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topics in biomedical engineering, namely clinical. Introduction to Biochemical Engineering dg rao. in ice cold (-4°C) PBS. The staining specificity in all three channels was verified by omitting the primary antibody (Supplementary Fig. [4](#MOESM1){ref-type="media"}). Proteome profiler array {#Sec15} ----- \*GFP\*-\*vimentin\*, \*GFP\*-\*α\*-\*actinin\* and \*GFP\*-\*slam\* constructs were generated in pCS2+ and transfected into \*X. laevis\* embryos. Two and a half days after injection, embryos were removed from the chorion and dechorionated in 0.8% (w/v) cysteine HCl (Sigma-Aldrich, Gillingham, UK) in nuclease-free (NF) water for one hour at room temperature. After two washes in NF water, embryos were homogenised in lysis buffer (30 mM Tris-HCl, 5 mM MgCl<sub>2</sub>, 0.2% (v/v) Triton X-100, 0.5 mM PMSF) supplemented with protease inhibitor cocktails (Sigma) on ice and then centrifuged for 10 minutes at 15,000 × \*g\* at 4 °C. The supernatant was

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