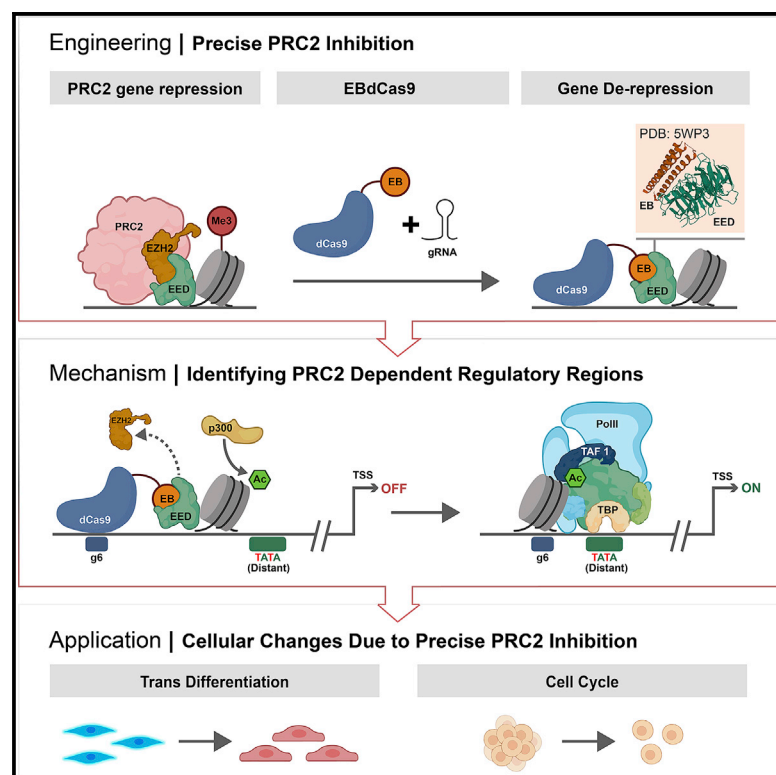


# dCas9 fusion to computer-designed PRC2 inhibitor reveals functional TATA box in distal promoter region

## Graphical abstract



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## In brief

Levy et al. fused a computationally designed protein, EED binder (EB), which competes with EZH2 and thereby inhibits PRC2 function, to dCas9 (EBdCas9). EBdCas9 represses PRC2 action in precise loci, remodels epigenomic marks, exposes transcriptional elements, and induces transdifferentiation.

## Highlights

- EBdCas9 inhibits PRC2 function in precise genomic locations
- EBdCas9 upregulates *TBX18*, *p16*, *CDX2*, and *GATA3* by de-repression
- EBdCas9 identifies PRC2-controlled, active TATA box >500 bp upstream of *TBX18* TSS
- EBdCas9 is sufficient to induce transdifferentiation and repress cancer cell cycle



## Article

# dCas9 fusion to computer-designed PRC2 inhibitor reveals functional TATA box in distal promoter region

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## SUMMARY

Bifurcation of cellular fates, a critical process in development, requires histone 3 lysine 27 methylation (H3K27me3) marks propagated by the polycomb repressive complex 2 (PRC2). However, precise chromatin loci of functional H3K27me3 marks are not yet known. Here, we identify critical PRC2 functional sites at high resolution. We fused a computationally designed protein, EED binder (EB), which competes with EZH2 and thereby inhibits PRC2 function, to dCas9 (EBdCas9) to allow for PRC2 inhibition at a precise locus using gRNA. Targeting EBdCas9 to four different genes (*TBX18*, *p16*, *CDX2*, and *GATA3*) results in precise H3K27me3 and EZH2 reduction, gene activation, and functional outcomes in the cell cycle (*p16*) or trophoblast transdifferentiation (*CDX2* and *GATA3*). In the case of *TBX18*, we identify a PRC2-controlled, functional TATA box >500 bp upstream of the *TBX18* transcription start site (TSS) using EBdCas9. Deletion of this TATA box eliminates EBdCas9-dependent TATA binding protein (TBP) recruitment and transcriptional activation. EBdCas9 technology may provide a broadly applicable tool for epigenomic control of gene regulation.

## INTRODUCTION

Chromatin modification mediated by the polycomb repressive complex 2 (PRC2) plays a critical role in developmental transitions. However, the specific genetic loci at which PRC2 function is critical for determining cell fate has remained unknown, as there has been no way to target PRC2 inhibition to specific chromosomal locations. A central question in epigenomics and developmental biology is the role of PRC2-dependent specific histone 3

lysine 27 methylation (H3K27me3) marks in cell fate decisions (Margueron and Reinberg, 2011). While broad upstream regions of developmental genes are decorated with H3K27me3 marks, it is not known which, if any, single nucleosomes require H3K27me3 marks for gene repression and cell-fate determination.

The two main complexes involved in polycomb-based repression are PRC1 and PRC2. PRC1 catalyzes monoubiquitylation of histone H2A Lys 119 (H2AK119ub), while PRC2 catalyzes the mono-, di-, and trimethylation of histone H3 Lys 27



(H3K27me1/me2/me3) (Lee et al., 2019). Recent structural analysis has revealed the mechanism of PRC1/PRC2 spatiotemporal regulation and H3K27me3 mark spreading (Banaszynski et al., 2013; Laugesen et al., 2016). Non-canonical PRC1 containing RYBP ubiquitinates H2K119 to recruit accessory proteins JARID2 and MTF2 (Coleman and Struhl, 2017; Laprell et al., 2017; Yu et al., 2019). JARID2 recruits PRC2 and contributes to the allosteric activation of PRC2 (Lee et al., 2018). PRC2 containing EED and EZH2 facilitates H3K27me3 marks by reading (EED) and writing/catalyzing (EZH2) new marks. H3K27me3 marks are then read by canonical PRC1 containing CBX, resulting in chromatin compaction, H2AK119ub, and gene repression (Brockdorff, 2017; Holoch and Margueron, 2017; Francis et al., 2009; Eskeland et al., 2010; Illingworth et al., 2015; Pengelly et al., 2015; Oksuz et al., 2018). During developmental transitions, the chromatin undergoes extensive epigenetic remodeling at the promoter and enhancer regions, where H3K27me3 marks are added or removed to produce the required decrease or increase in gene expression (Hawkins et al., 2010; Battle et al., 2019). Importantly, only the repressive marks, but not the activating histone tail marks, are instrumental for histone mark inheritance during development (Escobar et al., 2019). Recent studies suggest that proximal regulatory marks governing gene expression are associated with histone modification domains within 2.5 kb of known transcriptional start sites (Woo Jun Shim et al., 2019). However, it is not known if any specific H3K27me3-marked nucleosomes are critical for control of gene expression or if the entire broad 2.5 kb region is required for gene repression. This issue has been challenging to address as existing genetic and chemical methods eliminate all H3K27me3 marks without precision.

Recent studies that fuse catalytically dead Cas9 (dCas9) to proteins that promote or remove histone methyl groups provide tools for manipulating the epigenome. For example, the repressor KRAB recruits methyltransferases to increase H3K9me3 repression levels (Pengue and Lania, 1996; Groner et al., 2010), and its fusion to dCas9 results in 50%–99% gene repression in various cell lines, primary cultures, and pluripotent stem cells (Gao et al., 2013; Kearns et al., 2015; Shechner et al., 2015; THAKORE et al., 2015; Amabile et al., 2016; Pradeepa et al., 2016). Fusion of dCas9 to the transcriptional activators VP16, VP64, and VP65 promotes chromatin de-condensation, accumulation of H3K27ac and H3K4me3 marks, RNA polymerase II (RNA Pol II) recruitment, and transcriptional activation (Chavez et al., 2016; Gilbert et al., 2014). Likewise, gene expression was found to be regulated following the recruitment of histone acetyltransferase dCas9-p300 (Chen et al., 2019; Hilton et al., 2015). Similarly, fusion of dCas9 to the histone demethylase LSD1/KDM1A results in H3K4 and H3K9 methyl group removal and affects embryonic stem cell (ESC) self-renewal, differentiation, cancer cell proliferation, and development (Kearns et al., 2015; Adamo et al., 2011; Goodman and Smolik, 2000).

Despite the considerable array of tools available to modify the epigenome, currently there is no method to inhibit PRC2 function at a particular genomic locus and precisely at a single nucleosome in order to determine which specific H3K27me3 marks result in transcriptional repression. We have generated a computer-designed protein that binds EED (EED binder [EB]) and

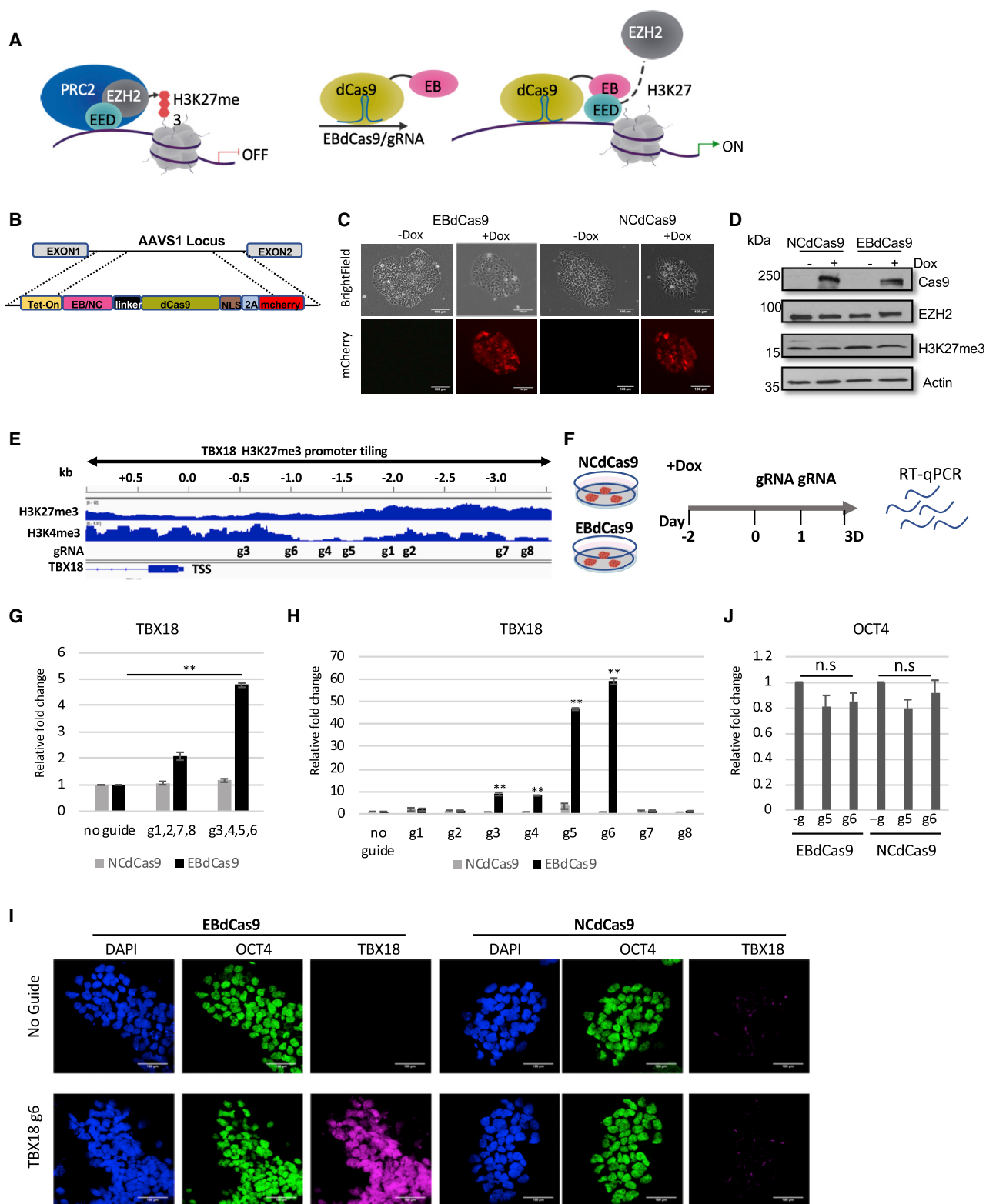
competes with EZH2 (Moody et al., 2017). By fusing EB to dCas9, we enable probing H3K27me3 function at precise gene loci. Here, we show that EBdCas9/gRNA is able to affect the PRC2 complex, remodel the histone marks of specific target sites, upregulate genes of interest, and promote epigenomic memory. We reveal precise loci for PRC2 requirement for transcription repression; in the *TBX18* gene, PRC2 activity normally represses the distal TATA box region. Using EBdCas9, we show that specific H3K27me3 marks are required for inner cell mass (ICM) versus trophectoderm (TE) cell-fate decision (Saha et al., 2013; Xiao et al., 2020) and proliferation control of the lethal pediatric CNS tumor diffuse midline glioma (DMG). These data suggest that EBdCas9 technology is broadly applicable for the regulation of histone marks at a single locus to control gene expression in different cell types.

## RESULTS

### EBdCas9/gRNA activates *TBX18* transcription

The catalytic and substrate recognition functions of PRC2, mediated by the SET domain containing EZH2 subunit and the trimethyl lysine-binding EED subunit, respectively, are coupled by binding of the N-terminal helix of EZH2 to an extended groove on EED (Kim et al., 2013). Disrupting PRC2 function can be achieved by chemical drugs or knocking out key PRC2 components (Kong et al., 2014; Knutson et al., 2013). We previously generated and characterized a computationally designed protein (EB) that binds to the EZH2 binding site on EED by two orders of magnitude higher affinity than EZH2 (Moody et al., 2017). As a control, we created an EB negative control (NC) where two amino acid mutations, F47E and I54E on the EED binding interface, abolish binding to EED (Moody et al., 2017). EB, but not NC, with subnanomolar affinity, forms tight complexes with EED, reduces EZH2 and JARID2 global levels, and exhibits a significant genome-wide reduction of H3K27me3 repressive marks in promoter regions (Moody et al., 2017).

To target EB to a specific chromatin locus, we fused EB to dCas9 (EBdCas9). This enables disruption of PRC2 function in specific regions within H3K27me3 domains to determine which, if any, control gene expression (Figure 1A). We fused EB to dCas9 with a 30 aa residue 6 × 5 (SGGGG) linker under the control of the AAVS1-TREG inducible promoter of the dCas9-NLS-2A-mCherry plasmid (Mandegar et al., 2016) (Figure 1B). Similarly, we fused the NC control to dCas9 to distinguish between dCas9 non-specific effects on chromatin and EB-specific effects on PRC2. The EB-linker-dCas9-NLS-2A-mCherry (EBdCas9) and NC-linker-dCas9-NLS-2A-mCherry (NCdCas9) constructs were transfected to induced pluripotent stem cells (iPSCs) (WTC) using TALENS to enforce recombinant homology at the AAVS1 site on chromosome 19. Following antibiotic selection, EBdCas9 and NCdCas9 mCherry expression were doxycycline (Dox) inducible, and stem cell morphology was maintained with no changes in EZH2 and H3K27me3 levels (Figures 1C, 1D, and S1A–S1D; Table S1) or gene expression in general (Figures S1E and S1F). Expression of EB alone has more global effects; the difference may reflect the 50-fold lower expression of EBdCas9 (Figure S1G) or the steric hindrance of the EB–EED interaction by the fused dCas9 when not bound to DNA (Cofsky Joshua



**Figure 1. EbdCas9 upregulates *TBX18* expression in precise loci**

(A) Model of EbdCas9 precise elimination of PRC2 activity in targeted loci.

(B) EbdCas9-mCherry and NCdCas9-mCherry construct under Tet-On operator.

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et al., 2021) (colocalization of EED and EBdCas9 increases 2.7× in the presence of +gRNA versus –gRNA [Figures S1H and S1I]).

We analyzed EBdCas9 or NCdCas9 in genes that were responsive to previous EB treatment (Moody et al., 2017). *TBX18* is a growth-promoting transcription factor required for embryonic development and the conversion of myocytes into sinoatrial cells (Kapoor et al., 2013; Wiese et al., 2009). *TBX18* was upregulated with reduced H3K27me3 marks after EB expression (Figures 1E and S2A) (Moody et al., 2017). In iPSCs, the *TBX18* promoter is bivalent—the upstream region is simultaneously decorated with H3K27me3 repressive marks and H3K4me3 active marks. We tiled the *TBX18* upstream promoter region, based on UCSC Genome Browser (GRCh36/hg36) Assembly, with guides specific to the region using CRISPRscan (Moreno-mateos et al., 2015). Guides were used with EBdCas9 and NCdCas9 to screen for loci sensitive for PRC2 (Figure 1E). EBdCas9 was induced at day –2 using Dox and transfected with *in-vitro*-synthesized gRNA on days 0 and 1, and the cells were collected on day 3 (Figure 1F).

To identify upstream regions that are more sensitive to EB action, we transfected the cells with a combination of guides. While guides 1.9–3.5 kb upstream from transcription start sites (TSSs) (guide 1 [g1], g2, g7, and g8) show no noticeable effects in *TBX18* transcription, guides 0–1.9 kb upstream of TSSs (g3, g4, g5, and g6) showed 5-fold *TBX18* upregulation compared with no guide treatment or with the NCdCas9 control (Figure 1G). Induction of EBdCas9 with individual *TBX18* gRNAs increased *TBX18* transcript levels by 10-fold (gRNAs 3 and 4) and 50–60 fold (gRNAs 5 and 6) compared with EBdCas9 with no guide or with g1,2,7,8 and NCdCas9 or KRABdCas9 (Figures 1H and S2B). *TBX18* protein was overexpressed with EBdCas9/g6 but not with NCdCas9/g6 (Figure 1I). gRNAs 3–6 (–0.5 to –1.5 kb) localized to chromatin domains where H3K4me3 marks are low and H3K27me3 marks and EZH2 are enriched (Figures 1E and S2C). To ensure tiled gRNAs have access to targeted *TBX18* DNA, we localized EBdCas9 after transfection with *TBX18* gRNAs (1–8) using Cas9 antibody (Ab) for chromatin immunoprecipitation (ChIP)-qPCR (Figure S2D). To test the specificity of EBdCas9, we monitored *POU5F1* (*OCT4*) gene expression and observed no significant changes in *OCT4* in cells treated with gRNAs targeting the *TBX18* promoter (Figure 1J); hence, *TBX18* upregulation is not a secondary effect of cellular differentiation. Assay for transposase-accessible chromatin sequencing (ATAC-seq) analysis of EBdCas9/g6- or NCdCas9/g6-transfected cells resulted in similar open chromatin regions. Since no significant changes in chromatin accessibility were detected, we conclude that transcript upregulation is dependent on local changes in the PRC2 complex (Figure S2A) and that the process

is PRC2-dependent since EBdCas9, but not NCdCas9, activates *TBX18* gene expression at the precise loci.

### EBdCas9/gRNA remodels epigenomic marks and memory in *TBX18* locus

To dissect the mechanism of EBdCas9 action at precise genomic loci, we used ChIP-qPCR (PIXUL-Matrix-ChIP [Bomsztyk et al., 2019]). As noted above, EBdCas9 or NCdCas9 expression was induced in iPSCs (WTC) using Dox followed by *TBX18* gRNA6 (g6) transfections on days 0 and 1 and analysis on day 3 by qRT-PCR and ChIP-qPCR (Figure 2A). As expected, while both constructs showed similar dCas9 expression, only EBdCas9/g6 showed a significant increase of *TBX18* transcription (Figure 2B). Using the ChIP-qPCR method, we quantified the abundance of EB/NCdCas9 (Cas9), histone 3 with K27me3 marks (H3K27me3), and EZH2 proteins at the *TBX18* g6 region by generating and quantifying the 150 bp amplicon flanking the g6 chromatin region. ChIP-qPCR using a Cas9 Ab confirmed that both EBdCas9 and NCdCas9 are recruited to the g6 locus (Figure 2C). However, only EBdCas9 results in a reduction of EZH2 (Figure 2C), most likely by directly blocking EZH2 binding to EED (Moody et al., 2017). This resulted in a reduction of H3K27me3 marks in EBdCas9/g6 at the g6 locus (Figure 2C). In contrast, while the ChIP-qPCR assay of EBdCas9/g7 confirmed recruitment of Cas9 and reductions of EZH2 and H3K27me3 marks at the g7 locus, *TBX18* transcription was not affected (Figures 1H, 2D–2E, and S3A). Thus, while EBdCas9 is able to disrupt EED-EZH2 interactions at many loci, transcription is only affected at certain locations.

Recent data have shown that PRC2-dependent marks are faithfully inherited during cell division (Escobar et al., 2019). To learn whether EBdCas9-dependent reduction of EZH2 occupancy and H3K27me3 marks are inherited from the cell from which they descended (epigenomic memory), we repeated the assay as before but allowed the cells to grow for an additional 2 days (3 cell cycles; Figure S3B) after eliminating EBdCas9/gRNA expression (Figure 2F). qRT-PCR shows that the EBdCas9 transcript is upregulated at day 3 post-transfection but is not detected 2 days after elimination of induction (day 5) (Figure 2G). In contrast, *TBX18* transcription increased 80-fold at day 3 and 50-fold at day 5 (Figure 2G). To test if *TBX18* transcription 2 days after EBdCas9 elimination correlated with epigenomic memory, we performed PIXUL-ChIP for the EZH2 protein and H3K27me3 marks at the g6 region at 3 and 5 days. ChIP-qPCR showed a depletion of both H3K27me3 and EZH2 at day 3, as expected. Importantly, this reduction of EZH2 and H3K27me3 persisted until day 5 (Figure 2H), showing an epigenomic inheritance of PRC2 elimination in the g6 region. These

(C) Generation of stable EBdCas9 or NCdCas9 transgenic iPSC (WTC) lines after 3 days of induction of (2 µg/ml) doxycycline (Dox).

(D) Immunoblot analysis of EBdCas9 and NCdCas9 whole-cell lysate after 3 days of Dox induction.

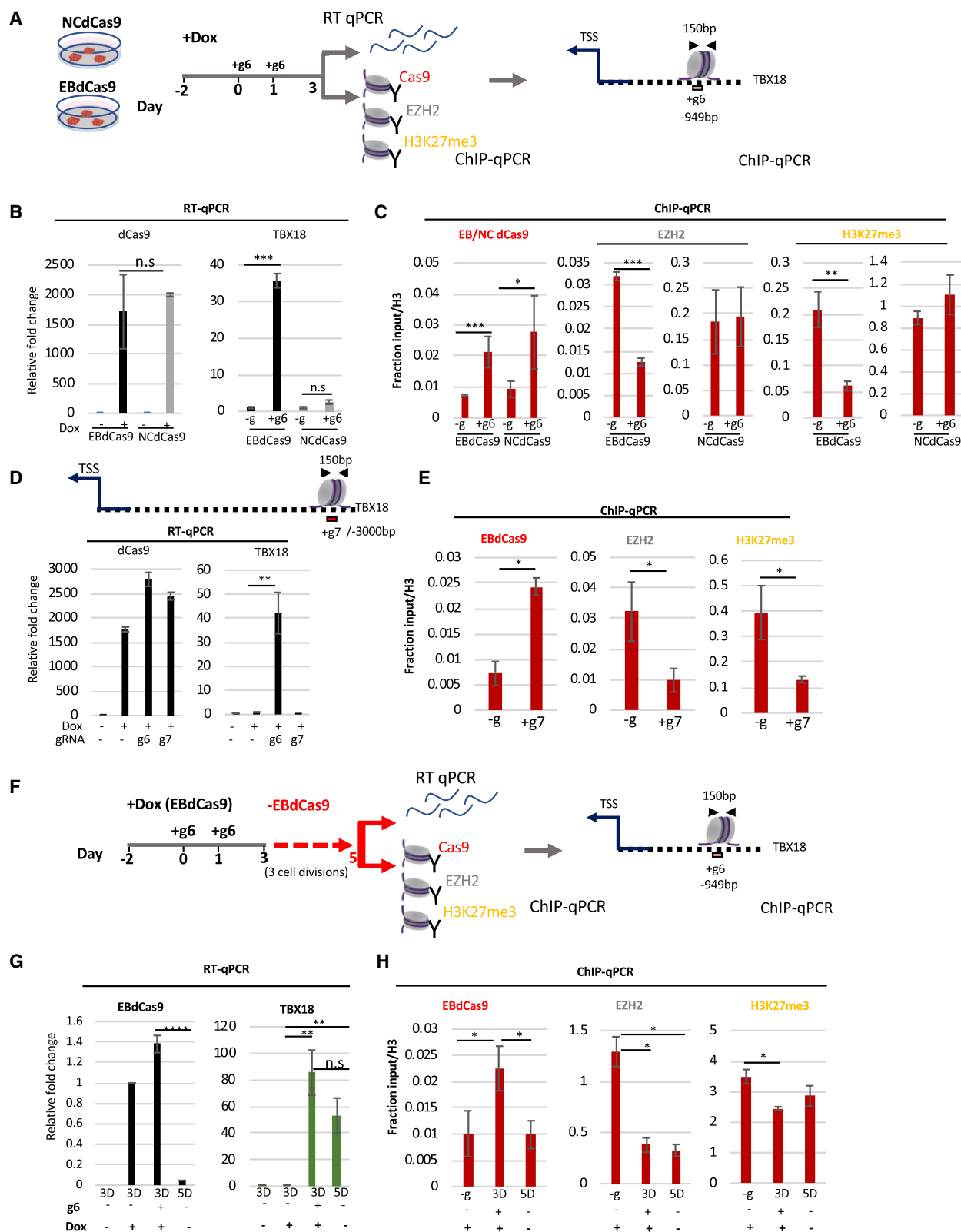
(E) Integrative genomic viewer of *TBX18* H3K27me3 and H3K4me3 promoter tiling.

(F) Timeline of EBdCas9 or NCdCas9 induction and gRNA transfection.

(G and H) qRT-PCR analysis of *TBX18* expression of EBdCas9 and NCdCas9 normalized to β-actin and calculated as the relative fold change compared with no guide (induced with Dox) of each respected cell line (G) after *TBX18* gRNAs cocktail transfection; g1, g2, g7, and g8 or g3, g4, g5, and g6 (H) after individual *TBX18* gRNA (1–8) transfection (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).

(I) Immunofluorescent imaging of EBdCas9 WTC and NCdCas9 WTC for either no guide or after transfection with *TBX18* gRNA 6 (*TBX18* g6). Blue, DAPI; green, *OCT4*; purple, *TBX18*; scale bar, 100 µm.

(J) qRT-PCR analysis of *OCT4* after individual transfection of *TBX18* gRNA g5 and g6.



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data reveal that EBdCas9 not only remodels the epigenome but also leads to inheritance of the changes.

### Spreading of EBdCas9 induced epigenomic changes

To learn whether (1) EBdCas9/*TBX18*g6 epigenomic changes are limited to the g6 chromatin locus and (2) the spread chromatin changes are inherited upon cell division, we tiled the *TBX18* genomic area with primer sets between the TSS and g6 and upstream of g6. As expected, at 3, but not at 5, days, EBdCas9 (Cas9) is localized to the g6 region (Figure 3A). At neither time point was EBdCas9 observed in any other loci upstream of the *TBX18* TSS. However, the EZH2 protein was significantly downregulated not only at g6 but also in loci between the TSS and g6 both at day 3 and 5 time points (Figure 3B). Since EBdCas9 was specifically localized to the g6 region at day 3, but not at day 5, EZH2 reduction in these regions suggest that PRC2 disruption by EBdCas9 at g6 at day 3 (1) spreads toward the TSS and (2) is inherited during cell division. Accordingly, disruption of the PRC2 complex also resulted in an ongoing reduction of H3K27me3 marks between the TSS and g6 (Figure 3C). The PRC2 component, JARID2, showed reduced binding to chromatin between the TSS and g6 at the day 3 time point (Figure 3D). Since JARID2 is critical for PRC2 homing and the stimulation of EZH2 catalytic activity, its reduction in this region is a further indication of reduced PRC2 activity (Kasinath et al., 2018). EED, on the other hand, binds to EBdCas9 and remains at g6 at day 3; however, it is depleted between the TSS and g6 (Figure 3D). Thus, PRC2 disruption and consequent reduction of H3K27me3 histone marks spread from g6 toward the TSS, and these changes persist after multiple cell divisions.

To test if the depletion of PRC2 results in epigenomic modifications indicative of transcription activation, we performed ChIP-qPCR using antibodies against H3K27ac, p300, RNA Pol II C-terminal domain (CTD), and RNA Pol II CTD Ser5P (indicative of RNA Pol II pause [Boehm et al., 2003]). We observed recruitment of RNA Pol II, histone acetyltransferase (p300), and acetylation marks (H3K27ac) at the *TBX18* g6 locus (Figures 3E and S3C), indicative of active transcription. RNA Pol II was not only highly enriched in the *TBX18* g6 locus at the day 3 time point but also at the day 5 time point, 2 days after the EBdCas9 was

eliminated (Figure 3F). Since RNA Pol II is recruited to the g6 region, we tested if *TBX18* nascent RNA is transcribed in that region. We observed increased nascent RNA amplification in EBdCas9 cells transfected with *TBX18*g6 compared with no guide using qRT-PCR (Figures 3G and 3H; upregulation on exon 1 but not on the 2.0 kb upstream region). There are two transcription start sites for *TBX18* (UCSC Genome Browser GRCh37/hg19); EBdCas9/g6 activates transcription in the TSS#2 region (532 bp downstream of g6; Figures 3G and 3H), which is the dominant start site based on cap analysis gene expression (CAGE)-seq data (Figures S3D and S3E).

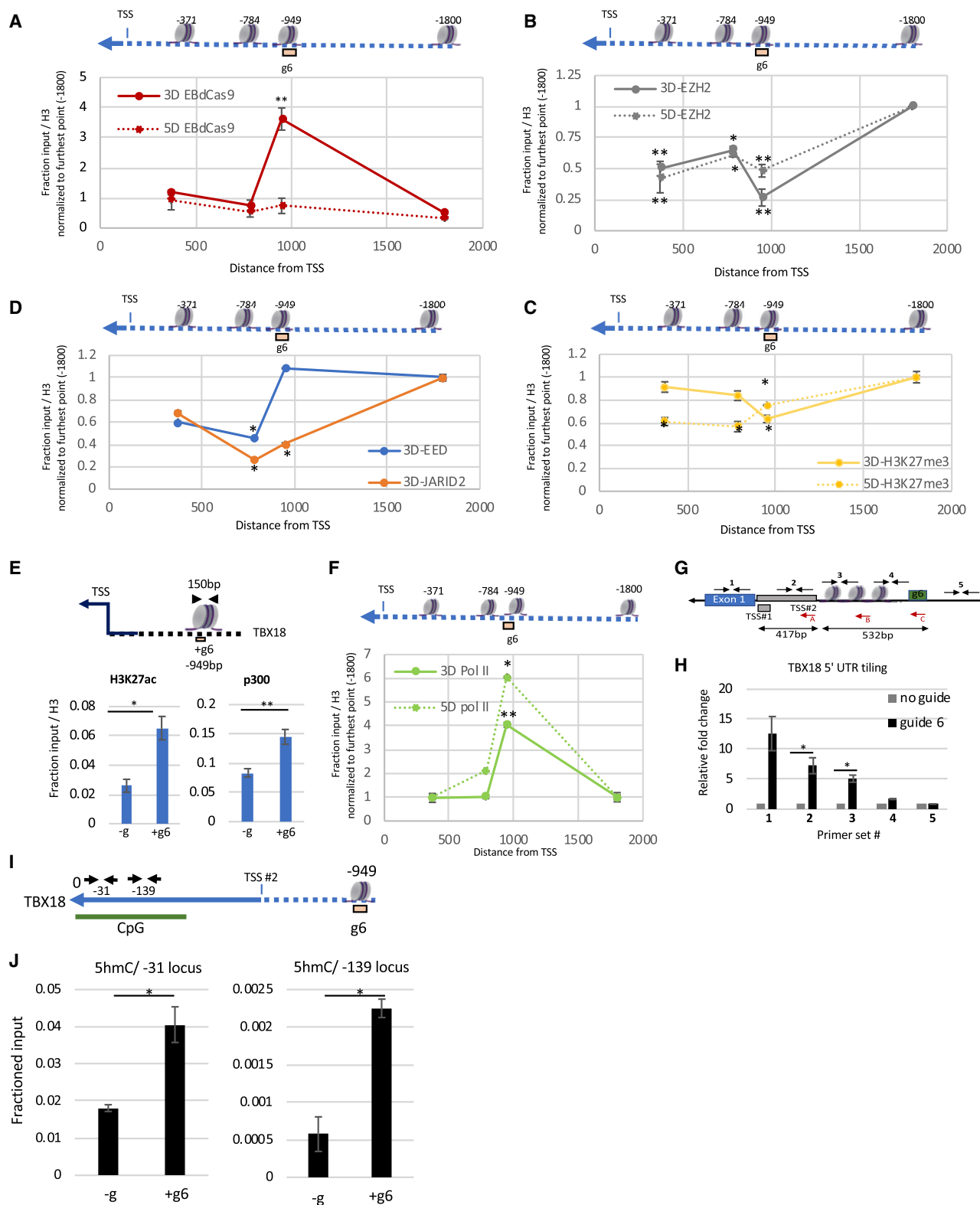
To test if DNA methyl marks are affected by EBdCas9/*TBX18*g6, we performed Matrix-MeDIP (Yu et al., 2011) using 5 methylcytosine (5mC) and 5 hydroxymethylcytosine (5hmC) antibodies (Figure 3I). While EBdCas9/*TBX18*g6 did not alter methylation (5mC) compared with no guide, it increased hydroxymethylation (5hmC; hydroxymethylations is actively involved in DNA de-methylation and transcription activation [Santiago et al., 2014]) (Figure 3J). Tiling regions outside of the CpG island on the *TBX18* promoter did not show significant 5hmC changes (Figures S3F–S3H).

### EBdCas9 reveals distal TATA box function

Since targeting the *TBX18*g6 site led to transcription activation, epigenetic remodeling, and epigenomic memory, we analyzed the chromatin region for transcription factor binding sites, core promoter sequence elements, and G-quadruplex patterns (Figures S4A–S4C). Quadruplex forming G-rich sequences (QGRSs) (Kikin et al., 2006) are structure motifs linked to increased transcriptional activity (Hansel-hertsch et al., 2016). In the 1 kb upstream from the TSS, we identified a combination of TATA box and mammalian initiator factor binding sites (INRs) near the *TBX18* g6 locus (Figures 4A and S4C; Table S2; TATA box 21 bp and INR 48–57 bp from g6). We tested if this distal TATA box is required for *TBX18* transcription by precisely deleting 40 bp surrounding it using CRISPR-Cas9-homology-directed repair (HDR) (Figure 4B). Two independent, pluripotent EBdCas9 cell lines with homozygous TATA box deletions were generated, TATA Δ #1 and TATA Δ #2 (see STAR Methods for details) (Figure 4B) and analyzed for *TBX18* transcription. In contrast to the

**Figure 2. EBdCas9 precisely eliminates EZH2 from the PRC2 complex, remodels epigenomic marks, and actuates epigenomic memory on *TBX18* locus**

- (A) EBdCas9 and NCdCas9 timeline for Dox induction, gRNA transfection, and analysis: qRT-PCR or ChIP-qPCR using the antibodies Cas9, EZH2, and H3K27me3 and analyzing *TBX18* g6 DNA region ~1.0 kb upstream of the TSS (150 bp).
- (B) qRT-PCR of dCas9 or *TBX18* relative fold change after 3 days of Dox induction (dCas9) or 3 days of *TBX18* g6 RNA transfection (*TBX18*) normalized to  $\beta$ -actin and compared with no guide (Dox induced for *TBX18*) of each respective cell line.
- (C) ChIP-qPCR of induced (+Dox) EBdCas9 and NCdCas9 after 3 days of transfection with *TBX18* g6 RNA (+g6) or no transfection (–g). Normalized as fraction of input/H3. Antibodies for ChIP are listed above the graphs, and the genomic region analyzed by qPCR includes *TBX18* g6 locus (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).
- (D) qRT-PCR of dCas9 and *TBX18* relative fold change after 3 days of Dox induction and *TBX18* g6 RNA (+g6) or g7 RNA (+g7) transfection normalized to  $\beta$ -Actin and compared with no guide (Dox induced) (–g) exactly as described in (A).
- (E) ChIP-qPCR of induced (+Dox) EBdCas9 after 3 days of transfection with *TBX18* g7 RNA (+g7) or no transfection (–g). Normalized as fraction of input/H3 and analyzed exactly as in (C).
- (F) EBdCas9 timeline for measuring memory. EBdCas9 induction and *TBX18* g6 RNA transfection exactly as in (A). On day 3, EBdCas9 media was replaced with no Dox (–EBdCas9) for 2 days and analyzed on day 5 for qRT-PCR and ChIP-qPCR.
- (G) qRT-PCR analysis of EBdCas9 and *TBX18* for 3 or 5 days while inducing with Dox (+) or not (–) and in the presence of *TBX18* g6 RNA (g6) (+) or not (–).
- (H) ChIP-qPCR of no guide (–g), 3- or 5-day EBdCas9 either induced with Dox (+) or not (–), or transfected with *TBX18* g6 RNA (+) or not (–). Normalized to fractioned Input/H3 and analyzed exactly as in (C).



**Figure 3. EBdCas9 causes epigenomic spreading and recruits activation marks**

(A–D) Tiling of EBdCas9/g6 ChIP (A) Cas9, (B) EZH2, (C) H3K27me3, (D) EED, and JARID 2 on *TBX18* genomic loci (bp) relative to the TSS (listed above nucleosome) using qPCR; solid red lines are day 3 and dashed red lines are day 5. Each point is the relative fold change *TBX18* g6RNA (+g6) versus no guide (–g), (legend continued on next page)



significant increase of *TBX18* transcription in wild-type Ebd-Cas9/g6 (WT), TATA  $\Delta$  #1 and TATA  $\Delta$  #2 EbdCas9 lines failed to upregulate *TBX18* transcription (Figure 4C). Similar levels of EbdCas9 mRNA expression after Dox induction were observed in all cell lines (Figure 4C). To test if other transcriptional elements may be responsible for *TBX18* expression, we transfected WT or TATA  $\Delta$  #1 with *TBX18*g5 (900 bp upstream of g6). Only WT was able to upregulate *TBX18* (Figures S4D and S4E). TATA  $\Delta$  #1 also reduced activation by the EZH2 chemical inhibitor EPZ-6438 (Knutson et al., 2013) (Figures S4F and S4G). We conclude that the distal, upstream, PRC2-regulated TATA box is essential for *TBX18* gene expression.

To explore why deletion of the distal TATA box affects *TBX18* gene expression, we examined recruitment of TATA binding protein (TBP), and RNA Pol II (Wasylyk et al., 1980), at the g6 region after EbdCas9/g6 treatment (Figure 4D). We found that TBP was recruited to the g6 region, showing that the TATA box 21 bp away from the g6 region and 900 bp away from TSS1 can bind TBP after local PRC2 inhibition (Figures 4D and 4E). EbdCas9 (Cas9) and RNA Pol II were also recruited to the g6 site, and the EZH2 protein was reduced, as observed earlier (Figures 2C, 3F, and S3C). However, when the TATA box region was deleted (TATA  $\Delta$  #1 and TATA  $\Delta$  #2 EbdCas9), TBP and RNA Pol II were not recruited (Figure 4E). Deletion of the TATA box did not alter EbdCas9 recruitment to, and EZH2 depletion at, the g6 region (Figure 4E). Thus, the TATA box >500 bp away from *TBX18* TSS can recruit TBP when PRC2 is inhibited at the locus (model in Figure 4F).

### EbdCas9 activates *CDKN2A* (*p16*) by epigenomic remodeling in iPSCs and DMG tumor cells

We studied EbdCas9/g action on PRC2 function at a second gene locus, *CDKN2A* (*p16*). *p16* is a critical regulator of cell division and a tumor suppressor that inhibits cyclin-D-dependent protein kinase activity and the G1-S transition (Piunti et al., 2017; Mohammad et al., 2017; Cordero et al., 2017; Chen et al., 2020). In rapidly dividing cells, such as in DMG, *p16* is repressed due to hypermethylation at the promoter area (Cordero et al., 2017). We searched for potential nucleation sites in the *p16* upstream region responsive to PRC2 reduction by Ebd-Cas9. We tiled the promoter area and gene body of *p16* with eight gRNAs ranging from 0.3 to 2.3 kb upstream of the TSS

and 0.2–0.7 kb downstream of the TSS (Figure 5A) and assessed DNA accessibility using the human ESC (hESC) E1 iCas9 cell line (Ferrecchio et al., 2018) (Figure S5A). WTC EbdCas9 or NCdCas9 were induced prior to transient transfection of the gRNAs followed by cell harvest at day 3 and *p16* transcript analysis (Figure 5B). EbdCas9 activated *p16* transcript expression on 6 out of the 8 gRNAs tested, but none activated transcription in the control NCdCas9-expressing line (Figure 5C). As observed with *TBX18* tiling, gRNAs that are within 0.5–1.5 kb of the TSS showed the highest *p16* transcript activation. g1, g2, g3, g4, g6, and g7 induced transcription of *p16* by 20- to 80-fold compared with no guide (–g) or NCdCas9 (Figures 5A–5C); g5 and g8 did not increase transcription significantly, even though EbdCas9/g had access to the DNA in these regions. TATA box and mammalian initiator factor binding sites in the *p16* –424 and –750 bp promoter regions, upstream from the TSS (Figure 5D), are in close proximity to g1. *p16* was also upregulated at the protein level by using immunofluorescence analysis (Figure 5E).

Since activation of *p16* expression results in cell-cycle arrest in gliomas (Cordero et al., 2017), we tested the effect of *p16* upregulation in iPSCs. Transfection of WTC EbdCas9 with g1(*p16*) resulted in a 2-fold cell number and colony size reduction compared with no-guide controls (–g) on day 3 (Figures S5B and S5C). Consistent with this, EbdCas9/g1(*p16*) reduced cell viability compared with no guide (–g) or NCdCas9 using a resazurin fluorescent (Alamar Blue) detection reagent (Figure 5F). Transfection of g1 resulted in 40-fold transcriptional upregulation of the *p16* gene with EbdCas9 but not the NC (Figures 5G and 5H). Similarly, EbdCas9/g1, but not NCdCas9/g1, reduced EZH2 protein and H3K27me3 marks at the g1 site (the two proteins were recruited equally to the locus) (Figure 5I) and increased H3K4me3 marks (Figures 5J [Skene and Henikoff, 2017] and S5D). H3K27ac marks also increased on EbdCas9/g1 compared with –g samples (Figure S5E).

DMG is an aggressive form of brain cancer that develops in glial cells in the pons and primarily affects children (Mohammad et al., 2017; Stafford et al., 2018). A lysine-to-methionine mutation in the histone H3.3 gene (H3.3K27M) defines DMG tumors (Mohammad et al., 2017; Stafford et al., 2018). Genome-wide studies of H3K27me3 marks in DMG cells have revealed that while a global loss of H3K27me3 is observed due to a

normalized to the respective fraction of input/H3 and compared with the relative fold change using –1,800 bp primer set as control (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).

(E) ChIP-qPCR of 3-days-induced EbdCas9 and *TBX18* g6 RNA (+g6) transfected or not (–g) using H3K27ac and p300 and normalized as a fraction of input/H3. Genomic region analyzed by qPCR is the *TBX18* g6 locus and is denoted above (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).

(F) Tiling of EbdCas9/+g6 ChIP (RNA Pol II CTD) on *TBX18* genomic loci (bp) relative to the TSS (listed above nucleosome) using qPCR exactly as in (A); solid green lines are day 3 and dashed green lines are day 5 (as described in A–D). Analyzed exactly as in (A).

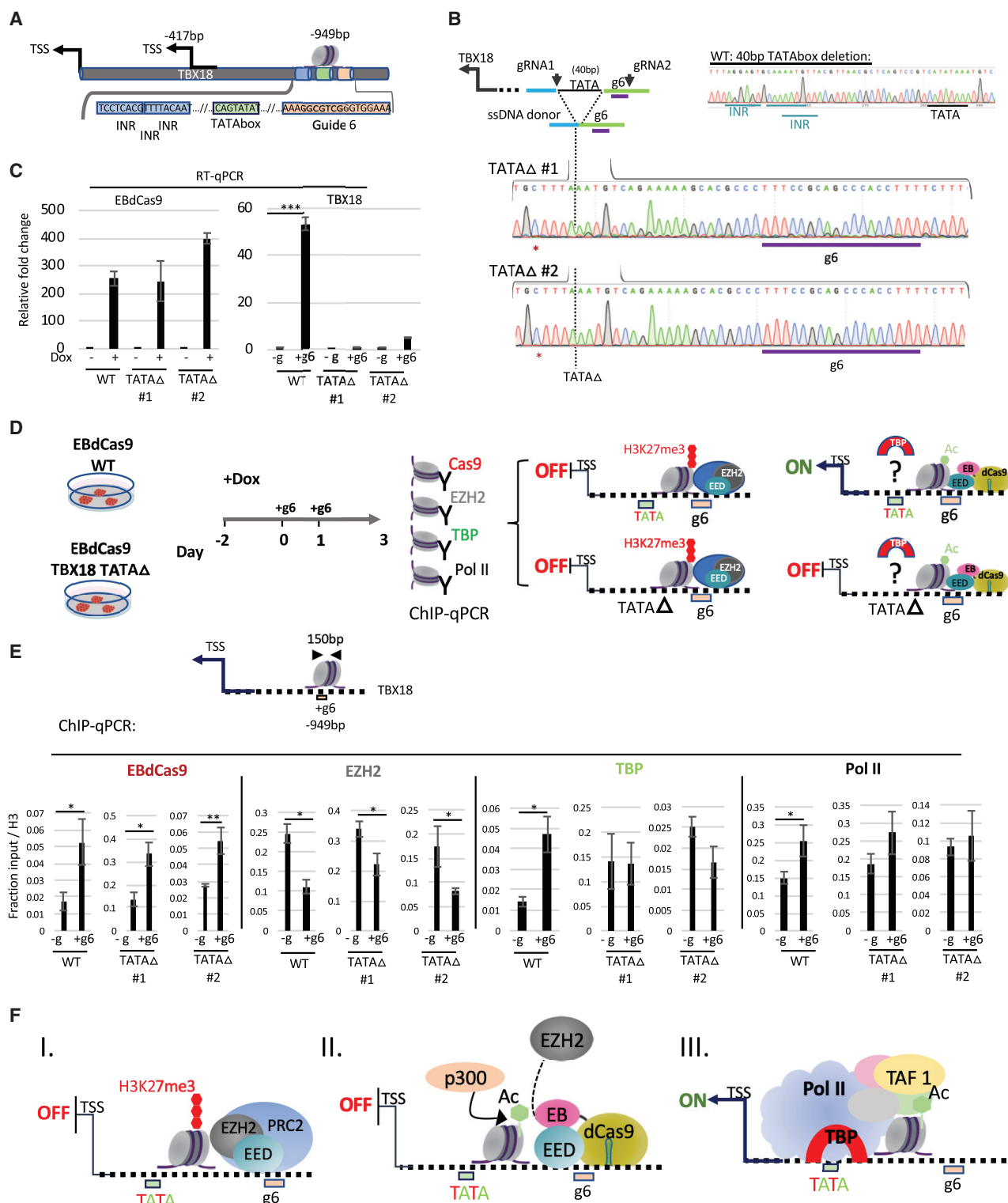
(G) Illustration of *TBX18* TSS peak position relative to the g6 region in bp.

(A–C) Red arrows denote the reversed primer used for cDNA reverse transcription. Internal black arrows with Arabic numbers denote the PCR amplicon generated following cDNA production: (C) corresponds to primer set #4; (B) corresponds to #3; and (A) corresponds to primer sets #2, #1, and #5.

(H) RNA expression of *TBX18* 5'UTR region using qRT-PCR (reverse primer in red) after 3 days of transfection with *TBX18* g6 RNA (g6) or no transfection (no guide). Regions amplified correspond to #1–#5 and are indicated with black arrows. Normalized to  $\beta$ -actin, and the relative fold change is compared with no guide (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).

(I) Illustration of *TBX18* CpG locus using UCSC Genome Browser (GRCh36/hg36) Assembly. “0” denotes first Met relative to the g6 region in bp. Small black arrows annotate the loci for qPCR amplification in (J).

(J) MedIP-qPCR of induced (+Dox) EbdCas9 after 3 days of transfection with *TBX18* g6 RNA (+g6) or no transfection (–g) using 5 hydroxymethylcytosine (5hmC) Abs. Normalized to fractionated input (n = 2, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).



**Figure 4. EbdCas9 reveals functional distal TATA box**

(A) TATA box and initiator (INR) representation with respect to g6 locus using the Element Navigation Tool for detection of core promoter elements. (B) Generation of TATA box deletion (TATA  $\Delta$ ) clones TATA  $\Delta$  #1 and TATA  $\Delta$  #2 using CRISPR-Cas9. gRNA1 and -2 excise an 88 bp fragment, including the g6 region. The g6 region was reconstructed using a 168 bp single-stranded DNA (ssDNA) donor. Depicted here is a deletion of 40 bp including the TATA box and 3

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H3K27me3 mutation, surprisingly, a sharp increase in H3K27me3 repressive marks is observed at certain genes (presumably due to other H3 variants than H3.3 being methylated) (Stafford et al., 2018), including the tumor suppressor *CDKN2A/p16/INK4a* (Mohammad et al., 2017). Previous work has shown that overexpression of p16 reduces DMG cell growth and viability (Cordero et al., 2017). We were able to transiently transfect both EBdCas9 and NCdCas9 into DMG cells with the p16-targeting guide g1 (Figure S6A), but only the former reduced cell viability and cell replication (by 50%; Figures S6B and S6C) and upregulated *p16* gene expression (Figure S6D). As in iPSCs, there was a reduction of EZH2 and H3K27me3, and an upregulation of H3K4me3, for EBdCas9 but not for NCdCas9 (Figure S6E). Thus, EBdCas9 can function in different cell types, and targeting EBdCas9 to specific loci produces similar epigenomic and transcriptional effects in different cell types.

### H3K27me3 remodeling induces trophoblast fate

The placental (trophectoderm [TE]) versus embryonic (inner cell mass [ICM]) cellular fate decision (the first in animal development) is dependent on PRC2 activity (Saha et al., 2013). While overexpression of H3K27me3 is associated with ICM lineage, genome-wide depletion of H3K27me3 marks is associated with TE lineage (Saha et al., 2013; Banaszynski et al., 2013; Liu et al., 2016; Yang et al., 2017a, 2017b; Yan et al., 2013) (Figure 6A). We first tested if ubiquitously expressed EB (Moody et al., 2017) can induce trophoblast fate (Figure 6A). Recently, two groups have generated culture conditions that enabled the establishment of extended (EPS) cells from either cleavage state of mouse embryos or hESCs (Yang et al., 2017a, 2017b). The EPS cell stage has a developmental potency to generate both embryonic (ICM) and extraembryonic placental tissue (TE cell lineage) (Yang et al., 2017a, 2017b). Moreover, EPS epigenetic analysis validated the enrichment of bivalent, H3K27me3, and H3K4me3 marks in developmental genes (Yang et al., 2017a, 2017b). We reprogrammed WTC EB-FLAG and WTC NC-FLAG (Moody et al., 2017) iPSC lines to the EPS stage using a hLIF, chir99021, (S)-(+)-dimethindene maleate (DiM), and minocycline hydrochloride (MiH) (LCDM) reprogramming cocktail (Yang et al., 2017b). Once established, we validated the EPS cell stage by colony dome-shaped morphology, single-cell colony forming efficiency, expression of pluripotency markers, and TE lineage differentiation capacity in trophoblast differentiation (TX) media containing tumor growth factor beta 1 (TGF- $\beta$ 1), fibroblast growth factor 4 (FGF4), and heparin (Kubaczka et al., 2014) (Figures 6B and S7A). EB-FLAG- but not NC-FLAG-expressing cells

lost EPS colony morphology and upregulated GATA3 and TBX3 expression at day 4, suggesting accelerated differentiation (Figures 6B–6D). We further showed that while confocal imaging confirmed the tight, dome-shaped morphology, the expression of nuclear stem cell transcription factor OCT4, and the absence of GATA3 expression for both EPS (EB-FLAG) and EPS(NC-FLAG) at day 0 (Figure 6E), TE differentiation (4 days) with EB-FLAG, but not NC-FLAG, resulted in a dramatic reduction of the OCT4 expression, loss of colony morphology, and upregulation of GATA3 (Figure 6E). Furthermore, immunoblot analysis revealed that 4 days of TE differentiation with EB-FLAG reduced H3K27me3 and EZH2, while GATA3 was upregulated (Figure S7).

To investigate these fate changes in more detail, we analyzed gene expression utilizing RNA sequencing (RNA-seq). We projected gene expression data for TE-differentiated EPS(EB-FLAG) cells (4 days, 6 days, and differentiated extravillous cytotrophoblast [EVT] [Okoe et al., 2018]) onto principal-component analysis (PCA) plots of single-cell transcriptomes of early cynomolgus monkeys (Nakamura et al., 2017) and found acceleration of TE differentiation in EB-FLAG-expressing cells (Figures 6F; Table S3); the projection is based on 773 highly variable genes (standard deviation >2) in the monkey single-cell dataset). TE-differentiated EPS EB-FLAG cells that were induced with Dox during differentiation migrate from post-implantation early or late epiblast (PostE-EPI or PostL-EPI), respectively, to post-implantation partial TE (Post-paTE) and pre-implantation late TE (PreL-TE) compared with no-Dox EB-FLAG-differentiated cells. The EB-FLAG samples are distant from ICM on the TE differentiation lineage, consistent with EB-FLAG accelerating TE differentiation. EVT EB-FLAG cells were first induced to express EB for 4 days and thereafter passaged 3 times (in trophoblast stem cell [TSC]-conditioned media) (Okoe et al., 2018) without Dox (and therefore without further EB expression) and were found to be closest to PreL-TE. Thus, elimination of H3K27me3 marks by induction of the EB-FLAG protein dramatically accelerates TE lineage differentiation.

### Targeted PRC2 inhibition at *CDX2* and *GATA3* loci induces trophoblast trans-differentiation

During mouse blastocyst formation, the relative levels of EED and KDM6B determine PRC2 complex recruitment and incorporation of H3K27me3 marks at the chromatin domains of target genes (Figure 6A). The TE-lineage-specific transcription factors *CDX2* and *GATA3* show PRC2-dependent repression in the ICM (Saha et al., 2013; Xiao et al., 2020). However, it is not known if PRC2 activity in the promoter regions of these two genes is

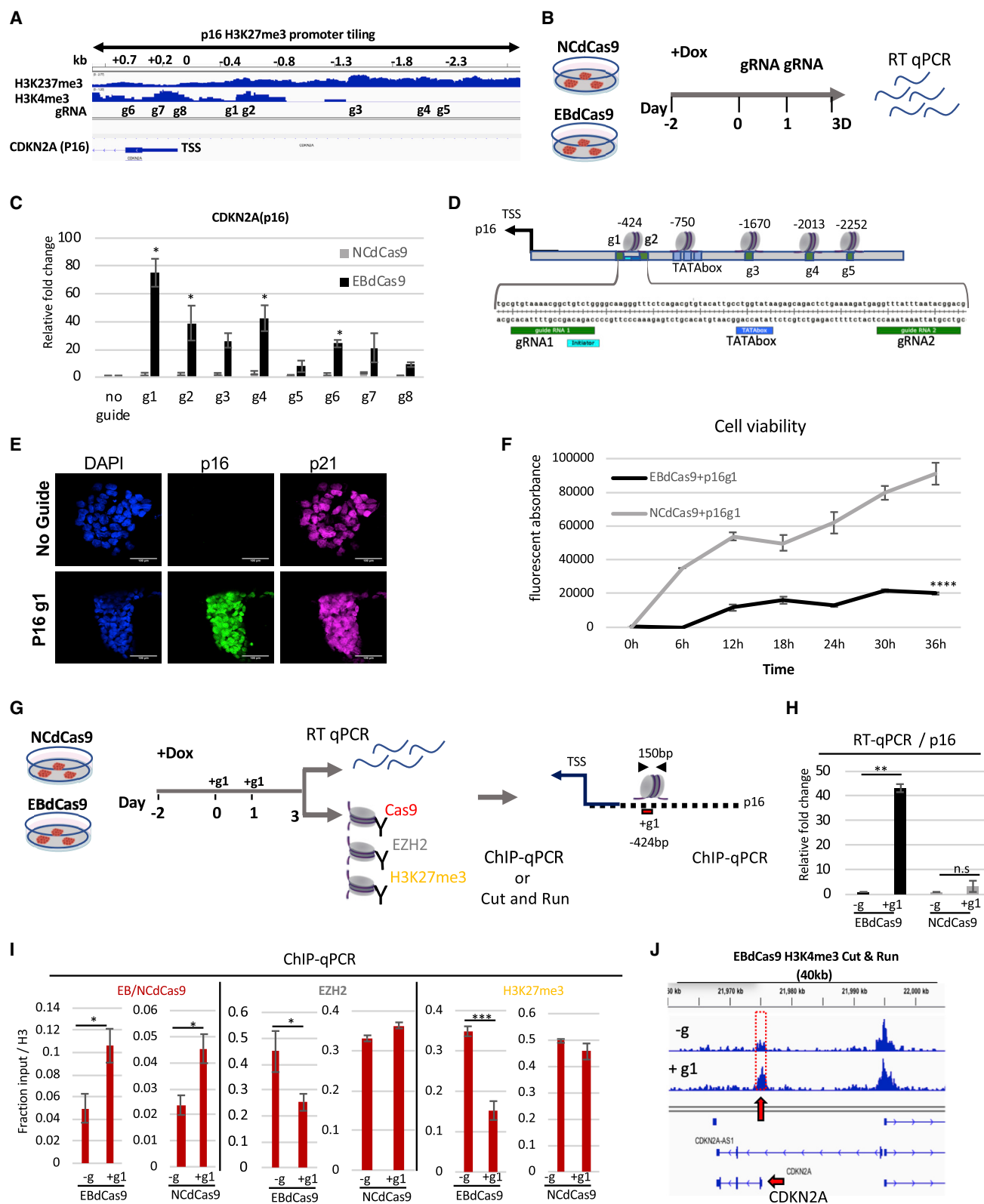
initiator elements (INR). Genomic DNA of TATA  $\Delta$  #1 or TATA  $\Delta$  #2 displaying homozygous clones of 40 bp TATA box deletion using Sanger sequencing. PAM-site-blocking mutation is denoted with a red asterisk.

(C) qRT-PCR of EBdCas9 and *TBX18* relative fold change after 3 days of Dox induction and *TBX18* g6 RNA transfection normalized to  $\beta$ -actin and compared with no guide for each respective cell line (WT: EBdCas9, TATA  $\Delta$  #1, TATA  $\Delta$  #2).

(D) EBdCas9 WT and EBdCas9 TATA  $\Delta$  #1 and TATA  $\Delta$  #2 timeline for Dox induction, gRNA transfection, and ChIP-qPCR analysis using the antibodies Cas9, EZH2, TATA binding protein (TBP), and RNA Pol II and analyzing for *TBX18* g6 DNA region. Right panel depicts 4 possible model scenarios for TATA box and TATA  $\Delta$  with and without EBdCas9.

(E) ChIP-qPCR of induced (+Dox) EBdCas9 (WT) and EBdCas9 TATA  $\Delta$  #1 and TATA  $\Delta$  #2 3 days following transfection with *TBX18* g6 RNA (+g6) or no transfection (–g). Normalized to fraction input/H3. Antibodies that were used for ChIP are listed above the graphs, and the genomic region analyzed by qPCR includes the *TBX18* g6 locus (n = 3, SEM; \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001, two-tailed t test).

(F) Model describing EBdCas9/TBX18 mode of operation: (I) The *TBX18* distal TATA box function is repressed by PRC2. (II) EBdCas9 de-represses the *TBX18* TATA box region. (III) TBP, TBP associated factors (TAFs), and RNA Pol II are recruited to the *TBX18* TATA box region to allow *TBX18* transcription.



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sufficient to induce the bifurcation between TE and ICM lineages. We tested if ICM-like cells, such as iPSCs, which have already passed the bifurcation point, are able to *trans*-differentiate to TE if *CDX2* and *GATA3* H3K27me3 epigenetic marks are reduced (Figure 6G). To do so, we targeted EBdCas9/gRNA to the gene-upstream regions of TE transcription factors *CDX2* and *GATA3*. After WTC(EBdCas9) induction and gRNA transfection (+Dox, 2 days, TeSR), the media was changed to TX-base media (+Dox) without factors (no TGF- $\beta$ 1, FGF4, and heparin) (Figure 6H). This created a permissive environment for TE differentiation, in which EBdCas9/gRNAs are the sole drivers for potential *trans*-differentiation. *CDX2* and *GATA3* were tiled across the promoter and gene body region with 5 different guides (Figure 6I). Since these two transcription factors are critical players in TE differentiation in mouse (Saha et al., 2013), we screened for the PRC2-responsive regions by co-transfecting guides covering upstream regions of *CDX2* and *GATA3* genes (g1/g1, g2/g2, ..., g5/g5). WTC EBdCas9, but not NCdCas9, with gRNA pairs g1 and g5 resulted in a significant gene activation (20- to 80-fold increase) for *CDX2* and *GATA3*, as well as the TE marker *TBX3*, and a reduction of *EZH2* and H3K27me3 at targeted loci (Figures 6J and S7C–S7F; the other gRNA pairs had little effect).

We analyzed the global transcriptomes of EBdCas9 g1/g1 or g5/g5 *CDX2* and *GATA3* samples by RNA-seq. PCA showed that the g1/g1 and the g5/g5 samples were more similar to TE samples than the no-guide controls (migration from PostE-EPI/PostL-EPI toward Post-paTE and PreL-TE state) (Figure 6K; Table S4). Furthermore, PCA projection of developmental genes indicated similarity to TE fate (Figure S7G; Krendl et al., 2017) in g1/g1 and g5/g5-treated WTC EBdCas9 cells but not in WTC EBdCas9 with no gRNA. After 3 additional days of differentiation using TGF- $\beta$ 1 and Neuregulin (Okoe et al., 2018), the guide-treated cells produced chorionic gonadotropin beta (CGB) consistent with the EVT fate, while in the presence of Forskolin, they produced mesenchyme-like cells with multi-nucleation morphology, consistent with the syncytiotrophoblast (ST) fate (Figures 6L–6N and S7H). Thus, targeting EBdCas9 to precise *CDX2* and *GATA3* loci results in *trans*-differentiation of iPSCs toward TE fate. To explore the effects on normal differentiation, we reprogrammed WTC EBdCas9 to the EPS state and

transfected with the g1/g1 and g5/g5 guides, which resulted in an increase in *GATA3* gene expression of 100- to 500-fold (Figures S7I–S7K; *CDX2* was less responsive) and loss of the pluripotency marker Oct4. Thus, inhibition of PRC2 activity at specific sites in only two genes can induce the bifurcation between TE and ICM fates.

## DISCUSSION

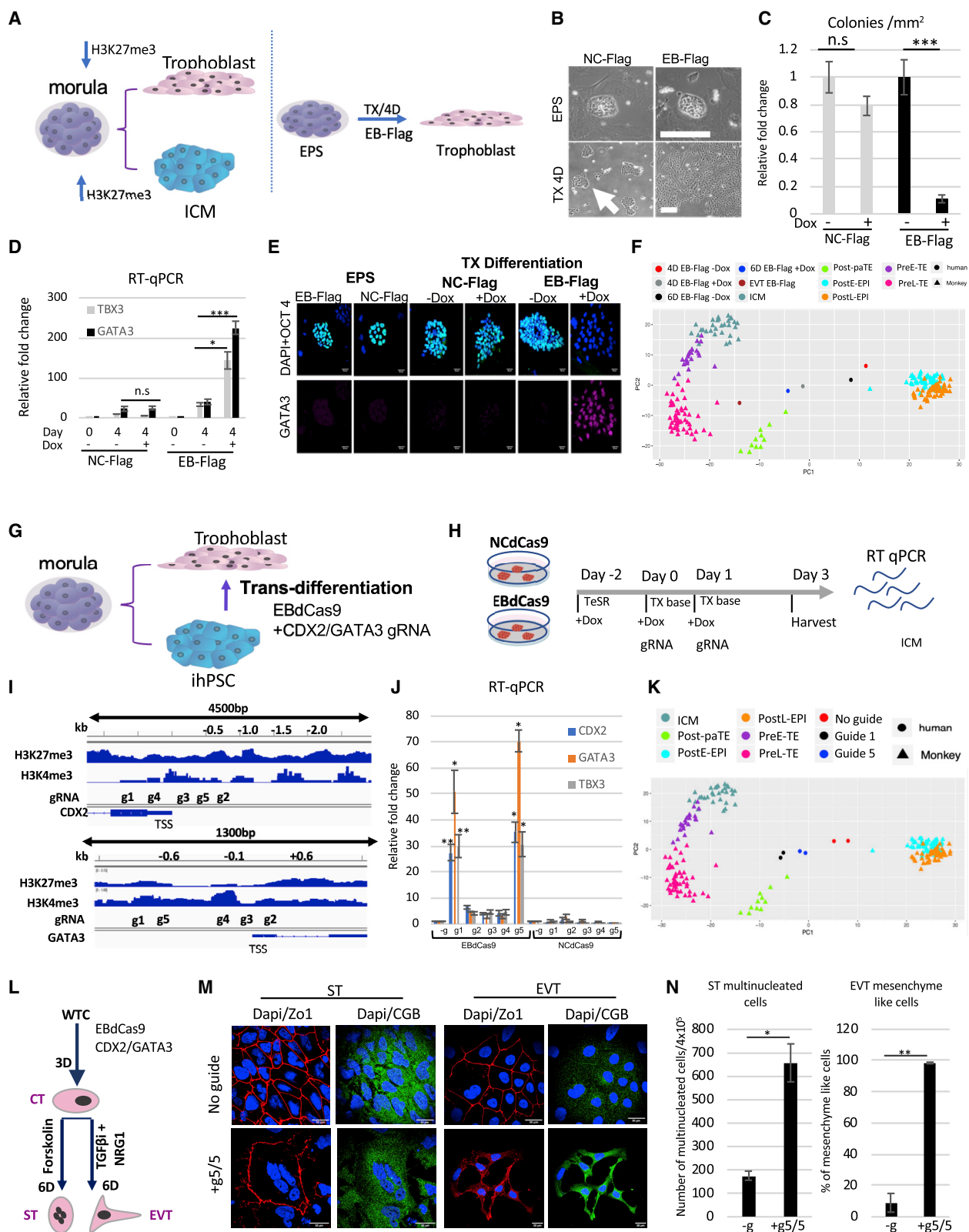
Control of epigenomic regulation holds immense therapeutic promise for human disease without manipulation of endogenous gene sequences. Here, we develop a system for the targeted inhibition of PRC2 that can identify the precise nucleosomes that require H3K27me3 marks for transcription repression. We identified a distal TATA box that is regulated by PRC2 repression and required for transcript activation. We show that this PRC2-inhibiting tool, EBdCas9, allows the study of epigenomic memory of PRC2 at specific loci. Furthermore, we show the general applicability of EBdCas9 by identifying the regions where PRC2 disruption induces transcription of bivalent genes (*TBX18*, *p16*, *CDX2*, and *GATA3*). In total, we have targeted 26 specific sites in promoter regions upstream of four different genes and observed significant PRC2-dependent transcriptional de-repression in 8 loci. NCdCas9, which differs from EB by two amino acids required for EED binding, does not result in transcriptional changes. This suggests that EBdCas9 functions by locally inhibiting PRC2 activity. When targeting PRC2 in the promoter regions of known TE transcription factors *CDX2* and *GATA3*, EBdCas9 application identified specific loci in which PRC2 elimination is sufficient to upregulate the gene transcription and therefore induce TE cell-fate commitment. Furthermore, EBdCas9/g-based epigenetic control of *p16* can result in reduced cell growth in DMG tumor cells.

dCas9 is the most promising method for targeted delivery of agents to chromatin. However, previous fusions of dCas9 to enzymes can result in off-target activity outside of the targeted locus. Existing histone modifying enzyme fusions are continuously active, potentially introducing or removing marks at off-target locations independent of the endogenous epigenetic modification machinery. In contrast, the computationally designed protein EB fused to dCas9 (EBdCas9) with appropriate gRNA presented

### Figure 5. EBdCas9 upregulates *CDKN2A* (*p16*) expression and compromises cell viability

- Integrative genomic viewer of *CDKN2A* H3K27me3 and H3K4me3 promoter tiling.
- Timeline of EBdCas9 or NCdCas9 induction and gRNA transfection.
- qRT-PCR analysis of EBdCas9 or NCdCas9 after 3 days of transfection with single gRNA (g1–g8) normalized to  $\beta$ -actin and relative fold change compared with no guide (–g).
- TATA boxes/initiator representation with respect to surrounding guide loci using the Element Navigation Tool for detection of core promoter elements.
- Immunofluorescent imaging of EBdCas9 after 3 days of transfection with single *p16* gRNA 1 (+g1) or no transfection (–g). Blue, DAPI; green, *p16*; far red, *p21*; scale bar, 100  $\mu$ m.
- Cell viability of EBdCas9 and NCdCas9 transfected with *p16* g1 (+g1) compared with no guide (–g). Time points were analyzed every 6 h using Alamar Blue ( $n = 3$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).
- EBdCas9 timeline for measuring epigenetic remodeling.
- qRT-PCR of *p16* relative fold change of EBdCas9 and NCdCas9 after 3 days of Dox induction and *p16* g1 transfection. Samples were normalized to  $\beta$ -actin and compared with no guide (–g) of each respective cell line.
- ChIP-qPCR of induced (+Dox) EBdCas9 and NCdCas9 after 3 days of transfection with *p16* g1 RNA (+g1) or no transfection (–g). Antibodies used for ChIP are listed above (Cas9, *EZH2*, H3K27me3). Normalized to fraction input/H3 ( $n = 3$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).
- H3K4me3 marks increase after EBdCas9/*p16* g1 (+g1) induction. CUT&RUN analysis of induced (+Dox) EBdCas9 after 3 days of transfection with *p16* g1 RNA (+g1) or no transfection (–g) for H3K4me3. H3K4me3 tracks are normalized to immunoglobulin G (IgG) and displayed on the Integrated Genome Viewer (IGV); 40 kb region of the *CDKN2A* (*p16*) gene displayed in the integrated genome viewer.





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here is able to precisely inhibit PRC2 at specific genetic loci, enabling (1) target and inhibition of PRC2 at a single nucleosome level; (2) reduced H3K27me3 at precise, targeted loci; (3) induced targeted transcription; (4) mediation of neighborhood spreading of remodeled epigenomics; (5) generation of epigenomic memory; (6) discovery of functional distal TATA box and TBP >500 bp of the TSS; (7) change of cell functionality; (8) trans-differentiation of one cell fate to another; and (9) reduction of tumor cells. Recently, several groups have activated targeted genes using synergistic activation mediator (SAM) such as dCas9-VP64, sgRNA, and MS2/p65/HSF1 (Konermann et al., 2015; Liao et al., 2017; Joung et al., 2017; Weltner et al., 2018) by recruitment of transcription factors to the promoter region. Our approach of targeted PRC2 inhibition utilizing EBdCas9/gRNA allows organic expression of targeted genes as the cell makes holistic decisions for transcriptional activation. EBdCas9 function is specific to PRC2-repressed genes.

One of the fundamental findings of this study is that most broad H3K27me3 marks on promoter regions appear indiscriminate. Using EBdCas9/gRNA, we identified key PRC2-dependent single nucleosomes on promoter regions that control gene expression. In eukaryotes, genes are classified as TATA-containing and TATA-less in their promoter regions (Basehoar et al., 2004). The TATA-containing core promoter, in a range of –150 to –1 relative to the TSS, constitute ~17% of the total promoters in yeast, ~10% in worms, ~14% in fruit flies, ~10% in zebrafish, and ~3% in human and mouse, making 97% of core promoters TATA-less (Yella and Bansal, 2017). If the –500 to +500 region spanning the TSS is considered, the number of predicted TATA-containing promoter sequences grows to 14% (Bansal et al., 2014). While TATA-containing promoters have a TATA sequence at region –30 to –1 relative to the TSS region,

TATA-less promoters have higher guanine-cytosine (GC) content in the TSS –150 to –1 region (Yella and Bansal, 2017; Yang et al., 2007). In addition, human TATA-less promoters have different structural motifs such as G-quadruplex (GNGNGNG) and CpG islands that are enriched at approximately ~42% at TATA-less promoter regions (Yella and Bansal, 2017). In the scenario of a distal TATA box, we hypothesize that a G-quadruplex structure contributes to active transcription as nucleosome-depleted regions in euchromatin associate with H3K4me3, RNA Pol II, and transcription factor recruitment (Hansel-hertsch et al., 2016). Here, we identify single nucleosomes on key promoter regions that are transcriptionally responsive to EBdCas9, suggesting that these regions are normally repressed due to PRC2 activity. In the case of *TBX18*, we show that PRC2-dependent sites co-localize with the distal TATA box and initiator binding elements. Furthermore, we show that the TATA box region is required for EBdCas9-based gene upregulation since TBP is recruited to the TATA box site after PRC2 de-repression by EBdCas9. CRISPR-Cas9-based deletion of the TATA box region eliminates TBP recruitment, which subsequently results in transcript inactivation. This suggests that remote TATA boxes are regulated by biological signals culminating in PRC2 activity. We now show that the EBdCas9 biological tool can reveal PRC2-dependent, masked regulatory elements on promoter regions through EBdCas9/gRNA multiplex promoter screens.

The combination of controlled epigenomic gain- and loss-of-function manipulations are the most desirable for elastic gene-expression-based epigenomic memory. In the future, EBdCas9 expression under tissue-specific promoters or by transfection of ribonucleoproteins (RNPs) will allow us to explore the *in vivo* applicability of this technology. The adaptive and efficient targeted PRC2 inhibition by EBdCas9 identifies functional

#### Figure 6. Trophoblast trans-differentiation using EBdCas9

- (A) Morula bifurcation to trophoblast and ICM is H3K27me3 dependent (left panel). EPS differentiation using EB-FLAG and TX media (Kubaczka et al., 2014) (right panel).
- (B) WTC EB-FLAG or WTC NC-FLAG as described previously (Moody et al., 2017) were reprogramed to EPS for 2 weeks (3 passages). Bright field of colony morphology of EPS EB-FLAG or NC-FLAG induced with Dox on MEFs for 4 days or plated on matrigel in TX media (TGF- $\beta$ 1, FGF4, and heparin) for 4 days.
- (C) Colony morphology count following 4 days in TX media EB-FLAG compared with NC-FLAG ( $n = 3$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).
- (D) qRT-PCR of EB-FLAG and NC-FLAG either in EPS stage or 4 days in trophoblast differentiation (TX media) with (+) or without (–) Dox. Normalized to  $\beta$ -actin and relative to EPS EB-FLAG stage ( $n = 3$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).
- (E) Immunofluorescence of EPS EB-FLAG or EPS NC-FLAG on MEF/LCDM media or differentiation using matrigel/TX media with (+) or without Dox (–). DAPI, blue; WGA, red; OCT4, green; GATA3, far red. Scale bar, 50  $\mu$ m.
- (F) PCA of EPS samples compared to monkey single-cell RNA-seq (Nakamura et al., 2017). EB-FLAG EPS cells were differentiating in TX media with or without Dox for 4 days or 6 days or passaged 3 times as extravillous cytotrophoblasts (EVTs). Cell types in the monkey single-cell data include the following: Post-paTE, post-implantation parietal trophoblast; PreL-TE, pre-implantation late TE; PreE-TE, pre-implantation early TE; ICM, inner cell mass; Pre-EPI, pre-implantation epiblast; PostE-EPI, post-implantation early epiblast; PostL-EPI, post-implantation late epiblast.
- (G) Model of WTC EBdCas9 trans-differentiation to trophoblasts using *CDX2* and *GATA3* gRNA cocktail.
- (H) Timeline of EBdCas9 or NCdCas9 induction and gRNA transfection.
- (I) Tiling of *CDX2* and *GATA3* promoter and gene body gRNA.
- (J) qRT-PCR analysis of *CDX2* and *GATA3* of co-transfected gRNAs in the presence of EBdCas9 or NCdCas9 induction (+Dox). g1–g5 correspond to g1/g1, g2/g2, g3/g3, g4/g4, and g5/g5 *CDX2/GATA3* co-transfection. Normalized to  $\beta$ -actin and compared with no guide (–g) ( $n = 3$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).
- (K) PCA of day 3 WTC EBdCas9 co-transfected with g5/g5 *CDX2/GATA3* and g1/g1 *CDX2/GATA3* or not transfected (–g) bulk RNA-seq compared with monkey single-cell RNA-seq (Nakamura et al., 2017). Cell types are exactly as in (F).
- (L) Timeline of EVT and syncytio trophoblast (ST) differentiation after EBdCas9:*CDX2/GATA3* g5,g5 transfection (cytotrophoblast [CT]-like stage) and the factors used.
- (M) Immunofluorescence of EBdCas9 3 days post-*CDX2/GATA3* g5,g5 RNA transfection compared with no guide and further 6-day differentiation to either EVT using differentiation factors (see STAR Methods). DAPI, blue; ZO-1, red; chorionic gonadotropin beta (CG $\beta$ ), green. Scale bar, 30  $\mu$ m.
- (N) Quantification of ST- (multinucleation) and EVT- (mesenchyme-like morphology) positive cell count. Area of total count is 18 mm<sup>2</sup> ( $n = 2$ , SEM; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , two-tailed  $t$  test).

H3K27me3 marks and mediates gene activation, which holds promise both as an epigenetic tool in biomedical research and as an approach for treating a wide range of human diseases, including cancer.

### Limitations of the study

In this study, we show that EBdCas9 inhibits local H3K27me3 and EZH2 levels. Since EBdCas9/gRNA targets 2 loci in each cell, local co-immunoprecipitation experiments of EED are technically limiting. Additionally, the gRNAs designed in this study were selected using CRISPR-Cas9 tools that are optimized for PAM sites recognition only. This limits the study to only those sites that are in close proximity to PAM sites.

### STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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### SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.celrep.2022.110457>.

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### AUTHOR CONTRIBUTIONS

S.L. and H.R.-B. conceived and designed the experiments. L.S., I.X.R., D.I.-M., A.P., S.S., and H.H. executed the molecular biology and biochemistry work. W.I.N. and A.A. analyzed the RNA-seq datasets. D.M. and N.M. analyzed the ChIP-seq datasets, J.D. performed the super-resolution microscopy. D.B., R.D.H., J.M., Y.W., N.V., K.B., and H.R.-B. provided materials and/or advice. S.L. and H.R.-B. wrote and edited the manuscript with input from the other authors.

### DECLARATION OF INTERESTS

S.L., H.R.-B., and D.B. are co-inventors on US patent application no. 17/434,832. S.L. is a founder and stockholder at Histone Therapeutics Corp., a company that aims to develop inventions described in this manuscript. K.B. is a co-founder, board member, and equity holder of Matchstick Technologies, Inc., and the developer and maker of the PIXUL instrument.

### INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in science. One or more of the authors of this paper self-identifies as a member of the LGBTQ+ community. One or more of the authors of this paper self-identifies as living with a disability. One or more of the authors of this paper received support from a program designed to increase minority representation in science.

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## STAR★METHODS

### KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
Anti-OCT4	Santa Cruz	Cat#sc-5279; RRID: AB_628051
Anti-TBX18	Santa Cruz	Cat#sc-514486
Anti-mCherry	Abcam	Cat#ab183628
Anti-EZH2	cell signaling	Cat#5246s
Anti-H3K27me3	active motif	Cat#39155
Anti-EED	EMD Millipore	Cat#09-774
Anti-JARID	cell signaling	Cat#13594s
Anti-H3K27ac	Active motif	39133
Anti-p300	Santa Cruz	sc-584
Anti-Pol II CTD	santa cruz	sc-47701
Anti-Pol II CTD PS5	covance	MMS-134R-200
Anti-p16	thermo fisher Scientific	MA517054
Anti-p21	cell signaling	Cat#2946
Anti-H3K4me3	abcam	ab272143
Anti-H3	abcam	AB1791
Anti-GATA3	cell signaling	Cat#5852
Anti-CGB	Invitrogen	Cat#PA558598
Anti-HLA	Novus Biologicals Inc	Cat#NBP1-43123
Anti-Actin	Cell Signaling	Cat#4970S
Anti-Flag	Sigma-Aldrich	Cat#F7425
Anti-Cas9	cell signaling	Ab #14697
Anti 5mC	Active Motif	AB_2793653
Anti 5hmC	Active Motif	AB_10013602
<b>Bacterial and virus strains</b>		
10-BETA Competent Cells	New England BioLabs	C3019H
DH5-alpha Competent Cells	New England BioLabs	C2987I
<b>Chemicals, peptides, and recombinant proteins</b>		
Matrigel	Corning	356231
Doxycycline	Millipore-Sigma	D9891-10G
Neurobasal	Thermo Fisher Scientific	21103-049
N2 supplement	Thermo Fisher Scientific	17502-048
B27 supplement	Thermo Fisher Scientific	12587-010
knockout serum replacement	Thermo Fisher Scientific	A3181502
hLIF	SPEED Biosystems, LLC	YSP1249
GS3Ki	Selleckchem	S7566 (2017), S2924 (2014), 72054 (2016)
Rocki Y27632	R & D Systems (R&D)	1254
(S)-(+)-Dimethindene maleate	santa Cruz Biotechnology Inc	sc-361329
Mino	Santa Cruz Biotechnology Inc	sc-203339
IWR	Selleckchem	S7086
DMEM/F12 w/o HEPES or L-Glu	Invitrogen (Gibco/BRL Life Tech)	21331-020
l-ascorbic acid 2-phosphate Mg	sigma	A8960-5G
Insulin-Transferrin-Selenium	Fisher Scientific	41400045
NaHCO3	sigma	S7277
FGF4	Life technologies	PHG0263
TGF-beta1	peprotech	100-21

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**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Heparin	Stem Cell Technologies	7980
ITS-X supplement	Thermo fisher	51500056
NRG1	Cell Signaling	5218SC
A83-01	Miltenyi Biotec Inc.	130-105-333
Forskolin	StemCell Technologies	72112
Alamar Blue	Thermo fisher Scientific	A50100
Click iT EdU	ThermoFisher Scientific	C10337

**Critical commercial assays**

qiagen PCR purification	Quiagen	28104
qiagen gel purification	Quiagen	28706
mini-prep	Quiagen	27104
maxi-prep	Quiagen	12662
T7 maxi invitro transcription	invitrogen	AM1314
Iscrip	biorad	1708891
RT	Applied Biosystems	205111
RNAiMax	invitrogen	13778030
spCas9	sigma	CAS9PROT
Cell Line Nucleofector™ Kit V	lonza	VCA-1003
Quick Extract	Lucigen	QE09050
ExoSAP-IT	ThermoFisher	78200.200.UL
GoTaq® DNA Polymerase	promega	M3001
pGEM®-T Easy Vector Systems	promega	A1360
Q5® High-Fidelity DNA Polymerase	NEB	M0491S

**Deposited data**

RNA-seq data	GEO	GSE155022
ChIP-seq data	GEO	GSE195552
Cut and Run data	GEO	GSE195553
ATAC-seq data	GEO	GSE195699
DEseq and bulk RNAseq	This study	<a href="#">Data S1</a>

**Experimental models: Cell lines**

WTC-11	Coriell Institute	GM25256
Elf1-iCas9	Ferreccio et al.	RRID:CVCL_VR49
IR-MEF	ISCRM hESC CORE	N/A
DIPG	gift from M Monje at Stanford University	N/A
WTC-11 EBdCas9	This study	N/A
WTC-11 NCdCas9	This study	N/A
WTC-11 EBdCas9 TBX18	This study	N/A
TATAdeletion clone 11		
WTC-11 EBdCas9 TBX18	This study	N/A
TATAdeletion clone 23		
Elf1 EBdCas9	This study	N/A
Elf1 NCdCas9	This study	N/A

**Oligonucleotides**

See <a href="#">Table S5</a> for oligonucleotide information	N/A
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**Recombinant DNA**

KRAB-dCas9	Concklin Lab	<a href="#">Mandegar et al., 2016</a>
EB-Flag	Our Lab	<a href="#">Moody et al., 2017</a>
NC-Flag	Our Lab	<a href="#">Moody et al., 2017</a>

(Continued on next page)

**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
EBdCas9	This paper	N/A
NCdCas9	This paper	N/A
<b>Software and algorithms</b>		
Element	Element	<a href="http://lifefaculty.biu.ac.il/gershon-tamar/index.php/element-description/element">http://lifefaculty.biu.ac.il/gershon-tamar/index.php/element-description/element</a>
Fantom	FANTOM 5	<a href="http://fantom.gsc.riken.jp/5/">http://fantom.gsc.riken.jp/5/</a>
R	The R Foundation	<a href="https://www-r-project-org.offcampus.lib.washington.edu/">https://www-r-project-org.offcampus.lib.washington.edu/</a>
ImageJ	NIH	<a href="https://imagej-nih-gov.offcampus.lib.washington.edu/ij/">https://imagej-nih-gov.offcampus.lib.washington.edu/ij/</a>
Microsoft office	Microsoft	<a href="https://products.office.com">https://products.office.com</a>
Snapgene	Snapgene	<a href="https://www.snapgene.com">https://www.snapgene.com</a>
Transcription factor prediction Tool		<a href="http://www.ifti.org/cgi-bin/ifti/Tfsitescan.pl">http://www.ifti.org/cgi-bin/ifti/Tfsitescan.pl</a>
QGRS Mapper Tool		<a href="http://bioinformatics.ramapo.edu/QGRS/analyze.php">http://bioinformatics.ramapo.edu/QGRS/analyze.php</a>
GEO DOI	This paper	<a href="https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE195555">https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE195555</a>

## RESOURCE AVAILABILITY

### Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact. Hannele Ruohola-Baker ([hannele@uw.edu](mailto:hannele@uw.edu))

### Materials availability

All unique constructs and stable cell lines generated in this study are available from the lead contact with a completed Uniform Biological Materials Transfer Agreement.

### Data and code availability

- RNA sequencing, ChIP sequencing, Cut and Run sequencing and ATAC sequencing data have been deposited at NIH GEO and are publicly available as of the date of publication. Accession numbers are listed in the key and resources table. Bulk RNAseq and DEseq values are found in [Data S1](#). Original western blot images, RT-qPCR Ct, ChIP-qPCR Ct, and plate reader values are available in [Data S2](#). Microscopy data reported in this paper will be shared by the lead contact upon request.
- All original code has been deposited at NIH GEO and is publicly available as of date of publication. DOI is listed in the [key resources table](#).
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

hiPSC line WTC #11, previously derived in the Conklin laboratory ([Kreitzer et al., 2013](#)), WTC line EB-Flag and NC-Flag, previously derived in our lab ([Moody et al., 2017](#)), Elf iCas9, previously derived in our lab ([Ferrecchio et al., 2018](#)) and DMG (DIPG), a gift provided by M Monje at Stanford University were used in this study.

## METHOD DETAILS

### hiPSC and hESC Cell culture

The hiPSC line WTC #11, previously derived in the Conklin laboratory ([Kreitzer et al., 2013](#)), was cultured on Matrigel (1:30) growth factor-reduced basement membrane matrix (Corning) in mTeSR media (StemCell Technologies). Cells were passaged using versene once reaching 70% confluency and plated at 1:6 density. For EPS conditions, cells were grown as previously described ([Yang et al., 2017b](#)). Briefly, WTC cells were reprogrammed in base medium containing 100 mL DMEM/F12, 100 mL Neurobasal, 1 mL N2 supplement, 2 mL B27 supplement, 1% GlutaMAX, 1% NEAA, 0.1 mM  $\beta$ -mercaptoethanol, penicillin-streptomycin and 5% KSR, and freshly supplemented with 10 ng/mL hLIF, GSK3i (1  $\mu$ M), ROCKi (2  $\mu$ M), (S)-(+)-Dimethindene maleate (2  $\mu$ M; Tocris), Minocycline

hydrochloride (2  $\mu$ M; Santa Cruz Biotechnology) and IWR-endo-1 (0.5–1  $\mu$ M; Selleckchem). Cells were adapted to EPS conditions for at least 3 passages before analysis. EPS cells were pushed toward differentiation using TX media (Kubaczka et al., 2014): TX medium formulation was DMEM/F12 without HEPES and L-glutamine (Life Technologies), 64 mg/l L-ascorbic acid-2-phosphate magnesium, 14 mg/L sodium selenite, 19.4 mg/L insulin, 543 mg/L NaHCO<sub>3</sub>, 10.7 mg/L holo-transferrin (all Sigma-Aldrich), 25 ng/mL human recombinant FGF4 (Reliatech), 2 ng/mL human recombinant TGF- $\beta$ 1 (PeproTech), 1  $\mu$ g/ml heparin (Sigma-Aldrich), 2 mM L-glutamine, 1% penicillin, and streptomycin (all PAN-biotech). Medium was prepared without growth factors (TX-growth factors) and stored at 4°C. To prepare complete TX, the growth factors: FGF4, heparin, and TGF-  $\beta$ 1 were added prior to use. Medium was changed every other day. All cells were cultured at 37°C in 5% CO<sub>2</sub>. For Ebf1 iCas9 conditions, cells were treated as described before (Ferrecio et al., 2018) but adapted to TeSR culture media and single cell passaging: the cells were grown on Matrigel (1:30) growth factor-reduced basement membrane matrix (Corning) in mTeSR media (StemCell Technologies) and passaged using versene once reaching 70% confluency and plated at 1:6 density.

### **EBdCas9 and NCdCas9 plasmid construction**

we used the AAVS1 TREG KRAB-dCas9 plasmid previously derived in the Conklin laboratory (Kreitzer et al., 2013), eliminated KRAB using PacI and AgeI restriction cut and then constructed and religated the EEDbinder-linker-dCas9-NLS-mCherry (EBdCas9) or EEDbinder Negative Control-linker-dCas9-NLS-mCherry (NCdCas9) to the cut plasmid, screened colonies and verified the sequence by Sanger sequencing (SI Table S5).

### **Insertion of inducible EBdCas9 and NCdCas9 into AAVS1 site of WTC cells**

1x10<sup>6</sup> cells of WTC p42 were transfected with 5mg AAVS1-TALEN R plasmid (Addgene #59026), 5 $\mu$ g AAVS1-TALEN L plasmid (Addgene #59025), and 5mg donor plasmid (AAVS1 TREG EBdCas9 or AAVS1 TREG NCdCas9) using the Amaxa Lonza Human stem cell Kit #2. The cells were then plated with 5 $\mu$ M of Rock inhibitor (ROCKi) onto 10cm with fresh media. Three days following the nucleofection, the cells were selected for neomycin resistance with Genetecin (50mg/ml) for four days. Clones that survived after selection were expanded as a pool. The clones were plated onto matrigel with or without Doxycycline (2 $\mu$ g/ml) and RNA was extracted in order to analyze the level of Cas9 expression by qPCR. Insertion of EBdCas9 or NCdCas9 into the AAVS1 site was confirmed by cellular genomic isolation, PCR amplification and Sanger sequencing.

### **Trophoblast differentiation using EB-Flag/NC-Flag EPS cells**

50k-100k EPS Cells grown on matrigel (1:30) coated plate in TX base medium (DMEM/F12 w/o HEPES or L-Glu (21331-020), 64mg/ml ascorbic acid phosphate Mg, 14ug/l sodium selenite, 19.4mg/L insulin, 10.7 mg/L holo-transferrin, 543mg/L NaHCO<sub>3</sub>, 1% Pen Strep and, 2mM Glutamax) and TX factors (25ng/ml FGF4, 2ng/ml TGF-  $\beta$ 1 1 mg/mL Heparin) TX media was changed every other day. 2 $\mu$ g/ml Doxycycline was added to the medium on day of differentiation.

### **Trophoblast trans-differentiation using EBdCas9/NCdCas9 WTC cells**

100k WTC cells were plated on matrigel plates (1:30) in the presence TeSR and Doxycycline (2mg/ml) for 48h. Prior to gRNA transfections the media was changed to TX media as described in derivation and maintenance of murine trophoblast stem cells under defined conditions (Kubaczka et al., 2014) (without factors). gRNAs were transfected using RNAiMAX for 2 consecutive days and were either harvest for RT-qPCR, ChIP-qPCR at day 3 or continued for further EVT/ST differentiation using the below media conditions for EVT or ST, without EB or NC induction.

### **EVT/ST differentiation**

3d EBdCas9 TX cells were grown to 80% confluency in TX medium (Kubaczka et al., 2014) and dissociated with 0.05% Trypsin-EDTA for 5 min at 37° and harvested as described in derivation of human trophoblast stem cells (Okoe et al., 2018). Briefly, trypsinization was stopped using EVT/ST base medium (DMEM/F12 0.1 mM 2 mercaptoethanol, 0.5% Penicillin-Streptomycin, 0.3% BSA, 1% ITS-X supplement, 2.5 mM Y27632 (Rock i), 4% KnockOut Serum Replacement) and cells were plated on matrigel (1:30) 0.75x10<sup>5</sup> density per well and cultured in 2 mL of EVT/ST medium with either EVT differentiation factors (100 ng/mL NRG1, 7.5 mM TGF- $\beta$ i A83-01) or ST differentiation factor (2mM forskolin). Cell were cultured for additional 6 days, with media changes every other day. On day 9 cells were fixed and permeabilized for immunofluorescent staining and imaging.

### **Guide RNA design, synthesis and transfection**

The gRNAs targeting *TBX18*, *p16*, *CDX2* and *GATA3* genes were designed using the CRISPRscan web prediction tools (Moreno-mateos et al., 2015) and ordered as T7-gRNA primers. The T7-gRNA forward primer and the reverse scaffold primer were used in primer extension reaction to synthesize a double stranded DNA fragment by using Q5 High Fidelity-based PCR (New England Biolabs) followed by PCR purification (Qiagen). The 120 bp dsDNA served as a template for IVT (MAXIscript T7 kit, applied Biosystems). The RNA was then purified using Pellet Paint Co-Precipitant (Novagen). For transfection, 1x10<sup>5</sup> WTC EBdCas9 or 1x10<sup>5</sup> NCdCas9 cells were seeded at day 0, and treated with Dox (2 $\mu$ g/ml) for 2 days before and during transfection. On day 2 cells were transfected with gRNAs using Lipofectamine RNAiMAX (Life Technologies). gRNA was added at a 40 nM final concentration when added alone or 20nM in co-gRNA transfection. A second transfection was performed after 24 h. All no guide controls were induced by Dox and harvested with



study groups. Two days after the last gRNA transfection, cells were harvest for either DNA, RNA, protein, ChIP-qPCR, or Cut and Run analysis (SI Table S5). N = 1 for gRNA transfection. p value was calculated based on 3 independent transfections.

### Elf iCas9 transfection

Elf iCas9 cell line (Ferreccio et al., 2018) was induced with Dox for 2 days prior to gRNA (SI Table S5) transient transfection. individual gRNAs were transfected transiently (800ng/ml) and cell were collected for genomic DNA isolation, amplification of designated regions and indel validation using Sanger sequencing.

### DNA extraction and sequencing

Genomic DNA was collected using DNAzol reagent (Invitrogen) according to manufacturer's instructions and quantified using Nano-drop ND-1000. Genomic regions flanking the AAVS1 were PCR amplified with the designed primers, purified by PCR Purification Kit (Invitrogen) and sent to Genewiz for sequencing.

### RNA extraction and RT-qPCR analysis

RNA was extracted using Trizol (Life Technologies) according to manufacturer's instructions. RNA samples were treated with Turbo DNase (ThermoFischer) and quantified using Nanodrop ND-1000. Reverse transcription was performed using iScript (BioRad). 10 ng of cDNA was used to perform qRT-PCR using SYBR Green, with suitable primers (SI Table S5) on an Applied Biosystems 7300 real-time PCR system with PCR conditions as stage 1 50°C for 2mins, stage 2 as 95°C for 10mis, 95°C for 15sec, 60°C for 1min(40 Cycles).  $\beta$ -actin was used as an endogenous control), related to Data S2.

### Protein extraction and western blot analysis

Cells were lysed directly on the plate with lysis buffer containing 20mM Tris-HCl pH 7.5, 150mM NaCl, 15% Glycerol, 1% Triton X-100, 1M  $\beta$ -Glycerolphosphate, 0.5M NaF, 0.1M Sodium Pyrophosphate, Orthovanadate, PMSF and 2.3% SDS. 25 U of Benzonase Nuclease (EMD Chemicals, Gibbstown, NJ) was added to the lysis buffer right before use. Proteins were quantified by Bradford assay (Bio-rad), using BSA as Standard using the EnWallac Vision. The protein samples were combined with the 4x Laemli sample buffer (900  $\mu$ L of sample buffer and 100  $\mu$ L  $\beta$ -Mercaptoethanol), heated (95°C, 5mins) and run on SDS-PAGE (protean TGX pre-casted gradient gel, 4%–20%, Bio-rad) and transferred to the Nitro-Cellulose membrane (Bio-Rad) by semi-dry transfer (Bio-Rad). Membrane was blocked for 1hr with 5% milk, and incubated in the primary antibodies overnight in 4°C. The antibodies used for Western blot were beta-Actin (Sigma A5441, 1:2000), Cas9 (Cell Signaling 1:1000), Oct-4 (Santa Cruz sc-5279, 1:1000, Novus Biologicals NB110-90606, 1:500), H3K27me3 (Active Motive 39,155 1:1000), EZH2 (Cell Signaling D2C9, 1:1000), GATA3 (Cell signaling D13C9, 1:1000), Flag (sigma F1804, 1:1000)The membranes were then incubated with secondary antibodies (1:10,000, goat anti-rabbit or goat anti-mouse IgG HRP conjugate (Bio-Rad) for 1hr and the detection was performed using the immobilon-luminol reagent assay (EMP Millipore) (SI Table S6).

### Immunostaining and confocal imaging

Cells were fixed in 4% paraformaldehyde for 15 min, washed with PBS (3  $\times$  5min), and blocked in the presence of 0.1% Triton X-100 and 2% BSA for 1h at room temp. Cells were then incubated in primary antibody overnight, washed with PBS (3  $\times$  5min), incubated with the secondary antibody in 2% BSA for 1hr, washed (4  $\times$  10mins, adding 1mg/ml DAPI at the second wash), mounted using Vectashield (Vector Labs) and imaged using a Leica TCS-SPE confocal microscope at 10x,40x or 63x. Antibodies use in immunostaining were: TBX18 (Santa Cruz sc-514486, 1:200), Oct 4 (Novus Biologicals, NB11- 1:150), p16 (Invitrogen MA5-14260, 1:250), p21 (Cell Signaling 2947S, 1:200), GATA3 (Cell signaling D13C9, 1:250), CGB (DAKO GA508; 1:200). Slides were rinsed with PBS-T and incubated with secondary antibodies: DAPI (0.02  $\mu$ g/mL, Molecular Probes), goat anti-rabbit 488, and goat anti-mouse 647 (1:500, Molecular Probes) for 2 h at room temperature in blocking buffer. (EBdCas9 or NCdCas9 were visualized using endogenous mCherry expression) (SI Table S6).

### Cell harvesting cross-linking and PIXUL sonication

PIXUL 96-well plate sonicator (Matchstick Technologies, Inc, Kirkland, WA and Active Motif, Carlsbad, CA) was used to fragment chromatin. All steps began from 1  $\times$  35 mm plate exactly as described in Guide RNA design, synthesis and transfection above. Cells were harvested using Versene (Thermo Fisher), followed by 3 min centrifugation (1.2 xg). Pelleted cells were washed with 1  $\times$  PBS. Cells were cross-linked by adding 500ul 1% formaldehyde in PBS for 20min in RT. Supernatant was removed and 500ul of PBS/glycine (125mM) was added for 5 min at RT for quenching. Supernatant was removed and the cells were washed with 500ul of PBS. PBS was removed and samples were stored in  $-80^{\circ}$ C. For shearing, cells were resuspended in 100ul chromatin sheering buffer (Active Motif) and transferred into wells of 96 well plate, sealed with PCR film (MiniAMP Optical Adhesive Film) and placed in PIXUL for shearing. PIXUL parameters were as follows: Cycles = 50; PRF = 1kHz; Burst = 20 for 6min  $\times$  4. Once sheered the cells were chromatin immunoprecipitated using matrix-ChIP. N = 1 for gRNA transfection and ChIP. p value was calculated based on 3 independent transfections.

### PIXUL-matrix-ChIP-qPCR (PIXUL-ChIP) analysis

PIXUL-ChIP was performed as previously described (Flanagan et al., 2008; Bomsztyk et al., 2019). Briefly, UV treated 96-well polypropylene microplates were incubated with protein A on a low-speed shaker at room temperature overnight. The next day, the wells of the 96-well plate-coated with protein A were blocked with blocking buffer containing 5% BSA, sheared salmon sperm DNA (10 µg/µL final) in immunoprecipitation buffer (150 mM NaCl, 50 mM Tris-HCl (pH 7.5), 5 mM EDTA, NP-40 (0.5% vol/vol), Triton X-100 (1.0% vol/vol)) on a shaker at room temperature for 60 min. Simultaneously, sheared chromatin samples (see above), blocking buffer and antibodies were added to a another polypropylene 96-well microplate (untreated) and incubated in ultrasonic bath for 60 min at 4°C. The blocking buffer was aspirated from the protein A-coated plate, and the chromatin + antibody mix from the untreated plate was transferred to the protein A-coated wells and incubated in the ultrasonic bath for 60 min at 4°C. The wells were then washed 3 times with immunoprecipitation buffer followed by 3 washes with TE buffer. Finally, elution buffer containing 25 mM Tris base, 1 mM EDTA (pH10) with proteinase K 200 µg/mL was added to the wells, nutated for 30 s at 1400 rpm and incubated for 45 min at 55°C and then 10 min at 95°C. The 96-well plates were then briefly agitated and centrifuged for 3 min at ~ 500 g at 4°C and were used for PCR. The antibodies (SI Table S6) used for Matrix ChIP were: mCherry (Abcam ab167453), H3K27me3 (Active motif 39,155), EZH2 (Cell Signaling D2C9), H3K27ac (Active motif 39,133), p300 (Santa Cruz sc-48343), Pol II 4H8 (Santa Cruz, sc-47701), Pol II Ser 5P+ (Santa Cruz), H3K4me3 (Active motif 39,159), Cas9 (Active motif 61,757). Matrix ChIP experiments were performed in triplicate followed by qPCR in 4–8 replicates. ChIP-qPCR analysis and calculation are described below.

### ChIP-qPCR calculations

All PCR reactions were run in quadruplicate using syber green using primer sets found in (SI Table S5). PCR calibration curves were generated for each primer set from a dilution series of total human genomic DNA. For each qPCR run the primer efficiency curve was fit to cycle threshold (CT) versus log (genomic DNA concentration) using an r-squared best fit. DNA concentration values for each ChIP and input chromatin DNA sample were calculated from their respective average Ct values as previously described (Flanagan et al., 2008; Bomsztyk et al., 2019). Final results are expressed as fraction of input DNA normalized to H3 antibody.

### PIXUL-matrix-ChIP-seq

Cells were dissociated using versene, washed once in cold PBS and counted. Approximately 6x10<sup>6</sup> cells were fixed and chromatin was PIXUL sonicated as described above. H3K27me3 (Active Motif Ab) Matrix-ChIP'ed DNA was purified using phenol-chloroform extraction and ChIP-seq libraries were prepared (Next Gen DNA Library Kit, Active Motif, Cat# 53,216, 53,264). Libraries were sequenced on NextSeq2000 in single end run and approximately 45 million raw reads were obtained for each sample. For analysis: reads were normalized via RPKM and aggregate read signal for all 22K genes was calculated in a 5k window centered at the TSS. Significance by first calculating average signal in 5k window for all TSS', then comparing means across conditions via t test.

### Cut and Run analysis

Cut and Run method and analysis was performed as previously described (Skene and Henikoff, 2017). 1million WTC EBdCas9 cells gRNA transfected or not were harvested by centrifugation (600 g, 3 min in a swinging bucket rotor) and washed in ice-cold phosphate-buffered saline (PBS). Nuclei were isolated by hypotonic lysis in 1 mL NE1 (20 mM HEPES-KOH pH 7.9; 10 mM KCl; 1 mM MgCl<sub>2</sub>; 0.1% Triton X-100; 20% Glycerol) for 5 min on ice followed by centrifugation as above. Nuclei were briefly washed in 1.5 mL Buffer 1 (20 mM HEPES pH 7.5; 150 mM NaCl; 2 mM EDTA; 0.5 mM Spermidine; 0.1% BSA) and then washed in 1.5 mL Buffer 2 (20 mM HEPES pH 7.5; 150 mM NaCl; 0.5 mM Spermidine; 0.1% BSA). Nuclei were resuspended in 500 µL Buffer 2 and 10 µL antibody was added and incubated at 4°C for 2 h. Nuclei were washed 3 x in 1 mL Buffer 2 to remove unbound antibody. Nuclei were resuspended in 300 µL Buffer 2 and 5 µL pA-MN added and incubated at 4°C for 1 h. Nuclei were washed 3 x in 0.5 mL Buffer 2 to remove unbound pA-MN. Tubes were placed in a metal block in ice-water and quickly mixed with 100 mM CaCl<sub>2</sub> to a final concentration of 2 mM. The reaction was quenched by the addition of EDTA and EGTA to a final concentration of 10 mM and 20 mM respectively and 1 ng of mononucleosome-sized DNA fragments from Drosophila DNA added as a spike-in. Cleaved fragments were liberated into the supernatant by incubating the nuclei at 4°C for 1 h, and nuclei were pelleted by centrifugation as above. DNA fragments were extracted from the supernatant and used for the construction of sequencing libraries. We have also adapted this protocol for use with magnetic beads (Skene and Henikoff, 2017).

### Matrix-MeDIP-qPCR

Single-stranded DNA (ssDNA) was isolated from iPSC (WTC) EBdCas9 +Dox cells either after 3 day treatment with TBX18 gRNA (+g6) or untreated (-g). DNA was sonicated for 6 min in PIXUL followed Matrix-MeDIP-qPCR protocol as described above and in (Flanagan et al., 2008; Bomsztyk et al., 2019; Yu et al., 2011). Note that cells were not fixed or quenched for MeDIP (SI Table S5).

### ATAC-seq data analysis

Protocol used for ATAC-seq is described in (Corces et al., 2017). Briefly, cells were dissociated using versene, washed once in cold PBS and counted. Approximately 500,000 cells were resuspended to final concentration of ATAC resuspension buffer (10mM Tris-HCl pH 7.4, 10mM NaCl, 3mM MgCl<sub>2</sub>) in the presence of 0.1% NP40, 0.1% Tween 20, and 0.01% Digitonin as previously describe in (Corces et al., 2017). Nuclei were pellet at 500 RCF for 10 min at 4°C and supernatant was aspirated. Cells were then resuspended to

final concentration of transposition mix (Tris-HCl 20mM, MgCl<sub>2</sub> 10mM, 20% Dimethyl Formamide, transposase 100nM, 1% digitonin, 10% Tween 20) and incubated at 37°C for 30 min in a thermomixer with 1000 RPM mixing. The reaction was cleaned up using Zymo DNA Clean and Concentrator-5 Kit (cat# D4014). The eluted DNA was amplified using NEBNext 2x MasterMix. KAPA Library Quantification kit (cat# KK4854) was used for final concentration determination.

### RNA-seq data analysis

RNA-seq samples were aligned to the custom human reference genome hg38 with EBdCas9 sequence inserted using Hisat2 (version 2.1.0) (Kim et al., 2019). Gene-level read counts were quantified using subread(v2.0.0) (Liao et al., 2014) using custom Ensembl GRCh38 gene annotations with EBdCas9 added. Custom reference genome and annotation files with EBdCas9 were generated as follows: 1) Genome sequence (GRCh38.p13) was downloaded from the GENCODE browser (07-08-2021) the transcript-level annotation was downloaded from the GENCODE browser (07-08-2021). 2) The EBdCas9 sequence and annotation were added at the end of the reference genome and annotation. Filtered estimated gene count matrices generated for each RNA-seq sample were normalized (median of ratios) with the R package DESeq2(v1.32.0) (Love et al., 2014) and subjected to differential expression analysis with significance cutoff of an actual fold-change of at least 2 (Log<sub>2</sub> FC > 1) with a false discovery rate (FDR) cutoff of 5% (adjusted p value < 0.05) was used to determine whether a gene was differentially expressed. Volcano plots were generated to display the list of DEG for each RNA-seq dataset using the R package EnhancedVolcano (v1.10.0) (Blighe et al., 2018).

For trophoblast transdifferentiation: RNA-seq samples were aligned to hg19 using Tophat (version 2.0.13) (Trapnell et al., 2009). Gene-level read counts were quantified using htseq-count using Ensembl GRCh37 gene annotations. Processed single cell RNA-seq data from Nakamura et al. (Nakamura et al., 2017) were used. Only genes expressed above 10 Reads Per Million in 3 or more samples were kept. t-SNE was performed with the *Rtsne* package, using genes with the top 20% variance across samples. Cluster labels from Nakamura et al. were used. A Principle Component Analysis (PCA) was performed using all of the cynomolgus monkey samples from Nakamura et al. (Nakamura et al., 2017) using R software. Genes used in the analysis were restricted to defined homologs expressed at non-zero Transcripts Per Million (TPM) in human *in vitro* cell lines, and in the preprocessed mouse and cynomolgus monkey single cell samples from Nakamura et al. RNA-seq data from human cell lines were corrected for batch effects using *ComBat* (Johnson et al., 2007). Human bulk RNA-seq samples were projected onto the PCA coordinate via matrix multiplication. Human, cynomolgus monkey and mouse RNA-seq data were separately centered and scaled within each species before PCA and projection was performed. These data can be found in the [Data S1](#).

### Cell count

Cells were lifted using Versene and resuspended in media for cell count using hemocytometer. 100ul of cell suspension were applied to hemocytometer using 10x objective. 4 most outward corners were counted, averaged, multiplied by 10<sup>4</sup>, and multiplied by the dilution factor to get the final number of cells.

### Cell viability

Cell viability assay was measured by AlamarBlue (thermofisher scientific, DAL1025). Cells were induced and transfected with gRNA as discussed above. 6h after last transfection, 10x of viability reagent was added to media. Time “0” was measured 4h after. Fluorescence measurements were read on PerkinElmer Envision plate reader after transferring 50ul of media from each condition to 96 well microplate reader.

### Cell proliferation and FACS analysis

Cell proliferation assay was measured using Click-iT EdU Cell Proliferation Kit (ThermoFisher Scientific, C10337). Approximately 0.5M DMG cells were transfected with either EBdCas9/p16 + g1, NCdCas9/p16 + g1, or untreated (-g) on day 0 in the presence of Dox (0.5ug/ml) following EdU incorporation (10uM) on day 1. On day 3 DMG were collected using accutase, washed twice with PBS and resuspended with 4% PFA. Subsequently, a Click-iT reaction was performed as directed by the manufacture’s protocol. Finally, cells were washed twice with PBS, resuspended in 500ul of PBS-5% FBS, analyzed by FACS and FlowJo software.

### CRISPR-based deletion of *TBX18* TATA box region

For WTC EBdCas9 *TBX18* TATA box deletion (40 bp deletion), we used CRISPR/Cas9 to delete an 88bp region by using 2 gRNAs flanking TATA box 5’ AND-3’ ends. Since gRNA 2 overlaps with *TBX18* g6 locus we reconstructed guide6 region using a ssDNA donor (168bp, IDT) that lacks the 40bp region around the TATA box (TATATGAC) but contains intact guide6 region and mutated PAM region. Two guides RNAs (Synthego) that are flanking TATA box region were designed using Benchling ([Table S5](#)). Briefly, 0.5μM of spCas9 (sigma) were mixed with 1μM of gRNAs and incubated at RT for 15min. 1x10<sup>6</sup> EBdCas9 cells were transfected with the above RNP complex and 100μM of ssDNA (HDR) ([Table S5](#)) using AMAXA electroporation kit (LONZA) and plated in TeSR media (supplemented with 5μM ROCKi). Several EBdCas9 colonies were picked, followed by chromosomal DNA isolation (Quick Extract/Lucigen), PCR amplification ([Table S5](#)) and PCR purification (ExoSAP-IT/ThermoFisher). PCR products were sequenced to identify potential 40bp TATA box region deletions and g6 reconstruction. To produce a homozygous deletion of *TBX18* TATA box and guide 6 reconstruction a second hit of CRISPR/Cas9 was introduced using the same exact 2 guide RNAs and ssDNA as describe above. Genomic DNA sequencing analysis of selected clones confirmed homozygous deletion of 40bp of TATA box region on *TBX18*. To validate that

the lines analyzed were homozygous TATA box deletion and guide6 reconstruction, we ligated PCR amplicons to pGEM-T (Promega) vector system and confirmed that all picked clones encompassed TATA box deletion and guide6 reconstruction using Sanger sequencing (Table S5).

#### ElemeNT Navigation Tool

TATA box prediction and other detection of core promoter elements were assayed using ElemeNT Navigation Tool (Sloutskin et al., 2015). Each analyzed sequence included 1000bp region. <https://www.juven-gershonlab.org/resources/element/run/>

#### Fantom5 cage tool

Functional annotation of the mammalian genome was used to evaluate *TBX18* transcription start site (TSS) against human RNAseq datasets. <https://fantom.gsc.riken.jp/5/ssstar/EntrezGene:9096>.

#### QGRS mapper tool

Distribution of putative Quadruplex forming G-Rich Sequences (QGRS) in nucleotide sequences was predicted using QGRS mapper too. <http://bioinformatics.ramapo.edu/QGRS/analyze.php>.

#### Transcription factor prediction tool

Prediction of transcription factor site scan on promoter region was analyzed using <http://www.ifti.org/cgi-bin/ifti/Tfsitescan.pl>.

#### SIM and image processing

Three-color, 3D-SIM images were acquired with a commercial OMX-SR system (GE Healthcare). Topica diode lasers with excitation at 405nm, 488nm, and 640nm were used. Emission was collected on three separate PCO.edge sCMOS cameras using an Olympus 60 × 1.42NA PlanApochromat oil immersion lens. 512 × 512 images (pixel size 6.5 μm) were captured with no binning. Acquisition was controlled with AcquireSR Acquisition control software. Z-stacks were collected with a step size of 125 nm and 15 images per optical slice (3 angles and 5 phases) using immersion oil with a refractive index 1.516, volumes varied between 2.0 and 3.5 μm per cell. Images were reconstructed using Wiener filter settings of 0.005 for the 405 nm (nuclei) channel and 0.001 for all other channels and optical transfer functions (OTFs) measured specifically for each channel with SoftWoRx 7.0.0 (GE Healthcare). Images from different color channels were registered using parameters generated from a gold grid registration slide (GE Healthcare). SIM data, including raw and reconstructed images, were subjected to SIMcheck FIJI Plugin (Ball et al., 2015) to ensure that image quality was within acceptable standards.

Cell images were analyzed using a combination of custom ImageJ/Fiji and python scripts. Images channels were separated followed by nuclear segmentation via Gaussian blur (sigma = 30) to even out illumination and auto thresholding using the Huang method (Liang-kai and Wang, 1995) and filling binary holes to create the final mask. This mask was then applied to each channel containing proteins of interest. The masked protein images were converted to 8-bit and background corrected (rolling ball radius = 2) followed by using the find maxima function in Fiji (prominence = 50) and measuring the mean intensity and location of each spot. A custom python script was used to determine puncta colocalization using a separation threshold distance of 0.25 μm as the cutoff. Colocalized puncta number and percentage relative to total puncta were recorded.

#### EZH2 chemical inhibitory drugs

Cells were plated on 6 well plates at day 0 and were treated with EPZ-6438 (Selleckchem, stock dilution 10mM) working solution of 1mM to final concentration of 5μM or untreated (DMSO) or treated on day 1. On day 3 the cells were harvested for RNA isolation and purification using Trizol. EPZ-6438, is a selective EZH2 inhibitor that is FDA approved for B-cell and follicular Lymphomas (NCT01897571, Epizyme). EPZ-6438 (EPZ) competes with the substrate S-adenosylmethionine (SAM) (Knutson et al., 2013).

#### QUANTIFICATION AND STATISTICAL ANALYSIS

All statistical methods for each experiment can be found in the Figure legends as well as in the [method details](#) section. p values and individual data points for the main figures can be found in the [Data S2](#).