- WHY ARE PLANTS GREEN ? CELL PLANTS (ALGAE) GREEN
(MARON SAYS THEY REFLECT GREEN AND ABSORB RED & BLUE LIGHT)

- WHY ARE RUBYS RED?

WHY ARE STOP LIGHTS RED?

BROADENING

(i) NATURAL

(li) COLLISION

(iii) DOPPLER

* LINES BECOME MORE NARROW AS A MATERIAL COOK

WE TAKE ADVANTAGE OF THE FACT THAT

Linkon

N = n+ix =1 : CAN USE LORENTZ MODEL

> K, "ABSORPTION COEFFICIENT"

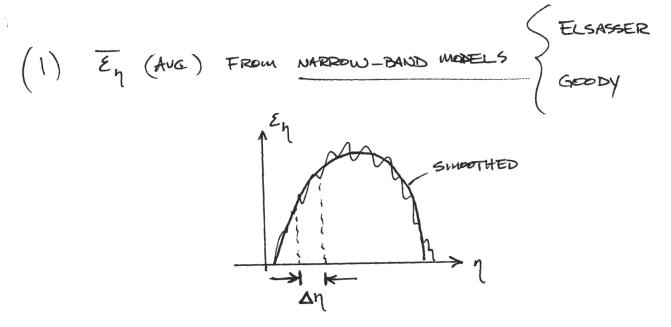
3

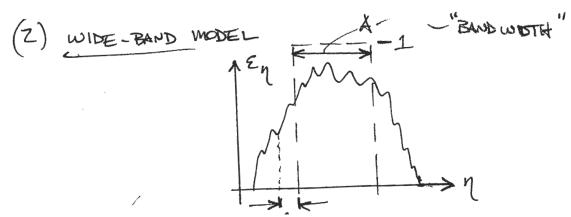
$$\frac{dI_{\eta}}{dx} = -K_{\eta}I_{\eta} \implies T_{\eta} = e^{-K_{\eta}X}$$

EX! BOX WITH A GAS

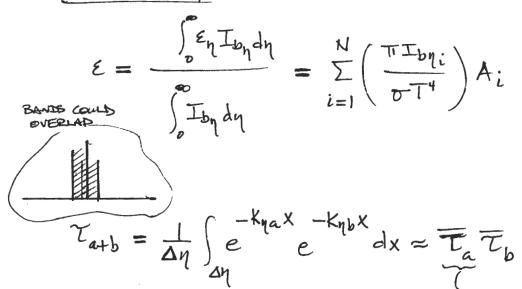
$$\Sigma_{\eta} = 1 - \tau_{\eta}$$
 $X=D$

* * WE CAN MODEL THE SPACING AND SHAPE OF EACH LINE TO GET En?





TOTAL EMISSIVITY

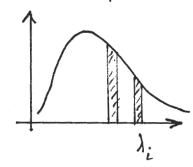


$$\mathcal{E}_{a+b} = 1 - \mathcal{T}_{a+b} = \mathcal{E}_a + \mathcal{E}_b - \mathcal{E}_a \mathcal{E}_b$$

PLANK MEAN ABSORPTION COEFFICIENT

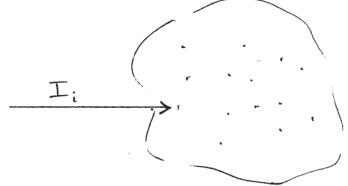
$$K_{p} \equiv \frac{\int_{0}^{\infty} K_{\eta} I_{b\eta} d\eta}{\int_{0}^{\infty} I_{b\eta} d\eta}$$

BLK-BODY SCLAR



2.58 4/13/06

EQUATION OF RADIATIVE TRANSFER REDRY OF PARTICLES (e.g., e, PHOTONS, MOLECULES



SCATTERING: CHANGES DIRECTION OF ENERGY FLOW

ABSORPTION: REDUCES ENERGY TRAVELING IN INCIDENT DIRECTION

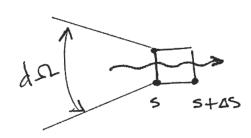
EMISSION: CAN OCCUR IN ALL DIRECTIONS AT

THE PARTICLES TEMPERATURE; AND THUS A PORTION

SO OF EMISSION CONTRIBUTES TO INCOMINE

DIRECTION

TAKE & CONTROL VOLUME



POWER IN POWER DON'T FOWER ABSORBED

I(s) d Ddy = I(s+ds) d Ddy + Ky I y d Ddy ds

$$(K_{\eta} = \frac{4\pi R}{1})$$
 $N = n + i \times$

... ADD EMISSION (ONLY PECTED TO OBJECTS TEMP. AND WITHOUT (N
ALL DIRECTIONS)

LOCAL Emissivity -
$$l^{oex}$$
 since $k_h \Delta s$ is small o.r.t. 1
$$E(x) = l - e \approx k_h \Delta s$$

TOTAL ABS. + EMISS. -

$$\frac{dI_{\eta}}{ds} = -K_{\eta}I_{\eta} + K_{\eta}I_{b\eta}(T)$$
ABSORPTION EMISSION

LEGISLATION

L

· OUT SCATTERING (SINGLE PARTICLE)

· GIZOUP OF PARTICLES

$$K_s = NC_s \left[\frac{1}{m}\right]$$
 $K_a = NC_a$



DISTRIBUTION OF SCATTERIN RECALL PHASE FUNCTION ACTUAL VS. ACTUAL * RELATED TO OUT INCOMING SCAFFERING IS THE RELATED TO A SPECTEL TURECTON NOW OUTGOING SCATTERING REDUCES THE INTENSITY .. our ERN. BECOMES abs outgoint toring emission dIy(a) - Kan In - Ksn In + Kan Ion(T) + in scattering NOW THE INCOMING SCATTERING $\frac{d \operatorname{In}(\Omega)}{ds} = -k_{a\eta} \operatorname{In} - k_{s\eta} \operatorname{In} + k_{a\eta} \operatorname{Ibn}(T) + \frac{k_{s\eta}}{4\pi} \int \operatorname{In}'(\Omega') \Phi(\Omega' \to \Omega) d\Omega'$ SWEE C IS SO LARGE, 1 dI + , BUT FOR MOST APPLICATIONS HAY COULD ADD TRANSIENT HERE ENGINEERING WE IGNORE THIS TERM

WE OBSERVE

$$I_{\eta} \left(\Omega, \vec{s}\right)$$

$$\Omega = \Omega(\Theta, \phi)$$

$$C \times \Theta \stackrel{?}{\leftarrow} \phi \Longrightarrow N_{\chi_1} N_{\gamma_1} N_{\overline{\gamma}_2}$$

=> PHASE SPACE

$$f(\vec{r}, \vec{r}, t)$$