

# CO: EXOTIC & NON-STANDARD EXCITATION

John H. Black  
Department of Earth and Space Sciences  
Chalmers University of Technology  
Onsala Space Observatory  
Sweden

2012 March 1  
Lorentz Center, Leiden

- Preliminaries: overlooked terms in the rate equations
- UV pumping of CO
- Excitation upon formation
- Triplet states of CO & electron-impact
- A digression on  $\text{OH}^+$

The usual rate equations for steady-state populations are

$$\frac{dn_i}{dt} = \sum_{j>i} n_j A_{ji} - \sum_{k<i} n_i A_{ik} + \sum_{\ell \neq i} (n_\ell C_{\ell i} - n_i C_{i\ell}) = 0$$

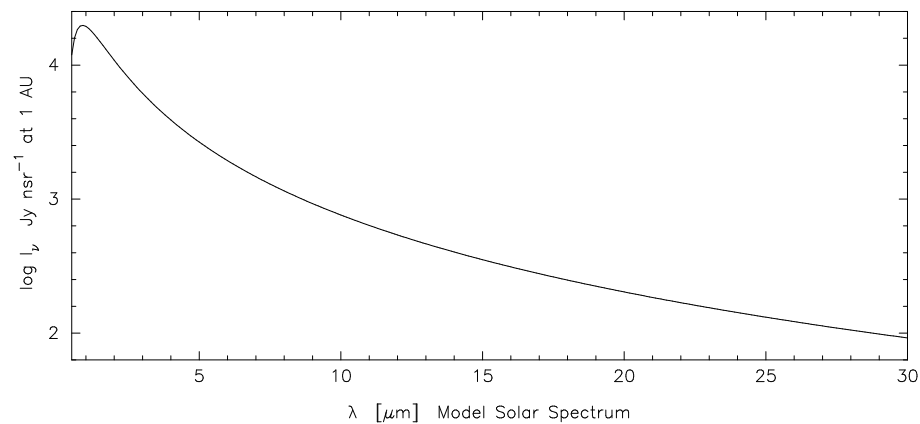
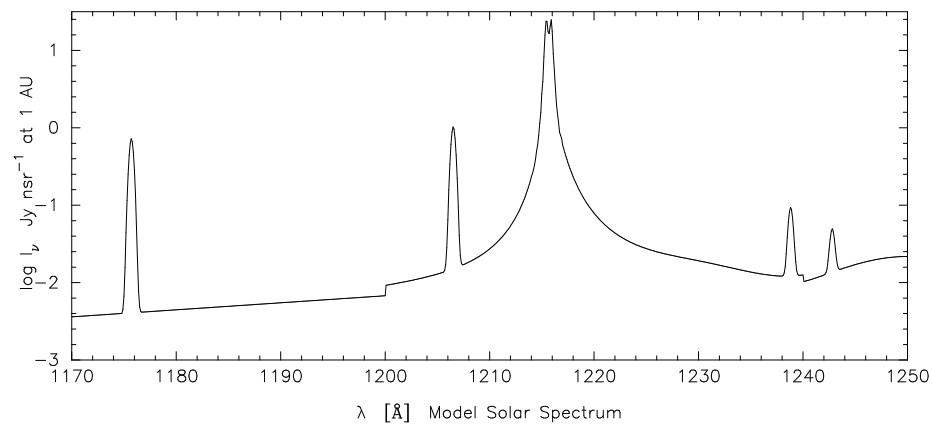
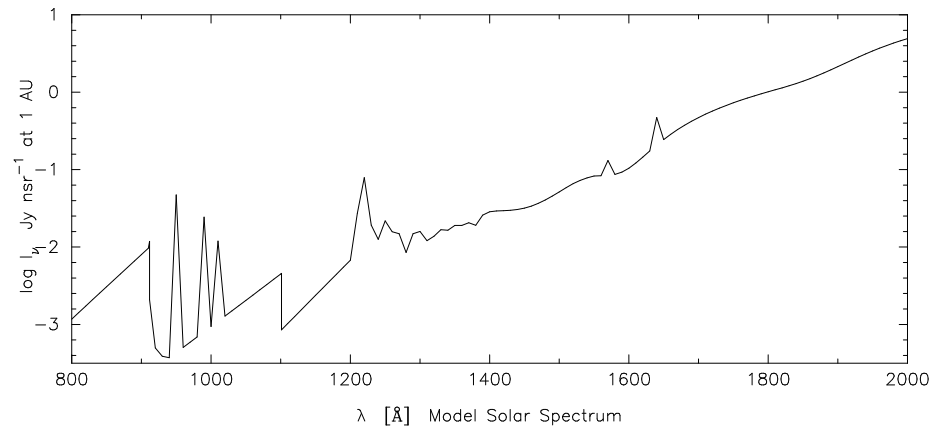
But this ignores source and sink terms as well as coupling to external radiation and it yields relative populations only.

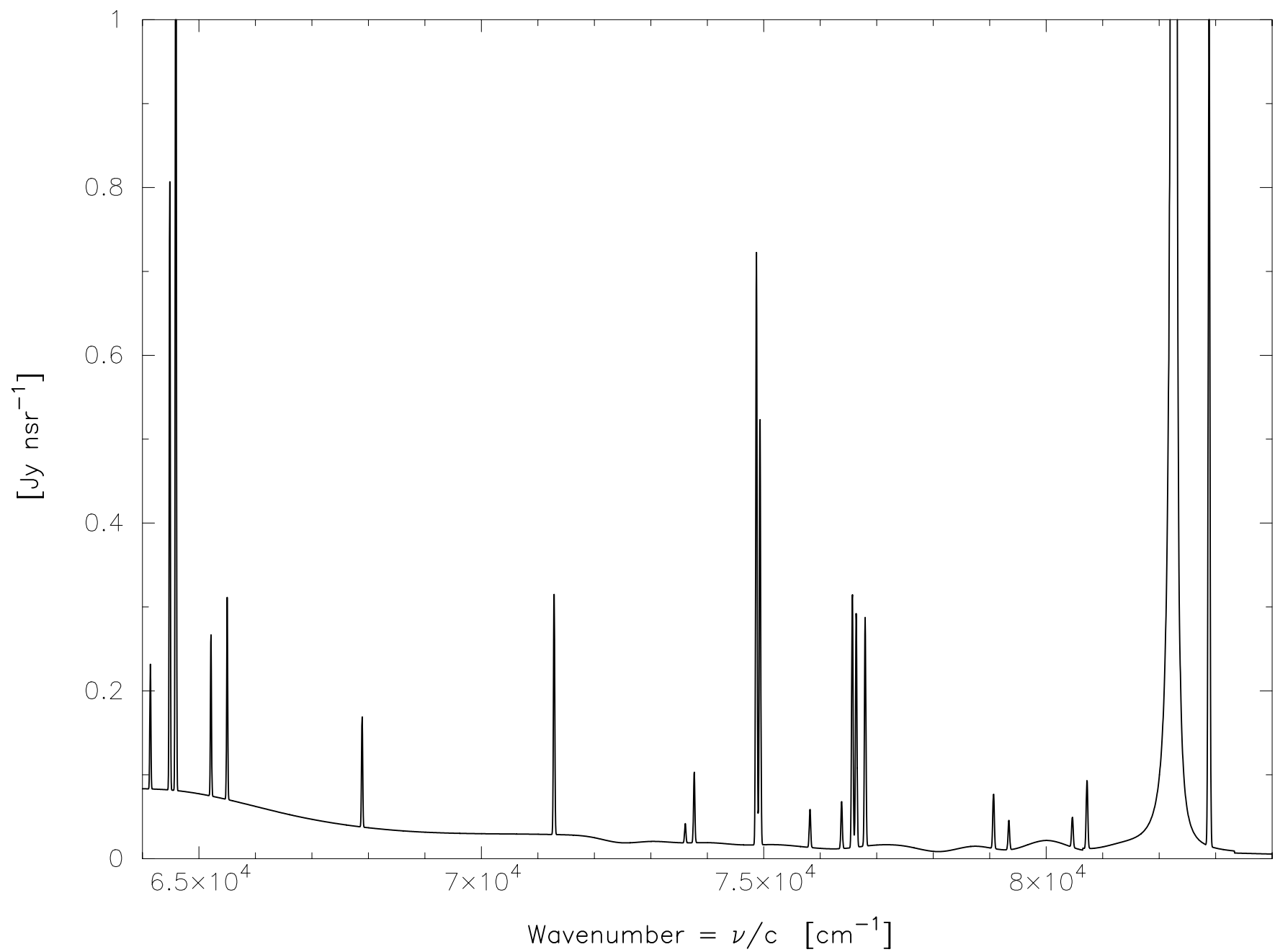
We recommend

$$\begin{aligned} \frac{dn_i}{dt} = & \sum_{j>i} n_j (A_{ji} + I_\nu B_{ji}) - \sum_{k<i} [n_i (A_{ik} + I_\nu B_{ik}) - n_k I_\nu B_{ki}] \\ & + \sum_{\ell \neq i} (n_\ell C_{\ell i} - n_i C_{i\ell}) = F_i(T_{\text{form}}) - n_i D_i \end{aligned}$$

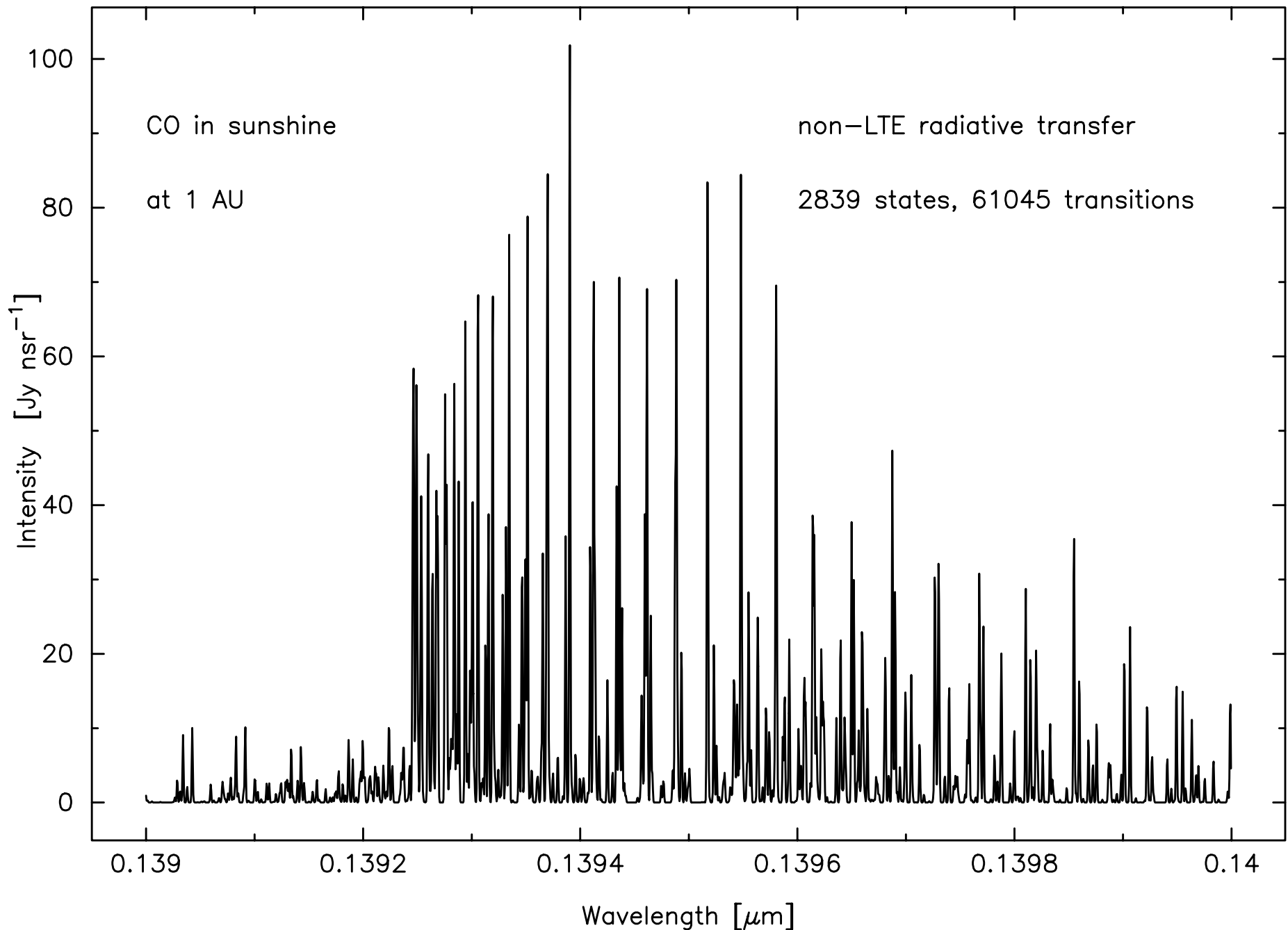
The state-specific formation rate  $F_i$  is a model of the formation process (e.g. a Boltzmann distribution at some formation temperature  $T_{\text{form}}$ ). It permits different spin-modifications to be treated together even without reactive interchange processes. For properly chosen  $F_i$  and  $D_i$ , the solutions are number densities.

Input solar spectrum  
UV from chromosphere  
and corona  
represents an enormous  
excess over the  
 $T_{\text{eff}}=5800\text{ K}$   
photosphere

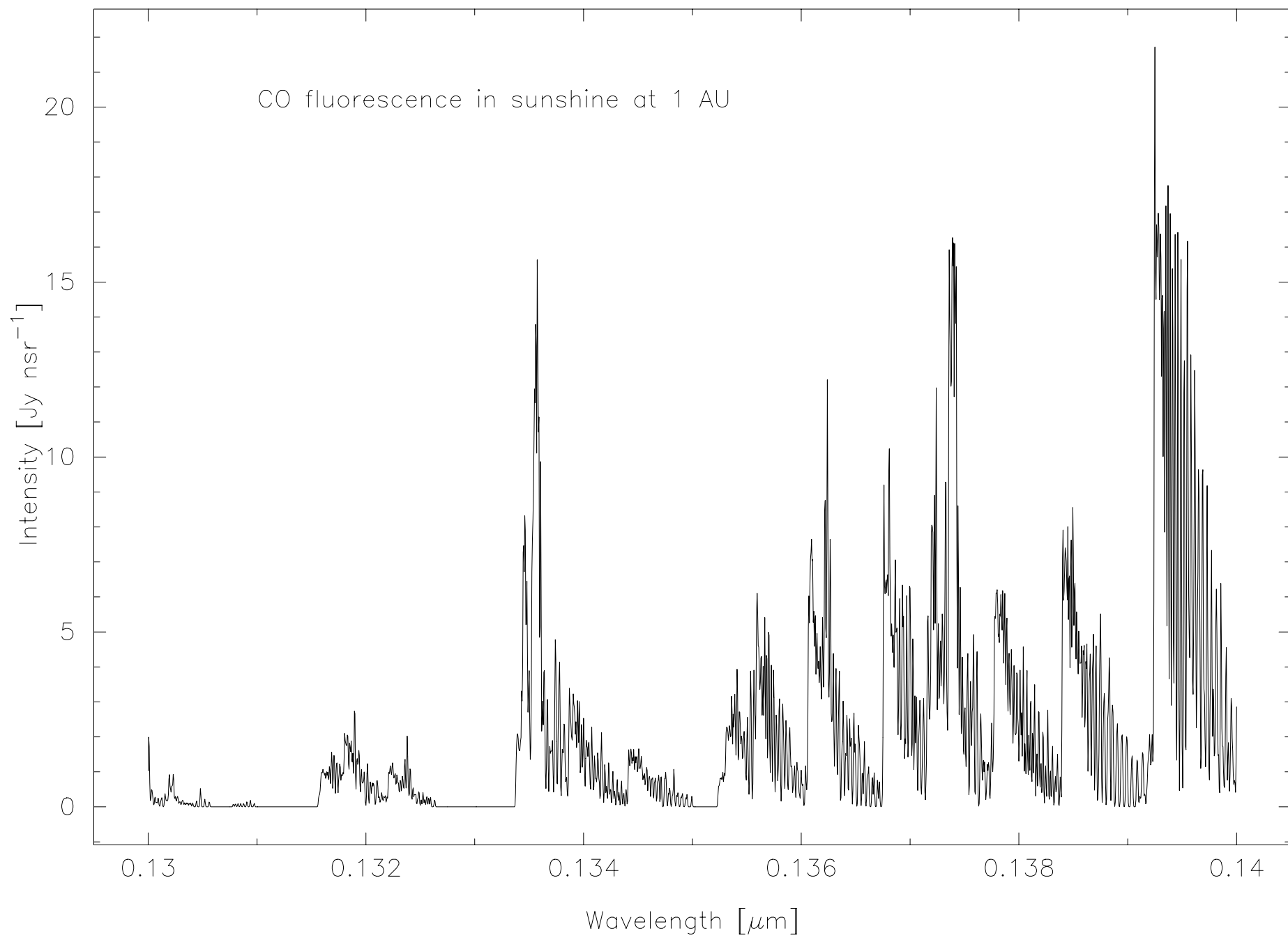


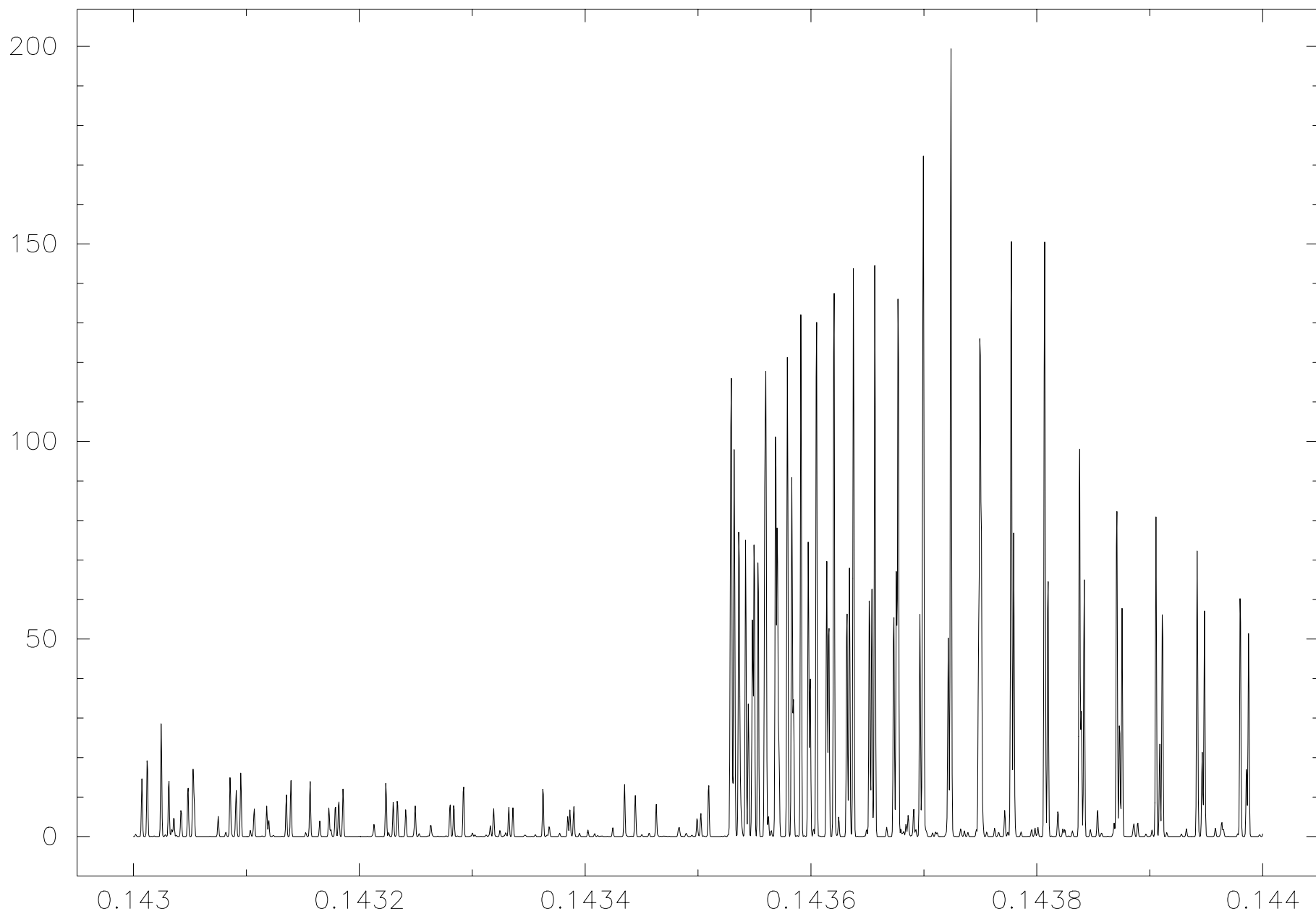


# Example: UV-pumped CO in sunlight



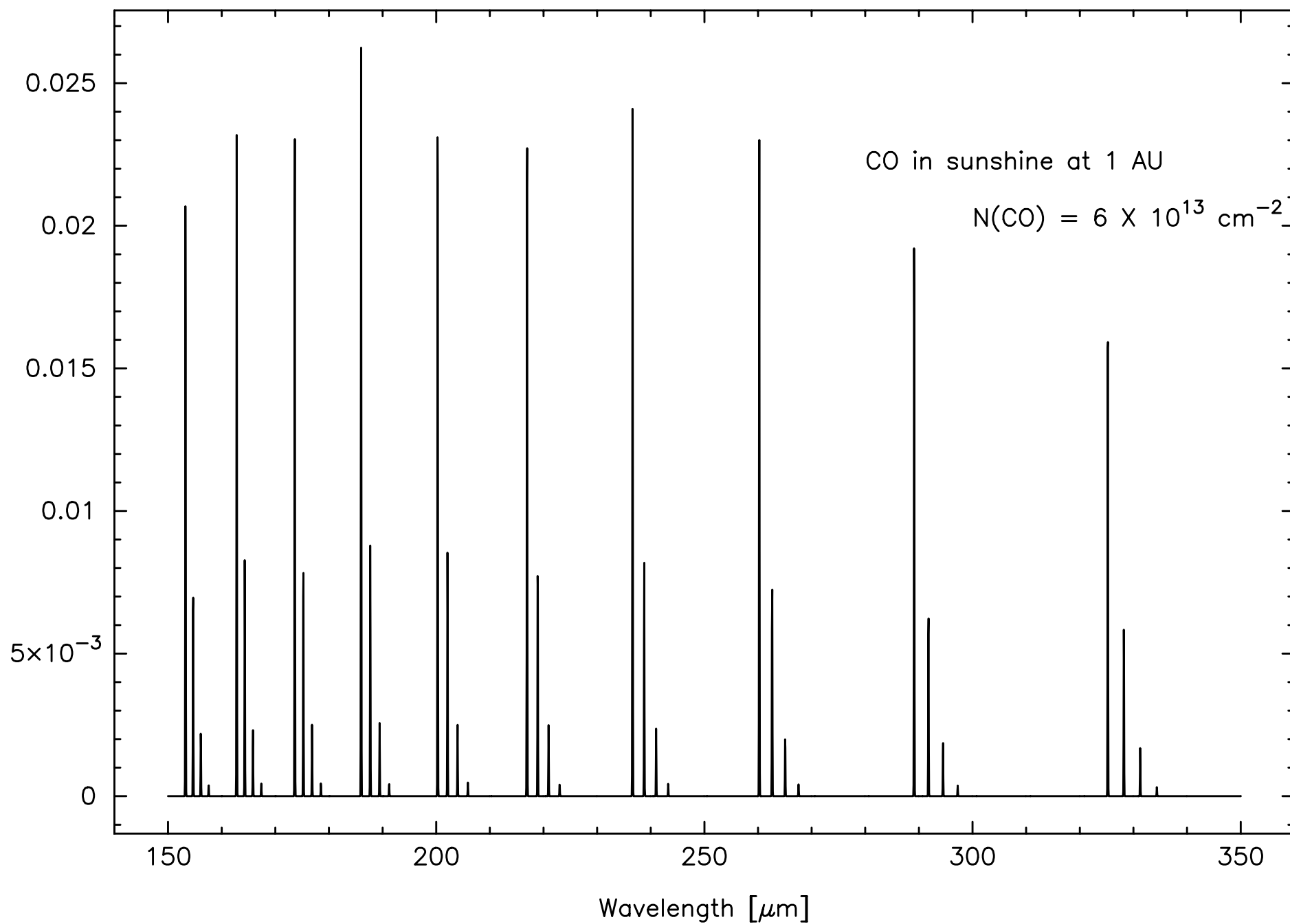
J.H. Black & J. L. Fox (in preparation)



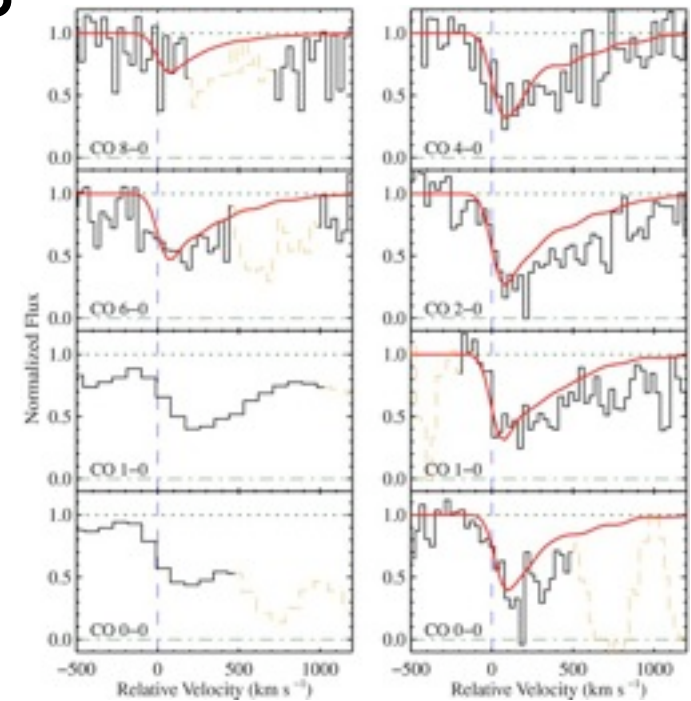
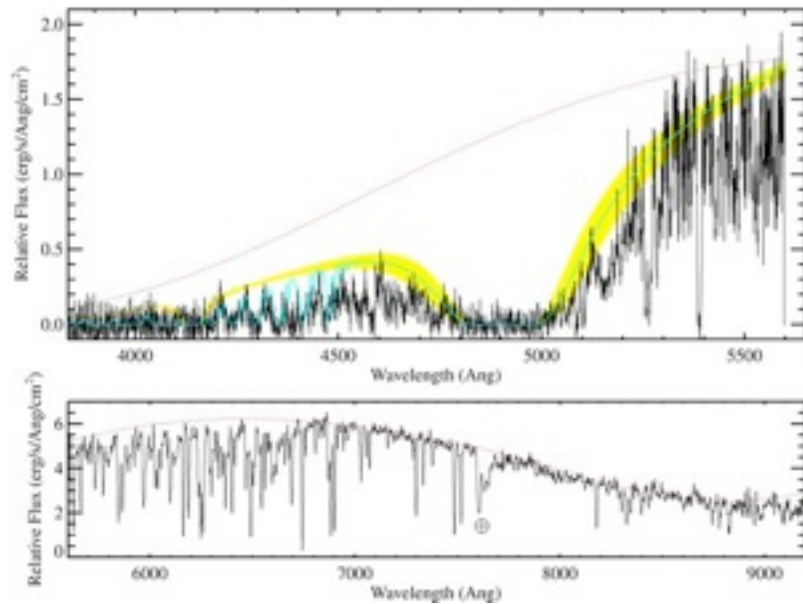




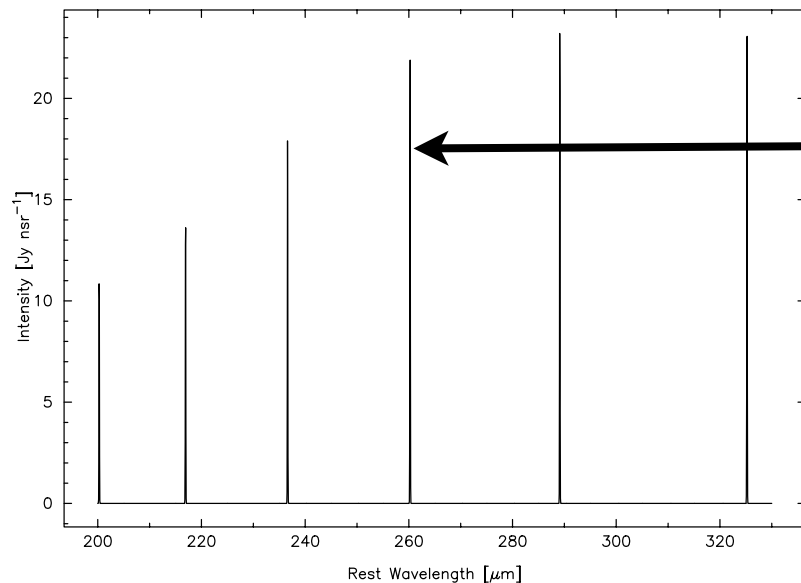
Intensity [ $\text{Jy nsr}^{-1}$ ] at  $0.1 \mu\text{m}$  resolution



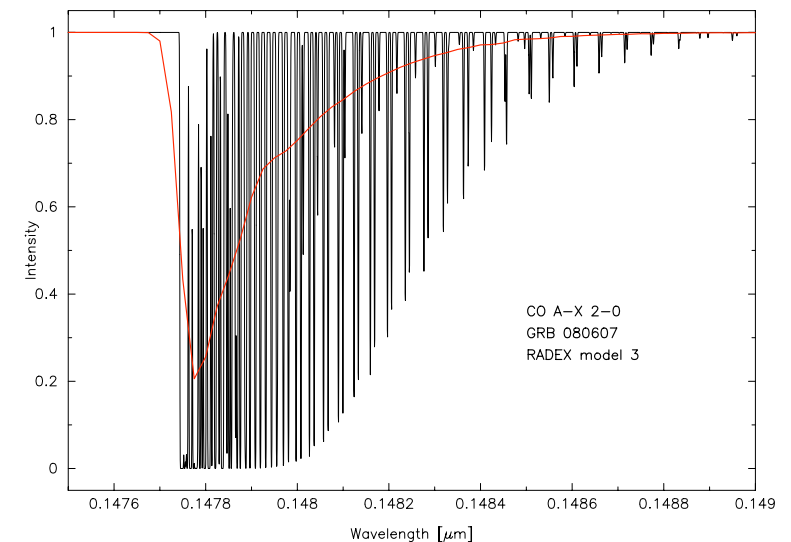
# GRB080607: H<sub>2</sub> and CO at z=3.0363



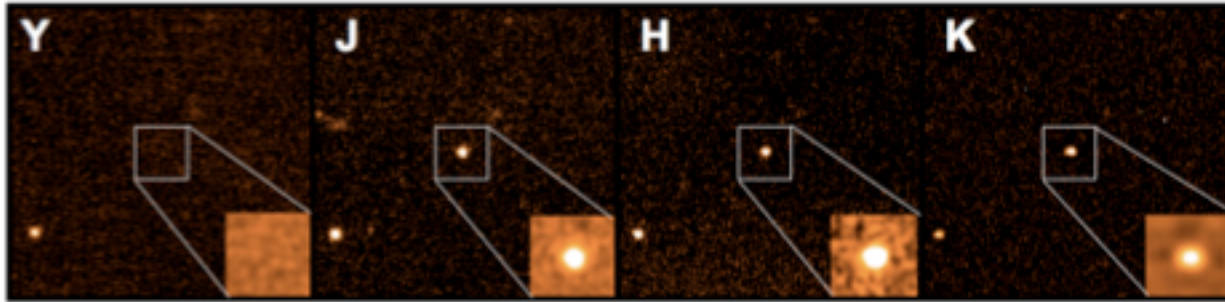
Prochaska et al. (2009, ApJ, 691, L27), Sheffer et al. (2009, ApJ, 701, L63) show that H<sub>2</sub> is pumped by the UV afterglow ↑  
Black (2009, in prep.) predicts the UV pumping in CO, which produces submm-wave emission, too ↓



CO J=12-11  
1.38 THz (rest)  
red-shifted to  
342.4 GHz  
at z=3.0363



# Afterglows of $\gamma$ -ray Bursts at High Redshift



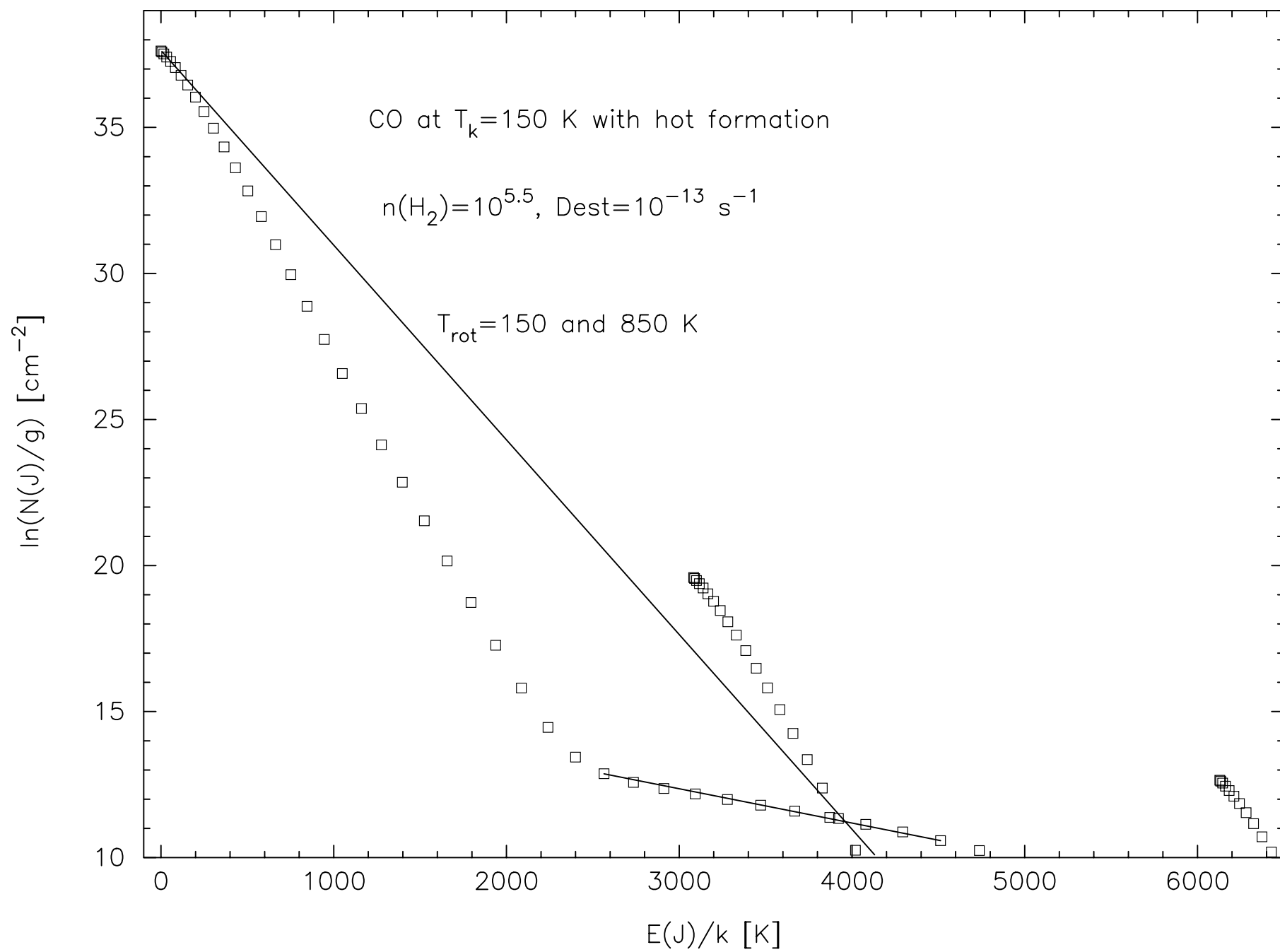
UKIRT discovery images  
from Tanvir et al.  
[astro-ph:0906.1577](#)

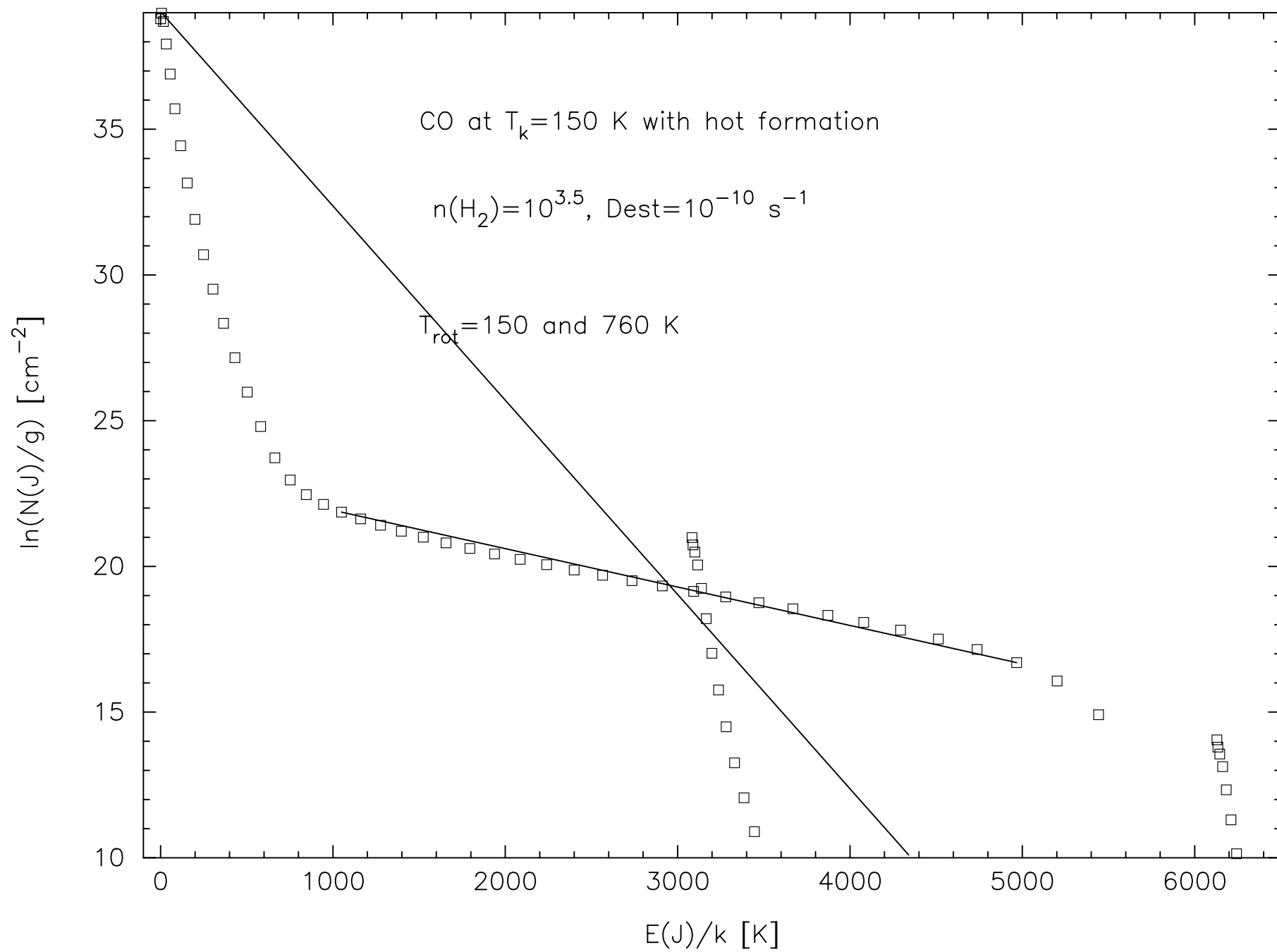
GRB090423 was the most distant known  
source in the Universe at  $z=8.26$

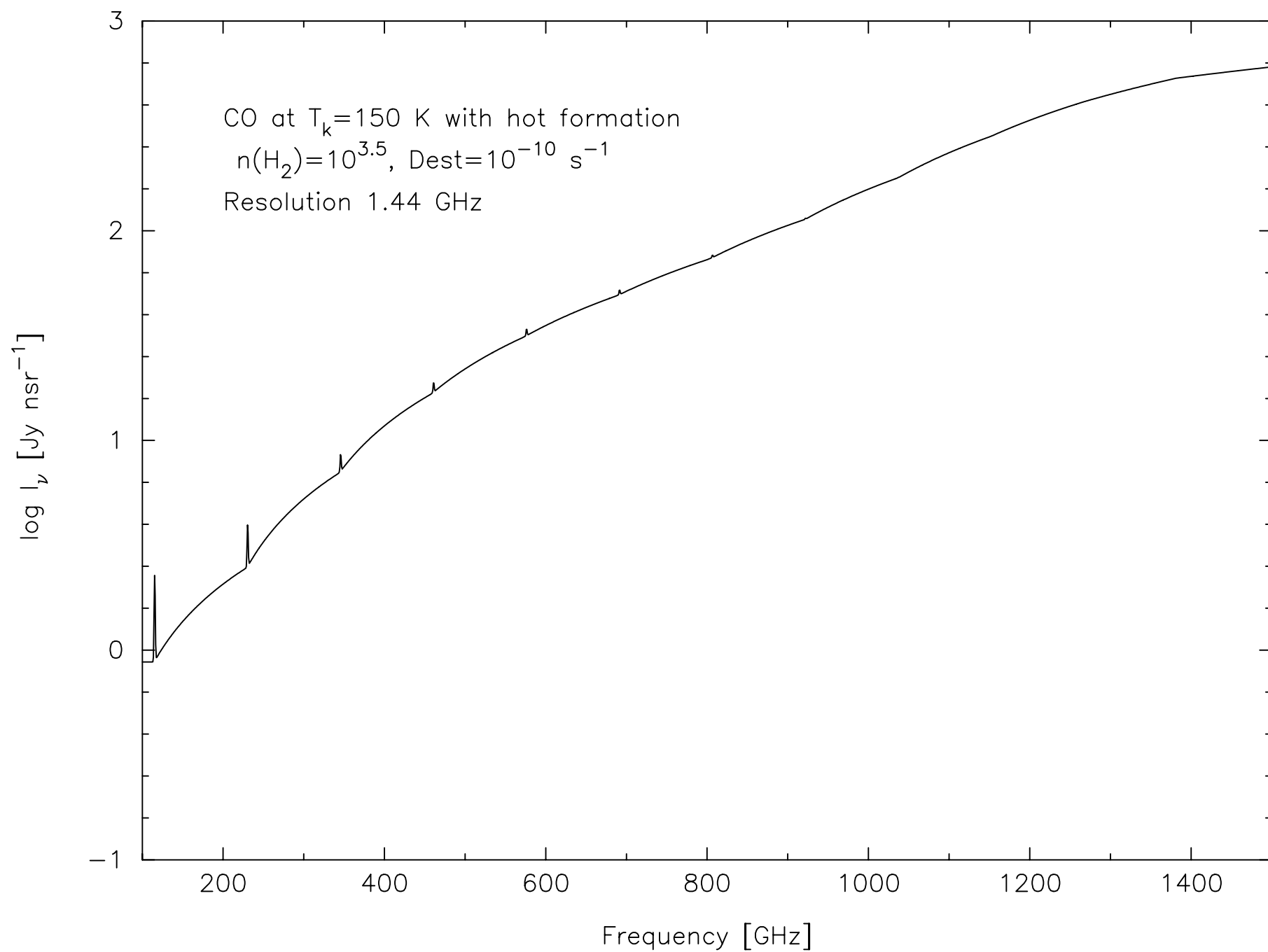
**mm/submm-wave afterglow?**

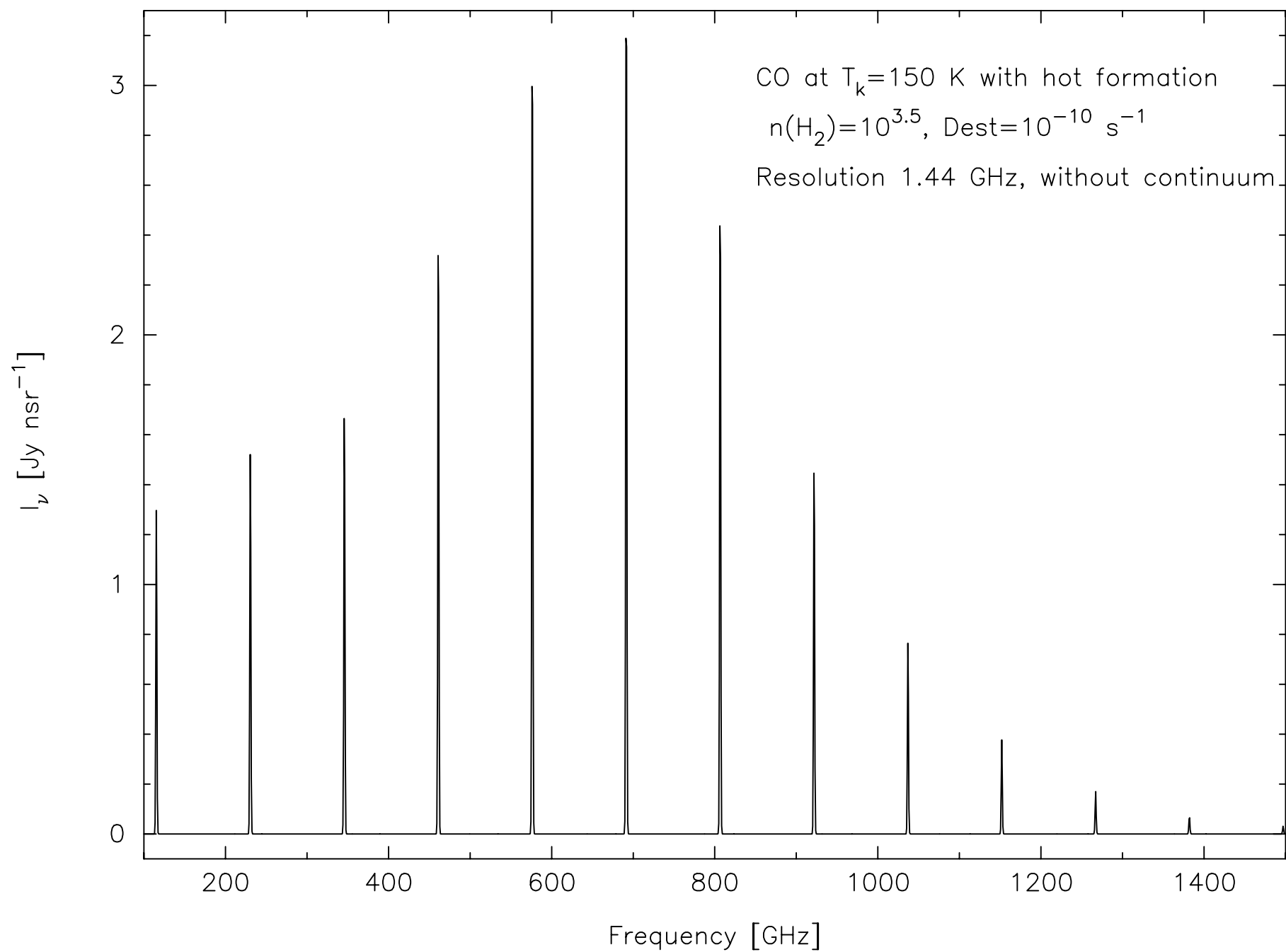
Castro-Tirado et al. report a  $\lambda=3$  mm  
source at the burst position with flux density 0.2 mJy  
(GCN Circular 9273, 09-04-28)  
consistent with Bock et al. upper limit of 0.7 mJy at CARMA  
 **$\Rightarrow$ Prospects for ALMA to probe high- $z$  galaxies**

- CO can be formed with considerable internal energy
- Example:  $\text{HCO}^+ + \text{e}^- \rightarrow \text{H} + \text{CO}(\text{v}, \text{J}) + 7.5 \text{ eV}$
- Set formation rate according to  $\text{HCO}^+$  and  $\text{e}^-$  abundances
- In PDR/XDR this could be  $F = 4 \times 10^{-19} n^2 \text{ [cm}^{-3} \text{ s}^{-1}]$  or higher
- Balance with a destruction rate that gives the needed CO abundance or column density
- Apply RADEX in all-singing, all-dancing mode





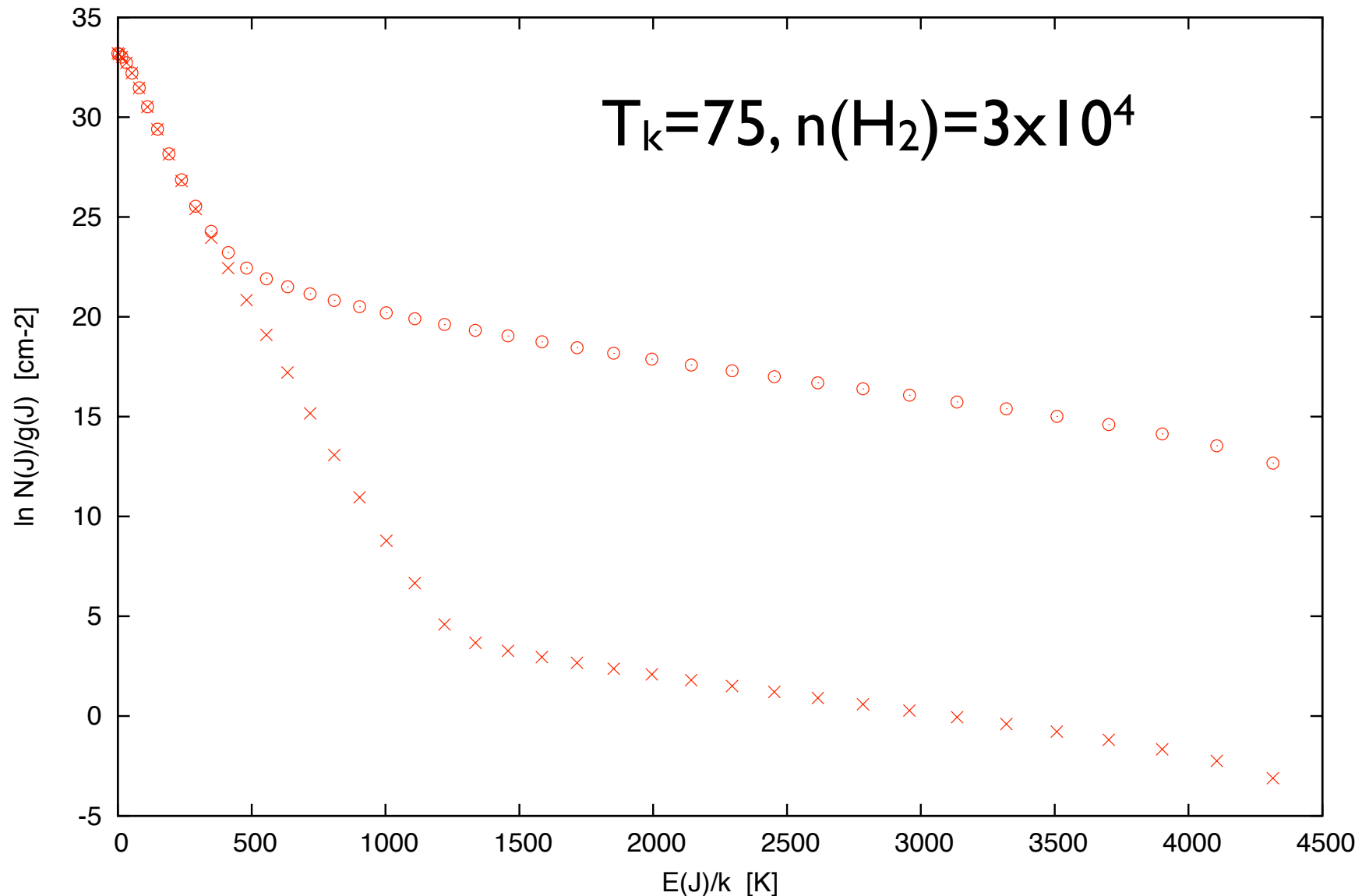






Formation/destruction may be important in common molecules if the rates are fast enough.  
For example,  $^{13}\text{CO} + ^{12}\text{C}^+ \rightarrow ^{12}\text{CO} + ^{13}\text{C}^+$  in PDR can mimic a high-temperature component.

$^{13}\text{CO}$  rotational excitation w/wo rapid destruction by  $^{12}\text{C}^+$  reaction



- Recent evidence from UV spectra of T Tau disks that CO is excited both by UV fluorescence and e-impact (France et al. 2011)
- Hot electrons are expected in XDR and in magnetospheres of Jupiter-like planets
- Electron collisions are not directly important in rotational excitation of CO (small dipole moment)
- Electron collisions can excite triplet electronic states

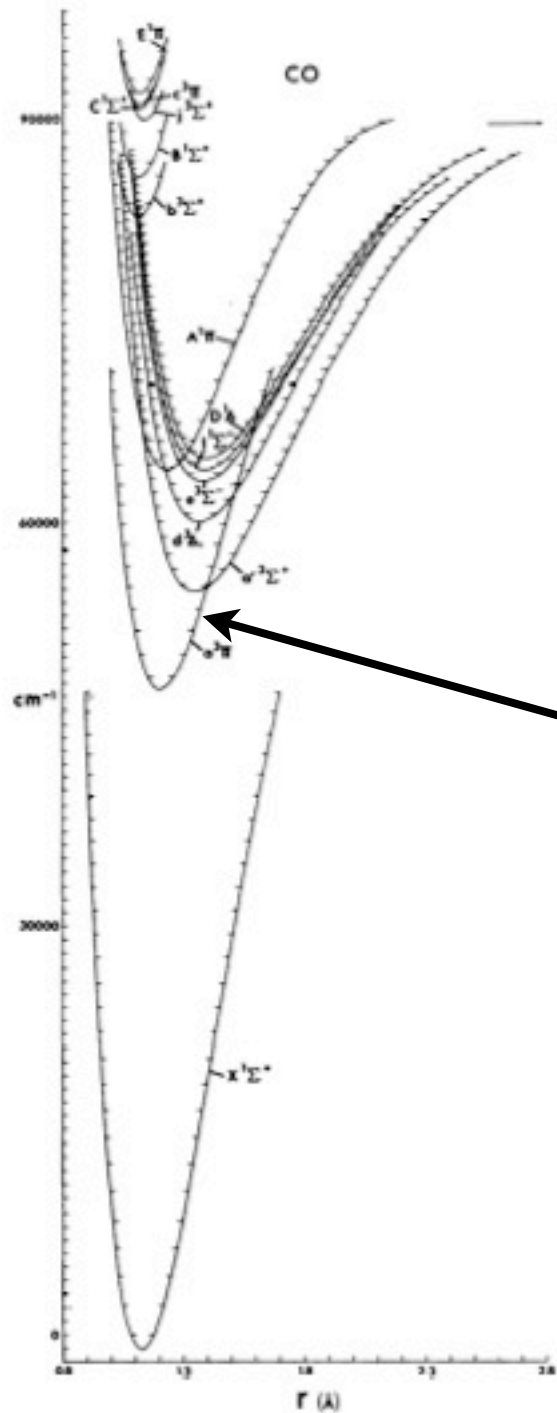
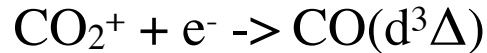


FIGURE 3. Potential energy curves for the observed states of CO below 11.77 eV.

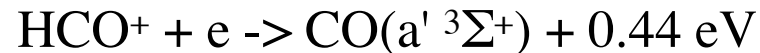
The arrow indicates the dissociation energy of CO. The dash

$a^3\Pi$  is the upper state of the  $a$ -X Cameron system with (0,0) band near 200 nm  $a^3\Pi$  has a well measured rotational spectrum; e.g.  $v=0, \Omega=0 J=1-0$  is a  $\Lambda$ -doublet at 92.5, 92.9 GHz

CO triplets can be excited dissociative recombination, e.g.



there is a yield of 0.29 for  $\text{CO}_2^+ + e^- \rightarrow \text{CO}(\text{a-X})$



and  $\text{HOC}^+ + e^-$  has even more accessible states in product CO  
(R. Johnsen et al. 2009)

J-dependence of  $\text{a}^3\Pi$  lifetimes suggests population inversions might easily occur

CO triplet bands show up in the UV spectrum of **HD 44179 (Red Rectangle)**  
while this is an AGB star in a binary system, the source of UV radiation  
has been identified as the hot, inner region of an accretion disk  
(Witt et al 2009)

in comets,  $\text{CO}(\text{a-X})$  comes from  $\text{CO}_2$  photodissociation and e-impact

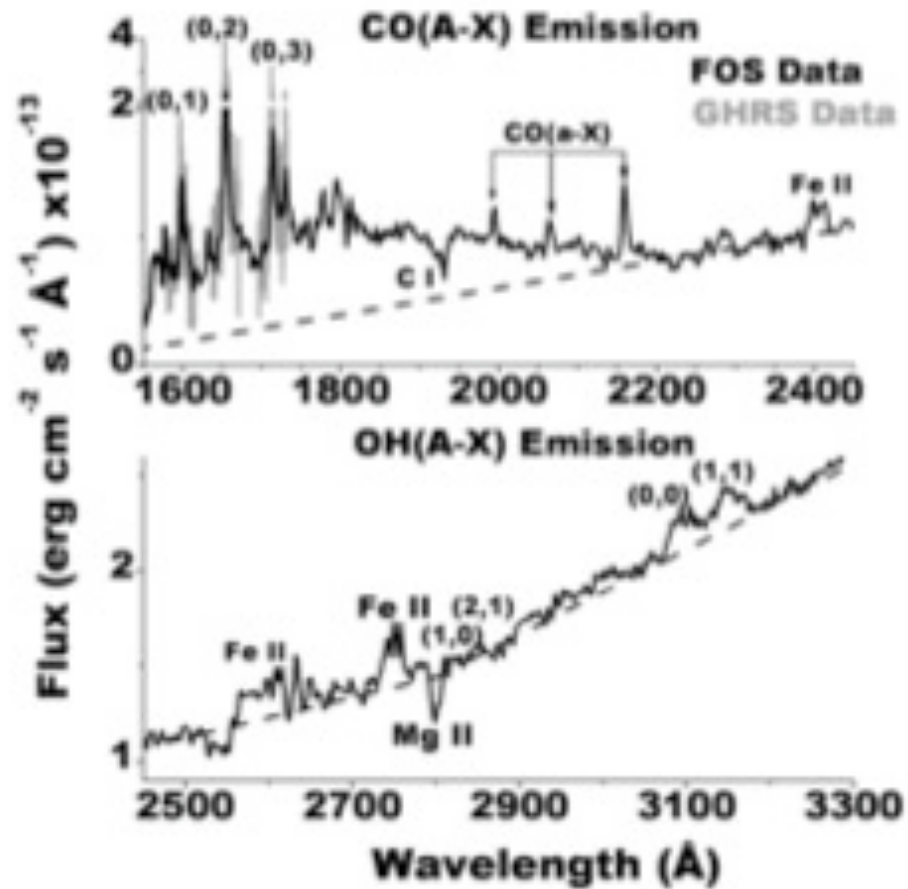
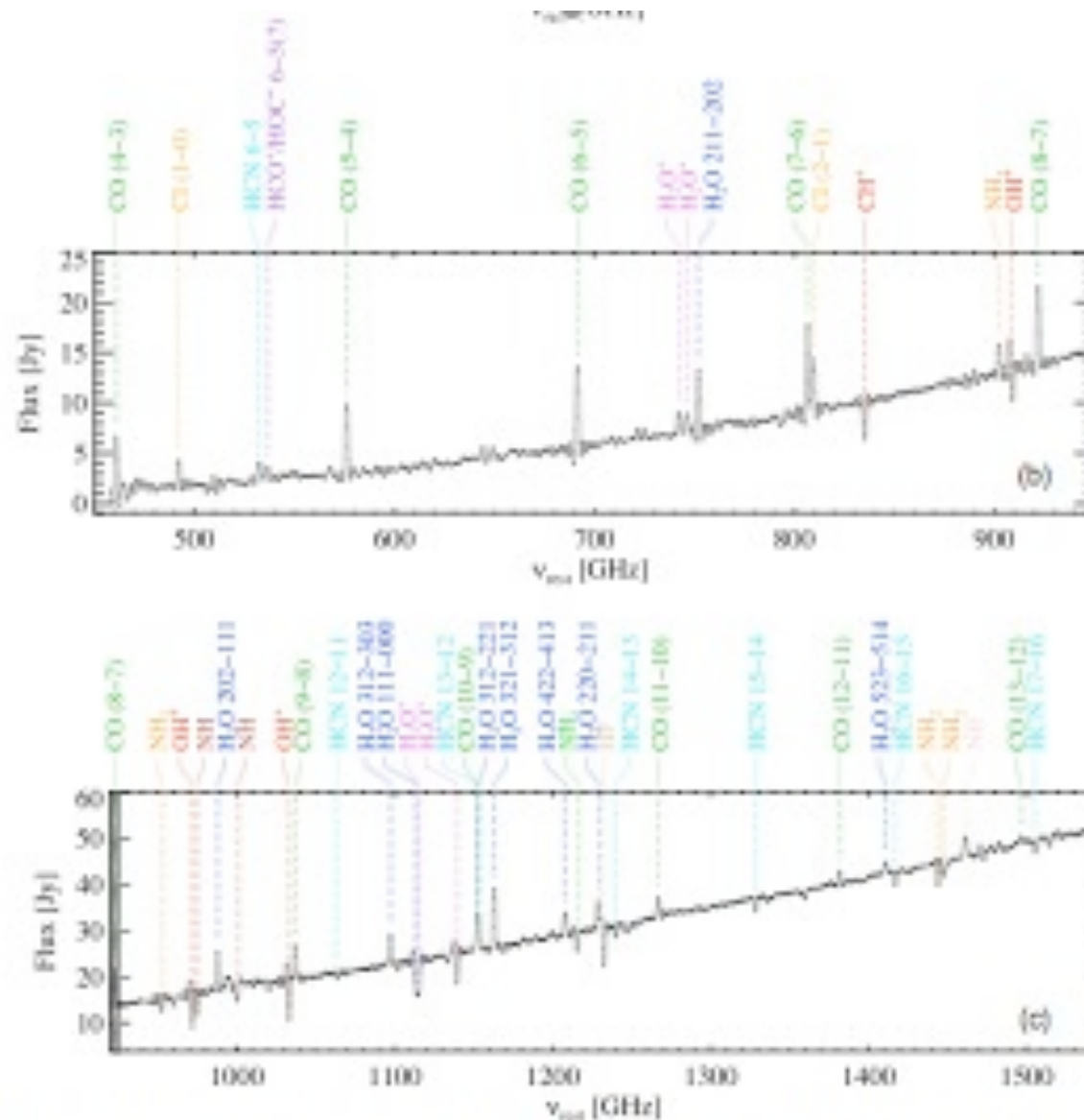


FIG. 3.—*Top:* FOS and GHRS data are compared in the CO(A-X) emission region. *Bottom:* FOS data in the OH(A-X) emission region. Also shown is the estimated underlying continuum for the FOS data.

## CO Cameron bands in the Red Rectangle



**Figure 1.** *Herschel* SPIRE-FTS spectrum of Arp 220. The spectrum shows the FIR continuum between 190 and 670  $\mu\text{m}$  in (a). Red solid points in (a) are the continuum measurements from the SPIRE photometer and the dotted curves show the photometer bandpasses with arbitrary normalization. Line identifications are shown for several molecular and atomic species in (b) and (c) with like colors for like species.

(A color version of this figure is available in the online journal.)

## OH<sup>+</sup> absorption/emission in Arp 220

### Rangwala et al. (2011)

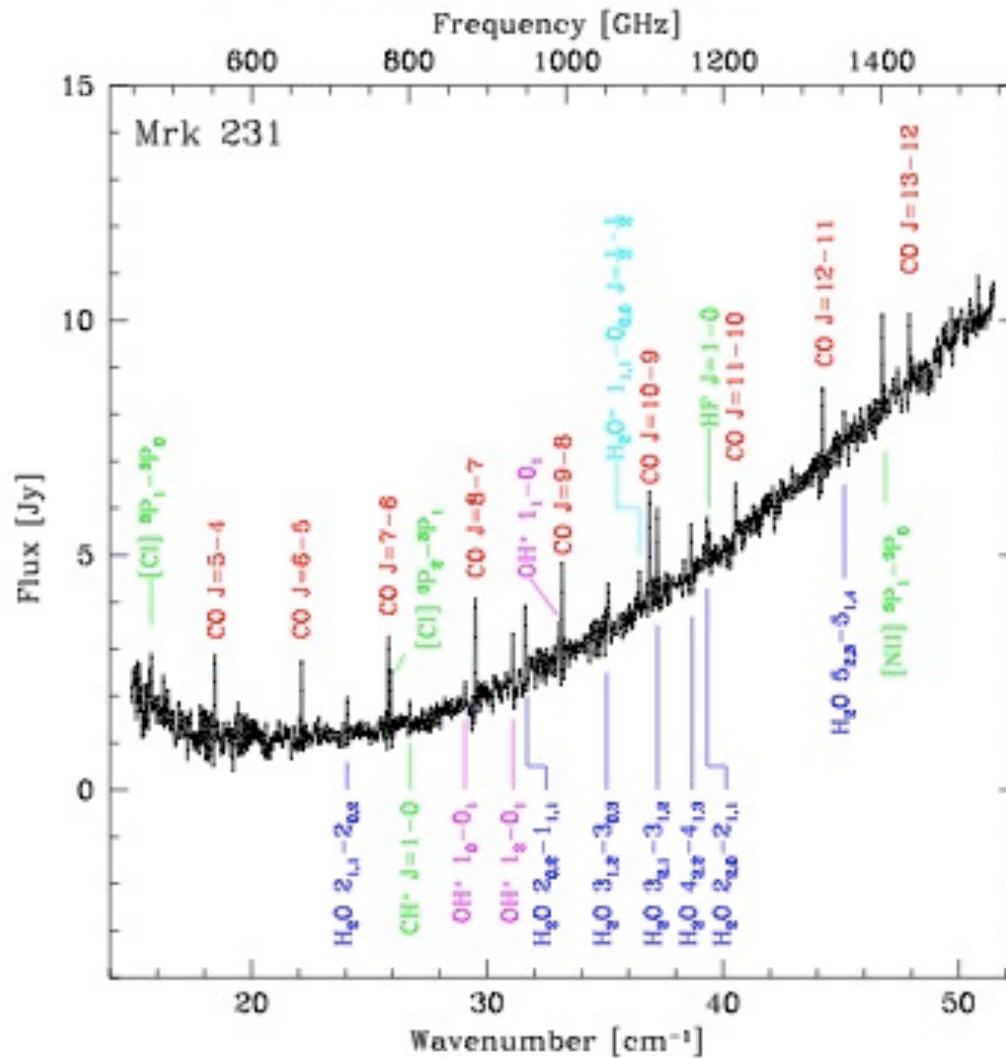


Fig. 1. SPIRE FTS spectrum of Mrk 231. Line identifications are given in red for CO lines, in blue for H<sub>2</sub>O, in magenta for OH<sup>+</sup>, in cyan for H<sub>2</sub>O<sup>+</sup>, and in green for the remaining lines.

OH<sup>+</sup> in  
Mrk 231  
(van der Werf et al. 2010)

# How does $\text{OH}^+$ work?

- 909, 971, 1033 GHz lines:  $A=0.02 \text{ s}^{-1}$
- Destruction: by  $\text{OH}^+ + \text{e}^- \rightarrow \text{O} + \text{H}$ ,  $k=0.4 \times 10^{-7} \text{ cm}^3 \text{ s}^{-1}$  at  $T=300$  and by  $\text{OH}^+ + \text{H}_2 \rightarrow \text{H}_2\text{O}^+ + \text{H}$ ,  $k=1.0 \times 10^{-9} \text{ cm}^3 \text{ s}^{-1}$
- Source:  $\text{H}_3^+ + \text{O} \rightarrow \text{OH}^+ + \text{H}_2$  exoergic by 5200 K
- At  $n(\text{H}_2)=10^{5.5} \text{ cm}^{-3}$ ,  $\text{Dest}=3 \times 10^{-4} \text{ s}^{-1}$ , faster than collisional excitation for  $n(\text{e}) < 400 \text{ cm}^{-3}$
- Far-IR pumping at 950 GHz  $\sim \text{Dest}$  where  $T_{\text{rad}}=11 \text{ K}$  ( $I=20 \text{ Jy nsr}^{-1}$ )
- SUMMARY:  $\text{OH}^+$  may naturally appear in emission where electron density is high enough (as in XDR), but the destruction rate must be included explicitly in rate equations. Emission/absorption: dilution of continuum -- relative sizes of central source & molecular gas



