Figure S1: Correlation of dry cell weight and BioLector online backscatter signal. DCW concentrations of samples from five microscale cultivation runs in BSM_{mod} (4% D-glucose, 150 mM PIPPS, pH 5.0, 0.8 mL, 1500 rpm, 30°C) were plotted against online backscatter values (620 nm, gain 15) and fitted to a third degree polynomial (solid line, R² = 0.99). DCW and backscatter were determined as triplicates, error bars show standard deviations.
Figure S2: Bioreactor cultivation of *P. pastoris* on D-glucose. Cultivations were performed in BSM 
mod (4% D-glucose, 0.8 L, pH = 5.0 (NH₄OH / H₂SO₄), DO = 30%, inoculated to OD₆₀₀ = 1.0). 
Results from one representative out of three cultivations are shown.
Figure S3: Online growth data for parallel cultivation of 47 transformants of AppA phytase secreting \textit{P. pastoris}::pGAPZ\textit{aB_appA}. DCW concentrations (A), DO (B) and pH (C) are shown for a clonal screening in BSM\textsubscript{mod} plus 4\% D-glucose (150 mM PIPPS, pH 5.0, 0.8 mL, 1500 rpm, 30°C). DCW concentrations were calculated from online backscatter measurements. Each clone was cultivated in triplicate, lines show mean values.
Figure S4: Specific growth rates for 47 transformants from a clonal library of AppA phytase secreting *P. pastoris*::pGAPZαB_appA. Values were calculated from the online biomass signal with the help of a non-linear correlation (figure S1). Each clone was cultivated in triplicate, error bars show standard deviations. Boxes show the 25 – 75 percentile, median (line) and mean value (square). Whiskers represent minimum and maximum values.
Figure S5: Biomass yields for 47 transformants from a clonal library of AppA phytase secreting *P. pastoris*::pGAPZαB_appA. Biomass yields ($Y_{X/S}$) were calculated from DCW concentrations determined experimentally at the end of the cultivation. Each clone was cultivated in triplicate, error bars show standard deviations. Boxes show the 25 – 75 percentile, median (line) and mean value (square). Whiskers represent minimum and maximum values.
Figure S6: SDS-PAGE analysis of phytase production during microscale cultivation under carbon-limited conditions. Cultivation was performed in 0.8 mL BSMmod at pH 5.0 with 2% glycerol as batch substrate and 10% dextrin (1300 rpm and 30°C). At three time points, 25 U/L amyloglucosidase was added. For all clones 20 µL of culture supernatant sampled after 72 h were analyzed.