



Super-enhancer LncRNA Microarrays

- The master gene regulators unmasked

Highlights

- Powerful SE-IncRNA collection to simultaneously profile all SE-IncRNAs, target genes, transcription factors and oncogenes.
- Detailed annotations of SE-IncRNAs, super-enhancers, and targets.
- Unambiguous and reliable SE-lncRNA isoform detection.
- Best solution for sensitive detection of labile SE-lncRNAs, powered by efficient labeling.

Introduction

Super enhancers are the transcription regulatory regions consist of constituent enhancers often clustered within 12.5 kb distance [1]. Super-enhancers produce long-noncoding RNAs (SE-lncRNAs) which are the hallmark of their activity. SE-lncRNAs, together with super-enhancers, activate the gene expression by multiple mechanisms. SE-lncRNAs are increasingly found to *trans*-activate target genes beyond *cis*-activation by typical enhancers [2](Fig. 1). SE-lncRNAs play master regulator roles in diverse gene expression programs that ultimately determine the cell types, cell identities, and diseases. SE-lncRNAs are also disease regulators and intimately involved in cancers.

Expression profiling of SE-lncRNAs provides a comprehensive view of the differential expressed SE-lncRNAs and what super-enhancers have turned active. Due to the very high tissue/cell/disease specificity restricted in only a few cell types, SE-lncRNAs can be an excellent class of biomarkers.

Powerful SE-IncRNA collection

Arraystar Super-Enhancer LncRNA microarrays are the first commercially available solution for sensitive and accurate SE-lncRNA profiling (Fig. 2). The SE-lncRNAs are compiled on the foundation of Arraystar Gold standard & Reliable lncRNA collections and authoritative super-enhancer databases. "Super lncRNAs", a special class of SE-lncRNAs forming RNA:DNA:DNA triplex with the super-enhancers, are also included [3].

To experimentally observe, analyze, and correlate with the superenhancer associated target gene expression, the nearest protein coding genes are simultaneously profiled on the same array. As super-enhancers are intimately related with transcription factors and oncogenes, all known transcription factors and oncogenes are collected, regardless in *cis* or *trans* with the super-enhancers.

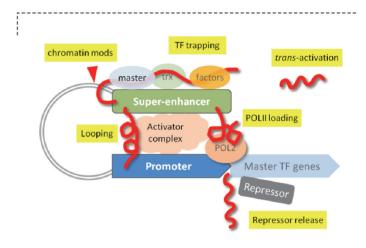


Fig. 1. Super-enhancer lncRNA activates the target gene promoter by multiple mechanisms.

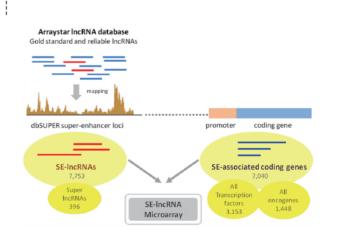


Fig. 2. Arraystar Human Super-enhancer LncRNA microarray features rich content and analysis of SE-lncRNAs, super lncRNAs, associated target genes, all transcription factors and oncogenes.





- Nearby target genes: To quickly locate nearby coding mRNA genes overlapping with or within 50-kb distance of the SE-lncRNAs as the potential *cis*-targets of the SE-lncRNAs.
- "Super-IncRNAs": A special class of SE-IncRNAs that form RNA:DNA:DNA triplex with the super-enhancers [3], which warrant special research on their mechanisms on super-enhancers.
- Subcellular localization: An important clue in knowing the functional sites and mechanisms of the SE-lncRNAs, which are tightly coupled with the subcellular localization.
- Tissue and cell type specificity: The key links to the high degrees of tissue or temporal specificity exhibited by super-enhancer activities.
- Cancer and disease SE-IncRNAs: Association with clinical or translational medicine sourcing databases such as Lnc2Cancer database and IncRNADisease.

Best and sensitive solution for labile SE-IncRNAs

Enhancer RNAs typically have short half-lives, high turnover rates and low abundance, which pose technical challenges if profiled by RNA-seq. Generic RNA-seq is simply not up for the task. Special techniques, such as GRO-seq are required. However, global run-on labeling of nascent transcripts in *vitro* cell culture is not feasible for tissues or clinical samples.

To sensitively detect and accurately quantify the transient, low level SE-IncRNAs, Arraystar has developed an efficient and robust linear amplification method to generate fluorescent cRNA for use with the microarrays, with a sensitivity > 100-fold higher than conventional methods.

SE-IncRNA Roadmap

As a roadmap to research forward, the differentially expressed SE-IncRNAs can be used to easily indicate the active state of the super-enhancers, study the gain or loss of functions, investigate the mechanisms of action, and explore biomarker applications. The super-enhancer IncRNAs, emerged from hidden just recently, are now a scientific gold mine in the research of any fields of biology and disease (Fig. 4).

References

- [1] Whyte W.A. et al. (2013) Cell [PMID: 23582322]
- [2] Alvarez-Dominguez J.R. et al. (2017) Cell Rep [PMID: 28636939]
- [3] Soibam B. (2017) RNA [PMID: 28839111]

SE-IncRNA						
TransID	Subcellular	Tier	Tissue specificity	Cancer		
ENST00000549804	Nucleus		2:pituitary_gland[6.29]			
ENST00000487240			25:nervous_system[275.20]	breast		
NR_126491		Gold	7:cardiac_chamber[49.69]			
ENST00000430027	Nucleus		19:nervous_system[196.11]	lung		
ENST00000502463	Nucleus		1:mammary_gland_epithelial_cell[5.30]	colon		
uc022bbo.1	Cytosol					
uc003ysb.3		Reliable				

SE-IncRNA		Super-Enhancer	Target Gene	
Disease	Super- LncRNA	Genomic	Overlapped	Proximal
Angelman syndrome	yes			SNRPN
Alzheimer's disease	yes		DNAJC19	
acute myocardial infarction		chr14:23867308:23891809	МҮН7	
split hand/split foot malformation		chr7:96630715:96657145	DLX6	
		chr8:128743538:128752334		MYC
AIDS		chr8:132815762:132873739		EFR3A
		chr8:128180346:128229384		POU5F1B

Fig. 3. A snapshot of comprehensive SE-lncRNA annotations.

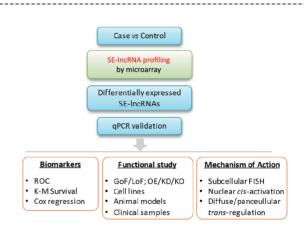


Fig. 4. Roadmap of super-enhancer lncRNA Research.

	Specifications	Human	Mouse
	Total Number of Probes	14,873	14,637
	SE-IncRNA probes -Arraystar Gold and Reliable IncRNA collections -Mapped to dbSUPER super-enhancer catalog	7,753	8,222
	target gene probes -Tfcheckpoint Transcription factors -Cancer genes (allOnco,SBCSSB)	7,040	6,385
	Array format	8 ×15K	8 × 15K

 $Fig.\ 5.\ Specifications\ of\ Arraystar\ Super-enhancer\ LncRNA\ Microarrays.$

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