

Fig S1. dsRNAi-mediated knockdown of small GTPases and Sra1. (A) qRT-PCR quantification of mRNA levels in control- or Arf79F-dsRNAi cells after either 3 or 6 days dsRNA incubation. qRT-PCR quantification of mRNA levels in (B) control-, Sra1-, Rac1- or Arf79F-dsRNAi cells or (C) control-, Arf79F-, Arf102F- or Arf51F-dsRNAi cells after 6 days dsRNA incubation.

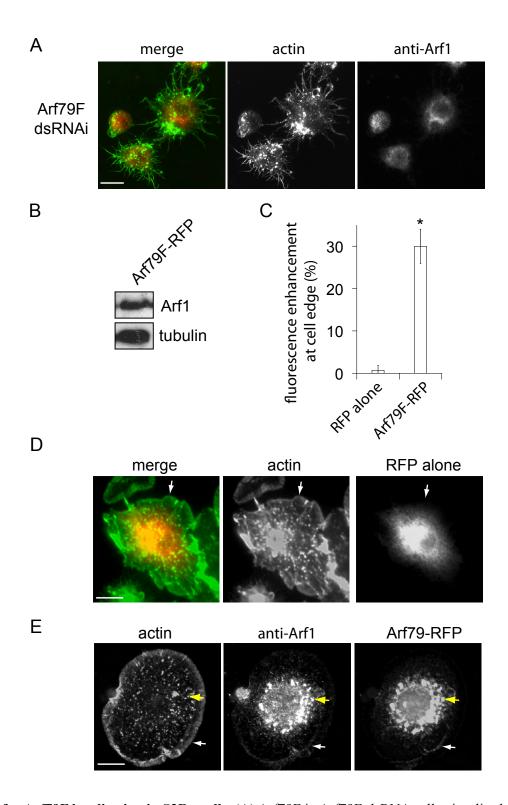
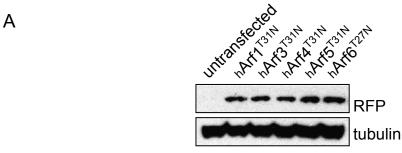


Fig S2. Controls for Arf79F localisation in S2R+ cells. (A) Arf79F in Arf79F-dsRNA cells visualised with Arf1 antibodies and actin as control (scale bars 10μm). (B) Immunoblot of Arf79F-RFP with antibodies to Arf1 and tubulin as control. (C) Percentage increase in RFP or Arf79F-RFP fluorescence at cell edge relative to cytoplasm. Statistics as Fig 1D. Lamellipodia (white arrows) visualised in cells expressing (D) RFP or (E) Arf79F-RFP by staining actin (scale bars 10μm). Arf79F-RFP stained with Arf1 antibodies confirming Arf79F localisation at the Golgi (yellow) and plasma membrane (white).



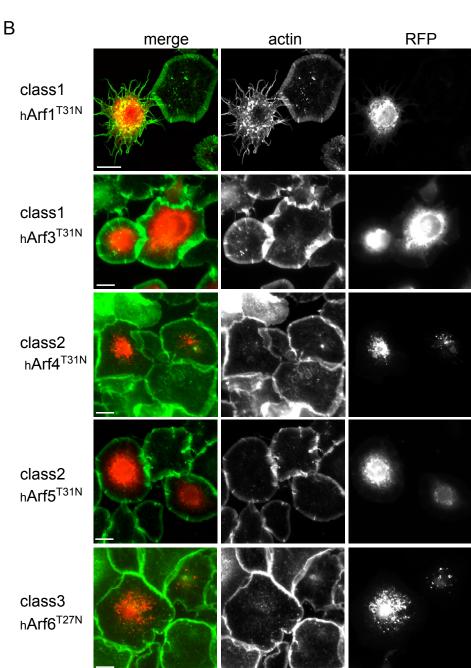
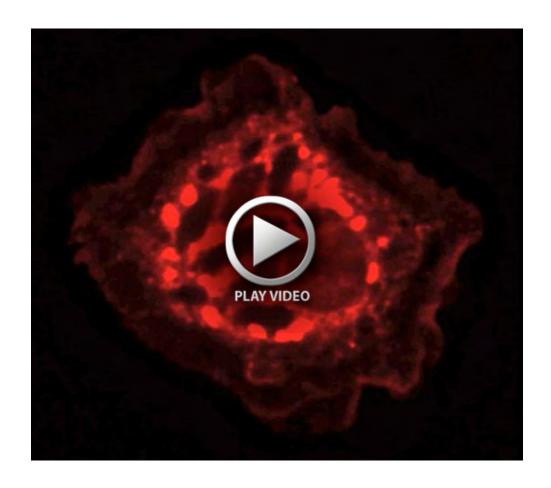


Fig S3. Influence of dominant negative Arf GTPases on lamellipodia formation. (A) Relative expression of indicated RFP-tagged human Arf dominant negatives determined by immunoblotting whole cell lysates with antibodies to RFP and tubulin as control. (B) Lamellipodia in cells expressing human dominant-negative Arf GTPases (hArf^{TN}) visualised by staining actin (scale bars $10\mu m$).



Movie 1. Arf79F localises to dynamic lamellipodia and macropinosomes. Live microscopy was used to image S2R+cells expressing Arf79F-RFP at a rate of 8 frames/second.