



Resonant waveguide grating biosensor-enabled label-free and fluorescence detection of cell adhesion

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ABSTRACT

Cell adhesion to extracellular matrix (ECM) is fundamental to many distinct aspects of cell biology, and has been an active topic for label-free biosensors. However, little attention has been paid to study the impact of receptor signaling on the cell adhesion process. We here report the development of resonant waveguide grating biosensor-enabled label-free and fluorescent approaches, and their use for investigating the adhesion of an engineered HEK-293 cell line stably expressing green fluorescent protein (GFP) tagged β_2 -adrenergic receptor (β_2 -AR) onto distinct surfaces under both ambient and physiological conditions. Results showed that cell adhesion is sensitive to both temperature and ECM coating, and distinct mechanisms govern the cell adhesion process under different conditions. The β_2 -AR agonists, but not its antagonists or partial agonists, were found to be capable of triggering signaling during the adhesion process, leading to an increase in the adhesion of the engineered cells onto fibronectin-coated biosensor surfaces. These results suggest that the dual approach presented is useful to investigate the mechanism of cell adhesion, and to identify drug molecules and receptor signaling that interfere with cell adhesion.

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1. Introduction

Cell adhesion to extracellular matrix (ECM) is critical to many different aspects of cell biology and human physiology, ranging from cell differentiation, migration, proliferation, and survival to tissue development and morphogenesis. Cell adhesion to the ECM is mediated through adhesion complexes that consist of specific integrin receptors, cytoskeletal elements and many distinct adaptor and signaling proteins [16,33,44]. These adhesion complexes permit cells to sense the composition, organization, density, rigidity, and biochemical and rigidity gradients of the ECM [2,3,23,28]. Abnormal cell adhesion has been implicated in many diseases including arthritis, cancer, cardiovascular diseases, inflammation, and osteoporosis [25,39].

Cell adhesion to a surface is also essential to the development of biomaterials for tissue engineering and of novel cell culture

vessels for cell therapy and other applications [20,21,37]. The ability to examine the spatial and temporal dynamics of cell adhesion to a surface is important to understand the biochemical, biological and physical aspects of cell adhesion [34]. Owing to their non-invasiveness and high temporal resolution in measurements various label-free biosensors have been used to characterize cell adhesion [5,7,15,30,32,42]. Optical biosensors including surface plasmon resonance imaging (SPRi) offers high spatial resolution, so it is possible to estimate the cell/substrate distance. For cultured primary goldfish glial cells, the cell/substrate distance was estimated using SPRi to be about 160 nm for most parts of the cells, and about 25 nm for the cell peripheral lamellipodia [17]. Optical biosensors including SPR and resonant waveguide grating (RWG) biosensor are to large degree sensitive to the changes in local mass density, so it is possible to examine the remodeling of sensor surfaces [9]. For vascular smooth muscle cells cultured on fibronectin, it was found that the cells secret and deposit proteins as much as 120 ng/cm² overnight [26] and undergo dynamic membrane ruffling with nanometer precision [27]. Both surface plasmon-based infrared spectroscopy [43] and resonant waveguide grating (RWG) biosensor [9,18] enable cell sensing with multiple sampling depths, so it is possible to resolve the multi-staged process of the adhesion and monolayer formation of Madin–Darby canine kidney (MDCK) cells [43], and to detect the inhomogeneity in refractive index within the distinct layers of a fibroblast cell perpendicular to the biosensor surface during cell adhesion [18]. Furthermore, these

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biosensors are also useful to study receptor signaling that leads to alteration in adhesion pattern of cultured cells [1,4,10,12,14], and to decode the impact of ECM coatings on receptor signaling and drug pharmacology [36]. However, few studies have been attempted to study the impact of the external stimuli-induced intracellular signaling, in particular, G protein-coupled receptor (GPCR) signaling, on the cell adhesion process *per se*.

We here report the development of RWG biosensor-enabled label-free and fluorescence detection, and its use to characterize the adhesion of an engineered human embryonic kidney HEK293 cell line on distinct ECM coated surfaces. HEK293 is one of the most widely used model cell line for basic research and drug discovery. We found that the activation of β_2 -adrenergic receptor (β_2 -AR) enhances cell adhesion.

2. Materials and methods

2.1. Materials

Carvedilol, collagen Type IV from human cell culture, epinephrine, fibronectin from human plasma, isoproterenol, pin-dolol, salbutamol, and salmeterol were obtained from Tocris Chemical Co. (St. Louis, MO, USA). Cytochalasin D, nocodazole, RGD peptide (Arg-Gly-Asp), and vinblastine were purchased from Sigma Chemical Co. (St. Louis, MO, USA). Except for two ECM proteins which were freshly dissolved in the assay buffer (1× Hank's balanced salt buffer, 20 mM Hepes, pH 7.1; HBSS), all compounds were stocked in dimethyl sulfoxide (DMSO) at 100 mM, and were diluted using the assay buffer to the indicated concentrations. Epic® 384-well biosensor cell culture compatible (tissue culture treated, TCT) and Epic® 384-well biosensor fibronectin-coated microplates were from Corning Inc., (Corning, NY, USA). Collagen IV coated biosensor microplates were freshly made by overnight drying 10 μ l 20 μ g/ml collagen IV solution per well under nitrogen inside an incubator.

2.2. Generation and culture of HEK- β_2 AR-GFP stable cell line

The parental HEK293 cell line was obtained from American Type Cell Culture (Manassas, VA, USA), and cultured in minimum essential medium having 2 mM glutamine, 4.5 g/L glucose, 2 mM glutamine, 10% fetal bovine serum (FBS), and antibiotics.

The engineered HEK- β_2 AR-GFP cells were made in house. Briefly, HEK293 cells were transfected with human pCMV- β_2 AR-GFP plasmid (OriGene Technologies, Inc., Rockville, MD, USA) using Lipofectamine™ LTX and Plus Reagent (Invitrogen) in a 6-well cell culture plate. The cells were treated with 500 μ g/ml G418 Geneticin® (Invitrogen, Grand Island, NY, USA) the next day. After transfection for 7 days in total, the survived cells were then diluted to 2–3 cell/well in a 96well cell culture plate to allow clones derive from single cell. Stable clones with homogeneous expression level of β_2 AR-GFP were selected by visualizing the GFP signal under a fluorescence microscope. The β_2 AR protein function of these stable cell lines were verified by using dynamic mass redistribution (DMR) assays. The stable cell lines were maintained in the complete medium (that is, DMEM medium containing 10% FBS, penicillin/streptomycin, L-glutamine, and 500 μ g/ml G418). The cells were passaged at 37 °C with 5% CO₂. Except for cell adhesion studies which used physical scraping approach to avoid damage of cell surface β_2 ARs, these cells were passed with trypsin/ethylenediamine-tetraacetic acid when approaching 90% confluence to provide new maintenance culture on T-75 flasks and experimental culture on the biosensor microplates.

2.3. Quantitative real time PCR

Total RNA was extracted from HEK293 cells using an RNeasy mini kit (Qiagen, Cat#74104). On-column DNase digestion was performed using RNase-free DNase set from Qiagen (Valencia, CA) to eliminate genomic DNA contamination. The concentration and quality of total RNA were determined using a Nanodrop 8000 from Thermo Scientific. Customized PCR-array plates for 352 GPCR genes and reagents were ordered from SABiosciences (Frederick, MD). About 1 μ g total RNA was used for each 96-well PCR-array. The PCR-array was performed on an ABI 7300 Real-Time PCR System following the manufacturer's instructions.

2.4. DMR assays

Label-free DMR assays enabled by RWG biosensor were performed using Corning® Epic® BT system, a swept wavelength interrogation system that is capable of imaging whole RWG biosensor microplates with a spatial resolution of 90 μ m [13]. This system uses a light beam from a tunable light source to illuminate whole 384well biosensor microplate, and a high speed complementary metal-oxide semiconductor (CMOS) digital camera to record the escaped and reflected resonant lights from the whole plate. The tunable light source sweeps the wavelength range from 825 to 840 nm in a stepwise fashion to identify the resonant wavelength at each location. Total 150 spectral images were acquired within a single sweeping cycle (3 s), and were then processed into sensor resonant wavelength or DMR image in real time. This system, due to its small footprint, enables cell assays inside a typical cell culture incubator.

For receptor signaling using DMR assays, cells were first harvested from T75 flask using trypsin/ethylenediamine-tetraacetic acid, and re-suspended in the completed medium. Cells were seeded onto the fibronectin coated biosensor microplates using a seeding density of 12,000 cells per well in 40 μ l the complete medium, and cultured at 37 °C with 5% CO₂ inside a cell culture incubator for about 22 h to reach a confluency of ~95%. Afterwards, the cells were washed three times using a plate washer (Bio-Tek Microplate Washers ELx405™, Bio-Tek, Winooski, VT), and maintained in the assay buffer (HBSS) for about 1 h to reach equilibrium at 26 °C. After establishing a 2 min baseline, DMR assays were then initiated by adding the HBSS buffered compound solution and the compound-induced DMR was recorded in real time.

For cell adhesion studies, the biosensor surface was first coated with a specific ECM protein, and pre-incubated with the complete medium solution (25 μ l per well) in the absence and presence of a compound until it reaches an equilibrium state. After a 2 min baseline, 25 μ l the cell suspension in the same medium in the presence of the same compound at equal amount were introduced to each well using a 384well liquid handling device (Epic®). The biosensor response is monitored over time. The cells were mechanically scraped off from T75 flask to avoid damage cell surface receptors, and then re-suspended in the completed medium. For inhibitors, the cells were pretreated with the inhibitors for about 1 h. For the β_2 -AR ligands, the cells were premixed with the ligand 1 min before added into the biosensor microplate. All studies were carried out with at least four replicates unless specifically mentioned.

2.5. Fluorescence microscopy

Epi-fluorescence microscopic imaging was obtained a Zeiss Axioplan fluorescence microscope. Total internal reflection fluorescence (TIRF) microscopic imaging was obtained using an in-house TIRF microscope. Briefly, a 488 nm diode laser (Continuous Wave 488 nm Solid-State Laser, JDSU, Milpitas, CA, USA) is used to launch a light beam into the glass substrate of an Epic®

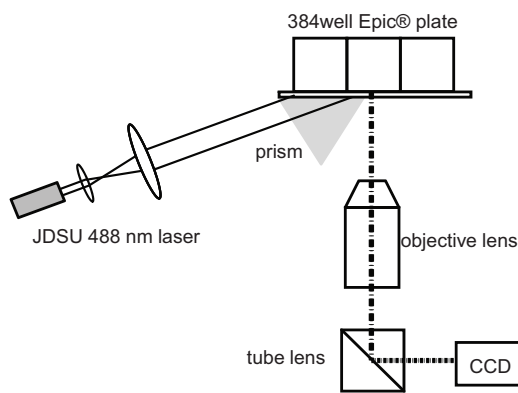


Fig. 1. Schematic drawing showing the setup of a resonant waveguide grating biosensor-enabled total internal reflection fluorescence microscopy. Here, a 488 nm laser light is used to illuminate the biosensor within a well of 384well microplate with the aid of a right angle prism and the biosensor diffractive grating. The evanescent wave excited fluorescence is collected using an objective lens, and focused via a tube lens onto a CCD camera.

biosensor microplate through a right angle prism (5 mm BBAR Coated Right Angle Prism, VIS 0° Coated, Edmund Optics Inc., Barrington, NJ, USA). When the laser incident angle exceeds the total internal reflection critical angle, total internal reflection is observed and the evanescent wave is obtained and used to excite green fluorescent protein-tagged β_2 -ARs located at the cell surface. Immersion oil is applied to fill the gap between the prism and the plate substrate. A Nikon 20 $\times$ (0.4 NA) objective, a tube lens with a 200 mm effective focal length, and a regular cooled CCD camera are used to record the TIRF images. The TIRF microscopy is calibrated by measuring the transmission image of an US Air Force target (USAF), showing that the imaging system has a field of view of 192 $\mu\text{m} \times 144 \mu\text{m}$ and a 1.38 μm spatial resolution.

2.6. Statistic analysis

DMR data were analyzed by using GraphPad Prism 5.0 (GraphPad Software Inc., San Diego, CA, USA). The EC_{50} or IC_{50} values were obtained by fitting the dose DMR response curves with nonlinear regression. The kinetic cell adhesion data were analyzed using paired student's *t* test.

3. Results

3.1. RWG biosensor-enabled fluorescence microscopy

We combined RWG biosensor-enabled label-free with fluorescence imaging approach to characterize cell adhesion. The label-free measurement was performed using a recently reported whole plate RWG imager [13]. The instrument setup for the RWG-enabled TIRF microscopy is illustrated in Fig. 1. This TIRF system uses a 488 nm diode laser to illuminate the RWG biosensors within a 384well biosensor microplate. The light is coupled into the biosensor substrate with the help of a right angle prism and the RWG diffractive grating when the laser incident angle exceeds the total internal reflection critical angle. The evanescent wave obtained is then employed to excite the GFP tagged to the C-terminal of the β_2 -AR within the sensing sampling depth. A CCD camera, coupled with an objective and a tube lens, is used to record the TIRF images of cells with a spatial resolution of 1.38 μm . The label-free and fluorescence measurements were performed independently.

3.2. Characterization of GFP tagged β_2 -ARs

To utilize the dual detection approach to characterize cell adhesion, we constructed an engineered HEK293 cell line that stably expresses GFP tagged β_2 -AR fusion protein ($\beta_2\text{AR-GFP}$). Conventional fluorescence microscopic imaging showed that the fusion protein is primarily located at the cell plasma membrane and the epi-fluorescence intensity is mostly concentrated at the peripheral area of the cells (Fig. 2a). RWG biosensor-enabled TIRF imaging showed that the TIRF intensity was higher in the center area of most individual cells than their peripheral area (Fig. 2b). Given that cell adhesion complexes are known to mostly locate at the cell peripheral area, this result suggests that the $\beta_2\text{AR-GFPs}$ are uniformly distributed in most parts of cells that are outside the adhesion complexes. The TIRF imaging also showed that stimulating the cultured cells with epinephrine led to marked reduction of fluorescence (Fig. 2c), suggesting that once activated the $\beta_2\text{AR-GFP}$ become internalized and move outside the TIRF sampling depth. DMR profiling showed that isoproterenol triggered a small DMR in the parental HEK293 cells (Fig. 2d), but a more robust DMR with distinct characteristics in the engineered cells (Fig. 2e). Isoproterenol is a full agonist for β -adrenergic receptors. Quantitative RT-PCR showed that the parental HEK293 cells predominately express the β_2 -AR, as evidenced by the cycle threshold value which was 34.4, 30.2, 33.3, 29.2, 31.3, 26.5, 27.2, 26.0, and 38.3 for α_{1A} , α_{1B} , α_{1D} , α_{2A} , α_{2B} , α_{2C} , β_1 , β_2 and β_3 -adrenergic receptor, respectively. The cycle threshold for the control gene, hypoxanthine phosphoribosyltransferase 1, was found to be 20.1. The isoproterenol DMR in both cell lines was completely blocked by the β -blocker propranolol, but not by the α_2 -AR-selective blocker yohimbine or the α_1 -AR selective blocker prazosin, suggesting that the isoproterenol DMR in both cell lines are primarily due to the activation of the β_2 -AR. Furthermore, isoproterenol gave rise to a single EC_{50} of $0.78 \pm 0.10 \text{ nM}$ ($n=4$) in the parental cells (Fig. 2f). In contrast, in the engineered cells isoproterenol displayed an assay time dependent potency; that is, the potency to reach the early peak response was found to be biphasic with two well-separated EC_{50} ($0.36 \pm 0.03 \text{ nM}$, and $94.0 \pm 7.6 \text{ nM}$, respectively; $n=4$), while the potency to reach the late plateau response was monophasic with an EC_{50} of $4.05 \pm 0.34 \text{ nM}$ (Fig. 2f). Epinephrine also behaved similarly; it gave rise to a single EC_{50} of $11.4 \pm 1.2 \text{ nM}$ ($n=4$) in the parental cells, and of $124 \pm 11 \text{ nM}$ ($n=4$) in the engineered cells when the DMR amplitudes at 50 min post-stimulation were analyzed. However, when the DMR amplitudes at 5 min poststimulation was analyzed, a biphasic dose response was also evident, leading to two separated EC_{50}s (0.58 ± 0.04 , and $69.5 \pm 7.1 \text{ nM}$, $n=4$). Several possibilities may contribute to this; overexpression of a GPCR may result in altered G protein coupling, receptor organization, and/or signaling pathways [6,29], as well as increased numbers of spontaneously (unliganded) active and inactive receptors [38]. Nonetheless, these results suggest that when overexpressed in HEK293 cells the $\beta_2\text{AR-GFP}$ is functional.

3.3. Temperature and ECM coating sensitive cell adhesion

The adhesion of the HEK- $\beta_2\text{AR-GFP}$ cells onto three distinct surfaces, tissue culture treated (TCT), fibronectin and collagen IV coated, was monitored in real time using the whole plate RWG imager under two different conditions. Results showed that under ambient condition the HEK- $\beta_2\text{AR-GFP}$ cells preferably attached onto the fibronectin coated surface with the fastest kinetics and the largest signal (Fig. 3a and b). Almost identical trend was found under physiological condition (Fig. 3c and d). Furthermore, regardless of surface chemistry cell adhesion became faster under physiological condition than that under ambient condition. Interestingly, once reaching equilibrium state during the assay time, the total signal arising from the adhesion of the same number of cells

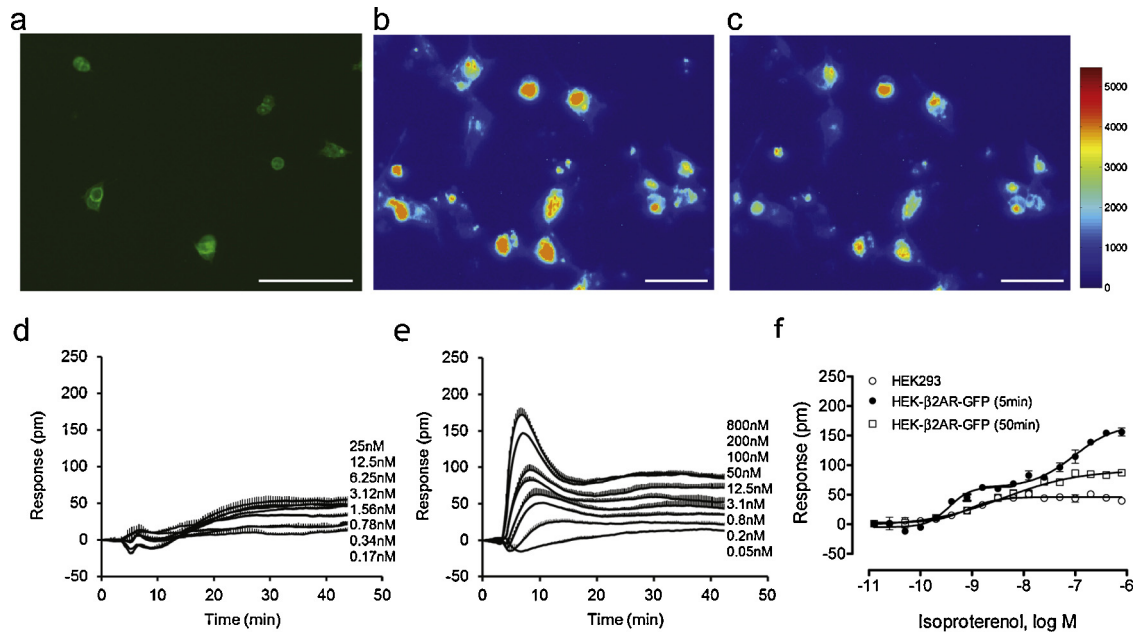


Fig. 2. Characterization of the engineered HEK- β_2 AR-GFP cells. (a)–(c) Representative fluorescence microscopic images of the engineered cells after overnight culture: (a) epifluorescence image; (b) TIRF image before any treatment; (c) TIRF image after treatment with 1 μ M epinephrine for 1 h. For (a)–(c) the image scale bar is 50 μ m. (d)–(f) The DMR dose responses of epinephrine in cells after overnight culturing to form monolayer: (d) the parental HEK293 cells; (e) the engineered cells; (f) comparison of the dose-dependent responses of epinephrine in both types of cells, wherein for the parental cells the DMR amplitude at 50 min post stimulation was plotted as a function of epinephrine dose, and for the engineered cells the DMR amplitudes at both 5 min and 50 min post stimulation were plotted. For (d)–(f), the data represents mean $\pm$ S.D. ($n = 4$).

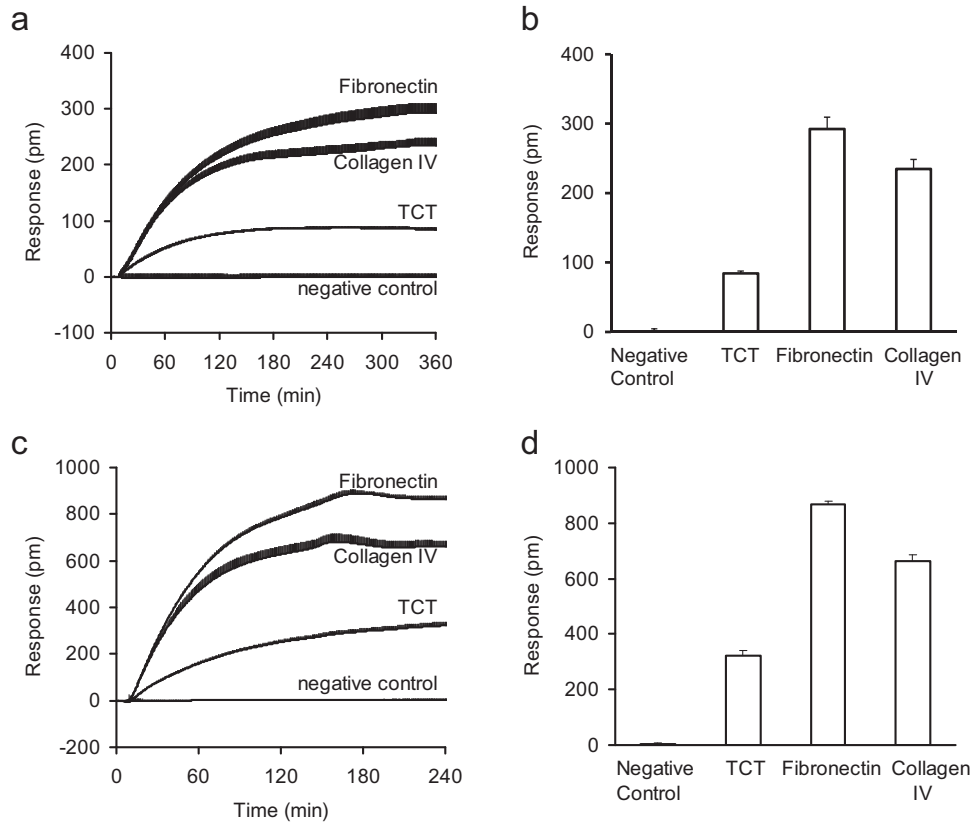


Fig. 3. DMR characterization of HEK- β_2 AR-GFP cell adhesion. (a) and (b) The DMR of the engineered cells adherent onto different surfaces under ambient condition: (a) real-time; (b) the DMR amplitudes at 5 h after cell seeding. (c) and (d) The DMR of the engineered cells adherent onto different surfaces under physiological condition: (c) real-time; (d) the DMR amplitudes at 4 h after cell seeding. The medium only was used as the negative control. For all, the total number of cells added were the same (18,000 cells per well). The data represents mean $\pm$ S.D. ($n = 12$).

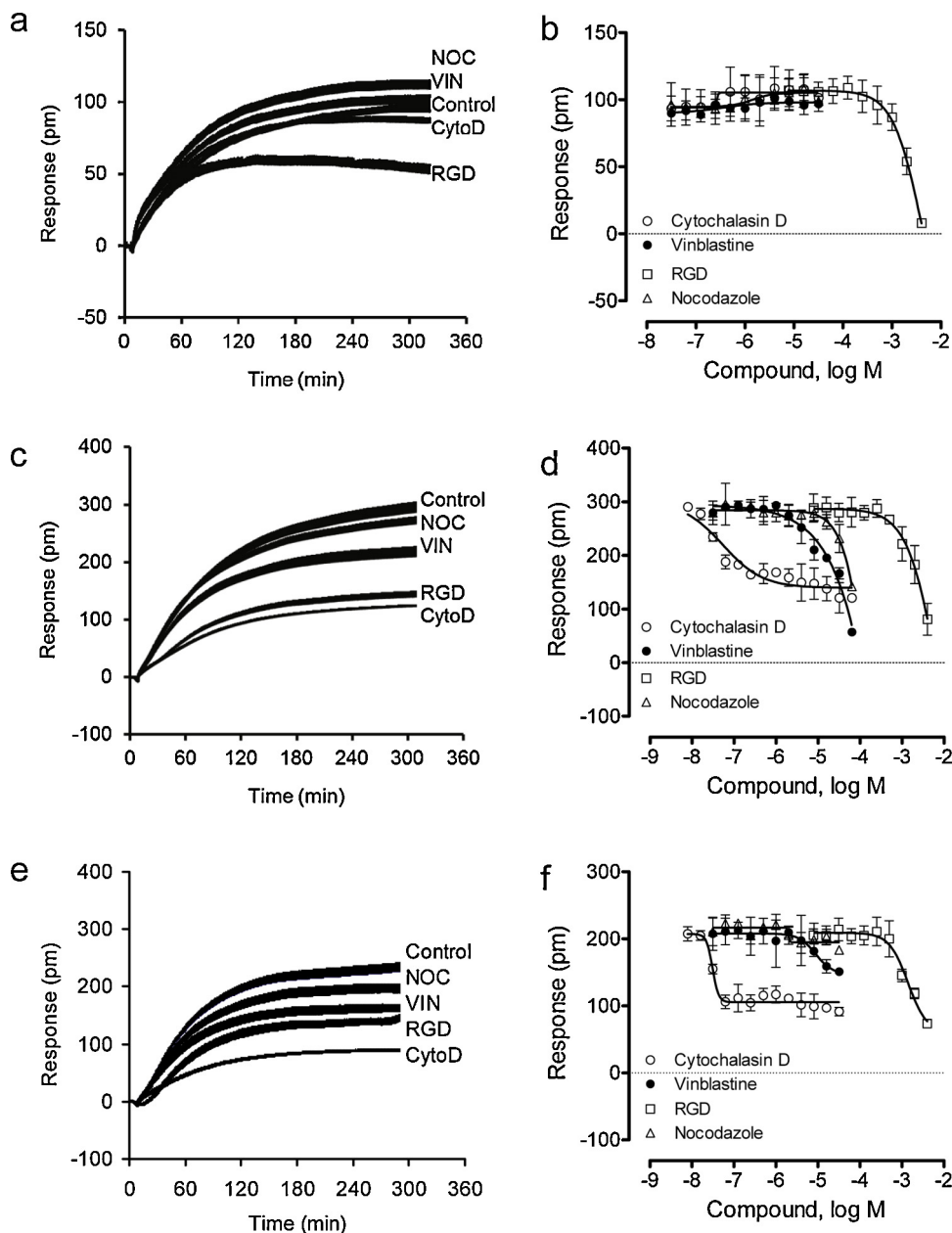


Fig. 4. DMR characterization of HEK-β₂AR-GFP cell adhesion in the presence of different inhibitors under ambient condition. (a)–(f) The DMR of the engineered cells adherent onto different sensor surfaces: tissue culture treated (a) and (b), fibronectin coated (c) and (d), and collagen IV coated (e) and (f). (a), (c) and (e) real-time DMR; (b), (d) and (f) the DMR amplitudes at 5 h after cell seeding as a function of inhibitor dose. For all, the total number of cells added were the same; that is, 18,000 cells per well. For (a), (c) and (e), the inhibitor dose was fixed to be 10 μM, 10 μM, 10 μM and 1 mM for nocodazole (NOC), vinblastine (VIN), cytochalasin D (CytoD), and RGD peptide, respectively. The data represents mean ± S.D. (*n* = 12 for a, c and e; *n* = 4 for b, d and f).

under physiological condition was found to be almost two folds higher than that under ambient condition for all surfaces. These results suggest that the adhesion of the HEK-β₂AR-GFP cells is dependent on temperature and ECM coating, and the temperature effect seems to be intrinsic to cell physiology and independent on ECM coating.

3.4. Mechanisms of cell adhesion under different conditions

To explore the cellular mechanisms governing adhesion process under different conditions, we examined the effects of a panel of inhibitors on the responses of cell adhesion. Cytochalasin D is a potent actin filament disrupting agent by binding to G-actin and thus preventing polymerization of actin monomers [22]. Vinblastine is a microtubule disrupter by binding to tubulin, and can

strongly depolymerize microtubules at minus ends, but does not significantly depolymerize microtubules at plus ends [24]. Nocodazole is a microtubule inhibitor that causes overall decrease in microtubule turnover without net microtubule disassembly [40]. Soluble RGD peptide is an integrin agonist that mimics the binding motif of common ECM proteins including fibronectin and collagen IV [19].

We first examined the cell adhesion mechanism under ambient condition. DMR inhibitor assays showed that on the TCT surface only RGD dose-dependently inhibited the adhesion signal, while the two microtubule inhibitors slightly increased the signal, and cytochalasin D had no effects (Fig. 4a and b). The little effect of cytochalasin D on the cell adhesion onto the TCT surface under ambient condition suggests that actin remodeling is not essential to cell adhesion to the TCT surface. Cytochalasin D is known to be

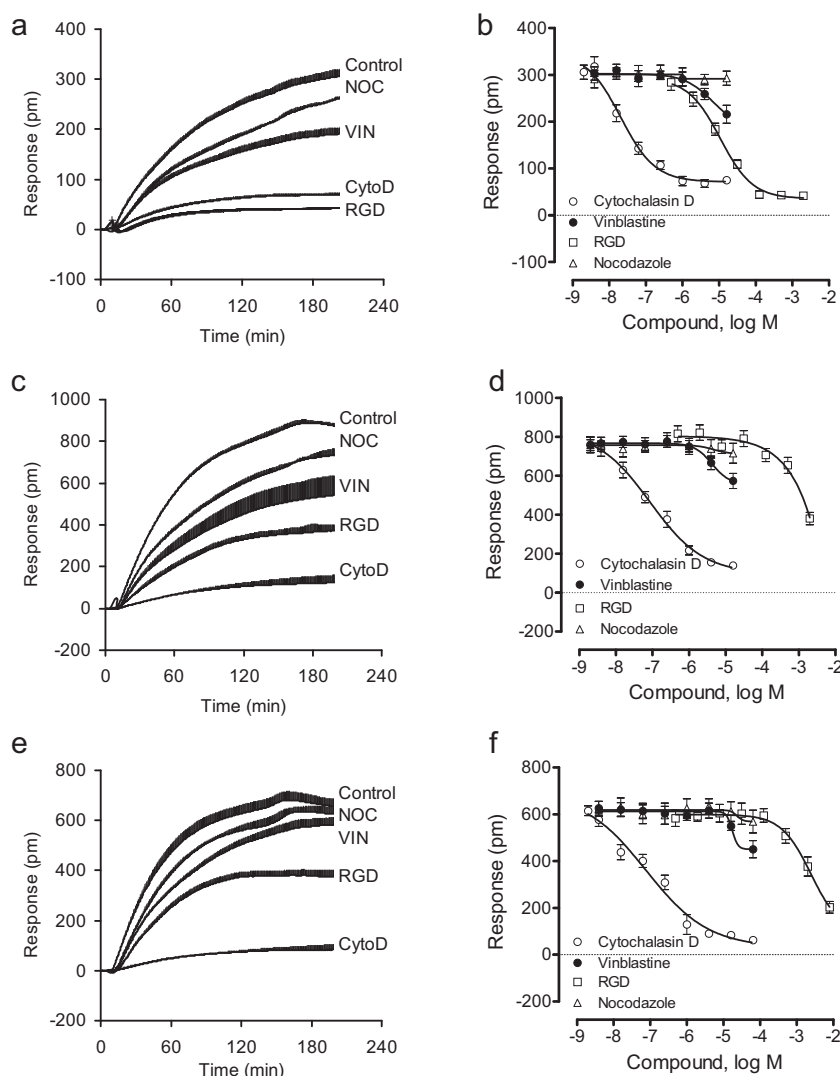


Fig. 5. DMR characterization of HEK- β_2 AR-GFP cell adhesion in the presence of different inhibitors under physiological condition. (a)–(f) The DMR of the engineered cells adherent onto different sensor surfaces: tissue culture treated (a) and (b), fibronectin coated (c) and (d), and collagen IV coated (e) and (f). (a), (c) and (e) real-time DMR; (b), (d) and (f) the DMR amplitudes at 4 h after cell seeding as a function of inhibitor dose. For all, the total number of cells added were the same; that is, 18,000 cells per well. For (a), (c) and (e), the inhibitor dose was fixed to be 10 μ M, 10 μ M, 10 μ M and 1 mM for nocodazole (NOC), vinblastine (VIN), cytochalasin D (CytoD), and RGD peptide, respectively. The data represents mean $\pm$ S.D. ($n = 12$ for a, c and e; $n = 4$ for b, d and f).

capable of disrupting cell adhesion (both cell–substrate and cell–cell) under ambient temperature. This result, together with the sensitivity to RGD treatment, suggests that the engineered cells may interact with the TCT surface through a simple integrin-mediated binding but without active actin remodeling under ambient temperature. This is also consistent with the low DMR signal obtained for the cell attachment to the TCT surface. However, on fibronectin-coated surfaces all inhibitors dose-dependently attenuated the cell adhesion (Fig. 4c and d). Cytochalasin D displayed the highest potency but only partially suppressed the adhesion. In contrast, on collagen IV coated surfaces, only cytochalasin D and RGD peptides partially suppressed the adhesion, while vinblastine had small effect, and nocodazole had not effect (Fig. 4e and f).

We then examined the cell adhesion mechanism under physiological condition. DMR inhibitor assays showed that on all three surfaces examined, similar modulation patterns were observed. In general, both cytochalasin D and RGD peptide dose-dependently and strongly attenuated the adhesion, whereas vinblastine had small effect, and nocodazole decreased the adhesion kinetics but had little on the total adhesion response (Fig. 5). Notably, the

RGD tripeptide displayed quite different potency to suppress cell adhesion; and the potency order was found to be tissue culture treated > collagen IV > fibronectin.

We further examined the TIRF images of HEK- β_2 AR-GFP cells 2 h after adhesion onto the fibronectin coated sensor surface at room temperature. Results showed that the TIRF of cells in the absence of any inhibitors tend to be more diffusive with relatively lower intensity than that in the presence of 1 μ M cytochalasin D, or 1 mM RGD peptide (Fig. 6). The presence of RGD peptide resulted in a TIRF pattern that has the highest intensity, the most uniform and the smallest size per cells. It is noting that the physical scrapping often leads to certain portions of cell clusters, rather than completely individual cells. Together, these results suggest that both surface chemistry and temperature influence the degree of cell attachment, and distinct mechanisms govern cell adhesion under different conditions. Specifically, integrin binding is involved in cell adhesion under all conditions. Except for the TCT surface under ambient condition, actin remodeling is also involved in the cell attachment under all other conditions. Furthermore, the contribution of integrin binding and actin remodeling to the overall cell

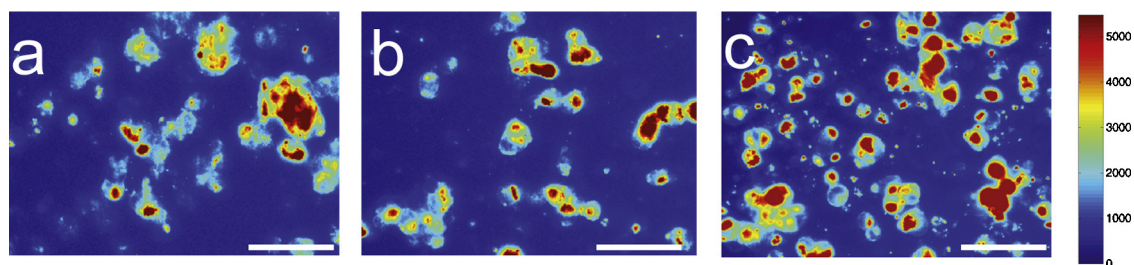


Fig. 6. TIRF images of HEK- β_2 AR-GFP cells on the fibronectin coated surface under ambient condition. The images were taken 2 h after cell adhesion onto the fibronectin-coated biosensor surface in the absence and presence of an inhibitor. (a) No any inhibitor; (b) 10 μ M cytochalasin D; (c) 1 mM RGD peptide. The inhibitors were present throughout the cell adhesion process. Image scale bar: 50 μ m.

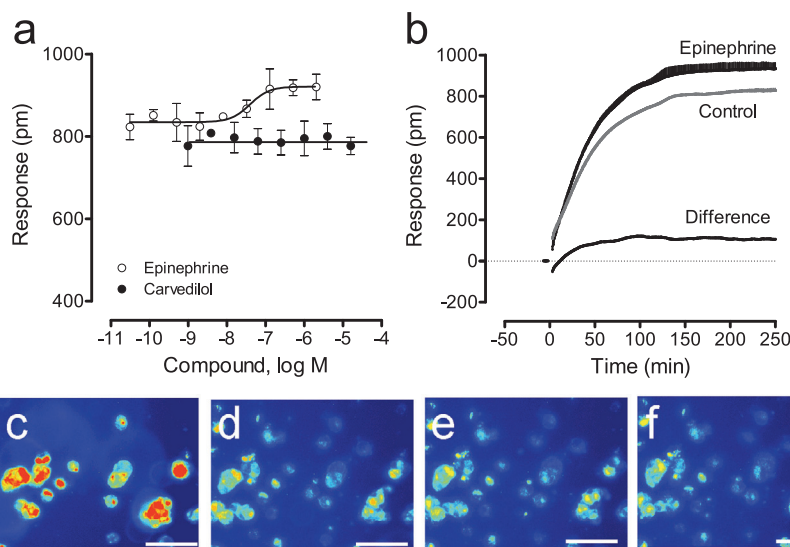


Fig. 7. Impact of the β_2 AR signaling on cell adhesion on the fibronectin coated surface under physiological condition. (a) Dose-dependent responses arising from the cell adhesion in the presence of different ligands (epinephrine or carvedilol) at different doses. (b) Comparison in real-time DMR arising from the cell adhesion in the absence and presence of 1 μ M epinephrine. (c)–(f) Representative time series TIRF images of the engineered cells in the presence of 1 μ M epinephrine throughout the cell adhesion: (c) 5 min; (d) 30 min; (e) 60 min; and (f) 120 min post cell adhesion. Image scale bar: 50 μ m. For (a) and (b) the data represents mean $\pm$ S.D. ($n = 4$ for a; $n = 12$ for b).

attachment appears to be sensitive to both surface chemistry and temperature.

3.5. The impact of β_2 -AR signaling on cell adhesion

To investigate the effect of β_2 -AR signaling on the adhesion of HEK- β_2 AR-GFP cells, we chose a panel of β_2 -AR ligands including its full agonist epinephrine, partial agonist salbutamol and salmeterol, and antagonist pinodol and carvedilol. DMR assays showed that pre-mixing of cells with the β_2 AR ligand, 1 min before cell adhesion, had distinct effect on the cell adhesion process (Fig. 7a). Salbutamol, salmeterol, pinodol and carvedilol had little effect on both the kinetics and total amount of cell adhesion signal. However, the full agonist epinephrine dose-dependently increased the total amount of cell adhesion signal, with an apparent EC_{50} of 40.0 ± 3.6 nM ($n = 4$) (Fig. 7a), which is similar to its EC_{50} to trigger DMR signal in pre-cultured cell monolayer. Student's t test suggests that the difference in the kinetic cell adhesion responses in the absence and presence of epinephrine is significant with a p value of <0.0001 (Fig. 7b). The net difference, by subtracting the adhesion DMR in the absence of epinephrine from its DMR in the presence of epinephrine, closely mimics the classic G_s -type DMR [8,9,11,12,35]. These results suggest that the activation of the β_2 AR may increase cell adhesion through classic G_s -pathway mediated signaling, a possibility worthy of further examination. Furthermore, TIRF imaging showed that the presence of 1 μ M epinephrine caused a time-dependent decrease in fluorescence

intensity (Fig. 7c–f), mostly due to epinephrine-induced internalization of the β_2 AR-GFP fusion proteins. Together, these results suggest that epinephrine activates the β_2 AR-GFP, thereby causing receptor internalization during the cell adhesion process, but also resulting in an increase in cell adhesion.

4. Discussion

Cell adhesion to a surface is an active and dynamic process. Cells use their adhesion receptors to recognize and bind to molecules in the existing ECM. Cells can also secrete and deposit their own ECM proteins to remodel the surface during cell adhesion. Cell signaling is part of cell adhesion process. The binding of cells to the ECM can activate signaling via integrins, resulting in an “outside-in” signal transduction so the cells can sense the composition, organization, density, rigidity, and biochemical and rigidity gradients of the ECM [2,3,23,28]. On the other hand, receptor signaling can occur during the cell adhesion process *in vivo*; certain signaling mediated through receptors such as GPCRs can also cause remodeling of cell adhesion, resulting in an “inside-out” signal transduction [31,41]. Here we used RWG biosensor-enabled label-free DMR assay and TIRF imaging approach to show that the activation of the β_2 AR during cell adhesion process increases the adhesion of the engineered HEK293 cells onto the fibronectin coated sensor surface under physiological condition. The engineered cell line stably expresses human native β_2 -AR whose C-terminal is fused with GFP. The β_2 AR-GFP fusion protein was used as a cell surface marker not

only to examine the cell adhesion process, but also to validate the receptor signaling and its impact on cell adhesion. The receptor activation in the presence of epinephrine was evidenced by the receptor internalization during the cell adhesion, as well as after cells form a monolayer.

Using RWG-enabled label-free DMR and TIRF imaging, we also showed that the cell adhesion is sensitive to both temperature and ECM coating, and distinct cellular mechanisms govern the cell adhesion under different conditions. Given that the complete serum media contains trace amount of ECM proteins such as fibrinogen, collagen, vitronectin, and fibronectin, and the cells can secrete and deposit their own ECM proteins during cell adhesion process, it is no surprise that under all conditions the RGD peptide displayed the greatest effect among all inhibitors examined. Furthermore, we also observed temperature and surface dependent inhibition potency of the RGD peptide which may be indicative of the density of RGD binding motif presenting by different surfaces. The lower potency the RGD peptide is the higher density is for the RGD binding motif on the surface. On the other hand, the modulation pattern of different inhibitors also suggest that under ambient condition the adhesion of HEK- β_2 AR-GFP cells onto the tissue culture treated surface is unique in that it does not involve actin and microtubule remodeling, meaning no or minimal active adhesion process taking place. Under all other conditions both actin and microtubule remodeling are required for cell adhesion, and actin remodeling is at least more important in early cell adhesion process.

5. Conclusions

We had developed a RWG biosensor-enabled TRIF microscopy with moderate spatial resolution. Together with RWG biosensor enabled label-free DMR assays, we had found that the adhesion of an engineered cell line containing fluorescent β_2 ARs at the cell surface is dependent on ECM coating and temperature, and the activation of the β_2 ARs during the cell adhesion process increases the cell adhesion.

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