

Supplementary Tables and Figures

Supp. Table S1. Optimized network parameters to obtain HUB network

Column1	Network parameter	Range
1	Degree	Between 30 and 93 inclusive
2	Betweenness Centrality	Between 0.003 and 1 inclusive
3	Closeness Centrality	Between 0.3 and 1 inclusive
4	Eigenvector	Between 0.009 and 0.219 inclusive

Supp. Table S2. Top 10 enriched pathways of Gene Ontology terms based on False discovery rate. Nanog is present in enriched pathways. Their row is marked by *

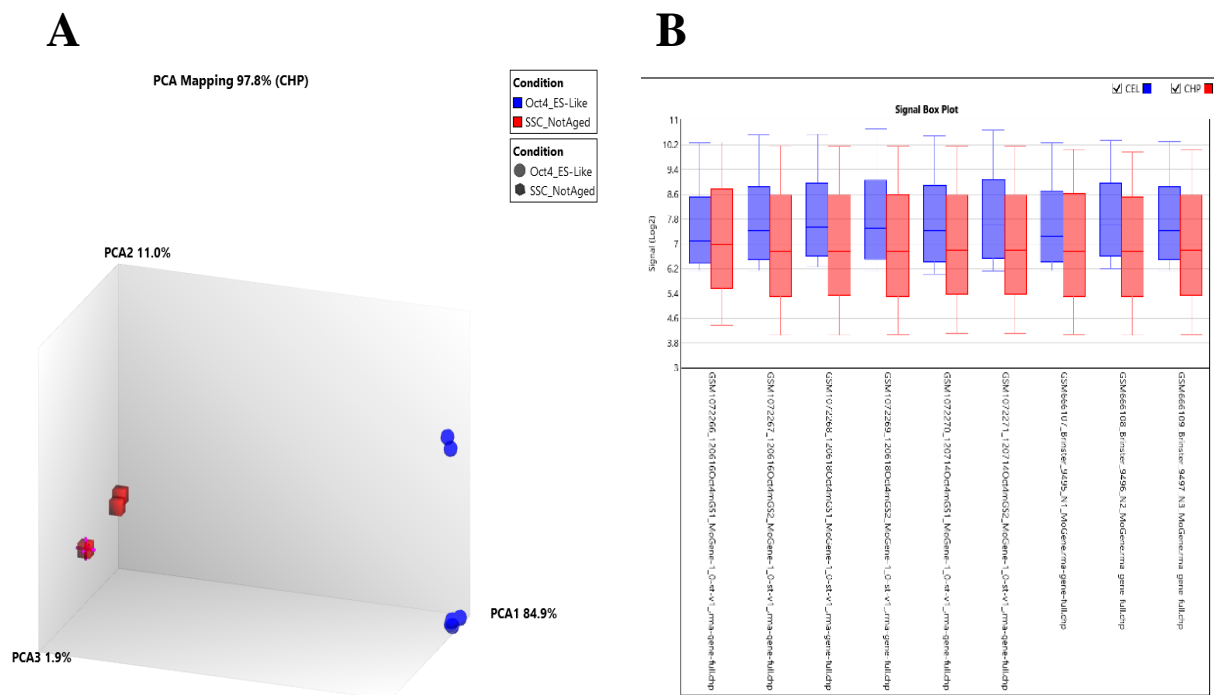
Column1	Category	Term ID	Term description	Observed gene count	Strength	False discovery rate
1*	GO Function	GO:0005515	Protein binding	41	0.37	5.90E-08
2*	GO Function	GO:0005488	Binding	50	0.21	6.74E-08
3	GO Function	GO:0005102	Signaling receptor binding	19	0.7	2.44E-06
4	GO Function	GO:0008083	Growth factor activity	8	1.36	3.46E-06
5	GO Function	GO:0008201	Heparin binding	7	1.25	0.00016
6*	GO Function	GO:0003682	Chromatin binding	11	0.86	0.00021
7	GO Function	GO:0044877	Protein-containing complex binding	16	0.64	0.00023
8	GO Function	GO:0050839	Cell adhesion molecule binding	8	1.05	0.00037
9	GO Function	GO:0005104	Fibroblast growth factor receptor binding	4	1.81	0.00039
10	GO Function	GO:0005178	Integrin binding	6	1.22	0.00087
1	GO Component	GO:0032991	Protein-containing complex	35	0.43	2.93E-07
2*	GO Component	GO:0000785	Chromatin	13	0.91	5.51E-06
3*	GO Component	GO:0005694	Chromosome	15	0.68	0.00019
4	GO Component	GO:0005615	Extracellular space	17	0.61	0.00019
5	GO Component	GO:0009986	Cell surface	13	0.72	0.00026
6	GO Component	GO:0009897	External side of plasma membrane	9	0.88	0.00089
7	GO Component	GO:0031012	Extracellular matrix	9	0.87	0.0009
8	GO Component	GO:0005583	Fibrillar collagen trimer	3	2.04	0.0011
9	GO Component	GO:0098552	Side of membrane	10	0.77	0.0011

10	GO Component	GO:0005576	Extracellular re- gion	18	0.48	0.0017
1*	GO Process	GO:0010033	Response to or- ganic substance	34	0.66	4.66E-13
2*	GO Process	GO:0070887	Cellular re- sponse to chemi- cal stimulus	32	0.69	5.68E-13
3*	GO Process	GO:0051173	Positive regula- tion of nitrogen compound meta- bolic process	33	0.64	4.10E-12
4*	GO Process	GO:0071310	Cellular re- sponse to or- ganic substance	28	0.75	4.71E-12
5*	GO Process	GO:0070848	Response to growth factor	17	1.15	6.10E-12
6	GO Process	GO:0030334	Regulation of cell migration	21	0.96	6.10E-12
7*	GO Process	GO:0010604	Positive regula- tion of macro- molecule meta- bolic process	34	0.6	6.10E-12
8*	GO Process	GO:0071363	Cellular re- sponse to growth factor stimulus	16	1.15	2.25E-11
9*	GO Process	GO:0009653	Anatomical structure mor- phogenesis	28	0.71	2.25E-11
10*	GO Process	GO:0048522	Positive regula- tion of cellular process	40	0.47	2.25E-11

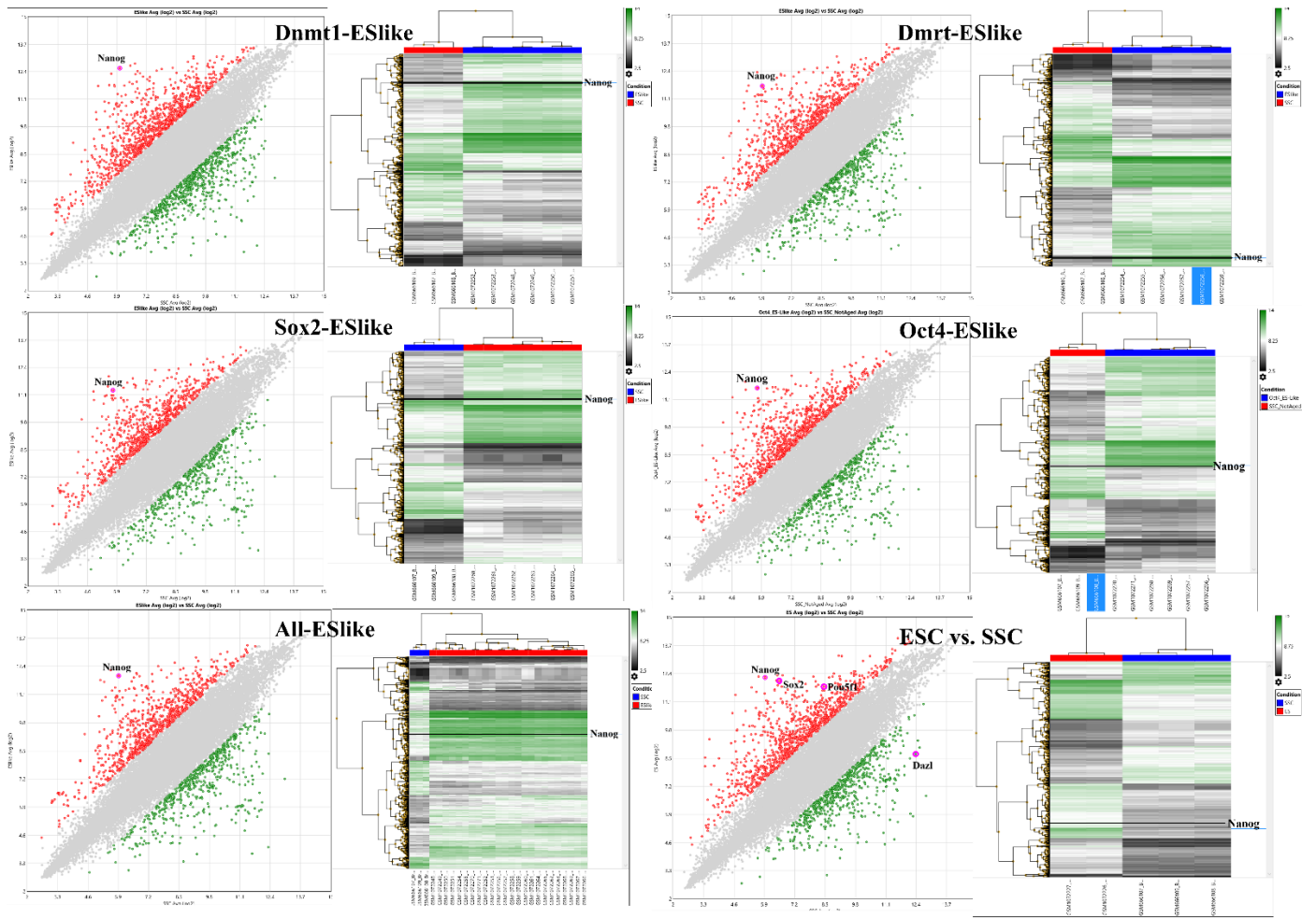
Supp. Table S3. Top 10 enriched TISSUES, KEGG, and Wikipathways pathways based on False discovery rate. Nanog is present in enriched pathways that their row is marked by *. The last row of Wikipathways (11**) represents an enriched pathway that wasn't among the top 10

TISSUES, KEGG and Wikipathways						
Col- umn1	Category	Term ID	Term description	Observed gene count	Strength	False discovery rate
1*	TISSUES	BTO:000232 2	Cell property	10	1.97	1.25E-13
2*	TISSUES	BTO:000607 8	Pluripotent stem cell	10	1.51	1.00E-09
3*	TISSUES	BTO:000000 0	Tissues, cell types and enzyme sources	45	0.34	1.00E-09
4*	TISSUES	BTO:000066 9	Embryonic cell line	9	1.56	3.11E-09
5*	TISSUES	BTO:000148 9	Whole body	44	0.33	3.11E-09
6	TISSUES	BTO:000148 6	Skeletal system	13	1.12	4.75E-09

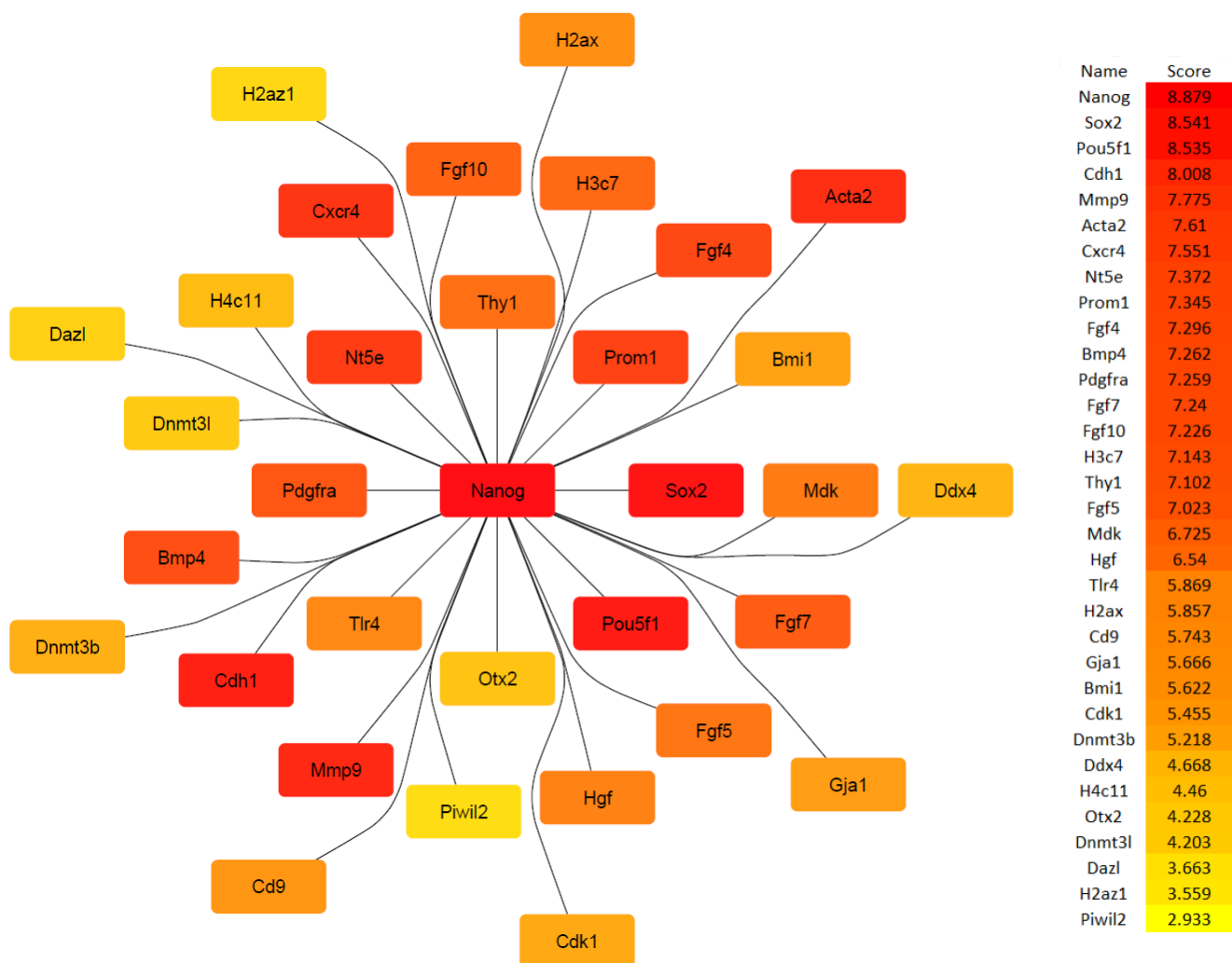
7	TISSUES	BTO:000205 0	Osteogenic cell	8	1.63	5.97E-09
8*	TISSUES	BTO:000028 4	Organism form	22	0.72	5.97E-09
9	TISSUES	BTO:000159 3	Osteoblast	7	1.75	1.60E-08
10*	TISSUES	BTO:000109 9	Blastocyst	6	1.97	2.32E-08
1	KEGG	mmu05218	Melanoma	7	1.63	1.62E-07
2	KEGG	mmu05200	Pathways in cancer	12	1	3.78E-07
3	KEGG	mmu04151	PI3K-Akt signaling pathway	10	1.09	8.76E-07
4	KEGG	mmu04015	Rap1 signaling pathway	8	1.23	2.46E-06
5*	KEGG	mmu05205	Proteoglycans in cancer	7	1.19	2.52E-05
6	KEGG	mmu05226	Gastric cancer	6	1.25	7.21E-05
7	KEGG	mmu05144	Malaria	4	1.53	0.00038
8	KEGG	mmu04810	Regulation of actin cytoskeleton	6	1.09	0.00041
9	KEGG	mmu04014	Ras signaling pathway	6	1.07	0.00051
10*	KEGG	mmu04550	Signaling pathways regulating plu- riipotency of stem cells	5	1.2	0.00062
1*	WikiPath- ways	WP1763	Mechanisms associated with pluripo- tency	11	1.22	1.15E-08
2	WikiPath- ways	WP2841	Focal adhesion: PI3K-Akt-mTOR sig- naling pathway	10	1.14	2.82E-07
3	WikiPath- ways	WP339	ESC pluripotency pathways	6	1.36	2.18E-05
4	WikiPath- ways	WP5242	Comprehensive IL-17A signaling	5	1.33	0.00021
5	WikiPath- ways	WP3632	Lung fibrosis	4	1.46	0.00055
6	WikiPath- ways	WP523	Regulation of actin cytoskeleton	5	1.16	0.00091
7	WikiPath- ways	WP458	Inflammatory response pathway	3	1.64	0.0017
8	WikiPath- ways	WP85	Focal adhesion	5	1.07	0.0018
9	WikiPath- ways	WP2432	Spinal cord injury	4	1.25	0.0019
10	WikiPath- ways	WP2375	miRNAs and TFs in iPS Cell Genera- tion	2	2.1	0.0036
11**	WikiPath- ways	WP723	Wnt signaling pathway and pluripo- tency	3	1.13	0.0206



Supp. Figure S1. (A) The results of the principal component analysis (PCA), indicate that 97.8% of the variance is accounted for by the principal components. (B) Data normalization results of the microarray data.



Supp. Figure S2. Confirmation analysis was conducted on samples using different combinations to test our main analysis. The results revealed consistent gene expression patterns across different combinations and samples. Specifically, Nanog and other crucial genes within the graphs exhibited similar patterns to those observed in the main figures of the analysis.



Supp. Figure S3. The hub gene network of Nanog and its connected nodes are visualized by the EPC method of the Cytohubba app. Darker colors represent higher scores and as a result, higher importance and fainter colors have lower scores, which represents lower importance in the network.