





Supplemental Figure legends

Supplemental Figure 1. Determination of the concentration of NmR in the culture supernatant of the *nrk1* Δ *urh1* Δ *pnp1* Δ mutant. A standard curve is established by supplementing a series of 8 mL culture of the *npt1* Δ *qpt1* Δ mutant (recipient cells, staring OD₆₀₀=0.05) with 50 μ L of 5 μ M, 10 μ M, 15 μ M or 20 μ M of enzymatically synthesized NmR following by determining the growth (OD₆₀₀) after 20 hr incubation at 30°C. The equation y = 0.1564x - 0.1512 is then generated to formulate the relationship between NmR concentrations (x) and growth (OD₆₀₀) of the recipient cells (y). The *npt1* Δ *qpt1* Δ mutant is also treated with 50 μ L of supernatant from the overnight culture supernatant of the *nrk1* Δ *urh1* Δ *pnp1* Δ mutant is therefore estimated to be 6.74 μ M (y = 0.903) using the equation generated with the standard set. One set of representative data conducted in triplicate is shown. Error bars denote standard deviations.

Supplemental Figure 2. Comparisons of the efficiency of NmR and NaR to support the growth of the *npt1* Δ *qpt1* Δ mutant. The *npt1* Δ *qpt1* Δ mutant (recipient cells, staring OD₆₀₀=0.05) is supplemented with indicated final concentrations of chemically synthesized NmR (left) or NaR (right). Growth of the recipient *npt1* Δ *qpt1* Δ mutant is determined (OD₆₀₀) after 20 hr incubation at 30°C. Results show the relationship between concentrations (the x axis) of NmR (left) or NaR (right) and growth (OD₆₀₀) (the y axis) of the recipient cells. NmR appears to be a more efficient NAD⁺ precursor (y = 6.722x + 1000).

0.0593) compared to NaR (y = 0.0696x + 0.1029). One set of representative data conducted in triplicate is shown. Error bars denote standard deviations.

Supplemental Figure 3. Analysis of the role of other Sir2 family members, Hst1 and Hst2, in NmR salvage. (A) Deletions of HST1 (left) or HST2 (right) do not cause severe growth defect in the $npt1\Delta nrk1\Delta$ mutant grown in rich medium. (B) NmR supplement supports the growth of the $npt1\Delta qpt1\Delta$ recipient cells, which is further enhanced by deletion of SIR2 but not by deletions of HST1 or HST2. One set of representative data conducted in triplicate is shown. Error bars denote standard deviations. For (B), NmR is supplemented to the growth media at a final concentration of 0.1 μ M.