## **Supplementary Figures:**

- Figure S1. HDLEC monolayers incubated with the indicated CM for 24 h were analysed
- 4 by immunofluorescence (IF) staining of VE-cadherin (red). Blue indicates the nucleus.

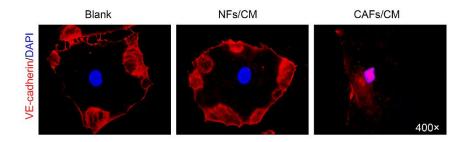


Figure S2. siLRP1 abolished the biological effects of PAI-1 on HDLECs, related to Fig 5C.

**A.** Representative micrographs of the Transwell assay with HDLECs. **B.** Representative micrographs of the tube formation assay. Scale bar, 50 μm. **C.** HDLEC monolayer following the indicated treatments described in **Fig. 5C** and quantification of SiHa-mCherry cell migration.

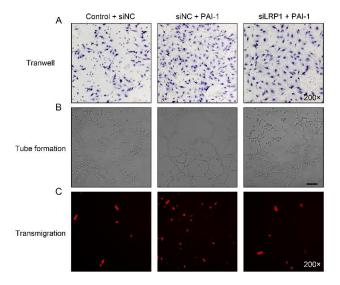


Fig S3. Suppression of ERK1/2 phosphorylation by U0126 in HDLECs affected the biological behaviour of PAI-1, related to Fig 5F.

**A.** Representative micrographs of the Transwell assay in HDLECs. **B.** Representative micrographs of the tube formation assay. Scale bar, 50 μm. **C.** HDLEC monolayer following the indicated treatments described in **Fig. 5G** and quantification of SiHa-mCherry cell migration.

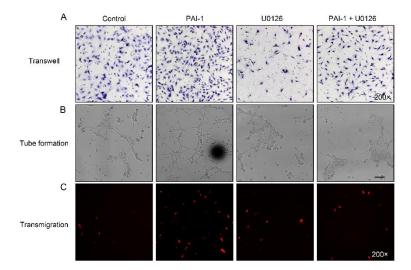


Fig S4. Suppression of AKT phosphorylation by MK2206 in HDLECs affected the biological behaviour of PAI-1, related to Fig 5G.

**A.** Representative micrographs of the Transwell assay with HDLECs. **B.** Representative micrographs of the tube formation assay. Scale bar, 50 μm. **C.** HDLEC monolayer following the indicated treatments described in **Fig. 5F** and quantification of SiHa-mCherry cell migration.

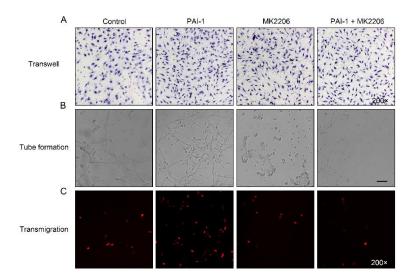


Fig S5. Validation and efficacy of LRP1 gene silencing was performed by qPCR.

Normalized gene expression over GAPDH is shown with the means ± SD of three

## 29 independent experiments. \*P < 0.05.

30

