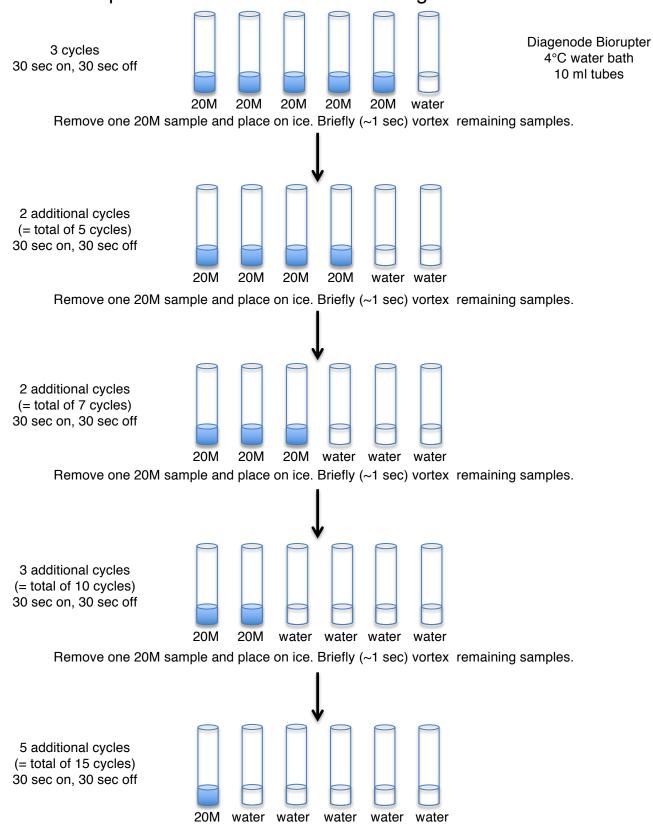
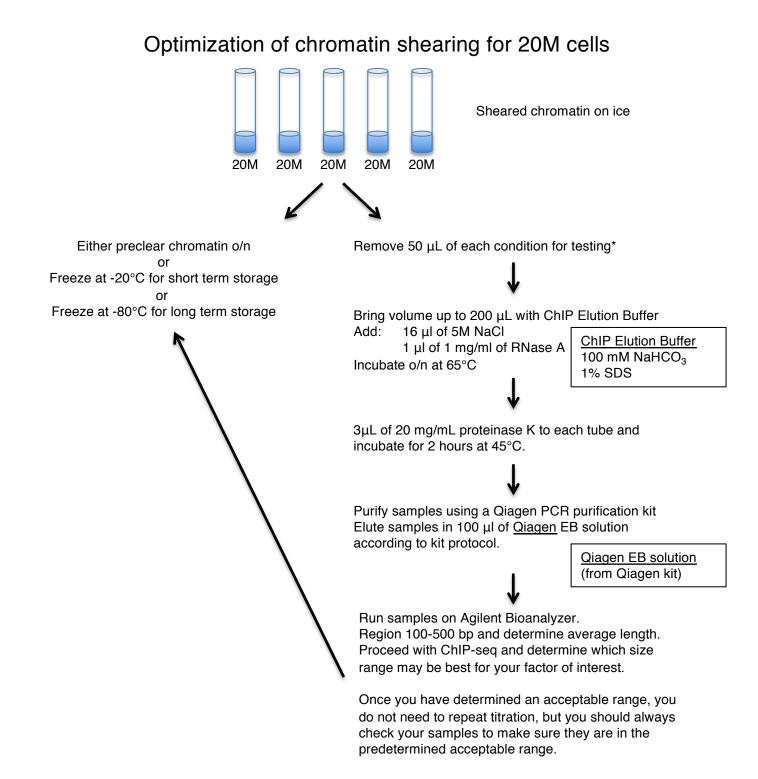
## **Supplemental Figure S1**

Optimization of chromatin shearing for 20M cells



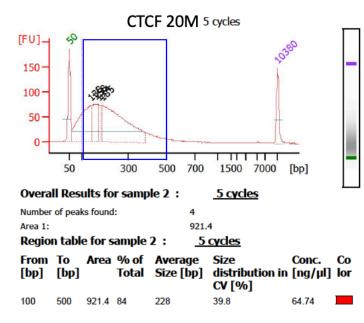
Remove last 20M sample and place on ice

## **Supplemental Figure S1 continued**



\*You can use less than 50  $\mu$ l, however, we always increase the volume to 200  $\mu$ l with ChIP elution buffer and process samples in the same way that we process ChIP samples for reverse cross-linking. If you sample less than 50  $\mu$ l, please also remember to decrease your elution volume from Qiagen column accordingly.

## **Supplemental Figure S2**



Peak table for sample 2 :				<u>5 cycles</u>
Peak		Size [bp] Conc. [ng/µl]		Molarity [nmol/l]
1	4	50	8.30	251.5
2		138	21.05	231.2
3		158	9.77	93.9
4		174	3.32	28.9
5		185	37.36	305.8
6	•	10,380	4.20	0.6

**Supplemental Figure S2:** Representative Aglient Bioanalzyer 2100 tracing of a chromatin sample using an Agilent DNA 7500 kit. Following sonication, the average size of each chromatin sample was determined by measuring the average length within a 100-500 bp window.