## Insights into the role of sulfated glycans in cancer cell adhesion and migration through use of branched peptide probe

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## Supplementary material

**Figure S1.** Movie of PANC-1 cell migration in wound healing assay without NT4 peptide (control). CytoSMART Lux 10x System was used to take pictures every 10 min for a total of 22 h.

Figure S2. Movie of PANC-1 cell migration in wound healing assay with 10  $\mu$ M NT4 peptide. CytoSMART Lux 10x System was used to take pictures every 10 min for a total of 22 h.

**Figure S3.** Movie of PANC-1 cell migration in a Matrigel wound healing assay without NT4 peptide (control). CytoSMART Lux 10x System was used to take pictures every 7.5 min for a total of 24 h.

**Figure S4.** Movie of PANC-1 cell migration in a Matrigel wound healing assay with 10  $\mu$ M NT4 peptide. CytoSMART Lux 10x System was used to take pictures every 7.5 min for a total of 24 h.

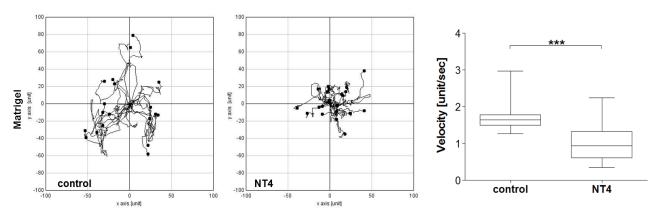


Figure S5. Directionality of cancer cell migration analyzed by time lapse microscopy using an in vitro wound healing assay in the presence of Matrigel. PANC-1 cancer cells were seeded on each side of a culture insert, covered with Matrigel and then incubated with and without 10  $\mu$ M NT4 (central panels and left panels, respectively). Cells were tracked every 15 min for 10 hours post-wounding and their paths plotted on a polar grid. Each plot represents 22 individual cell tracks. Velocity (unit/sec; where unit correspond to nm) of each analysed cell is reported in the box plot graph (right panels) where the median value is indicated by the line inside each box. \*\*\* p<0.001 calculated using one-tailed Student t-test.