DESCRIPTION OF TABLES S1-S18

https://doi.org/10.1093/nar/gkad339

Table S1:

Sheet Name Summary of Contents

BBBsamples Information on samples obtained from the Brisbane Breast Bank (BBB). FACSmethods Antibodies and immuno-staining conditions for FACS experiments.

Table S2:

Sheet Name Summary of Contents

Contains the identifiers for all sequenced samples, including the original Novogene file

SampleDescription name, short name for internal identification and descriptive name containing cell type

and patient origin.

ReadNumbers Overal number of reads sequenced per sample and final average of all samples.

Main metrics obtained by FastQC (version 0.11.5) for reads in R1 files, before any

processing or filtering (raw reads).

Main metrics obtained by FastQC (version 0.11.5) for reads in R2 files, before any

processing or filtering (raw reads).

Table S3:

MetricsFastQC R1

Sheet Name Summary of Contents

ReadNumbers Overal number of reads sequenced per sample and final average of all samples.

MetricsFastQC_R1 Main metrics obtained by FastQC (version 0.11.5) for reads in R1 files, after trimming.

MetricsFastQC R2 Main metrics obtained by FastQC (version 0.11.5) for reads in R2 files, after trimming.

Table S4:

Sheet Name Summary of Contents

Strandness Percentage of reads identified as RSeQC as being of the type "1+-,1-+,2++,2--".

CorrectedReads Percentage of reads for which Rcorrector attempted correction.

UncorrectableReads Percentage of reads FurPe deemed uncorrectable after Rcorrector attempts.

TrimmingSummary Percentage of reads surviving trimming either in a pair or individually. Percentages

are presented for reads before and after correction efforts, for comparison.

RibosomalReads

Percentage of reads coming from ribosomal RNAs (as annotated in the SilvaDB) in each sample in reads before and after correction, for comparison.

MappedReads Overall alignment rate between reads and the human genome (GENCODE hg38).

Table S5:

Sheet Name Summary of Contents

AssemblyMetrics Main assembly metrics calculated by TrinStats for the raw assembly and the

TransRate filtered assembly.

TransRateMetrics Trinity assembly statistics before and after TransRate filtering for optimisation.

BUSCOmetrics Results for BUSCO using both the Eukaryote and the Mammalian sets of orthologs.

Comparison with

Holzer and Marz Comparisons between our in-house assembly and data from Hölzer and Marz, 2019.

Table S6:

Sheet Name Summary of Contents

CodingPotential Summary of all coding potential calculators used in this work.

FEELnc Summary of FEELnc results for genes with supported lack of coding potential

(only 'best' targets are shown).

ClassesIncRNAs Number of IncRNAs per class (defined by their genomic context) used to create

the included pie chart.

Table S7:

Sheet Name

Summary of Contents

IncRNAs_with_LongReadSupport_A

IncRNAs_with_LongReadSupport_B

NumberOfLongReads per IncRNA

IncRNAs with TSS A

IncRNAs with TSS B

RT-PCR primers

Summary of BLAT alignment between NB-IncRNAs and publicly available long-reads (after applying coverage filter).

Summary of BLAT alignment between NB-IncRNAs and long-reads generated in-house (after applying coverage filter).

Number of long-reads that align to each IncRNA (after applying coverage filter).

Summary of Bedtools intersect between lncRNAs and TSS peaks from publicly available RAMPAGE data (after filtering for TSSs located <500bp upstream or <50bp internal to transcript start).

Summary of Bedtools intersect between lncRNAs and TSS peaks from in-house RAMPAGE experiments (after filtering for TSSs located

<500bp upstream or <50bp internal to transcript start).

Primers designed for RT-PCT of NB-IncRNAs presented at Fig. S1b

Table S8:

SheetName

Summary of Contents

NBdb

NBdb Statistics

NBdb_GeneAnnotation

InHouse_IncRNAdb

Telomere_geneDB

DNArepair_geneDB

EMT geneDB

ClaudinLow_geneDB HumanHousekeepingGenes

RibosomalRNA

Normal Breast DataBase of genes from the literature previously

assigned to each cell subpopulation.

Statistics of the database, including the number of genes per cell

subpopulation and genes per reference.

Genomic annotation of all genes in the database. Database of unannotated IncRNAs compiled from multiple sources.

Genes implicated in telomere maintenance listed in GSEA (Reactome

and Biocarta gene sets).

Genes implicated in DNA repair pathways listed in REPAIRtoire and/or

MD Anderson databases.

EMT-related genes from GSEA and/or EMTome.

Genes upregulated in claudin-low samples, according with Prat, 2010.

Human housekeeping genes, deposited at HK and/or HRT. Sequences for rRNA depletion (169 entries of rRNAs).

Table S9:

Sheet Name

Summary of Contents

GencodeAnnotated

SummaryOfGencodeAnnotated

ncRNAdbAnnotation Lnc2CancerMatches Lnc2CancerClasses IncRNAfuncMatches

MaTARs

List of all assembled lncRNAs annotated in hg38 human genome

assembly from GENCODE. Overall counts of genes from different types from GencodeAnnotated.

List of NB-IncRNAs annotated in the in-house database of ncRNAs. List of NB-IncRNAs annotated in the Lnc2Cancer database. Classification of Lnc2CancerMatches NB-IncRNAs in Lnc2Cancer. List of NB-IncRNAs annotated as genes featuring in IncRNAfunc.

Correspondence between MaTARs and NB-IncRNAs.

Table S10:

Sheet Name

349_elncRNAs_AndPairedENhancers

TargetsOf elncRNAs

NBmarkers TargetedBy elncRNAs 1968pancRNAs AndPairedPromoters

TargetsOf_parcRNAs

NBmarkers TargetedBy parcRNAs 825TALRs AndPairedUTRs

Summary of Contents

List of 349 lncRNAs that co-localize with known enhancer regions. Pairwise correlation between elncRNAs and the correspondent annotated enhancer element.

Normal breast marker genes targeted by the elncRNAs.

List of 1968 IncRNAs that co-localize with known promoter regions. Pairwise correlation between pancRNAs and the correspondent

annotated promoters.

Normal breast marker genes targeted by the parcRNAs. List of 825 TALRs that co-localize with known UTRs.

TargetsOf_TALRs

NBmarkers_TargetedBy_TALRs

Pairwise correlation between TALRs and the correspondent annotated UTRs.

Normal breast marker genes targeted by the TALRs.

Table S11:

Sheet Name Summary of Contents

ConsistentlyExpressed in C1 ConsistentlyExpressed_in_C2 $Consistently Expressed_in_C4$

NBmarkersOfInterest

NBmarkers TargetsOf CE in C1

NBmarkers_TargetsOf_CE_in_C2

NBmarkers TargetsOf CE in C4

NBmarkers_TargetsOf_UCE_in_C1

NBmarkers_TargetsOf_UCE_in_C2

NBmarkers_TargetsOf_UCE_in_C4

IncRNAs consistently expressed in the C1 population (above 1 TPM) IncRNAs consistently expressed in the C2 population (above 1 TPM) IncRNAs consistently expressed in the C4 population (above 1 TPM) Population-specific NB-lncRNAs that are partners of known proteincoding markers

Normal breast marker genes targeted by the IncRNAs consistently expressed in the C1 population

Normal breast marker genes targeted by the lncRNAs consistently expressed in the C2 population

Normal breast marker genes targeted by the lncRNAs consistently expressed in the C4 population

Normal breast marker genes targeted by the IncRNAs consistently expressed only in the C1 population

Normal breast marker genes targeted by the lncRNAs consistently expressed only in the C2 population

Normal breast marker genes targeted by the IncRNAs consistently expressed only in the C4 population

Table S12:

Sheet Name Summary of Contents

FACS CellLabels

ClusterAnnotation_FACS

PhysiologicalCharacteristics

Based on currently used laboratory clusters, how would each cluster ClusterAnnotation LabMarkers

BasalMarkers Pal2021

LumProgenitorMarkers Pal2021 LumMatureMarkers Pal2021

RefClusters_LiteratureMarkers

ClusterCorrespondence

Flow cytometry labels provided in Nguyen, 2018 for each cell.

Based only on information from the FACS labels, how we characterized each cluster and the corresponding confusion matrices.

Overlap between marker genes and genes in investigated physiological characteristics of basal and luminal cell types.

be annotated.

List of basal markers from Pal et al., 2021

List of luminal progenitor markers from Pal et al., 2021 List of luminal mature markers from Pal et al., 2021

List of literature markers from the in-house database and which cluster has each gene as Serurat marker.

How cells from A-clusters were divided into L-clusters.

Table S13:

Sheet Name Summary of Contents

List of markers assigned by Seurat for each cluster obtained based on SeuratMarkers_Aclusters GENCODE-annotated gene expression (A-clusters).

List of markers assigned by Seurat for each cluster obtained based on SeuratMarkers_Lclusters NB-IncRNA expression (L-clusters).

List of markers assigned by Seurat for each cluster obtained based on merged GENCODE-annotated gene expression and NB-IncRNA SeuratMarkers_Mclusters expression (M-clusters).

List of markers assigned by Seurat for each cluster obtained based on SeuratMarkers_Oclusters the expression of GENCODE-annotated genes that are neither confirmed protien-coding or IncRNAs (O-clusters).

List of markers assigned by Seurat for each cluster obtained based on SeuratMarkers AnnotatedLncRNAs GENCODE-annotated IncRNAs.

Table S14:

Sheet Name Summary of Contents

MarkersOfInterest_NBdb_FEELnc Seurat-assigned NB-lncRNA markers with FEELnc-predicted targets in the database of normal breast marker genes.

Seurat-assigned NB-IncRNA markers expressed in all subpopulations of

MarkersOfInterest_BroadCellType each cell type.

MarkersOfInterest_UniqToCluster

Seurat-assigned NB-lncRNA markers that have expression confined to each cluster.

Table S15:

Sheet Name Summary of Contents

WidespreadExpression_NBIncRNAs List of NB-IncRNAs expressed in more than 1/3 of the cells and their FEELnc-assigned targets.

WidespreadExpression_PCGs List of GENCODE-annotated genes expressed in more than 1/3 of the cells and their gene names.

Human Housekeeping Genes Human housekeeping genes, deposited at HK and/or HRT.

Overlap_WE_NBIncRNA_HK_GeneDB

Overlap_WE_NBIncRNA_HK_GeneDB

Overlap_WE_NBIncRNA_HK_GeneDB

Overlap between the list of NB-IncRNAs of widespread expression and housekeeping genes (Sheet 4). The overlaps are shown for genes co-expressed with NB-IncRNAs (left) and for predicted targets of the NB-IncRNAs (right).

Table S16:

Sheet Name Summary of Contents

Brain_SeuratMarkers_Aclusters

List of markers assigned by Seurat for each cluster obtained based on GENCODE-annotated gene expression (A-clusters).

Brain_SeuratMarkers_Lclusters

List of markers assigned by Seurat for each cluster obtained based on

NB-IncRNAs expression (L-clusters).

CellTypes_PerCluster

Distribution of cell types (defined based on provided labels from original files) in each cluster.

/-----Name mes procluster

KnownMarkers_PerCluster Number of known markers (in Darmanis, 2015) in each cluster.

MarkersOfInterest Seurat-assigned markers not necessarily mentioned in Darmanis, 2015

which we found to be of interest.

Table S17:

Sheet Name Summary of Contents

SelectedStemCellMarkers Contains the list of genes selected for root-state determination, based

on experimental data.

ClusterSRvalues SR values calculated by SCENT for each cell, averaged per cluster.

Table S18:

Sheet Name Summary of Contents

PearsonCorrelation_Cluster_TCGA For each L-cluster, the Pearson correlation coefficient and associated

p-value for its correspondence with each TCGA subtype.

subtype the top 300 most frequently assigned markers.

TCGAmarkers_GENCODEannotated GENCODE-annotated genes defined in-house as markers of TCGA subtypes (top 300 most frequently assigned markers per subtype).

P-values of overlaps between in-house markers of TCGA subtypes and Correspondence_wWuetalMarkers markers in Wu et al., 2021, for both GENCODE-annotated and NB-

IncRNA markers.
P-values of overlaps between in-house NB-IncRNA markers of TCGA

Correspondence_wSeuratlMarkers subtypes and Seurat-assigned markers of L-clusters.

List of MGFR-assigned NB-IncRNA markers used for subtype separation with connection to breast cancer or other features of interest. The top markers contributing with the first two principal components are

bolded.