## Probing the nanoparticles-Ago2 interaction for

## enhanced gene knockdown

Sonal Deshpande<sup>1</sup> and Neetu Singh<sup>1,2</sup>\*

<sup>1</sup>Centre for Biomedical Engineering, Indian Institute of Technology-Delhi, Hauz Khas, New Delhi-110016, India.

<sup>2</sup>Biomedical Engineering Unit, All India Institute of Medical Sciences, Ansari Nagar, New Delhi-110029, India.

## **Supplementary Information**

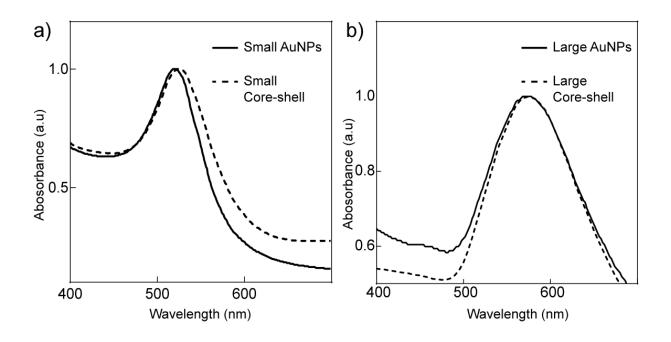
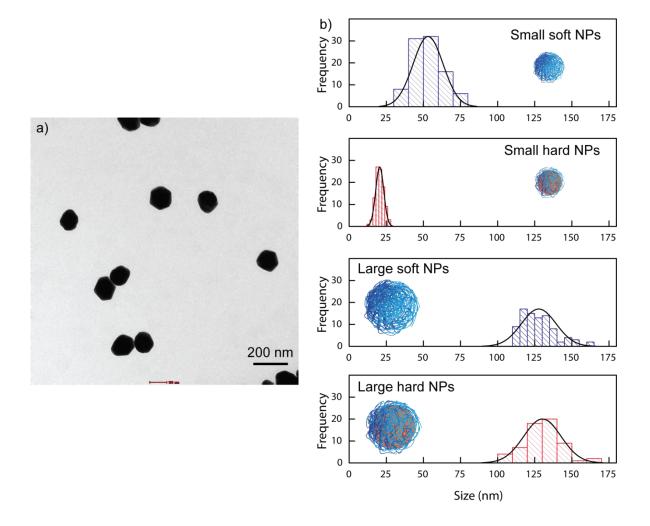


Figure S1: Absorbance spectra of AuNPs, before and after shell synthesis.



**Figure S2.** a) TEM micrograph of  $Hard_{Large}$  nanoparticles. b) Size distribution of nanoparticles by TEM micrograph analysis. Atleast 60 nanoparticles were used for analysis.

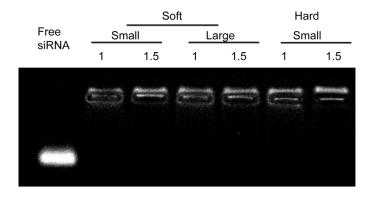


Figure S3. Optimization of siRNA loading in the nanoparticles by agarose gel electrophoresis

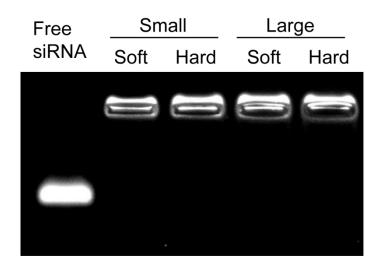
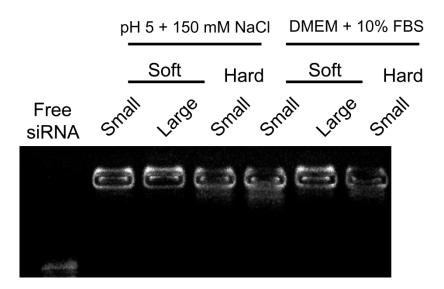


Figure S4: Loading of siRNA in the nanoparticles



**Figure S5.** Electrophoretic mobility assay stability of siRNA in FBS supplemented media and at higher ionic strength and low pH.

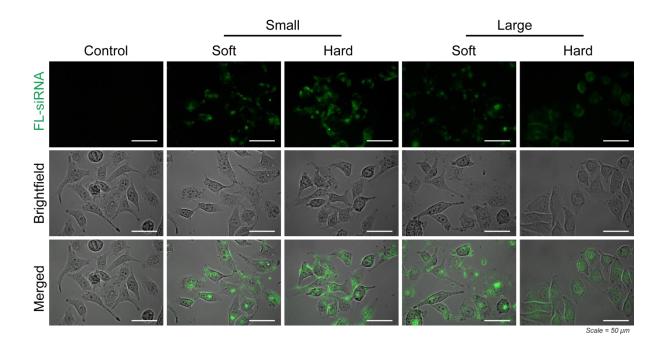
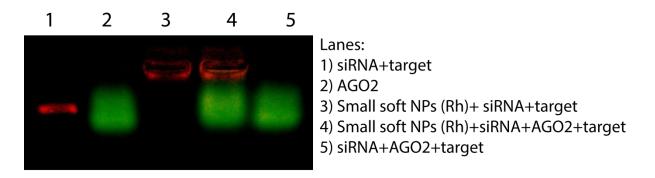
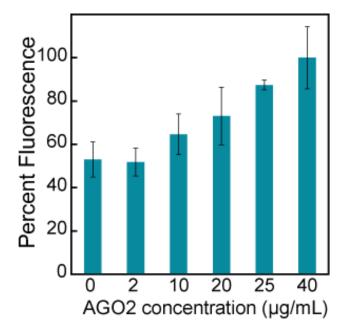


Figure S6: Uptake of FL-siRNA loaded nanoparticles by HeLa cells by fluorescence microscopy.



**Figure S7.** Colocalization of rhodamine (Rh) tagged small nanogel and fluorescein labelled AGO2 in presence of siRNA and target, by electrophoretic mobility assay.



**Figure S8**. Assay showing AGO2 concentration depended increase in fluorescence in AGO2siRNA interaction study.

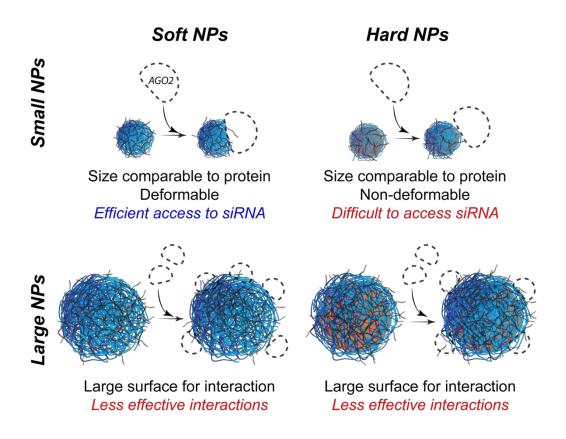


Figure S9. Schematic representation of the hypothesis for interaction of nanoparticles of different size and softness