

# Pairing Uniform Whole-Genome Amplification with Simple Single-cell Sorting

Aly Krasny,<sup>1</sup> Jon Zawistowski<sup>2</sup>

1. NanoCelect Biomedical, Inc., San Diego, CA. 2. BioSkryb, Inc. Durham, NC

## Introduction

### Whole-genome sequencing of single cells involves a multitude of technical challenges:

- Physically isolating single cells
- Obtaining sufficient genomic material for downstream analyses
- Low and variable genomic coverage
- Amplification artifacts such as allelic bias, mutations, and chimeras

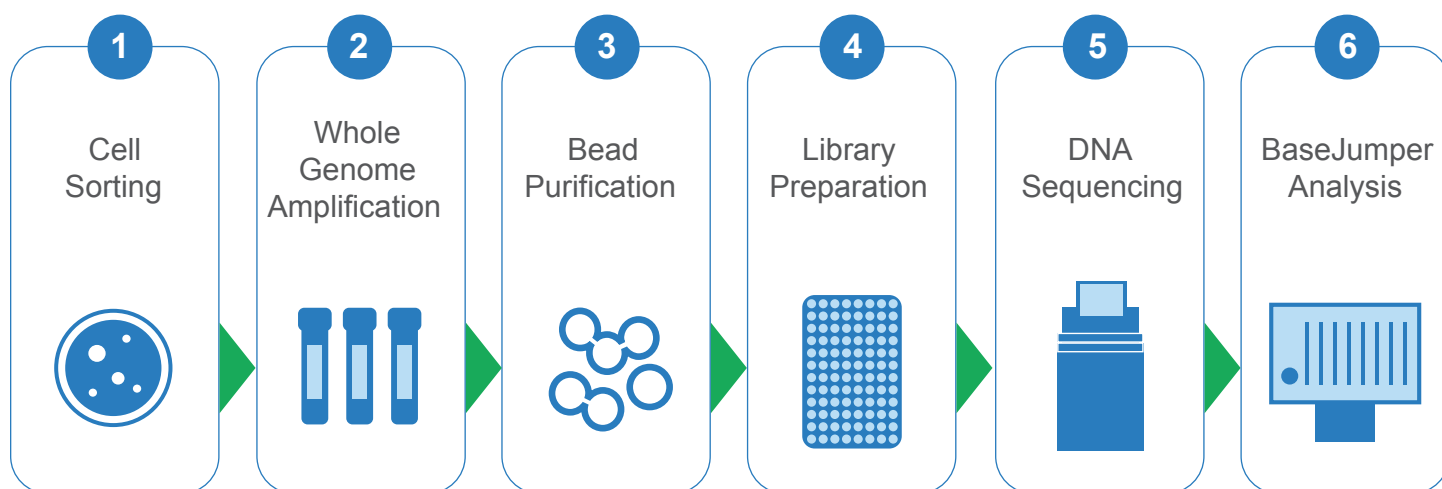
However, NanoCelect and BioSkryb have established a workflow to simplify this process and bypass these hurdles. This workflow profile highlights how next-generation cell sorting and sequencing technologies can be used to obtain high quality single-cell genomic data.

### Upstream, NanoCelect's WOLF Cell Sorter:

- Gently and accurately isolates single cells
- Confidently avoids contamination concerns
- Easily integrates into any workflow with a small footprint <2 ft<sup>3</sup> and a user-friendly interface
- Affordably fits any budget

### Downstream, BioSkryb's ResolveDNA™ platform:

- Directly copies single-cell genomes or low-input DNA with Primary Template-direct Amplification<sup>1</sup>
- Amplifies with unprecedented genomic coverage uniformity and breadth
- Precisely thwarts error propagation with high allelic balance
- Collectively, enables hitherto unachieved confidence in single nucleotide variant (SNV) and copy number variation (CNV) calling from single cells



**Figure 1. WOLF and ResolveDNA™ Workflow:** WOLF-sorted single cells undergo Primary Template-directed Amplification, followed by ligation or tagmentation-based library preparation, sequencing, and analysis with BaseJumper software.

## ▼ Isolate single cells with the WOLF

**EASILY** fit the entire instrument into the smallest biosafety cabinets with <2 ft<sup>3</sup> footprint. Master sorting quickly with user-friendly software, abundant training, and exceptional customer service.



**CONFIDENTLY** avoid contamination from previous experiments with disposable sterile cartridges and fluidics. When using pre-loaded well-specific reagents, switch from dispensing cells directly to the well bottom (Direct-Dispense Mode) to touching them off on the sidewall of each well (Sidewall-Dispense Mode) to avoid well-to-well crossover.

**GENTLY** sort cells-of-interest without shear force and with less than 2 PSI pressure. Avoid worrying about changes to cell viability and sequence quality.

**ACCURATELY** dispense up to 74% single cells in 8.5µL droplets with Standard Mode. Or for assays requiring lower volumes, dispense with similar efficiency in 2.2µL droplets using Low-Volume Mode (Table 1).

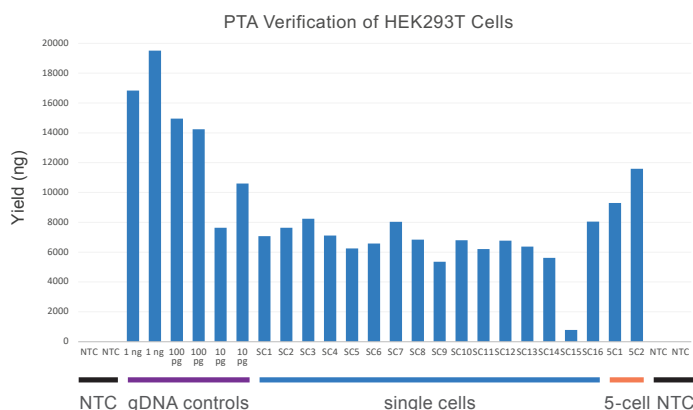
Table 1. Dispense Efficiency<sup>2</sup>

Dispense Mode	% Wells with Single Beads	
	8.5µL Droplet	2.2µL Droplet
Direct	74	69
Sidewall	72	56

Pure fluorescent beads were sorted into 3 x 96-well plates with each dispense mode combination. Plates were imaged and single-bead wells were counted. This was completed 3 times for a total of 9 plates/combination. Efficiency may be higher for cells since bead chemistry makes them likely to stick to plate sidewalls.

**Amplify genomic material with PTA**

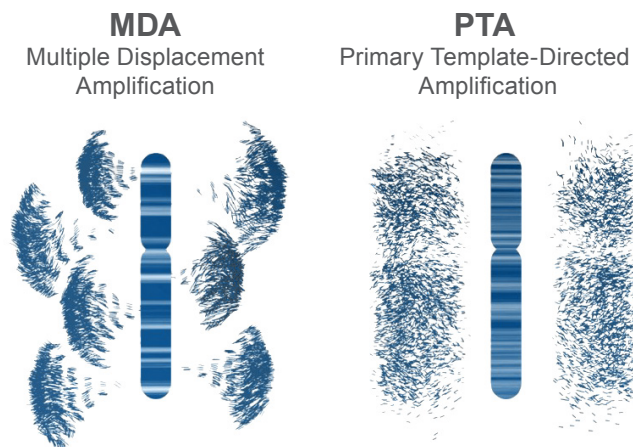
**PRECISELY** amplify low-input DNA and single-cell inputs to reproducibly capture >95% of the genome (Figure 2).



**Figure 2. Uniform single-cell yield with PTA:** Single-cell amplification reaction yield is shown relative to genomic DNA control and 5-cell input yields. Importantly, no-template control (NTC) reactions are not detectable by Qubit.

**DIRECTLY** copy the primary template with an isothermal polymerase and proprietary termination chemistry that attenuates the size of amplicons. The smaller amplicons do not efficiently amplify, so random primers are redirected to the primary template of interest.

**UNIFORMLY** amplify with high breadth of coverage, few replication errors, and low allelic bias, to accurately call single nucleotide variants (SNV) at the whole genome sequencing (WGS), whole exome sequencing (WES), and small-panel levels.



**Figure 3. Schematic of genomic amplification output obtained from MDA vs PTA:** PTA prevents exponential read pileup and error propagation and yields highly uniform coverage.

## ▼ Construct a library directly with PTA product

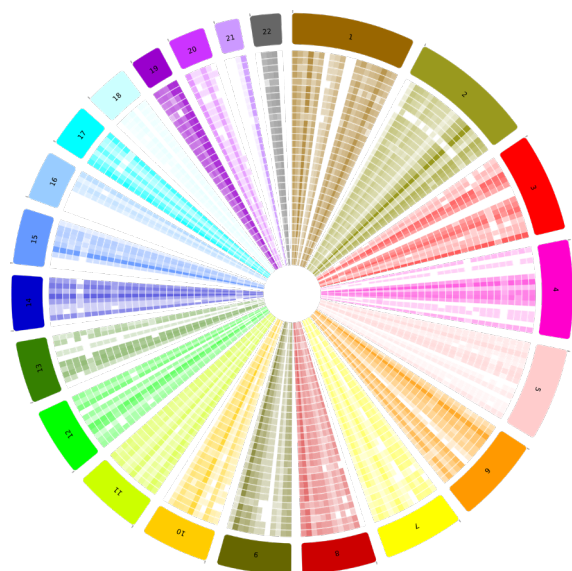
**ROBUSTLY** construct a library using PTA product as input with a choice of library construction protocols. Only 100 ng of unfragmented PTA product is required as input with the ResolveDNA™ Library Preparation Kit from BioSkryb. It uses a ligation-based workflow that does not require fragmentation of the input DNA and utilizes unique dual-index adapters that are compatible with Illumina sequencers.

Unfragmented product can also be used in the KAPA HyperPlus construction protocol, though at a higher 500 ng input. Alternatively, directly fragmented PTA products can be used with Illumina DNA Prep reagents. All workflows yield >400 ng of amplified library, facilitating downstream enrichments if necessary.

## ▼ Sequence and Analyze with BaseJumper

**QUICKLY** perform low-depth sequencing on any Illumina instrument to obtain QC sequencing metrics prior to performing alignment (Table 3).

**SIMPLY** analyze high-depth sequencing, align and call variants, and visualize variant call file data with BioSkryb's BaseJumper Bioinformatics platform (Figure 4). Single nucleotide variation can be explored in the context of a density map—to explore nucleotide differences between single cells at tunable levels of resolution.



**Figure 4. Variant density in BaseJumper:** Quickly visually discern genome-wide differences in variant density between single cells (concentric rings) or zoom in for high-resolution regional ascertainment of variant density.

## ▼ Verification with HEK293T cells

This workflow was verified by sorting live HEK293T cells into 96-well plates with each dispense mode, amplifying with PTA, and low-depth sequencing all wells with >500 ng PTA yield.

All dispense modes yielded even better amplification efficiency than expected from bead testing (Table 2). This is likely due to an improvement in analysis efficiency caused by a difference in particle chemistry - cells may centrifuge to the well bottoms more easily than their sticky bead counterparts. Doublet wells may also contribute to the higher values, since Table 1 bead data only includes singlets, but they do not account for the entire difference; 75-86% of wells contained any number of beads. Therefore amplification yield may often exceed the established bead range.

Table 2. Amplification Efficiency

Dispense Mode	% Wells with PTA Yield	
	8.5µL Droplet	2.2µL Droplet
Direct	100	81
Sidewall	94	63

Viable (SytoxGreen-) HEK293T cells were sorted into 96-well plates with BioSkryb Cell Buffer. 2 total plates were dispensed for each mode. For each plate, PTA was performed on 16 wells and any well with yield > 500 ng was counted.

All single cells yielded a preseq<sup>3</sup> library complexity estimate of greater than 3.0E9 bp, which indicates the likelihood of high sensitivity of variant calling upon high depth sequencing. Additional metrics indicate hg38 alignment of > 97% and low chimeric and duplication rate. The data presented here are using KAPA HyperPlus library construction.

Table 3. QC Analysis of NanoCollect-BioSkryb Workflow

QC Metric	Mean ± Standard Deviation			
	8.5µL Droplet		2.2µL Droplet	
Preseq Count	3.47 E09	± 4.44 E07	3.52 E09	± 1.22 E08
Alignment	0.978	± 0.001	0.979	± 0.002
Chimeras	0.046	± 0.002	0.042	± 0.004
Duplicates	0.004	± 0.001	0.003	± 0.001

## Conclusion

Easily amplify single-cell genomes with high breadth, uniform coverage, reduced error propagation, and low allelic dropout by combining NanoCelect's WOLF and BioSkryb's ResolveDNA workflow. Accurately call single nucleotide variants (SNV) and copy number variations (CNV) – at the whole-genome, exome, or small panel level – for diverse applications such as cancer genomics, prenatal genetic testing (PGT), and neurobiology.

### Acknowledgements

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For more information, visit [nanocelllect.com](https://nanocelllect.com)  
or email [info@nanocelllect.com](mailto:info@nanocelllect.com)

## References

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