



These products are sold for Research Use Only; not for use in diagnostic procedures



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Workflow Summary

- O1 Allow the Master Mix tubes to thaw; then gently vortex to mix.
- O2 In a containment hood or dead air box, pipette 45 μL of Master Mix into individual wells of a PCR plate (use a different indexed Master Mix for each sample and control).
- O3 Add 0.2 µL EagleTaq™ DNA polymerase to each Master Mix.
- O4 Add 5 μL of DNA (at a minimum of 10 ng/μL) from the unknown samples and controls to wells containing the respective Master Mix reactions then pipette up and down 5-10 times to mix.
- 05 Add 5 μL of molecular biology grade water to the well containing the Master Mix for the no template control then pipette up and down 5-10 times to mix.
- 06 Seal the plate and amplify target DNA using the following thermal cycler program:

Step	Temperature		Cycle
1	95 °C	7 minutes	1
2	95 °C	45 seconds	
3	60 °C	45 seconds	29x*
4	72 °C	90 seconds	
5	72 °C	10 minutes	1
6	15 °C	\sim	1

*LymphoTrack IGHV Leader Somatic Hypermutation Assay (32 cycles).

PRODUCTS

CATALOG #

CATALOG #	PRODUCTS
7-121-0129 7-121-0139	LymphoTrack® IGH FR1/2/3 Assay Kit A - MiSeq™ LymphoTrack® IGH FR1/2/3 Assay Panel - MiSeq™
7-121-0009	LymphoTrack [®] <i>IGH</i> FR1 Assay Kit A - MiSeq™
7-121-0039	LymphoTrack® <i>IGH</i> FR1 Assay Panel - MiSeq™
7-121-0149	LymphoTrack® <i>IGH</i> FR1 Assay Panel B - MiSeq™
7-121-0089	LymphoTrack® <i>IGH</i> FR2 Assay Kit A − MiSeq™
7-121-0099	LymphoTrack® <i>IGH</i> FR2 Assay Panel - MiSeq™
7-121-0109	LymphoTrack® <i>IGH</i> FR3 Assay Kit A − MiSeq™
7-121-0119	LymphoTrack® <i>IGH</i> FR3 Assay Panel - MiSeq™
7-121-0059	LymphoTrack® IGHV Leader Somatic Hypermutation Assay Kit A - MiSeq™
7-121-0069	LymphoTrack [®] IGHV Leader Somatic Hypermutation Assay Panel - MiSeq
7-122-0009	LymphoTrack® <i>IGK</i> Assay Kit A − MiSeq™
7-122-0019	LymphoTrack® <i>IGK</i> Assay Panel - MiSeq™
7-225-0009	LymphoTrack® <i>TRB</i> Assay Kit A - MiSeq™
7-225-0019	LymphoTrack® <i>TRB</i> Assay Panel - MiSeq™
7-227-0019	LymphoTrack® <i>TRG</i> Assay Kit A – MiSeq™
7-227-0009	LymphoTrack® <i>TRG</i> Assay Panel - MiSeq™
7-500-0009	LymphoTrack® Software – MiSeq™
7-500-0008	LymphoTrack® MRD Software

- O7 Purify the PCR products using the Agencourt® AMPure® XP PCR Purification system. When using the LymphoTrack *TRB* Assay add 35 μL of particles to each 50 μL reaction, for all other LymphoTrack Assays add 50 uL of particles to each 50 μL reaction; elute DNA in 25 μL of eluant.
 - Quantify amplicons using the KAPA™ library quantification kit according to the kit instructions. Dilute amplicons 1:4,000 before proceeding to qPCR.
 - O9 Pool equal amounts of amplicons from samples and positive and negative controls (do not include the no template control), dilute 1:1,000 then quantify the library using the KAPA library quantification kit.
 - Denature and dilute the library.
 - Load 600 μL of denatured and diluted library on the MiSeq[®] Reagent Cartridge.
 - 12 Set up a MiSeq[®] sample sheet using the Illumina[®] Experiment Manager or provided Sample Sheet **.csv* file.
 - 3 Start the MiSeq[®] run.
 - Analyze and visualize the acquired data using the LymphoTrack Software for the MiSeq®.

QUANTITY

indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 25-48 (5 sequencing reaction each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) indices 1-8 (5 sequencing reactions each) indices 1-24 (5 sequencing reactions each) 1 CD 1 CD

Storage Conditions: -85°C to -65°C (DNA controls may be seperated from kits and stored at 2°C to 8°C

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