Nuclei EXtraction by SONication - NEXSON

Scientific Relevance

- ChIP-Seq is a powerful tool to map DNA binding proteins as well as histone modifications across the genome (1, 2).
- To allow the comparison of different studies and different input materials standardization of the sample preparation steps are required, which has been challenging (3, 4).
- NEXSON provides a simple but highly reproducible technique for efficient nuclei isolation, ensuring the generation of comparable chromatin maps from different sample types (5).

Challenges

- Huge variations in sample preparation for ChIP make data comparison challenging.
- In particular, the chromatin shearing step is tedious to optimize and standardize for different input materials and across different laboratories.
- Insufficient chromatin shearing compromises the quality of chromatin and sequencing results.
- Shearing optimization is labor-intensive, material-consuming, expensive, and especially difficult when working with precious samples such as clinical specimens.

Workflow

Fixed Cells
Nuclei Extraction
Shearing
IP

Schematic representation of NEXSON workflow (6)
Nuclei from fixed cells are efficiently isolated applying tuneable AFA. Pure fractions of isolated nuclei are subjected to ChIP workflow. In brief, chromatin is sheared using AFA and subjected to IP for histone modifications or transcription factors of interest.

Advantages of Adaptive Focused Acoustics® (AFA®)

AFA technology is a very gentle, reproducible, and tuneable shearing method.

- The high efficiency of nuclei isolation allows the scale down of input material to 10,000 cells/histone ChIP and 100,000 cells/transcription factor ChIP.
- Increased robustness permits comparison across different tissues, cell types, disease stages, and laboratories.
- The highly reproducible and efficient isolation of nuclei allows for novel high throughput derivates of ChIP technology (7).

Suggested Covaris Products

- Covaris Focused-ultrasonicator (M-Series, S-Series, E-Series, or LE-Series)

Citations

- Arrigoni et al. Standardizing chromatin research: a simple and universal method for ChIP-seq. NAR. (2016)