## Supplementary Figure 1. Cell slippage.



**Supplementary Figure 1. Cell slippage.** (a) and (b) represent the schematic evolution of two neighbouring cells whose shapes are modelled by ellipses. The points C1 and C2 are the cell centres. (a) corresponds to the initial state, and (b) to the configuration of the same two cells slightly later. Cell centres evolve according to the tissue velocity gradients,  $L_t$ , whereas each individual cell's shape evolves according to the cell shape strain rate,  $L_c$ . The geometrical construction of the effective slippage velocity is shown (see text for details). (c) illustrates the relationship between the intercalation tensor,  $L_i$  (drawn as in **Fig. 3h**), and the mean slippage direction and velocity relative to neighbouring cells (arrows).







Supplementary Figure 3. Pipeline of the algorithms used to calculate strain rates from raw volume images.



IMAGE ANALYSIS & CELL TRACKING

## **STRAIN RATE ANALYSIS**

Supplementary Figure 3. Pipeline of the algorithms used to calculate strain rates from raw volume images.

Supplementary Figure 4. Strain rates at the zebrafish midline.



Supplementary Figure 4. Strain rates at the zebrafish midline. (a)-(d) show average strain rates for pre-neurulation (~10-11 hpf) zebrafish trunk ectoderm. Strain rates projected onto the AP axis are shown for the length of the movies presented in Figure 5g,h. Colors for average strain rates (d) are as for cumulative stretch ratios in Figure 5.