SUMMARY

Silencing receptor GUCY2C tumor suppressor signaling occurs widely in colorectal tumors, reflecting loss of GUCY2C ligand expression. This study identifies a novel β-catenin/TCF-sensitive locus control region that mediates ligand loss and can be targeted for gene reactivation with CRISPR activation.

BACKGROUND & AIMS: Sporadic colorectal cancers arise from initiating mutations in APC, producing oncogenic β-catenin/TCF-dependent transcriptional reprogramming. Similarly, the tumor suppressor axis regulated by the intestinal epithelial receptor GUCY2C is among the earliest pathways silenced in tumorigenesis. Retention of the receptor, but loss of its paracrine ligands, guanylin and uroguanylin, is an evolutionarily conserved feature of colorectal tumors, arising in the earliest dysplastic lesions. Here, we examined a mechanism of GUCY2C ligand transcriptional silencing by β-catenin/TCF signaling.

METHODS: We performed RNA sequencing analysis of 4 unique conditional human colon cancer cell models of β-catenin/TCF signaling to map the core Wnt-transcriptional program. We then performed a comparative analysis of orthogonal approaches, including luciferase reporters, chromatin immunoprecipitation sequencing, CRISPR/Cas9 (clustered regularly interspaced short palindromic repeats) knockout, and CRISPR epigenome editing, which were cross-validated with human tissue chromatin immunoprecipitation sequencing datasets, to identify functional gene enhancers mediating GUCY2C ligand loss.

RESULTS: RNA sequencing analyses reveal the GUCY2C hormones as 2 of the most sensitive targets of β-catenin/TCF signaling, reflecting transcriptional repression. The GUCY2C hormones share an insulated genomic locus containing a novel locus control region upstream of the guanylin promoter that mediates the coordinated silencing of both genes. Targeting this region with CRISPR epigenome editing reconstituted GUCY2C ligand expression, overcoming gene inactivation by mutant β-catenin/TCF signaling.

CONCLUSIONS: These studies reveal DNA elements regulating corepression of GUCY2C ligand transcription by β-catenin/TCF signaling, reflecting a novel pathophysiological step in tumorigenesis. They offer unique genomic strategies that could reestablish hormone expression in the context of canonical oncogenic mutations to reconstitute the GUCY2C axis and oppose transformation. (Cell Mol Gastroenterol Hepatol 2022;13:1276–1296; https://doi.org/10.1016/j.jcmgh.2021.12.014)
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In the healthy intestine, expression of the paracrine hormone Wnt at the crypt base promotes a transcriptional program driving stem cell renewal, while tapering expression up the crypt-surface axis supports epithelial differentiation. Wnt signaling destabilizes a cytosolic destruction complex, assembled on the scaffold protein APC, which targets the transcription cofactor β-catenin for proteasomal degradation. Disruption of the complex enables accumulation and nuclear translocation of β-catenin, which partners with TCF/LEF transcription cofactors to activate proliferation programs. Mutations in the APC-β-catenin-TCF axis are the most common drivers of colorectal tumorigenesis.1,2 Indeed, 80% of sporadic tumors arise from loss-of-function mutations in APC, in which acquired mutations produce allelic heterozygosity, and loss of the nonmutant allele (loss of heterozygosity) eliminates wild-type protein expression. These mutations liberate β-catenin from regulation by Wnt signaling, permitting oncogenic transcription, polypl formation, and accumulation of subsequent mutations (eg, p53, KRAS) driving tumorigenesis.3,4

While these mutational events are well defined, mechanisms contributing to tumor initiation and progression continue to evolve. Guanylyl cyclase C (GUCY2C) and its downstream effector, cyclic guanosine monophosphate (cGMP), have emerged as key intestinal tumor suppressors.4–12 GUCY2C, a member of the family of transmembrane receptor guanylyl cyclases, is activated by the diarrheagenic bacterial heat-stable enterotoxins (STs) and the structurally homologous endogenous hormones guanylin (GUCA2A) and uroguanylin (GUCA2B).5,13,14 While GUCA2B and GUCA2A quantitatively predominate in the small and large intestine, respectively, mechanisms regulating those levels are unknown. Peptide stimulation of GUCY2C drives cGMP accumulation, promoting electrolyte and fluid secretion, and the synthetic peptides linaclotide and tenacotide target the receptor throughout disease progression.22–24

Hormones regulating the GUCY2C axis are central to the pathophysiology of intestinal tumorigenesis. However, mechanisms regulating GUCA2A and GUCA2B levels in health and disease remain undefined. Here, comprehensive RNA sequencing (RNA-seq) analysis of unique conditional human colon cancer cell models of Wnt signaling29–32 demonstrate that GUCA2A and GUCA2B are 2 of the most sensitive targets of APC-β-catenin-TCF regulation, reflecting transcriptional repression. Comparative analysis using orthogonal approaches to identify gene enhancers, including luciferase reporters, chromatin immunoprecipitation sequencing (ChiP-seq), CRISPRi/Cas9 (clustered regularly interspaced short palindromic repeats/Cas9) knockout (KO), and CRISPR epigenome editing reveals a novel locus control region regulated by APC-β-catenin-TCF that silences GUCA2A and GUCA2B transcription. Moreover, these studies offer unique genomic strategies, including CRISPR epigenome editing, that could re-establish hormone expression in the context of mutant APC-β-catenin-TCF signaling to reconstitute the GUCY2C axis and oppose transformation.

Results

GUCY2C Ligands Are 2 of the Most Sensitive Targets of Wnt Signaling in Colon Cells

While GUCY2C is expressed by colonicocytes along the rostral-caudal and crypt-surface axes, GUCA2A is produced by differentiated cells bordering the lumen and is absent in crypts (Figure 1A).13,14,33 This gradient is the inverse of
gene sets driven by Wnt, suggesting reciprocal regulation. Indeed, compared with undifferentiated mouse colon spheroids maintained in Wnt, mouse colonoids differentiated in the absence of Wnt exhibited a 36- and 31-fold increase in \( \text{Guca2a} \) and \( \text{Guca2b} \) messenger RNA (mRNA), respectively (Figure 1B and C). Also, 80% of colon tumors exhibit aberrant Wnt signaling arising from either APC loss-of-function or \( \beta \)-catenin gain-of-function mutations.\(^1,2\) Transcriptomic analysis of colon tumors (\( n = 380 \)) and normal mucosa (\( n = 51 \)) from The Cancer Genome Atlas (TCGA) confirmed a 131- and 385-fold higher expression of \( \text{GUCA2A} \) and \( \text{GUCA2B} \) mRNA in normal tissue, respectively.

Figure 1. Wnt signaling silences GUCY2C ligand expression. (A) Immunofluorescence of \( \text{GUCA2A} \) (red), GUCY2C (green), and nuclei (blue) in mouse colon crypts. (B) Mouse colon organoids cultured in the presence (+) or absence (−) of Wnt3a, and (C) corresponding Guca2a and Guca2b mRNA. (D) \( \text{GUCA2A} \) and \( \text{GUCA2B} \) mRNA quantified by RNA-seq from human colorectal tumor (T) (\( n = 380 \)) and normal (N) (\( n = 51 \)) tissue from TCGA (COAD/READ datasets). (E) Colon cancer cells express 1 of 3 inducible inhibitors of Wnt signaling: zinc-inducible wild-type APC (HT29 cells), DOX-inducible \( \beta \)-catenin shRNA (LS174T cells), or DOX-inducible dominant negative TCF7L2 (DNTCF; DLD1 and LS174T cells). (F, G) HT29( APC) cells treated with 300 \( \mu \text{M} \) zinc for 24 hours express APC, resulting in loss of \( \beta \)-catenin and upregulation of \( \text{GUCA2A} \) protein, \( \text{GUCA2A} \) mRNA, and \( \text{GUCA2B} \) mRNA. (H, I) LS174T( sh\( \beta \)-catenin) cells treated with 1 \( \mu \text{g/mL} \) DOX for 72 hours lose \( \beta \)-catenin and upregulate \( \text{GUCA2A} \) protein, \( \text{GUCA2A} \) mRNA, and \( \text{GUCA2B} \) mRNA. (J, K) DLD1(DNTCF) cells treated with 1 \( \mu \text{g/mL} \) DOX for 24 hours express DNTCF, upregulating \( \text{GUCA2A} \) protein, \( \text{GUCA2A} \) mRNA, and \( \text{GUCA2B} \) mRNA. (L, M) LS174T(DNTCF) cells treated with 1 \( \mu \text{g/mL} \) DOX for 48 hours express DNTCF, upregulating \( \text{GUCA2A} \) protein, \( \text{GUCA2A} \) mRNA, and \( \text{GUCA2B} \) mRNA. (C, F–M) Data points represent the average of 3 wells of cells from a single experiment, with the mean of 2–8 independent experiments indicated. Significance was determined by Student’s t test with matched analysis for independent experiments on log2-transformed results. Data are presented relative to noninduced cells. \(* P < .05; \ ** P < .01; \ *** P < .001; \ **** P < .0001\).
(Figure 1D). Further, activating a zinc-inducible wild-type APC transgene in HT29 human colon cancer cells, which harbor mutant APC, degrades β-catenin, disrupts Wnt signaling, and reconstitutes GUCA2A protein (11-fold), GUCA2A mRNA (15-fold), and GUCA2B mRNA expression (18-fold) (Figure 1F and G). Activation of a doxycycline (DOX)-inducible β-catenin short hairpin RNA (shRNA) in LS174T human colon cancer cells, which harbor gain-of-function mutations in β-catenin, reduces β-catenin accumulation and reconstitutes GUCA2A protein, GUCA2A mRNA, and GUCA2B mRNA expression (Figure 1H and I). Activation of a DOX-inducible dominant negative TCF7L2 (DNTCF) transgene in DLD1 and LS174T human colon cancer cells, which harbor mutations in APC or β-catenin, respectively, also restores GUCA2A protein, GUCA2A mRNA, and GUCA2B mRNA expression (Figure 1J–M). Transcriptomic analysis of these cell lines confirmed downregulation of the Wnt-β-catenin signaling gene set upon transgene induction (Figure 2A). Indeed, MYC Targets and Wnt-β-catenin signaling were among the 5 most downregulated gene sets in all cell lines (DLD1 gene sets shown in Figure 2B and C). Across all cell lines, 1289 genes were differentially regulated (Figure 2D, center of overlap), representing the core Wnt transcriptome, including canonical β-catenin-TCF targets such as MYC, AXIN2, SP5, ASCL2, and LGR5 (Figure 2E). Conversely, GUCA2A and GUCA2B were the 7th and 12th most upregulated transcripts among the 1289 genes.

**Mutant Wnt Signaling Represses GUCA2A Nuclear Transcription**

The above results show that the silencing of GUCY2C hormone expression by β-catenin-TCF reflects steady state transcript loss, agnostic to the relative contributions of transcriptional or posttranscriptional processes. Recently, we revealed that β-catenin-TCF regulated the incorporation of labeled uridine into GUCA2A mRNA, suggesting a level of transcriptional control. Here, silencing Wnt signaling did not regulate expression of luciferase reporter constructs containing the GUCA2A 3′ untranslated region (UTR) (Figure 3A and B) or expression constructs containing the full-length GUCA2A mRNA coding region (Figure 3C and D), suggesting little posttranscriptional regulation of steady-state transcript levels. In contrast, silencing Wnt signaling fully reconstituted nuclear GUCA2A mRNA levels, both in whole cells and their cytoplasm (Figure 3E and F). Reconstituted RNA transcripts contained GUCA2A introns, representing precursor mRNA (preRNA) (Figure 3G and H). Thus, APC-β-catenin-TCF regulation of GUCY2C hormone expression occurs at the earliest steps of transcription in the nucleus.

**GUCY2C Hormones Occupy an Insulated Transcriptional Domain Regulated by Wnt Signaling**

GUCA2A and GUCA2B are convergently transcribed, with their promoters separated by 21 kb in mice and 11 kb in humans. ChIP-seq suggests that these genes occupy a chromatin domain bounded by CTCF sites, an architecture that is conserved across species (Figure 4A and B). Between these CTCF sites, DNase hypersensitive sites sequencing (DNase-seq) of normal human colon identified several sites of DNase hypersensitivity, hallmarks of transcriptional activity (Figure 4C). Two of these sites correspond to the CTCF sites, 2 correspond to the gene promoters, and the remaining represent putative enhancers, a structure confirmed by additional DNase-seq (Figure 4D) and formaldehyde-assisted identification of regulatory elements sequencing (Figure 4E) analyses. These putative enhancers correspond to regions with the greatest transcription factor occupancy, as defined by ChIP-seq (Figure 4F). Regions upstream of both genes also harbor the enhancer-associated epigenetic marks H3K27ac (Figure 4G) and H3K4me1 (Figure 4H). Greater H3K27ac and H3K4me1 enrichment was observed upstream of GUCA2A than GUCA2B, consistent with greater expression of GUCA2A mRNA in the colon. Importantly, these marks were observed in normal human colon, but not colon cancer tissue (Figure 4G and H), suggesting enhancer inactivation consistent with transcriptional silencing during transformation.

This locus architecture, in which DNase, H3K27ac, and H3K4me1 enrichment terminate at the CTCF sites between GUCA2A and GUCA2B and their adjacent genes, FOXJ3 and HIVEP3, respectively, is consistent with a role for CTCF in transcriptional insulation. RNA-seq in the 4 cell lines revealed that GUCA2A and GUCA2B were among the most upregulated genes upon silencing Wnt signaling, while FOXJ3 and HIVEP3 were not significantly changed (Figure 5A). Quantitative real-time polymerase chain reaction (qRTPCR) analysis in all 4 cell lines confirmed that silencing APC-β-catenin signaling did not alter FOXJ3 or HIVEP3 expression (DLD1 cells shown in Figure 5B and C). These results recapitulate those from human normal colon and tumor samples in TCGA, in which HIVEP3 and FOXJ3 expression does not correlate with GUCA2A (Figure 5D and E). In contrast, expression of GUCA2A and GUCA2B directly correlate, and both are lost in colon tumors (Figure 5F). Indeed, GUCA2B is the gene most correlated with GUCA2A in colon tumors in TCGA, and single cell transcriptomics reveal that these transcripts are coexpressed in normal colon epithelial cells as well (Figure 5G and H). Thus, GUCA2A and GUCA2B are transcriptionally coregulated within an insulated domain sensitive to APC-β-catenin-TCF signaling.

**Regulation of the GUCA2A-GUCA2B Locus by Wnt Signaling Is Mediated by an Upstream RNA Polymerase II–Rich Super-Enhancer**

There are 9 (#1–#9) DNase hypersensitive sites within the GUCA2A and GUCA2B insulated domain representing putative regulatory DNA (Figure 6A). ChIP-seq in DLD1(DNTCF) cells revealed enrichment of RNA polymerase II (Pol II) in sites #5–#8, extending from the GUCA2A promoter to 6 kb upstream (Figure 6B). In contrast, the GUCA2B 5′ upstream region was devoid of Pol II enrichment.
This differential Pol II enrichment is consistent with greater GUCA2A, compared with GUCA2B, transcript expression in this cell line (Figure 6D). While silencing β-catenin-TCF signaling induced GUCY2C hormone expression and Pol II loss at Wnt target genes such as LEF1 and ASCL2 (Supplementary Table 1), Pol II recruitment to the GUCA2A locus was unaffected, consistent with poised transcriptional machinery independent of Wnt status. Additionally,
H3K27ac was enriched at sites #5–#6, and increased 1.3-fold ($P = .008$) after inducing DNTCF expression, consistent with transcriptional activation of this enhancer region (Figure 6C). The existence of a discrete $\beta$-catenin-TCF-sensitive element within this enhancer region was tested by cloning DNA fragments encompassing individual sites #1–#9 into luciferase reporters (Figure 6E). Unexpectedly, although DNTCF silenced luciferase expression in cells expressing the TOPflash TCF-containing reporter, it did not activate expression in cells expressing any of the individual GUCA2A DNase site reporters (Figure 6F). In contrast, DNTCF induced the coordinated expression of GUCA2A and GUCA2B mRNA, as well as luciferase driven by a region extending from the GUCA2A 5' UTR (+15) to beyond the CTCF site (−10,000; Figure 6G and H). Truncation analysis revealed a region between −1000 and −6000 that conferred enrichment in sites #5–#6.

**Figure 3.** $\beta$-catenin-TCF signaling regulates GUCA2A nuclear transcription. (A, B) Luciferase constructs driven by a constitutive (SV40) promoter and containing the guanylin 3' UTR or no 3' UTR, were expressed in (A) DLD1(DNTCF) or (B) LS174T(DNTCF) cells. Luciferase was quantified with (+) or without (−) 1 μg/mL DOX for 24 hours in DLD1 cells or 48 hours in LS174T cells. (C, D) Expression constructs containing the entire human GUCA2A gene from 5' to 3' UTR, or the entire murine Guca2a gene under the control of a constitutive (ROSA) promoter were expressed in (C) DLD1(DNTCF) or (D) LS174T(DNTCF) cells. mRNA was quantified with (+) or without (−) 1 μg/mL DOX for 24 hours in DLD1 cells, or 48 hours in LS174T cells. (E) DLD1(DNTCF) cells were treated with 1 μg/mL DOX for 24 hours and (F) HT29(APC) cells were treated with 300 μM zinc for 24 hours. GUCA2A mRNA was quantified in whole cell (W), cytoplasmic (C), or nuclear (N) fractions, with fractionation confirmed by GAPDH (cytoplasmic) or histone H3 (nuclear) protein. (G, H) GUCA2A preRNA was quantified in the (G) DLD1 and (H) HT29 whole cell fractions. (A–H) Data points represent the average of 3 wells of cells from a single experiment, with the mean of 2–4 independent experiments indicated. Significance was determined by (A, B, E, F) 2-way analysis of variance or (C, D, G, H) Student’s t test with matched analysis for independent experiments on log2-transformed results. No significance was identified between any group in individual or combined experiments in A–D. Data are presented relative to noninduced cells. *$P < .05$; **$P < .01$; ***$P < .001$. 

**Figure 2.** (See previous page). The GUCY2C ligands are among the most sensitive targets of $\beta$-catenin/TCF signaling. HT29(APC) cells were treated with 300 μM zinc for 24 hours, LS174T(sh$\beta$-catenin) cells were treated with 1 μg/mL DOX for 72 hours, DLD1(DNTCF) cells were treated with 1 μg/mL DOX for 24 hours, and LS174T(DNTCF) cells were treated with 1 μg/mL DOX for 48 hours. RNA-seq and differential gene expression analysis was performed in cells treated with or without the respective inducing agent. Unbiased gene set enrichment analysis on each cell line compared the ranked log fold change of 17,229 genes to the 50 hallmark gene sets maintained by the Molecular Signatures Database (MSigDB). (A) The Wnt-$\beta$-catenin signaling gene set enrichment plot for each cell line, with the corresponding normalized enrichment score and false discovery rate (FDR) $q$ value. (B, C) The top 10 upregulated and downregulated gene sets for DLD1(DNTCF) cells, ranked by normalized enrichment score. Color indicates FDR $q$ value, and size of the data points indicates the number of genes in each gene set. (D) Differential gene expression in the 4 cell lines by RNA-seq reveals 782 upregulated (red) and 507 downregulated (blue) genes upon silencing Wnt signaling. (E) Heatmap of the 1289 differentially expressed genes ranked by log2 fold change. GUCA2A and GUCA2B are the 7th and 12th most upregulated transcripts. RNA-seq results represent the average of 3 replicates.
A 2683 bp Element Mediates β-Catenin-TCF-Sensitive Control of the GUCY2C Hormone Locus

Mutational analysis to define the β-catenin-TCF-sensitive element revealed that deletion of sites #6 (1000-2000) or #7 (3000-4238) only modestly decreased, while a 2683-bp deletion spanning both sites (1000-3683) abolished, DNTCF-regulated luciferase expression (Figure 7A). Sequential truncation of either edge of this region partially reduced, but did not eliminate, DNTCF sensitivity (Figure 7B), demonstrating the contribution of the entire region to maximum β-catenin-TCF regulation. Expression of a luciferase construct containing the 2683 bp region with a minimal GUCYA2A promoter conferred full β-catenin-TCF-sensitivity (Figure 7C). Moreover, while individual DNase sites failed to activate a constitutive (SV40) promoter (Figure 6F), the 2683 bp region was sufficient (Figure 7D), identifying this region as a bona fide β-catenin-TCF-sensitive enhancer. Finally, the 2683-bp enhancer conferred β-catenin-TCF-sensitivity to a minimal GUCY2B promoter (Figure 7E), supporting its role in coregulating GUCYA2A and GUCY2B expression.

To validate the role of this enhancer in mediating β-catenin-TCF-sensitivity to GUCYA2A and GUCY2B in the context of the chromosomal locus, DNase sites were eliminated with CRISPR-Cas9. Deletion of the GUCY2A promoter abolished GUCYA2A mRNA and protein expression, and deletion of single DNase sites only incompletely reduced DNTCF sensitivity (Figure 8A and B). In contrast, deletion of both sites almost completely abolished DNTCF sensitivity. Moreover, GUCY2B mRNA expression paralleled that of GUCYA2A, revealing that GUCYA2A promoter-proximal and
Figure 5. Transcriptional coregulation of GUCA2A and GUCA2B within an insulated domain sensitive to Wnt signaling. (A) RNA-seq volcano plots illustrate the fold change of 17,229 genes upon silencing Wnt signaling in the 4 inducible cell lines, with GUCA2A, GUCA2B, HIVEP3, and FOXJ3 indicated. Each gene is represented as a data point, dotted lines denote 2-fold change, and significance is indicated in red ($P > .01$) or blue ($P < .01$). (B–C) qRTPCR of (B) HIVEP3 and (C) FOXJ3 in DLD1(DNTCF) cells treated with (+) or without (−) 1 μg/mL DOX for 24 hours. (D–F) GUCA2A mRNA compared with that of (D) HIVEP3, (E) FOXJ3, or (F) GUCA2B, quantified by RNA-seq from human colorectal tumor (n = 380; black) and normal (n = 51; blue) tissue from TCGA (COAD/READ datasets). Significance was determined by 2-tailed Spearman rank correlation ($r_s$), and linear regression ($R^2$) is indicated by the red line. (G, H) Single-cell gene expression data retrieved from The Human Protein Atlas reveals coexpression of (G) GUCA2A and (H) GUCA2B in normal colon epithelial cell clusters (insets show corresponding colon UMAP plots of cell clusters). (B, C) Data points represent the average of 3 wells of cells from a single experiment, with the mean of 3 independent experiments indicated. Significance was determined by Student’s t test with matched analysis for independent experiments on log2-transformed results. Data are presented relative to noninduced cells. *$P < .05$; ****$P < .0001$.

upstream super-enhancer elements synchronize the expression of both genes (Figure 8C). Collectively, these data support a paradigm involving a 2683 bp Pol II-rich locus control region (LCR) that confers β-catenin-TCF-sensitivity to the GUCA2A and GUCA2B promoters (Figure 8D). This sensitivity is conveyed indirectly by β-catenin/TCF. ChIP analysis did not identify TCF enrichment in the GUCA2A LCR (Figure 8E), indicating that TCF does not directly interact with the GUCA2A LCR to regulate transcription. Furthermore, elimination of TCF expression with shRNA restored GUCA2A mRNA expression (Figure 8F), mimicking the effect of DNTCF expression and indicating that TCF is not required for GUCA2A expression.

Epigenetic Modifiers Targeted to the GUCA2A LCR Mediate Wnt-Independent Control of Gene Expression

For unbiased validation of the results with luciferase and Cas9 KO, monoclonal DLD1(DNTCF) cells were generated
stably expressing dCas9 fused to the SID4X and KRAB transcriptional repressor domains (dCas9.KRAB),\textsuperscript{42} or the VP64 transcriptional activation domain (dCas9.VP64).\textsuperscript{43} The ability to target dCas9.KRAB to the GUCA2A promoter was confirmed by ChIP, and knockdown of mRNA expression with guide RNA (gRNAs) targeting multiple genes was confirmed by qRTPCR (data not shown). Fifty-four gRNAs spanning the GUCA2A gene body, promoter, and enhancer regions were expressed individually in DLD1(DNTCF).dCas9.KRAB cells with DNTCF to identify sites controlling GUCA2A expression (Figure 9A). Indeed, individual gRNAs targeting the gene promoter, Dnase site #6, or Dnase site #7 repressed DNTCF-sensitive GUCA2A expression, supporting previous observations. Further, gRNA pools targeting the GUCA2A promoter or upstream enhancers also inhibited GUCA2A expression (Figure 9B). Importantly, pools targeting the enhancer (Dnase site #6) also repressed DNTCF-sensitive GUCA2B expression, consistent with a role for the LCR in gene coregulation (Figure 9C).

Finally, overcoming GUCA2A transcript loss in the context of mutant β-catenin-TCF-signaling represents a therapeutic goal to reconstitute hormone expression. As
RNA-seq FPKM quantification of gene expression in the context of mutant GUCY2C hormone loss represents a well-recognized point of potential therapeutic intervention, studies here add reconstitution of gene expression to GUCY2C agonist replacement and cGMP-dependent silencing switch controlling the events between Pol II transcription initiation. To our knowledge, this represents the first example of a β-catenin/TCF-dependent transcriptional silencing switch controlling the events between Pol II recruitment to an LCR and activation of transcription at a gene promoter. Interestingly, in contrast to results here and in normal colon, H3K27ac is depleted at the GUCY2A LCR in colorectal cancer (Figure 4). These observations suggest that at least 2 mechanisms contribute to GUCY2C hormone loss in tumorigenesis. Acutely, APC-β-catenin-TCF signaling dynamically regulates hormone mRNA transcription at the LCR, a mechanism likely operating in normal colon to elevate agents as strategies to transform a disease of reversible condition that restores cGMP-elevating agents as strategies to transform a disease of reversible condition that restores cGMP

discussion

The tumor suppressive axis controlled by the intestinal receptor GUCY2C and its paracrine ligands GUCA2A and GUCA2B is among the first pathways dysregulated in tumorigenesis. Loss of GUCY2C hormones is an early feature of tumorigenesis in mice and humans, occurring in dysplastic crypts in temporal proximity to APC loss.23 Here, we reveal that the GUCY2C hormones share an insulated genomic locus containing an LCR that mediates gene silencing by APC-β-catenin-TCF signaling. This mechanistic insights support the pathophysiologic hypothesis that silencing of GUCY2C hormone gene transcription is a key step in tumorigenesis, overcoming tumor suppressive signaling by the GUCY2C-cGMP axis.4,5,13 While hormone loss represents a well-recognized point of potential therapeutic intervention, studies here add reconstitution of gene expression to GUCY2C agonist replacement and cGMP-elevating agents as strategies to transform a disease of irreversible genetic mutations in APC or β-catenin into a reversible condition that restores cGMP signaling.4,5,11-13,18,19,21,28

To explore mechanisms linking mutant APC-β-catenin-TCF-signaling to GUCY2C ligand loss, we assembled human colon cancer cells engineered to define intestinal Wnt target genes.29-32,44,45 These cells express inducible transgenes for wild-type APC, β-catenin shRNA, or dominant negative TCF7L2, which interrupt Wnt signaling at 3 discrete steps. Silencing Wnt signaling at any step reconstituted GUCA2A and GUCA2B expression, consistent with ligand loss in tumors in mice and humans.16,21,22,25 Interestingly, these genes have not been identified previously as Wnt targets in these models,32,44-46 likely reflecting the insensitivity of array-based approaches to genes with low transcript abundance in the context of Wnt signaling. Here, RNA-seq, applied to the 4 cell models for the first time, identified a common set of 1289 differentially expressed genes in which GUCA2A and GUCA2B mRNA were among the most sensitive readouts of Wnt signaling.

RNA-seq identifies changes in steady-state mRNA levels, reflecting either (or both) transcriptional (synthesis) and posttranscriptional (stability) regulation. With respect to GUCY2C hormones, silencing Wnt signaling increased nascent transcripts,23 transcript levels in the nucleus, and preRNA transcripts. Further, APC-β-catenin-TCF sensitivity was recapitulated with luciferase reporters containing the LCR. These observations are consistent with primarily transcriptional, rather than posttranscriptional, regulation of GUCY2C hormones by Wnt signaling. In that context, stable enrichment of H3K27ac and Pol II in the LCR, seen in both the presence and the absence of Wnt signaling, suggest dynamic regulation of hormone mRNA synthesis beyond the histone modification and Pol II recruitment steps preceding transcription initiation. To our knowledge, this represents the first example of a β-catenin/TCF-dependent transcriptional silencing switch controlling the events between Pol II recruitment to an LCR and activation of transcription at a gene promoter. Interestingly, in contrast to results here and in normal colon, H3K27ac is depleted at the GUCY2A LCR in colorectal cancer (Figure 4). These observations suggest that at least 2 mechanisms contribute to GUCY2C hormone loss in tumorigenesis. Acutely, APC-β-catenin-TCF signaling dynamically regulates hormone mRNA transcription at the LCR, a mechanism likely operating in normal colon to...
establish the ascending gradient of hormone expression along the crypt-surface axis.14 Chronically, during the evolution of a tumor, removal of activating chromatin marks and closure of chromatin domains produces LCR atrophy, reflected by H3K27ac ChIP-seq.38 Elucidating these phases of regulation will inform therapeutic strategies focused on reconstitution of gene expression in existing tumors.

While mechanisms activating gene expression by Wnt signaling are defined, those silencing gene expression remain incompletely understood. In Wnt-activated genes, TCF at consensus DNA Wnt recognition elements (WREs) in target gene promoters or enhancers binds β-catenin, which recruits chromatin remodeling enzymes and coactivators required for transcription.2 Paradoxically, in Drosophila,
Figure 8. Cas9 deletion confirms a super-enhancer necessary for β-catenin-TCF-sensitive locus control. DLD1(DNTCF) cells expressing Cas9 and untargeted (clone #1) or GUCA2A-locus-targeted gRNAs harbor biallelic deletions encompassing the GUCA2A promoter (clone #2), DNase site #6 (clone #3), DNase site #7 (clone #4), or both DNase sites (clone #5). Expression of (A) GUCA2A protein, (B) GUCA2A mRNA, and (C) GUCA2B mRNA was quantified with (+) or without (−) 1 μg/mL DOX for 24 hours. (D) Hypothetical model of GUCA2A and GUCA2B promoter positioning relative to a Pol II-rich super-enhancer region upstream of GUCA2A, consisting of multiple DNase sites. (E) ChIP-PCR in HT29(APC) cells treated with (+) or without (−) 300 μM zinc for 24 hours reveals enrichment of TCF at the promoter of a Wnt target gene, SP5, with a TCF-specific antibody, but not with control IgG. In contrast, TCF was not detected at sites within 6kb of the GUCA2A TSS: DNase site #5 (GUCA2A promoter), site #6, site #7, or site #8. (F) GUCA2A mRNA expression in HT29(APC) cells stably expressing an untargeted (Ctr) or TCF-targeted shRNA, and treated with (+) or without (−) 300 μM zinc for 24 hours. GUCA2A mRNA expression is retained despite TCF knockdown, illustrated by Western blot. (A–C) Bars represent the average of (A) 2 or (B, C) 3 independent experiments ± SEM, and data are presented relative to noninduced cells. Significance was determined by 2-way analysis of variance with matched analysis for independent experiments on log2-transformed results. (E) Bars represent the mean ± SD of 3 IPs, and data are presented relative to input DNA. Significance was determined by 2-way analysis of variance. (F) Data points represent the average of 3 wells of cells from a single experiment, with the mean of 2 independent experiments indicated. Significance was determined by 2-way analysis of variance with matched analysis for independent experiments on log2-transformed results. Data are presented relative to noninduced cells receiving control shRNA. *P < .05; **P < .01; ***P < .001; ****P < .0001.
Wnt-repressed genes also include TCF binding sites, which differ from canonical WREs and are required for repression. In that context, there are no canonical WREs in the GUCA2A LCR (Supplementary File 1). Further, TCF was not enriched in the GUCA2A LCR by ChIP analysis (Figure 8E), ruling out binding to noncanonical WREs. Moreover, shRNA inhibition of TCF expression restored GUCA2A mRNA production (mimicking DNTCF), eliminating the possibility that

**Figure 9. Targeting the GUCA2A LCR for Wnt-independent gene control.** (A–C) CRISPRi in DLD1(DNTCF) cells stably expressing SID4X.dCas9.KRAB (dCas9.KRAB). (A) GUCA2A mRNA was quantified after stable transduction with 54 individual gRNAs targeting the GUCA2A locus and induction of DNTCF with 1 μg/mL DOX for 24 hours. gRNAs producing significant (P < .05) inhibition of GUCA2A mRNA relative to that of untargeted gRNAs are indicated in red. (B) GUCA2A mRNA and (C) GUCA2B mRNA were quantified after stable transduction with 3–5 gRNAs targeting nothing (sham), the GUCA2A promoter, DNase site #6, or DNase site #7, followed by treatment with 1 μg/mL DOX for 24 hours. (D–F) CRISPRa in DLD1(DNTCF) cells stably expressing dCas9.VP64. (D) GUCA2A mRNA was quantified after stable transduction with 54 gRNAs, as in panel A, in the absence of DOX. (E) GUCA2A mRNA and (F) GUCA2B mRNA were quantified after stable transduction with 3–5 gRNAs, as in panels B and C, in the absence of DOX. Data points represent the average of (A) 2 independent experiments ± SEM or (B) 1 experiment ± SEM. (B, C, E, F) Data points represent the average of 3 wells of cells from a single experiment, with the mean of 3 independent experiments indicated. Significance was determined by 1-way analysis of variance with matched analysis for independent experiments on log2-transformed results. Data are presented relative to cells expressing untargeted gRNAs. **P < .01; ***P < .001; ****P < .0001.
TCF directly regulates GUCY2C hormone transcription (Figure 8F). Thus, Wnt regulation of GUCY2C hormone mRNA synthesis appears indirect, wherein β-catenin-TCF controls expression of the transcriptional machinery required for hormone regulation. The 1289 Wnt target gene set revealed by RNA-seq includes 95 transcription factors (Supplementary Table 2), many of which are known intermediates of Wnt (ie, MYC, SP5, and ASCL2). Ongoing analyses are exploring interactions between this Wnt transcriptional network and the GUCYA2 LCR.

This is the first report to reveal transcriptional coregulation of GUCY2C hormones and its sensitivity to Wnt repression mediated by cis elements within an insulated genomic locus. This regulation is mediated by an LCR containing 4 DNase hypersensitive sites. Deletion of individual sites from luciferase constructs or from the genomic locus did not eliminate gene regulation, suggesting functional redundancy that might reflect evolutionary pressure to retain GUCY2C hormone expression in healthy intestine. Indeed, loss of GUCY2C hormone expression has been implicated in the pathophysiology of colorectal cancer, inflammatory bowel disease, irritable bowel disease, and obesity.4,5,13 DNase sites clustered in LCRs form phase-separated transcriptional condensates rich in transcriptional machinery.48 The region of Pol II and H3K27ac enriched upstream of GUCYA2 is consistent with this model. Further, LCRs integrate multiple gene control elements.49 Here, site #9 associates with CTCF and borders H3K27ac, supporting its role as an insulator, while sites #6 and #7 together form a 2683-bp Wnt-sensitive region. This 2683-bp region is a bona fide enhancer, with the ability to confer Wnt sensitivity to luciferase reporters driven by the GUCYA2A, GUCYA2B, or SV40 promoters. Further, deleting the enhancer with Cas9, or targeting it with the repressor dCas9.KRAB, eliminates Wnt sensitivity from both genes. Together, these observations are consistent with a model in which the LCR forms a transcriptional condensate which concentrates the machinery mediating Wnt-sensitive coregulation of GUCYA2A and GUCYA2B expression (Figure 8D). While speculative, ongoing chromosome conformation capture studies will clarify this working model.

Enhancer DNA sequences share common features, including sensitivity to DNase, enrichment with transcriptional machinery (eg, Pol II), and histone modifications (eg, H3K27ac). Here, putative enhancers with these characteristics were identified from public databases, and enhancer activity was validated with traditional reporter plasmids. Complementary analytic strategies included a candidate approach, incorporating individual DNase sites into luciferase reporters, and an unbiased approach, incorporating serial truncations of the 10-kb sequence spanning the GUCYA2 TSS and nearest CTCF site (Figure 6). While the candidate approach was not yielding, and no single DNase site regulated luciferase expression, the unbiased approach revealed regulatory activity corresponding to sites #6 and #7, underscoring a pitfall of overly reductionist approaches to examining DNA elements. These considerations highlight the importance of applying comprehensive strategies to capture regulatory elements that function as small units independent of native chromatin, as well as those that function as large interdependent units, which can include super-enhancers and LCRs capable of forming transcriptional condensates.

Complementary CRISPR approaches were used to examine Wnt sensitivity of the GUCYA2 A’ super-enhancer, including CRISPR/Cas9 KO, CRISPR inhibition (CRISPRi), and CRISPR activation (CRISPRa). In that context, there was concordance between luciferase reporter and CRISPR KO analyses. However, luciferase analyses only revealed regulation of reporter gene expression, while CRISPR KO analyses revealed regulation of both GUCYA2A and GUCYA2B mRNA expression, identifying the super-enhancer as an LCR. Pitfalls of CRISPR KO include unanticipated alterations to chromatin architecture and the creation of new DNA motifs at recombination sites. CRISPRi and CRISPRa tiling screens have emerged as a less disruptive tool to interrogate enhancers through epigenetic modulation. Thus, to validate CRISPR KO analyses, CRISPRi and CRISPRa were targeted to the GUCYA2 locus to identify sites contributing to hormone expression. CRISPRi targeting site #6 and site #7 repressed GUCYA2A and GUCYA2B expression, confirming LCR activity. Conversely, CRISPRa targeting site #6 reconstituted GUCYA2A mRNA expression. Interestingly, CRISPRa targeting the LCR failed to reconstitute GUCYA2B expression, likely reflecting the relatively weak transactivation capacity of the VP64 domain,50 and barriers imposed by active repressive Wnt signaling on interactions between the LCR and GUCYA2B promoter. Notably, this is the first example of GUCY2C hormone mRNA reconstitution in the context of repressive mutant APC-β-catenin-TCF signaling. Indeed, LCR-targeted CRISPRa produced greater reconstitution than CRISPRa targeted to the GUCYA2A promoter.

These observations provide a compelling rationale for pathologic enhancer identification and targeted epigenome modulation to overcome active oncogenic signaling, and proof-of-principle for reconstitution of GUCY2C ligand expression. In the context of colorectal cancer, β-catenin-TCF-mediated decommissioning of the GUCY2C hormone LCR represents a previously unrecognized pathophysiologic step in tumorigenesis. This mechanistic insight suggests several future pathways for clinical translation. Ongoing efforts to identify the intermediate factors conveying the β-catenin-TCF signal to the LCR should reveal novel targets for small molecule inhibitors, which could be employed for hormone reconstitution to oppose tumorigenesis. We also envision future applications for direct LCR targeting with CRISPRa, enabling gene activation without a priori knowledge of the transcription factor network acting on the locus. Indeed, CRISPRa has recently been employed to increase in situ gene expression in tumor stroma to augment antitumor immune responses.51 Given the sustained expression of the orphaned GUCY2C receptor on transformed cells, it is tempting to speculate that intratumoral reactivation of GUCY2C ligand expression with CRISPRa could have a therapeutic role. Ultimately, the present study reveals a mechanistic basis for GUCY2C hormone loss in colorectal cancer, creating a unique opportunity to reverse GUCY2C...
silencing and oppose tumorigenesis in the context of mutant Wnt signaling.

**Materials And Methods**

**Cell and Organoid Culture Reagents**

McCoy's 5A (#10050CV), Dulbecco's Modified Eagle Medium (DMEM) (#10013CV), DMEM/F12 (#10092CV), Matrigel (#354230), and cell recovery solution (#354253) were from Corning (Corning, NY). Ca²⁺/Mg²⁺-free Hank's Balanced Salt Solution (HBSS) (#14175095), GlutaMax supplement (#35050061), penicillin-streptomycin (#15140122), HEPES (#15630080), HyClone fetal bovine serum (#SH3007103), hygromycin B (#10687010), zeocin (#R25005), blasticidin hydrochloride (#A1113903), geneticin/G418 sulfate (#10131027), doxycycline hydrochloride (BP2653-1), puromycin (#A1113803), N-2 supplement (#CRL-3276) was from American Type Culture Collection (Manassas, VA), Mouse recombinant epithelial growth factor (EGF) (#315-09) and Noggin (#250-38) were from PeproTech (Rocky Hill, NJ). Human recombinant R-spondin-1 (#4645-5-S-025/CF) was from R&D Systems (Minneapolis, MN). CHIR99021 (#129830-38-2) and Y-27632 (#252917-06-9) were from StemCell Technologies (Vancouver, Canada).

**Mouse and Organoid Studies**

Animal protocols were approved by the Thomas Jefferson University Institutional Animal Care and Use Committee. Colon organoids from wild-type C57B/6(J) mice (6–12 weeks old) were cultured according a protocol modified from Fan et al. Mouse colon was dissected, flushed with cold HBSS, everted onto a gauze needle, and vortexed with six 5-second pulses in cold HBSS supplemented with 0.5% penicillin-streptomycin. The colon was incubated in HBSS supplemented with 20 mM EDTA at 37°C for 30 minutes, then vortexed in cold HBSS, with eight 5-second pulses to release crypts. Crypts were centrifuged at 125 g for 3 minutes and resuspended in cold complete medium (DMEM/F12 supplemented with 1% Glutamax, 1% penicillin-streptomycin, and 1% HEPES). A total of 500 crypts were plated in 50 μL of Matrigel in 24-well plates, incubated for 15 minutes at 37°C, and then overlaid with 500 μL of complete medium supplemented with Wnt3a, R-spondin-, and Noggin-conditioned medium (L-WRN, 1:1 dilution), 1× N-2, 1× B-27, 100 μM N-acetyl-l-cysteine, 50 ng/mL EGF, 500 nM A83-01, 10 μM SB202190, 10 mM nicotinamide, 10 μM CHIR99021, and 10 μM Y-27632. Medium was replaced every 3 days, and organoids were passaged weekly. For organoid differentiation studies, medium was replaced with differentiation medium (complete medium supplemented with 1 μg/mL R-spondin-1, 100 ng/mL Noggin, 1× N-2, 1× B-27, 100 μM N-acetyl-l-cysteine, 50 ng/mL EGF, 500 nM A83-01, and 10 μM Y-27632). Differentiation medium was replaced daily for 3 days. To harvest, Matrigel containing organoids was dissolved in cell recovery solution for 1 hour at 4°C. Organoid pellets were flash frozen and stored at −80°C for biochemical analyses.

**Cell Lines**

DLD1 and LS174T cells stably expressing Tet-inducible dominant negative TCF7L2 (DNTCF), and LS174T cells stably expressing Tet-inducible shRNA to β-catenin were provided by Dr. H. Clevers in November 2013. Cells were cultured in DMEM supplemented with 10% fetal bovine serum (FBS), 500 μg/mL zeocin, and 10 μg/mL blasticidin. Cells were induced with 1 μg/mL DOX for 24 hours (DLD1-DNTCF), 48 hours (LS174T-DNTCF), or 72 hours (LS174T-shβ-catenin) for mRNA analyses, or an additional 24 hours for protein analyses. HT29 cells stably expressing zinc-inducible wild-type APC were provided by Dr. B. Vogelstein in September 2013. HT29 cells were cultured in McCoy’s-5A supplemented with 10% FBS and 600 μg/mL hygromycin. Cells were induced with 300 μM zinc chloride for 24 hours for mRNA, or 48 hours for protein analyses. Conditional cell lines were authenticated at each use by testing their genetic inducibility, and all cell lines were confirmed to be free of mycoplasma semiannually.

**Immunofluorescence**

Following deparaffinization and rehydration in a xylene-ethanol-water gradient, 4μm sections of wild-type mouse C57BL/6(J)) intestines underwent antigen retrieval by heating at 100°C for 15 minutes in a pressure cooker in pH 9.0 DAKO antigen retrieval buffer (Agilent Technologies, Santa Clara, CA; S236784-2). The monoclonal antibody to GUCY2C (#2504, 1:1000), APC (#2504, 1:1000), and the anti-sera to GUCA2A (S236784-2) was kindly provided by Dr. M. Goy. GUCY2C and GUCA2A were detected by tyramine signal amplification, as described previously. Images were captured using an EVOS FL auto cell imaging system (Thermo Fisher Scientific).

**Immunoblots**

Nuclear and cytoplasmic cell fractions were separated with the Nuclei EZ Prep kit (Sigma-Aldrich; #NUC-101), per kit instructions. Protein was extracted from nuclear, cytoplasmic, or total cell lysates using M-Per (Thermo Fisher Scientific; #78501) supplemented with protease inhibitors (Thermo Fisher Scientific; #a32955). Lysates were prepared in 4X LDS Sample buffer (Invitrogen, Waltham, MA; #NP0007) supplemented with 2.75 mM 2-Mercaptoethanol (Thermo Fisher Scientific; #21985023), boiled at 100°C for 8 minutes, loaded onto 4%–12% Bis-Tris gels (20 μg/ lane; Thermo Fisher Scientific; #NP0336), and transferred to nitrocellulose membranes with an iBlot station (Thermo Fisher Scientific; #IB301031). Blots were blocked for 1 hour in phosphate-buffered saline (PBS) containing 0.1% Tween 20 and 10% milk, then incubated overnight at 4°C with primary antibodies at the indicated dilutions targeting GAPDH (#2118, 1:5000), APC (#2504, 1:1000), β-catenin (#8480,
with a Nanodrop 1000 (Thermo Fisher Scientific) according to kit instructions. RNA concentration and purity were measured with a Nanodrop 1000 (Thermo Fisher Scientific), and 200 ng of RNA was reverse transcribed to complementary DNA (cDNA) using the TaqMan Reverse Transcription kit according to kit directions (Thermo Fisher Scientific; #N8080234). Transcripts were quantified by densitometry normalized to that of GAPDH using FIJI software (ImageJ version 2.0.0-rc-49).

**RNA Analysis**

Samples were lysed in RLT buffer supplemented with 550 μM 2-mercaptoethanol, genomic DNA was removed, and RNA was purified on spin columns using the RNeasy Plus kit (Qiagen, Hilden, Germany; #74134) according to kit directions. RNA was amplified by qRTPCR using TopHat (v2.0.12) and reads mapping to each gene were counted with HTSeq (v0.6.1). Genes with counts per million >0.5 in at least 3 samples were included in subsequent analyses (17,728 genes). Differential gene expression (adjusted P value <.05) was determined between cells treated with or without respective inducing agents, using the limma (v3.40.6) and edgeR (v3.26.8) packages in R (R Foundation for Statistical Computing, Vienna, Austria, version 1.2.1335). Gene set enrichment analysis was performed for each cell line, comparing the 17,728 genes (ranked by log-fold change) to the H:Hallmark Gene Sets in MSigDB (v.7.2), and results were plotted using the ggplot2 (v3.3.0) package in R.

**Plasmids and Cloning**

For GUCA2A luciferase reporter assays, DNA fragments were amplified by PCR from the RP11-799L22 BAC (BAC-PAC Genomics), which contains the human GUCA2A/GUCA2B locus. Control inserts containing TCF or mutant TCF binding sites were amplified from the TOPflash and FOPflash luciferase reporter plasmids (Sigma-Aldrich; #21-170, #21-169). The PGL3-Basic and PGL3-Promoter (Promega, Madison, WI; #E1751, #E1761) luciferase reporter plasmids were linearized with Ncol (#R3193)/Xhol (#R0146) or ScaI (#R0156)/Xhol, respectively, from New England Biolabs. Gibson assembly of inserts and linearized plasmids was performed with the Geneart Seamless Cloning and Assembly kit (Thermo Fisher Scientific; #A1328B) per package instructions. CRISPR-Cas9 plasmids were obtained from Addgene (Watertown, MA): SID4X.dCas9.KRAB was a gift from Jason Gertz (#106399), pCDNA-dCas9-VP64 was a gift from Charles Gersbach (#47107), and Lentigui-Puro (52963) was a gift from Feng Zhang. The active Cas9 plasmid, pLV5-Cas9-Neo, was from Sigma (#CAS9NEO). Protopspacer-adjacent motifs in the GUCA2A locus were identified with the online tool, CHOPCHOP, and gRNAs without off-target binding sites were selected. Forward and reverse gRNA oligos were designed, annealed, and ligated into the Esp3I (New England Biolabs; #R0734)-digested Lentigui-Puro vector according to directions provided by the Zhang lab on Addgene. Plasmid DNA was prepared from overnight E.coli cultures using HiPure Plasmid Midiprep kits (Thermo Fisher Scientific; #K210005). Plasmids were validated by Sanger sequencing by Eurofins Genomics. All cloning primers and gRNA sequences are listed in Supplement Table 1.
Luciferase Reporter Assay

Cells were seeded in white 96-well plates at 20,000 cells/well. The following day, cells were cotransfected with 250 ng of Firefly luciferase plasmid and 50 ng of Renilla luciferase plasmid (pRLTK; Promega; #E2241), using Fugene HD (Promega; #E2311), according to kit instructions. DOX was added to induce cells for 24 hours (DLD1) or 48 hours (LS174T). A total of 48 hours after transfection, cells were lysed with the Dual-Glo Luciferase Assay kit (Promega; #E2940), following kit instructions. Luminescence was read on a PolarStar Omega Plate Reader (BMG Labtech, Ortenberg, Germany). Firefly luciferase activity was normalized to Renilla luciferase activity in each well.

CRISPR Cell Line Generation

DLD1(DNTCF) cells were transfected with SID4X.dCas9.KRAB or pcDNA-dCas9-VP64 using Fugene HD (Promega; #E2311), or were transduced with pLv5-Cas9-Neo lentiviral supernatants. After 48 hours, cells were selected with 1mg/mL Geneticin (Thermo Fisher Scientific; #10131027) until death of nontransduced cells. Stable cells were passaged into 96-well plates by limiting dilution. Single clones were identified, expanded for 1 month, and probed for Cas9 protein expression. A single subclone was selected for subsequent experiments. For CRISPRi and CRISPRa experiments, DLD1(DNTCF).dCas9.KRAB (clone E9) or DLD1(DNTCF).dCas9.VP64 (clone A4) cells were transduced with lentiviral gRNA supernatants, incubated for 48 hours, and selected with 1 μg/mL Puromycin for 72 hours. Cells were induced with or without 1 μg/mL DOX for 24 hours and collected for mRNA analysis. For CRISPR KO experiments, DLD1(DNTCF).Cas9 (clone C1) cells were transduced with lentiviral gRNA pools targeting the GUCA2A DNase site #6, DNase site #7, both DNase sites, or nothing (scrambled gRNAs). After 48 hours, cells were selected with 1 μg/mL puromycin until death of nontransduced cells. Stable cells were passaged into 96-well plates by limiting dilution. Single colonies were identified, expanded for 1 month, genomic DNA was harvested, and target loci were amplified by PCR to identify deletions. A single subclone from each group was selected for subsequent experiments, including a control clone identified with a deletion encompassing the GUCA2A promoter and a portion of the 5′ coding region. No clone was initially identified with a deletion spanning both DNase sites. The clone harboring a deletion of DNase site #6 was retransduced with gRNAs spanning both DNase sites. Subclones were again isolated and expanded, and 1 clone was identified with a deletion spanning both sites.

Lentiviral Production and Transduction

Lentiviral packaging and transfer plasmids were from Addgene: pCMV-dR8.2-dvpr (#8455) and pCMV-VSV-g (#8454) were gifts from Bob Weinberg; 62 and pPGS-DNTCF4(deltaN41) was a gift from E. Fearon (#19284). 63 TCF and β-catenin shRNA transfer plasmids were obtained as bacterial glycerol stocks from Sigma (sequences listed in Supplementary Table 1). Two million HEK293T cells were plated in 10-cm dishes and cultured in DMEM supplemented with 10% FBS. The following day, cells were cotransfected with 3 μg VSV-g, 6 μg dr8.2-dvpr, and 6 μg of lentiviral transfer plasmids using the Profection Mammalian Transfection reagents (Promega; #E1200) according to the kit protocol. Media was replaced the following day. Viral supernatants were collected 48 hours after transfection and filtered through 0.45 μm syringe filters (Sigma-Aldrich; #SLHV033RS). Cancer cell lines were transduced with viral supernatants supplemented with 0.8 μg/mL Polybrene (Sigma-Aldrich; #TR-1003-G), incubated for 48 hours, then selected with 1 μg/mL Puromycin (Thermo Fisher Scientific; #A1113803). Cells were collected for analysis 72 hours after selection (upon death of nontransduced cells), or propagated to produce stable cell lines.

Chromatin Immunoprecipitation

ChIP was performed according to a protocol modified from Schmidt et al.64 Cells grown to confluence in 15-cm dishes were fixed with 1% formaldehyde (Thermo Fisher Scientific; #28906) for 10 minutes at room temperature, then quenched with 180 mM glycin (Thermo Fisher Scientific; #15527013). Cells were washed with ice-cold PBS, collected, resuspended in lysis buffer 1 (50 mM HEPES-KOH pH 7.5, 140 mM NaCl, 1 mM EDTA, 10% glycerol, 0.5% NP-40, 0.25% Triton X-100, and protease inhibitors), and rotated for 10 minutes at 4°C. Cells were pelleted, resuspended in lysis buffer 2 (10 mM Tris-HCl pH 8.0, 200 mM NaCl, 1 mM EDTA, 0.5 mM EGTA, and protease inhibitors), and rotated for 5 minutes at 4°C. Cells were pelleted, resuspended in lysis buffer 3 (10 mM Tris-HCl pH 8.0, 100 mM NaCl, 1 mM EDTA, 0.5 mM EGTA, 0.1% Na-deoxycholate, 0.5% N-lauroylsarcosine, and protease inhibitors), and sonicated to an average length of 1kb in a Q800R2 sonicator (QSonica, Newtown, CT) at 4°C. Sheared chromatin was frozen at –80°C. Protein G Dynabeads (Thermo Fisher Scientific; #100003D) were washed 3 times in blocking buffer (0.5% bovine serum albumin in PBS; Sigma-Aldrich; #BP1605). A total of 25 μL of bead slurry was incubated with 5 μg of ChIP antibody per IP, rotating overnight at 4°C. ChIP antibodies targeted TCF7L2 (Sigma-Aldrich; #17-10109), Cas9 (Active Motif, Carlsbad, CA; #61757), or were IgG control (Abcam, Cambridge, United Kingdom; #ab171870). The following morning, unbound antibody was washed from the beads, and antibody-coated beads were incubated with 25 μg of chromatin diluted in lysis buffer 3 supplemented with 1% Triton X-100 and protease inhibitors. A 2% sample of input chromatin was set aside at –20°C, and beads were rotated with chromatin at 4°C overnight. The following morning, beads were washed 3 times with wash buffer 1 (0.1% SDS, 1% Triton X-100, 10mM EDTA, 150mM NaCl, 20mM Tris HCl, pH 8.0) and once with wash buffer 2 (0.1% sodium dodecyl sulfate, 1% Triton X-100, 10 mM EDTA, 500 mM NaCl, 20 mM Tris HCl, pH 8.0). Beads were eluted for 30 minutes at 65°C with frequent vortexing in elution buffer (1% sodium dodecyl sulfate, 100 mM NaHCO3), and the supernatants were collected in fresh, low-adherence tubes (Eppendorf,
Hamburg, Germany; #022431021). Input and IP samples were incubated at 37°C for 30 minutes with 5 μg RNase A (Thermo Fisher Scientific; #AM2271), followed by reverse crosslinking for 5 hours at 65°C with 100 μg proteinase K. ChIP DNA was purified with MinElute PCR Purification columns (Qiagen; #28004) according to kit instructions.

**ChIP Sequencing**

ChIP-seq samples were prepared as previously described, but chromatin was sonicated to an average size of 250–300 bp. IPs were scaled up to 50 μL of bead slurry, 10 μg of ChIP antibody, and 100 μg of chromatin per IP. ChIP antibodies targeting H3K27ac (#ab4729) and RNA Pol II (#ab26721) were from Abcam. Library preparation, sequencing, and analysis was performed by the MetaOmics Shared Resource Facility of the Sidney Kimmel Cancer Center of Thomas Jefferson University. Briefly, ChIP-seq libraries were prepared using Swift Biosciences ACCEL-NGS 2S Plus DNA library kit following the manufacturer’s protocol. Single-end 75-bp sequencing was performed using a high-output flowcell on the Illumina NextSeq 500 at a depth of 40–55M reads per sample. Sequence read fragments were aligned to the GRCh38 human genome using the BWA-MEM aligner. Samtools was used to sort the resulting alignments and merge replicate .bam files for consensus peak calling. Peak calling was performed for each replicate in addition to the merged alignments using macs2, first with default settings, and then using the –call-summits options to call subpeaks. Differential binding analysis was performed on the consensus peaks and again on the subpeak summits using the DiffBind package (v3.0.14) in R/Bioconductor. Consensus peaks and differentially bound summits were annotated for genomic features using the ChIPpeakanno package (v3.24.1) in R/Bioconductor.

**Dataset Acquisition and Analyses**

RNA-seq gene expression data from normal mucosa (n = 51) and primary colon tumors (n = 380) was downloaded from the TCGA COAD/READ dataset on Xenabrowser on January 30, 2021. Analyses of single cell RNA-seq gene expression data from human colon tissue were retrieved from The Human Protein Atlas on February 23, 2021. Expression data from human colon tissue were retrieved from The Human Protein Atlas on February 23, 2021.41,70

**Statistical Analysis**

Except where noted, experiments were performed with triplicate measures for each condition (ie, 3 wells of cells with independent treatment, collection of samples, and measurement of analytes). Experiments were repeated at least twice (on separate days), and results are presented as the mean of the experiments, where each data point represents the mean of the replicates from a single experiment. Where graph space is limited, bars represent the mean of independent experiments ± SEM. Statistical significance was determined by 2-tailed Student’s t test or 1- or 2-way analysis of variance (as appropriate), on log-transformed data, with matched analysis accounting for independent experiments. Statistical tests were calculated using GraphPad Prism (GraphPad Software, San Diego, CA, version 9.0) (*P < .05, **P < .01, ***P < .001, ****P < .0001).

All authors had access to the study data and have reviewed and approved the final manuscript.

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