Enrichment of circulating head and neck tumour cells using spiral microfluidic technology

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Whilst locoregional control of head and neck cancers (HNCs) has improved over the last four decades, long-term survival has remained largely unchanged. A possible reason for this is that the rate of distant metastasis has not changed. Such disseminated disease is reflected in measurable levels of cancer cells in the blood of HNC patients, referred to as circulating tumour cells (CTCs). Numerous marker-independent techniques have been developed for CTC isolation and detection. Recently, microfluidics-based platforms have come to the fore to avoid molecular bias. In this pilot, proof of concept study, we evaluated the use of the spiral microfluidic chip for CTC enrichment and subsequent detection in HNC patients. CTCs were detected in 13/24 (54%) HNC patients, representing both early to late stages of disease. Importantly, in 7/13 CTC-positive patients, CTC clusters were observed. This is the first study to use spiral microfluidics technology for CTC enrichment in HNC.

Head and neck cancers (HNCs) account for the 7th most common tumour type globally. Whilst there have been improvements in locoregional control of HNC, distant metastasis remains a challenge1. Circulating tumour cells (CTCs), are rare cancer cells shed into circulation, representing metastatic seeds and providing a window into metastasis. CTCs have the potential to provide critical information on the metastatic cascade, tumour heterogeneity and chemoresistance2–5. Whilst CTCs have been well studied in metastatic breast, prostate and colorectal cancer patients6,7, the HNC CTC field remains in its infancy1,7–9. Epidermal growth factor receptor (EGFR), which has been shown to be amplified in HPV-negative tumours compared to HPV-positive tumours, has previously been characterized in HNC CTCs7,9–14.

CTCs, when present, are extremely rare in comparison to the plethora of white blood cells in circulation15. The CellSearch system (Janssen Diagnostics) has FDA-approval for CTC enumeration in a number of tumour types and has been used to demonstrate the clinical relevance of EpCAM-positive CTCs6. This system has been robust in CTC enumeration through marker based CTC capture6,16. Numerous marker-based assays using microfluidic technologies are available and have been previously reported17,18. However, it has been shown that affinity based platforms have limitations due to poor sensitivity19,20. This is further compounded by varying levels of cell surface marker expression, CTC heterogeneity and specific processes such as epithelial-to-mesenchymal transition (EMT)1–3,21,22. The current view is that CTCs can show an intermediate EMT phenotype2,20,23 as well as express varying degrees of other markers such as PD-L1, CXCR4 and Plastin-32–24. Based on the heterogeneity found among CTCs, compounded with sometimes low-EpCAM expression, critical subpopulations may not be detected. Therefore, various marker-independent CTC enrichment strategies need to be tailored for each tumour

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Marker-independent enrichment relies on the physical properties of CTCs such as size, deformability, charge and density. Recently, there has been a demand in the field for the isolation of viable CTCs on which to perform functional assays. There has been a shift toward methodologies that allow for (i) marker-independent CTC capture (ii) CTC propagation (iii) minimal pre-enrichment processing and (iv) processing larger volumes of blood in a short time period. To encompass the above, the spiral microfluidics technology developed by Warkiani et al. was evaluated. The spiral chip is a marker-independent and high-throughput device that utilises hydrodynamic forces present in curvilinear microchannels for size-based cell sorting. The spiral chip has previously been reported for use in breast and lung cancers. CTC separation from lysed whole blood occurs based on CTC deformability and size. In this study, we enriched blood from 24 HNC patients using the spiral microfluidics platform and enumerated CTCs using immunocytochemistry. Our evaluation is the first study using the spiral technology for CTC enrichment from patients with HNCs, and shows positive results. Future, multi-centre trials are required and warranted before this technology can be implemented in a clinical setting.

Results

Spiral Chip enrichment. The spiral chip utilizes the inherent Dean vortex flows present in curvilinear microchannels under continuous flow, along with inertial lift forces, which focus the CTCs near the microchannel inner wall while driving the smaller hematologic cells toward the microchannel outer wall, allowing an efficient separation at the outlet. Typically, cells larger than 14 μm flow into the “CTC outlet” and cells smaller than this size flow into the “waste outlet.”

Recovery Studies. FaDu, CAL27, RPMI2650 and MDA-MB-468 cell line cells, which all have a mean diameter above 15 μm, were used for spike-in experiments. The percentage recovery of spiked-in cells ranged from 60–70% when experiments were performed with 100–150 cells spiked into 10 mL blood, trending to lower recover rates (40–60%) when less cells were spiked in. The overall recovery of the spiked cell lines were comparable to each other. We also assessed the decrease in background leukocytes and recovery of cancer cells after repeat passages through the spiral chip. When the samples were collected from the “CTC outlet” and run through the chip again, a 30% decrease in leukocytes was observed, however there was a 10–15% decrease in cancer cell recovery which was accounted for in the dead volumes of the setup. More than 1 repeat run did not significantly affect the overall recovery.
not further purify the sample. Single cells and cell clusters were able to be enriched by the spiral chip (Fig. 3ii).

Patient Characteristics. A total of 24 patients and 8 normal healthy controls were included in this study. Patient characteristics are shown in Table 1. The median age was 62 years (range 36–89); including men (n = 22), women (n = 2). The most distinctive histology was mucosal oral cavity SCC (63%). The majority of patients presented with tumour grading T2/T3 (71%) and either no nodal spread N0 (50%) or advanced nodal spread N2a-c (50%). Patients were assessed for distant metastases by CT/PET-CT and MRI scans. All patients were found to be radiographically M0 upon assessment. CTC status was determined at the same time point. The clinicopathological patient findings are presented in Table 2.

Enrichment of CTCs from controls and patients. 10 mL blood samples were obtained from 8 healthy volunteers and 24 head and neck cancer patients and processed using the spiral chip. All samples were enumerated for CTCs. Putative CTCs were identified on the stained cytospots as positive for: pan-CK, DAPI and negative for CD45. Further characterisation of head and neck CTCs was performed by EGFR ICC and DNA FISH. Leukocytes were identified as positive for CD45, DAPI and negative for pan-CK. No CTC-like events were observed in the 8 normal healthy volunteers (Fig. 5). CTCs were detected in 13/24 (54%) (Range 1–12 CTCs/10 ml blood sample) of the HNC patients screened in the study. CTC clusters were readily detectable in 7/13 (53.8%) of the CTC-positive patients (Fig. 6). The CTCs were further molecularly characterised and found to have EGFR gene amplifications (Fig. 7). Further observations made in the patient samples include double positive pan-CK and CD45 cells in 2/20 samples and 27 CK-negative, CD45-negative cells in 1 patient presenting with a CTC cluster. These events were not considered for enumeration in the study.

Discussion

HNC patients often present with advanced metastatic spread, predominantly to the lungs, liver and bone. The ability to assess “intact” metastatic precursors in the form of tumour cells in blood holds remarkable potential as a liquid biopsy. The number of CTCs, and molecular and genetic changes in the CTCs could be monitored.
Moreover, patients whose disease will potentially respond to specific therapies could be identified. Recent studies have shown that PD-L1 is frequently expressed on CTCs and that these are present in patients where anti-PDL1 therapies could be effective. Numerous CTC enrichment platforms have been developed in recent years, each with its own advantages and limitations. In this study, we tested and evaluated the spiral microfluidics chip for CTC enumeration and characterization. This platform has features which are of value to clinical research; CTCs are found in suspension and not immobilized allowing for functional assays and CTC clusters can be readily isolated with a leukocyte depletion of up to 4 log. Limitations of the spiral technology are that “small CTCs” or CTCs (<12 μm) in size will be lost.

Four cell lines representing a mean cell diameter range between 16–28 μm were chosen for the spiking experiments to get a decent coverage of known head and neck cancer cells. In our hands, the percentage recovery of the spiral chip is similar to that of other microfluidic CTC isolation platforms. By repeat passages of the spiral chip, a decrease of 30% background leukocytes was found at the expense of 10–15% recovery. However, repeated passages were not performed with patient samples so as not to risk losing CTCs. The range of detected CTCs in HNC patients remains low. Moreover, it was observed that the same population of leukocytes was present after 2 or more repeat runs and this could not be further purified. Furthermore, the spiral chip successfully sorted for cell clusters and tumour spheroids in the CTC outlet.

Figure 4. Day 24, Bioreactor grown (A) FaDu spheroid (B) MDA-MD-468 (cell cluster), (C) MDA-MD-468 Spiral sorted single and cell clusters (D) 600x image of Spiral sorted cell cluster. Scale bar represents 10 μm.

| Variables                        | N  |
|----------------------------------|----|
| Total                            | 24 (100%) |
| Gender                           |    |
| Male                             | 22 (91.7%) |
| Female                           | 2 (8.3%)  |
| Age, y                           |    |
| <60                              | 7  |
| >60                              | 17 |
| Anatomic site of primary         |    |
| Oral Cavity                      | 15 |
| Oropharyngeal                    | 9  |
| Tumour Size                      |    |
| T1                               | 3  |
| T2                               | 10 |
| T3                               | 7  |
| T4                               | 4  |
| Nodal spread                     |    |
| N0                               | 12 |
| N1                               | 0  |
| N2a-c                            | 12 |
| Distant metastases               |    |
| M0                               | 24 |
| M1                               | 0  |
| HPV status                       |    |
| HPV-positive                     | 12 |
| HPV-negative                     | 12 |
| CTC status                       |    |
| CTC-positive                     | 13 (Range from 1–12 CTCs/10 ml whole blood) |
| CTC-negative                     | 11 |

Table 1. Patient demographics (n = 24).

| Variables                          | N  |
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| Distant metastases                 |    |
| M0                                 | 24 |
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| HPV status                         |    |
| HPV-positive                       | 12 |
| HPV-negative                       | 12 |
| CTC status                         |    |
| CTC-positive                       | 13 (Range from 1–12 CTCs/10 ml whole blood) |
| CTC-negative                       | 11 |
This pilot study evaluated the use of a spiral chip for CTC enrichment in HNCs. HNC patients ranged from early stage (T1N0M0) to advanced stage patients (T4N2b). Patients with early stage disease present with low rates of distant metastasis and a favourable prognosis. Whilst patients with locally advanced disease are more likely to have CTCs\(^4\). CTCs were observed in 50% of the HNC samples, ranging from 1–12 CTCs/10 mL sample. This finding is consistent with previously published studies using marker-independent enrichment platforms\(^7,13,45\). Notably, post spiral enrichment involves transferring rare cells onto a glass slide and immunocytochemistry which can lead to cell loses\(^3\). Therefore, the actual CTC count per HNC patient could be much higher.

| Patient | Gender | Age | HPV status | T Staging | N Staging | Description | Lung findings | CTC findings | CTC description |
|---------|--------|-----|------------|-----------|-----------|-------------|--------------|--------------|-----------------|
| Micro_01 | m | 66 | positive | T2 | N0 | Oral Cavity lesion, query | Positive | 1 CTC cluster ~12 cells |
| Micro_02 | m | 54 | unknown | T1 | N0 | Oropharyngeal | Negative |
| Micro_03 | m | 63 | negative | T4 | N2c | Oropharyngeal lesion, query | Positive | 1 CTC |
| Micro_04 | m | 58 | positive | T3 | N2b | Oropharyngeal | Negative |
| Micro_05 | m | 62 | positive | T2 | N2c | Oropharyngeal | Negative |
| Micro_06 | m | 42 | negative | T3 | N2 | Oral Cavity | Positive | 1 CTC cluster ~5 cells, 27 nucleated single cells |
| Micro_07 | f | 72 | unknown | T1 | N0 | Oral Cavity | Negative |
| Micro_08 | m | 48 | unknown | T1 | N0 | Oral Cavity | Negative |
| Micro_09 | m | 63 | positive | T3 | N2c | Oropharyngeal | Positive | 1 CTC cluster ~6 cells |
| Micro_10 | m | 61 | positive | T2 | N0 | Oral Cavity | Negative |
| Micro_11 | m | 73 | unknown | T2 | N0 | Oral Cavity | Positive | 1 CTC cluster ~5 cells |
| Micro_12 | m | 36 | negative | T4 | N2b | Oropharyngeal lesion, query | Positive | 3 CTCs |
| Micro_13 | m | 56 | positive | T2 | N2b | Oropharyngeal | Positive | 2 CTCs |
| Micro_14 | m | 55 | positive | T4 | N2b | Oropharyngeal | Positive | 1 CTC |
| Micro_15 | m | 71 | negative | T2 | N0 | Oral Cavity | Negative |
| Micro_16 | m | 65 | negative | T3 | N2b | Oral Cavity | Positive | 1 cluster ~5 cells + 2 single cells |
| Micro_17 | m | 60 | positive | T3 | N2b | Oropharyngeal | Positive | 1 cluster ~5 cells + 4 single cells |
| Micro_18 | m | 67 | positive | T3 | N0 | Oral Cavity | Negative |
| Micro_19 | m | 60 | positive | T2 | N2b | Oropharyngeal | Negative |
| Micro_20 | m | 68 | positive | T4 | N0 | Oral Cavity | Negative | 6 CTCs |
| Micro_21 | m | 62 | negative | T2 | N0 | Oropharyngeal | Negative |
| Micro_22 | m | 68 | negative | T3 | N2b | Oral cavity | Positive | 2 CTCs |
| Micro_23 | m | 89 | negative | T2 | N0 | Oral Cavity | Negative |
| Micro_24 | m | 60 | negative | T2 | No | Oral Cavity | Positive | 3 CTCs |

Table 2. Clinicopathological findings including CTC status and Lung findings.

Figure 5. Box plot summary indicating the range of CTCs (pan-CK + CD45-DAPI + ) cells/10 mL. The box plot presents the median and all individual data points. NHV; normal healthy volunteers.
Over half of the CTC positive patients presented with clusters of cancer cells. These multicellular CTC clusters have shown 23–50 fold increased metastatic potential compared to single CTCs in breast cancers and associations with adverse clinical outcomes1,46–49, however the clinical significance in HNC is yet to be determined1,50–52. Moreover, these cell clusters have been reported to have a shorter circulation half-life in blood, consistent with rapid entrapment within the capillaries of distant organs46. Initial studies have shown that CTC clusters are potent initiators of distant organ metastasis, highlighting the need to develop strategies which disrupt cell-cell or cell-matrix adhesions in these clusters to reduce metastatic potential53. Further investigation is required to determine the clinical implications of CTC clusters in HNC patients. Importantly, CTCs were found upon presentation to clinic in 3/3 patients presenting with queried lung lesions by the MDT clinic (Fig. 8). These lung lesions were to be investigated by ultrasound guided fine-needle biopsy to determine the extent of distant metastasis. In these patients, a window for treatment escalation could become a possibility if the presence of CTCs predicts for patients at-risk of developing metastasis. Further studies assessing CTCs during the course of treatment and timing of blood sampling are warranted to investigate the role of monitoring CTCs in HNC50.

Figure 6. Patient “micro_01, micro_06”; CTC clusters detected in the blood. (a) 100x magnification showing the CTC cluster in the top of the cytospot. 1000x images of (b) DAPI (c) Cytokeratin/DAPI composite (d) DAPI (e) Cytokeratin (f) Cytokeratin/DAPI composite. Scale bar represents 50μm. Figure 6 continued. Patient “micro_12”; CTC doublet detected in blood. (g) DAPI (h) Cytokeratin/DAPI composite (i) EGFR/DAPI composite. Patient “micro_03” single CTCs detected in blood. (j) DAPI (k) Cytokeratin/DAPI composite (l) EGFR/DAPI composite. Scale bar 10μm (g–i). Scale bar 50μm (j–l).
Figure 7. EGFR gene amplification in CTCs derived from head and neck cancer patients. Dual colour FISH assay probes for EGFR (red) and chromosome seven (CEP7, green). Row 1 shows individual stains for DAPI, CEP7, EGFR and the composite image (DAPI, CEP7, EGFR). Row 2 shows further patient CTC composite images showing EGFR amplification. Scale bar represents 10 μm.

Figure 8. A FDG-PET study from a patient presenting with CTCs and had progressive disease to the lungs showing an 8 × 5 mm nodule in the left upper lobe.
Conclusion
We present the first study in HNCs using a microfluidics based approach for CTC enrichment. 24 HNC patient blood samples were enriched using the spiral microfluidics chip. We were able to identify single and clusters of CTCs and further characterise these cells by ICC and DNA FISH. This study highlights the use of the spiral microfluidic technology in HNC and provides future clinical use in the field of CTCs.

Materials and Methods

Patient cohort.
Ethics approval was obtained from the Metro South and North Health Service District Human Research Ethics Committee in accordance with the National Health and Medical Research Council's guidelines (HREC/12/PHAAH381) to collect blood from the Royal Brisbane and Women's Hospital (RBWH). All methods were performed in accordance with these ethical guidelines and regulations. This study also has Queensland University of Technology ethics approval (1400000617). After written informed consent was obtained from the participants, 10 mL blood samples were collected from patients presenting to the HNC clinic.

Cell lines and culture.
The cell lines FaDu (ATCC HTB43), CAL27 (ATCC CRL2095), RPMI2650 (ATCC CCL30) were obtained from ATCC (Manassas, VA). UD-SCC9 HNC cells (ATCC CRL1629) were a generous gift from A/Prof Nick Saunders (UQDI). The MDA-MB-468 breast cancer cells were originally from the ATCC transferred from the Lombardi Cancer Center and fluorescently tagged with pCherryN1, a modification of the Clonetech vector, kindly provided by Dr Alpha Yap, (UQ). Cells were cultured under standard conditions in humidified incubators at 37 °C, 5% CO2 in RPMI-1640-Glutamax (Life Technologies, Inc) supplemented with 10% fetal bovine serum (FBS) and 1% Penicillin/Streptomycin (Pen/Strep).

3D cell culture (bioreactor).
Cells were spun down to obtain cell pellets post-trypsinization. The cell pellets were resuspended in pre-warmed 1x Happy Cell media (Biocroi Ltd, Dublin, Ireland) supplemented with 10% FBS and 1% Pen/Strep. Optimal seeding densities for the bioreactors were determined by titrating cells from 10,000–500 cells/ml. 1,500 cells/ml, resuspended in 1x Happy Cell media were layered over 2x Happy Cell media FBS and 1% Pen/Strep. Optimal seeding densities for the bioreactors were determined by titrating cells from 10,000–500 cells/ml. 1,500 cells/ml, resuspended in 1x Happy Cell media were layered over 2x Happy Cell media FBS and 1% Pen/Strep. Optimal seeding densities for the bioreactors were determined by titrating cells from 10,000–500 cells/ml. 1,500 cells/ml, resuspended in 1x Happy Cell media were layered over 2x Happy Cell media FBS and 1% Pen/Strep.

Spiking of cell lines into bloods collected from normal healthy controls.
Blood was collected from normal healthy volunteers in 10 mL EDTA (BD-Plymouth, UK) tubes. The cell lines used for spiking were labelled with CellTracker™ Green (Life Technologies) as per manufacturer's instructions. The collected blood was spiked with clinically relevant numbers (10–150) of cells within 2 hours of collection. Collection of spheroids was transferred to a 96 well format and an inactivation buffer was used to collapse the hydrogel matrix for 1 hour at 37 °C.

Preparation of blood sample for the spiral Chip.
To reduce the cellular components passing through the spiral chip, an initial red blood cell (RBC) lysis was performed, which has been shown to have minimal effects on the recovered cells15. Briefly, RBC lysis buffer (Astral Scientifix) was added to the blood sample, after which cells were centrifuged and the pellet resuspended in 10 mL of sheath buffer (1xPBS, 2 mM EDTA, 0.5% BSA).

Setup and running of the spiral Chip.
Tygon® tubing was inserted into the inlet/outlets of the spiral chip, and the inlet tubing connected to a syringe pump. The spiral chip was positioned and fixed onto a phase contrast microscope (Olympus IX71). The outlet tubing was connected to two sterile 15 mL collection BD Falcon tubes. An initial priming run was performed using the sheath buffer at a flow rate of 2.0 mL/min for 5 minutes. To test patient samples, the sample was loaded carefully into a 10 mL syringe (Terumo) and pumped through the spiral chip using the syringe pump at a flow rate of 1.7 mL/min for all experiments. The outputs were collected and spun down at 300 x g for 5 minutes.

Immunostaining.
Prelabelling and immunocytochemistry (ICC) were used for the detection of spiked tumour cells and CTCs enriched by the spiral microfluidic chip. Enriched cells were cytospun onto a glass slide and CTCs identified by ICC using the CellSearch antibody cocktail (Cytokeratin/CD45/DAPI; Janssen Diagnostics). Cells were further characterised for cell surface EGFR using an anti-EGFR antibody (AY13, BioLegend, San Diego). Briefly, the cytospun sample was air dried overnight and incubated with a mixture of CellSearch reagents (20 μl staining reagent, 20 μl DAPI, 20 μl permeabilization buffer, 20 μl fixation buffer in 120 μl PBS). The slide was incubated for 1 hour at room temperature, washed 3 times in PBS, overslipped and imaged under a fluorescence microscope. The CellSearch criteria for CTCs were used for CTC identification: intact, cytokeratin-positive, DAPI-positive, CD45-negative cells that were larger than 4 μm were considered a CTC. Cells staining positive for CD45 and DAPI and negative for pan-cytokeratin were determined to be white blood cells.

DNA Fluorescence in situ hybridization.
Cytospots were placed in pretreatment solution for 10 mins at 98 °C, and digested with pepsin for 5 mins at 37 °C in the Dako hybridiser. After dehybridisation through a graded ethanol series (70%, 85%, 96%), dual colour, dual target DNA FISH assays were done with EGFR/CEN7FISH probe mix (DakoCytomation, Glostrup, Denmark). Cytospots were covered with 10 μl probe solution, sealed with rubber sealant and placed in the Dako Hybridizer for denaturation (5 mins, 85 °C) and hybridisation (18 hours, 45 °C). Post hybridisation stringency washes were performed at 65 °C for 10 mins. After further...
washing and dehydation steps, the cytoplots were air dried and mounting media containing DAPI added. The cytopsots was coupled and imaged on the Nikon Eclipse Ti inverted microscope fitted with a Nikon digital camera. **EGFR** was visualized as a red signal (tetramethylrhodamine isothiocyanate filter), CEP7 (fluorescein isothiocyanate filter) as a green signal and the nuclei as a blue signal with a DAPI filter. **EGFR** status was scored as the number of **EGFR** signals per nucleus and as the ratio of **EGFR** status was scored as

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Additional Information

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