

# Femtosecond pump-probe studies of zinc phthalocyanine in DMSO

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## INTRODUCTION

Phthalocyanines are new class of photosensitiser used for photodynamic therapy. These drugs are used to treat small and superficial tumours. In recent years, a particular attention has been given to phthalocyanine in order to search suitable photosensitisers that can be clinically used as photodynamic cancer drug. The zinc phthalocyanine (ZnPc) has been reported to have suitable photophysical properties in photodynamic therapy. Recent work done on zinc phthalocyanine dynamic has proven that the energy transfer from singlet state to triplet state occurs on ultrafast timescale and the determination of the ultrafast component remains a challenge.

## MECHANISM OF PHOTODYNAMIC THERAPY

The processes of light absorption and energy transfer are at the heart of the photodynamic therapy. It is based on introducing a tumour localising photosensitiser and then excites the photosensitisers with an appropriate wavelength. There be an energy transfer from singlet to triplet via inter-system crossing. By quenching of the triplet state of the photosensitiser, there will be an energy transfer which will generate an electronically-excited state of the singlet state of the oxygen. The excited state of the oxygen will lead to the destruction of the tumour.

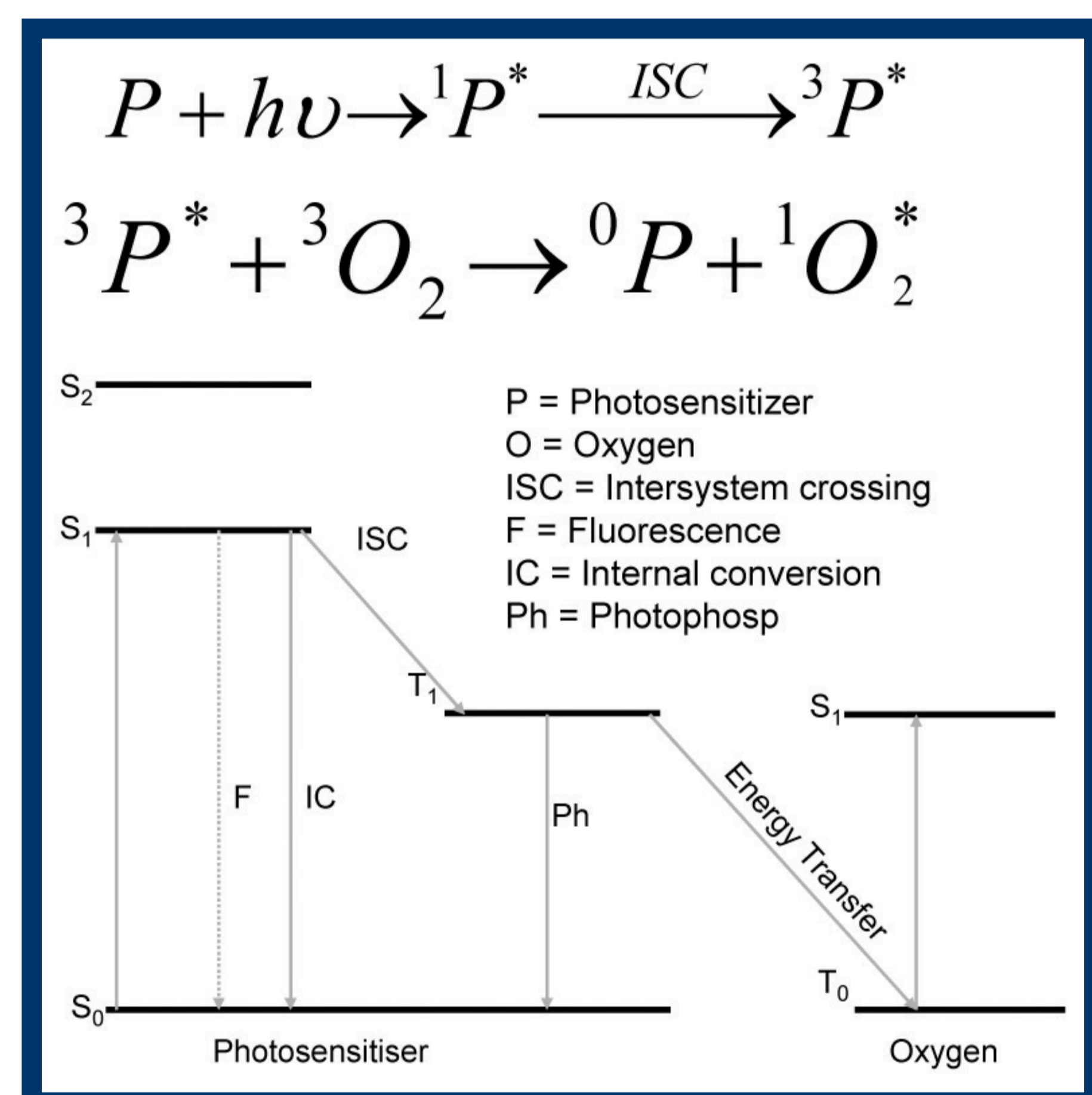


Figure 1: Energy transfer and energy level diagram of zinc phthalocyanine

## PUMP-PROBE TECHNIQUE

Figure 2 indicates the Femtosecond pump-probe technique used to investigate the ultrafast dynamics ZnPc.

- Start process with an intense ultrashort pump pulse.
- Probe process after  $t_D$  with a weaker ultrashort probe pulse. Slow detector.
- Repeat experiment with different delay times  $t_D$ .
- Follow the transient concentration.

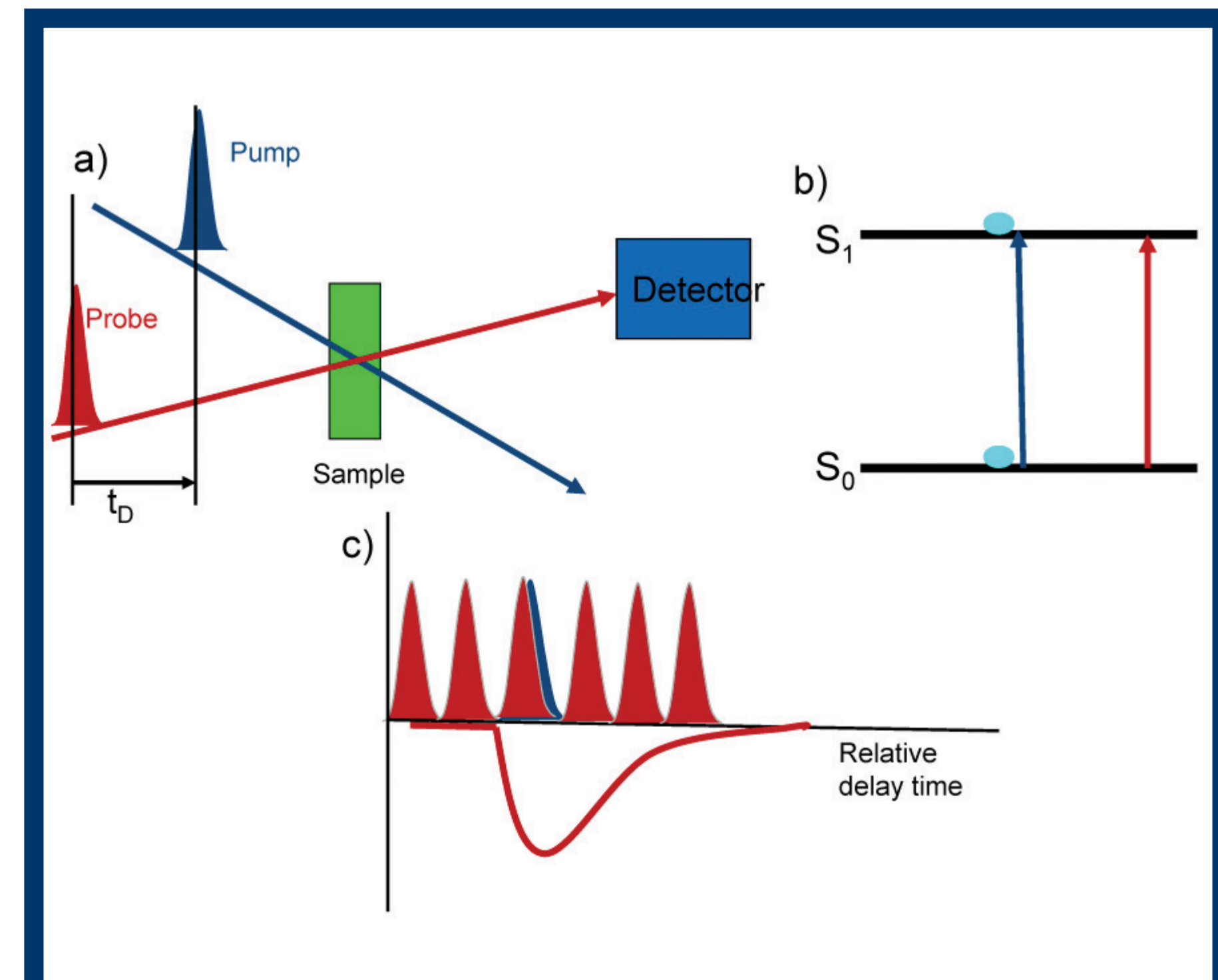


Figure 2: Femtosecond pump-probe technique : a) The overlap of pump and probe beams, b) Energy level diagram showing the pump beam exciting molecules from ground state to the first excited state and the probe. c) kinetics of the sample

## EXPERIMENTAL SETUP

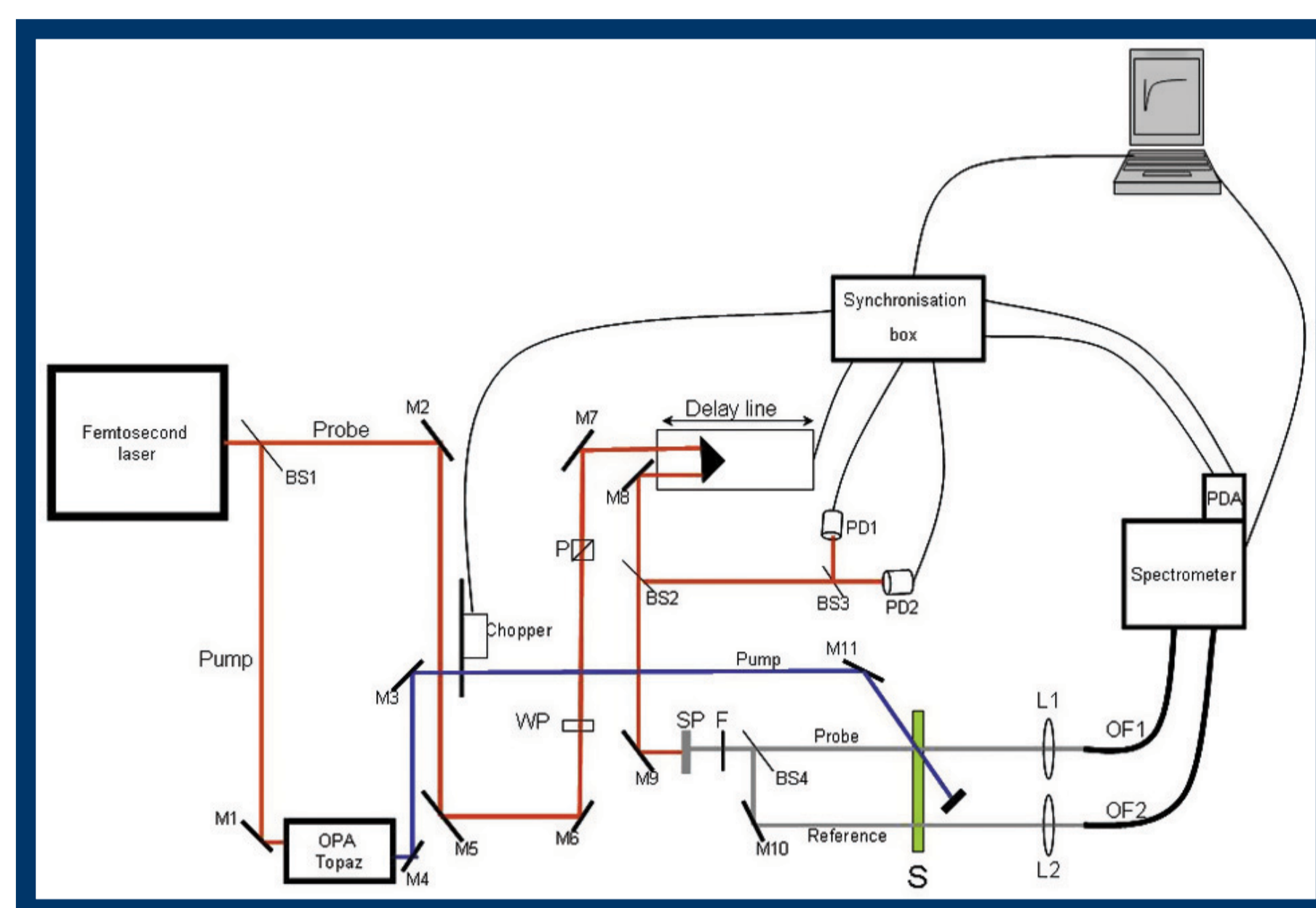


Figure 3: Experimental setup used to investigate the energy transfer in ZnPc

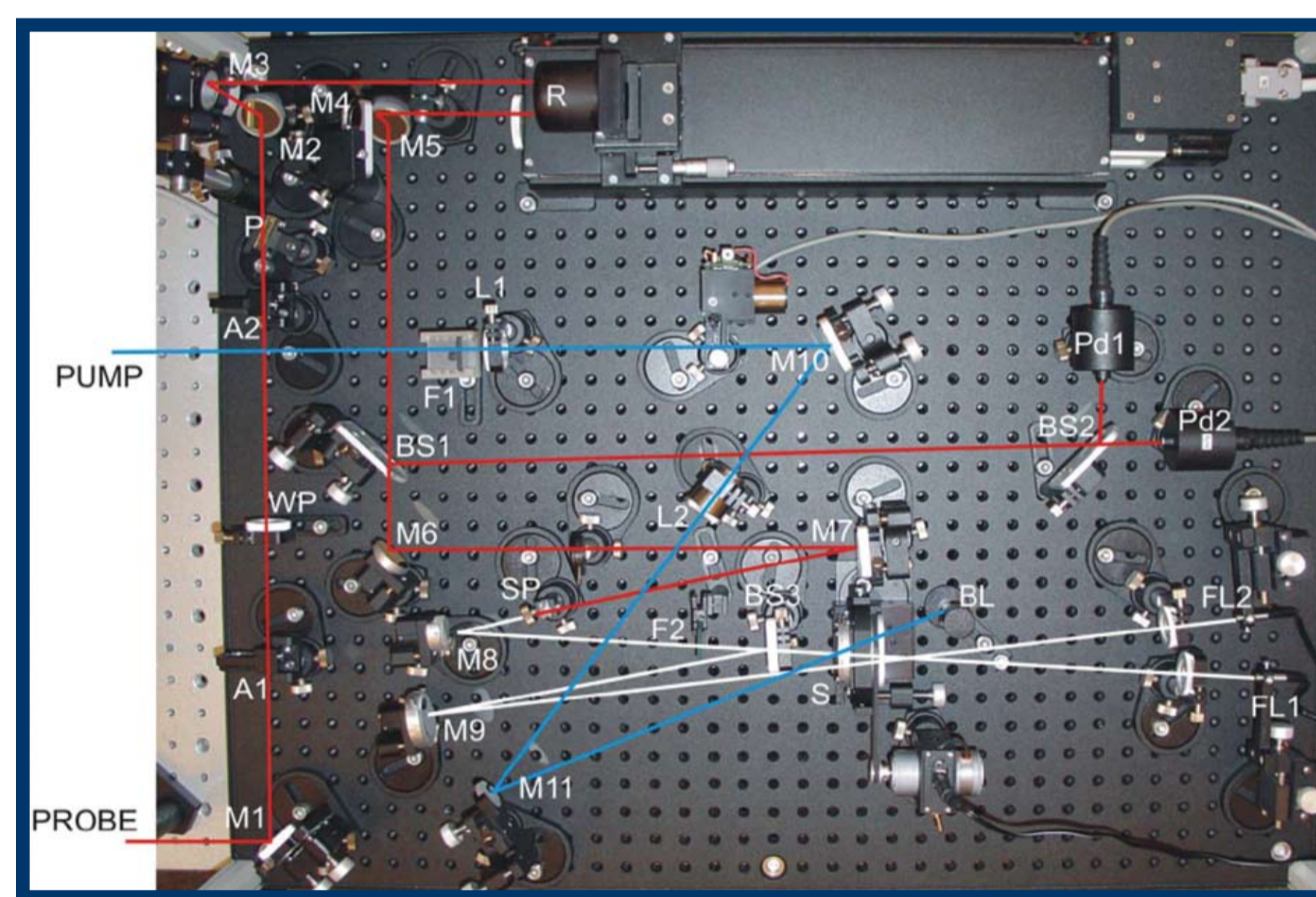


Figure 4: Image of the pump-probe setup

## RESULTS

The zinc phthalocyanine sample was pumped at wavelength 625 nm and probed by a white light continuum. The results obtained in our experiment show a good fit to a bi-exponential decay at the 675 nm band peak, comprising a fast process with a time constant of  $1.82 \pm 0.06$  ps and a slower process with a time constant of about  $38.9 \pm 2.6$  ps.

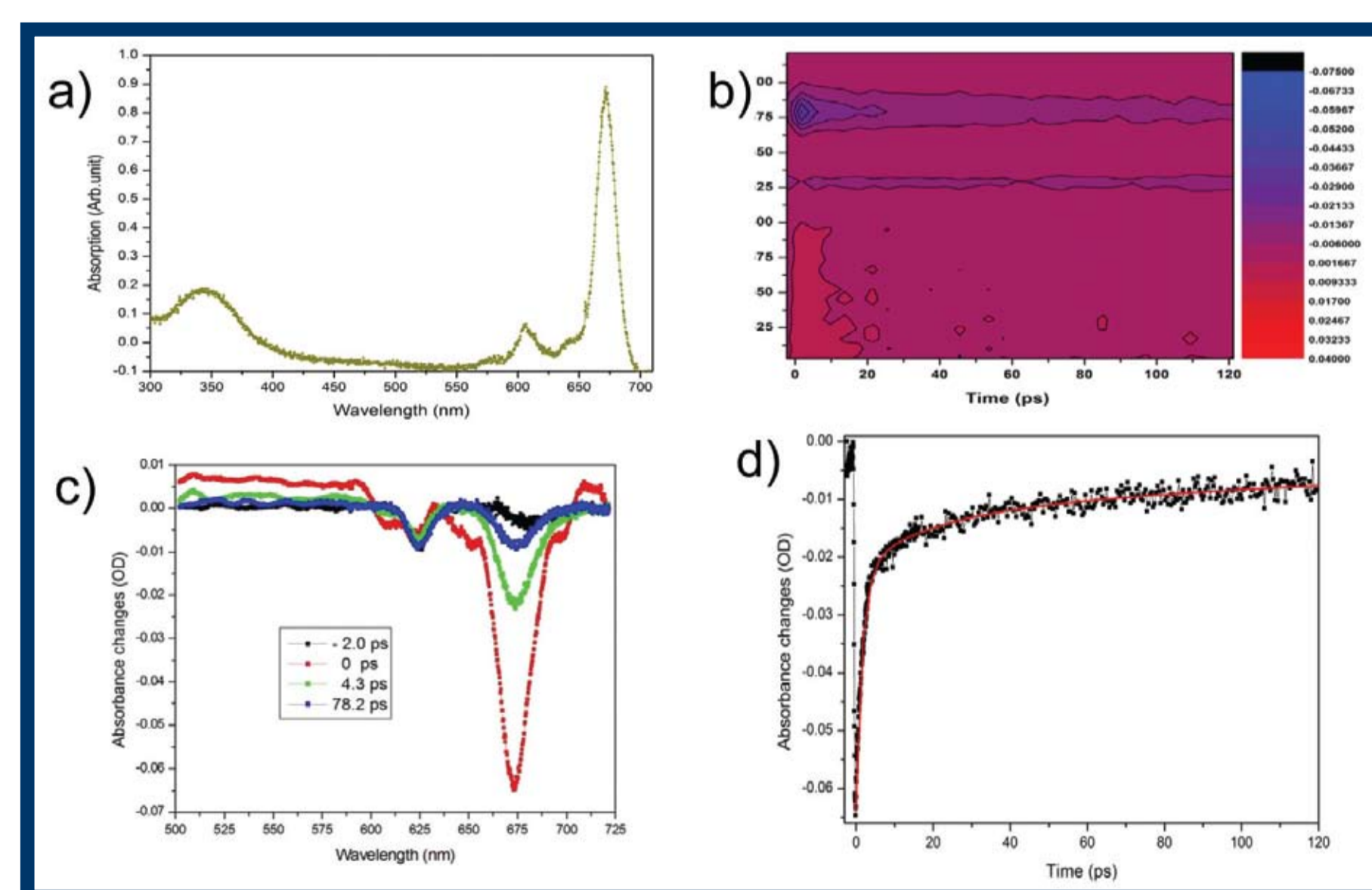
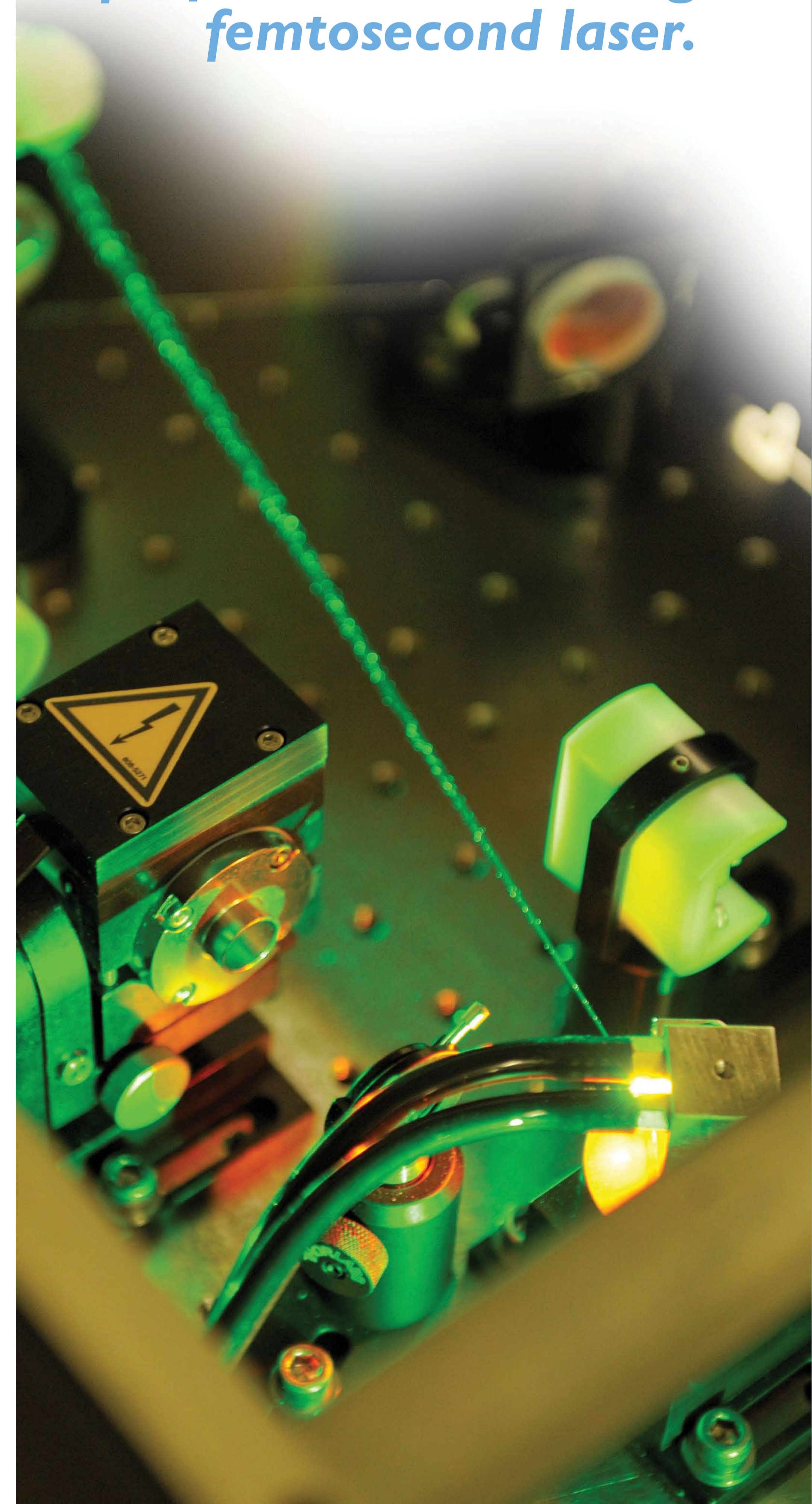


Figure 5: a) A steady state absorption spectrum of ZnPc in DMSO at a concentration of  $10 \mu\text{M}$  acquired using a Xe-Kr lamp combined with USB2000 ocean optics spectrometer Image of the pump-probe setup. b) Transient absorption contour graph indicating measured absorbance spectra for a range of delay times between pump and probe pulses. Pump was set at 625nm and probe was a white light continuum. c) Transient absorption spectra of ZnPc in the indicated wavelength interval and time delays with the pump wavelength set at 625 nm. d) Absorbance change kinetics as a function of relative delay time between pump and probe pulses for ZnPc at 675 nm

## CONCLUSION

We were able to measure the transient absorption of Zinc Phthalocyanine in DMSO and bi-exponential fit on the kinetics data allow us to extract time constants for a fast and slow processes. However, a difference between our results and those reported literature was noticed. This could be attributed to the fact that in our setup we could not resolve time constants below 300 fs.

CSIR researchers are investigating drugs that can treat small and superficial tumours using femtosecond laser.



## REFERENCES

1. Ogunsiye, A., Chen, J., and Nyokong, T., (2004). *Photophysical and Photochemical studies of zinc(II) phthalocyanine derivatives - effect of substituents and solvent*, New J. Chem. 28, 822-827.
2. Nunes, S.M.T., Sguilla, F.S., and Tedesco, A.C., (2004). *Photophysical studies of zinc phthalocyanine and chloroaluminum phthalocyanine incorporated into liposomes in the presence of additives*, Braz. J. Med. Biol. Res. 37, 273-284.
3. Savolainen, J., van der Linden, D., Dijkhuizen, N., and Herek, J.L., (2008). *Characterizing the functional dynamics of zinc phthalocyanine from femtoseconds to nanoseconds*, J. Photochem. Photobiol., A 196, 99-105.
4. Ombinda-Lemboumba, S., du Plessis, A., Sparrow, R.W., Molukanele, P., Botha, L.R., Rohwer, E.G., Steenkamp, C.M., and van Rensburg, L., (2009). *Femtosecond pump-probe spectroscopy for the study of energy transfer of light-harvesting complexes from extractions of spinach leaves*, S. Afr. J. Sci. 105, 376-386 (2009).
5. Howe, L., and Zhang, J.Z., (1997). *Ultrafast studies of excited-state dynamics of Phthalocyanine and zinc phthalocyanine tetrasulfonate in solution*, J. Phys. Chem. A 101, 3207-3213.
6. Rao, S.V., and Rao, D.N., (2002). *Excited state dynamics in phthalocyanines studied using degenerate four wave mixing with incoherent light*, J. Porphyrins Phthalocyanines 6, 233-237 (2002).