

Supporting Information

Development of fullerenes and their derivatives as semiconductors in field effect transistors: exploring the molecular design

Celebrating 50 years of Professor Fred Wudl's contributions to Organic Semiconductors

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1. Some relative papers based on C₆₀, C₇₀, PCBM and PC₇₁BM.

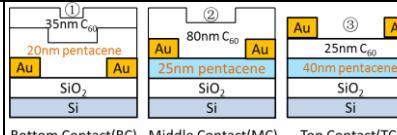
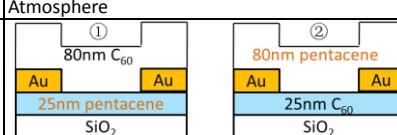
Abbreviation:

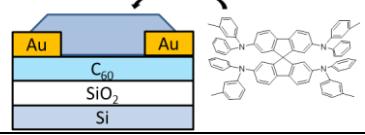
Alq3= tris(8-hydroxyquinoline) aluminum; **BCB**=divinyltetramethyldisiloxane-bis(benzocyclo-butene); **BCP**=bathocuproine; **Bphen**=bathophenanthroline; **b-PS**=polystyrene-based dimethylchlorosilane monolayer; **CB**=chlorobenzene; **CYTOP**=poly(perfluoroalkenyl vinyl ether); **CoTMPP**=5,10,15,20-tetrakis(4-methoxyphenyl) porphyrinato cobalt(II) hybrid nanosheets; **DABT**=4-(dimethylamino)benzenethiol; **DAE**=diarylethene; **DCB**=o-Dichlorobenzene; **DIP**=diindenoperylene; **DPTTA**=meso-diphenyltetrathia[22]-annulene[2,1,2,1]; **FP**=fullropyrrodine; **HMDS**=hexamethyldisilazane; **HWE**=hot-wall epitaxy; **NPs**=nanoparticles; **ODCB**=o-dichlorobenzene; **ODPA**=n-octadecylphosphonic acid; **ODS**=octadecyltrimethoxysilane; **ODT**=1-octadecanethiol; **OTS**=octadecyltrichlorosilane; **NW**=nano-whisker; **PAC**=poly(dimethylsiloxane) (PDMS)-assisted crystallization method; **PB-PyDI**=pyromellitic diimide based polymer; **Pc**=phthalocyanine; **PE**=polyethylene; **PEG**=poly(ethylene glycol); **PF**=poly(9,9-diethyl-fluorenyl-2,7-diyl) end capped with N,N-bis(4-methylphenyl)-4-aniline; **PFBT**=Pentafluorobenzenethiol; **PHDA**=phosphonohexadecanoic acid; **PMMA**=poly(methylmethacrylate); **PPV**=poly(phenylene vinylene); **PS**=polystyrene; **PTCDA**=3,4,9,10-perylenetetracarboxylic dianhydride; **PTmT**=poly(2,5-bis(3'-dodecyl-2,2'-bithiophen-5-yl)-3,6-dimethylthieno[3,2-*b*]thiophene); **PTS**=phenyltrimethoxysilane; **PVA**= poly(vinyl alcohol); **PVP**=poly(4-vinylphenol); **[RuCp*(mes)]₂**=ruthenium(pentamethylcyclopentadienyl)(1,3,5-trimethylbenzene) dimer; **SDS**=sodium dodecyl sulfate; **SU8**=(epoxy photosensitive commercial ink); **TCB**=1,2,4-trichlorobenzene; **UHV**=ultrahigh vacuum conditions; **ZSO**=zirconium-silicon oxide

Some relative papers are collected here by year. They are device engineering reports. Some of them are mentioned in the main text. In the main text, Figure 1 is extracted from the mobility presented in the second column in the following tables.

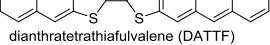
C₆₀

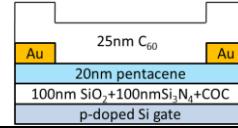
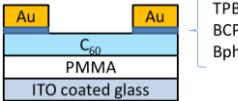
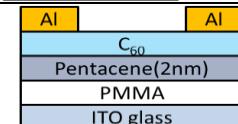
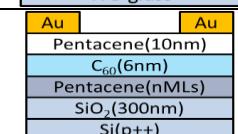
| Ref. | Mobility (cm ² V ⁻¹ s ⁻¹) | I _{on} /I _{off} | V _T (V) | Deposition Structure (Measured in) | Title |
|-------------|---|-----------------------------------|--------------------|---|---|
| 1993 | | | | | |
| 1 | Too low to be measured | | | Evaporating BGBC; Si/SiO ₂ ; Cr/Au Vacuum | Conduction mechanisms in undoped thin films of C ₆₀ and C _{60/70} |
| 2 | 10 ⁻⁴ | | | Evaporating BGBC; Si/SiO ₂ ; Cr/Au | Fullerene Field-Effect Transistors |
| 3 | Undoped: 4×10 ⁻⁵ In: 0.03 Sb: 0.04 | | | Evaporating BGBC; Si/SiO ₂ /C ₆₀ &dopant; Au Vacuum | Semiconductor-like carrier conduction and its field-effect mobility in metal-doped C ₆₀ thin films |
| 1995 | | | | | |
| 4 | 0.08 | 10 ⁶ | | Vacuum deposition BGBC; Si/SiO ₂ ; Au Vacuum | C ₆₀ thin film transistors |

| | | | | | |
|-------------|---|--|----------------------|--|---|
| 5 | $\mu_e=5\times 10^{-3}$ | | 40 | Evaporating BGBC; Si/SiO ₂ /α-hexathielenylene/C ₆₀ ; Au Vacuum | ① Organic Heterostructure Field-Effect Transistors; ② Organic field-effect bipolar transistors |
| 6 | 1996 | | | | |
| 7 | 4.8×10^{-5} | | 2.0 | Evaporating deposition BGBC; Si/SiO ₂ ; Au | Transport Mechanisms in Evaporated C ₆₀ Film Evaluated by Means of Field Effect |
| 8 | Air: 4×10^{-9} Vacuum: 2×10^{-3} | | | Vacuum deposition BGBC; Si/SiO ₂ /insulating/C ₆₀ ; Cr/Au Vacuum or Air | Transport studies in C ₆₀ and C ₆₀ /C ₇₀ thin films using metal-insulator-semiconductor field-effect transistors |
| 2002 | | | | | |
| 9 | 0.1 | | | Vacuum deposition BGBC; Si/SiO ₂ /C ₆₀ ; Au/Ti Vacuum | Passivation effects of alumina insulating layer on C ₆₀ thin-film field-effect transistors |
| 2003 | | | | | |
| 10 | 0.56 | $>10^8$ | 17 | Vacuum/molecular beam deposition BGBC; Si/SiO ₂ ; Ti/Au or Cr/Au Vacuum | C ₆₀ thin-film transistors with high field-effect mobility, fabricated by molecular beam deposition |
| 11 | 0.5-0.3 | $>10^8$ | | Vacuum/molecular beam deposition BGBC; Si/SiO ₂ ; Ti/Au or Cr/Au Vacuum | Fabrication and characterization of C ₆₀ thin-film transistors with high field-effect mobility |
| 2004 | | | | | |
| 12 | 0.085(linear) 0.22(photopolymerization) | | | Vacuum deposition BGBC; Si/SiO ₂ ; Cr/Au Vacuum | Accelerated photopolymerization and increased mobility in C ₆₀ field-effect transistors studied by ultraviolet photoelectron spectroscopy |
| 13 | Au: 5.6×10^{-7} (La@C ₈₂)/Au: 4.8×10^{-5} | 10^3 | 17.1 | Vacuum deposition BGBC; Au/Si/SiO ₂ ; (La@C ₈₂)/Au Vacuum | C ₆₀ field effect transistor with electrodes modified by La@C ₈₂ |
| 14 | Ag: 4.2×10^{-3} Mg/Ag: 0.064 | 9×10^3 1×10^5 | 18.8 18.9 | Vacuum deposition BGTC; Si/SiO ₂ /HMDS; Ag or Mg/Ag Vacuum | C ₆₀ thin-film transistors with low work-function metal electrodes |
| 15 | ① Only n-type after annealing ① Only p-type before annealing ② $\mu_e=1.3\times 10^{-3}$ ② $\mu_h=6.8\times 10^{-2}$ ③ not mention | 7.6×10^2 1.3×10^5 | 98 -15 |  | Fabrication of ambipolar field-effect transistor device with heterostructure of C ₆₀ and pentacene |
| 16 | ~ 0.11 | | | Molecular beam deposition generally fabricated Vacuum | Low-glancing-angle x-ray diffraction study on the relationship between crystallinity and properties of C ₆₀ field effect transistor |
| 2005 | | | | | |
| 17 | $\mu_e=2.6\times 10^{-4}(\text{sat})$ $\mu_h=6.4\times 10^{-4}$ | | 59 -82 | Vacuum deposition BGBC; Si/SiO ₂ /SAM; Cr/Au without exposing to air | Ambipolar operation of fullerene field-effect transistors by semiconductor/metal interface modification |
| 18 | $\mu_e=7\times 10^{-3}$ $\mu_h=1.7\times 10^{-2}$ | | 15.6 -2 | Vacuum deposition BGTC; Si/SiO ₂ /pentacene/C ₆₀ /LiF; Au Atmosphere | Ambipolar organic thin-film transistors using C ₆₀ /pentacene structure: Characterization of electronic structure and device property |
| 19 | ① $\mu_e=5.8\times 10^{-3}$ ① $\mu_h=3.7\times 10^{-2}$ ② $\mu_e=1.9\times 10^{-3}$ ② $\mu_h=3.1\times 10^{-5}$ | | |  | Fabrication of a logic gate circuit based on ambipolar field-effect transistors with thin films of C ₆₀ and pentacene |
| 20 | ① $7.1\times 10^{-3}(\text{lin})$ ① $1.2\times 10^{-2}(\text{sat})$ ② $4.1\times 10^{-5}(\text{without})$ ② $1.1\times 10^{-4}(\text{HMDS})$ | normally off 160 normally off | 7 2 3 -5 | thermal deposition ① BGBC; Au/polyimide/HMDS; Au ② BGBC; Si/SiO ₂ /Ba _{0.4} Sr _{0.6} Ti _{0.96} O ₃ /(HMDS); Au Vacuum | Fabrication of C ₆₀ field-effect transistors with polyimide and Ba _{0.4} Sr _{0.6} Ti _{0.96} O ₃ gate insulators |
| 21 | $\mu_e=2\times 10^{-3}\sim 9\times 10^{-3}$ $\mu_h=8\times 10^{-5}\sim 3\times 10^{-4}$ | | | Spin-coating/(di)chlorobenzene BGBC; Si/SiO ₂ /C ₆₀ &PPV; Ti/Au/cap Vacuum | Facile fabrication method for p/n-type and ambipolar transport polyphenylenevinylene-based thin-film field-effect transistors by blending C ₆₀ fullerene |
| 22 | 0h: 0.192 1h in ambient: 0.159 24 h in nitrogen 0.170 | 1.5×10^6 8.3×10^5 1.0×10^6 | 29.7 35.6 33.5 | Vacuum vapor deposition BGBC; Si/SiO ₂ ; Ti/Au Nitrogen or Air | Fullerene based n-type organic thin-film transistors |
| 23 | 0.4~1 | $>10^4$ | -35 | HWE BGBC; ITO/BCB/C ₆₀ ; LiF/Al Argon or Helium | High-mobility n-channel organic field-effect transistors based on epitaxially grown C ₆₀ films |
| 24 | 0.08 and 0.5(after annealing) oxygen exposure: $\mu_e=4\times 10^{-4}$ $\mu_h=4\times 10^{-5}$ | | | Molecular-beam deposition BGTC; Si/SiO ₂ /C ₆₀ ; Cr/Au UHV or oxygen exposure | Ultrapure C ₆₀ field-effect transistors and the effects of oxygen exposure |
| 2006 | | | | | |
| 25 | $5\times 10^{-4}(\text{undoped})$ 0.2(1.8mol% doped) | | | Vacuum deposition BGBC; Si/SiO ₂ /C ₆₀ &acridine orange; Au Vacuum | Acridine orange base as a dopant for n doping of C ₆₀ thin films |
| 26 | $\sim 10^{-3}$ | | ~ 70 | evaporated deposition BG; Si/SiO ₂ /C ₆₀ ; Au during film growth | Analysis of transient phenomena of C ₆₀ field effect transistors |

| | | | | | |
|----------------------------------|--|--|---|--|---|
| 27 | $\mu_e=0.23$ $\mu_h=0.14$ | | | Vacuum deposition BGBC; Si/SiO ₂ /pentacene; Au Nitrogen/ Vacuum/Air | Bottom Contact Ambipolar Organic Thin Film Transistors Based on C ₆₀ /Pentacene Heterostructure |
| 28 | 0.02 | normally-on | ~0 | NW from m-xylene/isopropyl alcohol BGBC; Si/SiO ₂ /C ₆₀ -NW; Ti/Au Vacuum | Electrical properties of field-effect transistors based on C ₆₀ nanowhiskers |
| 29 | ①6 ($T_{sub}=250^{\circ}\text{C}$) ①3 ($T_{sub}=120^{\circ}\text{C}$) ①0.6 ($T_{sub}=25^{\circ}\text{C}$) ②0.2 | | | Channel: HWE ①BGTC; ITO/BCB/C ₆₀ ; LiF/Al ②BGBC; Si/SiO ₂ /HMDS/C ₆₀ ; Ti/Au Vacuum | High performance n-channel organic field-effect transistors and ring oscillators based on C ₆₀ fullerene films |
| 30 | No pentacene: 0.25~1 With pentacene: 2.0~4.9 | | | Vacuum deposition BGTC; Al/Al ₂ O ₃ /(pentacene)/C ₆₀ ; Mg Vacuum | High-Mobility C ₆₀ Field-Effect Transistors Fabricated on Molecular-Wetting Controlled Substrates |
| 31 | 0.5~3 | | | Channel: HWE BGTC; ITO/BCB/C ₆₀ ; LiF/Al | Influence of film growth conditions on carrier mobility of hot wall epitaxially grown fullerene based transistors |
| 32 | Highest:0.28 | 7.3×10^6 | 18 | Vacuum deposition BGBC; Si/SiO ₂ ; Au Vacuum | Intrinsic transport and contact resistance effect in C ₆₀ field-effect transistors |
| 33 | $\mu_e=2.23 \times 10^{-3}$ $\mu_h=5.53 \times 10^{-4}$ | | 50.2 -37.1 |  | Organic heterostructure field-effect transistors using C ₆₀ and amorphous spirolinked compound |
| 34 | Eu: 0.5 Cu: 2.3×10^{-4} Pt: 2.4×10^{-2} | Normally-on Normally-off Normally-off | 34 | thermal deposition BGTC; Si/SiO ₂ /HMDS; Eu or Cu or Pt Vacuum | Output properties of C ₆₀ field-effect transistor device with Eu source/drain electrodes |
| 35 | 0.03 | | | Patterning BC; Si/SiO ₂ /PVP; Cr/Au Vacuum | Patterning organic single-crystal transistor arrays |
| 36 | ~0.6 | | | Channel: HWE BGTC; ITO/BCB/C ₆₀ ; LiF/Al | Switching in C ₆₀ -fullerene based field effect transistors |
| 2007 | | | | | |
| 37 | C ₆₀ : $\mu_e=0.32(\text{sat})$ C ₆₀ &CuPc: $\mu_e=2.2 \times 10^{-2} \sim 3.1 \times 10^{-4}$ C ₆₀ &CuPc: $\mu_h=1.6 \times 10^{-6} \sim 1.0 \times 10^{-4}$ | | 60.4 44.8~30.7 -24.0~-1.7 | thermal evaporation ring-type; Si/SiO ₂ /C ₆₀ (&CuPc); Ti/Au Vacuum | Ambipolar charge carrier transport in mixed organic layers of phthalocyanine and fullerene |
| 38 | 0.074(nonpolymerized) 0.068(polymerized) | | 19 | Vacuum deposition BCTC; Si/SiO ₂ ; Au dry box | C ₆₀ field-effect transistors: Effects of polymerization on electronic properties and device performance |
| 39 | 6 ($T_{sub}=250^{\circ}\text{C}$) | | | HWE | ①Characterization of highly crystalline C ₆₀ thin films and their field-effect mobility; |
| 40 | 3 ($T_{sub}=120^{\circ}\text{C}$) | | | BGTC; ITO/BCB/C ₆₀ ; LiF/Al glove box | ②Correlation of crystalline and structural properties of C ₆₀ thin films grown at various temperature with charge carrier mobility |
| Note: data is similar to Ref. 29 | 0.6 ($T_{sub}=25^{\circ}\text{C}$) | | | | |
| 41 | Untreated: $2.8 \times 10^{-3} \sim 6.6 \times 10^{-7}$ OTS: $1.7 \times 10^{-3} \sim 6.1 \times 10^{-5}$ | $10^3 \sim 10^5$ $10^4 \sim 10^6$ | 42~82 60~76 | Thermally evaporated BGBC; Si/SiO ₂ /(OTS); Cr/Au Helium or Air | Estimation of electron traps in carbon-60 field-effect transistors by a thermally stimulated current technique |
| 42 | 0.02 | | 45 | Thermally evaporated BGBC; Cr/PMMA; Au or Al in air | Fullerene thin-film transistors fabricated on polymeric gate dielectric |
| 43 | 1 | 10^5 (at 5V) | 1.13 | Thermally evaporated BGTC; Ti-Si/SiO ₂ /TiSiO ₂ /SiO ₂ /C ₆₀ ; LiF/Al Nitrogen | High performance n -channel thin-film transistors with an amorphous phase C ₆₀ film on plastic substrate |
| 44 | buffer layer BCB: 3.1 ± 0.2 (Max:5.0) PS: 2.1 ± 0.1 PMMA: 1.1 ± 0.1 OTS: 1.2 ± 0.1 | 1×10^7 6×10^6 6×10^6 2×10^6 | -0.1±0.4 1.2±1.3 2.1±0.4 1.8±0.6 | Thermally evaporated BGTC; /Si/SiO ₂ /buffer layer/C ₆₀ ; Al Nitrogen | High-performance and electrically stable C ₆₀ organic field-effect transistors |
| 45 | UV ozone: 0.061 HMDS: 1.04 ODS: 1.46 | 2×10^4 1×10^6 2×10^6 | 3.3 1.7 1.9 | Thermally evaporated BGTC; TiSi/SiO ₂ /ZSO/SiO ₂ /(SAM)/C ₆₀ ; LiF/Al Nitrogen | High-performance fullerene C ₆₀ thin-film transistors operating at low voltages |
| 46 | polymer dielectrics PVP: 0.27 PMMA: 0.66 | 1.6×10^5 3.9×10^5 | 2 5.5 | Thermally evaporated BGTC; ITO/polymer dielectrics/C ₆₀ ; Ba/Al Nitrogen | Influence of polymer dielectrics on C ₆₀ -based field-effect transistors |
| 47 | 0.68 | 1×10^6 | 0.80 | Thermally evaporated BGTC; TiSi/SiO ₂ /TiSiO/SiO ₂ /HMDS; Au Nitrogen | Low-voltage-operating complementary inverters with C ₆₀ and pentacene transistors on glass substrates |
| 48 | 5.9×10^{-3} | normally-off | 38 | Thermally evaporated BGBC; Si/SiO ₂ ;1-Alkanethiols/Cr/Au Vacuum | Output Properties of C ₆₀ Field-Effect Transistors with Au Electrodes Modified by 1-Alkanethiols |
| 49 | ITO: 0.16 Au: 0.096 Pt: 0.14 | 4.0×10^6 2.5×10^6 3.3×10^6 | 36 42 41 | Thermally evaporated BGBC; Si/SiO ₂ /C ₆₀ ; ITO or Au or Pt Vacuum | Output properties of C ₆₀ field-effect transistors with different source/drain electrodes |
| 2008 | | | | | |
| 50 | 0.07 | | 63 | Thermally evaporated BGBC; Si/SiO ₂ /OTS; Au Vacuum | Ambipolar charge carrier transport in organic semiconductor blends of phthalocyanine and fullerene |

| | | | | | |
|-------------|---|--|---|---|--|
| 51 | 6.8×10^{-2} | | | Thermally evaporated BGBC; Si/SiO ₂ ; Au Vacuum | Bipolar transport in organic field-effect transistors: organic semiconductor blends versus contact modification |
| 52 | 3.23-0.68 | 4×10^6 - 8×10^6 | 17.1~11.8 | Thermally evaporated BGBC; Si/SiO ₂ /HMDS; Au Nitrogen | Bottom-contact fullerene C ₆₀ thin-film transistors with high field-effect mobilities |
| 53 | Best: 0.41 | $\sim 10^7$ | | Thermally evaporated BGTC; Au/Cr/Si/SiO ₂ /parylene/C ₆₀ ; Au Helium | High-performance C ₆₀ thin-film field-effect transistors with parylene gate insulator |
| 54 | (1) Al(W/L=10): 1.7±0.1 Al(W/L=80): 1.4±0.05 Highest: 4.3(W=L=200μm, Ca) | $(0.4 \pm 0.1) \times 10^6$ $(1.0 \pm 0.2) \times 10^6$ | 0.3±0.1 0.2±0.1 | Thermally Vacuum evaporated (1) BGTC; Au/Ti/Si/SiO ₂ /Al ₂ O ₃ /BCB/C ₆₀ ; Al or Ca (2) BGTC; Si/SiO ₂ /Al ₂ O ₃ /BCB/C ₆₀ ; Ca Nitrogen | (1) High-performance C ₆₀ n-channel organic field-effect transistors through optimization of interfaces; (2) Low-voltage C ₆₀ organic field-effect transistors with high mobility and low contact resistance |
| 55 | (1) Ca(W/L=10): 2.3±0.2 Ca(W/L=80): 2.3±0.1 | $(1.0 \pm 0.3) \times 10^6$ $(4.0 \pm 0.3) \times 10^6$ | 0.2±0.1 0.1±0.1 | | |
| 56 | HfO ₂ /ODPA: 0.28(sat) HfO ₂ : 0.097(sat) | 10^5 10^3 | 0.35 0.40 | Vacuum deposition BGTC; Si/HfO ₂ /(ODPA); LiF/AI Vacuum | Low-voltage high-performance C ₆₀ thin film transistors via low-surface-energy phosphonic acid monolayer/hafnium oxide hybrid dielectric |
| 57 | 0.061 PTS: 1.22 HMDS: 1.04 ODS: 1.46 | 10^6 | 1.9 | Vacuum deposition BGTC; Ti-Si/Si/SiO ₂ /ZSO/SiO ₂ /insulator; LiF/AI Nitrogen | Low-voltage-operating fullerene C ₆₀ thin-film transistors with various surface treatments |
| 58 | Linear: 0.14 (highest) Sat: 0.26 (highest) | 2.9×10^7 1×10^7 | 33 32.9 | Vacuum deposition BGBC; Si/SiO ₂ ; Ti/Au Vacuum | Potential barriers to electron carriers in C ₆₀ field-effect transistors |
| 59 | 0.05~0.15 | | | BGBC/TC; Si/SiO ₂ /C ₁₂ H ₂₅ SH; Cr/Au Vacuum | Transport properties in C ₆₀ field-effect transistor with a single Schottky barrier |
| 60 | 0.15~0.55 | | 0.2~4 | Vacuum deposition BGBC; Al/ITO/polyaniline; Al Nitrogen | Vacuum-Processed Polyaniline-C ₆₀ Organic Field Effect Transistors |
| 2009 | | | | | |
| 61 | $\mu_e=0.04$ (sat) $\mu_h=0.2$ (sat) | 66 2.3 | | Vacuum deposition BGTC; Si/SiO ₂ /OTS/pentacene/C ₆₀ /pentacene; Au Nitrogen | Ambipolar pentacene/C ₆₀ -based field-effect transistors with high hole and electron mobilities in ambient atmosphere |
| 62 | 1.12-0.86(sat) | | 8.6-5.9 | Vacuum deposition BGTC; Al ₅ Si ₁ /SiO ₂ /HMDS; Au _{0.9} Ni _{0.1} /Au Nitrogen | Current-gain cutoff frequencies above 10 MHz for organic thin-film transistors with high mobility and low parasitic capacitance |
| 63 | spin-cast :4.7±0.41 vapor: 0.27±0.15 (Highest: 5.3) | $(3.5 \pm 1.2) \times 10^7$ $(7.5 \pm 6.3) \times 10^5$ | 35.6±6.33 39.8±7.5 | Vacuum /OTS by spin-cast or vapor BGTC; Si/SiO ₂ /OTS; Au Nitrogen | Crystalline Ultrasmooth Self-Assembled Monolayers of Alkylsilanes for Organic Field-Effect Transistors |
| 64 | ~6 | | ~11 | HWE BGTC; ITO/BCB; LiF/AI Vacuum | Electrical response of highly ordered organic thin film metal-insulator-semiconductor devices |
| 65 | Ca/Al:0.22 Al:0.21 Au:0.035 | 5×10^5 | -3 0.7 22 | Spin-cast/trichlorobenzene(1 wt%) BGTC; ITO/PVP; Al or Ca or Au Nitrogen | Flexible Fullerene Field-Effect Transistors Fabricated Through Solution Processing |
| 66 | Highest: 0.05 | $\sim 10^3$ | 20 | Liquid-liquid interface precipitation BGBC; Si/SiO ₂ ; Au/Ti Vacuum | Field-effect-transistor characteristics of solvate C ₆₀ fullerene nanowhiskers |
| 67 | 0.1-0.3 | | | Thermally deposition BGTC; Al/ITO/melamine/C ₆₀ ; Al Nitrogen | Small-molecule vacuum processed melamine-C ₆₀ organic field-effect transistors |
| 68 | $\mu_{lin}=0.0033$ ~1.3 $\mu_{sat} = 2.5$ ~2.8 | 4×10^5 ~ 5×10^7 | 5.1~16.8 | Vacuum deposition BGTC; Si/SiO ₂ /HMDS;/benzenethiol/Ni-Au/Au Nitrogen | Threshold voltage control of bottom-contact n-channel organic thin-film transistors using modified drain/source electrodes |
| 2010 | | | | | |
| 69 | 6.5(lin) | | | HWE BGTC; ITO/BCB/C ₆₀ ; LiF/AI glove box | Dependence of Meyer–Neldel energy on energetic disorder in organic field effect transistors |
| 70 | 3.2×10^{-2} (pristine) 5.2×10^{-4} (supersonic wave) 2.4×10^{-3} (ultraviolet light) | ~ 10 ~ 1000 ~ 10 | | NW from m-xylene/isopropyl alcohol BGBC; Si/SiO ₂ /C ₆₀ -NW; Ti/Au Vacuum or air(working) | Electron Transport Properties in Photo and Supersonic Wave Irradiated C ₆₀ Fullerene Nano-Whisker Field-Effect Transistors |
| 71 | Lactose:0.055 Glucose:0.085 Guanine:0.12 Cytosine:0.09 Adenine:5.5(HWE) Thymine:0.5 | | | Thermally deposition BGTC; Al/Dielectric/C ₆₀ ; Al or Au | Environmentally sustainable organic field effect transistors |
| 72 | BC: $\mu_e=5.28 \times 10^{-3}$ BC: $\mu_h=4.2 \times 10^{-2}$ MC: $\mu_e=5.5 \times 10^{-2}$ MC: $\mu_h=5.45 \times 10^{-2}$ TC: $\mu_e=2.44 \times 10^{-2}$ TC: $\mu_h=4.5 \times 10^{-3}$ | | 79 14.5 73 -4.3 68.5 -30 | Bottom Contact(BC) Middle Contact(MC) Top Contact(TC) OSC are deposited by thermal evaporation, test atmosphere is controlled with 0.1 ppm O ₂ , Poly(3,4-ethylene dioxythiophene):Poly(styrenesulfonate) (PEDOT:PSS) acts as Gate and Source/Drain electrodes. | Influence of device geometry in the electrical behavior of all organic ambipolar field effect transistors |

| | | | | | |
|-------------|--|---|---|---|--|
| 73 | Ba: 1.15 Al: 0.97 Au: 0.126 | 3.2×10^5 2.8×10^5 6.7×10^4 | 29 30 36 | Thermally deposition BGTC; ITO/polystyrene/C ₆₀ ; Ba/Al or Al or Au Nitrogen | Properties of C ₆₀ thin film transistor based on polystyrene |
| 2011 | | | | | |
| 74 | 100°C: ~1 250°C: ~5 | | | Thermally deposition BGBC; ITO/BCB; LiF/Al | Electric field and grain size dependence of Meyer–Neldel energy in C ₆₀ films |
| 75 | 0nm Alq3: 1.71×10^{-3} 5nm Alq3: 1.12×10^{-2} 10nm Alq3: 1.28×10^{-2} 15nm Alq3: 1.88×10^{-3} | 10^4 10^2 10^2 10^2 | 18 11 10 21 | thermally evaporated BGTC; Si/SiO ₂ /PMMA/C ₆₀ /Alq3; Al Ar atmosphere | Enhanced performance of C ₆₀ organic field effect transistors using a tris(8-hydroxyquinoline) aluminum buffer layer |
| 76 | BCB: 5.1 Parylene: 0.046 AlO _x –BCB: 3.5 AlO _x –PE: 2.9 AlO _x –Adenine: 3.2 | $>10^6$ $\sim 10^{2.5}$ $\sim 10^{2.5}$ $>10^3$ $\sim 10^3$ | 13.2 -3.5 -0.003 0.39 -0.25 | Channel: HWE BGTC; Al/BCB; Al TGBC; Al/Parylene-C; Al BGTC; Al/AlO _x /BCB or PE or Adenine; Al Nitrogen | High mobility, low voltage operating C ₆₀ based n-type organic field effect transistors |
| 77 | Ag: 2.74 LiF/Ag: 5.07 | | | Thermally deposition BGTC; ITO/PMMA/pentacene/C ₆₀ ; (LiF)/Ag Nitrogen | Mobility Improvement in C ₆₀ -Based Field-Effect Transistors Using LiF/Ag Source/Drain Electrodes |
| 78 | Solution: 0.6~0.8(highest:0.86) Vacuum: 0.7~0.9 | (4×10^6) | (3) | Drop cast and dry/Vacuum deposition BGTC; Si/SiO ₂ /PTS; LiF/Al Nitrogen | Novel Solution Process for High-Mobility C ₆₀ Fullerene Field-Effect Transistors |
| 79 | 1.5 0.0012 0.005 | | | evaporation BGTC; Si/SiO ₂ /OTS/C ₆₀ ; Au BGTC; Si/SiO ₂ /OTS/DATTF/C ₆₀ ; Au BGB; Si/SiO ₂ /OTS/C ₆₀ ; DATTF/C ₆₀ BGB; Si/SiO ₂ /OTS/C ₆₀ ; DATTF  dianthratetrathiafulvalene (DATTF) | Organic Electrodes Consisting of Dianthratetrathiafulvalene and Fullerene and Their Application in Organic Field Effect Transistors |
| 2012 | | | | | |
| 80 | 4.02 ± 0.35 | $(4.68 \pm 1.04) \times 10^6$ | | Vacuum deposition BGTC; Si/SiO ₂ , BCB; Au Nitrogen | 2-(2-Methoxyphenyl)-1,3-dimethyl-1H-benzimidazol-3-ium Iodide as a New Air-Stable n-Type Dopant for Vacuum-Processed Organic Semiconductor Thin Films |
| 81 | Only C ₆₀ : 0.38 C ₆₀ /Bphen: 0.50 Pentacene/C ₆₀ : 4.11 Pentacene/C ₆₀ /Bphen: 5.17 | 4.4×10^3 3.1×10^3 19.5 25 | 13 11 13 12.5 | Thermal evaporation BGTC; ITO/PMMA/pentacene/C ₆₀ ; Bphen/Ag Nitrogen | A high mobility C ₆₀ field-effect transistor with an ultrathin pentacene passivation layer and bathophenanthroline/metal bilayer electrodes |
| 82 | $0.17 \pm 0.02 \sim 0.11 \pm 0.02$ | | $1.4 \pm 0.1 \sim 3.8 \pm 0.1$ | Vacuum deposition BGTC; Si/SiO ₂ /HMDS; Au Nitrogen | Direct structuring of C ₆₀ thin film transistors by photo-lithography under ambient conditions |
| 83 | Ti: 1.94 Ti/Pt: 2.09 | 0.61×10^6 1.69×10^6 | 0.51 1.01 | Thermal evaporation BGTC; Ti/(Pt)/Si/SiO ₂ /AlO _x /SAM; Au Nitrogen | Engineering the metal gate electrode for controlling the threshold voltage of organic transistors |
| 84 | 10^{-5} (μ_e) 10^{-6} (μ_h) | | | Dropping BGBC; Si/SiO ₂ /HMDS/C ₆₀ &CoTMPP; Au Nitrogen | Fullerene/Cobalt Porphyrin Hybrid Nanosheets with Ambipolar Charge Transporting Characteristics |
| 85 | Needles: 5.2 ± 2.1 (average) ~11 (highest) Ribbons: 3.0 ± 0.87 (average) | $>10^5$ $>10^6$ | 15~43 36~85 | Solution Grown (Single Crystals) BGTC; Si/SiO ₂ /BCB/C ₆₀ ; Au Nitrogen | High-Mobility Field-Effect Transistors from Large-Area Solution-Grown Aligned C ₆₀ Single Crystals |
| 86 | Highest: 0.081 | $>10^4$ | | Drop-casting or dip-coating deposition BGB; Si/SiO ₂ /C ₆₀ ; Cr/Au Nitrogen | On the fabrication of crystalline C ₆₀ nanorod transistors from solution |
| 87 | 0.4 | 6×10^4 | 2.8 | Vacuum deposition BGTC; Si/SiO ₂ /CYTOP; Al Nitrogen | Organic nonvolatile memory transistors based on fullerene and an electron-trapping polymer |
| 88 | Unpurified C ₆₀ : No dopant: 0.38 ± 0.02 Dopant: $0.48 \pm 0.03 \sim 0.67 \pm 0.02$ purified C ₆₀ : No dopant: 1.62 ± 0.03 Dopant: $1.3 \pm 0.1 \sim 1.73 \pm 0.02$ | 1×10^6 $1 \times 10^5 \sim 1 \times 10^6$ 3×10^6 $70 \sim 3 \times 10^6$ | 17.9±0.8 4.7±0.6~15±1 4.7±0.3 -0.4±0.9~4.7±0.3 | Evaporation or co-evaporation with dopant of [Ru(Cp*) ₂ (mes)] ₂ BGTC; Si/SiO ₂ /BCB/C ₆₀ (&Dopant); Al Nitrogen | Passivation of trap states in unpurified and purified C ₆₀ and the influence on organic field-effect transistor performance |
| 89 | 1.4 | | | drop-casted to solution patterning SAM BGTC; Si/SiO ₂ /PTS; LiF/Al | Solution-Processed C ₆₀ Single-Crystal Field-Effect Transistors |
| 90 | Chlorobenzene: 0.16 m-Xylene: 0.083 Tetrahydronaphthalene: 0.18 1,2,4-Trichlorobenzene: 0.86 | 1.2×10^6 1.1×10^6 3.6×10^5 3.9×10^6 | 5.2 3.6 5.3 1.5 | Vacuum-drying from various solvents BGTC; Si/SiO ₂ /HMDS; LiF/Al Nitrogen | Solvent Dependence of Vacuum-Dried C ₆₀ Thin-Film Transistors |
| 91 | 0.58 | 10^5 | -0.1 | HWE TGBC; Al/Parylene-C/C ₆₀ ; Al Nitrogen | Strain induced anisotropic effect on electron mobility in C ₆₀ based organic field effect transistors |
| 2013 | | | | | |
| 92 | 0.08±0.01(sat) 0.05±0.01(lin) | | 1.2±0.08 | Evaporation deposition BGTC; Al/Al ₂ O ₃ /cellulose/C ₆₀ ; Al Glove box | Cellulose as biodegradable high-k dielectric layer in organic complementary inverters |

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|-------------|---|--|--|--|--|
| 93 | $\mu_e=0.18$ $\mu_h=0.28$ | | |  | Dual Channel Operation Upon n-Channel Percolation in a Pentacene-C ₆₀ Ambipolar Organic Thin Film Transistor |
| 94 | No buffer:0.19 TPBi:0.25 BCP:0.52 Bphen:0.65 | 0.14×10^5 0.34×10^5 0.66×10^5 1.25×10^5 | 35 26 27 25 |  | Effect of organic buffer layers on the performance of n-type organic field-effect transistor based on C ₆₀ active layer |
| 95 | No Pentacene: 0.213 Pentacene (2nm): 1.01 | 10^3 10^4 | 3 11 |  | Enhanced performance of C ₆₀ N-type organic field-effect transistors using a pentacene passivation layer |
| 96 | $\mu_e=2.8$ $\mu_h=0.3$ | | |  | Enhancing crystallinity of C ₆₀ layer by thickness-control of underneath pentacene layer for high mobility C ₆₀ /pentacene ambipolar transistors |
| 97 | Cocrystals of C ₆₀ -DPTTA $\mu_e=0.01$ $\mu_h=0.3$ | | | Drop-casting/chlorobenzene BGTC; Si/SiO ₂ ; Au Vacuum | Fullerene/Sulfur-Bridged Annulene Cocrystals: Two-Dimensional Segregated Heterojunctions with Ambipolar Transport Properties and Photoresponsivity |
| 98 | 2.4-2.2 | $10^7 \sim 10^8$ | 0.4~0.6 | Inkjet-printing and vacuum drying / TCB TC; Si/SiO ₂ /CYCLOTENE; Al or Au or Ag | High performance inkjet-printed C ₆₀ fullerene thin-film transistors: Toward a low-cost and reproducible solution process |
| 99 | 3.3×10^{-6} (dark); 1.5×10^{-4} (illumination) 7.2×10^{-5} (dark); 1.4×10^{-3} (illumination) 2.8×10^{-3} (dark); ----- (illumination) | | | evaporated BCTC; Si/SiO ₂ /PdPc/C ₆₀ ; Au BCTC; Si/SiO ₂ /C ₆₀ /PdPc; Au BCTC; Si/SiO ₂ /C ₆₀ ; Au Vacuum with or without illumination | Influence of donor-acceptor layer sequence on photoresponsive organic field-effect transistors based on palladium phthalocyanine and C ₆₀ |
| 100 | 1 day: 1.110 14 days: 0.669 | 10^3 10^4 | 0 7 | Thermally evaporated BGTC; Al/Aloe vera + 1.5wt%SiO ₂ NPs/C ₆₀ ; Al Exposure to open air | N-Type Organic Field-Effect Transistor Based on Fullerene with Natural Aloe Vera/SiO ₂ Nanoparticles as Gate Dielectric |
| 101 | Al: 1.82(highest) dimer/Al: 2.23(highest) | 1×10^6 2×10^6 | 4.6 5.0 | Thermally evaporated BGTC; Si/SiO ₂ /BCB; (rhodocene dimer)/Al Nitrogen | Reduction of contact resistance by selective contact doping in fullerene n-channel organic field-effect transistors |
| 102 | As-grown: $3.0 (\pm 2.9) \times 10^{-1}$ (Heighest: 1.01) Photo-exposed: $4.7 (\pm 3.9) \times 10^{-3}$ | 10^{-4} 10^{-3} | $21.5 (\pm 3.8)$ $20.4 (\pm 5.5)$ | Drop-casting/ nanorods in m-DCB + ethanol BGBC; Si/SiO ₂ /C ₆₀ ; Cr/Au Nitrogen | Solution-Based Phototransformation of C ₆₀ Nanorods: Towards Improved Electronic Devices |
| 103 | No pentacene: 0.014 Pentacene(Vacuum):1 Pentacene(Air): 10 | $\sim 5 \times 10^2$ $\sim 3 \times 10^2$ 1×10^3 | 25 36 5.9 | Thermally evaporated BGTC; Au/silk fibroin/(pentacene)/C ₆₀ ; Au Vacuum or Air | Solution-based silk fibroin dielectric in n-type C ₆₀ organic field-effect transistors: Mobility enhancement by the pentacene interlayer |
| 2014 | | | | | |
| 104 | Height: ~1 | | | HWE BGTC; Al/Parylene; Al(encapsulated) Air | Air stability of C ₆₀ based n-type OFETs |
| 105 | 0.32 | 6×10^3 | 2.2 | drop-casting/DCB BGTC; Si/SiO ₂ /BCB; Al Nitrogen | Comparative Study of the N-Type Doping Efficiency in Solution-processed Fullerenes and Fullerene Derivatives |
| 106 | BG: 0.1 TG: 0.2 dual gate: 0.9 | $\sim 1 \times 10^3$ $\sim 2 \times 10^3$ $\sim 1 \times 10^4$ | 20.7~34.6 14.3~0.1 11.5~8.5 | HWE under vacuum TG, BG or dual gate; Al/Parylene; Al Nitrogen | Geometrical Structure and Interface Dependence of Bias Stress Induced Threshold Voltage Shift in C ₆₀ -Based OFETs |
| 107 | 10^{-4} Pa: 0.199 ± 0.020 10^{-3} Pa: 0.204 ± 0.015 10^{-2} Pa: 0.195 ± 0.017 10^{-1} Pa: 0.090 ± 0.013 | $10^{5 \sim 10^7}$ $10^{5 \sim 10^7}$ $10^{5 \sim 10^7}$ $10^{5 \sim 10^7}$ | 11.9 ± 0.8 7.2 ± 1.0 8.7 ± 0.7 13.2 ± 1.2 | Deposition under different pressure BGTC; Si /SiO ₂ /C ₆₀ , Au Vacuum | Influence of Deposition Pressure on the Film Morphologies, Structures, and Mobilities for Different-Shaped Organic Semiconductors |
| 108 | No DIP: 0.21±0.10 DIP: 2.62±0.32(Max:2.92) | 3×10^4 4×10^5 | 17 5 | Deposition under different pressure BGTC; Si/SiO ₂ /(DIP)/C ₆₀ , Cu Vacuum | Interface optimization using diindenoperylene for C ₆₀ thin film transistors with high electron mobility and stability |
| 109 | ~1.6(lin) | | | HWE BGTC; ITO/BCB/C ₆₀ ; LiF/Al Vacuum | Origin of Electric Field Dependence of the Charge Mobility and Spatial Energy Correlations in C ₆₀ -Based Field Effect Transistors |
| 110 | needle crystals: 0.08±0.04 (highest): 0.34±0.13 ribbon crystals: 2.0±0.61 | $> 10^5$ $> 10^6$ | 15~43 36~85 | Crystals grown from m-xylene/(CCl ₄) BGTC; Si/SiO ₂ /BCB/C ₆₀ , Au Nitrogen | Solution-grown aligned C ₆₀ single-crystals for field-effect transistors |
| 2015 | | | | | |
| 111 | Highest: 0.38 | 5.3×10^5 | 21.2 | Vacuum deposition BGTC; Si/SiO ₂ /PVA/OTS/C ₆₀ /BCP, Al | A striking performance improvement of fullerene n-channel field-effect transistors via synergistic interfacial modifications |
| 112 | Average(Highest) m-xylene:0.07(0.155) CS ₂ :1.24(1.70) ODCB:0.3(1.06) | $10^4 \sim 10^5$ | 40~60 | PAC method /m-xylene, CS ₂ , or ODCB BGTC; Si/SiO ₂ ; Au Nitrogen | A Facile PDMS-Assisted Crystallization for the Crystal-Engineering of C ₆₀ Single-Crystal Organic Field-Effect Transistors |
| 113 | 3.9×10^{-2} (dark) 6.7×10^{-3} (air) | | 45 | Vacuum deposition BGTC; Si/SiO ₂ /OTS/C ₆₀ /PbPc, Au Air or dark box | Enhanced performance of isotype planar heterojunction photoresponsive organic field-effect transistors by using Ag source-drain electrodes |

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|-------------|--|--|--|--|---|
| 114 | thermally anneal 50°C: 0.002±0.001 90°C: 0.027±0.002 100°C: 0.055±0.004 | ~10 ⁴ ~10 ⁵ ~10 ⁶ | 2.93 3.20 2.39 | Spin-coating/ dichlorobenzene BGBC; Si/SiO ₂ /PTS/C ₆₀ ; graphene Nitrogen | Solution-processed n-type fullerene field-effect transistors prepared using CVD-grown graphene electrodes: improving performance with thermal annealing |
| 115 | ①with PbPc:2.77×10 ⁻¹ /6.64×10 ⁻¹ ②with PbPc:2.41×10 ⁻⁵ /6.93×10 ⁻⁴ ①without PbPc: 2.48×10 ⁻² /- ②without PbPc: 1.79×10 ⁻³ /- | | 6.80/6.77 28.7/14.3 14.1/- 29.5/- | Vacuum deposition ①BGTC; ITO/PVA/C ₆₀ /(PbPc), Au ②BGTC; Si/SiO ₂ /OTS/C ₆₀ /(PbPc), Au In the dark/under illumination | Ultrahigh near infrared photoresponsive organic field-effect transistors with lead phthalocyanine/C ₆₀ heterojunction on poly(vinyl alcohol) gate dielectric |
| 2016 | | | | | |
| 116 | ODT: 0.15/0.88 (lin/sat↓) PFBT: 0.24/1.27 DABT: 0.3/1.52 No thiol derivatives: 0.12/0.56 | 6×10 ⁴ 7×10 ⁴ 9×10 ⁴ 2.5×10 ⁵ | 10.1 8.5 6.3 10.5 | Vacuum deposition BGBC; Al/SU8/C ₆₀ ; thiol derivatives/Al Nitrogen | Improvement of n-type OTFT electrical stability by gold electrode modification |
| 117 | 1.05 | 5.65×10 ² | 2.96 | Vacuum deposition BGTC; Al/PVA/SDS/C ₆₀ /(encapsulation); Au Air | Poly(Vinyl Alcohol) Gate Dielectric Treated With Anionic Surfactant in C ₆₀ Fullerene-Based n-Channel Organic Field Effect Transistors |
| 2017 | | | | | |
| 118 | 5.6 | >10 ⁵ | 4.9 | Drop/CCl ₄ + <i>m</i> -xylene BGTC; Si/SiO ₂ /BCB/C ₆₀ ; Au Nitrogen | Enhanced performance of field-effect transistors based on C ₆₀ single crystals with conjugated polyelectrolyte |
| 119 | Pure C ₆₀ : 2.43×10 ⁻² (sat) Heterojunction:1.43×10 ⁻³ (sat) | | 37 | Evaporation deposition BGTC; Si/SiO ₂ /C ₆₀ /(PTCDA:AlClPc:PbPc); Au in the dark or illumination | Towards high performance broad spectral response fullerene based photosensitive organic field effect transistors with tricomponent bulk heterojunctions |

C₇₀

| Ref. | Mobility (cm ² V ⁻¹ s ⁻¹) | I _{on} /I _{off} | V _T (V) | Deposition Structure Measured in | Title |
|-------------|--|---|----------------------|---|--|
| 1996 | | | | | |
| 120 | 2×10 ⁻³ | 10 ⁵ | 27 | Vacuum deposition BGBC; Si/SiO ₂ ; Au/Cr Vacuum | C ₇₀ thin film transistors |
| 2005 | | | | | |
| 22 | 0h: 0.060 1h in ambient: 0.042 24 h in nitrogen 0.041 | 3.9×10 ⁵ 1.5×10 ⁵ 1.2×10 ⁵ | 34.7 40.1 42.2 | Vacuum vapor deposition BGBC; Si/SiO ₂ ; Ti/Au Nitrogen or Air | Fullerene based n-type organic thin-film transistors |
| 2013 | | | | | |
| 97 | Cocrystals of C ₇₀ -DPTTA $\mu_e=0.05$ $\mu_h=0.07$ | | | Drop-casting/chlorobenzene BGTC; Si/SiO ₂ ; Au Vacuum | Fullerene/Sulfur-Bridged Annulene Cocrystals: Two-Dimensional Segregated Heterojunctions with Ambipolar Transport Properties and Photoresponsivity |
| 2014 | | | | | |
| 105 | 0.94 | 3×10 ⁴ | 3.8 | drop-casting/DCB BGTC; Si/SiO ₂ /BCB; Al Nitrogen | Comparative Study of the N-Type Doping Efficiency in Solution-processed Fullerenes and Fullerene Derivatives |

PCBM

| Ref. | Mobility (cm ² V ⁻¹ s ⁻¹) | I _{on} /I _{off} | V _T (V) | Deposition Structure Measured in | Title |
|-------------|---|--------------------------------------|--------------------|--|---|
| 2003 | | | | | |
| 121 | Highest: 4.5×10 ⁻³ | | 41 | Spin-coating/ chloroform BGTC; Au/Ti/resin; Ca or Al or Au Under ambient conditions (sealed) | Solution-Processed Organic n-Type Thin-Film Transistors |
| 2004 | | | | | |
| 122 123 | $\mu_e=1\times10^{-2}$ $\mu_h=8\times10^{-3}$ | 10 ⁶ 10 ⁶ | | Spin-coating/ chlorobenzene BGBC; Si/SiO ₂ /HMDS/PCBM; Ti/Au Vacuum | ①Ambipolar Organic Field-Effect Transistors Based on a Solution-Processed Methanofullerene; ②Organic complementary-like inverters employing methanofullerene-based ambipolar field-effect transistors |
| 124 | 9×10 ⁻² | 10 ⁴ | | Spin-coating/ chlorobenzene BGTC; ITO/PVA/PCBM; Cr Argon | Nonvolatile organic field-effect transistor memory element with a polymeric gate electret |
| 125 | $\mu_e=1\times10^{-2}$ $\mu_h=8\times10^{-3}$ | ~10 ⁶ ~10 ⁶ | | Spin-coating/ chlorobenzene BGBC; Si/SiO ₂ /HMDS/PCBM; Ti/Au Vacuum | Organic complementary-like inverters employing methanofullerene-based ambipolar field-effect transistors |

| 2005 | | | | | |
|------|--|---|---|--|--|
| 126 | 0.02-0.1 | $\sim 4 \times 10^3$ | 13.6 | Spin-coating/ chlorobenzene BGTC or BGBC; Al-Nd/insulator; Ca, Mg, Al, Ag, Au; Air(with protector) | All-solution-processed n-type organic transistors using a spinning metal process |
| 127 | PVA: $\mu_e=5 \times 10^{-4}$ BCB-Au: $\mu_e=5 \times 10^{-5}$ BCB-Au: $\mu_h=1 \times 10^{-5}$ BCB-LiF/Al: $\mu_e=10^{-4}$ PVP: poor transport properties | 7 4 | | Spin-coating/ chlorobenzene BGTC; ITO/dielectric/PPV&PF&PCBM; LiF/Al or Au | Correlation between morphology and ambipolar transport in organic field-effect transistors |
| 128 | 0.05-0.2 | $7.5 \times 10^2 \sim 2 \times 10^3$ | -20~7 | Spin-coating/ chlorobenzene BGTC; ITO/BCB or PVA/PCBM; Cr or LiF/Al Argon | Fabrication and characterization of solution-processed methanofullerene-based organic field-effect transistors |
| 129 | 0.024 | | | Spin-coating/ chloroform BGTC; Si/SiO ₂ ; Au | Relation between carrier mobility and cell performance in bulk heterojunction solar cells consisting of soluble polythiophene and fullerene derivatives |
| 130 | PCBM : 10^{-3} PCBM&P3HT: 10^{-4} | | | Spin-coating/ chloroform BGTC; Si /SiO ₂ /insulator/PCBM(P3HT); Al Nitrogen | Study of field effect mobility in PCBM films and P3HT:PCBM blends |
| 2006 | | | | | |
| 131 | $\mu_e=2.0 \times 10^{-3}$ $\mu_h=1.7 \times 10^{-3}$ | | | Spin-coating/ chlorobenzene BGBC; Si /SiO ₂ /PCBM&P3HT; Al Nitrogen | Ambipolar organic field-effect transistors fabricated using a composite of semiconducting polymer and soluble fullerene |
| 132 | $\mu_e=\sim 10^{-6} \sim 10^{-2}$ $\mu_h=\sim 10^{-3} \sim 10^{-4}$ | | | Spin-coating/ chloroform BGTC; Si/SiO ₂ /PCBM(&P3HT); Au Vacuum | Field effect measurements on charge carrier mobilities in various polymer-fullerene blend compositions |
| 133 | $\sim 10^{-4} \sim 10^{-6}$ | | | Spin-coating/ chloroform BGBC; Si /SiO ₂ ; Au Vacuum | Investigations of electron injection in a methanofullerene thin film transistor |
| 134 | PCBM-Au:0.008 PCBM-Mg:0.01 PCBM&P3HT: $\mu_e=7 \times 10^{-5}$ PCBM&P3HT: $\mu_h=1 \times 10^{-4}$ | | | Spin-coating/ chloroform BGTC; Si /SiO ₂ /PCBM(&P3HT); Au or Mg Vacuum | Investigations of the effects of tempering and composition dependence on charge carrier field effect mobilities in polymer and fullerene films and blends |
| 135 | Insulator PVA: 10^{-2} BCB: 10^{-2} | | | Spin-coating/ chlorobenzene BGTC; ITO/insulator/PCBM&PPV; LiF/Al Argon | Photoresponse of Organic Field-Effect Transistors Based on Conjugated Polymer/Fullerene Blends |
| 2007 | | | | | |
| 136 | pure PCBM: 0.01 | $>10^4$ | 20~38 | Spin-coating/chloroform BGTC; Si/SiO ₂ /PCBM(&P3HT); Au Vacuum | Ambipolar Transport in Field-Effect Transistors Based on Composite Films of Poly(3-hexylthiophene) and Fullerene Derivative |
| 137 | 1.1×10^{-5} | | | Spin-coating/chloroform BG BC or TC; Si/SiO ₂ ; Au or Al Vacuum | Ambipolar Field-Effect Transistors Based on Poly(3-hexylthiophene)/Fullerene Derivative Bilayer Films |
| 138 | $10^{-3} \sim 10^{-4}$ (sat) | $\sim 10^4$ | ~ 20 (electron) ~ 5 (hole) | Spin-coating/chlorobenzene BGBC; Si/SiO ₂ /HMDS/PCBM(&PPV); Ti/Au Vacuum or Nitrogen | Electro-optical circuits based on light-sensing ambipolar organic field-effect transistors |
| 139 | o-Xylene: 2×10^{-2} Chlorobenzene: 8×10^{-3} | | | Spin-coating/solvent BGBC; Si/SiO ₂ /HMDS; Au Nitrogen | Organic Field-Effect Devices as Tool to Characterize the Bipolar Transport in Polymer-Fullerene Blends: The Case of P3HT-PCBM |
| 140 | 8.6×10^{-3} | 1.5×10^5 | 43.0 | Spin-coating/chloroform BGTC; Si/SiO ₂ /HMDS; Au Vacuum | Unipolarization of ambipolar organic field effect transistors toward high-impedance complementary metal-oxide-semiconductor circuits |
| 141 | 0.02~0.034 | $4.5 \times 10^5 \sim 1.42 \times 10^6$ | | Spin-coating/chloroform BGBC; Si/SiO ₂ /HMDS; Ti/Au; Nitrogen | Solution-Processed n-Type Organic Field-Effect Transistors With High ON/OFF Current Ratios Based on Fullerene Derivatives |
| 2008 | | | | | |
| 142 | Vaccum: 0.025 Air: Not active | 2×10^4 | 33 | Spin-coating/ chloroform BGTC; Si/SiO ₂ /HMDS; Au Vacuum/Air | High-Performance n-Type Organic Thin-Film Transistors Based on Solution-Processable Perfluoroalkyl-Substituted C ₆₀ Derivatives |
| 143 | 0.21 | $>10^4$ | 7 | Spin-coating/ chlorobenzene BGTC; ITO/BCB; Ca/Al Nitrogen | High mobility n-channel organic field-effect transistors based on soluble C ₆₀ and C ₇₀ fullerene derivatives |
| 144 | Au: 0.0081 Al: 0.0120 Ca/Al: 0.0227 Cs ₂ CO ₃ /Al: 0.0445 | 10^3 10^6 10^3 10^3 | 4.87 3.15 0.74 -2.30 | Spin-coating/ chloroform BGTC; ITO/PVP; S/D Nitrogen | Improved performance in n-channel organic thin film transistors by nanoscale interface modification |
| 145 | Dark: 2.9×10^{-5} Light: 3.7×10^{-5} | $\sim 10^4$ | 24 20 | Spin-coating/ chloroform BGBC or BGTC; Si/SiO ₂ /HMDS; Au Vacuum | Light illumination effects in ambipolar FETs based on poly(3-hexylthiophene) and fullerene derivative composite films |
| 146 | $\mu_e=8.9 \times 10^{-3}$ (lin) $\mu_h=5.7 \times 10^{-3}$ (lin) | | | Spin-coating/CB(PCBM)&chloroform(P3HT) BGMC; Au/Ti/Si/SiO ₂ /PCBM/TiO _x /P3HT; Al Nitrogen | Multilayer bipolar field-effect transistors |
| 2009 | | | | | |
| 147 | 2.8×10^{-2} | | | Spin-coating/ chloroform BGTC; Si/SiO ₂ /PCBM/TiO _x ; Al Nitrogen | Enhanced Performance of Fullerene n-Channel Field-Effect Transistors with Titanium Sub-Oxide Injection Layer |

| | | | | | |
|-------------|---|---|---|---|---|
| 148 | 0.028 | | | Spin-coating/chloroform BGTC; Si/SiO ₂ /OTS; Ca/Al Nitrogen | Heteroanalogues of PCBM: N-Bridged Imino-PCBMs for Organic Field-Effect Transistors |
| 149 | 0.10-0.14 | 1×10 ⁴ ~2×10 ⁵ | 0.27~0.38 | Spin-coating/ chlorobenzene BGTC; Ti/Au/Si/SiO ₂ /HfO ₂ /BCB;Ca Nitrogen or air | Low-voltage solution-processed n-channel organic field-effect transistors with high-k HfO ₂ gate dielectrics grown by atomic layer deposition |
| 150 | 0.03 | | 0.1 | Spin-coating/ chlorobenzene BGTC; Al/AlO _x /PHDA; Al vacuum | Solution processed low-voltage organic transistors and complementary inverters |
| 151 | 10 ⁻⁵ (lin) | | 0-10 | Spin-coating/ chlorobenzene BGTC; Al/BCB; Al | Studies of charge transfer processes across donor-acceptor interface using a field effect transistor geometry |
| 152 | Ca: N ₂ :0.12 ; Air:No Au: N ₂ :0.08 ; Air:0.04 Ca/Au: N ₂ :0.12 ; Air:0.06 | N ₂ :5×10 ⁻⁵ ;Air: No N ₂ :2×10 ⁻⁵ ;Air: 1×10 ⁵ N ₂ :1×10 ⁻⁶ ;Air: 3×10 ⁵ | N ₂ :2.1;Air: No N ₂ :4.6;Air: 6.7 N ₂ :2.1;Air: 5.4 | Spin-coating/ chlorobenzene BGTC; Ti/Au/Si/SiO ₂ /BCB;Ca or Au or Ca/Au Nitrogen or air | Study of electrical performance and stability of solution-processed n-channel organic field-effect transistors |
| 153 | μ _e =5.8×10 ⁻² (140°C anneal) μ _h =7.2×10 ⁻² (140°C anneal) | | | Spin-coating BGTC; Si/SiO ₂ /PCBM&PTmT; Ag | Thermal annealing induced bicontinuous networks in bulk heterojunction solar cells and bipolar field-effect transistors |
| 2010 | | | | | |
| 154 | PCBM/HMDS-Au: 6.0×10 ⁻⁶ PCBM/HMDS-LiF: 1.5×10 ⁻³ PCBM/OTS-Au: 1.0×10 ⁻⁴ PCBM/OTS-LiF: 8.6×10 ⁻² PCBM/ODTS-LiF: 4.6×10 ⁻² PCBM/P3HT: μ _e =1.5×10 ⁻³ PCBM/P3HT: μ _h =3.3×10 ⁻⁵ | 2.3 1.3×10 ² 1.6 -9 1.6×10 ⁵ 1.5×10 ⁵ | 35 40 38 29 45~74 | Spin-coating/chloroform BGMC; Si/SiO ₂ /SAM/PCBM/(P3HT); LiF/Au Vacuum | Ambipolar Transport in Bilayer Organic Field-Effect Transistor Based on Poly(3-hexylthiophene) and Fullerene Derivatives |
| 155 | 0.13 | 6×10 ⁴ | 25 | Spin-coating/chloroform BGTC; Si/SiO ₂ /OTS; Au Nitrogen | High-Performance Solution-Processed n-Channel Organic Thin-Film Transistors Based on a Long Chain Alkyl-Substituted C ₆₀ Derivative |
| 156 | 0.125 | 4.67×10 ⁶ | 19.10 | Spin-coating/chlorobenzene BGTC; Si/SiO ₂ /BCB; Au Nitrogen | Use of a 1H-Benzimidazole Derivative as an n-Type Dopant and To Enable Air-Stable Solution-Processed n-Channel Organic Thin-Film Transistors |
| 2011 | | | | | |
| 157 | 0.09 | | 9.0 | Spin-coating/ chlorobenzene BGTC; Si/SiO ₂ /BCB; Au or Al Nitrogen, Vacuum or Air | Soluble fullerene derivatives: The effect of electronic structure on transistor performance and air stability |
| 2012 | | | | | |
| 158 | ~0.01(pure PCBM) | | | Spin-coating/ o-dichlorobenzene BGBC; Si/SiO ₂ /HMDS/PCBM&polymer; Au Nitrogen | Ambipolar charge transport in polymer:fullerene bulk heterojunctions for different polymer side-chains |
| 159 | 0.024~0.054 | 8.0×10 ⁵ ~2.2×10 ⁶ | 2.18~2.9 | Spin-coating/ chloroform BGTC; Si/SiO ₂ /OTS/PCBM; Al Nitrogen | Effects of direct solvent exposure on the nanoscale morphologies and electrical characteristics of PCBM-based transistors and photovoltaics |
| 160 | Ca/Al: 0.104 Au: 0.051 | 3.5×10 ⁶ 4.6×10 ⁶ | 8 12 | Spin-coating / chloroform or DCB BGTC; Si/SiO ₂ /BCB; Ca/Al or Au Nitrogen | Evaluation of structure–property relationships of solution-processable fullerene acceptors and their n-channel field-effect transistor performance |
| 161 | Highest: 0.044 | 4.1×10 ⁴ | -2.61 | Spin-coating / chloroform BGTC; ITO/PVP/PCBM/PEG; Al Nitrogen | Simple source/drain contact structure for solution-processed n -channel fullerene thin-film transistors |
| 162 | 3×10 ⁻³ | 10 ³ | 20 | Spin-coating/ chlorobenzene BGTC; Si/SiO ₂ /HMDS/PCBM&PB-PyDI; Al Vacuum | Synthesis and Characterization of a Pyromellitic Diimide-Based Polymer with C- and N-Main Chain Links: Matrix for Solution-Processable n-Channel Field-Effect Transistors |
| 2013 | | | | | |
| 163 | 2×10 ⁻⁸ ~3×10 ⁻⁸ | | | Spin-coating BG; Si /SiO ₂ /HMDS/PCBM; Au Nitrogen | Microstructure and Optoelectronic Properties of P3HT-b-P4VP/PCBM Blends: Impact of PCBM on the Copolymer Self-Assembly |
| 164 | ~0.04 | | | Spin-coating/ chloroform BGTC; Si/SiO ₂ /HMDS/PCBM; Al | Organic [6,6]-phenyl-C ₆₁ -butyric-acid-methyl-ester field effect transistors: Analysis of the contact properties by combined photoemission spectroscopy and electrical measurements |
| 165 | 8.70×10 ⁻² (FP doped)2.2×10 ⁻² | 6.7×10 ⁶ | 21 | Spin-coating/ chloroform BGTC; Si /SiO ₂ /BCB/PCBM; Ag Nitrogen | Solution - Processible Highly Conducting Fullerenes |
| 2014 | | | | | |
| 105 | 0.057 | 2×10 ³ | 15.7 | Spin-coating/CB BGTC; Si/SiO ₂ /BCB; Al Nitrogen | Comparative Study of the N-Type Doping Efficiency in Solution-processed Fullerenes and Fullerene Derivatives |
| 166 | Highest: 0.12 | ~10 ³ | 10.5 | Spin-coating/ chlorobenzene TGBC; Al/CYTOP/PCBM/Interlayer; Au Nitrogen | Simultaneous Enhancement of Electron Injection and Air Stability in N-Type Organic Field-Effect Transistors by Water-Soluble Polyfluorene Interlayers |
| 2015 | | | | | |

| | | | | | |
|-------------|--|---|-------------------|---|---|
| 167 | No DAE: $1 \times 10^{-2} \sim 4.41 \times 10^{-2}$ & DAE: $0.61 \times 10^{-2} \sim 3.80 \times 10^{-2}$ | 1.8×10^7 $3.5 \times 10^5 \sim 4.2 \times 10^5$ | | Spin-coating/ chlorobenzene BGBC; Si/SiO ₂ /HMDS/PCBM(&DAE); Au Nitrogen | Optically switchable transistors comprising a hybrid photochromic molecule/n-type organic active layer |
| 2016 | | | | | |
| 168 | 0.04 | | 11 | Spin-coating/DCB BGBC; Si/SiO ₂ ; Au Air | Synthesis of Fullerene Derivatives for the Application to Organic Photovoltaic Cell and n-Channel Organic Thin-Film Transistors |
| 169 | BCB: 9.3×10^{-2} b-PS(8K): 7.3×10^{-2} b-PS(108K): 1.0×10^{-1} | $>10^5$ $>10^5$ $>10^5$ | 5.1 8.5 6.6 | Spin-coating/ chlorobenzene BGTC; Si /SiO ₂ /BCB or b-PS/PCBM; Au Nitrogen | Use of a cross-linkable or monolayer-forming polymeric buffer layer on PCBM-based n-channel organic field-effect transistors |
| 2017 | | | | | |
| 170 | Pure: 0.027 With P3HT: 0.01 | | | Spin-coating/ toluene BGTC; Si/SiO ₂ /HMDS/PCBM&P3HT; Au Nitrogen | Balanced Ambipolar Organic Field-Effect Transistors by Polymer Preaggregation |
| 171 | 8×10^{-3} | | | Spin-coating/ chlorobenzene TGBC; tungsten/water/CYTOP; Au Nitrogen | Water-Gated n-Type Organic Field-Effect Transistors for Complementary Integrated Circuits Operating in an Aqueous Environment |

PC₇₁BM

| Ref. | Mobility ($\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$) | I_{on}/I_{off} | V_T (V) | Deposition Structure Measured in | Title |
|-------------|---|--|----------------------|---|--|
| 2005 | | | | | |
| 172 | $\mu_e=1 \times 10^{-3}$ $\mu_h=2 \times 10^{-5}$ | $\sim 10^4$ $\sim 10^4$ | | Drop cast/ chlorobenzene BGBC; Si/SiO ₂ /HMDS; /Au Vacuum | Solution processible organic transistors and circuits based on a C ₇₀ methanofullerene |
| 2008 | | | | | |
| 143 | 0.1 | | | Spin-coating/ chlorobenzene BGTC; ITO/BCB; Ca/Al Nitrogen | High mobility n-channel organic field-effect transistors based on soluble C ₆₀ and C ₇₀ fullerene derivatives |
| 2009 | | | | | |
| 147 | 2.2×10^{-2} | | | Spin-coating/ chloroform BGTC; Si /SiO ₂ /PC ₇₁ BM/TiO _x ; Al Nitrogen | Enhanced Performance of Fullerene n-Channel Field-Effect Transistors with Titanium Sub-Oxide Injection Layer |
| 2012 | | | | | |
| 160 | Ca/Al: 0.066 Au: 0.018 | 2.6×10^6 1.1×10^6 | 8 22 | Spin-coating / chloroform or DCB BGTC; Si/SiO ₂ /BCB; Ca/Al or Au Nitrogen | Evaluation of structure–property relationships of solution-processible fullerene acceptors and their n-channel field-effect transistor performance |
| 2013 | | | | | |
| 173 | Pure PC ₇₁ BM: 0.016 | | | Spin-coating/CB BGTC; Si/SiO ₂ /HMDS/PC ₇₁ BM&donor; Al Nitrogen | Electron and hole mobility in solution-processed small molecule-fullerene blend: Dependence on the fullerene content |
| 2014 | | | | | |
| 105 | 0.046 | 1×10^3 | 19.9 | Spin-coating/CB BGTC; Si/SiO ₂ /BCB; Al Nitrogen | Comparative Study of the N-Type Doping Efficiency in Solution-processed Fullerenes and Fullerene Derivatives |
| 2016 | | | | | |
| 174 | $\mu_e=1.3 \times 10^{-3}$ $\mu_h=2.7 \times 10^{-3}$ | | | Spin-coating TGBC; tungsten/electrolyte/PC ₇₁ BM&polymer; Au Air | An organic water-gated ambipolar transistor with a bulk heterojunction active layer for stable and tunable photodetection |
| 175 | HMDS 7°C: 3.47×10^{-3} HMDS 25°C: 2.5×10^{-3} HMDS60°C: 4.27×10^{-3} | 10^4 10^3 10^6 | 22.2 10.0 14.8 | Drop cast/ chlorobenzene BGBC; Si/SiO ₂ /HMDS; Au Nitrogen | PC ₇₀ BM n-type thin film transistors: Influence of HMDS deposition temperature on the devices properties |

2. LUMO converting

The LUMO of IC₇₀MA and IC₇₀BA comes from Yongfang Li group report ¹⁷⁶ (CV was measured in a 0.1 mol/L tetrabutylammonium hexafluorophosphate (Bu₄NPF₆) in o-dichlorobenzene/acetonitrile (5:1) solution). The original data are LUMO₁(IC₇₀MA)=−3.85 eV, LUMO₁(IC₇₀BA)=−3.72 eV , LUMO₁(PC₆₀BM)=−3.91 eV . In Hojeong Yu et al. report ¹⁷⁷ (CV was measured in a 0.1 mol/L tetrabutylammonium tetrafluoroborate (NBu₄BF₄) in o-dichlorobenzene solution), LUMO₂(PC₆₀BM)=−3.85 eV. In order to compare, we choose PC₆₀BM as standard to calculate as follow:

$$LUMO(IC_{70}MA) = -3.85eV \times \frac{-3.85}{-3.91} = -3.79eV$$

$$LUMO(IC_{70}BA) = -3.72eV \times \frac{-3.85}{-3.91} = -3.66eV$$

3. Converting unit of solubility

Solubility of C₆₀, PCBM and C₇₀, PC₇₁BM in various solvents

| Solvent | Solubility | | Ref. | Solubility | | Ref. |
|-------------------|-----------------|--------------------------|----------------|---------------------|--------------------------|----------------|
| | C ₆₀ | | | PCBM | | |
| | mg/ml | 10 ⁻³ mmol/ml | | mg/ml | 10 ⁻³ mmol/ml | |
| chloroform | 0.16(r.t.) | 0.22 | ¹⁷⁸ | 28.8(25°C) | 31.6 | |
| chlorobenzene | 5.7(r.t.) | 7.9 | ¹⁸⁰ | 59.5(25°C) | 65.4 | ¹⁷⁹ |
| o-dichlorobenzene | 24.6(r.t.) | 34.2 | | 42.1(25°C) | 58.4 | |
| | C ₇₀ | | | PC ₇₁ BM | | |
| o-dichlorobenzene | 36.2(303K) | 43.1 | ¹⁸¹ | 225.2(25°C) | 218.6 | ¹⁸² |

The date in the left is original data that come from the corresponding references, in order to give an intuitive comprehension and compare, the unit is converted from mg/ml to 10⁻³mmol/ml. The more solubility can be see references: ¹⁸³⁻¹⁸⁵.

1. J. Paloheimo, H. Isotalo, J. Kastner and H. Kuzmany, *Synthetic Metals*, 1993, **56**, 3185-3190.
2. J. Kastner, J. Paloheimo and H. Kuzmany, Berlin, Heidelberg, 1993, **113**, 512-515.
3. H. Katsunori, F. Shigeo, F. Shizuo and F. Shigeo, *Japanese Journal of Applied Physics*, 1993, **32**, L1070.
4. R. C. Haddon, A. S. Perel, R. C. Morris, T. T. M. Palstra, A. F. Hebard and R. M. Fleming, *Applied Physics Letters*, 1995, **67**, 121.
5. A. Dodabalapur, H. E. Katz, L. Torsi and R. C. Haddon, *Science*, 1995, **269**, 1560.
6. A. Dodabalapur, H. E. Katz, L. Torsi and R. C. Haddon, *Applied Physics Letters*, 1996, **68**, 1108-1110.
7. Keiichi Kaneto, Kazuya Yamanaka, Kouichi Rikitake, Takahiro Akiyama and Wataru Takashima, *Japanese Journal of Applied Physics*, 1996, **35**, 1802.
8. C. P. Jarrett, K. Pichler, R. Newbould and R. H. Friend, *Synthetic Metals*, 1996, **77**, 35-38.
9. K. Horiuchi, K. Nakada, S. Uchino, S. Hashii, A. Hashimoto, N. Aoki, Y. Ochiai and M. Shimizu, *Applied Physics Letters*, 2002, **81**, 1911-1912.
10. S. Kobayashi, T. Takenobu, S. Mori, A. Fujiwara and Y. Iwasa, *Science and Technology of Advanced Materials*, 2003, **4**, 371-375.
11. S. Kobayashi, T. Takenobu, S. Mori, A. Fujiwara and Y. Iwasa, *Applied Physics Letters*, 2003, **82**, 4581.
12. T. Shimada, T. Suetsugu, T. Miyadera, Y. Yamamoto, A. Koma, K. Saiki and K. Kudo, *Applied Physics Letters*, 2004, **84**, 2439-2441.
13. K. Tanigaki, R. Kumashiro and H. Ohashi, *Chemical Physics Letters*, 2004, **400**, 235-238.
14. M. Chikamatsu, S. Nagamatsu, T. Taima, Y. Yoshida, N. Sakai, H. Yokokawa, K. Saito and K. Yase, *Applied Physics Letters*, 2004, **85**, 2396-2398.
15. E. Kuwahara, Y. Kubozono, T. Hosokawa, T. Nagano, K. Masunari and A. Fujiwara, *Applied Physics Letters*, 2004, **85**, 4765-4767.
16. H. Ohashi, K. Tanigaki, R. Kumashiro and S. Sugihara, *Applied Physics Letters*, 2004, **84**, 520-522.
17. T. Nishikawa, S.-I. Kobayashi, T. Nakanowatari, T. Mitani, T. Shimoda, Y. Kubozono, G. Yamamoto, H. Ishii, M. Niwano and Y. Iwasa, *Journal of Applied Physics*, 2005, **97**, 104509.
18. S. J. Kang, Y. Yi, C. Y. Kim, K. Cho, J. H. Seo, M. Noh, K. Jeong, K.-H. Yoo and C. N. Whang, *Applied Physics Letters*, 2005, **87**, 233502.
19. E. Kuwahara, H. Kusai, T. Nagano, T. Takayanagi and Y. Kubozono, *Chemical Physics Letters*, 2005, **413**, 379-383.
20. Y. Kubozono, T. Nagano, Y. Haruyama, E. Kuwahara, T. Takayanagi, K. Ochi and A. Fujiwara, *Applied Physics Letters*, 2005, **87**,

21. Y. Hayashi, H. Kanamori, I. Yamada, A. Takasu, S. Takagi and K. Kaneko, *Applied Physics Letters*, 2005, **86**, 052104.
22. J. N. Haddock, X. Zhang, B. Domercq and B. Kippelen, *Organic Electronics*, 2005, **6**, 182-187.
23. T. B. Singh, N. Marjanović, G. J. Matt, S. Günes, N. S. Sariciftci, A. Montaigne Ramil, A. Andreev, H. Sitter, R. Schwödiauer and S. Bauer, *Organic Electronics*, 2005, **6**, 105-110.
24. A. Tapponnier, I. Biaggio and P. Günter, *Applied Physics Letters*, 2005, **86**, 112114.
25. F. Li, M. Pfeiffer, A. Werner, K. Harada, K. Leo, N. Hayashi, K. Seki, X. Liu and X.-D. Dang, *Journal of Applied Physics*, 2006, **100**, 023716.
26. T. Miyadera, M. Nakayama and K. Saiki, *Applied Physics Letters*, 2006, **89**, 172117.
27. S. Wang, K. Kanai, Y. Ouchi and K. Seki, *Organic Electronics*, 2006, **7**, 457-464.
28. K. Ogawa, T. Kato, A. Ikegami, H. Tsuji, N. Aoki, Y. Ochiai and J. P. Bird, *Applied Physics Letters*, 2006, **88**, 112109.
29. T. D. Anthopoulos, B. Singh, N. Marjanovic, N. S. Sariciftci, A. Montaigne Ramil, H. Sitter, M. Cölle and D. M. de Leeuw, *Applied Physics Letters*, 2006, **89**, 213504.
30. K. Itaka, M. Yamashiro, J. Yamaguchi, M. Haemori, S. Yaginuma, Y. Matsumoto, M. Kondo and H. Koinuma, *Advanced materials*, 2006, **18**, 1713-1716.
31. A. M. Ramil, T. B. Singh, N. T. Haber, N. Marjanović, S. Günes, A. Andreev, G. J. Matt, R. Resel, H. Sitter and S. Sariciftci, *Journal of Crystal Growth*, 2006, **288**, 123-127.
32. Y. Matsuoka, K. Uno, N. Takahashi, A. Maeda, N. Inami, E. Shikoh, Y. Yamamoto, H. Hori and A. Fujiwara, *Applied Physics Letters*, 2006, **89**, 173510.
33. T. P. I. Saragi and J. Salbeck, *Applied Physics Letters*, 2006, **89**, 253516.
34. K. Ochi, T. Nagano, T. Ohta, R. Nouchi, Y. Kubozono, Y. Matsuoka, E. Shikoh and A. Fujiwara, *Applied Physics Letters*, 2006, **89**, 083511.
35. A. L. Briseno, C. Reese, F. Wudl, M. M. Ling, M. E. Roberts, R. J. Tseng, S. Liu, S. C. B. Mannsfeld, Y. Yang and Z. Bao, *Nature*, 2006, **444**, 913-917.
36. G. J. Matt, T. B. Singh, N. S. Sariciftci, A. M. Ramil and H. Sitter, *Applied Physics Letters*, 2006, **88**, 263516.
37. A. Opitz, M. Bronner and W. Brütting, *Journal of Applied Physics*, 2007, **101**, 063709.
38. A. Dzwilewski, T. Wågberg and L. Edman, *Physical Review B*, 2007, **75**, 075203.
39. T. Birendra Singh, H. Yang, B. Plochberger, L. Yang, H. Sitter, H. Neugebauer and N. S. Sariciftci, *physica status solidi (b)*, 2007, **244**, 3845-3848.
40. T. B. Singh, N. S. Sariciftci, H. Yang, L. Yang, B. Plochberger and H. Sitter, *Applied Physics Letters*, 2007, **90**, 213512.
41. T. Matsushima, M. Yahiro and C. Adachi, *Applied Physics Letters*, 2007, **91**, 103505.
42. J. Puigdollers, C. Voz, S. Cheylan, A. Orpella, M. Vetter and R. Alcubilla, *Thin Solid Films*, 2007, **515**, 7667-7670.
43. J. H. Na, M. Kitamura and Y. Arakawa, *Applied Physics Letters*, 2007, **91**, 193501.
44. X. H. Zhang, B. Domercq and B. Kippelen, *Applied Physics Letters*, 2007, **91**, 092114.
45. M. Kitamura, Y. Kuzumoto, M. Kamura, S. Aomori and Y. Arakawa, *Applied Physics Letters*, 2007, **91**, 183514.
46. J. Zhou, F. Zhang, L. Lan, S. Wen and J. Peng, *Applied Physics Letters*, 2007, **91**, 253507.
47. M. Kitamura and Y. Arakawa, *Applied Physics Letters*, 2007, **91**, 053505.
48. Takayuki Nagano, Michiko Tsutsui, Ryo Nouchi, Naoko Kawasaki, Yohei Ohta, Yoshihiro Kubozono, Nobuya Takahashi and A. Fujiwara, *Journal of Physical Chemistry C*, 2007, **111**, 7211-7217.
49. N. Takahashi, A. Maeda, K. Uno, E. Shikoh, Y. Yamamoto, H. Hori, Y. Kubozono and A. Fujiwara, *Applied Physics Letters*, 2007, **90**, 083503.
50. M. Bronner, A. Opitz and W. Brütting, *Physica Status Solidi*, 2008, **205**, 549-563.
51. O. Andreas, K. Michael, B. Markus, W. Julia and B. Wolfgang, *New Journal of Physics*, 2008, **10**, 065006.
52. M. Kitamura, S. Aomori, J. H. Na and Y. Arakawa, *Applied Physics Letters*, 2008, **93**, 033313.
53. Y. Kubozono, S. Haas, W. L. Kalb, P. Joris, F. Meng, A. Fujiwara and B. Batlogg, *Applied Physics Letters*, 2008, **93**, 033316.
54. X.-H. Zhang and B. Kippelen, *Journal of Applied Physics*, 2008, **104**, 104504.
55. X. H. Zhang and B. Kippelen, *Applied Physics Letters*, 2008, **93**, 133305.

56. O. Acton, G. Ting, H. Ma and A. K.-Y. Jen, *Applied Physics Letters*, 2008, **93**, 083302.
57. M. Kitamura, Y. Kuzumoto, M. Kamura, S. Aomori, J. H. Na and Y. Arakawa, *physica status solidi (c)*, 2008, **5**, 3181-3183.
58. A. Konishi, E. Shikoh, Y. Kubozono and A. Fujiwara, *Applied Physics Letters*, 2008, **92**, 173302.
59. Y. Ohta, Y. Kubozono and A. Fujiwara, *Applied Physics Letters*, 2008, **92**, 173306.
60. M. Irimia-Vladu, N. Marjanovic, A. Vlad, A. M. Ramil, G. Hernandez-Sosa, R. Schwoödauer, S. Bauer and N. S. Sariciftci, *Advanced Materials*, 2008, **20**, 3887-3892.
61. H. Yan, T. Kagata and H. Okuzaki, *Applied Physics Letters*, 2009, **94**, 023305.
62. M. Kitamura and Y. Arakawa, *Applied Physics Letters*, 2009, **95**, 023503.
63. Y. Ito, A. A. Virkar, S. Mannsfeld, J. H. Oh, M. Toney, J. Locklin and Z. Bao, *Journal of the American Chemical Society*, 2009, **131**, 9396-9404.
64. M. Ullah, D. M. Taylor, R. Schwödauer, H. Sitter, S. Bauer, N. S. Sariciftci and T. B. Singh, *Journal of Applied Physics*, 2009, **106**, 114505.
65. C. F. Sung, D. Kekuda, L. F. Chu, Y. Z. Lee, F. C. Chen, M. C. Wu and C. W. Chu, *Advanced materials*, 2009, **21**, 4845-4849.
66. Y. Ochiai, K. Ogawa, N. Aoki and J. P. Bird, *Journal of Physics: Conference Series*, 2009, **159**, 012004.
67. M. Irimia-Vladu, N. Marjanovic, M. Bodea, G. Hernandez-Sosa, A. M. Ramil, R. Schwödauer, S. Bauer, N. S. Sariciftci and F. Nüesch, *Organic Electronics*, 2009, **10**, 408-415.
68. M. Kitamura, Y. Kuzumoto, S. Aomori, M. Kamura, J. H. Na and Y. Arakawa, *Applied Physics Letters*, 2009, **94**, 83310.
69. M. Ullah, I. I. Fishchuk, A. Kadashchuk, P. Stadler, A. Pivrikas, C. Simbrunner, V. N. Poroshin, N. S. Sariciftci and H. Sitter, *Applied Physics Letters*, 2010, **96**, 213306.
70. D. Tatsuya, K. Kyouhei, C. Yasuto, T. Hajime, U. Misaki, C. Shih-Ren, A. Nobuyuki, B. Jonathan Paul and O. Yuichi, *Japanese Journal of Applied Physics*, 2010, **49**, 04DN12.
71. M. Irimia-Vladu, P. A. Troshin, M. Reisinger, G. Schwabegger, M. Ullah, R. Schwoedauer, A. Mumyatov, M. Bodea, J. W. Fergus and V. F. Razumov, *Organic Electronics*, 2010, **11**, 1974-1990.
72. P. Cosseddu and A. Bonfiglio, *Applied Physics Letters*, 2010, **97**, 203305.
73. Z. Jianlin and N. Qiaoli, *Chinese Physics B*, 2010, **19**, 77305.
74. M. Ullah, A. Pivrikas, Fishchuk, II, A. Kadashchuk, P. Stadler, C. Simbrunner, N. S. Sariciftci and H. Sitter, *Synth Met*, 2011, **161**, 1987-1990.
75. H. Zheng, X. Cheng, H. Tian and G. Zhao, *Journal of Semiconductors*, 2011, **32**, 094005.
76. G. Schwabegger, M. Ullah, M. Irimia-Vladu, M. Baumgartner, Y. Kanbur, R. Ahmed, P. Stadler, S. Bauer, N. S. Sariciftci and H. Sitter, *Synthetic Metals*, 2011, **161**, 2058-2062.
77. C. Xinyang, Y. Junsheng, Z. Jianlin, Y. Xinge and J. Yadong, *Japanese Journal of Applied Physics*, 2011, **50**, 124203.
78. K. Woogun, K. Masatoshi and A. Yasuhiko, *Applied Physics Express*, 2011, **4**, 121602.
79. K. Takuji, O. Chikako, S. Masato and A. Chihaya, *Japanese Journal of Applied Physics*, 2011, **50**, 050202.
80. P. Wei, T. Menke, B. D. Naab, K. Leo, M. Riede and Z. Bao, *Journal of the American Chemical Society*, 2012, **134**, 3999-4002.
81. Z. Jian-Lin, Y. Jun-Sheng, Y. Xin-Ge and C. Xin-Yang, *Chinese Physics B*, 2012, **21**, 027305.
82. H. Kleemann, A. A. Zakhidov, M. Anderson, T. Menke, K. Leo and B. Lüssem, *Organic Electronics*, 2012, **13**, 506-513.
83. Y. Chung, O. Johnson, M. Deal, Y. Nishi, B. Murmann and Z. Bao, *Applied Physics Letters*, 2012, **101**, 063304.
84. T. Wakahara, P. D'Angelo, K. I. Miyazawa, Y. Nemoto, O. Ito, N. Tanigaki, D. D. C. Bradley and T. D. Anthopoulos, *Journal of the American Chemical Society*, 2012, **134**, 7204.
85. H. Li, B. C. Tee, J. J. Cha, Y. Cui, J. W. Chung, S. Y. Lee and Z. Bao, *Journal of the American Chemical Society*, 2012, **134**, 2760-2765.
86. C. Larsen, H. R. Barzegar, F. Nitze, T. Wagberg and L. Edman, *Nanotechnology*, 2012, **23**, 344015.
87. T. T. Dao, T. Matsushima and H. Murata, *Organic Electronics*, 2012, **13**, 2709-2715.
88. S. Olthof, S. Singh, S. K. Mohapatra, S. Barlow, S. R. Marder, B. Kippelen and A. Kahn, *Applied Physics Letters*, 2012, **101**, 253303.
89. K. Woogun, K. Masatoshi, I. Tetsuji and A. Yasuhiko, *Japanese Journal of Applied Physics*, 2012, **51**, 11PD06.
90. K. Woogun, K. Masatoshi, K. Masakazu, A. Shigeru and A. Yasuhiko, *Japanese Journal of Applied Physics*, 2012, **51**, 02BK10.

91. A. Nigam, G. Schwabegger, M. Ullah, R. Ahmed, I. I. Fishchuk, A. Kadashchuk, C. Simbrunner, H. Sitter, M. Premaratne and V. R. Rao, *Applied Physics Letters*, 2012, **101**, 083305.
92. A. Petritz, A. Wolfberger, A. Fian, M. Irimia-Vladu, A. Haase, H. Gold, T. Rothländer, T. Griesser and B. Stadlober, *Applied Physics Letters*, 2013, **103**, 153303.
93. S. J. Noever, S. Fischer and B. Nickel, *Advanced materials*, 2013, **25**, 2147-2151.
94. Q. Li, X. Yu, W. Shi and J. Yu, *Synthetic Metals*, 2013, **163**, 57-60.
95. X. Liang, X. Cheng, B. Du, X. Bai and J. Fan, *Journal of Semiconductors*, 2013, **34**, 084002.
96. K. Ahn, J. Beom Kim, H. Park, H. Kim, M. Hyung Lee, B. Joon Kim, J. Ho Cho, M. Sung Kang and D. Ryeol Lee, *Applied Physics Letters*, 2013, **102**, 043306.
97. J. Zhang, J. Tan, Z. Ma, W. Xu, G. Zhao, H. Geng, C. Di, W. Hu, Z. Shuai, K. Singh and D. Zhu, *Journal of the American Chemical Society*, 2013, **135**, 558-561.
98. W. Kang, M. Kitamura and Y. Arakawa, *Organic Electronics*, 2013, **14**, 644-648.
99. D. Chen, B. Yao, G. Fan, W. Lv, P. Gao, M. Zhou and Y. Peng, *Applied Physics Letters*, 2013, **102**, 163303.
100. L. Qian Khor and K. Yew Cheong, *ECS Journal of Solid State Science and Technology*, 2013, **2**, P440-P444.
101. S. Singh, S. K. Mohapatra, A. Sharma, C. Fuentes-Hernandez, S. Barlow, S. R. Marder and B. Kippelen, *Applied Physics Letters*, 2013, **102**, 153303.
102. H. R. Barzegar, C. Larsen, L. Edman and T. Wågberg, *Particle & Particle Systems Characterization*, 2013, **30**, 715-720.
103. L.-S. Tsai, J.-C. Hwang, C.-Y. Lee, Y.-T. Lin, C.-L. Tsai, T.-H. Chang, Y.-L. Chueh and H.-F. Meng, *Applied Physics Letters*, 2013, **103**, 233304.
104. R. Ahmed, C. Simbrunner, G. Schwabegger, M. A. Baig and H. Sitter, *Synthetic Metals*, 2014, **188**, 136-139.
105. S. Rossbauer, C. Müller and T. D. Anthopoulos, *Advanced Functional Materials*, 2014, **24**, 7116-7124.
106. R. Ahmed, A. Kadashchuk, C. Simbrunner, G. Schwabegger, M. A. Baig and H. Sitter, *ACS applied materials & interfaces*, 2014, **6**, 15148-15153.
107. Y. Li, S. Chen, Q. Liu, Y. Li, Y. Shi, X. Wang, J. Ma and Z. Hu, *The Journal of Physical Chemistry C*, 2014, **118**, 14218-14226.
108. J.-P. Yang, Q.-J. Sun, K. Yonezawa, A. Hinderhofer, A. Gerlach, K. Broch, F. Bussolotti, X. Gao, Y. Li, J. Tang, F. Schreiber, N. Ueno, S.-D. Wang and S. Kera, *Organic Electronics*, 2014, **15**, 2749-2755.
109. I. I. Fishchuk, A. Kadashchuk, S. V. Novikov, M. Ullah, J. Genoe, N. S. Sariciftci, H. Sitter and H. Bässler, *Molecular Crystals & Liquid Crystals*, 2014, **589**, 18-28.
110. H. Li, C. Fan, M. Vosgueritchian, B. C. K. Tee and H. Chen, *Journal of Materials Chemistry C*, 2014, **2**, 3617-3624.
111. D. Lili, L. Xiao, W. Zhanwei, Z. Jianping, S. Lei, L. Wenli, L. Yao, Z. Feiyu, Z. Junkang, R. Qiang, H. Fobao, X. Hongquan and P. Yingquan, *Journal of Physics D: Applied Physics*, 2015, **48**, 405105.
112. K. Y. Wu, T. Y. Wu, S. T. Chang, H. Chain - Shu and C. L. Wang, *Advanced materials*, 2015, **27**, 187-194.
113. L. Yao, L. Wenli, L. Xiao, S. Lei, Z. Maoqing, Z. Jianping, Z. Feiyu, Z. Junkang and P. Yingquan, *EPL (Europhysics Letters)*, 2015, **110**, 17006.
114. Y. J. Jeong, D. Yun, J. Jang, S. Park, T. K. An, L. H. Kim, S. H. Kim and C. E. Park, *Physical Chemistry Chemical Physics*, 2015, **17**, 6635-6643.
115. L. Sun, J. Zhang, F. Zhao, X. Luo, W. Lv, Y. li, Q. Ren, Z. Wen, Y. Peng and X. Liu, *Nanotechnology*, 2015, **26**, 185501.
116. M. Robin, M. Harnois, Y. Molard and E. Jacques, *Organic Electronics*, 2016, **39**, 214-221.
117. A. Nawaz, C. De Col, I. A. Hümmelgen, A. Nawaz, C. De Col and I. A. Hümmelgen, *Mat Res*, 2016, **19**, 1201.
118. Q. Li, J. Wu, R. Wu, Y. Liu, H. Chen, F. Huang and H. Li, *Science China Chemistry*, 2017, **60**, 490-496.
119. F. Huang, Y. Li, H. Xia, J. Zhang, K. Xu, Y. Peng and G. Liu, *Carbon*, 2017, **118**, 666-674.
120. R. Haddon, *Journal of the American Chemical Society*, 1996, **118**, 3041-3042.
121. C. Waldauf, P. Schilinsky, M. Perisutti, J. Hauch and C. J. Brabec, *Advanced materials*, 2003, **15**, 2084-2088.
122. T. D. Anthopoulos, C. Tanase, S. Setayesh, E. J. Meijer, J. C. Hummelen, P. W. M. Blom and D. M. De Leeuw, *Advanced materials*, 2004, **16**, 2174-2179.
123. T. D. Anthopoulos, D. M. d. Leeuw, E. Cantatore, S. Setayesh, E. J. Meijer, C. Tanase, J. C. Hummelen and P. W. M. Blom, *Applied Physics Letters*, 2004, **85**, 4205-4207.

124. T. B. Singh, N. Marjanovic, G. J. Matt, N. S. Sariciftci, R. Schwodiauer and S. Bauer, *Applied Physics Letters*, 2004, **85**, 5409-5411.
125. T. D. Anthopoulos, D. M. De Leeuw, E. Cantatore, S. Setayesh, E. J. Meijer, C. Tanase, J. C. Hummelen and P. W. M. Blom, *Applied Physics Letters*, 2004, **85**, 4205-4207.
126. T.-W. Lee, Y. Byun, B.-W. Koo, I.-N. Kang, Y.-Y. Lyu, C. H. Lee, L. Pu and S. Y. Lee, *Advanced materials*, 2005, **17**, 2180-2183.
127. T. B. Singh, S. Günes, N. Marjanović, N. S. Sariciftci and R. Menon, *Journal of Applied Physics*, 2005, **97**, 114508.
128. T. B. Singh, N. Marjanovic, P. Stadler, M. Auinger, G. J. Matt, S. Gunes, N. S. Sariciftci, R. Schwodiauer and S. Bauer, *Journal of Applied Physics*, 2005, **97**, 83714.
129. J. Nakamura, K. Murata and K. Takahashi, *Applied Physics Letters*, 2005, **87**, 3693.
130. E. Von Hauff, V. Dyakonov and J. Parisi, *Solar Energy Materials and Solar Cells*, 2005, **87**, 149-156.
131. S. Cho, J. Yuen, J. Y. Kim, K. Lee and A. J. Heeger, *Applied Physics Letters*, 2006, **89**, 153505.
132. E. von Hauff, J. Parisi and V. Dyakonov, *Thin Solid Films*, 2006, **511-512**, 506-511.
133. E. v. Hauff, J. Parisi and V. Dyakonov, *Journal of Applied Physics*, 2006, **100**, 073713.
134. E. v. Hauff, J. Parisi and V. Dyakonov, *Journal of Applied Physics*, 2006, **100**, 043702.
135. N. Marjanović, T. B. Singh, G. Dennler, S. Günes, H. Neugebauer, N. S. Sariciftci, R. Schwödiauer and S. Bauer, *Organic Electronics*, 2006, **7**, 188-194.
136. M. Shibao, T. Morita, W. Takashima and K. Kaneto, *Japanese Journal of Applied Physics*, 2007, **46**, L123-L125.
137. K. Kaneto, M. Yano, M. Shibao, T. Morita and W. Takashima, *Japanese Journal of Applied Physics*, 2007, **46**, 1736-1738.
138. T. D. Anthopoulos, *Applied Physics Letters*, 2007, **91**, 113513.
139. M. Morana, P. Koers, C. Waldauf, M. Koppe, D. Muehlbacher, P. Denk, M. Scharber, D. Waller and C. Brabec, *Advanced Functional Materials*, 2007, **17**, 3274-3283.
140. W. Takashima, T. Murasaki, S. Nagamatsu, T. Morita and K. Kaneto, *Applied Physics Letters*, 2007, **91**, 071905.
141. S. Tiwari, E. Namdas, V. R. Rao, D. Fichou and S. Mhaisalkar, *Electron Device Letters, IEEE*, 2007, **28**, 880-883.
142. M. Chikamatsu, A. Itakura, Y. Yoshida, R. Azumi and K. Yase, *Chemistry of Materials*, 2008, **20**, 7365-7367.
143. P. H. Wöbkenberg, D. D. C. Bradley, D. Kronholm, J. C. Hummelen, D. M. de Leeuw, M. Cölle and T. D. Anthopoulos, *Synthetic Metals*, 2008, **158**, 468-472.
144. C.-W. Chu, C.-F. Sung, Y.-Z. Lee and K. Cheng, *Organic Electronics*, 2008, **9**, 262-266.
145. M. Shibao, T. Morita, W. Takashima and K. Kaneto, *Thin Solid Films*, 2008, **516**, 2607-2610.
146. S. Cho, J. Yuen, J. Y. Kim, K. Lee, A. J. Heeger and S. Lee, *Applied Physics Letters*, 2008, **92**, 063505.
147. S. Cho, J. H. Seo, K. Lee and A. J. Heeger, *Advanced Functional Materials*, 2009, **19**, 1459-1464.
148. C. Yang, S. Cho, A. J. Heeger and F. Wudl, *Angewandte Chemie*, 2009, **48**, 1592-1595.
149. S. P. Tiwari, X. H. Zhang, W. J. P. Jr and B. Kippelen, *Applied Physics Letters*, 2009, **95**, 313.
150. J. M. Ball, P. H. Wöbkenberg, F. Colléaux, M. Heeney, J. E. Anthony, I. McCulloch, D. D. C. Bradley and T. D. Anthopoulos, *Applied Physics Letters*, 2009, **95**, 103310.
151. M. Rao and K. S. Narayan, *Applied Physics Letters*, 2009, **95**, 183306.
152. S. P. Tiwari, X. H. Zhang, W. J. Potsavage and B. Kippelen, *Journal of Applied Physics*, 2009, **106**, 253.
153. H. Kong, J. S. Moon, N. S. Cho, I. H. Jung, M.-J. Park, J.-H. Park, S. Cho and H.-K. Shim, *Applied Physics Letters*, 2009, **95**, 173301.
154. M. Takeomi, S. Vipul, O. Shinya, N. Shuichi, T. Wataru, H. Shuzi and K. Keiichi, *Japanese Journal of Applied Physics*, 2010, **49**, 041601.
155. Y. Horii, K. Sakaguchi, M. Chikamatsu, R. Azumi, K. Yase, M. Kitagawa and H. Konishi, *Applied Physics Express*, 2010, **3**, 101601.
156. P. Wei, J. H. Oh, G. Dong and Z. Bao, *Journal of the American Chemical Society*, 2010, **132**, 8852-8853.
157. J. M. Ball, R. K. M. Bouwer, F. B. Kooistra, J. M. Frost, Y. Qi, E. B. Domingo, J. Smith, D. M. de Leeuw, J. C. Hummelen, J. Nelson, A. Kahn, N. Stingelin, D. D. C. Bradley and T. D. Anthopoulos, *Journal of Applied Physics*, 2011, **110**, 014506.
158. S. Fall, L. Biniek, N. Leclerc, P. Lévéque and T. Heiser, *Applied Physics Letters*, 2012, **101**, 123301.
159. S. Nam, J. Jang, H. Cha, J. Hwang, T. K. An, S. Park and C. E. Park, *Journal of Materials Chemistry*, 2012, **22**, 5543.

160. C.-Z. Li, C.-C. Chueh, H.-L. Yip, J. Zou, W.-C. Chen and A. K. Y. Jen, *Journal of Materials Chemistry*, 2012, **22**, 14976.
161. F. C. Chen, T. H. Tsai and S. C. Chien, *Organic Electronics*, 2012, **13**, 599-603.
162. S. Kola, N. J. Tremblay, M. L. Yeh, H. E. Katz, S. B. Kirschner and D. H. Reich, *Acs Macro Letters*, 2012, **2012**, 136-140.
163. V. Gernigon, P. Lévêque, F. Richard, N. Leclerc, C. Brochon, C. H. Braun, S. Ludwigs, D. V. Anokhin, D. A. Ivanov, G. Hadzioannou and T. Heiser, *Macromolecules*, 2013, **46**, 8824-8831.
164. S. Scheinert, M. Grobosch, J. Sprogies, I. Hörselmann, M. Knupfer and G. Paasch, *Journal of Applied Physics*, 2013, **113**, 174504.
165. C. Z. Li, C. C. Chueh, H. L. Yip, F. Ding, X. Li and A. K. Jen, *Advanced materials*, 2013, **25**, 2457-2461.
166. J. Kim, D. Khim, R. Kang, S. H. Lee, K. J. Baeg, M. Kang, Y. Y. Noh and D. Y. Kim, *ACS applied materials & interfaces*, 2014, **6**, 8108.
167. K. Börjesson, M. Herder, L. Grubert, D. Duong, A. Salleo, S. Hecht, E. Orgiu and P. Samori, *J.mater.chem.c*, 2015, **3**, 4156-4161.
168. J.-M. Yun, D.-Y. Kim and Y.-Y. Noh, *Science of Advanced Materials*, 2016, **8**, 450-457.
169. K. Kim, H. J. Jeong and F. S. Kim, *Polymer Bulletin*, 2016, **73**, 2493-2500.
170. L. Janasz, A. Luczak, T. Marszalek, B. G. R. Dupont, J. Jung, J. Ulanski and W. Pisula, *ACS applied materials & interfaces*, 2017, **9**, 20696-20703.
171. R. Porrazzo, A. Luzio, S. Bellani, G. E. Bonacchini, Y. Y. Noh, Y. H. Kim, G. Lanzani, M. R. Antognazza and M. Caironi, *Acs Omega*, 2017, **2**, 1-10.
172. T. D. Anthopoulos, D. M. de Leeuw, E. Cantatore, P. van 't Hof, J. Alma and J. C. Hummelen, *Journal of Applied Physics*, 2005, **98**, 054503.
173. A. K. K. Kyaw, D. H. Wang, H.-R. Tseng, J. Zhang, G. C. Bazan and A. J. Heeger, *Applied Physics Letters*, 2013, **102**, 163308.
174. H. Xu, Q. Zhu, T. Wu, W. Chen, G. Zhou, J. Li, H. Zhang and N. Zhao, *Applied Physics Letters*, 2016, **109**, 213301.
175. M. R. Fiorillo, C. Diletto, P. Tassini, M. G. Maglione, E. Santoro, F. Villani, R. Liguori, P. Maddalena, A. Rubino and C. Minarini, *Materials Today Proceedings*, 2016, **3**, 720-726.
176. Y. He, G. Zhao, B. Peng and Y. Li, *Advanced Functional Materials*, 2010, **20**, 3383-3389.
177. H. Yu, H. H. Cho, C. H. Cho, K. H. Kim, D. Y. Kim, B. J. Kim and J. H. Oh, *ACS applied materials & interfaces*, 2013, **5**, 4865-4871.
178. R. S. Ruoff, D. S. Tse, R. Malhotra and D. C. Lorents, *Journal of Physical Chemistry*, 1993, **97**, 3379-3383.
179. F. Machui, S. Langner, X. Zhu, S. Abbott and C. J. Brabec, *Solar Energy Materials & Solar Cells*, 2012, **100**, 138-146.
180. W. A. Scrivens and J. M. Tour, *J. Chem. Soc., Chem. Commun.*, 1993, **0**, 1207-1209.
181. N. Sivaraman, R. Dhamodaran, I. Kaliappan, T. G. Srinivasan, P. R. P. V. Rao and C. K. C. Mathews, *Fullerene Science & Technology*, 1994, **2**, 233-246.
182. X. Guo, M. Zhang, W. Ma, S. Zhang, J. Hou and Y. Li, *Rsc Advances*, 2016, **6**, 51924-51931.
183. M. V. Korobov and A. L. Smith, *Fullerenes: Chemistry, physics, and technology*, 2000, 53-90.
184. V. N. Bezmel'nitsyn, A. V. Eletskii and M. Okun', *Physics-Uspekhi*, 1998, **41**, 1091-1114.
185. M. T. Beck and G. Mandi, *Fullerenes, Nanotubes, and Carbon Nanostructures*, 1997, **5**, 291-310.