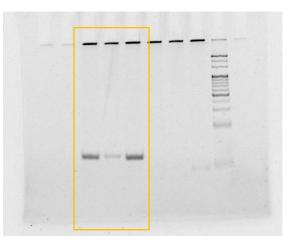
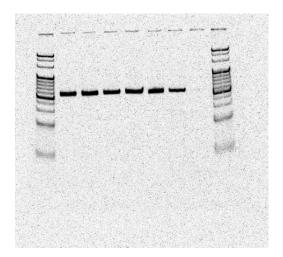


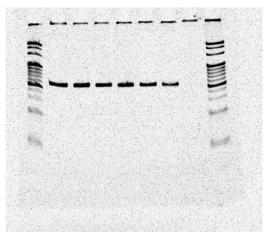
**Figure 1j.** Evaluation of the integrality of DNA strands by polyacrylamide gel electrophoresis.



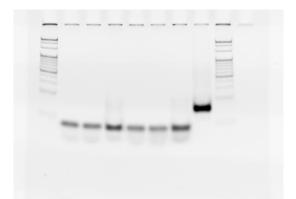
**Figure 1j.** Evaluation of the integrality of RNA strands by polyacrylamide gel electrophoresis.



**Supplementary Figure 7b.** Gel electrophoresis analysis of PCR products amplified by different concentrations of DNA strand with or without MW-assisted heating-dry treatment.



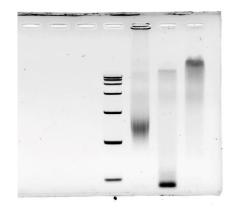
**Supplementary Figure 7b.** Gel electrophoresis analysis of PCR products amplified by different concentrations of DNA strand with or without MW-assisted heating-dry treatment.



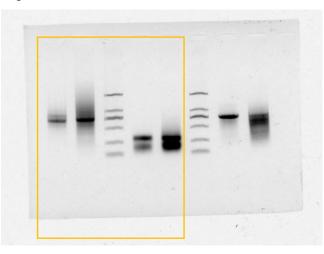
**Supplementary Figure 8b.** Gel electrophoresis analysis of short-chain ssDNA with or without MW-assisted heating-dry treatment in the absence/present of endonuclease.

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**Supplementary Figure 8b.** Gel electrophoresis analysis of long-chain EGFP plasmid DNA with or without MW-assisted heating-dry treatment in the absence/present of endonuclease.



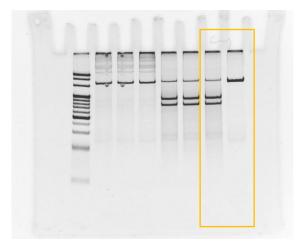
**Figure 6e.** Gel electrophoresis analysis of RCA products with perfectly matched or singlebase mismatched target DNA.



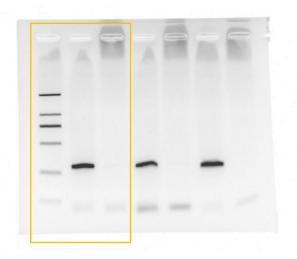
**Figure 6I.** Gel electrophoresis analysis of the in vitro transcribed sgRNA and N-RNA, and sgRNA and N-RNA displaced from AuNPs surface by ME.

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**Figure 6m.** Electrophoresis analysis of the cleavage ability of the assembled Cas9/sgRNA-AuNP complex.



**Supplementary Figure 19a. Gel electrophoretic analysis.** Cleavage ability evaluation using designed poly (U)-tagged sgRNA and Cas9 protein.



**Supplementary Figure 19b. Gel electrophoretic analysis.** Gel electrophoretic analysis of the PCR products of *B646L* (VP72) gene.