# Photoinduced phenomena in fullerene-doped PDLC: potentials for optoelectronic applications

N.V. KAMANINA\*

Vavilov State Optical Institute, 12 Birzhevaya Line Str., St. Petersburg, 199034 Russia

Laser-induced change in refractive index and optical limiting effect have been studied in the liquid crystal systems based on fullerene-doped 2-cyclooctylamino-5-nitropyridine, polyimide, N-(4-nitrophenyl)-(L)-prolinol compounds. Experiments have been made under nano-, pico- and femtosecond pulsed laser irradiation at the wavelength of 532 nm and 805 nm. From the results obtained, both the optical limiting level and nonlinear refractive index  $n_2$  as well as the third-order susceptibility  $\chi^{(3)}$  have been determined. Potentials of the systems studied to attenuate laser irradiation and to record thin amplitude-phase hologram over the visible and near-infrared spectral ranges have been evaluated to solve optoelectronic problems more efficiently.

Keywords: liquid crystal, fullerenes, optical limiting, laser-induced change in refractive index, optoelectronics.

## 1. Introduction

At present, the polymer-dispersed liquid crystals (PDLCs) are extensively being studied [1–3] for various optoelectronic applications. These systems combine film-forming capability and mechanical strength of polymers and unique electro-optical properties of liquid crystals mesophase.

Generally, PDLC is formed from liquid crystal and polymer. In this case the ordinary refractive index of the liquid crystal  $n_0$  and the refractive index of polymer  $n_p$  are close together [1]. Initially, as a result of random LC director orientation, there is a gradient of the refractive index at the interface between an LC drop and polymer that causes drastic light scattering by the composite. On application of an electrical field or under laser irradiation, the gradient becomes small, i.e., the condition  $n_0 \cong n_p$  is met. In this case, the LC director aligns itself along the electric field or the electric vector of a light wave. As the result, the light scattering is not observed and the system clarification occurs. After switching off the electrical or light action, the composite reverts to the original state.

Until now the different fullerene-doped photosensitive components have been used to develop PDLC structures. Poly(vinyl alcohol), poly(methyl methacrylate) [4,5], 2-cyclooctylamino-5-nitropyridine (COANP) [6,7], polyimide [8], phthalocyanine nanocrystals [9], etc. have been treated for this purpose. A fullerene introduction has been used to control PDLC systems to good advantage. The fullerene-doped systems allowed them to be applied to process optical information, to attenuate a laser irradiation and duplicate and triple the laser frequency. Moreover, these systems could be used as laser valves over visible and infrared spectral range due to good sensitisation ability of fullerenes.

In this paper the photoinduced phenomena have been investigated in the C<sub>70</sub>-COANP-LC system, C<sub>60</sub>-N-(4-nitrophenyl)-(L)-prolinol (NPP)-LC compound, C<sub>70</sub>-polyimide--LC structure under nano- and pico-second pulsed laser irradiation over the visible spectral range. From the results obtained, the nonlinear refractive index  $n_2$  and the third-order susceptibility  $\chi^{(3)}$  have been determined.

# 2. Experiment

Fullerene free COANP were thoroughly studied in Refs. 10 and 11. When COANP was doped with a dye, an additional absorption peak was found [10]. It should be noticed that the absorption edge of pure COANP lies in the range of 420–430 nm. When COANP was doped with fullerenes, the absorption edge was shifted to long wave region [12]. The PDLC cells based on fullerene-doped COANP allowed the laser irradiation to be modulated and attenuated efficiently [13]. Additional bands, which were close to 532 and 810 nm, were obtained. The absorption spectra for films of the pure COANP composite and the C<sub>70</sub>-doped COANP composite, as well as pure C<sub>70</sub> were presented in Ref. 14. Fullerene influence on spectral and energy ranges of the polyimide 6B [15] and NPP [16] molecules was discussed recently.

With UV stirring, the initial mixture of polymer PI6B and nematic liquid crystal (NLC) was prepared in the ratio of 3 polymer parts to 2 NLC ones. 3% solutions of PI6B in 1,1,2,2-tetrachloroethane (TCIE) were used. PDLCs based on COANP and NPP structures were made in the ratio of 1 photosensitive component to 2 NLC one. A 2.5% solution

<sup>\*</sup>e-mail: kamanin@ffm.ioffe.ru

of non-photosensitive polyimide 81A in TCIE was used as a film-forming base for COANP and NPP compounds. The relation between COANP and NPP compounds and the film-forming base was 2:1. The C<sub>70</sub> and C<sub>60</sub> concentration in COANP and NPP materials was 5 wt.% and 1 wt.%, respectively. 0.2–0.5 wt.% C<sub>70</sub> was used to dope polyimide 6B. 10 µm films were prepared from emulsion of fullerene-doped photosensitive component and LC. Two types of NLC with positive optical and dielectric anisotropy: ZhK1289 ( $\Delta n = 0.168$ ,  $\Delta \varepsilon = 10$ ) and E7 BDH ( $\Delta n = 0.224$ ,  $\Delta \varepsilon > 0$ ) were used.

A sinusoidal holographic grating was recorded by the second harmonic ( $\lambda = 532$  nm) of a pulsed Nd laser with the pulsewidth of both 20 ns and 400 ps. Two laser beams formed a spot of 4–5 mm in diameter on the film surface. The recording energy density was 0.5–40 mJcm<sup>-2</sup>. The spatial frequency was 100 mm<sup>-1</sup>. The grating was read out under the self-diffraction conditions. OL experiments were made at the wavelength of 532 nm in the nanosecond pulsed regime and using femtosecond pulsed irradiation of a quasi-CW Ti-sapphire laser at  $\lambda = 805$  nm.

# 3. Results and discussion

## 3.1. Holographic recording study

In this part of the paper, the emphasis will be given to investigate the PDLC structure based on COANP. The dependence of the response in the first diffraction order  $\eta^*$  on the input laser energy density  $W_{in}$  is shown in Fig. 1.

It should be noticed that the response obtained under nanosecond laser irradiation (curve 1) is more than the one obtained under picosecond laser irradiation (curve 2). However, the reversible process in the picosecond range has been obtained at the higher energy density than the one observed in the nanosecond range. This effect is associated with rejection of the thermal nonlinearity and determines the potentials to irradiate the media in the picosecond range at the higher  $W_{in}$ . It should be noticed that the response of the non-sensitized system was at least two orders of magni-



Fig. 1. Dependence of the response in the first diffraction order  $\eta^*$ on the input laser energy density  $W_{in}$  for PDLC based on C<sub>70</sub> (5 wt.%)-doped COANP at the laser pulsewidth: (1) 20 ns and (2) 400 ps,  $\lambda = 532$  nm.

tude less than the one of the fullerene-doped system. The increase in the diffractive efficiency and hence in the induced refractive index could be explained from the spectral and thermal peculiarities of the samples, as well as in the framework of the photorefractive effect under the laser irradiation.

From the data presented in Fig. 1, the induced change in refractive index could be estimated. The first diffraction order response was changed from  $2 \times 10^{-4}$  to  $3.7 \times 10^{-3}$  mJ and from  $10^{-4}$  to  $2 \times 10^{-3}$  mJ as the incident energy density increased from 2.5 to 17.5 mJcm<sup>-2</sup> (nanosecond irradiation) and from 4.2 to 35 mJcm<sup>-2</sup> (picosecond irradiation), respectively. Therefore, using equation [17]

$$\eta = I_1 / I_0 = (\pi \Delta n_i d / 2\lambda)^2, \tag{1}$$

the light-induced change in refractive index could be estimated to be  $7.3 \times 10^{-4} - 1.4 \times 10^{-3}$  and  $5.1 \times 10^{-4} - 7.2 \times 10^{-4}$  un-

Material	Impurity concentration (wt.%)	Wavelength (nm)	Light intensity (Wcm <sup>-2</sup> )	Incident energy density (Jcm <sup>-2</sup> )	Laser pulsewidth (ns)	Change of refractive index $(\Delta n)$	Ref.
COANP + TCNQ*	0.1	676	2.2			2×10 <sup>-5</sup>	[10]
Pure COANP**	0	532		0.9	20	~10 <sup>-5</sup>	[14]
$COANP + C_{70}$	5	532		0.9	20	6.89×10 <sup>-3</sup>	[14]
PDLC based on COANP + $C_{70}$	5	532		17.5×10 <sup>-3</sup>	20	1.4×10 <sup>-3</sup>	[6]
Liquid crystal	0	514.5	0.2			0.16×10 <sup>-3</sup>	[18]

Table 1. Light-induced change of refractive index for materials based on COANP.

\* 7,7.8.8-tetracyanoquinodimethane.

\*\* Polyimide 81A was used to prepare the thin COANP films as a film-forming base.

der nanosecond and picosecond laser irradiation, respectively. Here  $\eta$  is the diffraction efficiency,  $\Delta n_i$  is the change in a refractive index of the medium, *d* is the thickness of the sample,  $\lambda$  is the wavelength,  $I_1$  and  $I_0$  are the light wave intensities in the first order and incident one, respectively, The values obtained was more than the thermal component typical of LCs (~10<sup>-5</sup>).

The light-induced changes of refractive index for different materials based on COANP are presented in Table 1.

It should be noticed that the  $\Delta n$  value obtained in the present paper for the fullerene-doped PDLC is one order of magnitude more than that for LC given in Ref. 18. Therefore, the fullerene introduction is likely strengthening the alignment of the LC molecule director. The possible reorientation mechanism of LC director using both the fast rotation ability of fullerenes and the complex formation process in conjugated organic structures has been recently discussed in Refs. 6, 7, and 9.

The investigation on the rotational behaviour of fullerenes [19,20] demonstrated that the reorientation time depended on temperature. The rotational reorientation of  $C_{60}$  and  $C_{70}$  in various solvents exhibited the following values of the reorientation time: in toluene  $-7 \pm 1.5$  ps, decalin  $-3.5 \pm 1.5$  ps, and in chlorobenzene  $-8 \pm 2$  ps. Therefore, the response of fullerene on changes of the light wave is very fast. Moreover, fullerene introduction in the COANP compounds provokes the charge-transfer complex with the donor fragment of CONP molecule [14,21]. It should be noticed that fullerene has electron affinity energy of 2.65 eV that is more than the one for most organic compound acceptors. For example, for COANP compounds, NO<sub>2</sub> group (as an acceptor fragment) has electron affinity only of 0.54 eV. Therefore fullerene dominates under acceptor fragment of COANP and creates the complex with its donor fragment. The possible scheme of donor-acceptor interaction in the fullerene-doped organic structures is shown in Fig. 2. The quantum-chemical calculation of the charge-transfer in COANP-fullerene structure has been made in the Ref. 21. To indicate this complex experimentally the mass-spectrometry and photoconductive data have

been done in Ref. 22. This complex has larger dipole moment, which interacts with light wave to align LC director with good advantage.

It should be noticed that from large induced change in refractive index for PDLC based on fullerene-doped COANP it predicts more nonlinear refraction index  $n_2$ . Both  $n_2$  and the nonlinear susceptibility  $\chi^{(3)}$  could be calculated from the following equations [23]

$$n_2 = \frac{\Delta n_i}{I},\tag{2}$$

$$\chi^{(3)} = \frac{n_2 n_0 c}{16\pi^2},\tag{3}$$

where  $n_0$  and  $n_2$  are linear and nonlinear refractive indices, respectively, *c* is the velocity of light; and *I* is the intensity of the light wave. Thus,  $n_2$  and  $\chi^{(3)}$  for PDLC based on 5 wt.% C<sub>70</sub>-doped COANP are  $1.6 \times 10^{-9}$  cm<sup>2</sup>W<sup>-1</sup> and  $4.86 \times 10^{-8}$  esu, respectively, at the incident energy density of 17.5 mJcm<sup>-2</sup> and  $\Delta n = 1.4 \times 10^{-3}$ . The data obtained are close to those for silicon ( $10^{-10}$  and  $10^{-8}$ , respectively [23]), but less than the ones for pure liquid crystals. However, the nonlinear effect in the structures under present investigation could be accumulated and hence increase significantly. The nonlinear coefficients obtained demonstrate potentials of PDLC based on fullerene-doped COANP for its optoelectronic applications as efficient nonlinear optical media.

Under the Raman-Nath diffraction conditions the thin amplitude-phase holograms in the PDLC structures based on the fullerene-doped NPP compounds have been recorded for comparison. The corresponding curves are presented in Fig. 3. The concentration of fullerene  $C_{70}$  in COANP was 5 wt.% and the concentration of  $C_{60}$  in NPP was 1 wt.%. Note, although the concentration of fullerene  $C_{70}$  in COANP is greater than that of  $C_{60}$  in NPP, curves 1 and 2 almost coincide. This is probably explained by the



Fig. 2. Possible scheme of donor-acceptor interaction in the fullerene-doped organic structures.



Fig. 3. Dependence of the first diffraction order response ( $\eta^*$ ) on the input laser energy density ( $W_{in}$ ) for PDLC based on COANP with 5 wt.% C<sub>70</sub> (1) and NPP with 1 wt.% C<sub>60</sub> (2). Spatial frequency was 100 mm<sup>-1</sup>,  $\lambda = 532$  nm.

fact that nonlinear absorption of radiation by the medium in the visible spectral range is determined not only by the reverse saturated absorption but also by a close location of the resonance line of fullerene  $C_{60}$  (565 nm) with respect to the 532 nm laser wavelength.

It should be mentioned that twice increase in the laser-induced refractive index has been observed as the laser energy density increased tenfold. The large laser-induced change in refractive index influences the OL effect in PDLCs based on photosensitive molecules mentioned above (due to energy loses by diffraction), predicts the large increase in nonlinear coefficient  $n_2$  and  $\chi^{(3)}$  and therefore gives an opportunity to apply these systems for conversion of laser radiation. Moreover, the change in refractive index could increase by an electrical control.

# 3.2. Optical limiting study

The study of the OL effect allows the laser-matter interaction to be investigated and the search for the new materials for optoelectronic problems (such as the protection of human eyes and technical devices against high laser radiation) to be made.

The dependence of the output energy density  $(W_{out})$  on the input one  $(W_{in})$  for PDLC structures is shown in Fig. 4.

More than 10-fold attenuation of the laser beam has been obtained for the PDLC cell based on polyimide 6B at the incident energy density of 0.15–0.2 Jcm<sup>-2</sup>. Twice attenuation of the laser energy density has been found in the fullerene-doped PDLC based on COANP and NPP compounds that is close to the OL level for 1-µm pure fullerene film. The difference in the nonlinear transmission of PDLC based on COANP structure and NPP one is associated with different matching between COANP and NLC refractive indexes as compared with that between NPP and NLC re-



Fig. 4. Dependence of the output energy density ( $W_{out}$ ) on the input one ( $W_{in}$ ) for PDLC structures: 1 – COANP-5 wt.%C<sub>70</sub>-LC (E7), 2 – NPP-1 wt.%C<sub>60</sub>-LC (E7), 3 – polyimide-(0.25 wt.% C<sub>60</sub>+C<sub>70</sub>)-LC (ZK999), 4 – polyimide-(0.5 wt.% C<sub>60</sub>+C<sub>70</sub>)-LC (ZK999),  $\lambda$  = 532 nm.

fractive indexes. Reverse saturable absorption, complex formation and laser-induced scattering could be discussed to explain OL results in the PDLC structures studied.

The first OL experiments have been carried out for PDLC based on COANP-C70 and E7 under femtosecond pulsed irradiation of a quasi-CW Ti-sapphire laser ( $\lambda$  = 805 nm). Taking into account the broad absorption band in 800-820 nm associated with charge transfer in the COANP-fullerene structure [21], more than 10-fold attenuation of the laser beam has been detected. The experiment allowed dynamics of photo-induced processes to be investigated at the upper excitation levels, which were characterized by the transition times of 1 fs, and the reverse saturable absorption to be revealed in the fullerene-doped organic structures based on COANP, that did not contradict the results of Ref. 24. In that paper it was found in pure fullerene films that a photo-induced absorption increase occurred at the forbidden transition of 1.6 eV due to the electron state hybridization in the electric field of the high laser beam.

# 4. Conclusions

To summarize, the photo-induced change  $\Delta n$  in refractive index was studied by the dynamic holographic technique in the fullerene-doped PDLC system under nano- and pico--second pulsed laser irradiation. Maximum  $\Delta n$  was established to be observed in the system with 5 wt.% of C<sub>70</sub> in COANP, the value was  $1.4 \times 10^{-3}$  at  $W_{in}$  of  $17.5 \times 10^{-3}$  Jcm<sup>-2</sup> and  $7.2 \times 10^{-4}$  at  $W_{in}$  of  $25 \times 10^{-3}$  Jcm<sup>-2</sup> in nano- and picosecond pulsewidth region, respectively. The results obtained for different composites based on COANP and NPP were compared. The nonlinear refractive index  $n_2$  and the third-order susceptibility  $\chi^{(3)}$  have been determined for fullerene-doped PDLC.

The data predict optoelectronic potentials of PDLC applications for modulation and generation of the laser irradiation over the visible spectral range. The spectral peculiarities of the system studied propose to carry out the investigations over the near-infrared spectral range.

Optical limiting results have been shown for fullerene-doped PDLCs at wavelength of 532 and 805 nm. 10-fold attenuation of the laser beam has been obtained that predicts to use fullerene-doped PDLCs to protect the human eyes and technical devices against high laser irradiation.

## Acknowledgements

The author would like to thank Dr. V.I. Berendyaev (Karpov Research Physical-Chemical Institute, Moscow, Russia), Dr. S.E. Putilin (Vavilov State Optical Institute, St.Petersburg, Russia) as well as Dr. A. Barrientos (Department of Physics, UPR-RUM, Mayagüez, PR, USA) for their help in this study. This work was partially supported by RFBR grant # 01-03-33162 and Russian Program "National Technology Base".

# References

- 1. G.M. Zharkova and A.S. Sonin, *Liquid Crystal Compositions*, Nauka, Novosibirsk, 1994. (in Russian)
- F. Simoni, G. Cipparrone, C. Umeton, G. Arabia, and G. Chidichimo, "Optical nonlinearities induced by thermal effects in polymer dispersed liquid crystals", *Appl. Phys. Lett.* 54, 896–897 (1989).
- H. Ono, I. Saito, and N. Kawatsuki, "Orientational photorefractive effects observed in poly(vinyl alcohol)/liquid crystal composites", *Appl. Phys.* B66, 527–529 (1998).
- 4. H. Ono and N. Kawatsuki, "Orientational photorefractive gratings observed in polymer dispersed liquid crystals doped with fullerene", *Jap. J. Appl. Phys.* Part 1, **36**, 6444–6448 (1997).
- H. Ono and N. Kawatsuki, "Response characteristics of high-performance photorefractive liquid crystals", *Jap. J. Appl. Phys.* Part 1, 38, 737–740, 1999.
- N.V. Kamanina, "Optical investigations of a C<sub>70</sub>-doped 2-cyclooctylamino-5-nitropyridine–liquid crystal system", *J. Opt. A: Pure Appl. Opt.* 4, 571–574 (2002).
- L.P. Rakcheeva and N.V. Kamanina, "Prospects of the use of fullerenes for the orientation of liquid-crystalline compositions", *Techn. Phys. Lett.* 28, 457–460 (2002).
- N.V. Kamanina, "Nonlinear optical study of fullerene--doped conjugated systems: new materials for nanophotonics applications", *Proceed. NATO Advanced Research Workshop on Organic Nanophotonics* II/100, 177–192 (2003).
- N.V. Kamanina and I.Yu. Denisyuk, "Study of the dynamic characteristics of polymer-dispersed liquid crystal compositions: Prospects for optoelectronic applications of fullerene complexes with phthalocyanine nanocrystals", *Tech. Phys. Lett.* 30, 36–39 (2004).
- K. Sutter, J. Hulliger, and P. Günter, "Photorefractive effects observed in the organic crystal 2-cyclooctylamino--5-nitropyridine doped with 7,7,8,8,-tetracyanoquinodimethane", *Sol. St. Commun.* 74, 867–870 (1990).
- Y. Cui, J. Wu, N. Kamanina, A. Pasaje, A. Leyderman, A. Barrientos, M. Vlasse, and B.G. Penn, "Dielectric study of dynamics of organic glasses", *J. Phys. D: Appl. Phys.* 32, 3215–3221 (1999).
- 12. N. Kamanina, A. Barrientos, A. Leyderman, Y. Cui, V. Vikhnin, and M. Vlasse, "Effect of fullerene doping on

the absorption edge shift in COANP", Mol. Mater. 13, 275–280 (2000).

- N.V. Kamanina, S. Putilin, and D. Stasel'ko, "Nano-, picoand femtosecond study of fullerene-doped polymer-dispersed liquid crystals: holographic recording and optical limiting effect", *Synth. Met.* **127**, 129–133 (2002).
- N.V. Kamanina, "Peculiarities of optical limiting effect in -conjugated organic systems based on 2-cyclooctylamino-5-nitropyridinedoped with C<sub>70</sub>", *J. Opt. A: Pure Appl. Opt.* 3, 321–325 (2001).
- N.V. Kamanina, "Mechanisms of optical limiting in π-conjugated organic system: fullerene-doped polyimide", *Synth. Met.* 127, 121–128 (2002).
- N.V. Kamanina, V.N. Sizov, and D.I. Stasel'ko, "Recording of thin phase holograms in polymer-dispersed liquid-crystal composites based on fullerene-containing π-conjugated organic systems", *Opt. Spectrosc.* **90**, 1–3 (2001).
- 17. R.J. Collier, C.B. Burckhardt, and L.H. Lin, *Optical Holog-raphy*, Academic Press, New York and London, 1971.
- I.C. Khoo, H. Li, and Y. Liang, "Observation of orientation photorefractive effects in nematic liquid crystals", *Opt. Lett.* 19, 1723–1725 (1994).
- I.V. Rubtsov, D.V. Khudiakov, V.A. Nadtochenko, A.S. Lobach, and A.P. Moravskii, "Rotational reorientation dynamics of C<sub>60</sub> in various solvents. Picosecond transient grating experiments", *Chem. Phys. Lett.* **229**, 517–523 (1994).
- I.V. Rubtsov, D.V. Khudiakov, A.P. Motavskii, and V.A. Nadtochenko, "Orientational behavior of C<sub>70</sub> molecules in chlorobenzene", *Chem. Phys. Lett.* 249, 101–104 (1996).
- N.V.Kamanina and E.F. Sheka, "Optical limiters and diffraction elements based on a COANP-fullerene system: Nonlinear optical properties and quantum-chemical simulation", *Opt. Spectrosc.* 96, 599–612 (2004).
- 22. N.V. Kamanina and A.I. Plekhanov, "Mechanisms of optical limiting in fullerene-doped  $\pi$ -conjugated organic structures demonstrated with polyimide and COANP molecules", *Opt. Spectrosc.* **93**, 408–415 (2002).
- 23. S.A. Akhmanov and S.Yu. Nikitin, *Physical Optics*, Oxford Press, Oxford, 1997.
- A.V. Bazhenov, A.V. Gorbunov, M.Yu. Maksimuk, and T.N. Fursova, "Photoinduced light absorption by C<sub>60</sub> films in the 0.08–4.0-eV spectral range", *J. Exp. Theor. Phys.* 85, 135–140 (1997).

# EUROCONFERENCE Photovoltaic Devices



# **EUROPV 2004**



Kranjska Gora, Slovenia, October 15-20, 2004

### SCOPE OF THE CONFERENCE

The aim of the conference is to apply the results from the first conference Photovoltaics and Environment (Granada, 2003) and focus on the large-scale production and application of PV in the future. Ecological and economical aspects of mass production of the different cell types in the GW-scale will be examined and speakers with a broad experience in today's PV-production will give their views on existing production methods, possible future improvements and up-scaling. The mass application of PV as a major source of electricity will be scrutinised with respect to the impact on the grids, usage of existing buildings and open space. The conference will provide an intense forum to researchers from industry and academic institutions to exchange their views on new developments and strategies as well as to form new collaborations. The set up of the programme is multidisciplinary in order to promote the cross-fertilisation of different disciplines, technologies and developments. Leading scientists and representatives from companies will refer about the latest research results and developments. In addition there will be poster sessions, informal working group sessions and round table discussions. To assure an efficient exchange and transfer of the knowledge of the photovoltaics and environment the number of the participants will be limited to about 100. Grants will be available for younger scientists and INCO scientists waiving the registration fee.

LANGUAGE

English is the official language. CONFERENCE SITE

Hotel Larix, (http://www.htp-gorenjka.si/)

Borovška cesta 99, Si-4280 Kranjska Gora, Slovenia

**REGISTRATION FEE** 

Registration fees include Accommodation & Full Board, Excursion and bus transfer from/to airport Ljubljana

### INVITED SPEAKERS

Dr. Jesus Alonso (Isofoton, Spain) Dr. Derk Blätzner (ETH Zürich, Switzerland)

Artur Buechel (Unaxis, Lichtenstein)

Dr. Net Daukes (Imperial College London, UK)

Dr. Alessandro Fraleoni-Morgera (University of Bologna, Italy)

Dr. Jonathan Halls (CDT, Cambridge (UK)

Dr. Arnulf Jäger Waldau (European Commission JRC Ispra, Italy)

Dr. Gert Jan Jorgenden (Akzo Nobel, The Netherlands)

Prof. Daniel Lincot (Laboratoire d'Electrochimie ENSCP, France)

Dr. Tobias Meyer (Solaronix SA, Switzerland)

Pietro Menna (European Comission DG TREN, Belgium)

Dr. Ersin Ozsan (Teksolar, UK)

Dr. Jef Poortmans (IMEC, Belgium)

Dr. Volker Probst (Shell Solar, Germany)

Dr. Bernd Rech (Forschungszentrum Jülich, Germany)

Prof. Wim Sinke (ECN, The Netherlands)

Dr. Jozef Szlufcik (Photovoltech, Belgium)

Dr. Karsten Wambach (Deutsche Solar, Germany) Prof. Peter Wuerfel (Univ. of Karlsruhe, Germany)

Prof. Peter wuerter (Univ. of Karlsrune, Germ

## CALL FOR PAPERS – ABSTRACTS

Contributed abstracts as poster presentations are encouraged. Abstracts must be no more than one page in length. The abstract should be typed in English, single spaced using Times New Roman style 12pt. A poster presentation is obligatory for grant applicants. The abstract deadline for grant applicants is June 15, 2004 and for regular participants July 15, 2004. The abstracts should be uploaded via conference web page or sent as an e-mail attachment to europv2004@paris.fe.uni-lj.si in MS Word format. Please note that there will be no proceedings from this research conference – only a book of abstracts. New ideas and results discussed at this conference should not be cited in publications without the explicit permission of their author.

#### DEADLINES

Deadline for grant applications: June 15, 2004 Grant notification: June 30, 2004 Deadline for pre-registration: July 15, 2004

ON-LINE REGISTRATION http://www.pv-net.net/europv2004.html