β-Catenin/CBP inhibition alters epidermal growth factor receptor fucosylation status in oral squamous cell carcinoma†

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Epidermal growth factor receptor (EGFR) is a major driver of head and neck cancer, a devastating malignancy with a major sub-site in the oral cavity manifesting as oral squamous cell carcinoma (OSCC). EGFR is a glycoprotein receptor tyrosine kinase (RTK) whose activity is upregulated in >80% OSCC. Current anti-EGFR therapy relies on the use of cetuximab, a monoclonal antibody against EGFR, although it has had only a limited response in patients. Here, we uncover a novel mechanism regulating EGFR activity, identifying a role of the nuclear branch of the Wnt/β-catenin signaling pathway, the β-catenin/CBP axis, in control of post-translational modification of N-glycans on the EGFR. Genomic and structural analyses reveal that β-catenin/CBP signaling represses fucosylation on the antennae of N-linked glycans on EGFR. By employing nUPLC-MS/MS, we determined that malignant human OSCC cells harbor EGFR with a paucity of N-glycan antennary fucosylation, while indolent cells display higher levels of fucosylation at sites N420 and N579. Additionally, treatment with either ICG-001 or E7386, which are both small molecule inhibitors of β-catenin/CBP signaling, leads to increased transcriptional expression of fucosyltransferases FUT2 and FUT3, with a concomitant increase in EGFR N-glycan antennary fucosylation. In order to discover which fucosylated glycan epitopes are involved in the observed effect, we performed in-depth characterization of multiply-fucosylated N-glycans via tandem mass spectrometry analysis of the EGFR tryptic glycopeptides. Data are available via ProteomeXchange with identifier PXD017060. We propose that β-catenin/CBP signaling promotes EGFR oncogenic activity in OSCC by inhibiting its N-glycan antennary fucosylation through transcriptional repression of FUT2 and FUT3.

Introduction

Head and neck squamous cell carcinoma (HNSCC) is a debilitating disease and the sixth most common malignancy in the world, with an estimated incidence of greater than 650,000 cases and an annual toll of 350,000 deaths worldwide. Major risk factors associated with head and neck cancer include alcohol and tobacco use and human papillomavirus (HPV) infection; men are more likely to be diagnosed with oral cavity cancer. Within the head and neck region, squamous cell carcinomas can develop in multiple subsites, with the majority of cases arising in the oral cavity and oropharynx. Oral squamous cell carcinoma (OSCC) accounts for the majority of HNSCC cases diagnosed on a yearly basis. In the United States, a large disparity exists in the survival of black and white individuals diagnosed with cancers of the oral cavity and pharynx; overall survival is 18% lower for black patients compared to white patients, and this difference is even more pronounced among black men. Understanding dysregulated signaling in oral cancer at the molecular, genetic, and epigenetic levels is critical to develop effective therapies to treat patients with OSCC. Major advances in the understanding of the molecular basis of oral cancer include the roles of Wnt/β-catenin signaling, NOTCH1, PI3K, and IL-6/JAK/STAT3 signaling, as well as epidermal growth factor receptor (EGFR), Cetuximab and pembrolizumab/nivolumab, monoclonal antibodies targeting EGFR and PD-1, respectively, are the only FDA-approved targeted therapies available for the
treatment of this malignancy, and even they have had only limited success (<20%) in the clinic.²

A major driver of OSCC is EGFR, a receptor tyrosine kinase (RTK) whose activity is aberrantly upregulated in > 80% of tumors.¹⁴ EGFR is highly modified with N-linked glycans, a structurally diverse class of carbohydrates that share a tri-mannosyl chitobiose (Man₃GlcNAc₂) core and vary in monosaccharide composition and degree of branching. Changes in cell surface protein glycosylation have been shown to alter receptor signaling and frequently accompany malignant transformation and metastasis.¹⁵,¹⁶ For instance, RTK N-glycosylation has been shown to regulate cell proliferation and differentiation, with RTK N-linked glycans being implicated in presenting binding sites for glycan-binding proteins that oppose receptor endocytosis, leading to prolonged signaling.¹⁷–²³ Accordingly, treatment of tumor cells with tunicamycin (to inhibit N-linked glycosylation) reduces RTK signaling.²⁴–²⁸ Both the monosaccharide types and linkages within the N-linked glycans regulate receptor dimerization. In particular, sialylation and antennary fucosylation of N-glycans at specific sites on the extracellular domain of the receptor have been demonstrated to interfere with EGFR dimerization and activation in lung adenocarcinoma.²¹ In contrast, core fucosylation on the glycans favors EGFR dimerization.²⁰,²¹,²⁹ Collectively, studies to date suggest that post-transcriptional modification of EGFR by N-glycans has a large impact on receptor activity.

In OSCC, EGFR signaling converges on Wnt/β-catenin activity, known to play pivotal roles in the pathobiology of this malignancy through the interaction of nuclear β-catenin with the histone acetyltransferase CREB-binding protein (CBP).³⁰ The protein N-glycosylation pathway interacts with Wnt/β-catenin signaling to promote OSCC pathogenesis.³¹–³³ N-glycosylation is regulated at multiple steps in this pathway that impact stem cell maintenance, proliferation and survival. We have shown that a small molecule inhibitor of the β-catenin–CBP interaction, ICG-001 (Fig. 1D), interferes with OSCC proliferation and aggressive features in cellular, zebrafish and murine models.⁵ Given that EGFR is a major driver of OSCC, we examined whether inhibition of β-catenin/CBP signaling also impacted EGFR function. We now show that OSCC-cell line induced mouse tumor xenografts exhibit reduced EGFR abundance, and genomic analyses reveal a positive correlation between ICG-001 and EGFR inhibition. Interestingly, whereas EGFR transcript levels were not affected by ICG-001 treatment, EGFR protein abundance decreased significantly, suggesting that EGFR levels were regulated on a post-translational level. Gene set expression analyses (GSEA) revealed significant increases in the expression of multiple glycosyltransferases, including fucosyltransferases FUT2 and FUT3, following inhibition of β-catenin/CBP signaling. Given that modification of EGFR N-glycans with antennary fucose residues has been reported to inhibit receptor dimerization and activity, we examined whether changes in EGFR fucosylation were associated with different aggressive characteristics of OSCC cells. Our studies suggest that β-catenin/CBP signaling in OSCC suppresses FUT2 and FUT3 expression coincident with reduced EGFR outer chain fucosylation, and that inhibition of β-catenin–CBP interaction relieves FUT2 and FUT3 suppression and promotes fucosylation on the antennae of the N-glycans of EGFR and attenuation of its oncogenic signaling. To our knowledge, this is the first report that aligns changes in EGFR fucosylation with aggressive OSCC phenotypes.

**Results**

**Inhibition of nuclear β-catenin/CBP signaling leads to decreased EGFR protein levels and higher expression of fucosyltransferases FUT2 and FUT3 in human HSC-3-induced mouse xenografts**

Our recent findings provided evidence that inhibition of β-catenin–CBP interaction with the small molecule ICG-001 abrogates OSCC cell proliferation and reduces mesenchymal traits by promoting E-cadherin adhesion.⁶ As EGFR is a major driver of OSCC, we sought to explore the impact of nuclear β-catenin/CBP inhibition on EGFR oncogenic activity in vivo. Thus, we evaluated human HSC-3 cell-derived orthotopic oral tongue tumor xenografts from nude mice treated with either the small molecule inhibitor ICG-001 or vehicle (DMSO) control, since we have reported that these cells form metastatic tumors.⁶,³⁰,³⁴–³⁶ Immunofluorescence analyses revealed lower levels of membranous EGFR in tumors from ICG-001-treated mice compared to tumors from control mice (Fig. 1A, left). This was in striking contrast to increased levels of junctional E-cadherin in tumors from ICG-001-treated mice (Fig. 1A, right).⁵ Furthermore, immunoblot analyses of total tissue lysates demonstrated a statistically significant (p < 0.0001) decrease in EGFR abundance in tumors from mice treated with ICG-001, compared to tumors from control mice (Fig. 1B). At the same time, total tissue levels of E-cadherin increased in response to ICG-001 treatment (Fig. 1B).

In this context, increased nuclear β-catenin/CBP signaling correlates with higher EGFR abundance and lower E-cadherin adhesion. This conclusion was supported by computational analyses comparing the ICG-001 treatment signatures and EGFR inhibition signatures in TCGA OSCCs which revealed a very strong positive correlation between ICG-001 and EGFR inhibition (Fig. 1C and Fig. S1A, B, ESI†), further suggesting that the ICG-001 treatment inhibits genes involved in EGFR signaling in primary oral cancer tumor tissues.

Upon disruption of nuclear β-catenin/CBP signaling with E7386, a novel β-catenin/CBP modulator that displays an activity profile that closely overlaps with that of ICG-001 but exhibits ~ 50–100-fold lower EC₅₀ in OSCC cell lines than ICG-001, the ICG-001 inhibition-associated transcriptional signatures tracked with tumor grade and poor human OSCC patient overall survival.⁶ Likewise, the E7386 inhibition signature was associated with advanced TCGA OSCC tumor grade that is associated with poor overall patient survival (Fig. S3A, ESI†). Lastly, similar to ICG-001, the E7386 inhibition signature was highly correlated with EGFR inhibition in TCGA OSCC (Fig. S3B, ESI†).

The observed downregulation of EGFR abundance in response to the inhibition of β-catenin/CBP activity was not a consequence of transcriptional attenuation, as treatment with either ICG-001
Fig. 1 β-Catenin/CREB-Binding Protein (CBP) inhibition alters EGFR abundance and fucosyltransferase expression in a mouse tongue squamous cell carcinoma orthotopic xenograft model. (A) We used a previously established mouse tongue squamous cell carcinoma orthotopic xenograft model.\textsuperscript{36} Inhibition of β-catenin/CBP with the small molecule inhibitor ICG-001 in HSC-3-induced mouse orthotopic tongue tumor xenografts led to decreased EGFR levels (left, lower panel) compared to the vehicle control (DMSO; left, upper panel). In contrast, β-catenin/CBP inhibition resulted in an increase in E-cadherin at cell–cell interfaces (right, lower panel) compared to the control (DMSO; right, upper panel). (B) Western blots of EGFR and E-cadherin from total tissue lysates (TTLs) harvested from mouse HSC-3-derived orthotopic tongue tumors treated with vehicle (DMSO) or ICG-001. GAPDH shown as a loading control. Quantification of EGFR and E-cadherin abundances are also shown, ***p < 0.0001. Results of triplicate analyses are shown, ± one standard deviation. (C) Heat map comparing gene expression in CAL27 cells and HSC-3 cells treated either with vehicle (DMSO) and ICG-001. Red indicates an increase in expression, while blue indicates a decrease in expression. (D) The chemical structures of ICG-001 and E7386 are shown.\textsuperscript{34,35} (E) Schematic of an N-linked glycan attached to an asparagine (Asn) residue, with potential linkages catalyzed by FUT2, FUT3 and FUT8 indicated. N-Acetylgalactosamine (blue squares), mannose (green circles), galactose (yellow circles); N-acetylnearaminic acid (purple diamonds), and fucose (red triangles) are indicated.
Table 1. Altered fucosyltransferase expression in response to β-catenin/ 
CBP inhibition. Expression change of fucosyltransferases FUT2, FUT3, 
FUT8, the epidermal growth factor receptor (EGFR), and E-cadherin 
(CDH1) after treatment with ICG-001 (top) or E7386 (bottom) compared 
to vehicle control (DMSO) in indolent CAL27 and metastatic HSC-3 
cells. The fold change of the treatment compared to the DMSO 
control (fold Δ) and q-value (q, FDR adjusted) are shown for each condition 
tested.

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<td>CDH1</td>
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or E7386 did not affect EGFR transcript levels (Table 1). Since 
EGFR exhibits high glycoform diversity at multiple N-glycosylation 
sites, and since modifications of the N-glycan structures affect 
protein folding, targeting and activity, we postulated that ICG-001 
and E7386 had impacted EGFR abundance on a post-translational 
level. Therefore, we assessed the impact of ICG-001 and E7386 
hinhibition on the expression of genes that function in the protein 
N-glycosylation pathway. Indeed, treatment of CAL27 and HSC-3 
cells with either ICG-001 or E7386 increased expression of two 
anterary fucosyltransferases, FUT2 and FUT3 (Table 1 and 
Fig. 1C), but did not alter the level of fucosyltransferase FUT8, which 
catalyzes core fucosylation of N-glycan structures (Table 1). While 
α,1,6-linked (core) fucosylation of EGFR catalyzed by FUT8 has 
been shown to increase EGFR signaling, α,1,3-linked antenary 
fucosylation has been reported to suppress EGFR signaling in 
lung adenocarcinoma. α,1,4-fucosylation of GlcNAc in Type-1 chains (Gal[b1-3GlcNAc), while FUT3 catalyzes 
α,1,4-fucosylation of GlcNAc in Type-1 chains (Gal[b1-3GlcNAc) and 
α,1,3-fucosylation of GlcNAc in Type-2 chains (Gal[b1-4GlcNAc) 
(Fig. 1E). These results suggested that fucosyltransferases FUT2 
and/or FUT3 affected EGFR protein levels in OSCC.

MS/MS analyses of EGFR from OSCC cell lines identify multiple 
sites with fucosylated N-glycans

Next, we sought to determine the extent to which EGFR derived 
from untreated OSCC cell lines displayed fucosylated N-glycans. 
EGFR contains 12 N-glycosylation sequons in its extracellular 
domain, divided among the receptor’s four subdomains, I–IV 
(Fig. 2A). To determine the site-specific patterns of EGFR 
N-glycosylation, we immunoprecipitated EGFR from indolent 
CAL27 and metastatic HSC-3 human tongue oral squamous 
carcinoma cell lines. Following proteolysis of EGFR, glycopeptides 
were enriched on an Amide-80 trapping column, separated 
via C18 reversed-phase chromatography, and analyzed via 
Q-TOF MS/MS. Additional analyses were performed without

Fig. 2. N-Glycosylation of the Epidermal Growth Factor Receptor (EGFR). (A) A schematic of EGFR is shown, with N-glycosylation sites labeled. EGFR 
contains four sub-domains (II–IV) within the extracellular domain, a transmembrane (TM) region, the cytoplasmic tyrosine kinase (TK) region and the 
C-terminal tail. (B) Schematic of the analytical workflow for the analysis of EGFR derived from OSCC cell lines. *PNGase F treatment was performed in the 
presence of H₂¹⁸O for peptide analyses and site-occupancy determination. For glycan analyses, H₂O was used. (C) Crystal structure of the EGFR 
extracellular domain (Protein Data Bank ID: 4KRO).64 Based on our survey of EGFR N-glycopeptides derived from OSCC cell lines, N-glycosylated 
asparagine residues bearing complex glycans are shown in red, and those bearing high-mannose type glycans are shown in white. N104 is not shown 
because it is not resolved in this structure determination – this is likely due to the higher flexibility of this region.
glycopeptide enrichment, using data-dependent MS/MS methods on an Orbitrap Tribrid MS, including glycopeptide oxonium ion-triggered Higher-energy Collisional Dissociation (HCD) and Electron-Transfer Dissociation (ETD) combined with gentler HCD (EThcD) fragmentation (Fig. 2B). In total, we detected glycopeptides representing 11 of the 12 reported EGFR N-glycosylation sites (Table S1 and Fig. S4–S14, ESI†). To determine which sites were modified with high mannose, hybrid, or complex N-glycans, we surveyed the glycan compositions associated with these 11 sites.Sites N172, N328, N337, and N599 displayed high mannose N-linked glycans in both CAL27 and HSC-3 cells, indicating that these sites could not be modified by fucosyltransferases FUT2 and FUT3. In contrast, sites N32, N104, N151, N389, N420, N544, and N579 displayed complex glycans or a combination of complex and high mannose glycans in both cell lines (Fig. 2C and Tables S1–S6). Glycopeptides corresponding to site N504 were not detected. Furthermore, treatment with ICG-001 did not significantly alter N-glycosylation site occupancy (Table S7, ESI†). Next, we performed a quantitative comparison of the fucosylation status of N-glycans from indolent CAL27 cells and metastatic HSC-3 cells. To achieve this, EGFR glycopeptides were analyzed in triplicate. Extracted ion chromatograms were generated for each EGFR glycopeptide/glycoform observed, and the area of each was calculated. At EGFR site N389, all glycopeptides detected in both cell lines displayed at least one fucose on complex glycans (Fig. 3A and Tables S2–S6, S9, ESI†). Singly-fucosylated glycoforms at site N389 accounted for 85.0 ± 2.5% of the area in CAL27 cells and 91.1 ± 1.2% of the area in HSC-3 cells, a small but statistically significant difference (p = 3.4 × 10⁻²). The doubly-fucosylated and sialylated N-glycan composition HexNAc₂Hex₅Fuc₂NeuAc₁(N₄H₅F₂S₁) had higher relative abundance in CAL27 cells compared to HSC-3 cells (p = 1.4 × 10⁻²). Triply-fucosylated glycoforms were 4.5-fold higher in CAL27 cells compared to HSC-3 cells (p = 1.1 × 10⁻²) (Fig. 3D). At site N420, 40.3 ± 1.8% of glycoforms (by area) were fucosylated in CAL27 cells, whereas fucosylated glycans accounted for roughly half as much of the total (20.9 ± 3.8%) in EGFR derived from HSC-3 cells, and this difference was highly statistically significant (p = 1.2 × 10⁻³) (Fig. 3B and Table S10A, B, ESI†). Singly-, doubly-, and triply-fucosylated glycoforms were
higher in CAL27 cells compared to HSC3 cells; singly-fucosylated glycans were 1.5-fold higher (p = 7.1 × 10⁻³) and doubly-fucosylated glycans were 3.2-fold higher (p = 2.3 × 10⁻⁴) (Fig. 3E). While triply-fucosylated glycopeptides accounted for 1.4 ± 0.2% of the area of N420 glycopeptides in CAL27 cells, no triply-fucosylated glycopeptides were detected in HSC-3 cells. At site N579, singly-fucosylated glycoforms had lower relative abundances in CAL27 cells compared to HSC-3 cells at 66.5 ± 4.6% and 88.3 ± 0.2% of total N579 glycopeptide area, respectively (Fig. 3C and Table S11, ESI†). However, the relative levels of doubly-fucosylated glycans were 2.3-fold higher in CAL27 cells compared to HSC-3 cells (p = 2.3 × 10⁻²), accounting for 20.2 ± 4.5% and 8.6 ± 0.2% of N579 glycopeptide area, respectively (Fig. 3F). At sites N328, N337, and N599 displaying high mannose glycans, and sites N32, N104, and N544, where a limited distribution of glycoforms were detected, we did not observe any changes in N-glycan fucosylation. In summary, multiple (but not all) EGFR N-glycosylation sites displayed higher proportions of multiply-fucosylated N-linked glycans in indolent CAL27 cells compared to metastatic HSC-3 cells.

**Inhibition of β-catenin–CBP signaling in metastatic HSC-3 cells results in increased fucosylation of EGFR N-glycans**

Given that expression of FUT2 and FUT3 increased in HSC-3 cells compared to HSC-3 cells at 66.5

| Table S10A, B, ESI† | Inhibition of β-catenin/CBP signaling in metastatic HSC-3 cells leads to higher levels of EGFR N-glycan fucosylation. EGFR N-glycan heterogeneity at sites (A) N151, (B) N389, (C) N420, and (F) N579, in EGFR derived from HSC-3 cells treated with either a vehicle control (DMSO) or ICG-001, a small molecule inhibitor of nuclear β-catenin/CBP. Comparison of the proportional area of fucosylated glycan compositions to total glycans at site N420 in CAL27 and HSC-3 cells in response to β-catenin/CBP inhibition (D). Proportional area of singly-, doubly-, and triply-fucosylated N-glycans at site N420 in HSC-3 cells in response to β-catenin/CBP inhibition. (E) At site N420, the proportion of fucosylated N-glycans increased in response to β-catenin/CBP inhibition in HSC-3 cells, and almost matched the levels of fucosylation observed in CAL27 cells. For all sites, nUPLC-MS analyses were performed in triplicate (technical replicates), and the area of each glycopeptide precursor was extracted (based on m/z): N: N-acetylhexosamine, H: hexose (mannose or galactose), F: fucose, S: N-acetylneuraminic acid (sialic acid). Multiply-fucosylated glycan compositions are outlined (box with dashed line). *p ≤ 0.05; **p ≤ 0.01; ***p ≤ 0.005.
analyses of released permethylated MS analyses of immunoprecipitated EGFR, MALDI-TOF MS analyses of EGFR, MALDI-TOF MS analyses of immunoprecipitated EGFR, MALDI-TOF MS analyses of released permethylated N-linked glycans derived from the whole lysates of HSC-3 and CAL27 cells treated with either DMSO or ICG-001 showed no significant changes in fucosylation (Fig. S15A–C, ESI†), though the dynamic ranges of these analyses were limited, with the result that we did not detect multiply-fucosylated species.

Inhibition of β-catenin/CBP signaling shifts the balance of fucosylated EGFR N-linked glycans toward antennary-fucosylation

Having observed an increase in fucosylation in response to β-catenin/CBP inhibition in HSC-3 cells, we next made determinations of the location of the fucose residue(s) on EGFR N-linked glycans. We observed multiple values for the retention times of fucosylated glycopeptides sharing the same precursor m/z values in HSC-3 cells after treatment with ICG-001 (Fig. 5A), and hypothesized that this diversity might be due to the presence of variations in the linkage positions of the fucose residues. Because fucose residues are frequently cleaved off during fragmentation with higher-energy collisional dissociation (HCD), we performed additional analyses with (a) low-energy HCD and (b) electron transfer dissociation with supplemental activation (EThcD) to favor the generation of glycopeptide fragment ions while minimizing glycosidic bond fragmentations. This enabled us to distinguish between antennary- and core-fucosylated N-glycopeptides. An HCD MS/MS spectrum of the early-eluting glycopeptide corresponding to EGFR site N420,406TKQHGQFSLAVVSLNITSLGLR427 + HexNAc4Hex5dHex1 ([M + 4H] 4+/z 1035.2483), provided evidence for a core-fucosylated isomer. This assignment was supported by the presence of the product ion consisting of the intact triply protonated peptide + HexNAc,dHex3 (pNF) ion at m/z 906.8231 (3+), as well as low levels of the HexNAc3Hex4dHex3 (NHF+) oxonium ion at m/z 512.1954 (Fig. 5B). Such a fragment could be generated with low efficiency by rearrangement of fucose during fragmentation.40–43 In contrast, the product ion scan of the late-eluting isomer showed very high levels of the HexNAc3Hex4dHex3 (NHF+) oxonium ion observed at m/z 512.1953, consistent with the presence of outer-arm fucosylation. As expected on the basis of this assignment, we detected high relative levels of the peptide + HexNAc1 peak without fucose at m/z 858.1374, and higher intensities of product ions associated with loss of components that would include antennary fucose, including [M − NHF]3+ observed at m/z 1209.5911 and [M − NH2F]3+ at m/z 1155.5740 (Fig. 5C). We have documented additional examples of such spectral differences in core and antennary fucosylation (Fig. S16–S18, ESI†). In addition, we examined multiply-fucosylated species. EThcD fragmentation of the dominant doubly-fucosylated EGFR site N420 glycopeptide species in HSC-3 cells after treatment with ICG-001, 41,42406TKQHGQFSLAVVSLNITSLGLR427 + HexNAc4Hex5dHex1 ([M + 4H] 4+/z 1071.7526), indicated that both fucose residues were located on the N-glycan antennae, based on the presence of product ions in the MS2 spectrum, including the triply protonated fragment ions that lack fucose: peptide + HexNAc3Hex3 (pN3H3), observed at m/z 1088.2133 (3+) and m/z 1155.5729 (3+) respectively (Fig. 6 and Fig. S17A, B, ESI†). These findings were consistent with fucosyl transferase expression level results.

![Fig 5](image-url)

**Fig. 5** β-Catenin/CBP inhibition shifts EGFR N-glycan fucosylation from “Core” fucosylation to a signal suppressing “Outer-Arm” fucosylation. (A) Extracted ion chromatograms (EICs) of the EGFR site N420 tryptic glycopeptide 406TKQHGQFSLAVVSLNITSLGLR427 + HexNAc4Hex5dHex1 ([M + 4H] 4+/z 1035.2483), in CAL27 and HSC-3 cells treated with either DMSO or ICG-001. Note two partially-resolved peaks in the chromatograms centered at approximately 32.2 min and 32.4 min with the same m/z value. Selected regions of the low-energy (10% NCE) HCD product ion spectra of the ‘early’ eluting form with annotated peaks demonstrating evidence of core fucosylation (B), and the ‘late’ eluting form with annotated peaks demonstrating evidence of antennary fucosylation (C). p = peptide, N = HexNAc, H = hexose, F = dHex.
Discussion

The Wnt/β-catenin signaling pathway and EGFR signaling are known to cross-talk at multiple levels and to enhance each other's activities, although the details of these interactions have not been well defined in OSCC. The Wnt/β-catenin pathway, the β-catenin/CBP axis, impacts EGFR protein abundance by modulating its N-glycosylation status via transcriptional repression of two fucosyltransferases, FUT2 and FUT3, shown to fucosylate outer chain N-glycans on EGFR and inhibit its activity. Our detailed structural analyses provide evidence that antenary fucosylation of EGFR N-glycans is associated with indolent OSCC phenotypes and that suppression of EGFR fucosylation aligns with malignant features. We have shown previously that HSC-3 cells displayed high nuclear β-catenin/CBP activity and that disruption of the β-catenin–CBP interaction reverted the mesenchymal phenotype of HSC-3 cells to an epithelial morphology similar to indolent CAL27 cells. Accordingly, disruption of the β-catenin–CBP interaction effectively relieved β-catenin/CBP-driven suppression of FUT2 and FUT3. These effects of nuclear β-catenin/CBP signaling on EGFR N-glycosylation reveal a novel level of EGFR regulation that includes transcriptional suppression of specific post-translational N-glycan remodelling enzyme activities.

Results showed that inhibition of β-catenin/CBP activity with ICG-001 in HSC-3-induced mouse tumor xenografts caused a decrease in total tissue levels of EGFR but not in EGFR transcript levels. Instead, there was an increase in the expression of FUT2 and FUT3 (Fig. 1A-C). Both of these fucosyltransferases transfer fucose to N-glycan antennae: FUT2 catalyzes α1,2-linkage of fucose residues to N-linked glycans, while FUT3 can catalyze the attachment of fucose via α1,3- and α1,4-linkages. The expression of an inactive form of FUT2 is associated with the development of oral cancer, suggesting that the active form of the enzyme has a protective role in OSCC. In contrast, FUT3 has the potential to contribute to the expression of sialyl-Lewis x (SLex) glycan epitopes and has been shown to play a role in establishing the invasive and metastatic properties of oral cancer stem cells. Thus, different fucosyltransferases may have opposing roles in oral cancer. On the other hand, the Lewis y (Le(y) antigen (with both α1,2- and α1,3-linked fucose) is expressed in OSCC cell lines and tissues, but disappears in invasive regions, suggesting that fucosylated glycans may be involved in the suppression of aggressive features. Moreover, EGFR antenary-fucosylated glycans are associated with patient survival, and suppress receptor dimerization and signaling in lung adenocarcinoma. Further studies are required to elucidate specific roles of FUT2 and FUT3 in modulating EGFR and potentially other oncogenic glycoproteins in OSCC (Fig. 7A).

We identified multiple EGFR sites bearing fucosylated N-glycans (Fig. 2C). Several EGFR N-glycosylation sites appear to play key roles in receptor dimerization and signaling, including site N420 which alters EGFR dimerization, and N579 which plays a role in EGFR signaling by preventing the receptor from becoming activated (auto-inhibition) in the absence of ligand stimulation. Using nUPLC-MS/MS, we tracked the glycosylation of EGFR across multiple sites, including sites N420 and N579. While EGFR fucosylation from CAL27 and HSC3 cells differs modestly at certain sites, we observed dramatic changes in EGFR fucosylation at sites N420 and N579. EGFR from metastatic HSC-3 cells displayed low levels of fucosylated N-glycans, while EGFR from non-metastatic CAL27 cells displayed higher levels of fucosylation at multiple sites, including sites N420 and N579.
Post-translational regulation of RTKs has been implicated in cell proliferation and differentiation, and inhibition of N-glycosylation has been shown to disrupt RTK oncogenic signaling in tumor cells. Moreover, other post-translational mechanisms, such as reduced proteolytic shedding, have been implicated in RTK-mediated cancer treatment resistance. Our present studies suggest that the β-catenin/CBP axis modulates EGFR signaling through downregulation of FUT2 and FUT3 expression and activity (Fig. 7B). Given that decrease in fucosylation has been reported in the invasive regions of OSCC tumors, it is possible that inhibition of this modification by FUT2 and FUT3 relieves EGFR from suppression and drives aggressive OSCC phenotypes. Likewise, fucosylated epitopes reported in OSCCs may serve to suppress EGFR signaling. Thus, given that ICG-001 and E7386 have overlapping activities, inhibition of β-catenin/CBP signaling with the more soluble E7386 may serve as a therapeutic approach not only to inhibit downstream targets of β-catenin/CBP that drive disease progression, but also to downregulate EGFR pro-tumorigenic activity in OSCC.

Experimental procedures

Reagents and cell lines
PNGase F was purchased from New England Biolabs (Ipswich, MA). MS grade trypsin (tosyl phenylalanine chloromethyl ketone treated) was purchased from Thermofisher Scientific (Waltham, MA). Pierce C18 Tips (100 µL), dithiothreitol (DTT), ammonium bicarbonate (NH₄HCO₃), and iodoacetamide were purchased from Sigma Aldrich (St. Louis, MO, USA). Small (10 µL) C18 ZipTips were purchased from Millipore (Billerica, MA). Heavy water (H₂¹⁸O, 99%) was purchased from Cambridge Isotope Laboratories (Andover, MA). CAL27 cells were purchased from ATCC while HSC-3 cells were obtained from XenoTech (Tokai, Japan). CAL27 and HSC-3 cell lines were routinely authenticated by short tandem repeat profiling every 6 months, either by ATCC or Genetica. ICG-001 was purchased from Selleckchem (Houston, TX) and E7386 was provided by Eisai Inc (Woodcliff Lake, NJ). For immunofluorescence and western blot analyses, rabbit monoclonal antibody to EGFR [EP39Y; ab52894] and mouse monoclonal antibody to E-cadherin [4A2; ab231303] were purchased from Abcam (Cambridge, MA). DAPI and goat anti-rabbit and anti-mouse IgG derivatized with either Texas Red or FITC secondary antibodies were obtained from Molecular Probes (Eugene, OR). For western blot analyses, anti-GAPDH antibody (mouse monoclonal, G8795) was purchased from Sigma-Aldrich (St. Louis, MO).

Immunofluorescence and western blot analyses
HSC-3 cell-derived mouse tumor xenografts were generated and processed for immunofluorescence and immunoblot analyses as previously described. Tumor sections (5 µm) were immunostained with antibodies for EGFR and E-cadherin, counter-stained for nuclei with DAPI, followed by incubation with secondary Texas Red- and FITC-conjugated antibodies. Negative controls lacked primary antibodies. The slides were mounted in ProLong Gold Antifade and optical sections (0.5 µm intervals) were analyzed by confocal microscopy using a Zeiss...
For Western blot analyses, total tumor tissue lysates were prepared by grinding tumors to a fine powder in liquid nitrogen, followed by extraction of total tissue proteins with Triton/β-ocetylglucoside buffer. Protein concentrations were determined using the BCA assay (Pierce). Tissue lysates (20–30 μg of total protein) were fractionated via 4–12% SDS–PAGE, transferred onto polyvinylidene difluoride membranes, blocked with 5% nonfat dry milk, and incubated sequentially with primary antibodies to EGFR, E-cadherin and GAPDH. Protein-specific detection was carried out with horseradish peroxidase-labeled secondary antibodies conjugated to horseradish peroxidase (Bio-Rad), and Enhanced Chemiluminescence Plus (Amersham Biosciences). Signal intensities were normalized to GAPDH.

β-Catenin/CBP inhibition and EGFR immunoprecipitation

CAL27 and HSC-3 cells were treated with 10 μM ICG-001/DMSO for 50–55 h. Cells were cultured to approximately 60% confluence, then lysed using Triton-X-100/β-ocetylglucoside buffer (10 mM imidazole, 100 mM NaCl, 1 mM MgCl₂, 5 mM Na₂EDTA, 1% Triton-X-100, 0.87 mg mL⁻¹ β-ocetylglucoside) and MS-SAFE (Sigma) protease and phosphatase inhibitor cocktail. Protein levels of total cell lysates from CAL27 and HSC3 cells were quantified by absorption (280 nm), and 700 μg of lysate was used per immunoprecipitation. Immunoprecipitation was performed according to the manufacturer’s guidelines using a Pierce Cross-link ImmunoPrecipitation Kit from Thermo Scientific (#26147) for Western blot analysis. Following the trapping step, peptides were separated on the analytical column according to the following conditions: 0–1 min: 2% B, 1–3 min: 2–5% B, 3–43 min: 5–40% B (Solvent B: 99% acetonitrile, 0.1% formic acid). MS scans were acquired with the following settings: 70 000 resolution at 1600 m/z, maximum injection time of 60 ms, isolation window of 1.2%, and dynamic exclusion of 10 s. Profile data were acquired at 4 μL min⁻¹ for 4 min with 1% acetonitrile and 0.1% formic acid (Solvent A). Following the trapping step, peptides were separated on the analytical column according to the following settings: 0–1 min: 2% B, 1–3 min: 2–5% B, 3–43 min: 5–40% B (Solvent B: 99% acetonitrile and 0.1% formic acid). MS scans were acquired with the following settings: 70 000 resolution at m/z 400, scan range m/z 370–1880, 1 μscan per MS, AGC target 1 × 10⁶, and a maximum injection time of 100 ms. MS2 scans were acquired with the following settings: 17 500 resolution at m/z 400, AGC target of 5 × 10⁵, maximum injection time of 60 ms, isolation window 2.0 m/z, isolation offset of 0.4 m/z, normalized collision energy (NCE) of 27%, exclusion of charge states 1 and >2, underfill ratio of 1.2%, and dynamic exclusion of 8 s. Profile data were recorded for MS and MS2 scans.

Glycopeptide nanoLC-MS/MS

For the initial characterization of glycosylation, EGFR glycopeptides were enriched, separated, and analyzed using a 6550 Q-TOF MS with a 1200 series nanoflow HPLC-Chip-ESI source fitted with a custom HPLC-Chip with a 360 nL TSK Gel Amide-80 5 μm trapping and enrichment column and a 150 mM × 75 μm Polaris C18-A 3 μm analytical column (all from Agilent Corp., Santa Clara, CA). EGFR digests were initially dissolved in 50% acetonitrile with 0.1% TFA. Immediately prior to treatment, the digested peptides were fractionated by IMAC on a Trap column fitted with a custom HPLC-Chip with a 360 nL TSK Gel Amide-80 5 μm trapping and enrichment column and a 150 mM × 75 μm Polaris C18-A 3 μm analytical column (all from Agilent Corp., Santa Clara, CA). EGFR digests were initially dissolved in 50% acetonitrile with 0.1% TFA. Immediately prior to
injection, the concentration of acetonitrile was adjusted to 80% acetonitrile and 0.1% TFA. After each injection of the sample onto the Amide-80 enrichment column, the column was washed at a flow rate of 1.5 μL min⁻¹ for 4 min using 80% acetonitrile and 0.1% TFA, followed by elution of the sample onto the C18 analytical column with 1% acetonitrile and 0.1% formic acid. Finally, glycopeptides were separated on the analytical column using a gradient from 1% to 40% acetonitrile with 0.1% formic acid at a flow rate of 0.2 μL min⁻¹. The 6550 Q-TOF mass spectrometer was operated in positive mode using the high-resolution, extended dynamic range (2 GHz) setting. MS spectra were collected over the range m/z 150-2000, and MS2 spectra were collected from m/z 50-3000. The ion source gas temperature was set to 225 °C, and the flow was set at 11 L min⁻¹, with a capillary voltage of 1900 V. Precursors ≥1200 counts and charge states ≥2 were selected for fragmentation, and the collision energy was set according to the equation y = mx + b, with y being the collision energy, slope m = 5, x representing the charge state, and the offset b = -4.8. Spectra were collected in profile mode. The m/z measurement error for all peaks with signal-to-noise (S-to-N) ratio of >10 was within 5 ppm.

**Glycopeptide analysis for MS (only), HCD oxonium ion-triggered low-energy HCD, and EThcD**

For MS-only, low-energy HCD, and EThcD analyses, EGFR tryptic peptides were analyzed on an Orbitrap Fusion Lumos Tridrib mass spectrometer (Thermo Scientific) equipped with an ACQUITY UPLC M-Class system (Waters) and a TriVersa NanoMate (Advion). For chromatographic separation, a nanoEase Symmetry C18 UPLC Trap Column (100 Å, 5 μm, 180 μm × 20 mm, Waters) was used for trapping, and a nanoEase MZ HSS C18 T3 UPLC Column (100 Å, 1.8 μm, 75 μm × 100 mm, Waters) was used for separation. The peptide trapping and separation were performed as described above for peptide/glycopeptide analysis. All analyses were performed in positive mode, with the RF lens set to 30%. For MS-only analyses, initial MS scans were acquired with the following settings: isolation window m/z 1.6, 30% collision energy, 120 000 resolution at m/z 400, scan range m/z 500-2000, 100 ms maximum injection time, AGC target 4 × 10⁵, and 2 μscan per MS. For low-energy HCD analyses, initial MS scans (30% collision energy) were acquired with the following settings: 30 000 resolution at m/z 400, scan range m/z 100-2000, 1 μscan per MS, AGC target 5 × 10⁴, and a maximum injection time of 60 ms. Based on the initial HCD (30%) MS2 scans, a second low-energy fragmentation step was triggered if at least two glycopeptide oxonium ions were detected (at m/z 204.0867, HexNac; m/z 138.0545, HexNac–CH₃O; and m/z 366.1396, HexNacHex) within a 15 ppm mass tolerance. EThcD MS2 spectra were acquired after isolation in the quadrupole with the following settings: isolation window 1.6 m/z, a first mass of m/z 100, AGC target 1 × 10⁶, 30 000 resolution at m/z 400, 2 μscan per MS, and 150 ms maximum injection time.

**Analysis of peptide and glycopeptide nUPLC-MS/MS data**

To confirm protein identity, nano LC-MS/MS data were processed using Mascot search algorithm with the UniProtKB/Swiss-Prot *Homo sapiens* protein sequence database (modified September 29, 2019). For all data sets, cleavage rules were applied for each specific protease (trypsin: K, R, P*, F*) and peptides with up to two missed cleavages were considered. The following peptide modifications were considered: methionine oxidation (variable), deamidation (variable), deamidation with ¹⁵O (variable), and carbamidomethylation (fixed). To assign glycopeptide tandem mass spectra, nanoLC-MS/MS data were processed using Byonic (Protein Metrics), with a custom protein sequence database consisting of the forward and reverse protein sequence of human EGFR. Cleavage rules were consistent with those used in Mascot searches (trypsin). The following peptide modifications were considered: methionine oxidation (variable), carbamidomethylation (fixed), and an N-glycan database consisting of 300+ mammalian N-glycans, including multiply-fucosylated glycans. Following the glycopeptide assignment step, extracted ion chromatograms (EICs) were generated manually for each glycopeptide precursor, using a +/- 0.01 m/z window for extraction, and the area under the curve was determined using Xcalibur 4.1 (Thermo Scientific).

**N-Glycan release, permethylation, and MALDI-TOF MS analysis**

Three-hundred μg of whole cell lysates from CAL27 and HSC-3 cells treated either with DMSO (vehicle control) or 10 μM ICG-001, were subjected to N-glycan release via filter-aided N-glycan separation with 500 units of PNGase F (NEB) in 50 mM ammonium bicarbonate buffer overnight at 37 °C. Samples were passed through SepPak C18 (1 cc) cartridges (Waters, Milford, MA, USA), and the flow-through was dried. Samples were resuspended in DMSO, then subjected to permethylation with periodic addition of methyl iodide (Sigma-Aldrich, St. Louis, MO, USA) according to an established protocol. Following permethylation, glycans were dried, then dissolved in 10% acetonitrile/90% water with 0.1% formic acid, then desalted using a C18 ZipTip™ (Millipore, Billerica, MA). Glycans were eluted into 10 μL of 60% acetonitrile/40% water containing
Billerica, MA, USA) using 10% laser power and summing spectrometry data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository with the dataset identifier PXD017060, in compliance with MIRAGE standards.

Data availability

Peptide, glycopeptide and formerly-N-glycosylated peptide mass spectrometry data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository with the dataset identifier PXD017060, in compliance with MIRAGE standards.

Statement of contributions


Animal studies

All experiments involving mice were approved by the Boston University Medical Center Institutional Animal Care and Use Committee. All institutional and national guidelines for care and use of laboratory animals were followed.

Conflicts of interest

Disclosure: Trustees of Boston University (“Boston University”) and Eisai, Inc. (“Eisai”) have executed a Material Transfer Agreement (“MTA”) in which Dr Kukuruzinska is a named investigator and a Sponsored Research Agreement (“SRA”) in which Dr Kukuruzinska is a named investigator. All other authors declare that they have no conflicts of interest.

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References


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