

Supplementary Material

*Joseph M. Slocik, Rajesh R. Naik, Morley O. Stone, Trevor Douglas, Mark Young
and David W. Wright**

- S-1.** UV-Vis spectra of gold reduced by (HRE)-SubE virus over 4 hours sampled every 30 seconds in a septum sealed 1.5 mL quartz cuvette in water.
- S-2.** Time plots of absorbance at 550 nm for reduction of AuCl_4^- with (HRE)-SubE, Wild type virus, and SubE virus expressed in yeast in water.
- S-3.** Time plots of absorbance at 550 nm for AuCl_4^- with HRE peptide, YHRE peptide, Y_4HRE peptide, and the tripeptide of YYY in water.
- S-4.** Time plots of fluorescence over four hours of (HRE)-SubE with AuCl_4^- in water. Excited at 270 nm and 295 nm, while fluorescence was monitored at 352 nm. Excitation slit set at 10 nm, emission slit at 5 nm, 30 second cycles, average time 1 second, PMT voltage 800 V.
- S-5.** Spatial relationship between tyrosine and tryptophan residues within CCMV subunit.
- S-6.** Time plot of fluorescence for $\text{AuCl}_4^-/\text{YYY}$ tripeptide in water with same parameters as S-4.
- S-7.** Gold titration plot of (HRE)-SubE virus. 20 μL of purified (HRE)-SubE (stock concentration 0.2 $\mu\text{g}/\mu\text{L}$) was added to 750 μL of a 50 mM HEPES buffer pH 7.5 in a septum sealed 1.00 mL quartz cuvette. A 5 μL aliquot of a 1 mM stock solution of HAuCl_4 (Strem) was added to (HRE)-SubE and incubated at room temperature for 1 hour at which time the UV-Visible spectrum was obtained on a Hewlett Packard Agilent 8453 photodiode array spectrophotometer. Subsequent 5 μL additions followed with 1 hour incubation period between additions and monitored until the absorbance at 550 nm became constant.

S-8. Time plots of absorbance for the metal precursors of Ag^+ , PtCl_6^{2-} , and PdCl_6^{2-} with (HRE)-SubE virus. Absorbance monitored at 300 nm for PdCl_6^{2-} /SubE, at 410 nm for Ag^+ , and at 600 nm for PtCl_6^{2-} .

S-9. Time plots of absorbance at 550 nm of (HRE)-SubE/ AuCl_4^- buffered over the pH range of 4.0 – 9.0 with acetate, phosphate, borate, and Tris buffers prepared at 50 mM and adjusted with HCl or NaOH to desired pH.

S-10. pH profile of (HRE)-SubE reduction of gold vs absorbance at 550 nm at 4 hours from time plots.

S-11. TEM micrographs of native viral capsids negatively stained with 2 % uranyl acetate. (A) (HRE)-SubE capsid. (B) Wild type capsid. (C) SubE capsid expressed in yeast.

S-12. TEM histograms of gold reduced by virus in water. (A) (HRE)-SubE/ Au^0 . (B) Wild type/ Au^0 . (C) SubE (yeast)/ Au^0 .

S-13. EDS spectra of viral/ Au^0 nanoparticle structure (negatively stained with 2 % uranyl acetate). Copper peaks are from TEM grid.

S-14. TEM image of AuCl_4^- reduced by YYY tripeptide in water.

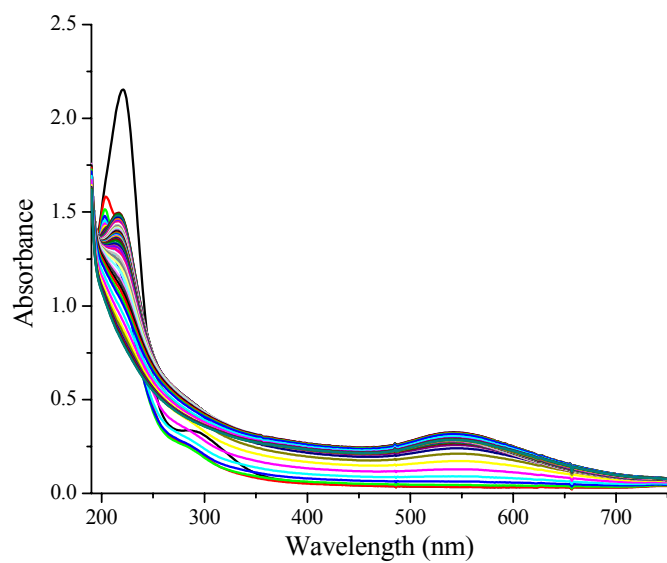
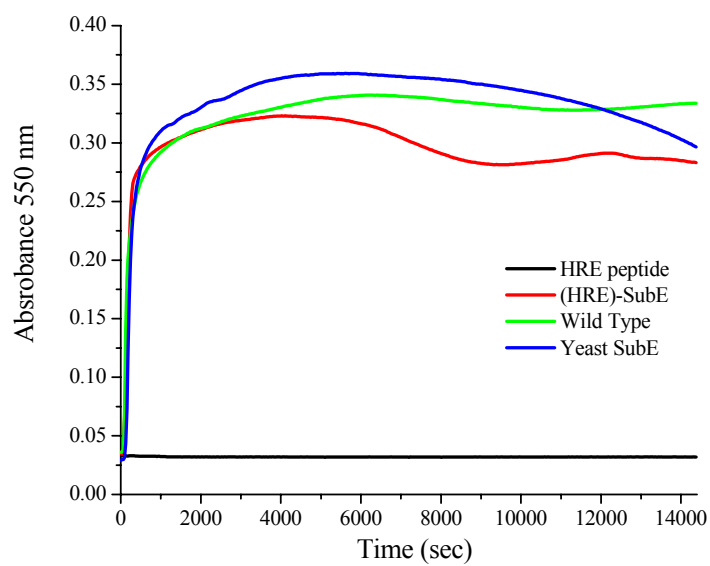
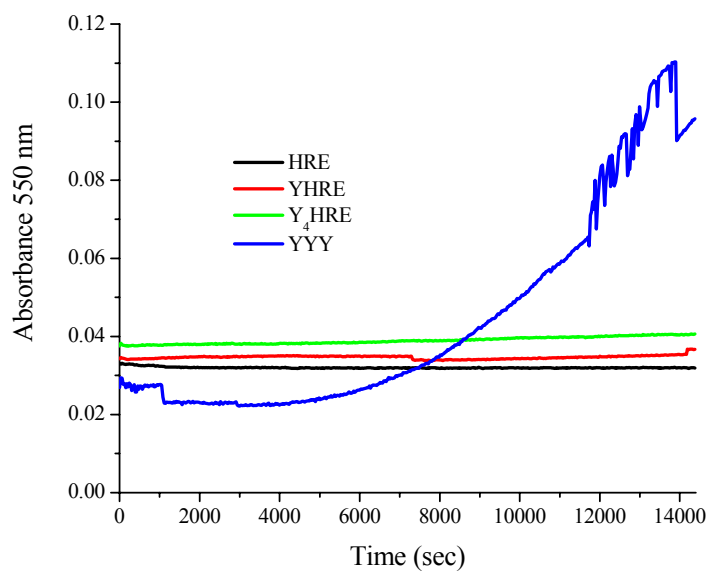
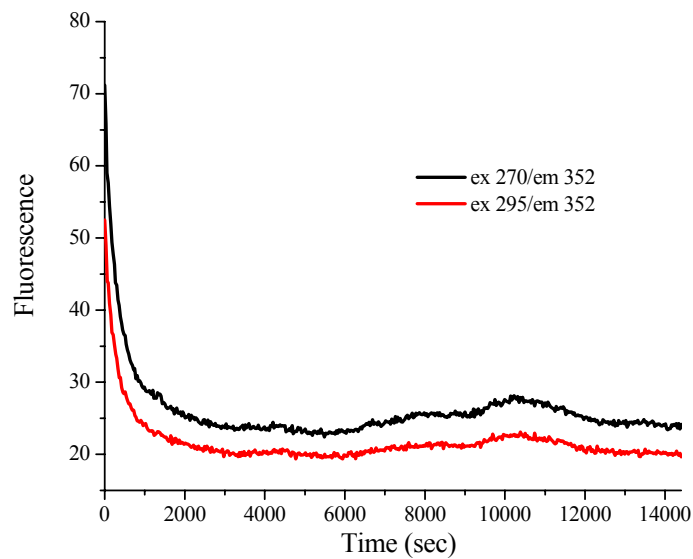
S-15. TEM micrographs of Au^0 /virus structures from borohydride reduction of virus saturated with $\text{AuClP}(\text{CH}_3)_3$. (A) SubE (yeast)/ Au^0 . (B) Wild type virus/ Au^0 .

S-16. TEM histograms of Au^0 from S-15. (A) Au^0 /(HRE)-SubE. (B) Au^0 /Wild type. (C) Au^0 /SubE (yeast).

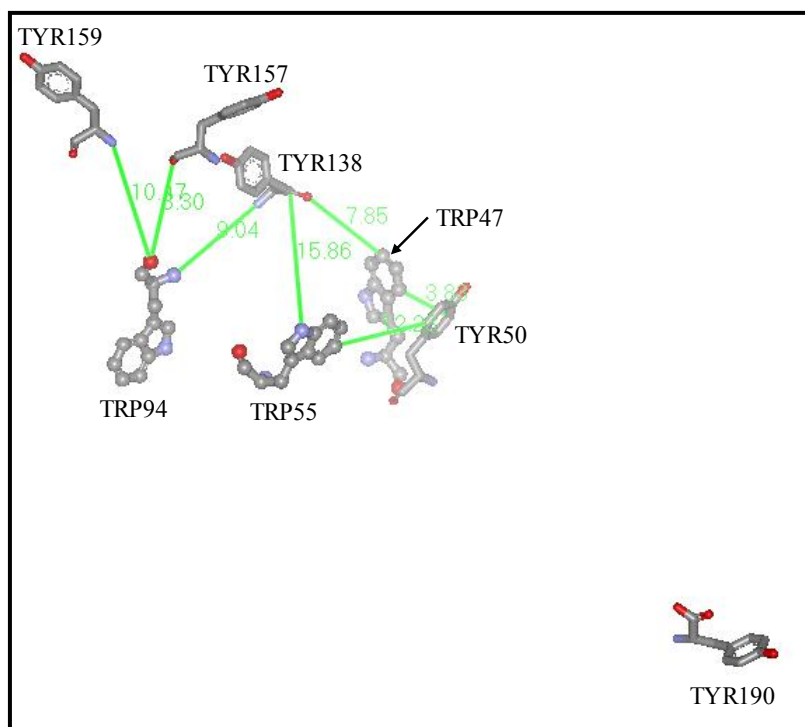
S-17. TEM images of Au^0 /SubE (yeast) from borohydride reduction of $\text{AuClP}(\text{CH}_3)_3$ tilt at -40° , -20° , 0° , $+20^\circ$, and $+40^\circ$ with respect to image plane.

S-18. TEM of Au^0 /SubE (yeast) diluted 10 fold from concentrated sample.

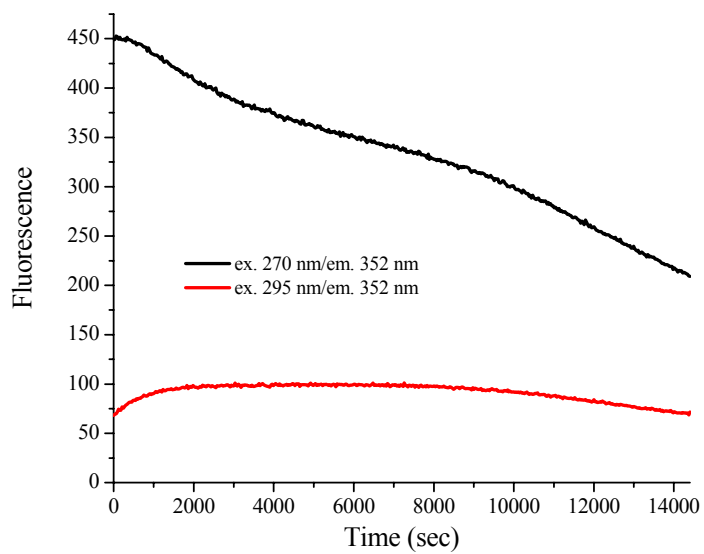
S-19. UV-Vis absorbance spectra of Au⁰/SubE (yeast) from AuClP(CH₃)₃ (diluted 10 fold) and Au⁰ from virus modified with diethylpyrocarbonate (undiluted).

S-1.**S-2.****S-3.****S-4.**

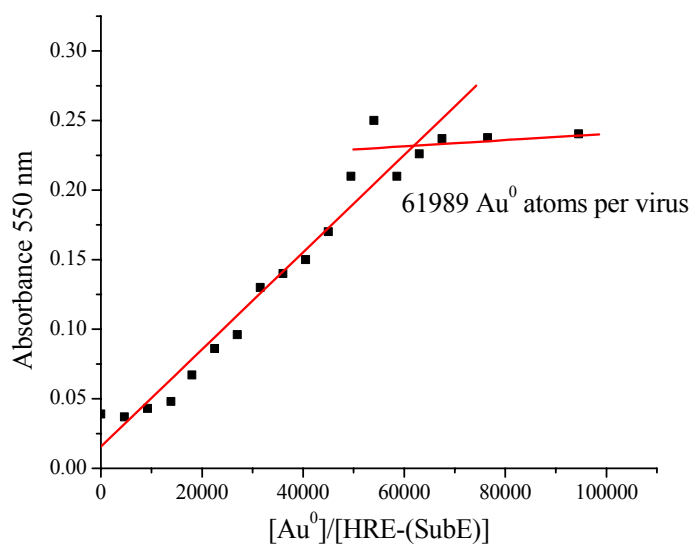
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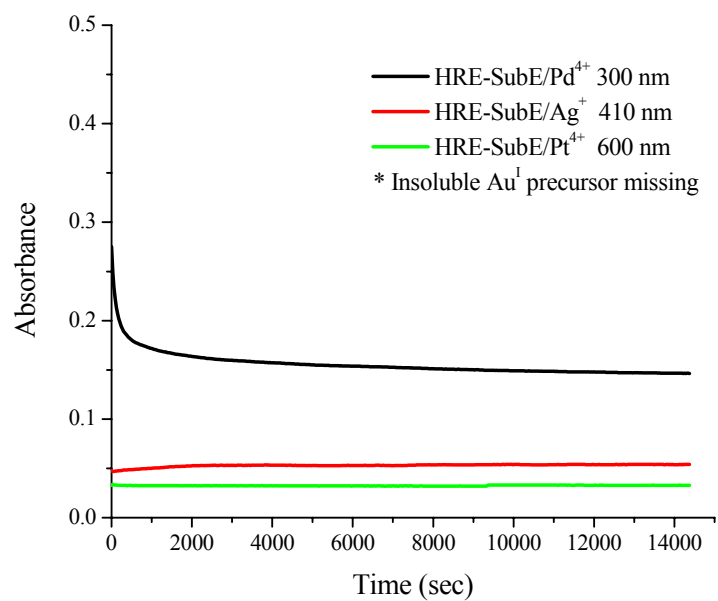
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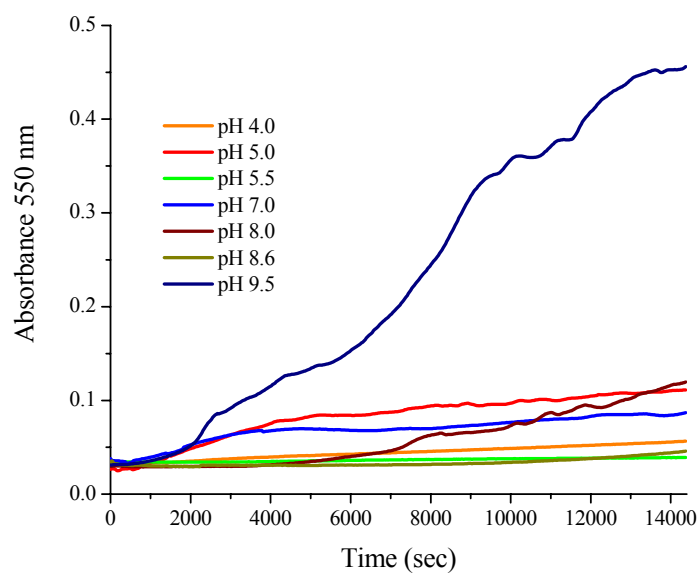
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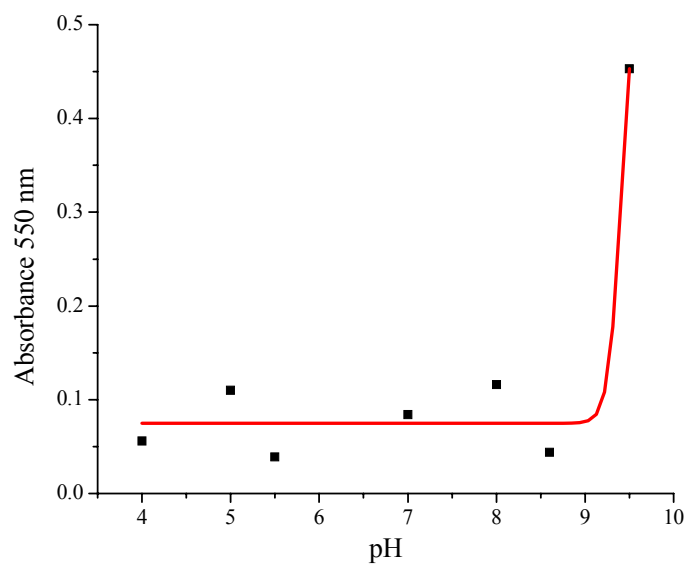
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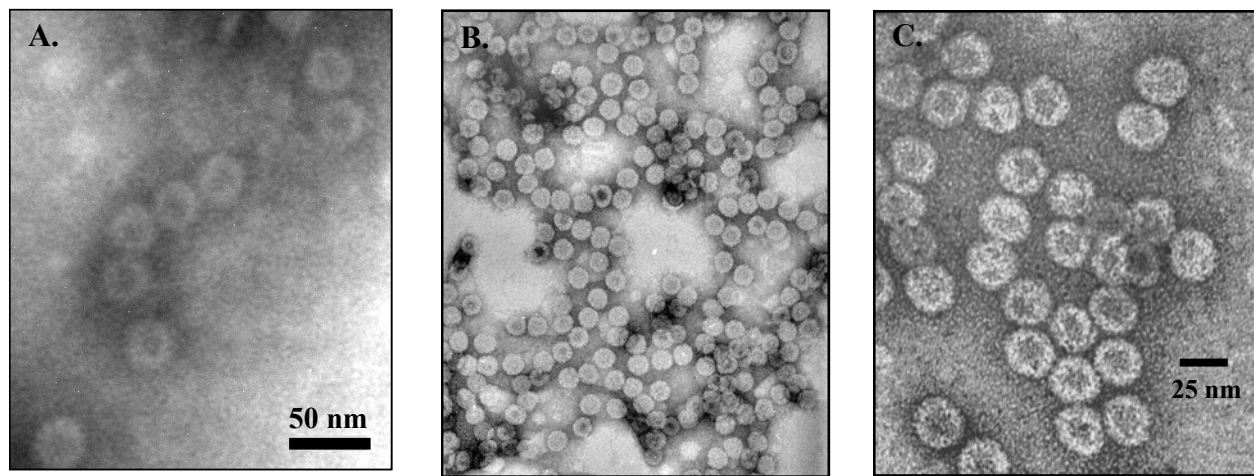
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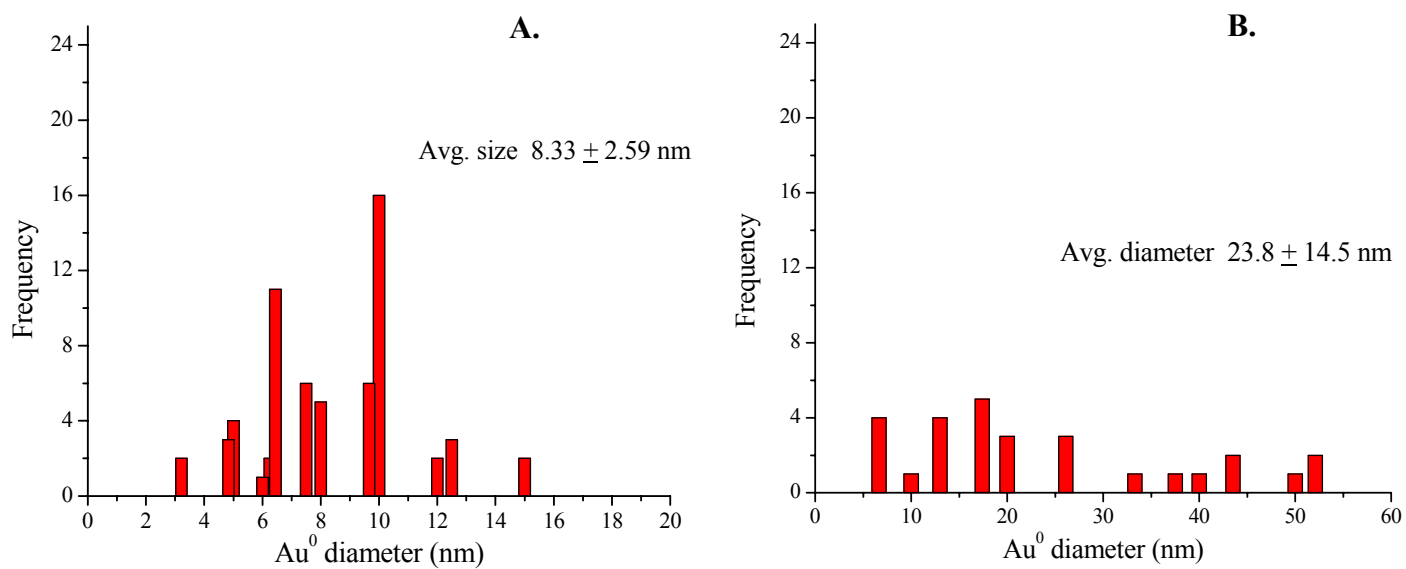
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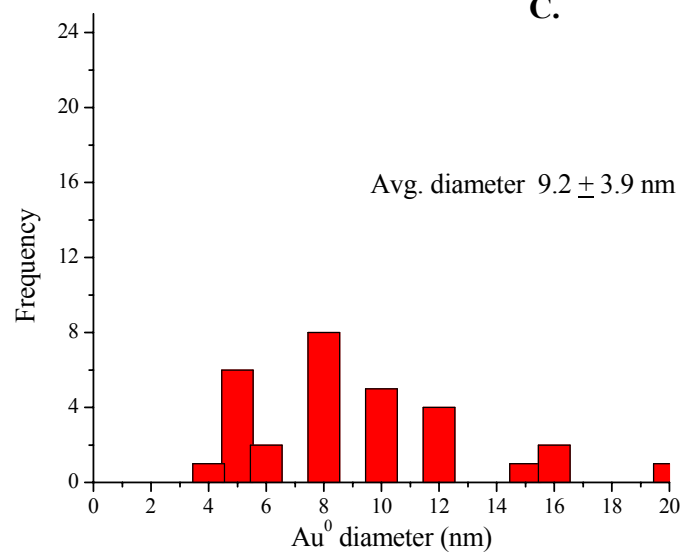
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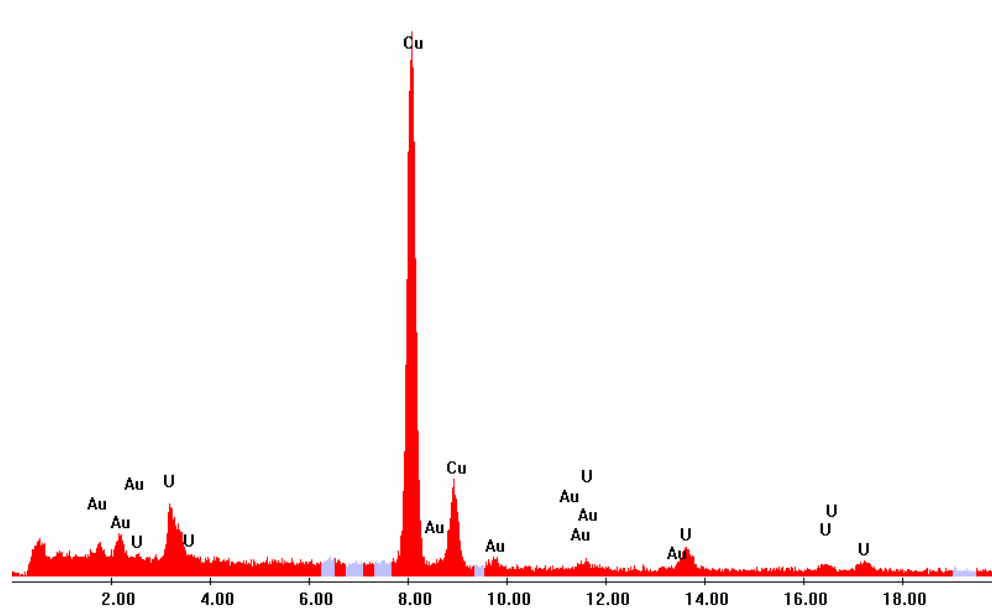
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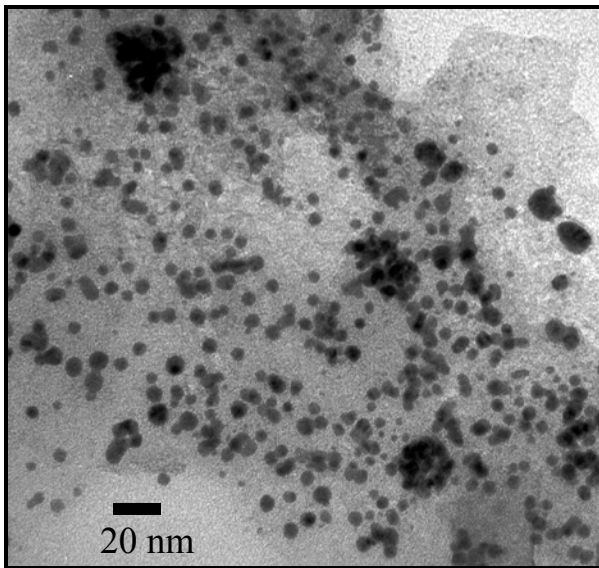
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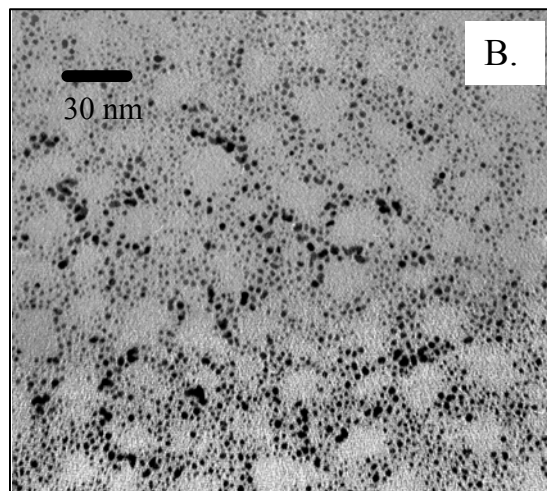
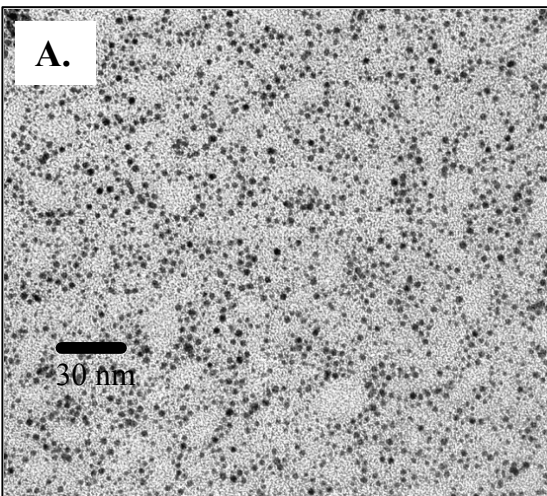
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S-14.

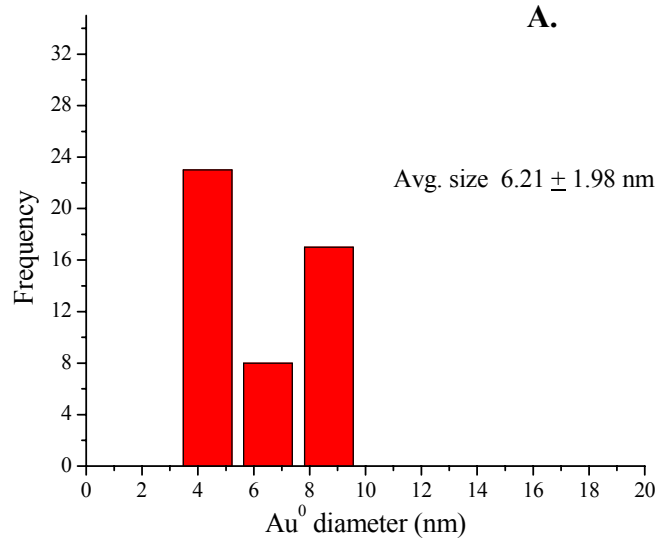


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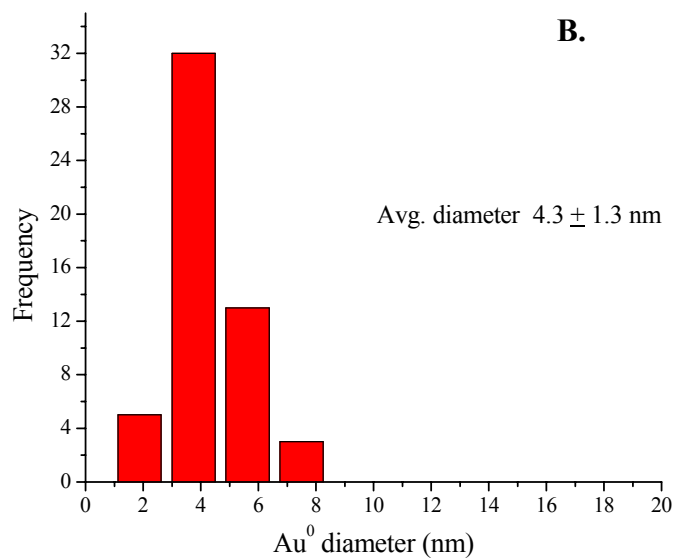


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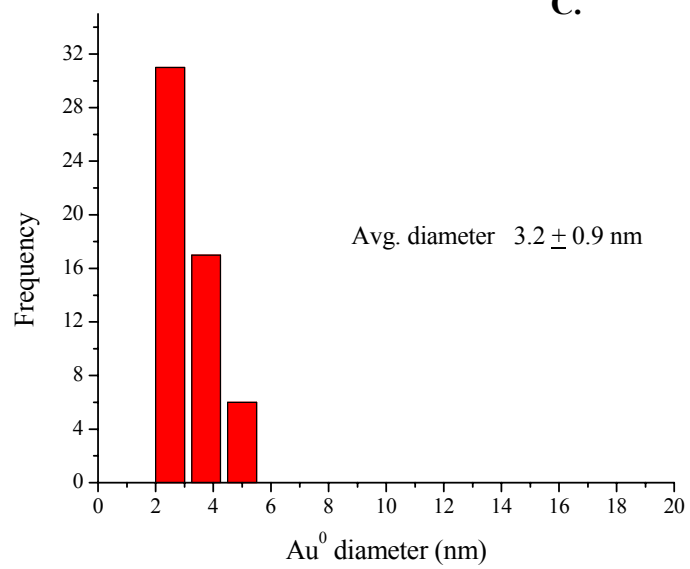
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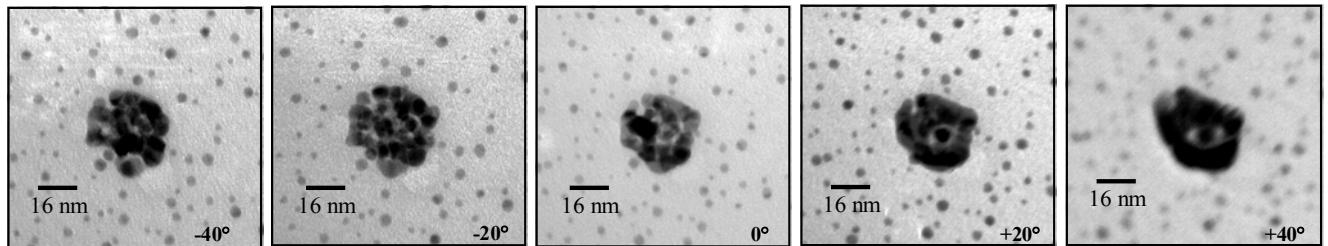
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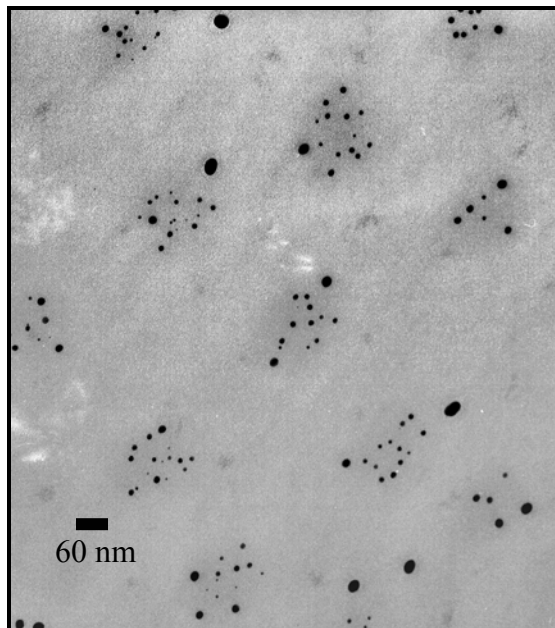
C.



S-17.



S-18.



S-19.

