



1 μm

4800 10.0kV x15.0k SE(M)

1 μ**m**

Supplementary Figure 1 Water dispersibility of carbon nanotubes (CNTs)





Cellulose: 6.3 wt%











100 μm

Supplementary Figure 2 Manufacturing process of CNT composite sheet (CNT sheet)



Supplementary Figure 3 Characteristics of CNT composite sheet (CNT sheet)



Supplementary Figure 4 Another manufacturing process of CNT/gel composite 1



Supplementary Figure 5

Another manufacturing process of CNT/gel composite 2



Supplementary Figure 6

Transmission electron microscopy (TEM) observation of CNT/rotaxane composite

Α

Procedure of colony forming assay



В

Control (polyethylene sheet)



Polyrotaxane-based gel Dilute concentration 20%

40% 60% 80% 100%

Polyrotaxane-based + CNT

Dilute concentration 20%



80%



Supplementary Figure 7

Colony-forming assay for cytotoxic evaluation

A Implanted electrodes



- B Condition for implanting and measurement
 - Each electrode was implanted by 6 pairs.



 Current with < 0.1 mA to avoid inflammation by continuous impedance measurements

Supplementary Figure 8 Implant assay for biocompatibility evaluation



Supplementary Figure 9

Magnified cross-sectional images of the graft pathology by staining a subcutaneous tissue after an electrode was explanted

4-week implantation



Supplementary Figure 10

Graft pathologies of the four different samples after implantation for four weeks



Supplementary Figure 11

Procedures of perfusion fixation, tissue-cleaning method, and evaluation using multiphoton microscopy



Supplementary Figure 12 In vivo implant tolerance of organic transistors





Supplementary Figure 13 1.2-µm-thick organic circuits



Capacitor

Selector TFT



Α



Supplementary Figure 14

Material profiling and cross-sectional image of organic transistors on a 1.2-µm-thick substrate using ultrahigh resolution scanning TEM (STEM) and EDX analysis system





Supplementary Figure 15

Material profiling and cross-sectional image of organic transistors on a 1.2-µm-thick substrate using ultrahigh resolution scanning TEM (STEM) and EDX analysis system

DNTT morphology on SAM gate dielectric with SiO₂ substrate



50 μm x 50 μm scan

DNTT morphology on SAM gate dielectric with 1-µm-thick PEN substrate



50 µm x 50 µm scan

0	100	200	(nm)
			Height

Supplementary Figure 16 Morphology of DNTT on SAM



Supplementary Figure 17

Optimization of the manufacturing process on a 1-µm-thick substrate



Supplementary Figure 18 Bending test



Design of organic transistor for higher amplifier gain

Supplementary Figure 20 Pseudo-CMOS inverter

Signal gain and frequency response of organic amplifiers

Ischemia-induced myocardial infarction

myocardial infarction

Supplementary Figure 22 Detection of ischemic state due to myocardial infarction

Supplementary Figure 23 Effects of the CNT-sheet/gel-composite electrode

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(I) Colony forming ability (%) : T

(II) Rate of relative colony formation (%) :

The number of colonies with processed well

The number of colonies with control well × 100

(III)		
(11)	IC ₅₀ value	Cell toxicity
	Negligible small change	None
	Higher than values from positive control B	Slight
	Values between positive control A and B	Moderate
	Lower than values from positive control A	Severe

В

Rate of relative colony formation and IC_{50} value of V79 cells processed with the following extraction liquids

	Concentration	Т	he number (of colonies	(count / wel	1)	Rate of relative	IC 50 value
Sample	(%)	Well 1	Well 2	Well 3	Mean	S.D.	colony formation (%)	(%)
Negative control	0	101	90	80	03	7	100.0	
(Fresh MO5 culture media)	0	101	90	07	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	1	100.0	-
Gel	20	92	97	92	94	3	101.1	
	40	97	92	82	90	8	96.8	
	60	97	87	80	88	9	94.6	
	80	94	104	91	96	7	103.2	
	100	75	86	89	83	7	89.2	
Gel + ionic liquid	20	98	101	90	96	6	103.2	
	40	73	91	91	85	10	91.4	
	60	82	97	98	92	9	98.9	— ^a
	80	86	80	89	85	5	91.4	
	100	83	80	82	82	2	88.2	
Gel + ionic liquid + CNT	20	99	88	74	87	13	93.5	
	40	83	86	78	82	4	88.2	
	60	72	92	91	85	11	91.4	
	80	95	84	89	89	6	95.7	
	100	97	77	76	83	12	89.2	

a: No decreases were observed on the rate of relative colony formation, as a result that IC₅₀ value cannot be estimated.

Rate of relative colony formation and IC_{50} value of V79 cells processed with the following control extraction liquids

	Concentration	Т	he number (ofcolonies	(count / we	1)	Rate of relative	IC 50 value
Control sample	(%)	Well 1	Well 2	Well 3	Mean	S.D.	colony formation (%)	(%)
Negative control (Fresh MO5 cultivate media)	0	101	90	89	93	7	100.0	-
Negative control	25	99	83	95	92	8	98.9	
(Defection of fine)	50	104	87	81	91	12	97.8	а
(Polyetnylene film)	75	106	83	89	93	12	100.0	_
	100	95	81	72	83	12	89.2	
Positive control A	0.5	42	53	49	48	6	51.6	
(Polyurethane film with	1	0	0	0	0	0	0.0	
0.1%-zinc	2	0	0	0	0	0	0.0	0.52
diethyldithiocarbamate)	5	0	0	0	0	0	0.0	
	10	0	0	0	0	0	0.0	
Positive control B	40	96	101	93	97	4	104.3	
(Polyurethane film with	50	92	97	88	92	5	98.9	
0.25%-zinc	60	53	51	46	50	4	53.8	61
dibutyldithiocarbamate)	80	1	0	0	0	1	0.0	
	100	0	0	0	0	0	0.0	

a: No decreases were observed on the rate of relative colony formation, as a result that IC 50 value cannot be estimated.

Supplementary Table 1

Colony-forming assay for cytotoxic evaluation

ROTA→ Polyrotaxane-based gel

CNT→ Multiwall carbon nanotube

IL→ Ionic liquid (N,N-Diethyl-N-methyl-N-(2-methoxyethyl) ammonium tetrafluoroborate) HEMA→ Poly-hydroxyethyl methacrylate gel

Appendix 1 One-week implantation

Cell tune/response		RC	TA			ROI	A-IL			ROTA	-CNT			ROTA-	IL-CNT	
con typortspuise	00101	00102	00103	00104	00101	00102	00103	00104	00101	00102	00103	00104	00101	00102	00103	00104
Polymorphoraclear cells	2	2	3	3	2	1	3	3	4	2	3	3	3	2	3	2
Lymphocytes	1	1	1	1	1	1	1	1	0	1	1	1	1	1	1	1
Plasma cells	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Macrophages	3	3	2	2	2	2	2	2	2	3	3	3	2	3	3	3
Giant cells	0	0	0	0	0	1	0	0	1	1	1	0	1	1	0	1
Necrosis	1	2	2	2	1	2	2	3	2	1	2	2	1	1	2	1
SUB TOTAL (X2)	14	16	16	16	12	14	16	18	18	16	20	18	16	16	18	16
Neovascularisation	1	1	0	0	0	0	0	0	0	0	1	0	1	0	0	0
Fibrosis	2	2	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Fatty infiltrate	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
SUB TOTAL	3	3	1	1	1	1	1	1	1	1	2	1	2	1	1	1
TOTAL	70				64				77				71			
AVERAGE	17.5				16.0				19.3				17.8			

Cell trae/response		HE	MA			HEMA	A-CNT			HEMA-	IL-CNT	1		Au				Negative control			
con typortsponso	00101	00102	00103	00104	00101	00102	00103	00104	00101	00102	00103	00104	00101	00102	00103	00104	00101	00102	00103	00104	
Polymorphoraclear cells	3	4	3	1	2	2	4	3	3	4	3	4	3	3	4	3	2	2	1	1	
Lymphocytes	1	1	1	0	1	1	1	1	1	1	1	1	0	1	0	0	0	1	1	1	
Plasma cells	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Macrophages	2	3	2	2	3	3	3	2	3	3	3	3	2	3	3	2	3	3	2	2	
Giant cells	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Necrosis	1	2	2	1	2	2	2	1	2	2	2	1	1	2	2	2	2	2	1	2	
SUB TOTAL (X2)	14	20	16	8	16	16	20	14	18	20	18	18	12	18	18	14	14	16	10	12	
Neovascularisation	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Fibrosis	1	1	1	2	1	1	1	1	1	1	1	1	1	2	1	1	0	1	0	1	
Fatty infiltrate	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
SUB TOTAL	1	1	1	2	2	1	1	1	1	1	1	1	1	2	1	1	0	1	0	1	
TOTAL		6	3			7	1			7	8			6	7			5	4		
AVERAGE		15	.8			17	.8			19	.5			16	.8			13	3.5		

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Appendix 2 4-week implantation

Cell type/resmose		RO	TA			ROT	A-IL			ROTA	-CNT			ROTA-IL-CNT				
con type to sponse	00105	00106	00107	00108	00105	00106	00107	00108	00105	00106	00107	00108	00105	00106	00107	00108		
Polymorphonuclear cells	1	2	1	2	1	0	1	1	0	2	0	2	1	1	0	0		
Lymphocytes	1	2	3	2	2	1	2	3	2	3	1	2	3	1	2	1		
Plasma cells	0	1	1	0	0	0	1	0	0	0	0	0	0	1	0	0		
Macrophages	3	3	4	2	3	3	3	3	3	3	3	3	3	2	3	3		
Giant cells	1	0	0	0	0	1	0	0	2	1	1	1	1	0	1	1		
Necrosis	0	0	1	0	1	1	0	1	0	1	1	0	1	1	0	0		
SUB TOTAL (X2)	12	16	20	12	14	12	14	16	14	20	12	16	18	12	12	10		
Neovascularisation	1	2	1	1	0	0	1	0	1	2	0	1	2	1	0	0		
Fibrosis	2	2	2	1	1	1	2	1	2	2	1	2	2	1	1	2		
Fattyinfiltrate	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0		
SUB TOTAL	3	4	3	2	1	1	3	1	3	4	1	3	4	2	1	2		
TOTAL	72			62					7	3		61						
AVERAGE	18.0			15.5					18	.3		15.3						

Cell tupe/restmise		HE	MA			HEMA-CNT				HEMA-IL-CNT			Ац				Negative control			
con type response	00105	00106	00107	00108	00105	00106	00107	00108	00105	00106	00107	00108	00105	00106	00107	00108	00105	00106	00107	00108
Polymorphonuclear cells	2	1	0	1	2	2	1	0	1	3	1	2	2	1	2	2	0	0	0	0
Lymphocytes	2	2	2	1	2	3	1	2	3	3	3	3	3	3	3	2	0	0	0	0
Plasma cells	1	1	0	0	1	0	0	0	0	1	0	0	3	2	1	1	0	0	0	0
Macrophages	3	3	3	1	3	3	2	3	3	3	3	3	2	2	3	3	1	1	2	1
Giant cells	1	0	1	0	0	0	0	0	1	1	1	0	0	0	0	0	0	0	0	0
Necrosis	1	0	0	0	1	0	0	0	1	2	0	1	1	0	1	0	0	0	0	0
SUB TOTAL (X2)	20	14	12	б	18	16	8	10	18	26	16	18	22	16	20	16	2	2	4	2
Neovascularisation	1	0	0	0	2	2	0	0	2	2	2	2	2	1	0	0	0	0	0	0
Fibrosis	1	1	1	2	1	3	1	1	2	2	3	3	3	3	2	3	1	1	2	2
Fattyinfiltrate	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
SUB TOTAL	2	1	1	2	3	5	1	1	4	4	5	5	5	4	2	3	1	1	2	2
TOTAL		5	8			6	2			9	6			8	8			1	6	
AVERAGE		14	.5			15	5.5			24	.0			22	2.0			4	.0	

Supplementary Table 2

More quantitative data analysed from the pathology grafts and a summary

of the results 1

ROTA → Polyrotaxane-based gel

CNT→ Multiwall carbon nanotube

IL→ Ionic Iiquid (N,N-Diethyl-N-methyl-N-(2-methoxyethyl) ammonium tetrafluoroborate) HEMA→ Poly-hydroxyethyl methacrylate gel

			Non-irritant (0.0 up to 2.9)					
			Slight irritant (3.0 up to 8.9)					
	Average irritant	∠ between test	Moderate irritant					
			(9.0 up to 15.0)					
			Severe irritant (>15)					
1-week implantation	•	•						
Negative control	13.5	-	-					
ROTA	17.5	4.0	Slight irritant					
ROTA-IL	16.0	2.5	Non-irritant					
ROTA-CNT	19.3	5.8	Slight irritant					
ROTA-IL-CNT	17.8	4.3	Slight irritant					
HEMA	15.8	2.3	Non-irritant					
HEMA-CNT	17.8	4.3	Slight irritant					
HEMA-IL-CNT	19.5	6.0	Slight irritant					
Au	16.8	3.3	Slight irritant					
	•	•	•					
4-week implantation								
Negative control	4.0	-	-					
ROTA	18.0	14.0	Moderate irritant					
ROTA-IL	15.5	11.5	Moderate irritant					
ROTA-CNT	18.3	14.3	Moderate irritant					
ROTA-IL-CNT	15.3	11.3	Moderate irritant					
HEMA	14.5	10.5	Moderate irritant					
HEMA-CNT	15.5	11.5	Moderate irritant					
HEMA-IL-CNT	24.0	20.0	Severe irritant					
Au	22.0	18.0	Severe irritant					
a: Negative difference is rec	orded as zero.		-					

Supplementary Table 3

more quantitative data analysed from the pathology grafts and a summary

of the results 2