The Abdus Salam
International Centre for Theoretical Physics

United Nations
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1938-15

Workshop on Nanoscience for Solar Energy Conversion

27 - 29 October 2008

Molecular Engineering of Sensitisers for Solar Cells Applications

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Molecular Engineering of Sensitizers for Solar Cell Applications

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Joint ICTP-KFAS Workshop on Nanoscience for Solar Energy Conversion
27-29 October 2008

Acknowledgment

Prof. M. Graetzel

Synthesis

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Dr. P. Péchy

Dr. Barolo (Italy)
Prof. Viscardi (Italy)

Prof. David L. Officer
University of Wollongong
Australia

Photovoltaic measurements

Dr. J-H Yum
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Dr. Ines Raabe
Takeru Bessho
Pascal Comte

Prof. Tomas Torres
Universidad Autonoma de Madrid

Prof. Jaejung Ko
Korea University

DFT Computational study

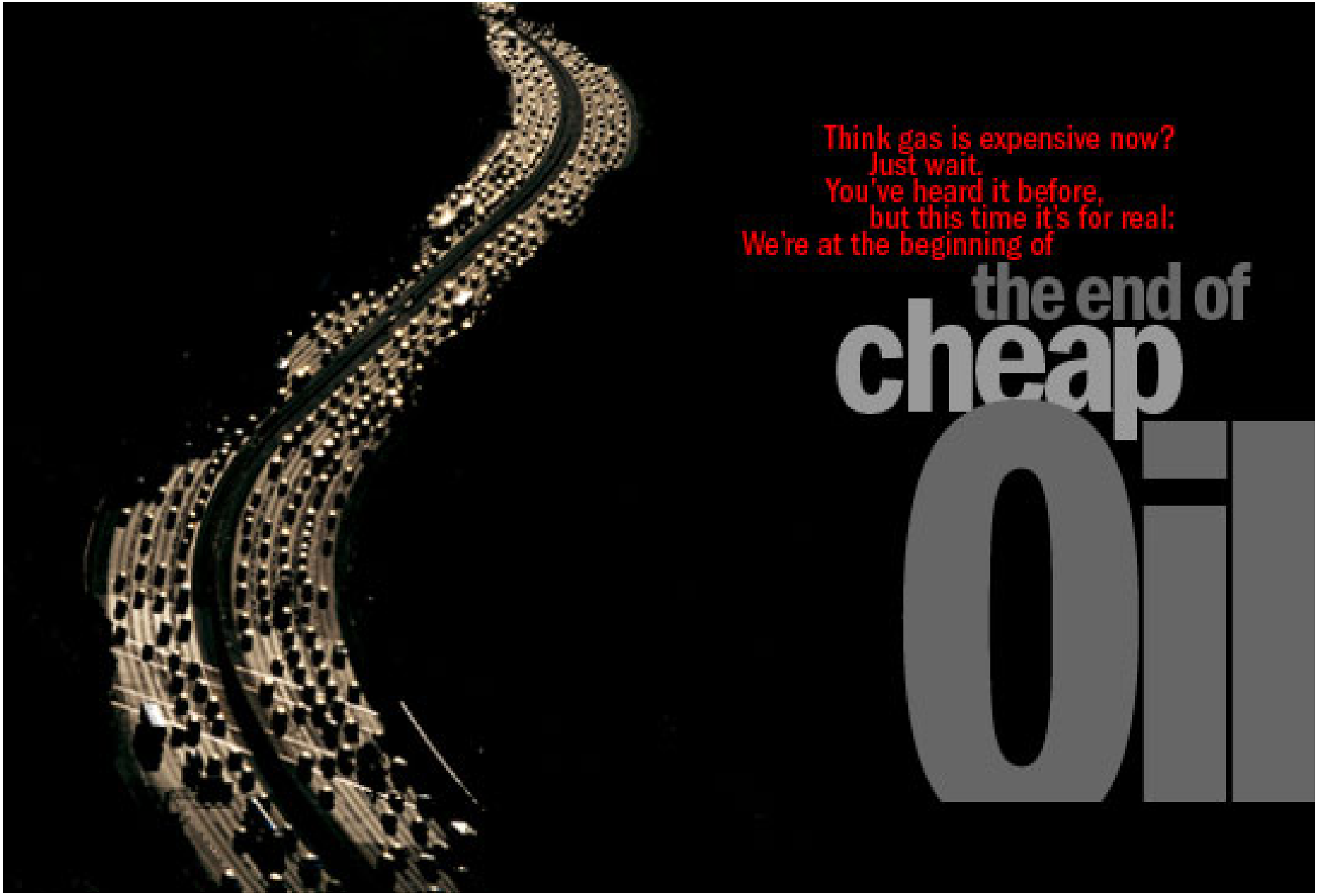
Prof. Filippo De Angelis,
Perugia, Italy

Prof. Ursula Roethlisberger,

EPFL

Financial support

Swiss Federal Institute of Technology
Swiss Federal Office for Energy (OFEN)



National Geography, June 2004

Real Outdoor Test of DSC Modules

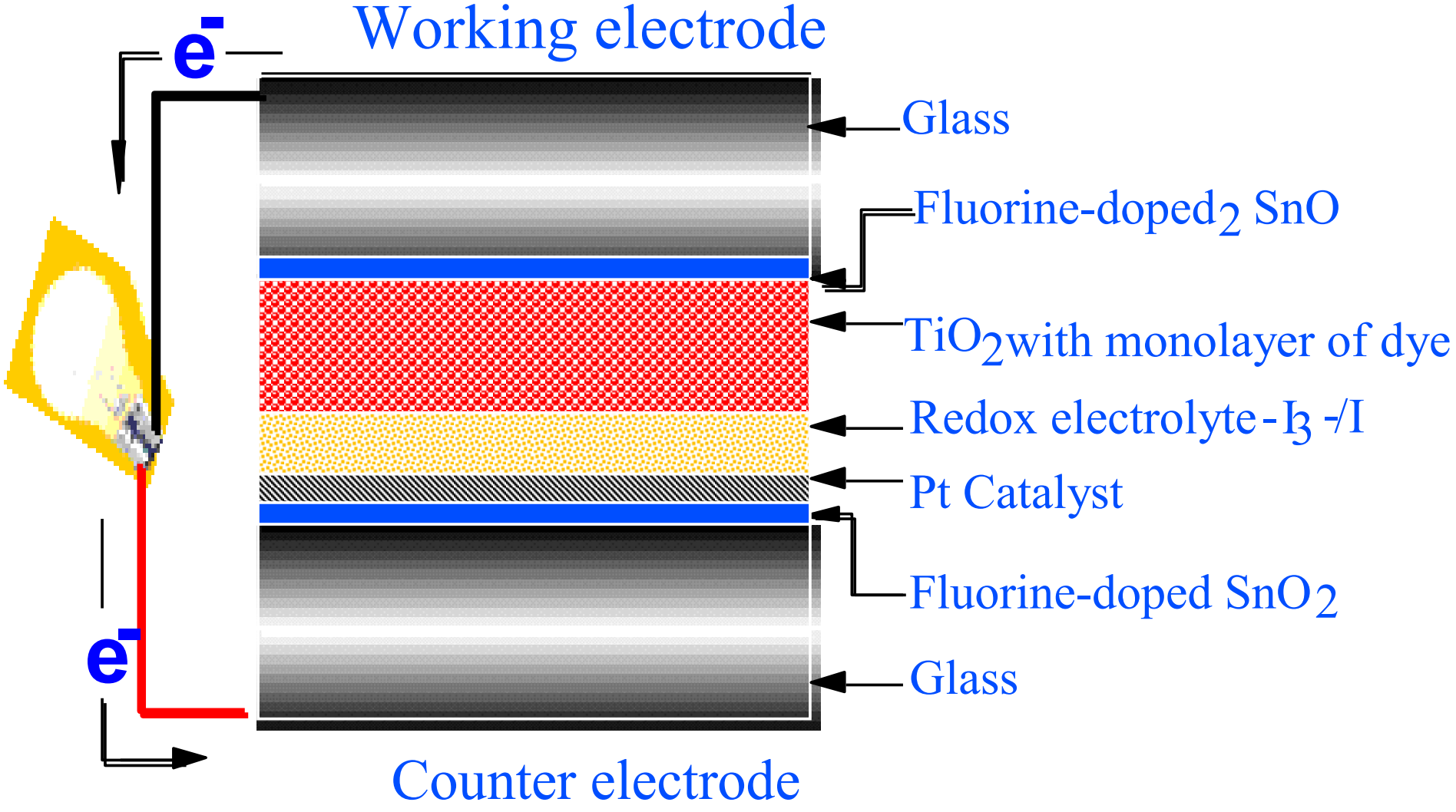
- Module Unit

Series connected
64 DSC cells

- Outdoor Test

Kariya City at lat. 35°10’N,
Asimuthal angle: 0°
Facing due south, Tilted at 30°

A cross section of the dye sensitized solar cell



Requirements of the Sensitizers

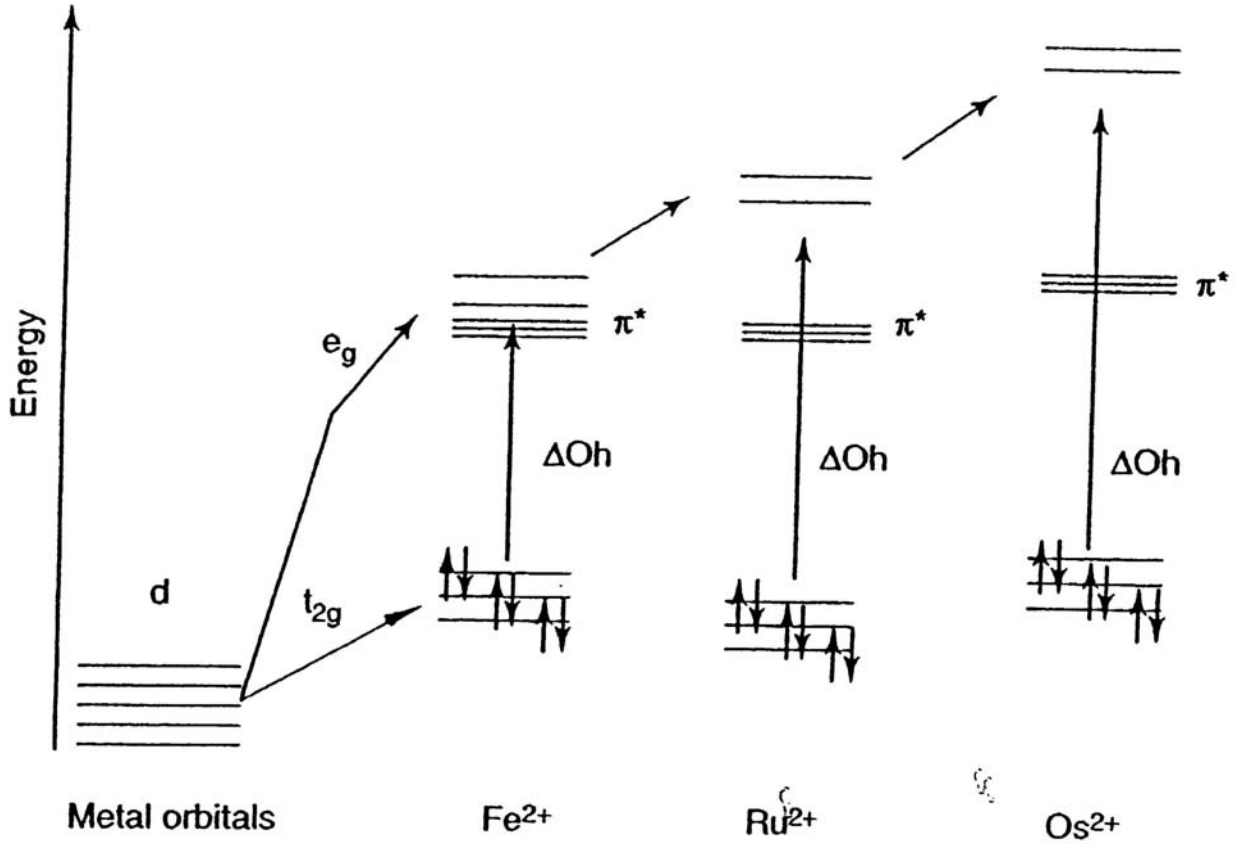
+ The optimal sensitizer for the dye sensitized solar cell should be panchromatic, i.e. absorb visible light of all colors.

+ It should possess suitable ground- and excited state redox properties (0.5 and -0.8 V vs.SCE)

+ It should exhibit thermal and photochemical stability

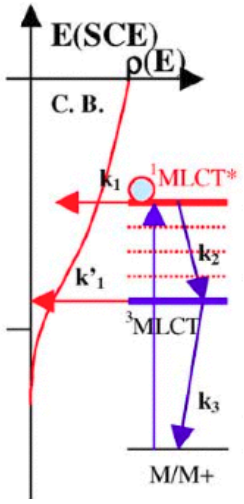
+ It must be firmly grafted to the semiconductor oxide surface and inject electrons into the conduction band with a quantum yield of unity.

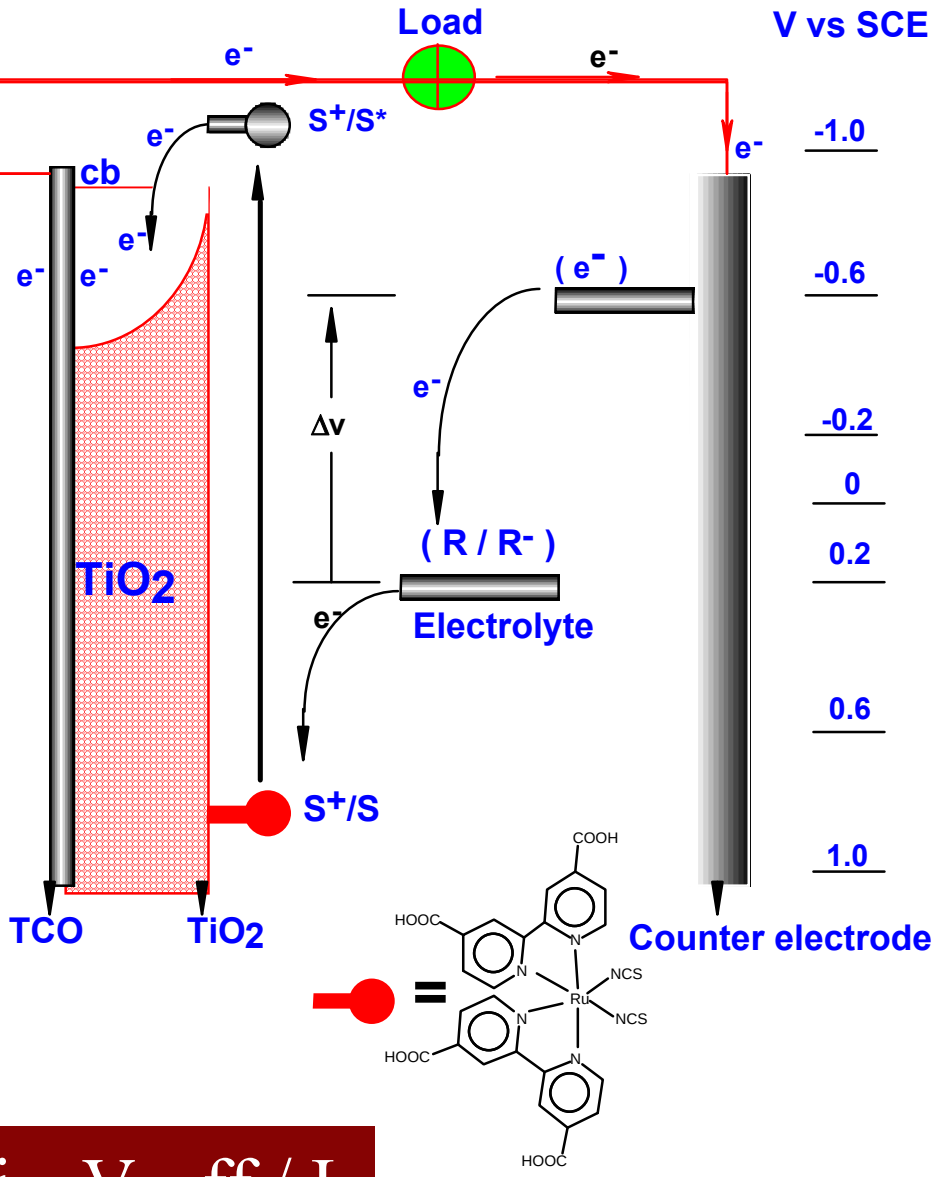
Splitting pattern of d-orbital in an octahedral field for Fe^{2+}, Ru^{2+} and Os^{2+}



VIIB 8	VIIB 9	VIII 10
26 Fe +2 +3 +6	27 Co +2 +3	28 Ni +2 +3
55,85 Geležis	58,93 Kobaltas	58,69 Nikelis
44 Ru +2 +3 +4 +6 +8	45 Rh +2 +3 +4 +5	46 Pd +2 +4
101,07 Rutenis	102,91 Rodis	106,42 Paladis
76 Os +3 +4 +6 +8	77 Ir +2 +3 +4 +6	78 Pt +2 +4 +6
190,2 Osmis	192,22 Iridis	195,08 Platina

Operating Principles of the Dye-Sensitized Solar Cell





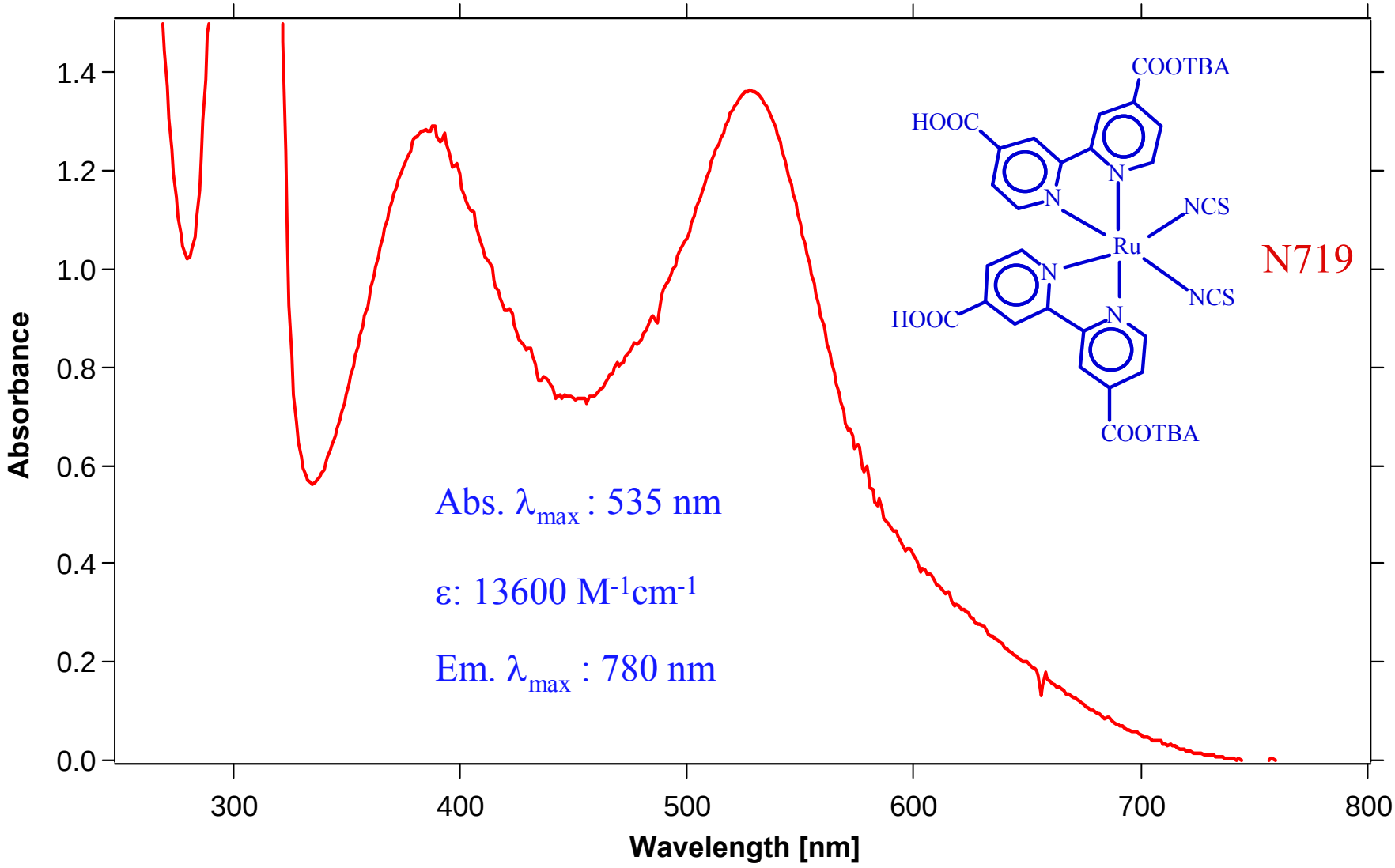
Electron injection from the excited dye into the TiO2 conduction band (CB) is a very fast process *femto*second scale $\tau 1 < 20$ fs.

The reduction of the oxidized dye by the redox electrolyte's I- ions occurs in about 10^{-8} seconds.

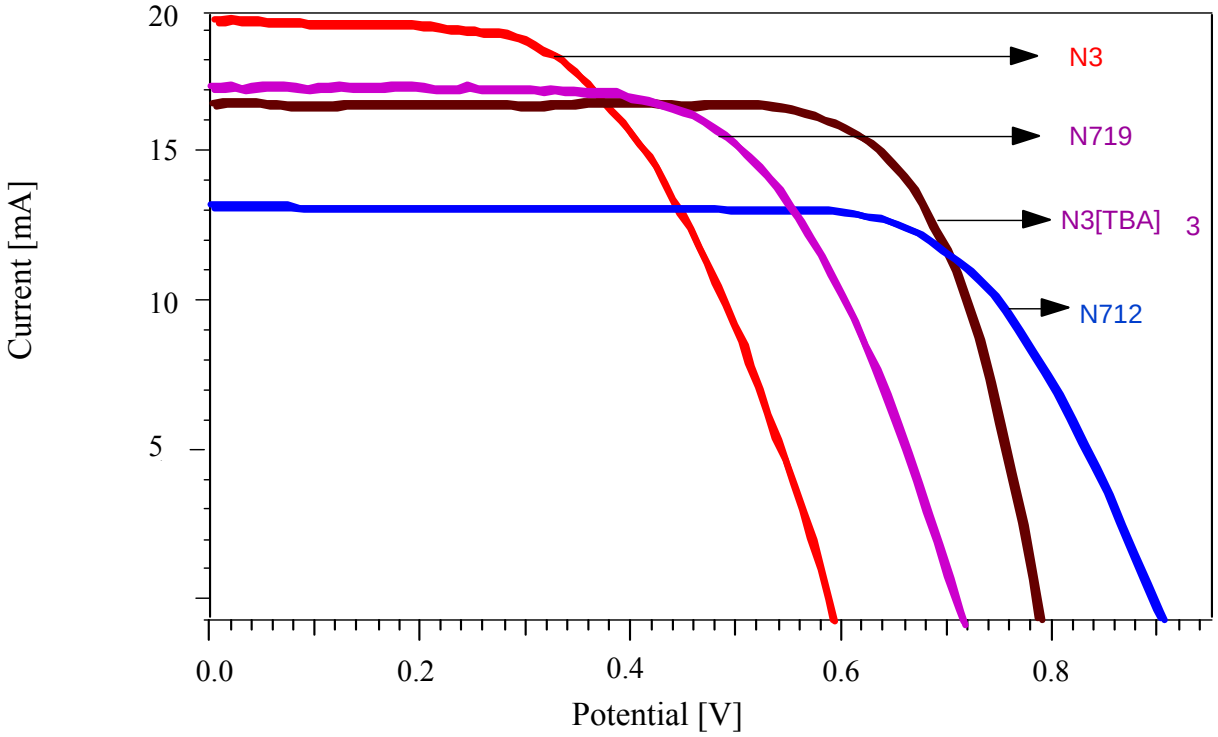
Recombination of photoinjected CB electrons with oxidized dye molecules or with the oxidized form of the electrolyte redox couple (I_3^- ions) occurs in *micro*seconds.

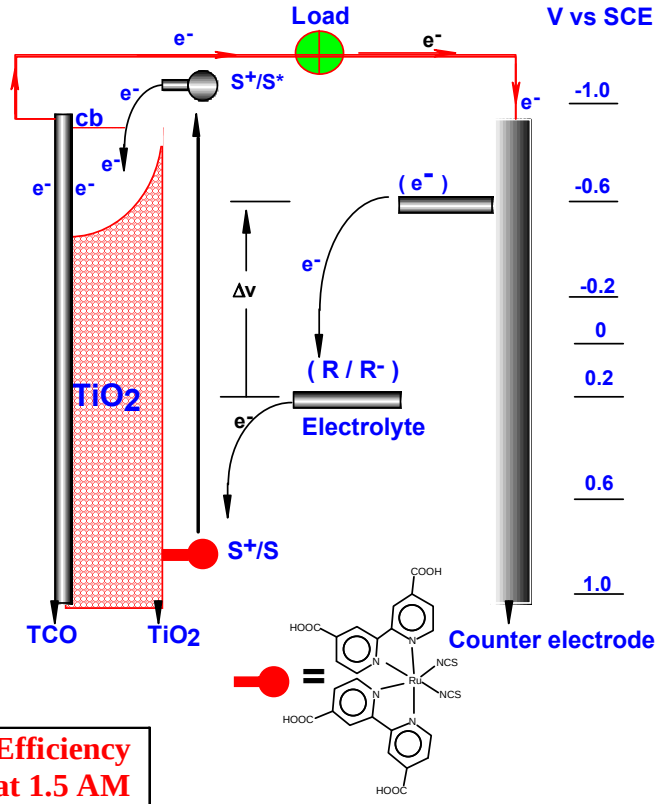
$$\eta = i_{ph}\, V_{oc}\, ff / I_s$$

Absorption Spectra of N719 Sensitizer



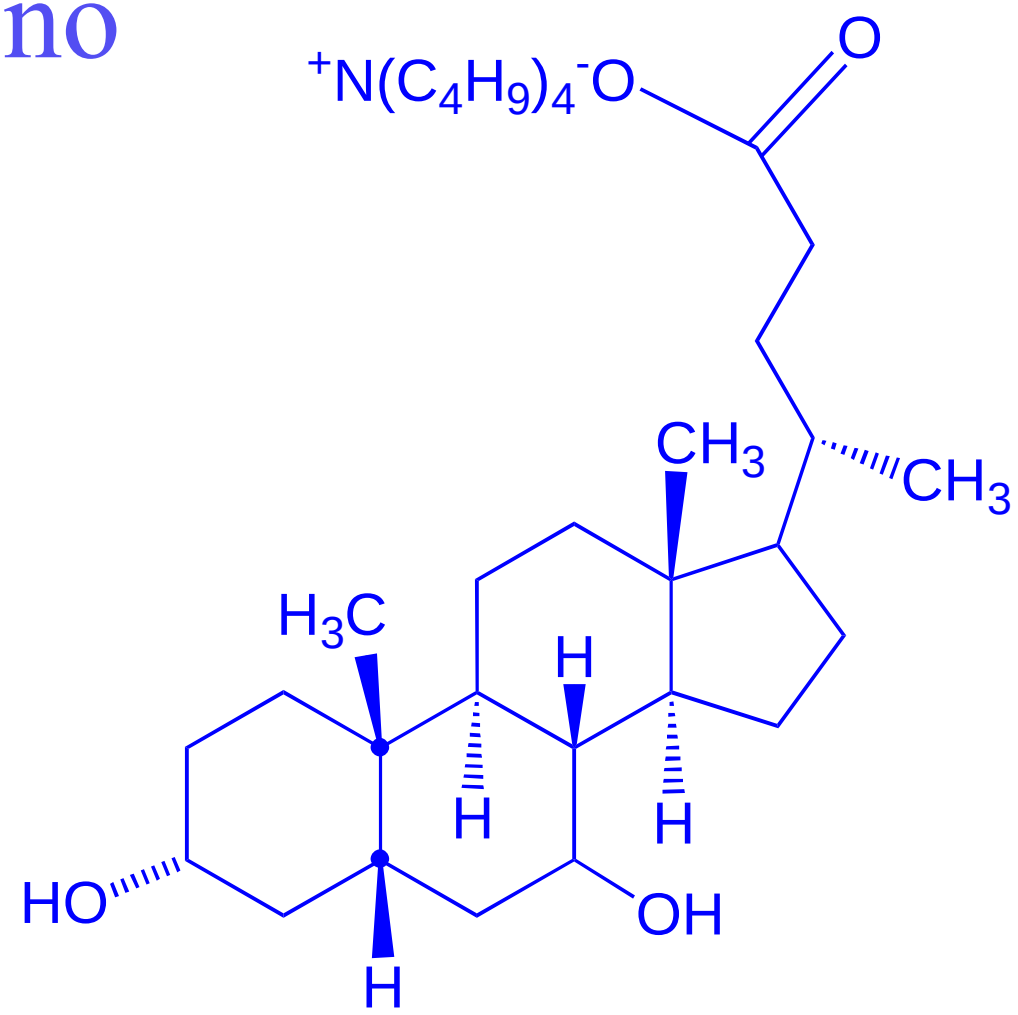
Photocurrent-voltage characteristics of nanocrystalline TiO_2 cell sensitized with N3 (4 protons), N719 (2 protons), $N3[TBA]_3$ (1 proton) and N712 (zero proton) dyes measured under AM 1.5 sun.





Sensitizer	Number of protons	Solvent for dye adsorption	Current mA/cm^2	Potential (mV)	Fill Factor	Efficiency at 1.5 AM
N3	4	1:1 CH_3CN+ *tert*-BuOH	19 ± 0.5	600 ± 30	0.65 ± 0.05	7.4
N719	2	1:1 CH_3CN+ *tert*-BuOH	16 ± 0.5	730 ± 30	0.70 ± 0.05	8.2
N712	0	C_2H_5OH	13 ± 0.5	900 ± 30	0.7 ± 0.05	8.2
$N3[TBA]_3$	1	5:95 CH_3CN +*tert*-BuOH	17±0.5	770 ±20	0.73±0.05	9.56
N3[TBA]	3	1:1 CH_3CN+ *tert*-BuOH	17 ± 0.5	700 ± 20	0.65 ±0.05	7.7

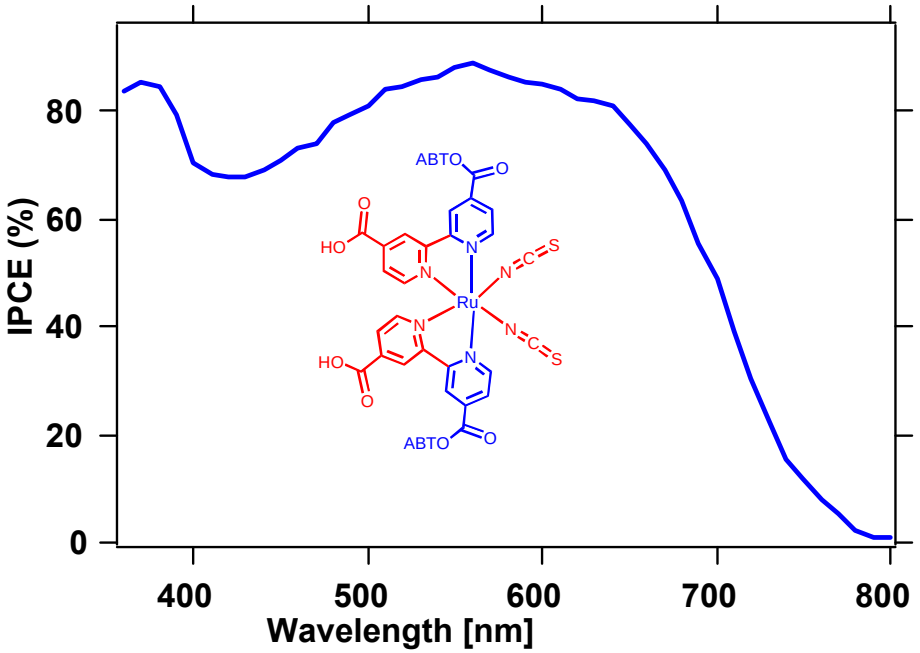
TBA Cheno

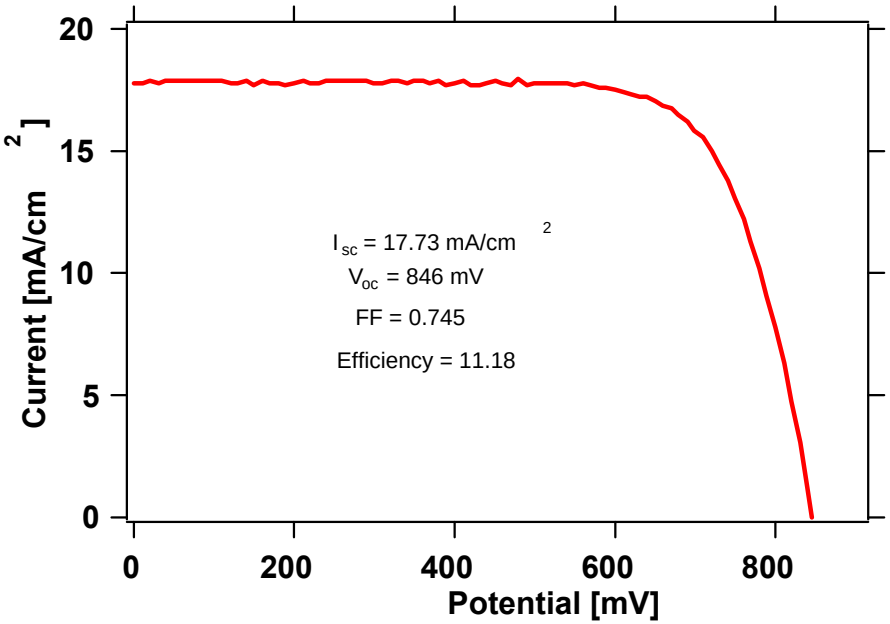


Chenodeoxycholic acid

3α,7α-dihydroxy-5β-cholani cacid

Conversion efficiency of 11.2 % have been reached under AM 1.5 sunlight

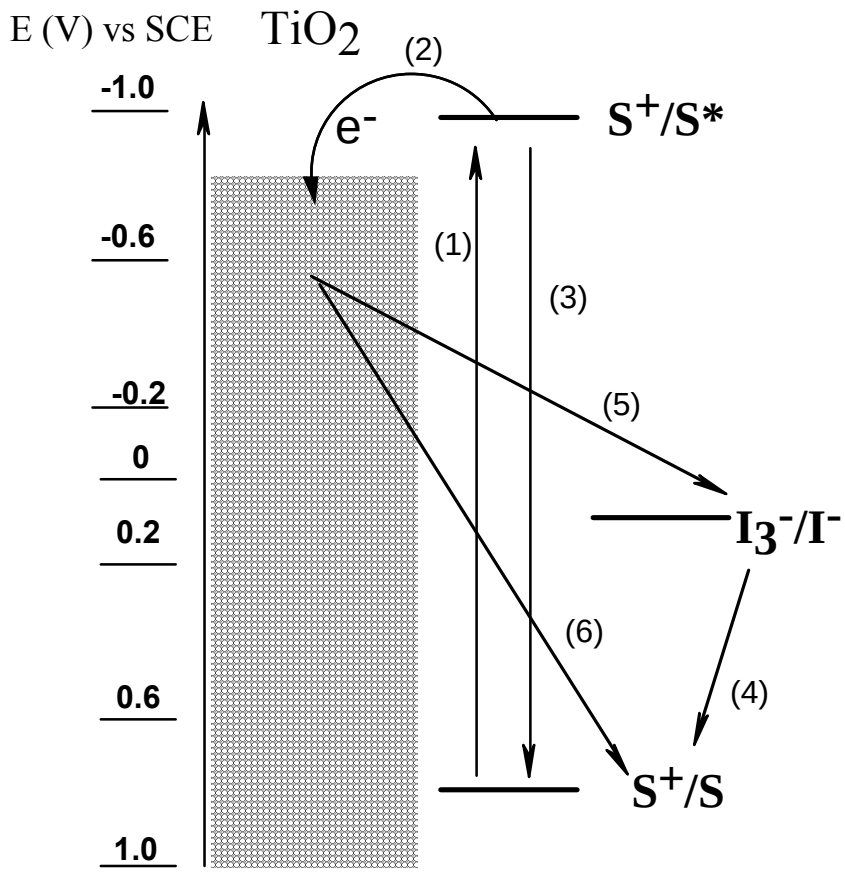




$$\eta = i_{ph}\, V_{oc}\, ff / I_s$$

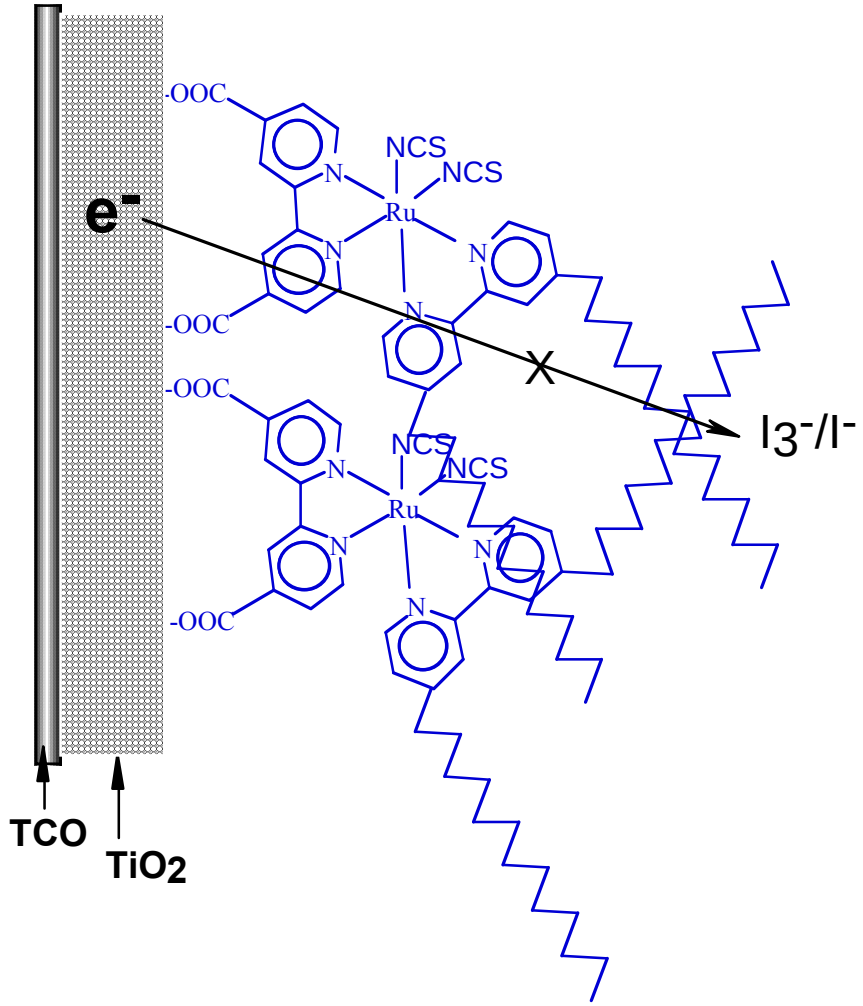
Nazeeruddin, Mohammad K.; De Angelis, Filippo; Fantacci, Simona; Selloni, Annabella; Viscardi, Guido; Liska, Paul; Ito, Seigo; Takeru, Bessho; Graetzel, Michael. JACS, 127, 16835, 2005.

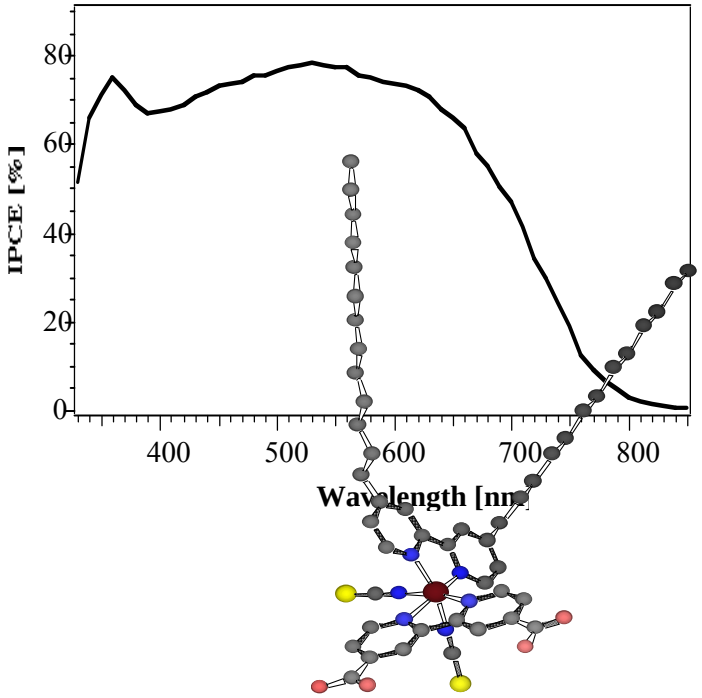
Illustration of the interfacial charge transfer processes in nanocrystalline dye sensitized solar cell.



(1) An excited state. (2) electron injection onto the conduction band of TiO_2. (4) The oxidized sensitizer gets reduced by I^- / I_3^- redox couple. (5) The injected electrons into the conduction band may react either with the oxidized redox couple or with oxidized dye molecule (6).

Pictorial representation of blocking of the oxidized redox couple I_3^- reaching onto the surface of TiO_2 for conduction band electrons using hydrophobic sensitizers, which forms aliphatic net work.



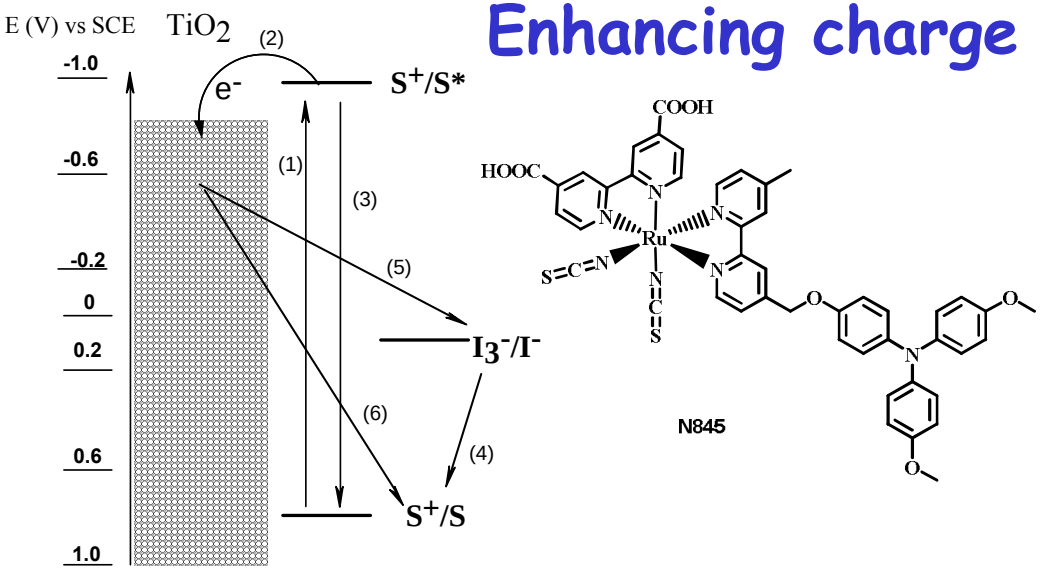


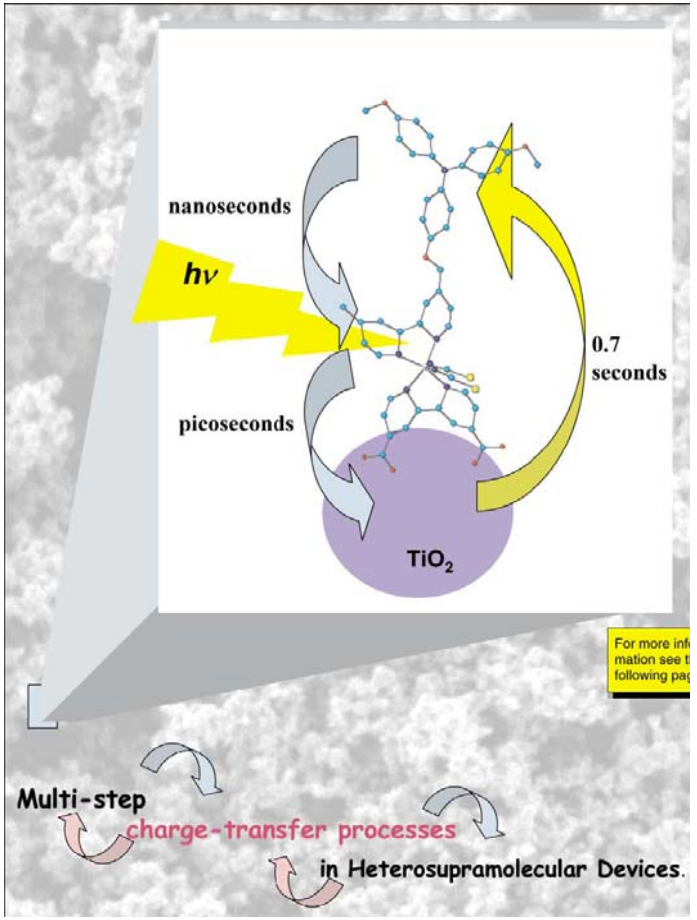
S. M. Zakeeruddin, M. K. Nazeeruddin, R. Humphry-Baker, P. Pechy, P. Quagliotto, C. Barolo, G. Viscardi, M. Grätzel, *Langmuir* 2002, **18**, 952

P. Wang, S. M. Zakeeruddin, J. E. Moser, M. K. Nazeeruddin, T. Sekiguchi, M. Grätzel, *Nat. Mater.* 2003, **2**, 402

M. K. Nazeeruddin, S. M. Zakeeruddin, J-J. Lagref, P. Liska, P. Comte, C. Barolo, G. Viscardi, K. Schenk, M. Graetzel *Coord. Chem. Rev.* 248 (13-14): 1317-1328 (**2004**)

Enhancing charge separation





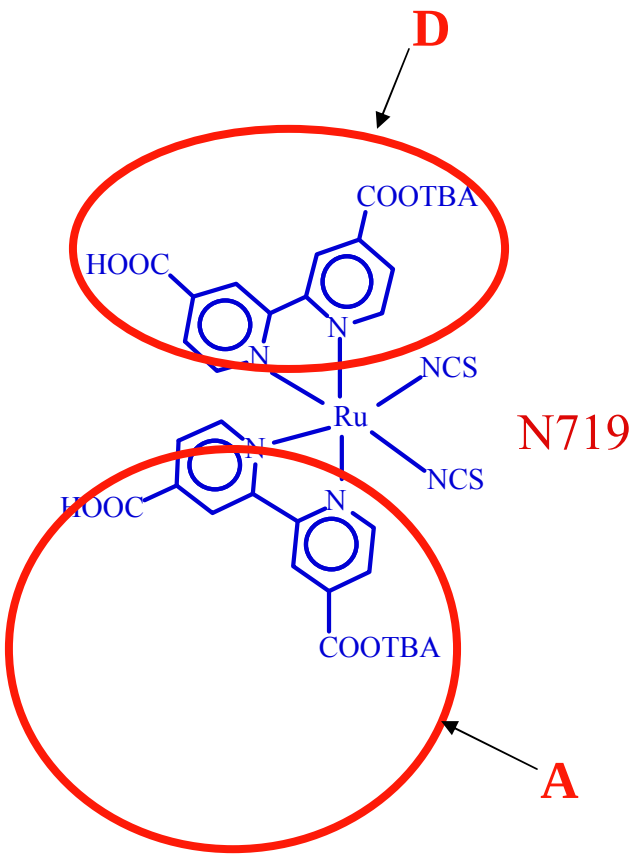
triarylamine moiety

1000-fold retardation of the recombination dynamics in comparison with N-719!

4 Å increase in distance between the cationic center of charge and the TiO_2 surface respect to N-719

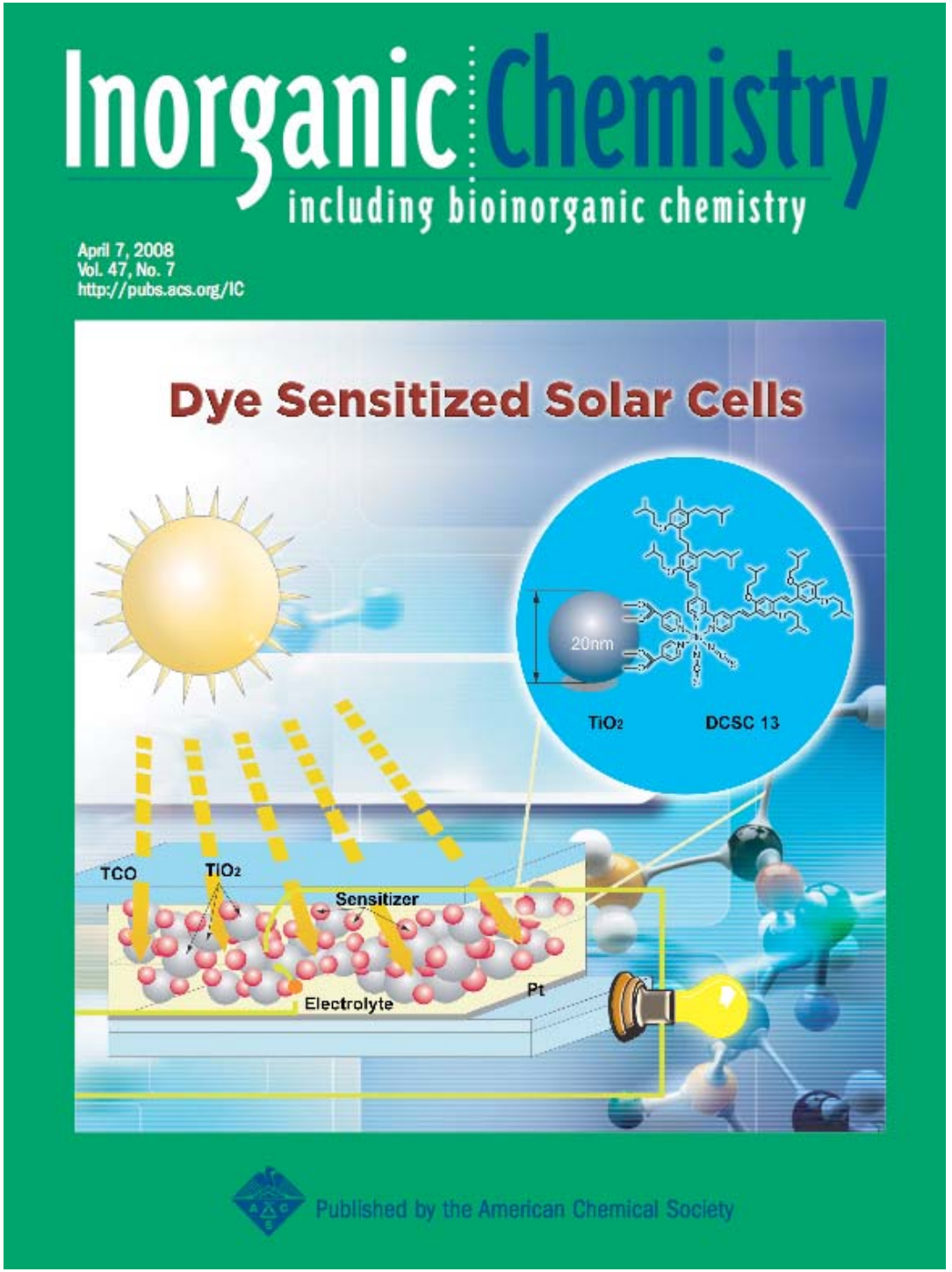
N. Hirata, J.-J. Lagref, E. J. Palomares, J. R. Durrant, M. K. Nazeeruddin, M. Graetzel, D. Di Censo, *Chem. Eur. J.* **2004** 10 595

New Sensitizers with π-extended donor ligands

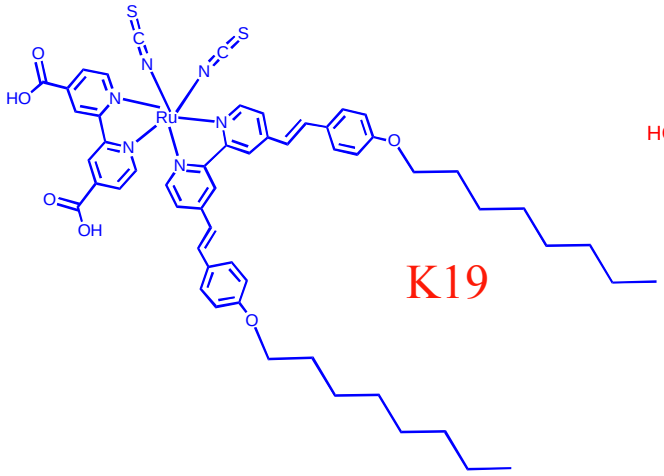


N719

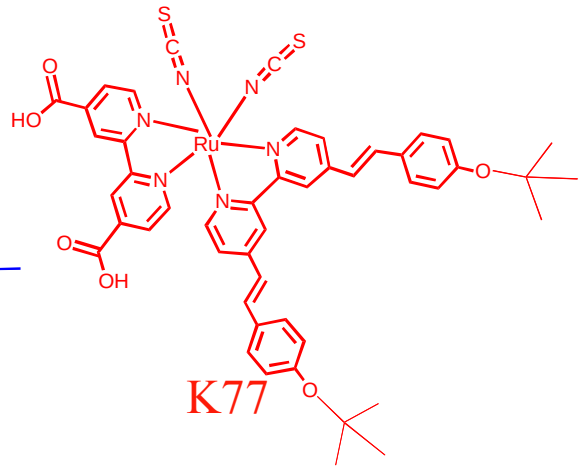
Ko et al. Inorganic Chemistry, Vol. 47, No. 7, **2008**, 2267



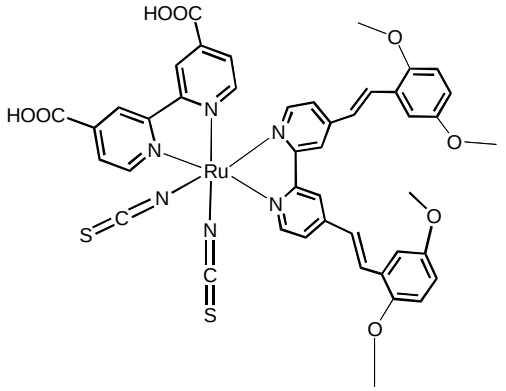
Sensitizers with π-extended donor ligands



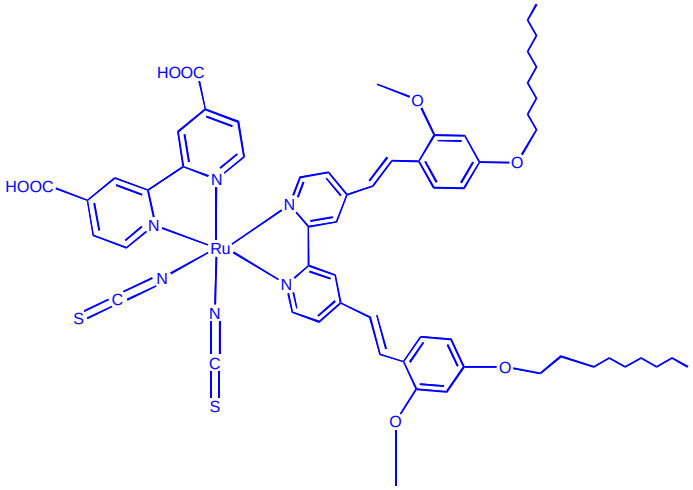
K19



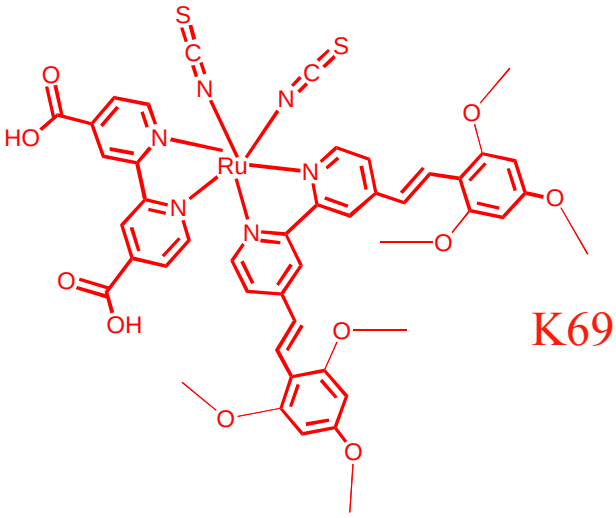
K77



N945



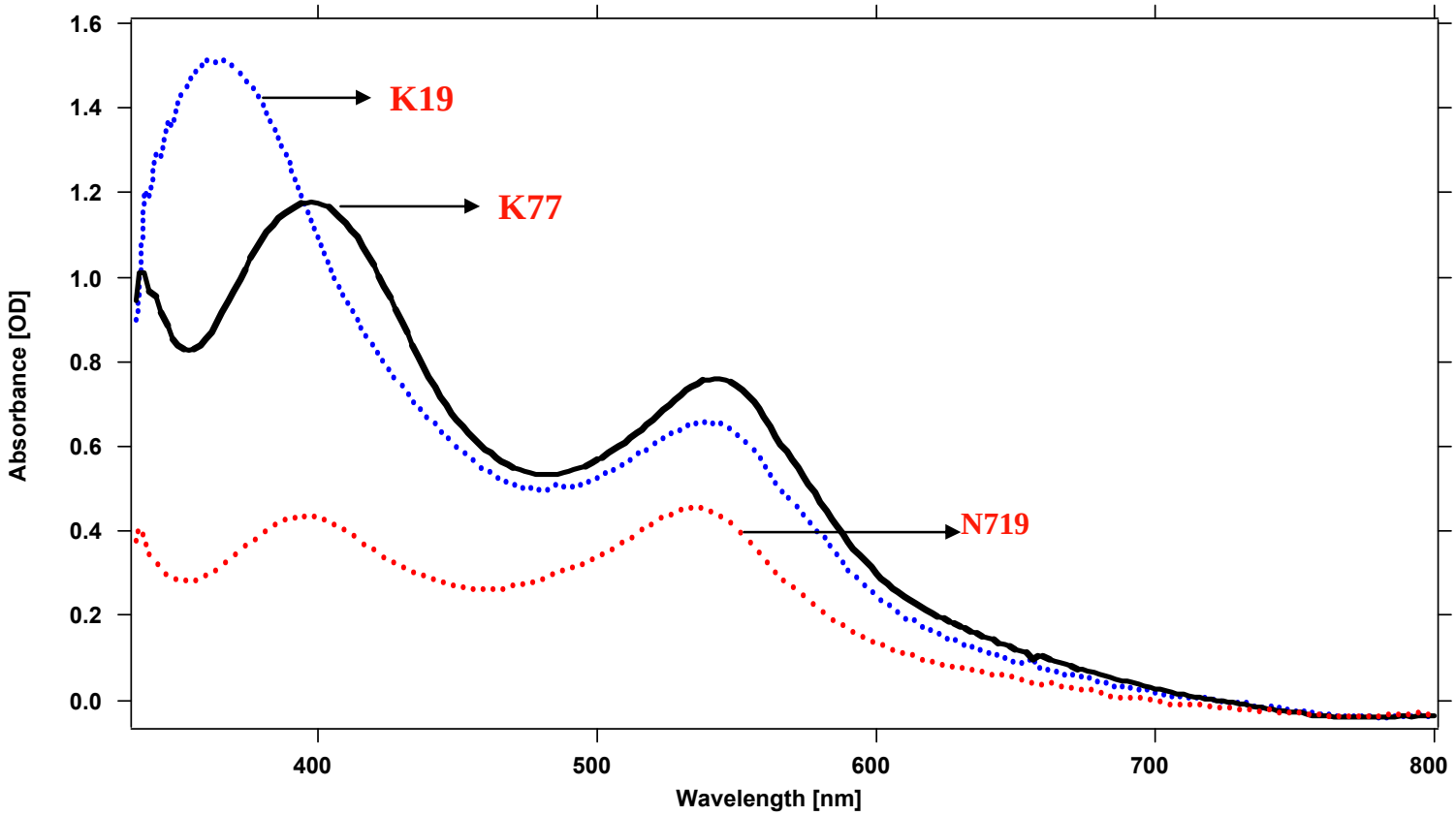
K66



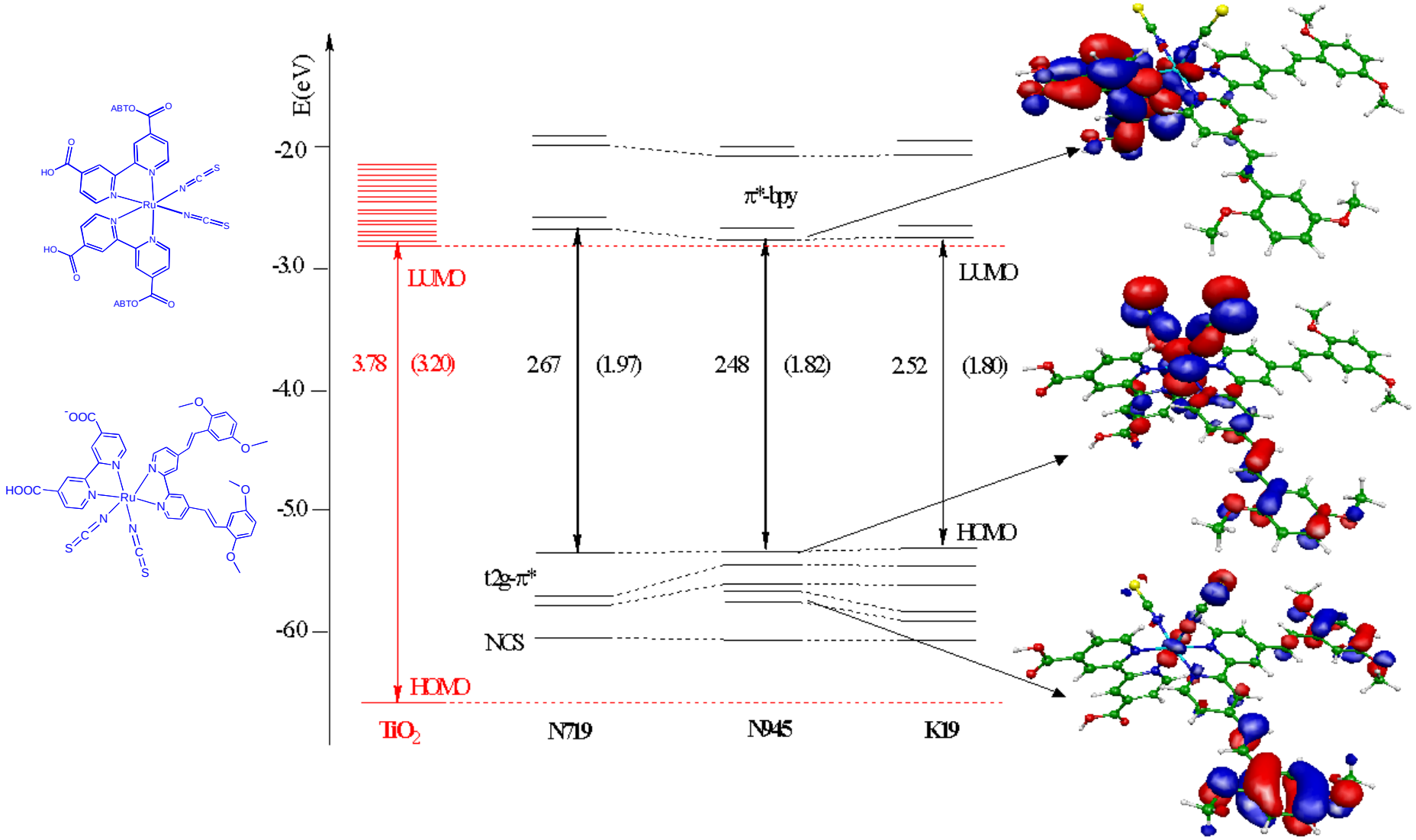
K69

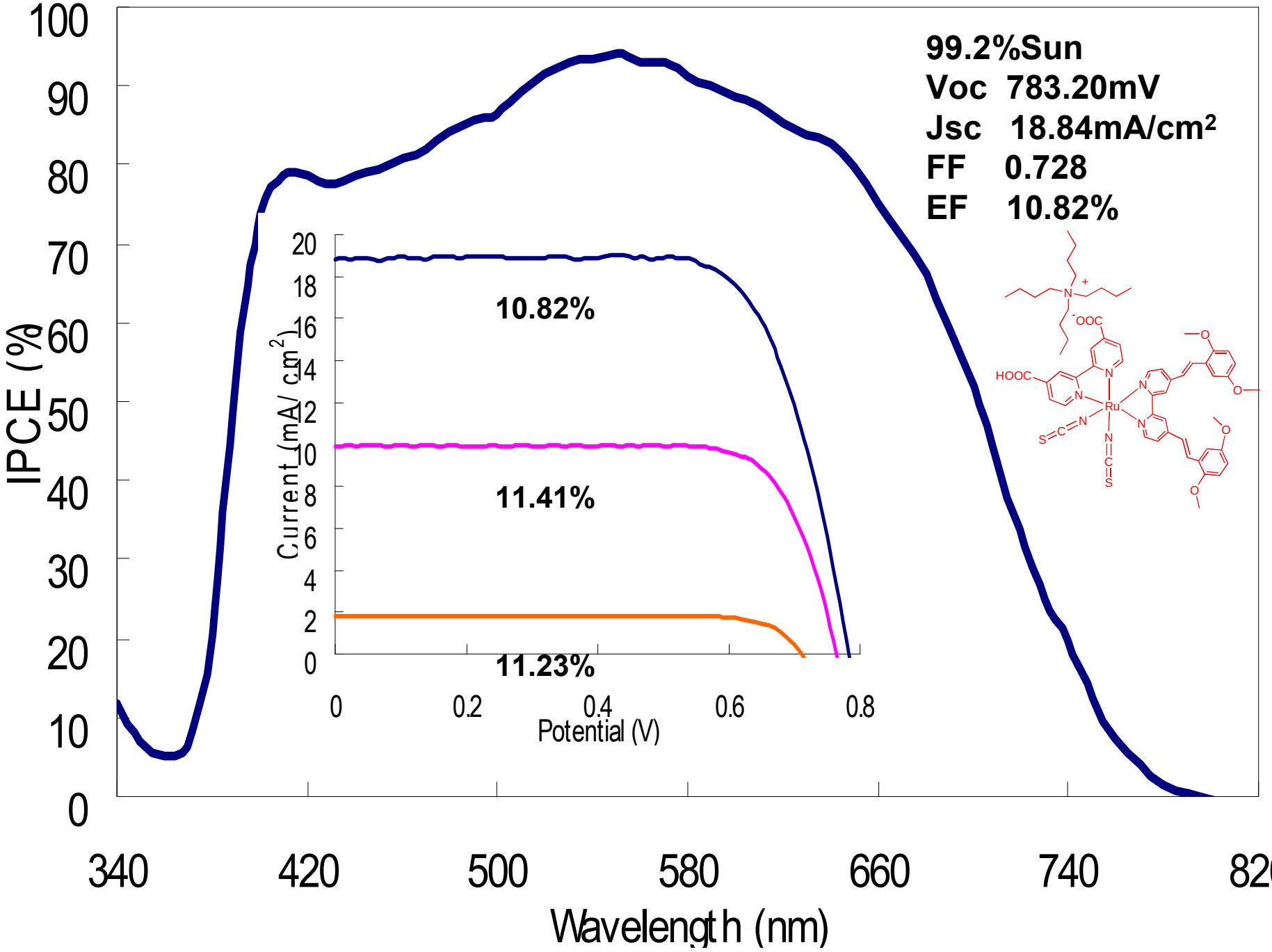
Inorg. Chem. 45, 787-797, 2006.

Comparison UV-Vis spectra of N719 (red), K19 (blue) and K77 (black)

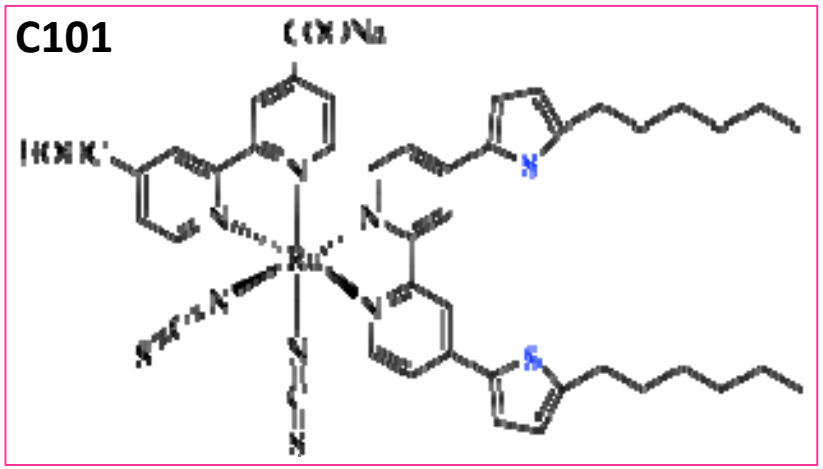


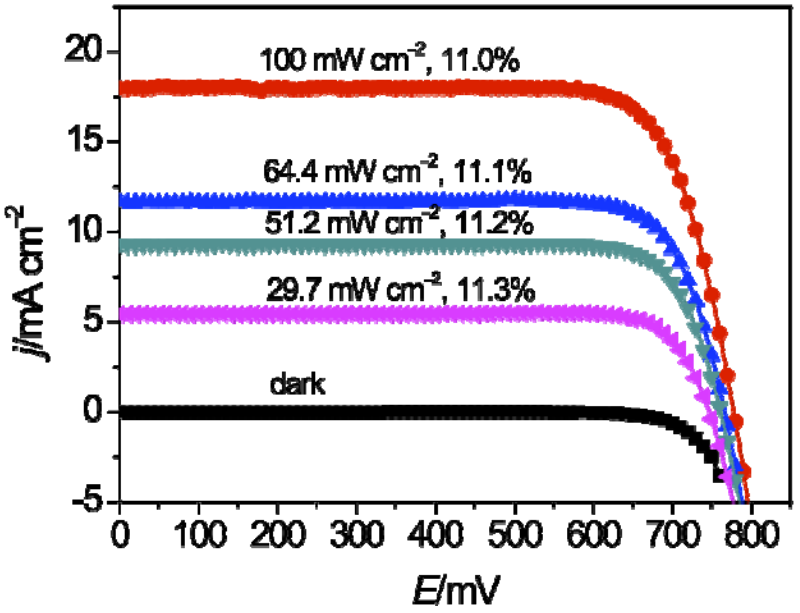
Molecular orbital energy diagram of N719, N945 and K19 compared to that of a TiO_2 nanoparticle model




99.2%Sun
Voc 783.20mV
Jsc 18.84mA/cm²
FF 0.728
EF 10.82%
IPCE (%)
Wavelength (nm)
Current (mA/ cm²)
Potential (V)
10.82%
11.41%
11.23%
HOOC
-OOC
Ru

C101



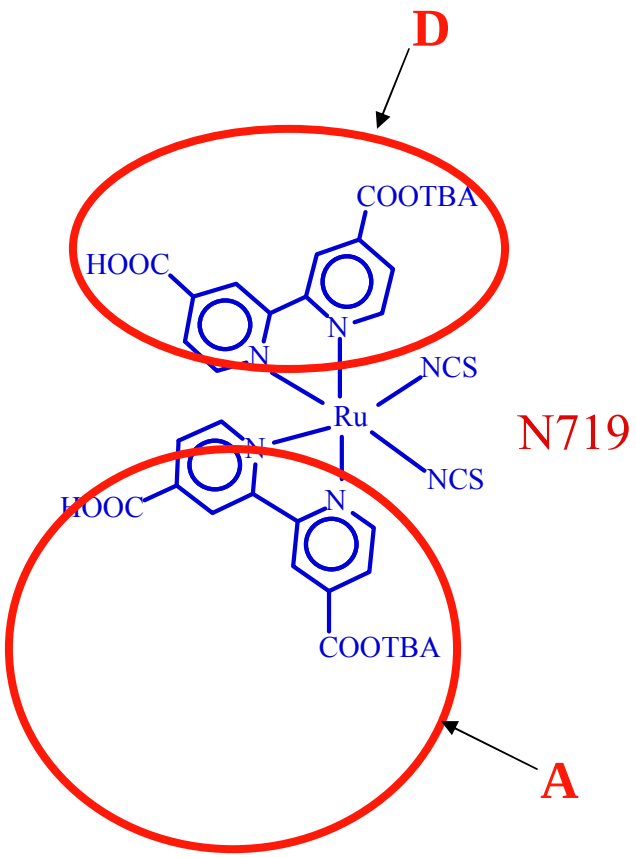


Note: 7+5 film, overnight
Dye solution: 300 ⌠M C101
and 300 ⌠M cheno in AN/t-BuOH
Electrolyte: Z960

J_{sc}=17.94 mA cm□2
V_{oc}=778 mV
ff=0.785
⌡=11.0%

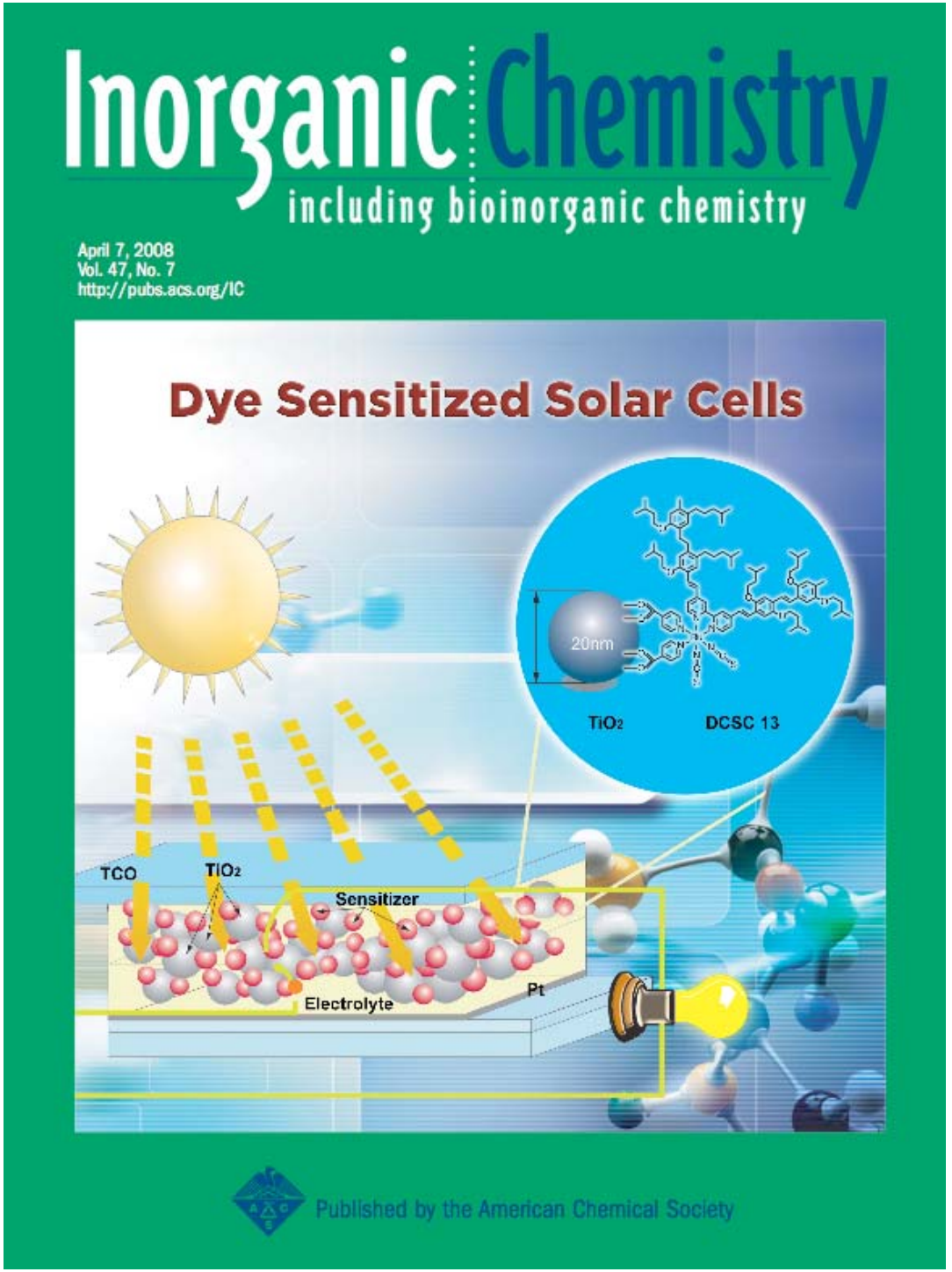
Peng Wang et al. J. AM. CHEM. SOC. 2008, *130*, 10720–10728

New Sensitizers with π-extended acceptor ligands

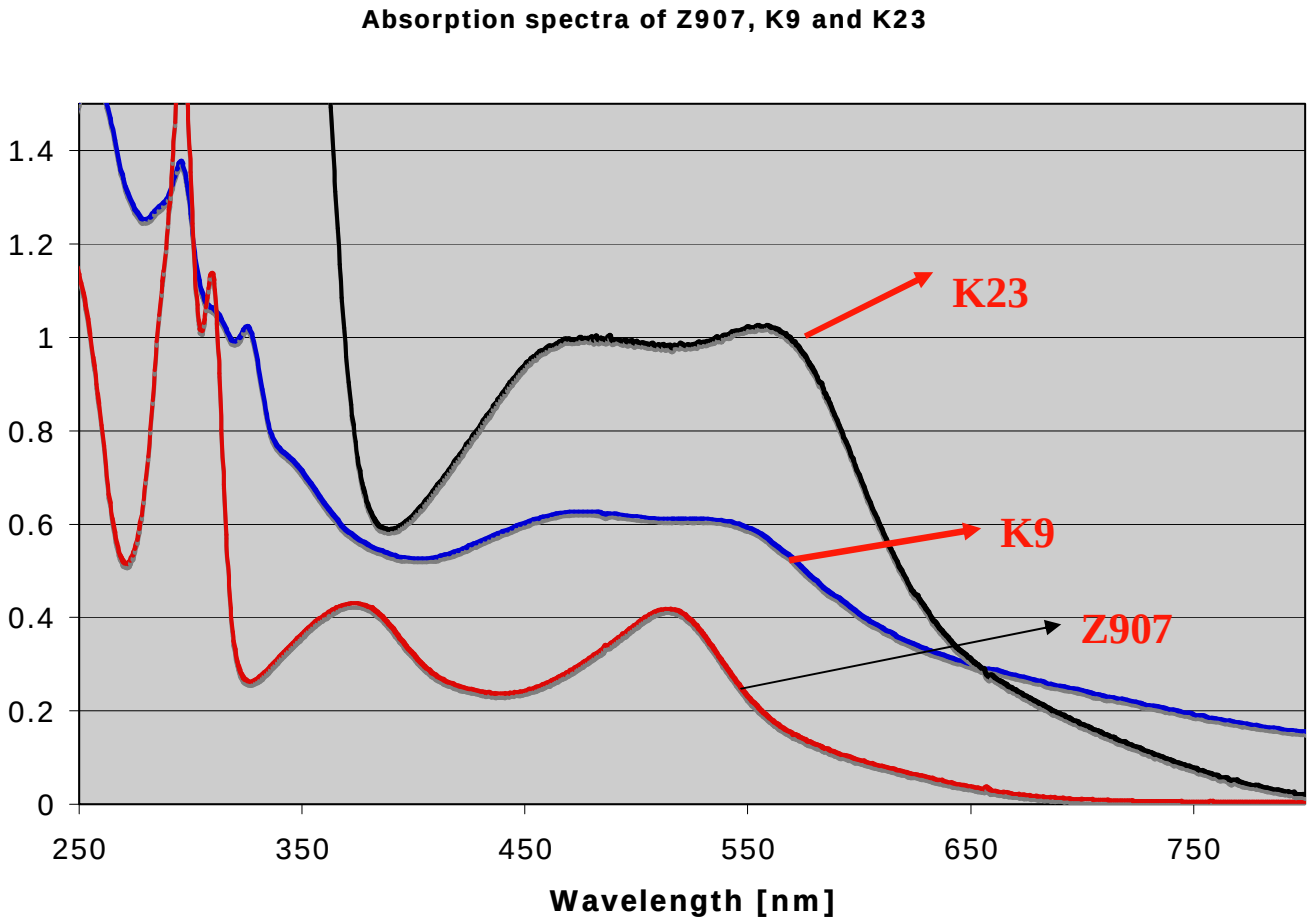


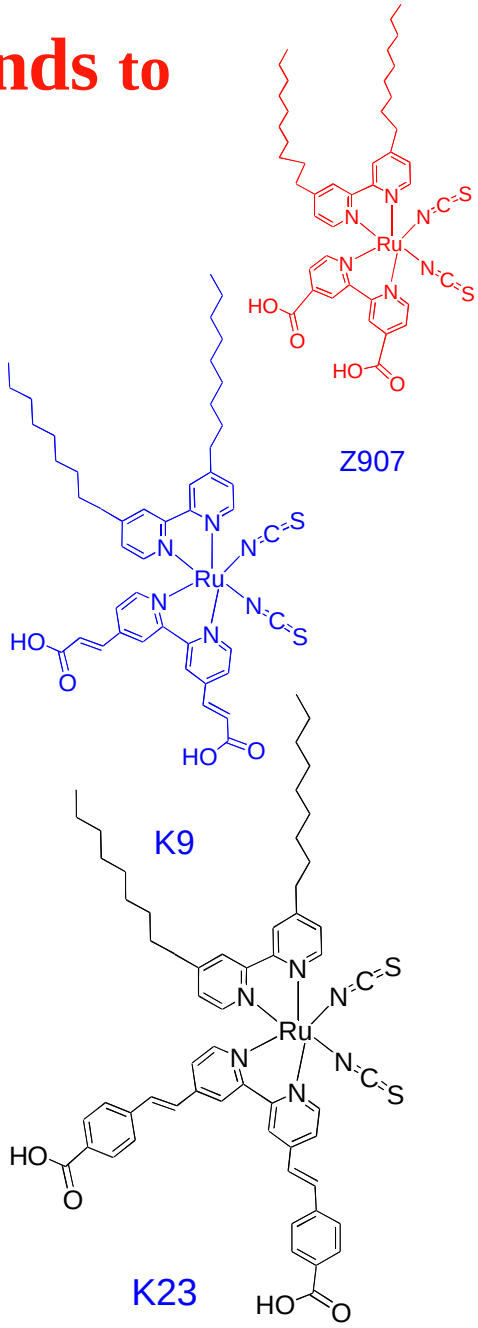
N719

Ko et al. Inorganic Chemistry, Vol. 47, No. 7, **2008**, 2267

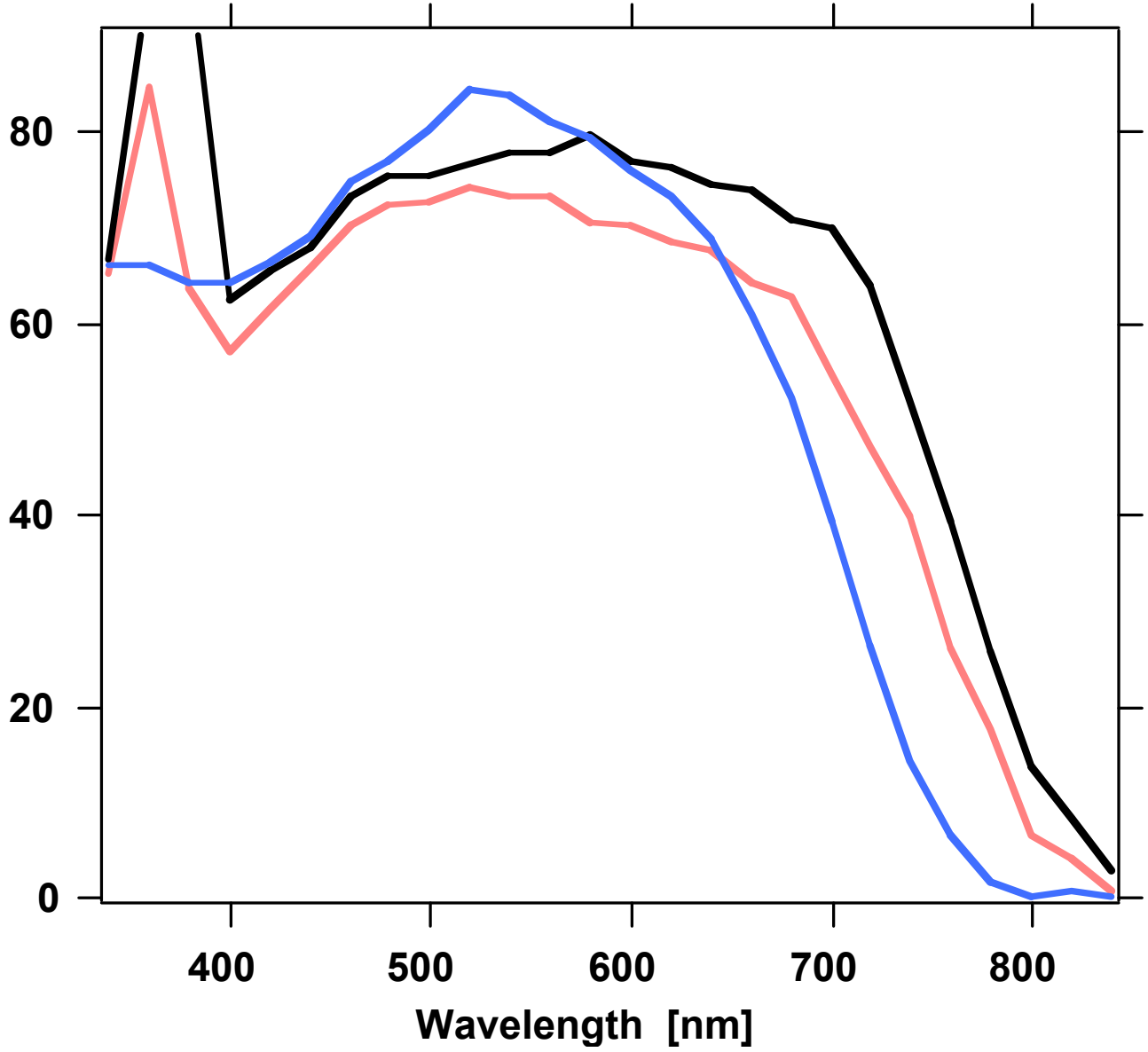


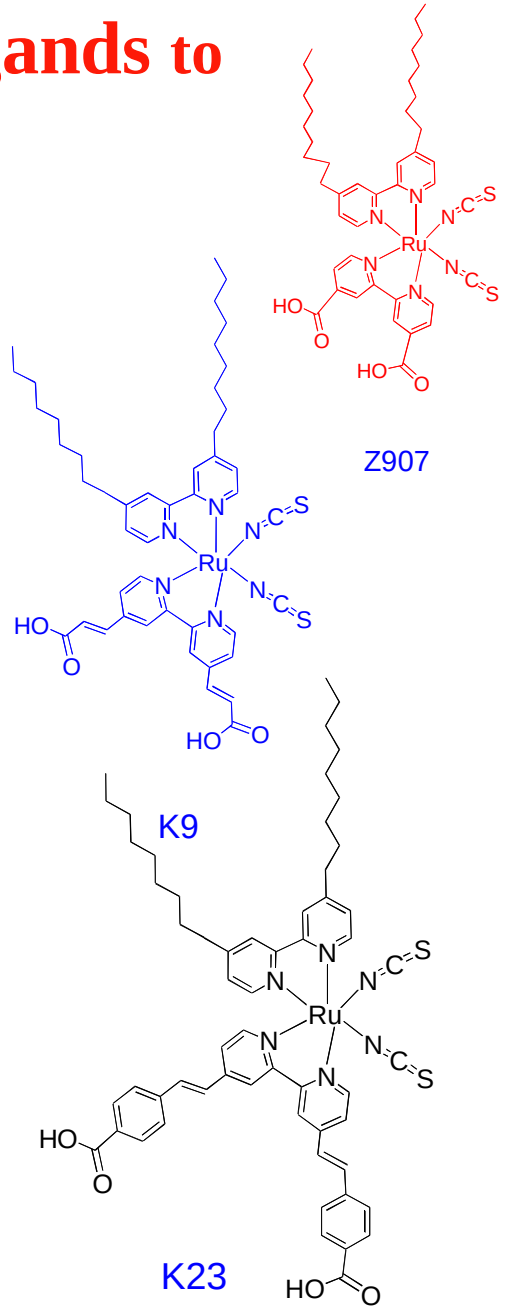
Sensitizers with π-extended acceptor ligands to enhance spectral response



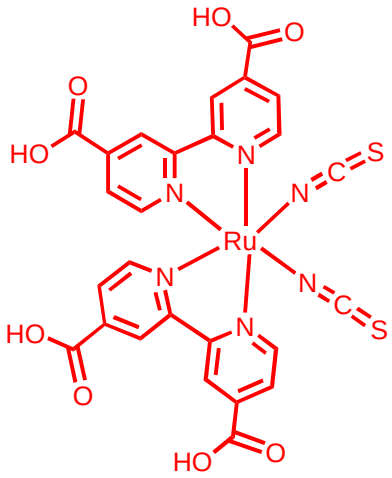


Sensitizers with π-extended acceptor ligands to enhance spectral response





New Sensitizers with extended π-system



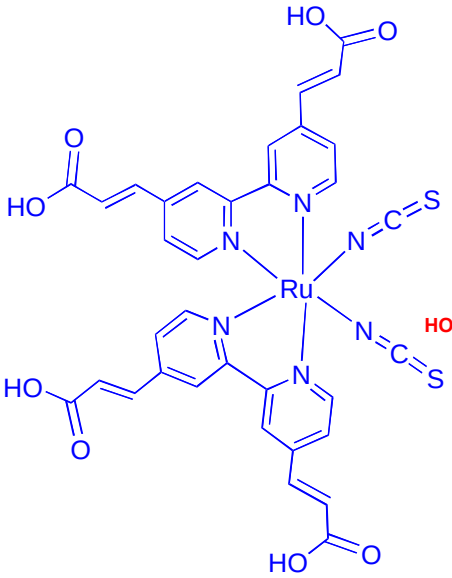
N3

λmax 535 nm

ε= : 13800 $M^{-1}cm^{-1}$

Em. λ_{max} : 780 nm

E_{ox} = 0.85 (irrev)



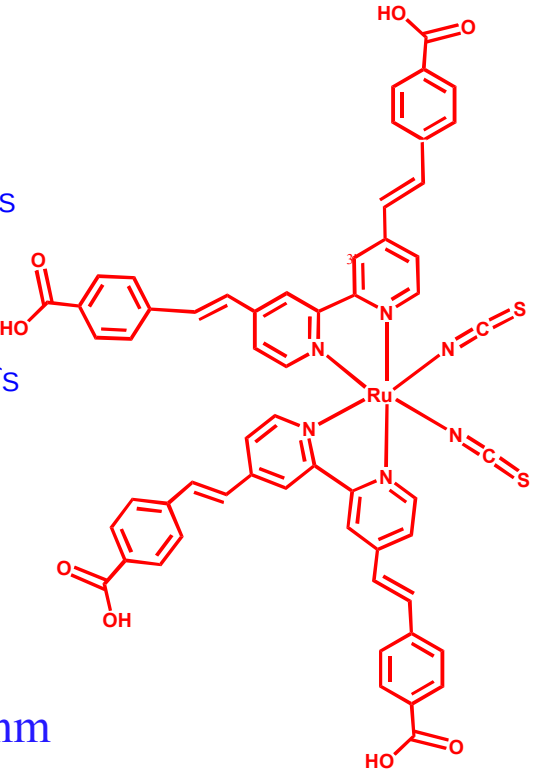
K-8

Abs. λ_{max} : 555 nm

ε: 17600 $M^{-1}cm^{-1}$

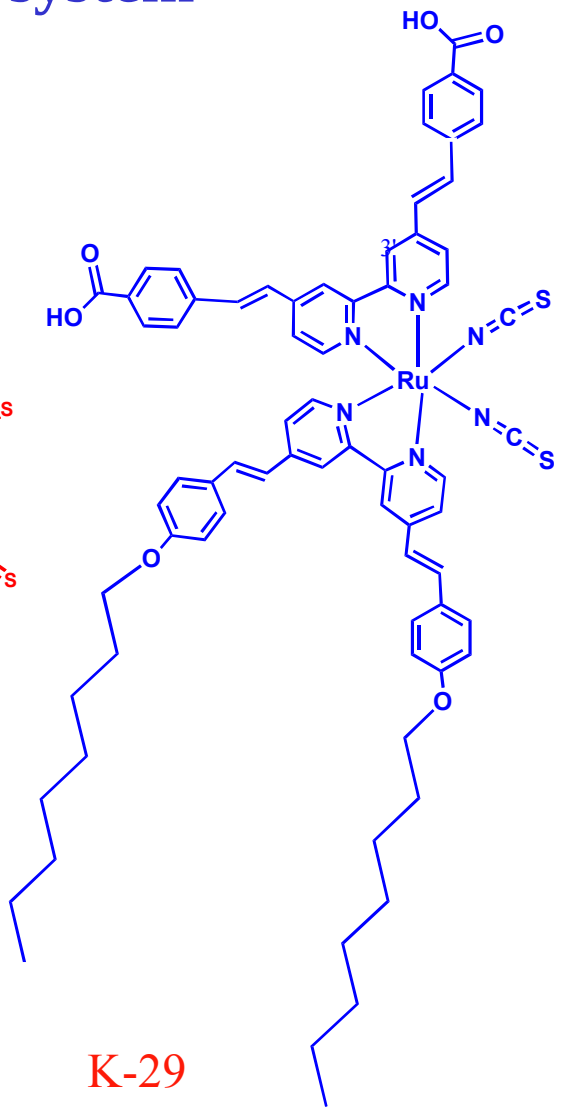
Em. λ_{max} : 840 nm

E_{ox} = 0.77 (rev)



K-27

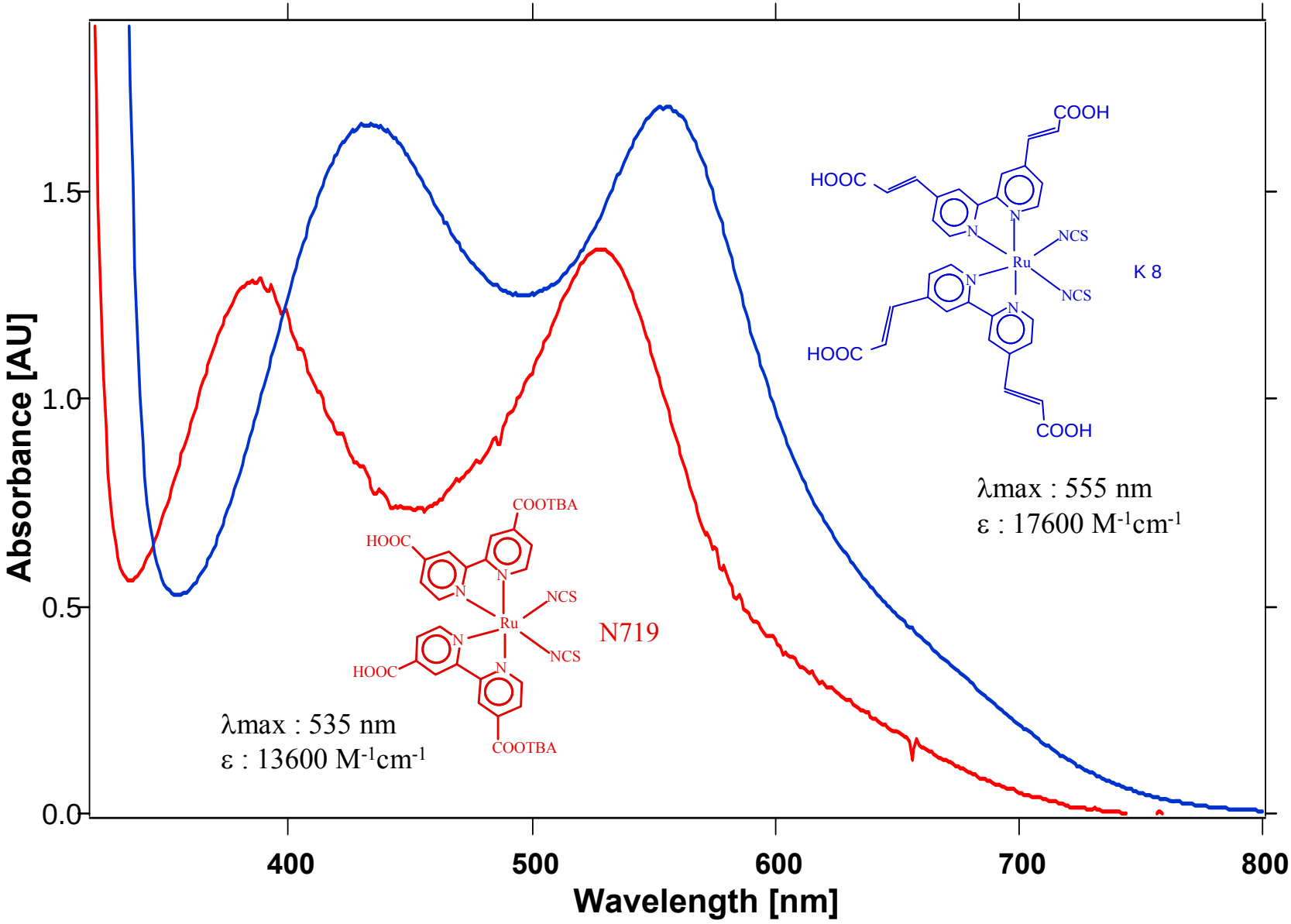
Abs. λ_{max} : 566 nm



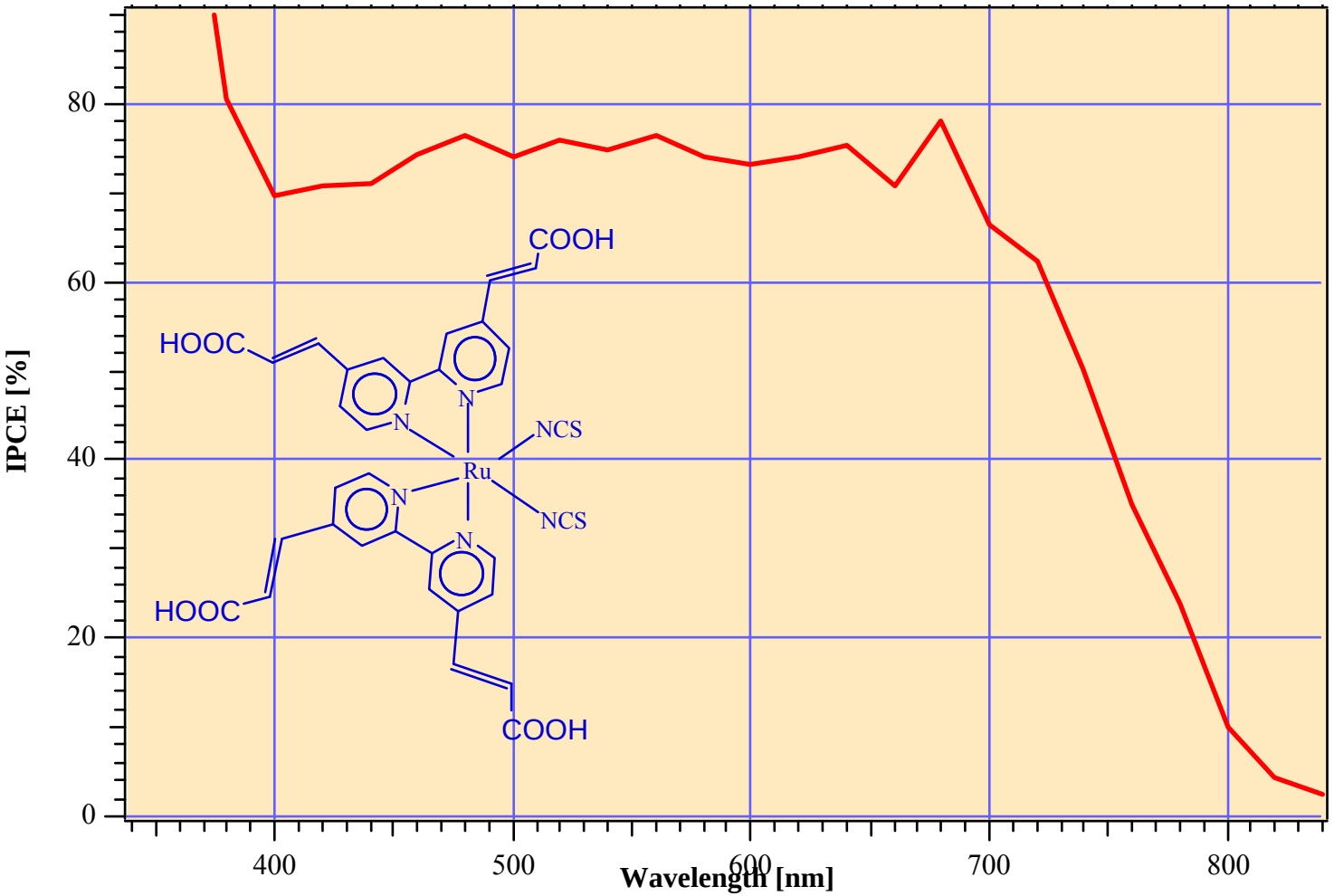
K-29

Abs. λ_{max} : 575 nm

UV/Vis Spectra of N719 and K8 Sensitizers



Incident Photon to Current Conversion Efficiency of K 8 Sensitizer

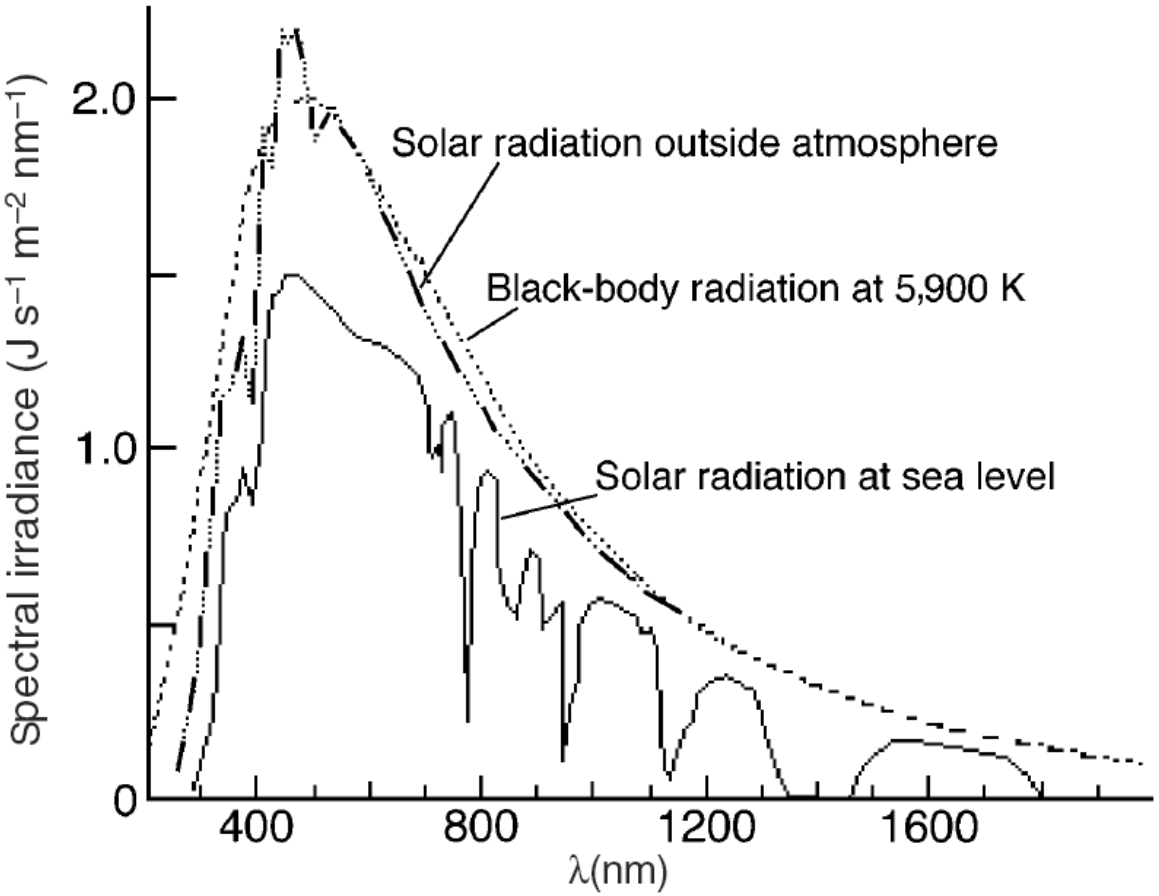


Solar AM 1.5 (1000 W/cm^2) is 18 - 19 mA/cm^2

M. K. Nazeeruddin, C. Klein, P. Liska, M. Graetzel, *Coord. Chem. Rev.* **2005**, 249, 1460

C. Klein, M. K. Nazeeruddin, P. Liska,D. Di Censo, N. Hirata, E. Palomares, J. R. Durrant, M. Graetzel, *Inorg. Chem.* **2005**, 44, 178

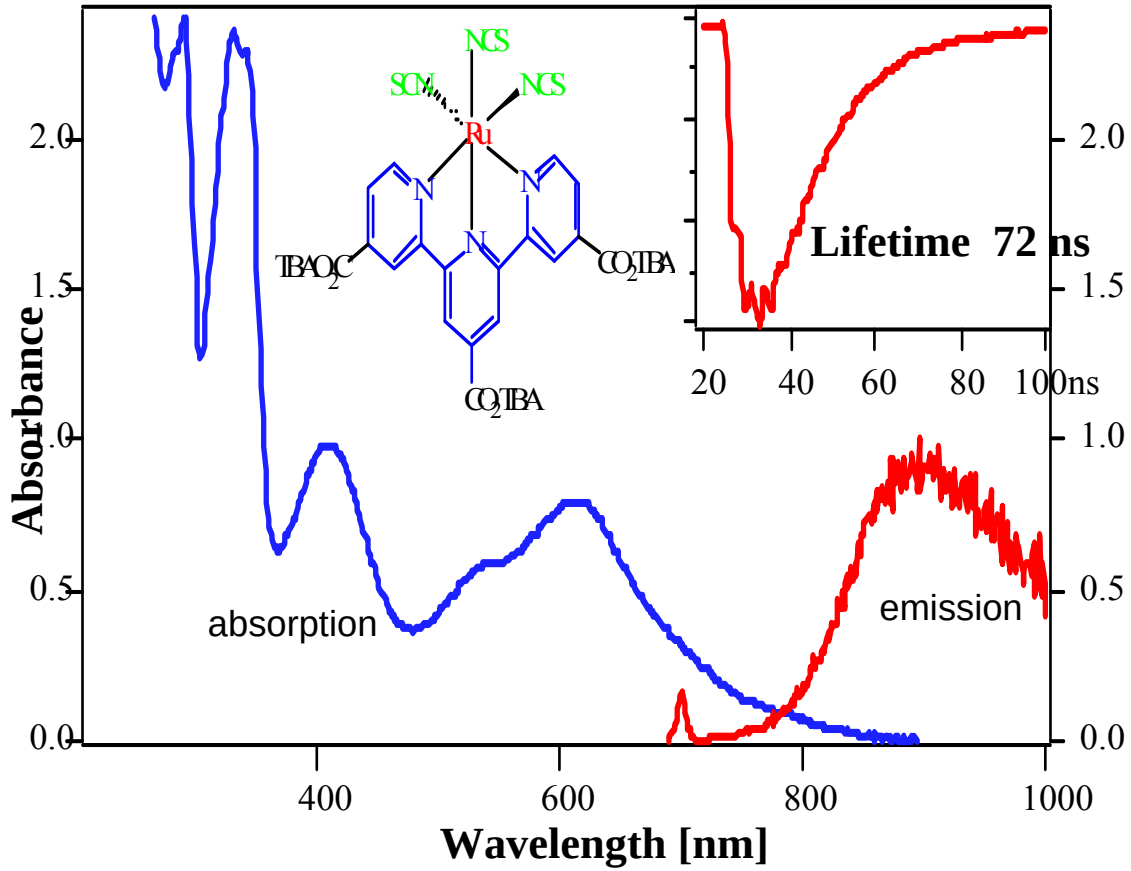
Spectral irradiance of the Sun at mean Earth-Sun separation



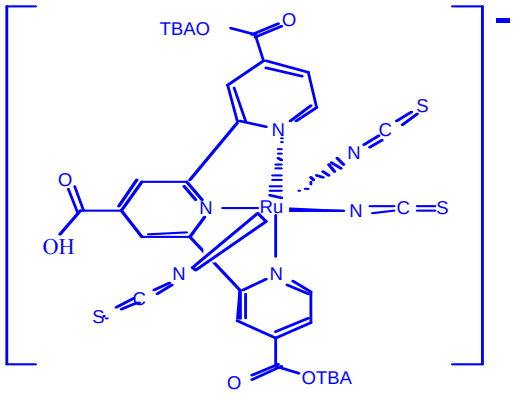
One factor that limiting further improvement of DSC is lack of energy capture by dyes in the IR region.

Half of the sun's energy reaching earth's lies above 700 nm and one third beyond 1000 nm

Absorption and Emission (ex = 700 nm) of Trithiocyanato (4,4',4''-tricarboxy-2,2';6,2''-terpyridine)Ruthenium(II) complex in MeACN



Separation of Linkage Isomers of Trithiocyanato (4,4',4''-tricarboxy-2,2';6,2''-terpyridine)Ruthenium(II)



Isomer 1

pH = 5

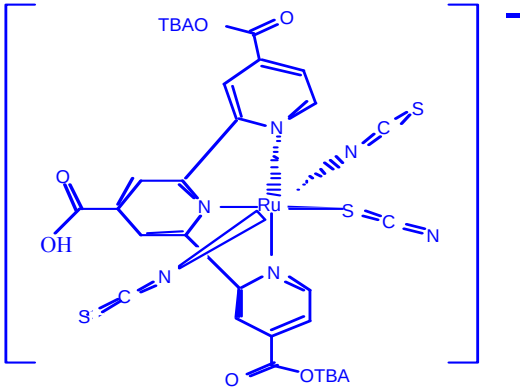
$\{(C_4H_9)_4N\}_2[Ru(H_2tcterpy)(NCS)_3]_2$

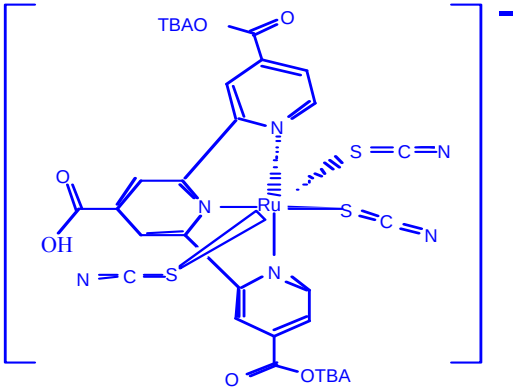
pH = 4.5

pH = 3.8

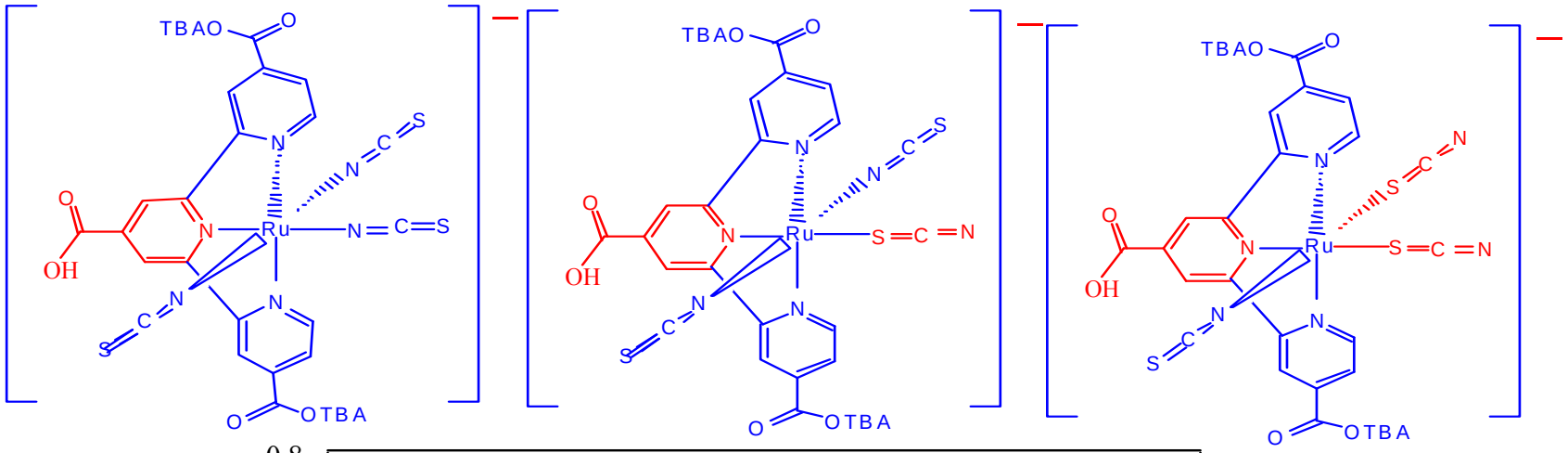
Isomer 2

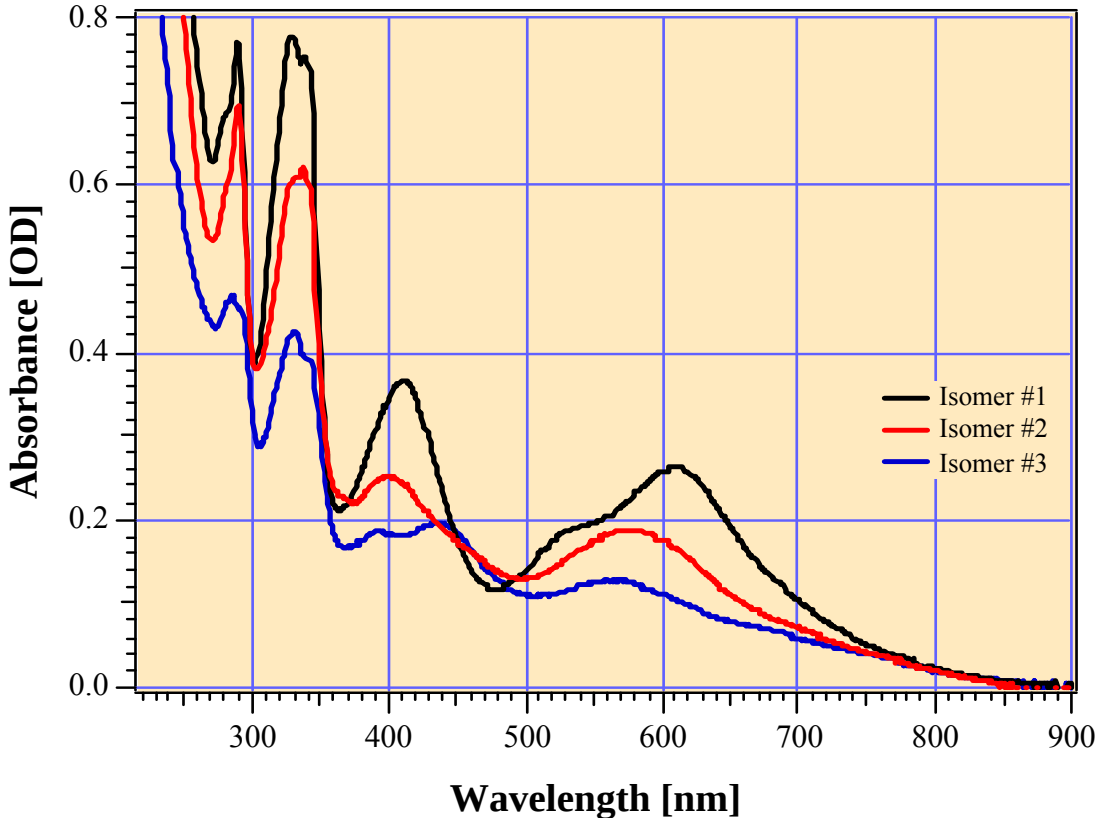
Isomer 3



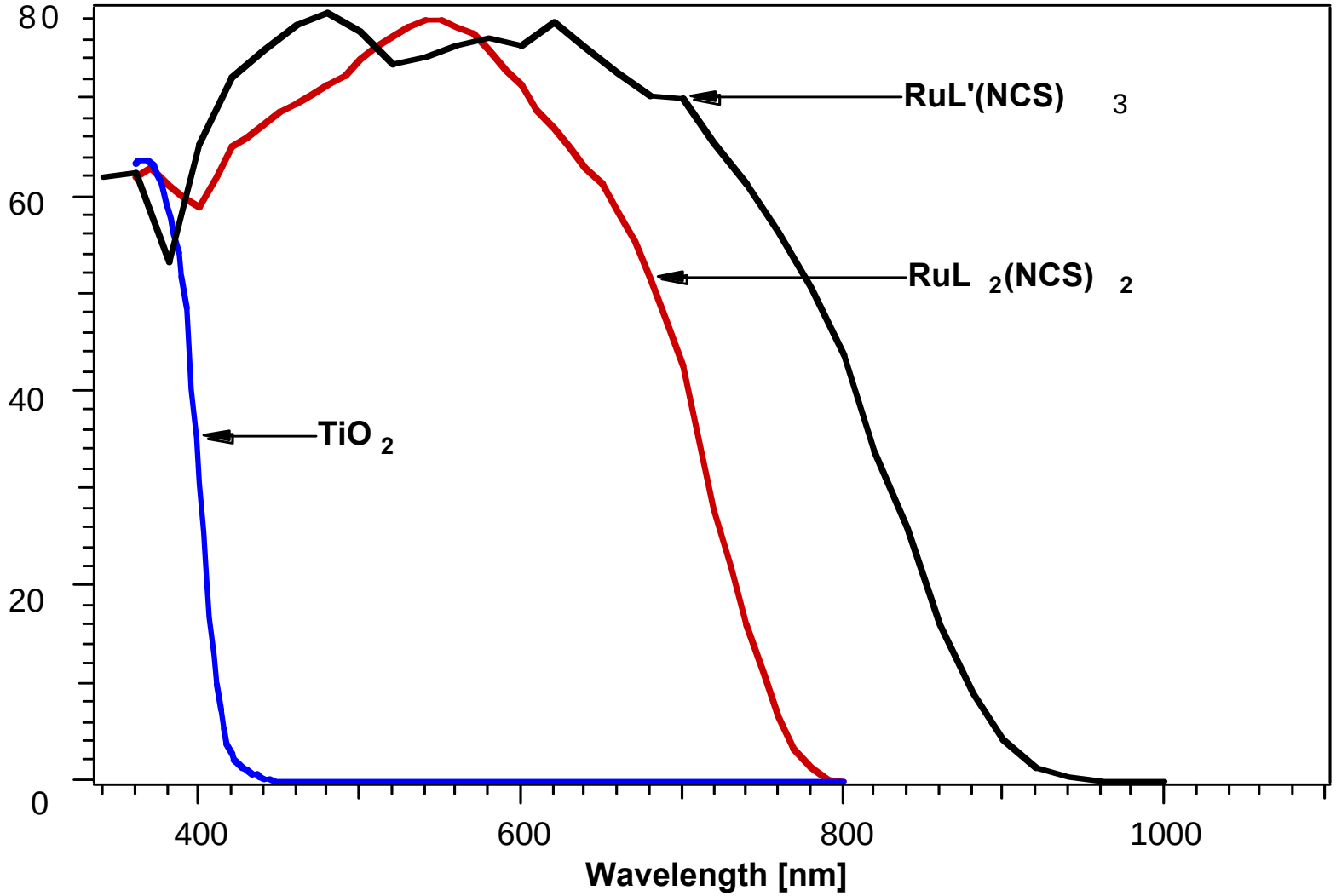


Isomers of the Black dye





Photocurrent action spectrum of different ruthenium complexes attached to nanocrystalline TiO_2 films

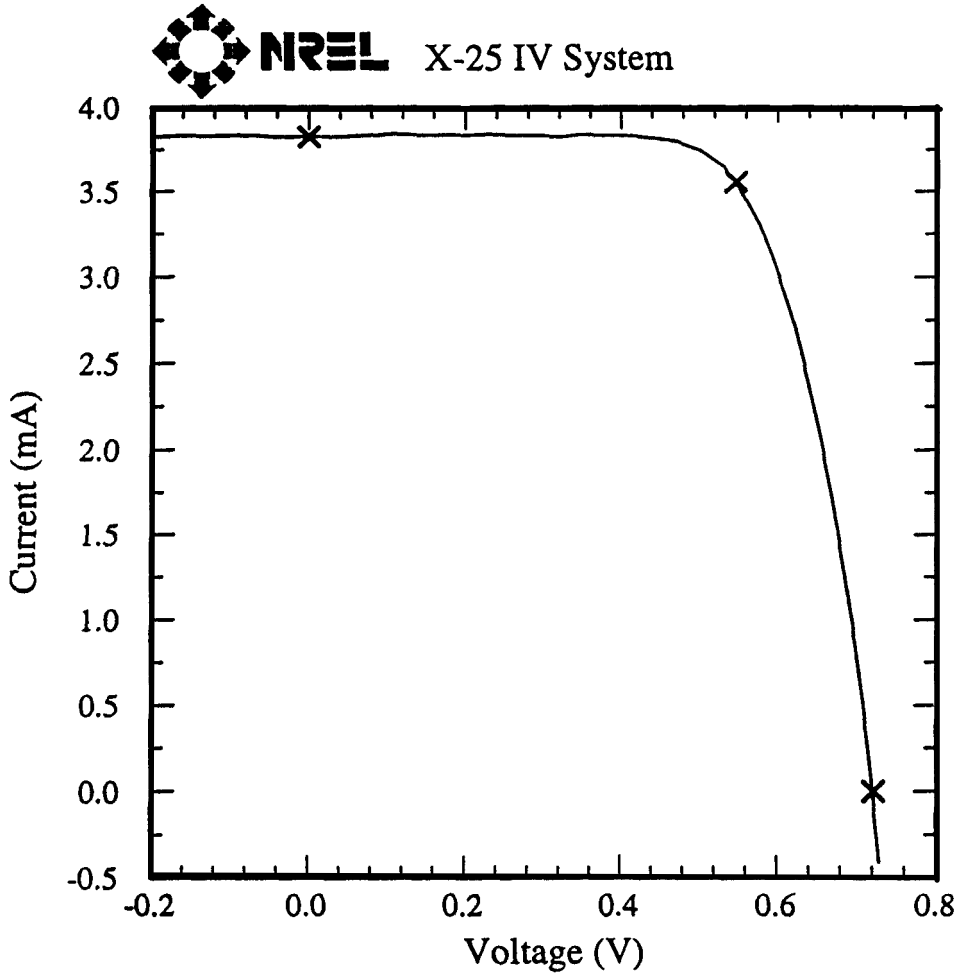


L = 4,4'-COOH-2,2'-bipyridine

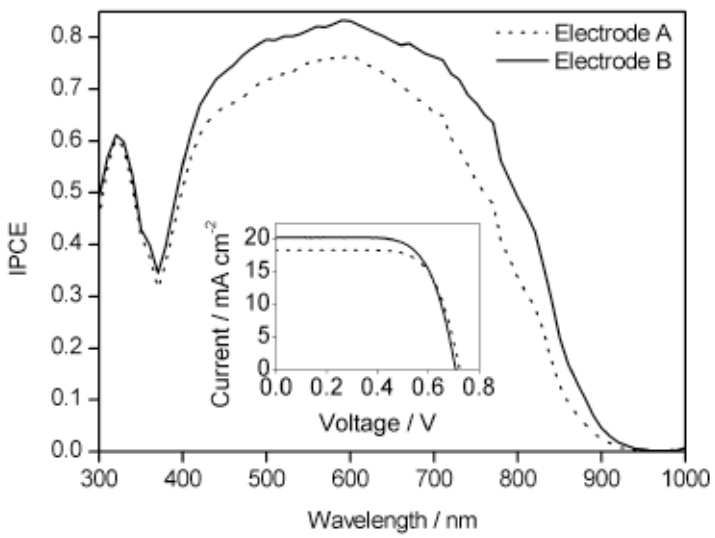
L' = 4,4',4''-COOH-2,2':6',2''-terpyridine

JACS, 2001, 123, 1613

Photocurrent-voltage characteristics of a nanocrystalline photoelectrochemical cell sensitized with the Black Dye



The results were obtained at the NREL calibration laboratory measured with an area of 0.1863 cm2 and irradiance of 1000 Wm-2. V_{oc} = 0.72 V, J_{sc} = 20.53 mAcm-2; fill factor = 70.41%; the efficiency = 10.4.



Data from Prof. Arakawa's lab:

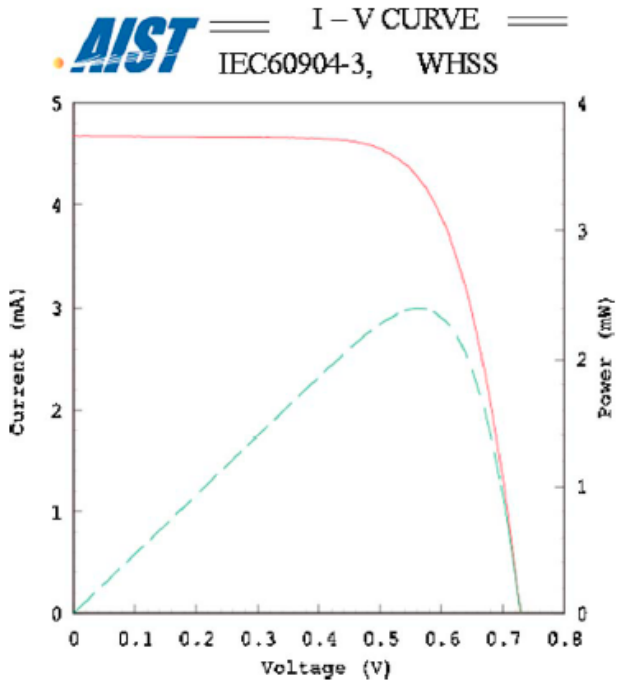
Irradiance of 1000 Wm-2.

V_{oc} = 700 mV

J_{sc} = 21.49 mAcm-2

Fill factor = 69.9%

Eficiency = 10.5.



Data from Sharp:

Irradiance of 1000 Wm-2.

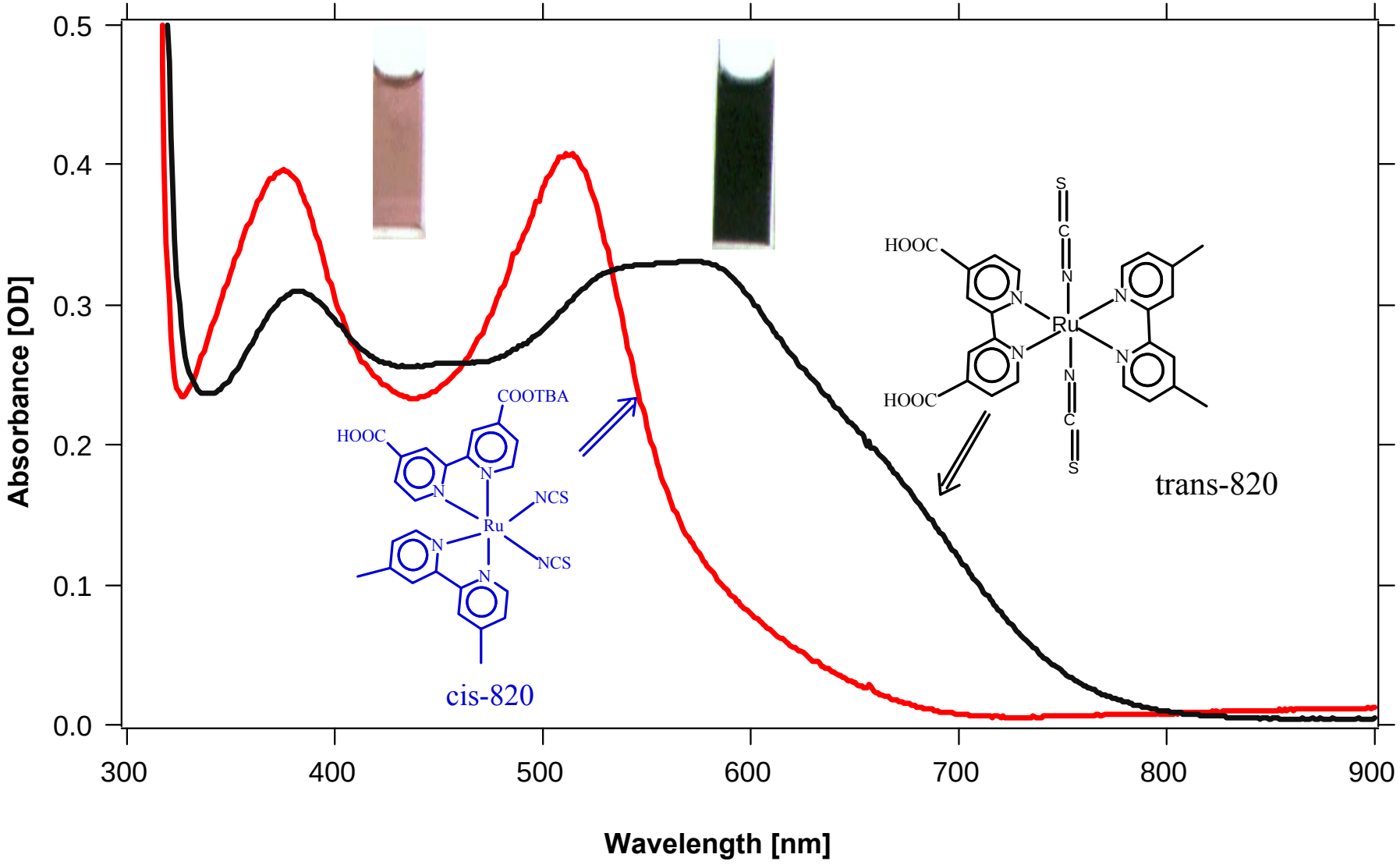
V_{oc} = 729 mV

J_{sc} = 21.0 mAcm-2

Fill factor = 70.4%

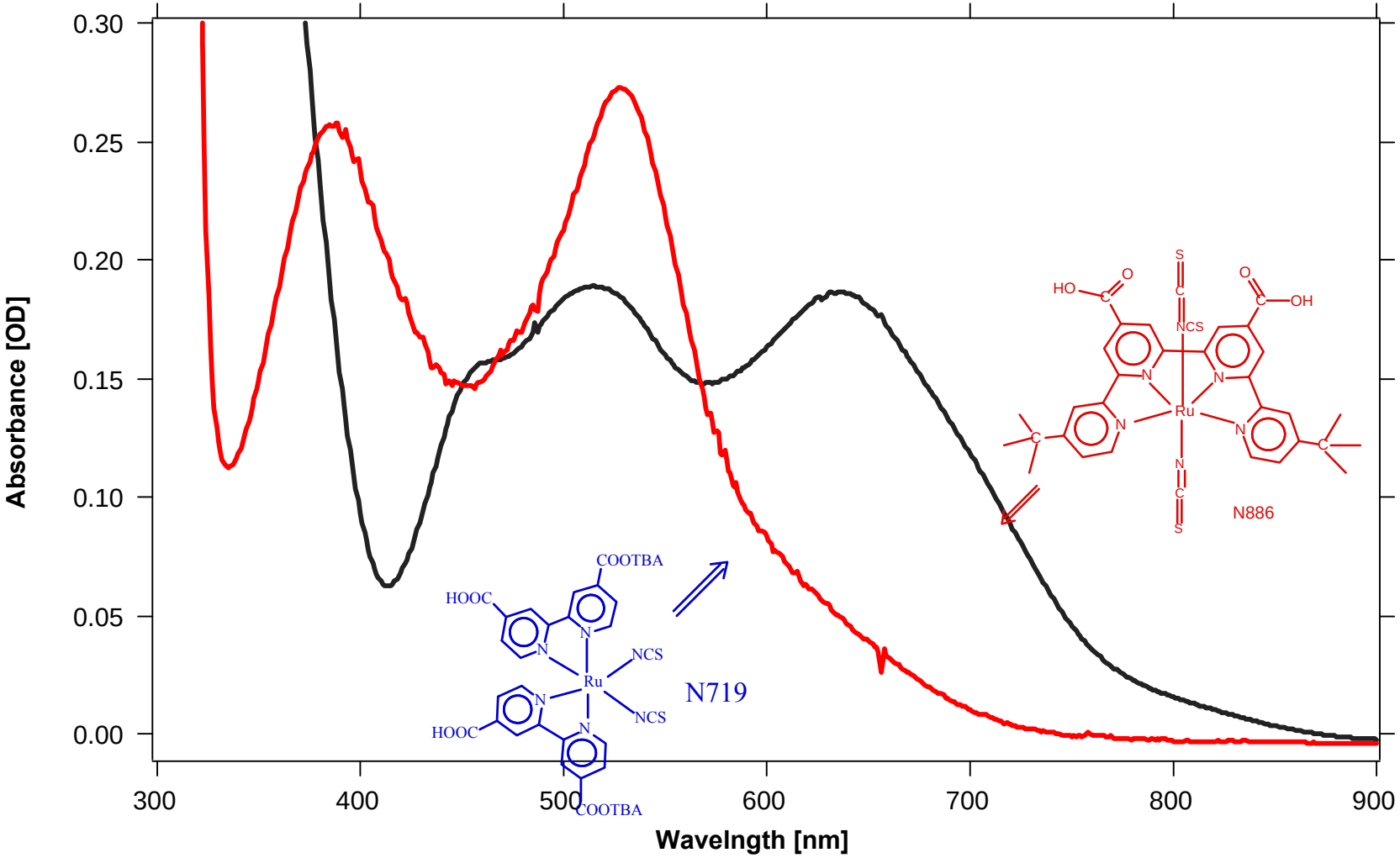
Eficiency = 10.8 **(11.2)**

Cis- and trans-Isomers of N820



Md. K. Nazeeruddin et al. *Coord. Chem. Rev.* 208 213 (2000)

Comparison of UV/Vis Absorption Spectra of N719 and trans-[Ru(L)(NCS)$_2$]

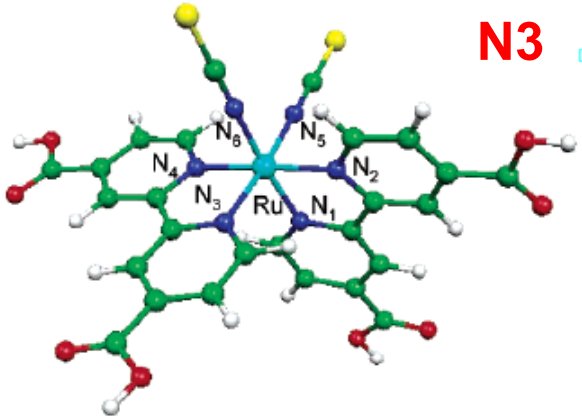


C. Barolo, Md. K. Nazeeruddin, Simona Fantacci, D. Di Censo, P. Comte, P. Liska, G. Viscardi, P. Quagliotto,Filippo De Angelis, S. Ito, and M. Graetzel *Inorg. Chem.* **2006**, 45, 4642-4653

Tuning the properties of Ru(II) sensitizers

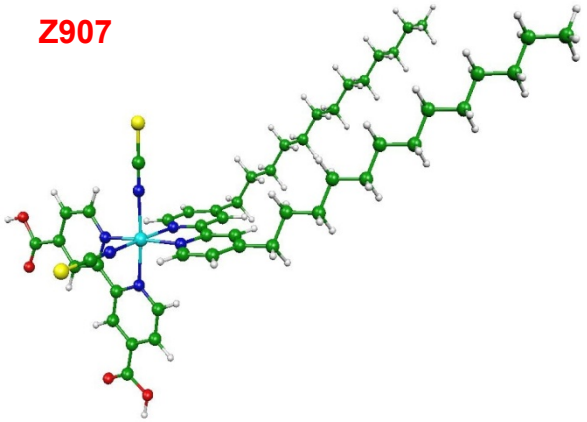
Effect of deprotonation

N3

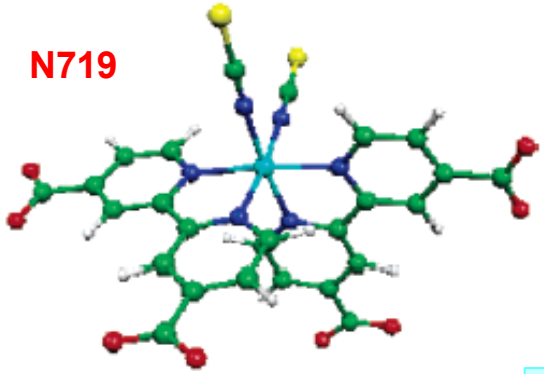


Bypyridine functionalization

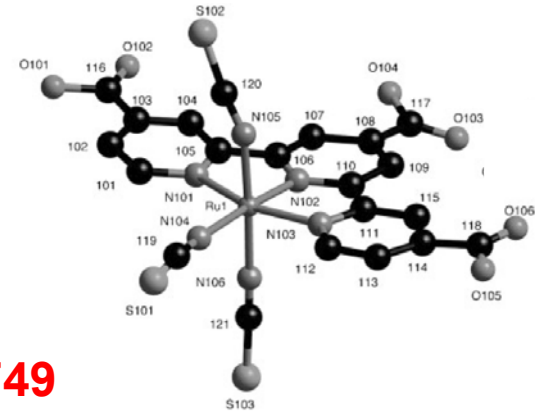
Z907



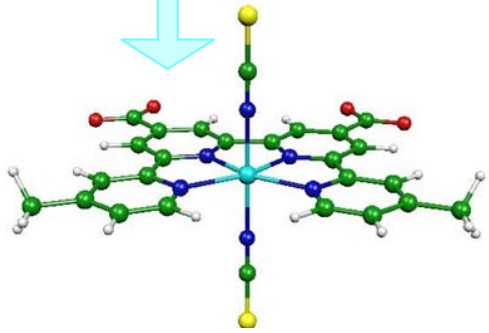
N719



Ligand engineering

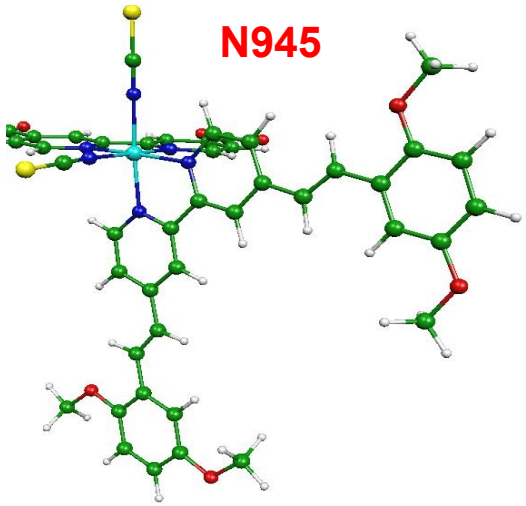


749



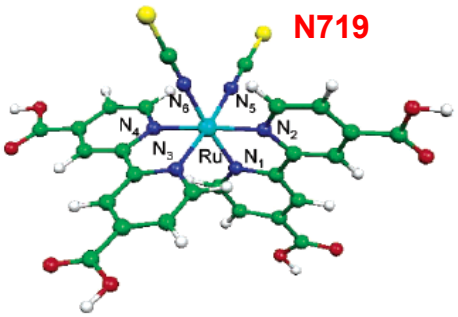
N866

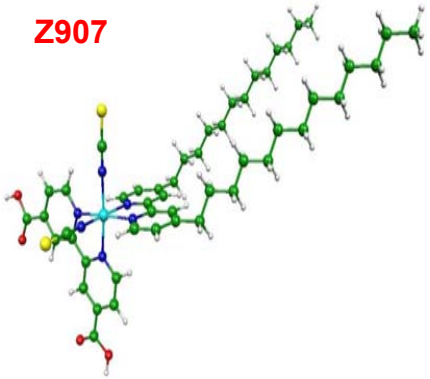
N945

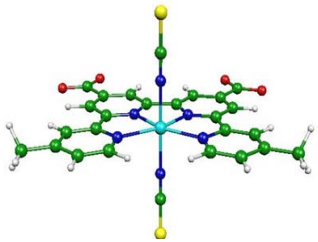


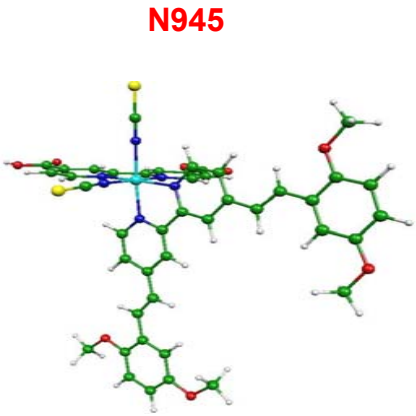
Mater. Chem. 2006, *16*, 4468.
J. Photochem. Photobiol. A 2007, 1*85*, 331.
Inorg. Chem. 2006,45,4642

Heteroleptic Ru(II) TiO_2 sensitizers







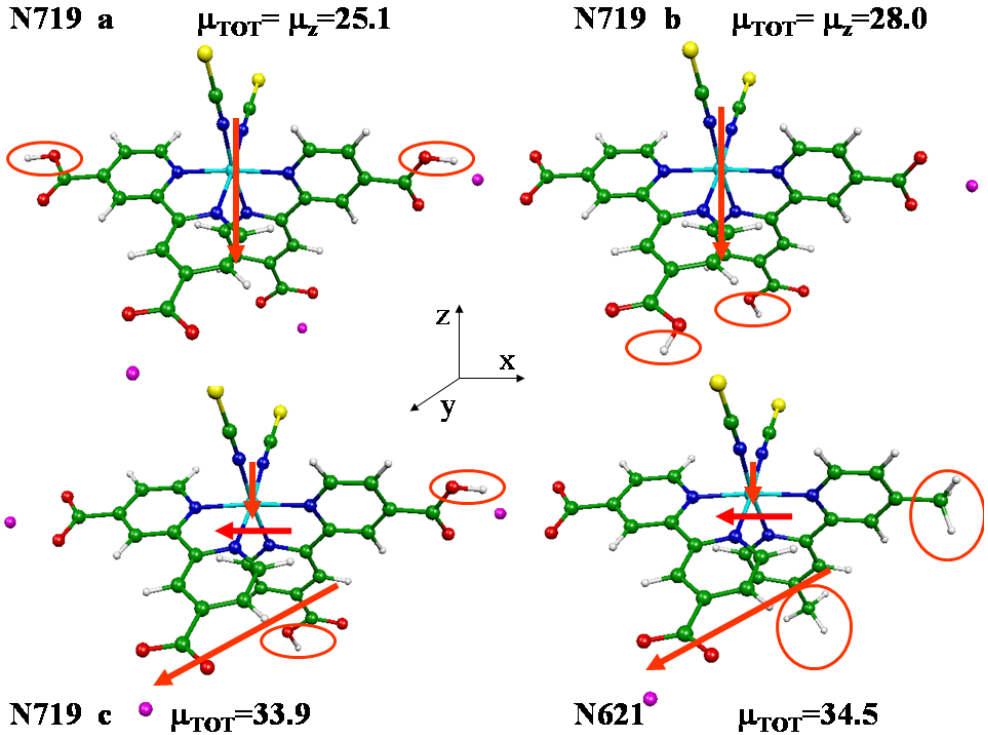


Sensitizer	Number of protons	Current mA/cm^2	Potential (mV)	Fill Factor	Efficiency at 1.5 AM
N719	2	16.66	846	0.73	10.28
N621	1	16.22	766	0.70	8.69
K19	1	16.40	768	0.73	9.19
N945	1	17.25	759	0.73	9.55
N886	1	11.80	700	0.73	6.02
N749	1	21.00	729	0.72	10.80

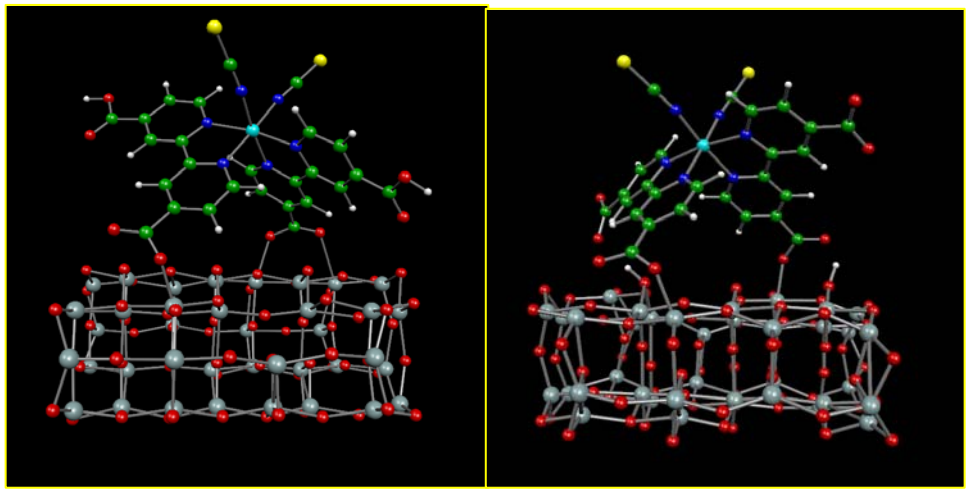
A considerable reduction of the open circuit potential (ca. 100 mV) is observed with heteroleptic and black sensitizers

J. AM. CHEM. SOC., 129, 14156, 2007
Nano Lett. 3189, 7, 2007.

Homoleptic/Heteroleptic comparison



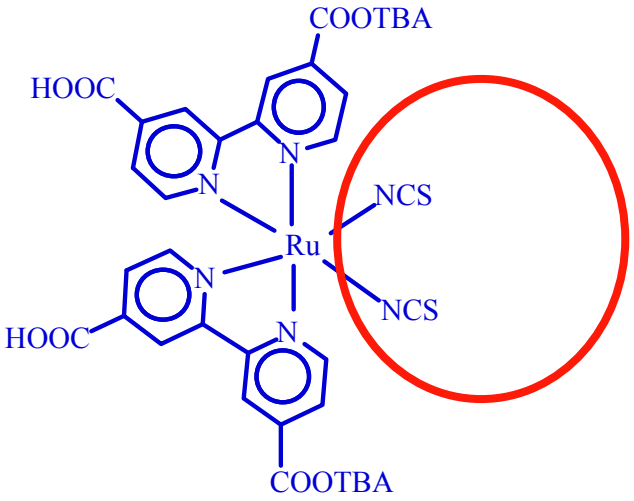
Homoleptic (N719 c) and heteroleptic (N621) sensitizers, adsorbed via a single bipyridine ligand, exert strong dipolar fields at the TiO_2 surface, causing a down-shift of the TiO2 conduction band.



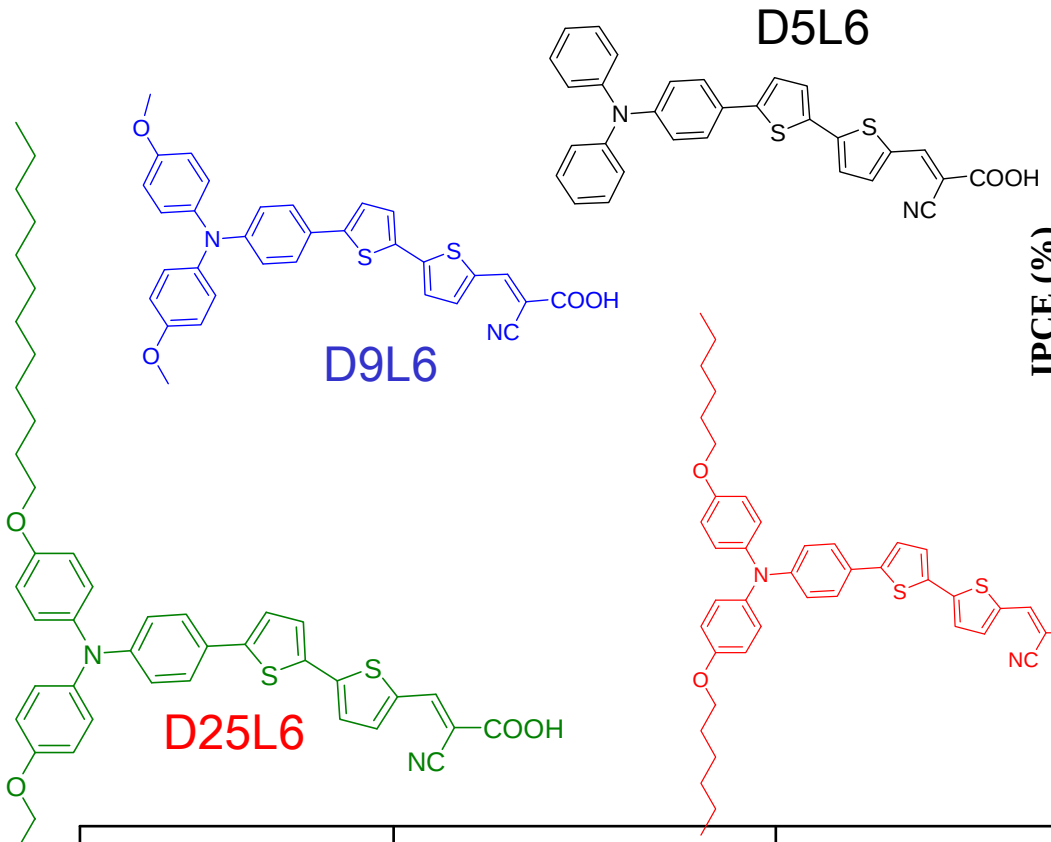
Two prototypical configurations of N719/TiO_2

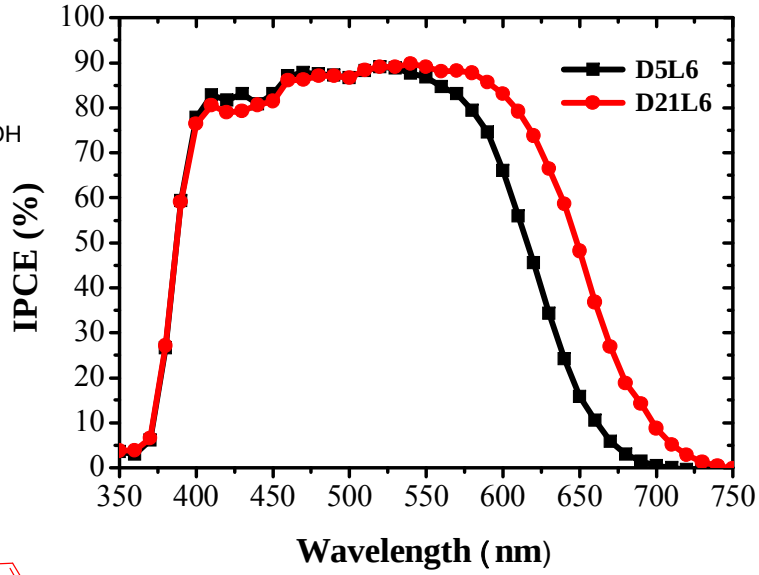
The two protons are located on the dye (left) or on the TiO_2 (trans to NCS)

Sensitizers with new **donor ligands**



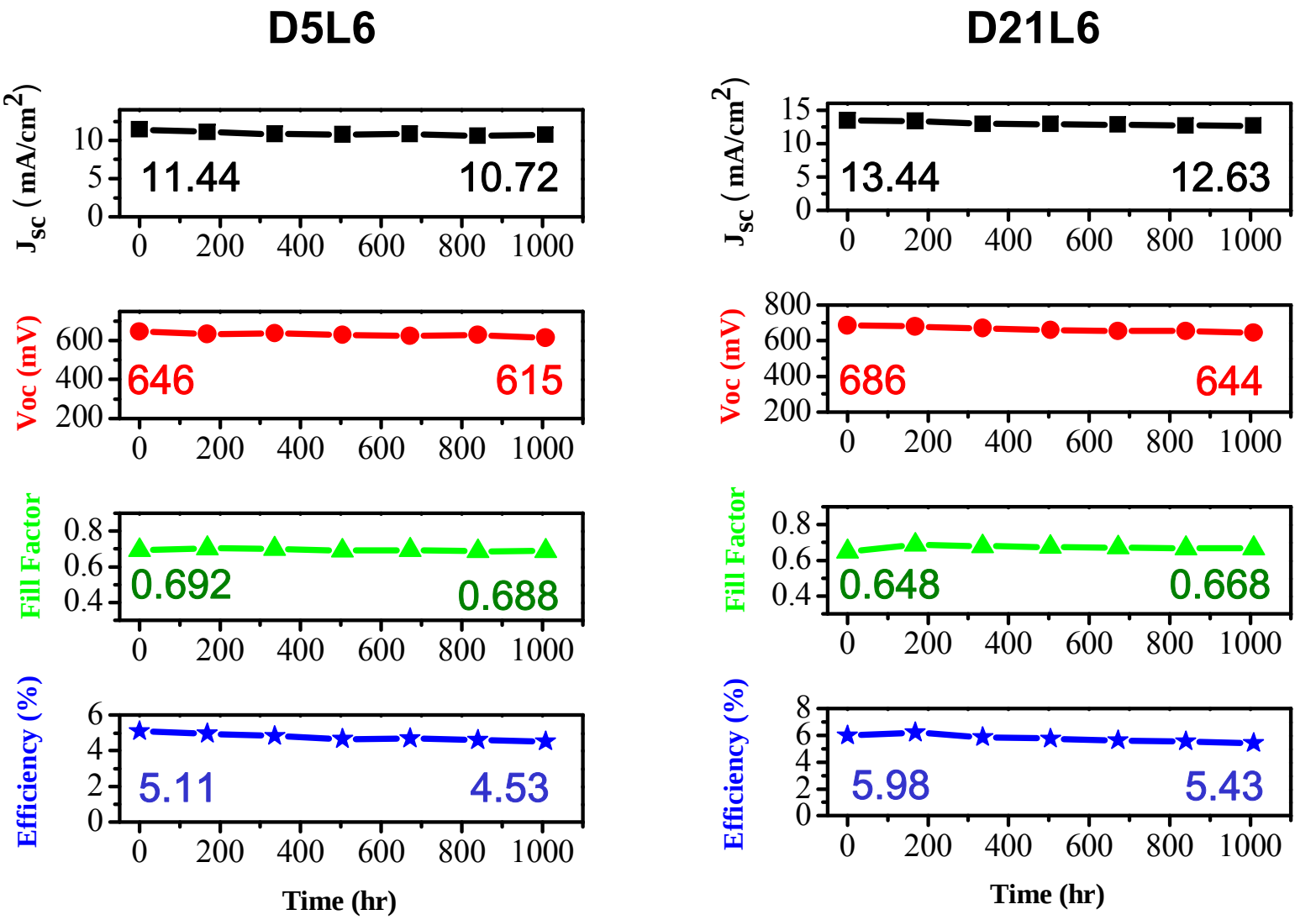
Organic Dyes



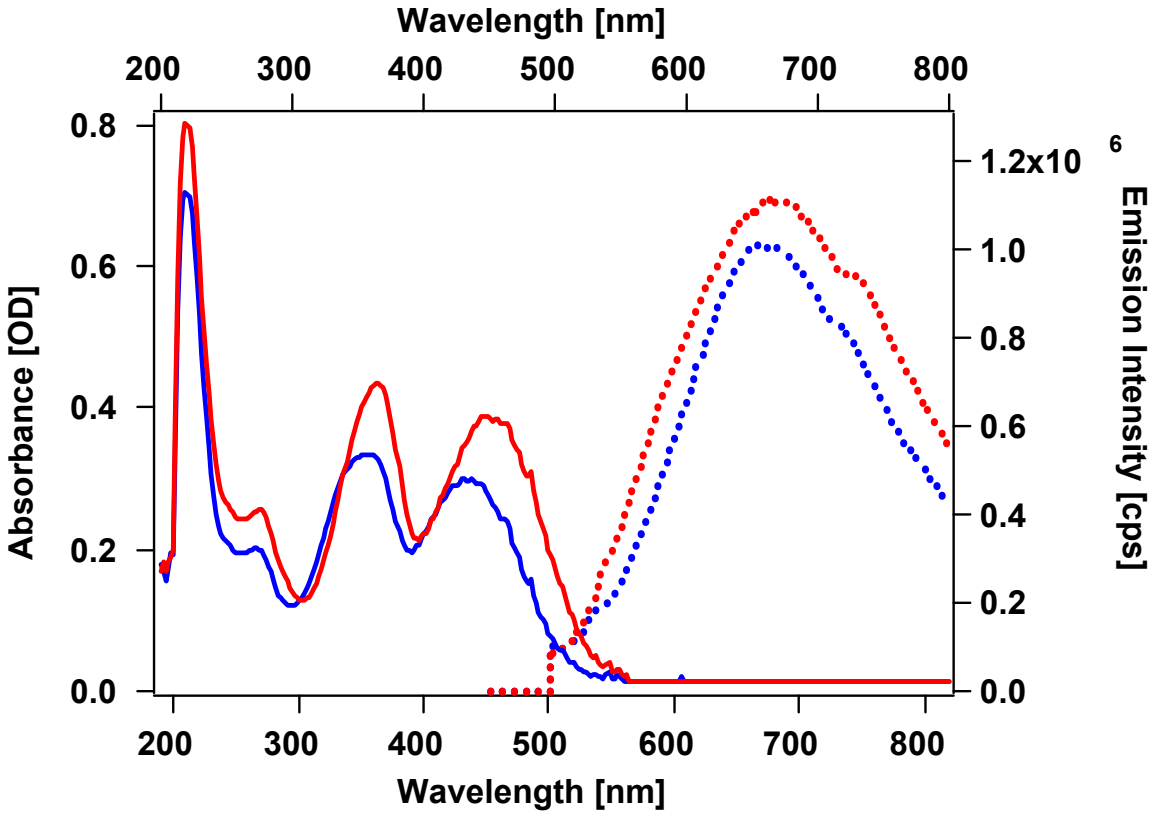


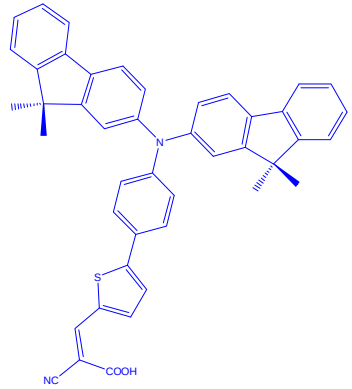
Dye	J (mA/cm2)	V (mV)	FF	efficiency
D5L6	12.50	685	0.74	6.29
D9L6	13.36	641	0.70	6.08
D21L6	13.70	733	0.72	7.25
D25L6	13.73	745	0.69	7.05

Stability with IL z655 under light soaking + 60 degree

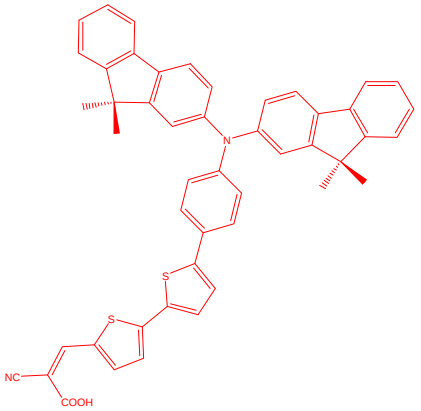


Absorption spectrum



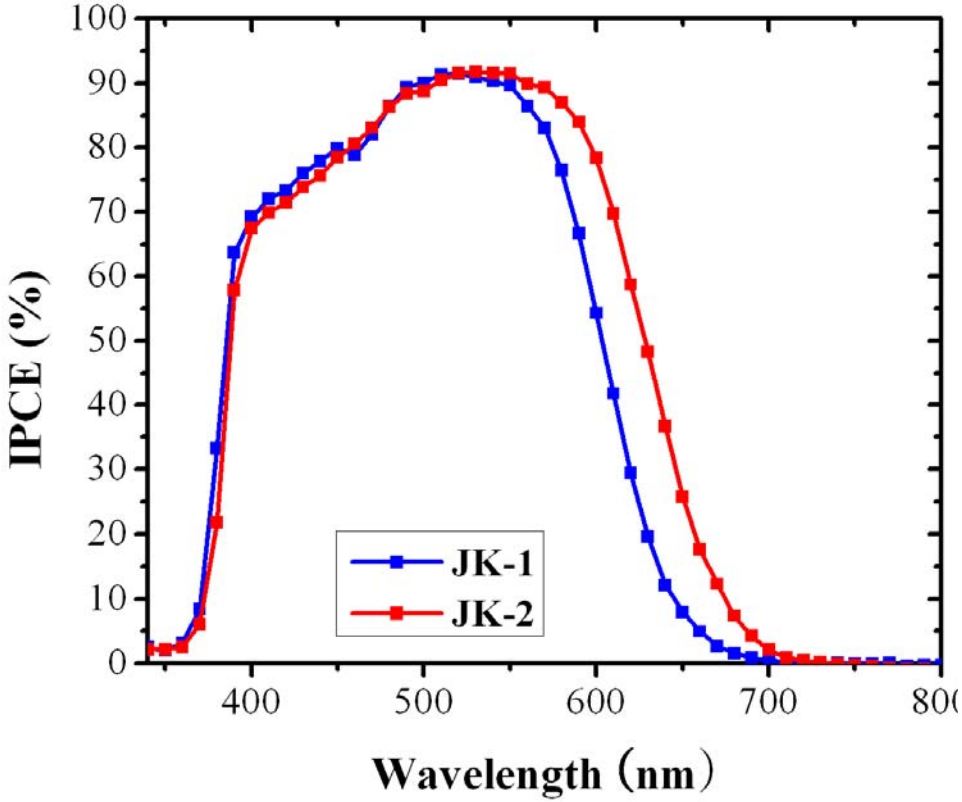


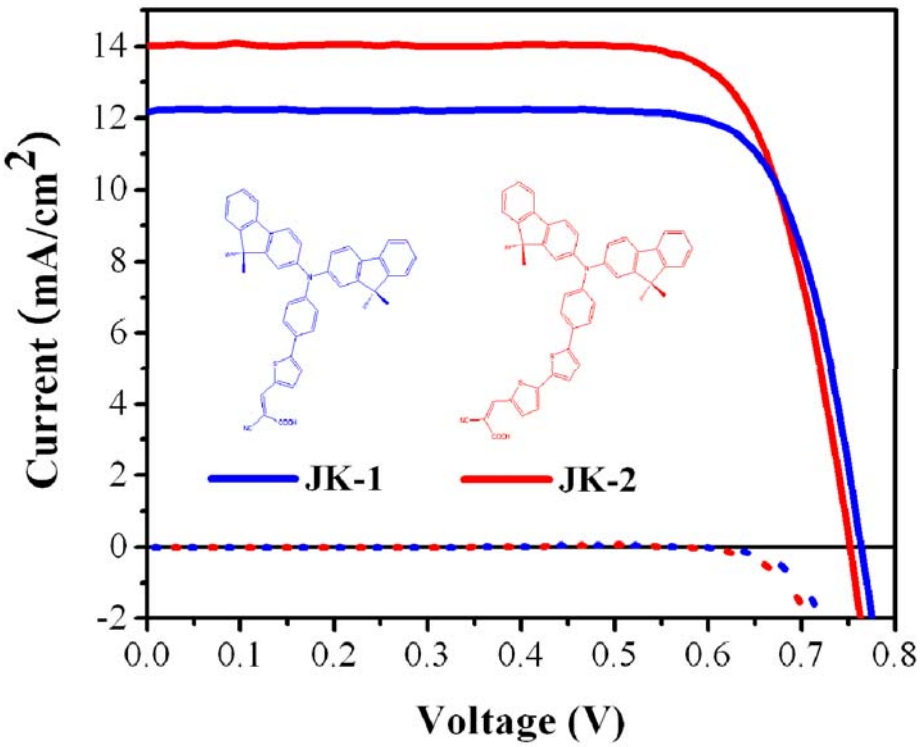
354 nm (ε = **34,000** $dm^3mol^{-1}cm^{-1}$)
436 nm (ε = **30,000** $dm^3mol^{-1}cm^{-1}$)



364 nm (ε = **44,000** $dm^3mol^{-1}cm^{-1}$)
452 nm (ε = **39,000** $dm^3mol^{-1}cm^{-1}$)

IPCE and IV data of JK1 and JK2 dyes

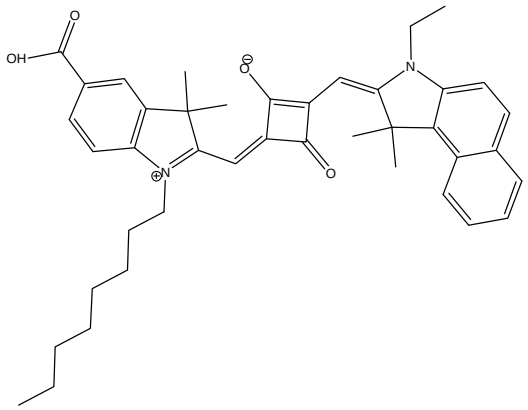


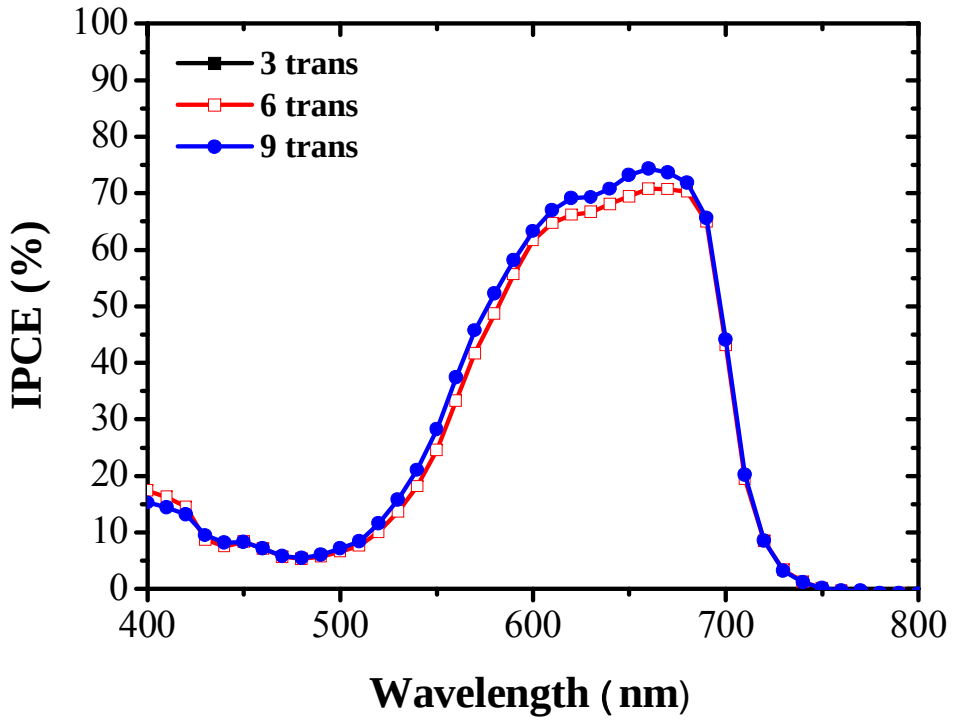


91 % IPCE

power conversion 7.20% (JK-1)

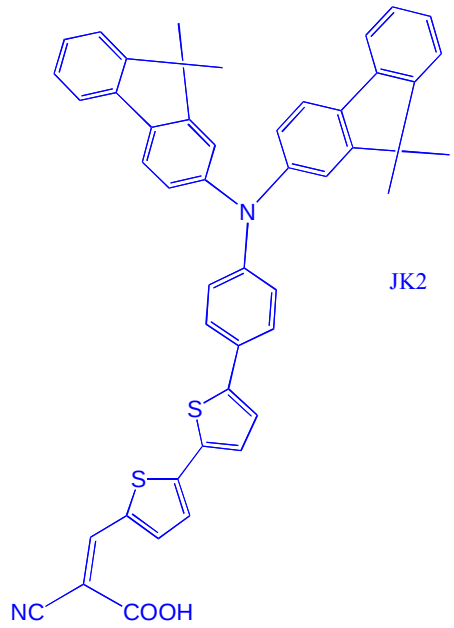
and 8.01% (JK-2)

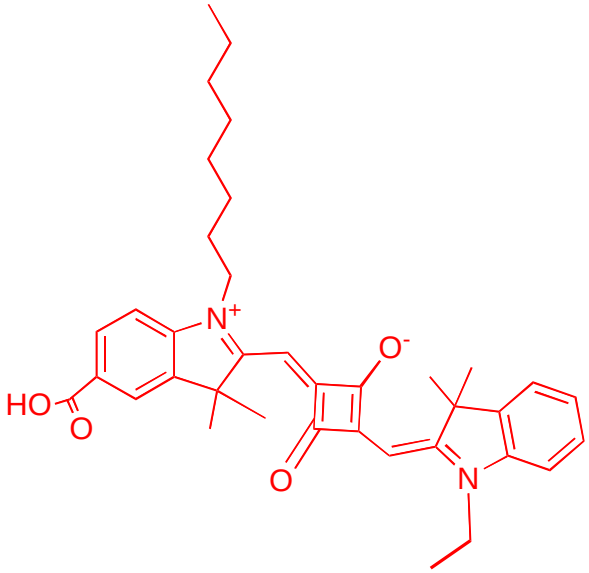


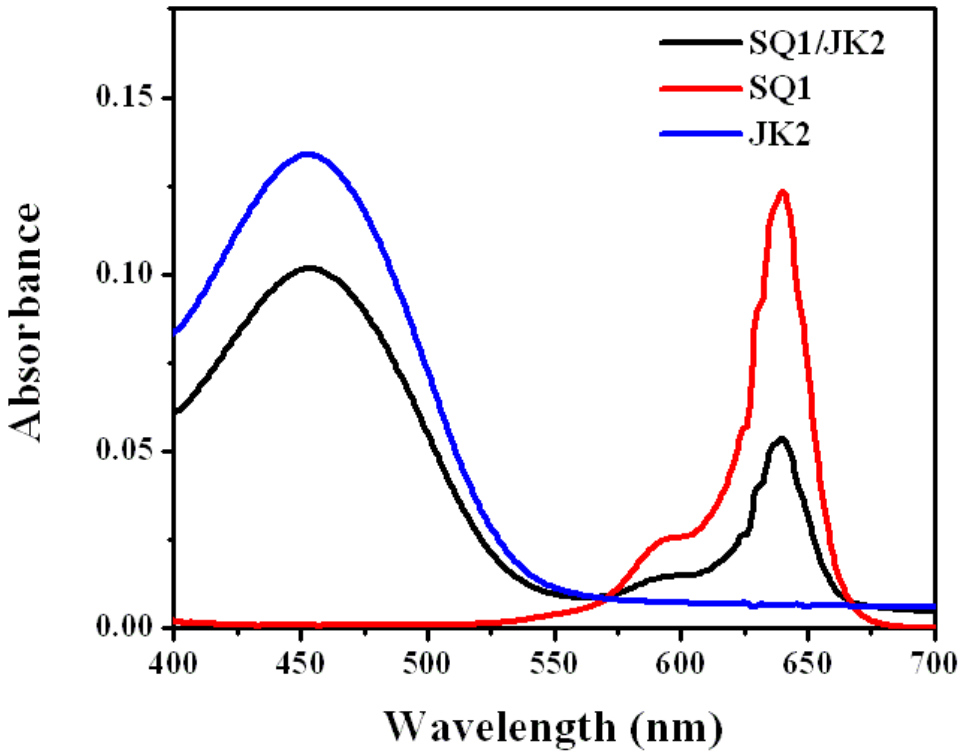


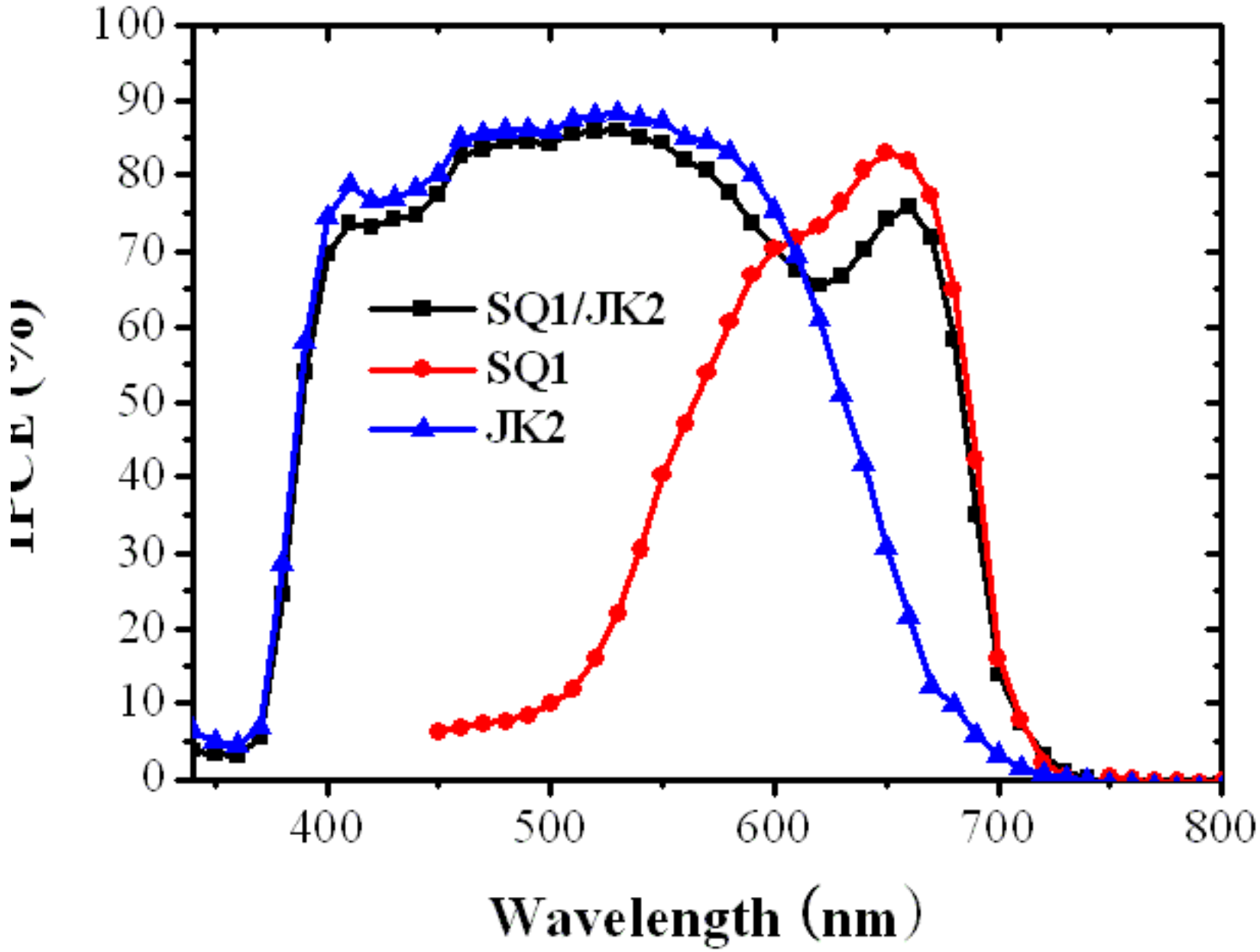
Thickness effect

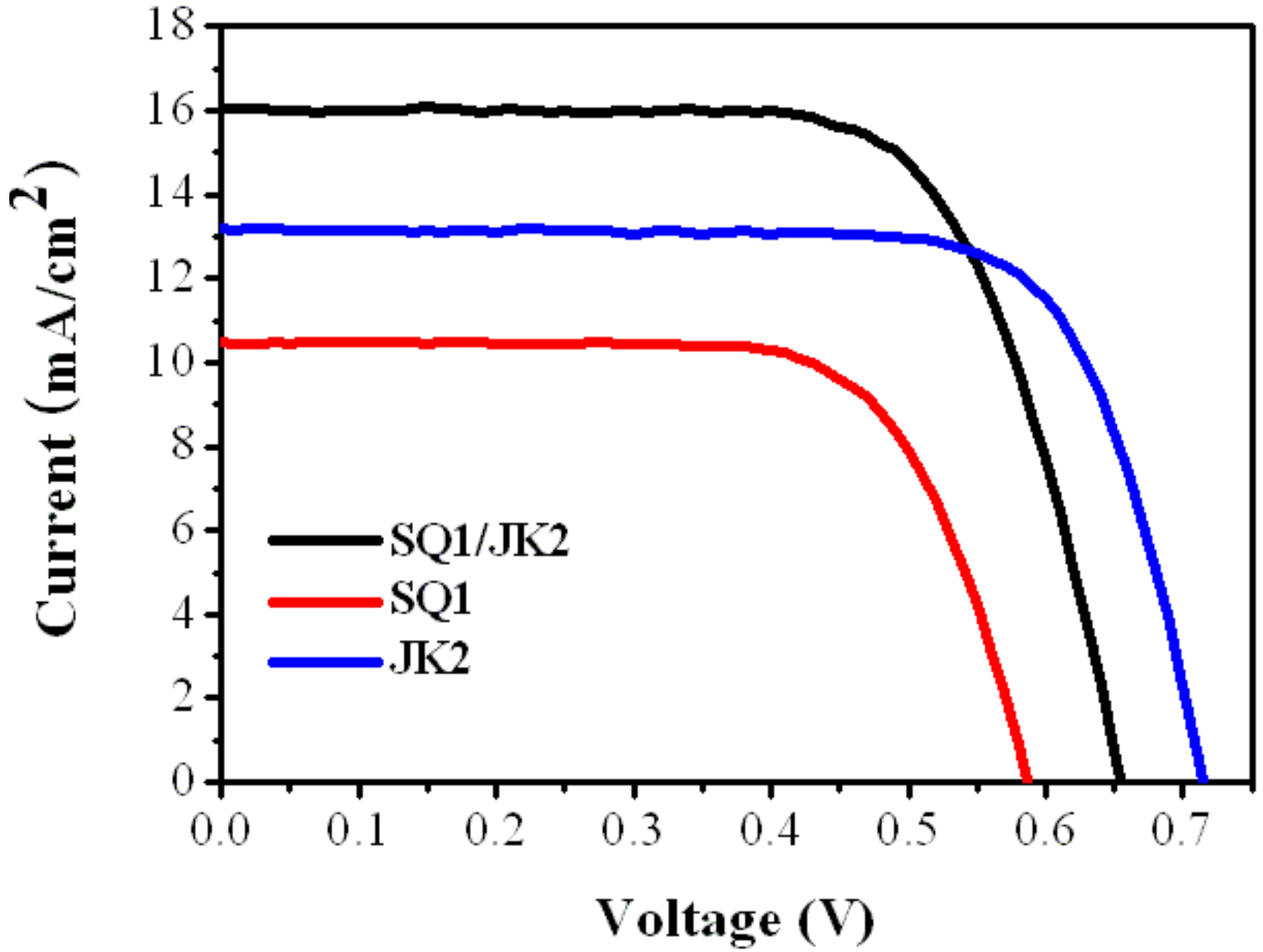
Film (μm)	J (mA/cm2)	V (mV)	FF	efficiency
6 trans	9.31	645	0.661	4.00
9 trans	10.50	621	0.654	4.28
9 +5	11.4	667	0.72	5.40


JK2
N
S
S
NC
COOH

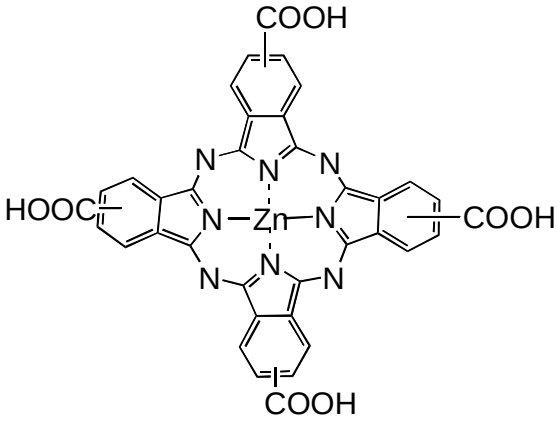

HO
O
N+
O-
O
N

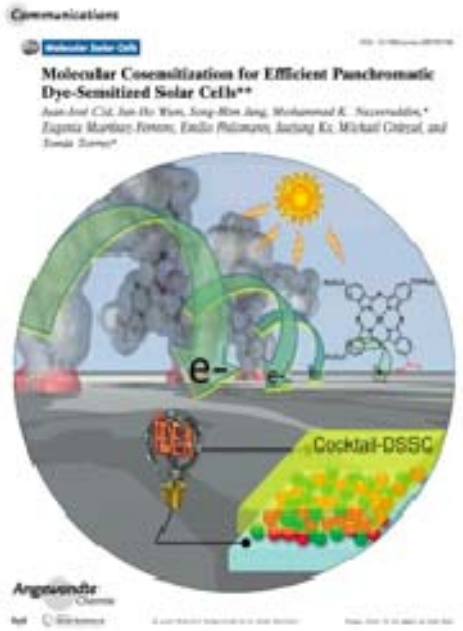

SQ1/JK2
SQ1
JK2
Absorbance
0.15
0.10
0.05
0.00
400
450
500
550
600
650
700
Wavelength (nm)

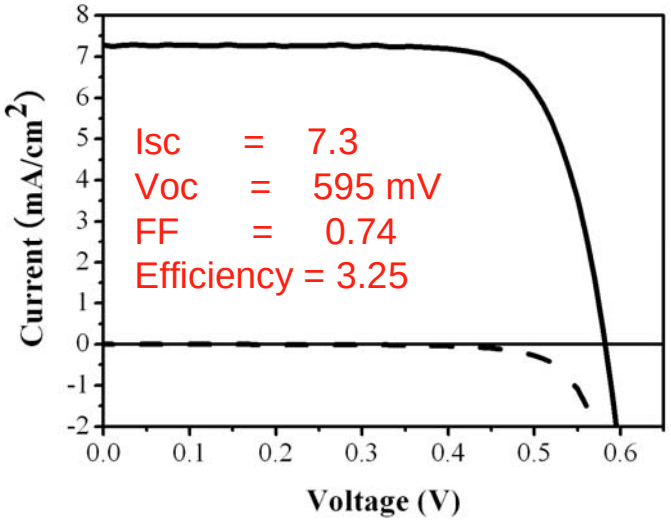

IPCE (%)
100
90
80
70
60
50
40
30
20
10
0
400
500
600
700
800
Wavelength (nm)
SQ1/JK2
SQ1
JK2

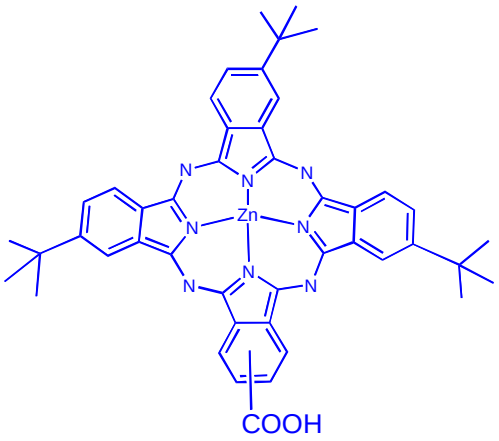

Current (mA/cm^2)
18
16
14
12
10
8
6
4
2
0
0.0
0.1
0.2
0.3
0.4
0.5
0.6
0.7
Voltage (V)
SQ1/JK2
SQ1
JK2

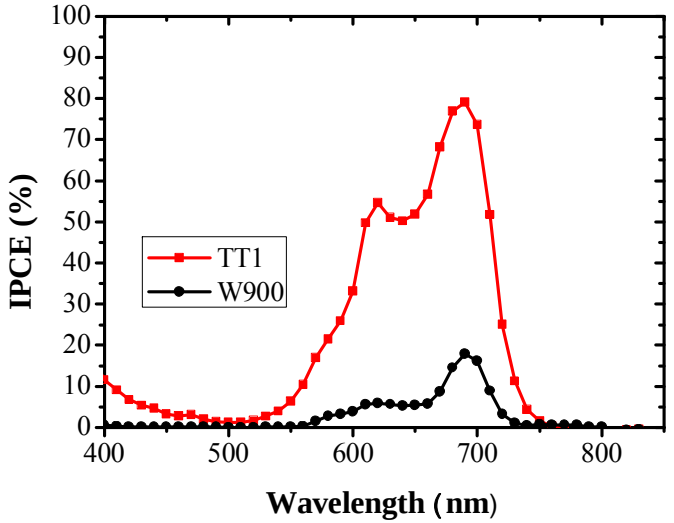
Zinc Phthalocyanine

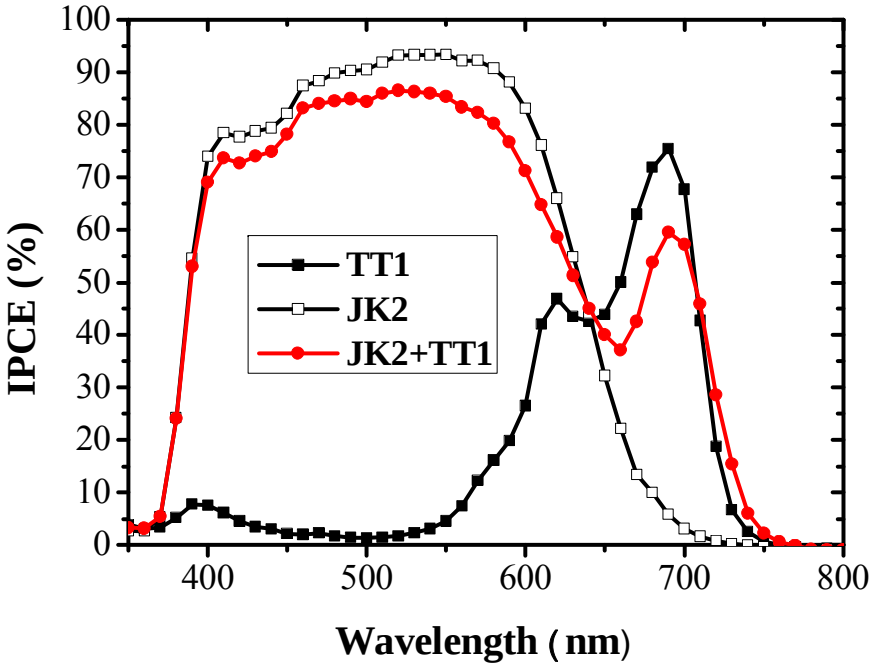


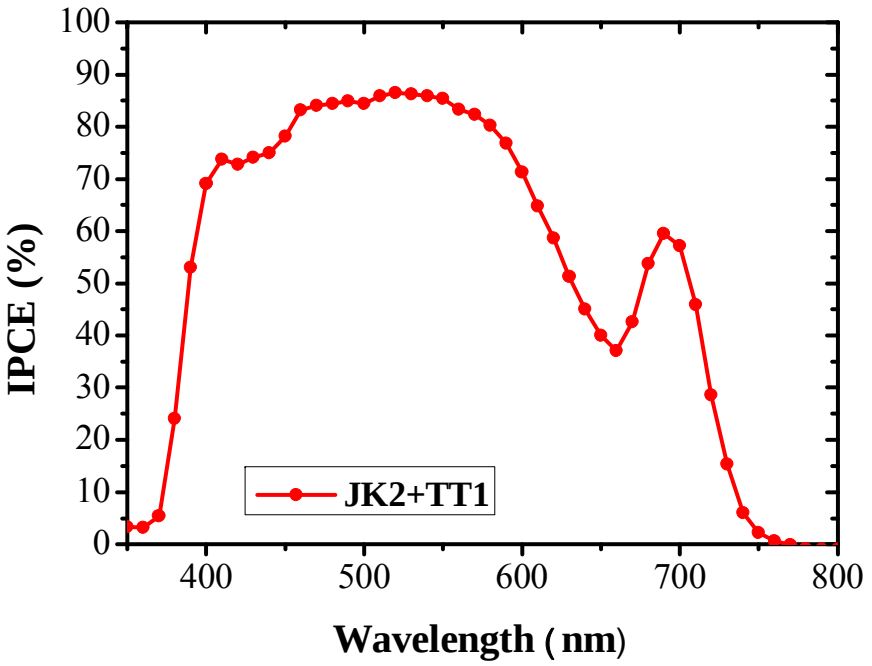




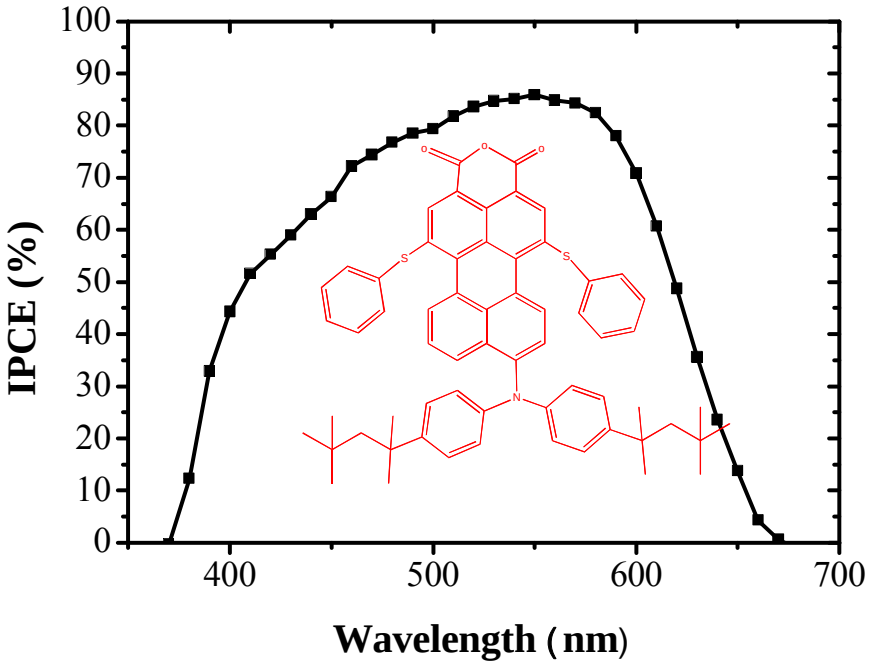




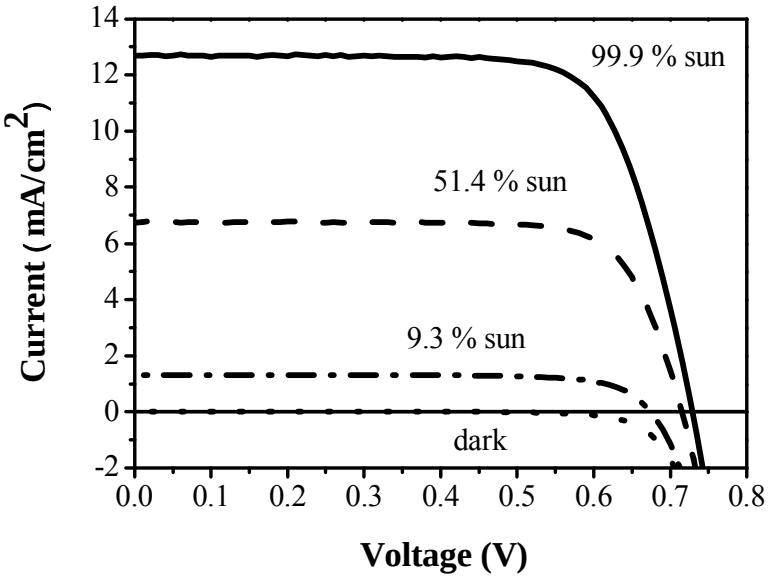




Dye	J (mA/cm2)	V (mV)	FF	efficiency
TT1	6.55	601.5	0.746	2.94
JK2	14.45	690.03	0.709	7.08
JK2+TT1	16.11	650.02	0.7	7.33
after 1 day	15.96	676.82	0.72	7.77



Light Intensity	J (mA/cm2)	V (mV)	FF	efficiency
99.9 %	12.68	728	0.736	6.80
51.4 %	6.74	715	0.766	7.18



Effect of solvent on performance of BASF_ID94 at 1 sun
TIO2: 6+4 μm; electrolyte A7117

Solvent	J (mA/cm2)	V (mV)	FF	efficiency
EtOH	3.75	641.0	0.740	1.77
DCM:AcN	10.30	651.0	0.723	4.85
CB	12.57	725.0	0.708	6.45
DCB	10.52	707.5	0.696	5.18
DMB	2.39	5079	0.725	0.88
BiCH	12.68	728.6	0.722	6.80
toluene	11.55	696.1	0.720	5.79
CCl_4	12.28	718.9	0.714	6.31

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