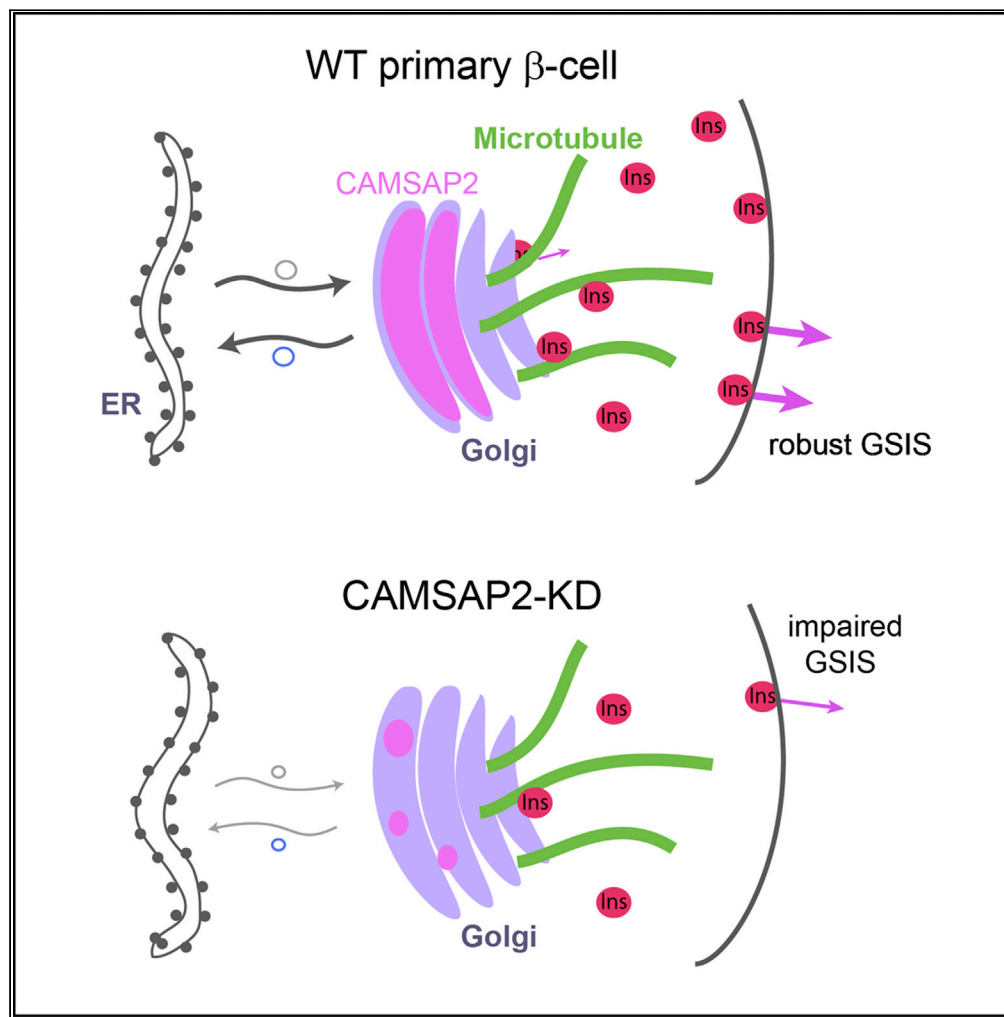


Article

CAM SAP2 localizes to the Golgi in islet β -cells and facilitates Golgi-ER trafficking

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Highlights

CAMSAP2 localizes to the Golgi apparatus in primary β -cells independent of microtubules

The localization of CAMSAP2 at the Golgi is specific to primary β - but not α -cells

CAMSAP2 facilitates Golgi-ER trafficking to support efficient insulin production

Knockdown of CAMSAP2 reduces total insulin and attenuates GSIS

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Article

CAMSP2 localizes to the Golgi in islet β -cells and facilitates Golgi-ER trafficking

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SUMMARY

Glucose stimulation induces the remodeling of microtubules, which potentiates insulin secretion in pancreatic β -cells. CAMSP2 binds to microtubule minus ends to stabilize microtubules in several cultured clonal cells. Here, we report that the knockdown of CAMSP2 in primary β -cells reduces total insulin content and attenuates GSIS without affecting the releasability of insulin vesicles. Surprisingly, CAMSP2 knockdown does not change microtubule stability. Unlike in cultured insulinoma cells, CAMSP2 in primary β -cells predominantly localizes to the Golgi apparatus instead of microtubule minus ends. This novel localization is specific to primary β - but not α -cells and is independent of microtubule binding. Consistent with its specific localization at the Golgi, CAMSP2 promotes efficient Golgi-ER trafficking in primary β -cells. Moreover, primary β -cells and insulinoma cells likely express different CAMSP2 isoforms. We propose that a novel CAMSP2 isoform in primary β -cells has a non-canonical function, which promotes Golgi-ER trafficking to support efficient production of insulin and secretion.

INTRODUCTION

Pancreatic islets are key regulators of glucose homeostasis. After a meal, glucose transported into pancreatic β -cells is metabolized to induce exocytosis of insulin secretory vesicles already existing in the cytoplasm.^{1–3} Insulin promotes the mobilization and usage of glucose in peripheral tissues to clear blood sugar.^{4,5} Thus, efficient biogenesis of insulin vesicles and proper stimulation-secretion coupling in β -cells are the prerequisite for avoiding diabetes, a disease featured by sustained high glucose in circulation.

The synthesis of insulin starts with the translation of its mRNA into preproinsulin, which is imported into the ER and processed to generate proinsulin.⁶ After proper folding and the formation of three intramolecular disulfide bonds, proinsulin is transported to the Golgi apparatus [referred to as Golgi in short].⁷ In the Golgi, proinsulin assembles into hexamers and is packed into secretory vesicles together with other vesicular proteins such as Ca^{2+} sensors.^{8,9} Further proteolytic cleavage generates C-peptide and mature insulin inside the secretory vesicles, which are thereafter stored in β -cell cytoplasm awaiting release.^{10–12}

Islet β -cells store thousands of insulin vesicles, a portion of which is released during a biphasic response to glucose stimuli.¹³ The first phase of glucose-stimulated insulin secretion (GSIS) that lasts a few minutes utilizes mostly vesicles that are pre-docked or very close to the plasma membrane and can readily respond to calcium influx [the readily releasable pool, RRP].^{14–16} The second phase secretion utilizes both vesicles from RRP and ones stored deeper in the cytoplasm that needed mobilization to reach cell periphery [the reserved pool, RP].^{17,18} In both phases, “young” insulin granules generated within several hours before the stimulation are most efficiently utilized.^{19,20} Thus, renewal of stored insulin vesicles is critical to the ability of β -cells to function and survive under metabolic stress. To provide sustainable β -cell functions, glucose metabolism triggers insulin biosynthesis that replenishes and renews the insulin pools.²¹ Efficient processing of insulin through the ER-Golgi trafficking is necessary for this process.

Besides metabolic signals, GSIS is tightly regulated by the cytoskeleton. Both actin filaments and microtubules mediate the transportation of insulin vesicles by molecular motors and can restrict their docking to

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the plasma membrane.^{22–25} Microtubules are biological polymers comprise of directionally aligned α -/ β -tubulin heterodimers forming an α -tubulin exposed (minus) end and a β -tubulin exposed (plus) end.²⁶ The orientation of microtubules and positioning of their ends inside a cell determine the routes of vesicle transportation by plus or minus end-directed motors. Microtubules regulate both phases of GSIS by controlling the transportation of insulin vesicles toward/away from secretion sites and the selectivity of young over aged vesicles.^{23,27–30} Additionally, microtubule nucleation regulated by cAMP/EPAC signaling facilitates insulin vesicles budding from the Golgi.³¹ Although some mechanistic details of microtubule-mediated insulin vesicle biogenesis are known, the roles of many specific microtubule-binding proteins, which regulate fundamental properties of microtubules, have not been explored yet.

Microtubule plus ends can polymerize at relatively low tubulin concentrations, which allows its efficient elongation and stochastic dynamics (dynamic instability) in cells.³² Minus ends, in contrast, have a slower polymerization kinetics and require a higher tubulin concentration for growth. Accordingly, at low physiological tubulin concentrations in cells, minus ends are prone to depolymerization, unless protected. For long-lasting microtubules, minus ends are either capped by a γ -tubulin ring complex or stabilized by minus end-targeting proteins [-TIPs].^{33,34}

CAMSAPs (CAIModulin-regulated Spectrin-Associated Proteins) are -TIPs presented by three paralogues in mammals, CAMSAP1, 2, and 3,³⁵ which stabilize microtubule minus ends.^{36,37} CAMSAP1 tracks growing minus ends, whereas CAMSAP2 and CAMSAP3 remain bound to microtubule lattice of growing minus ends, which recruit more CAMSAPs and overtime form short stretches of CAMSAP-decoration.^{38,39} These properties lead to essential roles of CAMSAPs in organizing non-centrosomal microtubule arrays in epithelial cells and neurons.^{40–43} Specifically, CAMSAP2 promotes the anchoring of microtubule minus ends to the Golgi in retinal pigment epithelial cells through interacting with Golgi proteins AKAP450 and myomegalin.⁴⁴ Intriguingly, polymorphisms in the CAMSAP2 locus are associated with type 1 diabetes in human.⁴⁵ These findings, together with the established importance of microtubules in GSIS, led us to pursue its roles in islet β -cells. We report that CAMSAP2 exhibits a novel subcellular localization at the Golgi in primary β -cells independent of binding to microtubules. Further analyses revealed CAMSAP2 is required for efficient Golgi-ER trafficking, which facilitates robust insulin biosynthesis and GSIS.

RESULTS

Knockdown of CAMSAP2 in primary islet β -cells impairs insulin secretion

To test whether CAMSAP2 is required for primary β -cells secretion, we used lentivirus to deliver CAMSAP2-targeting shRNAs for specific knockdown and examined stimulated secretion of these cells. To maximize the amount of islet cells accessible to lentivirus, isolated mouse islets were dissociated into individual cells, infected with lentivirus, and re-aggregated to form pseudoislets (Figures 1A and 1B). This process allows efficient infection of islet cells by lentivirus (60-70% of cells in a pseudoislet). The cell-cell interactions inside pseudoislets provide a niche similar to inside an intact islet and is a good platform to examine the secretion of β -cells.⁴⁶ Compared to those infected with non-targeting control shRNA (control cells), islet cells infected with CAMSAP2-targeting shRNA (CAMSAP2-KD cells) show reduced CAMSAP2 immunofluorescence signal (Figures 1C and S1). These results demonstrate the specificity of CAMSAP2 antibodies while showing the feasibility of knocking down CAMSAP2 in primary islet β -cells. Importantly, pseudoislets comprise cells expressing either of the two CAMSAP2-targeting shRNAs that do not have altered basal insulin secretion, but exhibit significantly impaired GSIS when compared to the control (Figure 1D). These results suggest that CAMSAP2 is required for robust GSIS.

We next tested whether insulin vesicles in CAMSAP2-KD pseudoislets lack essential components for secretion. Because acute increase in cAMP or Ca^{2+} levels leads to the release of only secretion-competent insulin vesicles,^{9,47} we analyzed secretion responses to cAMP and Ca^{2+} in CAMSAP2-KD cells. We stimulated pseudoislets with Liraglutide, a GLP-1 analog that promotes GSIS mainly through the increasing cAMP level, and found that it enhances GSIS of both control and CAMSAP2-KD pseudoislets (Figure 1D). Elevated K^{+} depolarizes β -cell membrane to induce Ca^{2+} influx. We found that KCl induces similar levels of secretion in CAMSAP2-KD and in control pseudoislets (Figure 1D). These results suggest that CAMSAP2 does not regulate the responsiveness of insulin vesicle to cAMP or Ca^{2+} influx. We hypothesized that CAMSAP2 may influence GSIS by regulating the biogenesis of insulin vesicles and/or the replenishment of the releasable pool of insulin during secretion through modulating microtubule dynamics and tested them later in discussion.

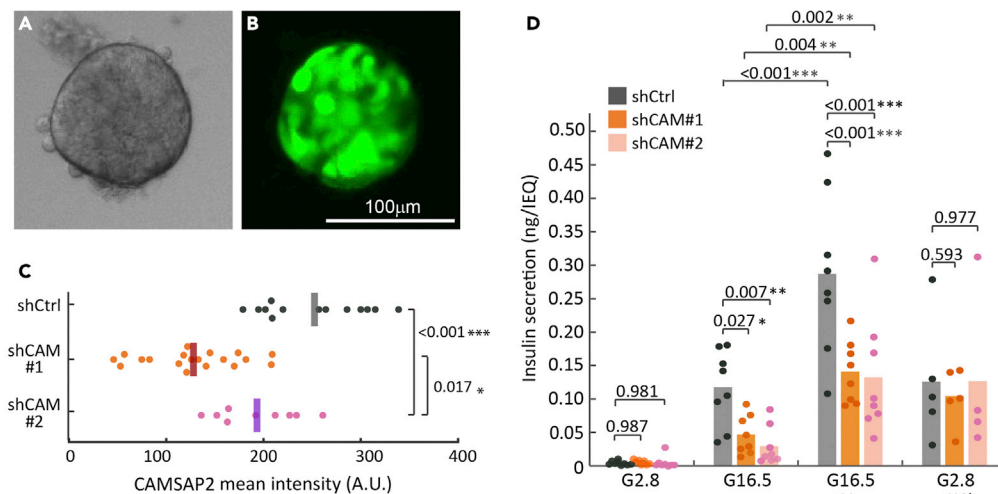


Figure 1. Knockdown of CAMSAP2 in primary β -cells impairs glucose-stimulated insulin secretion but not cAMP- or potassium-stimulated secretion

(A and B) Representative fluorescence images of a pseudoislet generated from dissociated islet cells transduced with lentivirus containing shRNA and a tGFP expression cassette. (C) Quantification of CAMSAP2 intensity of transduced primary β -cells in mouse islets. Dots represent individual β -cells. * $p < 0.05$, *** $p < 0.001$ (Dunnnett's multiple comparisons test). (D) Insulin secretion of pseudoislets expressing non-targeting (shCtrl) or CAMSAP2-targeting (shCAM) shRNA, shown as ng insulin secretion per IEQ within a 45 or 30 minutes window. Dots represent individual wells containing multiple pseudoislets. Columns represent the average of three animals * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ (multiple comparisons corrected by the false discovery rate of Benjamini and Hochberg). Pseudoislets were incubated in 2.8 glucose for 60 min, in 16.5 mM glucose with or without 100 nM Liraglutide for 45 min, or in 2.8 mM glucose plus 30 mM KCl for 30 min. Secretion was normalized by the total pseudoislet volume presented in IEQ (defined as $1,767,146 \mu\text{m}^3$).

CAMSAP2 expressed in primary islet β -cells does not regulate microtubule stabilization and architecture

CAMSAP2 is known as a microtubule minus end-binding protein and regulates microtubule dynamics and positioning.³³ We first tested whether CAMSAP2 regulates the amount and stability of microtubules in islet β -cells. We incubated intact mouse islets with lentivirus carrying shRNAs and individual transduced β -cells on the surface of each islet were analyzed. To evaluate the level of microtubules in transduced cells, islets were first extracted with cold methanol to remove free tubulin in the cytosol and then fixed with paraformaldehyde. We compared CAMSAP2-KD to control β -cells and did not observe a clear difference in the organization of the microtubule network or a significant change in the amount of polymerized tubulin evaluated by mean tubulin intensity after methanol extraction (Figures 2A-2G).

We next tested whether the lifetime of microtubules, a readout for their stability, is affected by CAMSAP2-KD. We measured the level of deetyrosinated tubulin, which has the C-terminal tyrosine of α -tubulin removed, in microtubules. This modification occurs only to polymerized tubulin and its accumulation indicates long-lived microtubules.⁴⁸ We found no difference in the levels of deetyrosinated tubulin between CAMSAP2-KD and control β -cells incubated in low (2.8 mM) or high (11 mM) glucose concentrations (Figures 2H-2T). Moreover, CAMSAP2-KD does not affect the glucose-induced microtubule turnover, which we previously reported to be essential for robust GSIS,²⁷ as evidenced by a significant reduction of deetyrosinated tubulin level in both CAMSAP2-KD and control β -cells when stimulated with 11 mM glucose (Figures 2H-2T). These results suggest that CAMSAP2 is not involved in glucose-dependent microtubule destabilization and indicate that CAMSAP2 does not noticeably contribute to microtubule organization, lifetime, and stability in primary β -cells.

CAMSAP2 is enriched at the Golgi in primary β -cells

In contrast to the reported function of CAMSAP2 in stabilizing microtubule minus ends, CAMSAP2-KD in primary β -cells does not alter microtubule abundance and stability. We next examined the subcellular

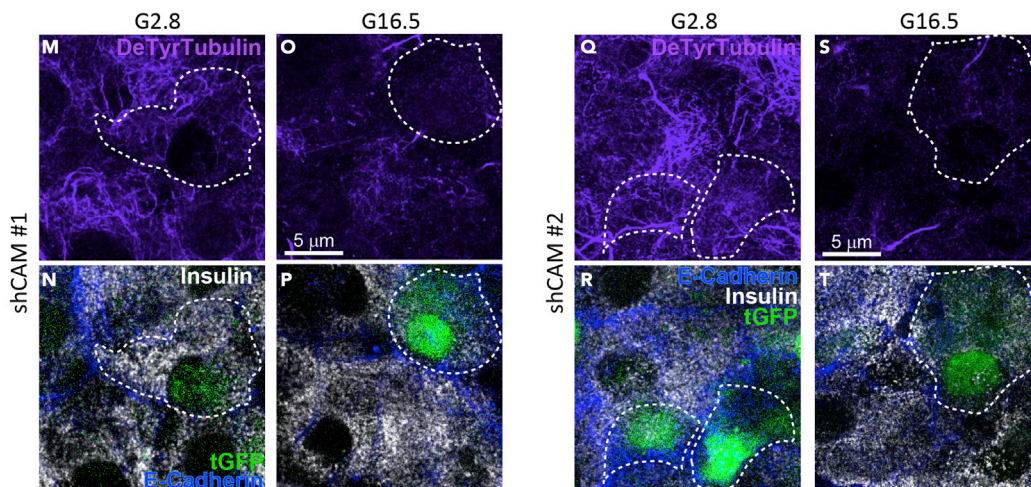
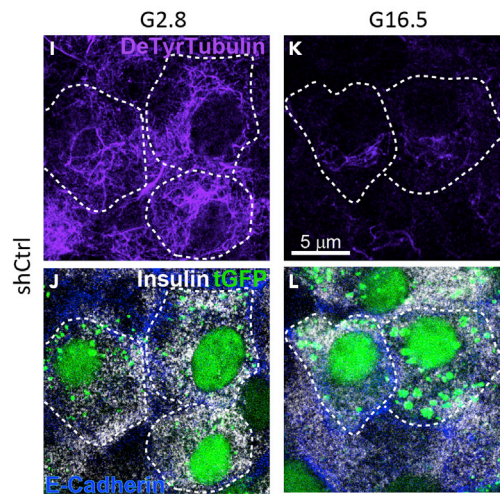
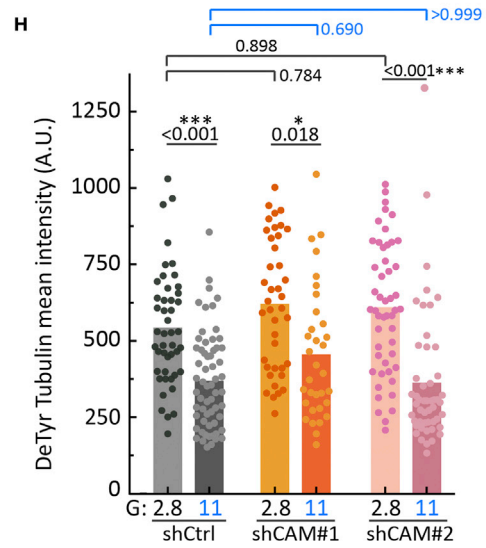
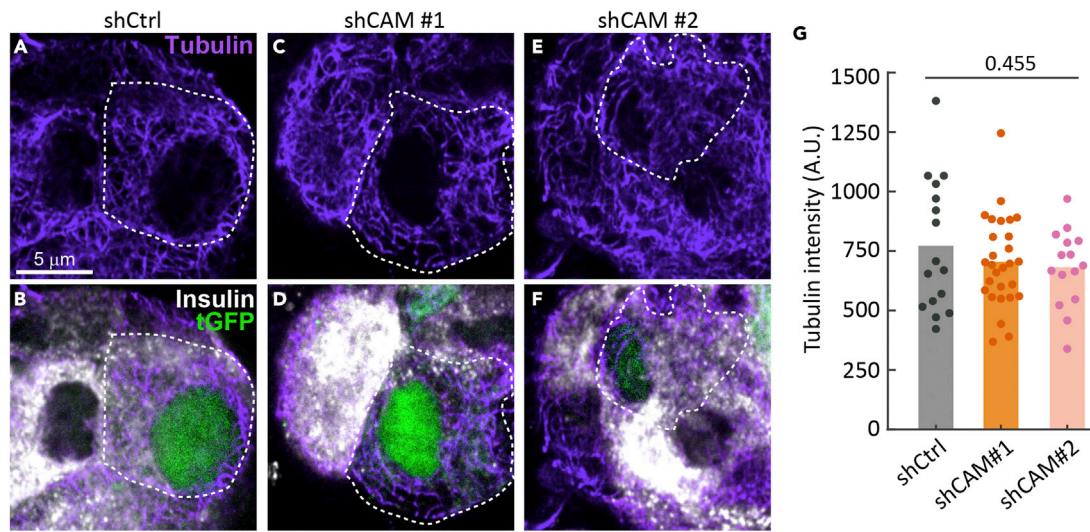


Figure 2. Knockdown of CAMSAP2 does not alter the amount and dynamics of microtubules in primary β -cells

(A-F) Immunofluorescence staining of tubulin (magenta) and insulin (white) in CAMSAP2-KD and control β -cells in mouse islets. Green, tGFP to indicate successful transduction. Dashed lines delineate transduced β -cells.
(G) Quantification of tubulin immunofluorescence intensity of CAMSAP2-KD and control β -cells in mouse islets. A.U., Arbitrary unit. Islets were extracted with methanol, which removed cytosolic free tubulin and the measured tubulin intensity represents polymerized tubulin. Dots represent individual β -cells. Bars represent the average of three animals (Dunnett's multiple comparisons test).
(H) Quantification of deetyrosinated tubulin immunofluorescence intensity in CAMSAP2-KD and control β -cells in mouse islets. Dots represent individual β -cells. Bars represent the average of three animals. * $p < 0.05$, *** $p < 0.001$ (Dunnett's multiple comparisons test).
(I-T) Immunofluorescence staining of deetyrosinated tubulin (magenta), E-Cadherin (blue), and insulin (white) in CAMSAP2-KD and control β -cells in mouse islets incubated in media containing 2.8 or 11 mM glucose. Green, tGFP to indicate successful transduction. Dashed lines delineate transduced β -cells.

localization of CAMSAP2 in β -cells. MIN6 and INS1 cells are transformed β -cells with detectable but compromised GSIS. In interphase MIN6 cells, CAMSAP2 forms short stretches or puncta at microtubule ends (Figures S2A-S2D). In dividing MIN6 cells at telophase, we observed the CAMSAP2 signal at the distal ends of the mid-body, confirming the binding of CAMSAP2 to microtubule minus ends in MIN6 cells (Figure S2E). In INS-1 cells, we observed a similar pattern of CAMSAP2 puncta/stretchers in the cytoplasm (S2F-G). These results are consistent with previously reported CAMSAP2 localization in cultured non- β -cells.³⁸

Strikingly, we did not detect CAMSAP2 puncta/stretchers in the cytoplasm of primary β -cells. Instead, the CAMSAP2 signal overlaps with Golgi markers, such as GM130 and GCC185 (Figures 3A-3F). This localization is specific to CAMSAP2 in contrast to the other two CAMSAP paralogues, CAMSAP1 and CAMSAP3, which are scattered in the cytoplasm of primary β -cells (Figures S3A and S3B). To confirm specific immunodetection of CAMSAP2, we ectopically expressed EGFP-tagged CAMSAPs in MIN6 cells and compared the fluorescence signal with that from immunofluorescence staining. We observed strong co-localization of CAMSAP2 immunofluorescence signal with EGFP-CAMSAP2, but not with EGFP-CAMSAP1 or 3 (Figures S3C-S3E). These results, together with the reduced CAMSAP2 signals in CAMSAP2-KD cells, demonstrate the specificity of utilized antibodies.

Interestingly, the high level of CAMSAP2 at the Golgi is specific to primary β -cells but not to α -cells in the same islets (Figures 3G-3N). These results and the findings that CAMSAP2-KD compromises GSIS suggest that the localization of CAMSAP2 at the Golgi may be associated with its function to promote robust insulin secretion. Because CAMSAP2 exhibit different subcellular localization in primary β -cells and in MIN6 cells, we also tested whether CAMSAP2-KD in MIN6 affects GSIS. Unlike in primary β -cells, CAMSAP2-KD does not significantly impair GSIS in MIN6 (Figures 1D and S2H). Because CAMSAP2 in MIN6 cells localizes to microtubule minus ends, it is likely other minus end-binding proteins, such as CAMSAP3, may compensate for CAMSAP2-KD in MIN6 cells. It is noteworthy that MIN6 cells expressing one of the shCAMSAP2 constructs (#1) exhibit elevated basal secretion and significantly reduced fold change of stimulated over basal secretion when compared to control cells (Figure S2I). This phenotype is distinct from that in CAMSAP2-KD primary β -cells, which do not have altered basal secretion but significantly reduced stimulated secretion (Figure 1D). These different phenotypes of insulin secretion in CAMSAP2-KD primary β -cells and MIN6 cells support the hypothesis that the Golgi-localized CAMSAP2 in primary β -cells and microtubule minus end-binding CAMSAP2 in MIN6 may have differential functions.

Golgi association of CAMSAP2 in primary islet β -cells is microtubule-independent

We next asked why primary β -cells and immortalized β -cells have different CAMSAP2 localization. MIN6 and INS-1 do not have cell-cell interactions with other types of islet endocrine cells, which has been shown to influence β -cells secretion and cytoskeletal configuration.^{49,50} To test whether cell-cell interactions underlie the observed difference, we examined the subcellular localization of CAMSAP2 in primary β -cells dissociated from mouse islets. We found that CAMSAP2 still localizes to the Golgi in dissociated primary β -cells cultured for one day (Figures 3O-3Q). These results indicate that the unique localization of CAMSAP2 at the Golgi is intrinsic to primary β -cells.

The Golgi is the major microtubule nucleation site in primary β -cells.²⁷ We next tested whether the accumulation of CAMSAP2 at the Golgi represents highly organized microtubule minus ends anchored to the Golgi. To test this possibility, isolated islets were treated with nocodazole or low temperature to induce microtubule disassembly. We found that neither perturbation alters the preferential association of CAMSAP2 with the Golgi, although the absence of microtubules led to the disassembly of the Golgi

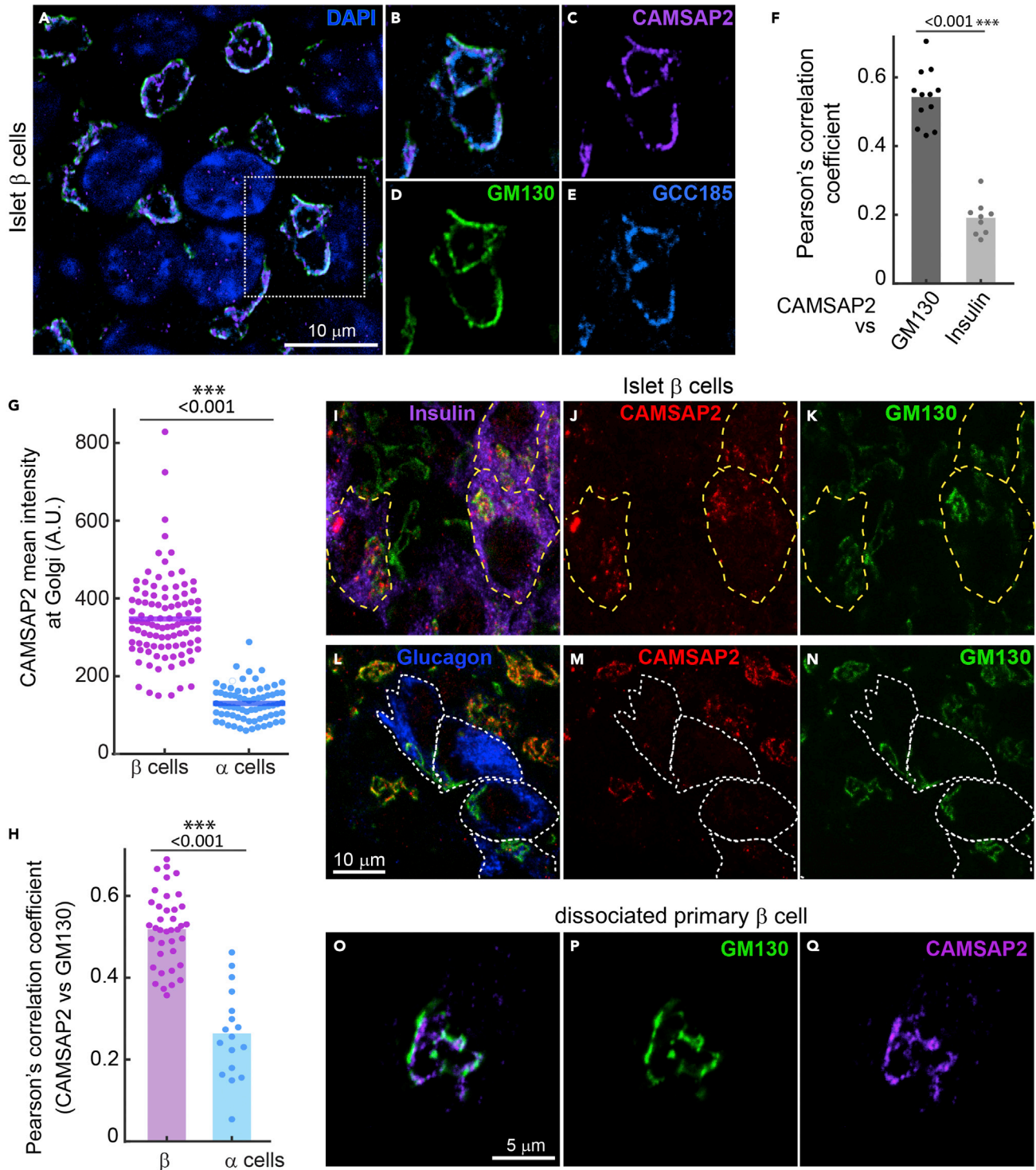


Figure 3. CAMSAP2 localizes to the Golgi in primary β-cells
(A-E) Immunofluorescence staining of GM130 (green), CAMSAP2 (magenta), and GCC185 (cyan) in primary β-cells in mouse islets. Blue, DAPI.
(F) Pearson's correlation coefficient of the immunofluorescence signal of CAMSAP2 with GM130 (dark gray) and CAMSAP2 with insulin (light gray). Dots represent individual images of mouse islets. Columns represent the average of three animals. *** $p < 0.01$ (t-test).
(G) Quantification of CAMSAP2 immunofluorescence intensity at the Golgi in mouse islets. Dots represent individual β-cells. Bars represent the average of three animals. *** $p < 0.001$ (t test). A.U., Arbitrary unit.
(H) Quantification of Pearson's correlation coefficient of CAMSAP2 with GM130 in mouse islets. Dots represent individual β-cells. Bars represent the average of three animals. *** $p < 0.001$ (t test).

Figure 3. Continued

(H) Pearson's correlation coefficient of the immunofluorescence signal of CAMSAP2 with GM130 in primary β - and α -cells in mouse islets. Dots represent individual islet cells. Columns represent the average of three animals. *** $p < 0.01$ (t test).

(I-N) Immunofluorescence staining of insulin (magenta), glucagon (blue), CAMSAP2 (red), and GM130 (green) in mouse islets.

(O-Q) Immunofluorescence staining of GM130 (green) and CAMSAP2 (magenta) in primary β -cells dissociated from mouse islets and cultured for one day.

into smaller mini-stacks (Figures 4A-4F, S4A, and S4B). The removal of nocodazole allows microtubules to emerge from the Golgi, but it also does not change the localization of CAMSAP2 (Figures 4N-O). These results indicate that the localization of CAMSAP2 at the Golgi is independent of microtubules. In contrast, treating islets with Brefeldin A, which disrupts the Golgi due to inhibited ER-to-Golgi anterograde trafficking,⁵¹ abolishes detectable CAMSAP2 localization to a particular organelle, while not affecting microtubules (Figures 4G-4I and 4P-4Q). When we removed Brefeldin A in the presence of nocodazole to allow the anterograde trafficking and Golgi re-formation without microtubules, CAMSAP2 accumulates at nascent Golgi mini-stacks at a level comparable to that of untreated islet β -cells (Figures 4J-4M). We examined the protein levels of CAMSAP2 in MIN6 and in primary β -cells and found no change after incubation with different glucose concentrations, nocodazole, or Brefeldin A (Figures S4C-S4E). These results suggest that the disappearance of accumulated CAMSAP2 at the Golgi after Brefeldin A treatment is not due to its degradation. Collectively, these results indicate that microtubules do not influence CAMSAP2 localization at the Golgi in primary β -cells and suggest a yet-unknown function for CAMSAP2 at this location.

CAMSAP2 facilitates the Golgi-to-endoplasmic reticulum trafficking

Our data show that CAMSAP2 in primary β -cells does not regulate microtubule stability and its localization to the Golgi is independent of microtubules. We, therefore, explored other mechanisms by which CAMSAP2 may regulate GSIS. We first examined the total insulin content of CAMSAP2-KD β -cells. The average total insulin in CAMSAP2-KD β -cells is significantly lower than that in control β -cells (Figures S5A-S5H). In contrast, no difference in the level of proinsulin was observed amongst these cells (Figures S5I-S5P). These data suggest that CAMSAP2-KD impairs insulin biosynthesis after the stage of translation and production of proinsulin.

The conversion of proinsulin to insulin depends on ER-Golgi trafficking. Because CAMSAP2 is enriched at the Golgi in primary β -cells, we hypothesized that CAMSAP2 may regulate trafficking between the Golgi and ER. To test this hypothesis, we used Brefeldin A to inhibit the ER-to-Golgi anterograde trafficking. This is expected to reduce Golgi volume over time because of the constant retrograde transport and fusion of Golgi-derived vesicles to the ER. The *cis*-Golgi matrix protein, GM130, is not transported to the ER as the Golgi collapses and is released into the cytoplasm.⁵² We followed the level of GM130 in the cytoplasm as a readout of the rate of Golgi-to-ER retrograde trafficking in the presence of Brefeldin A (Figures 5A and 5B). The removal of Brefeldin A restores anterograde trafficking, which allows the incorporation of GM130 to the newly assembled Golgi and the reduction of its level in the cytoplasm (Figure 5C). Therefore, monitoring the disappearance of GM130 from the cytoplasm after Brefeldin A removal will reveal the rate of anterograde trafficking.

In the presence of Brefeldin A, the increase of cytoplasmic GM130 intensity in CAMSAP2-KD cells is significantly slower than that in control β -cells, suggesting a defective retrograde transport when CAMSAP2 level is reduced (Figures 5D-5J). When Brefeldin A is removed, CAMSAP2-KD β -cells exhibit a slower reduction of cytoplasmic GM130 intensity compared to that in control β -cells, suggesting that CAMSAP2-KD compromises efficient anterograde ER-to-Golgi trafficking (Figures 5K-5M).

We next examined the level of β -COP, a subunit of COPI, which mediates Golgi to ER retrograde as well as anterograde transport in CAMSAP2-KD cells. Compared to controls, primary β -cells expressing CAMSAP2-targeting shRNAs have significantly increased β -COP (Figures 5N-5Q). These results are consistent with our model that CAMSAP2-KD impairs the trafficking between Golgi and ER leading to the accumulation of β -COP. Together, these results suggest that CAMSAP2-KD compromises the trafficking between the Golgi and ER, which likely contributes to the reduced insulin content without changing proinsulin levels.

Primary β -cells produce different isoforms of CAMSAP2 compared to MIN6 cells

We last examined why CAMSAP2 localizes to the Golgi in primary islet β -cells but at microtubule minus ends in insulinoma cells. When compared to MIN6 cells, CAMSAP2 from primary β -cells exhibits increased electrophoretic mobility on an acrylamide gel with an estimated molecular weight difference of around 10.6 kDa

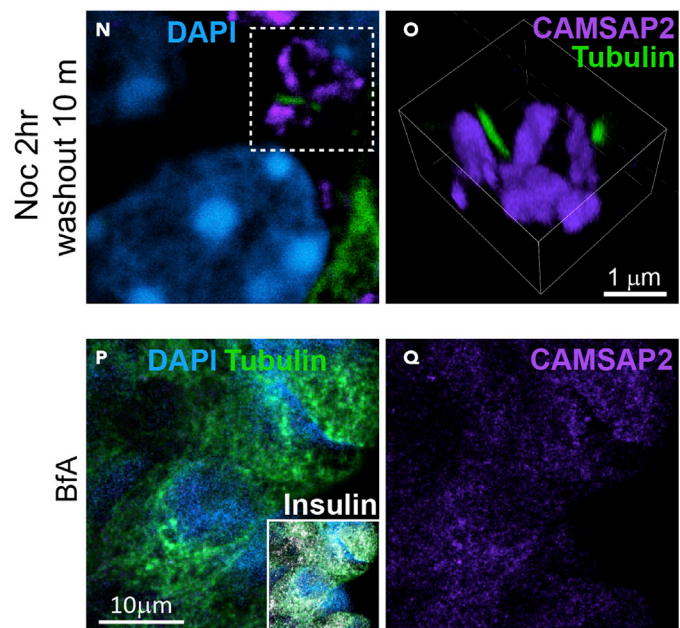
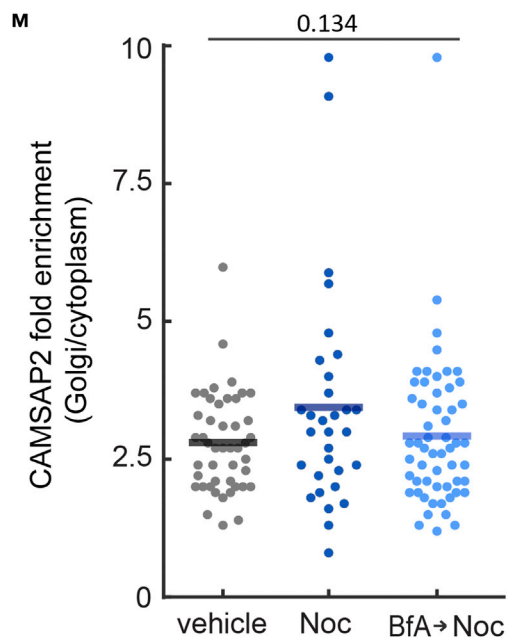
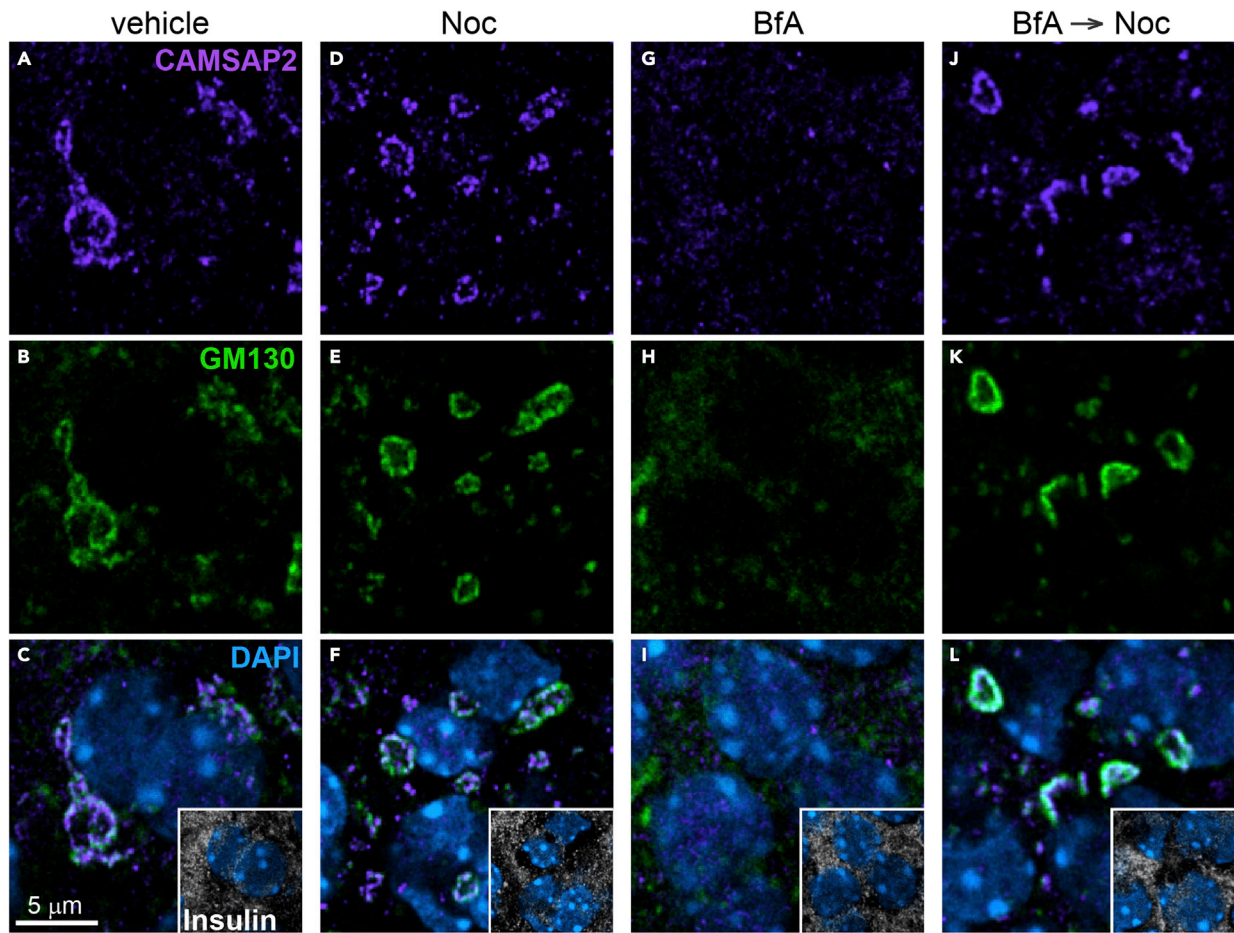


Figure 4. The localization of CAMSAP2 to the Golgi in primary β -cells is independent of binding to microtubules

(A-L) Immunofluorescence staining of GM130 (green), CAMSAP2 (magenta), and insulin (white) in mouse islets treated with nocodazole (D-F), Brefeldin A [BfA] (G-I), or BfA and then removed BfA in the presence of nocodazole (J-L). Cyan, DAPI.

(M) Quantification of CAMSAP2 immunofluorescence intensity at the Golgi over that in the cytosol in primary β -cells in mouse islets. Dots represent individual β -cells. Columns represent the average of three animals (One-way ANOVA).

(N and O) Immunofluorescence staining of CAMSAP2 (magenta) and tubulin (green) in mouse islets. Islets treated with nocodazole were then incubated in media without nocodazole to allow microtubule growth. Panel O is a 3D reconstituted image of Golgi mini-stacks. Cyan, DAPI.

(P and Q) Immunofluorescence staining of tubulin (green), CAMSAP2 (magenta), and insulin (white) in mouse islets treated with Brefeldin A. Cyan, DAPI.

(Figure 6A). These results suggest that primary β -cells may express a different CAMSAP2 variant or have different post-translational modifications on CAMSAP2 protein, which confers its unique localization to the Golgi.

To distinguish these possibilities, we ectopically expressed an EGFP-tagged human CAMSAP2 in mouse primary islet β -cells. We used a full-length cDNA that is the longest one reported in GenBank (NM001297707.3, human CAMSAP2 transcript variant 1). It differs from other reported full-length cDNAs with two extra exons coding for 11 and 9 amino acid residues (residues 216-226 and 381-396 out of the 1489 amino acids in the CAMSAP2 protein), respectively. The homolog of this protein is reported in mice (NP001334038.1). The ectopically expressed EGFP-CAMSAP2 forms small stretches in the cytoplasm of primary islet β -cells, suggesting that it localizes to microtubule minus ends (Figure 6B). A large portion of these CAMSAP2 stretches localizes to the cell periphery instead of near the cell center, suggesting that mouse primary β -cells have abundant microtubule minus ends in the cell periphery. To further examine the subcellular localization of EGFP-CAMSAP2, we stained for GM130 in mouse primary β -cells. We found that a minor population of CAMSAP2 stretches is associated with the Golgi, representing microtubule minus ends anchored to it (Figures 6C-6F). The localization and pattern of the human CAMSAP2 variant 1 expressed in mouse primary β -cells are very similar to that of endogenous CAMSAP2 in MIN6 cells (S2A-D). Because the expressed EGFP-CAMSAP2 from the human variant 1 cDNA does not localize to the Golgi, we conclude that mouse primary β -cells do not post-translationally modify CAMSAP2 protein to alter its subcellular localization from microtubule minus ends to the Golgi. These data favor the hypothesis that the different electrophoretic mobility between the endogenous CAMSAP2 in mouse primary β -cells and in MIN6 is likely due to alternatively splicing, which may confer the differential subcellular localization of CAMSAP2.

The linear structure of CAMSAP2 includes an N-terminal CH (calponin homology) domain and a C-terminal CKK (The C-terminal domain common to CAMSAP1, KIAA1078, and KIAA1543) domain. The CKK domain is essential for the recognition and binding to microtubule minus ends.³⁵ We hypothesized that the putatively longer MIN6 CAMSAP2 variant (Compared to the variant in primary β -cells) may have extra residues that allow its localization to microtubule minus ends but not being recruited to the Golgi. Although the removal of the C-terminal 683 residues, including the entire CKK domain, compromises its localization to microtubule minus ends, the truncated protein is not specifically recruited to the Golgi in MIN6 cells (Figures 6G-6M). The function of the CH domain is not fully understood, but the human CAMSAP2 is reported to have N-terminal alternatively spliced variants.^{38,53} We tested an N-terminal truncation missing 292 residues, without the majority of the CH domain, but it still localizes to microtubule ends and is not restricted to the Golgi (Figures 6G and 6N-6O). Unfortunately, we are not able to obtain the full-length cDNA of CAMSAP2 from mouse primary β -cells, likely due to the rareness and expected long length (>8 kb) of its message that lessens the success of reverse transcription, PCR, and RACE (rapid amplification of cDNA ends). Nevertheless, our data indicate the likely existence of novel CAMSAP2 cDNA isoform(s) in primary islet β -cells that localize to the Golgi and have a function in the ER-Golgi trafficking.

DISCUSSION

Regulated microtubule polymerization, disassembly, and anchorage are essential for the proper function of many cells. CAMSAP2 was shown to regulate cell morphology and organelle distribution through modulating microtubule organization and anchorage in specialized cell types such as neurons, glial, and polarized epithelia cells.^{40,43,54} Here, we report a novel function of CAMSAP2 in primary β -cells in facilitating the trafficking between the Golgi and ER and therefore promoting efficient insulin vesicle biogenesis and robust GSIS. Our studies, in combination with the established association between CAMSAP2 polymorphism and type 1 diabetes⁴⁵ underscore the unique roles of this protein in pancreatic islet β -cell function. Our findings also establish an example wherein a well-established microtubule-associated protein localizes

Figure 5. Knockdown of CAMSAP2 impairs the ER-to-Golgi trafficking in primary islet β -cells

(A-C) Schematic of the dynamics of the Golgi and GM130 (magenta) localization before (A), after Brefeldin A addition (B), and after its removal (C). (D-I) Immunofluorescence staining of GM130 (magenta) in Brefeldin A (BfA)-treated islets β -cells expressing shRNAs. Dashed lines delineate transduced cells. Cyan, *Ins2* promoter-driven H2B-mApple to label β -cells in islets; Green, tGFP to indicate successful transduction. (J-K) Quantification of cytoplasmic GM130 immunofluorescence intensity of transduced β -cells in mouse islets treated with Brefeldin A (J) and after Brefeldin A removal (K). The average GM130 intensity of the entire cell was used to normalize data across different time points. Dots represent the average of three animals. Bars represent SD. ** $p < 0.01$, *** $p < 0.001$ (Dunnett's multiple comparisons test). A.U., Arbitrary unit. (L and M) Immunofluorescence staining of GM130 (magenta) in BfA-treated islets β -cells expressing shRNAs incubated in media without BfA for 90 min. Dashed lines delineate transduced cells. Cyan, *Ins2* promoter-driven H2B-mApple to label β -cells in islets; Green, tGFP to indicate successful transduction. (N) Quantification of β -COP immunofluorescence intensity of CAMSAP2-KD and control β -cells in mouse islets. Dots represent individual β -cells. Bars represent the average of three animals. *** $p < 0.001$ (Dunnett's multiple comparisons test). (O-Q) Immunofluorescence staining of β -COP (magenta) in islets β -cells expressing shRNAs. Dashed lines delineate transduced cells. Blue, DAPI; green, tGFP to indicate successful transduction.

to a previously unreported subcellular localization primarily through microtubule-independent mechanisms to regulate cellular functions.

The function of CAMSAP in stabilizing microtubules, which has been studied in detail in cells and *in vitro*, are based on its recruitment to uncapped microtubule minus ends.^{36,53} CAMSAP2 and CAMSAP3 bind to elongating minus ends and form short stabilized microtubule stretches.⁴⁰ These CAMSAP-coated regions promotes the positioning of minus ends to specific cellular structures including the Golgi,⁴⁴ apical membrane in epithelial cells,⁴¹ and glial varicosities.⁵⁴ To date, all cellular functions of CAMSAP2 have been attributed to microtubule stabilization and anchorage. Accordingly, we initially propose that CAMSAP2 may contribute to the regulation of β -cell function through modulating microtubule.

In contrast to our hypothesis, we found that CAMSAP2 does not noticeably regulate microtubule stability and abundance in primary β -cells. This finding might have been a manifestation of the redundancy of CAMSAP2 function, because in some specialized cells, CAMSAPs act redundantly with their paralogues or other microtubule stabilizing and anchoring proteins.^{55,56} Despite undisrupted microtubule network, we have determined that CAMSAP2-KD in primary β -cells compromises the trafficking between the Golgi and ER, which results in reduced total insulin content and impaired GSIS. This indicates that CAMSAP2 in primary β -cells has a unique function that cannot be compensated by its paralogues. Corresponding to this novel function and unlike other CAMSAP paralogues, CAMSAP2 in primary β -cells is largely restricted to the Golgi and this localization is independent of the presence of microtubules.

Intriguingly, the Golgi-localization of CAMSAP2 is only observed in primary β -cells, but not in MIN6, INS1, or non- β islet cells. The underlying reason for this difference is not known but it is likely related to the functional status of these cells. MIN6 and INS1 are immortalized insulinoma cells. They maintain detectable but relatively low levels of GSIS compared to primary β -cells^{57,58} and may not have efficient insulin vesicle biogenesis. While primary non- β islet cells have glucose-regulated hormone secretion, their response to glucose differs from β -cells (e.g., glucagon secretion is inhibited by high glucose). In addition, non- β islet cells do not experience the constant ER stress as β -cells caused by a high level of insulin mRNA translation and unfolded/misfolded proinsulin.^{59,60} Thus, it will be interesting in the future to examine whether the different CAMSAP2 localization is a cause for compromised insulin secretion in MIN6/INS1 cells and a factor that determines the biosynthesis and secretion properties of glucagon in α -cells.

We found that the knockdown of CAMSAP2 compromises both retrograde and anterograde Golgi-ER trafficking, which likely facilitates insulin biosynthesis at the stage of proinsulin transition to the Golgi. Because the anterograde transport requires tight coordination with the retrograde trafficking from the Golgi to the ER, CAMSAP2 localization to the Golgi supports the model that it facilitates ER-to-Golgi trafficking. Rab GTPases and their regulatory guanine nucleotide exchange factors (GEF) are essential for the targeting of vesicles in the Golgi-to-ER trafficking.⁶¹ Two large scale studies using affinity purification and proximity-dependent biotinylation respectively have identified several Rab GEF as CAMSAP2-interaction partners, including DENND1A, DENND4C,⁶² and DENND6A.⁶³ We speculate that CAMSAP2 may facilitate the localization and/or functions of these GEFs at the Golgi to promote efficient Golgi-to-ER trafficking in primary β -cells, which can be explored in future studies.

Consistent with our model, we also found that the knockdown of CAMSAP2 in mouse primary β -cells increases the level of β -COP in the cytoplasm, suggesting a slower ER-Golgi trafficking. Importantly,

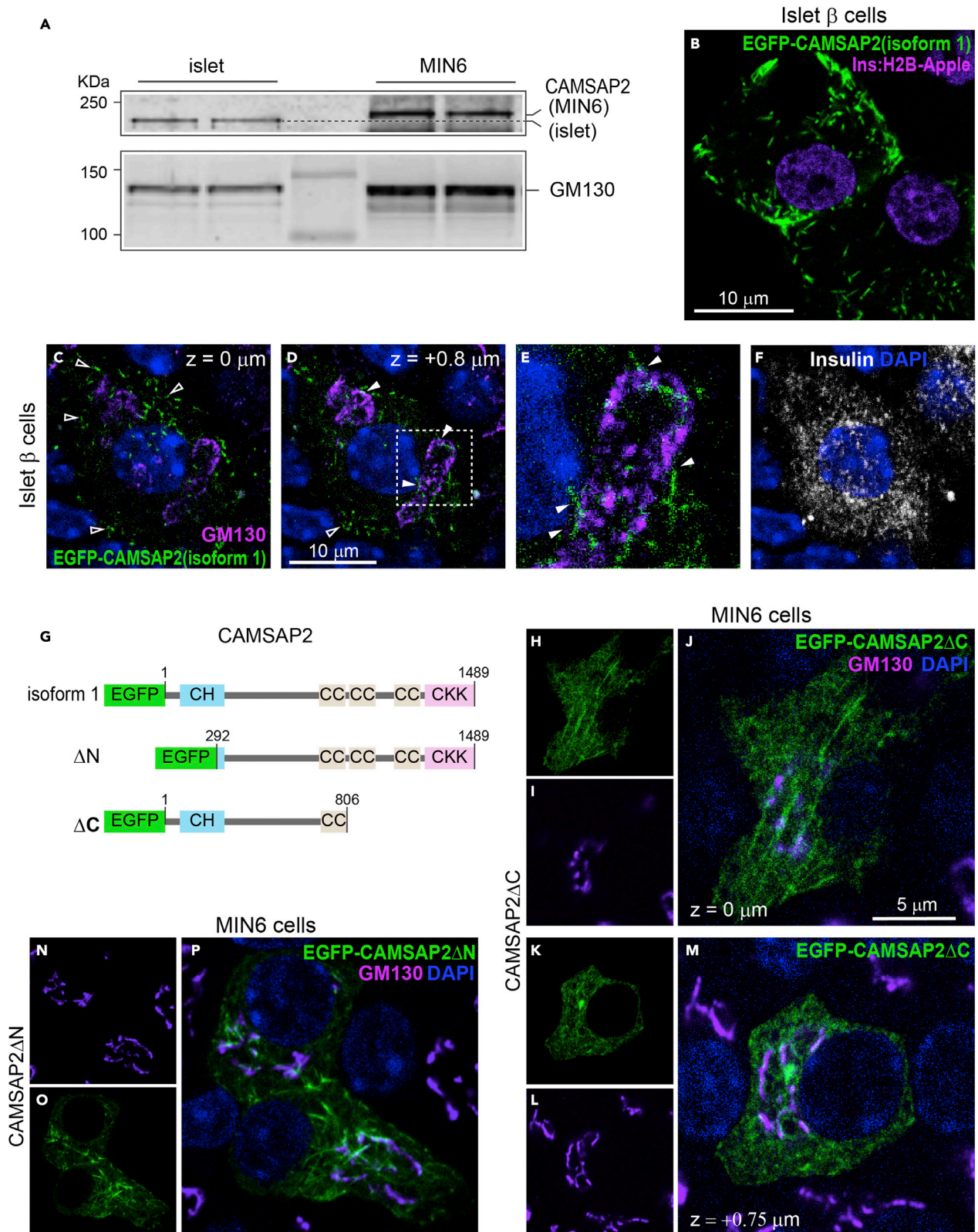


Figure 6. Primary islet β -cells express a different CAMSAP2 isoform compared to MIN6 cells

(A) Immunoblotting against CAMSAP2 and GM130 in lysates from islets and from MIN6. The dashed line indicates the position of CAMSAP2 from islet lysates.

(B) A fluorescence image of ectopically expressed EGFP-CAMSAP2 (human CAMSAP2 isoform 1, green) in mouse islets. Magenta, *Ins2* promoter-driven H2B-mApple to label β -cells in islets.

(C-F) Immunofluorescence staining of GM130 (magenta) and insulin (white) in primary β -cells expressing EGFP-CAMSAP2 (human CAMSAP2 isoform 1, green) in mouse islets. Blue, DAPI. Open arrow heads, CAMSAP2 stretches in cell periphery; closed arrow heads, CAMSAP2 stretches associated with the Golgi. Panel C and D are at the same x-y position and 0.8 μ m apart on the z axis.

(G) Schematic of the tested EGFP fusions of full-length or truncated CAMSAP2.

(H-P) Immunofluorescence images of GM130 (magenta) in MIN6 cells ectopically expressed truncated EGFP-CAMSAP2 (green). Blue, DAPI. Panel H-J and K-L are confocal images of the same cells at focal planes 0.75 μ m apart on the z axis.

COPI-mediated ER-Golgi trafficking depends on microtubules. Our data indicate that microtubules are not required for CAMSAP2 localization to the Golgi and CAMSAP2-KD does not significantly alter the amount and dynamics of total microtubules. However, because of the abundance of microtubules in β -cells, we may not be able to detect a significant reduction of microtubules even if CAMSAP2-KD destabilizes a sub-population of microtubules required for ER-Golgi trafficking. Therefore, we cannot rule out the possibility that CAMSAP2 may regulate the stability or positioning of a sub-population of microtubules between ER and Golgi required for trafficking. If this is the case, CAMSAP2 knockdown could compromise the Golgi-ER trafficking through both microtubule-independent and -dependent mechanisms.

Taking all findings together, we propose that CAMSAP2 is required for efficient insulin vesicle generation and secretion. Its knockdown impairs the Golgi-ER trafficking, reduce total insulin content and long-term β -cell functions. As a result, CAMSAP2-KD cells showed reduced secretion in response to glucose stimulation. This mechanism could contribute to the molecular connection between the reported CAMSAP2 polymorphism and human diabetes.⁴⁵ Additionally, in some differentiated cell types, CAMSAP loss-of-function did not cause significant microtubule phenotypes,⁵⁵ which may suggest unidentified microtubule-independent functions of CAMSAP in non- β -cells.

Limitations of the study

The underlying reason for the unique CAMSAP2 localization in primary β -cells is not known. Ectopically expressed human CAMSAP2 variant 1 cDNA construct in mouse primary β -cells forms stretches/puncta in the cytoplasm, suggesting this version of CAMSAP2 localizes to microtubule minus ends. These results also suggest that mouse primary β -cells do not post-translationally modify CAMSAP2 to alter its subcellular localization to the Golgi. Our western blot results show that CAMSAP2 from primary β -cells and from MIN6 cells have different electrophoretic mobility. Collectively, these data suggest that mouse primary β -cells may express an alternative CAMSAP2 transcript that is different from the reported human CAMSAP2 variant 1. Several versions of CAMSAP2 mRNAs have been reported in GenBank. Yet, testing the subcellular localization of truncated CAMSAP2 proteins could not identify the sequences that account for the Golgi-specific localization. Neither were we able to obtain the full-length CAMSAP2 cDNA from mouse primary islet cells. Thus, it remains possible that different mRNAs were transcribed or spliced to produce the Golgi-localized CAMSAP2 protein in primary β -cells. Although we favor this hypothesis, we cannot rule out the possibility that certain CAMSAP2-interacting proteins specifically expressed in primary β -cells may contribute to its localization to the Golgi. It is also unclear whether human primary islet β -cells express alternative CAMSAP2 variants localizing to the Golgi. A study compared human and mouse β -cells transcriptomes reports differential read coverage toward the 3' end of CAMSAP2, suggesting the CAMSAP2 transcript in primary β -cells may also be different in these two species.⁶⁴ While we don't yet understand the mechanisms underlying the unique localization of CAMSAP2 in primary β -cells, it is likely to support the specialized function of CAMSAP2 to facilitate insulin secretion. Pursuing such a possibility in the future will be important to fully understand how cells use different isoforms of CAMSAP2 to perform different functions.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.isci.2023.105938>.

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AUTHOR CONTRIBUTIONS

KHH, IK, and GG conceptualized the work and designed the study. KHH designed and conducted the majority of experiments. AJ assisted the immunofluorescence, microscopy, and analysis of the islet detyrosinated tubulin assay. MY, AN, and GG conducted RT-PCR and RACE. ABO and MAM generated the *Ins2^{H2B.Apple}* mice. KHH drafted the article. All authors contributed to writing the article.

DECLARATION OF INTERESTS

The authors declare no conflicts of interest.

INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a gender minority in their field of research.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit polyclonal anti- α -Tubulin	Abcam (Cambridge, UK)	Cat# ab18251; RRID: AB_2210057
Mouse monoclonal anti- α -Tubulin (clone DM1A)	MilliporeSigma, Burlington, MA	Cat# T9026; RRID: AB_477593
Rabbit polyclonal anti-Detyrosinated tubulin	MilliporeSigma	Cat# AB3201; RRID: AB_177350
Mouse monoclonal anti-E-Cadherin (clone 36)	BD Biosciences (San Jose, CA)	Cat# 610181; RRID: AB_397580
Mouse monoclonal anti-Glucagon (clone K79bB10)	MilliporeSigma	Cat# G2654; RRID: AB_259852
Guinea pig polyclonal anti-Insulin	Dako (Santa Clara, CA)	Cat# A0564; RRID: AB_10013624
Mouse monoclonal anti-GM130 (Clone 35)	BD Biosciences	Cat# 610823; RRID: AB_398142
Mouse monoclonal anti-Proinsulin	Developmental Studies Hybridoma Bank, Iowa City, IA	Cat# GS-9A8-C; RRID: AB_532383
Rabbit polyclonal anti-CAMSAP1	Novus Biologicals, Centennial, CO	Cat# NBP1-26645; RRID: AB_1852845
Rabbit polyclonal anti-CAMSAP2	Novus Biologicals	Cat# NBP1-21402; RRID: AB_1659977
Rabbit polyclonal anti-CAMSAP3	Thermo Fisher, Waltham, MA	Cat# PA5-48993; RRID: AB_2634449
Guinea pig polyclonal anti-GCC185	Covance research product Inc, Denver, PA	Cat# VU-140 (customized)
Mouse monoclonal anti- β -COP	MilliporeSigma	Cat# G6160; RRID: AB_477023
Chemicals, peptides, and recombinant proteins		
0.25% Trypsin-EDTA	Gibco (Waltham, MA)	Cat# 25200
Accumax	Innovative Cell Technologies (San Diego, CA)	Cat# AM-105
Brefeldin A	MilliporeSigma, Burlington, MA	Cat# B5936
CaCl ₂	MilliporeSigma	Cat# C1016
collagenase P	MilliporeSigma	Cat# C5138
cOmplete™ protease inhibitor cocktail	Roche (Basel, Switzerland)	Cat# 04 693 124 001
DAPI	Thermo Fisher, Waltham, MA)	Cat# 62247
DMEM, high glucose medium	Thermo Fisher	Cat# 11995-065
DMEM, no glucose medium	Thermo Fisher	Cat# 11966-025
DMSO	MilliporeSigma	Cat# D8418
Donor horse serum	Atlanta Biologicals (Flowery Branch, GA)	Cat# S12195
Fetal Bovine Serum	Atlanta Biologicals	Cat# S11550
Fibronectin	MilliporeSigma	Cat# F0162
HBSS with Ca ²⁺ & Mg ²⁺	Corning (Corning, NY)	Cat# 21-020-CV
HBSS without Ca ²⁺ & Mg ²⁺	Corning	Cat# 21-022-CV
HEPES	Corning	Cat# 25-060-CI
Human extracellular matrix	Corning	Cat# 354237
KCl	MilliporeSigma	Cat# PX1405
KH ₂ PO ₄	Thermo Fisher	Cat# BP363
Liraglutide	MedChemExpress (Monmouth Junction, NJ)	Cat# HY-P0014
LiCl	Thermo Fisher	Cat# L121-100
Methanol, anhydrous 99.8%	MilliporeSigma	Cat# 322415
β -Mercaptoethanol	MilliporeSigma	Cat# M7522
MgSO ₄	MilliporeSigma	Cat# M7506
NaCl	MilliporeSigma	Cat# S9625
NaHCO ₃	HyClone	Cat# SH30173.04
Nocodazole	MilliporeSigma	Cat# M1404
Opti-MEM® medium	Thermo Fisher	Cat# 31985-070

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Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Paraformaldehyde	MilliporeSigma	Cat# P6148
Penicillin-Streptomycin (10,000 U/mL)	Thermo Fisher	Cat# 15140-122
Precision Plus Protein™ Dual Color Standards	Bio-Rad (Hercules, CA)	Cat# 1610374
RPMI 1640 medium	Thermo Fisher	Cat# 11875-093
RPMI 1640, no glucose medium	Thermo Fisher	Cat# 11879-020
Saponin	MilliporeSigma	Cat# S4521
SDS	MilliporeSigma	Cat# L3771
TransIT®-Lenti	Mirus Bio (Madison, WI)	Cat# MIR 6606
Triton X-100	MilliporeSigma	Cat# T8787
Trizma®	MilliporeSigma	Cat# T1503
Tween® 20	MilliporeSigma	Cat# P9416
Vectashield mounting medium	Vector Laboratories (Burlingame, CA)	Cat# H-1000
Critical commercial assays		
Coomassie Plus (Bradford) Assay	Thermo Fisher	Cat# 23236
Mouse Ultrasensitive Insulin ELISA kit	ALPCO (Salem, NH)	Cat# 80-INSMSU
Deposited data		
Original western blot and microscopy images	Mendeley Data	https://data.mendeley.com/datasets/x69hns9ym/1
Experimental models: Cell lines		
INS1	Asfari et al. ⁵⁸	N/A
MIN6	Miyazaki et al. ⁵⁷	N/A
Experimental models: Organisms/strains		
CD-1 (ICR) mice	Charles River Laboratories (Wilmington, MA)	N/A
<i>Ins2^{H2B.Apple}</i> mice	Stancill et al. ⁶⁵	N/A
Oligonucleotides		
shCAM no #1 TRCN0000252418, 5'-TTGTCCGGCTAGAGGATATTT-3'	MilliporeSigma	Gene ID: 67886
shCAM no #2 TRCN0000252419, 5'-AGTTTCTCTGTCGGATTTAAA-3'	MilliporeSigma	Gene ID: 67886
shCtrl 5'-CCTAAGGTTAAGTCGCCCTCGCTC-3']	Addgene	1864
Recombinant DNA		
pEGFP-C1-CAMSAP1	This study	N/A
pEGFP-C1-CAMSAP2	This study	N/A
pEGFP-CAMSAP3	Jiang et al. ³⁸	N/A
PCDH-EGFP-CAMSAP2	This study	N/A
Software and algorithms		
Nikon Elements	Nikon	N/A
Prism 9	GraphPad	N/A
Other		
0.45 µm Cellulose Acetate Low Protein Binding filter	Corning	Cat# 430768
Amaxa nucleofector II	Lonza (Walkersville, MD)	N/A
Amicon Ultra-15 Centrifugal Filter, Ultracel-100 regenerated cellulose membrane	MilliporeSigma	Cat# UFC910024
GravityPLUS™ hanging drop 96-well microplate	PerkinElmer (Waltham, MA)	Cat# ISP-06-010

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Irina Kaverina irina.kaverina@vanderbilt.edu.

Materials availability

This study did not generate new unique reagents.

Data and code availability

- This paper does not report original code.
- This study did not generate new unique reagents, cell lines, or mouse lines.
- Original western blot and microscopy images have been deposited at Mendeley and are publicly available as of the date of publication. The DOI is listed in the [key resources table](#).
- Plasmids generated in this study and any additional information required to reanalyze the data reported in this paper are available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mouse models

All Mouse experimentation followed protocols approved by the Vanderbilt University Institutional Animal Care and Use Committee. Euthanasia follows the standard protocol set by the AALAC and Isoflurane inhalation was used. Mice were fed a regular chow diet *ad libitum* and maintained on a 12-12h dark-light cycle. Wild-type CD-1 (ICR) mice were from Charles River Laboratories (Wilmington, MA). *Ins2*^{H2B.Apple} mice have been described previously.⁶⁵

Cell lines and primary cell culture

MIN6⁶⁷ and INS1⁵⁸ cells were grown in DMEM with 25 mM glucose (Gibco), 0.071 mM β -mercaptoethanol (MilliporeSigma, St Louis, MO), 10% HI-FBS, and 100 U/mL penicillin-100 μ g/mL streptomycin. Primary β -cells were harvested from dissociation of isolated islets (see later in discussion in the islet isolation and in the pseudoislet generation procedure) and culture in islet media [RPMI-1640 media with 11 mM glucose (Gibco, Dublin, Ireland) plus 10% HI-FBS and 100 U/mL penicillin-100 μ g/mL streptomycin (Gibco)] in a glass-bottom 35-mm dish (MatTek, Ashland, MA) coated with human extracellular matrix [ECM] (Corning). Primary β -cells were cultured over one night before fixation for imaging.

METHOD DETAILS

Small molecules, chemicals, and treatment

Liraglutide was from MedChemExpress (Monmouth Junction, NJ). Nocodazole, Brefeldin A, DMSO, KCl and other chemicals were from MilliporeSigma. For nocodazole treatment, isolated islets were incubated in islet media plus 5 mg/mL nocodazole for 2 h. For Brefeldin A treatment, isolated islets were incubated in islet media plus 10 μ g/mL Brefeldin A for 2 h. To washout Brefeldin A, treated islets were transferred to islet media plus 5 mg/mL nocodazole three times and incubated for 2 h.

Plasmids, shRNA sequence, and lentivirus production

The CAMSAP2-targeting shRNA (shCAM) no #1 [TRCN0000252418, 5'-TTGTCGGCTAGAGGATATTT-3'] and no #2 [TRCN0000252419, 5'-AGTTTCTGTCCGATTTAAA-3'] are in the plasmid pLKO.1-CMV-tGFP and are from MilliporeSigma. The non-targeting shRNA control (shCtrl) [5'-CCTAAGGTTAAGTCGCCCTCGCTC-3'] plasmid, pLKO.1-shCtrl-puro, is from Addgene (cat. # 1864). For expression of EGFP-CAMSAP fusion proteins, pEGFP-C1-CAMSAP2 and -CAMSAP1 plasmids were kind gifts from Dr. Stephen Norris (Calico Life Sciences) and pEGFP-CAMSAP3 was a kind gift from Dr. Anna Akhmanova (Utrecht University). For lentiviral packaging of EGFP-CAMSAP2, its cDNA was amplified with high-fidelity PCR from pEGFP-C1-CAMSAP2 and cloned into the multi-cloning site of pCDH-EF1 (Addgene, #72266). For EGFP fusion proteins with CAMSAP2 truncation, corresponding PCR fragments were obtained and ligated into the multi-cloning sites of pCDH-EF1 using the assembly approach. For lentivirus production, the package plasmid pMDL, pRSV, pVSV-G and the transfer plasmid were mixed in a 4:1:1:6 ratio and

incubated with TransIT-Lenti reagent (Mirus Bio, Madison, WI) in Opti-MEM (Gibco). The assembled transfection complex was added to HEK293 cells at 80-95% confluency and incubated for 48 h. The harvested supernatant was precleared by filtering through a 0.45 μm -pore cellulose acetate membrane (Corning) and concentrated using Amicon Ultra-15 centrifugal filters (MilliporeSigma).

Islet isolation and culture

Islets were isolated from 8- to 12-week-old mice followed the published procedure.⁶⁶ Briefly, pancreas was perfused through the common bile duct with 2 mL of 0.8 mg/mL collagenase P (Roche, Indianapolis, IN) in Hanks' balanced salt solution [HBSS] with Ca^{2+} and Mg^{2+} (Corning, Corning, NY). The dilated pancreas was digested at 37°C for 20 min. The homogenate was washed with HBSS plus 10% heat-inactivated Fetal Bovine serum [HI-FBS] (Atlanta Biologicals, Flowery Branch, GA) four times and islets were handpicked into islet media (details see above in [cell lines and primary cell culture](#)) and cultured at 37°C with 5% CO₂.

Islet dissociation, pseudoislets generation and transduction

To dissociate islets to primary cells, isolated mouse islets were recovered in islet media overnight, washed with HBSS without Ca^{2+} and Mg^{2+} for three times and incubated for 20 min, digested with Accumax (Innovative Cell Technologies, San Diego, CA) at 30°C for 10 min, and gently pipetted up and down for 10 times to dissociate cells. Dissociated islet cells were not sorted and the ratio of β and non- β endocrine cells were maintain as that in mouse islets. The cell suspension was spun at 1000g for 3 min, re-suspended in islet media, and mixed with concentrated lentivirus and 8 $\mu\text{g}/\text{mL}$ DEAE-Dextran hydrochloride (MilliporeSigma). The mixture was loaded to the GravityPLUS hanging drop 96-well microplate (InSphero, Schlieren, Switzerland) and incubated for 4 days to generate pseudoislets. Pseudoislets were transferred to an ECM-coated 96-well plate and cultured for 1 day before analyses. For total pseudoislet volume calculation, two perpendicular diameters of each pseudoislet were measured and its volume was calculated using the equation $4/3\pi*r1*r2*(r1+r2)/2$, where $r1$ and $r2$ represent the two radii calculated from the measured diameters. The volume is converted to IEQ, which is defined as the volume of a spherical islet with a diameter of 150 μm and is equal to 1,767,146 μm^3 .⁶⁸

GSIS assay

Pseudoislets were washed and conditioned in KRB buffer (111 mM NaCl, 4.8 mM KCl, 1.2 mM MgSO_4 , and 1.2 mM KH_2PO_4 , 2.3 mM CaCl_2 , 25 mM NaHCO_3 , 10 mM HEPES and 0.2% BSA) plus 2.8 mM glucose (KRB-G2.8) for 1 h and incubated with fresh KRB-G2.8 for 1 h to assess basal secretion. After that, pseudoislets were incubated with KRB buffer plus 16.5 mM glucose (KRB-G16.5) for 45 min to assess glucose-stimulated secretion. Pseudoislets were then incubated with KRB-G16.5 plus 100 nM Liraglutide for 45 min, washed and conditioned in KRB-G2.8 for 45 min, and stimulated with KRB-G2.8 plus 30 mM KCl for 30 min. For GSIS in MIN6, cells were conditioned in KRB-G2.8 for 2 h, incubated with fresh KRB-G2.8 for 45 min and stimulated with KRB buffer plus 25 mM glucose for 45 min. Secreted insulin was quantified using ELISA kits from ALPCO (Salem, NH).

Immunofluorescence staining, microscopy, and quantification

For CAMSAP2 detection, samples were fixed with methanol at -20°C for 10 min. For the co-staining of CAMSAP2 with insulin or glucagon, samples were briefly fixed with 4% paraformaldehyde plus 0.1% saponin (MilliporeSigma) for 5 min and then fixed with methanol at -20°C for 10 min. For all other antigens, samples were fixed with 4% paraformaldehyde plus 0.1% saponin. Antibodies used are as follows: anti- α -tubulin (1:200, cat. #T9026, MilliporeSigma), anti-E-cadherin (1:250, #610181, BD Biosciences, San Jose, CA), CAMSAP1 (1:1000, #NBP1-26645, Novus Biologicals, Centennial, CO), CAMSAP2 (1:1000, #NBP1-21402, Novus Biologicals), CAMSAP3 (1:200 #PA5-48993, Thermo Fisher Scientific, Waltham, MA), anti-detyrosinated tubulin (1:250, #AB3201, MilliporeSigma), anti-glucagon (1:200, #G2654, MilliporeSigma), anti-GM130 (1:300, #610823, BD Biosciences), anti-insulin (1:500, #A0564, Dako, Santa Clara, CA), anti-proinsulin (1:30, #GS-9A8-C, Developmental Studies Hybridoma Bank, Iowa City, IA), anti-GCC185 (1:100, #VU-140 [customized], Covance research product Inc., Denver, PA), anti- β -COP (1:200, #G6160, MilliporeSigma). Secondary antibodies are from Invitrogen (1:200, Grand Island, NY). VECTASHIELD Mounting Medium was from Vector Labs (Burlingame, CA). All images, except in [Figures 1A](#) and [1B](#), were captured using Nikon Eclipse A1R laser scanning confocal microscope equipped with a CFI Apochromat TIRF 1003/1.45 oil objective. Images in [Figures 1A](#) and [1B](#) were captured using EVOS FL microscope equipped with an LPlan PH2 203/0.4 objective. Images were quantified using Nikon

NIS-Elements. The mean intensity (average fluorescence intensity per pixel after background subtraction) of the epitope in the region of interest (ROI) was measured. Background subtraction used the mean fluorescence intensity in a region without cells. For quantification of cytoplasmic GM130 immunofluorescence intensity in primary β -cell, a cell boarder ROI excluded the nucleus was first created to delineate a transduced β -cell on the tGFP (the transduction marker) channel. A second ROI was created on the GM130 channel to delineate GM130 positive area that is above a predetermined threshold that was kept consistent across cells expressing different shRNAs. Last, two small circular (measure) ROIs, 3 μm in diameter, were created between the first (cell boarder) and the second (Golgi) ROI on the tGFP channel. The average of the two measure ROIs on the GM130 channel was exported to represent the cytoplasmic GM130 intensity. For each β -cell, the average of three z-frames (+0.6 μm /-0.6 μm above and below the z-frame that shows the largest nuclear cross-section on the DAPI channel) was calculated for the statistical analysis.

Immunoblotting

Isolated mouse islets and MIN6 cells were lysed in Tris-lysis buffer (10 mM Tris-Cl pH 7.5, 100 mM NaCl, 1% Triton X-100, 10% glycerol, and cOmplete protease inhibitor cocktail [Roche, Basel, Switzerland]) on ice. For SDS-PAGE, 75 μg of total protein in the lysate was loaded per lane and was resolved in a 10% acrylamide gel. Antibodies used are as follows: anti-CAMSAP2 (1:500, #NBP1-21402, Novus Biologicals), anti- α -tubulin (1:1000, #ab18251, Abcam), anti-GM130 (1:200, #610823, BD Biosciences), IRDye 800 goat anti-rabbit IgG (1:5000, Rockland Immunochemicals, Pottstown, PA) and IRDye 700DX goat anti-mouse IgG (1:5000, LI-COR, Lincoln, NE). Blots were imaged using the Odyssey CLx imager (LI-COR).

QUANTIFICATION AND STATISTICAL ANALYSIS

All data represent three or more replicates of animals from separate experiments. Image quantification was performed in Nikon Elements and described in [method details](#) above. Statistical analyses were performed in GraphPad Prism. Student's *t* test or ANOVA followed by multiple comparisons tests were used and specified in figure legends, with $p < 0.05$ considered significant.