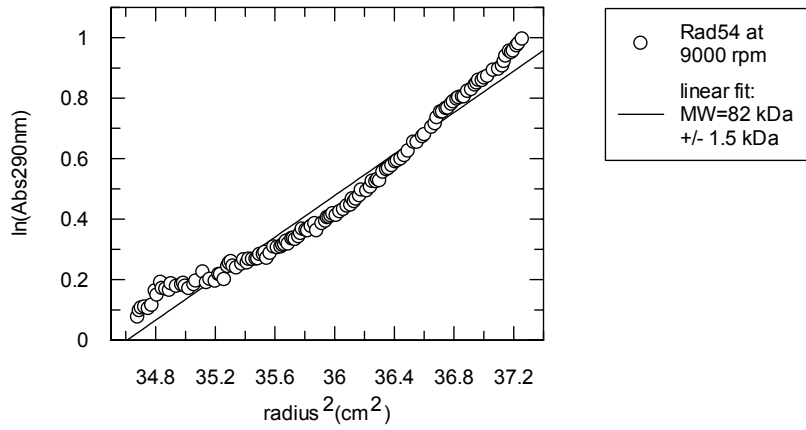
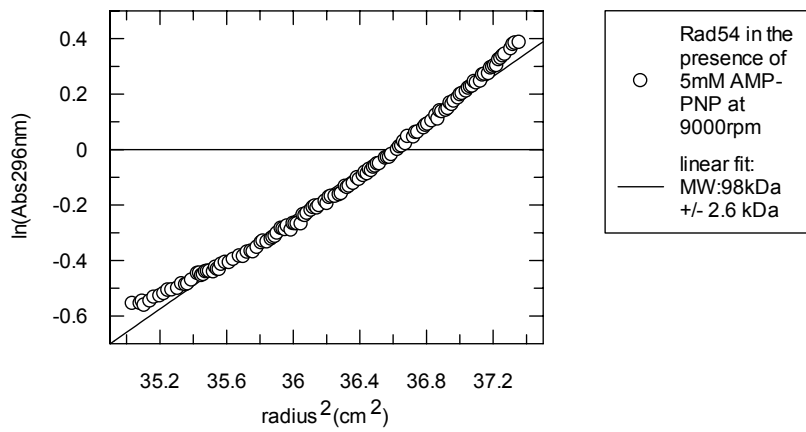


Supplementary Figure 5 Analytical ultra-centrifugation indicated that Rad54 is monomeric in solution.

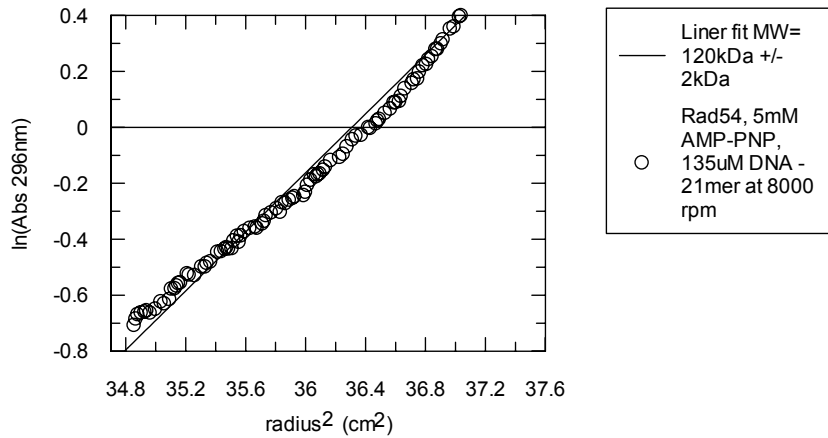
A



B



C



Analytical ultracentrifugation was carried out with 125 μ M dnRad54 Δ N in 50mM Tris 8.0, 2mM DTT, 300mM salt. The calculated molecular weight for dnRad54DN is \sim 83 kDa. The observed value for dnRad54DN was $MW_{\text{obs}} \sim 82 \pm 1.5$ kDa (A). Addition of 5mM AMP-PNP and 10mM $MgCl_2$ gave a $MW_{\text{obs}} \sim 98 \pm 2.7$ kDa (B). Addition of a double stranded 21mer DNA fragment at 135 μ M ($MW \sim 13.6$ kDa) in the presence of AMP-PNP (5mM) and Mg^{2+} resulted in a species of $MW_{\text{obs}} \sim 120 \pm 2$ kDa (C). Similar results were also obtained in dynamic light scattering (data not shown). On the basis of these experiments it appears that dnRad54 Δ N is a monomer in the presence and absence of AMP-PNP (A,B). Furthermore, the dsDNA fragment appears bound by the enzyme, but does not change the aggregation state (C).