

# Time-resolved photoluminescence spectroscopy of Si nanocrystals prepared by SiO<sub>x</sub>/SiO<sub>2</sub> superlattices technique

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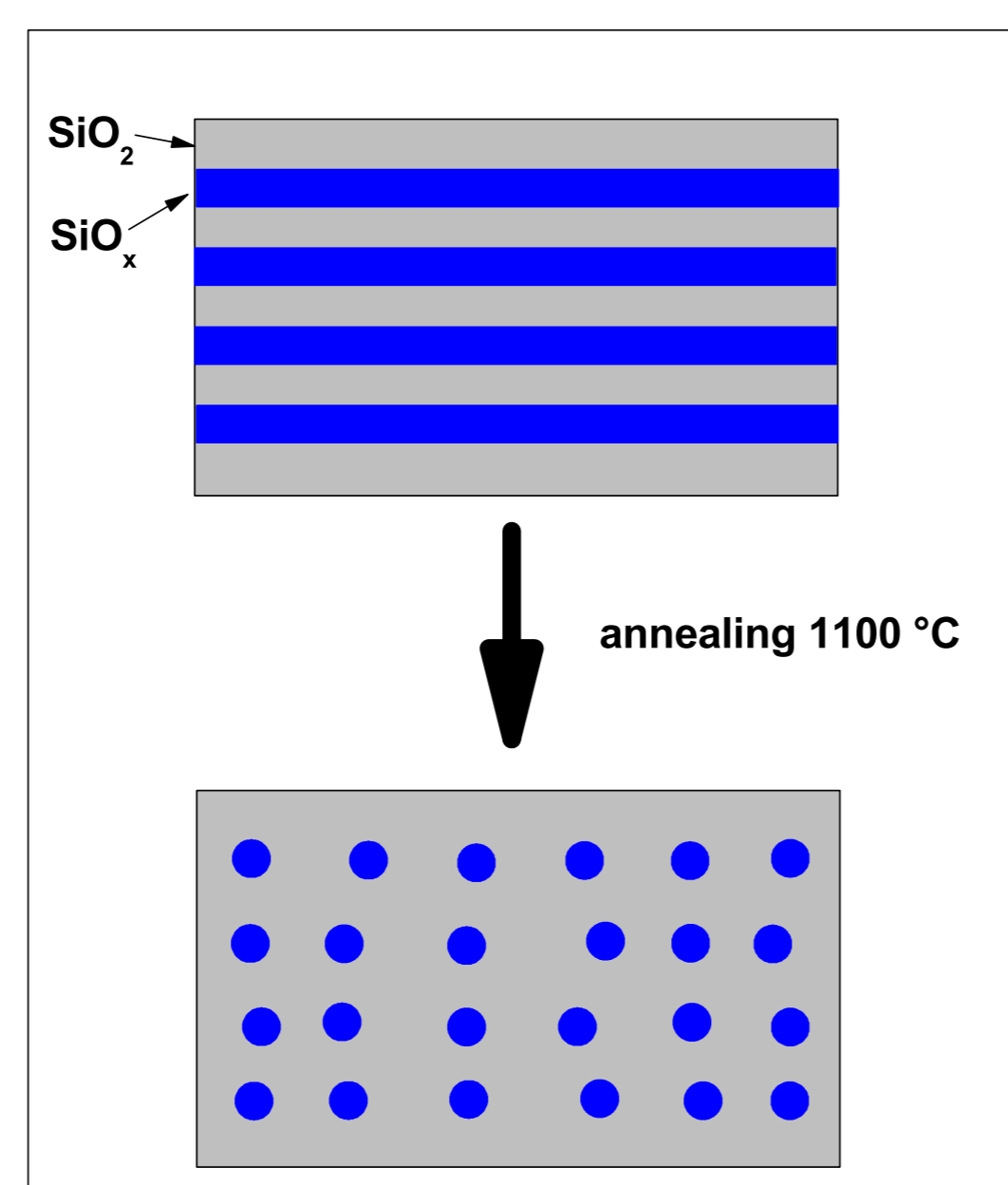


## Introduction

- ▶ **Si nanocrystals** (Si-NCs) are promising material for application in optoelectronics [1], biophysics [2] and photovoltaics (**the third generation solar cells** [3]).
- ▶ It is generally accepted that the unique properties of Si-NCs result from both **the quantum confinement effect** and presence of **the surface states**. The predominantly used matrix material for Si-NCs is SiO<sub>2</sub> due to its high band gap and the well understood properties of the bulk Si/SiO<sub>2</sub> interface [4].
- ▶ The aim of this study is **time-resolved photoluminescence spectroscopy** of Si-NCs prepared by SiO<sub>x</sub>/SiO<sub>2</sub> superlattices technique for photovoltaic application.

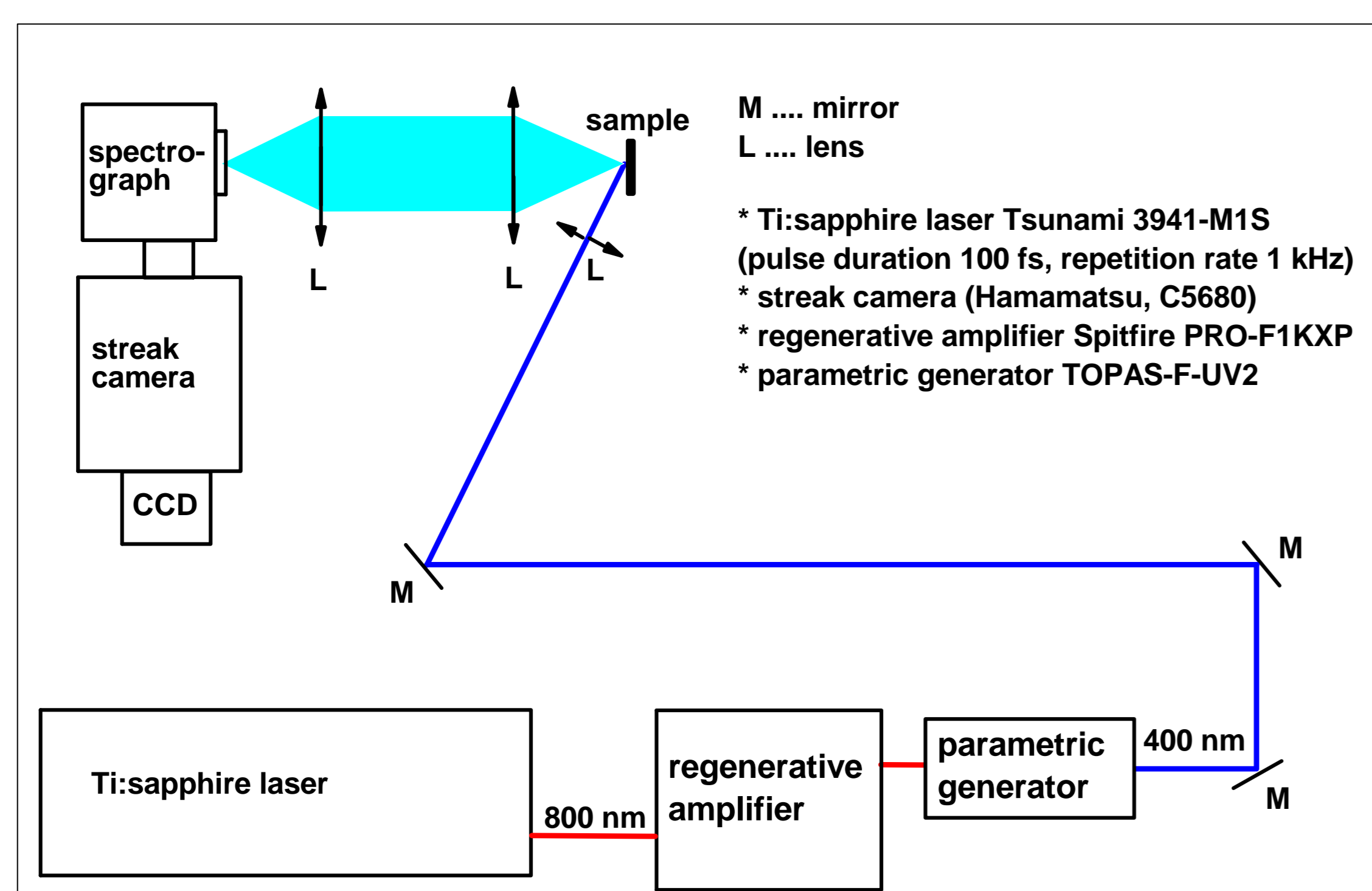
## Samples

- ▶ samples Da125H2 and Da125 fabricated by IMTEK (Freiburg)
- ▶ amorphous SiO<sub>x</sub>/SiO<sub>2</sub> superlattices prepared by reactive evaporation of SiO powders in oxygen atmosphere
- ▶ annealing at 1100 °C in N<sub>2</sub> for 1 hour
- ▶ sample Da125H2 additionally post-annealed in H<sub>2</sub> at 450 °C
- ▶ the high temperature annealing results in phase separation  
 $\text{SiO}_x \rightarrow \frac{x}{2} \text{SiO}_2 + (1 - \frac{x}{2}) \text{Si}$  [5]
- ▶ 30 layers of ~ 3.5 nm nanocrystals



Two samples with well defined nanocrystal size, density and stoichiometry which differ in the surface quality.

## Experimental setup



## Results

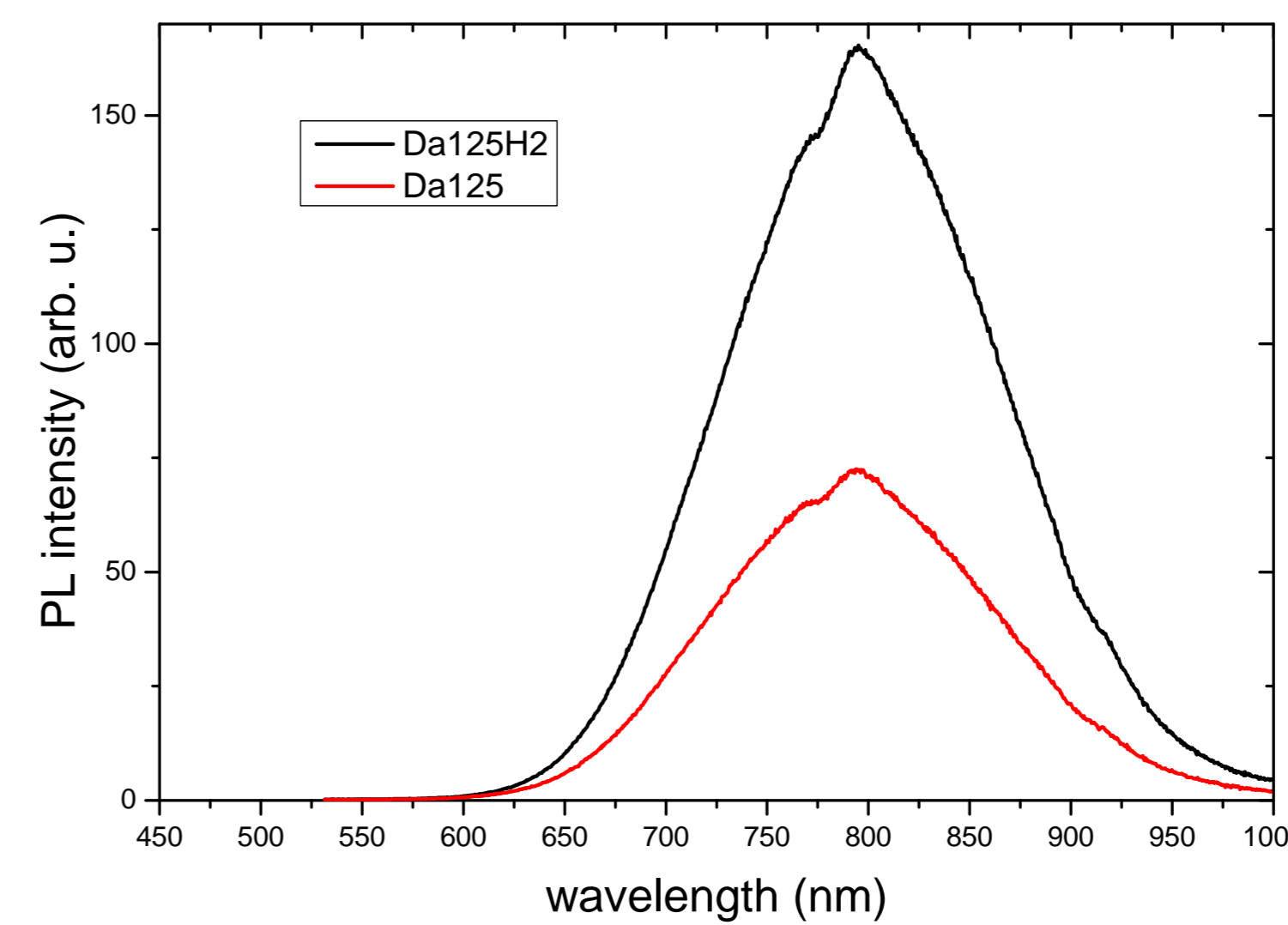


Fig.1: Time-integrated PL spectra of the investigated samples, excited by a 325 nm cw He-Cd laser.

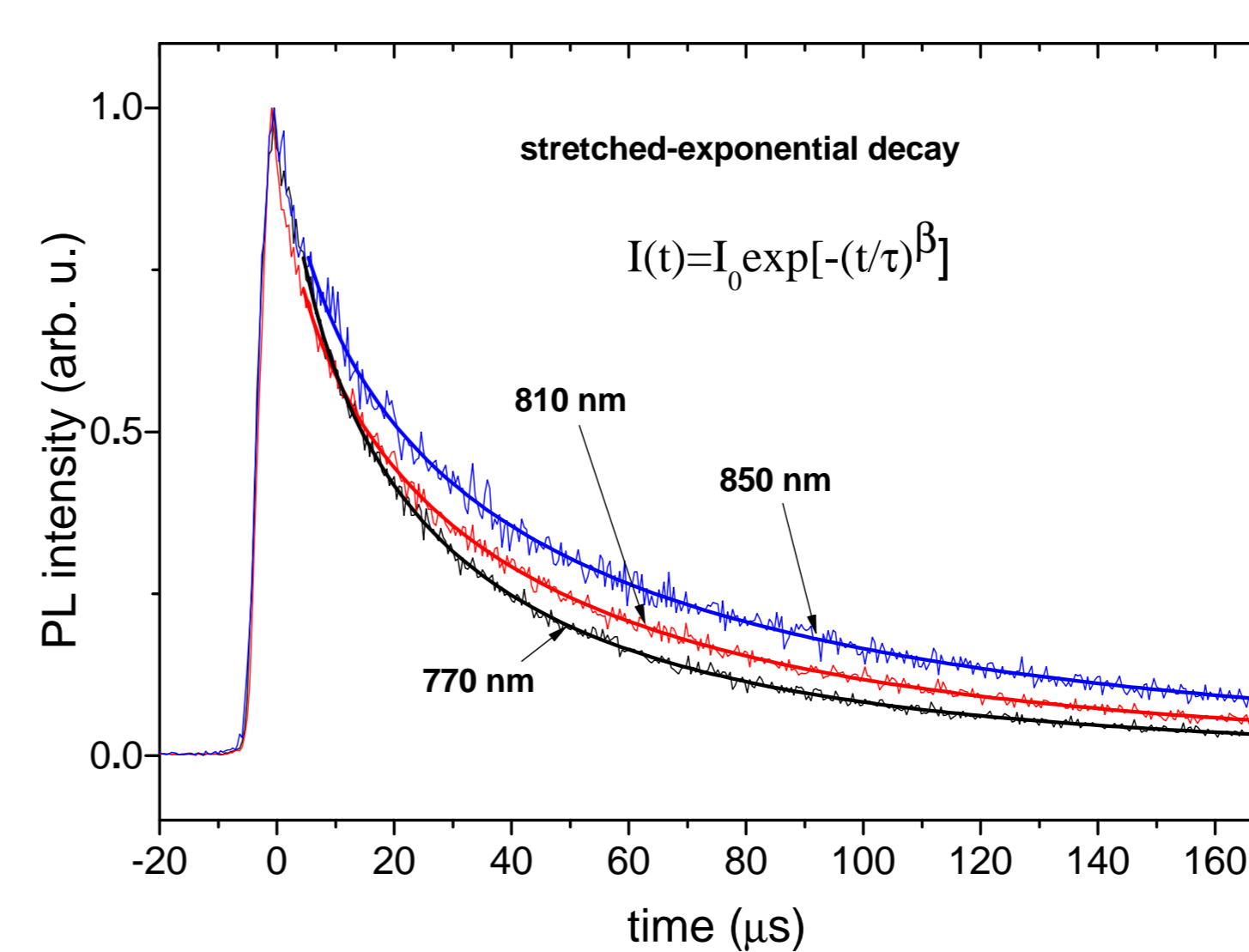


Fig.2: Dynamics of microsecond PL of the sample Da125H2.

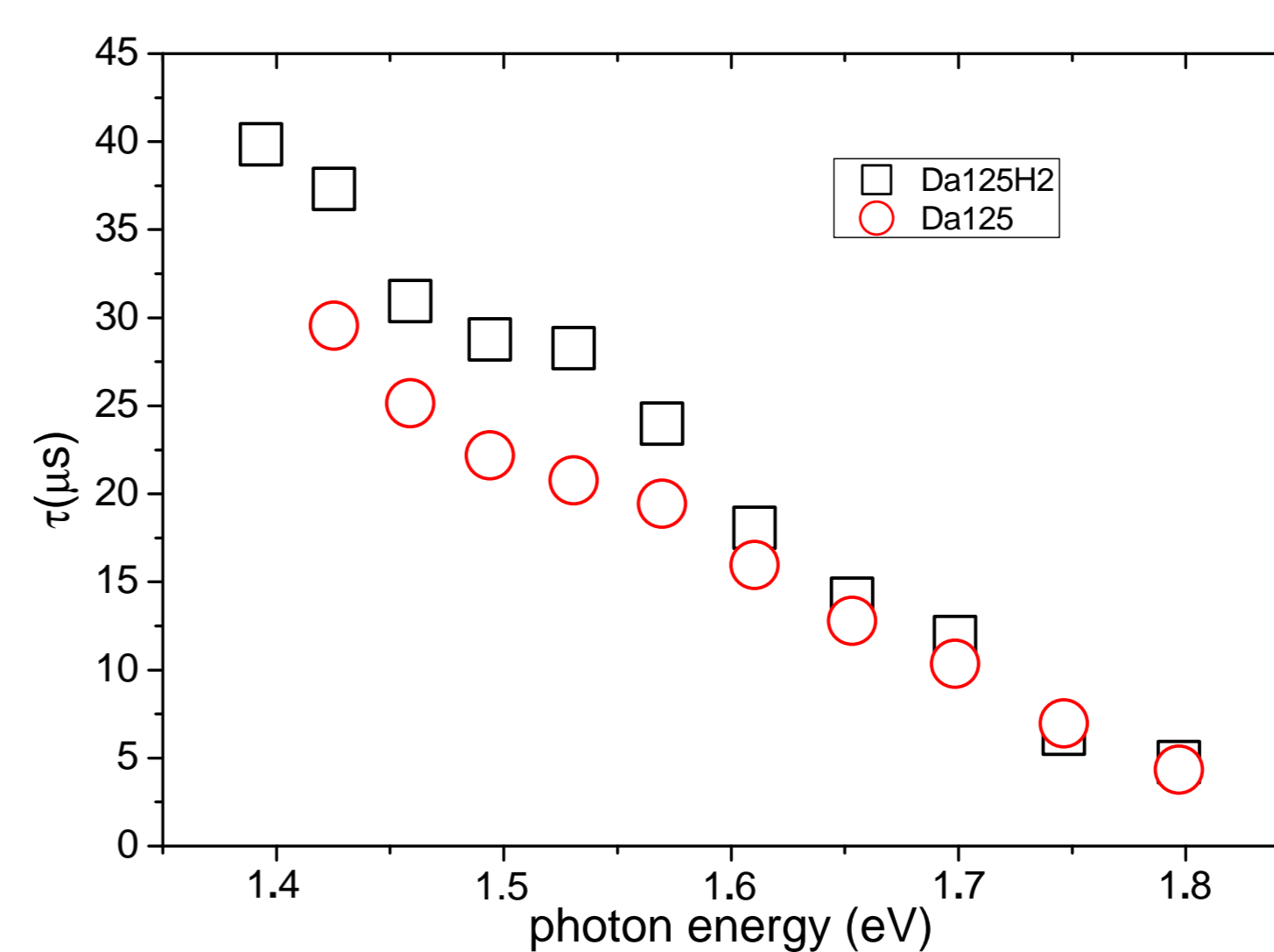


Fig.3: Parameter  $\tau$  of stretched-exponential function vs the PL photon energy.

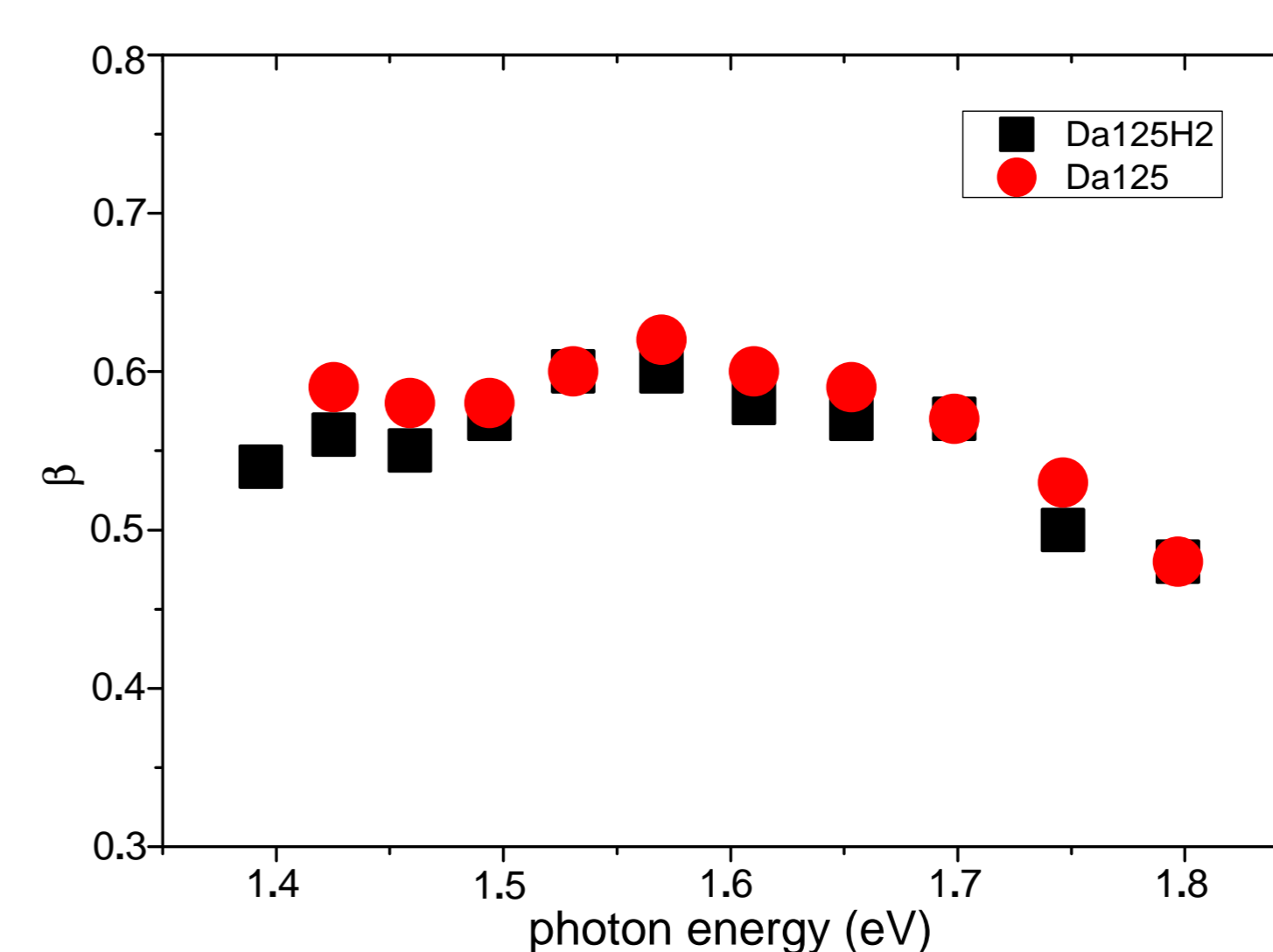


Fig.4: Parameter  $\beta$  of stretched-exponential function vs the PL photon energy.

## Proposed explanation

- ▶ H<sub>2</sub> annealing = passivation of nonradiative defects at the NC/oxide interface (Si dangling-bond defect [6])  
⇒ PL intensity increase  
⇒ time constant  $\tau$  increase
- ▶ exciton migration (hopping) between different NCs [7]  
⇒ stretched-exponential time decay (dispersion factor  $\beta$  depends on the spatial distribution of localized states)

## Acknowledgement

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## References

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