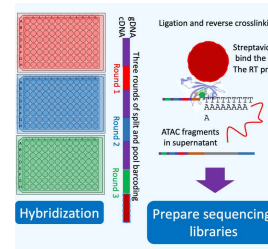


Sep 25, 2024 Version 3

SHARE-seq protocol v2.2 V.3

This protocol is a draft, published without a DOI.



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Protocol status: Working

We use this protocol and it's working

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Disclaimer

Updates in progress, however this version is stable and reflects how we process cells/nuclei in the lab.

Abstract

An updated version of the protocol SHARE-seq, as used by the Epigenomics Platform and Gene Regulation Observatory at the Broad Institute in the service of data production for the IGVF project. Link to the original paper and protocol here:

<https://www.sciencedirect.com/science/article/pii/S0092867420312538>

Materials



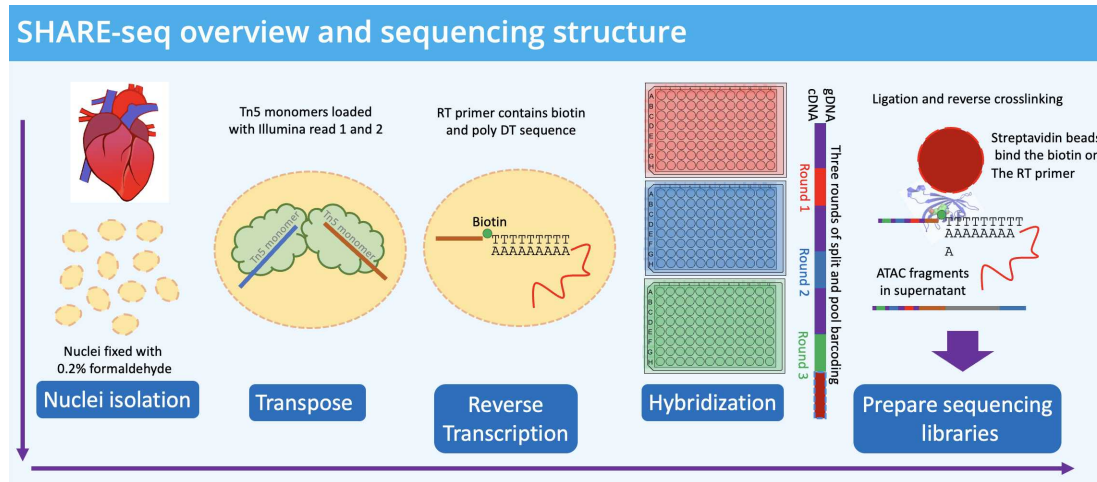
SHARE_seq_ordering_reagents.xlsx 13KB

Before start

Some general notes and guidelines (many thanks to Liz Rebboah, Ryan Weber, and Ali Mortazavi for their thoughtful comments and suggestions). A slightly irreverent way to describe SHARE would be "punch a bunch of holes in a bag (the cell/nucleus), manipulate it a lot, hope it doesn't fall apart." That is to say: this is a lossy protocol! Whatever you start with, you'll probably only retain 10-25% of the starting material by the end of the first day. One of the biggest challenges of SHARE is keeping the RNA quality high (the ATAC is more stable). To support this, at the Broad we generally aliquot the following reagents: DTT, 10x SYBR, proteinase K, PIC, BSA (in 5ml tubes). We also aliquot buffer stocks (like Tris, MgCl₂, NaCl) into 50mL or 15mL conical tubes. If there are any issues with the RNA capture or quality (as revealed by running gels on the second day of SHARE), throw out your current in use aliquot or stocks! Then you can balance a need for RNase free reagents with avoiding waste.

1.1 Ordering Oligo Plates and oligos for plate production

1 A visual overview of SHARE-seq:



This image shows a workflow for SHARE-seq

2 One of the more involved aspects of SHARE-seq (and SPLiT-seq) is properly making the oligo hybridization 96 well plates - this section covers that in detail before moving into any of the day to day aspects of this protocol (those start at step 24, in section 2).

In summary, this section will walk you through ordering deepwell 96 well oligo plates for all 3 rounds of hybridization in SHARE-seq (or SPLiT-seq), with an additional set of 96 oligos that can be ordered ("plate 2"), plus linker oligos.

In the [original SPLiT-seq paper](#) the authors calculated a barcode collision rate above ~5% with greater than around 25,000 cells or nuclei in a cellular sub pool (these are generated at the end of the first day of SHARE-seq). So for experiments using a singular set of three rounds of 96 barcodes, don't create cellular sub pools greater than 25,000 cells - the rate of barcode collisions will be high. If a second plate of 96 barcodes is used (we refer to these as "plate2"), and the two plates are pooled together and intermixed after each round of hybridization, we determined that up to 100,000 cells or nuclei can be pooled together safely, with low (<5%) risk of barcode collision.

3 Ordering oligo deepwell plates from IDT:


Using the attached Excel file, batch order the oligos on the tabs labeled "Plate1 R1 96", "Plate1 R2 96", "Plate1 R3 96" from IDT in a 96 well "deepwell" plate. If 192 barcodes are desired (2 plates), then order the oligos on the tabs labeled "Plate2 R1 96", "Plate2 R2 96", "Plate2 R3 96." Order at 25nM yield, 100uM concentration, resuspended in IDTE buffer, in a 96 well plate format. An example of what the plate upload looks like is shown below. Note that the R3

plate(s) will be less expensive than the R1 or R2 plates, since there is no 5' phosphate base needed in these molecules.

 2023_06_27_SHARE_seq_oligo_plat...

File Upload ×

Plate Upload Instructions

- Only one plate can be entered per tab
- Empty wells do not need to be indicated
-  Download a sample Excel plate ordering template.

1. Please specify an excel file.

No file selected.

2. Review and select plates you would like to add

	Plate	# Wells
<input checked="" type="checkbox"/>	Plate1_R1 96	96
<input checked="" type="checkbox"/>	Plate1_R2 96	96
<input checked="" type="checkbox"/>	Plate1_R3 96	96
<input type="checkbox"/>	Plate2_R1 96	96
<input type="checkbox"/>	Plate2_R2 96	96
<input type="checkbox"/>	Plate2_R3 96	96

ADD PLATES

What it looks like to select plates from the above excel file for IDT batch plate ordering.

Name

Plate1_R1 96

Normalization Type

Full Yield

Scale

25 nmole DNA Oligo

Concentration(μM)

100

Min: 50 Max: 200

Purification

Standard Desalting

Volume (μL)

0

Min: 20 Max: 800

Plate Type

Deep Well

Ship Option

Wet

Buffer

IDTE 8.0 pH



DUPLICATE

APPLY SETTINGS TO ALL PLATES



REMOVE

Ordering specifications for all plates ordered

- 4 You'll also need linkers to anneal with the oligos in each plate to create the unique cellular barcode during the hybridization of SHARE. Order the oligos below from IDT, order the linker oligos at 1mM (1000uM) concentration, and the blocking oligos at 100uM; order oligos resuspended in IDTE.

Note: the 1uM is the "scale" of what you're ordering from IDT, not the concentration! The concentration is "formulation" from IDT, and that's 100uM in IDTE for the blocking oligos, and 1000uM in IDTE for the linker oligos.

A	B	C	D	E	F	G
Name	LEN GTH	Sequence	Sca le	Pur ific ati on	Spec ific o ligos	order from IDT
Round 1 lin ker	30	CCGAGCCCACGAG ACTCGGACGATCAT GGG	1u m	ST D		Round 1 linker,CCGAGCCCACG AGACTCGGACGATCATGGG,1u m,STD

A	B	C	D	E	F	G
Round 2 linker	30	CAAGTATGCAGCGC GCTCAAGCACGTGG AT	1u m	ST D		Round 2 linker,CAAGTATGCAGC GCGCTCAAGCACGTGGAT,1um, STD
Round 3 linker	30	AGTCGTACGCCGAT GCGAAACATCGGCC AC	1u m	ST D		Round 3 linker,AGTCGTACGCC GATGCGAAACATCGGCCAC,1u m,STD
Round 1 blocking	30	CCCATGATCGTCCG AGTCTCGTGGGCTC GG	1u m	ST D		Round 1 blocking,CCCATGATC GTCCGAGTCTCGTGGGCTCGG, 1um,STD
Round 2 blocking	30	ATCCACGTGCTTGA GCGCGCTGCATACT TG	1u m	ST D		Round 2 blocking,ATCCACGTG CTTGAGCGCGCTGCATACTTG,1 um,STD
Round 3 blocking	30	GTGGCCGATGTTTC GCATCGGCGTACGA CT	1u m	ST D		Round 3 blocking,GTGGCCGAT GTTTCGCATCGGCGTACGACT,1 um,STD

Table of linkers and blockers for construction of the SHARE-seq hybridization plates

1.2 Annealing Oligo plates - making oligo plates

- This part occurs only after you have all of the oligo deepwell plates and the Round 1,2,3 linker oligos from IDT. **If you don't have these yet, don't proceed with platemaking!**

The recipe below is for eighteen 96 well plates for each round of barcoding (54 total plates per set of 96 oligos, that's 108 if you're making both plate1 and plate2). We recommend that you use different color PCR plates for each round of each plate. Conveniently, ThermoFisher offers 96 well skirted plates in 6 different colors, search for part number AB2396N (where N=R is red, G=green, Y=yellow, B=blue, O=orange - the base model is clear). Please be sure you purchase part number AB2396, and not AB3396 - those plates have white wells that will make it hard to tell how much fluid is in the well.

- Thaw deepwell plates at RT for 20-40 minutes at RT, and spin down before usage to avoid oligo cross-contamination. **Can speed the thaw by setting on a thermomixer set to 28°C if needed.**
- Make 50mL of STE, recipe below:

A	B	C
Oligo Annealing buffer (STE)	Volume (mL)	1x Concentration
1M Tris pH 8.0	0.5	10 mM
5M NaCl	0.5	50 mM
0.5 M EDTA	0.1	1 mM



A	B	C
H2O	48.9	
Total	50	

recipe for STE

- 8 Using a 12 channel multipipettor:
- 9 Dilute 180µl Round 1 linker oligo (1 mM) with 7920 µl STE buffer. Mix 80 µl diluted Round 1 linker oligo with 20 µl Round 1 oligo (100 µM) in a 96 well PCR plate.
- 10 Dilute 259.2 µl Round 3 linker oligo (1 mM) with 7047 µl STE buffer. Mix 72 µl diluted Round 3 linker oligo with 28 µl Round 3 oligo (100 µM) in a 96 well PCR plate.
- 11 Dilute 216µl Round 2 linker oligo (1 mM) with 7371 µl STE buffer. Mix 76 µl diluted Round 2 linker oligo with 24 µl Round 2 oligo (100 µM) in a 96 well PCR plate.
- 12 Seal the plates - we use an **BioRad PX1 plate sealer**, seal at 180 degrees C for 5 seconds. We use the **specific foil seals** for that sealer as well (part number 1814045). If you don't have a heat sealer, we recommend BioRad Microseal F foil seals (part number MSF1001), which we use for sealing the final oligo plates as well.
- 13 Anneal Round 1, Round 2, and Round 3 plates using the PCR cycling conditions below. **The slow ramp is critical for this step!!** The plates won't work without the exact cycling conditions below!



A	B
95°C	2 min
Slow ramp, -1°C/cycle, 1 min per cycle	
20°C	2 min
4°C	Forever
Total	~1h 26 min

PCR cycling conditions for the oligo plate annealing



- 14 After annealing, check if there is significant water evaporation for the wells at the corners/sides of the wells. With the PX1 heat sealing, we don't find that this occurs, but if you used more standard foil seals, there may be some evaporation on the sides.
- 15 Add 100µl STE to each well of the annealed plate, mix 7-8x and transfer 100µl to another 96 well plate. These two plates are your "stock plates" from which you will aliquot into the multi-colored plates for each round of SHARE.
- 16 Aliquot 10ul from your stock plates (you'll have either 3 or 6 sets) to make an oligo plate. We use the Agilent Bravo to make this go faster, but it's fairly quick with a 12 channel pipette as well. Don't forget to: label the plates! The rounds have to go in order for hybridization to work! Also don't forget to put each "round" of oligo in a separate colored 96 well plate to reduce errors when selecting plates for the experiment.
- 17 Seal plates with foil and label using labels or marker (printed labels are faster if you've got them). Store the plates in rack at -30°C, they are good for at least 6 months, and probably 12 months.

1.3 Ordering and annealing oligos for cDNA tagmentation

- 18 SHARE-seq generates long cDNA (presuming no nuclease contamination or other harms befall the RNA). In this state it is overlong to sequence, so we must tagment it using Tn5 complexed with Illumina Read 1 only. To do this, we use commercial Tn5 from Diagenode, and incubate it with annealed oligos that include Illumina Read 1, and block the ME_Comp region of that sequence.
- 19 Order the following oligos from IDT resuspended at 100uM in IDTE

A	B	C	D	E	F	G
Name	LENGTH	Sequence	Scale	Purification	Specific oligos	order from IDT
Read1	33	TCGTCGGCAGCGTCAG ATGTGTATAAGAGACAG	1um	HP LC	*	Read1,TCGTCGGCAGCGTCAG ATGTGTATAAGAGACAG,1um, HPLC
Blocked_ME_Comp		/5Phos/C*T*G* T*C*T* C*T*T* A*T*A* C*A*/3ddC/	1um	HP LC	*	Blocked_ME_Comp,/5Phos/C* T*G* T*C*T* C*T*T* A*T*A* C* A*/3ddC/ ,1um,HPLC

Read1 and Blocked_ME_Comp to order

Wait for them to arrive before proceeding!

20

Prepare tagmentation adapter mix in PCR tubes.

A	B
Tagmentation Adapter	ul



A	B
100uM Read1	26
100uM Blocked_ME_Comp	26
1M Tris pH 8.0	0.52
5M NaCl	0.52
Total	53

tagmentation adapter mix, ideally make several aliquots, the above recipe is for a single aliquot.

- 21 Anneal oligos in a thermal cycle as follows.



A	B
85°C	2 min
Slow ramp, -1°C/cycle, 1 min per cycle	
20°C	2 min
4°C	Forever
Total	~1h 14 min

PCR cycling conditions for annealing the tagmentation adapter oligos

- 22 Heat a 150ul aliquot of RNase free 100% glycerol to 65°C in a heat block or thermomixer (RNase zap the surfaces). Make two 50ul aliquots and equilibrate to RT. Mix the annealed tagmentation adapter with an aliquot of glycerol, respectively (i.e. mix each 50ul aliquot of adapter with a separate 50ul aliquot of glycerol). The annealed adapters can be immediately used.

1.4 Essential SHARE-seq buffers (make the day before performing an experiment)

- 23 Make each of the following buffers - note that for large experiments you may need more than one aliquot of these buffers.



A	B	C
Dilution buffer	Volume (ul)	Final Concentration
100% glycerol (wide-bore tips)	500	50%
1M Tris pH 7.5	50	50 mM
5M NaCl	20	100 mM
5mM EDTA	20	0.1 mM
1M DTT	1	1 mM
10% NP-40	10	0.1%
H2O	390	
Total	1000	

Dilution buffer recipe. This buffer is used to dilute Tn5 enzyme for transposition and tagmentation

A	B	C
5x SMART RT buffer (adapted from smart-seq-3)	Volume (ul)	Final Concentration (of 5x)
1000mM (1M) DTT	40	40 mM
1M Tris pH 8.3	125	125 mM
100 mM GTP	50	5 mM
5M NaCl	30	150 mM
1M MgCl ₂	12.5	12.5 mM
H2O	742.5	
Total	1000	

SMART reverse transcription buffer - adapted from the smart-seq-v3 protocol.

You can also make the NI (see step 28) the day before SHARE. We recommend making a fresh 50mL aliquot for every experiment as a safeguard against RNase contamination.

1.5 Ordering SHARE-seq oligos

- 24 All of the below oligos are necessary for performing SHARE-seq (mainly amplifying libraries on Day 2 of SHARE). Wait until these arrive to start the SHARE-seq protocol. All should be ordered formulated at 100uM in IDTE.



A	B	C	D	E	F	G
Name	LENGTH	Sequence	Scale	Purification	Specific oligos	order from IDT
RT_primer	61	/5Phos/GTCTCGTGGGCTC GGAGATGTGTATAAGAGA CAGNNNNNNNNNN/iBiod T/TTTTTTTTTTTTTTTVN	1u m	RNA SE	*	RT_primer,/5Phos/GTCTCGTG GGCTCGGAGATGTGTATAAGA GACAGNNNNNNNNNN/iBiod T/TTTTTTTTTTTTTTTVN,1um, RNASE
P7	24	CAAGCAGAAGACGGCATA CGAGAT	100 nm	STD	*	P7,CAAGCAGAAGACGGCATA CGAGAT,100nm,STD
RNA_PCR_primer	23	AAGCAGTGGTATCAACGC AGAGT	100 nm	STD	*	RNA_PCR_primer,AAGCAGTG GTATCAACGCAGAGT,100nm, STD

DNA oligos for SHARE to order from IDT


When the RT primer arrives, aliquot into single use amounts (ex: for 1e6 cells, you would need ~300ul of the RT primer). This oligo is long and somewhat unstable, so minimize freeze thaw cycles.

A	B	C	D	E	F	G
Name	LENGTH	Sequence	Scale	Purification	Specific oligos	order from IDT
TSO	30	AAGCAGTGGTATCAACG CAGAGTGAATrGrG+G	1u mR	RNASE	*	TSO,AAGCAGTGGTATCAAC GCAGAGTGAATrGrG+G,1u mR,RNASE


TSO (Template Switching Oligo) RNA oligo for SHARE-seq

2.0 Washing and Counting Fixed cells or nuclei (time: 30-60m)

- 25 This section presumes cells or isolated nuclei have been fixed in formaldehyde (0.2% for cultured cells/nuclei, 1% for primary PBMCs) and have been frozen at -80 degrees as a dry pellet. For more details on nuclei isolation, see the attached files (will be made into protocols.io links as time permits):

 2023_06_12_cell_Fixation_and_Free... #fixation and freezing guide for primary/cultured cells.

 GRO nuclear prep for SHARE-seq (u... 116KB #nuclear preparations from post-mortem tissue

 Nuclear Prep Protocol (edited for low ... 119KB #nuclear preparation protocol optimized for small amounts of tissue (25mg). Both protocols include an iodixanol gradient step and fixation instructions.

Important note: if your samples are PBMCs or HSCs, you'll want to use the optional polyAdenylation module (section 4) and perform centrifugation at 1000xg.

- 26 BEFORE YOU BEGIN!!! Spray down your bench with RNase zap! Spray your pipettors with RNase Zap! Put on a surgical mask! Spray your gloves with RNase Zap all the time! I know this seems annoying, but RNA will degrade easily in this protocol, and taking precautions early will really help!
- 27 Put swinging bucket centrifuge rotor (Eppendorf # S-24-11-AT) in the refrigerable benchtop Eppendorf centrifuge (model #5430R) and cool down to 4°C.
- 28 Prepare RNase free, sterile NIB-2RI, NI-2RI, NID-2RI (as well as NIB and NI). NIB and NI are stable for at least 6 months at room temperature. We always make a fresh aliquot of NI when making the 5x SMART RT buffer and dilution buffer.



A	B	C
Nuclei Isolation Buffer (NIB)	Volume (mL)	Final Concentration
1M Tris-HCl pH 7.5	0.5	10mM
5M NaCl	0.1	10mM
1M MgCl ₂	0.15	3mM
10% NP40	0.5	0.1%
H ₂ O	48.75	
Total	50	

NIB buffer recipe

Nuclei Isolation (NI)	Volume (mL)	Final Concentration
-----------------------	-------------	---------------------



1M Tris-HCl pH 7.5	0.5	10mM
5M NaCl	0.1	10mM
1M MgCl ₂	0.15	3mM
H ₂ O	49.25	
Total	50	

NI buffer recipe

A	B
NIB-2RI	Volume (ml)
NIB	1000
Enzymatic RI	7
SUPERase RI	7

NIB-2RI recipe. Around 100uL per sample is used for cells/nuclei

A	B	C
NI-2RI	Volume (ml)	Volume (ml)
NI (per expt)	1000	50000
Enzymatic RI	6.25	125
SUPERase RI	6.25	125
7.5% BSA	5.33	266.67

NI-2RI recipe - around 7mL per sample is used for cells/nuclei

- 29 Transfer fixed cells to a new 1.5mL tube coated with 7.5% BSA. If starting from a fixed pellet, use 500ul NI-2RI to resuspend and transfer to a BSA coated tube.

Preparing 1.5ml BSA coated tubes: Prepare a 1.5mL Eppendorf tube by adding 150ml 7.5%



- BSA, vortex, remove supernatant, let tube sit on ice briefly, remove any collected BSA with a P20.
- 30 Spin 750xg, 4°C, 5 minutes. Gently remove supernatant.
- 31 Wash with 500ul NID-2RI without disturbing pellet.
- 32 Spin 750xg, 4°C, 5 minutes. Gently remove supernatant.
- 33 Resuspend in 50ul NID-2RI.
- 34 Add 100ul NIB-2RI to each sample, incubate for 5 minutes on ice to permeabilize the nuclei.
- 35 Add 900ul NI-2RI (NI-RI: **1200ul NI + 3ul Enzymatic RI**) to each sample, mix by inverting
- 36 Spin 750xg, 4°C, 5 minutes. Gently remove supernatant, using a P20 to remove as much NIB-2RI as possible.
- 37 Turn on a thermomixer (Eppendorf ThermoMixer C) for the ATAC Tn5 transposition, set to 37 degrees and put the 1.5mL block on the thermomixer (you should have a 96 well block for the hybridization step).
- 38 Resuspend in 50ul NI-RI (NI-RI: **1200ul NI + 3ul Enzymatic RI**). SUPERase RI inhibits the Tn5 transposition, so using RI instead of 2RI is very important here.
- 39 Use 2.5ul cells and 7.5ul HCB-RI plus 10ul Trypan blue to count nuclei using the BioRad TC20 (part number 1450102) or a manual hemocytometer (we like the Reichert Bright-Line: http://hauserscientific.com/products/reichert_bright_line.html). Determine cell counts per mL and per ul, so the volume needed to get to 10000 nuclei is known. Multiply the count by 4, since the cells/nuclei are diluted 1:4 before adding trypan blue. The TC20 takes trypan blue dilutions into account, so divide the count per mL by 1000 to get the count per ul.
- 40 Determine the volume of cell suspension that is 10,000 cells (one ATAC transposition), and the total number of ATAC reactions (10,000 cells/reaction). We typically do SHARE using 25 ATAC reactions (250K cells) as the basic "unit" of SHARE.



3.0 ATAC Tn5 transposition (time: ~45m)

41 Assemble transposome by mixing assembled Tn5, and dilution buffer. Here we use volumes of ASSEMBLED Tn5 from the SeqWell produced Tn5 (Tagify: <https://seqwell.com/tagify-umi-reagents/>)

41.1 We use 0.6ul per reaction (10000 cells) (enzyme concentration is 10mM).

42

A	B	C
Transposome	Volume (ml)	Volume N=25
1x SeqWell Tn5 (volume as of 01/27/2024)	0.6 × N reactions	15ul
Dilution buffer	1.875 × N reactions	46.9ul
Total	2.5 × N reactions	61.9ul

Recipe for assembling the Tn5 for tubes of ATAC reactions (25 reactions per 1.5mL tube max)

For every 250K of cells, prepare one of the above tubes. You could also prepare a master mix using 10% overage as well, but we do a single tube for every ATAC reaction tube to use a bit less of the Tagify reagent.

43 Calculate the total number of ATAC reactions for this experiment. Prepare 1x TB accordingly to the recipe below. Note that 1x TB buffer must be made fresh.

A	B	C
1x TB	Volume (ml) × samples (N)	Volume N=27.5 (per tube plus overage)
0.2M Tris-acetate	8.25 × N	226.875
5M K-acetate	0.66 × N	18.15
1M Mg-acetate	0.5 × N	13.75
100% DMF	8 × N	220
H2O	24.04 × N	661.1



A	B	C
PIC	$0.2 \times N$	5.5
Enzymatic RI	$0.85 \times N$	23.375
Total	$42.5 \times N$	1168.75

Recipe for 1xTB buffer

- 44 The total volume of cells for the N number of ATAC reactions should fit into a 50ul reaction. Determine the total volume of cells needed for the ATAC reactions, and then use NI to bring the total volume up to 50ul. Example: with 25 ATAC reactions (250K cells), and 13,920 cells per ul, use 250K/13920 to get 17.95ul of cells needed. Add 50ul-17.95ul = 32.05ul NI-RI to get 50ul total volume of cells.
- 45 Add $42.5 \times N$ ul of 1xTB to cells + NI (in 50ul total volume): 1062.5ul for 25 rxns/sample, incubate at RT for 10 minutes (this is a permeabilization step).
- 46 Add $2.5 \times N$ ul (61.9ul for 25 reactions) of assembled Tn5 to sample, mix well by pipetting.
- 47 In the 1.5mL tube, seal, shake in a thermomixer at 500rpm for 30 minutes, 37°C.
- 47.1 Every 5 minutes, take the entire block off the thermomixer and invert to mix
This resets the timer on the thermomixer! So use an external timer
- 48 Add 500ul NID-2RI to each tube, spin down at 750xg, RT, 5 minutes.
- 49 Remove supernatant, wash with 500ul NI-2RI, spin down at 750xg, RT, 5 minutes.
- 50 Remove supernatant, resuspend cells in 60ul NI-2RI (from a single ATAC reaction tube).
- 51 Set thermomixer(s) to 23°C for the hybridization reaction (after the RT step) and put the 96 well blocks on.





4.0 Optional PolyAdenylation module (time: ~45m)

- 52 If your samples are from hematopoietic compartment (PBMCs, HSCs), you'll want to use the optional exogenous polyAdenylation module in this section before Reverse Transcription to improve RNA capture (since PBMCs have a lot of nucleases). If not, skip ahead to section 5.0 Reverse Transcription (starting at step 57).
- 53 For each sample, mix 60ul cells/nuclei in NI-2RI from the previous step with 240ul PolyA master mix (recipe below), and incubate at 37°C for 15 minutes in a thermomixer without mixing. (each 1.5mL ATAC tube is a single polyA reaction).



A	B	C
polyA mix	Volume (ml) × samples (N)	Volume per sample (60ul NI-2RI + nuclei)
H2O	21.1xN	126.6
5x SMART RT buffer	10xN	60
Enzymatic RI	0.3xN	1.8
SUPERase RI	0.6xN	3.6
rATPs (NEB:P0756S)	5xN	30
E coli PolyA (NEB: M0276 S)	3xN	18
Cells/NI-2RI	10xN	60
Total	50xN	300

Polyadenylation master mix recipe

- 54 Add 200ul NI-2RI, spin down at 1000xg, 5 minutes, RT.
- 55 Remove supernatant and add 200ul NI-2RI, mix by inverting.
- 56 Spin down at 1000xg, 5 minutes, RT, remove as much supernatant as possible.
- 57 Resuspend so the total volume is 60ul NI-2RI.
- 58 Proceed to step 57 of the protocol below and continue with the RT reaction. Each polyA tube becomes a single RT reaction (7 0.2mL PCR tubes).



5.0 Reverse Transcription (time: 50m-1h)

59 Make the following master mix for reverse transcription:

A	B
Reverse transcription (RT) mix	Volume/ATAC tube (ul)
5x SMART RT buffer	70
Enzymatic RI	2.19
SUPERase RI	4.38
dNTPs	17.5
100uM RT primer	35
H2O	10.94
50% PEG	105
Maxima H Minus Reverse Transcriptase (add right before RT reaction)	35
Total	280

recipe for the RT mix.

For each ATAC 25 reaction tube, add 280ul RT mix to 60ul cells/Ni-2RI, aliquot 50ul to 7 wells in a 96 well plate. Then, run the RT thermocycler protocol as described below.

A	B	C
50°C	10 min	hold to start
8°C	12 s	3 cycles
15°C	45 s	cycle
20°C	45 s	cycle
30°C	30 s	cycle
42°C	2 min	cycle



A	B	C
50°C	3 min	cycle
50°C	5 min	hold after cycles
Total	~41 min	

Cycling conditions for the reverse transcription reaction: set the lid temp to 60°C

- 60 Pool all reactions for a given sample, add 300ul NID-2RI **per 7 PCR tubes**, spin down at 750xg, 5minutes, RT in a 1.5mL Eppendorf tube coated with 7.5% BSA. Note: the pellet may look larger than before - this is due to BSA dropping out of solution.
- 61 Wash with 500ul NI-2RI, spin down at 750xg, 5minutes, RT

6.0 Hybridization

- 62 Remove supernatant and resuspend each sample in NI-2RI. The amount each sample is resuspended in depends on the number of samples, and plates, per experiment. For a single sample in a single plate (96 well): resuspend in 1152ul NI-2RI. For four samples on a single plate, resuspend each **sample** in 288ul NI-2RI.

fraction of plate	ul NI-2RI
2 plates	2304
1 plate	1152
1/2 plate	576
1/3 plate	384
1/4 plate	288
1/8 plate	144

NI-2RI volumes per sample by experiment size

- 63 Note: it is very important to thaw oligo plates to RT (~23°C) before hybridization. Spin down plates before using. **Coat all reservoirs used for pooling with 7.5% BSA.**

Make the hybridization buffer:



A	B	C	D
Hybridization buffer	Volume (ml)	Stock concentration	Final Concentration
H ₂ O	5510	--	--
T4 ligase buffer (10x)	1152	10x	1x
Enzymatic RI	28.8	20 U/ml	0.05 U/ml
SUPERase RI	92.16	40 U/ml	0.32 U/ml
7.5% BSA	184.32	7.50%	0.10%
Total	6967.28		

Hybridization buffer recipe for two plates

A	B	C	D
Hybridization buffer	Volume (ml)	Stock concentration	Final Concentration
H ₂ O	2755	--	--
T4 ligase buffer (10x)	576	10x	1x
Enzymatic RI	14.4	20 U/ml	0.05 U/ml
SUPERase RI	46.08	40 U/ml	0.32 U/ml
7.5% BSA	92.16	7.50%	0.10%
Total	3483.64		

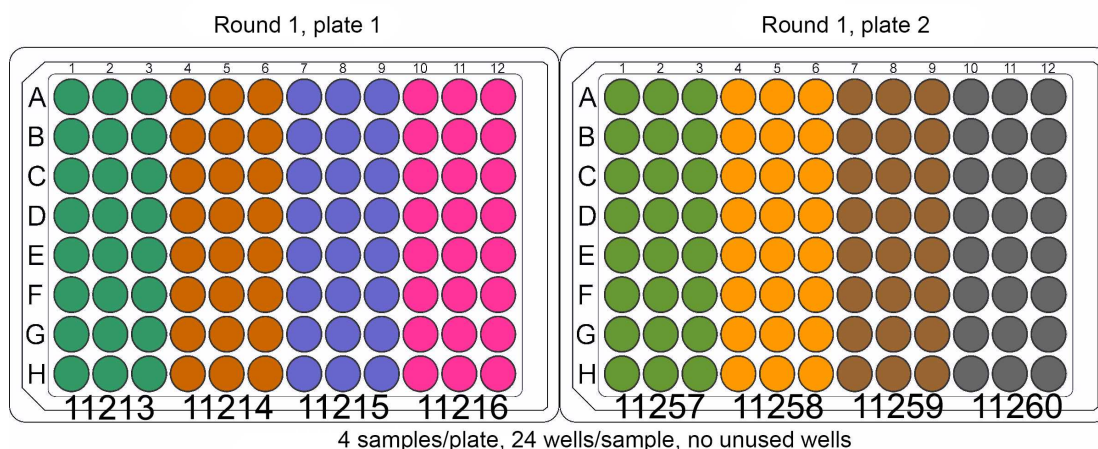
Hybridization buffer recipe for a single plate experiment

- 64 If you have a single sample: transfer the cells in NI-2RI into a BSA-coated reservoir suitable for an 8 or 12 channel multi-channel pipette. Add all of the hybridization buffer to the reservoir (adjust volume for 1 or 2 plates).
- 65 If you have multiple samples, put each sample in a **separate BSA-coated reservoir**, suitable for an 8-channel pipette (this makes loading samples across the 96 well plate easier). Add the hybridization buffer to each sample according to the table below:

fraction of plate	ul Hybridization buffer
2 plates	6967.28
1 plate	3483.64
½ plate	1741.82
⅓ plate	1161.21

¼ plate	870.91
⅛ plate	435.455

- 66 We find that using a 96 well plate map helps a lot for sample mapping! An example image is below, along with an open-access svg file of a 96 well plate.



An example of a plate to sample mapping file for the first round of hybridization. The five digit identifiers are lab-internal identifiers for a given aliquot of biological material.

96-Well_plate.svg 38KB

2x96_well_plates_share_template.psd 3.5MB

96-Well_plate.psd 1.4MB

For those who have access to Photoshop or Gimp, the *.psd files are editable templates. We recommend a making a layer to color the wells by sample identity.

- 67 Transfer sample(s) to Round 1 plate(s) as follows for each row of wells:
- Set a 12-channel pipette to 45ul, mix 2-3x
 - Transfer 45ul of mixture to Round 1 plates and mix by pipetting gently 5-7x.
 - Change tips between columns loaded!
- 68 Seal and shake at 300rpm for 25 minutes at RT (23 deg C).
- 69 Prepare Blocking oligo 1 as below and transfer to reservoir.

A	B	C
Blocking oligo 1	Volume (ul)	Volume (ul)



A	B	C
Round_1_blocking (100mM)	266.07	532.14
T4 ligase buffer (10x)	221.76	443.52
H2O	721.77	1443.54
Total	1209.6	2419.2

Column B is for a single plate, and column C is for two plates.

- 70 Add blocking oligo mix as follows:
- Using a 12-channel pipette, transfer 10ul Blocking oligo 1 to each well, pipetting up and down to mix thoroughly, changing tips between rows.
 - Seal and shake at 300rpm for 15 minutes at RT
- 71 Pool samples as follows:
- Set a 12-channel pipette to 70ul, pipette each row up and down 2-3 times, pipette up and down 2-3 times as ejecting each row into a **BSA-coated reservoir** (avoid bubbles).
 - After all rows have been pooled, pipette up and down 4-5 times to mix the reservoir
- 72 Open Round 2 oligo plates, and transfer 55ul of pooled mixture to each well, pipetting to mix each row. Shake at 300rpm for 25 minutes at RT.
- 73 Prepare Blocking oligo 2 as below and transfer to reservoir.

A	B	C
Blocking oligo 2	Volume (ul)	Volume (ul)
Round_2_blocking (100mM)	319.305	638.61
T4 ligase buffer (10x)	221.76	443.52
H2O	668.535	1336.86
Total	1209.6	2419.2

Column B is for a single plate, and column C is for two plates.

- 74 Add blocking oligo mix as follows:
- Using a 12-channel pipette, transfer 10ul Blocking oligo 2 to each well, pipetting up and down to mix thoroughly, changing tips between rows.



b. Seal and shake at 300rpm for 15 minutes at RT

75 Pool samples as follows:

- a. Set a 12-channel pipette to 80ul, pipette each row up and down 2-3 times, pipette up and down 2-3 times as ejecting each row into a **BSA-coated** reservoir (avoid bubbles).
- b. After all rows have been pooled, pipette up and down 4-5 times to mix the reservoir

76 Open Round 3 oligo plate, and transfer 65ul of pooled mixture to each well, pipetting to mix each row. Shake at 300rpm for 25 minutes at RT.

77 Prepare Blocking oligo 3 as below and transfer to reservoir.

A	B	C
Blocking oligo 3	Volume (ul)	Volume (ul)
Round_3_blocking (100mM)	278.25	556.5
H2O	931.35	1862.7
Total	1209.6	2419.2

Column B is for a single plate, and column C is for two plates.

78 Add blocking oligo mix as follows:

- a. Using a 12-channel pipette, transfer 10ul Blocking oligo 3 to each well, pipetting up and down to mix thoroughly, changing tips between rows.
- b. Seal and shake at 300rpm for 15 minutes at RT.

7.0 Ligation (time: ~45m)

79 Using either a P1000, or a 12-channel pipette and a BSA coated reservoir, aspirate the contents of each well and pool into a 15mL conical tube coated with 7.5% BSA.

- a. Total volume should be ~14mL (for a two plate experiment), but will fit in a single tube.
- b. For a single plate experiment, the total volume would be 7mL

80 Spin down at 750xg for 5 minutes at RT in a swinging bucket rotor.

81 Discard supernatant and add 1mL NID-2RI.

82 Spin down at 750xg for 5 minutes at RT in a swinging bucket rotor.



- 83 Discard supernatant and add 1mL NI-2RI, without disturbing pellet (let flow down side of tube slowly)
- 84 Spin down at 750xg for 5 minutes at RT in a swinging bucket rotor.
- 85 Discard supernatant and resuspend in 80ul NI-2RI. It's ok if there is a little bit of extra volume here - you'll just resuspend all of the cells into the ligation mix.
- 86 Mix sample with 320ul ligation mix (recipe below) and aliquot 50ul to 8 PCR tubes.

A	B
Ligation mix	Volume (ul)
H2O	253.7
Enzymatic RI	3.2
SUPERase RI	1
T4 ligase buffer (10x)	40
T4 ligase 400U /ul	20
7.5% BSA	2.13
Total	320

Ligation mix recipe. Note that if you have multiple single plate experiments, you'll need one ligation mix per experiment. If you're doing a double plate experiment (192 barcodes), you just need the single ligation mix recipe.

- 87 Shake at 300rpm for 30 min at RT.
- 88 Pool samples in a 1.5mL Eppendorf tube coated with 7.5% BSA, spin down 750xg, 5 minutes, RT, discard supernatant.
- 89 Gently resuspend pellet with 1mL NI-2RI

- 90 Spin at 750xg, 5 minutes, RT.
- 91 Presume 80% loss of nuclei over the course of the protocol, so for every 20,000 retained nuclei, resuspend in 10ul (goal is ~2000 cells or nuclei/ul).
For example: With 1 million starting cells, presume 100K are retained, and resuspend in 50ul NI-2RI.
- 92 Using 2ul cells and 8ul NI-2RI, mix with 10ul trypan blue. Count intact nuclei in a manual hemocytometer.
To get a count per mL, multiply hemocytometer counts by 100K (dilution factors: 2x = trypan blue, 5x = cell:NI dilution, 10,000x = hemocytometer dilution).
To get a count per ul, multiply counts by 100 (conversion: 100 = 100,000 (dilution factor for hemocytometer plus dilutions)/1,000 (microliters in one mL)).

8.0 Reverse crosslinking-pull down (time: 1h15m)

- 93 Here you will aliquot cells into cellular or nuclear "sub pools", and then digest the remaining proteins away to leave only barcoded nucleic acids. The number of cells per pool strongly depends on the experimental setup: a 96 barcode (single plate) experiment should not contain more than 25,000 cells or nuclei - the risk of barcode collisions (two different cells with the same randomly generated barcode) increases substantially (>2.5%). The math here is $96 \times 96 \times 96 = 884,000$, and $884K \times 0.025 = 22,118$.
- A 192 barcode (two plate) experiment has a maximum cell count of 100,000 cells per cellular or nuclear sub pool, though theoretically 200,000 should be possible (however we have not tested cellular/nuclear sub pools of this size as of 2024). Other important points: nuclease rich cells benefit from extra RNase inhibitors at the proteinase K stage as well as on the second day of SHARE. We have increased these numbers to working with pools around 50,000-100,000 cells or nuclei, but you may need to adjust these numbers depending on your source material. Replicates are also important: if you return 50,000 cells or nuclei for a two plate experiment, doing two cellular/nuclear sub pools of 25,000 cells is better than a single pool - it allows for concordance checking across replicates as well as providing some insurance for handling samples on the second day of SHARE.
- 94 Bring the volume of each sub-library up to 50ul using NI-2RI
- 95 For proteinase K digestion, add 50ul 2x RCB, 2ul proteinase K (20mg/mL), 2ul SUPERase RI

A	B	C
2x RCB	Volume (mL)	2x Concentration
1M Tris pH 8.0	1	100 mM



A	B	C
5M NaCl	0.2	100 mM
20% SDS	0.2	0.40%
H ₂ O	8.58	
Total	10	

2x RCB buffer - stable for 6-12 months at room temperature.

A	B	C	D	E
ProtK mix	Volume (ul) × samples (N+1)	5 samples: N+1=6	10 samples N+1=11	20 samples N+1=21
RCB	50	300	550	1050
Proteinase K (20mg/mL)	2	12	22	42
SUPERase RI	2	12	22	42
Cell in NI-2RI	50			
Total	104	324	594	1134

Proteinase K mix volumes. Add 54ul ProtK mastermix to each sample in 50ul NI-2RI

96 Incubate at 55°C for one hour on the thermal cycler with lid is set to 65°C

97 Can stop and store proteinase K digestions at -80°C for up to a week.



98 If coming from -80°C, thaw samples on ice first. Mix each reaction with 2.5ul 100mM PMSF in IPA and incubate at RT for 10 minutes.

98.1 To make 1mL 100mM PMSF in 100% IPA, use 17mg PMSF and weigh into a 1.5mL Eppendorf tube.



99 Prepare the following buffers:

A	B	C
2x BW	Volume (mL)	2x Concentration
1M Tris pH 8.0	0.5	10 mM
5M NaCl	20	2 M
0.5M EDTA	0.1	1 mM
H ₂ O	29.4	
Total	50	

2x BW recipe - stable at room temperature

A	B	C
1x BW-T	Volume (mL)	1x Concentration
1M Tris pH 8.0	0.25	5 mM
5M NaCl	10	1 M
0.5M EDTA	0.05	0.5 mM
10% Tween 20	0.25	0.05%
H ₂ O	39.675	
Total	50	

1x BW-T recipe - stable at room temperature

A	B	C
Oligo Annealing buffer (STE)	Volume (mL)	1x Concentration
1M Tris pH 8.0	0.5	10 mM
5M NaCl	0.5	50 mM
0.5 M EDTA	0.1	1 mM



A	B	C
H ₂ O	48.9	
Total	50	

STE recipe - stable at room temperature

100 Now prepare the above buffers with RNase inhibitors.

A	B
2x BW/RI	Volume/ul
2x BW	110x(N+1)
SUPERase RI	2x(N+1)

2x BW/RI - make this fresh. Note that the RNase inhibitor concentration is adjustable - this concentration works well for 25,000-100,000 cells or nuclei.

A	B
1x BW-T/RI	Volume/ul
1x BW-T	500x(N+1)
SUPERase RI	10x(N+1)

1x BW-T/RI - make this fresh. Note that the RNase inhibitor concentration is adjustable - this concentration works well for 25,000-100,000 cells or nuclei.

A	B
1x STE/RI	Volume/ul
STE	200x(N+1)
SUPERase RI	N+1



1x STE/RI - make this fresh. Note that the RNase inhibitor concentration is adjustable - this concentration works well for 25,000-100,000 cells or nuclei.

- 101 In a separate tube, mix 10xN ul of MyOne C1 Dynabeads with 100xN ul of 1xBW-T and place on a magnetic rack
- 102 Remove supernatant, wash twice with 100xN ul BW-T without RI
- 103 Remove supernatant, wash once with 100xN ul BW-T/RI, and resuspend beads in 110xN ul 2x BW/RI.
- 104 Add 100ul of washed beads to each sample, nutate at RT for 60 minutes
- 105 Put on a magnetic rack, transfer supernatant containing transposed ATAC chromatin fragments to a new tube for future library preparation. The supernatant is stable at RT for several hours (but we prepare the ATAC libraries right away, using a paired system where one person handles the cDNA, and the other can handle the ATAC, see step #102 below).
- 106 Wash cDNA/RNA-bound beads with 100ul 1xBW-T/RI, three times.
- 107 Place sample on magnetic rack, wash with 100ul 1xSTE/RI, without resuspending beads

9.0 RNA library preparation: template switching (time: 2h, ATAC prep during incubations)

- 108 Make the template switch mix using the recipe below:

A	B	C
Template switch mix	Volume (ul)	Final Concentration
H2O	1.25x(N+2)	
50% PEG 6000	15x(N+2)	15%
5x SMART RT buffer	10x(N+2)	1x
Ficoll PM-400 (20%)	10x(N+2)	4%



A	B	C
10 mM dNTPs, each	5x(N+2)	1 mM
RNase inhibitor (Lucigen)	10x(N+2)	0.2x
100 mM TSO	1.25x(N+2)	2.5 mM
Maxima H Minus Reverse Transcriptase (add right before RT reaction)	2.53x(N+2)	
Total	55x(N+2)	

Template Switch Mix recipe. Note that we have doubled the amount of the lucigen RNase inhibitor to better preserve RNA.

- 109 Remove all supernatant from the beads, resuspend beads in 55ul Template switch mix, be careful to avoid beads drying out after removing supernatant
- 110 Rotate/nutate samples at RT for 30 minutes
- 111 Incubate at 42°C for 90 minutes, shaking at 300rpm, resuspend beads at start, 30 minutes, 60 minutes, and at the end of the incubation by flicking the tubes to mix. We find RNase contamination can happen when the samples are pipetted, so we recommend mixing the settled beads by flicking for this reason.

10.0 ATAC library preparation (time: 2h50m, variable depending on qPCR results)

- 112 Using the supernatant with ATAC fragments from step 95, purify the samples using a Qiagen MinElute PCR purification kit (Qiagen cat #: 28006) as follows:
 - 112.1 In new tube, mix sample and 5x volume Qiagen PB buffer (1000ul), then add to MinElute spin column in two batches 2x600ul, spinning, then discarding flowthrough after each addition (as below).
 - 112.2 Spin down the tube at 13,000rpm for 1 minute and discard the flowthrough
 - 112.3 Add 750ul of PE Wash Buffer to the column, spin down at 13,000rpm for 1 minute, and discard flowthrough.
 - 112.4 Spin the samples at 13,000rpm for 1 minute to capture any excess buffer.



- 112.5 Place the spin column into a new tube and add 11ul of EB (10mM Tris, pH 8.0) directly to the column matrix. Incubate for 1 minute.
- 112.6 Spin down the tube at 13,000rpm for 1 minute to elute the DNA.
- 112.7 Add 11ul EB to spin column, incubate for 1 minute, and spin again for total of ~20ul eluate.
- 113 Mix ~20ul sample with 29ul PCR mix (recipe below), 1ul of 25uM IDT8_i5 01-96 primer and H₂O, run PCR for 5 cycles (as below)

A	B	C
ATAC PCR temp	Time	Cycles
72°C	5 min	1 cycle
98°C	30 s	1 cycle
98°C	10 s	5 cycles ~22 min
65°C	30 s	
72°C	60 s	

ATAC initial PCR amplification program setup

A	B
ATAC PCR mix	Volume (ul)
NEBnext 2x PCR mastermix	25x(N+1)
P7 primer 25uM	1x(N+1)
H ₂ O	3x(N+1)
Total	29ul

ATAC PCR mastermix recipe. Don't forget the 1ul of IDT8_i5_xx primer for each reaction!

- 114 qPCR: using 5ul sample and 10ul qPCR mix (NEBnext, Ad1.xx, P7 primer, recipe below), run the qPCR cycle program below, and calculate the number of cycles needed to each 1/3 of the



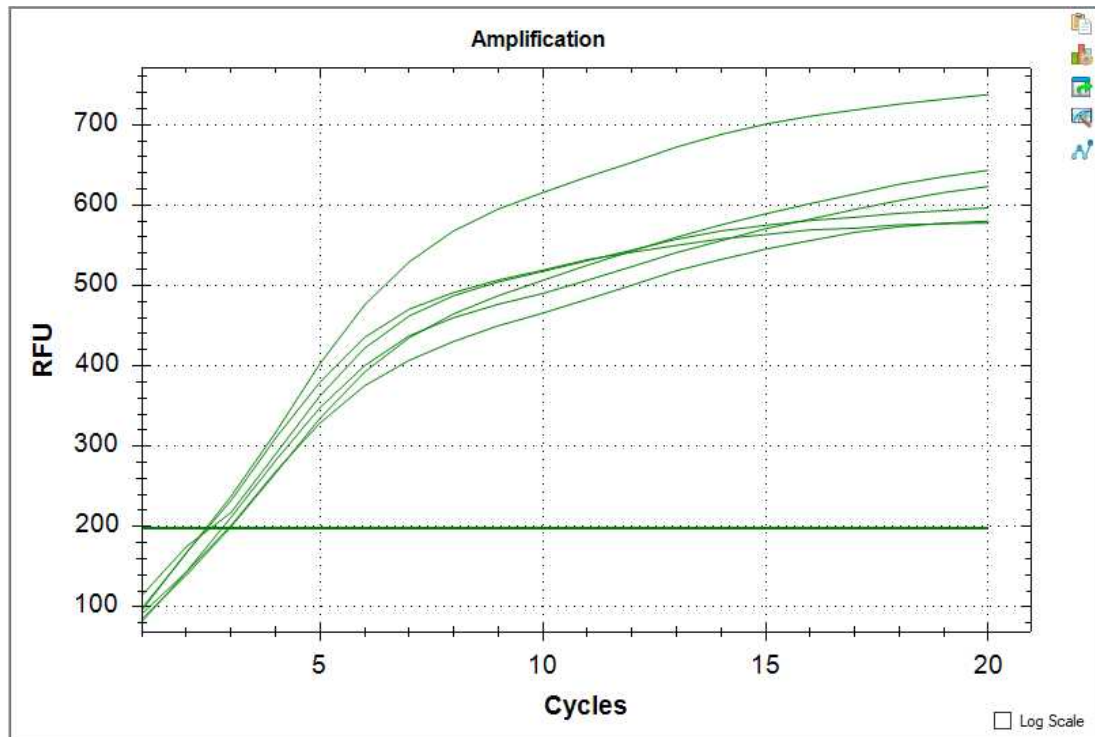
plateau fluorescence (0.33 Ct). Don't forget to use a no template control (NTC) to check for PCR mastermix contamination!

A	B
ATAC qPCR mix	Volume (ul)
NEBnext 2x PCR mastermix	5x(N+2)
Ad1.01 25uM	0.2x(N+2)
P7 primer 25uM	0.2x(N+2)
10x SYBRgreen	0.9x(N+2)
H2O	3.7x(N+2)
Total	10x(N+2)

ATAC qPCR recipe mixture - the 10x SYBR green is made from a 10,000x stock (Thermo), and is stored in small aliquots at -20 degrees C.

A	B	C
qPCR	cycles	
98°C	30 s	1 cycle
98°C	10 s	20 cycles ~1 hour
65°C	30 s	
72°C	60 s	

program for qPCR cycling



example ATAC qPCR curves

We amplified these libraries for 3 cycles - plateau around 600 RFU, 1/3 is 200, see an intersect at 3 cycles!

- 115 Save the qPCR plate to check qPCR product size on an e-gel. This can be combined with checking the cDNA after amplification. Expect a smear with an average size of 400bp. Nucleosomal banding may be present. A smaller MW band at ~270bp is likely non-nucleosomal, and does not correlate with poor quality.



An e-gel of several ATAC libraries

We recommend using an e-gel, 2%, and we run these using the ThermoFisher/Invitrogen Snap apparatus. We use the e-gel 50bp ladder.

- 116 Run additional PCR cycles for the rest of the sample (45ul), using the PCR cycle schema above, where $N=0.33$ Ct cycles. The number of additional cycles expected is complex and based on cell type as well as cell number. Generally libraries with a large number of cells (25000-50000+) will need only 2-4 additional cycles of amplification. Smaller numbers of cells (2000-10000) will need more like 4-10 additional cycles of amplification. These are generalities, your samples may vary, especially for non-diploid samples.
- 117 After PCR reamp, add 1ul of Exonuclease I (NEB cat # M0293L) to each library and place libraries back in thermocycler for "ExoCleanup" program (parameters below) to get rid of excess primer.

ExoCleanup	
37°C	15 mins



80°C	15 mins
4°C	Forever
Total	~30 min

incubation parameters for exonuclease I digestion

- 118 Clean sample using Qiagen MinElute DNA clean up kit (225ul binding buffer, 250ul if qPCR stage was skipped), elute in 10ul EB, then repeat elution with 10ul EB, for a total of ~20ul eluate. Libraries can be stored at -20°C or -80°C.

11.0 cDNA amplification and cleanup (time 3h30m)

- 119 Mix each cDNA sample with 100ul H₂O, magnetize and discard supernatant, wash with 200ul STE; don't resuspend beads.
- 120 Mix 50ul PCR mix with beads, keep in existing tubes

A	B
cDNA PCR mix	Volume (ul)
Kapa Hifi 2x mastermix	25x(N+1)
RNA PCR primer 25uM	0.8x(N+1)
P7 primer 25uM	0.8x(N+1)
H ₂ O	23.4x(N+1)
Total	50x(N+1)

cDNA PCR amplification mix

- 121 Run PCR for 5 cycles

A	B	C
PCR	Cycles	
95°C	3 min	



A	B	C
98°C	20 s	5 cycles ~32 min
65°C	45 s	
72°C	3 min	

cDNA amplification program

122 With samples in magnetic rack, transfer supernatant to new PCR wells.

123 qPCR: for each sub-library, mix 2.5ul sample with 7.5ul qPCR mix, run qPCR protocol below and calculate 0.33 Ct.

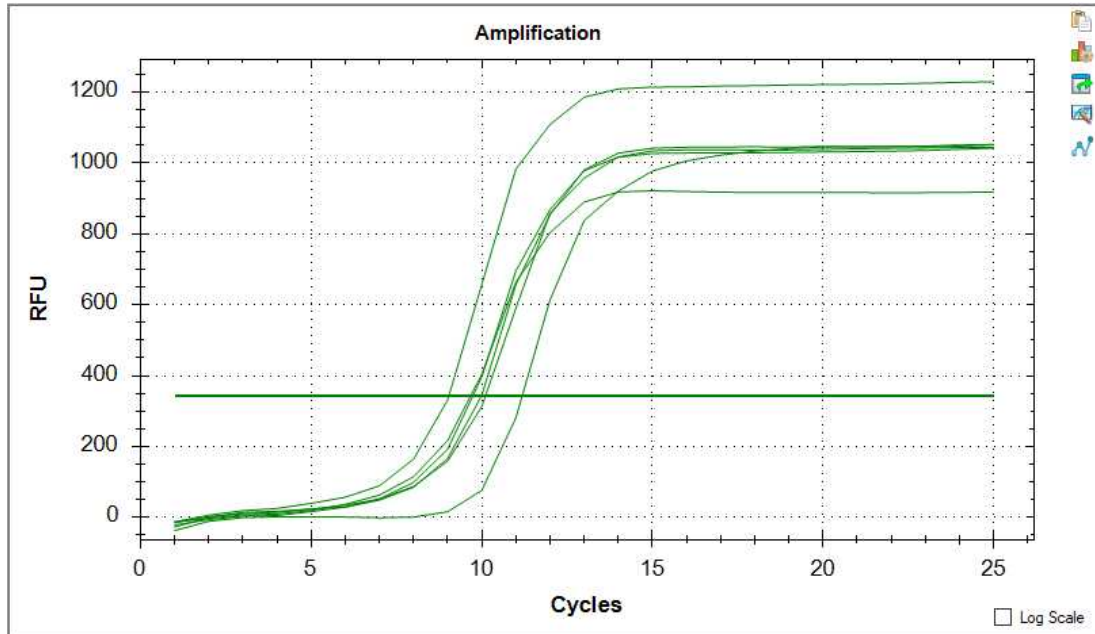
A	B
cDNA qPCR mix	Volume (ul)
Kapa Hifi 2x mix	3.75x(N+2)
RNA PCR primer 25 mM	0.12x(N+2)
P7 primer 25mM	0.12x(N+2)
EVAgreen 20x	0.5x(N+2)
H2O	3.01x(N+2)
Total	7.5x(N+2)

qPCR master mix for cDNA

A	B	C
qPCR		
95°C	3 min	
98°C	20 s	20 cycles ~1h 38m
65°C	20 s	

A	B	C
72°C	3 min	

qPCR cycling conditions for cDNA



example cDNA qPCR curves

Plateau at 1000 RFU, $1/3 = 333$, see an intersect at around 10 cycles.

- 124 Run additional PCR cycles for the rest of the sample (~50ul), $N=0.33$ Ct cycles, see step 106 above. Cycle timing for cDNA is as multifaceted as ATAC, but has the additional complexity of possible nuclease degradation. Generally, the more cells, the lower the cycle number, but this is highly variable across input material. For cultured cell lines, we often see a 50,000+ cell sample will require [6-8] additional cycles of amplification. This increases substantially for nuclei, PBMCs and hematopoietic compartment cells especially - even for samples with 50,000 cells or more, we still generally require at least 10 cycles of amplification.

Below are some generalities, but your results may vary.

for 10000 cells:

low RNA yield (like PBMCs): 14-18 cycles

medium RNA yield: 10-14 cycles

high RNA yield: 8-12 cycles

for 25000 cells:

low RNA yield (like PBMCs) 10-15 cycles

medium RNA yield (post-mortem human nuclei from colon, adrenal glands, others) 8-12 cycles

higher RNA yield (like cultured cells, mouse brain nuclei) 6-10 cycles

124.1 Run the qPCR from the cDNA on an e-gel, an example is below:



ATAC on the left and cDNA on the right (lanes 6-10). The top band in the ladder is 2.5kb

We recommend using an e-gel, 2%, and we run these using the ThermoFisher/Invitrogen Snap apparatus. We use the e-gel 50bp ladder.

- 125 Purify each sample (47.5ul) with 38ul AMPure beads (0.8x, equilibrated to RT) for cell line samples, or 28.5ul AMPure beads (0.6x) for primary samples. Elute cDNA to 20ul EB (10 mM Tris, pH 8.0). Can stop and store samples at -80°C here. Ampure XP protocol is below, as substeps:
- 125.1 Bind samples and beads for 5 minutes at RT
- 125.2 Magnetize for 5 minutes
- 125.3 Wash 2x with 200ul fresh 80% ethanol with samples on magnet (~30s incubate with ethanol on beads)



- 125.4 Dry for ~5 minutes (can place in fume hood to speed things up). Don't let the beads look cracked, they should look glossy, and just at the point of turning matte.
- 125.5 Resuspend in 20.5ul EB
- 125.6 Incubate 5 minutes at RT
- 125.7 Magnetize until clear
- 125.8 Pull 20ul eluate and store in a new tube at -20 degrees.
- 126 ATAC libraries are ready to sequence, cDNA requires further processing (see the next section below). This BioA and tagmentation typically occurs on a third day.



12.0 cDNA tagmentation, amplification, and indexing

- 127 Quantify cDNA concentration using BioAnalyzer using 1ul cDNA diluted 1:10 in water (or 4.5ul water and 0.5ul DNA). Ideally, dilute 50ng cDNA to 5ng/ul in ddH₂O. Note: expect > 50ng cDNA, if cDNA amount is low, you can use as little as 20ng cDNA for tagmentation. Adjust the volume of DNA and H₂O accordingly.
- 128 Use Diagenode Tn5 and assemble transposome by mixing a 1:100 dilution (in dilution buffer) of Tn5, dilution buffer and annealed oligo, and incubate the mix at RT for 30 minutes (see recipe below). See section 1.3 above, steps 17-21 for an overview of annealing the adapter oligos in preparation for this step. **Always perform this step, and the adapter oligo annealing, on the day of tagmentation.**



Transposome	Volume (ul)
1x Tn5	1.25x(N+1)
Dilution buffer	1.25x(N+1)
Annealed tagmentation adapter in glycerol	2.5x(N+1)
Total	5x(N+1)



Assembling the transposome

- 129 Mix 10ul of 5ng/ul cDNA with 10ul H₂O, 25ul TD buffer (recipe below), and 5ul assembled Tn5. Incubate at 55 °C for 5 minutes on a thermal cycler.

2x TD buffer: aliquot and store at -20C	Volume (ul)	2x Final conc
H ₂ O	7700	
1M Tris pH 7.5	200	20mM
1M MgCl ₂	100	10mM
Dimethylformamide (DMF)	2000	20%
Total	10000	

2X TD buffer recipe - many thanks to Rachel Savage for this homegrown recipe!

- 130 Purify tagmented library with Qiagen column (250ul binding buffer), elute twice with 11ul EB (22ul total eluate volume)
- 131 Add 29ul of post tagmentation PCR mix with 1ul tube specific (sample specific) IDT8_i5_xx primer, amplify for 7 cycles.

A	B
Post-tagmentation PCR mix	Volume (ul)
Sample in EB	20 per sample
NEBnext 2x PCR mastermix	25x(N+1)
P7 primer 25mM	1x(N+1)
H ₂ O	3x(N+1)
Total	50x(N+1)



Post tagmentation PCR mix. Don't forget to add 1ul of the i5 index primer!

A	B	C
Tagmentation PCR		
72°C	5 min	
98°C	30 s	
98°C	10 s	7 cycles ~27 min
65°C	30 s	
72°C	60	

tagmentation PCR cycling conditions

- 132 After PCR reamp, add 1ul of Exonuclease I to each library and place libraries back in thermocycler for “ExoCleanup” program to get rid of excess primer.

ExoCleanup	
37°C	15 mins
80°C	15 mins
4°C	Forever
Total	~30 min

exonuclease cycling conditions

- 133 Purify each sample with 35ul AMPure beads (0.7x), elute library to 50ul EB, **then repeat the 0.7x purification and elute in 20ul EB.**

This double elution is to remove small MW fragments (less than ~250bp) which represent tagmentation events where the Illumina R2 is likely destroyed, and there is no actual cDNA retained (the SHARE scaffold takes up around 234bp). Leaving these fragments present drastically reduces the efficiency of RNA sequencing, so we highly recommend you perform these steps as written!



- 133.1 Bind samples and beads for 5 minutes at RT
- 133.2 Magnetize for 5 minutes
- 133.3 Wash 2x with fresh 80% ethanol with samples on magnet
- 133.4 Dry for ~5 minutes (can place in fume hood to speed things up). Don't let the beads look cracked, they should look glossy.
- 133.5 Resuspend in 50ul EB
- 133.6 Incubate 5 minutes at RT
- 133.7 Magnetize until clear
- 133.8 Pull 50ul eluate and store in a new tube.
- 133.9 **Repeat with 35ul AMPure beads**, elute in ~20ul EB and store at -20 degrees C.

12.0 Library quantification and sequencing

- 134 We recommend running all of the cDNA libraries on a bioanalyzer or tapestation to assess the average molecular weight, and the presence/absence of low MW molecules (less than 250bp), which may affect sequencing quality negatively.
- 135 After confirmation of satisfactory bioanalyzer or tapestation traces, quantify all libraries to be sequenced together (in the same independently addressable lane) using a Kapa quantification kit (Roche cat number: 7960140001).

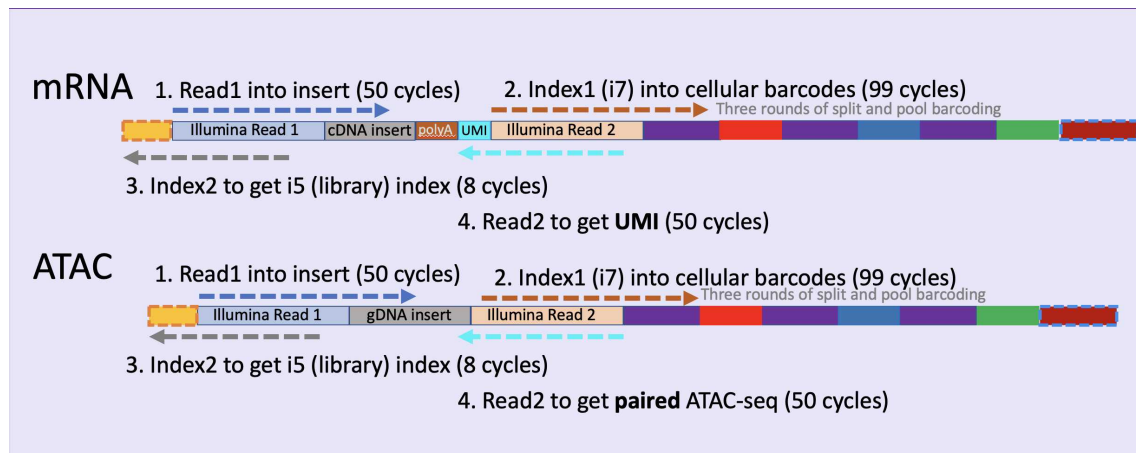
We recommend running reactions in duplicate, using 10ul reactions, loading standards in order of ascending concentration (load NTC, then std6, std5..... std1), and using two dilutions. We first dilute a library 1:1000 in water, then serially dilute 1:1000 again to get a 1:10⁶ dilution. We also dilute the 1:1000 again at 1:100 to get a 1:100,000 dilution. Duplicates of both 1:10⁶ and 1:100K are used to calculate the concentration of the libraries (these values will be different, that is OK).



136 Quantify using the Kapa Quant qPCR kit for Illumina, and pool libraries to 4-10nM concentration. Expect > 5-100nM, average library size of 600bp for cDNA, and 1-2kb for ATAC with minimal adapter dimers.

137 The Epstein lab typically sequences libraries on the NovaSeq X 10B, as it has the lowest cost per base. The sequencing read configuration is as below, for separated(!!) cDNA and ATAC-seq libraries. We find separating the libraries allows for better quality sequencing data, likely due to the differences in molecular weights between cDNA and ATAC.

138



The recommended read structure for SHARE-seq