Advanced Optics (PHYS690)

HEEDEUK SHIN

Lecture 20

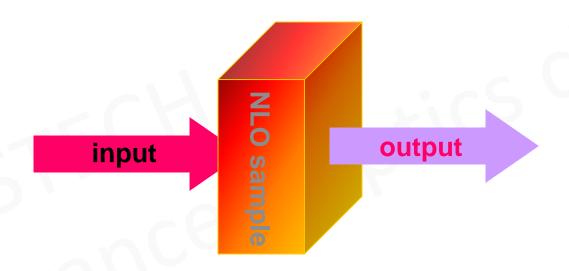
POHANG UNIVERSITY OF SCIENCE AND TECHNOLOGY, KOREA





Question:

Is it possible to change the color of a monochromatic light?



Answer:

Not without a laser light



Nonlinear Optics



• Dielectric media characterized by a linear relation between polarization and E-field. Polarization: the total induced dipole moment per unit volume

$$P = \varepsilon_0 \chi E$$

Media characterized by a nonlinear relation between E and P.

$$P = a_1 E + \frac{1}{2} a_2 E^2 + \frac{1}{6} a_3 E^3 + \dots$$

$$= \varepsilon_0 \left(\chi E + \chi^{(2)} E^2 + \chi^{(3)} E^3 + \dots \right)$$

$$= \varepsilon_0 \chi E + 2\varepsilon_0 dE^2 + \varepsilon_0 \chi^{(3)} E^3 + \dots$$

$$d = \frac{1}{4} a_2 = \frac{1}{2} \chi^{(2)}$$
1st order 2nd order 3rd order

d: second order nonlinear coefficient



Why polarization?



From Maxwell's equations

$$\nabla \cdot \vec{D} = \rho$$

Gauss's law for electric field

$$\nabla \cdot \vec{B} = 0$$

Gauss's law for magnetic field

$$\nabla \times \vec{E} = -\frac{\partial \vec{B}}{\partial t}$$

Ampere's law

$$\nabla imes ec{H} = ec{J} + rac{\partial ec{D}}{\partial t}$$

Faraday's law

where
$$\vec{D}$$
 electric displacement (C/m^2)
 \vec{E} electric field (V/m)
 \vec{B} magnetic field $(T \text{ or } V \cdot s/m^2)$
 \vec{H} magnetic field intensity (A/m)
 ρ volume charge density (C/m^3)
current density (A/m^2)

 μ_0 : permeability ϵ_0 : permittivity

$$\mu_0 \epsilon_0 = \frac{1}{c^2}$$

Free space wave equation

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = 0$$

$$\vec{E}(\vec{r}) = \hat{n}E_0 e^{-j\vec{k}\cdot\vec{r}}$$

$$\left|\vec{k}\right| = c\omega$$

• In a dielectric medium

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \epsilon \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = 0$$

$$\left| \vec{k} \right| = \frac{c}{n} \omega$$

$$n = \sqrt{\epsilon}$$



Why polarization?



In a dielectric medium

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \epsilon \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = 0$$

$$\left| \vec{k} \right| = \frac{c}{n} \omega$$

$$n = \sqrt{\epsilon}$$

Note that

$$\vec{D} = \epsilon_0 \vec{E} + \vec{P}$$
 $\vec{P} = \epsilon_0 \chi \vec{E}$
 $\vec{D} = \epsilon_0 (1 + \chi) \vec{E} = \epsilon_0 \epsilon \vec{E}$

Then

$$\nabla^2 \vec{E}(\vec{r}) = -\mu_0 \epsilon_0 \epsilon \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = -\mu_0 \epsilon_0 \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) - \mu_0 \epsilon_0 \chi \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r})$$

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = -\mu_0 \frac{\partial^2}{\partial t^2} \vec{P}(\vec{r}) \qquad - \Delta \text{ source term}$$

Propagating wave



Nonlinear wave equation

From Maxwell's equation and now considering

$$\vec{D} = \epsilon_0 \vec{E} + \vec{P} \qquad \qquad \vec{P} = \epsilon_0 \chi \vec{E}$$

We obtain a nonlinear wave equation

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = -\mu_0 \frac{\partial^2}{\partial t^2} \vec{P}(\vec{r})$$

where P is usually written as

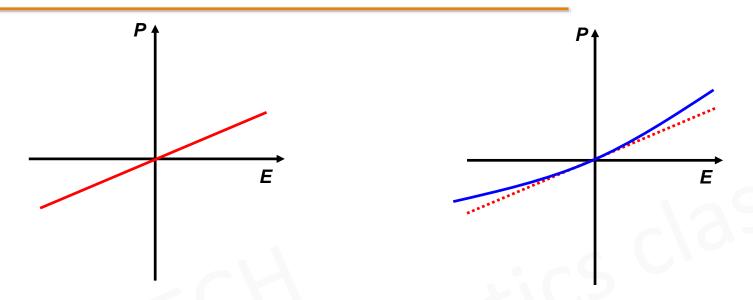
$$P = \epsilon_0 \chi E + 2\epsilon_0 dE^2 + \epsilon_0 \chi^{(3)} E^3 + \cdots$$

$$P = P_L + P_{NL}$$

$$P_L = \epsilon_0 \chi E, \qquad P_{NL} = 2\epsilon_0 dE^2 + \epsilon_0 \chi^{(3)} E^3 + \cdots$$

In centrosymmetric media, d vanish, and the lowest order nonlinearity is of third order





P-E relation for (a) a linear dielectric medium, and (b) a nonlinear medium.

https://www.youtube.com/watch?v=anwl6OZ1UuQ

Typical values

$$d \sim 10^{-3} - 10^3 \left[\frac{pm}{V}\right]$$

$$\chi^{(3)} \sim 10^{-25} - 10^{-8} \left[\frac{m^2}{V^2} \right]$$





 $\chi^{(2)}$

Second order nonlinear effects



Second harmonic generation (SHG) I

Assume higher order than second order are negligible

$$P_{NL} = 2\epsilon_0 dE^2$$

Take

$$\begin{split} E &= \frac{1}{2} \Big(E(\omega) e^{i\omega t} + c.c. \Big) \\ \mathcal{E}(t) &= \text{Re}\{ E(\omega) \exp(j\omega t) \} = \frac{1}{2} [E(\omega) \exp(j\omega t) + E^*(\omega) \exp(-j\omega t)] \end{split}$$

Corresponding nonlinear polarization density is

$$P_{NL} = 2d\frac{1}{4} \Big(E(\omega)e^{i\omega t} + c.c. \Big) \Big(E^*(\omega)e^{-i\omega t} + c.c. \Big)$$

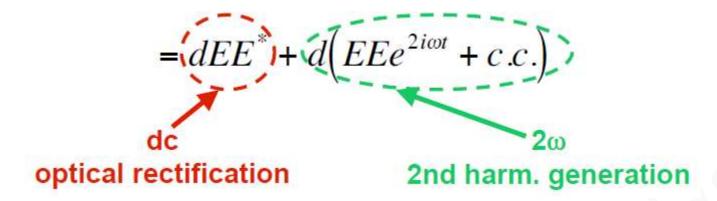
$$\mathcal{P}_{\mathrm{NL}}(t) = P_{\mathrm{NL}}(0) + \operatorname{Re}\{P_{\mathrm{NL}}(2\omega) \exp(j2\omega t)\}$$

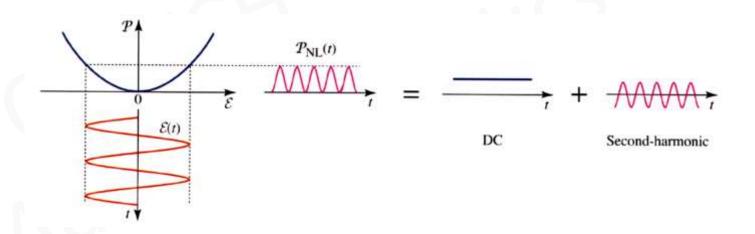
$$P_{\mathrm{NL}}(0) = \operatorname{d}E(\omega)E^*(\omega)$$

$$P_{\mathrm{NL}}(2\omega) = \operatorname{d}E^2(\omega). \longrightarrow E(2\omega)$$



Second harmonic generation (SHG) II







First SHG report



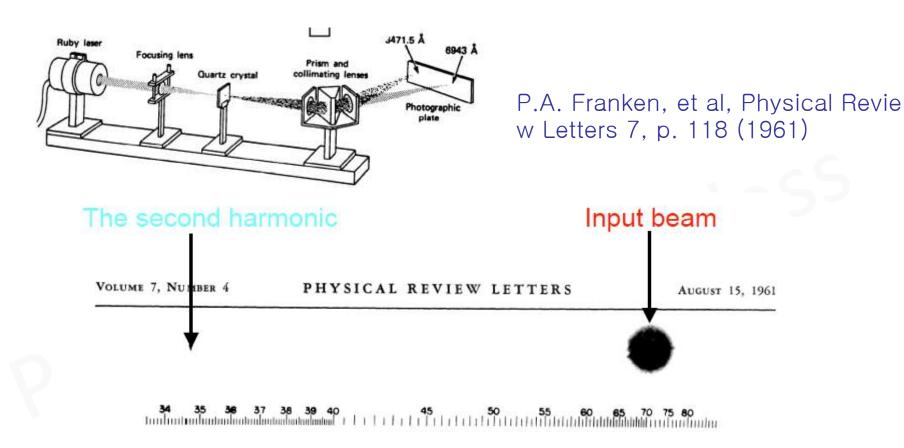


FIG. 1. A direct reproduction of the first plate in which there was an indication of second harmonic. The wavelength scale is in units of 100 A. The arrow at 3472 A indicates the small but dense image produced by the second harmonic. The image of the primary beam at 6943 A is very large due to halation.



Phase Matching



$$E(\omega_1) = A_1 \exp(-j\vec{k_1} \cdot \vec{r})$$

$$E(\omega_2) = A_2 \exp(-j\vec{k_2} \cdot \vec{r})$$

$$P_{NL}(\omega_3) = 2dE(\omega_1)E(\omega_2) = 2dA_1A_2 \exp(-j\vec{k}_3 \cdot \vec{r})$$

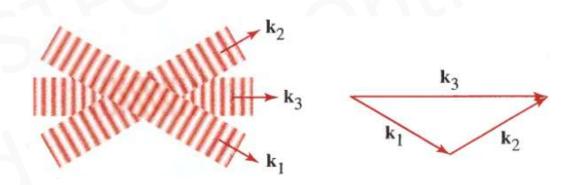
where

$$\left(\omega_3 = \omega_1 + \omega_2\right)$$

Frequency-Matching Condition

$$\vec{k}_3 = \vec{k}_1 + \vec{k}_2$$

Phase-Matching Condition



The phase-matching condition



Phase matching I



Consider a plane wave propagating along z, and write the paraxial Hemh oltz equation

$$\left[\nabla_{\perp}^{2}E(2\omega)-2ik\partial_{z}E(2\omega)\right]e^{-k(2\omega)z}=\mu_{0}\partial_{t}^{2}\left(dE^{2}(\omega)\right)e^{-k(\omega)z}$$

Can be approximated as

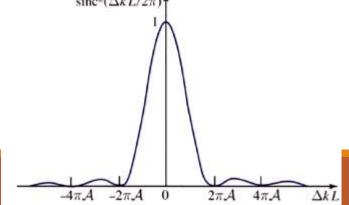
$$\partial_z E(2\omega) = AdE^2(\omega)e^{\Delta kz}$$

where $\Delta k = k(2\omega) - k(\omega)$ so finally

$$E(2\omega) \propto \frac{\sin \Delta kz}{\Delta kz}$$

need Δk =0 to achieve maximum E(2 ω) electric field (this was not taken c are in the 1961 experiment) $\frac{\sin^2(\Delta k L/2\pi)}{\uparrow}$

This is referred to as "phase matching"





Phase matching II



So we're creating light at $\omega_{sig} = 2\omega$.

So we're creating light at
$$k_{sig} = \frac{\omega_{sig}}{c_0} n(\omega_{sig}) = \frac{(2\omega)}{c_0} n(2\omega)$$

And the k-vector of the polarization is: $k_{pol} = 2 k = 2 \frac{\omega}{c_0} n(\omega)$

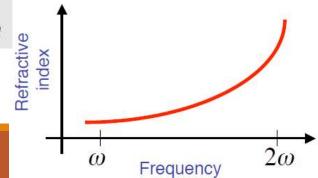
The phase-matching condition is: $k_{sig} = k_{pol}$

which will only be satisfied when:

$$n(2\omega) = n(\omega)$$

$$\omega_1 + \omega_2 = \omega_3, \quad \omega_1 n_1 + \omega_2 n_2 = \omega_3 n_3,$$

Unfortunately, dispersion prevents this from ever happening!

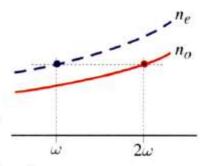




Phase matching III



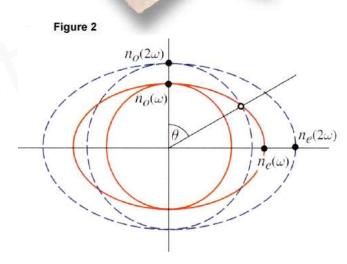
- Birefringent materials different refractive indices for different polarizations.
- Ordinary and extraordinary refractive indices can be different by up to ~0.1 for SHG crystals.
- We can now satisfy the phase-matching condition.



Use the extraordinary polarization for ω and the ordinary for 2ω .

$$n_o(2\omega) = n_e(\omega)$$

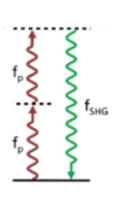
 n_e depends on propagation angle, so we can tune for a given ω . Some crystals have $n_e < n_o$, so the opposite polarizations work.

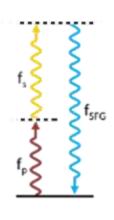


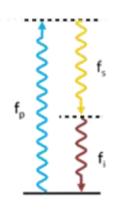


Phase Matching





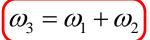




Second Harmonic

Sum Frequency

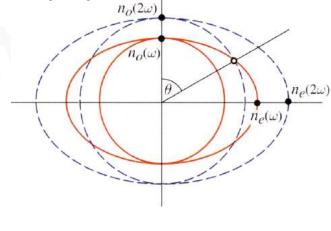
Difference Frequency



Frequency-Matching Condition

$$\vec{k}_3 = \vec{k}_1 + \vec{k}_2$$

Phase-Matching Condition



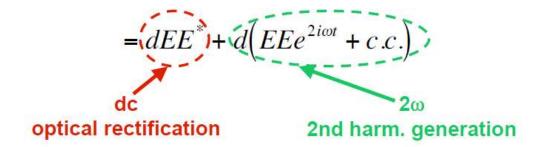
$$\omega$$
 2ω

$$n_o(2\omega) = n_e(\omega)$$



Optical Rectification





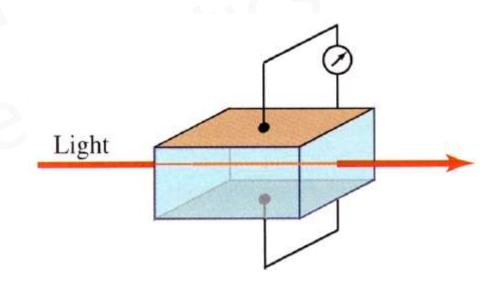
$$\mathcal{P}_{\rm NL}(t) = P_{\rm NL}(0) + \text{Re}\{P_{\rm NL}(2\omega) \exp(j2\omega t)\}$$

$$P_{\rm NL}(0) = d E(\omega) E^*(\omega)$$

$$P_{\rm NL}(2\omega) = d E^2(\omega).$$

Generation of DC voltage

MW → uV





Electro-Optic Effect



• E-field consists of harmonic component at optical frequency and zero frequency.

$$\mathcal{E}(t) = E(0) + \text{Re}\{E(\omega) \exp(j\omega t)\}\$$

$$\mathcal{P}_{\rm NL}(t) = P_{\rm NL}(0) + \text{Re}\{P_{\rm NL}(\omega)\exp(j\omega t)\} + \text{Re}\{P_{\rm NL}(2\omega)\exp(j2\omega t)\},$$

$$P_{\rm NL}(0) = d \left[2E^2(0) + |E(\omega)|^2 \right]$$

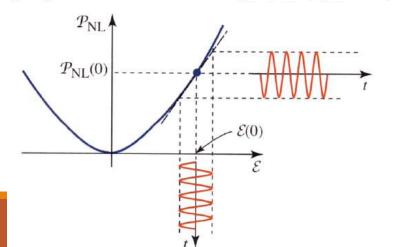
$$P_{\rm NL}(\omega) = 4d E(0)E(\omega)$$

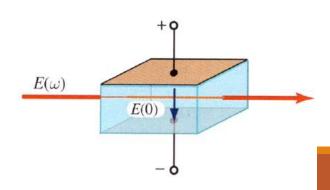
$$P_{\rm NL}(2\omega) = d E^2(\omega),$$

$$|E(\omega)|^2 \ll |E(0)|^2$$

$$\Delta n = \frac{2\mathbf{d}}{n\epsilon_o} E(0).$$

Pockels effect







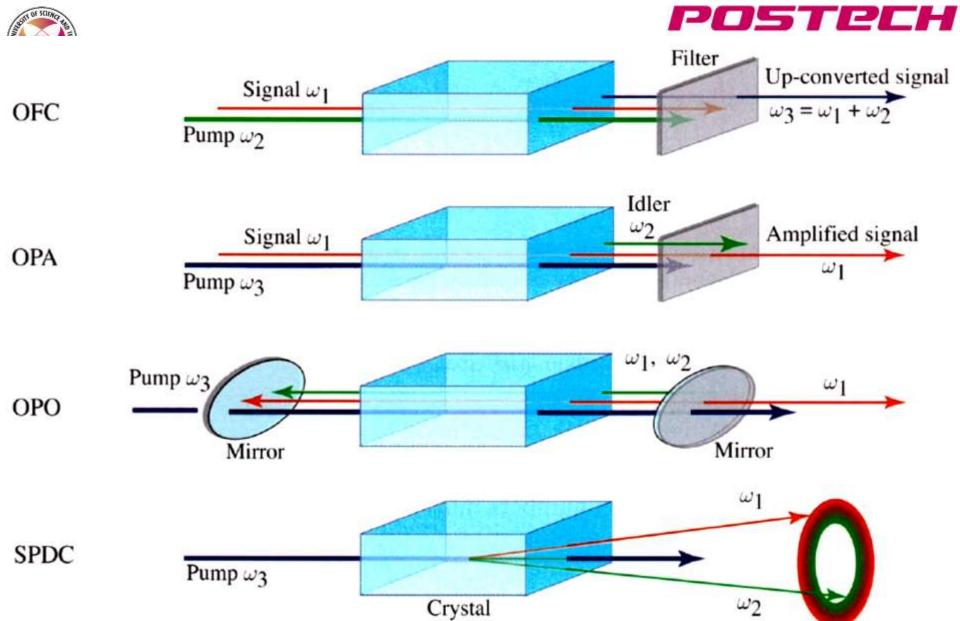
Three-wave mixing



E-field

$$\mathcal{E}(t) = \operatorname{Re}\{E(\omega_1) \exp(j\omega_1 t) + E(\omega_2) \exp(j\omega_2 t)\}$$

$$\begin{split} P_{\mathrm{NL}}(0) &= \mathrm{d} \left[|E(\omega_1)|^2 + |E(\omega_2)|^2 \right] \\ P_{\mathrm{NL}}(2\omega_1) &= \mathrm{d} \ E(\omega_1) E(\omega_1) \\ P_{\mathrm{NL}}(2\omega_2) &= \mathrm{d} \ E(\omega_2) E(\omega_2) \\ P_{\mathrm{NL}}(\omega_+) &= 2 \mathrm{d} \ E(\omega_1) E(\omega_2) \\ P_{\mathrm{NL}}(\omega_-) &= 2 \mathrm{d} \ E(\omega_1) E^*(\omega_2). \end{split}$$





POSTECH POHANG UNIVERSITY OF SCIENCE AND TECHNOLOGY

Application of noncollinear phase matching

 Noncollinear phase BBO matching can be used to measure the duration of ultra-short pulse 1000 800 600 FWHM=2.89 mm ∆t=6.2 ps Signal (a.u.) 400 200 0 -200 12 16

Distance (mm)



 $\chi^{(3)}$

Third order nonlinear effects



Third order nonlinear polarization

$$P_{NL}^{\delta} = 4\chi^{(3),\alpha\beta\gamma\delta} E_{\alpha} E_{\beta} E_{\gamma}$$

• Let's consider the simple case of third harmonic generation (THG).

$$E = E_0 \cos(\omega t)$$

Then the third order polarization is

$$P_{NL} = 4\chi^{(3)} E_0^3 \cos^3(\omega t)$$

$$= 4\chi^{(3)} E_0^3 \left[\frac{1}{4} \cos(3\omega t) + \frac{3}{4} \cos(\omega t) \right]$$

3rd order generation

Nonlinear contribution that affects the fundamental frequency

ullet The ω term in the polarization is of the form

$$P_{NL}(\omega) = \chi^{(1)} E(\omega) + 3\chi^{(3)} |E(\omega)|^2 E(\omega)$$
$$= \chi_{eff} E(\omega)$$



Third order nonlinear effects

• The general form of the $\chi^{(3)}$ tensor is:

$$P_i^{(3)}(\omega) = \epsilon_0 \chi_{iikl}^{(3)}(\omega : \omega_1, \omega_2, \omega_3) E_i(\omega_1) E_k(\omega_2) E_l(\omega_3)$$

- $\chi^{(2)}$ effects are only observed in materials without inversion symmetry.
- All materials have nonzero $\chi^{(3)}$ coefficients.
- The $\chi^{(3)}$ coefficient gives rise to an intensity dependent index of refraction.

$$P = \chi^{(1)}E + \chi^{(3)}E^3$$
 Optical Kerr effects
= $(\chi^{(1)} + \chi^{(3)}|E|^2)E$ $n = n_0 + n_2I$

- In Gaussian laser beams, the intensity is highest in the center of the beam.
- Hence the index of refraction is highest in the center.



Nonlinear index of refraction of science and technology

- Real part of index is best described as a power series
- $n = n_1 + n_2 (P/A_{eff})$ $n = n_0 + n_2 I$
- For silica fiber, $n_2 \cong 2.6 \times 10^{-11} \, \mu m^2 / mW$

Interaction Length

$$L_{eff} = \frac{1 - e^{-\alpha L}}{\alpha} \approx \frac{1}{\alpha}$$
 If L >> 1/ α

where α (in cm⁻¹) is the loss coefficient of the fiber. 0.1 dB/km=2.3x10⁻⁷ cm⁻¹.

Nonlinear parameter

$$\gamma = \frac{2\pi n_2}{\lambda A_{eff}} \qquad \beta_{NL} = \beta + \gamma P$$

Propagation constant is power-dependent.



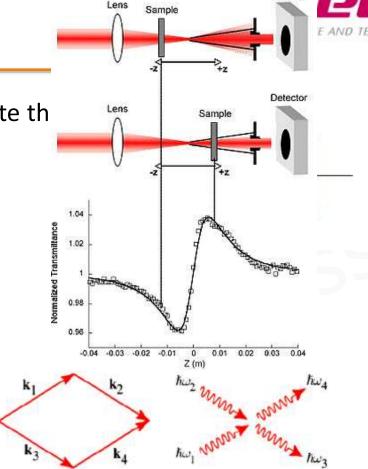
Self Focusing

From Fermat's principle we can estimate th

$$(n_0 + n_2 I) z_{SF} = \frac{n_0 z_{SF}}{\cos \theta_{SF}} \Rightarrow \theta_{SF} = \sqrt{\frac{2n_2 I}{n_0}}$$
$$z_{SF} = w_0 \theta_{SF} = w_0 \sqrt{\frac{n_0}{2n_2 I}}$$

4-wave mixing

It couples 4 E-fields together.



Similar to 3-wave mixing in 2nd order crystal (SHG is a degenerate case of 3-wave mixing), the process follow the conservation laws.

$$\omega_1 + \omega_2 = \omega_3 + \omega_4.$$
 $\mathbf{k}_1 + \mathbf{k}_2 = \mathbf{k}_3 + \mathbf{k}_4.$ $P_{\text{NL}}(\omega_2) = 6\chi^{(3)}E(\omega_3)E(\omega_4)E^*(\omega_1),$



Self-phase modulation



Self-phase modulation is due to the intensity dependent refractive index

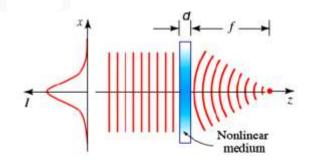
$$\varphi = -n(I)k_oL = 2\pi n(I)L/\lambda_o = -2\pi(n + n_2P/A)L/\lambda_o$$

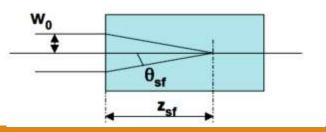
- P and A are respective the beam power and cross-sectional area.
- The induces self phase modulation is

$$\Delta \varphi = -2\pi n_2 \frac{L}{\lambda_o A} P$$

3rd order effects: Self Focusing

- Assume a beam with intensity larger at its center.
- When passing through a 3rd order medium, the me dium acts as an index graded element and results in focusing of the beam.
- In too long media, self focusing can locally leads to intensity above damage threshold.







Self-phase modulation in commission in commission in the commissio

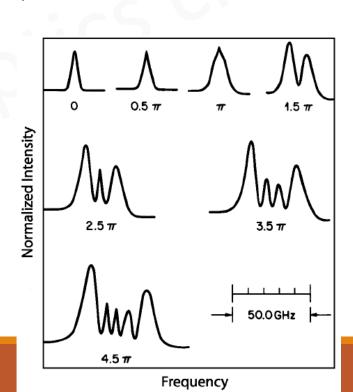
Refractive index depends on optical intensity as (Kerr eect)

$$n(\boldsymbol{\omega},I)=n_0(\boldsymbol{\omega})+n_2I(t)$$

Intensity dependence leads to nonlinear phase shift

$$\phi_{\rm NL}(t) = (2\pi/\lambda)n_2I(t)L.$$

- An optical field modifies its own phase (SPM).
- Phase shift varies with time for pulses.
- Each optical pulse becomes chirped.
- As a pulse propagates along the fiber,
 its spectrum changes because of SPM.





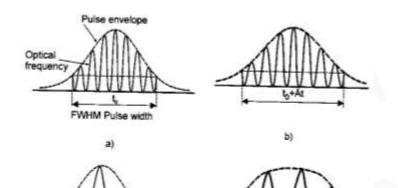
Phase modulation



Self-modulation: $\phi_{NL} = \gamma PL_{eff}$

Cross-modulation: $\phi_{NL} = 2\gamma P_{other} L_{eff}$

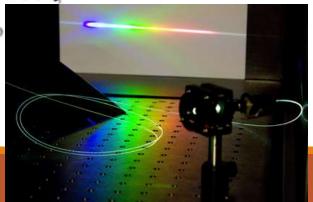
Effect of these phase changes is a frequency chirp (frequency changes during pulse), broadening pulse and reducing bit rate-length product.

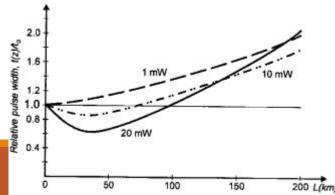


Self-phase modulation effect:

Spreading of chirped pulse:

- (a) Regular unchirped pulse entering the link;
- (b) The same pulse distorted after traveling distance *L* along the fiber;
- (c) Chirped pulse entering the link;
- (d) Chirped pulse broadens after traveling distance L.







Cross-Phase Modulation

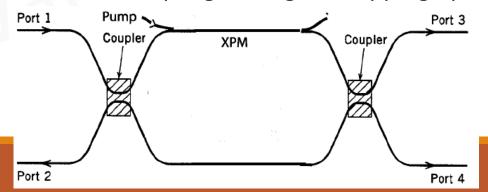


- Consider two optical fields propagating simultaneously.

• Total nonlinear phase shift in a fiber of length L:

$$\phi_{\rm NL} = (2\pi L/\lambda) n_2 [I_1(t) + bI_2(t)]$$

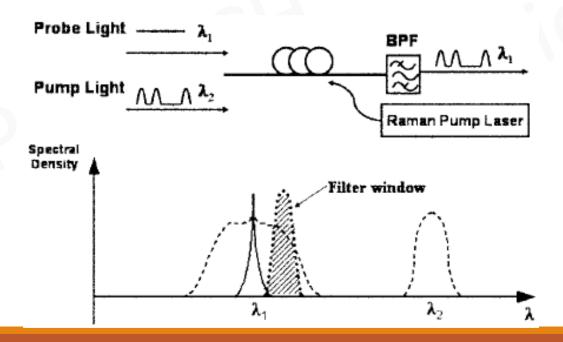
- An optical beam modifies not only its own phase but also of other co-propagating beams (XPM).
- XPM induces nonlinear coupling among overlapping optical pulses.





XPM-Based Wavelength Converter

- WDM channel at λ_2 requiring conversion acts as a pump.
- A CW probe is launched at the desired wavelength λ_1 .
- Probe spectrum broadens because of pump-induced XPM.
- An optical filter blocks pump and transfers data to probe.
- Raman amplification improves the device performance.

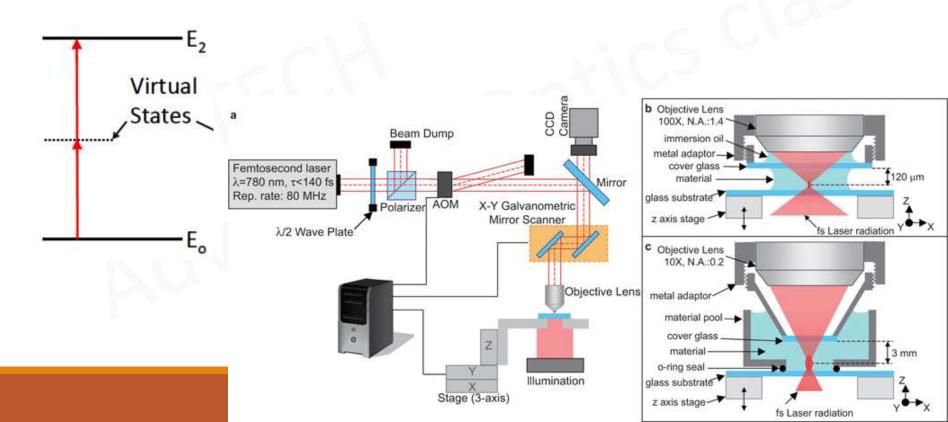




Two Photon Absorption



- •Two photons absorbed simultaneously to bring atom to high-lying energy level.
- •Absorption enhanced by enhanced by allowed transitions near sin gle photon resonance.
- •Can limit the performance of high-powered lasers.

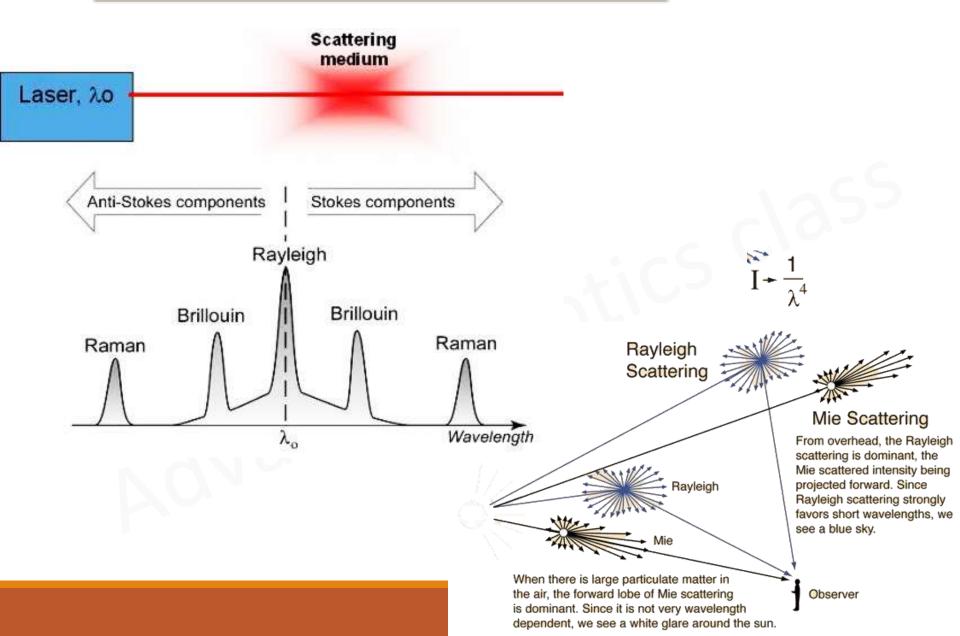






Scattering







Stimulated Raman Scattering (SRS)

Interaction of light with an acoustic phonon in the material.

Ω

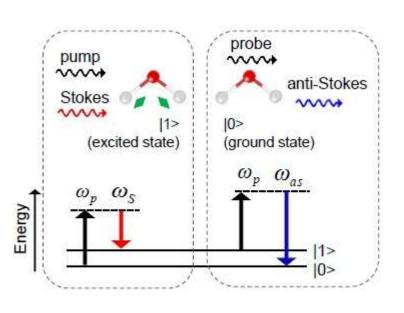
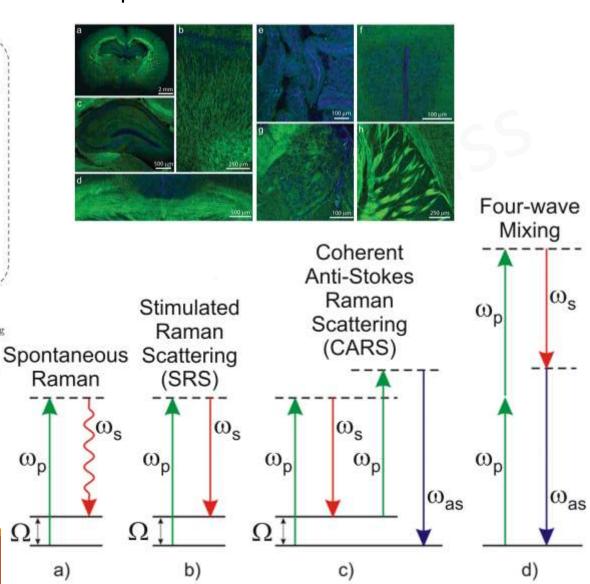


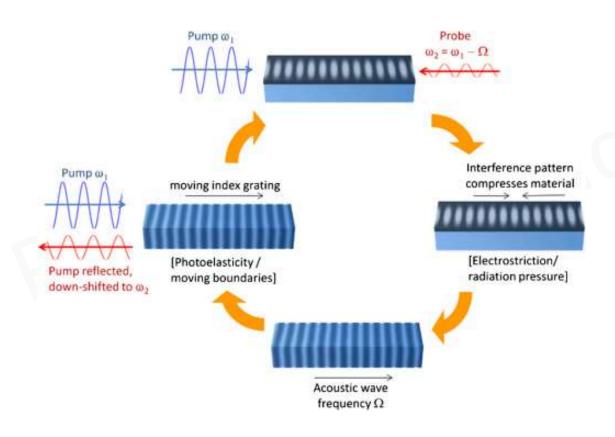
Figure 1: Energy level diagram for coherent anti-Stokes Raman scattering





Stimulated Brilliouin Scattering (SBS)

- Interaction of light with an acoustic phonon in the material.
- The scattered wave is always(?) backwards propagating.





Concluding Remarks



- Nonlinear effects are sources of noise in many systems and not easy to understand.
- Don't be afraid of it.
- Nonlinearities can be managed thorough proper system d esign.
- By combining two or more nonlinear effects, unpreceden ted behavior properties and applications can be designed.



POSTECH Optics class



Nonlinear wave equation Marketta

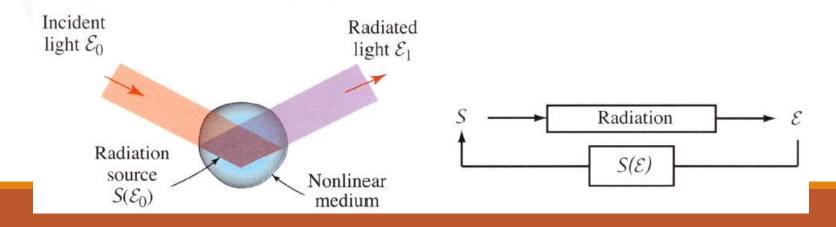
• Using
$$c = c_0/n$$

 $n^2 = 1 + \chi$

The nonlinear wave equation can be rewritten

$$\nabla^2 \vec{E}(\vec{r}) + \mu_0 \epsilon_0 \epsilon \frac{\partial^2}{\partial t^2} \vec{E}(\vec{r}) = -\mu_0 \frac{\partial^2}{\partial t^2} \vec{P}_{NL}(\vec{r})$$

• The fist Born approximation. An incident optical field \mathcal{E}_0 creates a sourc e $\mathcal{S}(\mathcal{E}_0)$, which radiates an optical field \mathcal{E}_1 .







Coupled- Wave Equations for Second-Harmonic Generation.

$$\frac{da_1}{dz} = -jga_3a_1^* \exp(-j\Delta kz)$$

$$\frac{da_3}{dz} = -j\frac{g}{2}a_1a_1 \exp(-j\Delta kz)$$

$$\frac{da_3}{dz} = -j\frac{g}{2}a_1a_1 \exp(-j\Delta kz)$$

where
$$\Delta k = k_3 - 2k_1$$
 $g^2 = 4\hbar \omega^3 \eta^3 d^2$

$$g^2 = 4\hbar\omega^3\eta^3d^2$$

perfect phase matching

$$\Delta k = 0$$

$$\frac{da_1}{dz} = -jga_3a_1^*$$

$$\frac{da_3}{dz} = -j\frac{g}{2}a_1a_1$$

Coupled Equations (Second-Harmonic Generation)





the solution

$$a_1(z) = a_1(0) \sec h \frac{ga_1(0)z}{\sqrt{2}}$$

$$a_3(z) = -\frac{j}{\sqrt{2}} a_1(0) \tan h \frac{g a_1(0) z}{\sqrt{2}}$$

Consequently, the photon flux densities

$$\phi_1(z) = \phi_1(0) \sec h^2 \frac{\gamma z}{2}$$

$$\phi_3(z) = -\frac{1}{2}\phi_1(0) \tan h^2 \frac{\gamma z}{2}$$





complex amplitude

$$S(2\omega) = 4\mu_o \omega^2 \, dE(\omega) E(\omega)$$

SHG intensity

$$I(2\omega) \propto |\mathcal{S}(2\omega)|^2 \propto |I(\omega)|^2 \propto d^2 \propto L^2$$

Efficiency of second-harmonic generation

$$I(\omega) = P/A$$

$$\eta_{\rm SHG} = I(2\omega)/I(\omega) \propto L^2 I(\omega)$$





The efficiency of second-harmonic generation for an interaction region of I ength L is

$$\frac{I_3(L)}{I_1(0)} = \frac{\hbar \omega_3 \phi_3(L)}{\hbar \omega_1 \phi_1(L)} = \frac{2\phi_3(L)}{\phi_1(0)} = \tanh^2 \frac{\gamma L}{2}$$

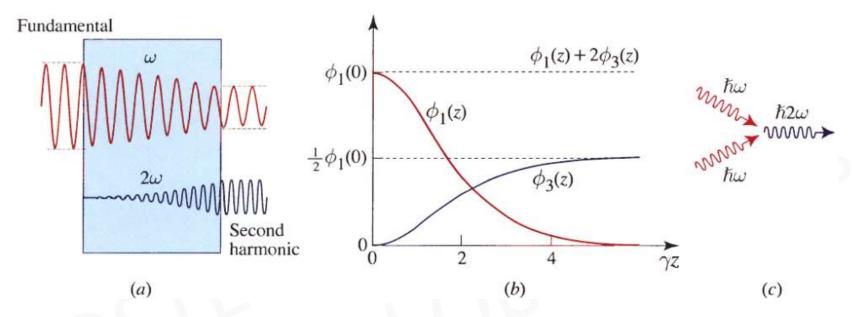
For large γL (long cell, large input intensity, or large nonlinear parameter), the efficiency approaches one. This signifies that all the input power (at frequency w) has been transformed into power at frequency 2w; all input photons of frequency w are converted into half as many photons of frequency 2w.

For small γL (small device length L, small nonlinear parameter d, or small inp ut photon flux density $\phi_1(0)$), the argument of the tanh function is small and th erefore the approximation tanhx=x may be used. The efficiency of second-harmonic generation is then

$$\frac{I_3(L)}{I_1(0)} = 2\eta_0^3 \omega^2 \frac{d^2}{n^3} \frac{L^2}{A} P$$



Second-harmonic generation of Science AND TECHNOLOGY



- •Two photons of frequency w combine to make one photon of frequency 2w.
- As the photon flux density $\phi_i(z)$ of the fundamental wave decreases, the photon flux density $\phi_i(z)$ of the second-harmonic wave increases.
- Since photon numbers are conserved, the sum $\phi_1(z)+2\phi_3(z)=\phi_1(0)$ is a constant.



Effect of Phase Mismatch



$$\frac{da_1}{dz} = -jga_3a_1^* \exp(-j\Delta kz)$$

$$\frac{da_3}{dz} = -j\frac{g}{2}a_1a_1 \exp(-j\Delta kz)$$

$$\Delta k \neq 0$$

$$\frac{da_3}{dz} = -j\frac{g}{2}a_1a_1 \exp(-j\Delta kz)$$

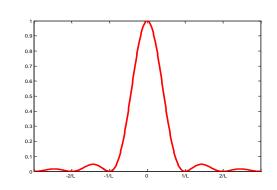
$$\Delta k \neq 0$$

Solution

$$a_3(L) = -j\frac{g}{2}a_1^2(0)\int_0^L \exp(-j\Delta kz')dz' = -(\frac{g}{2\Delta k})a_1^2(0)[\exp(-j\Delta kL) - 1]$$

Efficiency

$$\frac{I_3(L)}{I_1(0)} = \frac{2\phi_3(L)}{\phi_1(0)} = \frac{1}{2}g^2L^2\phi_1(0)\sin c^2\frac{\Delta kL}{2\pi}$$





Phase matching I



Consider a plane wave propagating along z, and write the paraxial Hemh oltz equation

$$\left[\nabla_{\perp}^{2}E(2\omega)-2ik\partial_{z}E(2\omega)\right]e^{-k(2\omega)z}=\mu_{0}\partial_{t}^{2}\left(dE^{2}(\omega)\right)e^{-k(\omega)z}$$

Can be approximated as

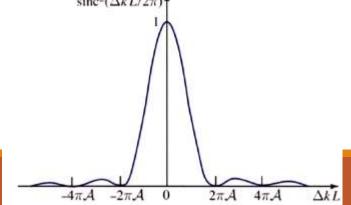
$$\partial_z E(2\omega) = AdE^2(\omega)e^{\Delta kz}$$

where $\Delta k = k(2\omega) - k(\omega)$ so finally

$$E(2\omega) \propto \frac{\sin \Delta kz}{\Delta kz}$$

need Δk =0 to achieve maximum E(2 ω) electric field (this was not taken c are in the 1961 experiment) $\frac{\sin^2(\Delta k L/2\pi)}{\uparrow}$

This is referred to as "phase matching"



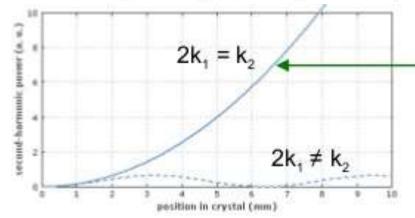


Quasi-phase matching



- Only when $2k_1 = k_2$, SHG will be efficient.
- Only when phase matching is achieved, these contributions add up con structively, and a high power conversion efficiency is achieved.
- The direction of energy transfer changes periodically according to the change in the phase relation between the interacting waves.
- The energy then oscillates between the waves rather than being transferred in a constant direction.
- The effect on the power conversion is illustrated.

$$\frac{I_{2\omega_1}}{I_{\omega_1}} = \frac{A_2^* A_2}{A_1^* A_1} = \left(\frac{2\omega_1 d_{eff}}{n_2 c \Delta k}\right)^2 |A_1|^2 \sin^2\left(\frac{\Delta k L}{2}\right)$$



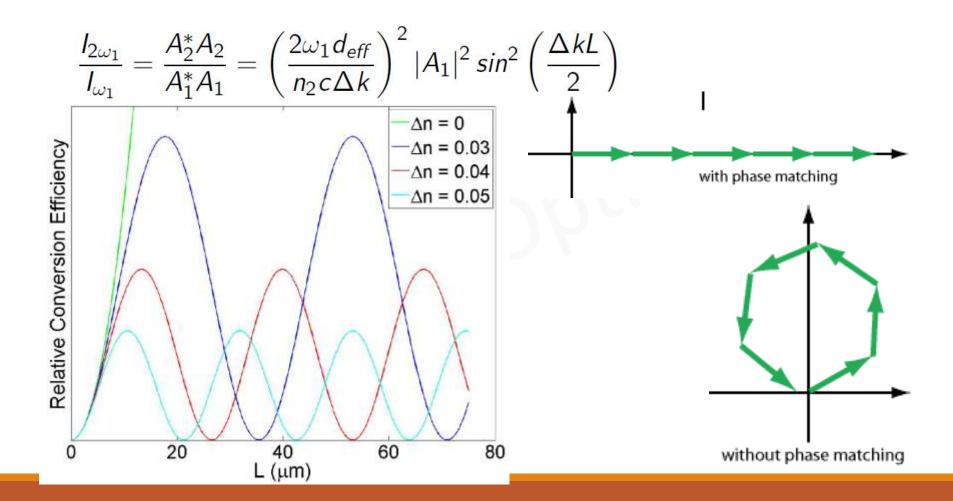
~ 100% SHG conversion efficiency is possible by optimizing phase matching!



Conversion efficiency



• Wavevector mismatch Δk may cause significant power reduction of generated parametric wave.

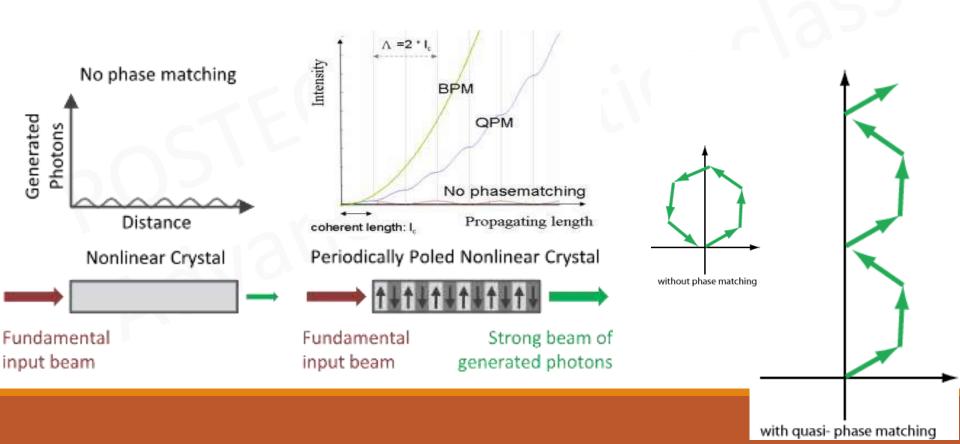




Quasi-phase matching



- Perfect phase matching can be difficult to achieve.
- Alternate way for high efficient generation is periodic nonlinearity.
- Such periodicity introduces an opposite phase bringing back the phases of the distributed radiation elements into better alignment.





POHANG UI

An = 0

An = 0.03

An = 0.04

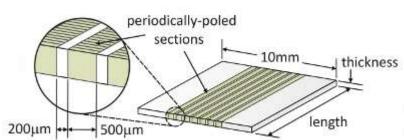
An = 0.04

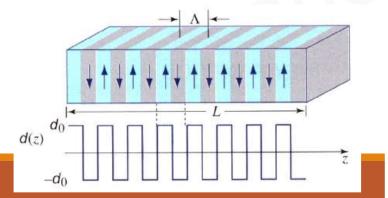
An = 0.05

Ce length or one of the control of the control

- Consider constructively adding distance, coherence length $l_c = \pi/\Delta k$
- It is useful to think of the periodic structure as adding a grating vector of magnitude $2m\pi/\Lambda$ to the interaction.
- The grating allows momentum to be conserved in the nonlinear process.

$$\Delta k - \frac{2\pi m}{\Lambda} = 0 \qquad \Lambda < 10\mu m$$





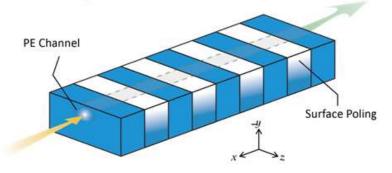
Crystal	Periodically poled LiNbO3	
Coercive Field	21 kV/mm	
Maximum Poling Depth	1 mm	
Maximum Crystal Length	50 mm	
$d_{ m eff}$	17 pm/V	
Transparency Range	0.4 - 5.5μm	

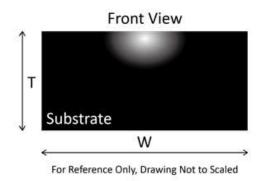


PP crystal waveguides

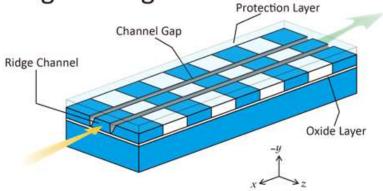


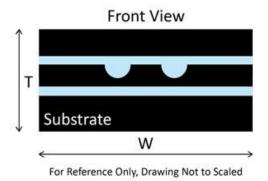
PE Waveguide





Ridge Waveguide

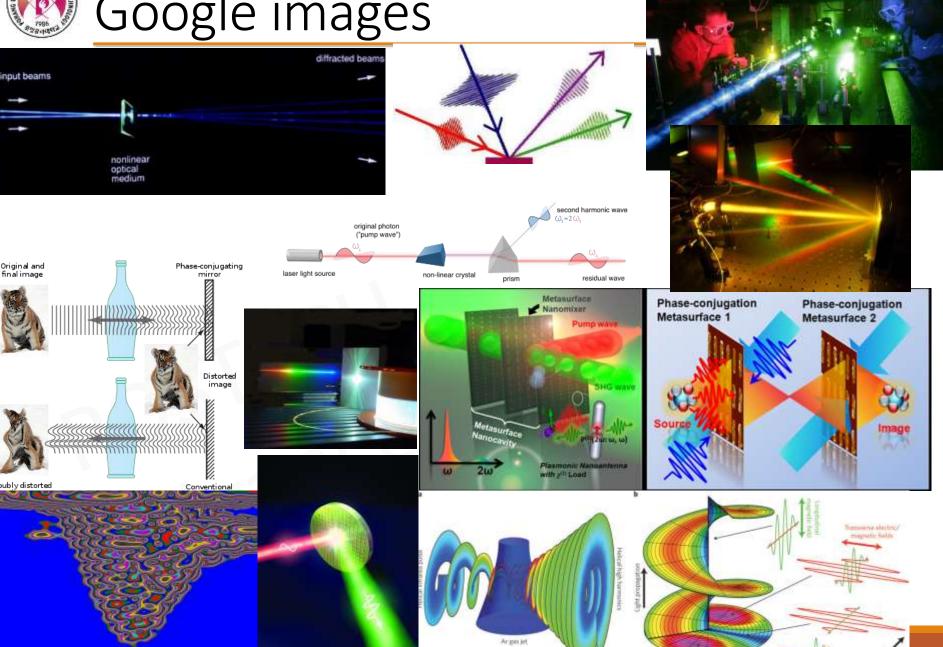




	PE WG	Ridge WG
Fundamental Wavelength for SHG (nm)	750~2100	780 ~ 1580
Power Handling (mW)	~200	>1000



Google images





Optical phase conjugation (OPC) LE AND TECHNOLOGY

 OPC is a totally degenerated case of the 4-wave mixing and is sometime refer to as degenerate 4-wave mixing:

$$\omega_1 = \omega_2 = \omega_3 = \omega_4 = \omega$$
.

• Consider two plane waves with $\mathbf{k}_4 = -\mathbf{k}_3$,

$$E_3(\mathbf{r}) = A_3 \exp(-j\mathbf{k}_3 \cdot \mathbf{r}), \qquad E_4(\mathbf{r}) = A_4 \exp(-j\mathbf{k}_4 \cdot \mathbf{r}),$$

The corresponding polarization is

$$P_{NL} = 6\chi^{(3)}A_3A_4E_1^*(\mathbf{r})$$

• And thus the E-field is

$$E_2 = 6\chi^{(3)} A_3 A_4 E_1^*(\mathbf{r})$$

- The E-field E₂ is conjugate of field E₁.
- The wave E_3 and E_4 are called pumps.

