

Effective Temperature Determination

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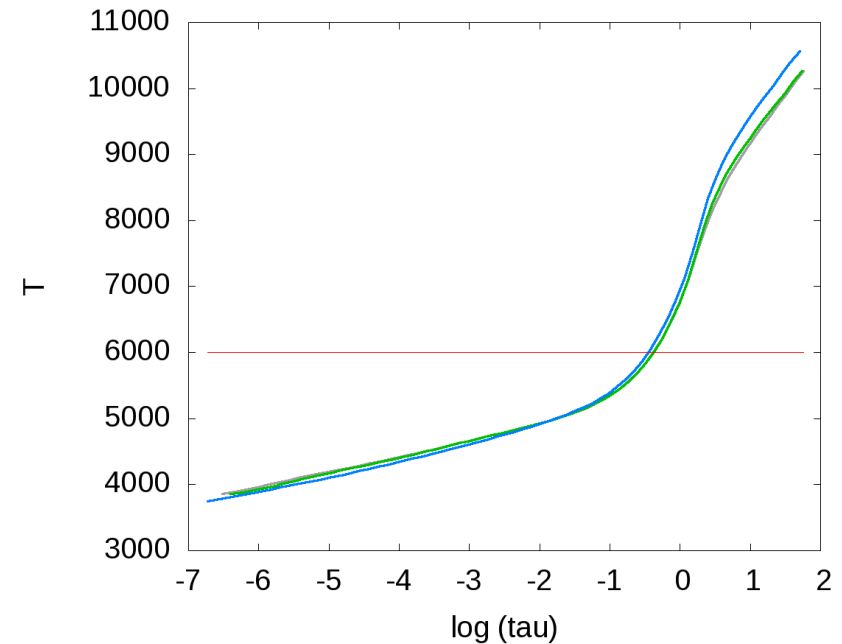
Effective Temperature

$$\sigma T_{eff}^4 \equiv \int_0^\infty F_\nu d\nu = F_\star = \frac{L}{4\pi R^2}$$

- It is the temperature of a black body that gives the same total power per unit area.
- Physically related to F_\star total radiant power per unit area at stellar surface.
- **Directly** given by stellar luminosity and radius.

Depth Dependence

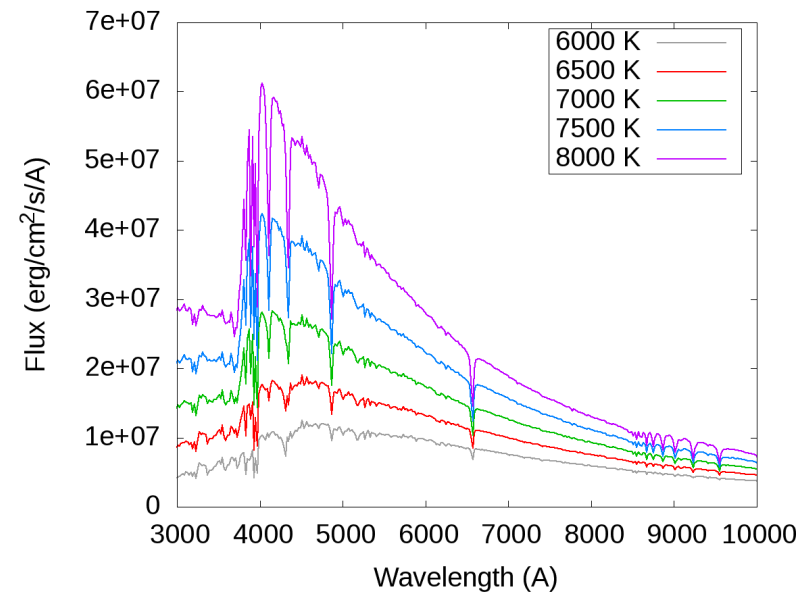
- Temperature of line forming region is lower than T_{eff}
- Spectral lines are formed at different depths and temperatures.



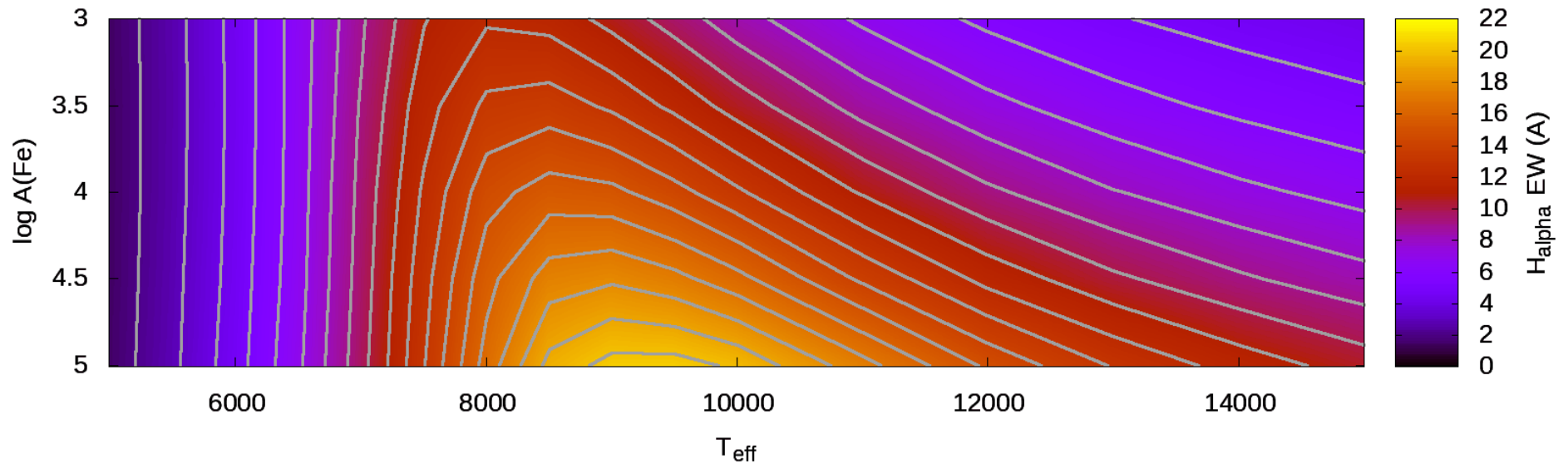
$T_{\text{eff}} 6000, \log g 4.5$

Paschen Continuum

- Using stellar fluxes to determine T_{eff}
- Requires accurate moderate-resolution flux-calibrated spectra or spectrophotometry.
- Basis for T_{eff} – colour calibrations
 - More in later lecture



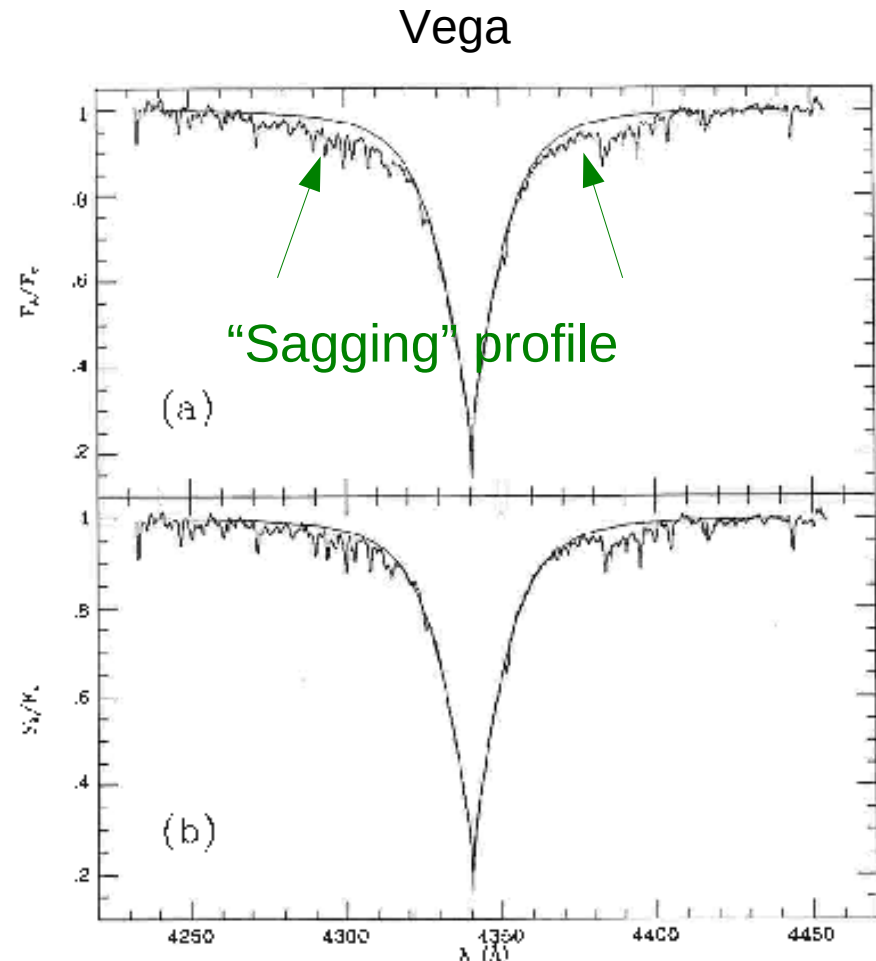
Balmer Profiles



- The Balmer lines provide good T_{eff} diagnostic below around 8000K due to low sensitivity to surface gravity.
- For hotter stars sensitivity to both T_{eff} and $\log g$.

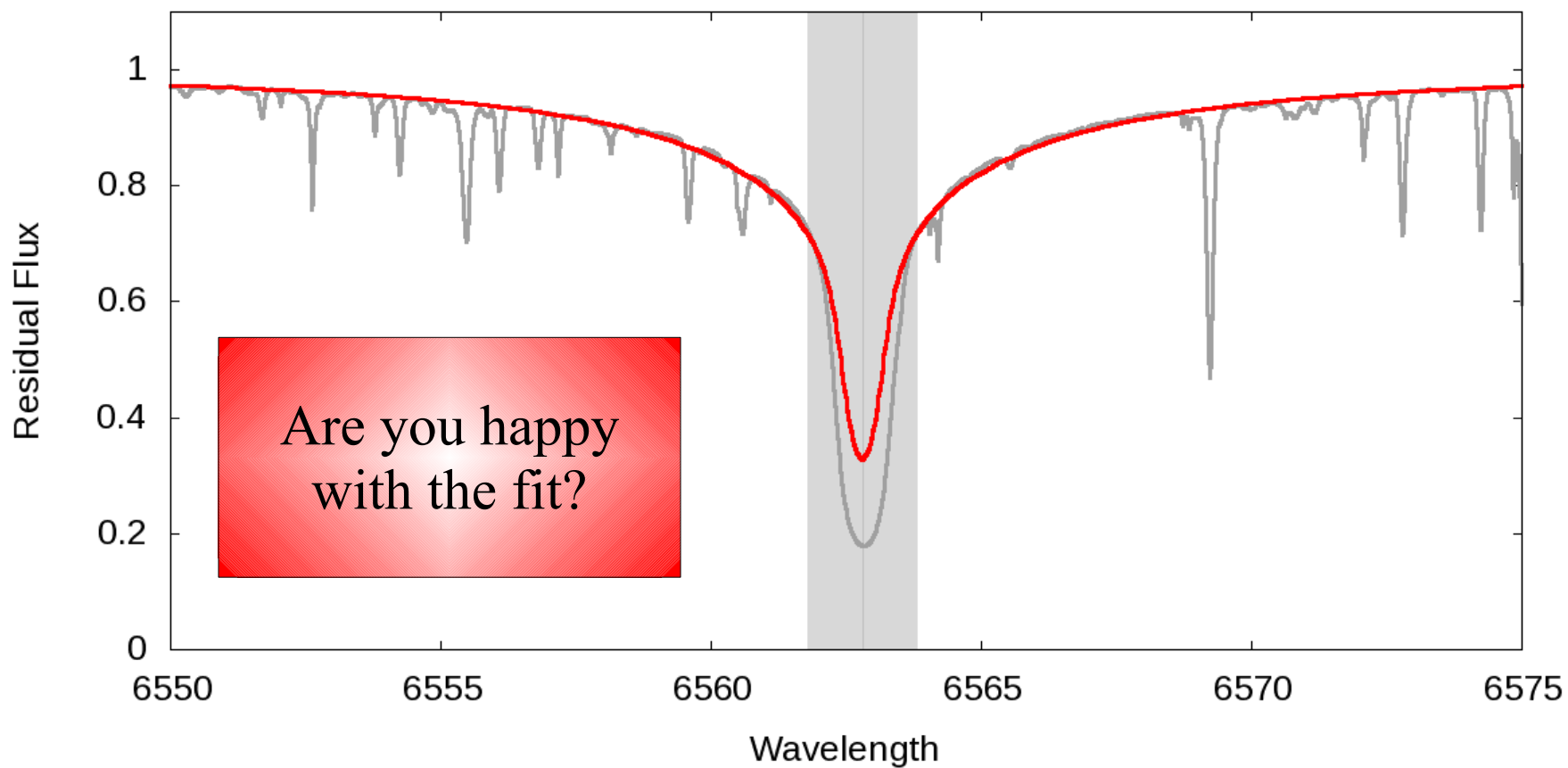
Balmer Line Normalization

- Broad lines require careful continuum determination
 - Must preserve true profile shape
- Echelle spectra usually give poor profiles due to curved orders
- Medium-resolution single order spectra preferred



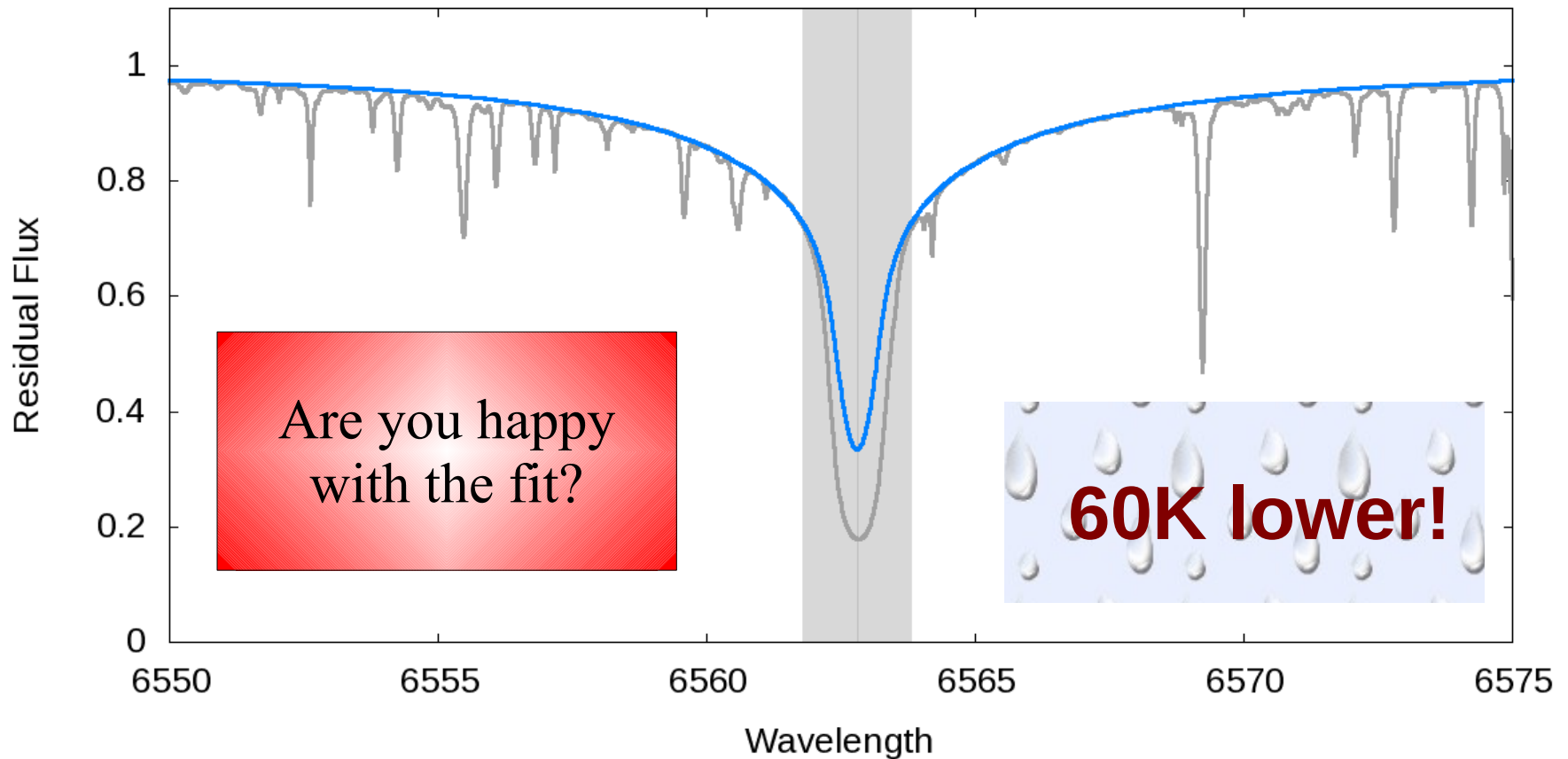
Smalley 1992 PhD Thesis

Solar H α



$T_{\text{eff}} 5777 \log g 4.44$

Solar H α

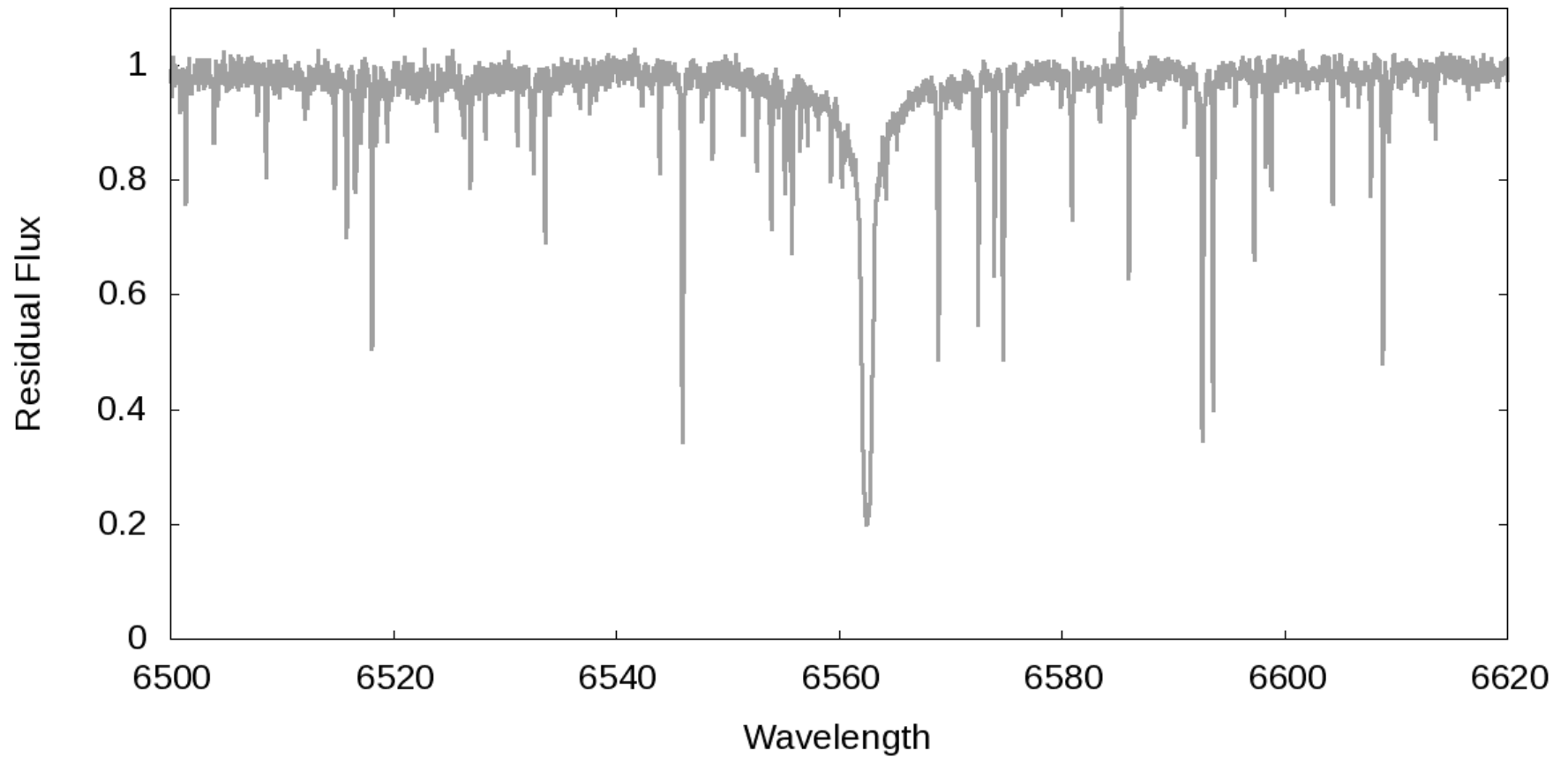


3-d, Convection,
Line broadening

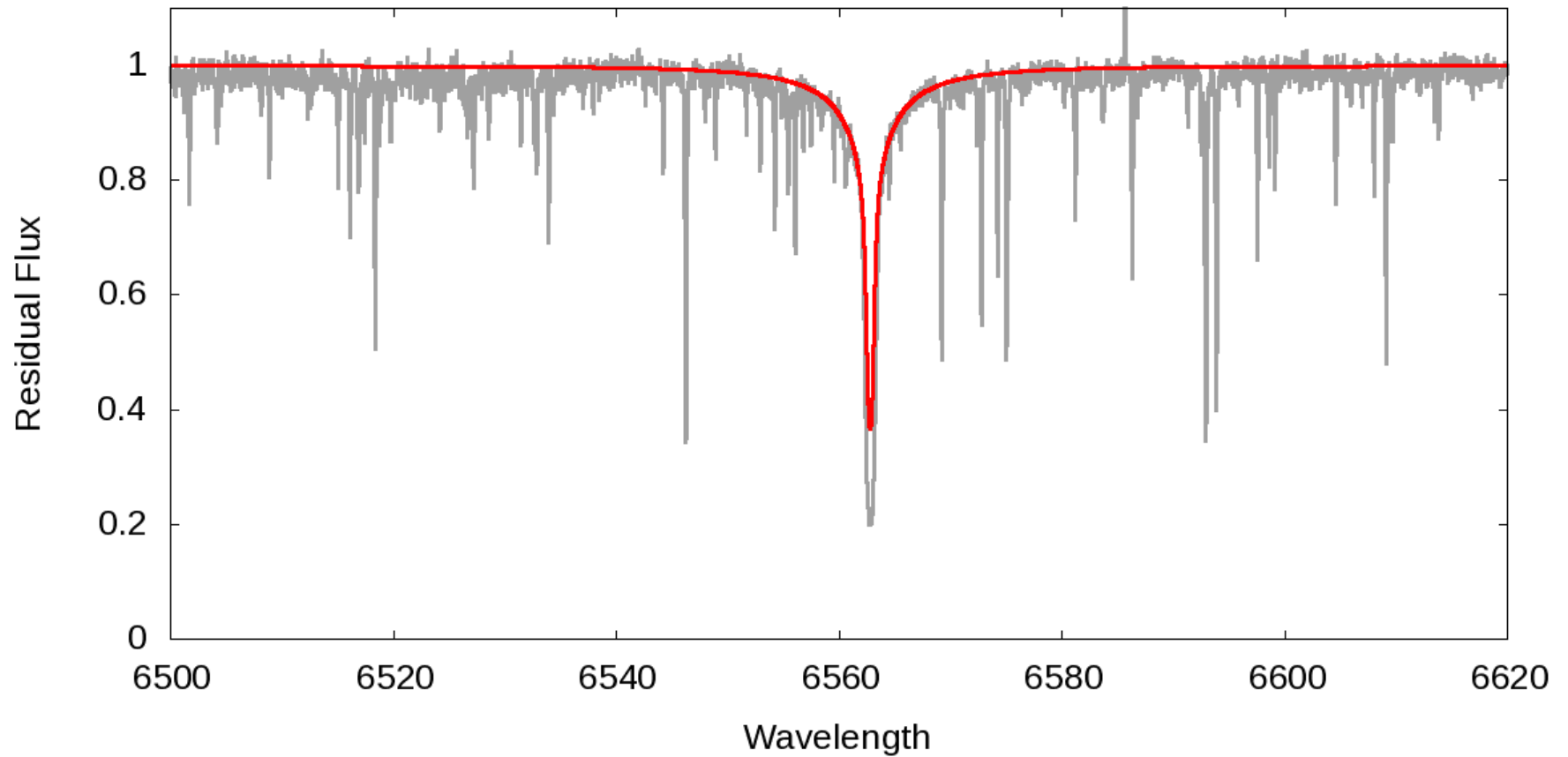
$T_{\text{eff}} 5720 \log g 4.44$

See for example Cayrel et al., 2011, A&A, 531, 83

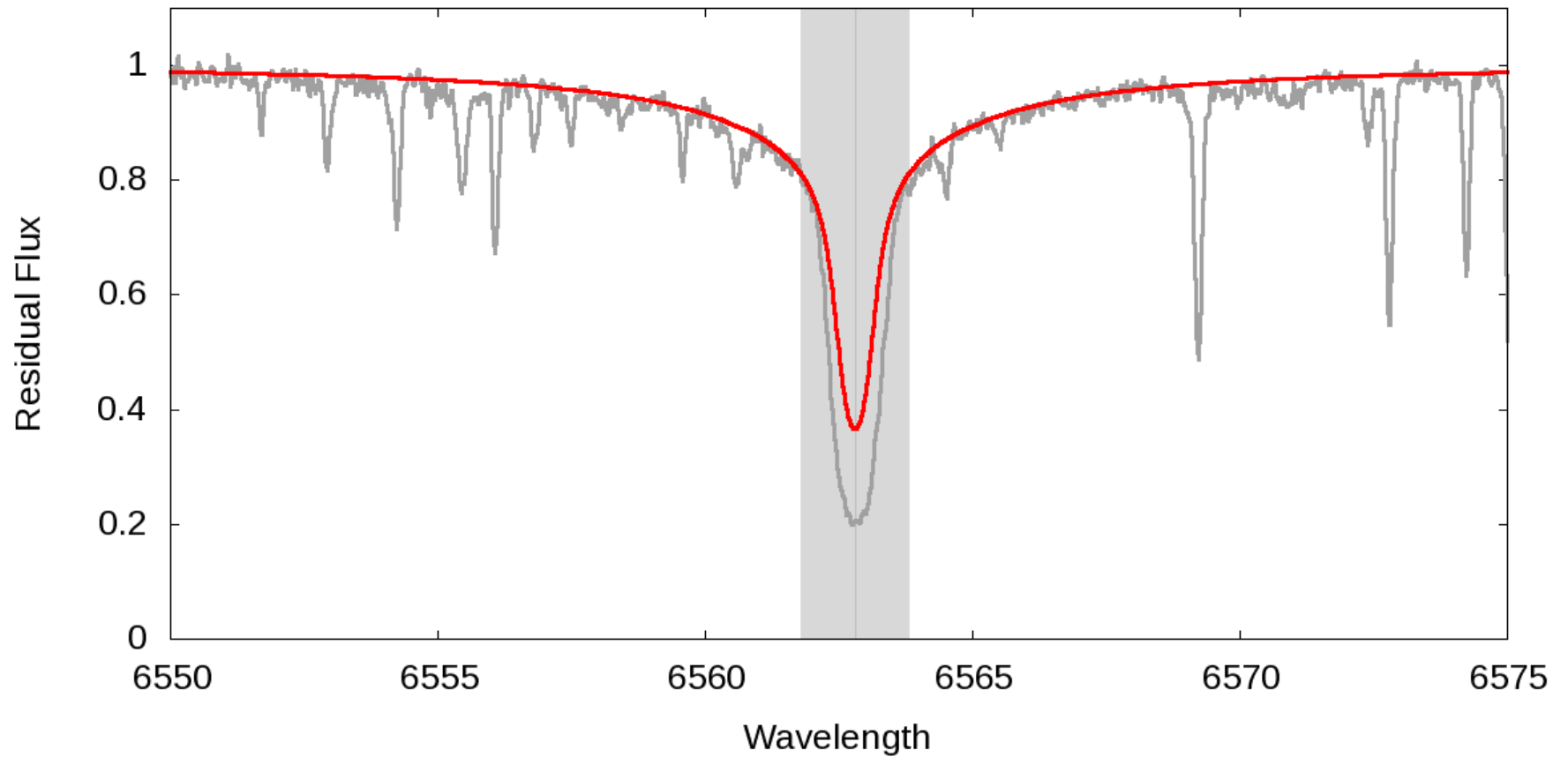
KIC 11772920



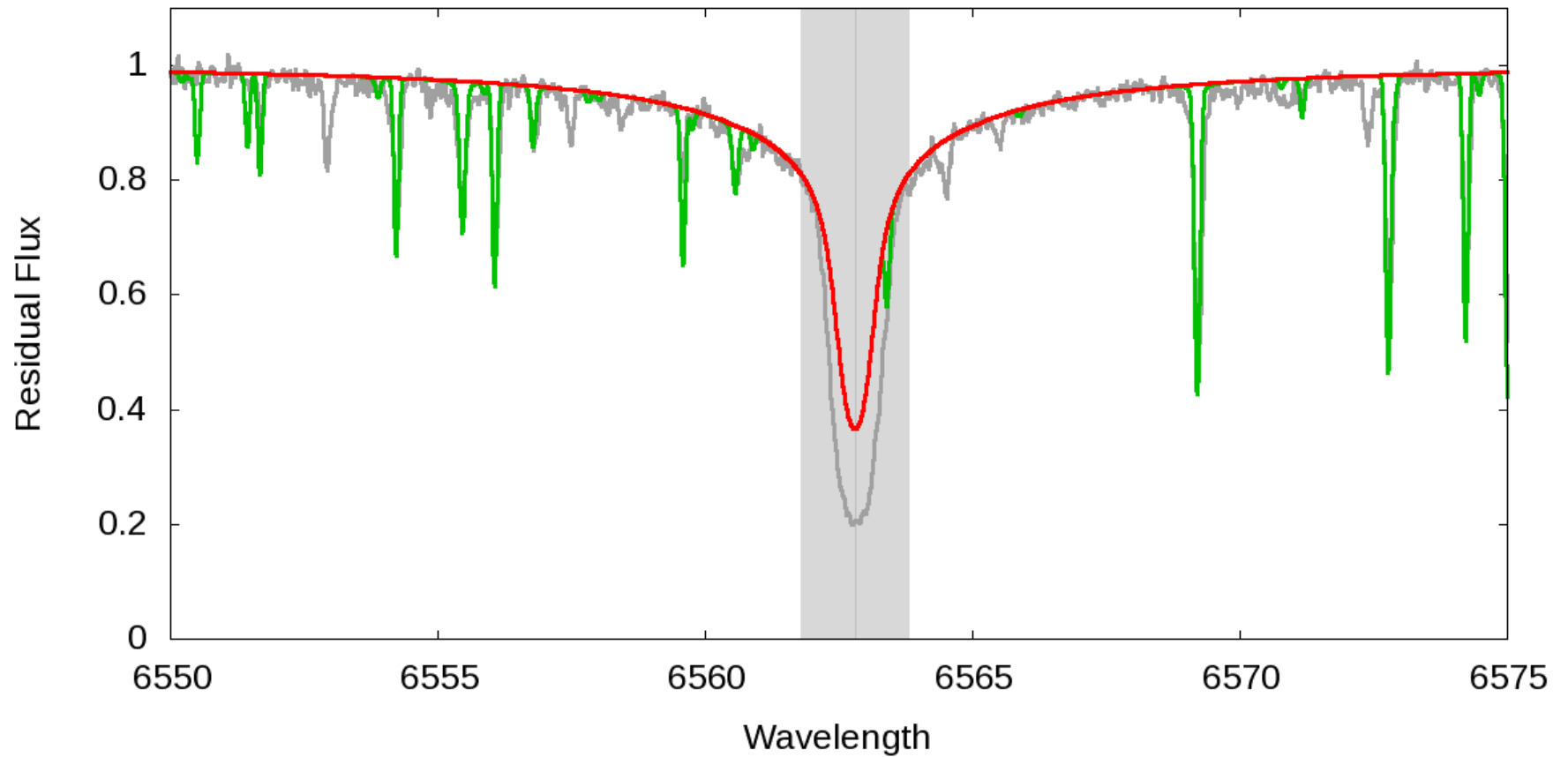
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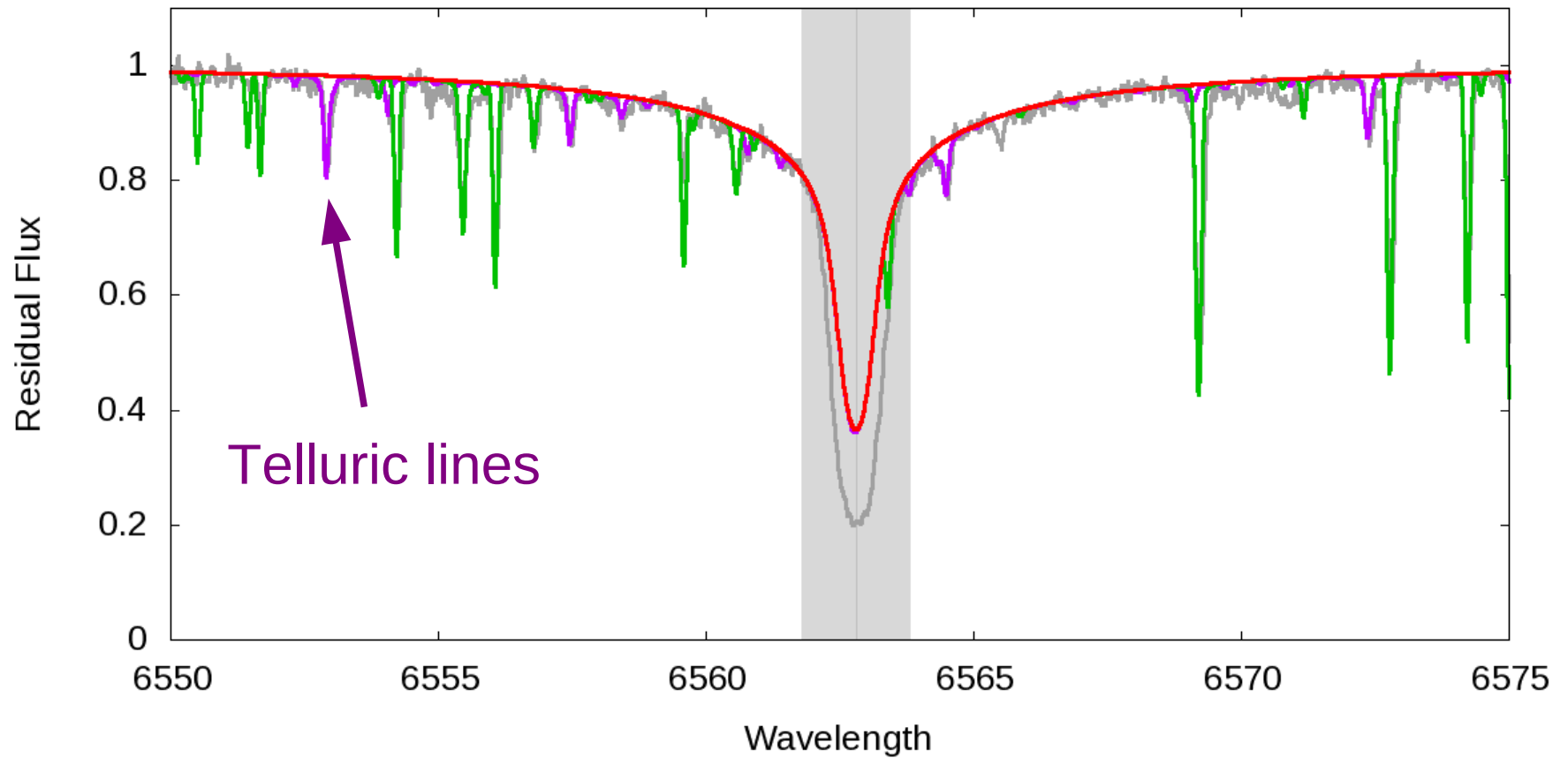
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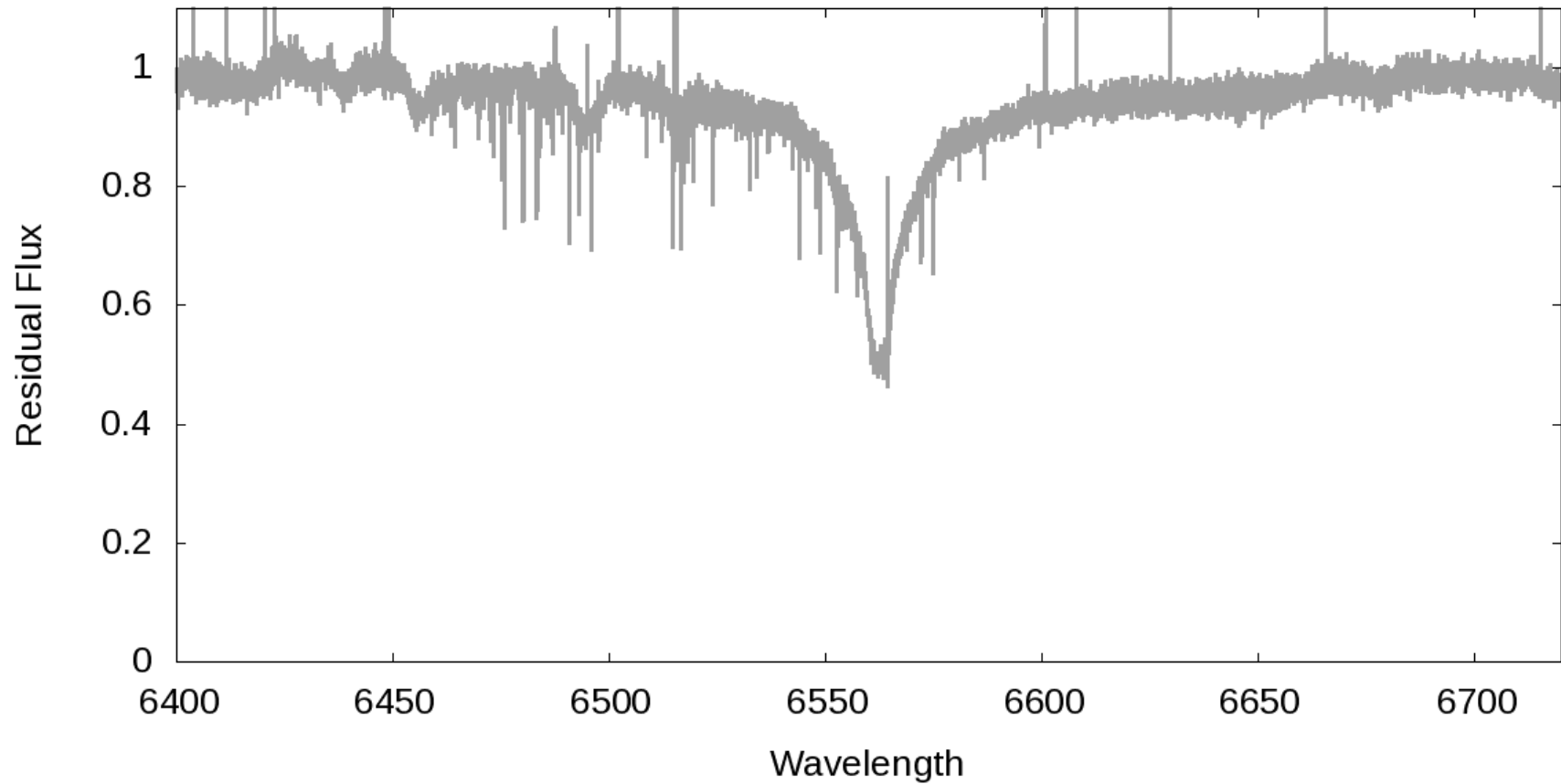
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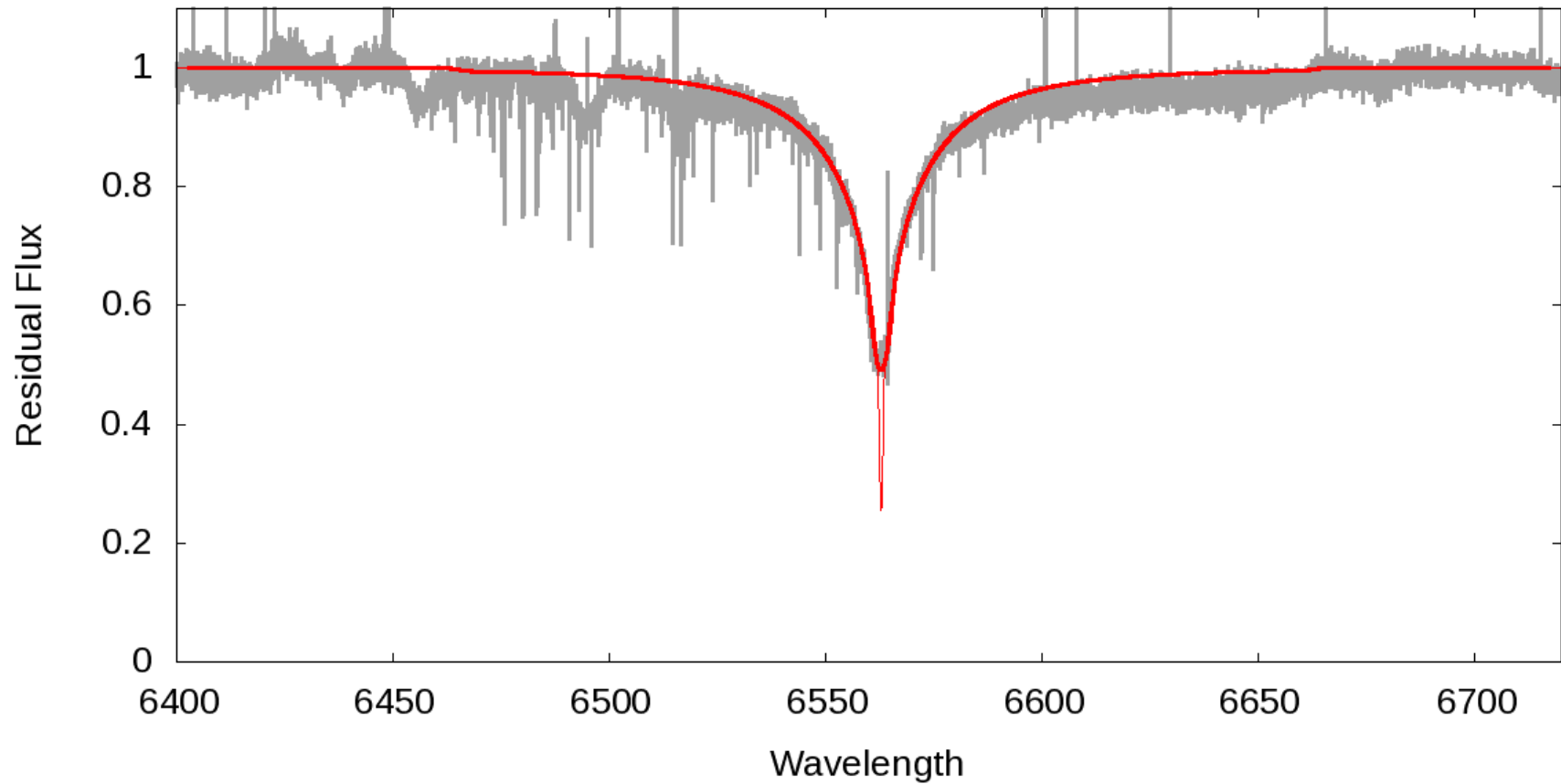
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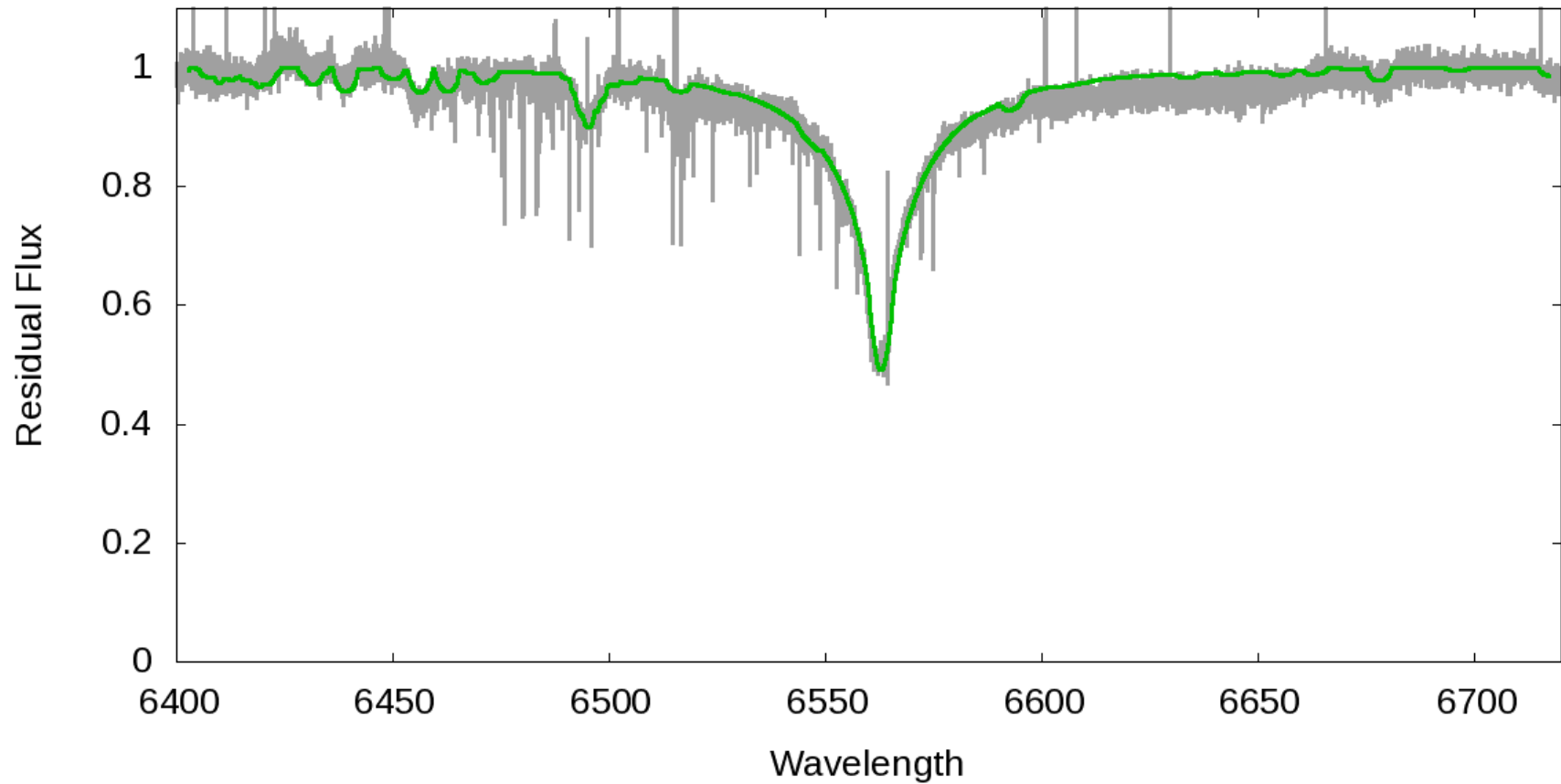
KIC 11090405



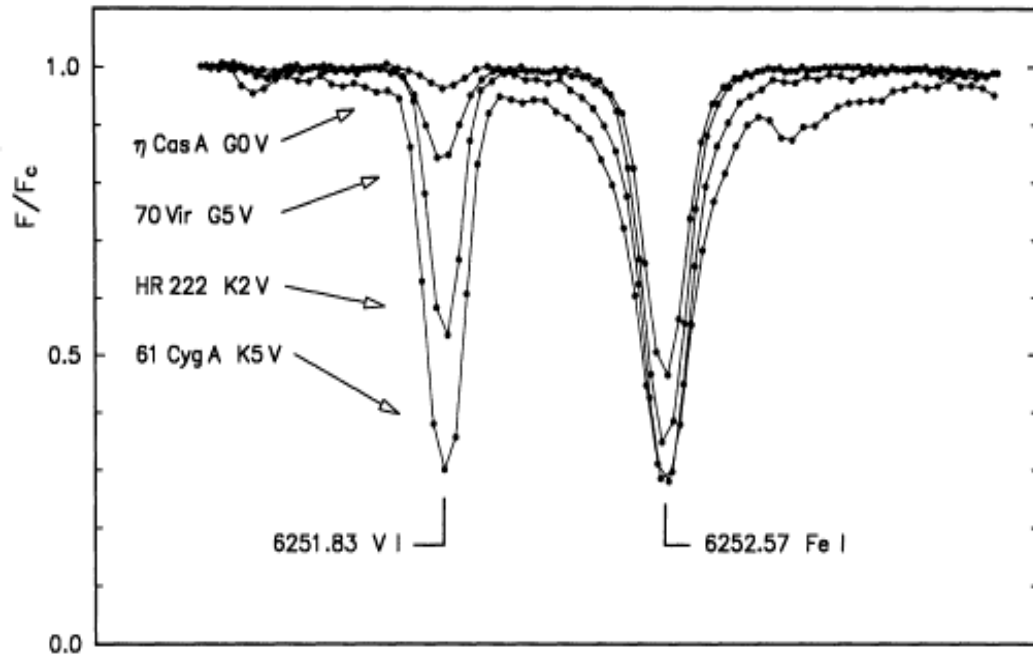
KIC 11090405



KIC 11090405



Spectral Line depth ratios



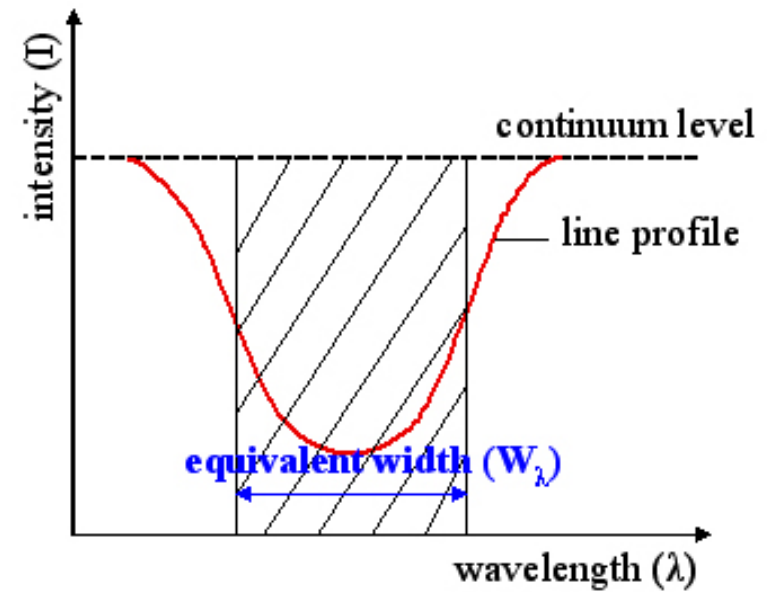
Gray & Johanson, 1991, PASP, 103, 439

Good for looking for T_{eff} variations in a given star

- Line depth ratios
 - Differing excitation potentials
 - Precise to $\pm 10\text{K}$
 - A measure of temperature in line forming regions
 - Model dependent
- Tied to T_{eff} scale by empirical calibrations.

Equivalent Width

- Measure of number of absorbers
 - Abundance
- BUT
 - No information on profile shape
 - Wings can affect measurement



$$W_{\lambda} = \int (1 - F_{\lambda}/F_0) d\lambda$$

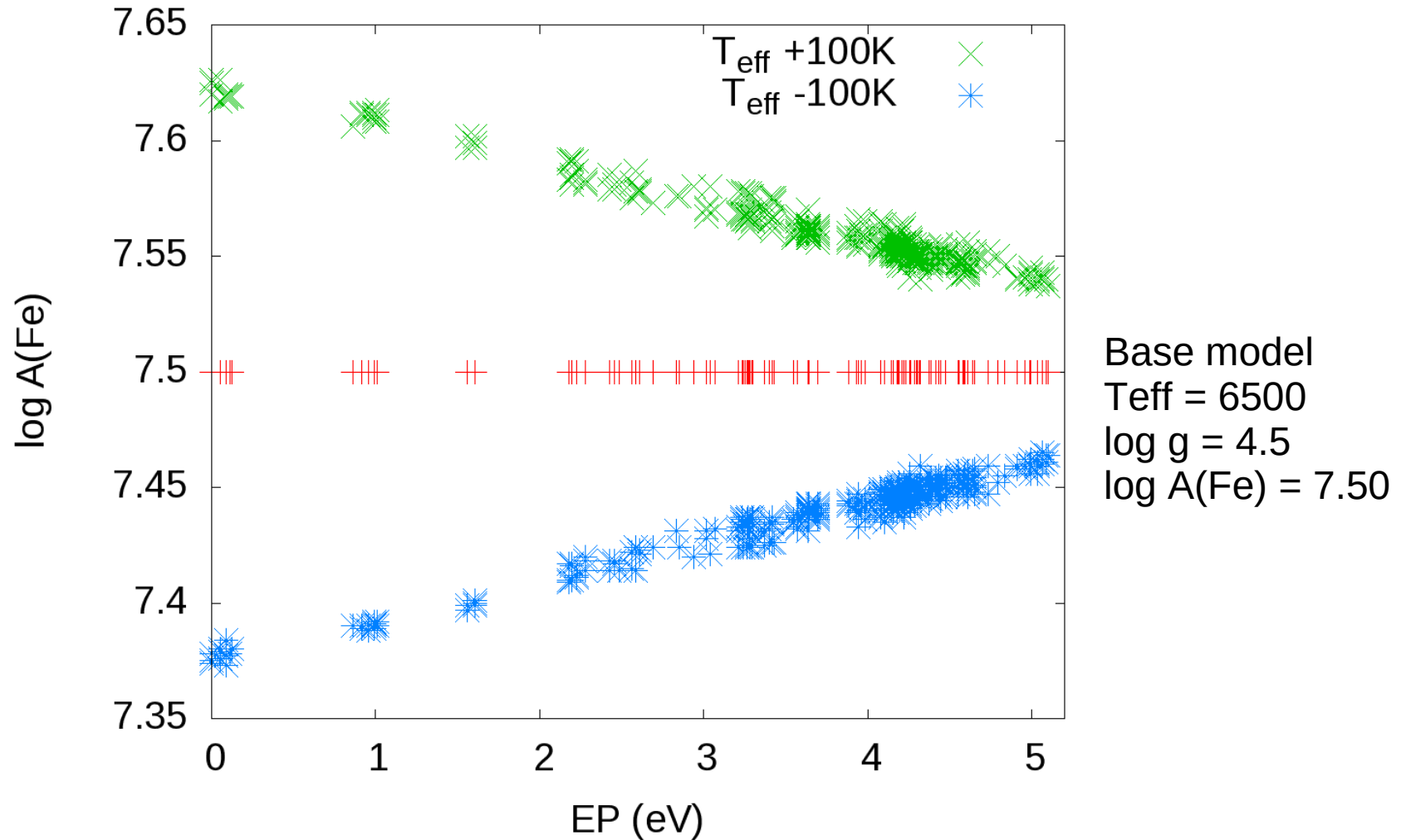
Metal Line Diagnostics

- Excitation Potential
 - Abundances from the same element should agree for all excitation potentials
- Ionization Balance
 - The abundances obtained from differing ionization stages of the same element must agree
 - Fe I/Fe II ratio can be used as a $T_{\text{eff}} - \log g$ diagnostic

Exploring Excitation Potential

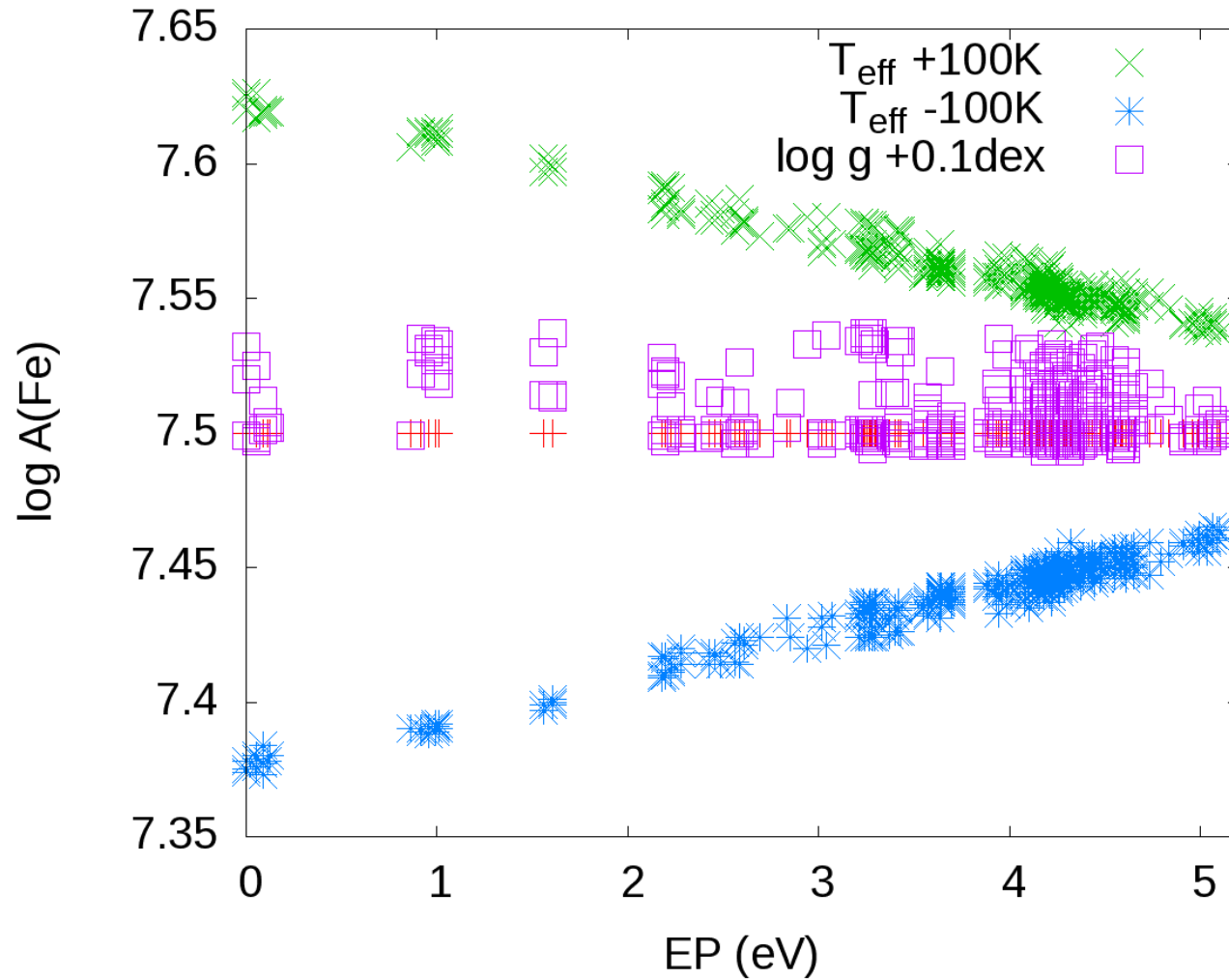
- Use lines that are not pressure sensitive
 - Solar-type stars, use neutral metals
 - In hotter stars, use ionised metals
 - Weak lines to avoid saturation effects
- Generated a simulated set of “observed” equivalent widths (W_0/λ).
 - Model: T_{eff} 6000 K, $\log g$ 4.5, $\log A(\text{Fe})$ 7.50
 - Take Fe I lines taken from Kurucz gfall linelist in wavelength range 5000 – 6000 Å
 - Select those with $5 < \text{EW} < 100 \text{ mÅ}$

Change Temperature



Note change in mean $\log A(\text{Fe})$ by ± 0.18 dex

Change Surface Gravity



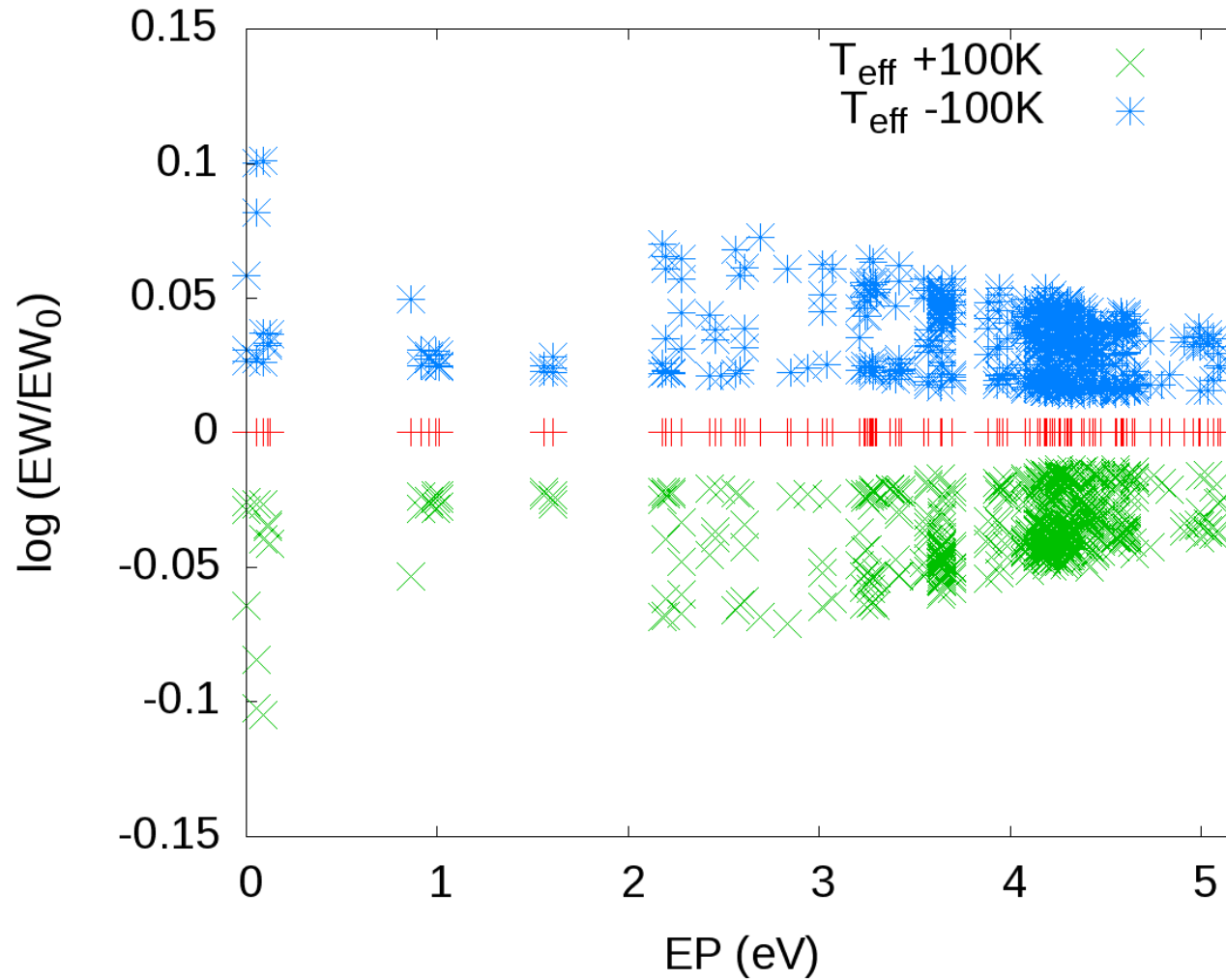
A lack of abundance correlation

- Analysis method searches for zero correlation between $\log A$ and EP:
 - Regression fit and find where gradient is zero
 - Could use where correlation coefficient is zero
- Error estimates
 - 1-sigma error on gradient or correlation coefficient
 - 1% error in W_λ gives $\sim 10\text{K}$ in T_{eff}
 - 5% , 50K; 10%, 100K
 - $\pm 0.1\text{dex}$ in $\log g$ gives $\pm 20\text{K}$ in T_{eff}

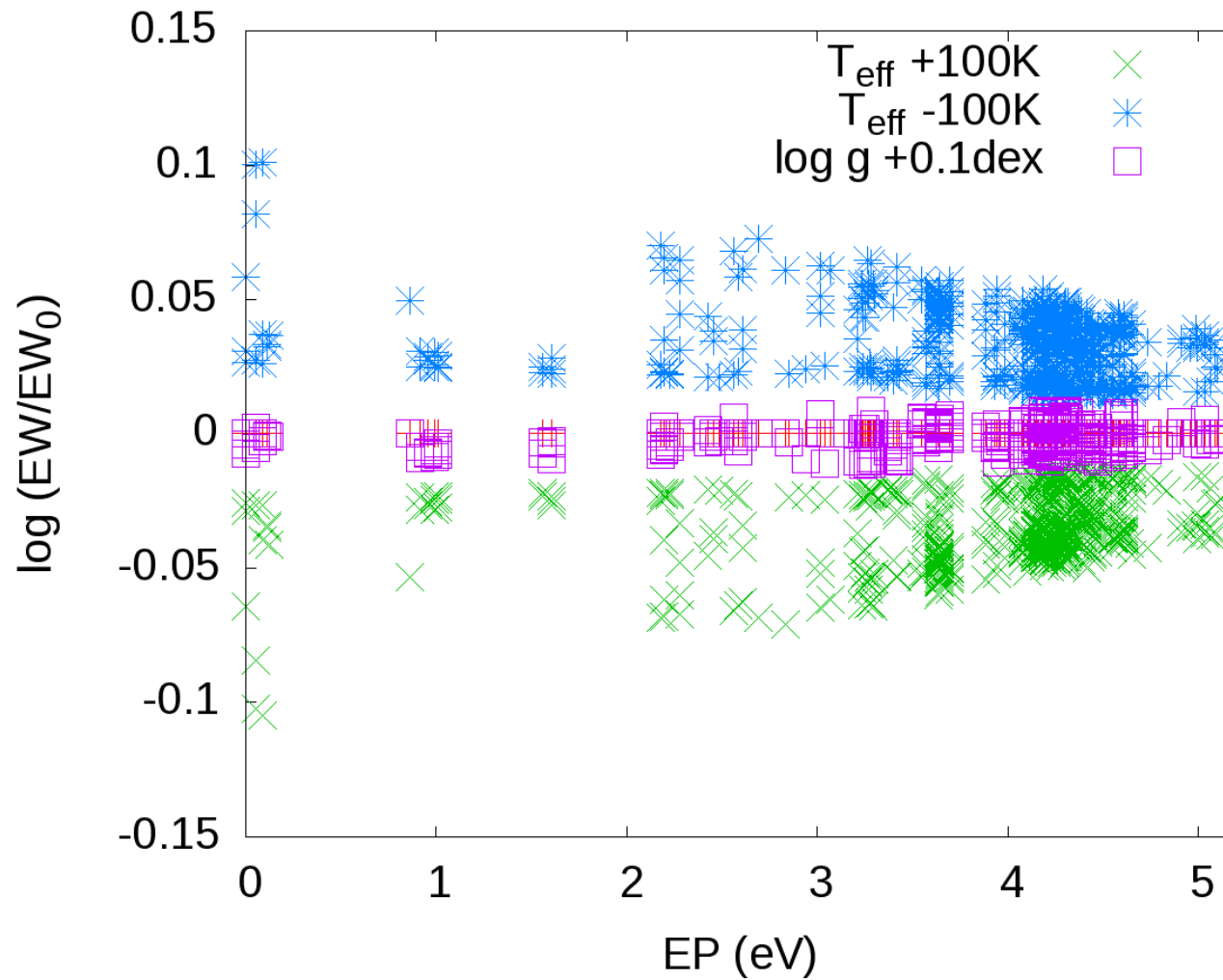
Equivalent width correlation

- From a “*philosophical*” viewpoint all Fe I lines in the atmosphere should have the same abundance.
- Since $\log\left(\frac{w}{\lambda}\right) \propto \log A$ we can change the procedure to using equivalent width differences.
- Let us use $\log\left(\frac{w}{\lambda}\right) - \log\left(\frac{w_0}{\lambda}\right) = \log\left(\frac{w}{w_0}\right)$
- Now plot this against EP
 - Assume an abundance: $\log A(\text{Fe}) = 7.50$

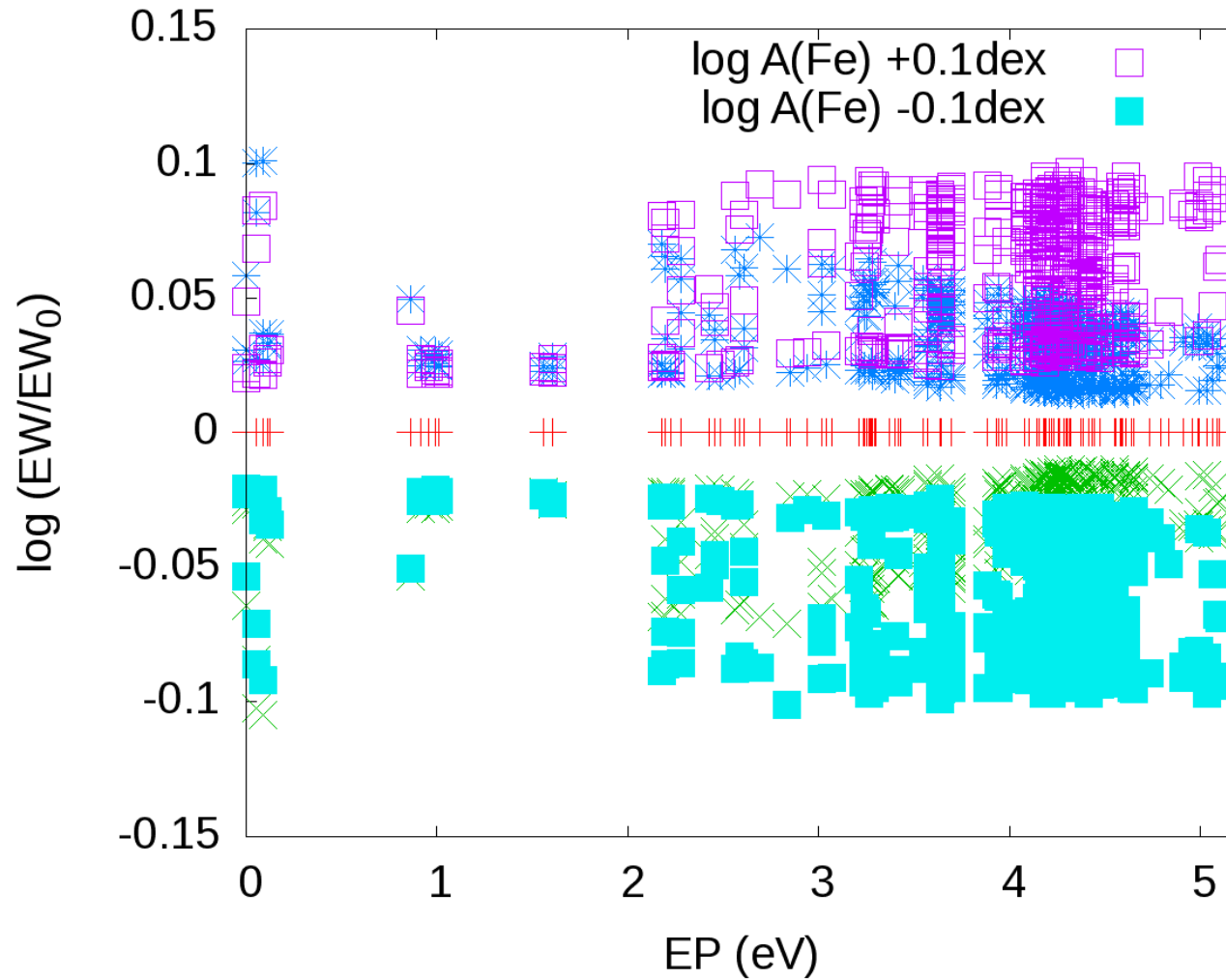
Change Temperature



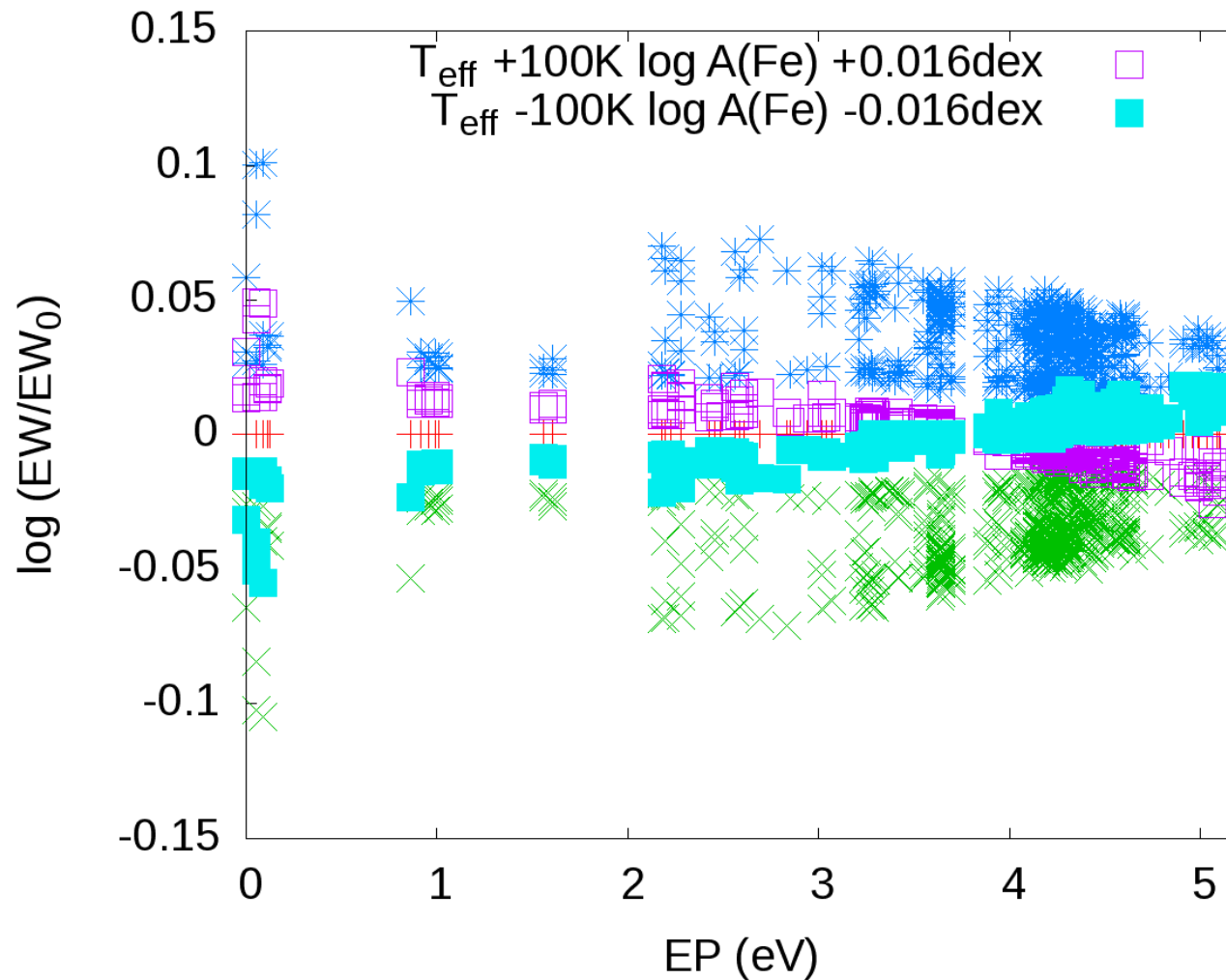
Change surface gravity



Change Abundance



Change T_{eff} and $\log A(\text{Fe})$



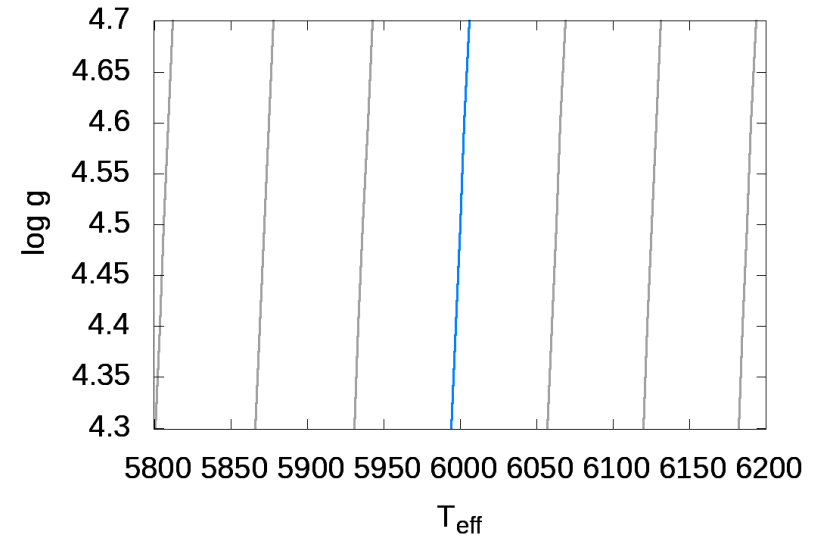
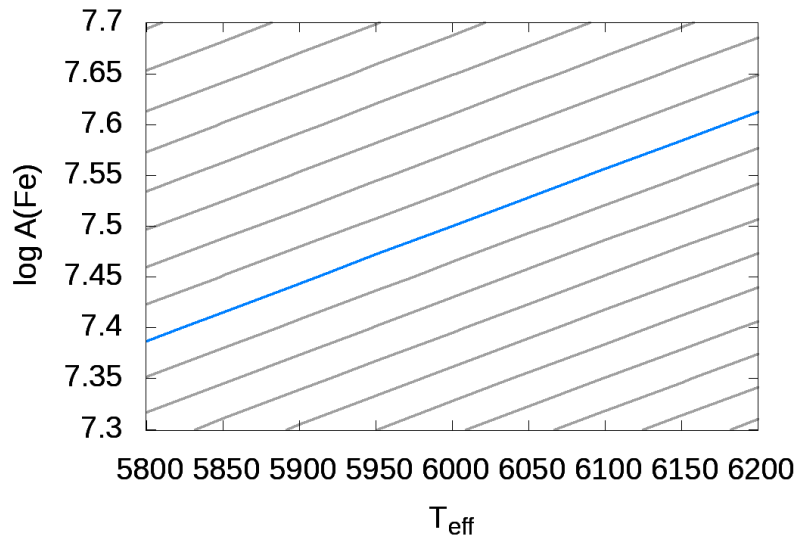
“Total” Equivalent Width

- Rather than using individual equivalent width differences, now use sum of the differences:

$$\sum \log\left(\frac{W}{W_0}\right)$$

- Rather than trend with EP, just have one number
 - Would appear to be a loss of information?

Total EW Trends

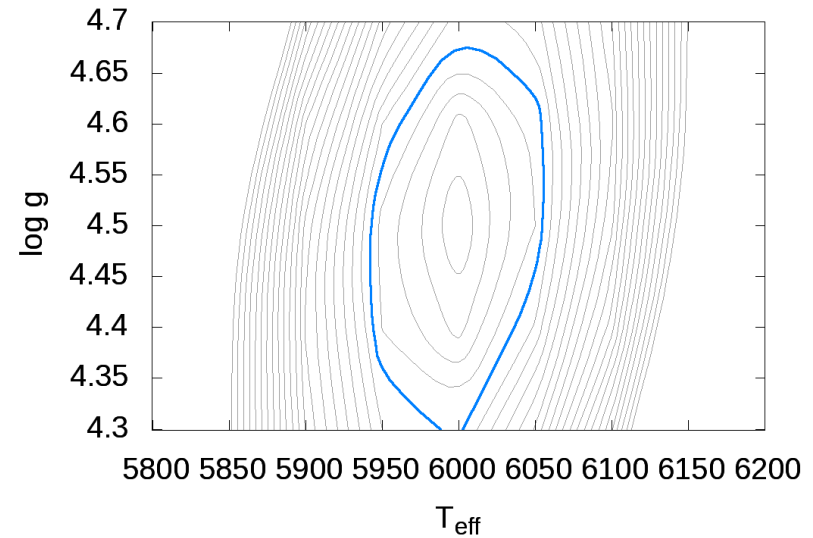
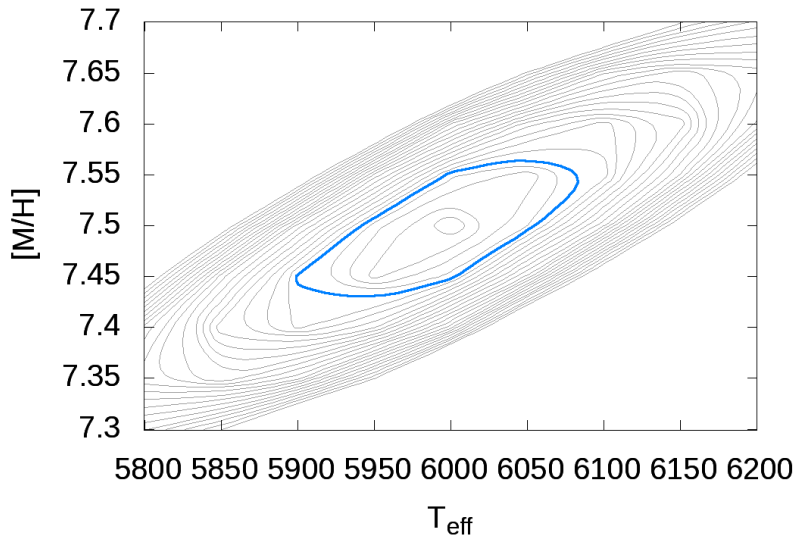


- Trend in $\log A(\text{Fe})$ with T_{eff}
- T_{eff} virtually insensitive to $\log g$

Using χ^2 instead

- Measuring equivalent widths might not always be practical.
 - Blending, high rotation, etc.
- Now consider T_{eff} determination using χ^2 fitting to spectrum
 - Generated with **all** lines $>5\text{m}\text{\AA}$ in range 5000-6000 \AA
 - Same Model: T_{eff} 6000 K, $\log g$ 4.5, $\log A(\text{Fe})$ 7.50
 - Assume S/N 100:1 for χ^2 calculation

x² Correlations



- Correlation between [M/H] and T_{eff}
- Weak sensitivity to log g

Conclusions on Simulation

- Using $\log A(\text{Fe})$ versus EP provides a good determination of T_{eff} visual diagnostic diagram
 - Low sensitivity to $\log g$
- Using EW has $\log A(\text{Fe})$ complication
 - “Total” EW has T_{eff} correlated with $\log A(\text{Fe})$
- Using χ^2 gives T_{eff} might be coupled with $\log A(\text{Fe})$, but not (significantly) with $\log g$.

At least in this simulated example!

Global Spectral Fitting

- Take a large grid of synthetic spectra with varying T_{eff} , $\log g$, $[M/H]$, etc.
- Locate the best-fitting synthesis
 - Hence obtain T_{eff} , $\log g$, $[M/H]$, etc.
- Issues to consider
 - How reliable are these parameters?
 - What are the hidden dangers?
 - What are realistic error estimates, over and above the internal precision?

Do we care about T_{eff} ?

- The effective temperature of a star is not important!
- It is the $T(\tau_0)$ relationship that determines the spectral characteristics.
 - The parameters obtained from spectroscopic methods alone **may not** be consistent with the true values
 - Not necessarily important for abundance analyses
 - Important when interested in fundamental properties

Summary

- Currently T_{eff} determinations are generally accurate to no more than 1~2% (50~100K)
- Possible to get an internal precision on measurements that are sensitive to temperature variations of $\pm 10\text{K}$
 - but accuracy on the true T_{eff} scale is *significantly* less.

Beware T_{eff} values without errorbars!