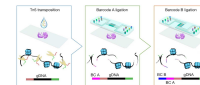


Jan 14, 2025

Yale University Co_Profile Spatial-ATAC and RNA Sequencing via Deterministic Barcoding in Fresh Frozen Human Lymph Node Tissue



Forked from [Yale University_Spatial ATAC Sequencing for Fixed Fresh Frozen Human Lymph node Tissue via DBiT-seq](#)

DOI

<https://dx.doi.org/10.17504/protocols.io.3byl4wrezvo5/v1>

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Cellular Senescence Net...



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DOI: <https://dx.doi.org/10.17504/protocols.io.3byl4wrezvo5/v1>

External link: <https://www.nature.com/articles/s41586-022-05094>

Protocol Citation: Negin Farzad, Archibald Enniful, Di Zhang, Rong Fan 2025. Yale University Co_Profile Spatial-ATAC and RNA Sequencing via Deterministic Barcoding in Fresh Frozen Human Lymph Node Tissue. **protocols.io**
<https://dx.doi.org/10.17504/protocols.io.3byl4wrezvo5/v1>

Manuscript citation:

Deng, Y., Bartosovic, M., Ma, S. *et al.* Spatial profiling of chromatin accessibility in mouse and human tissues. *Nature* **609**, 375–383 (2022). <https://doi.org/10.1038/s41586-022-05094-1>

Farzad N, Enniful A, Bao S, Zhang D, Deng Y, Fan R. Spatially resolved epigenome sequencing via Tn5 transposition and deterministic DNA barcoding in tissue. *Nat Protoc.* 2024 Jun 28. doi: 10.1038/s41596-024-01013-y. Epub ahead of print. PMID: 38943021.

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Protocol status: Working

We use this protocol and it's working

Created: January 10, 2025

Last Modified: January 14, 2025

Protocol Integer ID: 118089

Keywords: epigenome profiling of cellular senescence, fixed fresh frozen human lymph node tissue sample, fresh frozen human lymph node tissue sample, transcriptome atlas, epigenome profiling, fresh frozen human lymph node tissue, cellular senescence, cell resolution, tissue for spatial atac, rna, single cell

Funders Acknowledgements:

Yale TMC for Cellular Senescence in Lymphoid Organs

Grant ID: 1U54AG076043-01

Abstract

This protocol describes the use of Deterministic Barcoding in Tissue for spatial ATAC and RNA sequencing to construct epigenome and transcriptome atlases of fixed fresh frozen human lymph node tissue samples. This approach uses a microfluidic-based method to introduce combinatorial DNA oligo barcodes directly to the cells in a tissue section fixed on a glass slide using DBiT-seq device. This method spatially resolved transcriptome and epigenome profiling of cellular senescence in fresh frozen human lymph node tissue samples. This technique does not directly resolve single cells but can achieve a near-single-cell resolution.

Materials

KEY RESOURCES TABLE

	A	B	C
	Name	Catalog number	Vender
	Formaldehyde solution	PI28906	Thermo Fisher Scientific
	Glycine	50046	Sigma-Aldrich
	NaCl	AM9760G	Thermo Fisher Scientific
	Digitonin	G9441	Promega
	MgCl ₂	AM9530G	Thermo Fisher Scientific
	NP40	11332473001	Sigma-Aldrich
	EDTA Solution pH 8.0	AB00502	AmericanBio
	Bovine Serum Albumin (BSA)	A8806	Sigma-Aldrich
	Triton X-100	T8787	Sigma-Aldrich
	T4 DNA Ligase	M0202L	New England Biolabs
	T4 DNA Ligase Reaction Buffer	B0202S	New England Biolabs
	NEBuffer 3.1	B7203S	New England Biolabs
	DPBS	14190144	Thermo Fisher Scientific



	A	B	C
	Proteinase K	EO0491	Thermo Fisher Scientific
	Ampure XP beads	A63880	Beckman Coulter
	NEBNext High-Fidelity 2X PCR Master Mix	M0541L	New England Biolabs
	SYBR Green I Nucleic Acid Gel Stain	S7563	Thermo Fisher Scientific
	DNA Clean & Concentrator-5	D4014	Zymo Research
	Tn5 Transposase - unloaded	C01070010	Diagenode
	Tagmentation Buffer (2x)	C01019043	Diagenode
	Sodium dodecyl sulfate	71736	Sigma-Aldrich
	Maxima H Minus Reverse Transcriptase (200 U/L)	EP0751	Thermo Fisher Scientific
	dNTP mix	R0192	Thermo Fisher Scientific
	SUPERased In RNase Inhibitor	AM2694	Thermo Fisher Scientific
	Ampure XP beads	A63880	Beckman Coulter
	Dynabeads MyOne C1	65001	Thermo Fisher Scientific
	RNase Inhibitor	Y9240L	Enzymatics
	Kapa Hotstart HiFi ReadyMix	KK2601	Kapa Biosystems
	Nextera XT DNA Preparation Kit	FC-131-1024	Illumina

Table S1A. **List of reagents**

Oligo Name	Sequence
Ligation Linker	CGAATGCTCTGGCCTCTCAAGCACGTGGAT
P5 oligo	AATGATACGGCGACCAACGAGATCTACACTAGATCGCTCGTCGGCAGCGTCAGATGTGTATAAGAGACAG
P7 oligo (701)	CAAGCAGAAGACGGCATAACGAGATTTCGCCTTAGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT
P7 oligo (702)	CAAGCAGAAGACGGCATAACGAGATCTAGTACGGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT
P7 oligo (703)	CAAGCAGAAGACGGCATAACGAGATTTCTGCCTGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT
P7 oligo (704)	CAAGCAGAAGACGGCATAACGAGATGCTCAGGAGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT
P7 oligo (705)	CAAGCAGAAGACGGCATAACGAGATAGGAGTCCGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT
P7 oligo (706)	CAAGCAGAAGACGGCATAACGAGATCATGCCTAGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGCAAGCGTTGGCTTCTCGCATCT

Table S2A. **List of PCR Oligo.**

A	B
Barcode A-1	/5Phos/AGGCCAGAGCATTCTGAACGTGATTTTTTTTTTTTTTTTV N
Barcode A-2	/5Phos/AGGCCAGAGCATTCTGAAACATCGTTTTTTTTTTTTTTTV N
Barcode A-3	/5Phos/AGGCCAGAGCATTCTGATGCCTAATTTTTTTTTTTTTTTTV N
Barcode A-4	/5Phos/AGGCCAGAGCATTCTGAGTGGTCATTTTTTTTTTTTTTTTV N
Barcode A-5	/5Phos/AGGCCAGAGCATTCTGACCACTGTTTTTTTTTTTTTTTV N
Barcode A-6	/5Phos/AGGCCAGAGCATTCTGACATTGGCTTTTTTTTTTTTTTTTV N
Barcode A-7	/5Phos/AGGCCAGAGCATTCTGCAGATCTGTTTTTTTTTTTTTTTV N
Barcode A-8	/5Phos/AGGCCAGAGCATTCTGCATCAAGTTTTTTTTTTTTTTTV N



A	B
Barcode A-9	/5Phos/AGGCCAGAGCATTGCGCTGATCTTTTTTTTTTTTTTV N
Barcode A-10	/5Phos/AGGCCAGAGCATTGACAAGCTATTTTTTTTTTTTTTV N
Barcode A-11	/5Phos/AGGCCAGAGCATTGCTGTAGCCTTTTTTTTTTTTTTV N
Barcode A-12	/5Phos/AGGCCAGAGCATTGAGTACAAGTTTTTTTTTTTTTV N
Barcode A-13	/5Phos/AGGCCAGAGCATTGGAACAACCATTTTTTTTTTTTTTV N
Barcode A-14	/5Phos/AGGCCAGAGCATTGGAACCGAGATTTTTTTTTTTTTTV N
Barcode A-15	/5Phos/AGGCCAGAGCATTGGAACGCTATTTTTTTTTTTTTTV N
Barcode A-16	/5Phos/AGGCCAGAGCATTGGAAGACGGATTTTTTTTTTTTTTV N
Barcode A-17	/5Phos/AGGCCAGAGCATTGGAAGGTACATTTTTTTTTTTTTTV N
Barcode A-18	/5Phos/AGGCCAGAGCATTGACACAGAATTTTTTTTTTTTTTV N
Barcode A-19	/5Phos/AGGCCAGAGCATTGACAGCAGATTTTTTTTTTTTTTV N
Barcode A-20	/5Phos/AGGCCAGAGCATTGACCTCCAATTTTTTTTTTTTTTV N
Barcode A-21	/5Phos/AGGCCAGAGCATTGACGCTCGATTTTTTTTTTTTTTV N
Barcode A-22	/5Phos/AGGCCAGAGCATTGACGTATCATTTTTTTTTTTTTTV N
Barcode A-23	/5Phos/AGGCCAGAGCATTGACTATGCATTTTTTTTTTTTTTV N
Barcode A-24	/5Phos/AGGCCAGAGCATTGAGAGTCAATTTTTTTTTTTTTTV N
Barcode A-25	/5Phos/AGGCCAGAGCATTGAGATCGCATTTTTTTTTTTTTTV N
Barcode A-26	/5Phos/AGGCCAGAGCATTGAGCAGGAATTTTTTTTTTTTTTV N
Barcode A-27	/5Phos/AGGCCAGAGCATTGAGTCACTATTTTTTTTTTTTTTV N



A	B
Barcode A-28	/5Phos/AGGCCAGAGCATTGATCCTGTATTTTTTTTTTTTTTV N
Barcode A-29	/5Phos/AGGCCAGAGCATTGATTGAGGATTTTTTTTTTTTTTV N
Barcode A-30	/5Phos/AGGCCAGAGCATTGCAACCACATTTTTTTTTTTTTTV N
Barcode A-31	/5Phos/AGGCCAGAGCATTGGACTAGTATTTTTTTTTTTTTTV N
Barcode A-32	/5Phos/AGGCCAGAGCATTGCAATGGAATTTTTTTTTTTTTTV N
Barcode A-33	/5Phos/AGGCCAGAGCATTGCACTTCGATTTTTTTTTTTTTTV N
Barcode A-34	/5Phos/AGGCCAGAGCATTGCGAGCGTTATTTTTTTTTTTTTTV N
Barcode A-35	/5Phos/AGGCCAGAGCATTGCGATACCAATTTTTTTTTTTTTTV N
Barcode A-36	/5Phos/AGGCCAGAGCATTGCGCAGTTCATTTTTTTTTTTTTTV N
Barcode A-37	/5Phos/AGGCCAGAGCATTGCGCGAAGTATTTTTTTTTTTTTTV N
Barcode A-38	/5Phos/AGGCCAGAGCATTGCGCGTGAGATTTTTTTTTTTTTTV N
Barcode A-39	/5Phos/AGGCCAGAGCATTGCGCTCCTGATTTTTTTTTTTTTTV N
Barcode A-40	/5Phos/AGGCCAGAGCATTGCGGAACTTATTTTTTTTTTTTTTV N
Barcode A-41	/5Phos/AGGCCAGAGCATTGCGGACTGGATTTTTTTTTTTTTTV N
Barcode A-42	/5Phos/AGGCCAGAGCATTGCGGCATACATTTTTTTTTTTTTTV N
Barcode A-43	/5Phos/AGGCCAGAGCATTGCTCAATGATTTTTTTTTTTTTTV N
Barcode A-44	/5Phos/AGGCCAGAGCATTGCTGAGCCATTTTTTTTTTTTTTV N
Barcode A-45	/5Phos/AGGCCAGAGCATTGCTGGCATATTTTTTTTTTTTTTV N
Barcode A-46	/5Phos/AGGCCAGAGCATTGGAATCTGATTTTTTTTTTTTTTV N

A	B
Barcode A-47	/5Phos/AGGCCAGAGCATTCTGCAAGACTATTTTTTTTTTTTTTTTV N
Barcode A-48	/5Phos/AGGCCAGAGCATTCTGGAGCTGAATTTTTTTTTTTTTTTTV N
Barcode A-49	/5Phos/AGGCCAGAGCATTCTGGATAGACATTTTTTTTTTTTTTTTV N
Barcode A-50	/5Phos/AGGCCAGAGCATTCTGGCCACATTTTTTTTTTTTTTTTV N

Table S2B. List of DNA Barcode A Sequences.

A	B
Barcode B-1	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAACGTGATATCCAC GTGCTTGAG
Barcode B-2	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAACATCGATCCAC GTGCTTGAG
Barcode B-3	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNATGCCTAAATCCAC GTGCTTGAG
Barcode B-4	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGTGGTCAATCCAC GTGCTTGAG
Barcode B-5	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACCACTGTATCCAC GTGCTTGAG
Barcode B-6	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACATTGGCATCCAC GTGCTTGAG
Barcode B-7	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNCAGATCTGATCCAC GTGCTTGAG
Barcode B-8	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNCATCAAGTATCCAC GTGCTTGAG
Barcode B-9	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNCGCTGATCATCCAC GTGCTTGAG
Barcode B-10	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACAAGCTAATCCAC GTGCTTGAG
Barcode B-11	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNCTGTAGCCATCCAC GTGCTTGAG
Barcode B-12	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGTACAAGATCCAC GTGCTTGAG



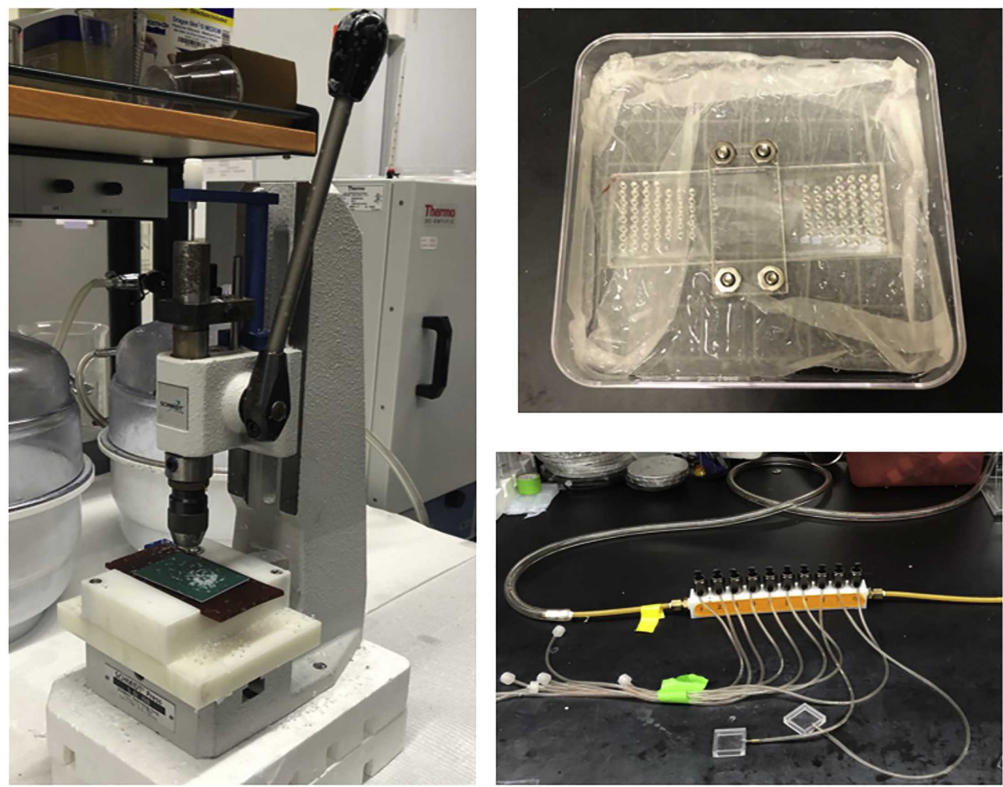
A	B
Barcode B-13	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAACAACCAATCCAC GTGCTTGAG
Barcode B-14	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAACCGAGAATCCAC GTGCTTGAG
Barcode B-15	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAACGCTTAATCCAC GTGCTTGAG
Barcode B-16	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAAGACGGAATCCAC GTGCTTGAG
Barcode B-17	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAAGGTACAATCCAC GTGCTTGAG
Barcode B-18	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACACAGAAATCCAC GTGCTTGAG
Barcode B-19	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACAGCAGAATCCAC GTGCTTGAG
Barcode B-20	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACCTCCAAATCCAC GTGCTTGAG
Barcode B-21	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACGCTCGAATCCAC GTGCTTGAG
Barcode B-22	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACGTATCAATCCAC GTGCTTGAG
Barcode B-23	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNACTATGCAATCCAC GTGCTTGAG
Barcode B-24	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGAGTCAAATCCAC GTGCTTGAG
Barcode B-25	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGATCGCAATCCAC GTGCTTGAG
Barcode B-26	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGCAGGAAATCCAC GTGCTTGAG
Barcode B-27	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNAGTCACTAATCCAC GTGCTTGAG
Barcode B-28	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNATCCTGTAATCCAC GTGCTTGAG
Barcode B-29	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNATTGAGGAATCCAC GTGCTTGAG
Barcode B-30	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNCAACCACAATCCAC GTGCTTGAG
Barcode B-31	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNNGACTAGTAATCCAC GTGCTTGAG



A	B
Barcode B-32	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCAATGGAAATCCAC GTGCTTGAG
Barcode B-33	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCACTTCGAATCCAC GTGCTTGAG
Barcode B-34	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCAGCGTTAATCCAC GTGCTTGAG
Barcode B-35	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCATAACCAATCCAC GTGCTTGAG
Barcode B-36	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCCAGTTCAATCCAC GTGCTTGAG
Barcode B-37	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCCGAAGTAATCCAC GTGCTTGAG
Barcode B-38	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCCGTGAGAATCCAC GTGCTTGAG
Barcode B-39	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCCTCCTGAATCCAC GTGCTTGAG
Barcode B-40	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCGAACTTAATCCAC GTGCTTGAG
Barcode B-41	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCGACTGGAATCCAC GTGCTTGAG
Barcode B-42	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCGCATACAATCCAC GTGCTTGAG
Barcode B-43	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCTCAATGAATCCAC GTGCTTGAG
Barcode B-44	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCTGAGCCAATCCAC GTGCTTGAG
Barcode B-45	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCTGGCATAATCCAC GTGCTTGAG
Barcode B-46	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNGAATCTGAATCCAC GTGCTTGAG
Barcode B-47	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNCAAGACTAATCCAC GTGCTTGAG
Barcode B-48	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNGAGCTGAAATCCAC GTGCTTGAG
Barcode B-49	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNGATAGACAATCCAC GTGCTTGAG
Barcode B-50	/5Biosg/CAAGCGTTGGCTTCTCGCATCTNNNNNNNNNNGCCACATAATCCAC GTGCTTGAG

Table S2C. List of DNA Barcode B Sequences.

- This protocol requires using a microfluidic device fabricated with PDMS using soft lithography.
- A hole punching machine (SCHMIDT® Manual Press) is needed to punch the inlet and outlet holes in the PDMS chip after its fabrication.
- The tissue of interest should be placed on the center of a poly-L-lysine coated glass slide. (CatLog no. 63478-AS, electron microscopy sciences).
- A custom-designed acrylic clamp with screws is needed to hold the PDMS device and the glass slide together firmly.
- The silicon wafer used for fabricating the PDMS mold is purchased from WaferPro (CatLog. No. C04004).
- The photoresist is purchased from MicroChem Laboratory (CatLog. No. SU-8 2010).
- A homemade laboratory vacuum system(See Figure Below) is needed to applying suction to move fluid in the microfluidic channels.
- Microscopy – The tissue of interest can be scanned and imaged using EVOS (Thermo Fisher EVOS fl), typically at a magnification of 10×. Any suitable optical microscope can be used.
- A Laser Engraver Cutter is used to cut out the acrylic barcode and lysis clamps.
- A “humidified chamber” (See Figure Below) to prevent reagent evaporation during incubation.



Key equipment used in DBiT-seq

Hole punch machine, the key device for DBiT-seq, and the house vacuum lines. (Left) SCHMIDT® Manual Press used for hole punching 2-mm diameter holes for the inlets and outlets of Chip-A and Chip-B. (Top Right) Example of a “humidified chamber” to prevent reagent evaporation during the incubation steps. (Bottom Right) House vacuum used for pulling reagents through the microchannels.

Recipes

Critical: Prepare the RT mix, template switch mix, ligation mix, PCR mix, 1X lysis solution, flow wash buffer, PBS-RI, 80% ethanol on the same day as usage. 2X lysis buffer, 1X B&W Buffer with 0.05% Tween-20, and 2X B&W Buffer can be stored for up to 6 months at room temperature.

PBS-RI

	Reagent	Volume (μL)
	1X PBS	5000
	RNase Inhibitor (40 U/μL) (Enzymatics)	7
	Total	5007

80% Ethanol



	Reagent	Volume(ml)
	RNase-free water	1
	100% Ethanol	4
	Total	5

RT Mixture

	Reagent	Volume (μL)
	5X Maxima RT buffer	50
	RNase-free water	32.8
	RNase Inhibitor (Enzymatics)	1.6
	Suprase In RNase Inhibitor (Ambion)	3.2
	dNTPs (10 mM stock)	12.5
	Maxima H Minus Reverse Transcriptase	25
	PBS-RI	100
	Total	225.1

Ligation Mix

	Reagent	Volume (μL)
	10X T4 Ligase Buffer	27
	1X NEB buffer 3.1 with 1% RI (Enzymatics)	115.8
	5% Triton-X100	5.4
	RNase Inhibitor (Enzymatics)	2.2
	RNase-free water	69.5
	SupraseIn RNase Inhibitor (Ambion)	0.7
	T4 DNA Ligase (400 U/μL)	11
	Total	231.6

Template switch mix

	Reagent	Volume (μL)
	20% Ficoll PM-400	44
	5X Maxima RT buffer	44



Reagent	Volume (μ L)
dNTPs (10 mM stock)	22
RNase Inhibitor (Enzymatics)	5.5
Maxima H Minus Reverse Transcriptase	11
Template Switch Primer (100 μ M stock)	5.5
RNase-free water	88
Total	220

PCR mix

Reagent	Volume (μ L)
2X Kapa Hifi HotStart master mix	110
Primer1 BC_0062 (10 μ M)	8.8
Primer2 BC_0108 (10 μ M)	8.8
RNase-free water	92.4
Total	220

Flow Wash Buffer

Reagent	Volume (mL)
1X PBS	4
10 % Triton X-100	0.04
Suprase In RNase Inhibitor	0.01
Total volume	4.05

1X Lysis Solution

Reagent	Volume (μ L)
1X PBS	50
2X Lysis Buffer	50
Proteinase K (20mg/mL) (Thermo)	10
Total	110

2X Lysis Buffer

	Reagent	Stock Concentration	Final Concentration (2X)	Volume (mL)
	Tris, pH 8.0	1 M	20 mM	0.5
	NaCl	5 M	400 mM	2
	EDTA, pH 8.0	0.5 M	100 mM	5
	SDS	10%	4.4 %	11
	RNase-free Water	NA	NA	6.5
	Final Volume			25

1X B&W Buffer with 0.05% Tween-20

	Reagent	Volume
	1M Tris-HCl pH 8.0	100 µL
	EDTA, 0.5M	20 µL
	5M NaCl	4 mL
	Tween 20 10%	100 µL
	RNase-free water	15.78 mL
	Total	20 mL

2X B&W Buffer

	Reagent	Volume
	1M Tris-HCl pH 8.0	500 µL
	EDTA, 0.5M	100 µL
	5M NaCl	20 mL
	RNase-free water	29.4 mL
	Total	50 mL

CRITICAL: Handle all in an RNase-free area.



Before start

The protocol below describes the reagents, equipment, and specific experimental steps for using the spatial-ATAC-seq and RNA-seq platform on fixed fresh frozen tissue slides. The 20 μm microfluidic channel width were designed and validated to provide spatial epigenomic and transcriptome profiles on human lymph node tissue sections.

CRITICAL: Work in an RNase-free environment when the microfluidic device is not on the tissue slide. Use RNaseZapTM or other commercially available cleaner solution and filter-tips. Clean surfaces and gloves with RNaseZapTM.

CRITICAL: Keep reagents on ice at all times.

Fabricating the Silicon Wafer Device Mold

- 1 Prepare chrome photomasks of the microfluidics chip by printing the high-resolution CAD files onto a glass substrate (Figure 1).

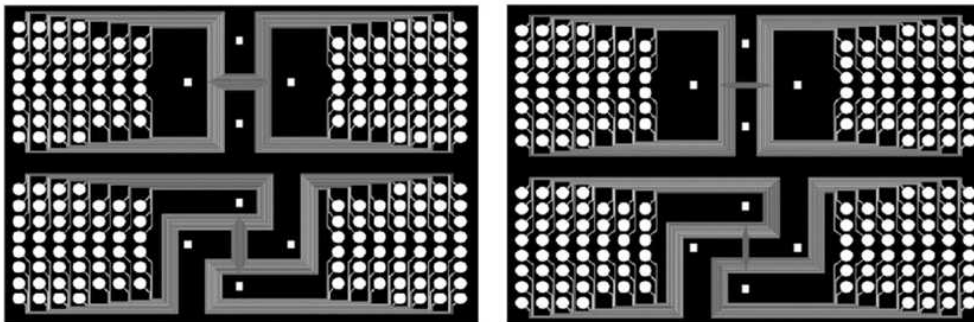


Figure 1: Microfluidic Photomask Design.

(Left) AutoCAD design for the 20 micron-width fifty channel Chip-A and Chip-B. (Right) AutoCAD design for 10 micron-width channel Chip-A and Chip-B.

NOTE: We have outsourced this step (Front Range Photomask, USA).

Using these masks, prepare a replica mold as follows:

- 2 Start the process by cleaning a 4-inch silicon wafer with 100% acetone (Aldrich) and then 100% isopropanol (Aldrich), then dry with compressed air.

NOTE: Acetone and isopropanol are mildly toxic. Use proper PPE when handling and discard waste in the appropriate containers.
- 3 Prepare a high-resolution computer-aided-design (CAD) file with the desired microfluidics chip design. A CAD file is also available in the link provided. (<https://ars.els-cdn.com/content/image/1-s2.0-S2666166721002392-mmc4.zip>)
- 4 Bake the wafer at 180°C for 10 min on a hot plate to dry it out.
- 5 Use a spin coater to evenly spread SU-8 2010 photoresist (MicroChem) for 10 μm device or SU-8 2025 (MicroChem) for 25 μm and 50 μm device onto the wafer at 500 rpm for 5 seconds followed by 1100 rpm (10 μm device), 3500 rpm (25 μm device) or 1750 rpm (50 μm device) for 40 seconds.
- 6 Soft bake the wafer for 3 min at 65°C (25 μm and 50 μm device) and 4 min (10 μm device) or 6 min (25 μm and 50 μm device) at 95°C.

- 7 Expose the SU-8 on the wafer through the photomask using Mask Aligner with a dose of 150 mJ/cm² UV.
- 8 For the post exposure bake, bake the wafer for 1 min at 65°C (25 µm and 50 µm device) and 5 min (10 µm and 25 µm device) or 6 min (50 µm device) at 95°C.
- 9 Develop the SU-8 for 4 min (10 µm and 25 µm device) or 5 min (50 µm device) in a bath of ~50 mL SU-8 developer (MicroChem).
- 10 Rinse the wafer with 100% isopropanol and dry with compressed air.
- 11 Perform a hard bake by baking the wafer for 10 min at 180°C.

CRITICAL: This process needs to be carried out in a microelectronics cleanroom.

Pause Point: The replica mold can now be stored and reused indefinitely.

Creating the Acrylic Clamps

30m

- 12 Prepare the pattern and dimensions for the barcoding clamp and lysis clamp.
- 13 Peel covering of acrylic sheet and place it in the laser cutting machine.
- 14 Select the program dimensions and cut two pieces for the top and bottom of the barcoding clamp.
- 15 Place the top blank piece into the laser cutter and select the pattern to cut out the four holes in the corner for the screws and nuts.
- 16 Remove cut-out scraps with a pipet tip or a similar pointed tool.
- 17 Repeat steps 15 and 16 for the bottom blank piece.
- 18 Repeat steps 13 to 17 for the lysis clamp using the correct dimensions and pattern with an additional hole in the center of the top piece (Example shown in Figure 2).

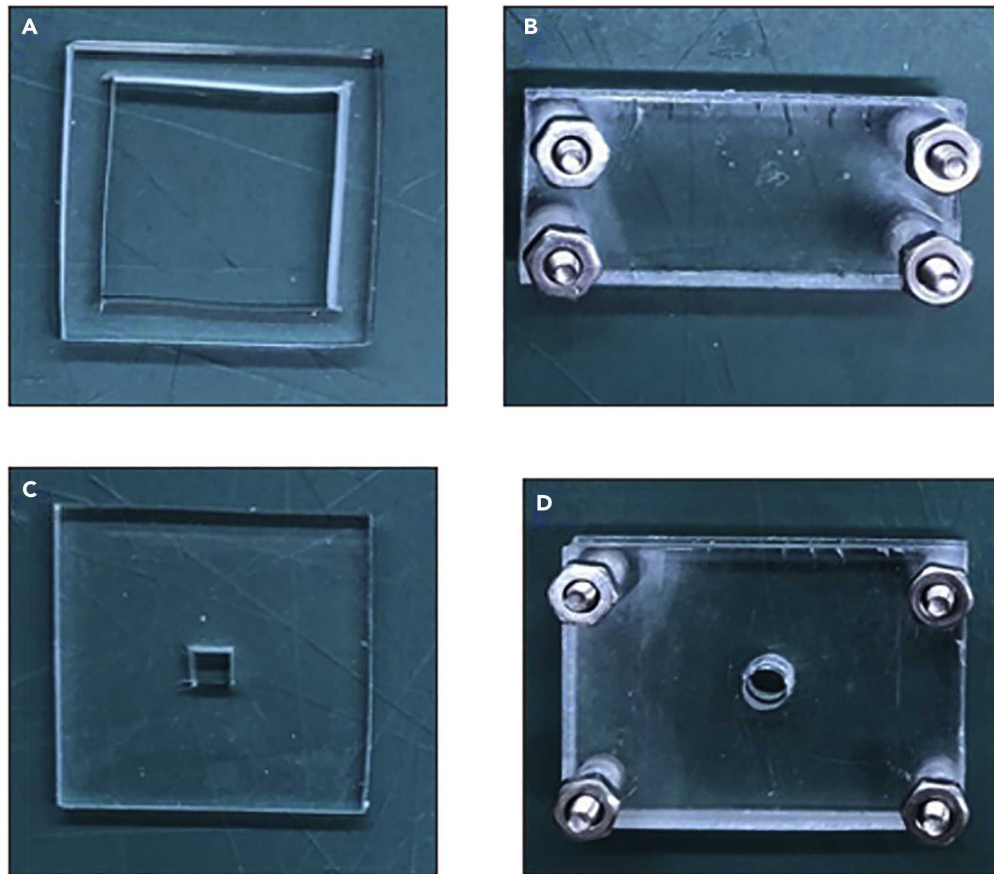


Figure 2. Key parts for setup. PDMS reservoirs and acrylic clamps used in DBiT-seq. (A) Inlet Reservoir. (B) Barcoding Clamp. (C) Lysis Reservoir. (D) Lysis Clamp.

NOTE: The acrylic clamps can be reused indefinitely.

Preparing the Microfluidic Device and reagent reservoirs

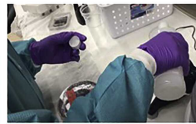
3h

- 19 Thoroughly mix polydimethylsiloxane (PDMS) elastomer base and curing agent (these come together) at a 10:1 ratio (See Figure 3).

Chronological steps for preparing the PDMS chip from left to right. The liquid PDMS mix is poured into the replica mold, degassed and baked. The solidified PDMS is then cut out and assembled into the microfluidics chip by punching inlet and outlet holes and attaching to a glass slide. Detailed steps are available in the text.



Materials needed to make the PDMS device



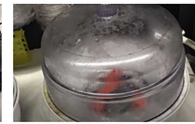
Weigh out PDMS base and curing agent in a 1:10 mass ratio



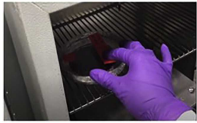
Mix PDMS base and curing agent thoroughly



Transfer PDMS mixture to the mold



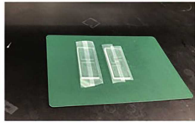
Degas PDMS mixture with vacuum desiccator



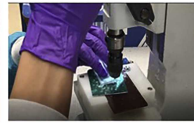
Bake at 65-70 ° C for 2 hours



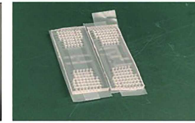
Cut out PDMS using scalpel and gently peel from the mold



Cover the PDMS device with scotch tape to protect from dust and debris



Punch inlet and outlet holes using SCHMIDT press hole puncher



Cover the final PDMS device with scotch tape

Figure 3: Step-by-step visual guide for making Chip-A and Chip-B.

20 Pour the mixture into the silicon device mold.

NOTE: Aim for a chip height of about 5mm.

21 Place in a vacuum desiccator until all bubbles dissipate from the mixture about 30-60 min.

22 Cure in an oven at 65-70°C for a minimum of 2 hours and up to overnight.

NOTE: Make sure to place on an even surface to prevent uneven curing!

23 Cut out the cured device and hole punch each of the 50 inlets and outlets for the channels.

NOTE: Cut both Chip-A and Chip-B in the approximate size of a glass slide.

NOTE: Make sure there are no PDMS pieces left in the inlets or outlets.

24 Thoroughly clean the surface of the device with scotch tape.

25 Pour mixture in a container large enough to cut out two roughly 25×25×5mm PDMS pieces.

26 For the barcoding reservoir, cut out a 20×20mm piece in the center, barely large enough to surround the inlets of the chips when placed above.



- 27 For the lysis reservoir, depending on the device size used, cut out the center to barely surround the barcoded region of the tissue (1×1mm for the 10µm channel device, 2.5×2.5mm for the 25µm channel device).

NOTE: The reservoirs are reusable. Thoroughly wash with 70% ethanol after each use.

CRITICAL: Be careful when cutting out the Chips A and B. Make sure not to cut across any patterned areas of the silicon wafer mold, otherwise it will need to be replaced and re-fabricated.

Preparing Barcodes (A and B).

1h

- 28 Thoroughly mix the ligation linker 1 with each barcode A₁₋₅₀ at and ligation linker 2 with each barcode B₁₋₅₀ at a 1:1 ratio.
- 29 Place the 50 mixes in a thermal cycle and heat to 97°C to anneal.
- 30 Slowly cool to room temperature at a rate of -0.1°C/sec.
- Store at -20°C for up to 6 months.

Preparing the Tissue Slide

1d

- 31 25% sucrose overnight bath.
- 32 Embed in OCT frozen on dry ice.
- 33 Section at ~7 µm thickness for the 25µm channel device and ~5 µm thickness for the 10 µm channel device onto Poly-L-lysine slides.
- 34 Store slides at -80°C for up to 6 months until use.

CRITICAL: Ensure that the tissue is sectioned as evenly as possible, otherwise the risk of cross-flow between the microfluidic channels will be increased. Tissue section should be placed at the center of a glass slide to the greatest extent.

Transposome assembly preparation



35

Prepare the transposome assembly annealing buffer (The Tn5 annealing buffer contains 400 μl of 1 M Tris-HCl pH 8.0, 100 μl of NaCl 5 M and 9.5 ml water. This annealing buffer is used to resuspend each oligos of Tn5-Rev, Tn5-A and Tn5-B in this annealing buffer to stock concentrations of 100 μM .

36 Resuspend the oligos (Tn5Erev, Tn5ME-A , and Tn5ME-B) in Annealing Buffer to stock concentration of 100 μM .

37 In a PCR tube, mix 10 μl of oligo Rev with 10 μl of oligo A.

38 In a separate PCR tube, mix 10 μl of oligo Rev with 10 μl of oligo B

39 Vortex the tubes and place them in a thermocycler, and run the thermocycler with the following program:

	Temperatu re	Time
	95 °C	5 min
	Cool to 65 °C	-0.1C/ secon d
	65 °C	5 min
	Cool to 4 °C	-0.1 C/sec ond

The Annealed linker oligos can be stored at -20 °C.

- 40 Mix 5 µl of each prepared oligo A/oligo Rev and oligo B/oligo Rev at the ratio of 1:1 in a pcr tube.
- 41 Add 10 µl of unloaded Tn5 transposome (Diagnode Tagmentase) to the mixture of oligo A/oligo Rev and oligo B/oligo Rev.
After vortexing the mixture briefly, incubate the tube at 23 °C for 30 minutes in a thermocycler. The product can be used the same day or stored in -20 °C with 10 µl glycerol up to 60 days.

Tissue Preparation

1h

- 42 Remove stored sections from the freezer and allow to warm to room temperature for 10 minutes.
- 43 Clean sections by pipetting 2 mL of PBS-R1 across the tissue.
- 44 If sections are not yet fixed, then fix here with 1 mL 0.2% PFA, without methanol in 1X PBS,
by applying on top of the tissue and incubate for 5 min at room temperature.
- 45 Remove the formaldehyde by pipetting it off and quench the tissue with 1 mL 1.25M glycine for 5 min at room temperature.
- 46 Wash the slides with 1 mL of PBS for 1 minute and then dip the slide in a 50 mL falcon tube with DI water.
- 47 Dry the section with gentle air flow.
- 48 Take a full pre-scan image of the section at the desired optical resolution (Figure 4A).

Recommended 10X resolution.

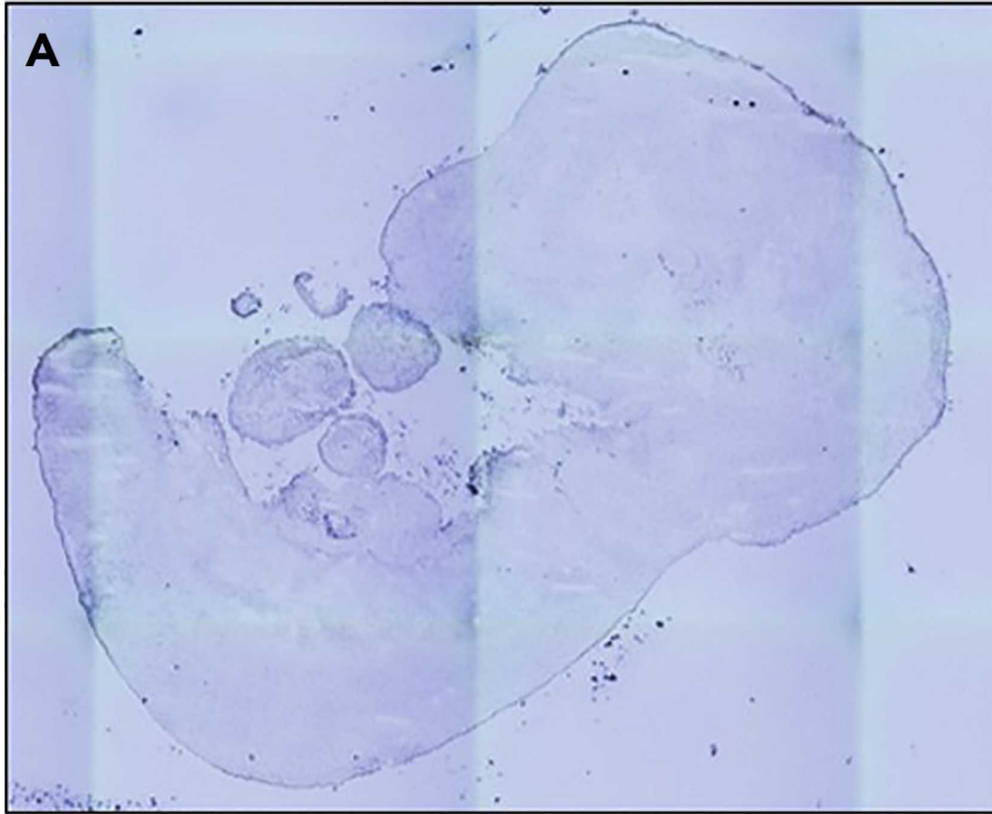


Figure 4 : Example scanning of the tissue slide.

(A) Initial full scan of the tissue section prior to DBiT-seq.

49 Attach the Antibody PDMS reservoir and 12mm diameter clamp to the tissue.

50 Permeabilize the tissue with 500uL 0.1 Xlysis buffer and incubate at room temperature for 15 min

Make sure to prepare 0.1x lysis buffer fresh each time. using the following reagents:

1X Lysis Buffer (10 mM Tris-HCl, pH 8.0; 10 mM NaCl; 3

mM MgCl₂; 0.1% Tween-20; 0.1% NP-40; 0.01% Digitonin; 1% BSA)

Dilute down to 0.1X with lysis dilution buffer

(10 mM Tris-HCl, pH 8.0; 10 mM NaCl; 3 mM MgCl₂; 1% BSA)



- 51 Wash the tissue with 500uL wash buffer for 5 min
make sure to prepare wash buffer fresh each time:

Wash Buffer (10 mM Tris-HCl pH 8.0; 10 mM NaCl; 3 mM MgCl₂; 1% BSA; 0.1% Tween-20)

- 52 Prepare the transposition mix with the loaded Tn5 transposon and add 100uL to the tissue.

make sure to prepare the Tn5 mix fresh each time

	A	B	C	D
	Volume (200 ul/chip)	Volume (300 ul)	Volume (400 ul)	Reagent (50 ul/reaction)
	100 ul	150 ul	200 ul	2X TD Buffer (25 ul)
	66 ul	99 ul	132 ul	1X PBS (16.5 ul)
	2 ul	3 ul	4 ul	10% Tween-20 (final 0.1% v/v) (0.5 ul)
	2 ul	3 ul	4 ul	1% Digitonin (final 0.01% v/v) (0.5 ul)
	10 ul	15 ul	20 ul	Transposome (2.5 ul)
	20 ul	30 ul	40 ul	Nuclease-free H ₂ O (5 ul)

- 53 Incubate the tissue at 37 °C for 30 min
- 54 Stop the tagmentation reaction by adding 500 µL of 40 mM EDTA onto the tissue and incubating at room temperature for 5 min.



55 Dip in a 50 mL falcon tube with DI water and gently dry with air flow.

Bulk Tissue Reverse Transcription

56 Primer sequences: 15T Biotin RT Primer:
/5Phos/CATCGGCGTACGACTNNNNNNNNNN/iBiodT/TTTTTTTTTTTTTTTTTVN

57 Reverse transcription mix as below

	A	B	C	D
1	Reagent name	Concentration	For 1 chip	For 2 chips
2	0.5X PBS-RI		20µL	40µL
3	5X Maxima RT buffer	5X	12µL	24µL
4	30T Biotin RT Primer	25uM	10µL	20µL
5	Maxima H Minus RT	200U/µL	6µL	12µL
6	Nuclease-free Water		8µL	16µL
7	dNTPs (10mM)	10mM	3µL	6µL
8	SupraseIn RNase Inhibitor	20U/µL	0.8µL	1.6µL
9	RNase Inhibitor	40U/µL	0.4µL	0.8µL

Reverse transcription mix

58 The 60µL RT Mix was loaded on the sample with PDMS reservoir and sealed with parafilm. Then, incubated at room temperature for 30 min, and 42°C for 90 min, in a wet box.

59 1X NEB buffer 3.1: 100µL 10X NEB buffer + 900µL Nuclease-free Water.

60 After RT reaction, samples were flow washed with 1 mL Nuclease-free Water for 5 times, then incubated in 500µL 1X NEB buffer 3.1 for 5 min at room temperature, followed by air drying.

DBiT-seq assembly and Barcoding



- 61 Refer to <https://www.protocols.io/view/spatial-multi-omics-sequencing-for-fixed-tissue-vi-3byl4j2kolo5/v1>

Lysis

- 62 Take a small PDMS solution reservoir and place it on the tissue slide so that the reservoir opening is directly over the barcoded ROI.
- 63 Load the slide into a lysis clamping device and tighten screws as much as possible by hand.
- 64 Image the lysis chip on the ROI using a 10x objective lens.
- 65 Prepare the reverse crosslinking lysis solution.

Reagent	Volume	Stock Concentration	Final Concentration
Tris-HCl (pH 8.0)	7.5 μ L	1M	50 mM
EDTA	0.3 μ L	0.5M	1 mM
SDS	15 μ L	10%	1%
NaCl	6 μ L	5M	200 mM
proteinase K	3 μ L	20mg/mL	0.4 mg/mL
dH2O	118.2 μ L	-	-

- 66 Add approximately 50- 100 μ L of the reverse crosslinking lysis solution into the lysis chip. CRITICAL STEP Make sure to record the volume added of the lysis solution for the remaining steps.
- 67 Cover the ROI hole using parafilm and load the slide into a humidifying chamber.

- 68 Incubate tissue the slide with PDMS reservoir and Lysis Solution at 58 °C for 2 hours to reverse formaldehyde crosslinks.
- 69 After incubation, collect the lysate in a 200µL PCR tube (or 1.5mL tube) and seal the tube with parafilm.
- 70 Incubate the lysate at 65 °C with rotation overnight.

Purification

- 71 If necessary, transfer the lysate to a 1.5mL micro-centrifuge tube.
- 72 During this transfer, carefully use a P200 pipette to measure the volume of the lysate, for an accurate PCR product: DNA fragment ratio in the Zymo purification.
- 73 Purify the lysate with the Zymo DNA Clean & Concentrator-5 according to the kit protocol:
 - a. Follow the 5:1 DNA Binding Buffer: DNA Sample
Use a pipette to accurately determine the initial volume of the PCR Product
 - b. For the final elution, elute with 100 uL of DNA Elution buffer

cDNA Purification

- 74 The cDNA is purified and bound to DynabeadsTM MyOneTM Streptavidin C1 beads in this step.
Prepare 40 µL DynabeadsTM MyOneTM Streptavidin C1 beads (Thermofisher) per sub-library.
- 75 Wash 3 times with 800 µL of 1X B&W buffer with 0.05% Tween-20.
- 76 Resuspend beads in 100 µL 2X B&W buffer + 2 µL SUPERase[•] InTM RNase Inhibitor.
- 77 Purify lysate following the DNA Clean & Concentrator with Zymo-spin IC Columns (RPI Research Products) protocol. Use 100 µL water to elute DNA.



- 78 Add 100 μL of resuspended DynabeadsTM MyOneTM Streptavidin C1 magnetic beads to each lysate.
- 79 Rotate at 30 RPM and room temperature for 60 min for binding to occur.
- 80 Save the supernatant for the spatial-ATAC-seq library
- 81 Wash twice with 400 μL of 1X B&W buffer with 5 min rotation after resuspending beads.
- 82 Wash once more with 400 μL 10 mM Tris containing 0.1% Tween-20 with 5 min rotation after resuspension.

Template Switch

- 83 This step performs template switching and adds the second PCR handle. Resuspend the streptavidin beads bound with cDNA in solution containing 44 μL of 5X Maxima RT buffer (ThermoFisher), 44 μL of 20% Ficoll PM-400 solution, 22 μL of 10 mM dNTPs each (ThermoFisher), 5.5 μL RNase Inhibitor (Enzymatics), 11 μL of Maxima H Minus Reverse Transcriptase (ThermoFisher), 5.5 μL of 100 μM of template switch primer, and 88 μL of RNase-free water.
- 84 Rotate beads at 30 RPM and room temperature for 30 min followed by rotation at 42°C for 90 min.

PCR fro

- 85 Wash beads and resuspend.
- 86 Wash once with 400 μL of 10 mM Tris and 0.1% Tween-20 solution and once more with 400 μL of RNase-free water.
- 87 Resuspend in the PCR mix solution containing 110 μL of 2X Kapa HiFi HotStart Master Mix (Kapa Biosystems), 8.8 μL each of 10 μM stocks of primers 1 and 2, and 92.4 μL of RNase-free water. Transfer 200 μL to four PCR tubes with 50 μL in each.
- 88 Perform PCR to detach cDNA from beads.



	Steps	Temperature	Time	Cycles
1	Initial Denaturation	95°C	3 min	1
2	Denaturation	98°C	20 sec	5
3	Annealing	65°C	45 sec	
4	Extension	72°C	3 min	

PCR Cycling Conditions

- 89 Remove the Dynabeads™ from the PCR solution using a magnetic tube holder.
- 90 Add Evagreen (Biotium) at a 1X concentration.
NOTE: This step is optional.
- 91 Perform PCR again with the following thermocycling conditions. Cycling can also be halted once the qPCR signal begins to plateau.

	Steps	Temperature	Time	Cycles
1	Initial Denaturation	95°C	3 min	1
2	Denaturation	98°C	20 sec	15 cycles
3	Annealing	65°C	20 sec	
4	Extension	72°C	3 min	
5	Final Extension	72°C	5 min	1
6	Hold	4°C	forever	

cDNA Purification

- 92 Add 80 µL KAPA pure beads for 100 µL of the cDNA sample.
- 93 Mix thoroughly by pipetting or vortexing.
- 94 Incubate the tubes at room temperature for at least 10 min to bind the cDNA to the beads.



- 95 Keep the tubes on the magnet and add 200 μ L of 80% ethanol.
- 96 Carefully remove and discard the supernatant.
- 97 Incubate the tubes at room temperature for at least 30 sec.
- 98 Carefully remove and discard the ethanol.
- 99 Repeat steps 102-104.
- 100 Remove the tubes from the magnet and resuspend the beads in 15-20 μ L of PCR-grade water or elution buffer depending on downstream application.
- 101 Incubate the tubes at room temperature for at least 10 min to elute the cDNA off the beads.
- 102 Place the tubes back on the magnet and incubate until the liquid is fully clear.
- 103 Transfer the clear supernatant to a new tube.
- 104 Perform Bioanalyzer QC to obtain cDNA length profile and concentration.

Pause Point: Stop here for the second day. Store at 4°C for 1-2 weeks or up to 6 months at -20°C.

Library Preparation

- 105 PCR product from previous step can be sequenced.

Purification of spatial-ATAC-seq product

- 106 If necessary, transfer the lysate to a 1.5mL microcentrifuge tube.
- 107 During this transfer, carefully use a P200 pipette to measure the volume of the lysate, for an accurate PCR product: DNA fragment ratio in the Zymo purification.
- 108 Purify the lysate with the Zymo DNA Clean & Concentrator-5 according to the kit protocol:
 - a. Follow the 5:1 DNA Binding Buffer: DNA Sample
Use a pipette to accurately determine the initial volume of the PCR Product
 - b. For the final elution, elute with 21 uL of DNA Elution buffer

PCR for spatial-ATAC-seq

2h

- 109 Prepare the PCR solution:
 - a. 21uL purified DNA
 - b. 2.5uL 25uM Universal or barcoded i5 primer
 - c. 2.5uL 25uM Universal or barcoded i7 primer
(using different indices if needed)
 - d. 25uL 2X NEBNext Master Mix

Pipette up and down to mix once you have added everything.
Briefly vortex and spin down the tubes.
Place the tube in the NEW thermocycler and run the following protocol with a heated lid:

 - a. Lid should be heated at 105°C
 - b. Used default ramp rate

	Steps	Temperature	Time	Cycles
	Initial Denaturation	72°C	5 min	1
	Denaturation	98°C	30 sec	
	Denaturation	98°C	10 sec	5
	Annealing	63°C	30 sec	
	Extension	72°C	1 min	

PCR Cycling Conditions



110 Determine Additional Cycles

1. Prepare preliminary qPCR solution :

- Pre-amplified library , 5uL
- Pre-amplified solution, 0.5uL
- 25uM Universal or barcoded i5 primer, 0.5uL
- 25uM Universal or barcoded i7 primer (using different indices if needed), 5uL
- NEBNext Master Mix, 3.25uL
- Nuclease-free water, 0.75uL
- For a final concentration of 1X EvaGreen Dye, 0.75 uL

Pipette up and down to mix once you have added everything.

Briefly vortex and spin down the tubes.

Place the tube in the NEW thermocycler and run the following protocol with a heated lid:

- Lid should be heated at 105°C
- Use default ramp rate

Denaturation	98°C	30 sec	
Denaturation	98°C	10 sec	20 cycles
Annealing	63°C	30 sec	
Extension	72°C	1 min	

2. Use the qPCR data in Excel to determine the cycle where you reach 1/3 of the saturated signal in qPCR

- Graph the qPCR fluorescence values
- It's a good run if it plateaus in less than 10 cycles
- $((\text{fluorescence value at cycle 20} - \text{fluorescence value at cycle 1})/3) + \text{fluorescence value at cycle 1} = 1/3 \text{ Saturated Signal Fluorescence Value}$
- Determine how many cycles it takes to get to this 1/3 Saturated Signal Fluorescence Value. If in the middle of two cycles, always take the lower.

Note: Sometimes, all the graphed values may be negative if the amount of DNA is very high. This is fine, and just proceed to determine the number of cycles.

111 Repeat step 68 PCR protocol to amplify the remaining product with the cycles determined in step 69



Ampure Purification and Library preparation for spatial-ATAC-seq

40m

- 112 Perform a 1X SPRI purification by adding 45uL Ampure XP beads to the sample. Mix thoroughly by pipetting or vortexing.
- 113 Mix thoroughly by pipetting.
- 114 Keep the tubes on the magnet and add 200 μ L of 80% ethanol.
- 115 Carefully remove and discard the supernatant.
- 116 Incubate the tubes at room temperature for at least 30 sec.
- 117 Carefully remove and discard the ethanol.
- 118 Repeat steps 102-104102-.
- 119 Remove the tubes from the magnet and resuspend the beads in 15-20 μ L of PCR-grade water or elution buffer depending on downstream application.
- 120 Incubate the tubes at room temperature for at least 10 min to elute the cDNA off the beads.
- 121 Place the tubes back on the magnet and incubate until the liquid is fully clear.
- 122 Transfer the clear supernatant to a new tube.
- 123 Perform Bioanalyzer QC to obtain cDNA length profile and concentration.

Pause Point: Stop here for the second day. Store at 4°C for 1-2 weeks or up to 6 months at -20°C.