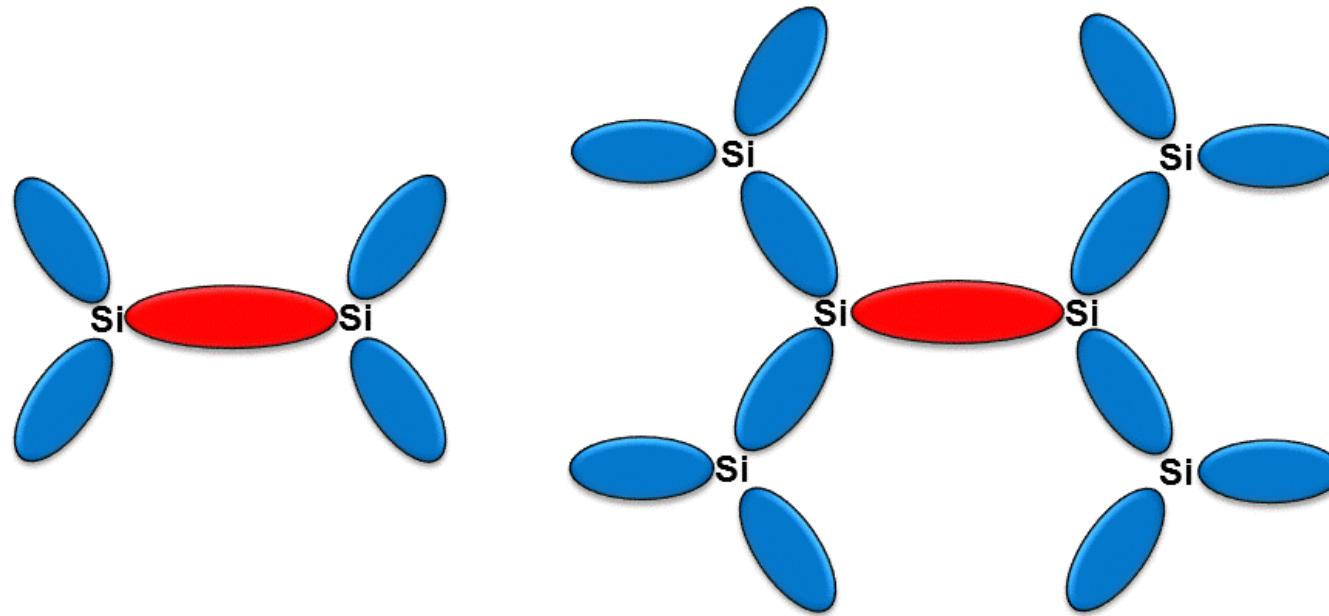


Nanostructured Organosilicon Luminophores as effective wavelength shifters for Cherenkov light and elementary particles detectors



Oleg V. Borschev^{a), b)}, Nikolay M. Surin^{a), b)}, Maxim S. Skorotetcky^{a), b)},
Bayarto K. Lubsandorzhev^{c)}, Sergey A. Ponomarenko^{a), d)}

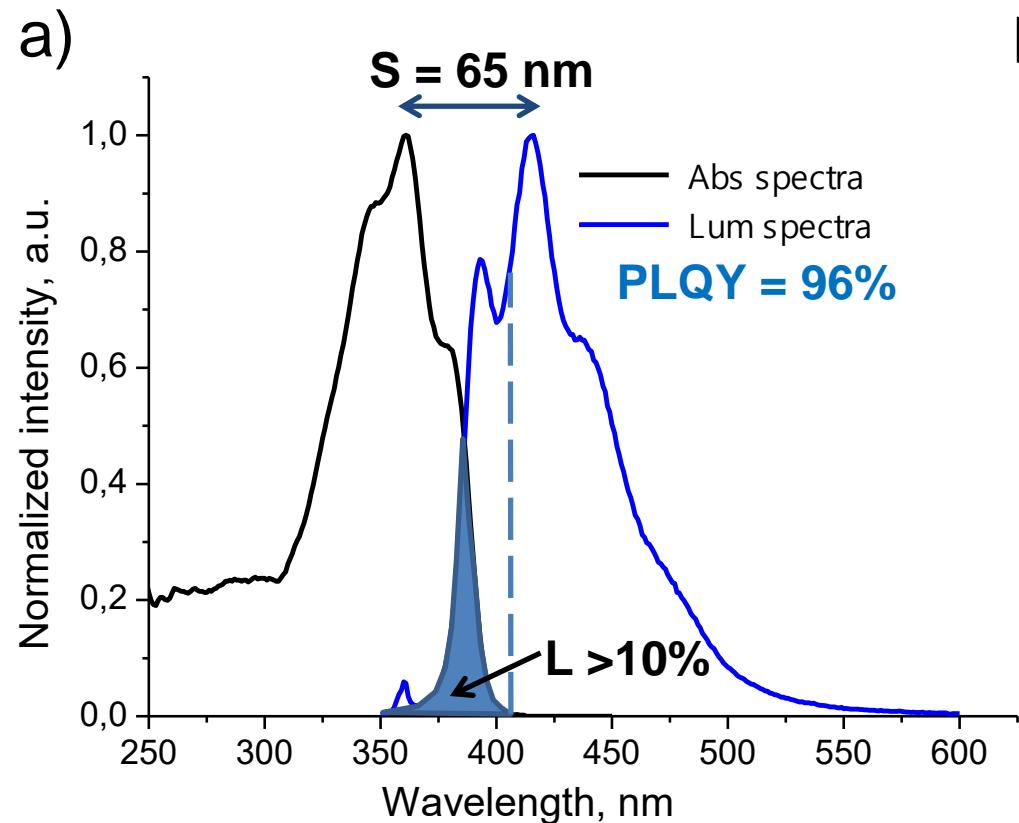
- a) Enikolopov Institute of Synthetic Polymer Materials of Russian Academy of Sciences,
Moscow, Russia; b) Luminescent Innovation Technologies LLC, Moscow, Russia;
c) Institute for Nuclear Research of Russian Academy of Sciences, Moscow, Russia;
d) Moscow State University, Chemistry Department, Moscow Russia*

Outline

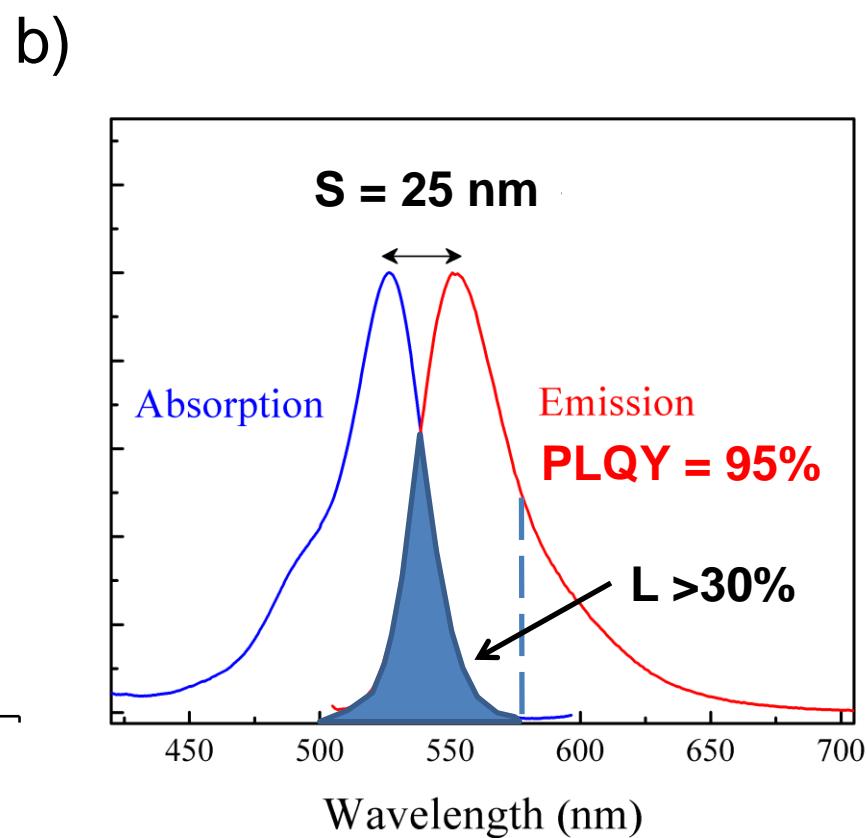
1. What are organosilicon nanostructured luminophores (NOLs)?
2. Tuning the optical properties of NOLs
3. Applications of NOLs:
 - a) plastic scintillators
 - b) scintillating fibers
 - c) VUV spectral shifters for noble gas detectors
 - d) UV spectral shifters for pure CsI detectors
 - e) Cherenkov light detectors
5. Conclusions

Optical properties of the best organic luminophores

POPOP



Rhodamine 6G

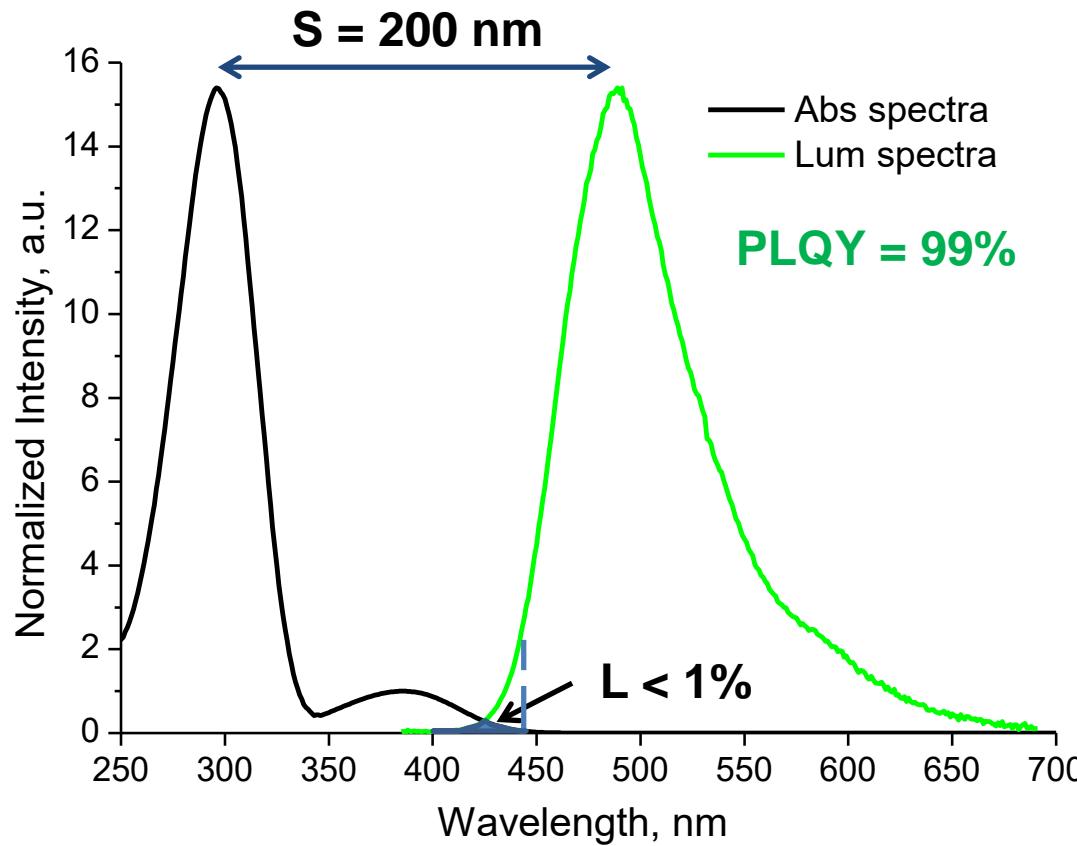


Main Drawbacks of Organic luminophores:

- 1) Small Stocks shift (S), 2) Large losses (L) of the luminescence due to self-absorption,
- 3) Luminescence quenching due to aggregation.

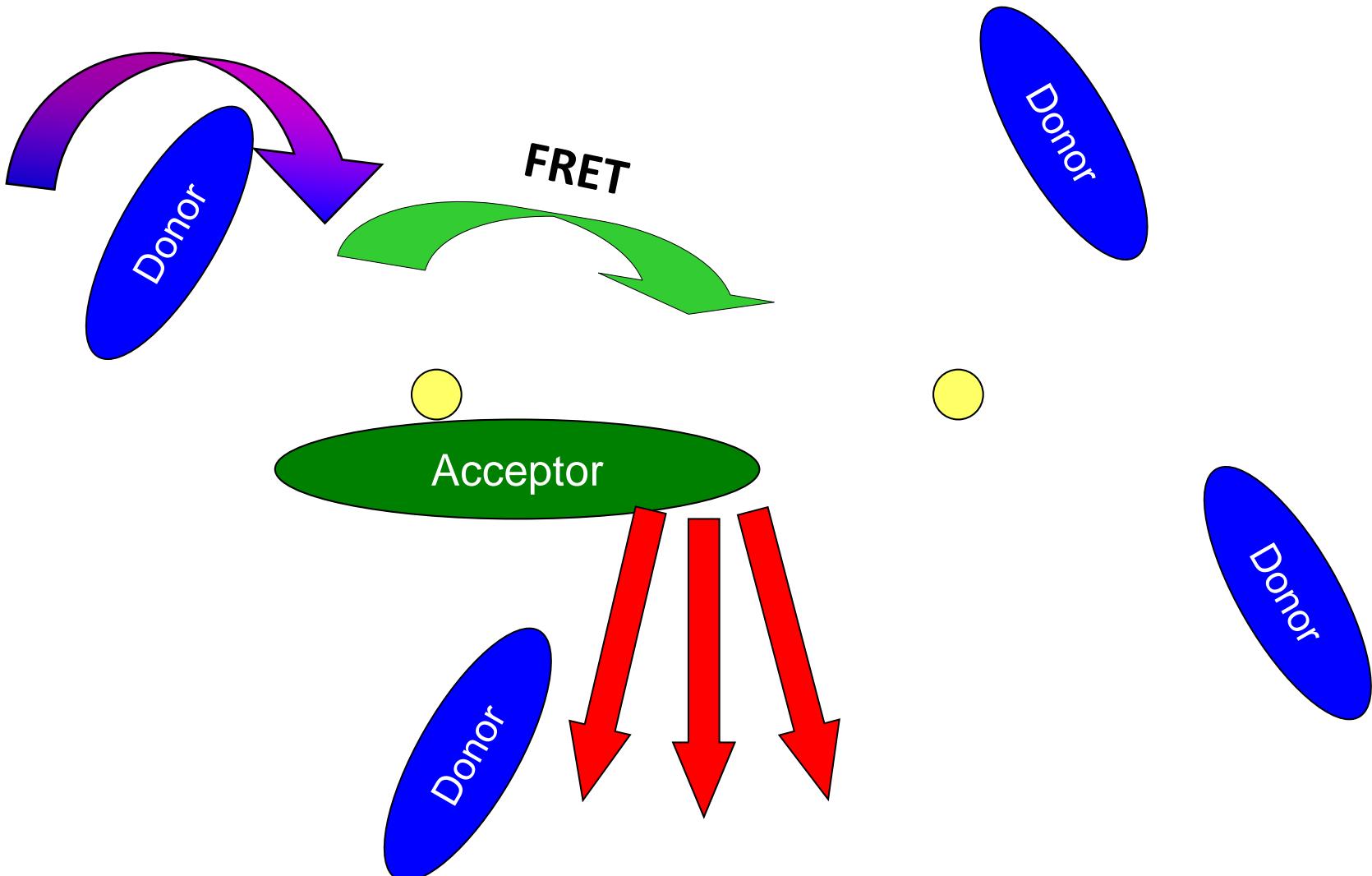
Can we improve the situation?

Can we improve the situation?
YES!

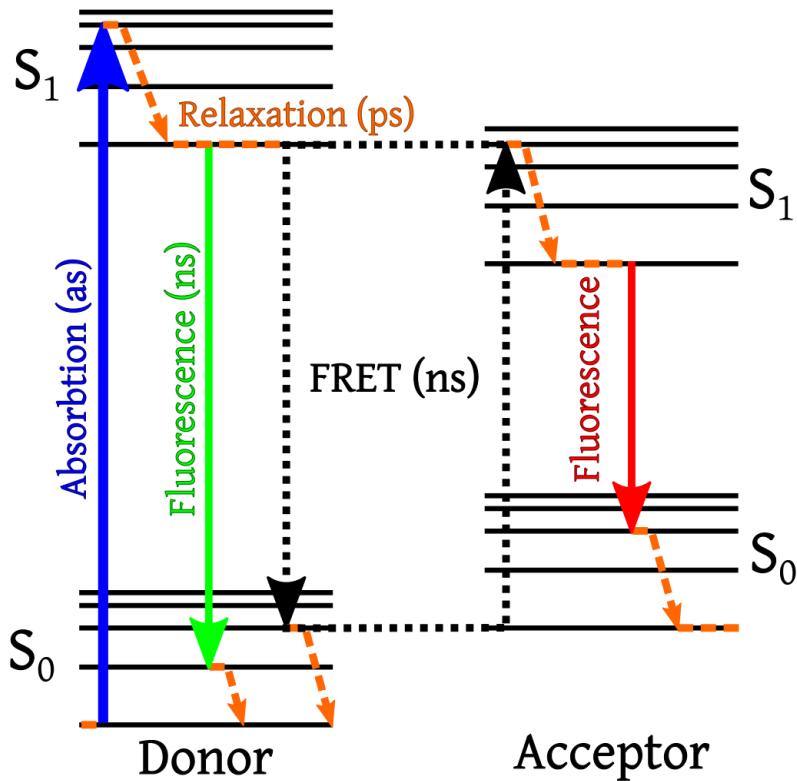


Spectral luminescence properties of
nanostructured organosilicon luminophore (NOL)

NOL works as “Molecular antennae” based on efficient intramolecular energy transfer



Förster resonance energy transfer (FRET)



FRET conditions:

1) small **distance** between the luminophores

$$\left(\frac{\vec{R}_{DA}^0}{\vec{R}_{DA}} \right)^6$$

2) **spectra overlap**;

$$\left(\int_0^{\infty} f_D(v) \cdot \varepsilon_A(v) \frac{dv}{v^4} \right)^{1/6}$$

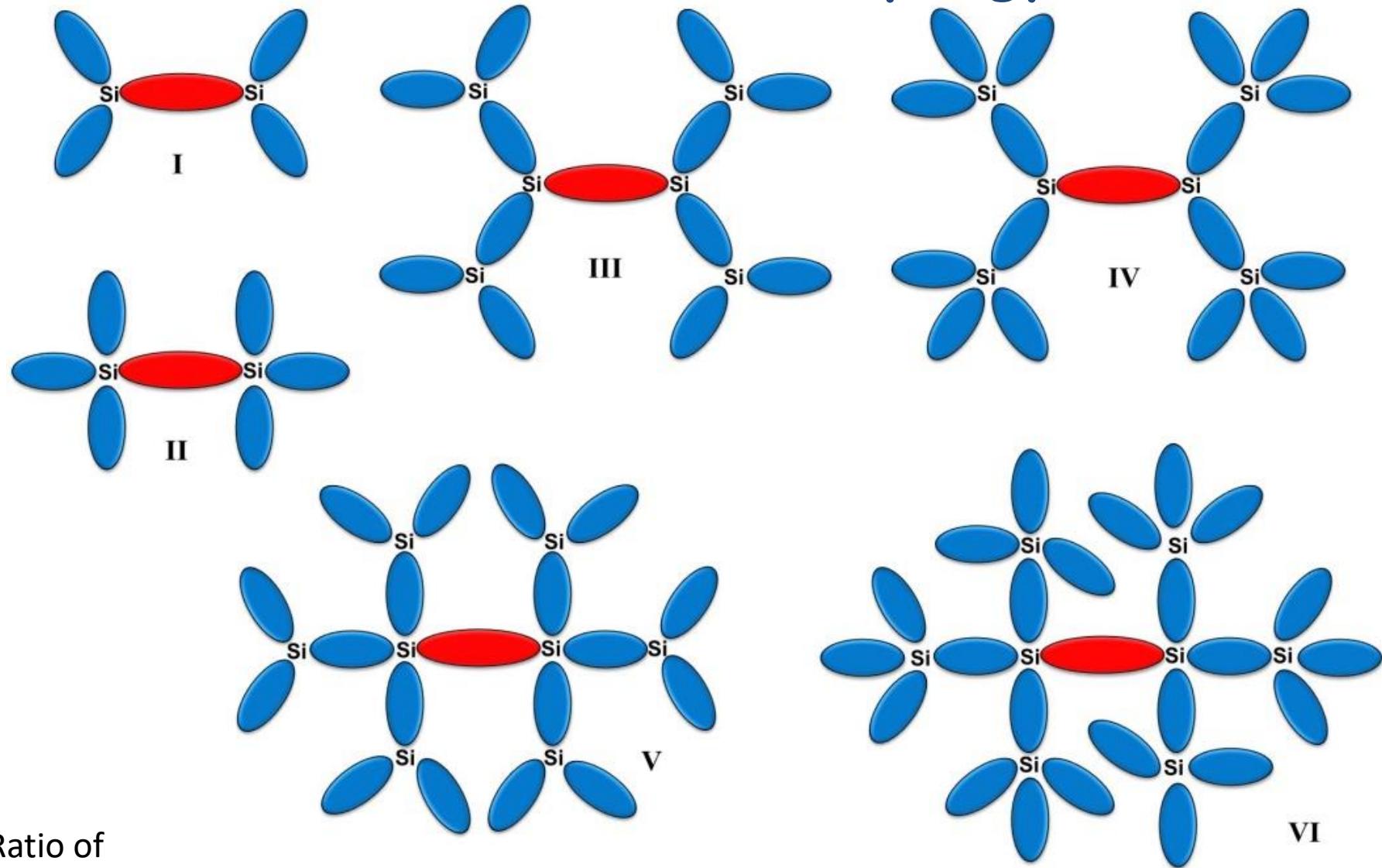
3) **orientation** of the luminophores.

\vec{R}_{DA} – distance between the centers of the donor and acceptor fragments

\vec{R}_{DA}^0 – critical radius of the inductive-resonance energy transfer

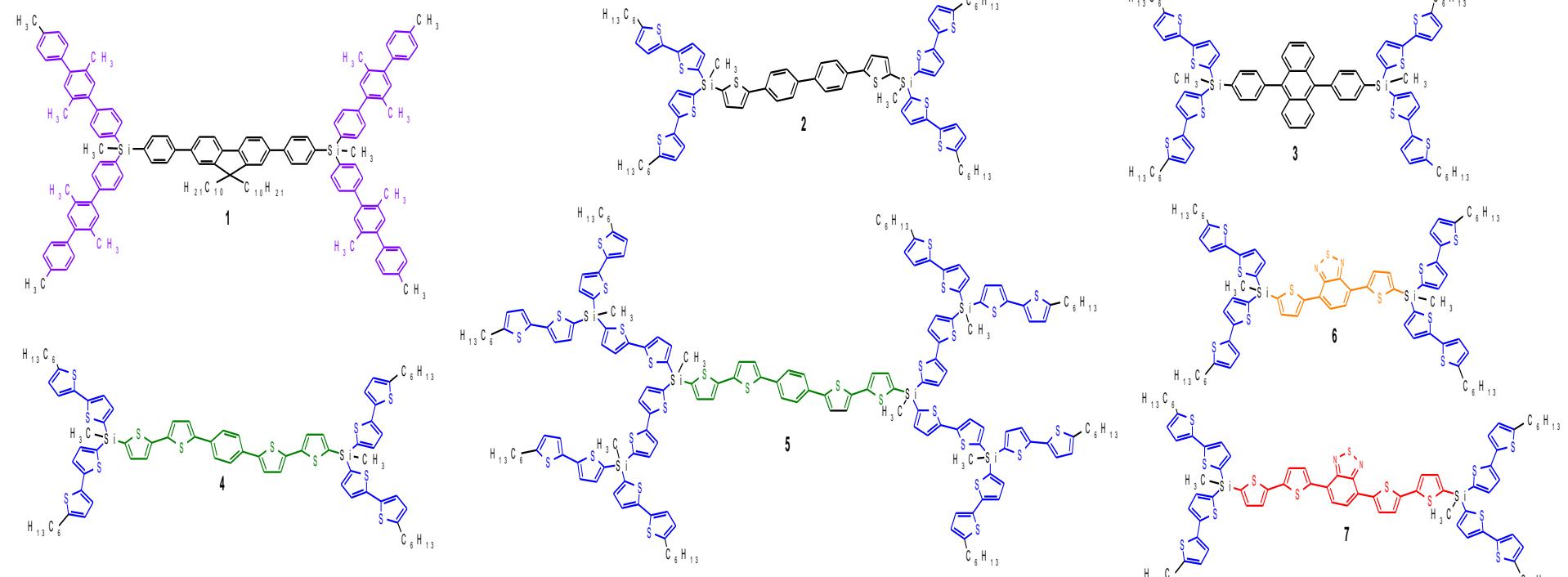
$\varepsilon_A(v)$ – acceptor absorption spectra, $f_D(v)$ – donor luminescence spectra

Schematic representation of NOLs with different topology

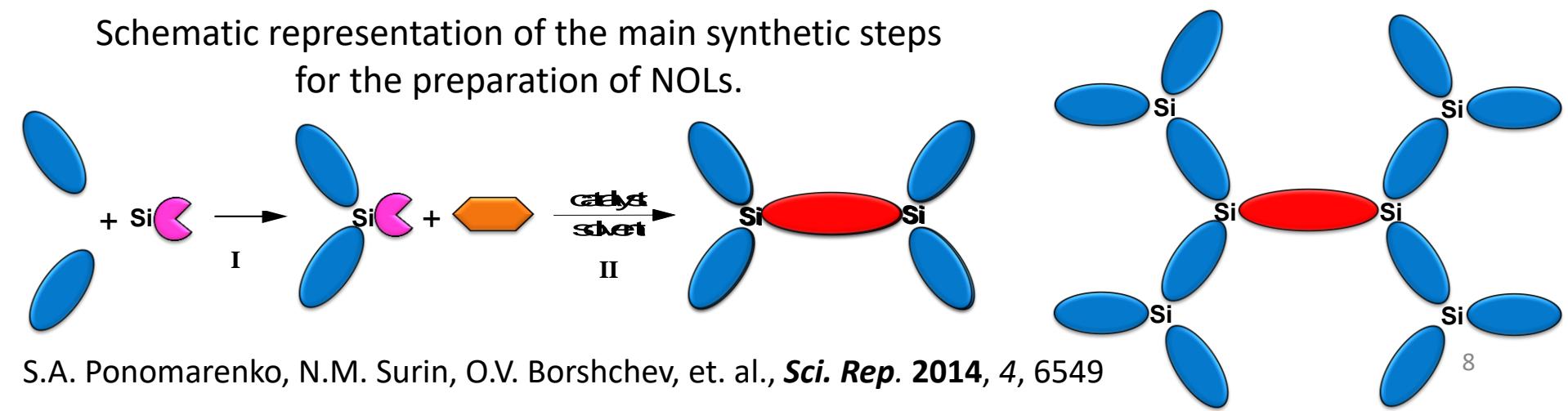


Ratio of
Donor (blue) to Acceptor (red) luminophores: 4 (**I**), 6 (**II**), 12 (**III**), 16 (**IV**), 18 (**V**) or 24 (**VI**)

Chemical structures of the first NOLs

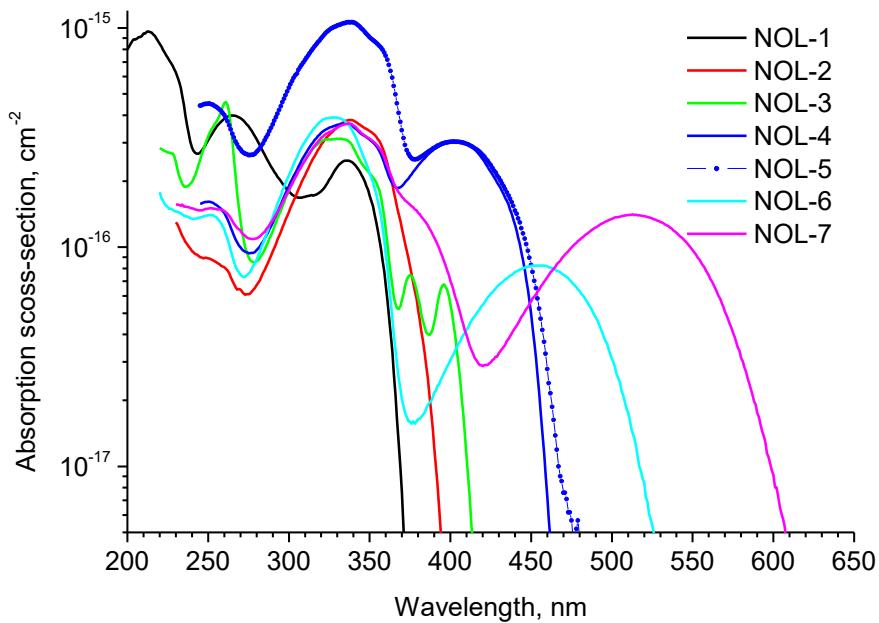


Schematic representation of the main synthetic steps
for the preparation of NOLs.

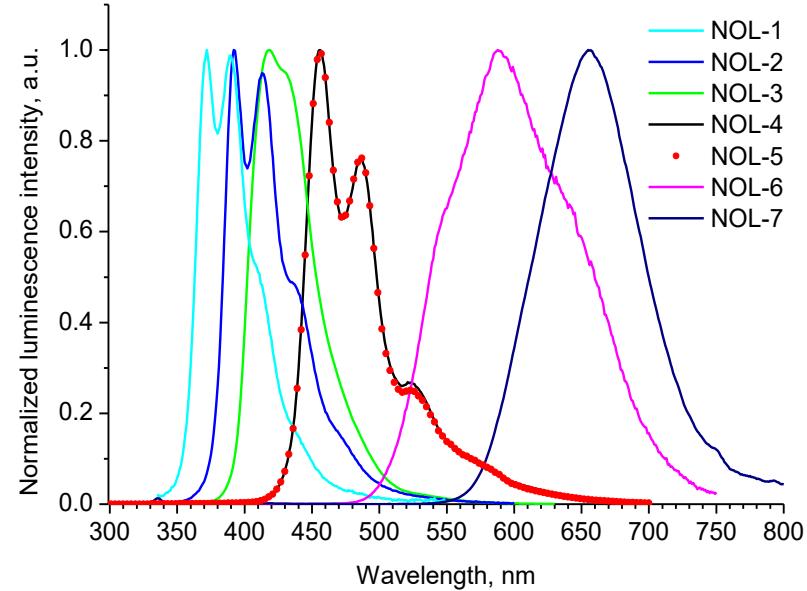


Optical spectra and main advantages of NOLs

Spectral distributions
of absorption cross-sections



Luminescence spectra

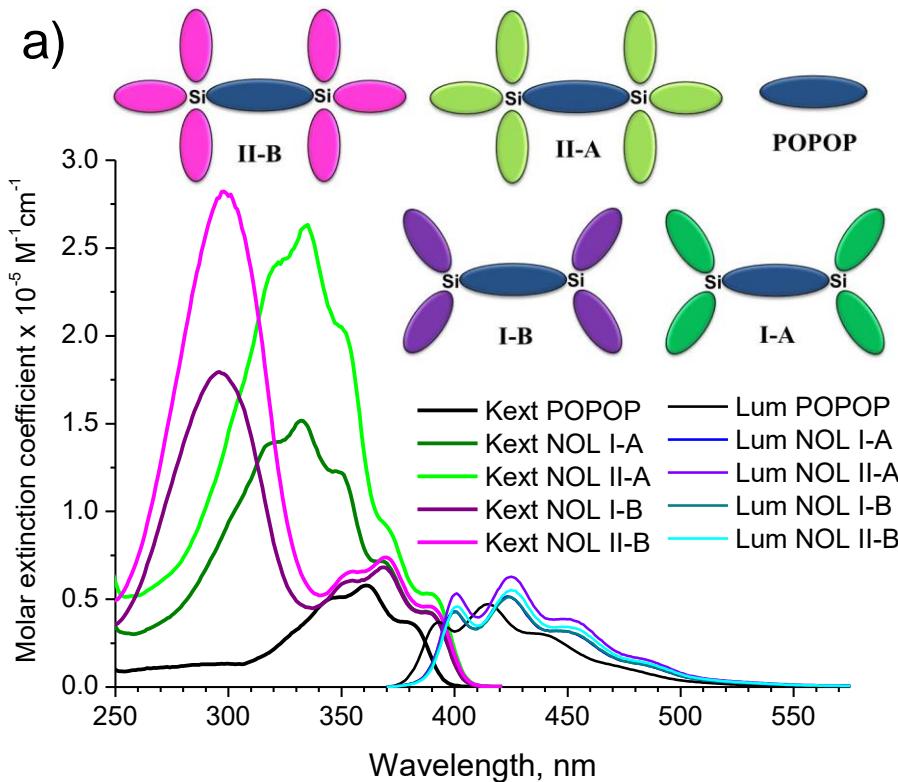


Main advantages of NOLs:

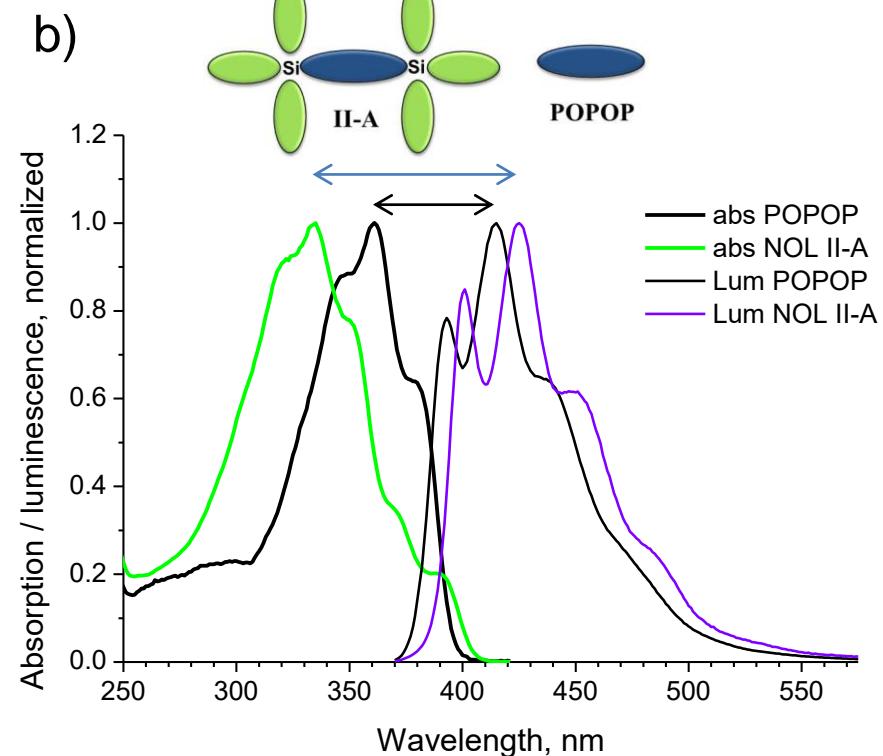
- ✓ absorption in a wide optical spectral region that can be tuned;
- ✓ 5-10 times higher absorption cross-sections as those of the best organic luminophores;
- ✓ high photoluminescence quantum yield (up to 99%);
- ✓ large pseudo Stokes shift (up to 250 – 330 nm);
- ✓ luminescence spectra in the required wavelength region;
- ✓ short luminescence lifetime;
- ✓ good solution processability.

Tuning the absorption spectra of NOLs

Molar extinction coefficients



Normalized absorption



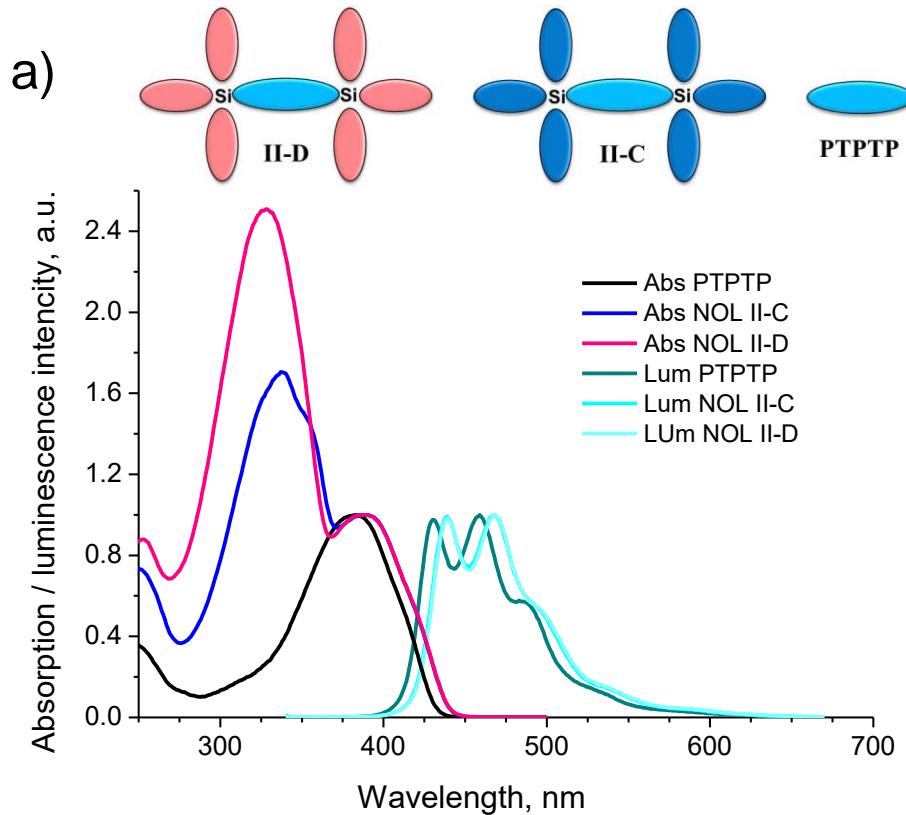
The higher is a donor-acceptor ratio in the NOL, the larger is its molar absorption coefficient.

NOLs absorb the UV light more efficiently and can be used in smaller concentrations.

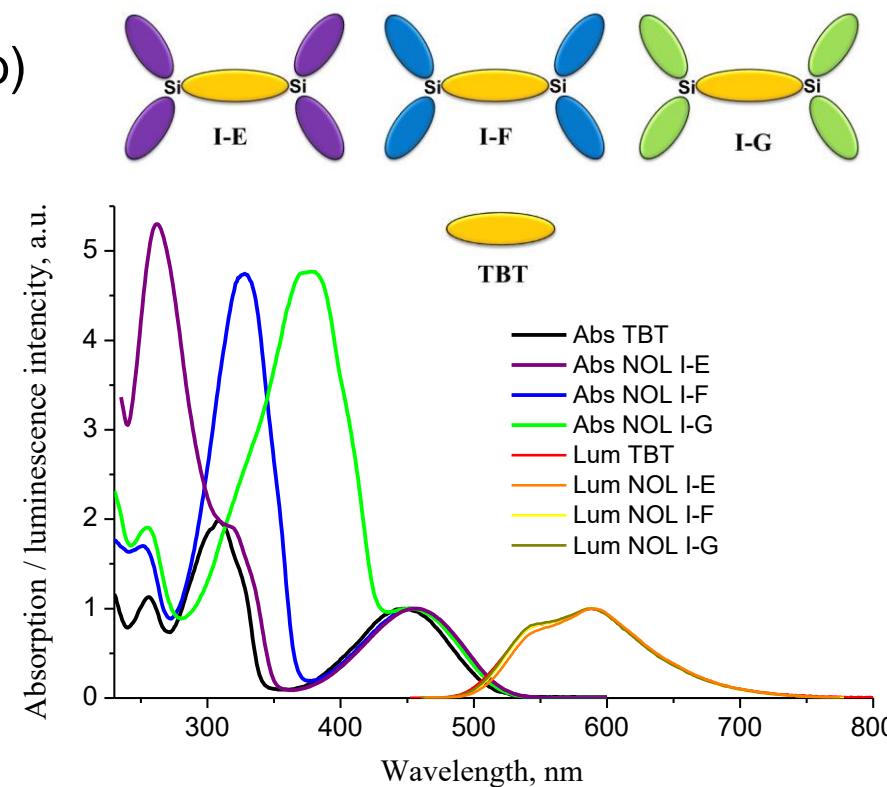
NOLs have increased Stokes shifts from $S = 65 \text{ nm}$ for POPOP to 115 nm for NOL I-A.

Tuning the absorption spectra of NOLs

Blue-emitting NOLs



Yellow-emitting NOLs



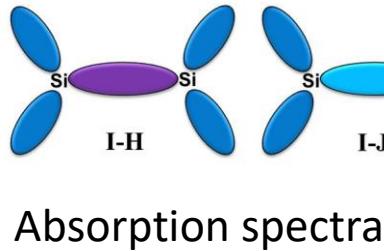
NOLs absorb the UV light more efficiently than the acceptor luminophores themselves.

Blue-emitting NOLs have the pseudo Stokes shifts up to 110 nm.

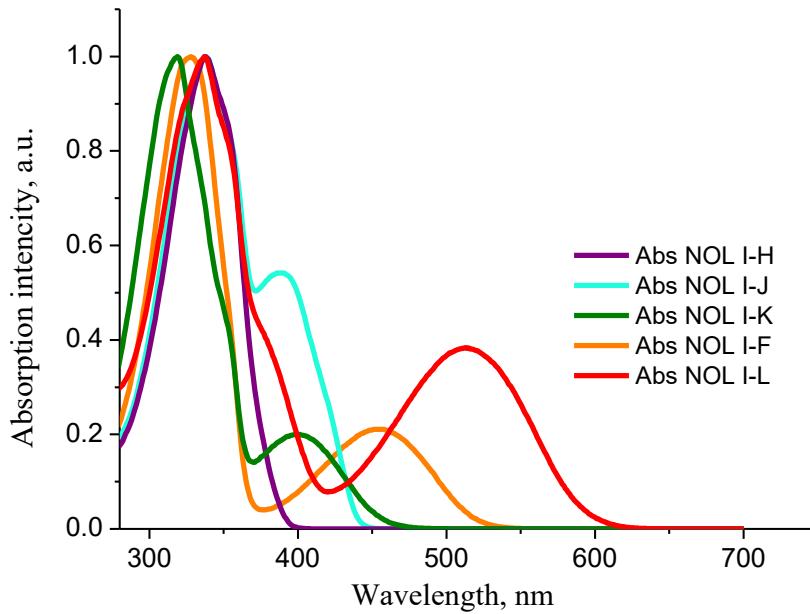
Yellow-emitting NOLs have the pseudo Stokes shifts up to 330 nm!

Tuning the luminescence spectra of NOLs

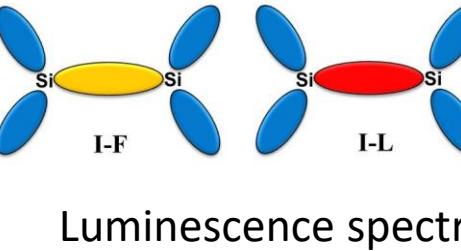
a)



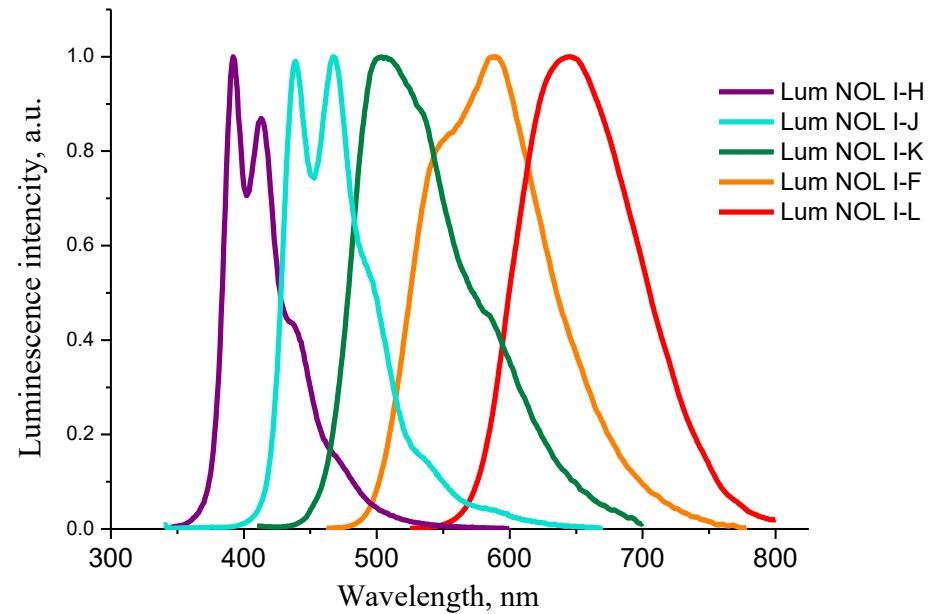
Absorption spectra



b)



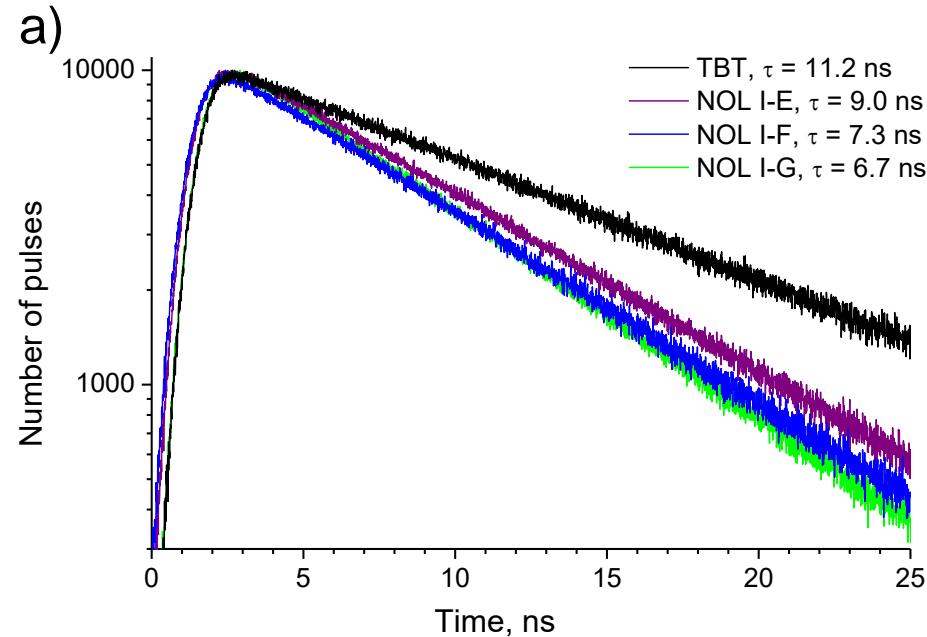
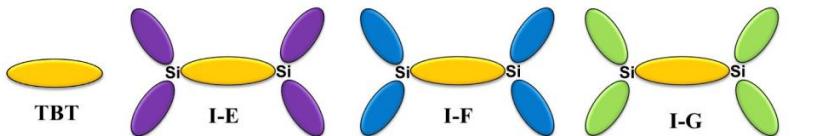
Luminescence spectra



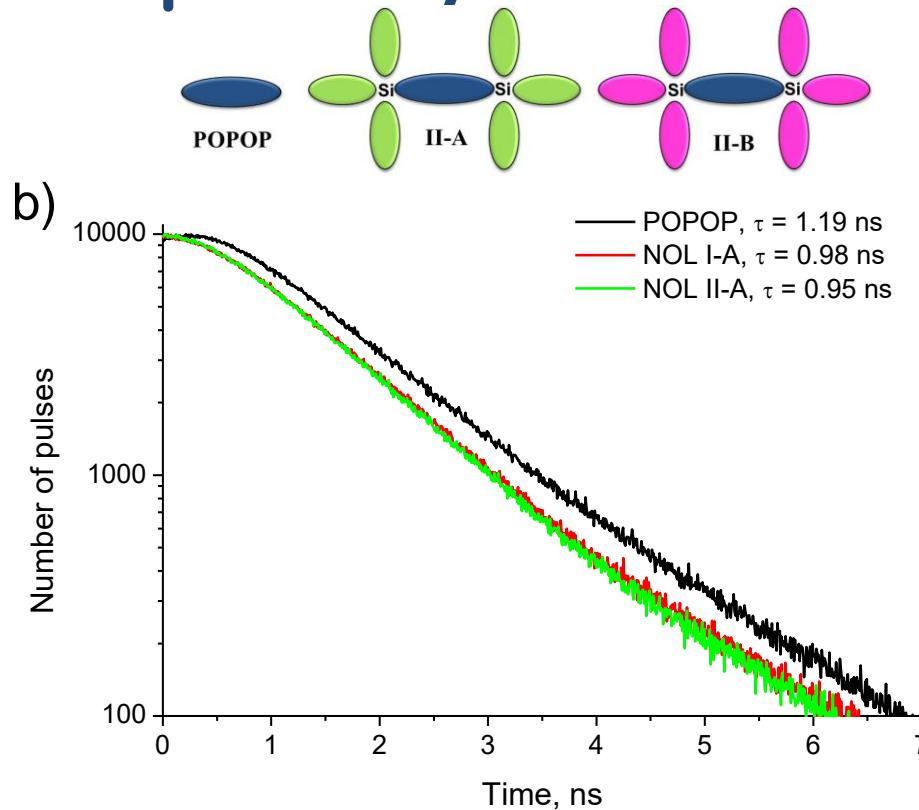
The main absorption maxima of NOLs are determined by the donor and lies in the UV range.

The luminescence spectra of NOLs are determined by the acceptor luminophore emissions, and can cover the whole visible spectral range – from violet to red.

Improving the luminescence decay time of the acceptor luminophore by NOLs



Yellow NOLs have 1.25 – 1.67 times faster luminescence than the TBT itself. Moreover, NOLs I-E and I-F have PLQY of 85 and 92%, respectively, which exceed 74% measured for TBT under the same conditions.



Blue NOLs have 1.21 – 1.25 times faster luminescence than the POPOP itself. Nevertheless, PLQY of these NOLs is essentially the same as POPOP, 96-98%.

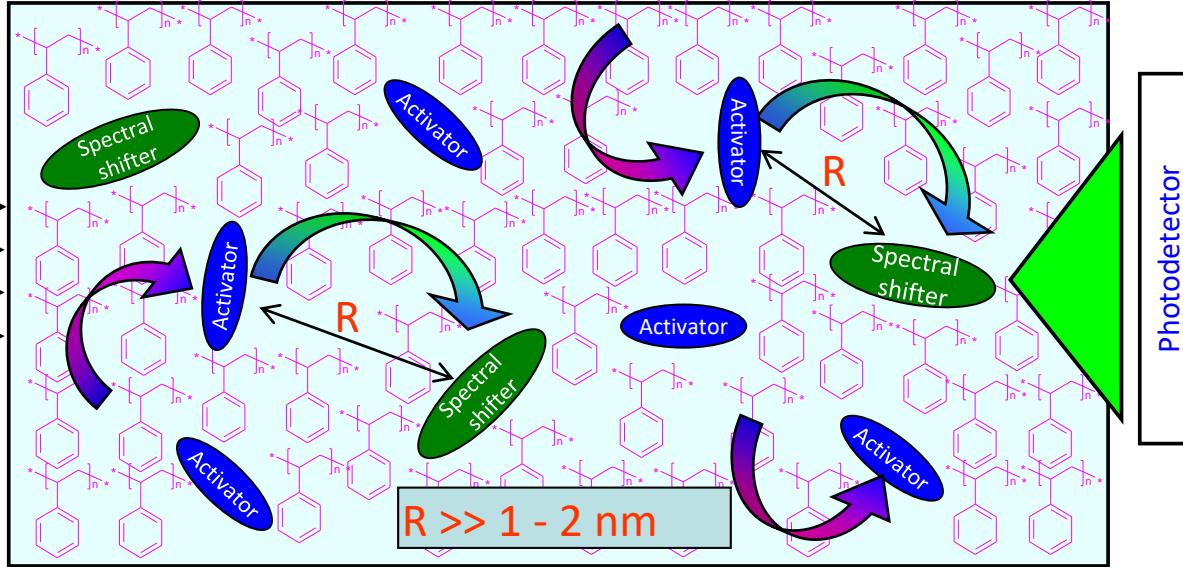
Application of NOLs in plastic scintillators

a)

classical plastic scintillator

E

(α , β or γ -radiation)



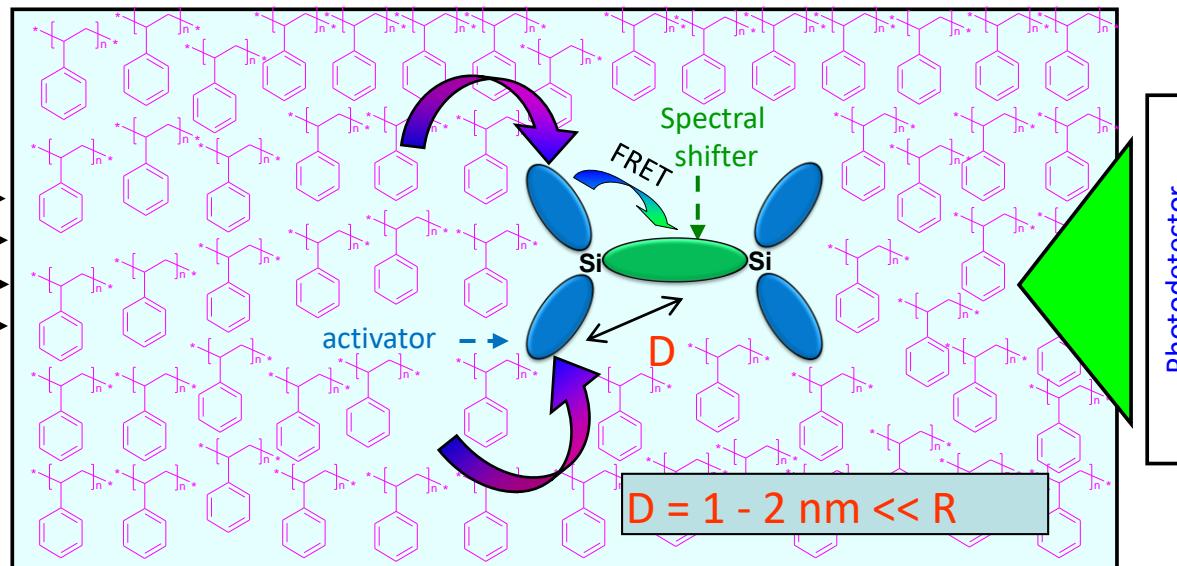
radiative energy transfer between the activator and spectra shifter

b)

plastic scintillator based on NOLs

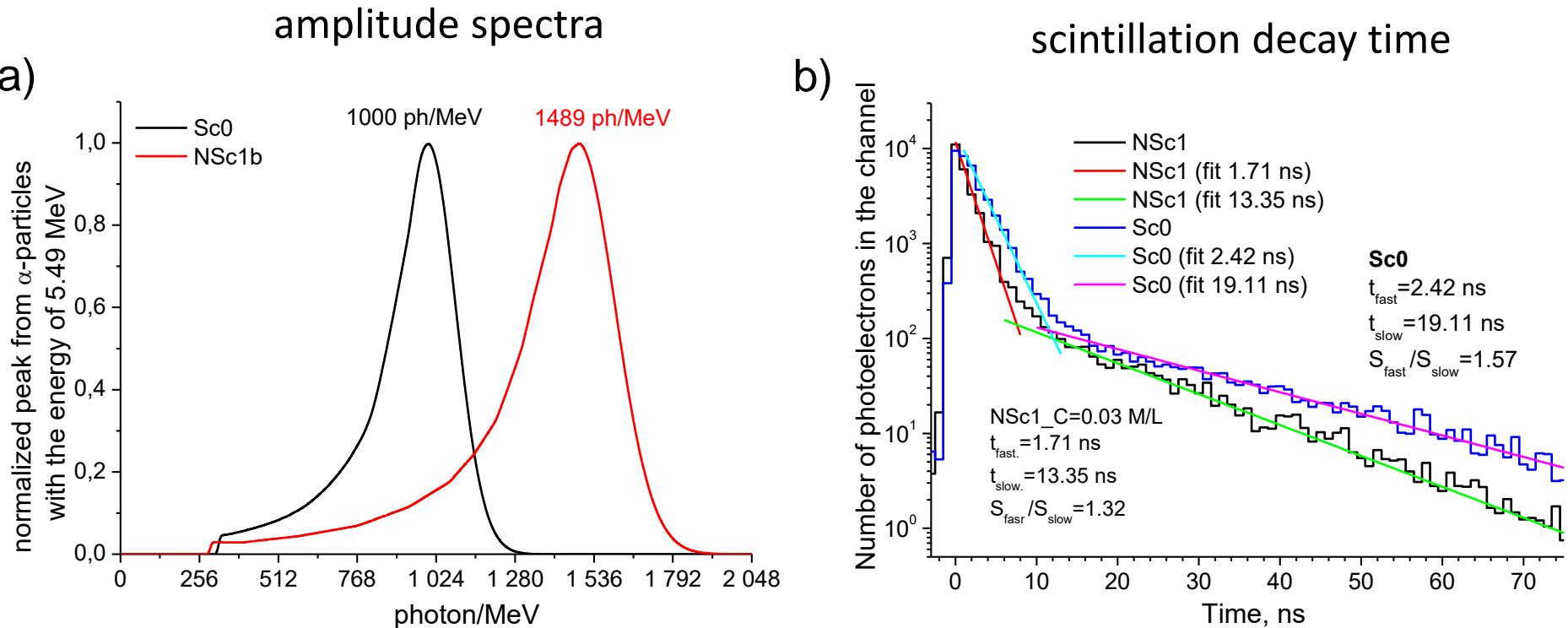
E

(α , β or γ -radiation)



highly efficienct and fast FRET between the activator and spectral shifter!

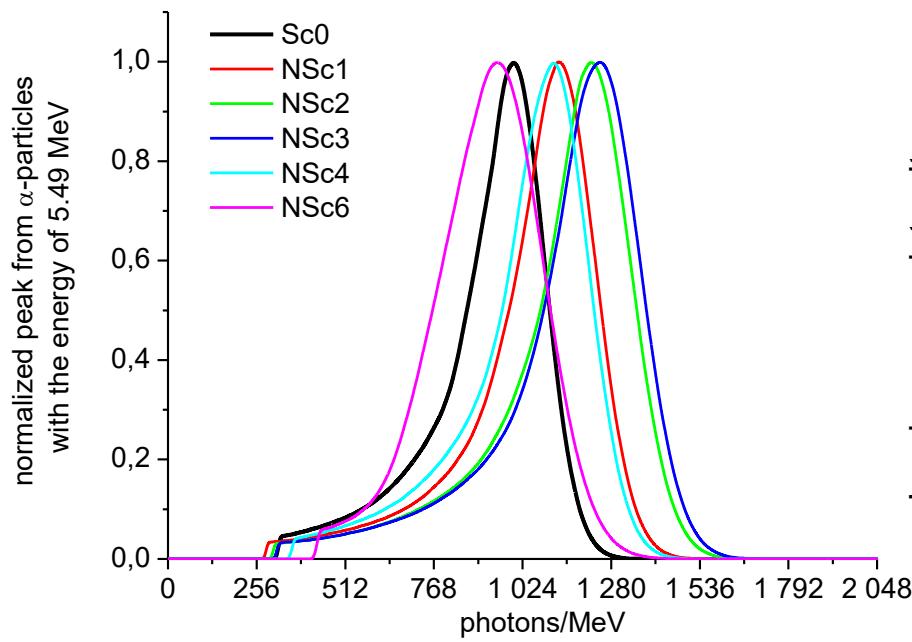
Comparison of the standard (Sc0) and NOL-based (NSc1) plastic scintillators



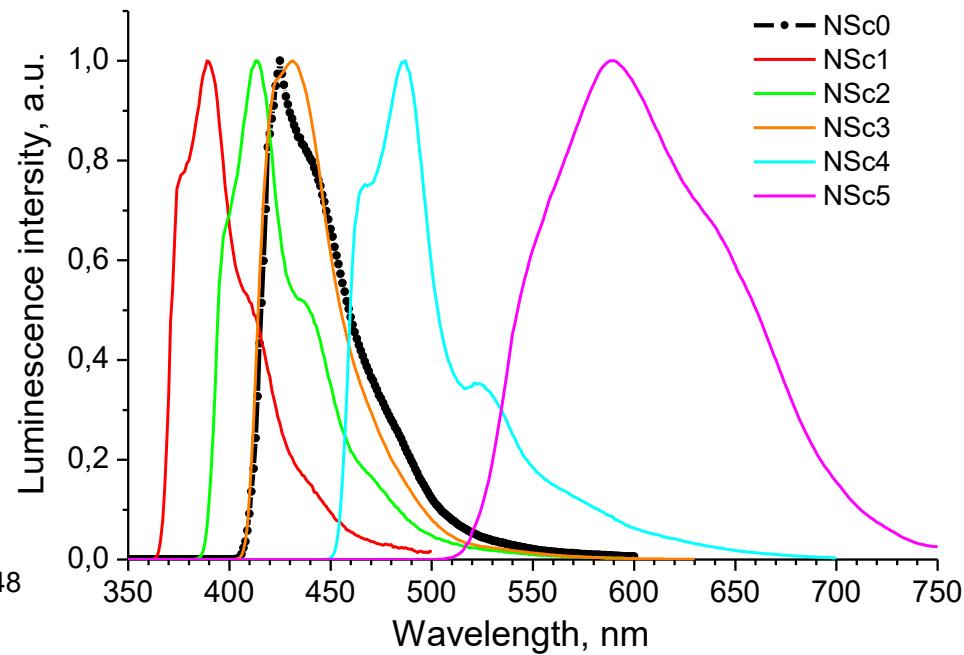
Using NOLs in plastic scintillators allows raising their scintillation light output on 50% as compared to the standard one, while their scintillation decay time can be improved on 40%.

Plastic scintillators with NOLs having different emission range

Amplitude spectra



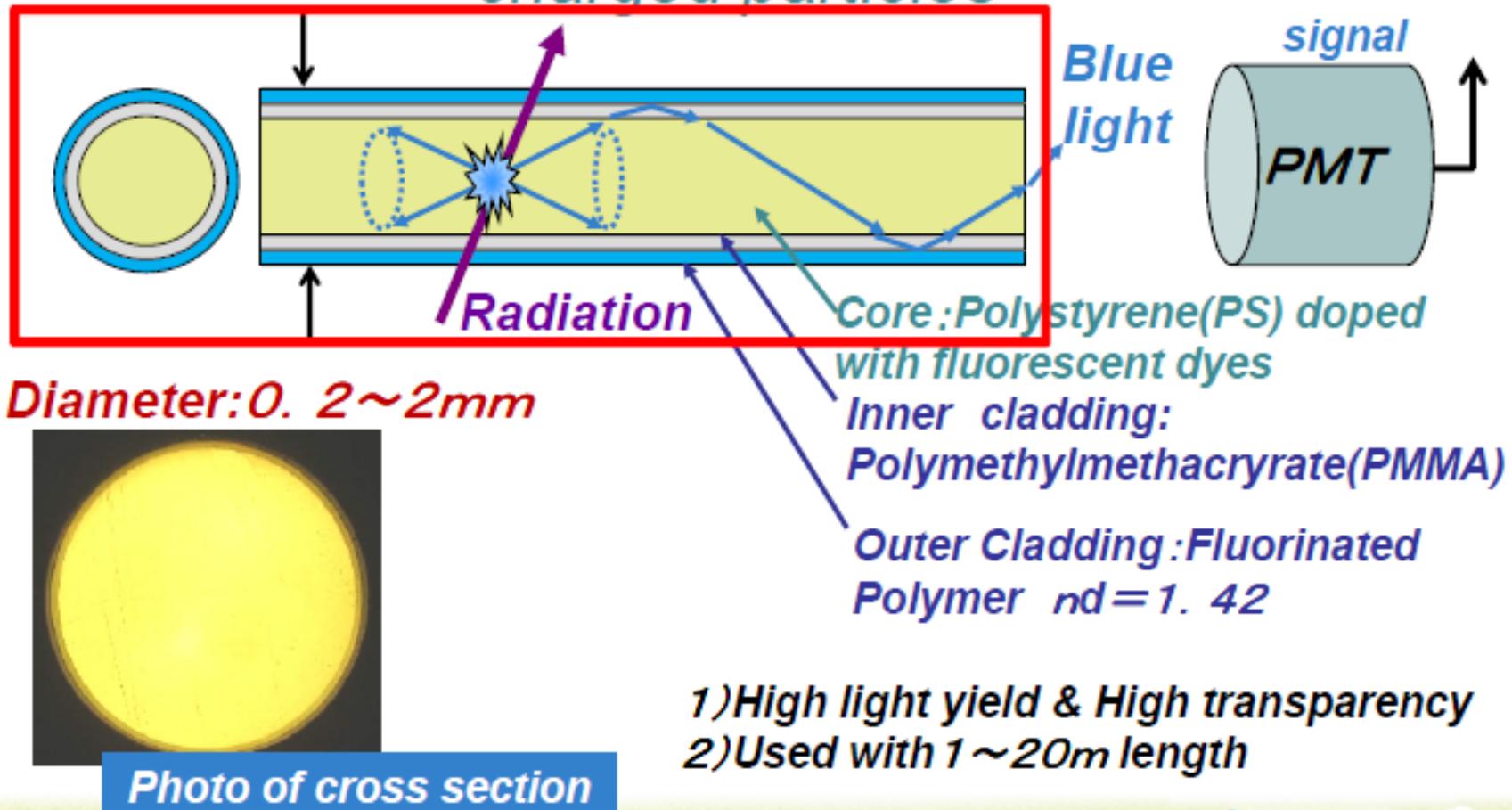
Scintillation spectra



Plastic scintillators with NOLs are able emit light in any possible visible region, which allows to adopt plastic scintillators for any desired photodetector to reach the maximal efficiency of the elementary particles detections.

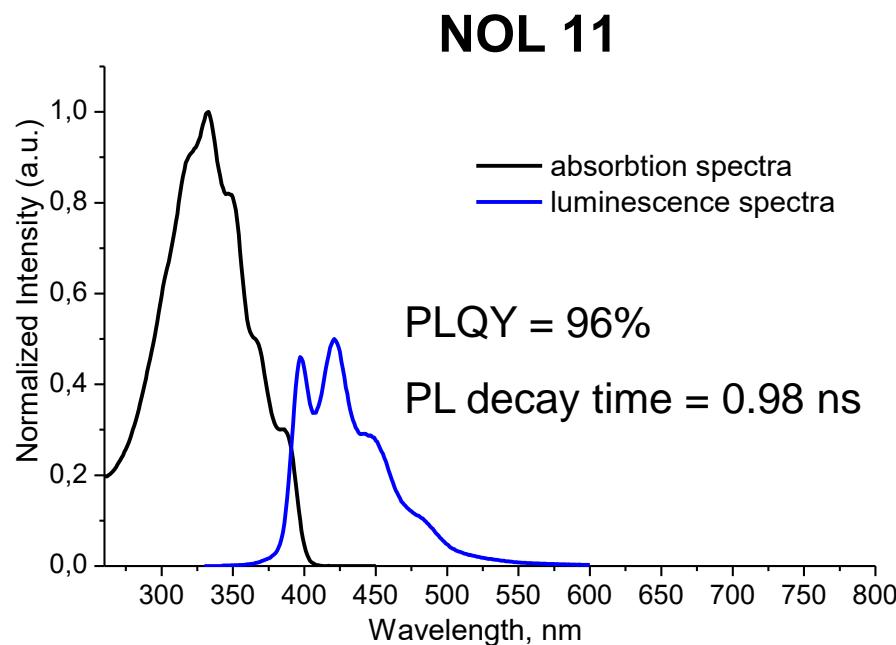
Plastic scintillating fibers

A plastic sensor fiber for detecting radiations and charged particles

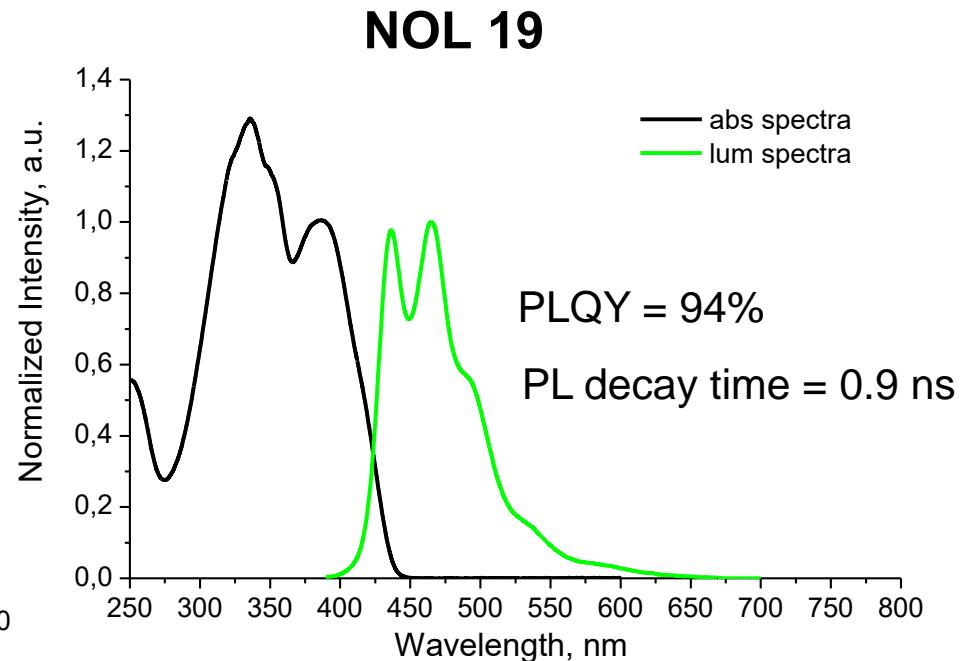


Absorption and emission spectra of NOLs used for initial preform synthesis

Blue-emitting NOL

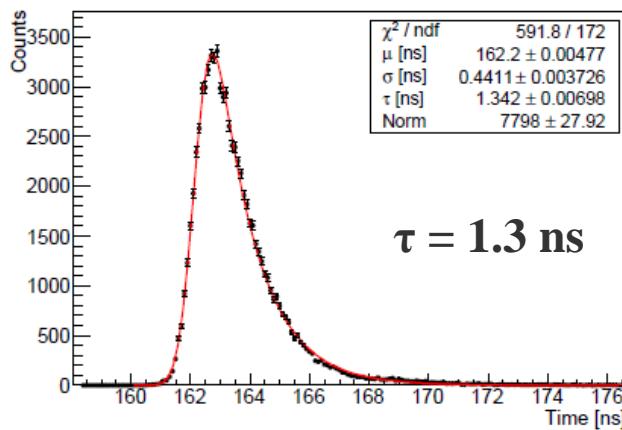


Green-emitting NOL

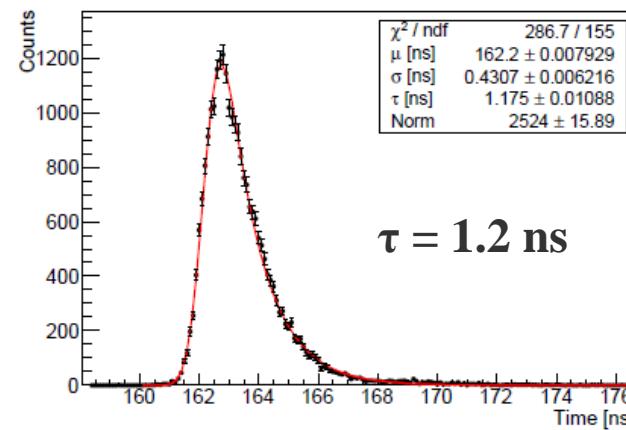


NOLs have good solubility in standard organic solvents, monomers used for plastic scintillation synthesis and in the polymers utilized as matrixes in plastic scintillators as well as in scintillating and wavelength shifting fibers.

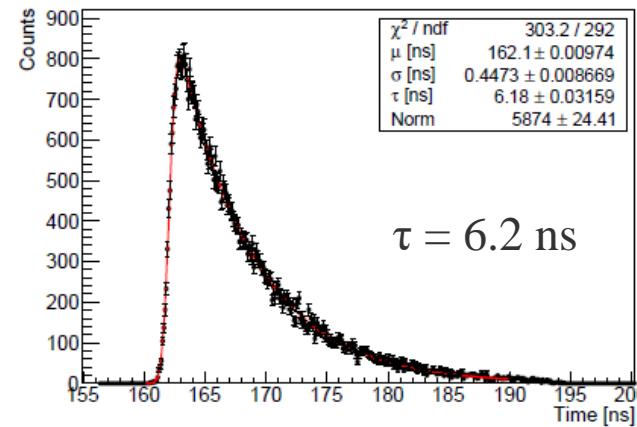
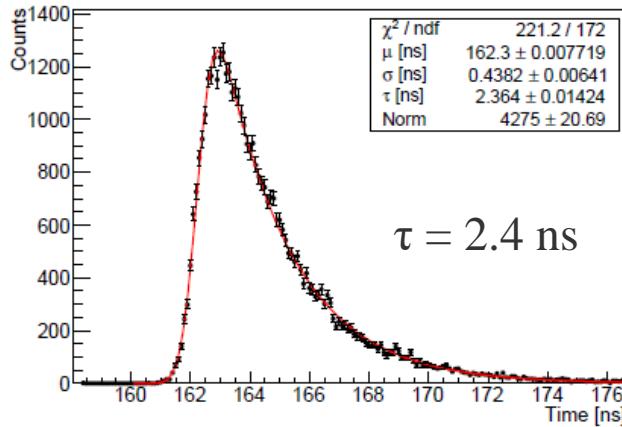
Ultrafast plastic scintillators fibers



Blue-emitting fibers

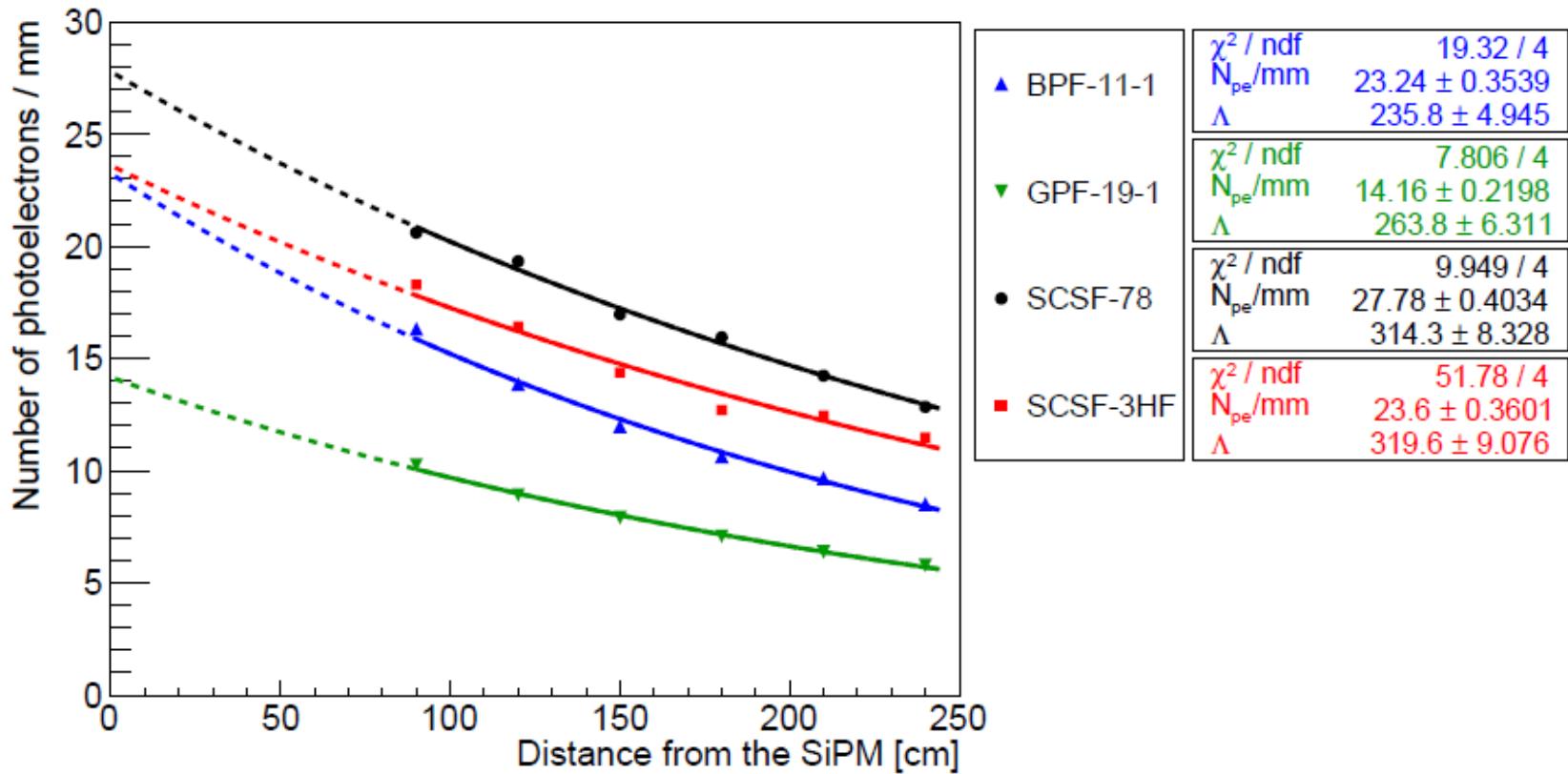


Green-emitting fibers



Decay time of NOL-based prototype fibers BPF-11-1 (a) and GPF-19-1 (b) as well as reference fibers SCSF-78 (c) and SCSF-3HF (d).

Ultrafast plastic scintillators fibers



Light yield of NOL-based prototype fiber samples and comparison to Kuraray standard fibers SCSF-78 and SCSF-3HF.

NOL-based VUV wavelength sifters for noble gas detectors

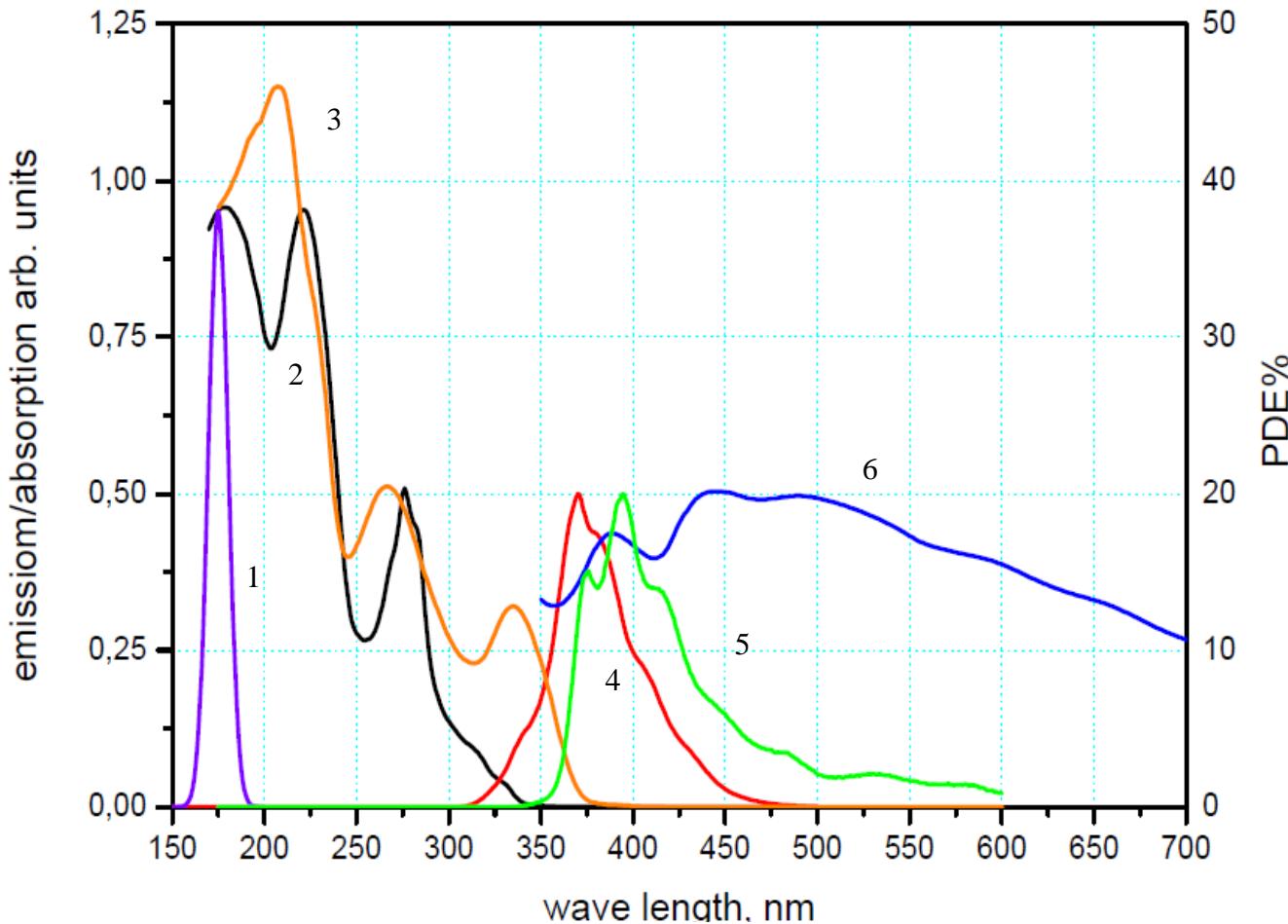
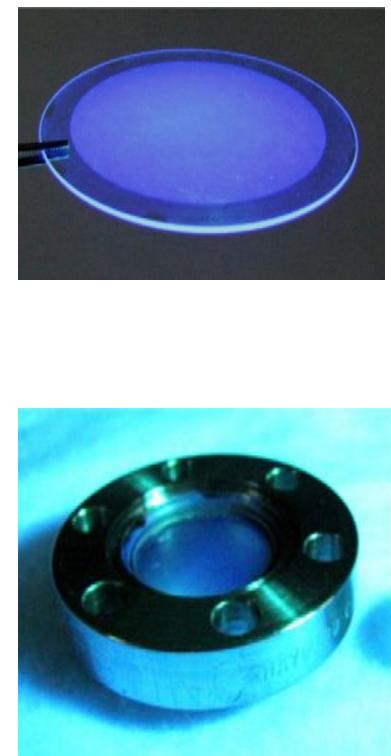
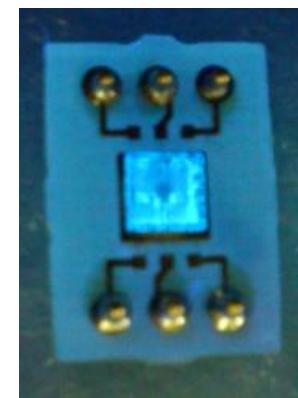
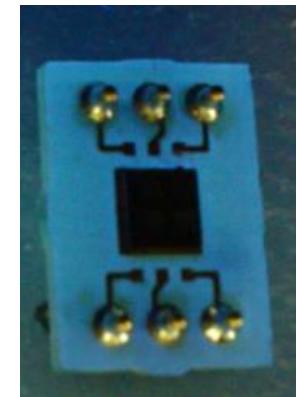
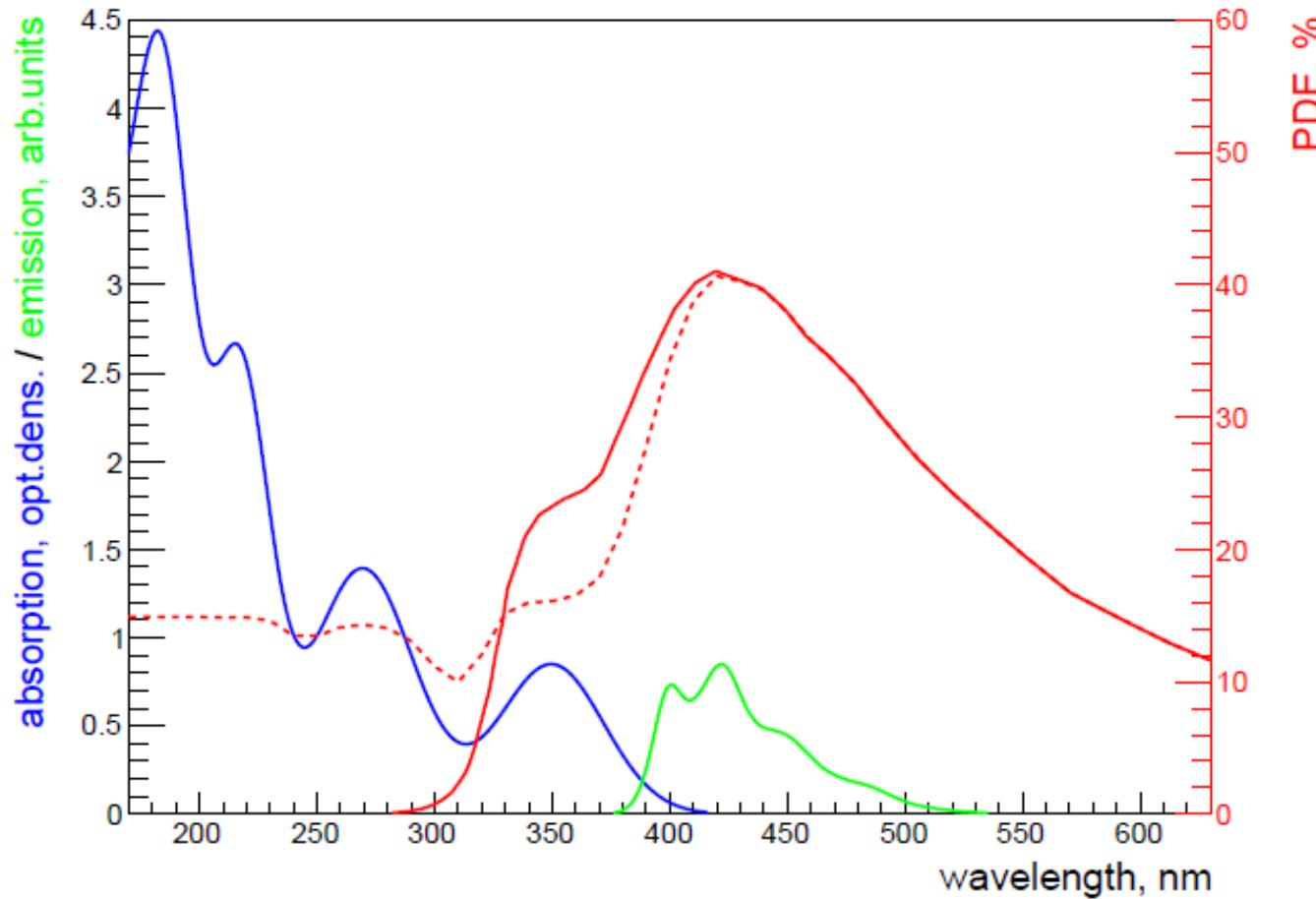


Fig 1. Emission spectrum of LXe (1), absorption spectrum of p-terphenyl (2), absorption spectrum of new WLS (3), emission spectrum of p-terphenyl (4), emission spectrum of new WLS (5), photon detection efficiency (PDE) of the CPTA “blue-sensitive” photodiode (6), right axis.



NOL-based VUV wavelength shifters for noble gas detectors

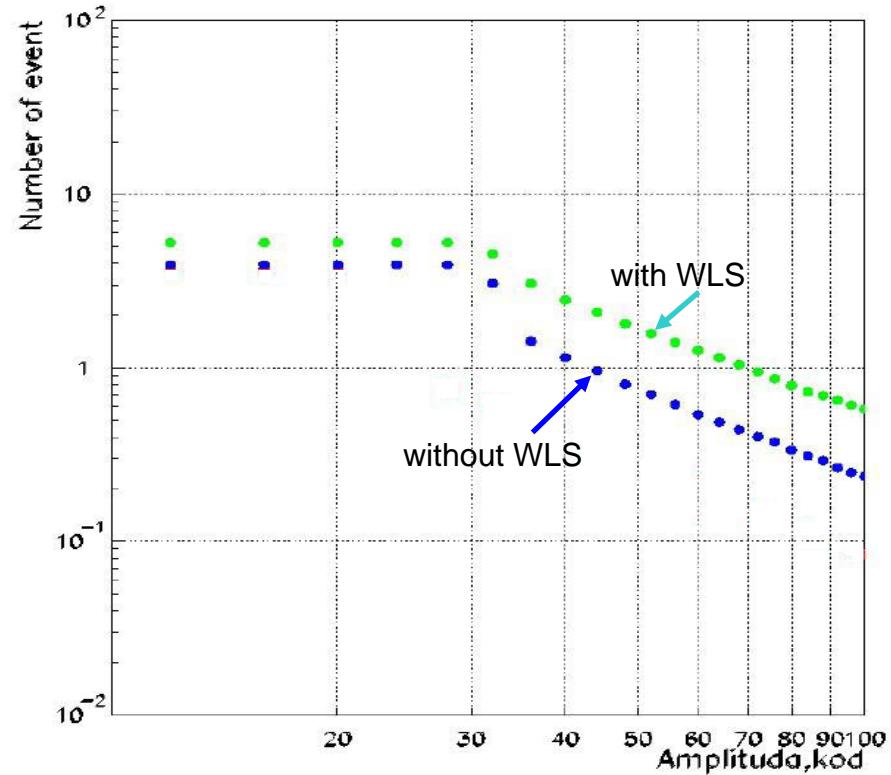
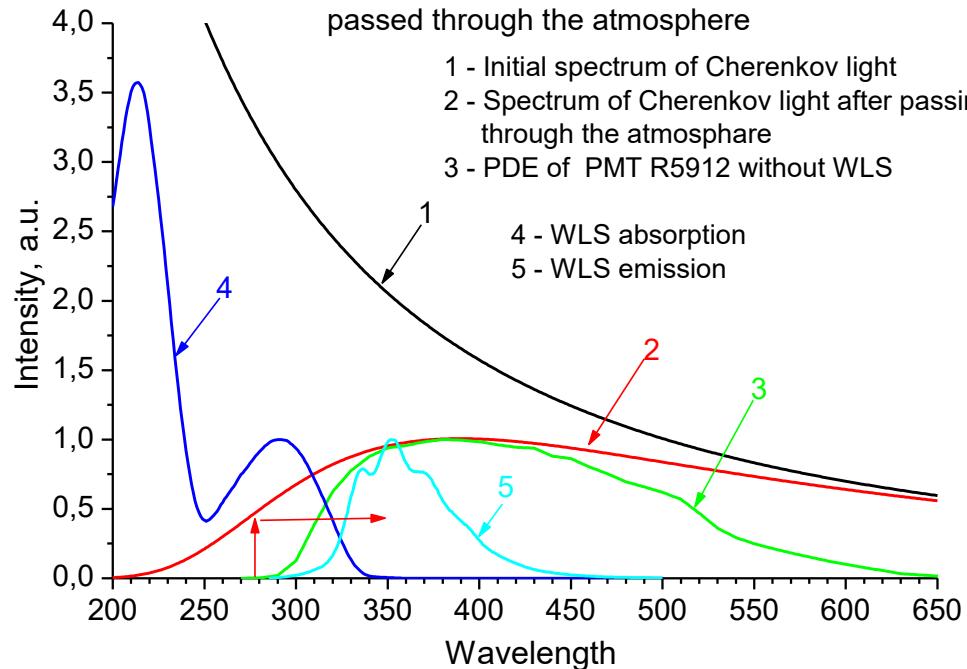


Absorption (blue) and luminescence (green) spectra of NOL-1, as well as spectral dependence of SiPM PDE without (red solid) and with (red dashed) 200 nm layer of NOL-1

Akimov D.A., et al., *J. Instrum.* 2017, 12, P05014

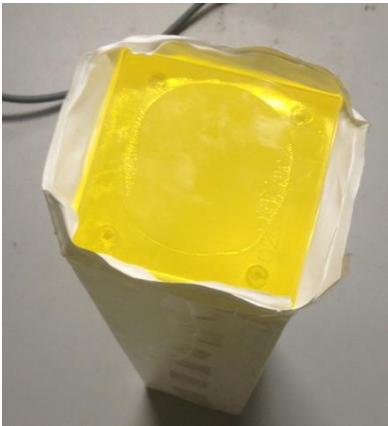
NOLs as effective wavelength shifters for TUNKA

Specral scheme of work
of the wavelength shifters of Cherenkov light
passed through the atmosphere

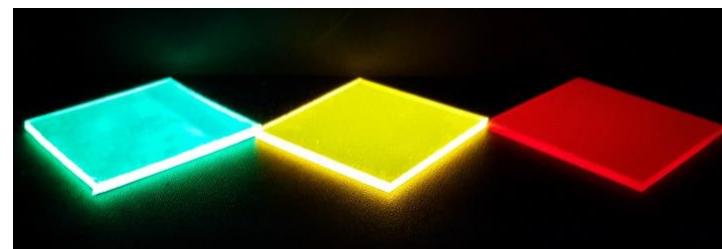
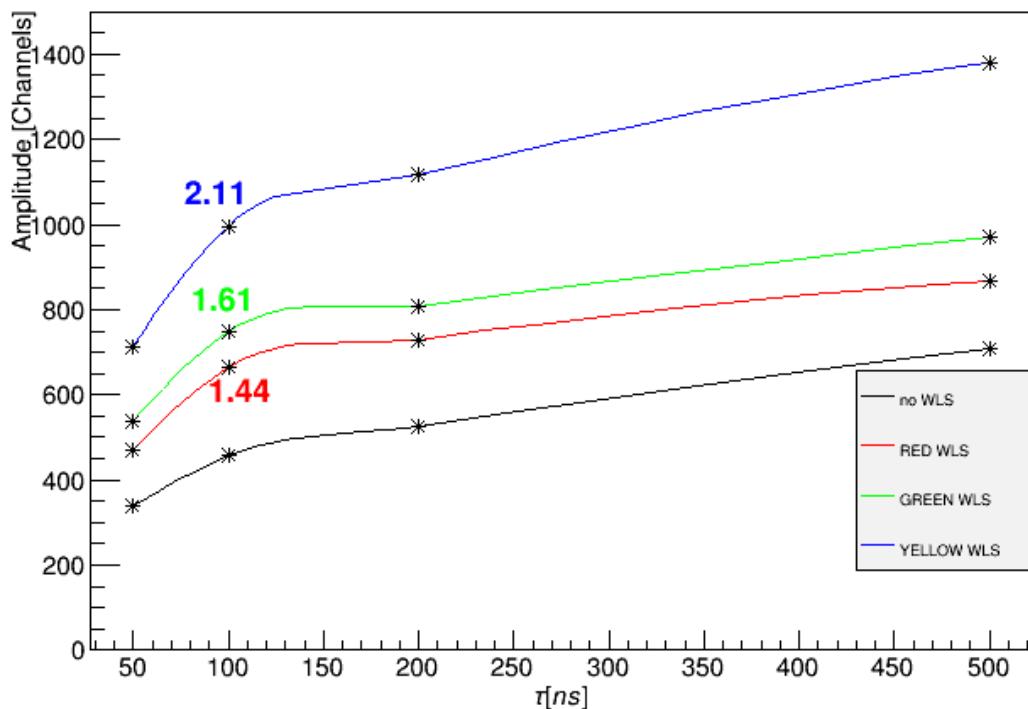
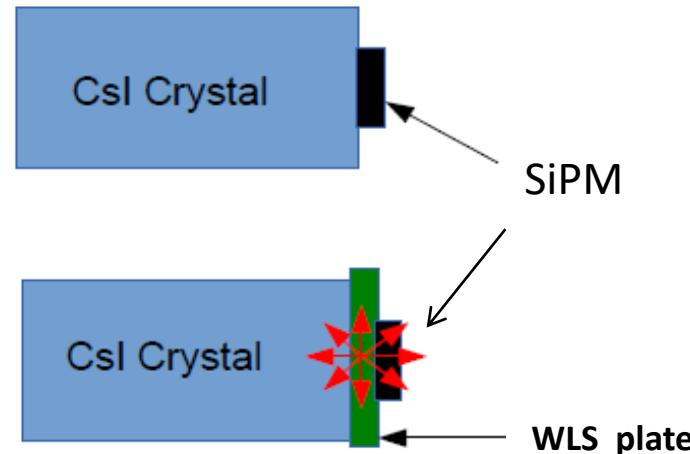


Photos of the optical station of
TUNKA-HiSCORE and PMT
R5912-TUNKA

NOL-based VUV WLS plates for CsI single crystals



signal-shaping time



Usage of WLS plates based on NOLs allows **2-3 times** to increase the detection efficiency

Jin, Y., Aihara, H., Borshchev, O.V., Epifanov, D.A., Ponomarenko, S.A., Surin, N.M., *Nucl. Instr. Meth. Phys. Res. A*, 2016, 824, 691-692

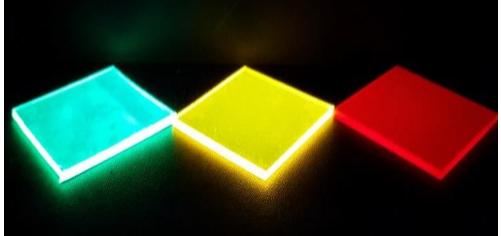
Summary of various application of NOLs

VUV wavelength shifters for noble gas detectors



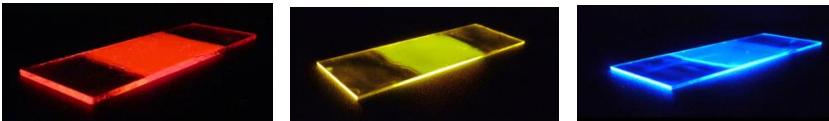
NIM A, 2012, 695
J. Instrum. 2017, 12,
P05014

UV wavelength shifters for pure CsI detectors

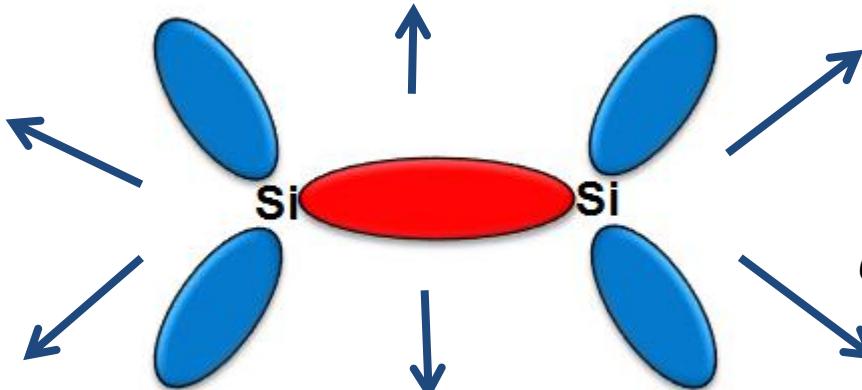


NIM A, 2016,
824, 691-692

Organosilicon scintillators



Silicon, 2015, 7, 191-200

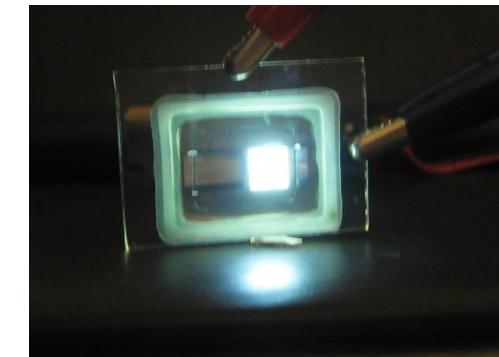


Plastic scintillators,
scintillating fibers



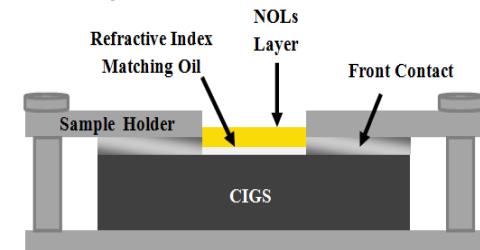
Sci. Rep. 2014, 4, 6549
J. Instrum. 2017, 12, P05013

Wavelength shifters in OLEDs



Org. Photonics Photovolt.,
2015, 3, 148-155

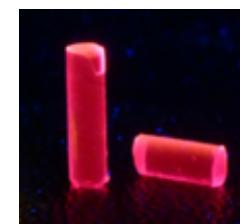
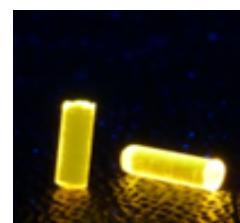
Wavelength shifters
for photovoltaics



Sol. Energy Mater. Sol. Cells
2016, 155, 1-8

A library of nanostructured organosilicon luminophores (NOLs)

	Absorption, λ_{max} , nm	Luminescence, λ_{max} , nm	Luminescence quantum yield Q_F , %
NOL 1	213, 262, 342	390, 412	73
NOL 2	213, 262, 366	390, 420	78
NOL 3	213, 262, 335	373, 390	85
NOL 4	213, 262, 367	396, 420	96
NOL 5	213, 262, 375, 396	416, 436	82
NOL 6	213, 262, 316, 457	588	87
NOL 7	296, 367	396, 419	96
NOL 8	337, 348	396, 419	85
NOL 9	327, 455	588	95
NOL 10	337, 513	655	78
NOL 11	332	396, 420	93
NOL 12	213, 262, 342	396, 420	99
NOL 13	319, 400	502	83
NOL 14	302, 402	502	90
NOL 15	375, 455	588	65
NOL 16	296, 386	486	94
NOL 17	340, 385	438, 468	94
NOL 18	328, 388	438, 468	93



Conclusions

Nanostructured Organosilicon Luminophores (NOLs) are a new class of nanomaterials with a combination of unique properties:

- ✓ large and tunable absorption in a wide optical spectral region
- ✓ tunable luminescence spectra in the visible spectral range
- ✓ high photoluminescence quantum yield (up to 99%)
- ✓ large «Stokes shift» (up to 330 nm)
- ✓ short luminescence lifetime
- ✓ good solution processability

Usage of NOLs can significantly increase the efficiency of different photonics devices used in high energy physics, such as Cherenkov detectors, plastics scintillators and scintillating fibers, Noble gas detectors, pure CsI scintillation detectors and other optoelectronic devices.

Asknowledgements

ISPM RAS (Moscow, Russia)

Yury Luponosov

Eugenia Svidchenko

Marina Polinskaya

CERN (Geneva, Switzerland)

Christian Joram

Lukas Grube

ITEP (Moscow, Russia)

Dmitry Akimov

Alexander Akindinov

Ivan Alexandrov

Thank you for your attention!

IPAC RAS (Chernogolovka, Russia)

Sergey Pisarev

IPCP RAS (Chernogolovka, Russia)

Prof. Pavel Troshin

Diana Susarova

Kuraray (Tokyo, Japan)

Osamu Shinji

University of Tokyo (Japan)

Denis Epifanov

Yi-Fan Jin

H. Aihara

G. E. Simakov

Viktor Stekhanov

O. Ya. Zeldovich

ZAE Bayern (Germany)

Prof. Christoph Brabec

Anastasia Solodovnyk

Taylor Uekert