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Primer ID MiSeq Library Prep for HIV-1 DR and diversity

Forked from [Primer ID MiSeq Library Prep](#)

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ABSTRACT

This is the protocol to prepare Primer ID MiSeq sequencing library. Viral RNA was first extracted using QIAamp viral RNA extraction kit. The block of random nucleotides (Ns) in the cDNA primers served as the Primer ID. The Superscript III kit was used for the cDNA synthesis. We used two rounds of PCR to amplify the cDNA and incorporate Illumina indexed adapters with KAPA2G Robust and KAPA HiFi PCR kits, respectively.

ATTACHMENTS

[amplicon](#)
[protocol_v1.6.docx](#)

GUIDELINES

Viral RNA extraction using QIAamp vRNA mini kit.

Primers (NOTE: HIV-1 ENV REGION AS AN EXAMPLE, SUBSTITUTE THE BLUE Sequences in the forward primer and cDNA primer)

OPEN ACCESS

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Protocol status: Working
We use this protocol and it's working

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26884

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V1F(forward) 5'-3'	GCCTCCCTCGCGCCATCAGAGATGTGTATAAGAGACAGNNNNTTATGGGATC AAAGCCTAAAGCCATGTGTA
BV3R Uni (cDNA Primer) 5'-3'	GTGACTGGAGTTCAGACGTGTGCTCTTCCGATCTNNNNNNNNNCAAGTCCATT TGCTCTACTAATGTTACAATGTGC
Universal Adapter	AATGATACGGCGACCACCGAGATCTACACGCCTCCCTCGCGCCATCAGAGATGT G
Indexed Adapter	CAAGCAGAAGACGGCATACGAGATNNNNNNGTGACTGGAGTTCAGACGTGT GCTC
ADPT_2a	GTGACTGGAGTTCAGACGTGTGCTC

Note: Primer ID primer and forward primer use random bases. Indexed primers have 24 fixed barcodes.

Table of Indexed Primers

Indexed Primer	Index	Index Sequence	Sequence
PCR Primer, Index 1	1	ATCACGA	CAAGCAGAAGACGGCATACGAGATCGTGATGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 2	2	CGATGTA	CAAGCAGAAGACGGCATACGAGATACATCGGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 3	3	TTAGGCA	CAAGCAGAAGACGGCATACGAGATGCCTAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 4	4	TGACCAA	CAAGCAGAAGACGGCATACGAGATGGGTCAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 5	5	ACAGTGA	CAAGCAGAAGACGGCATACGAGATCACTGTGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 6	6	GCCAATA	CAAGCAGAAGACGGCATACGAGATATTGGCGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 7	7	CAGATCA	CAAGCAGAAGACGGCATACGAGATGATCTGGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 8	8	ACTTGAA	CAAGCAGAAGACGGCATACGAGATCAAGTGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 9	9	GATCAGA	CAAGCAGAAGACGGCATACGAGATCTGATCGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 10	10	TAGCTTA	CAAGCAGAAGACGGCATACGAGATAAGCTAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 11	11	GGCTACA	CAAGCAGAAGACGGCATACGAGATGTAGCCGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 12	12	CTTGTA	CAAGCAGAAGACGGCATACGAGATTACAAGGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 13	13	TCCATAA	CAAGCAGAAGACGGCATACGAGATTATGGAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 14	14	GTACTAA	CAAGCAGAAGACGGCATACGAGATTAGTACGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 15	15	ACAGTAA	CAAGCAGAAGACGGCATACGAGATTACTGTGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 16	16	CTCATGA	CAAGCAGAAGACGGCATACGAGATCATGAGGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 17	17	ACGATAA	CAAGCAGAAGACGGCATACGAGATTATCGTGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 18	18	TGCAGAA	CAAGCAGAAGACGGCATACGAGATCTGCGAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 19	19	TTCATAA	CAAGCAGAAGACGGCATACGAGATTATGAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 20	20	TGCTGTA	CAAGCAGAAGACGGCATACGAGATACAGCAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 21	21	TATCACA	CAAGCAGAAGACGGCATACGAGATGTGATAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 22	22	TGGATAA	CAAGCAGAAGACGGCATACGAGATTATCCAAGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 23	23	CGCATTA	CAAGCAGAAGACGGCATACGAGATAATGCGGTGACTGGAGTTCAGACGTGTGCTC
PCR Primer, Index 24	24	GCCTTAA	CAAGCAGAAGACGGCATACGAGATTAAGCGGTGACTGGAGTTCAGACGTGTGCTC

MATERIALS

MATERIALS

- ☒ QIAamp vRNA mini kit **Qiagen Catalog #51304**
- ☒ Agencourt RNAClean XP **Beckman Coulter Catalog #A63987**
- ☒ 70% ethanol **Contributed by users**
- ☒ DNase-free water **Contributed by users**
- ☒ AmpureXP PCR cleanup kits **Beckman Coulter Catalog #A63880**
- ☒ QIAquick gel extraction kit **Qiagen Catalog #28704**
- ☒ Qubit dsDNA BR Assay kit **Invitrogen - Thermo Fisher Catalog #Q32850**

Prepare Primer Mix (Optional, only for multiplexed Primer ID library prep)

- For multiplexing sequencing, first, prepare Primer Mix.
Example (For HIV drug resistance pipeline).

Regions	DR cDNA primer	DR F primer
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PR	R261 4_PID	F2163 AD
RT	R328 4_PID 11	F2620 _AD
IN	R475 2_PID 11	F4383 _AD
V3	R720 9_PID 11	V1F- AD

Make 10 μ M primer mix: mix 10 μ L of each primer in one set and 60 μ L of dH₂O.

Primer Tables

prime r	seque nce (5'-3')
R261 4_PID	GTGA CTGG AGTT CAGA CGTG TGCT CTTC CGAT CTNN NNN NNN NCAG TTTA ACTT TTGG GCCA TCCA TTCC
R328 4_PID 11	GTGA CTGG AGTT CAGA CGTG TGCT CTTC CGAT CTNN NNN NNN NNN CAGT CACT ATAG GCTG TACT GTCC ATTT ATC
R475 2_PID 11	GTGA CTGG AGTT CAGA CGTG TGCT CTTC CGAT CTNN NNN NNN NNN ATCG AATA CTGC CATT TGTA CTGC
R720 9_PID 11	GTGA CTGG AGTT CAGA CGTG TGCT CTTC CGAT CTNN NNN NNN NNN CAGT CCAT TTTG CTYT AYTR ABVT TACA ATRT GC

F2163 AD	GCCT CCCT CGCG CCAT CAGA GATG TGTA TAAG AGAC AGNN NNTC AGAG CAGA CCAG AGCC AACA GCCC CA
F2620 _AD	GCCT CCCT CGCG CCAT CAGA GATG TGTA TAAG AGAC AGNN NNGG CCAT TGAC AGAA GAAA AAAT AAAA GC
F4383 _AD	GCCT CCCT CGCG CCAT CAGA GATG TGTA TAAG AGAC AGNN NNAA AAGG AGAA GCCA TGCA TG
V1F_ AD	GCCT CCCT CGCG CCAT CAGA GATG TGTA TAAG AGAC AGNN NNTT ATGG GATC AAAG CCTA AAGC CATG TGTA

"N" in this Primer table is a random nucleotide.

P1	AATG ATAC GGCG ACCA CCGA GATC TACA CGCC TCCC TCGC GCCA TCAG AGAT GTG
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Index ed Adapt er	CAAG CAGA AGAC GGCA TACG AGAT NNN NNN GTGA CTGG AGTT CAGA CGTG TGCT C
ADPT _2a	GTGA CTGG AGTT CAGA CGTG TGCT C

The 6 "N"s are not random nucleotides. They are a set of 24 pre-designed indexed sequences.

2 Mix well.

cDNA synthesis

3 Pipette the following components into a 0.5 ml RNase-free tube:

μl/tube		[stock]	[final]	[mastermix]
2.0	dNTP Mix	10 mM each	0.5	
1.0	cDNA primer	10 μM	0.25 μM	
23.0*	RNA template			
26.00	Total volume			

* The input volume of vRNA depends on the viral loads. We recommend 1,000 to 20,000 copies, but it can work with samples as little as 200 copies. If copy number unknown, use the maximum volume of 23 ul.

🧪 2 μL dNTPMix

🧪 1.0 μL cDNA primer

🧪 23.0 μL RNA template

4 Place tube in 65°C heat block for 3-5'.

🕒 00:05:00 65°C heat block

5 Place the tube on ice for 1'.

🕒 00:01:00 on ice

6 Add the following components:

μl/tube		[stock]	[final]	[mastermix]
8.0	5x buffer	5x	1x	
2.0	DTT	100 mM	5	
2.0	RNaseOUT	40 u/μl	2	
2.0	SSIII RT	200 u/μl	10	

14.0	Per tube			
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8 µL 5x buffer

2 µL DTT

2 µL RNaseOUT

2 µL SSII RT

- 7 Mix and incubate at 50°C for 1 hr.

01:00:00 Incubation at 50°C

- 8 Increase to 55°C and incubate for 1 hr.

01:00:00 Incubation at 55°C

- 9 Inactivate SSIII RT by heating at 70°C for 15'.

00:15:00 Heating at 70°C

- 10 To each tube, add 1 µL RNase H, incubate at 37°C for 20'.

0.5 µL RNase H

00:20:00 Incubation at 37°C

Purification

- 11 Purify cDNA using Agencourt RNAClean XP.

Resuspend the beads and take an aliquot out. Keep at room temperature for at least **30 minutes** before use. (Should be in 1ml aliquots)

00:30:00 Room temperature

- 12 Transfer the cDNA reactions into 1.7 mL RNase-free tubes.

- 13 Resuspend the beads (Vortex). Add **28 µl of beads to 40 µl cDNA** Agencourt RNAClean XP beads to each cDNA reaction.

28 µL beads

40 µL cDNA

- 14 Mix the Agencourt RNAClean XP and sample thoroughly by pipette mixing 15 times. No **vortexing**. Let the tube incubate at room temperature for **20 minutes** before proceeding to the next step.

00:20:00 Room temperature

- 15 Place the tube onto the magnetic tube rack for **5 minutes** to separate the beads from solution.

00:05:00 Magnetic tube rack

- 16 Slowly aspirate the cleared solution from the tube and discard. This step should be performed while the tube is situated on the rack. Do not disturb the magnetic beads, which have formed a spot on the side of the tube.

- 17 Dispense 400 µL of 70% ethanol into the tube and incubate for **30 seconds** at room temperature. Aspirate out the ethanol and discard. (1/3)

It is important to perform this step with the tube situated **on the rack**. **Do not disturb the separated magnetic beads**. Be sure to remove all of the ethanol from the bottom of the well as it may contain residual contaminants.

⌚ 00:00:30 Incubation at room temperature

- 18 Dispense 400 µL of 70% ethanol into the tube and incubate for **30 seconds** at room temperature. Aspirate out the ethanol and discard. (2/3)

It is important to perform this step with the tube situated **on the rack**. **Do not disturb the separated magnetic beads**. Be sure to remove all of the ethanol from the bottom of the well as it may contain residual contaminants.

⌚ 00:00:30 Incubation at room temperature

- 19 Let the reaction tube air-dry **10 minutes** on the rack with the cap open. The tube(s) should air-dry until the last visible traces of ethanol evaporate. Over drying the sample may result in a lower recovery.

⌚ 00:10:00 Air-drying

- 20 Remove the tube from the rack and resuspend beads in 40 µl DNase-free water by pipetting up and down. Place tube back on the rack and leave for **3 minutes**.

🧪 40 µL DNase-free water

⌚ 00:03:00 Magnetic rack

- 21 Pipette the eluant from the tube while it is situated **on the magnetic tube rack**. Transfer the eluant into the tube with 28 µl of RNACleanup XP beads.

- 22 Remove the tube from the rack and resuspend beads in 24 µl DNase-free water by pipetting up and down. Place tube back on the rack and leave for **3 minutes**.

🧪 24 µL DNase-free water

⌚ 00:03:00 Magnetic rack

- 23 Pipette the eluant from the tube while it is situated **on the magnetic tube rack**.

PCR 1

- 24 Complete thaw and vortex KAPA reagents (except for enzyme) before use.

	µl/tube		[stock]	[final]	[mastermix]
	10.0	5x Buffer A	5x	1x	
	10.0	Enhancer	5x	1x	
	1.0	dNTPs	10 mM	0.2 mM	
	2.5	Forward primer	10 µM	0.5 µM	
	2.5	ADPT _{2a}	10 µM	0.5 µM	
	0.5	KAPA Robust polymerase	5 U/µl	2.5 U	
	23.5	Template cDNA			

Prepare mastermix in cold box and use repeater pipette to add to each tube in the cold box. Add template cDNA to each tube and pipette up and down to mix.

🧪 10 µL 5x Buffer A

🧪 10 µL Enhancer

🧪 1 µL dNTPs

🧪 2.5 µL Forward primer

2.5 µL ADPT_2a

0.5 µL KAPA Robust polymerase

23.5 µL Template cDNA

26 Cycle

95°C	1 min
95°C	15 s
58°C	1 min
72°C	30 s
15-25 cycles	
72°C	3 min
4°C	On hold

Purification PCR products

27 Purify PCR products using AmpureXP PCR cleanup kits.

Vortex the 1 ml aliquot and remove the needed volume. Keep at room temperature for at least **30 minutes** before use.

00:30:00 Room temperature

28 Transfer the PCR1 reactions into 1.7 mL RNase-free tubes.

29 Resuspend the beads. Add **40 µl (Ratio: 0.6 – 0.8: 1, 36µl – 48µl)** Ampure XP beads to each cDNA reaction.

40 µL Ampure XP beads

30 Mix the Ampure XP and sample thoroughly by **vortexing**. Let the tube incubate at room temperature for **5 minutes** before proceeding to the next step (incube off the rack).

00:05:00 Incubation at room temperature

31 Place the tube onto the magnetic tube rack for **5 minutes** to separate the beads from solution.

00:05:00 Incubation magnetic rack

32 Slowly aspirate the cleared solution from the tube and discard. This step should be performed while the tube is situated on the rack. Do not disturb the magnetic beads, which have formed a spot on the side of the tube.

33 Dispense 500 µL of 70% ethanol into the tube and incubate for **30 seconds** at room temperature. Aspirate out the ethanol and discard. (1/2)

Note

It is important to perform this step with the tube situated **on the rack**. **Do not disturb the separated magnetic beads**. Be sure to remove all of the ethanol from the bottom of the well as it may contain residual contaminants.

500 µL 70% ethanol

00:00:30 Incubation at room temperature

34 Dispense 500 µL of 70% ethanol into the tube and incubate for **30 seconds** at room temperature. Aspirate out the ethanol and discard. (2/2)

Note

It is important to perform this step with the tube situated **on the rack**. **Do not disturb the separated magnetic beads**. Be sure to remove all of the ethanol from the bottom of the well as it may contain residual contaminants.

🧪 500 µL 70% ethanol

🕒 00:00:30 Incubation at room temperature

- 35 Let the reaction tube air-dry **10 minutes** on the rack with the cap open. The tube(s) should air-dry until the last visible traces of ethanol evaporate. Over drying the sample may result in a lower recovery.

🕒 00:10:00 Air-drying

- 36 Remove the tube from the rack and resuspend beads in 50 µl DNase-free water by pipetting up and down. Place tube back on the rack and leave for **3 minutes**.

🧪 50 µL DNase-free water

🕒 00:03:00 Magnetic rack

- 37 Pipette the 45 µl eluant from the tube while it is situated **on the magnetic tube rack**.

PCR 2

- 38 Complete thaw and vortex KAPA reagents (except for enzyme) before use.

39

µl/tube		[stock]	[final]	[mastermix]
5.0	5x KAPA HiFi Fidelity Buffer	5x	1x	
1.0	dNTP Mix	10 mM	0.4 mM	
1.0	Uni Adapter (ADPT_P1)	10 µM	0.4 µM	
0.5	KAPA HiFi polymerase	1 U/µl	0.5 U	
1.0	Indexed Adapter	10 µM	0.4 µM	
2.0	Purified template DNA			
14.5	Water			

Prepare mastermix in cold box and use repeater pipette to add to each tube in the cold box. Add Indexed Adapter to each tube.

🧪 5 µL 5x KAPA HiFi Fidelity Buffer

🧪 1 µL dNTP Mix

🧪 1 µL Uni Adapter (ADPT_P1)

🧪 0.5 µL KAPA HiFi polymerase

🧪 1 µL Indexed Adapter

🧪 2 µL Purified template DNA

🧪 14.5 µL Water

- 40 **Cycle** (PCR machine #5 SZ -> ILM2):

95°C	2 min
98°C	20 s
63°C	15 s
72°C	30 s
25 - 35 cycles	
72°C	3 min
4°C	On hold

Gel Purification and quantification

- 41 Before gel purification, run 2 µl products on 1% agarose gel to check the bands.
- 42 **Gel purification.** (Qiagen QIAquick gel extraction kit)
Run 2nd round PCR products on 1.2% agarose gel. E = 4 V/cm, T = 60 min.
- 43 Excise DNA fragment.
- 44 Weight the gel; add 3 volume of Buffer QG to 1 volume of gel.
- 45 Incubate at 50 °C for **10 minutes** to completing dissolve. Vortex every **2-3 minutes** to help dissolve.
00:10:00 Incubation at 50 °C
00:02:30 Vortex
- 46 Check the color of gel solution (should be yellow, otherwise add 10 µl 3M sodium acetate).
- 47 Place **MinElute** column, apply the sample to the column and centrifuge for **1 minute**.
00:01:00 Centrifugation
- 48 Add 500 µl buffer QG and centrifuge for **1 minute**.
500 µL Buffer QG
00:01:00 Centrifugation
- 49 Add 0.75 ml buffer PE, **incubate for 5 minutes** at room temperature, centrifuge for **1 minute**.
00:05:00 Incubation at room temperature
00:01:00 Centrifugation
- 50 Discard the fluid, centrifuge for additional **3 minutes**.
00:03:00 Centrifugation

- 51 Put the column in a new 1.7 ml tube, add **10** μ l buffer EB. Stand for **4 minutes**, centrifuge for **2 minutes**.
00:04:00 Stand
- 52 Quantification using Invitrogen Qubit dsDNA BR Assay kit. See Qubit dsDNA BR assay protocol. **Don't use Nanodrop to quantify!**
- 53 After quantification, pool libraries in equal amount. Use AMPure XP beads to purify pooled libraries (2:3, two washes), elute in 20 to 30 μ L Elution Buffer.
- 54 Quantify the pooled library using Qubit dsDNA BR assay kit. Check the quality of the library by Agilent Bioanalyzer or Bio-rad Experion. Repeat purification if primer dimer is present (~200bp).