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LIDAR - Parts I and II -

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LIDAR 1 & 2

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LIDAR: What and Why?

LIDAR stands for Light Detection and Ranging, commonly known as Laser Radar.

Lidar is not only replacing conventional sensors, but also creating new methods with unique properties that could not be achieved before.

Lidar is extremely useful in atmospheric and environmental research as well as space exploration. It also has wide applications in industry, defense, and military.

Earth Atmosphere and Space





Fundamentals of the LIDAR as Remote Sensing Tool

Physical Picture of Lidar Equation

Fundamental Lidar Equation

Different Forms of Lidar Equation

Illustration of Lidar Equation



Picture of LIDAR in its Environment



Physical Picture of Lidar Equation



Considerations for Lidar Equation

In general, the interaction between the light photons and the particles is a scattering process.

- The expected photon counts are proportional to the product of the
- (1) transmitted laser photon number,
- (2) probability that a transmitted photon is scattered,
- (3) probability that a scattered photon is collected,
- (4) light transmission through medium, and
- (5) overall system efficiency.
- Background photon counts and detector noise also contribute to the expected photon counts.

Basic Assumptions in Lidar Equation

- Independent scattering: particles are separated adequately and undergo random motion so that their contributions to the total scattered energy have no phase relation. Thus, the total intensity is simply a sum of the intensity scattered from each particle.
- Single scattering implies that a photon is scattered only once and multiple scattering is excluded.

Fundamental Lidar Equation

 $N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R)\Delta R\right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R)T(\lambda, R)\right] \cdot \left[\eta(\lambda, \lambda_{L})G(R)\right] + N_{B}$

1st Term: Transmitted Photon Number

$$N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R) \Delta R \right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R) T(\lambda, R) \right] \cdot \left[\eta(\lambda, \lambda_{L}) G(R) \right] + N_{B}$$

$$N_L(\lambda_L) = \left(\frac{P_L(\lambda_L)\Delta t}{hc/\lambda_L}\right)$$

Laser Power x time bin length

Planck constant x Laser frequency

Transmitted laser energy within time bin

Single laser photon energy

— Transmitted laser photon number within time bin length

2nd Term: Probability to be Scattered

 $N_{S}(\lambda,R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda,\lambda_{L},\theta,R) \Delta R \right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L},R)T(\lambda,R) \right] \cdot \left[\eta(\lambda,\lambda_{L})G(R) \right] + N_{B}$

Angular scattering probability – the probability that a transmitted photon is scattered by scatters into a unit solid angle.

Angular scattering probability = volume scatter coefficient β x scattering layer thickness ΔR

$$\beta(\lambda,\lambda_L,R) = \sum_{i} \left[\frac{d\sigma_i(\lambda_L,\theta)}{d\Omega} n_i(R) p_i(\lambda) \right] \quad (m^{-1}sr^{-1})$$

3rd Term: Probability to be Collected

$$N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R) \Delta R\right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R)T(\lambda, R)\right] \cdot \left[\eta(\lambda, \lambda_{L})G(R)\right] + N_{B}$$

The probability that a scatter photon is collected by the receiving telescope, i.e., the solid angle subtended by the receiver aperture to the scatterer.



4th Term: Light Transmission

$$N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R) \Delta R\right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R)T(\lambda, R)\right] \cdot \left[\eta(\lambda, \lambda_{L})G(R)\right] + N_{B}$$

The atmospheric transmission of laser light at outgoing wavelength λ_{L} and return signal at wavelength λ

Transmission
for laser light
$$T(\lambda_L, R) = \exp\left[-\int_0^R \alpha(\lambda_L, r) dr\right]$$
Transmission
for return signal $T(\lambda, R) = \exp\left[-\int_0^R \alpha(\lambda, r) dr\right]$

Where $\alpha(\lambda_L, R)$ and $\alpha(\lambda, R)$ are extinction coefficients (m⁻¹)

$$T(\lambda_L, R)T(\lambda, R) = \exp\left[-\left(\int_0^R \alpha(\lambda_L, r)dr + \int_0^R \alpha(\lambda, r)dr\right)\right]$$

when $\lambda = \lambda_L$
= $\exp\left[-2\int_0^R \alpha(\lambda, r)dr\right]$

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Extinction Coefficient $\boldsymbol{\alpha}$

$$\alpha(\lambda, R) = \sum_{i} \left[\sigma_{i, ext}(\lambda) n_i(R) \right]$$

 $\sigma_{i,ext}(\lambda)$ is the extinction cross-section of species i $n_i(R)$ is the number density of species i

Extinction = Absorption + Scattering (Integrated)

$$\sigma_{i,ext}(\lambda) = \sigma_{i,abs}(\lambda) + \sigma_{i,sca}(\lambda)$$

Total Extinction = Aerosol Extinction + Molecule Extinction

$$\alpha(\lambda, R) = \alpha_{aer, abs}(\lambda, R) + \alpha_{aer, sca}(\lambda, R) + \alpha_{mol, abs}(\lambda, R) + \alpha_{mol, sca}(\lambda, R)$$

5th Term: Overall Efficiency

$$N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R) \Delta R\right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R)T(\lambda, R)\right] \cdot \left[\eta(\lambda, \lambda_{L})G(R)\right] + N_{B}$$

 $\eta(\lambda,\lambda_L) = \eta_T(\lambda_L) \cdot \eta_R(\lambda)$ is the lidar hardware optical efficiency e.g., mirrors, lens, filters, detectors, etc

G(R) is the geometrical form factor, mainly concerning the overlap of the area of laser irradiation with the field of view of the receiver optics



6th Term: Background Noise

$$N_{S}(\lambda, R) = N_{L}(\lambda_{L}) \cdot \left[\beta(\lambda, \lambda_{L}, \theta, R) \Delta R\right] \cdot \frac{A}{R^{2}} \cdot \left[T(\lambda_{L}, R)T(\lambda, R)\right] \cdot \left[\eta(\lambda, \lambda_{L})G(R)\right] + N_{B}$$

 N_{B} : Background noise, detector and circuit shot noise.

Up-looking lidar: background noise comes from solar scattering, solar straight radiation, star/city light.

Down-looking lidar: besides above noise, it could have three extra backgrounds: (1) Specular reflection from water/ice surface; (2) Laser reflectance from ground; (3) Solar reflectance from ground.

Illustration for LIDAR Equation



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Brief Review of Physical Processes in Lidars

Physical Processes in LIDAR

- Interaction between light and objects
- (1) Scattering (instantaneous elastic & inelastic):

Mie, Rayleigh, Raman, Brillouin scattering

- (2) Absorption and differential absorption
- (3) Laser induced fluorescence
- (4) Resonance fluorescence
- (5) Doppler shift and Doppler broadening
- (6) Boltzmann distribution
- (7) Reflection from target or surface

Light propagation in atmosphere or medium: transmission vs extinction

Extinction = Scattering + Absorption

$$T(\lambda, R) = \exp\left[-\int_0^R \alpha(\lambda, r) dr\right] \qquad \alpha(\lambda, R) = \sum_i \left[\sigma_{i,ext}(\lambda)n_i(R)\right]$$

 $T(\lambda,R)$ represents the fraction of light lost on the way from the lidar to the scattering volume and back!

(1) Elastic and Inelastic Scattering

Elastic scattering:

scattering with no apparent change of wavelength Sum of elastic scattering from atmospheric molecules and elastic scattering from aerosol particles and cloud droplets Rayleigh scattering and Mie scattering

Inelastic scattering:

scattering with apparent change of wavelength Raman scattering Pure rotational Raman and vibration-rotational Raman

Elastic and Inelastic Scattering (cont'd)

Rayleigh scattering is referred to the elastic scattering from atmospheric molecules (particle size is much smallerthan the wavelength), i.e., scattering with no apparent change of wavelength

With lower detection resolution, Rayleigh scattering can consist of the Cabannes scattering (really elastic scattering from molecules) and pure rotational Raman scattering.

Scattering still undergoes Doppler broadening and Doppler shift.

Temp.; Press. and collective motion of molecules lead to spectral broadening. Rayleigh scattering is therefore the sum of cabannes lines and the rotational Raman bands



Rayleigh Backscatter Coefficient

In lidar field, a common practice is to use the equation

$$\beta_{Rayleigh}(\lambda, z, \theta = \pi) = 2.938 \times 10^{-32} \frac{P(z)}{T(z)} \cdot \frac{1}{\lambda^{4.0117}} \left(m^{-1} s r^{-1} \right)$$

where P is the atmosphere pressure in mbar and T is the temperature in Kelvin at altitude z, λ is the wavelength in meter, and $\beta_{Rayleigh}$ is the backscatter coefficient (angular).

<u>Rayleigh scattering is proportional to λ^{-4} and dominates elastic</u> <u>backscatter at short laser wavelength</u>.

Rayleigh Backscatter Cross Section

□ It is also common in lidar field to calculate the Rayleigh backscatter cross section using the following equation

$$\frac{d\sigma_m(\lambda)}{d\Omega} = 5.45 \cdot \left(\frac{550}{\lambda}\right)^4 \times 10^{-32} \left(m^2 s r^{-1}\right)$$

Elastic and Inelastic Scattering (cont'd)

Mie (Aerosol/Cloud) Scattering

In lidar field, Mie scattering is referred to the elastic scattering from spherical particles whose size is comparable to or larger than the wavelength of the laser.

Furthermore, Mie scattering is generalized to elastic scattering from overall aerosol particles and cloud droplets, i.e., including non-spherical particles.

Polarization in Scattering

Backscattering from spherical particles does not change the polarization state of the radiation (Mie). The backscattered light has polarization parallel to that of the transmitted beam (usually linearly polarized).

Large non-spherical particles lead to a depolarization of backscattered radiation, i.e., partial backscattered light has polarization perpendicular to that of the transmitted beam.

And hence the range-resolved linear depolarization ratio is:

$$\delta(R) = \left[\beta_{\perp}(R) / \beta_{\parallel}(R)\right] \exp(T_{\parallel} - T_{\perp})$$

Elastic and Inelastic Scattering (cont'd)

Raman Scattering

Inelastic scaterring process involves the change of the vibrationalrotational energy level of the molecule.

Frequency shift corresponds to energy difference between initial and final molecular states and hence is specific for the interacting molecule.

Intensity distn. within the Raman bands contains information on temperature in scattering volume (energy levels population follows Boltzamann's distn. law)



Scattering Form of Lidar Equation

Rayleigh, Mie, and Raman scattering processes are instantaneous scattering processes, so there are no finite relaxation effects involved, but infinitely short duration.

□ For Rayleigh and Mie scattering, there is no frequency shift when the atmospheric particles are at rest. The lidar equation is written as

$$N_{S}(\lambda, R) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\beta(\lambda, R)\Delta R\right) \left(\frac{A}{R^{2}}\right) T^{2}(\lambda, R) \left(\eta(\lambda)G(R)\right) + N_{B}$$

□ For Raman scattering, there is a large frequency shift. Raman lidar equation may be written as

$$N_{S}(\lambda,R) = \left(\frac{P_{L}(\lambda_{L})\Delta t}{hc/\lambda_{L}}\right) \left(\beta(\lambda,\lambda_{L},R)\Delta R\right) \left(\frac{A}{R^{2}}\right) \left(T(\lambda_{L},R)T(\lambda,R)\right) \left(\eta(\lambda,\lambda_{L})G(R)\right) + N_{B}(\lambda,\lambda_{L})G(R) + N_{B}(\lambda,\lambda_{L})G(R)\right) + N_{B}(\lambda,\lambda_{L})G(R) + N_{B}(\lambda$$

(2) Differential Absorption

In the lidar equation for DIAL, the influence of molecular species is in the extinction (atmosphere transmission) part, not in the backscatter part of the lidar equation.

Scattering of laser light by air molecules and aerosols constitutes a distributed Reflector of the Laser beam.



Differential Absorption/Scattering Form of Lidar Equation

 \square For the laser with wavelength $\lambda_{\mbox{\tiny on}}$ on the molecular absorption line

$$N_{S}(\lambda_{on}, R) = N_{L}(\lambda_{on}) \Big[\beta_{sca}(\lambda_{on}, R) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{z} \overline{\alpha}(\lambda_{on}, r') dr' \Big] \\ \times \exp \Big[-2 \int_{0}^{z} \sigma_{abs}(\lambda_{on}, r') n_{c}(r') dr' \Big] \Big[\eta(\lambda_{on}) G(R) \Big] + N_{B}$$

 \square For the laser with wavelength λ_{off} off the molecular absorption line

$$\begin{split} N_{S}(\lambda_{off},R) &= N_{L}(\lambda_{off}) \Big[\beta_{sca}(\lambda_{off},R) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{z} \overline{\alpha}(\lambda_{off},r') dr' \Big] \\ &\times \exp \Big[-2 \int_{0}^{z} \sigma_{abs}(\lambda_{off},r') n_{c}(r') dr' \Big] \Big[\eta(\lambda_{off}) G(R) \Big] + N_{B} \end{split}$$

Differential absorption cross-section

$$\Delta \sigma_{abs}(R) = \sigma_{abs}(\lambda_{ON}, R) - \sigma_{abs}(\lambda_{OFF}, R)$$

(3) Laser Induced Fluorescence

Strong intermolecular interactions in solids and liquids cause broad absorption and emission spectra.

A fixed frequency laser can be used for the excitation due to the broad absorption.

Following the excitation, a very fast (ps) radiationless relaxation down to the lowest sub-level of the excited state, where the molecules remain for a typical excited state (fluorescence lifetime).

A decay to different sublevels of the ground state gives rise to a distribution of fluorescence light, which reflect the lower-state level distribution.

1- Fixing the excitation wavelength, we can obtain fluorescence spectra. 2- While fixing the detection channel and varying the excit. wavelength, an excitation spectrum can be recorded.



(4) Resonance Fluorescence

Resonance fluorescence is obtained if energy of incoming photon coincides with energy of a transition in an atom, ion, or molecule from one level into another energy level.

In the middle and upper atmosphere, there exist some metal atoms (Na) and atomic ions with large absorption cross section. Quenching is not a problem in that region.

Therefore, laser tuned to the resonance frequency of the absorption lines can excite resonance fluorescence from these atoms and ions.


Fluorescence Form of Lidar Equation

The lidar equation in fluorescence form is given by

$$N_{S}(\lambda,R) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{eff}(\lambda,R)n_{c}(z)R_{B}(\lambda)\Delta R\right) \left(\frac{A}{4\pi R^{2}}\right) \left(T_{a}^{2}(\lambda,R)T_{c}^{2}(\lambda,R)\right) \left(\eta(\lambda)G(R)\right) + N_{B}(\lambda)G(R)$$

 \Box Here, T_c(R) is the transmission caused by the constituent absorption.

$$T_c(R) = \exp\left(-\int_{R_{bottom}}^R \sigma_{eff}(\lambda, r') n_c(r') dr'\right) = \exp\left(-\int_{R_{bottom}}^R \alpha_c(\lambda, r') dr'\right)$$

 \Box Here, $\alpha(\lambda,R)$ is the extinction coefficient caused by the absorption.

$$\alpha_c(\lambda, R) = \sigma_{eff}(\lambda, R) n_c(R)$$

Backscatter Cross-Section Comparison

Physical Process	Backscatter Cross-Section	Mechanism
Mie (Aerosol) Scattering	10-8 - 10-10 cm ² sr-1	Two-photon process
		Elastic scattering, instantaneous
Resonance Fluorescence	10 ⁻¹³ cm ² sr ⁻¹	Two single-photon process (absorption and spontaneous emission)
		Delayed (radiative lifetime)
Molecular Absorption	10 ⁻¹⁹ cm ² sr ⁻¹	Single-photon process
Fluorescence from	10 ⁻¹⁹ cm ² sr ⁻¹	Two single-photon process
molecule, liquid, solid		Inelastic scattering, delayed (lifetime)
Rayleigh Scattering	10 ⁻²⁷ cm ² sr ⁻¹	Two-photon process
		Elastic scattering, instantaneous
Raman Scattering	10-30 cm ² sr-1	Two-photon process
		Inelastic scattering, instantaneous

(5) Doppler Shift and Broadening

Doppler Technique – Doppler linewidth broadening and Doppler frequency shift are temperature-dependent and wind-dependent, respectively (applying to both Na, K, Fe resonance fluorescence and molecular scattering)



(6) Boltzmann Distribution

Boltzmann distribution is the law of particle population distribution according to energy levels



 N_1 and N_2 - particle populations on energy levels E_1 and E_2 g₁ and g₂ - degeneracy for energy levels E_1 and $\mathsf{E}_2, \Delta\mathsf{E} = \mathsf{E}_2 - \mathsf{E}_1$ $k_{\scriptscriptstyle B}$ - Boltzmann constant, T - Temperature, N - total population

Population Ratio ⇒ **Temperature**

Fe Boltzmann temperature lidar

By measuring the Fe signal level at the two wavelengths, one can derive the temperature provided that the ratios of the extinction and the effective backscatter cross sections are known.

$$T(z) = \frac{\Delta E/k_B}{\ln \left[\frac{g_2}{g_1} \frac{R_{B374}}{R_{B372}} \left(\frac{\lambda_{374}}{\lambda_{372}}\right)^{4.0117} \frac{R_E^2(z)R_\sigma}{R_T(z)}\right]} \\ = \frac{598.44}{\ln \left[\frac{0.7221R_E^2(z)R_\sigma}{R_T(z)}\right]}$$
(K).

Where

$$\begin{split} R_{E}(z) &= \frac{E(\lambda_{374}, z)}{E(\lambda_{372}, z)}, \\ R_{\sigma} &= \frac{\sigma_{\text{eff}}(\lambda_{374}, T, \sigma_{L374})}{\sigma_{\text{eff}}(\lambda_{372}, T, \sigma_{L372})} \end{split}$$

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Solutions of Some Lidar Equations

Solution for scattering form lidar equation
 Solution for fluorescence form lidar equation
 Solution for differential absorption lidar equation
 Solution for resonance fluorescence lidar

Solution for Scattering Form Lidar Equation

Scattering form lidar equation

$$N_{S}(\lambda,R) = \left[\frac{P_{L}(\lambda_{L})\Delta t}{hc/\lambda_{L}}\right] \cdot \left[\beta(\lambda,\lambda_{L},R)\Delta R\right] \cdot \left[\frac{A}{R^{2}}\right] \cdot \left[T(\lambda_{L},R)T(\lambda,R)\right] \cdot \left[\eta(\lambda,\lambda_{L})G(R)\right] + N_{B}$$

Solution for scattering form lidar equation

$$\beta(\lambda,\lambda_L,R) = \frac{N_S(\lambda,R) - N_B}{\left[\frac{P_L(\lambda_L)\Delta t}{hc/\lambda_L}\right] \Delta R \left(\frac{A}{R^2}\right) \left[T(\lambda_L,R)T(\lambda,R)\right] \left[\eta(\lambda,\lambda_L)G(R)\right]}$$

Solution for Fluorescence Form Lidar Equation

Fluorescence form lidar equation

$$N_{S}(\lambda,R) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{eff}(\lambda,R)n_{c}(R)R_{B}(\lambda)\Delta R\right) \left(\frac{A}{4\pi R^{2}}\right) \left(T_{a}^{2}(\lambda,R)T_{c}^{2}(\lambda,R)\right) \left(\eta(\lambda)G(R)\right) + N_{B}(\lambda)G(R)$$

Solution for fluorescence form lidar equation

$$n_{c}(R) = \frac{N_{S}(\lambda, R) - N_{B}}{\left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{eff}(\lambda)R_{B}(\lambda)\Delta R\right) \left(\frac{A}{4\pi R^{2}}\right) \left(\eta(\lambda)T_{a}^{2}(\lambda, R)T_{c}^{2}(\lambda, R)G(R)\right)}$$

Differential Absorption/Scattering Form

 $\hfill \label{eq:laser}$ For the laser with wavelength λ_{on} on the molecular absorption line

$$\begin{split} N_{S}(\lambda_{on},R) &= N_{L}(\lambda_{on}) \big[\beta_{sca}(\lambda_{on},R) \Delta R \big] \bigg(\frac{A}{R^{2}} \bigg) \exp \Big[-2 \int_{0}^{R} \overline{\alpha}(\lambda_{on},r') dr' \Big] \\ &\times \exp \Big[-2 \int_{0}^{R} \sigma_{abs}(\lambda_{on},r') n_{c}(r') dr' \Big] \big[\eta(\lambda_{on}) G(R) \big] + N_{B} \end{split}$$

 $\hfill \label{eq:laser}$ For the laser with wavelength λ_{off} off the molecular absorption line

$$\begin{split} N_{S}(\lambda_{off},R) &= N_{L}(\lambda_{off}) \Big[\beta_{sca}(\lambda_{off},R) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{R} \overline{\alpha}(\lambda_{off},r') dr' \Big] \\ &\times \exp \Big[-2 \int_{0}^{R} \sigma_{abs}(\lambda_{off},r') n_{c}(r') dr' \Big] \Big[\eta(\lambda_{off}) G(R) \Big] + N_{B} \end{split}$$

Differential Absorption/Scattering Form

The ratio of photon counts from these two channels is a function of the differential absorption and scattering:

$$\frac{N_{S}(\lambda_{on},R) - N_{B}}{N_{S}(\lambda_{off},R) - N_{B}} = \frac{N_{L}(\lambda_{on})\beta_{sca}(\lambda_{on},R)}{N_{L}(\lambda_{off})\beta_{sca}(\lambda_{off},R)} \frac{\eta(\lambda_{on})}{\eta(\lambda_{off})} \\ \times \exp\left\{-2\int_{0}^{R} \left[\overline{\alpha}(\lambda_{on},r') - \overline{\alpha}(\lambda_{off},r')\right]dr'\right\} \\ \times \exp\left\{-2\int_{0}^{R} \left[\sigma_{abs}(\lambda_{on},r') - \sigma_{abs}(\lambda_{off},r')\right]n_{c}(r')dr'\right\}$$

$$\Delta \sigma = \sigma_{abs}(\lambda_{on}) - \sigma_{abs}(\lambda_{off})$$

Solution for Differential Absorption Lidar Equation

Solution for differential absorption lidar equation

$$n_{c}(R) = \frac{1}{2\Delta\sigma} \frac{d}{dR} \begin{cases} \ln \left[\frac{N_{L}(\lambda_{on})\beta_{sca}(\lambda_{on},R)}{N_{L}(\lambda_{off})\beta_{sca}(\lambda_{off},R)} \frac{\eta(\lambda_{on})}{\eta(\lambda_{off})} \right] \\ -\ln \left[\frac{N_{S}(\lambda_{on},R) - N_{B}}{N_{S}(\lambda_{off},R) - N_{B}} \right] \\ -2\int_{0}^{R} \left[\overline{\alpha}(\lambda_{on},r') - \overline{\alpha}(\lambda_{off},r') \right] dr' \end{cases}$$

$$\Delta \sigma = \sigma_{abs}(\lambda_{on}) - \sigma_{abs}(\lambda_{off})$$

Solution for Resonance Fluorescence Lidar Equation

■ Resonance fluorescence and Rayleigh lidar equations $N_{S}(\lambda,z) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{eff}(\lambda,z)n_{c}(z)R_{B}(\lambda)\Delta z\right) \left(\frac{A}{4\pi z^{2}}\right) \left(T_{a}^{2}(\lambda)T_{c}^{2}(\lambda,z)\right) \left(\eta(\lambda)G(z)\right) + N_{B}$ $N_{R}(\lambda,z_{R}) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{R}(\pi,\lambda)n_{R}(z_{R})\Delta z\right) \left(\frac{A}{z_{R}^{2}}\right) T_{a}^{2}(\lambda,z_{R}) \left(\eta(\lambda)G(z_{R})\right) + N_{B}$

Rayleigh normalization

$$\frac{n_c(z)}{n_R(z_R)} = \frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2} \cdot \frac{4\pi\sigma_R(\pi, \lambda)}{\sigma_{eff}(\lambda, z)R_B(\lambda)} \cdot \frac{T_a^2(\lambda, z_R)G(z_R)}{T_a^2(\lambda, z)T_c^2(\lambda, z)G(z)}$$

Solution for resonance fluorescence

$$n_c(z) = n_R(z_R) \frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2} \cdot \frac{4\pi\sigma_R(\pi, \lambda)}{\sigma_{eff}(\lambda, z)R_B(\lambda)} \cdot \frac{1}{T_c^2(\lambda, z)}$$

Solution for Rayleigh and Mie Lidars

Rayleigh and Mie (middle atmos) lidar equations

$$N_{S}(\lambda,z) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\beta_{R}(z) + \beta_{aerosol}(z)\right) \Delta z \left(\frac{A}{z^{2}}\right) T_{a}^{2}(\lambda,z) \left(\eta(\lambda)G(z)\right) + N_{B}$$
$$N_{R}(\lambda,z_{R}) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\beta_{R}(z_{R})\Delta z\right) \left(\frac{A}{z_{R}^{2}}\right) T_{a}^{2}(\lambda,z_{R}) \left(\eta(\lambda)G(z_{R})\right) + N_{B}$$

Rayleigh normalization

$$\frac{\beta_R(z) + \beta_{aerosol}(z)}{\beta_R(z_R)} = \frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2} \cdot \frac{T_a^2(\lambda, z_R)G(z_R)}{T_a^2(\lambda, z)G(z)}$$

For Rayleigh scattering at z and z_R

$$\frac{\beta_R(z)}{\beta_R(z_R)} = \frac{\sigma_R(z)n_{atm}(z)}{\sigma_R(z_R)n_{atm}(z_R)} = \frac{n_{atm}(z)}{n_{atm}(z_R)}$$

Solution (Continued)

Solution for Mie scattering in middle atmosphere

$$\beta_{aerosol}(z) = \beta_R(z_R) \left[\frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2} - \frac{n_{atm}(z)}{n_{atm}(z_R)} \right]$$

$$\beta_R(\lambda, z_R, \pi) = 2.938 \times 10^{-32} \frac{P(z_R)}{T(z_R)} \cdot \frac{1}{2^{4.0117}} \left(m^{-1} s r^{-1} \right)$$

Rayleigh normalization when aerosols not present

$$\frac{\beta_R(z)}{\beta_R(z_R)} = \frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2} \cdot \frac{T_a^2(\lambda, z_R)G(z_R)}{T_a^2(\lambda, z)G(z)}$$

 $T(z_R) \ \overline{\lambda^{4.0117}} \langle^m$

Solution for relative number density in Rayleigh lidar

$$RND(z) = \frac{n_{atm}(z)}{n_{atm}(z_R)} = \frac{\beta_R(z)}{\beta_R(z_R)} = \frac{N_S(\lambda, z) - N_B}{N_R(\lambda, z_R) - N_B} \cdot \frac{z^2}{z_R^2}$$

Review of Lidar Architecture

Basic Lidar Architecture

Configurations vs. Arrangements
 Lidar Classifications

Physical Picture in Lidar Equation



Basic Architecture of LIDAR



Transmitter (laser pulses that meet some requirements): wavelength, frequency accuracy, bandwidth, pulse duration time, pulse energy, repetition

rate, divergence angle, etc

Receiver (receiver is to collect and detect returned photon signals while compressing background noise): it consists of telescopes, filters, collimating optics, photon detectors, discriminators, etc.

Data acquisition and control system (record returned data and corresponding time-offlight, provide system control and coordination to transmitter and receiver.): it usually consists of multi-channel scaler which has very precise clock so can record time precisely, discriminator, computer and software.

Basic Configurations of LIDAR Bistatic and Monostatic



Monostatic Configuration

Bistatic Configuration

Monostatic Configuration Example



Coaxial vs. Biaxial Arrangements

□ In a coaxial system, the axis of the laser beam is coincident with the axis of the receiver optics.

In the biaxial arrangement, the laser beam only enters the field of view of the receiver optics beyond some predetermined range.

Biaxial arrangement helps avoiding near-field backscattered radiation saturating photo-detector.

The near-field backscattering problem in a coaxial system can be overcome by either gating of the photo-detector or use of a fast shutter or chopper.

Example of Biaxial Arrangement



Example of Coaxial Arrangement



Classifications of Lidar

There are several different classifications on lidars e.g., based on the physical process; (Mie, Rayleigh, Raman, Res. Fluorescence, ...) based on the platform; (Groundbased, Airborne, Spaceborne, ...) based on the detection region; (Atmosphere, Ocean, Solid Earth, Space, ...) based on the emphasis of signal type; (Ranging, Scattering, ...) based on the topics to detect; (Aerosol, Constituent, Temp, Wind, Target, ...)

••• •••

Classification on Physical Process



Classification on Platform

Spaceborne lidar	Satellite, Space Shuttle. Space Station		
Airborne lidar	Jet, Propeller Airplanes Unmanned Aerial Vehicle (UAV) Kite		
Groundbased lidar	Stationary Contanerized moved with truck		
Shipborne lidar	Icebreaker, Ships		
Submarine lidar	Submarine		

Classification on Detection Regions

Atmosphere lidar

Various types From various platforms

Hydrosphere lidar

Various types From various platforms

Solid Earth lidar

Airborne or Spaceborne Laser altimeter

Target lidar

Various type With or without Imaging function

LIDAR 1 & 2

Taieb Gasmi College of Science & Engineering Saint Louis University in Spain



International Centre for Theoretical Physics (ICTP) International College on Optics in Environmental Science

LIDAR 2: Classes of LIDARS

- 1. Temperature LIDAR
- 2. Wind LIDAR (review)
- 3. Aerosol LIDAR
- 4. Constituent LIDAR

Temperature Measurement LIDAR

Use temperature-dependent effects or phenomena

Doppler Technique - Doppler broadening (not only for Na, K, and Fe, but also for Rayleigh scattering, as long as Doppler broadening dominate and can be detected)

Boltzmann Technique – Boltzmann distribution of atomic populations on different energy levels (not only for Fe, but also for molecular spectroscopy in optical remote sensing)

□ Integration Technique (Rayleigh or Raman) – integration lidar technique using ideal gas law and assuming hydrostatic equilibrium (not only for modern lidar, but also for cw searchlight and rocket falling sphere – some way to measure atmosphere number density)

Rotational Raman Technique – temperature dependence of population ratio, similar to Boltzmann technique

Boltzmann Distribution

Boltzmann distribution is the law of particle population distribution according to energy levels



 N_1 and N_2 - particle populations on energy levels E_1 and E_2 g₁ and g₂ - degeneracy for energy levels E_1 and $\mathsf{E}_2, \Delta\mathsf{E} = \mathsf{E}_2 - \mathsf{E}_1$ $k_{\scriptscriptstyle B}$ - Boltzmann constant, T - Temperature, N - total population

Population Ratio ⇒ **Temperature**



Temperature Measurement Principle of the Fe Boltzmann Lidar Technique

There are four natural isotopes of Fe atoms: 54 Fe, 56 Fe, 57 Fe and 58 Fe. Among them 56 Fe is the most abundant isotope with natural abundance of 91.754%. The isotopic data of Fe atoms are summarized in the following table.

	$^{54}\mathrm{Fe}$	$^{56}\mathrm{Fe}$	57 Fe	$^{58}\mathrm{Fe}$
Z	26	26	26	26
A	54	56	57	58
Nuclear spin	0	0	1/2	0
Natural abundance	5.845%	91.754%	2.119%	0.282%

Table 5.4 Fe Resonance Line Parameters

Transition wavelength λ	372.0993 nm	373.8194 nm
Degeneracy for ground state	$g_1 = 9$	$g_2 = 7$
Degeneracy for excited state	$g_1' = 11$	$g_2' = 9$
Radiative lifetime of excited state (ns)	61.0	63.6
Einstein coefficient $A_{\rm ki}~(10^8~{ m s}^{-1})$	0.163	0.142
Oscillator strength f_{ik}	0.0413	0.0382
Branching ratio $R_{\rm B}$	0.9959	0.9079
$\sigma_0 \ (10^{-17} \ { m m}^2)$	9.4	8.7



-The isotope shifts of 54 Fe, 57 Fe and 58 Fe with respect to 56 FE are ~ 1 Ghz for the two resonance lines 372 nm and 374 nm.

- Energy difference between the lowest ground states is about 416 cm⁻¹ which is almost 1,25 x10⁴ GHz, which means that the contribution of these isotope shifts (⁵⁴Fe, ⁵⁷Fe and ⁵⁸Fe) to the Boltzmann factor (exp(- $\Delta E/K_BT$) is negligible (less than 1x10⁻⁴).

- The Fe Boltzmann temperature Lidar relies on the unique energy level distribution diagram of the ground state manifold of Fe atom. In thermodynamic equilibrium, the ratio of the populations in the J=3 and J=4 sublevels in the ground state manifold is given by Maxwell-Boltzamnn distribution law.

Boltzmann Technique



p(372) and p(374) are the associated number desities. ΔE is the energy difference between the two level= 416 cm⁻¹. Subscripts 372 and 374 refers to the excitation wavelengths of the two transitions.

The Fe densities at those two states can be measured using resonance fluorescence lidar technique.

The Fe photon count is therefore given by the lidar equation:

$$N_{\rm Fe}(\lambda, z) = \left(\frac{P_{\rm L}\Delta t T_{\rm a}}{hc/\lambda} E(\lambda, z)\right) \sigma_{\rm eff}(\lambda, T, \sigma_{\rm L}) R_{\rm B\lambda} \rho_{\rm Fe}(\lambda, z) \\ \times \Delta z \left(E(\lambda, z) T_{\rm a} \frac{A_{\rm R}}{4\pi z^2} \eta\right)$$
(5.92)
RFLE

- Z is the altitude; P_L laser power; ΔT temporal resolution; ΔZ is the vertical resolution; T_a is the lower transmitance of the atmosphere; $E(\lambda.z)$ is the extinction coefficient of the signal associated with the Fe absorption; $\sigma_{eff}(\lambda, T, \sigma_L)$ is the effective scattering cross-section; R_B is the branching ratio.

- The system uses two separate lasers because of the two Fe lines which are far apart in λ to be tuned easily with a single laser.

- To compensate for the signal variations in each of these systems, number densities of Fe layers at each wavelength are normalized by the Rayleigh count (N_R) over an altitude range of 50 Km.

$$N_{R}(\lambda, z_{R}) = \left(\frac{P_{L}(\lambda)\Delta t}{hc/\lambda}\right) \left(\sigma_{R}(\pi, \lambda)n_{R}(z_{R})\Delta z\right) \left(\frac{A}{z_{R}^{2}}\right) T_{a}^{2}(\lambda, z_{R}) \left(\eta(\lambda)G(z_{R})\right)$$

$$70$$

The Rayleigh-normalized background-corrected Fe signal count is hence defined:

$$egin{aligned} N_{ ext{norm}}(\lambda,z) &= rac{N_{ ext{Fe}}(\lambda,z) + N_{ ext{B}}(\lambda,z) - \hat{N}_{ ext{B}}(\lambda)}{N_{ ext{R}}(\lambda,z_{ ext{R}}) + N_{ ext{B}}(\lambda,z_{ ext{R}}) - \hat{N}_{ ext{B}}(\lambda)} \ &= rac{z_{ ext{R}}^2 E^2(\lambda,z) R_{ ext{B}\lambda} \sigma_{ ext{eff}}(\lambda,T,\sigma_{ ext{L}})
ho_{ ext{Fe}}(\lambda,z)}{z^2 \sigma_{ ext{R}}(\lambda)
ho_{ ext{atmos}}(z_{ ext{R}})} \end{aligned}$$

 $N^B(\lambda)$ is the estimated background noise count.

The Fe densities p(372) and p(374) at each wavelength can be derived from the last equation.

The ratio of the normalised signal counts at two wavelengths (R_T) can be related to the temperature as follows:
$$T(z) = rac{\Delta E/\!k_{
m B}}{\ln\left[\!rac{g_2}{g_1}rac{R_{
m B374}}{R_{
m B372}}\!\left(\!rac{\lambda_{374}}{\lambda_{372}}\!
ight)^{4.0117}rac{R_E^2(z)R_\sigma}{R_{
m T}(z)}
ight]}$$

At 200 K, (372nm) Fe signal is ~ 30 times stronger than (374nm) Fe signal so $R_T \sim 1/30$. So the final result is:



Thus, by measuring the Fe signal level at the two wavelengths and computing RT(z), one can derive the temperature, provided that the ratios of the extinction and the effective backscatter cross sections are known.



Fig. 6. (a) Temperature profiles measured by Fe Boltzmann temperature lidar over the North Pole on 21 June 1999 and over the South Pole on 24 December 2000 and 29 December 2000 along with MSISE-00 model data for 21 June 1999 at the North Pole. (b) The 372-nm ground-state Fe density profiles corresponding to the temperature profiles plotted in Fig. 6(a).

$$T = \frac{\Delta E / k_B}{\ln \left[\frac{g_2}{g_1} \frac{\rho_{Fe}(372)}{\rho_{Fe}(374)}\right]}$$

$$\rho_{Fe}^{372} \text{ and } \rho Fe^{374} \text{ are the associated Fe number densities, } \Delta E \text{ is the energy difference between the two levels}$$

$$Fe Boltzmann Temperature Lidar System$$

$$Fe Boltzmann System$$

$$Fe Boltzmann System$$

$$Fe Boltzmann System$$

$$Fe Boltz$$

Diagram of Univ. of Illinois Fe Boltzmann Temperature Lidar System J.Appl. Opt. 41, 2002.

Fe Boltzmann Lidar Schematic



Based on injection-seeded, frequency-doubled, pulse Alexandrite laser systems (372 and 374 nm output)

Fe Lidar Seeder Laser System

Seeder Laser Structure and Wavelength Control



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Aerosol Lidar

Motivations to Study Aerosols

Atmospheric aerosols play an important role in many atmospheric processes: - Aerosols have appreciable influence on the Earth's radiation budget, air quality and visibility, clouds, precipitation, and chemical processes in the troposphere and stratosphere.

Their occurrence, residence time, physical properties, chemical composition as well as the resulting climate-relevant optical properties are subject to large diversity especially in the troposphere because of their widely different sources and meteorological processes.

Therefore, vertically resolved measurements of physical and optical properties of particles such as the mean particle size, and the volume extinction coefficient are of great interest.

Routine (long-term), range-resolved observations of these parameters can only be carried out with lidar.

(3) Aerosol LIDAR: <u>Elastic-Scattering</u>, Raman, HSRL

Elastic-Scattering for Lower Atmos

Lidar equation for elastic backscatter by air molecules and aerosols

$$P(R) = \frac{E_o \eta}{R^2} O(R) \beta(R) \exp\left[-2\int_0^R \alpha(r) dr\right]$$

O(R) - overlap between laser and FOV $\beta(R)$ - backscatter coefficient (km⁻¹sr⁻¹)

 $\alpha(R)$ - extinction coefficient (km⁻¹)

Backscatter and extinction are contributed by aerosols and molecules

$$\begin{split} \beta(R) &= \beta_{aer}(R) + \beta_{mol}(R) \\ \alpha(R) &= \alpha_{aer}(R) + \alpha_{mol}(R) \end{split}$$

Index aer refer to aerosol particles Index mol refer to air molecules

Combining above equations, we obtain

$$P(R)R^{2} = E_{0}\eta_{L} \Big[\beta_{aer}(R) + \beta_{mol}(R)\Big] \exp \left[-2\int_{o}^{R} \Big[\alpha_{aer}(r) + \alpha_{mol}(r)\Big]dr\right]$$

Note: backscatter is local effect while extinction is integrated effect \Box Recall the Rayleigh scattering from air molecules $\beta_{mol}(R)$ can be determined from atmosphere temperature and pressure

$$\beta_{mol}(\lambda, z_R) = 2.938 \times 10^{-32} \frac{P(z_R)}{T(z_R)} \cdot \frac{1}{\lambda^{4.0117}} \qquad z_R \text{ is altitude for range R}$$

In general, both the volume extinction coefficient a and backscattering coefficient β_T are unknown. it is therefore necessary to assume some kind of relation between the extinction a and the backscattering coefficient β_T (extinction-to-backscattering ratio); where the volume backscaterring coefficient is $\beta = \beta_T / 4\pi$ (m⁻¹.Sr⁻¹)

Extinction-to-backscattering ratio:

Let S(R)= Ln[R².P(R)] and let S(Ro) be the signal at the reference range R₀, we have: S(R) -S(RO)= Ln[$\beta(R)/\beta(R_0)$]-2 $\int_{R_0}^{r} \alpha dr$

In differential form:

 $dS(R)/d(R)=1/\beta(R).d\beta(R)/dR - 2\alpha(R)$

Slope method of inversion: assumes that the scatterers are homogeneously

Distributed along the lidar path so $d\beta(R)/dR \sim 0$

Thus $dS/dR = 2\alpha(R)$ is estimated from the slope of the plot S vs. R Limitations: applicable for a homogeneous path only.

Lidar Detection of Aerosols/Clouds

Aerosols and clouds are shown as distinct peaks above the Rayleigh scattering background in range-resolved lidar profiles.

The common way to detect aerosols and clouds is to use elastic scattering lidar with Rayleigh and Mie scattering detection capability.

Virtually this can be done by any lidars that receive scattering signals, including resonance fluorescence, DIAL, Raman, Rayleigh, & Mie.



Polarization Lidar Detection

Backscattering from a spherical particle does not change the polarization, but the non-spherical particle changes the polarization. Thus, by monitoring the polarization status of the scattered light, information on the shape of the aerosol particles can be obtained.





(2) Wind Lidar Motivations for Wind Measurements

Global atmospheric wind profiles from ground to 120 km are important for validation of the output of global atmosphere models, and for study of the atmosphere dynamics, as wave information can be inferred from the wind measurements.

Wind/velocity measurements have much more applications in industry, environment, and defense business. For example, (1) Aircraft true airspeed, aircraft wake vortices

- (2) Clear air turbulence, wind shear, gust fronts
- (3) Air pollution monitoring
- (4) Vibration of objects
- (5) Laboratory, machine shop, production facility, wind tunnel

Techniques for Wind Measurements

Use wind-dependent effects or use definition of wind

 Direct Motion Detection Technique: (using the definition of velocity)

$$\vec{v} = \frac{d\vec{r}(t)}{dt}$$

(1) Tracking aerosol/cloud motions
 (2) Laser Time-of-Flight Velocimetry
 (3) Laser Doppler Velocimetry

For non-resonant scattering

The Doppler frequency shift is given by

$$\Delta \omega_{scattering} = \omega_2 - \omega_1 = -(\vec{k}_1 \cdot \vec{\mathbf{v}}_1 - \vec{k}_2 \cdot \vec{\mathbf{v}}_2)$$

□ Therefore, for forward scattering, $\vec{k}_2 \approx \vec{k}_1, \vec{v}_2 \approx \vec{v}_1$, so $\Delta \omega = 0$

D For backward scattering,
$$\vec{k}_2 \approx -\vec{k}_1, \vec{v}_2 \approx \vec{v}_1$$
, so $\Delta \omega = -2\vec{k}_1 \cdot \vec{v}_1$

(4) Constituent Lidar

 Motivations to study atmosphere constituents
 Lidar detection of atmospheric constituents (spectroscopic signatures to distinguish species)
 Metal atoms by resonance fluorescence lidar
 DIAL detection of molecules and pollutant (DIAL Equation and Solution)
 Raman Lidar detection of molecules and pollutant (Raman Equation and Solution)

Motivations to Study Constituents

Atmospheric constituents are referred to the trace gases that occur in relatively small but sometimes highly variable concentrations. In our class, constituents mean the atomic or molecular trace gases in the atmosphere, excluding aerosols and clouds.



Molecule	Typical Wavelength	Laser (Examples)	Absorption Cross Sectio (10 ⁻¹⁸ cm ²)
Nitric oxide, NO	226.8 nm	Dye	4.6
Benzene, C ₆ H ₆	252.9 nm	Dye	5.7
Mercury, Hg	253.65 nm	Dye	3.3×10^4
Toluene, C ₇ H ₈	266.9 nm	Dye, Ti:Sa	2.8
Ozone, O ₃	266.0 nm	Dye, Nd:YAG-IV	9.49
	289.0 nm	Raman D ₂	1.59
	299.0 nm	Raman H ₂	0.42
Formaldehyde, CH ₂ O	286.5 nm	Dye, Ti:Sa	0.068
Sulfur dioxide, SO ₂	300.0 nm	Dye, Ti:Sa	1.3
Chlorine, Cl ₂	$330.0\mathrm{nm}$	Dye, XeCl + Raman cell	0.26
Nitrous acid, HONO	354.0 nm	Dye, Ti:Sa	0.50
Nitrogen dioxide, NO ₂	448.1 nm	Dye	0.69
Methane, CH4	$3.270 \mu m$	OPO	2.0
	$3.391 \mu m$		0.6
Propane, C ₃ H ₈	$3.391 \mu m$	OPO	0.8
Hydrogen chloride, HCl	$3.636 \mu m$	DF, OPO	0.20
Methane, CH ₄	$3.715 \mu m$	DF, OPO	0.002
Sulfur dioxide, SO ₂	$3.984 \mu m$	DF, OPO	0.42
Carbon monoxide, CO	$4.709 \mu m$	CO_2	2.8 Ta
	$4.776\mu\mathrm{m}$		0.8 at
Nitric oxide, NO	$5.215 \mu m$	CO	0.67 Th
	$5.263 \mu m$	CO_2	0.6
Propylene, C ₃ H ₆	$6.069 \mu m$	CO	0.09
1,3-Butadiene, C ₄ H ₆	$6.215 \mu m$	CO	0.27
Nitrogen dioxide, NO ₂	$6.229 \mu m$	CO	2.68
Sulfur dioxide, SO ₂	$9.024 \mu{ m m}$	CO_2	0.25 M
Freon-11, CCl ₃ F	$9.261 \mu m$	CO_2	1.09
Ozone, O ₃	$9.505 \mu m$	CO_2	0.45 Tr
	$9.508 \mu m$		0.9 1
Fluorocarbon-113, C ₂ Cl ₃ F ₃	9.604 µm	CO_2	0.77 11
Benzene, C ₆ H ₆	$9.621 \mu m$	CO_2	0.07
MMH, CH ₃ N ₂ H ₃	$10.182 \mu m$	CO_2	0.06 V1
Ethyl mercaptan, C ₂ H ₅ SH	$10.208\mu m$	CO_2	0.02 U
Chloroprene, C ₄ H ₅ Cl	$10.261 \mu{\rm m}$	CO_2	0.34 Fl
Monochloroethane, C2H5Cl	$10.275\mu m$	CO_2	0.12 Pc
Ammonia, NH ₃	$10.333 \mu m$	CO_2	1.0
Ethylene, C ₂ H ₄	$10.533 \mu m$	CO_2	1.19 1
Sulfur hexafluoride, SF ₆	$10.551 \mu m$	CO_2	30.3 Pe

Table 4.1 Absorption Cross Section of Pollutant Gases at Wavelengths Suitable for DIAL Measurement in the Troposphere

Pollutant Gases in Earth Atmosphere

Note: the absorption cross-sections of molecules are about 6 orders of magnitude smaller than the metal atoms such as Na, Fe, K.

Yable 4.1 Absorption Cross Section of Pollutant Gases t Wavelengths Suitable for DIAL Measurement in the proposphere—(Continued)

Molecule	Typical Wavelength	Laser (Examples)	$\begin{array}{c} {\rm Absorption}\\ {\rm Cross} \ {\rm Section}\\ (10^{-18}{\rm cm}^2) \end{array}$
Trichloroethylene, C ₂ HCl ₃	$10.591\mu\mathrm{m}$	CO_2	0.49
1,2-Dichloroethane, C ₂ H ₄ Cl ₂	$10.591 \mu m$	CO_2	0.02
Hydrazine, N ₂ H ₄	$10.612 \mu m$	CO_2	0.18
Vinyl chloride, C ₂ H ₃ Cl	$10.612 \mu m$	CO_2	0.33
UDMH, $(CH_3)_2N_2H_2$	$10.696 \mu m$	CO_2	0.08
Fluorocarbon-12, CCl ₂ F ₂	$10.719 \mu m$	CO_2	1.33
Perchloroethylene, C ₂ Cl ₄	$10.742 \mu m$	CO_2	0.18
1-Butene, C ₄ H ₈	$10.787 \mu m$	CO_2	0.13
Perchloroethylene, C ₂ Cl ₄	$10.834 \mu{ m m}$	CO_2	1.14

Differential Absorption Lidar

□ For the molecular species to be detected by DIAL, usually they have strong absorption, however, nearly none resonance fluorescence due to strong relaxation processes other than fluorescence (e.g., frequent collisions with surrounding atmosphere molecules can make molecules decay from excited states to ground state without giving fluorescence).

□ Thus, in the lidar equation for DIAL, the influence of molecular species is in the extinction (atmosphere transmission) part, not in the backscatter part. In other words, the molecular absorption contributes to the extinction of light when incident light and scattered light propagate through atmosphere, while the return signals are from the scattering of laser light by air molecules and aerosols.



Illustration of the DIAL Concept



<u>Case I:</u> DIAL Elastic scattering from diffuse reflector (Mie & Rayleigh Scattering)

Elastic scattering is when the scattering frequency is the same as the frequency of the incident light (e.g.,Rayleigh scattering and Mie scattering)

$$N(R) = \frac{1}{2\Delta\sigma} \left[\frac{d}{dR} \left\{ \ln\left[\frac{P(\lambda_{off}, R)}{P(\lambda_0, R)}\right] - \ln\left[\frac{\beta(\lambda_{off}, R)}{\beta(\lambda_0, R)}\right] \right\} + \overline{\kappa}(\lambda_{off}, R) - \overline{\kappa}(\lambda_0, R) \right]$$

Where



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<u>Case II</u>: Elastic scattering from a topographical target

$$\int_{0}^{R_{T}} N(R) dR = \frac{1}{2\Delta\sigma} \left[\ln \left[\frac{E(\lambda_{off}, R_{T})}{E(\lambda_{0}, R_{T})} \right] + \overline{\kappa}(\lambda_{off}, R) - \overline{\kappa}(\lambda_{0}, R) \right]$$

Lidar Equation for DIAL

DIAL lidar equation

$$P_{S}(\lambda,R) = P_{L}(\lambda) \Big[\beta_{scatter}(\lambda,R) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{R} \overline{\alpha}(\lambda,r) dr \Big] \\ \times \exp \Big[-2 \int_{0}^{R} \sigma_{abs}(\lambda,r) n_{c}(r) dr \Big] \Big[\eta(\lambda) G(R) \Big] + P_{B} \Big]$$

Extinction caused by interested constituent absorption

Extinction caused by other molecules and aerosols

□ Compared to resonance fluorescence, the main difference in DIAL is that the backscatter coefficient is from the elastic-scattering from air molecules and aerosols, not from the fluorescence of interested molecules.

 $\beta_{Scatter}(\lambda,R) = \beta_{aer}(\lambda,R) + \beta_{mol}(\lambda,R)$

The parameter in the DIAL equation

$$\overline{\alpha}(\lambda,r) = \alpha_{aer}(\lambda,r) + \alpha_{mol}(\lambda,r) + \sigma_{IG}(\lambda,r)n_{IG}(r)$$

Aerosol Extinction Air Molecule Extinction Interference gas absorption

DIAL Equation Cont'd

□ Notice that the constituent information is already included in the extinction part. If the lidar parameters and aerosol/air molecule backscatter and extinction are known, we can derive the constituent number density from the lidar equation directly.

□ Unfortunately, most of these parameters are unknown. So the DIAL technique is to utilize two wavelengths - one tuned to strong molecular absorption and another one detuned off the absorption line. Thus, most of the parameters can be cancelled or estimated to a much better degree, allowing the constituent information to be derived more truthfully.

$$P_{S}(\lambda_{ON}, R) = P_{L}(\lambda_{ON}) \Big[\left(\beta_{aer}(\lambda_{ON}, R) + \beta_{mol}(\lambda_{ON}, R) \right) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{R} \left(\alpha_{aer}(\lambda_{ON}, r) + \alpha_{mol}(\lambda_{ON}, r) \right) dr \\ \times \exp \Big[-2 \int_{0}^{R} \sigma_{IG}(\lambda_{ON}, r) n_{IG} dr \Big] \exp \Big[-2 \int_{0}^{R} \sigma_{abs}(\lambda_{ON}, r) n_{c}(r) dr \Big] \Big[\eta(\lambda_{ON}) G(R) \Big] + P_{B}$$

$$P_{S}(\lambda_{OFF}, R) = P_{L}(\lambda_{OFF}) \Big[\Big(\beta_{aer}(\lambda_{OFF}, R) + \beta_{mol}(\lambda_{OFF}, R) \Big) \Delta R \Big] \Big(\frac{A}{R^{2}} \Big) \exp \Big[-2 \int_{0}^{R} \Big(\alpha_{aer}(\lambda_{OFF}, r) + \alpha_{mol}(\lambda_{OFF}, r) \Big) dr \\ \times \exp \Big[-2 \int_{0}^{R} \sigma_{IG}(\lambda_{OFF}, r) n_{IG} dr \Big] \exp \Big[-2 \int_{0}^{R} \sigma_{abs}(\lambda_{OFF}, r) n_{c}(r) dr \Big] \Big[\eta(\lambda_{OFF}) G(R) \Big] + P_{B}$$

Solution for DIAL Equations

To derive solution for DIAL equations, we take the ratio between the ON and OFF wavelength equations and take the logarithm of the ratio:

$$\ln\left[\frac{P_{S}(\lambda_{ON}, R) - P_{B}}{P_{S}(\lambda_{OFF}, R) - P_{B}}\right] = \ln\left[\frac{P_{L}(\lambda_{ON})(\beta_{aer}(\lambda_{ON}, R) + \beta_{mol}(\lambda_{ON}, R))\eta(\lambda_{ON})}{P_{L}(\lambda_{OFF})(\beta_{aer}(\lambda_{OFF}, R) + \beta_{mol}(\lambda_{OFF}, R))\eta(\lambda_{OFF})}\right]$$
$$-2\int_{0}^{R} \left[\alpha_{aer}(\lambda_{ON}, r) - \alpha_{aer}(\lambda_{OFF}, r)\right]dr$$
$$-2\int_{0}^{R} \left[\alpha_{mol}(\lambda_{ON}, r) - \alpha_{mol}(\lambda_{OFF}, r)\right]n_{IG}dr$$
$$-2\int_{0}^{R} \left[\sigma_{IG}(\lambda_{ON}, r) - \sigma_{IG}(\lambda_{OFF}, r)\right]n_{IG}dr$$
$$-2\int_{0}^{R} \left[\sigma_{abs}(\lambda_{ON}, r) - \sigma_{abs}(\lambda_{OFF}, r)\right]n_{c}(r)dr$$

□ Note: several factors have been eliminated from the equation: ΔR , A/R^2 , G(R), as the ON and OFF wavelength laser lights are transmitted and received through the save transmitter and receiver, so share the same geometry factor and the same receiver aperture.

Solution for DIAL Cont'd

Take the differentiation of the equation, we obtain the number density of the constituent that we are interested in

 $\hfill Here,$ the $\Delta\sigma_{abs}$ is called the differential absorption cross-section

$$\Delta \sigma_{abs}(R) = \sigma_{abs}(\lambda_{ON}, R) - \sigma_{abs}(\lambda_{OFF}, R)$$
$$= \sigma_{abs}(\lambda_{ON}) - \sigma_{abs}(\lambda_{OFF})$$

DIAL Solution in Simple Case

□ In the case that the ON and OFF wavelengths have small separation, the laser power, lidar efficiency, aerosol/molecule backscatter coefficient and extinction coefficient are close to be the same for the ON and OFF wavelengths. Therefore, terms B-E are zero. Also, assume there is no interference gas present, so term F is zero. Then we have

$$n_{c}(R) = \frac{1}{2\Delta\sigma_{abs}} \frac{d}{dR} \left\{ \ln \left[\frac{P_{S}(\lambda_{OFF}, R) - P_{B}}{P_{S}(\lambda_{ON}, R) - P_{B}} \right] \right\}$$

In practice, lidar signals are not recorded or analyzed as continuous functions, but rather as values in discrete range bins. Thus, the above solution should be expressed in terms of a range increment ΔR :

$$n_{c}(R) = \frac{1}{2\Delta\sigma_{abs}\Delta R} \ln \left[\frac{P_{S}(\lambda_{OFF}, R + \Delta R) - P_{B}}{P_{S}(\lambda_{ON}, R + \Delta R) - P_{B}} \cdot \frac{P_{S}(\lambda_{ON}, R) - P_{B}}{P_{S}(\lambda_{OFF}, R) - P_{B}} \right]$$

Number density average between range R₁ and R₂

$$n_{c} = \frac{1}{2\Delta\sigma_{abs}\Delta R} \ln \left[\frac{P_{S}(\lambda_{OFF}, R_{2}) - P_{B}}{P_{S}(\lambda_{ON}, R_{2}) - P_{B}} \cdot \frac{P_{S}(\lambda_{ON}, R_{1}) - P_{B}}{P_{S}(\lambda_{OFF}, R_{1}) - P_{B}} \right]$$



Schematic diagram of the Dial system:

1) Photo-detection and signal boosting stage, 2) Signal calibration,

3) Signal digitization, **4)** Data processing and storage, M2-M3: Steering mirrors, BS: Beam splitter, **5)** Topographical target.















View of the 3-D scanner





1- Incoming beam is focussed by lens 1 to an intensity very close to gas breakdown

- 2- High voltage pulse triggers the breakdown3- Plasma once generated absorbs the remaining energy of the laser pulse
- 4- Lens 2 collimates the beam





Parameters	Specifications	
Transmitter		
Laser	Tunable, pulsed CO ₂ -TEA Lumonics 370XT upgraded at 20 Hz	
Pulse width	Unclipped pulse: 3.5 µs Clipped with plasma shutter:<70 ns	
Energy	1.2 J (gain-switched spike at P20)	
Repetition rate	2-20 Hz	
Wavelength	9.2 to 10.8 μm	
Tuning	Scanner computer controlled	
Receiver module		
Telescope	Newtonian φ=40cm. F=183cm, F#/4.5	
Field of view	<5 mrad.	
Detector	EG&G LN cooled: J15-D12	
Data acquisition		
Computer	PC 3 GHz, 1Gb Ram	
Software language	C++	
Digitization	Tektronix TDS 540-200 MHz	
Interfaces	IEEE-488	



Acrosol volume backscattering coefficient as a function of wavelength (Wright et al

Acrosol extinction coefficient as a function of wavelength (Wright et

Aerosol Volume Backscattering and Extinction Coefficients in the NIR (CO₂ laser wavelengths and others)


Backscattered signal measured with the ir lidar system.

Logarithmic Amplification of IR LIDAR Return Signal. SNR~300

Atmospheric Ammonia

It is produced by a combination of natural and anthropogenic sources. Concentration of ambient NH_3 varies between 3ppb and 10ppb.

Detection and measurement of Ammonia are important:

Concentration of NH₃ is critical parameter in the rates of formation, transport, transformation and removal of sulfur and nitrogen oxides.
Nitrogen aerosols play an important role in the generation of acid rain.



Ammonia Measurements Near a Corn Field

 α (P30) = 1.1 (atm. cm)⁻¹ α (P32) = 12 (atm. cm)⁻¹

Time variation of LIDAR returns of 10.69- μ m P(30) and 10.71- μ m P(32) radiation passing through the tank and reflected from a topographic target at a range of 2.7 km after injecting 0.2 cm³ of 28% aqueous NH₃





Ammonia Measurements Near a Corn Field

Single-laser DIAL measurement of ambient atmospheric NH₃ using 9.24- μ m R(26) and 9.22- μ m R(30) CO₂ laser lines over a 2.7-km range with 26% relative humidity

> DIAL measurement of ambient atmospheric NH₃ using 9.24- μ m R(26) and 9.22- μ m R(30) CO₂ laser lines over a 2.7-km range showing the effect of a change in relative humidity from 54 to 41%

Ozone and Ozone Layer:

-Ozone is a naturally occurring gas found in the Earth's atmosphere that absorbs certain wavelengths of the sun's UV radiation.

-Ozone Layer: around 15-35 Km above the earth's surface.

-10% of the total ozone is found in the troposphere.

-The troposphere contains 90% of the atmosphere and nearly all the atmosphere's water vapor.

Tropospheric Ozone:

-the layer between the earth's surface and tropopause (border between troposphere and stratosphere) contains only a fraction of the ozone. Due to multiple scattering (dust and aerosols) of rays within this layer, the solar UV is absorbed there more efficiently than within an equal amount of stratospheric ozone.



-In spite of this benefit, tropospheric ozone is often referred to as "bad ozone" or "bad Nearby" because of its adverse effects.

Ozone in the Troposphere "Bad Nearby"

VOCs = volatile organic compounds

 $Oightharpoonup NO_X = nitrogen oxides$ VOC + NOx + Heat + Sunlight = Ozone

VOCs and NO_X result primarily from combustion of fossil fuels







Effect of UVB Radiation on the Persistence Time of Ozone



Inter-relation between NO₂, O₃, UVB and NO₂*UVB



Atmospheric Ethylene as a Hydrocarbon Representative and its Relation with Ozone

 \succ It is an important urban air pollutant.

> It is emitted as a portion of motor vehicle exhaust

> Plants exhibit symptoms of ethylene toxicity at concentrations as low as 10 ppb.

Typical effects:

Reduction of plant growth

> Reduction of flowering and hence fruit production

> Premature aging of plants.

Motivations:

> Study the ethylene-NO_X-Ozone inter-correlation on one-day scale as a function of UV.



Ethylene DIAL measurements:

 C_2H_4 with 7 ppb-correction and ozone point monitored concentration.

 C_2H_4 starts at a background level of 15-20 ppb and reaches a peak of 80 ppb. Ozone, showed the same behavior namely, a slow increase then reached a maximum at around 15:00 and back to 40-50 ppb level.



7 ppb-correction ethylene and Ozone measurements of May 13.

Summary and Outlook

Lidar remote sensing is an advanced technology that is not only replacing conventional sensors in science study, environmental research, and industry application, but also creating new methods with unique properties that could not be achieved before.

- □ Lidar technology has been advanced dramatically in the past 20 years, owing to the new availability of lasers, detectors, creative people involved, and the demanding needs from various aspects.
- Potential growing points at this stage include
- (1) Solid-state resonance fluorescence lidar for mobile deployment globally
- (2) Extend measurement range into thermosphere and lower mesosphere
- (3) Doppler, DIAL, HSRL, and Raman lidar for lower atmosphere research
- (4) Target lidar for novel applications

Material used in preparing the lectures:

-Laser Remote Sensing, Edited by Takshi Fujii and Tetsuo Fukuchi, Published by CRC Press, Taylor & Francis Group, ISBN: 0-8247-4256-7, 2005.

-Lidar: Range-resolved optical remote sensing of the atmosphere, edited by Claus Weitkamp, published by Springer, ISBN: 0-387-40075-3, 2005.

-Laser Remote Sensing: Fundamentals and Applications, by Raymond M. Measures, Wiley- Interscience, New York, ISBN: 0-89464-619-2, 1984.

- Lecture notes of Professor X. Chu; CIRES, Colorado University, USA

Thank you for your attention!