Supplementary Figure 1. Expression of FAHD1 in breast cancer tissue and verification of regular FAHD1 levels in control cells. 

A) The abundance of FAHD1 in breast cancer tissue from patient samples were analyzed by immuno-histochemical staining and confirmed high levels of FAHD1 in malignant tissue. 

B) Mitochondrial PEPCK expression in breast cancer cell lines analyzed by Western Blot. 

C) Western blot analysis of infected MCF-7 cells shows similar FAHD1 levels in uninfected (WT) and control (SCR) cells while FAHD1 KD cells do not express the enzyme at physiological levels.
Supplementary Figure 2

**A)** Western blot depicting upregulation of mPEPCK in MCF-7 cells under limiting conditions such as glutamine only or glucose only media. **B)** MCF-7 cultured in regular media containing glucose and glutamine (full media) with no apparent morphological alterations. **C)** Protein levels of succinate dehydrogenase (SDH, mitochondrial complex II) were analyzed in control (SCR) and FAHD1 KD MCF-7 cells via western blot. **D)** Expression levels of NADH:ubiquinone-oxidoreductase (mitochondrial complex I, NDUF9) were analyzed in SCR and FAHD1 KD MCF-7 cells.

**Supplementary Figure 2.** MCF-7 cells cultivated in limiting conditions and in full media

A) Western blot depicting upregulation of mPEPCK in MCF-7 cells under limiting conditions such as glutamine only or glucose only media. B) MCF-7 cultured in regular media containing glucose and glutamine (full media) with no apparent morphological alterations. C) Protein levels of succinate dehydrogenase (SDH, mitochondrial complex II) were analyzed in control (SCR) and FAHD1 KD MCF-7 cells via western blot. D) Expression levels of NADH:ubiquinone-oxidoreductase (mitochondrial complex I, NDUF9) were analyzed in SCR and FAHD1 KD MCF-7 cells.
Supplementary Figure 3. High resolution respirometry for complex II activity

A) Schematic overview of the way of action for the compounds and substrates that were added during the measurement in the Oxygraph-2k. Details can be found at bioblast.at (SUIT-010) showing the respiratory states which were analyzed during high resolution respirometry.

B) Representative measurement of MCF-7 breast cancer cells in full media showing absolute values for oxygen flux in the Oxygraph-2k.

C) Absolute extracellular acidification rates (ECAR) in MCF-7 during the glycolytic stress test in the Seahorse HS Mini; n = 6 with SEM.

D) Glycolytic rate assay with MDA-MB-231 cells in the Seahorse HS Mini; n = 3 with SEM. Significance threshold for students t-test: * p ≤ 0.05; ** p ≤ 0.01; *** p ≤ 0.001
Supplementary Figure 4 – Part 1. Cell death and expression of GLS in FAHD1 KD BT-20 cells

A) Gating strategy for flow-cytometric evaluation of apoptosis in basal BT-20 cells. Representative figure for day 7 post-infection. Percentages of Annexin V+ cells (early apoptotic) from the living cell population (SSC-A/FSC-A) were obtained from n = 3 biological replicates of each control (SCR) and FAHD1 KD cells.

B) Evaluation of GLS isoform GAC expression and C) KGA expression in BT-20 cells upon FAHD1 KD. Mean of n = 8 biological replicates with SEM, also see Figure 5E.

D) Levels of mitochondrial phosphoenolpyruvate-carboxykinase (mPEPCK) were measured via western blot in SCR and FAHD1 KD BT-20 cells which were culture for 48 h in limiting conditions.

E) Protein expression of succinate-dehydrogenase (Complex II, SDH) and NADH:ubiquinone-oxidoreductase (mitochondrial complex I, NDUFA9) were analyzed in SCR and FAHD1 KD BT-20 cells. Significance threshold for students t-test: * p ≤ 0.05; ** p ≤ 0.01; *** p ≤ 0.001
Supplementary Figure 4 – Part 2. Cell death and expression of GLS in FAHD1 KD BT-20 cells

A) Gating strategy for flow-cytometric evaluation of apoptosis in basal BT-20 cells. Representative figure for day 7 post infection. Percentages of Annexin V+ cells (early apoptotic) from the living cell population (SSC-A/FSC-A) where obtained from n = 3 biological replicates of each control (SCR) and FAHD1 KD cells.

B) Evaluation of GLS isoform GAC expression and C) KGA expression in BT-20 cells upon FAHD1 KD. Mean of n = 8 biological replicates with SEM, also see Figure 5E.

D) Levels of mitochondrial phosphoenolpyruvate-carboxykinase (mPEPCK) were measured via western blot in SCR and FAHD1 KD BT-20 cells which were culture for 48 h in limiting conditions. E) Protein expression of succinate-dehydrogenase (Complex II, SDH) and NADH:ubiquinone-oxidoreductase (mitochondrial complex I, NDUFA9) were analyzed in SCR and FAHD1 KD BT-20 cells. Significance threshold for students t-test: * p ≤ 0.05; ** p ≤ 0.01; *** p ≤ 0.001