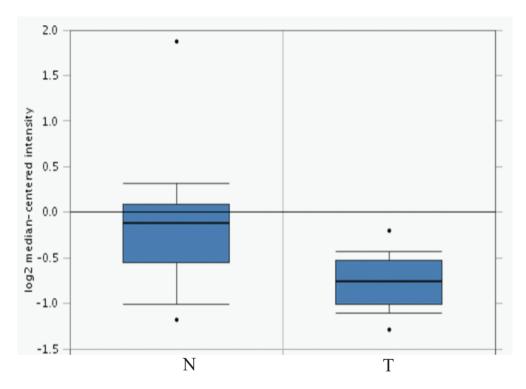
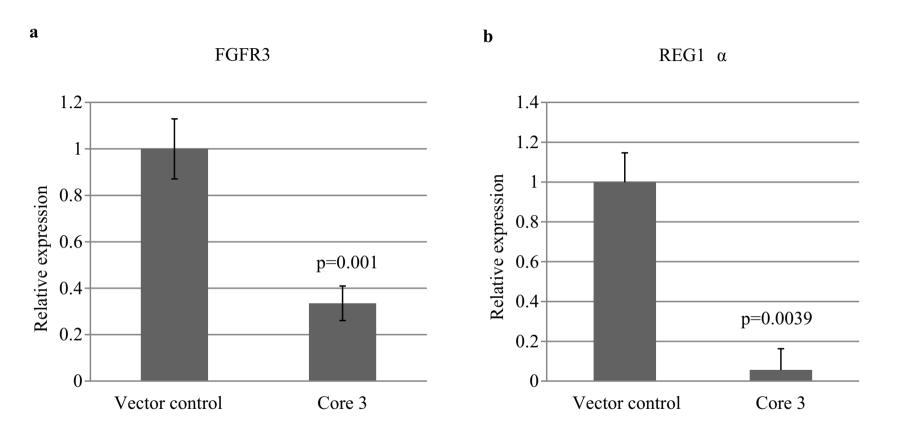
Supplementary Figure S1



Expression of core 3 synthase in paired normal pancreas (N; n=39) and Pancreatic Ductal Adenocarcinoma (T; n=39) was extracted from deposited gene expression data sets (n=36 patients, 3 patients were analyzed in duplicates) (Badea et al., Hepato-Gastroenterology 2008: 55:2015-2026), analyzed in oncomine and presented as box and whisker plots. Core 3 synthase expression was down regulated in tumor specimens compared to normal pancreas. The log2 median-centered intensity for normal and tumor specimen is (-0.0115) and (-0.758), respectively. Oncomine™ (Compendia Bioscience, Ann Arbor, MI) was used for analysis and visualization.

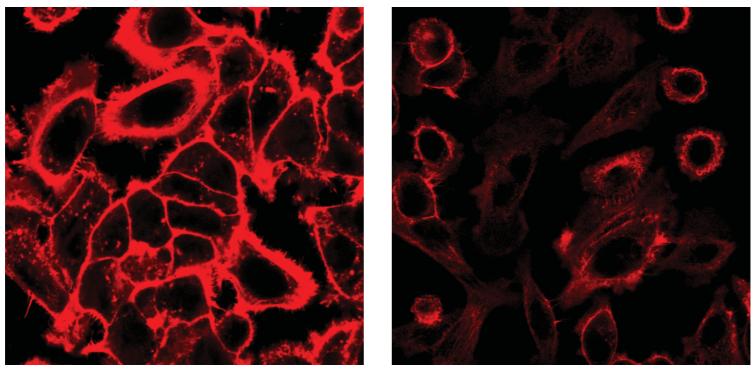
Supplementary Figure S2



Core 3 synthase downregulates FGFR3 and REG1 α . Real-time PCR analysis of FGFR3 (b) and REG1 α (b) expression in core 3 synthase and vector control expressing FG cells. The values are the average of the three experimental means obtained and error bars represents S.D of mean. A p-value less than 0.05 was considered to be statistically significant.

Supplementary Fiugre S3

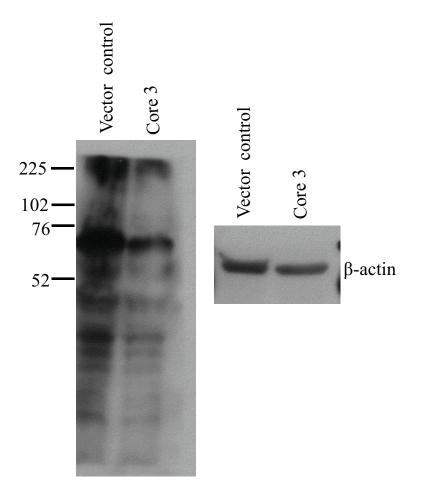
Vector control



Core 3

Core 3 and vector control FG cells were stained with Rhodamine conjugated Phalloidin for detection of F-actin polymerization. Core 3 expressing FG cells show reduced Phalloidin staining intensity.

Supplementary Figure S4



FG cell lysates (control or expressing core 3) were treated with neuraminidase (Millipore, Billerica, MA, USA) and then western blotted with T-antigen specific monoclonal antibody 3C9 (a kind gift from Dr. Henrik Clausen, Department of Cellular and Molecular Medicine, University of Copenhagen, Denmark). We observed decreased T antigen expression in core 3 synthase expressing cells compared to vector control cells. β-actin is used as internal control.