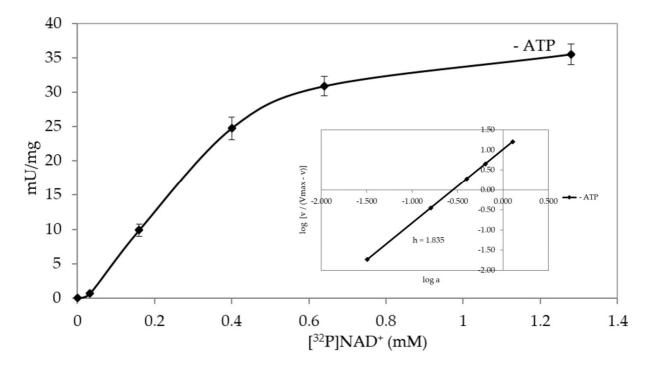
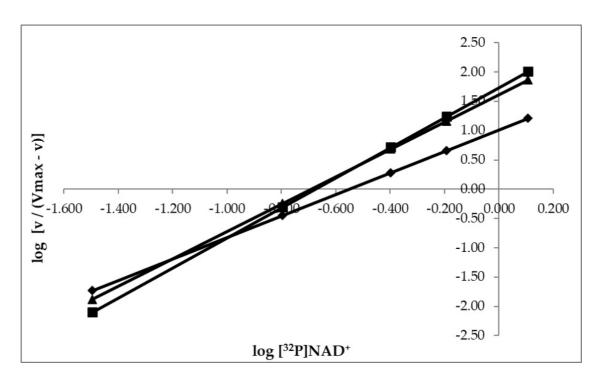
## **Supplementary Material**



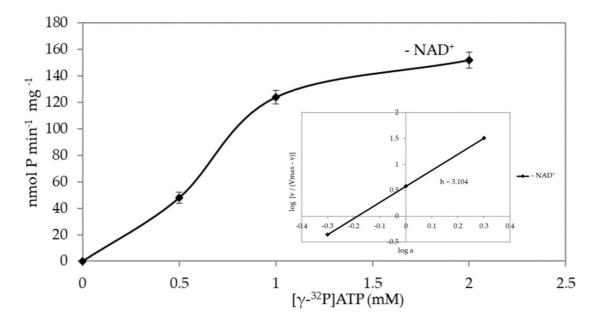
**Figure S1.**PARPSso Activity. The sigmoid curve was determined by measuring purified PARPSso activity at increasing concentrations of [32P]NAD+ (10,000 cpm/nmol). The reported values are means of 4 different assays with two enzyme preparations. The inlet shows the linearization of the curve by Hill equation:

$$\log [v/(V-1)] = h \log a - h \log K_{0.5}$$

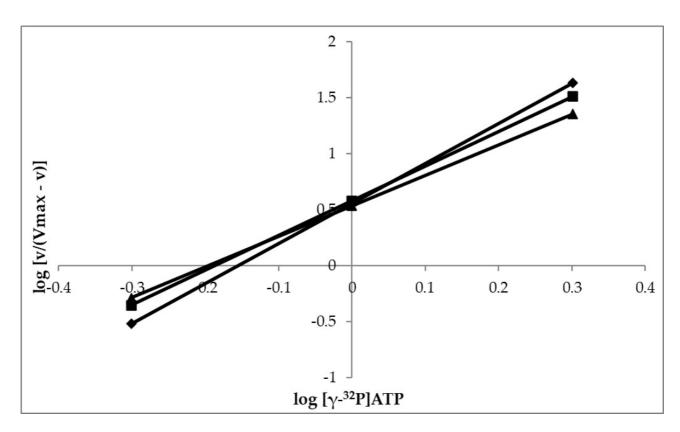
Where V= Vmax; a= substrate concentration; K  $_{0.5}$  = substrate concentration at half of Vmax; h= nH= Hill coefficient measured by the slope of the plot.nH=1.8.



**Figure S2.** Linearized plots of ADPribosylating activity in presence of different ATP concentrations ( $\mu$ M: 0,  $\spadesuit$ ; 5,  $\blacksquare$ ; 10,  $\blacktriangle$ ). At 0.4mM [ $^{32}$ P]NAD $^{+}$  Ki was -7.9·10 $^{-3}$  (at 5  $\mu$ M ATP), and -1.710 $^{-2}$  (at 10  $\mu$ M ATP).



**Figure S3.**ATPase Activity of PARPSso. The sigmoid curve was determined by measuring purified PARPSso activity at increasing concentrations of  $[\gamma^{-32}P]$ ATP (10,000 cpm/nmole). The reported values are means of 4 different assays with two enzyme preparations. The inlet shows the linearization of the curve by Hill equation as described in the legend of Figure S1. nH=3.1.



**Figure S4.** Linearized plots of ATPase activity in the presence of different NAD+ concentrations ( $\mu$ M: 0,  $\blacksquare$ ; 10,  $\spadesuit$ ; 100,  $\blacktriangle$ ). At 0.5mM [ $^{32}$ P]ATP Ki was 7.3·10-3 (at 10  $\mu$ M NAD+), and 1.910-1 (at 100  $\mu$ M NAD+).

Ki was calculated by the following equation: