

Accurate tracking of chemotactic cancer cell migration – basic insights in metastasis

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Introduction

Worldwide, cancer is responsible for around 10 million mortalities every year [1]. The main cause for cancer-related death is not the primary tumor, but cancer metastasis: malignant cells invade tissues other than where the primary tumor is located and spread in those tissues [1]. Fundamental cancer research is one of the key factors in the World Health Organization's strategy to reduce cancer mortality [1], and investigating the underlying mechanisms for metastasis can provide important information for the main clinical problem.

In cancer metastasis, the migration of single cells is largely affected by chemical cues, which provide direction for migration to the cells via chemotaxis [2]: the single cells migrate along a concentration gradient of a chemoattractant [3]. Various chemoattractants have been described for commonly investigated cell lines. HeLa cells (cervical cancer cells) have been reported to migrate towards higher concentrations of for instance fibronectin [4] and stromal cell-derived factor 1 α [5]. The rat glioma cell line C6 has been described to be attracted by bradykinin [6]. A commonly-used supplement in cell culturing that contains all of the aforementioned chemoattractant molecules is fetal bovine serum (FBS; the blood serum of an unborn calf) [7-9]. Therefore, FBS provides an interesting and straightforward model system for the chemical cues provided to cancer cells.

However, in order to study the behavior of cancer cells in response to the chemical cues, accurate cell tracking in an environment with a chemoattractant gradient is required. Multiple cell culture vessel designs have been made to provide a chemoattractant gradient to the cells [10-12], but the imaging

and monitoring of cells in these vessels can provide practical issues. Currently, the most commonly-used live-cell imaging setup is a microscope with sufficient magnification and a stage-top incubation box to regulate the culture conditions. Although these microscopes have the required optical properties for accurate imaging and cell tracking, there are practical issues when using such a setup for live-cell imaging. The regulation of the culture conditions in the incubation box is more sensitive to variations compared to a dedicated incubator [13,14]. Besides that, images are only captured at certain time points, but the microscope is unavailable for (end-point) imaging by other users during the entire live imaging experiment. A system that could overcome these issues is the CytoSMART® Lux3 BR Duo Kit. This dedicated system for live-cell imaging fits in a regular incubator, and therefore enables culture monitoring in a constant and optimal culture environment. The optical properties of the devices provide high-quality imaging, as well as accurate cell tracking. By connecting two devices within the same incubator to a single laptop, two cultures can be monitored simultaneously and compared side-by-side, without occupying a microscope for other lab members.

In this proof-of-concept study, we determine the effect of an FBS gradient on directed migration of cancer cell lines, using side-by-side high-quality live-cell imaging. The CytoSMART® Lux3 BR Duo Kit was used to monitor two cancer cell lines exposed to an FBS gradient or constant FBS concentration. Cells were tracked over time, and this provided fundamental insight into the chemotactic migration of cancer cells, which may ultimately be related to cancer metastasis.

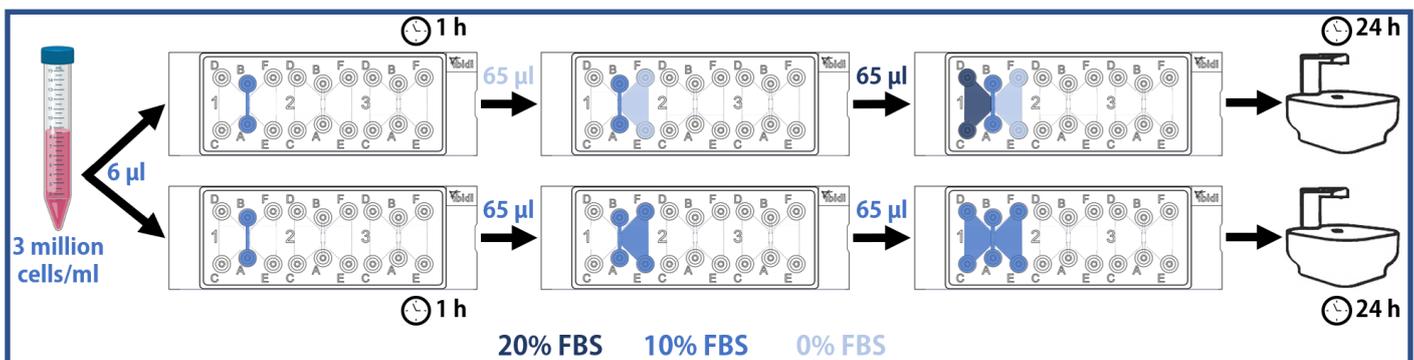


Fig. 1. Experimental setup. Cancer cell lines were seeded in Ibidi µ-Slide Chemotaxis and culture medium was added, either resulting in an FBS gradient or a constant FBS concentration. Cultures were monitored for 24 h using the CytoSMART® Lux3 BR Duo Kit, with an imaging interval of 5 min.

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Materials and methods

HeLa cells (Innoprot P20107) and C6 cells (ATCC CCL-107) were cultured to sub-confluency in DMEM (Gibco) supplemented with 10% FBS (Gibco) and 1% pen-strep (Gibco), under standard culture conditions (37°C; 5% CO₂). Each of the cancer cell lines was seeded in the Ibidi μ-Slide Chemotaxis at 18,000 cells per channel (6 μl cell suspension; 3 million cells/ml; medium with 10% FBS). 65 μl culture medium (DMEM, FBS, pen-strep) was added to the reservoirs: either 20% FBS in one reservoir and

0% FBS in the other or 10% FBS in both reservoirs (Fig. 1). High-quality images of the cultures were made using the CytoSMART® Lux3 BR Duo Kit (37°C; 5% CO₂): cultures were monitored for 24 h, taking a snapshot every 5 min. Afterwards, images were exported from the CytoSMART® Cloud, and single cells were tracked with FIJI-plugin TrackMate [15]. Tracked paths were converted to chemotaxis plots with FIJI-plugin Chemotaxis and Migration Tool [16].

Results

Images from the time-lapses of the cultures with FBS gradient or constant FBS concentration are displayed in Figure 2 (HeLa cells) and Figure 3 (C6 cells). The HeLa cells migrated a larger distance from their origin in a constant FBS concentration, but showed a slight preference to migrate towards a higher FBS

concentration when exposed to a gradient. The C6 cells seemed to migrate a smaller distance from the origin in the constant FBS concentration. However, these cells also displayed less directed migration – and therefore no clear chemotactic migration – in an FBS gradient.

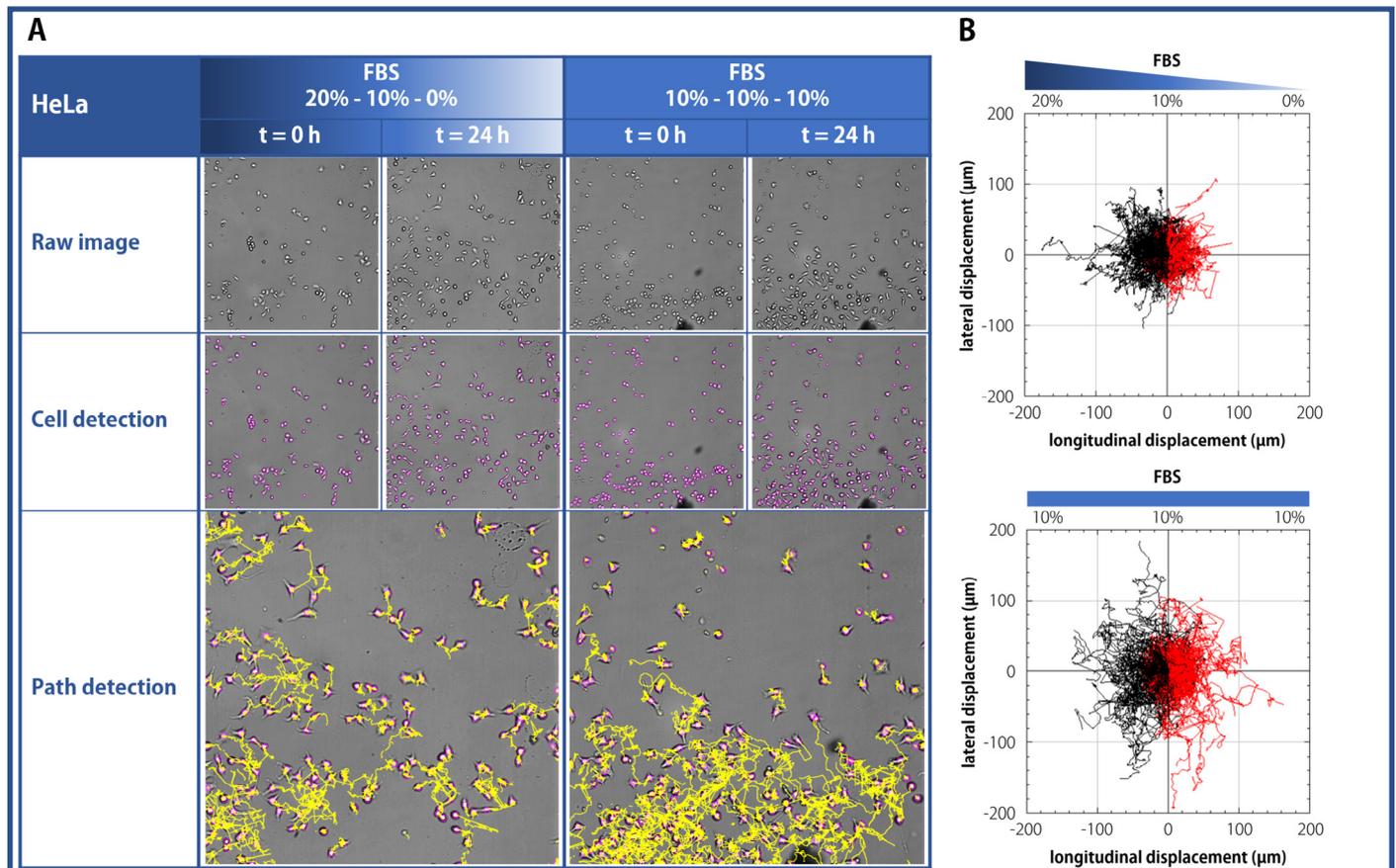


Fig. 2. HeLa cells cover a larger distance in constant FBS concentration, but prefer to migrate towards a higher FBS concentration when exposed to a gradient. A) Raw images of HeLa cells made using CytoSMART® Lux3 BR Duo Kit, single cell detection (purple) and path detection (yellow) with FIJI-plugin TrackMate. B) Chemotactic displacements of HeLa cells visualized using FIJI-plugin Chemotaxis and Migration Tool, with longitudinal and lateral displacements defined with respect to the FBS gradient.

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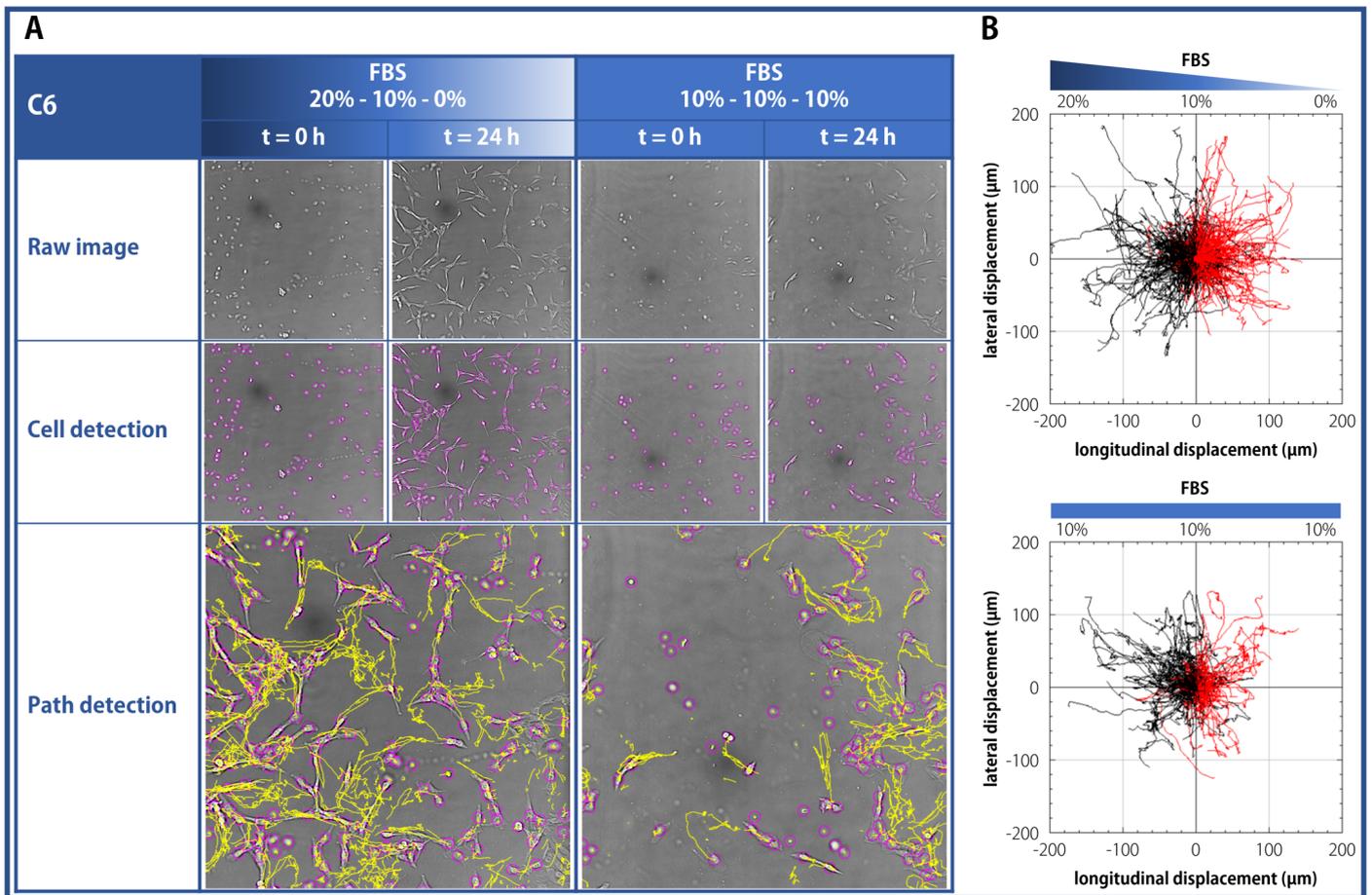


Fig. 3. C6 cells display no clear chemotactic migration in an FBS gradient. A) Raw images of C6 cells made using CytoSMART® Lux3 BR Duo Kit, single cell detection (purple) and path detection (yellow) with FIJI-plugin TrackMate. B) Chemotactic displacements of C6 cells visualized using FIJI-plugin Chemotaxis and Migration Tool, with longitudinal and lateral displacements defined with respect to the FBS gradient.

Discussion

Cancer metastasis is the main cause of cancer-related mortalities, and therefore cell migration is an important topic in cancer research. High-quality live-cell imaging is a prerequisite for accurate cell tracking, which provides fundamental insight into mechanisms driving metastatic cell migration. One of these mechanisms is chemotaxis, where cells migrate along an increasing gradient of a chemoattractant. This research aimed to determine the effect of an FBS gradient on directed migration of cancer cell lines, using side-by-side high-quality live-cell imaging with the CytoSMART® Lux3 BR Duo Kit. These devices provided high-quality images that were immediately applicable in the analysis software, enabling easy and accurate single cell tracking. From this tracking, it was observed that HeLa cells preferred to migrate towards the higher FBS concentration, whereas this was less clear for the C6 cells.

Cervical cancer is amongst the most commonly metastasizing primary tumors, with metastases found in e.g. bone, liver and lung tissue [17]. Glioma metastases, however, are a very rare phenomenon [18,19]. The clinical prevalence of each of these metastases could potentially be related to how much the blood serum – modeled by the FBS in this research – acts as a chemoattractant to the cells. Since spreading via the bloodstream is a prominent mechanism for metastasis [17], directed and possibly chemotactic migration of the tumor cells towards a blood vessel can initiate metastasis. The cervical cancer cell line HeLa displaying more chemotactic migration in this research compared to the glioma cell line C6 seems to be in line with the clinical prevalence of the respective metastases. The FBS that was applied as chemoattractant in this research provided an experimentally straightforward model system.

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Besides that, directed migration towards blood vessels was mimicked with this setup. However, FBS has a variable and complex composition [20], therefore providing little insight in the specific molecule(s) responsible for directed migration of the investigated cells. It should also be considered that FBS has other properties that affect cell behavior besides chemotactic migration: the FBS concentration in a cell culture for example influences cell proliferation [21]. Although no clear difference in proliferation rate in the high and low FBS concentration could be observed, the cell migration could still have been affected by other FBS-related processes. Therefore, follow-up experiments in the same setup using individual chemoattractants can

provide more detailed information on the cellular behavior. The high-quality images made by the CytoSMART® Lux3 BR Duo Kit were directly suitable for the TrackMate plugin. Therefore, automated cell tracking could be performed easily and quickly: results were obtained within minutes. Lower-quality images would have required manual adjustments to the cell tracking – which can lead to inaccurate results. This also would have slowed down the analysis, probably to hours or even days, since tracking on hundreds of images should have been checked and manually adjusted. Therefore, the image quality of the CytoSMART® Lux3 BR Duo Kit proved its added value to this research.

Conclusion

In this study, we successfully showed the possibilities of using high-quality live-cell imaging for accurate cell tracking. The CytoSMART® Lux3 BR Duo Kit enabled side-by-side monitoring of experimental conditions, and provided images directly applicable for cell tracking. This revealed the chemotactic

migration of HeLa cells in an FBS gradient, while C6 cells displayed no clear chemotactic migration. The high-quality imaging and corresponding results can ultimately be relevant for fundamental research regarding cancer metastasis.

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