Super-enhancers modulate interleukin-6 expression and function in cancers

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**Background:** It is widely accepted that inflammatory cytokine, interleukin 6 (IL-6), was not only elevated in cancer but also important in carcinogenesis. But how did IL-6 be produced in tumor microenvironment remains to be addressed.

**Methods:** Both bioinformatics tools and quantitative real time polymerase chain reaction (RT-PCR) were used to examine the expression of IL-6 in cancer cells. To map super-enhancers of IL-6, sgRNAs were constructed. Stable knockout cells were established and subsequently used for cell proliferation and colony formation assay. The correlation between mapped super-enhancers and IL-6 expression was studied by ATAC-seq analysis.

**Results:** The expression of IL-6 was high in multiple cancers, including pancreatic cancer (PAAD). The elevated expression of IL-6 in PAAD was further confirmed by transcriptional data and in a panel of pancreatic cancer cell lines (one immortal HPDE6-C7 cell line and four PDAC cell lines: BxPC-3, PANC-1, AsPC-1 and CFPAC-1). When treated with JQ-1 and I-BET-762, two inhibitors of super-enhancers, the expression of IL-6 in multiple cancer cells including CFPAC-1, HeLa and SUM-159 cells was significantly reduced. By analyzing the H3K27Ac profiling, BRD4 binding, Med1 binding and DNA conservation in CFPAC-1, HeLa and SUM-159 cells, we identified a potential super-enhancer (IL6-SE) that might be important for IL-6 expression in cancer. The super-enhancer (IL6-SE) can be further divided into two elements (IL6-SEA and IL6-SEb). Genetic deletion of IL6-SEA in cancer cells greatly reduces the expression of IL-6. IL6-SEA deficient cells also showed low proliferation and colony formation ability. In patients, the epigenetic activation (ATAC signal) of IL6-SEA is correlated with the expression of IL-6.

**Conclusions:** In summary, we not only provide convincing experimental evidence to demonstrate that IL-6 expression in cancer is dependent on super-enhancers but also identified IL6-SEA as a critical DNA regulatory element. Our findings provide new insights to understand the epigenetic regulation of IL-6 expression in cancers.

**Keywords:** IL-6; super-enhancers; BRD4; pancreatic cancer

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Introduction

Cancer is the leading cause of death in developed countries. Pancreatic cancer contributes largely to cancer-related death and becomes the fourth leading cause of cancer mortality, although it is the tenth most commonly diagnosed cancer (1). Therefore, it is an extremely malignant tumor and its overall 5-year survival is only 5% (2). Pancreatic cancer is almost resistant to all current therapeutic regimens and this is likely due to broad heterogeneity of genetic mutations and dense stromal environment (3). So far, no molecular-targeted therapies have given satisfactory results and gemcitabine which approved at 1997, was still the first-line drug for pancreatic cancer patients at a good performance status (4,5). Inflammatory conditions such as pancreatitis not only represent a well-known risk for pancreatic cancer development but also as a consequence for pancreatic cancer (6). This is supported by epidemiological evidence which shows that patients suffering hereditary autoimmune pancreatitis or patients with chronic pancreatitis carry 40% of lifetime risk or a 13-fold higher risk for PDAC development, respectively (7,8). Furthermore, the inflammatory tumor microenvironment contributes significantly to the chemotherapy resistance and cancer relapse (9). Inflammatory tumor microenvironment-inducing oncogenic signal pathways such as nuclear factor-kB (NF-kB) and STAT3 contribute to not only inherent resistance but also acquired resistance (10). However, how inflammatory microenvironment was generated and how it affects carcinogenesis and chemotherapy resistance are largely unknown.

The encoding gene for interleukin 6 (IL-6) locates on chromosome 7p21 and its protein product has pleiotropic effect on immune response and inflammation. In the acute phase of stimulation such as infection and tissue injuries, IL-6 was promptly and transiently produced. Besides its critical role in immunity, IL-6 widely involved in cancer progression and therapeutic resistance. IL-6 regulates multiple cancer-associated processes such as proliferation, survival, apoptosis, invasiveness and metastasis, angiogenesis as well as cell metabolism (11-14). Its high expression was frequently observed in cancer both in patients’ serum and tumor tissues (15-17). However, the molecular mechanism that drives IL-6 production in cancer remains to be understood.

The elevated expression of IL-6 in tumor microenvironment can be induced by many stimuli that associated with tissue damage and stresses such as ionizing radiation, viruses, reactive oxygen species and other pro-inflammatory cytokines (18). Its expression is mainly induced through transcriptional regulation via key transcriptional factors such as NF-kB, CCAAT/enhancer-binding protein A and AP-1 (activator protein-1) (18). There are several small cis-acting elements that locating in a 1.2-kb fragment of the 5'-flanking region, which cooperatively with transcriptional factors to induce IL-6 expression (19,20). Super-enhancers are recently defined large cis-acting elements that have powerful ability to induce much high gene expression (10). However, it is unclear whether do super-enhancers play critical role in the induction of IL-6 in cancer cells.

Methods

Cell culture and transfection

HPDE6-C7, PANC-1, BXPC-3, AsPC-1, CFPAC-1, HeLa and SUM-159 cells were maintained in Dulbecco Modified Eagle Medium containing 10% fetal bovine serum at 37 °C in a 5% CO2 condition. HeLa cells were transfected with Lipofectamine 2000 (Invitrogen) according to the manufacturer’s manuals. The BRD4 inhibitors JQ1 (A1910) and I-BET-762 (B1498) were bought from ApexBio.

Quantitative real time polymerase chain reaction (RT-PCR)

For RNA extraction, cells will be seeded into 6-cm dishes and harvested after achievement of 80% confluence. Trizol (Ambion) was used to extract the total RNA from indicated cells. The synthesis of cDNA was performed in accordance with the instructions for reverse transcription kits supplied by Thermo. The primers for IL-6 and GAPDH were designed by the primer design program. Primers sequences as follows: human IL-6: 5’-AGCCAGAGCTGTGCAGATGAG-3’ (forward), 5’-GTTGTCATGTCCTGCAGCCAC-3’ (reverse); human GAPDH: 5’-GTGAACCATGAGAAGTATGAC-3’ (forward), 5’-AGTGATGGCATGGACTGTGGT-3’ (reverse).

Construction of sgRNA

The DNA sequence of IL6-SE was subjected to sgRNA finder. The oligos for designed sense and antisense were synthesized and annealed in the annealing buffer: 10mM Tris (pH 7.5), 1 mM EDTA and 50 mM NaCl. The epiCRISPR vector was digested with BSPQ1 (New...
England Biolads) and purified by Gel purification kits. The information of epiCRISPR vector was published previously (21). The annealed oligonucleotides were inserted into the CRISPR/Cas9 vector via T4 DNA ligase (New England Biolads) overnight at 16 ℃. The plasmid colonies were examined by digestion of BSPQ1. The sgRNA sequences were confirmed by direct sequencing. The cloned sgRNA sequences are shown as follows:

SgRNA-1: GGCACAGTTATATACATATT;
SgRNA-2: GCTTTGCTTTGTGTTT.

Transfection of sgRNA and genotyping

For knockout of interested super-enhancers, HeLa cells were placed in 6-well plates and were transfected with indicated combination of sgRNAs using Lipofactamine 2000. Empty epiCRISPR vector was transfected parallel as a control. After 48 hours, cells were killed by puromycin (InvivoGen) and remaining cells were cultured with puromycin for additional two weeks. Then, total RNA from selected cells were extracted by Trizol and followed by RT-PCR examination. To examine the deletion of indicated DNA region, the cells were digested in protease mixture (bimake.com) to obtain the genomic DNA according to the manufacturer's manuals. The genomic PCR were performed via 2XM-PCR OPTI Mix (bimake.com) based on the manufacturer's manuals. The PCR products were then sent for DNA sequencing to confirm the deletion. The sequences of genotyping primers are as follows: forward: 5’-GAGTCTAGCAAGAGTTGTCTT-3’, reverse: 5’-TGCCAGTAGAGCAGGTCTGCC-3’.

Cell proliferation and colony formation assay

For cell proliferation assay, cells were inoculated into 96-well plates with the number of 2×10^3 cells for each well. Cell Counting Kit-8 (CCK-8) was used to evaluate cell number with indicated days. For colony formation assay, stable cell lines were inoculated on six-well plates with a density of 5×10^3 per well. Cell colonies were allowed to grow for 2 weeks and then captured by microscopy.

RNA-seq analysis

We download the raw fastq file by using Aspera (https://asperasoft.com/). We next align reads to the human genome (Version: hg19) by using STAR (https://github.com/alexdobin/STAR) (22) and perform the quality control by using multiqc (https://multiqc.info/) (23). We quantify the gene reads by using FeatureCounts (24) (http://subread.sourceforge.net/). We use Deseq2 (25) to perform the differential gene expression analysis (https://bioconductor.org/packages/release/bioc/html/Deseq2.html). We them calculate the TPM expression of each genes, where we normalize for gene length first and then normalize for sequencing depth second. As for the RNA-seq analysis using The Cancer Genome Atlas data, we fetched the level 3 RNA-seq raw counts from The Cancer Genome Atlas (https://portal.gdc.cancer.gov/cart). We then use limma package (https://bioconductor.org/packages/release/bioc/html/limma.html) to perform the differential expression analysis. We fetched the clinical annotation data of patients from The Cancer Genome Atlas (https://portal.gdc.cancer.gov/cart). The log-rank test was performed to examine the prognostic significance of IL-6.

ChIP-seq analysis and super-enhancer identification

The fastq file including raw ChIP-seq data was downloaded by using Aspera (https://asperasoft.com/). We next align reads to the human genome (Version: hg19) by using Bowtie2 (26) (http://bowtie-bio.sourceforge.net/bowtie2/index.shtml). We use Samtools (27) (http://samtools.sourceforge.net/) to convert files in to sam and use bedtools (28) (https://bedtools.readthedocs.io/en/latest/) to convert files in to bed file type. We next convert the bed file into bigwig file by using bedGraphToBigWig (http://hgdownload.cse.ucsc.edu/admin/exe/) and ATAC-seq analysis, we download peak data from The Cancer Genome Atlas (https://portal.gdc.cancer.gov/cart) and perform the correlation analysis by using ggpubr (https://github.com/kassambara/ggpubr) in R. To identify the potential super-enhancers, we conduct the peak calling by using macs14 (29) (http://liulab.dfc.iharvard.edu/MACS/00README.html). We next rank enhancers and identify super-enhancers by using ROSE (10) (http://younglab.wi.mit.edu/super_enhancer_code.html).

Results

High expression of IL-6 predicts poor outcome in multiple cancers

IL-6 has been widely reported to associate with cancer,
but its regulation is still not fully addressed. To fully understand the expression of IL-6 across different cancer, we systematically analyzed the IL-6 expression using TCGA data. As shown in Figure 1A, the expression of IL-6 was significantly upregulated in cancers including pancreatic cancer (PAAD), testicular germ cell tumors (TGCT), esophageal carcinoma (ESCA) and glioblastoma (GBM). The upregulation of IL-6 in PAAD was further confirmed by transcriptional data from individual patients (Figure 1B). Furthermore, high expression of IL-6 in PAAD predicts poor outcome (Figure 1C). To confirm the finding from public data, we examined the expression of IL-6 in one immortal HPDE6-C7 cell line and four PDAC cell lines including BxPC-3, PANC-1, AsPC-1 and CFPAC-1. Compared to control cells, PDAC cells show elevated expression of IL-6 in generally (Figure 1D). Interestingly, the CFPAC-1 cells show extremely high expression of IL-6. Together, our results reveal that IL-6 was up-regulated in several cancer and its expression associated with patients’ outcome.

**IL-6 expression in cancers is dependent on super-enhancers**

The molecular mechanism that drives high expression of IL-6 in cancer is still not fully addressed. We tested whether IL-6 expression in cancer was dependent on super-enhancers. We treated the PDAC cells (CFPAC-1) using inhibitors (JQ-1 and I-BET-762) of super-enhancers. JQ-1 treatment greatly reduces the expression of IL-6 in CFPAC-1 cells (Figure 2A). Using another super-enhancer inhibitor (I-BET-762), we got similar result (Figure 2A). Next, we determined whether super-enhancer-dependent IL-6 expression is cancer type dependent. We treated HeLa and SUM-159 cells with JQ-1 and I-BET-762. We found that both JQ-1 and I-BET-762 significantly reduces IL-6 expression in both HeLa and SUM-159 cells (Figure 2B,C). Our results suggest that super-enhancers drive high expression of IL-6 is a general mechanism in cancer. Our results were further supported by published GEO dataset regarding SUM-159 cells treated with or without JQ-1 (Figure 2D). RNA-seq clearly demonstrates the reduced expression of IL-6 after JQ-1 treatment in SUM-159 cells (Figure 2D). ChIP-seq also reveals that JQ-1 treatment greatly reduces the BRD4 and P300 binding in super-enhancer regions (Figure 2E,F,G,H). Collectively, our finding suggests a critical role of super-enhancers in the upregulation of IL-6 in cancers.

**Potential super-enhancers around IL-6 gene (IL6-SE) in cancers**

Next, we tried to identify the critical super-enhancers that important for IL-6 expression in cancer. Because, the expression of IL-6 in CFPAC-1 is significantly higher than in PANC-1 cells. Super-enhancers are large DNA regulatory elements enriched with H3K27Ac. Thus, we reasoned that the potential large DNA elements enriched with H3K27Ac were exist in CFPAC-1 cells but not PANC-1 cell. Therefore, we compared the H3K27Ac modification around IL-6 gene locus (Figure 3A). Interestingly, we identified a region at Chr7: 22,615,000–Chr7: 22,630,000 (IL6-SE) in the 5’ end of IL-6 gene was enriched with H3K27Ac at CFPAC-1 cells but not PANC-1 (Figure 3A). In HeLa cells, this DNA region was also enriched with H3K27Ac (Figure 3B). The DNA sequence of this region was conserved between human and mouse (Figure 3C). To further characterize this region, we analyzed in detail by different markers including BRD4, MED1 and H3K27Ac in SUM-159 cells (Figure 3C). Based on BRD4 enrichment and DNA conservation, this DNA region can be further divided into two sub-elements that we called IL6-SEa (Chr7: 22,625,000–Chr7: 22,630,000) and IL6-SEb (Chr7: 22,615,000–Chr7: 22,619,000) (Figure 3C). Collectively, we identified a potential important super-enhancer (IL6-SE) that might be important for IL6 expression in cancers. To verify that the IL-6-SE is a super-enhancer, we ranked the DNA elements based on the signal of H3K27Ac according to published protocol (ROSE, Ranking Of Super Enhancer) (Figure 3D). The exceptionally high amounts of H3K27Ac modification clearly demonstrates that IL6-SE is a super-enhancer in SUM-159, HeLa and L3-6 cells (Figure 3D). Consistently, JQ-1 treatment could significantly reduce BRD4 and P300 binding at IL-6-SE locus (Figure 3E). It suggests that IL6-SE is a functional super-enhancer. In summary, we identified a potential important super-enhancer that is IL6-SE in the upregulation of IL-6.

**Deletion of a super-enhancer greatly reduces IL-6 expression**

Next, we determined whether identified super-enhancers are important for IL-6 expression in cancer cells. We designed sg-RNAs to delete IL6-SEa and IL6-SEb in cancer cells (Figure 4A). The transfection efficiency was extremely low in CFPAC-1 and we failed to generate knockout colony in such cells. However, we successfully generated
Figure 1 The expression of IL-6 in cancer tissues and cell lines. (A) The raw data of IL-6 expression in 31 cancers was downloaded from TCGA database. Comparing to normal tissues, significant higher expression of IL-6 in pancreatic cancer (PAAD), testicular germ cell tumors (TGCT), esophageal carcinoma (ESCA) and glioblastoma (GBM) was observed. The other TCGA Cancer Codes are as following: STAD (stomach adenocarcinoma), DLBC (lymphoid neoplasm diffuse large B-cell lymphoma), CESC (cervical squamous cell carcinoma and endocervical adenocarcinoma), OV (ovarian serous cystadenocarcinoma), THYM (thymoma), CHOL (cholangiocarcinoma), UCEC (uterine corpus endometrial carcinoma), UCS (uterine carcinosarcoma), LIHC (liver hepatocellular carcinoma), KIRC (kidney renal clear cell carcinoma), LGG (brain lower grade glioma), SKCM (skin cutaneous melanoma), PRAD (prostate adenocarcinoma), READ (rectum adenocarcinoma), COAD (colon adenocarcinoma), ACC (adrenocortical carcinoma), LAML (acute myeloid leukemia), KIRP (kidney renal papillary cell carcinoma), THCA (thyroid carcinoma), KICH (kidney chromophobe), HNSC (head and neck squamous cell carcinoma), PCPG (pheochromocytoma and paraganglioma), SARC (sarcoma), BLCA (bladder urothelial carcinoma), LUSC (lung squamous cell carcinoma), LUAD (lung adenocarcinoma). (B) The expression of IL-6 in individual PAAD patients and controls were present. The expression of IL-6 is significantly higher in PAAD patients when compared to it in controls. (C) In PAAD patients, high expression of IL-6 predicts poor survival. (D) The mRNA level of IL-6 was measured by RT-PCR in HPDE6-C7, BxPC-3, PANC-1, AsPC-1 and CFPAC-1 cells. GAPDH was used as a loading control. *, P<0.05.
Figure 2 JQ-1 and I-BET-762 reduce IL-6 expression in cancer cells. (A) The mRNA level of IL-6 was measured by real-time polymerase chain reaction (RT-PCR) in CFPAC-1 cells treated with or without JQ-1 (or I-BET-762) with indicated doses. (B) The mRNA level of IL-6 was measured by RT-PCR in HeLa cells treated with or without JQ-1 (or I-BET-762) with indicated doses. (C) The mRNA level of IL-6 was measured by RT-PCR in SUM-159 cells treated with or without JQ-1 (or I-BET-762) with indicated doses. (D) RNA-sequencing data of SUM-159 cells treated with or without JQ-1 from GEO dataset was downloaded and presented. (E,F) Average ChIP-seq signal (BRD4 and P300) on all super-enhancers of SUM-159 cells treated with or without JQ-1. The data from GEO Number (GSE131097; GSE87418) was downloaded and re-analyzed. (G,H) Heatmap of ChIP-seq signal (BRD4 and P300) on all super-enhancers of SUM-159 cells treated with or without JQ-1.
Figure 3 The H3K27Ac profile and BRD4 binding around IL-6 locus. (A) The distribution of H3K27Ac around IL-6 gene in PANC-1 and CFPAC-1 cells was visualized through UCSC genome browser. The original H3K27Ac data of PANC-1 (GSM1254196) and CFPAC-1 (GSM2439222) cells was downloaded from GEO database. (B) The distribution of H3K27Ac around IL-6 gene in HeLa cells was visualized through UCSC genome browser. (C) The distribution of H3K27Ac modification, BRD4 and MED1 binding around IL-6 gene in SUM-149 cells was visualized through UCSC genome browser. The DNA conservation between human, Rhesus, Mouse, Dog and elephant was analyzed and visualized by VISTA tool (http://genome.lbl.gov/vista/index.shtml). (D) Distribution of H3K27Ac ChIP-seq density across all identified DNA elements. The super-enhancer is a subset of enhancers containing exceptionally high amounts of H3K27Ac modification. The IL6-SE is a potential super-enhancer based on H3K27Ac density using the published protocol (ROSE) in SUM-159, HeLa and L3-6 cells. (E) Precise analysis of H3K27Ac modification at the DNA region around IL-6 locus. The change of H3K27Ac modification before and after JQ-1 treatment were also analyzed. The core region (IL6-SE, Chr7: 22,615,000–22,630,000) of potential super-enhancer was highlighted by yellow color.
IL6-SEa knockout cells in HeLa cells. Genotyping and DNA sequencing clearly demonstrate that the IL6-SEa DNA region was deleted (Figure 4B,C). Deletion of IL6-SEa region greatly reduces IL-6 expression in HeLa cells (Figure 4D). The proliferation ability of IL6-SEa-deficient cells was significantly reduced compared to sgVec control cells (Figure 4E). Furthermore, the colony formation ability of IL6-SEa-deficient cells was greatly reduced compared to sgVec control cells (Figure 4F,G). Our results clearly demonstrate that IL6-SEa is functional important for IL-6 expression and function. To further support such conclusion, we analyzed the relationship between the chromatin status of IL6-SEa and IL-6 expression in patients (Figure 4H). The ATAC signal which indicating a loose and active status of IL-6-SEa is positively associated with IL-6 expression (Figure 4H). It suggests that the regulation of IL6-SEa on IL-6 expression is patient relevant. Taken together, we demonstrate that IL6-SEa was critical important for IL6 expression by genetic tools.

Discussion

IL-6 is a glycosylated polypeptide chain and was discovered as a B cell differentiation factor to induce their antibody production (30). IL-6 triggers signal transduction via a transmembrane receptor (mbIL-6R, also known as IL-6Ra) and a secretory soluble receptor (sIL-6R) (31,32). The mbIL-6R with a restricted expression and predominately expressed on hepatocytes, some epithelial cells and certain leukocytes (33). Upon IL-6 binding to mbIL-6R, the complex recruits gp130 and causes gp130 homodimerization to form an activated IL-6 receptor complex. GP130 is expressed ubiquitously and can also be activated by IL-6/sIL-6R in various cells including cancer cells (34,35). Enhanced levels of IL-6 have been found in many cancers and its high expression inversely related with the response to chemotherapy (36). Comparing to primary tumors, recurrent tumors and metastatic lesions show higher expression of IL-6 (37). However, the molecular mechanism inducing IL-6 expression in cancer is not fully understood. It is believed that IL-6 production is predominantly regulated by various transcription factors such as NF-xB, CCAAT/enhancer-binding protein-a and activator protein-1. Here, our results demonstrate a critical role of epigenetic mechanism that is super-enhancer in the up-regulation of IL-6 expression. The activation of super-enhancer DNA might be an initiation step to allow TFs binding to DNA. However, the relationship between identified IL-6 super-enhancer and reported TFs such as AP1 needs to be further studied.

The human IL-6 gene locates on chromosome 7p21 and has three transcription start sites and three TATA boxes (38). There is a 1.2-kb fragment of the 5’-flanking region that has multiple cis-acting elements (19). This DNA region including promoter has critical role in the regulation of IL-6 transcription. For example, there are two pairs of glucocorticoid response elements (GREs) locate at the positions –557 to –552 and –466 to –461 respectively (39,40). Besides GREs sites, this region tightly occupied by other identified regulatory sites such as AP-1 binding site (TGAGTCG) and cyclic AMP (cAMP)-response element (CRE: ACGTCG) (41). Many trans-acting factors have been identified to specifically regulate IL-6 expression through such DNA region. These trans-acting factors such as NF-xB, C/EBP-b, activator protein-1, interferon regulatory factor and estrogen receptor are closely associated with cancer (42-44). This DNA region and its trans-acting factors are extensively studied. Here, we identified a novel DNA region, IL6-SE, which is around 15 kb. IL6-SE is far away to promoter and the distance between IL6-SE and promoter is about 120 kb. In ATAC profiling, which shows chromatin accessibility, indicates IL6-SE is active in cancer cells and the status of IL6-SE correlated with IL-6 expression. It is interesting to know how IL6-SE changes its status from silence in normal cells to be active in cancer cells. One possibility is that the abnormal pioneer transcriptional factors in cancer cells specifically bind to IL6-SE region and make it available to transcriptional machine. However, the abnormal pioneer transcriptional factors remain to be explored. Based on DNA conservation and BRD4 binding, IL6-SE can be further divided into IL6-SEa and IL6-SEb. Although, we tried our best, we only established IL6-SEa knockout cells. Our results indicate that IL6-SEa is critical important for IL-6 expression and cancer cell growth. The role of IL6-SEb in IL-6 expression needs to be further studied. It is also interesting to known the relationship between IL6-SEa and IL6-SEb. Furthermore, the trans-acting factors act with IL6-SE DNA region are also largely unknown.

In summary, we provide convince evidence to show that super-enhancers play critical role in the induction of IL-6 in cancer. We also identified a novel DNA region (IL6-SE) that is important for IL-6 expression in cancer. Our results provide new insights to understand the epigenetic
Figure 4 Deletion of IL6-SEa impairs IL-6 expression in cancer cells. (A) Schematically presentation of the design of CRISPR/Cas9 mediated deletion of the IL6-SEa locus. The location of sgRNAs and primers that used to validate the deletion was indicated. (B) Genotyping the deletion of IL6-SEa locus by PCR. (C) Cloning and direct sequencing of the joint DNA sequences after deletion of the IL6-SEa super-enhancer. (D) Real-time PCR was used to determine the mRNA level of IL-6 in IL6-SEa deficient and control cells. (E) Cell proliferation of IL6-SEa deficient and control cells was evaluated by CCK-8 assay. (F,G) Anchorage-dependent colony formation of IL6-SEa deficient and control cells. (G) Crystal violet staining and the magnification ×2. (H) The correlation between ATAC signal at IL6-SEa region and the expression of IL-6 in breast cancer patients was analyzed. *, P<0.05; **, P<0.01.
regulation of IL-6 transcription.

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Footnote

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Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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