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HEK3/LINC01509 Library Preparation

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ABSTRACT

Library preparation for HEK3 recording site (LINC01509) using PCR and digestion.

MATERIALS

- NGS primer F ("CTGGCCTGGGTCAATCCTTG")
- NGS primer R ("CCCAGCCAAACTTGTCAACC")
- Q5 HiFi polymerase 2X master mix (NEB)
- BclI (NEB)
- LMW agarose
- SYBR Safe gel stain
- Generuler 1kb PLUS
- SPRI beads (Nucleomag)

OPEN ACCESS



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Protocol status: In development
We are still developing and optimizing this protocol

Created: May 24, 2023

HEK293T Cell Lysis

1 Make up **1:50** Thermolabile Proteinase K (NEB) in Viagen (cell).

2 Resuspend cells in **25 μ L** of Viagen/Proteinase K.



Quickly transfer cells into a PCR strip to avoid viscosity.

3 Perform lysis in a PCR machine:

1h 30m



37 °C **01:00:00**

55 °C **00:30:00**

On ice

Store lysates at **-20 °C** in the pre-PCR freezer.

PCR Amplification of Genomic DNA

4 Make up a **100 μ L** PCR reaction per sample (Q5) using **10 μ L** of cell lysate.



4.1 **98 °C** **00:00:30**

30s

4.2 **98 °C** **00:00:05**

35s

🌡️ 68 °C ⌚ 00:00:10
🌡️ 72 °C ⌚ 00:00:20

4.3 ➡️ go to step #4.2 24 X

4.4 🌡️ 72 °C ⌚ 00:02:00

2m

SPRI Bead Concentration

5 Purify amplicons by **1X SPRI** bead cleanup, eluting in 🧴 10 µL of water.



Digestion of Zero-Length Recordings

6 Digest zero-length recordings with 🧴 5 µL BclI in **50 uL** volume:

1h 20m



🌡️ 37 °C ⌚ 01:00:00
🌡️ 65 °C ⌚ 00:20:00
🌡️ On ice

Gel Separation and Extraction of Recordings

7 Perform gel electrophoresis in **4% LMP agarose** at **3 V/cm** for ⌚ 01:00:00 .



1h

8 Extract the region from **300 bp to 500 bp** and elute in 🧴 12 µL of water.



Store gel products at 🌡️ -20 °C in the post-PCR freezer.







Indexing PCR of Recombinant DNA

9 Using  1 μL of gel product, perform indexing PCR in  12 μL volume with **Nextera** primers.




9.1  98 °C  00:00:30

30s

9.2  98 °C  00:00:05
 67 °C  00:00:10
 72 °C  00:00:20


35s

9.3  go to step #9.2 9 X

9.4  72 °C  00:02:00

2m

SPRI Bead Purification

10 Purify amplicons by **1X SPRI** bead cleanup, eluting in  10 μL of water.



D1000 Tapescreen

11 Take  1 μL of each indexed amplicon to D1000 Tapescreen.



Expected result

Recordings should be above $158 + 67$ (adaptors) + 69 (indexes) = **294 bp**.

- 12 Pool indexed amplicons according to desired relative molarities for sequencing.