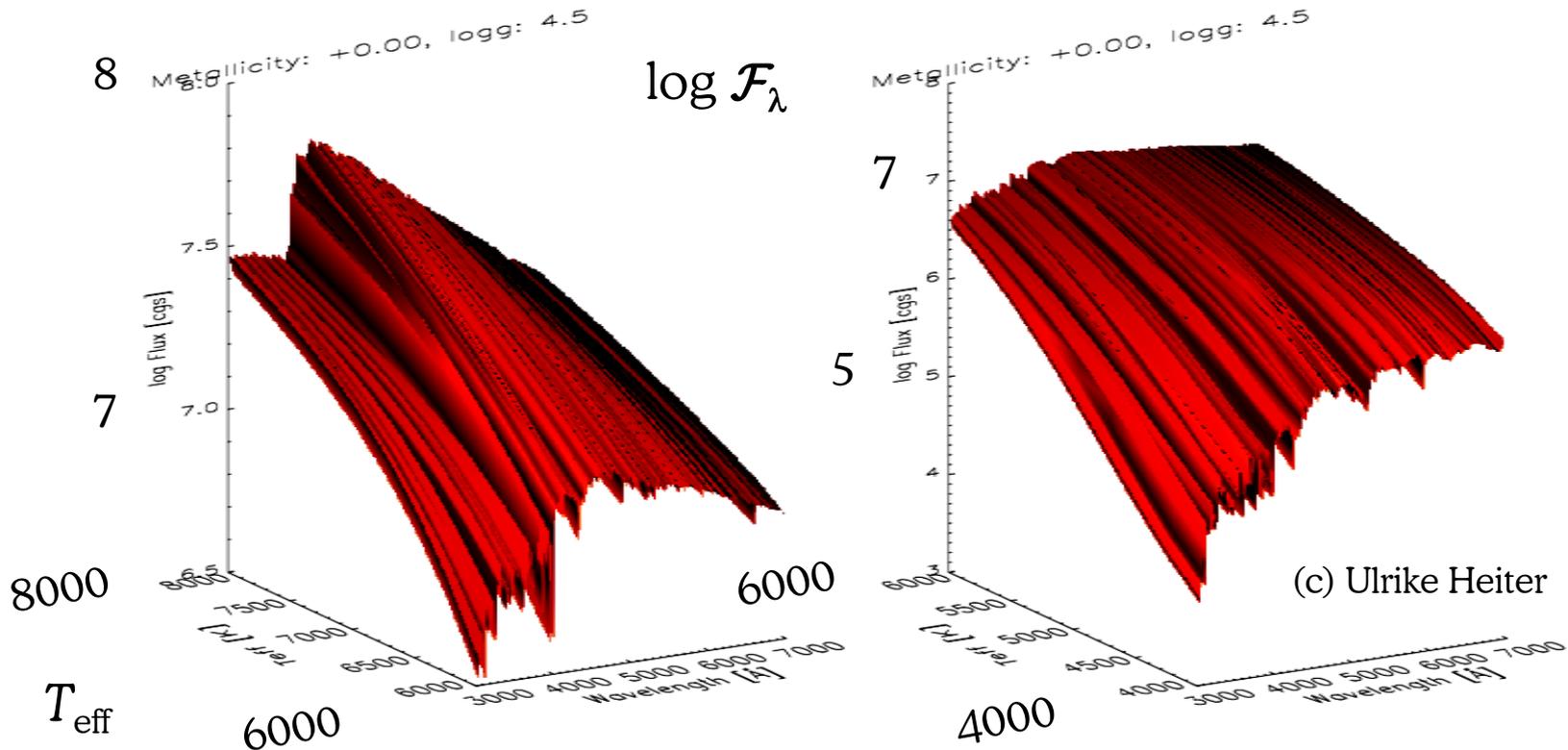


**Warning:** there is often more than one filter set for one system!

# Photometry: $T_{\text{eff}}$ dependence

$T_{\text{eff}}$  variations dominate the flux variations of cool stars.

In the BB approximation to stellar fluxes, it suffices to measure the flux at two points to uniquely determine  $T$ . In reality,  $[m/\text{Fe}]$  and reddening complicate the derivation of photometric stellar parameters.

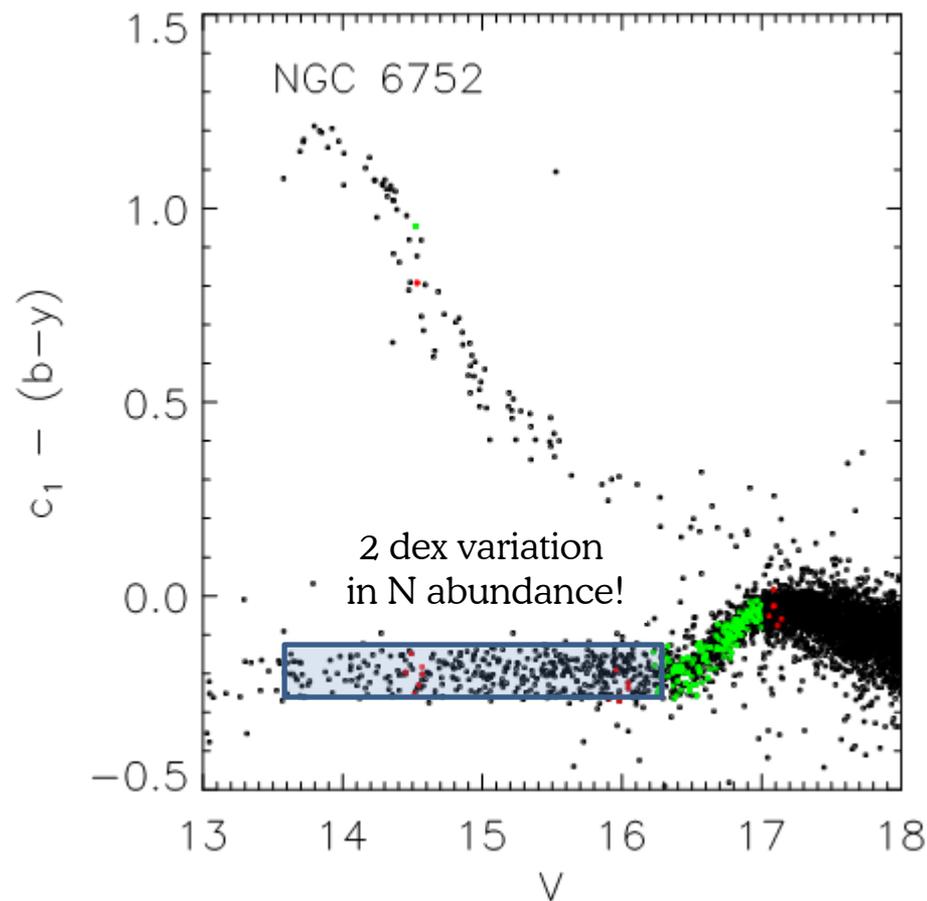


# Photometry: metallicities

After  $T_{\text{eff}}$ , the *global* metallicity has the largest influence on stellar fluxes (with the potentially disastrous exception of **reddening!**).

But the **precision** with which metallicities can be determined **is limited** (of order 0.3 dex). In addition, it is difficult to determine metallicities for stars with  $[\text{Fe}/\text{H}] < -2$ , as classical indicators like  $\delta(U - B)$  lose sensitivity.

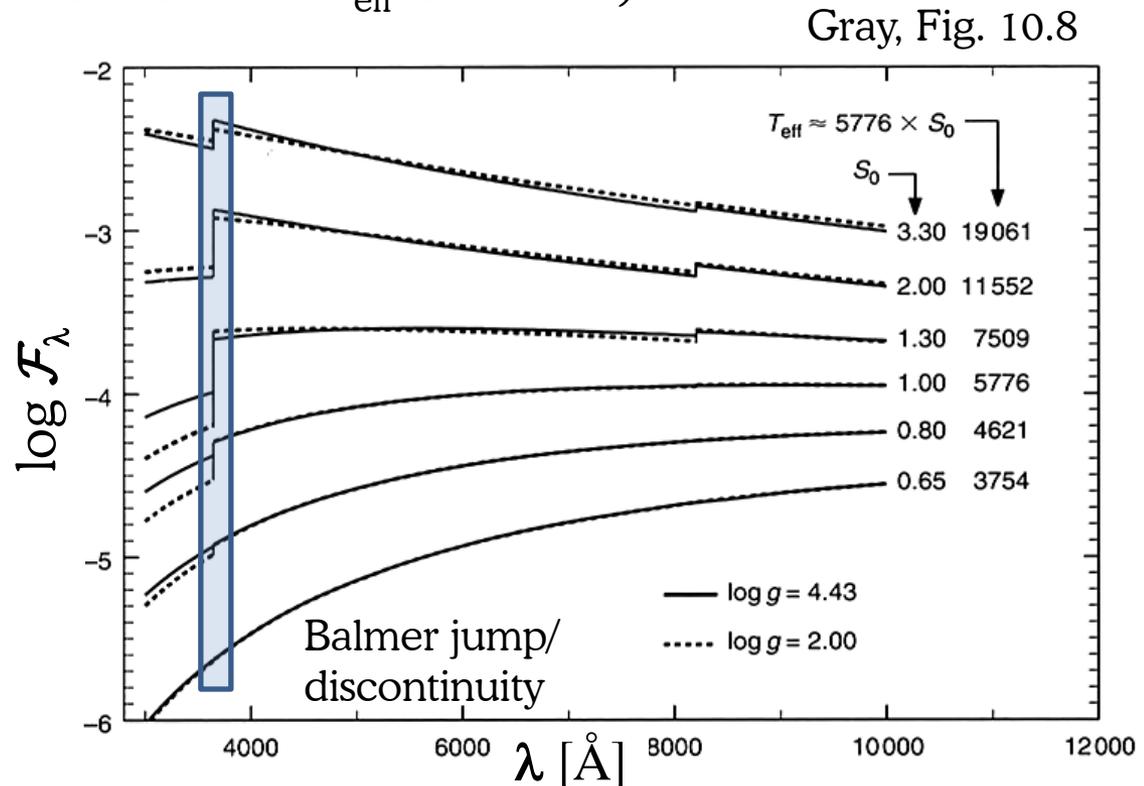
On the other hand, there are narrow-band indices which allow one to measure abundance variations (e.g. via molecular bands).



# Photometry: gravity dependence

The only feature that has a sufficiently (?) large gravity sensitivity to be exploited by photometry is the **Balmer jump at 3647 Å** (in hot stars it can be used as a sensitive  $T_{\text{eff}}$  indicator).

Colours like  $(U - B)$  or  $(u - y)$  measure the Balmer discontinuity, but the usefulness as a precise gravity indicator is hampered by the high line density in this spectral region (missing opacity problem), the difficulties with ground-based observations in the near-UV and a proper treatment of the overlapping Balmer lines.



The  $c_1$  index ( $\equiv (u - b) - (b - y)$ ) works well for metal-poor giants (Önehag *et al.* 2008).

# IRFM: a semi-fundamental $T_{\text{eff}}$ scale

Basic idea of the infrared-flux method:

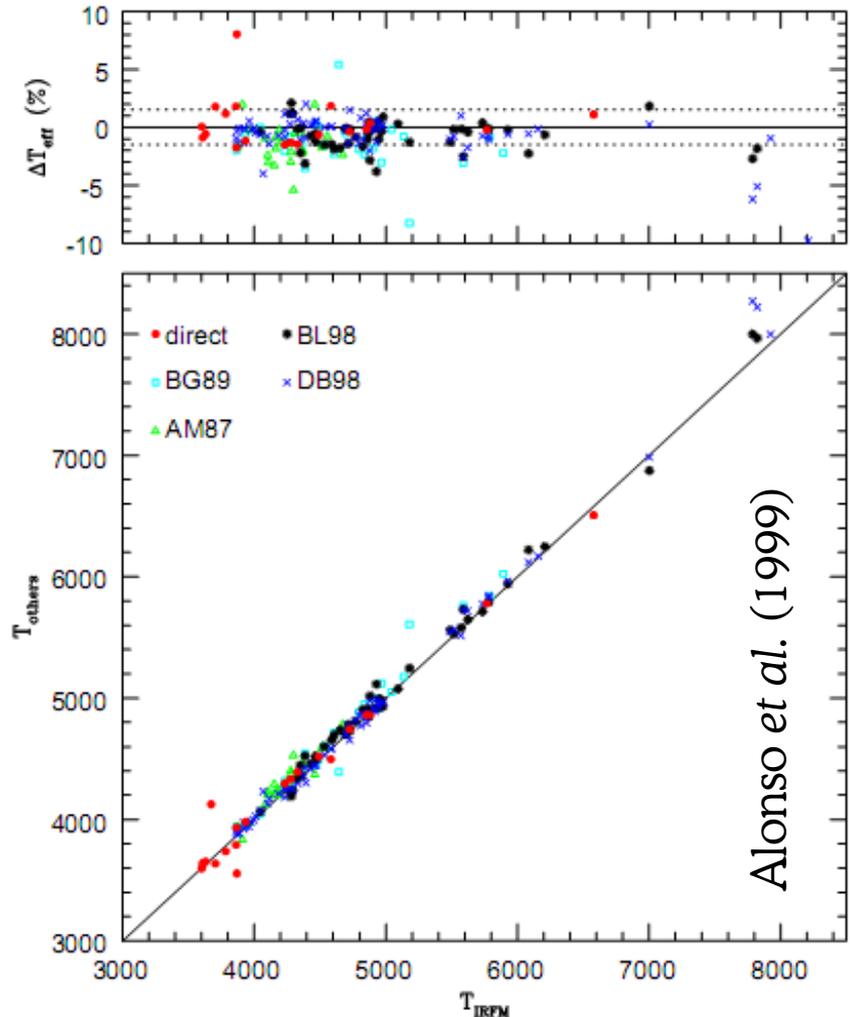
$$\frac{\mathcal{F}_{\text{surface}}}{\mathcal{F}_{\lambda_{\text{IR}}}(\text{Earth})} = \frac{\sigma T_{\text{eff}}^4}{\mathcal{F}_{\lambda_{\text{IR}}}(\text{model})}$$

$\mathcal{F}_{\lambda_{\text{IR}}}(\text{model})$  is said to be only weakly model dependent (but cf. Grupp 2004).

Once calibrated on stars with known diameters, any colour index can be calibrated on the IRFM.

**Direct** sample:  $\Delta T_{\text{eff}} = 0.06 \pm 1.25\%$

Comparing different IRFM calibrations (Blackwell *et al.*, Ramírez & Meléndez, Casagrande *et al.*), the **zero point** proves to be **uncertain by  $\pm 100$  K**, in particular for metal-poor stars.



$$\mathcal{F}_{\lambda_{\text{IR}}}(\text{Earth}) = q(\lambda_{\text{IR}}) \mathcal{F}_{\lambda_{\text{IR}}}^{\text{std}}(\text{Earth}) 10^{-0.4(m_{\text{IR}} - m_{\text{IR}}^{\text{std}})}$$

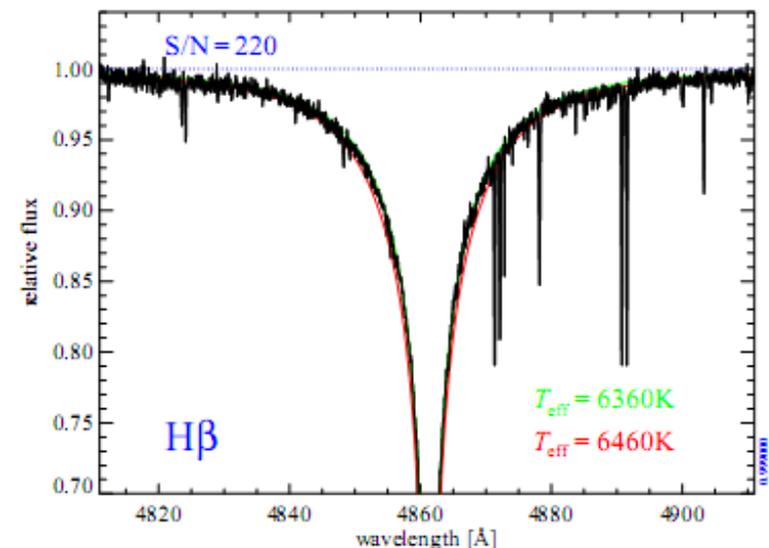
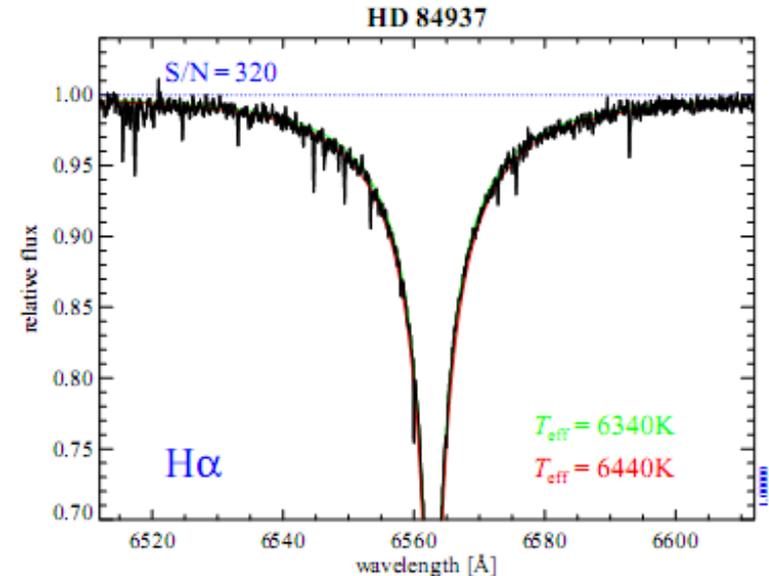
# Spectroscopic $T_{\text{eff}}$ indicators: H lines

Above 5000 K, the **wings of Balmer lines** are a sensitive  $T_{\text{eff}}$  indicator, **broadened by H + H collisions** (mainly  $H\alpha$ ) **and the linear Stark effect** ( $H + e^-$ ).

In cool stars, the  $\log g$  sensitivity is low (line and continuous opacity both depend on  $P_{\rho}$ ), as is the metallicity dependence. There is some dependence on the mixing-length parameter ( $H\beta$  and higher).

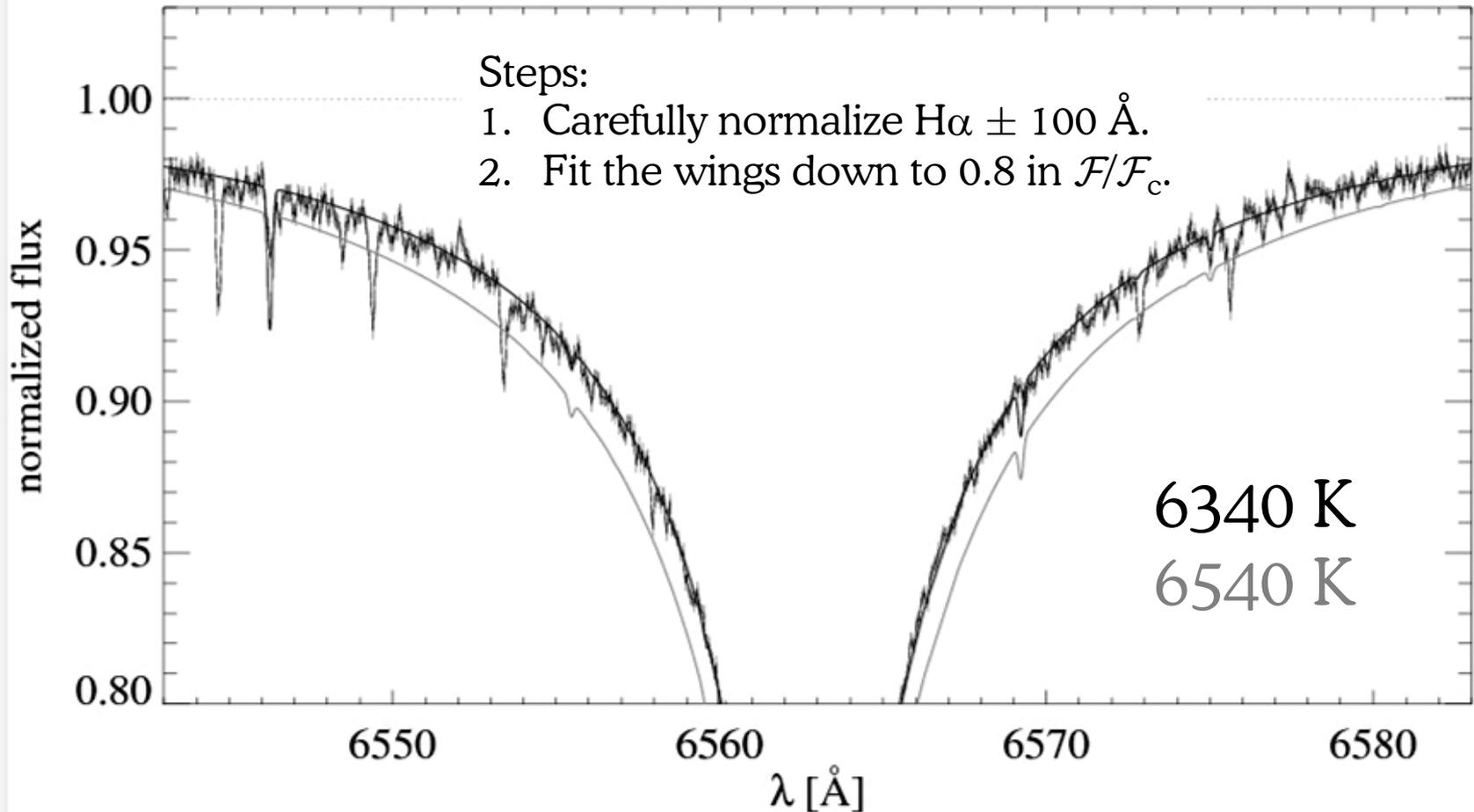
Main **challenge** (apart from the surprisingly complex broadening): **recovering the intrinsic line profiles** from (echelle) observations.

In hot stars, Balmer lines can constrain the surface gravity.



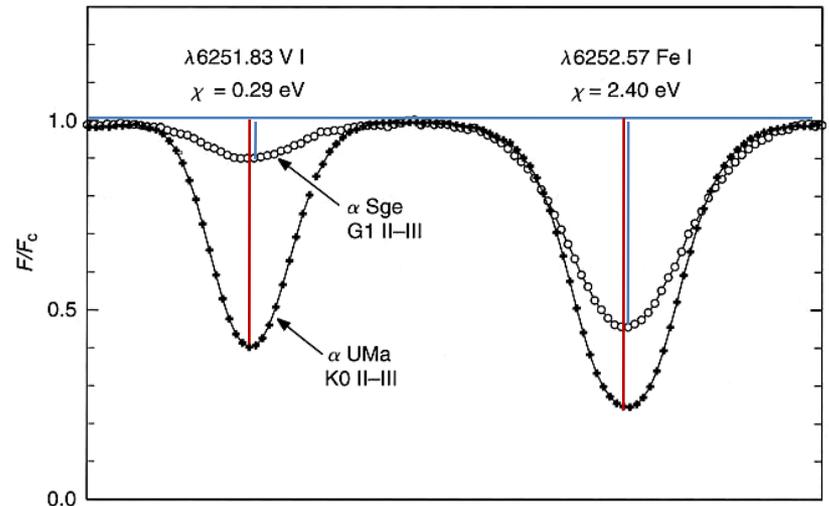
# H $\alpha$ as a function of $T_{\text{eff}}$

## HD84937 FOCES



# Line-depth ratios (LDRs)

Using the **ratio of two lines'** central depths (rather than  $W_\lambda$ ) can be a **remarkably sensitive temperature indicator** (precision as high as 5 K!), if the lines are chosen to have different sensitivities to  $T$ . Ideally, the LDR is close to 1 and the lines should not be too far apart.



Gray Fig.14.7

The main **challenge** lies in a **proper  $T_{\text{eff}}$  calibration** across a usefully large part of the HRD.

# Gravity sensitivity of ionized lines

Recall that ionized lines of an element that is mainly ionized have a  $P_e^{-1}$  sensitivity via the continuous opacity of  $H^-$ .

Integrating the hydrostatic equation, we find

$$P_g \propto g^{2/3}$$

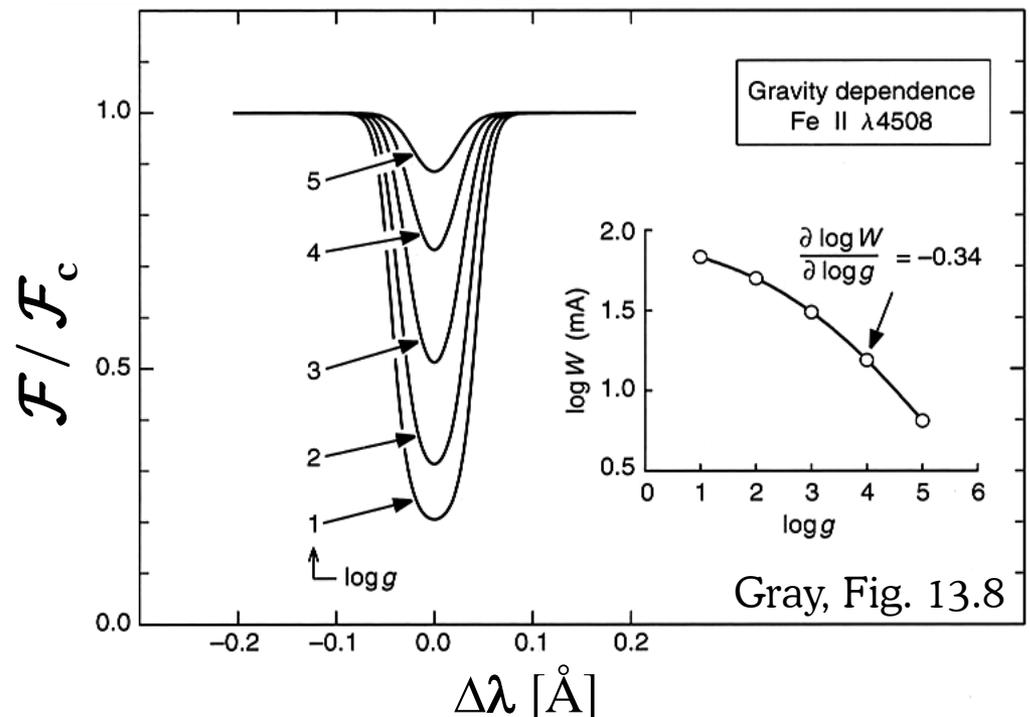
and together with  $P_e \propto \text{sqrt}(P_g)$  we expect

$$I_\nu / \kappa_\nu \propto g^{-1/3}.$$

This is borne out by actual calculations.

## Hydrostatic equilibrium

$$dP/d\tau_\nu = g / \kappa_\nu$$



# Practicalities of ionization equilibria

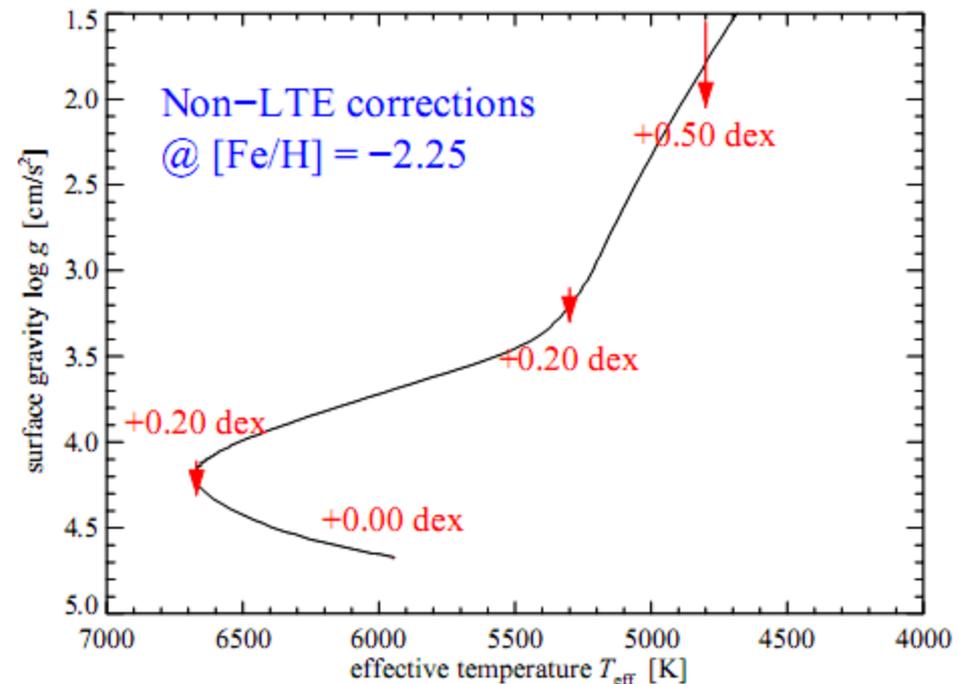
A change of **0.1 dex in  $\log \epsilon$**   
translates to a change of **0.3 dex  
in  $\log g$** .

Consequences:

A line-to-line scatter of 0.1 dex  
means that  $\log g$  is known to  
within 0.3 dex.

Relatively small changes in  $\log \epsilon$ ,  
e.g. because of a change in  $T_{\text{eff}}$   
or NLTE effects, can lead to  
factor-of-two changes in the  
surface gravity.

**Astrometry can help** to establish  
the correct surface-gravity scale.



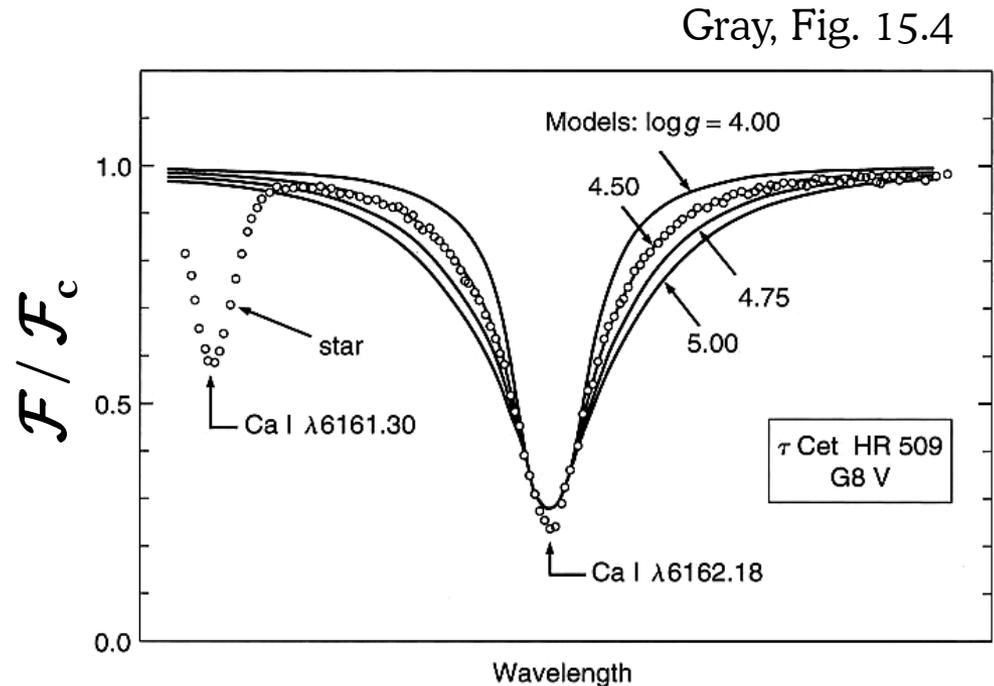
Korn (2004),  
Carnegie Observatories Centenary (2003)  
<http://www.ociw.edu/ociw/symposia/series/symposium4/proceedings.html>

# The strong line method

Damped (neutral) lines show a strong gravity sensitivity, because

$$I_{\nu} \propto \gamma_6 \propto P_g \propto g^{2/3}.$$

Like with ionization equilibria,  $\log \varepsilon$  needs to be known. This is to be obtained from weak lines of the same ionization stage, preferably originating from the same lower state (no differential NLTE effects).



Examples: Ca I 6162 (see above), Fe I 4383, Mg I 5183, Ca I 4226. Below  $[\text{Fe}/\text{H}] \approx -2$ , there are no optical lines strong enough to serve as a surface-gravity indicator.

# Spectroscopy of the Solar neighbourhood

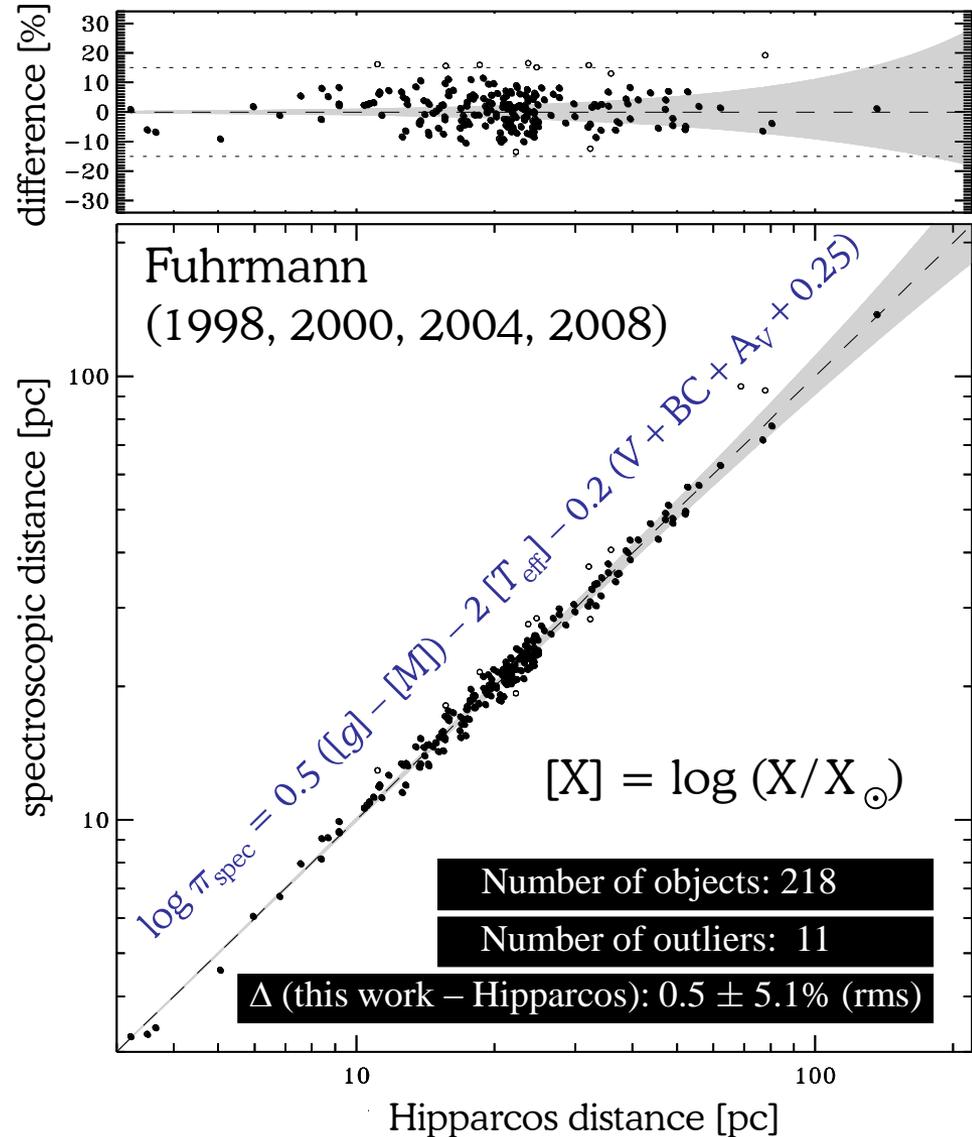
## Aim:

Derive precise stellar parameters and chemical abundances of FGK stars within  $d = 25$  pc.

## Example:

The strong-line method as a surface-gravity indicator for not too metal-poor, not-too-evolved stars coupled with  $T_{\text{eff}}$  values from Balmer lines.

## Benchmark: Hipparcos

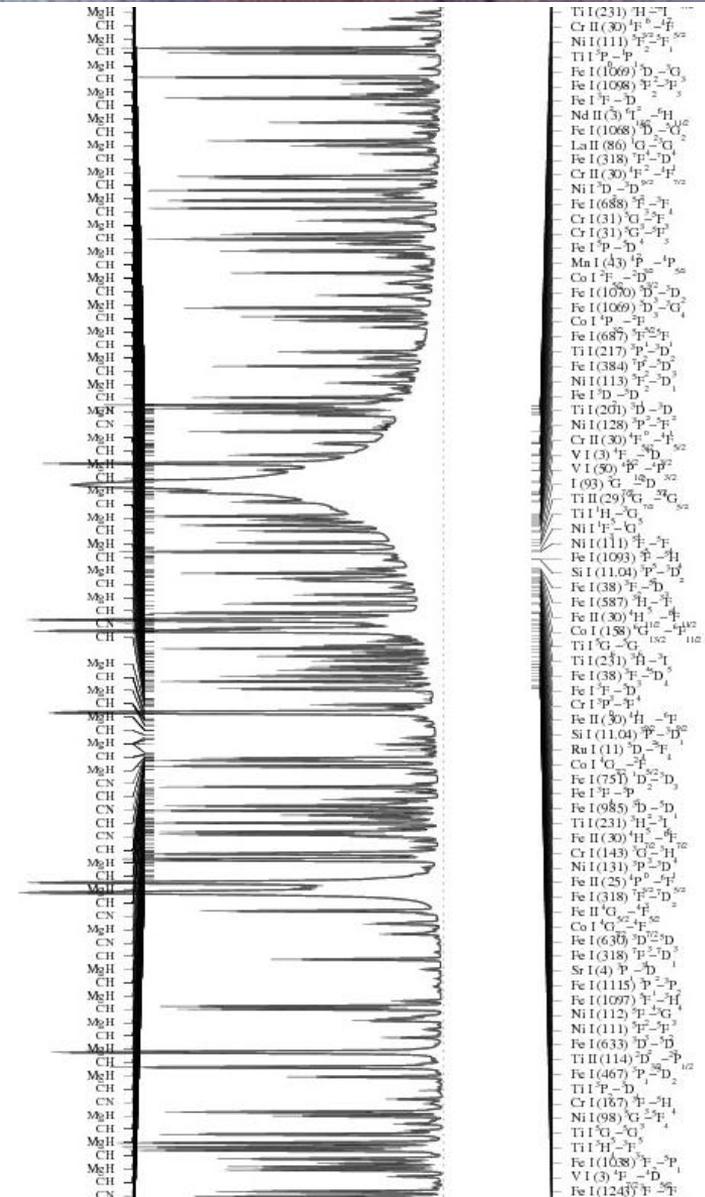


# Abundances from H to U

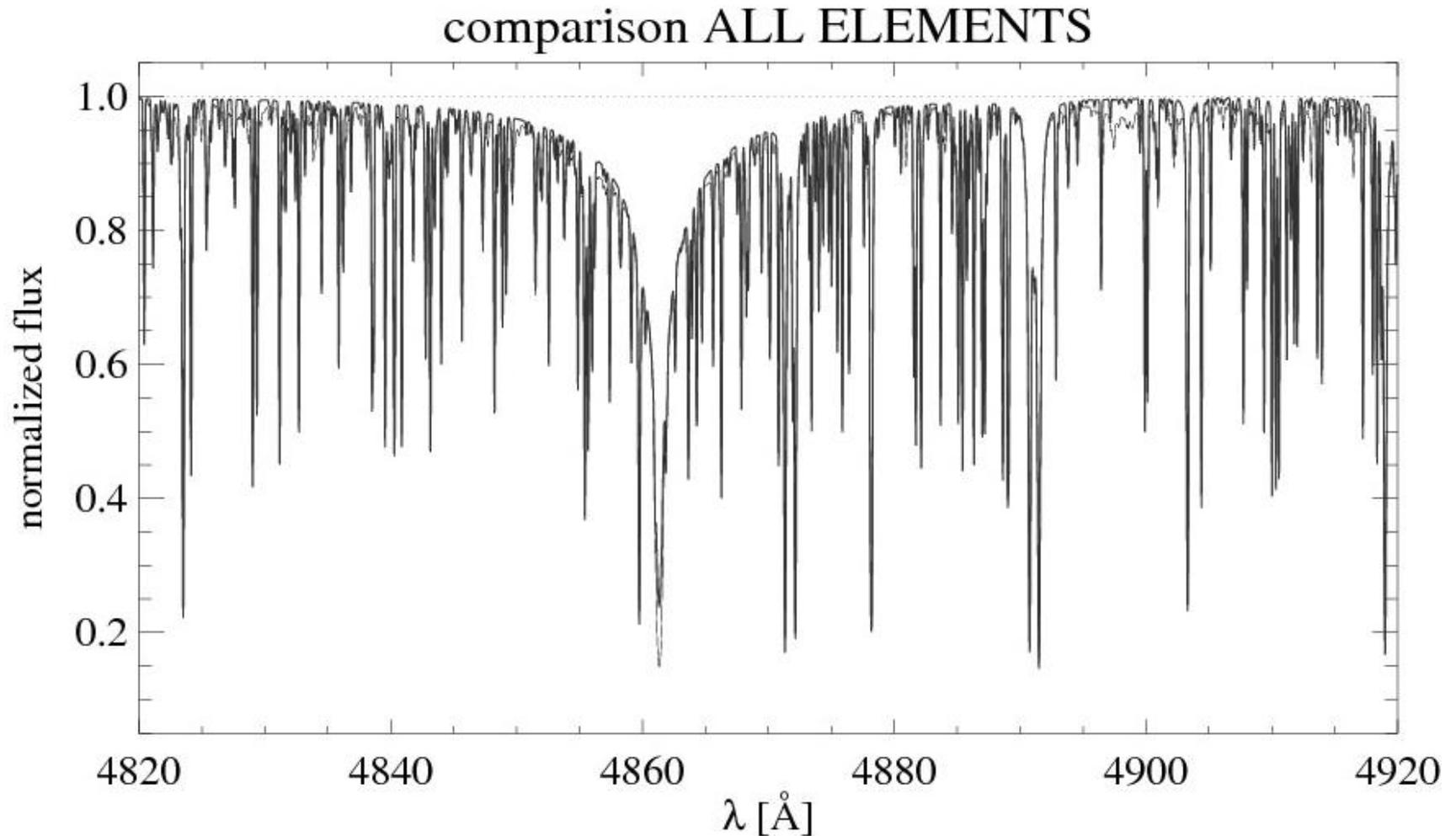
Once you have good stellar parameters, it is relatively easy to determine chemical abundances for your favourite element(s).

## Caveats

- ❑ some elements are not visible, e.g. noble gases in cool stars
- ❑ lines may lack or have inaccurate atomic data
- ❑ lines can be blended leading to overestimated abundances
- ❑ lines can be subject to effect you are unaware of, e.g. 3D and NLTE effects, hfs, isotopic and Zeeman splitting
- ❑ ...



# Quantitative spectroscopy: the Sun



# Spectroscopy: pros *vs.* cons

Spectroscopy is

- ✓ a way of determining a great number of stellar parameters,
- ✓ the key technique for obtaining detailed chemical abundances,
- ✓ (usually) reddening-free.

However, hi-res spectroscopy is

- ❑ comparatively costly at the telescope,
- ❑ currently limited to 18<sup>m</sup> in *V*,
- ❑ more difficult to master than photometry.



...especially when they accept photometry as a source of valuable information.

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