

lubio science



2020 Product Catalogue



Reagents for NGS Library Preparation

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Contact us for ordering or additional information info@lubio.ch - www.lubio.ch LubioScience GmbH - Baumackerstrasse 24 - 8050 Zürich - 041 417 02 80

Overview of Vazyme

Vazyme: InoVation in Enzyme Technology

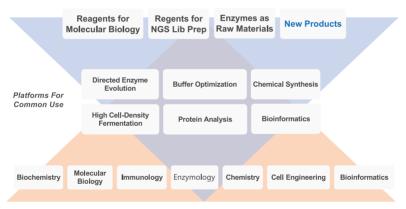
With the faith of "InnoVation in Enzyme Technology", Vazyme Biotech Co., Ltd. has passionately focused on developing enzyme and antibody technologies and products for years. The Vazyme Biotech is now staffed by more than 1,000 employees. The headquarter is located in Nanjing of China with a R&D / manufacturing base that covers 25,000 m² and a GMP workshop of 4,000 m². Vazyme has developed a powerful sales network in China and is expanding into international markets.



Vazyme Technologies and Products

With years of experience, Vazyme has developed six technology platforms that can be commonly used for R&D and manufacturing, including (1) directed enzyme evolution, (2) buffer optimization, (3) chemical synthesis, (4) high cell-density fermentation, (5) protein analysis, and (6) bioinformatics.

Based on the above platforms, Vazyme now provides a variety of products, solutions, and services, which fall into three major product lines, including (1) solutions for molecular biology research, (2) solutions for Next-Generation Sequencing (NGS) library preparation, and (3) enzymes as raw materials for industrial use.



Developing Technologies to Improve Human Health

Fascinated by the enzyme and antibody technologies, we regard enzymes and antibodies as the key factor of the biotechnology industry. Vazyme's vision is to develop technology to improve human health.

Expert I for Expert I

2020 Vazyme Product Catalogue

Reagents for NGS Library Preparation

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DNA-Seq

DNA Library Prep (for Illumina®)

| | Application | Product Name | Cat. No.# | Size |
|-----|--|--|-------------------------|-----------------------|
| | | TruePrep DNA Library Prep Kit V2 for IIIumina® (50 ng) | TD501 | 24 rxn/96 rxn |
| HOT | Transposase-based DNA | TruePrep DNA Library Prep Kit V2 for IIIumina® (5 ng) | TD502 | 24 rxn/96 rxn |
| | Lib Prep Kits | TruePrep DNA Library Prep Kit V2 for Illumina® (1 ng) | TD503 | 24 rxn/96 rxn |
| | | TruePrep Homo-N7 DNA Library Prep Kit for IIIumina® | TD511/TD512/TD513 | 96 rxn |
| | Single-Indexed Adapters | TruePrep Index Kit V4 for Illumina® | TD204/TD205/TD206/TD207 | 192 rxn |
| | Dual-Indexed Adapters | TruePrep Index Kit V2 / V3 for Illumina® | TD202/TD203 | 96 kinds/384 kinds |
| HOT | | VAHTS Universal DNA Library Prep Kit for Illumina® V3 | ND607-01/02 | 24 rxn/96 rxn |
| | Universal DNA Lib Prep Kits | VAHTS Universal DNA Library Prep Kit for Illumina® V3 (PCR-Free) | ND607-03/04 | 24 rxn/96 rxn |
| New | | VAHTS Universal DNA Library Prep Kit for Illumina® V5 | ND610-01/02 | 24 rxn/96 rxn |
| | DNA Lib Prep Kits for Enzymatic Fragmentation | VAHTS Universal Plus DNA Library Prep Kit for Illumina® | ND617-01/02 | 24 rxn/96 rxn |
| | FFPE DNA Lib Prep Kits | VAHTS Universal Pro DNA Library Prep Kit for Illumina $^{\otimes}$ | ND608-01/02 | 24 rxn/96 rxn |
| New | ssDNA Lib Prep Kits | VAHTS ssDNA Library Prep Kit for Illumina® | ND620-01/02 | 24 rxn/96 rxn |
| | MP DNA Lib Prep Kits | VAHTS Mate Pair Library Prep Kit for IIIumina® | ND104-01 | 48 rxn |
| | Single-Indexed Adapters | VAHTS DNA Adapters Set1 / Set 2 for Illumina® | N801/N802-01/02 | 10 µl each/40 µl each |
| | Single-Indexed Adapters | VAHTS DNA Adapters Set 3 - Set 6 for Illumina® | N805/N806/N807/N808 | 20 µ l each |
| | Dual-Indexed Adapters | VAHTS Multiplex Oligos Set 4 / Set 5 for Illumina® | N321/N322 | 192 rxn |
| | UMI Adapters | VAHTS Dual Index UMI DNA Adapters Set 1 - Set 4 for Illumina® | N331/N332/N333/N334 | 20 μ l each |
| | A sector of the December of the | VAHTS AmpSeq Library Prep Kit V2 | NA201-01/02 | 24 rxn/96 rxn |
| New | Amplicon Lib Prep Kits | VAHTS AmpSeq Library Prep Kit V3 | NA210-01/02 | 24 rxn/96 rxn |
| | Amplicon Lib Prep Adapters | VAHTS AmpSeq Adapters 1 - 24 for Illumina® | NA111-01/02 | 12 ×10 rxn |
| | Ampicon cio i Tep Adapters | VAHTS AmpSeq Adapters 25 - 96 for Illumina® | NA111-03/04/05 | 24 ×10 rxn |
| | Cancer Panels | VAHTS AmpSeq Cancer HotSpot Panel | NA102-01 | 24 rxn |
| | | | | |

DNA Library Prep (for Ion Torrent®)

| | Application | Product Name | Cat. No.# | Size |
|-----|-----------------------------|---|----------------|---------------|
| | Universal DNA Lib Prep Kits | VAHTS Universal DNA Library Prep Kit for Ion Torrent® | ND701-01/02 | 24 rxn/96 rxn |
| New | Amplicon Lib Prep Kits | VAHTS AmpSeq Library Prep Kit V2 | NA201-01/02 | 24 rxn/96 rxn |
| | | VAHTS AmpSeq Library Prep Kit V3 | NA210-01/02 | 24 rxn/96 rxn |
| | Amplicon Lib Prep Adapters | VAHTS AmpSeq Adapters 1 - 24 for Ion Torrent® | NA121-01/02 | 12 ×10 rxn |
| | Amplicon Lib Frep Adapters | VAHTS AmpSeq Adapters 25 - 96 for Ion Torrent® | NA121-03/04/05 | 24 ×10 rxn |

DNA Library Prep (for MGI[®])

| Application | Product Name | Cat, No,# | Size |
|--|---|--------------|-----------------------|
| Universal DNA Lib Prep Kits | VAHTS Universal DNA Library Prep Kit for MGI® | NDM607-01/02 | 24 rxn/96 rxn |
| Oniversal Drive Lib Trep Mis | VAHTS Universal DNA Library Prep Kit for MGI® V5 | NDM610-01/02 | 24 rxn/96 rxn |
| DNA Lib Prep Kits for Enzymatic Fragmentation | VAHTS Universal Plus DNA Library Prep Kit for MGI® | NDM617-01/02 | 24 rxn/96 rxn |
| FFPE DNA Lib Prep Kits | VAHTS Universal Pro DNA Library Prep Kit for MGI [®] | NDM608-01/02 | 24 rxn/96 rxn |
| Single-Indexed Adapters | VAHTS DNA Adapters Set 8 for MGI® | NM108-01/02 | 10 µl each/40 µl each |
| Dual-Indexed UMI Adapters | VAHTS Dual UMI Adapters for MGI® | NM301-01/02 | 10 µl each/40 µl each |
| Circularization Kit | VAHTS Circularization Kit for MGI® | NM201-01/02 | 16 rxn/48 rxn |
| Amplification Module | VAHTS HiFi Amplification Mix for MGI® | NM616-01/02 | 24 rxn/96 rxn |
| Amplicon Lib Prep Kits | VAHTS AmpSeq General Library Prep Kit for MGI® | NAM203-01/02 | 24 rxn/96 rxn |

Vazyme Biotech. Co. Ltd. Innovation in Enzyme Technology



Modules for DNA Library Prep

| Application | Product Name | Cat. No.# | Size |
|----------------------------|---|-------------|---------------|
| TurePrep Lib Amplification | TruePrep Amplify Enzyme | TD601 | 96 rxn |
| FFPE Repair | VAHTS DNA Damage Repair Kit | N208-01/02 | 24 rxn/96 rxn |
| Enzymatic Fragmentation | VAHTS Universal Plus Fragmentation Module | N209-01/02 | 24 rxn/96 rxn |
| | VAHTS Universal End preparation Module for Illumina® | N203-01/02 | 24 rxn/96 rxn |
| DNA Lib Prep Modules | VAHTS Universal Adapter Ligation Module for Illumina® | N204-01/02 | 24 rxn/96 rxn |
| | VAHTS HiFi Amplification Mix | N616-01/02 | 24 rxn/96 rxn |
| AmpSeq Multi-PCR | VAHTS AmpSeq Multi-PCR Module V2 | NA205-01/02 | 24 rxn/96 rxn |
| T4 DNA Polymerase | T4 DNA Polymerase | N101-01 | 2,000 U |
| T4 Polynucleotide Kinase | T4 Polynucleotide Kinase | N102-01 | 10,000 U |
| T4 DNA Ligase (Rapid) | T4 DNA Ligase (Rapid) | N103-01 | 600,000 U |
| Klenow DNA Polymerase | DNA Polymerase Klenow Fragment | N104-01 | 5,000 U |
| Klenow DNA Polymerase | DNA Polymerase I Klenow Fragment exo- | N105-01 | 5,000 U |
| Phi29 DNA Polymerase | Phi29 MAX DNA Polymerase | N106-01/02 | 250 U/1,250 U |
| Amplification Module | Phanta Uc Super-Fidelity DNA Polymerase for Library Amplification | P507-01/02 | 100 U/500 U |
| | | | |

RNA-Seq

RNA Library Prep (for Illumina[®])

| | Application | Product Name | Cat, No,# | Size |
|-----|---|--|---------------------|-----------------------|
| | Total RNA Lib Prep Kits (rRNA Depletion) | VAHTS Total RNA-Seq (H/M/R) Library Prep Kit for Illumina® | NR603-01/02 | 24 rxn/96 rxn |
| НОТ | Ultra Fast & Universal RNA Lib | VAHTS Universal V6 RNA-Seq Library Prep Kit for Illumina® | NR604-01/02 | 24 rxn/96 rxn |
| New | Prep Kits | VAHTS Universal V8 RNA-Seq Library Prep Kit for Illumina® | NR605-01/02 | 24 rxn/96 rxn |
| | Single-Indexed Adapters | VAHTS RNA Adapters Set 1 / Set 2 for Illumina® | N803/N804-01/02 | 10 μl each/40 μl each |
| | engle indexed radptere | VAHTS RNA Adapters Set 3 - Set 6 for Illumina® | N809/N810/N811/N812 | 20 µl each |
| | Dual-Indexed Adapters | VAHTS RNA Multiplex Oligos Set 1 / Set 2 for Illumina® | N323 / N324 | 192 rxn each |
| | Smal RNA Lib Prep Kits | VAHTS Small RNA Library Prep Kit for Illumina® | NR801-01/02 | 24 rxn/96 rxn |
| | Small RNA Lib Prep Adapters | VAHTS Small RNA Index Primer Kit for Illumina® | N813/N814/N815/N816 | 48 rxn each |

RNA Library Prep (for MGI[®])

| Application | Product Name | Cat. No.# | Size |
|---|---|--------------|-----------------------|
| Non-Stranded mRNA Lib Prep Kits | VAHTS mRNA-Seq V3 Library Prep Kit for MGI® | NRM611-01/02 | 24 rxn/96 rxn |
| Total RNA Lib Prep Kits (rRNA Depletion) | VAHTS Total RNA-Seq (H/M/R) Library Prep Kit for MGI® | NRM603-01/02 | 24 rxn/96 rxn |
| Ultra Fast & Universal RNA Lib | VAHTS Universal V6 RNA-Seq Library Prep Kit for MGI® | NRM604-01/02 | 24 rxn/96 rxn |
| Prep Kits | VAHTS Universal V8 RNA-Seq Library Prep Kit for MGI® | NRM605-01/02 | 24 rxn/96 rxn |
| Single-Indexed Adapters | VAHTS RNA Adapters Set 8 for MGI® | NM208-01/02 | 10 µl each/40 µl each |

Modules for RNA Library Prep

| | Application | Product Name | Cat. No.# | Size |
|-----|---------------------|---|------------|-----------------|
| | mRNA Capture Beads | VAHTS mRNA Capture Beads | N401-01/02 | 24 rxn/96 rxn |
| | RNA Fragmentation | VATHS 2 × Frag / Prime Buffer | N402-01 | 96 rxn |
| _ | | Ribo-off rRNA Depletion Kit (Human / Mouse / Rat) | N406-01/02 | 24 rxn / 96 rxn |
| НОТ | rRNA Depletion Kits | Ribo-off rRNA Depletion Kit (Bacteria) | N407-01/02 | 12 rxn / 24 rxn |
| New | | Ribo-off rRNA Depletion Kit (Plant) | N409-01/02 | 12 rxn / 24 rxn |

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Epigenetics

| Application | Product Name | Cat. No.# | Siz |
|----------------------------|---|----------------|----------------|
| Methylation Bisulfite Kit | EpiArt DNA Methylation Bisulfite Kit | EM101-01/02 | 50 rxn/200 rx |
| Amplification Kit | 2 × EpiArt HS Taq Master Mix | EM201-01/02/03 | 1 ml/5 ml/15 n |
| 7 mpinouon ru | 2 × EpiArt HS Taq Master Mix (Dye Plus) | EM202-01/02/03 | 1 ml/5 ml/15 r |
| Methylation Lib Prep Kit | EpiArt DNA Methylation Library Kit | NE101-01/02 | 24 rxn/96 rx |
| pG/pA-Tn5 for CUT&Tag | Hyperactive pG-Tn5 Transposase for CUT&Tag | S602-01/02 | 10 ug/20 ι |
| popArtibilition contenag | Hyperactive pA-Tn5 Transposase for CUT&Tag | S603-01/02 | 10 ug/20 u |
| pG/pA-Tn5 for CUT&Tag | Hyperactive pG-Tn5 Transposon for Illumina (4 µM) | S612-01/02 | 10 ug/20 ι |
| (Pre-loaded with Adapters) | Hyperactive pA-Tn5 Transposon for IIIumina (4 µM) | S613-01/02 | 10 ug/20 ι |
| CUT&Tag Lib Prep Kit | Hyperactive In-Situ ChIP Library Prep Kit for Illumina (pG-Tn5) | TD901-01/02 | 12 rxn/48 r |
| (Pre-loaded with Adapters) | Hyperactive In-Situ ChIP Library Prep Kit for Illumina (pA-Tn5) | TD902-01/02 | 12 rxn/48 r |
| CUT&RUN | Hyperactive pA-MNase for CUT&RUN | S701-01/02 | 10 ug/20 ι |
| | | | |

Single Cell-Seq

| Application | Product Name | Cat. No.# | Size |
|-----------------|--|---------------|----------------------|
| Single Cell WGA | Discover-sc Single Cell Kit | N601-01/02 | 24 rxn/96 rxn |
| Single Cell WGA | Discover-sc Single Cell WGA Kit | N603-01/02 | 24 rxn/96 rxn |
| Single Cell WTA | Single Cell Full Length mRNA-Amplication Kit | N712-01/02/03 | 12 rxn/24 rxn/96 rxn |

Beads & Library Quantification

| | Application | Product Name | Cat. No.# | Size |
|-----|-------------------------------|--|-------------------|-------------------|
| HOT | DNA Clean-up & Size-Selection | VAHTS DNA Clean Beads | N411-01/02/03 | 5 ml/60 ml/450 ml |
| | RNA Clean-up | VAHTS RNA Clean Beads | N412-01/02/03 | 5 ml/40 ml/450 ml |
| | | VAHTS Library Quantification Kit for Illumina® | NQ101/102/103/104 | 500 rxn |
| | Library Quantification Kits | DNA Standard 1-6 | NQ105 | 8 rxn |
| | | Library Dilution Buffer | NQ106 | 50 ml |
| | | Equalbit dsDNA HS Assay Kit | EQ111-01/02 | 100 / 500 assays |
| New | | Equalbit 1x dsDNA HS Assay Kit | EQ121-01/02 | 100 / 500 assays |
| | Equalbit Assay Kits | Equalbit RNA HS Assay Kit | EQ211-01/02 | 100 / 500 assays |
| | | Equalbit RNA BR Assay Kit | EQ212-01/02 | 100 / 500 assays |
| | | | | |

cfDNA

| Application | Product Name | Cat. No.# | Size |
|--------------------|--|------------|------------------|
| cfDNA Preservation | VAHTS Blood Collection Tube for Cell-Free DNA Preservation | N901 | 10 ml tube |
| cfDNA Isolation | VAHTS Serum / Plasma Circulating DNA Kit | N902-01/02 | 50 rxn / 200 rxn |

2020 Product Catalogue

TruePrep DNA Library Prep Kit V2 for Illumina (Vazyme, #TD501/502/503)

Transposase (Tn5)-based Ultra-Fast DNA Library Prep Kit

- **Easy to Use** One-step enzymatic reaction.
- Ultra-Fast Library prep within 90 min.
- Versatile Applicable for genomic DNA, single cell-seq, and epigenetics (i.e. ATAC-Seq).
- **Reliable** Optimized polymerase and buffer to achieve high efficiency and uniformity.

VAHTS Universal Plus DNA Library Prep Kit for Illumina (Vazyme, #ND617)

DNA Library Prep Kit with Enzymatic Fragmentation

- **Universal** Applicable for 100 pg 1 µg of input DNA from many species.
- **Enzymatic fragmentation with no need for physical shearing / sonication.**
- **Time-Saving** Fragmentation, end repair, dA-tailing are performed in one step.
- **Reliable** Generate high-quality DNA libraries with high yields.

VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina (Vazyme #NR604)

Fast & Effective RNA Library Prep Kit

- Fast The 2nd strand cDNA synthesis, end-repair, and dA-Tailing in one reaction! High-quality library prepared from enriched RNA within 4 hours.
- Flexible Compatible with a variety of RNA enrichment modules.
- Universal Options for the preparation of stranded or non-stranded RNA-Seq libraries.

Hyperactive In-Situ ChIP Library Prep Kit for Illumina (pG/pA-Tn5) (Vazyme, #TD901/902)

An revolutionary technology that can replace CUT&RUN and ChIP-Seq.

- Easy to Operate DNA tagmentation and adapters ligation in one step. Sequencing-ready libraries can be generated from live cells within only 9 hours.
 Hyperactive pG/A & Tn5 transposase with hyper activity results in high efficiency of DNA
- tagmentation.
- **Low Input** Starting from 60 cells or even single cells.
- **High Purity** Proteins with high purity and extra low residual amount of nucleic acid.



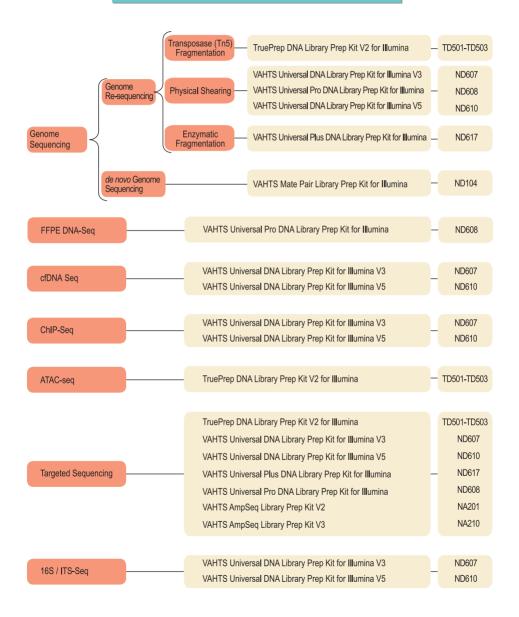








DNA-Seq Library Prep for Illumina®



DNA-Seq Library Prep for MGI®

| Genome Re-Sequencing | Physical Shearing – Enzymatic Fragmentation | VAHTS Universal DNA Library Prep Kit for MGI VAHTS Universal Pro DNA Library Prep Kit for MGI VAHTS Universal DNA Library Prep Kit for MGI V5 VAHTS Universal Plus DNA Library Prep Kit for MGI | NDM607 NDM608 NDM610 NDM617 |
|-------------------------|---|--|--------------------------------------|
| FFPE DNA-Seq | | VAHTS Universal Pro DNA Library Prep Kit for MGI | NDM608 |
| cfDNA Seq | | VAHTS Universal DNA Library Prep Kit for MGI VAHTS Universal DNA Library Prep Kit for MGI V5 | NDM607 NDM610 |
| ChIP-Seq | | VAHTS Universal DNA Library Prep Kit for MGI VAHTS Universal DNA Library Prep Kit for MGI V5 | NDM607 NDM610 |
| | | VAHTS Universal DNA Library Prep Kit for MGI VAHTS Universal DNA Library Prep Kit for MGI V5 | NDM607 NDM610 |
| Targeted Sequencir | ng | VAHTS Universal Plus DNA Library Prep Kit for MGI VAHTS Universal Pro DNA Library Prep Kit for MGI VAHTS AmpSeq General Library Prep Kit for MGI | NDM617 NDM608 NAM203 |
| 16S / ITS-Seq | | VAHTS Universal DNA Library Prep Kit for MGI VAHTS Universal DNA Library Prep Kit for MGI V5 | NDM607 NDM610 |

TruePrep DNA Library Prep (Transposase-based)

→ TruePrep DNA Library Prep Kit V2 for Illumina[®] (#TD501, #TD502, #TD503)

| Rapid & Easy | Time-Saving: Library prepared within 90 min. Easy-to-Use: one-step enzymatic reaction, no need for physical shearing / sonication. |
|-----------------------------------|---|
| High Adaptability to Input DNA | Applicable for genomic DNA, cDNA, and amplicons from multiple species. Input DNA: 1 ng - 50 ng. |
| High Amplification Uniformity | Optimized polymerase and buffer to achieve high efficiency and uniformity in library amplification. |

Selected Product Citations of TruePrep Kits (#TD501, #TD502, #TD503)

Wang J, et al. Asymmetric expression of LincGET biases cell fate in two-cell mouse embryos. *Cell*, 2018, 175(7):18871-1901.

Han X, et al. Mapping the Mouse Cell Atlas by Microwell-Seq. Cell, 2018, 172(5):1091-107.

Zhang L, et al. Lineage tracking reveals dynamic relationships of T cells in colorectal cancer. *Nature*, 2018, 564(7735):268-72.

Wu J, et al. Chromatin analysis in human early development reveals epigenetic transition during ZGA. *Nature*, 2018, 557(7704):256-60.

Zheng C, et al. Landscape of Infiltrating T Cells in Liver Cancer Revealed by Single-Cell Sequencing. Cell, 2017, 169(7):1342-56.

Huang X, et al. Genomic architecture of heterosis for yield traits in rice. Nature, 2016, 537(7622):629-33.

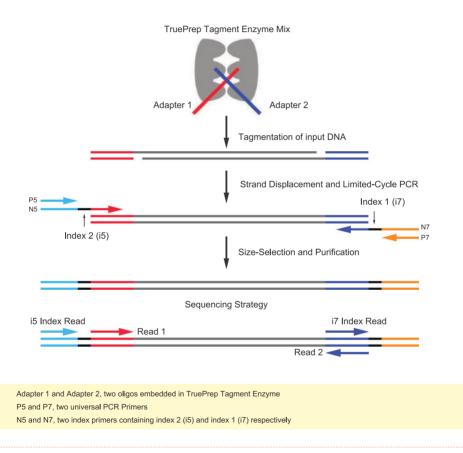
Wu J, et al. The landscape of accessible chromatin in mammalian preimplantation embryos. Nature, 2016, 534(7609):652-7.

Guo X, et al. Global characterization of T cells in non-small-cell lung cancer by single-cell sequencing. *Nature Medicine*, 2018, 24(7):978-85.



Mechanism of TruePrep DNA Library Prep

TruePrep Tagment Enzyme Mix (TTE Mix) contains transposase and two kinds of adapters (Adapter 1 and Adapter 2) with equal molar. Input DNA are fragmented and linked with adapters on both ends just by mixing with TTE Mix, followed by a 10-minute incubation at 55°C. The tagged DNA fragments can be further amplified with two pairs of primers N5 (N5XX) / N7 (N7XX) and P5 / P7 (PCR Primer Mix, PPM). After size selection and clean-up, the library is ready for sequencing on Illumina platforms.



 Index 2 (i5) 5'-AATGATACGGCGACCACCGAGATCTACACIIIIIIITCGTCGGCAGCGTCAGATGTGTATAAGAGACAG-NNNNNN-CTGTCTCTTATACACATCTCCGAGCCCACGAGACIIIIIIIIITCGTCGTATGCCGTCTTCTGCTTG-3' Index 1 (i7)

 IIIIIII: Index 2 (i5), 8 bases
 IIIIIII: Index 1 (i7), 8 bases
 -NNNNN-: insert sequence

DNA-Seq Library Preparation

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VAHTS Universal Plus DNA Library Prep Kit for Illumina[®] (#ND617) (for Enzymatic Fragmentation)

| Universal | Applicable for 100 pg-1 µg of |
|-------------|-------------------------------|
| Easy | Enzymatic fragmentation with |
| Time-Saving | Fragmentation, end repair, dA |
| Reliable | Generate high-guality DNA lib |

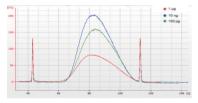
Applicable for 100 pg-1 µg of input DNA (e.g. genomic DNA, FFPE DNA) from many species.

Enzymatic fragmentation with a single protocol, with no need for physical shearing / sonication.

Fragmentation, end repair, dA-tailing are performed in one step. No clean-ups needed before adapter ligation.

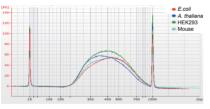
Generate high-quality DNA libraries with high yields.

1. Broad Input Amount Compatibility



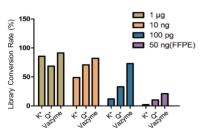
Size distribution of library fragments constructed with different DNA input. For different input mounts of salmon sperm gDNA (100 pg, 10 ng, 1 ug, respectively) with the same fragmentation time, the size distribution of these libraries were identical.

2. Extensive Species Compatibility

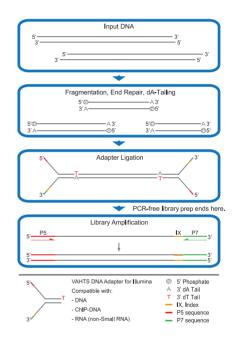


Size distribution of library fragments constructed by gDNA of different species. For the same input of 100 ng, the size distribution of these libraries from different species were identical.

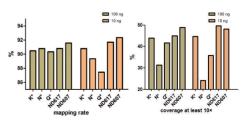
3. Excellent Library Conversion Rate



Comparison of library conversion rates. For different input amount of gDNA libraries, Vazyme #ND617 has a higher library conversion rate than that of other competitors.



4. High Sequencing Data Quality

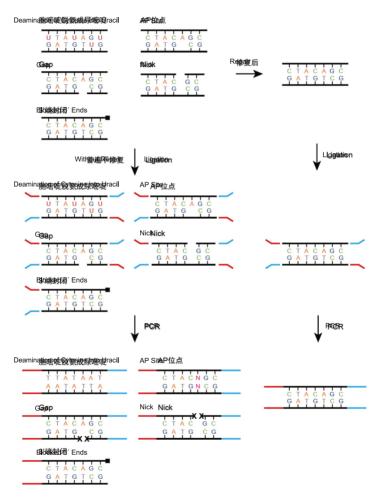


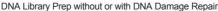
Comparison of sequencing data quality. The Arabidopsis thaliana gDNA library constructed using Vazyme ND617, followed by sequencing on Hiseq X10 PE150.



Mechanism of FFPE DNA Library Prep

VAHTS Universal Pro DNA Library Prep Kit for Illumina (Vazyme, #ND608) is specially designed for library preparation from 100 pg - 1 µg of input DNA for NGS on Illumina® platforms. This kit contains a DNA damage repair module that can effectively repair DNA damage caused by formalin-fixed paraffin-embedded (FFPE), including deamination of cytosine, nicks and gaps, oxidized bases, blocked 3' ends, compatible with common DNA samples without affecting the quality of normal DNA sample libraries. The overall optimization of the module of end-repair, ligation and library amplification leads to excellent library conversion rate and amplification output. It is widely applicable to PCR or PCR-Free library construction of multiple samples, and is compatible with targeted capture process.







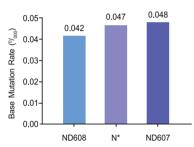
\rightarrow VAHTS Universal Pro DNA Library Prep Kit for Illumina[®] (#ND608)

Repairable Types of DNA Damage

| Types of DNA Damage in FFPE Samples | Deamination of Cytosine into Uracil | Nicks & Gaps | Oxidized Bases | Blocked 3' Ends | DNA Fragmentation | DNA-Protein Crosslinks |
|-------------------------------------|--|--------------|----------------|--------------------|----------------------|---------------------------|
| Whether Can Be Repaired by #ND608 | YES | YES | YES | YES | NO | NO |

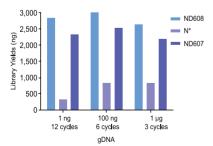
Validation Data

1. Efficient Repair of Base Damage



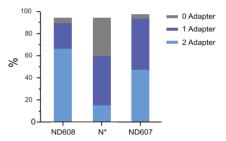
The DNA Repair Module of the ND608 has the ability to repair base damage efficiently, significantly reducing the number of base aberrant mutations introduced during FFPE sample preparation and storage.

3. Excellent Library amplification Efficiency



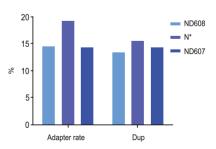
For amplification, the yield of the library constructed by ND608 is increased by 3-9 times compared with other kits under the same cycle number; the number of cycles required for competing products is reduced by 2-3 cycles under the same yield.

2. Efficient Library Conversion



The library conversion rate of ND608 is higher than that of N* company and VAHTS Universal Plus DNA Library Prep Kit for Illumina (Vazyme, #ND607, without FFPE repair module), and the library conversion rate is as high as 67%.

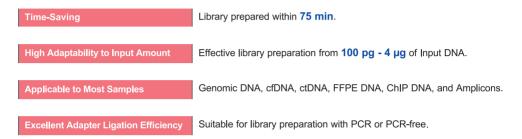
4. Excellent Raw Data Quality



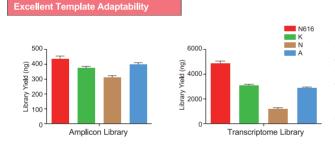
ND608 can effectively improve the data quality of samples, with lower Adapter rate and Duplication rate.



VAHTS Universal DNA Library Prep Kit for Illumina[®] V3 (#ND607) (for physical fragmentation)

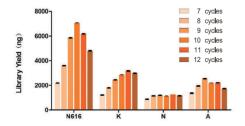






The amplification performance of VAHTS HiFi Amplification Mix (Vazyme, #N616) in multiple samples is significantly better than that of competitors. Amplicon library (80 ng) and transcriptome library (5 ng) was used as template and amplified for 4 and 13 cycles, respectively. The library concentrations were determined by Qubit.

Super High Plateau of Amplification



The amplification plateau of VAHTS HiFi Amplification Mix (Vazyme, #N616) can reach 7 µg, which is significantly superior to that of competitors. Fragmented DNA (50 ng, approximately 350 bp) was ligated with adapters and then amplified. The library concentrations were determined by Qubit.

RNA-Seq Library Prep for Illumina®

| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina VAHTS Universal V8 RNA-seq Library Prep Kit for Illumina | NR604 NR605 |
|-----------------------------------|--|------------------------------------|
| Ultra Fast & Universal Adapter | VAHTS RNA Adapters Set 1-Set 6 for Illumina VAHTS RNA Multiplex Oligos Set 1-Set 2 for Illumina | N803/N804 & N809-N812 N323/N324 |
| Module | VAHTS mRNA Capture Beads Ribo-off rRNA Depletion Kit (Human/Mouse/Rat) Ribo-off rRNA Depletion Kit (Bacteria) Ribo-off rRNA Depletion Kit (Plant) | N401 - N406 N407 N409 |
| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina VAHTS Universal V8 RNA-seq Library Prep Kit for Illumina | NR604 NR605 |
| Non-Stranded mRNA-Seq Adapter | VAHTS RNA Adapters Set 1-Set 6 for Illumina VAHTS RNA Multiplex Oligos Set 1-Set 2 for Illumina | N803/N804 & N809-N812 N323/N324 |
| Module | - VAHTS mRNA Capture Beads | - N401 |
| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina VAHTS Universal V8 RNA-seq Library Prep Kit for Illumina | NR604 NR605 |
| Stranded mRNA-Seq Adapter | VAHTS RNA Adapters Set 1-Set 6 for Illumina VAHTS RNA Multiplex Oligos Set 1-Set 2 for Illumina | N803/N804 & N809-N812 N323/N324 |
| Module | - VAHTS mRNA Capture Beads | - N401 |
| Kit | VAHTS Total RNA-Seq (H/M/R)Library Prep Kit for Illumina - VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina VAHTS Universal V8 RNA-seq Library Prep Kit for Illumina | NR603 - NR604 NR605 |
| Total RNA-Seq (rRNA Depletion) | VAHTS RNA Adapters Set 1-Set 6 for Illumina VAHTS RNA Multiplex Oligos Set 1-Set 2 for Illumina | N803/N804 & N809-N812 N323/N324 |
| Module | Ribo-off rRNA Depletion Kit (Human/Mouse/Rat) - Ribo-off rRNA Depletion Kit (Bacteria) Ribo-off rRNA Depletion Kit (Plant) | N406 - N407 N409 |
| Small RNA-Seg | - VAHTS Small RNA Library Prep Kit for Illumina | - NR801 |
| Adapter | - VAHTS Small RNA Index Primer Kit for Illumina | - N813-N816 |



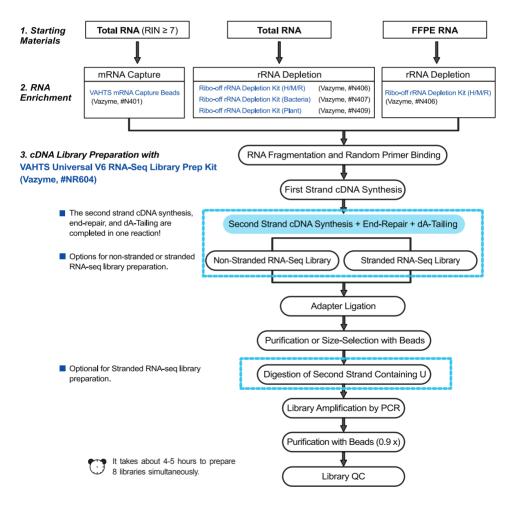
RNA-Seq Library Prep for MGI®

| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for MGI VAHTS Universal V8 RNA-seq Library Prep Kit for MGI | NRM604 NRM605 |
|-----------------------------------|---|--------------------------------|
| Ultra Fast & Universal RNA-Seq | - VAHTS RNA Adapters Set 8 for MGI | - NM208 |
| Module | VAHTS mRNA Capture Beads Ribo-off rRNA Depletion Kit (Human/Mouse/Rat) Ribo-off rRNA Depletion Kit (Bacteria) Ribo-off rRNA Depletion Kit (Plant) | N401 _ N406 N407 N409 |
| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for MGI VAHTS Universal V8 RNA-seq Library Prep Kit for MGI | NRM604 NRM605 |
| Non-Stranded mRNA-Seq Adapter | - VAHTS RNA Adapters Set 8 for MGI | - NM208 |
| Module | - VAHTS mRNA Capture Beads | - N401 |
| Kit | VAHTS Universal V6 RNA-seq Library Prep Kit for MGI VAHTS Universal V8 RNA-seq Library Prep Kit for MGI | NRM604 NRM605 |
| Stranded mRNA-Seq Adapter | - VAHTS RNA Adapters Set 8 for MGI | - NM208 |
| Module | - VAHTS mRNA Capture Beads | - N401 |
| Kit | VAHTS Total RNA-Seq (H/M/R)Library Prep Kit for MGI - VAHTS Universal V6 RNA-seq Library Prep Kit for MGI VAHTS Universal V8 RNA-seq Library Prep Kit for MGI | NRM603 - NRM604 NRM605 |
| Total RNA-Seq (rRNA Depletion) | - VAHTS RNA Adapters Set 8 for MGI | - NM208 |
| Module | Ribo-off rRNA Depletion Kit (Human/Mouse/Rat) - Ribo-off rRNA Depletion Kit (Bacteria) Ribo-off rRNA Depletion Kit (Plant) | N406 - N407 N409 |



Solutions of RNA-Seq Library Preparation

VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina® (Vazyme, #NR604) is specially designed for the preparation of transcriptome libraries for NGS platforms of Illumina®. This kit combines 2nd Strand cDNA synthesis, end-repair, and dA-Tailing into one step, with no need of clean-ups, which greatly simplifies the process of library construction and shortens the operation time. The kit also can be used for non-stranded or stranded transcriptome analysis.



Mechanism of VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina® (Vazyme, #NR604)

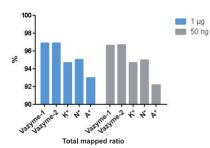


\rightarrow VAHTS Universal V6 RNA-seq Library Prep Kit for Illumina[®] (#NR604)

| Fast | The second strand cDNA synthesis, end-repair, and dA-Tailing are completed in one reaction! High-quality library prepared from enriched RNA within 4 hours . |
|-----------|--|
| Flexible | Compatible with a variety of RNA enrichment modules. |
| Universal | Options for the preparation of stranded or non-stranded RNA-Seq libraries. |

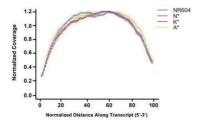
Validation Data

1. High Mapped Rate



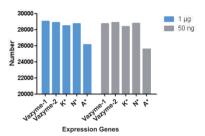
Compared with several similar kits, Vazyme, #NR604 performed better in mapped ratio.

3. Excellent Uniformity



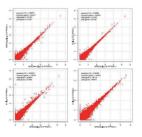
The data of library prepared using Vazyme, #NR604 was evenly distributed, with no 5'- or 3'- preference, and is highly consistent with several similar kits.

2. High Gene Detection Number



Compared with several similar kits on the market, Vazyme, #NR604 showed higher gene detection number.

4. High Expression Repeat Correlation

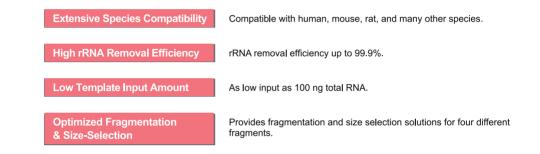


The correlation coefficient *r*² between Vazyme, #NR604 and several similar kits on the market is higher than 0.94.



Total RNA-Seq Library Prep

\rightarrow VAHTS Total RNA-seq (H/M/R) Library Prep Kit for Illumina[®] (#NR603)





He Y, et al. Transitory presence of myeloid-derived suppressor cells in neonates is critical for control of inflammation. *Nature Medicine*, 2018, 24(2):224-31.

LETTERS

medicine

Transitory presence of myeloid-derived suppressor cells in neonates is critical for control of inflammation

Yu-Mei He^{1,2,9}, Xing Li^{1–3,9}, Michela Perego^{4,9}, Yulia Nefedova⁴, Andrew V Kossenkov⁴, Erik A Jensen⁵, Valerian Kagan⁶, Yu-Feng Liu¹, Shu-Yu Fu¹, Qing-Jian Ye³, Yan-Hong Zhou⁷, Lai Wei⁸, Dmitry I Gabriovich^{1,2,1,0}, B₁ iz Zhou^{1,2},¹⁰

PMN-MDSCs and M-MDSCs from spleens of neonatal (7 d old) and adult mice (6 to 8 weeks old) were enriched with CD11b-beads and then sorted on a FACSAria cell sorter (BD Bioscience), on the basis of the CD11b+Ly6CintLy6Grhenotype for PMN-MDSCs and CD11b+Ly6ChighLy6G- for M-MDSCs. The sorting purity was >95%. RNA sequencing was performed on the Illumina Hiseq 2500 platform. A VAHTS Total RNA-Seq Library Preparation Kit (Vazyme Biotech) was used for library preparation. Single-end-read runs were used with read lengths up to 50 bp in high-output mode and 30 million total read counts. Data were analyzed in RSEM v1.2.12 software30 against the mm10 genome, and genelevel read counts and RPKM values on the gene level were estimated for the ensemble transcriptome.



Ribo-off rRNA Depletion Kits

| Ribo-off Modules | Cat# | In-put Total RNA | Species |
|--|-------|------------------|--------------------------|
| Ribo-off rRNA Depletion Kit (H/M/R) | #N406 | 50 ng - 1 µg | Human, mouse, rat, etc.* |
| Ribo-off rRNA Depletion Kit (Bacteria) | #N407 | 1 µg - 5 µg | Bacteria. |
| Ribo-off rRNA Depletion Kit (Plant) | #N409 | 1 µg - 5 µg | Plants. |

* Validated species of #N406: human, mouse, rat, cattle, dog, horse, chicken, monkey, pig, zebrafish, etc.

25 min

MCRNA

MRN

30 min

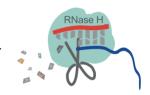
mRNA

Complete rRNA removal within 2 hours.



1. rRNA (> 80% of total RNA). 2. rRNA probe hybridization.





3. RNase H digestion to remove rRNA.



4. DNase I digestion to remove probes.

5. mRNA / IncRNA enriched after rRNA removal.

ncRNA

MCRNA

6. Library preparation for RNA-seq.

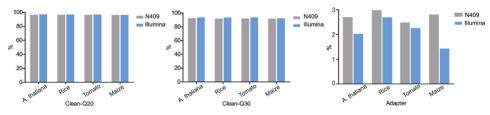


| Extensive Species Compatibility | Compatible with Arabidopsis, cotton, corn, soybean, rice, tomato, peanut, apple, wheat, Selaginella, Cuscuta, etc. |
|-----------------------------------|--|
| High rRNA Removal Efficiency | Effective removal of cytoplasmic rRNA, mitochondrial rRNA, and chloroplast rRNA. |
| Wide Range of Input Compatibility | 1 μg - 5 μg total RNA. |
| Easy & Fast Procedures | Using with #NR604, library preparation and quality control can be completed within one day. |

Validation Data

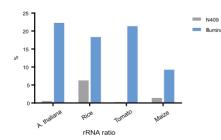
The transcriptome libraries of Arabidopsis thaliana, rice, tomato, and maize were prepared using Vazyme, #N409 and RNA-Seq library prep kit, respectively. The quality-controlled libraries were sequenced using an Illumina Hiseq X10 platform for PE150 sequencing. High-quality clean reads obtained from the raw data was analyzed.

1. Excellent Data Quality



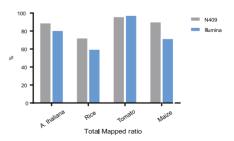
Compared with several similar kits on the market, Vazyme, #NR604 showed higher gene detection number.

2. High rRNA Removal Efficiency



For different species, N409 can effectively remove rRNA and minimize the waste of data caused by rRNA residues.

3. High Mapped Ratio



The data obtained by sequencing the library prepared using Vazyme, #N409 and RNA-Seq library prep kit showed high mapped ratio.

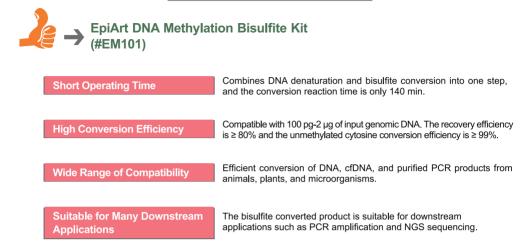


Library Prep for Epigenetics

| Bisulfite Sequencing | EpiArt DNA Methylation Bisulfite Kit 2 × EpiArt HS Taq Master Mix 2 × EpiArt HS Taq Master Mix (Dye Plus) EpiArt DNA Methylation Library Kit | | EM101 EM201 EM202 NE101 |
|----------------------|--|---|--|
| CUT&Tag | Hyperactive pG-Tn5 Transposase for CUT&Tag Hyperactive pA-Tn5 Transposase for CUT&Tag Hyperactive pG-Tn5 Transposon for Illumina (4 μ M) Hyperactive pA-Tn5 Transposon for Illumina (4 μ M) Hyperactive In-Situ ChIP Library Prep Kit for Illumina(pG-Tn5) Hyperactive In-Situ ChIP Library Prep Kit for Illumina(pA-Tn5) | _ | S602 S603 S612 S613 TD901 TD902 |
| CUT&RUN | Hyperactive pA-MNase for CUT&RUN | _ | S701 |

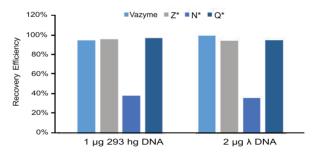


Methylation Bisulfite Kit

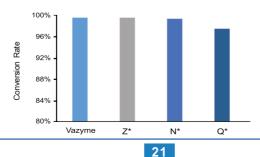


Validation Data

1. High Recovery Efficiency



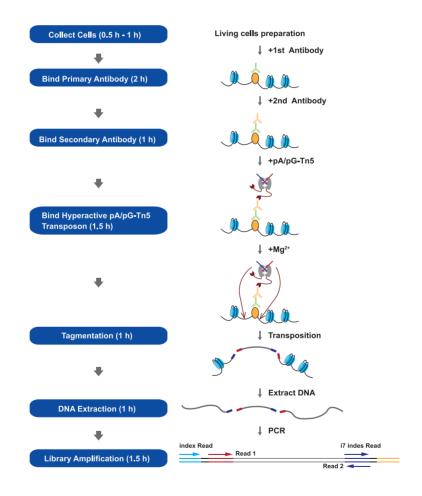
2. High Conversion Efficiency





CUT&Tag

Cleavage Under Targets and Tagmentation (CUT&Tag) is a specific technology designed for the study of protein-genomic interaction that can seamlessly replace the traditional ChIP-Seq. Compared with ChIP-Seq, the CUT&Tag has several significant advantages, including high signal-to-noise ratio, excellent repeatability, short operation time (generate sequencing-ready libraries beginning with live cells within one day), and low cell input. This technology will facilitates the research in epigenetics, tumors, and stem cells.



Workflow of CUT&Tag

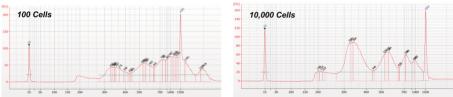
Epigenetics

→ Hyperactive In-Situ ChIP Library Prep Kit for Illumina (pG-Tn5/pA-Tn5) (#TD901 / #TD902)

| Easy to Operate | DNA tagmentation and adapters ligation can be achieved in one step. |
|---------------------------|---|
| Low Input Amount | Starting from 60 cells or even single cells. |
| Hyperactive | Protein G/A & Tn5 transposase with hyper activity results in high efficiency of DNA tagmentation. |
| High S/N Ratio | Higher signal-to-noise ratio than ChIP-Seq. |
| Excellent Reproducibility | Good reproducibility among sample replicates. |
| Time-Saving | Sequencing-ready libraries can be generated from live cells within only 9 hours. |

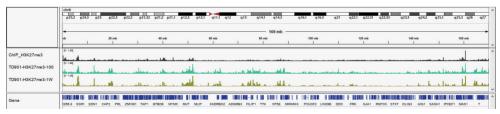
Validation Data

1. Low Input Amount of Cells



Size distribution of CUT&Tag library fragments prepared from 100 cells (left) or 10,000 cells (right). One hundred or 10,000 cells of HEK293 was used for CUT&Tag and library preparation using **Vazyme, #TD901**, respectively. The final concentration of pG-Tn5 transposon was 0.04 µM. The 1st antibody was H3K27me3 (CST, #9733), the 2nd antibody was Goat anti Rabbit (Bioworld, #BS13271). Size distribution of library fragments was detected using an Agilent 2100 Bioanalyzer. The results showed that, using **#TD901**, CUT&Tag libraries were prepared successfully from both 100 and 10,000 cells.

2. High Signal-to-Noise Ratio

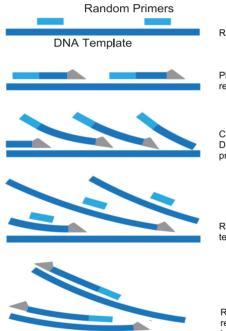


Peak enrichment of libraries that compared with different methods and different cell inputs. (The 1st row - ChIP-Seq; the 2nd row - CUT&Tag from 100 cells using #TD901; the 3rd row - CUT&Tag from 10,000 cells using #TD901). CUT&Tag libraries were sequenced on a Hiseq X10 PE150, respectively. The raw data was filtered and then subjected to Peak Calling analysis. The results showed that the peak enrichment of CUT&Tag library from 100 cells is similar to that of 10,000 cells, and the signal-to-nose ratio (S/N) of both CUT&Tag libraries was significantly better than that of traditional ChIP-Seq library.



Single Cell Whole Genome Amplification (WGA) Kit

Discover-sc Single Cell WGA Kit (#N603) is an isothermal amplification system based on the multiple displacement amplification (MDA) using Phi29. The Phi29 DNA polymerase is cloned from phage and has extremely strong stranddisplacement activity and can be used for in vitro MDA polymerization at a constant temperature, with no need for thermal cycling. One single polymerization reaction using Phi29 can achieve continuous polymerization extension up to 100 kb. This kit is suitable for the whole genome amplification (WGA) from a single cell, small amount of tissues, or even trace purified genomic DNA, to obtain a large amount of genomic DNA with high coverage.



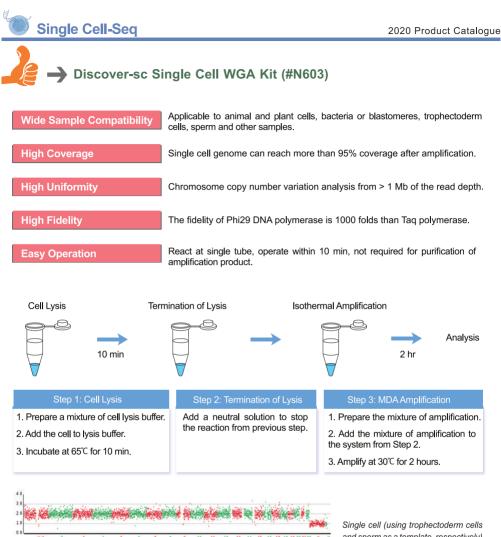
Random primers bind to multiple sites of the DNA template.

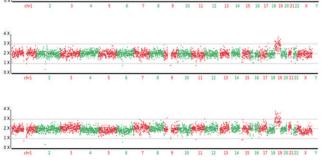
Phi29 DNA polymerase simultaneously initiates DNA replication at multiple primer binding sites.

Chain synthesis reaction in process replaces encountered DNA chain in synthesis and continue elongation, and produces replaced single strand DNA.

Replaced single strand DNA is bind by random primers as template.

Random primers initiate new DNA synthesis and chain replacement reaction, to synthesize double strand DNA of high molecular size.





Single cell (using trophectoderm cells and sperm as a template, respectively) genomic amplification was performed using Vazyme #N603. Then libraries were prepared for sequencing, which was performed in Illumina MiniSeq after pooling according to the effective concentration and the 0.01X of sequencing depth. The data showed that the distribution of the reads was uniform in all parts of the genome, indicating that the amplification uniformity of Vazyme #N603 was excellent.

Scatter plots of whole genome copy numbers.

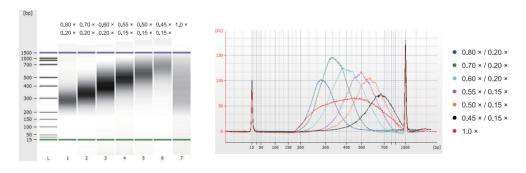


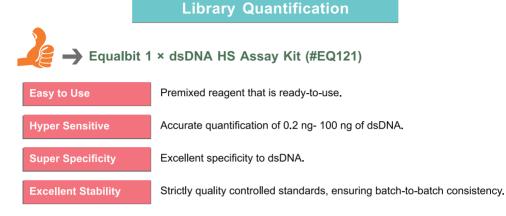
Beads VAHTS DNA Clean Beads (#N411) ♥ Applicable for DNA purification and size selection in NGS library preparation. Compatible with almost all library prep protocols provided by all manufacturers. VAHTS DNA Clean Beads Transfer Supernatant VAHTS DNA Separation DNA Fragments Separation Ethanol Rinse Elution Size-selected DNA Clean Beads 13333 Discard Larger Fragements Binding on Beads Discard Smaller Fragements in Supernatant

Validation Data

A DNA library (200 bp - 1,500 bp) was prepared using TruePrep DNA Library Prep Kit V2 for Illumina (**Vazyme, #TD501**). Size-selection was performed using VAHTS DNA Clean Beads (Vazyme, #N411) according to the different parameters in the following table, respectively. Size distribution was detected using an Agilent 2100 Bioanalyzer.

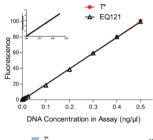
| Ratio of 1st-round (Beads: DNA) | 0.80 × | 0.70 × | 0.60 × | 0.55 × | 0.50 × | 0.45 × | 1.0 × |
|---------------------------------|--------|--------|--------|--------|--------|--------|----------|
| Ratio of 2nd-round (Beads: DNA) | 0.20 × | 0.20 × | 0.20 × | 0.15 × | 0.15 × | 0.15 × | |
| Average Size (bp) | 300 | 350 | 400 | 500 | 600 | 700 | 200-1500 |

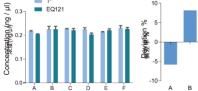




Validation Data

1. Hyper Sensitive

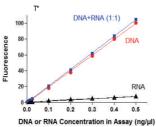




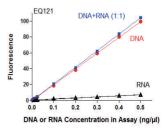
Linearity Analysis (0.2 ng-100 ng): dsDNA samples with 12 different concentrations were tested using **Vazyme**, **#EQ121** or a similar reagent from vendor T*, respectively. Fluorescence signals were read using a QubitTM Fluorometer 3.0. The result showed a excellent linear relationship between fluorescence and DNA concentration when the amount of dsDNA ranged from 0.2 ng to 100 ng.

2. Super Specificity to dsDNA

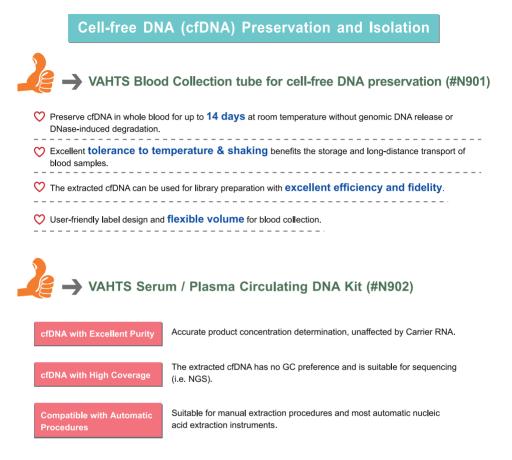
Samples of dsDNA, RNA, and DNA + RNA (1:1) were tested with **Vazyme**, **#EQ121** and a similar reagent from vendor T*. The results showed that Similar to products from T*, **Vazyme**, **#EQ121** specifically binds to dsDNA, even in the presence of RNA.



Results and deviation rate obtained from different operators: the standard of **Vazyme**, **#EQ121** was serial-diluted to 0.2 ng / μ l (theoretical concentration), and then tested by 6 different operators (A, B, C, D, E, F), respectively. The results are showed on the left, and the deviation rate, calculated as [result of EQ121 - result of T^{*} x 100%, are all within 10%.

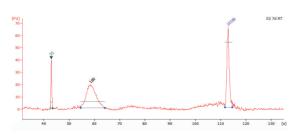






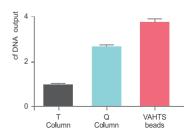
Validation Data

1. Products with High Purity



Electropherogram of separated cfDNA in Agilent 2100 analyzer.

2. High Yield



Actual yield of cfDNA detected by qPCR.





Wang J, Wang L, et al. Asymmetric Expression of LincGET Biases Cell Fate in Two-Cell Mouse Embryos. Cell, 2018, 175(7):18871-1901. (Vazyme #TD502)

Han X, Wang R, *et al*. Mapping the Mouse Cell Atlas by Microwell-Seq. Cell, 2018, 172(5):1091-107 (Vazyme #TD513)

Zhang L, Yu X, et al. Lineage tracking reveals dynamic relationships of T cells in colorectal cancer. *Nature*, 2018, 564(7735):268-72. (Vazyme #TD502)

Wu J, Xu J, et al. Chromatin analysis in human early development reveals epigenetic transition during ZGA. *Nature*, 2018, 557(7704):256-60 (Vazyme, #TD502)

Zheng C, Zheng L, *et al.* Landscape of Infiltrating T Cells in Liver Cancer Revealed by Single-Cell Sequencing. *Cell*, 2017, 169(7):1342-56 (Vazyme, #TD503)

Huang X, Yang S, et al. Genomic architecture of heterosis for yield traits in rice. Nature, 2016, 537(7622):629-33. (Vazyme, #TD501)

Wu J, Huang B, et al. The landscape of accessible chromatin in mammalian preimplantation embryos. Nature, 2016, 534(7609):652-7. (Vazyme, #TD501)

Guo X, Zhang Y, et al. Global characterization of T cells in non-small-cell lung cancer by single-cell sequencing. *Nature Medicine*, 2018, 24(7):978-85. (Vazyme, **#TD501**)

Liu Z, Yan M, et al. Nucleoporin Seh1 Interacts with Olig2/Brd7 to Promote Oligodendrocyte Differentiation and Myelination. *Neuron*, 2019, pii: S0896-6273(19)30153-9. (Vazyme, #TD502)

Li X, Meng D, et al. Single nucleus sequencing reveals spermatid chromosome fragmentation as a possible cause of maize haploid induction. *Nature Communications*, 2017, 8(1):991. (Vazyme, #TD501, #TD503)

Yang L, Wang W, et al. A single-cell transcriptomic analysis reveals precise pathways and regulatory mechanisms underlying hepatoblast differentiation. *Hepatology*, 2017, 66(5):1387-401. (Vazyme, **#TD502**, **#411**)

VAHTS Universal DNA Library Prep Kits for Illumina®

Zhang M, Dong Y, et al. Transcription factor Hoxb5 reprograms B cells into functional T lymphocytes. Nature Immunology, 2018, 19(3):279-90. (Vazyme, #ND604, #N601)

Cao S, Yu S, et al. Chromatin accessibility dynamics during chemical induction of pluripotency. Cell Stem Cell, 2018, 22(4): 529-42. (Vazyme, #ND102, #NQ101)

🖞 VAHTS Total RNA-seq (H/M/R) Library Prep Kit for Illumina®

He Y, Li X, et al. Transitory presence of myeloid-derived suppressor cells in neonates is critical for control of inflammation. *Nature Medicine*, 2018. 24(2):224-31. (Vazyme, #NR603)



VAHTS mRNA-seq Library Prep Kits for Illumina®

Tian X, He G, et al. Cryptococcus neoformans sexual reproduction is controlled by a quorum sensing peptide. *Nature Microbiology*, 2018: 3(6):698-707. (Vazyme, #NR601)

Han X, Chen H, et al. Mapping human pluripotent stem cell differentiation pathways using high throughput single-cell RNA-sequencing. *Genome Biology*, 2018, 19(1):47. (Vazyme, #NR601)

Qin X, Liu S, *et al*. Heterotrimeric G Stimulatory Protein α Subunit Is Required for Intestinal Smooth Muscle Contraction in Mice. *Gastroenterology*, 2017, 152(5):1114-25. (Vazyme, #NR601)

Shen C, Wang J, *et al.* Transcriptome Analysis of Differentially Expressed Genes Induced by Low and High Potassium Levels Provides Insight into Fruit Sugar Metabolism of Pear. *Frontiers in Plant Science*, 2017, 8:938. (Vazyme, #NR601)



VAHTS Stranded mRNA-seq Library Prep Kit for Illumina®

Song L, Ma Q, *et al.* Molecular Link between Leaf Coloration and Gene Expression of Flavonoid and Carotenoid Biosynthesis in Camellia sinensis Cultivar 'Huangjinya'. *Frontiers in Plant Science*, 2017, 8:803. (Vazyme, #NR602)



Discover-sc Single Cell Kit

Zhang M, Dong Y, et al. Transcription factor Hoxb5 reprograms B cells into functional T lymphocytes. *Nature Immunology*, 2018, 19(3):279-90. (Vazyme, #ND604, #N601)

VAHTS DNA Clean Beads

Yang L, Wang W, et al. A single-cell transcriptomic analysis reveals precise pathways and regulatory mechanisms underlying hepatoblast differentiation. *Hepatology*, 2017, 66(5):1387-401. (Vazyme, #TD502, #N411)

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