Featuring work from the Hagiwara Lab, at Nanoscience and Nanotechnology Research Center at Osaka Prefecture University, Osaka Japan, whose work probes the mechanisms of multicellular pattern formation.

An *in vitro*–*in silico* interface platform for spatiotemporal analysis of pattern formation in collective epithelial cells

The fusion of an experiment *in vitro* and *in silico* supplies a powerful tool to uncover the system underlying pattern formation. This article presents dynamic analysis of 2D pattern formation in bronchial epithelial cells by controlled cell culture and a reaction–diffusion model.

As featured in:

An *in vitro–in silico* interface platform for spatiotemporal analysis of pattern formation in collective epithelial cells†

M. Hagiwara

A multicellular organization is a complex resulting from the coordinated migration of cells to form a specific pattern. The directionality of migration is governed by the mechanical and molecular dynamics of factors secreted from the cells. The mechanism underlying pattern formation is too complex to unveil by culture experiments alone. A mathematical model could provide a powerful tool for elucidating the mechanism of pattern formation by computing the molecular dynamics, which are difficult to visualize by culture experiments. However, there tends to be a gap between mathematical models and experimental research due to incongruity between the idealized conditions of the model and the experimental results. This paper presents an *in vitro–in silico* interface platform for elucidating the logic of multicellular pattern formation. Two-dimensional collective cell pattern formation was developed using normal human bronchial epithelial cells. Then, geometrical control of collective cells followed by feedback iteration was used to bridge the gap between the mathematical model and *in vitro* experiments. The mechanisms underlying the pattern formation of bronchial epithelial cells were evaluated using a reaction-diffusion model. The results indicated that differences in the diffusion rates of the activator and inhibitor determine the direction of collective cell migration to form a specific pattern.

**Introduction**

Cells form a specific pattern by themselves in order to function as a tissue. Pattern formation *in vivo* involves a sophisticated process coordinated by a number of communications via a range of secreted molecular factors, yet the pattern developed is always the same, with little variation.1–3 Collective cell migration largely contributes to pattern formation,4–8 but the mechanisms involved in the development of multicellular organization are still poorly understood due to the complexity of the underlying system.

It is not known how cells sense their position within the tissue structure or determine the direction of development. Understanding these mechanisms is crucial for pathological study and the development of regenerative medicine. Lung branching is one of the most complex tissue patterns.9 Research during the past few decades has helped to identify some of the key morphogens involved in forming the branching structure,10–13 but the logic of pattern formation is still not clear. In fact, various mathematical models have been proposed to simulate the branching formation;14 some models are based on mechanical interactions among epithelial cell sheets,15,16 while others are based on chemical reactions.17–19 One major reason for the enduring lack of clarity regarding the mechanisms of pattern formation is that there have been few successful *in vitro* experimental models. Unlike *in vivo* experiments, *in vitro* experiments can use a limited number of cell types to focus on individual physiological processes,20,21 but the...
considerable experimental noise due to procedural variation during operation obscures the underlying system. It is impossible to decode the mechanisms of pattern formation using experimental systems in which the pattern developed varies from one experiment to the next.

Here, an \textit{in vitro} two-dimensional (2D) experimental model of pattern formation in bronchial epithelial cells was developed. In 2D culture, the initial culture conditions can be controlled by microfabrication, and directional collective cell movements were generated under specific conditions. Using the developed \textit{in vitro} experimental model, an \textit{in vitro–in silico} interface platform was introduced to bridge the gap between biological experiments and mathematical theory (Fig. 1). Controlling the initial culture conditions \textit{in vitro} can reduce the experimental noise to improve the reproducibility of the developed pattern formation. Then the remaining inherent experimental errors such as cell viability variations can be compensated by repetitive experiments. Multiple initial conditions are examined to avoid accidental matching of \textit{in vitro} results and \textit{in silico} results in order to validate the mathematical model itself as well as the model parameter. This approach can allow the \textit{in vitro} conditions to mimic the idealized conditions \textit{in silico} and enhances the agreement of computational simulations with \textit{in vitro} experiments. Spatiotemporal analysis of morphogens was performed to uncover the logic of pattern formation by collective cells of the bronchial epithelia. The mechanisms that determine the directionality of collective cell migration were revealed based on a reaction-diffusion (RD) model starting from various collective cell geometries.

**Materials and methods**

**Cell preparation**

Normal human bronchial epithelial cells (NHBEs) were obtained from LONZA (Walkersville, MD) and cultured using the BEGM™ Bullet Kit (LONZA) supplemented with 50 IU ml$^{-1}$ penicillin and 50 μg ml$^{-1}$ streptomycin (Thermo Fisher Scientific Inc., MA). NHBEs were only used up to passage 7 in order to reduce the amount of variation in the developed patterns.

**Initial cell culture control**

To develop branched pattern formation of NHBEs, Matrigel growth factor reduced (Corning Inc., Corning, NY) was used to
coat cells with a high cell concentration gradient. The Matrigel formed a hydrogel after 25 minutes of incubation and then BEGM medium was added.

Photolithography was employed to control the initial collective cell geometry and concentration (Fig. S1(a), ESI†). Before the fabrication process, a silicon substrate was soaked with a mixture of hydrogen peroxide and sulfuric acid (1:2, v/v) for 10 min to clean the substrate, which was followed by rinsing with deionized water for 5 min twice. Then, the substrate was dehydrated in an oven at 140 °C for 20 min. A negative photoresist coating (SU-8 2100; MicroChem Corp., MA) was spin-coated onto the silicon substrate with approximately 150 μm thickness. Ultraviolet light irradiation was then conducted over a pattern. PDMS was poured onto the mold at a lower thickness of 10 μm, and the PDMS membrane was cured. Once the ethanol was evaporated in an oven at 80 °C for 10 min, the adhesion of the PDMS membrane to the dish was irreversible transition of Y from the low state to the high state. At this point, Y is not affected by morphogen concentration, and the collective cell geometry is fixed in the area. Morphogens A, H, and S diffuse to the surrounding area at rates $D_A$, $D_H$, and $D_S$, respectively. The parameters $\rho_A$, $\rho_H$, and $\rho_M$ are determined based on the genetic programming of cells, while $d$, $e$, and $f$ are adjustment parameters for switching the value of Y. Other parameters are determined by environmental factors such as temperature, pressure, and extracellular matrix (ECM) stiffness. For example, if the stiffness of the ECM is increased, the mesh size inside the ECM becomes smaller and $D_A$, $D_H$, and $D_S$ become lower.

Parameter refinements were conducted so that the pattern similarity between the in vitro experimental results and the RD model simulation results increased. The calculation methods of pattern similarity are described in the following section. Each parameter differently contributes to the pattern shape; for example, $\rho_A$ and $\rho_H$ are change the spatial interval of branches, and $\varepsilon$ has correlation with the branch mode. All parameters are dimensionless, and after a number of refinements, the final parameters were set as follows: $\epsilon = 0.04 \pm 7\%$, $\mu = 0.1315$, $\nu = 0.04$, $\rho_A = 0.01$, $\rho_H = 0.00015$, $c_0 = 0.02$, $\gamma = 0.02$, $\varepsilon = 0.08$, $d = 0.013$, $e = 0.1$, $f = 10$, $D_A = 0.02$, $D_H = 0.18$, $D_S = 0.12$.

**Calculation of branching pattern similarity**

Normalized cross-correlation (NCC) was calculated after image processing to quantify the degree of similarity in the developed branching pattern formation. To determine the pattern, image processing was conducted using Imagej (Fig. S2(a–f), ESI†). First, the edges of individual cells in the phase contrast image were detected by finding the maximum colour gradient points using the “Find edges” function in Imagej. Then, the image was changed to a binary image (“Make binary”). The edge line was dilated followed by erosion (“Close”) and the small holes were filled (“Fill holes”) to identify the pattern area. Finally, the erosion operation was conducted again, followed by dilation (“Open”) to reduce signal noise.

The similarity defined by NCC (RNCC) was then calculated between the template with $M \times N$ pixel size and the target image as follows:

$$R_{\text{NCC}} = \frac{\sum_{j=0}^{M-1} \sum_{i=0}^{N-1} I(i,j)T(i,j)}{\sqrt{\sum_{j=0}^{M-1} \sum_{i=0}^{N-1} I(i,j)^2 \times T(i,j)^2}}$$

where $I(i,j)$ and $T(i,j)$ are the intensity values of the target and template images, respectively.
where $T(i, j)$ indicates the pixel value of the template image at position $(i, j)$, and $I(i, j)$ indicates the pixel value of the target image at position $(i, j)$. With increasing similarity between the two patterns, RNCC moves closer to 1. The template was moved to all areas of the targeted image with a raster scan, and RNCC was calculated at each position (Fig. S2(g), ESI†). The maximum RNCC was defined as the pattern similarity between the template and target images.

### Results

**In vitro experiments of 2D pattern formation in bronchial epithelial cells**

In our previous study, we succeeded in developing a hollow branching structure *in vitro* that is commonly observed in the lung airspace, using only NHBEs. However, it is difficult to precisely repeat the same experimental conditions in three-dimensional (3D) culture due to operational variation and this variation is amplified with cell growth. These uncontrollable conditions give rise to a large amount of experimental noise and obscure the logic underlying the system. In addition, it was difficult to observe the patterns of branches widely spreading three-dimensionally. On the other hand, the current study found that collective cells formed specific patterns in 2D when NHBEs were set on a dish with a high cell concentration gradient covered by the extracellular matrix (ECM) (Fig. 2(a) and Video 1, ESI†). At the beginning, cells collectively migrated in a dynamic manner to form a finger-shaped structure similar to that formed by epithelial Madin–Darby canine kidney (MDCK) cells under specific conditions. Then the pattern of collective NHBEs bifurcated after the finger reached a certain length to form a “branch” structure. Unlike MDCK cells, the collective NHBEs continuously migrated without any leader cell. In contrast, cells grew without direction (Fig. 2(b)) when there was no ECM or when cells were homogeneously distributed (Fig. 2(c)).

Unlike 3D culture, the initial culture conditions can be controlled with relative ease for 2D culture, and observation of the developed pattern can be achieved using a standard microscope.

**Fig. 2** 2D experimental model to generate pattern formation in bronchial epithelial cells. (a) When cells were placed with a high concentration gradient and were covered by the ECM, they showed 2D branching morphogenesis. (b) When cells were placed at a high concentration without any ECM, they proliferated and migrated without direction. (c) When cells were homogeneously distributed with the ECM, they barely migrated. Scale bars, 500 μm. (Video ESI†).

Besides, the system does not require heterotypic cell–cell interactions; epithelial cells can produce branches by themselves and therefore it is suitable for analysing the mechanisms underlying branching pattern formation. A simplified 2D platform also has great advantages in computational simulation. The calculation time in 2D can be significantly reduced comparing to 3D and it leads to prompt feedback between *in vitro* and *in silico* experiments. Control over the collective cell geometry can be achieved for 2D culture by simple photolithographic technology (Fig. S1, ESI†). Through employing this engineering technique, the initial collective cell conditions, including the collective cell geometry, and the cell concentration can be controlled to reduce the experimental noise. The initial size of the collective cells was fixed at 1 mm² for all geometrical shapes. As a result, the branching pattern becomes repeatable (Fig. 3(a)). With controlled initial conditions, the similarity of the pattern formation from one experiment to the next is higher than that without controlled conditions (Fig. 3(b) and Fig. S3, ESI†). When the initial conditions were not controlled, the collective cell size and concentration showed large variations among experiments, regardless of the control initial culture conditions by photolithography and from non-controlled culture at 0 hours, 24 hours, 48 hours, and 72 hours. Branch shapes were generated at all tips of the triangle starting from triangle shapes at 0 hours (upper left) and 3 samples after 72 hours of culture. Branch shapes were generated at all tips of the triangle shapes and the characteristics of the pattern are similar for all samples. Scale bar, 300 μm. (b) The similarity of the developed pattern starting from controlled initial culture conditions by photolithography and from non-controlled culture at 0 hours, 24 hours, 48 hours, and 72 hours. The repeatability of the pattern formation was higher under the controlled conditions than without control at all points ($p < 0.01$). The similarity was calculated by normalized cross-correlation after the image changed to the binary image (Fig. S5, ESI†). (c) Pattern formation starting from aligned collective cells in a long straight line. Scale bar, 500 μm. (d) Histogram of the initial branching angle starting from a long straight line. The direction tended to be perpendicular to the original straight line ($n = 50$).
which increased daily. On the other hand, when the collective cell geometry was triangular, which cannot be achieved without microfabrication, it was observed that a branched shape occurred at each tip of the triangle. The direction of the branches could be observed more quantitatively when the initial collective cells were set as a long line (Fig. 3(c and d)). The branches clearly showed a tendency to grow in a direction perpendicular to the original straight edge line.

**In silico experiments involving the RD model for branching pattern formation**

The advantage of controlling the initial culture conditions is that not only can the experimental noise be reduced but the experimental conditions can also be matched with the simulation conditions. This allows the model to obtain fair feedback from experiments to refine the model parameters. If the model can achieve reasonable agreement with the experimentally obtained pattern formation under multiple initial conditions with fixed parameters, it can be justified that the physical phenomena accounted for in the model are weighted more among the potential factors. Once the model is experimentally validated, dynamic simulations can be conducted to decode the logic underlying the system of multicellular pattern formation.

The RD model is a mathematical model used to simulate self-organized pattern formation by computing the spatial distributions and dynamic reaction of secreted morphogens. However, there tends to be a gap between mathematical models and in vitro experimental research due to incongruity between the idealized conditions of the theoretical model and the actual experimental conditions. Now that the experimental conditions can be controlled and only epithelial cells are used for epithelial pattern formation, the conditions of in vitro experiments can step much closer to the idealized in silico experiments. The in vitro experimental system is suitable for the validation of whether the RD model can describe the mechanisms that underlie pattern formation by collective cell migration in bronchial epithelia.

The RD model proposed by H. Meinhardt was examined to simulate the pattern formation developed by the collective cells of bronchial epithelia. To express the inherent variation in the system, 7% random fluctuation was included in the morphogen’s reaction rate parameter ($c$ in eqn (1) and (2)). After repetitive feedback from experimental patterns starting from multiple geometries to refine the parameters, the simulated patterns reached good agreement with actual branch formation under all conditions. The characteristics of the initial branched pattern, such as the position and direction, corresponded to those in experiments under all conditions, and branching occurred at the tips of the initial geometries. The direction tended to be perpendicular to the original edge line. The same tendency could be observed by RD simulations. The static and dynamic results indicate that the branching was initiated from tips (Video 3, ESI†). The static and dynamic dynamics of the biomarker $Y$ value. The same tendency as in (b) can be observed by RD simulations. The $Y$ value was higher at the tips and at the edge of the triangle at the beginning, which indicates that cells move to the edge at first. Then, the triangle center became higher and branching was initiated from tips (Video 3, ESI†). Scale bars, 500 μm.

**Fig. 4** Comparison of branch shape pattern formation between in vitro experimental results and the simulated pattern by the reaction-diffusion model. (a) Comparison of the branched pattern starting from multiple initial conditions. After repetitive feedbacks from in vitro experiments to refine the parameters, the simulated patterns reached good agreement with actual branch formation under all conditions. The characteristics of the initial branched pattern, such as the position and direction, corresponded to those in experiments under all conditions, and branching occurred at the tips of the initial geometries. The direction tended to be perpendicular to the original edge line. (b) Time lapse imaging of branching morphogenesis in vitro, starting from the triangle shape. Cells migrated to the tips of the triangle shape at the beginning, and there were lesser number of cells at the center of the initial triangle shape. Then, the cells reconstructed the triangle shape and developed branches at tips (Video 2, ESI†). (c) Simulated dynamics of the biomarker $Y$ value. The same tendency as in (b) can be observed by RD simulations. The $Y$ value was higher at the tips and at the edge of the triangle at the beginning, which indicates that cells move to the edge at first. Then, the triangle center became higher and branching was initiated from tips (Video 3, ESI†). Scale bars, 500 μm.

**Spatiotemporal analysis of morphogen distribution for branching pattern formation**

Once the mathematical model is verified and the simulation accuracy is improved by refining the parameters, the model can be used for the spatiotemporal analysis of morphogen
distribution, which cannot be done in cell culture experiments, in order to uncover the logic underlying the system determining pattern formation. In the RD model, the activator and inhibitor are the main factors involved in the self-organization of pattern formation. The activator drives cells to migrate and/or proliferate and increases the amount of the activator by an autocatalytic function as well as increasing the amount of inhibitor, while the inhibitor represses activator function. To develop self-organized pattern formation, short-range activation and long-range inhibition are required,\(^\dagger\) that is, the activator diffusion rate must be lower than the inhibitor diffusion rate.

First, the reason why 2D pattern formation of collective bronchial epithelial cells can be developed only when cells are set on a dish with a high cell concentration gradient covered by an ECM can be explained by the RD model (Fig. 5(a–c)). When cells are set on a dish without an ECM, the diffusion rates of the activator and inhibitor are inversely proportional to their molecular sizes and the molecular weights are usually in the same order. If the diffusion rate of the inhibitor is not high enough to spatially restrict the activator function, the cells grow homogeneously (Fig. 5(a) and Video 4, ESI\(^\dagger\)). On the other hand, if the cells are covered by the ECM, the fiber network structure in the ECM hinders molecular diffusion and increases differences between the diffusion rates of the activator and inhibitor. However, although the ECM was coated onto the cells, the inhibitor secreted from the cells interrupted the growth of the surrounding cells when they were homogeneously distributed (Fig. 5(b)). Under these conditions, the inhibitor was present in the entire culture area and therefore the cells could not undergo directional migration. By contrast, the effect of short-range activation can be enhanced by setting the cells at a single point while the interruption of long-range inhibition from neighbour cells is kept the same. As a result, branches can be developed under the limited condition that cells are set with a high concentration gradient covered by the ECM (Fig. 5(c) and Video 5, ESI\(^\dagger\)).

Next, to investigate the logic of why initial collective cells tend to migrate in a perpendicular direction at a specific position, a dynamic simulation of morphogens was performed using the RD model (Fig. 6(a) and Video 6, ESI\(^\dagger\)). Molecules, in general, diffuse from an area of higher concentration to an area of lower concentration. Therefore, when the collective cells are aligned in a straight line, as shown in Fig. 4(d), morphogens secreted from cells diffused in a direction perpendicular to the edge of the collective cells, which leads cells to likewise grow in a direction perpendicular to the edge of the collective cells. In addition, the relationship between activator and inhibitor diffusion rates results in determining the position of collective cell migration to form a branch shape. The activator concentration slightly varied along the edge of the collective cells due to the inherent variation in cell viability. Once the activator concentration increased at a certain spot, the concentration rapidly increased further because of autocatalysis at that spot (Fig. 6(b), Fig. S4 and Video 7, ESI\(^\dagger\)). The inhibitor concentration was also increased by the activator at that spot.

Because of long-range inhibition and short-range activation, the inhibitor suppressed the activator concentration within a broad range. As a consequence, the activator concentration near the activator peak was lower than that at other positions. Therefore, cells could not grow near a branch and the subsequent branches were generated while maintaining a certain distance from the other branches.

The logic of why branches were generated at the tips of the collective cell geometries and why the cells faded away from the center of collective cells at the beginning (Fig. 4(b and c)) was

---

**Fig. 5** Morphogen distribution computed by the RD model for analyzing why 2D branching morphogenesis can be developed only when NHBEs are set on a dish with a high cell concentration gradient covered by an ECM. (a) When cells are placed at a high concentration gradient without the ECM, the differences in diffusion rates of the activator and inhibitor are too small to generate branch patterns (Video 4, ESI\(^\dagger\)). (b) When cells are homogeneously distributed, they are interrupted by long-range inhibition from surrounding cells and collective cell migration is restricted. (c) When cells placed at a high concentration gradient were covered by the ECM, branch patterns could be generated (Video 5, ESI\(^\dagger\)).
The inhibitor, and directional migration of the collective cells was generated at the tips first, as shown in Fig. 4(b and c). Inhibitor geometries to avoid areas containing a high concentration of cells tend to move to either the tips or edges of the collective cell due to the low concentration of the inhibitor. Then, followed by an increase in the concentration of the activator at the activator concentration there increases at the beginning, relatively higher at the tips of the collective cell geometries and surrounding cells. As a result, the effect of the activator is lesser number of surrounding cells. On the other hand, the effect of the inhibitor is spatially limited due to its low diffusion rate. Therefore, the inhibitor concentration is higher at the center of the triangle but relatively lower at the tips of the triangles. The effect of the activator then relatively increased to generate branches at the tips (Video 8, ESI†).

The experiments described in this paper focused on pattern formation resulting from the collective cell migration of NHBEs, while the pattern formation of lung branching in the body is based on 3D morphogenesis. In 3D branch formation, various physical phenomena such as the folding process of epithelial sheets occur to form branching morphogenesis. Therefore, the mechanisms of monolayer pattern formation presented represent the fundamental drivers of the sophisticated physiological pattern of lung branching, but the developed experimental system and results cannot be directly applied to the actual lung pattern. On the other hand, it is interesting that the pattern that developed in 2D still had a “branched” shape with multiple bifurcations. The results indicate the possibility that the collective cell migration governed by an activator–inhibitor system has a certain contribution to the 3D pattern formation of lung branches. The key morphogens in the RD system of lung branching pattern formation have not been identified yet, but many previous studies have indicated that sonic hedgehog (SHH), bone morphogenetic protein 4 (BMP4), fibroblast growth factor 10 (FGF10), and sprouty 2 (SPRY2) are the morphogen candidates. By further elucidating the key morphogens, we can move forward to a fuller understanding of lung branch pattern formation.

Conclusions

It is possible to validate whether the physical dynamics described in the mathematical model are dominant determinants of pattern formation by using an interface that brings the in vitro experimental environment closer to the in silico experimental environment. Simple but robust engineering technology
significantly helps to bridge the in silico and in vitro experiments by reducing experimental variation and noise in vitro. In addition, the accuracy of the simulation can be improved considerably by repetitive feedback from pattern formation developed in vitro under multiple different initial conditions. Then, spatiotemporal analysis of molecular dynamics would become the most powerful tool for elucidating the logic of pattern formation, as presented in this paper. Moreover, an accurate mathematical model has the potential to aid in designing initial culture conditions so as to lead cells toward specific multicellular synthesis of different approaches such as those involving biology, mathematics, physics, and engineering would aid in resolving complex questions in biological studies and bring us forward to the study of new fields.

Acknowledgements

This work was financially supported by a Grant-in-Aid for Young Scientists (A), Challenging Exploratory Research from the Japan Society for the Promotion of Science (15K13325, 15H05512), The Nakajima Foundation and the Program to Disseminate Tenure Track System, MEXT, Japan.

Notes and references

23 H. Meinhardt, Differentiation, 1976, 6, 117–123.