## Supplementary data

## HIF stabilization inhibits renal epithelial cell migration and is associated with cytoskeletal alterations

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#### Supplementary Fig. S1: Faster Migration of E-cadherin-positive distal tubules

hPTEC were seeded in Ibidi barriers in 8-well slides and grown to confluence. After removal of the barriers, cells were allowed to migrate into the open space for 7 h. Cells were stained for E-cadherin (red) and N-cadherin (green). Arrrows point to faster migrating cells. Arrowheads marked E- and N-cadherin positive cells. Scale bar: 40 µm

#### Supplementary Fig. S2: DMOG-induced morphological alterations.

Sub-confluent hPTEC were incubated with DMOG (1 mM) for 24 h. Paxillin and F-actin were detected by immunofluorescence. Scale bar: 10  $\mu$ m

#### Supplementary Fig. S3: Variable expression of K18 in renal epithelial cells

A: Monolayers of hPTEC were stained with antibodies against N-cadherin and K18. Scale bar: 30  $\mu m$ 

B: K18 was detected in a monolayer of distal tubular cells. Scale bar: 30 µm

C: K18 was detected in human kidney sections by immunohistochemistry. Scale bar: 30µm

Supplementary Fig. S4: DMOG does not alter expression of K7 or K19 in hPTEC

hPTEC were incubated with DMOG (1 mM) for 24 h and 72 h. Expression of K7 and K19 was detected by Western blotting using specific antibodies. Blots obtained from cells incubated for 24 h were reprobed with Vinculin (Vinc). The graphs summarize data obtained by densitometric quantification of n=3 experiments with different isolations (means  $\pm$  SD). Data obtained with control cells were set to 100.

### Supplementary Fig. S5: Original blots

All blots used for the manuscript figures are shown without any modifications.

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## Α



С

В

Keratin 18



Keratin 18





0

Co DMOG

24 h

Co DMOG

72 h



Blots from Figure 5 B



Blots from Suppl. Figure 4A

### Blots from Suppl. Figure 4 B

