#### Enhanced infrared absorption of spatially ordered quantum dot arrays

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**Presentation Outline** 

(1), Introduction
(a), advantages of QDIP
(b), disadvantages of QDIP

(2), Enhanced infrared absorption of spatially ordered QDs
 (a), growth
 (b), absorption measurements

(3), Summary

### Advantages of QDIP

- 1, Allowed normal incidence intersubband transition
- $2\,$  ,  $\,$  Longer electron lifetime  $\,$

background limited performance:  $\Phi > n_{th}/\alpha\tau$ ,

 $\alpha$ : absorption coefficient,  $\tau$ : electron lifetime,

 $n_{th}\!\!:$  thermal generated carrier density,  $\Phi\!\!:$  photon flux

M.A. Kinch, J. Electronic Materials, 29, 809 (2000)

M.A. Kinch and A. Yariv, Appl. Phys.Lett. 55, 2093 (1989)

0.75 ns, InGaAs/GaAs QDs,

J. Urayama et al, Phys. Rev. Lett. 86, 4930 (2001)

# **Disadvantages of present QDIP**

1, relatively low areal density of QDs (<10<sup>11</sup>cm<sup>-2</sup>) e.g. S. Charkrabarti *et al*, J. Phys. D: Appl. Phys. 38, 2135(2005)

2, normal incidence absorption efficiency is very low due to

disk-like dot shape

wave function coupled QDs,



e.g. A.M. Adawi et al, Appl. Phys. Lett. 82, 3415(2003)

3, large size fluctuation

# Theoretical calculation of detectivity, dark current for MCT, QWIP, QDIP



J. Phillips, J. Appl. Phys. 91, 4590 (2002)

#### **QDIP** sample

#### Reference sample







#### Hexagonally ordered QD arrays





# **X-sectional TEM image**



# Non-ordered QDs



 $1 \, \mu m$ 

# FTIR spectra using air as reference under nornal incidence geometry



# FTIR spectra using undoped sample as reference under nornal incidence geometry



# Absorption spectra under waveguide geometry



### **RT PL spectra**



## Symmetric (400) reflection of XRD



# Growth modes



Growth mode of self-assembled QDs

# InGaAs/GaAs QDs





- Best developed QD system
- Everywhere direct
- Lasers to 1.3 µm

# Why a hexagonal lateral ordering?

Vertical island-island interaction is attractive while lateral Island-island interaction is repulsive. Therefore, degree of the lateral ordering depends on the complicated tradeoff of the two counteracting elastic interactions.

(1), Multilayer growth  $\longrightarrow$  vertical stacking of islands.

(2), anisotropic adatom migration  $\rightarrow$  anisotropic strain relaxation  $\rightarrow$  islands lined up along the [0-11] direction.

(3), the lateral island-island interaction energy is at minimum for a hexagonal arrangement.

(4), to realize the lateral ordering, the adatoms need to respond to the small energy barrier change in a short enough time.

High T, small growth rate, small V/III ratio



# Linear chains



3 µm

# Summary

# Conclusions

- Enhanced infrared absorption is observed for spatially ordered QDs.
- Enhanced absorption is attributed to enhanced uniformity of QDs.

• For FTIR measurement under normal incidence geometry, using a undoped sample as a reference can remove the multiple reflection effect.

### Problem

Smaller areal density for spatially ordered QDs

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$$\begin{split} E &= \frac{f_{OA}}{OA^3} + \frac{f_{OB}}{OB^3} + \frac{f_{OC}}{OC^3}, \\ OM &\equiv r_0 \text{ and } OA \equiv \eta r_0, \text{ then } OB = r_0 \cos^{-1} \alpha \text{ and } OC = r_0 \sigma^{\frac{1}{2}}, \text{ where } \sigma \equiv 1 + (\eta - \tan \alpha)^2. \\ E &= (\eta^{-3} f_{OA} + \cos^3 \alpha \cdot f_{OB} + \sigma^{-\frac{3}{2}} f_{OC}) r_0^{-3} \Rightarrow \frac{\partial E}{\partial \alpha} = -3r_0^{-3} (f_{OB} \cos^2 \alpha \sin \alpha + \frac{1}{2} f_{OC} \sigma^{-\frac{5}{2}} \frac{\partial \sigma}{\partial \alpha}). \end{split}$$
  
For a hexagonal ordering where  $\tan \alpha = \frac{1}{2}\eta$ , we get

$$\frac{\partial E}{\partial \alpha} = 3\cos^2 \alpha \sin \alpha (1 - f_{OB}/f_{OC}) \cdot f_{OC} \cdot r_0^{-3}.$$
 (1)

