Biological soliton in multicellular movement

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Solitons have been observed in various physical phenomena. Here, we show that the distinct characteristics of solitons are present in the mass cell movement of non-chemotactic mutants of the cellular slime mould Dictyostelium discoideum. During starvation, D. discoideum forms multicellular structures that differentiate into spore or stalk cells and, eventually, a fruiting body. Non-chemotactic mutant cells do not form multicellular structures; however, they do undergo mass cell movement in the form of a pulsatile soliton-like structure (SLS). We also found that SLS induction is mediated by adhesive cell-cell interactions. These observations provide novel insights into the mechanisms of biological solitons in multicellular movement.

Results

Non-chemotactic mutants of Dictyostelium discoideum exhibit a soliton-like structure in multicellular movement. In wild-type cells of the slime mould Dictyostelium discoideum, a starvation signal triggers a developmental programme. Following exhaustion of their bacterial food source, the erstwhile random movement of the amoeboid cells is orchestrated to yield an aggregation of cells, mediated by the self-secreted extracellular chemoattractant, cAMP. The resultant migratory, multicellular structure is known as a slug. Transformation of the slug occurs within 8 h and results in a fruiting body that consists of two differentiated cell types, spore cells and stalk cells⁶ (Fig. 1a, Supplementary video S1).

We have isolated D. discoideum mutants that lack all chemotactic activities, leaving them unable to proceed down developmental pathways that require cell aggregation⁷⁻⁸. Soliton-like structures (SLSs) were observed in the KI-5 and KI-10 mutants within 6 h following the consumption of bacteria (Fig. 1b, Supplementary video S2). Since the features of the SLSs were not distinguishable between the two mutants, we only describe the KI-5 phenotype in this manuscript. SLSs emerged around 6 h after exhaustion of the bacterial food supply and persisted for 48 ± 3 h in 5 independent experiments. (See a typical time course of SLS in Supplementary Fig. S1 and Supplementary video S3). Maintenance of the SLS structure despite collision with other SLSs is characteristic of a soliton wave (Fig. 1c, Supplementary video S4). The formation of SLSs is dependent on cell density such that the higher the density of the cells, the larger the SLS formed (Fig. 1d); however, an excessively high
density prevents SLS formation (Fig. 1d). Moreover, as the duration of cell starvation increased, SLS size and number decreased (Supplementary Fig. S1, Supplementary video S3). Despite the variability in size, SLSs moved at a constant velocity of approximately 20 μm/min, which is twice the rate (10 μm/min) of starved wild-type cells.9,10 Since SLS velocity is comparable to that of prespore or pre-stalk cells in the slug, SLS cell motility may represent a distinct aspect of differentiated cell movement.

SLSs depend on the cAMP signalling genes but are independent of extracellular cAMP and DIF-1. In D. discoideum, the sensory signalling pathway used during development involves cAMP synthesis and signalling through the serpentine cell surface receptor, trimeric G protein12. Therefore, mutants that lack the genes responsible for these processes—carA; the cAMP receptor, gapB; the trimeric G protein α subunit, and acaA; an adenylyl cyclase—are defective in cell aggregation and dependent processes due to lack of cAMP production13-15. These null mutants showed few signs of SLS formation, indicating that the cAMP signalling pathway is required for SLS formation (Fig. 2a, Supplementary video S5, S6, S7). We have shown that KI-5 and KI-10 are defective in cGMP metabolism in response to extracellular cAMP stimuli despite possessing the machinery for cAMP receptor-dependent cAMP production7,16. During cell aggregation, extracellular cAMP concentrations of an order of magnitude less than the micromolar level play a critical role such that cAMP concentrations of a millimolar order of magnitude substantially inhibit morphogenesis18. Next, in order to determine if extracellular cAMP is required for SLS formation, KI-5 cells were exposed to extremely high concentration and depletion of cAMP. Under the condition with the high concentration of 1 mM cAMP, formation of SLSs was not disturbed (Fig. 2b). Furthermore, KI-5 cells were plated on non-nutrient agar with 4 mM caffeine and conditioned medium prepared from the supernatant of wild-type cells in starvation (see Methods). Since caffeine is an adenylyl cyclase inhibitor in D. discoideum and the conditioned medium contains enough phosphodiesterase activity to degrade most of the physiological amount of extracellular cAMP19,20, little cAMP was measured in the extracellular fraction of KI-5 cells starved for 12 h with 5 mM caffeine and conditioned medium (Table I; see Supplementary methods). With 5 mM caffeine and the conditioned medium, SLSs were normally formed without disturbance (Fig. 2c). These observations indicate that extracellular cAMP is not required for SLS formation. Another morphogen, DIF-1, is a stalk cell-inducing factor and a modulator of chemotactic activity21,22. In the presence of high DIF-1 concentrations, SLSs developed normally (Fig. 2d).

SLSs are a dynamically stabilized structure. Cell motion in SLSs was investigated at a higher magnification and with RFP (Red fluorescent protein)-expressing KI-5 cells generated by transformation with...
monomeric RFP expression vector. Motile cells in front of the wave moved randomly and did not appear to migrate into the SLS but instead were incorporated passively (Fig. 3a, 3b & Supplementary video S8, S9, S10). At the same time, some cells at the rear of the SLS were left behind and reverted to random motion. The number of RFP-expressing incorporated cells was comparable to the number of cells left behind the SLS (Fig. 3b). In 3 independent SLSs, the number of highly RFP-expressing incorporated cells and the number of cells left behind were nearly the same (Table II). Furthermore, removal of cells in front of a moving SLS showed that the thickness of the SLS was reduced and eventually disappeared (Fig. 3c, Supplementary video S11). These results indicate that SLSs do not constitute a static, independent cluster of cells but are dynamically stabilized structures that continuously incorporate cells in front and leave cells trailing behind.

Collision of two independent SLSs. The SLS structure is stable regardless of collision other SLSs (Fig. 1c, Supplementary video S4). In order to investigate cell behaviour in colliding SLSs, SLS collisions were recorded at high magnification. In head-on collisions, the SLSs appeared to coalesce as clusters and glide in a circular motion (Supplementary video S11). They did not always maintain their original orientation, with some switching orientation to the opposite direction. This was confirmed in RFP-expressing KI-5 cells (Fig. 4, Supplementary video 12, 13). Some cells in each soliton maintained (arrow in Fig. 4) or reversed (arrowhead in Fig. 4) direction after the collision. These results indicate that, in SLS collisions, cells merge into each other as a cluster and glide, but separate in 2 directions independent of the original direction.

End-to-end adhesion may play a major role in the distinct and stable directional locomotion of SLS. Next, cells comprising SLS were moderately dispersed and placed under an agar overlay (Fig. 5a, Supplementary video 14). A population of cells contacted each other in an end-to-end manner and moved in a tandem structure, similar to cell movement in slugs. Within this structure, two or more cells were often found to concatenate although they sometimes formed a broken line (arrows in Fig. 5a). These end-to-end concatenations and

Table I | cAMP concentration in the extracellular fraction of KI-5 cells with 5 mM caffeine and conditioned medium

| Addition          | cAMP concentration in supernatant (pmol/mL) |
|-------------------|--------------------------------------------|
| No caffeine and no CM | 1.70 ± 0.33                                |
| 5 mM caffeine and CM | 0.09 ± 0.06                                |

The results are shown as the mean ± S.D. of triplicate determinations of 3 independent experiments. CM = conditioned medium (see Materials and methods).

Table II | The number of highly RFP-expressing cells that are incorporated into and left behind a SLS

|                | incorporated cells | cells behind | ratio |
|----------------|--------------------|--------------|-------|
| Experiment I   | 17                 | 17           | 1.0   |
| Experiment II  | 17                 | 20           | 0.8   |
| Experiment III | 18                 | 20           | 0.9   |

Ratio indicates the relative value of the number of RFP-expressing incorporated cells to cells behind a SLS per movie.
subsequent tandem structure movements were observed on occasions when one cell encountered other cells (arrowheads in Fig. 5a). Furthermore, SLS formation was never inhibited by 5 mM EDTA at which concentration calcium-dependent cell adhesion is abolished and, thus, cell aggregation in wild-type cells is blocked by inhibition of side-by-side cell adhesion24 (Fig. 5b). These observations suggest end-to-end adhesion may play a major role in the distinct and stable directional locomotion of SLS, and calcium-dependent cell adhesion is not involved in SLS formation.

Discussion
In this manuscript, we show that the distinct characteristics of solitons are observable in the mass cell movement of non-chemotactic mutants of a cellular slime mould. Upon starvation, wild-type cells of the cellular slime mould \textit{D. discoideum} stop growing and initiate spontaneous formation of multicellular structures, accompanied by chemotaxis-dependent cell aggregation. This results in cell differentiation. In contrast, non-chemotactic mutants do not undergo aggregation; however, they do undergo a mass cell movement that possesses the characteristics of a soliton. They form pulsatile SLSs that retain their shape and move at a constant velocity. Furthermore, the shape of each SLS remains unchanged after collisions with other SLSs (Supplementary video S4, S12, S13). These observations provide the first insight into soliton-related behaviours in eukaryotic multicellular movement.

In myxobacteria, starved cells exhibit a highly organized periodic pattern of accumulations that move as traveling waves called ripples25,26. Two sets of waves appear to move in opposite directions with the same wave length and speed, like SLSs; however, the mechanism of wave movement in collisions is rather different from that of SLSs. When two ripple wave fronts collide, cells move in the reverse direction. Consequently, the outgoing waves consist of a combination of individuals from both incoming waves. This gives the appearance of waves passing through one another. In the collision of SLSs, each cell maintains its direction forward or reverses. This feature is distinctly different from that of SLSs.

Null mutants that lack \textit{carA}, \textit{gpaB}, and \textit{acaA} are defective in SLS formation. These observations indicate that cAMP signalling is responsible for SLS formation. On the other hand, extracellular cAMP and exhaustion of cAMP by caffeine had no effect on SLS formation. Thus, cAMP signalling is important for the transition from vegetative to starved stage, but is not necessary for SLS maintenance. Furthermore, since KI-5 is defective in intracellular cGMP metabolism in response to cAMP stimuli, cGMP is not necessary for SLS emergence and maintenance.

In \textit{D. discoideum}, cell adhesion molecules mediate cell-to-cell contact during development. DdCAD-1 and gp80 are responsible for spatio-temporal cell adhesion. DdCAD-1 localizes primarily on cells at the tip and outer margin of cell streams, whereas gp80 localizes to contact regions between cells within well-developed streams27,28. DdCAD-1 is enriched in filopodial structures at the cell contact region, suggesting DdCAD-1 plays an important role at the leading edge of migrating cells. However, since we found that SLS formation and maintenance are EDTA-resistant phenomena, the adhesion molecule is not likely to be responsible for SLS, because DdCAD-1 is not adhesive under calcium-depleted conditions in the presence of EDTA. In the end-to-end adhesion in SLS, calcium-resistant adhesion molecules may serve as a target marker for contact. Based on these observations, the explanation for dynamically stable movement of SLS may be elucidated through the specific localization of calcium-resistant adhesive molecules such as gp80.

Based on our observations, we speculate that the formation of SLSs is initiated by random collisions in early-stage starvation. Since SLS is only observed under starvation conditions, we suggest cell polarization is important for SLS. Once cells become polarized, their movement becomes aligned and propagates as a cluster by end-to-end adhesion. To understand if the characteristics of biological solitons reported in this study can apply to multicellular movement in not only \textit{D. discoideum} but also general developmental morphogenesis, further mathematical treatment providing theoretical validation of the soliton nature of the SLSs is required.

Methods
Culture conditions and SLS observation. The parental strain of wild-type \textit{Dictyostelium discoideum} XP55 and the non-chemotactic mutants KI-5 and KI-10...
cells were cultivated as described. For each mutant, 1.0 × 10^5 cells were inoculated on a 9-cm 1/3 SM plate (0.3% glucose, 0.3% bactopeptone, 1.5% agar, and 40 mM KH2PO4/Na2HPO4, pH 6.0) with an aliquot of Klebsiella aerogenes suspension. At 2 days incubation at 21 °C, bacterial exhaustion was observed. Time-lapse videos were then taken at 21 °C with a digital stereo microscope (Olympus SZX12). Conditioned medium was prepared from the extracellular supernatant of wild-type cells (1.0 × 10^5 cells/mL) incubated in phosphate buffer (PB: 10 mM Na/K phosphate buffer, pH 6.5) for 12 h at 21 °C. To look at the effect of cell density on SLS, cells were cultivated on 1/3 SM plates at 21 °C. They were harvested and washed 3 times with PB using a repeated centrifugation procedure (350 g for 2 min 3 times). Conditioned medium was added to the wild-type suspensions to adjust the cell density indicated in the figure legend. Higher-magnification time-lapse videos were taken using a modified agar overlay method as described. Pictures and videos were taken using a contrast and fluorescence microscope (Zeiss Axiosvert 200M).

Transformation of KI-5 cells. Transformation of KI-5 cells was performed as described previously with a few modifications. Details are provided in the Supplementary methods.

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