

Effects of the Media Conditioned by Various Macrophage Subtypes Derived from THP-1 Cells on Tunneling Nanotube Formation in Pancreatic Cancer Cells

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Abstract

Background

Tunneling nanotubes (TNTs) are special membrane structures for intercellular communications. Vital cargoes (such as mitochondria) could be delivered from healthy cells to rescue damaged ones through TNTs. The TNTs could be utilized for the purpose of systematic delivery of therapeutic agents between cells. However, there are insufficient studies on the controlled enhancement of TNT formations. The purpose of this study is to understand how macrophages influence the TNT formation in cancer cells.

Results

Here we compared the capabilities of inducing TNTs in human pancreatic cancer cells (PANC-1) of the media conditioned by M0, M1 and M2 macrophages derived from THP-1 cells. The M0 and M1 macrophage conditioned media promoted TNT formation. Using a focused ion beam to cut through a TNT, we observed tunnel-like structures inside dense cytoskeletons with scanning electron microscopy. The TNT formation correlated with raised motility, invasion, and epithelial–mesenchymal transition in the PANC-1 cells. Mitochondria and lysosomes were also found to be transported in the TNTs.

Conclusions

These results suggest the TNT formation could be one of the responses to the immune stress in pancreatic cancer cells caused by M0 and M1 macrophages. This finding is valuable for the development of macrophage-targeting cancer therapy.

Background

Tunneling nanotubes (TNTs) are particular membrane structures for short- and long-distance (up to hundreds of micrometers) cell–cell communications. Two cells connected by a TNT can exchange various materials, including viruses [1], proteins [2], microRNAs [3], mitochondria [4], etc. The TNT formation is also thought to be one kind of stress response. For example, the PC12 cells treated by ultraviolet light formed TNTs with the untreated cells, and mitochondria were observed to be delivered from the healthy cells to the ultraviolet light-treated ones [4]. Moreover, as pancreatic cancer cells were treated by doxorubicin, the formation of TNTs were enhanced and the drug could be transferred between the cancer cells via the TNTs [5]. It seems that cells under various types of stresses can produce TNTs for delivering beneficial or detrimental components among nearby cells. It is straightforward to suspect that TNTs may also be involved in the complicated interactions between cancer cells in a tumor.

Macrophages are important constituents of the tumor microenvironment. The macrophages in a tumor microenvironment can be differentiated into various subtypes according to their T helper type 1 and T

helper type 2 (TH1–TH2) polarizations [6]. The M1 (pro-inflammatory) macrophage can enhance drug sensitivities or decrease tumor activities [7]. On the contrary, the M2 (anti-inflammatory) macrophage tend to promote tumorigenesis or malignancy in some cancers [8, 9]. It is thus required to explore how these different types of macrophages influence the TNT formation in cancer cells.

In the present work, we studied the effects of conditioned media (CMs) of macrophage subtypes M0, M1, and M2, on the TNT formation in PANC-1 human pancreatic cancer cells. The M0 and M1 CMs made the PANC-1 cells produce more TNTs and increased the cell migration speeds. In contrast, the M2 CM was less effective on the TNT formation in the PANC-1 cells. We employed confocal microscopy and focused ion beam-scanning electron microscopy (FIB-SEM) to observe dense cytoskeletal networks inside a TNT. We also checked the correlation between the TNT formation and the epithelial–mesenchymal transition (EMT) of PANC-1 cells. In addition, bidirectional transportation of mitochondria between two cancer cells in the macrophage CM was observed.

Results

M0 and M1 macrophage CMs induced the formation of TNTs in PANC-1 cells

In these experiments, the PANC-1 cells were originally cultured in DMEM for 24 h, and then we replaced DMEM with the macrophage CMs for another 48 h of culture. In the M0 and M1 CMs, the PANC-1 cells produced significantly more TNTs in comparison with those cultured in DMEM and the M2 CM, as shown in Fig. 1(a). We also observed morphological changes in the PANC-1 cells caused by the macrophage CMs. We used the images of PANC-1 cells in the M0 CM as the representative. As shown in Fig. 1(b), the PANC-1 cells cultured in DMEM formed compact colonies, while those in the M0 CM [Figs. 1(c) and 1(d)] were well separated and demonstrated TNT formation. Figures 1(e) and 1(f) show the redistribution of cytoskeletons (F-actin in orange and microtubule in green) caused by the M0 CM, and the TNT was filled with both F-actin and microtubules as indicated by the arrow in Fig. 1(f).

It was proposed that epidermal growth factor (EGF) might play an important role in the TNT formation in rat breast cancer cells (MTLn3) [10, 11]. In our experiments, however, the addition of 200 ng/mL EGF into DMEM induced much less TNTs than the M0 and M1 CMs did [Fig. 1(a)]. We also found very low concentrations of EGF in the macrophage CMs [Fig. S2(a) in the Supplementary Information]. Therefore, we conjectured that the TNT formation in PANC-1 cells induced by the macrophage CMs might be irrelevant with EGF.

Ultrastructures inside TNTs visualized with FIB-SEM

TNTs are not empty membrane tubes but filled with different cytoskeletons in different types of cells [12]. We observed that the TNTs between PANC-1 cells contained both actin filaments and microtubules, as shown in the fluorescence image in Fig. 1(f). The TNT looked like an open-ended tube extending from the cell membranes, and both cytoskeletons inside the TNT were consecutive with those in the cell body. However, how the cytoskeletons are arranged in such a narrow space and how they are capable of

delivering cargos were not resolvable by using optical microscopy. We therefore fixed the PANC-1 cells and dissected the TNTs by FIB, and then visualized the inner structures with SEM, as illustrated in Fig. 2(a). The SEM image of a longitudinally cut TNT in Fig. 2(b) reveals that the TNT was filled with densely packed cytoskeletons. We also employed FIB-SEM to produce a series of cross-section images along one TNT [Fig. 2(c)–(g)]. We observed some tunnel-like structures in the cross sections, surrounded by cytoskeletons. These inner tunnels suggest that cargos could be transported in the TNT in a squeezed manner. In other words, the cargos might be pushed by the contraction force provided by actin–myosin interaction and passed through small the tunnels inside cytoskeletal networks.

TNT formation is correlated with enhanced cell motility and EMT

In addition to the formation of TNTs, we also found that the M0, M1, and M2 CMs significantly increased the migration speed of PANC-1 cells [Fig. 3(a)], while only the M1 CM enhanced the cell invasiveness [Fig. 3(b)]. We thus conjectured that the enhanced formation of TNTs might result from the dissociation of two cells originally in contact, known as the “cell dislodgement” TNT formation process [13], as shown in Fig. S3. It is straightforward to correlate the increase of cell migration speeds with EMT, which generally leads to higher cell motility and invasiveness [14, 15]. We used qPCR to check the mRNA levels of two EMT-relevant proteins, vimentin and Zeb-1 [16]. The mRNA of Zeb-1 was increased by M0 and M1 CMs, although the difference was statistically insignificant [Fig. 3(c)]. In addition, the mRNA level of connexin 43 was also raised by M0 and M1 CMs [Fig. 3(d)]. Because connexin 43 is involved with the p-38 mediated cell migration [17], this result suggests this signaling pathway might be a part of the mechanisms in the migration acceleration by the M0 and M1 CMs in Fig. 3(a). In contrast, the M2 CM did not cause the variation of vimentin, Zeb-1, and connexin 43.

We also monitored the spatial distributions of two cell–cell adhesion proteins, ZO-1 and b-catenin [14, 18], in the PANC-1 cells by using confocal fluorescence microscopy. The images in Fig. 3(e) show that ZO-1 and b-catenin moved from cell membranes into the cytoplasm when the cells were in M0 and M1 CMs. This process was consistent with EMT as well as the cell-dissociation behavior in Fig. 1(c). The translocation of ZO-1 and b-catenin correlated with the TNT formation induced by M0 and M1 CMs in Fig. 1(a). On the other hand, the M2 CM did not induce similar variations of ZO-1 and b-catenin.

Mitochondria and lysosomes were delivered between PANC-1 cells through TNTs

A variety of cellular components (cargos) have been found to be delivered through TNTs [19]. Because the transfer of mitochondria through TNTs was thought to be a rescue of apoptotic cells [4], we wanted to understand if mitochondrion transportation occurs in macrophage CM-induced TNTs. Here we used the M0 CM as a representative treatment. Two groups of PANC-1 cells were sub-cultured and labeled with MitoTracker™ Green FM and MitoTracker™ Orange CMTMRos, separately. In Fig. 4(a), we overlapped fluorescence and bright-field images to show the relative positions of the two mitochondria and the TNT connecting the two PANC-1 cells in the M0 CM. The time-lapse images in Figs. 4(b)–4(f) show that two mitochondria moved head to head in a TNT, collided, and then moved together into the upper cell. (Please also see Video S1 in the Supplementary Information.) Such active transportation of mitochondria is

thought to involve motor proteins [20]. Similar bidirectional transportation of vesicles containing carboxyl-modified quantum dots has been observed in TNTs formed in mouse tissues [21]. We also confirmed the co-localization of kinesin with the mitochondria within the TNT (Fig. S4 in the Supplementary Information). In addition to mitochondria, lysosomes were also found in the TNT connecting two cancer cells, as shown in Figs. 4(g) and 4(h). Wong et al. pointed out that lysosomes could regulate mitochondrial functions by membrane contacts [22]. We conjectured that the co-transportation of mitochondria and lysosomes through TNTs could also be one of the PANC-1 cells' responses to the macrophage CMs.

Discussion

Macrophages play complicated roles in different kinds of cancers by secreting various cytokines [6, 23, 24]. They may also be involved in the interactions between cancer cells and other types of cells, such as fibroblasts [25]. It is generally recognized that M1 macrophages tend to perform pro-inflammatory activities and can be detrimental to tumor cells. On the other hand, M2 macrophages in the tumor microenvironment can support tumor progression and malignancy [6, 7]. Therefore, for clinical applications, we must be very careful to evaluate the pros and cons before introducing additional macrophages into the tumor microenvironment.

In the present work, in order to avoid direct contacts between macrophages and cancer cells, we studied the effects of macrophage CMs on the human pancreatic cancer cell line PANC-1. We found that M0 and M1 CMs induced the formation of TNTs and increased the migration speeds of the cancer cells, while only the M1 CM enhanced the invasiveness of PANC-1 cells. These effects were correlated with raised expression of the EMT-relevant protein Zeb-1, as well as the translocation of ZO-1 and β -catenin from membranes to the cytoplasm in the PANC-1 cells. Connexin 43, which is related to the p-38 mediated cell migration, was also increased by M0 and M1 CMs. In contrast, although the M2 CM increased the migration speed of PANC-1 cells, it did not induce identifiable variations in TNTs, Zeb-1, and connexin 43. We thus suspected that, the THP-1 cell-derived M2 macrophages might not resemble those polarized in the tumor microenvironment, or the macrophages co-cultured with cancer cells [26]. Patheja and Sahu reported that the CM from M0 macrophages derived from U937 monocytes also induced the formation of TNTs between human breast cancer cells MCF-7 [27]. Interestingly, the TNTs could connect the MCF-7 cells and cytoplasmic fragments called microplasts, and transported mitochondria and vesicles from the parent cell to the microplast.

We employed FIB-SEM to reveal the cytoskeletons and tunnel-like structures embedded in the crowded space in a TNT. Sartori-Rupp et al. reported individual tunneling nanotubes (iTNTs) bundled in a TNT between neuronal cells by using cryo-transmission electron microscopy [28]. These iTNTs could serve as transportation channels for vesicles and mitochondria. We suspected that mitochondria or lysosomes might be squeezed through the TNT by the contraction force generated from actin–myosin interactions and the thrust force from co-localized motor proteins such as kinesin.

Macrophages have been proposed as novel cell-based carriers for nanomedicines because of their internalization capability and tumor- and hypoxia-approaching tendency [29]. The nanoparticle-carrying M1 macrophages were demonstrated to be able to migrate through the endothelial barrier into tissues surrounding a tumor [30]. The efficacy of drugs carried by macrophages could be enhanced by the TNTs formed between macrophages and tumor cells [31]. Therefore, manipulating cancer cells to produce TNTs in a controllable manner may pave a new way in cancer therapy.

Conclusion

In summary, we have shown that M0 and M1 CMs induced the formation of TNTs in PANC-1 pancreatic cancer cells. The formation of TNTs was correlated with higher cell motility, EMT, and reduced cell viability of the PANC-1 cells. We also observed bidirectional transportation of mitochondria in the TNTs, and the mitochondrial transportation could facilitate the distribution of superoxides among the PANC-1 cells.

The findings may pave a way for the development of macrophage-targeting cancer therapy.

Methods

Cell culture

The human pancreatic cancer cell line PANC-1 (ATCC CRL-1469) was obtained from the Bioresource Collection and Research Center (Hsinchu, Taiwan). We cultured PANC-1 cells in Dulbecco's Modified Eagle Medium (DMEM, Gibco®, Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS) and 1% antibiotic pen-strep-ampho. The human monocyte cell line THP-1 (ATCC TIB-202) was also obtained from the Bioresource Collection and Research Center. We cultured the THP-1 cells in RPMI supplemented with 10% FBS and 1% antibiotic pen-strep-ampho.

Preparation of macrophage-conditioned media

In order to collect the macrophage CMs, we harvested 6×10^6 THP-1 cells into 10 mL DMEM with 0.32 mM phorbol 12-myristate 13-acetate (PMA, Sigma-Aldrich, St. Louis, MO, USA). After treatment with PMA for 24 h, most of the monocytes attached onto the bottom of the culture dish and were considered to be differentiated into M0 macrophages [32, 33]. Further incubating the M0 macrophages with 10 pg/mL LPS + 20 ng/mL IFN- γ for 24 h or 20 ng/mL IL-4 + 20 ng/mL IL-13 for 72 h, we polarized the M0 macrophages into M1 and M2 macrophages, respectively [33]. The expression levels of relevant markers of M1 and M2 macrophages are shown in Fig. S1 in the Supplementary Information. We replaced the culture medium of each macrophage mentioned above with fresh DMEM, and conducted another 24 h of cultivation. The media were then collected and used as the macrophage CMs in the experiments.

TNT quantification

We seeded PANC-1 cells at 1×10^5 cells/mL in a μ -Dish (35 mm, high Grid-500, ibidi GmbH, Gräfelfing, Germany) under normal cultivation conditions (5% CO₂ at 37 °C) overnight. Then the cells were cultured in the macrophage-CM for another 48 h. The cells were washed with phosphate-buffered saline (PBS, pH = 7.4) three times and then fixed with 3.7% formalin for 15 min at room temperature. We removed the fixative and washed the cells with PBS three times, and then used 1 μ g/mL DAPI (Thermo Fisher Scientific) to stain the nuclei. After 10 min of incubation, the fixed cells were thoroughly washed with PBS and then imaged with a 20 \times phase-contrast objective installed on an inverted microscope (ECLIPSE Ti, Nikon, Tokyo, Japan). We counted the TNTs following these two criteria: (1) the tube connected two cells, and (2) the tube length was longer than 50 μ m.

FIB-SEM imaging

PANC-1 cells were cultured on well-cleaned and sterilized cover slips (22 mm \times 22 mm). After treatment with the macrophage CMs, the cells were rinsed with PBS and then fixed with 3.7% formaldehyde for 15 min and later post-fixed with 1% OsO₄ in PBS for 2 h. After triple PBS flushes, we replaced the medium with ethanol gradually from 10% (v/v in H₂O) to 99.9%. The final samples were dried with a critical point dryer (EM CPD300, Leica Microsystems, Wetzlar, Germany). Before the FIB-SEM operation, the sample surface was sputtered with 10 nm gold to raise the conductivity. FIB-SEM operations were conducted with the FEI Helios NanoLab 660 (Thermo Fisher Scientific). The target area was firstly deposited with a 500 nm platinum protection layer via ion beam-induced deposition, rough-milled with 40 pA and then fine-milled with 7.7 pA gallium ion beams at a 30 kV acceleration voltage. All the SEM images were taken right after the FIB manipulations, acquired with a 25–50 pA beam current at a 1–5 kV acceleration voltage and an “in-lens” detector for the secondary electrons.

Cell migration speed analysis

We seeded PANC-1 cells at 5×10^4 cells/mL in a 35 mm dish under the normal cultivation condition overnight. The culture medium was then changed to the macrophage-CMs. We used a 10 \times phase-contrast objective to capture the time-lapse images of cells every 10 min for 48 h. In each experiment, the migration speeds of 45 cells were calculated using ImageJ.

Cell invasion assay

A sample of $\sim 3 \times 10^5$ PANC-1 cells was re-suspended in a 6-well plate with the normal cultivation condition overnight. The culture media were changed to the macrophage CMs for another 48 hour culture. Then cells were starved for 24 hours in the serum-free medium prior to the invasion assay. About 5×10^4 cells were seeded into a well of the Transwell chamber (ab235697, Basement Membrane, 8 μ m, Abcam) containing the normal culture medium in the bottom chamber. Cells were cultured for 24 hours then labeled with the fluorescence dye (Ex. 530 nm/ Em. 590 nm) for 60 mins according to the kit's instruction. The fluorescence intensities of invading cells were measured by using a plate reader (Synergy 2, BioTek Instruments, Winooski, VT, USA) to evaluate the cell invasion percentage.

Quantitative polymerase chain reaction (qPCR)

Total RNA was purified using an RNeasy mini kit (217604, Qiagen, Hilden, Germany) and reverse-transcribed to cDNA with the High-Capacity RNA-to-cDNA™ Kit (4387406, Applied Biosystems™, Thermo Fisher Scientific). qPCR was performed with PowerTrack™ SYBR Green Master Mix (A46110, Applied Biosystems™) and LightCycler 480 (Roche, Basel, Switzerland). The results were normalized with actin as internal control and evaluated by using the $2^{-\Delta\Delta Ct}$ method. Primers used in this study were synthesized by Genomics (New Taipei City, Taiwan) and listed in Table 1.

Table 1. Sequences of the primers for real-time PCR analysis.

Gene	Primer sequence (5'→3')
<i>VIM</i>	forward: AATGACCGCTTCGCCAACT reverse: ATCTTATTCTGCTGCTCCAGGAA
<i>ZEB1</i>	forward: TGAGCAAACCATGATCCTAATGT
<i>M-Sec</i>	reverse: CAGGTGCCTCAGGAAAAATGA
<i>Cx43</i>	forward: TGCTCCAGAACCTGCATGAGGA
<i>ZO-1</i>	reverse: AACTCAGGCAGCCTCGTGTCTA
<i>CTNNB1</i>	forward: GACAAGGTTCAAGCCTACTCAACTG
<i>IL1B</i>	reverse: TGTCCCCAGCAGCAGGAT
<i>TLR2</i>	forward: CACGCAGTTACGAGCAAG
<i>CCL22</i>	reverse: TGAAGGTATCAGCGGAGG
<i>ACTB</i>	forward: AGCCGACACCAAGAAGCAGAGATG reverse: CGGCGCTGGGTATCCTGATGT forward: ATGATGGCTTATTACAGTGGCAA reverse: GTCGGAGATTCGTAGCTGGA forward: GCTCGGAGTTCTCCCAGTTTC reverse: GAGCTGCCCTTGCAGATAC forward: ATTACGTCCGTTACCGTCTG reverse: TAGGCTCTTCATTGGCTCAG forward: TGACGGGGTCACCCACACTGTGCCCATCTA reverse: CTAGAAGCATTTGCGGTGGACGATGGAGGG

Confocal fluorescence imaging

For imaging the junction proteins, the PANC-1 cells were washed with PBS for three times and then fixed with anhydrous methanol for 10 min. The cells were permeabilized with 0.1% Triton X-100 for 10 minutes and blocked with 1% BSA in 1× PBS for 1h at room temperature. Cells were then incubated with the antibodies of ZO-1 (MA3-39100-A488, 1:100, Thermo Fischer Scientific) and b-catenin (ab32572, 1:250, Abcam) overnight at 4°C, and followed by a further incubation at room temperature for 1h with the Goat Anti-Rabbit (A11008, 1:1000, Invitrogen, Alexa Fluor®488) secondary antibody. Nuclear DNA was labeled with DAPI (62248, 1:1000, Thermo Fisher Scientific). Images were acquired with a confocal microscope (SP5, Leica Microsystems).

Cell viability test

PANC-1 cells were seeded at 5×10^3 cells/mL into a 96-well plate under the normal cultivation condition overnight. The culture medium was changed to the macrophage CMs for another 48 h of culture. The cell viability was quantified using the alamarBlue assay (DAL1100, Thermo Fisher Scientific).

Statistical analysis

All the data are presented as mean \pm standard deviation. Comparisons between two groups were made by using a two-tailed Student's *t*-test. Comparisons among data of more than two groups were conducted with one-way analysis of variance (ANOVA). Then the differences between specific two groups were checked with post hoc Tukey's test.

Abbreviations

VIM

vimentin

ZEB1

Zinc Finger E-Box Binding Homeobox 1

M-Sec

TNF α -inducible protein 2

Cx43

Connexin 43

ZO-1

zonula occluden-1

CTNNB1

Catenin Beta 1

IL1B

Interleukin 1 Beta

TLR2

Toll Like Receptor 2

CCL22
C-C Chemokine type 22
ACTB
Actin Beta
TNTs
tunneling nanotubes
CMs
conditioned media
FIB-SEM
focused ion beam-scanning electron microscope
EMT
epithelial–mesenchymal transition
EGF
epidermal growth factor

Declarations

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Authors' contributions

CHL and CNS conceived, designed the experiments, and supervised the project. CWL conducted most of the confocal microscopy and FIB-SEM imaging experiments. CCK performed most of the cell invasion, ELISA, and qPCR. CCK and HJP, prepared the samples and quantified the TNTs. CJL conducted the flow cytometry analysis. CWL analyzed the data. CWL and CHL wrote the manuscript. All authors read and approved the final manuscript.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article and its Additional files.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Competing interests

The authors declare that no conflicts of financial and non-financial interests exist.

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Figures

Figure 1

Macrophage conditioned media induce formation of TNTs in PANC-1 cells. (a) Average numbers of TNTs/cell in PANC-1 cells under the treatments of macrophage CMs and 200 ng/ml EGF. ***, $P < 0.005$ in comparison with the PANC-1 cells in DMEM (post hoc Tukey's test). (b), (c) Typical morphology of PANC-1 cells in DMEM and M0 CM, respectively. (d) Typical SEM image of PANC-1 cells cultured in M0 CM. (e), (f)

Fluorescence images of cytoskeletons (green, microtubules, orange, F-actin) in PANC-1 cells in DMEM and M0 CM, respectively. The blue color represents DAPI-stained nuclei.

Figure 2

FIB-SEM images of the inner structures of TNTs. (a) Schematic of the FIB-SEM dual beam system. (b) SEM image of a longitudinally cut TNT between two PANC-1 cells cultured in M0 CM. (c) SEM image of a PANC-1 cell cultured in M0 CM. The region to be cut by FIB is marked by the yellow dashed rectangle. (d)–(g) Series of cross sections along the TNT in (c). Each section was separated from the previous one by ~500 nm.

Figure 3

The TNT formation induced by macrophage CMs is correlated with enhanced cell motility and EMT. (a) Migration speeds of PANC-1 cells in DMEM and macrophage CMs. In each experiment, we measured the speeds of 45 cells. (b) Invasion in 24 h of PANC-1 cells in DMEM and macrophage CMs. (c) mRNA levels of vimentin (VIM) and Zeb-1 (ZEB1) in PANC-1 cells cultured in DMEM and macrophage CMs. For Zeb-1, the *P*-value between PANC-1 cells in DMEM and M0 CM is 0.17, while that between PANC-1 cells in DMEM and M1 CM is 0.34. (d) mRNA levels of connexin 43 (Cx43) in PANC-1 cells cultured in DMEM and macrophage CMs. All data were obtained from three independent experiments. ***, $P < 0.005$, **, $P < 0.01$, *, $P < 0.05$ (post hoc Tukey's test) in comparison with the PANC-1 cells in DMEM. (e) Confocal fluorescence images of ZO-1 and b-catenin of PANC-1 cells in DMEM and macrophage CMs.

Figure 4

Mitochondria and lysosomes can be transported between PANC-1 cells in M0 CM connected by a TNT. (a) Overlapped bright-field and fluorescence images showing two PANC-1 cells connected by a TNT. (b)–(f) Time-lapse confocal fluorescence images of mitochondrial trafficking within the TNT in (a). The two PANC-1 cells were sub-cultured and labeled with MitoTracker™ Green FM and MitoTracker™ Orange CMTMR, individually. Two mitochondria from each cell moved forward in opposite directions, collided, and then moved together into the upper cell. (g) Fluorescence image of lysosomes in PANC-1 cells. The lysosomes were labeled with LysoTracker® Green. (h) Magnified view of the area enclosed by a red dashed square in (g).

Supplementary Files

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- [VideoS1.mp4](#)
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