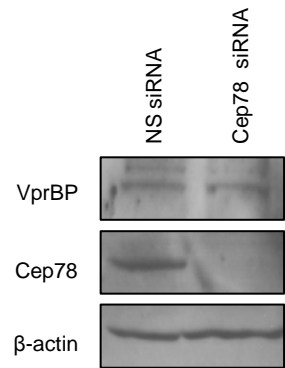


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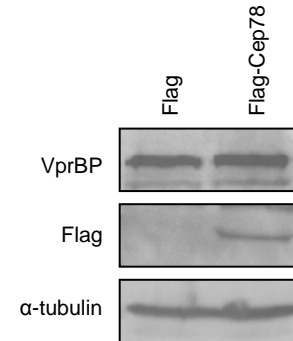
Table of Contents

Appendix Figure S1	pg 2
Appendix Figure S2	pg 3
Appendix Figure S3	pg 4
Appendix Figure S4	pg 5
Appendix Figure S5	pg 6
Appendix Figure S6	pg 7
Appendix Figure Legends	pg 8

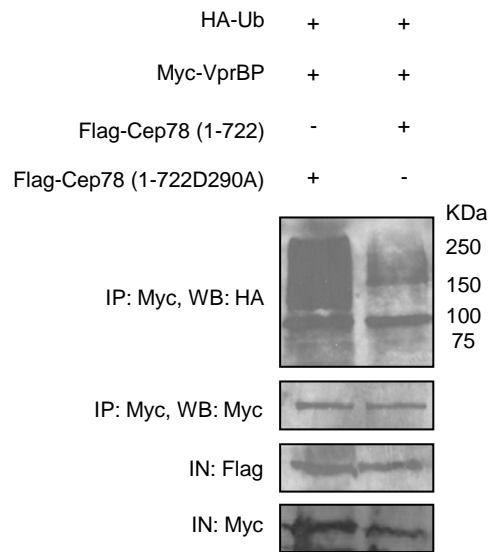
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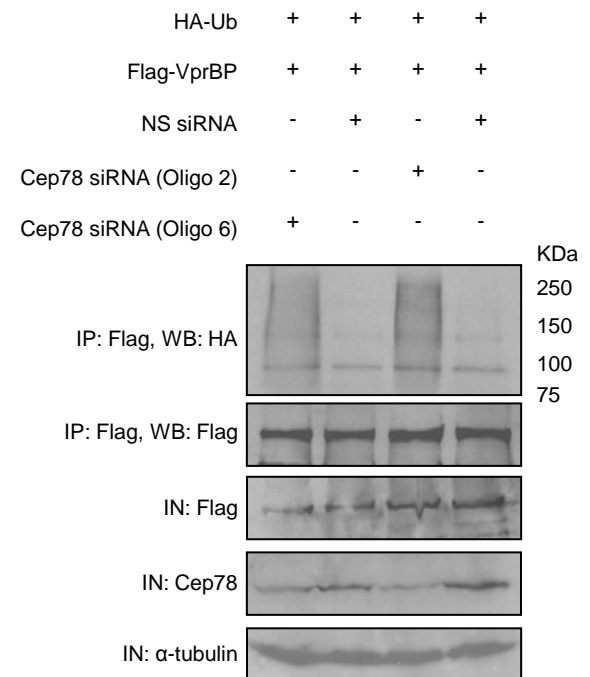
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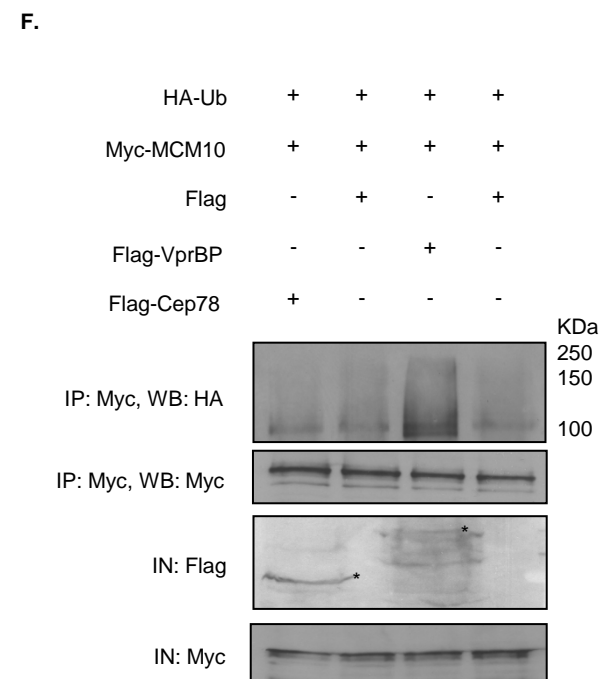
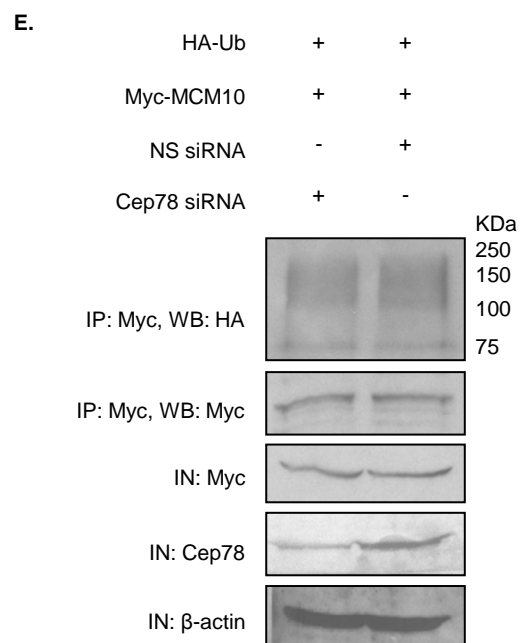
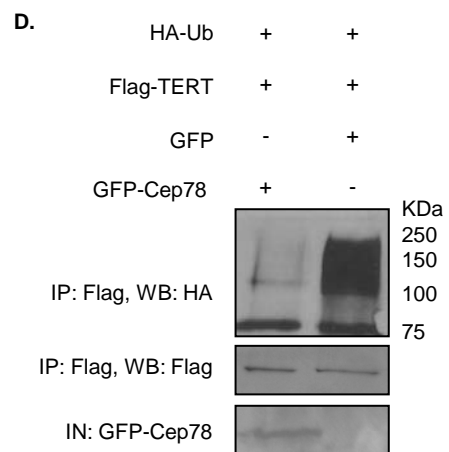
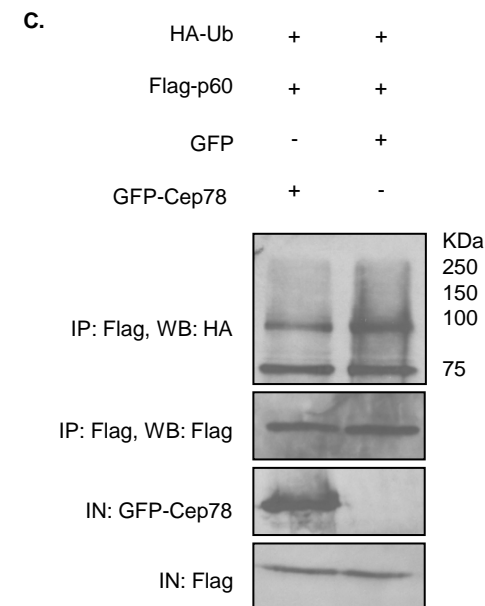
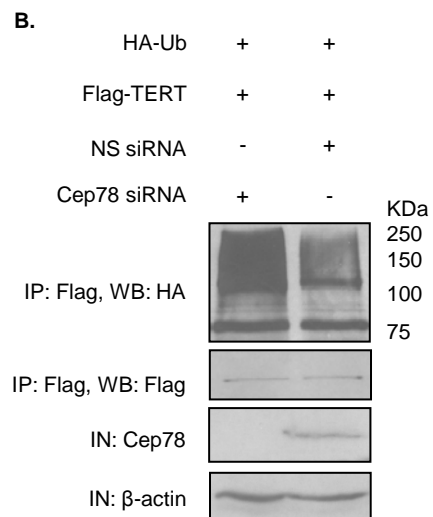
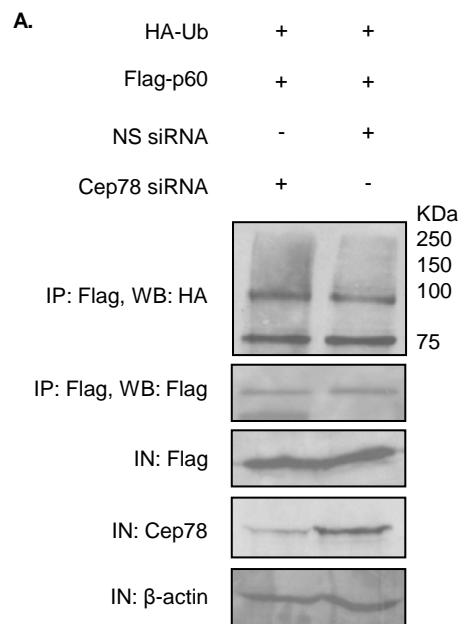
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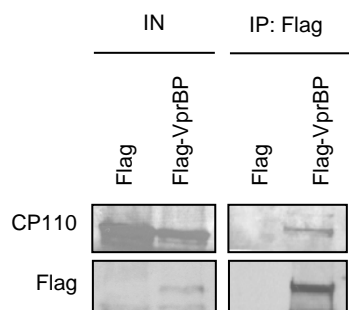
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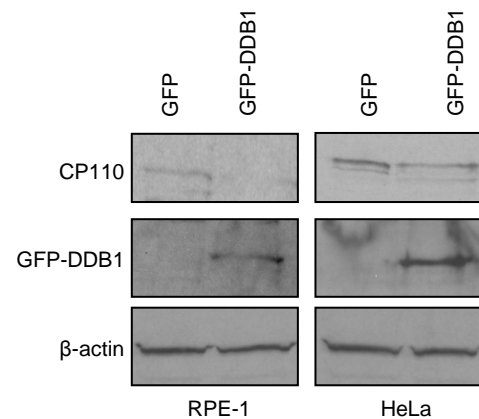
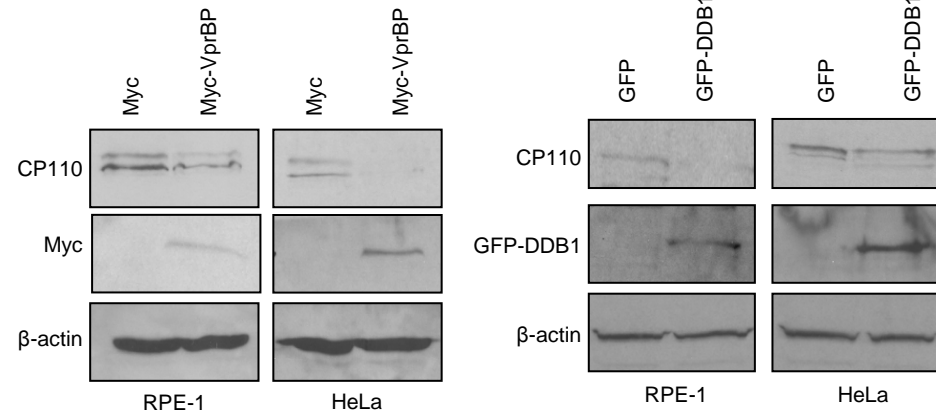
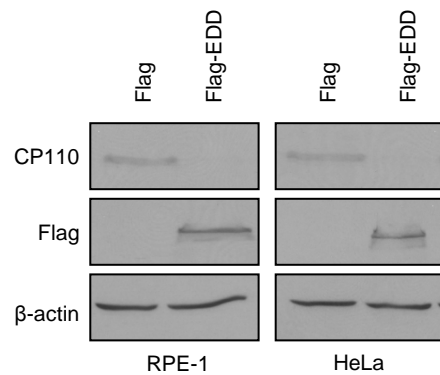
Appendix Figure S2



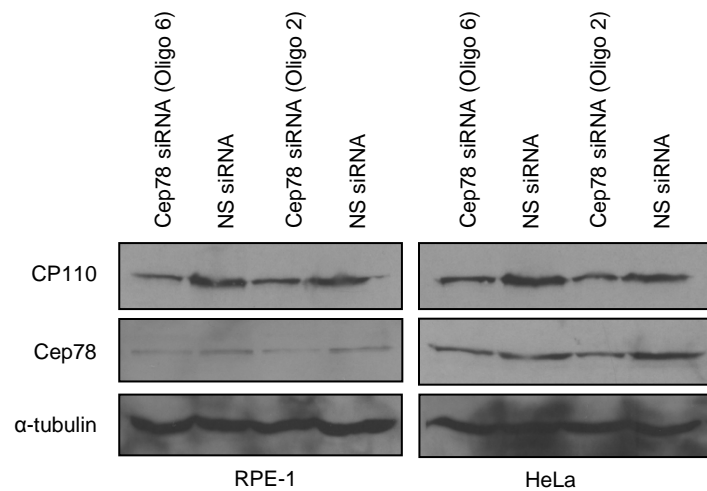
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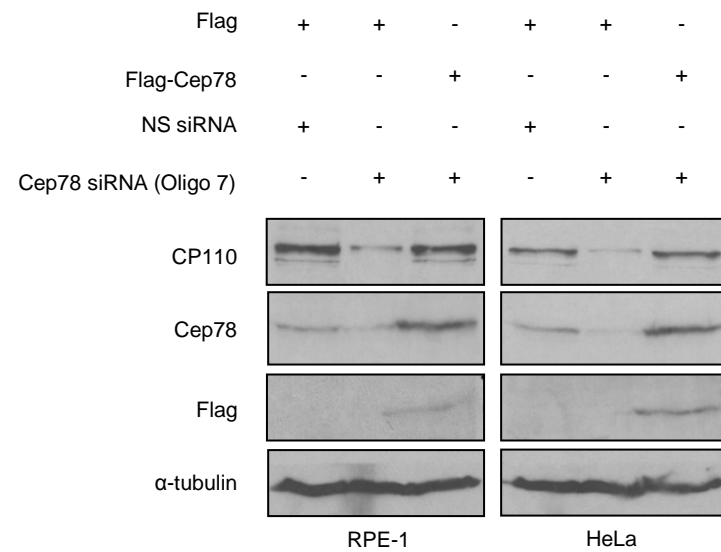
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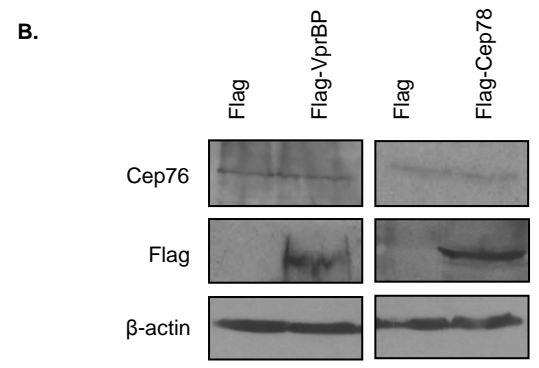
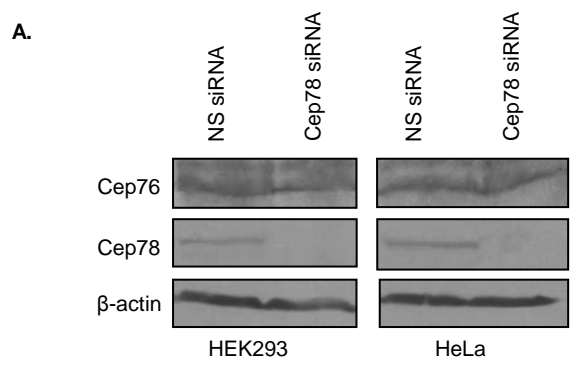


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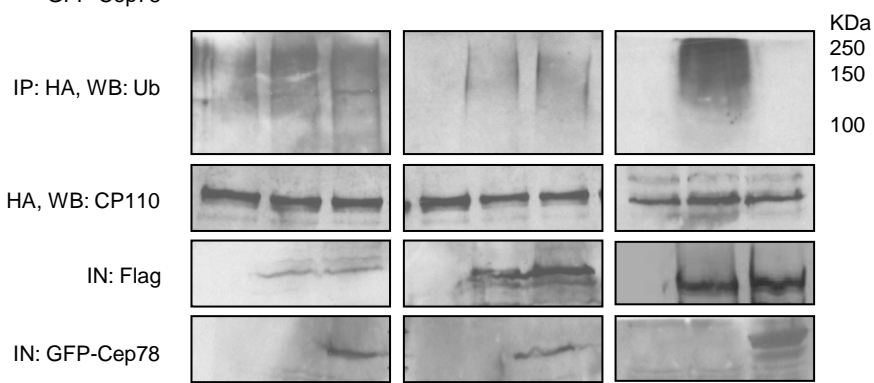
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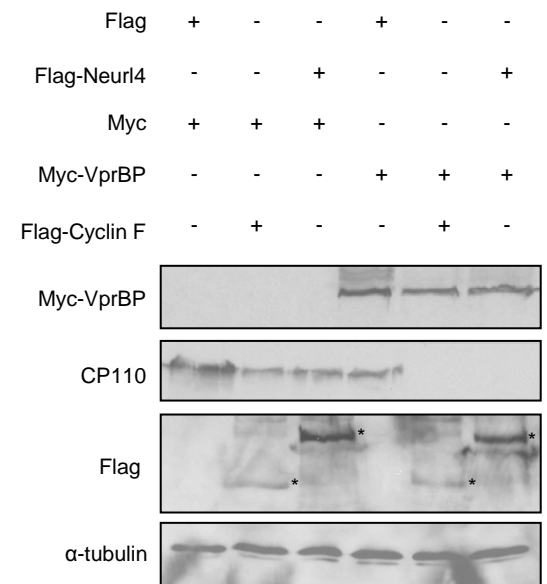


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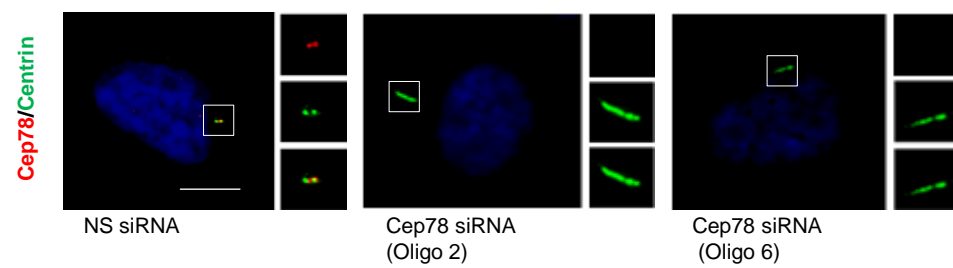
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Flag-VprBP	-	-	-	-	-	-	-	+	+
Flag-Cyclin F	-	+	+	-	-	-	-	-	-
HA-CP110	+	+	+	+	+	+	+	+	+
GFP	+	+	-	+	+	-	+	+	-
GFP-Cep78	-	-	+	-	-	+	-	-	+



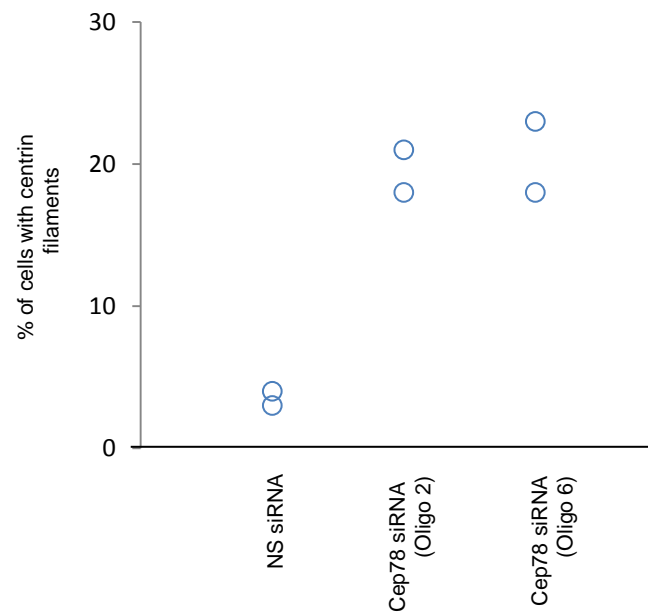
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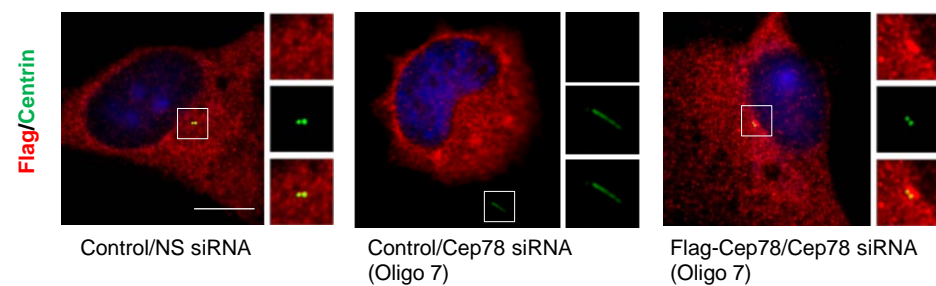
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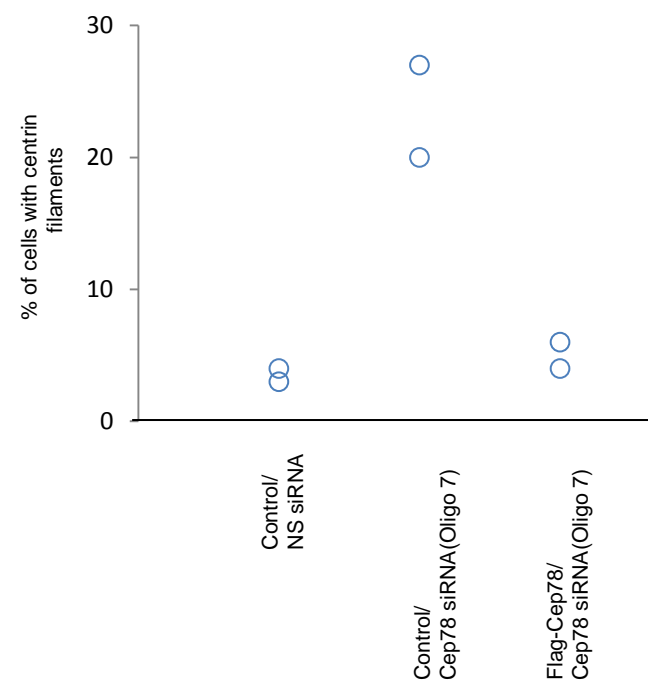
B.



C.



D.



Appendix Figure S1: Depletion or expression Cep78 had no effect on VprBP protein levels and Cep78 inhibits EDD-DYRK2-DDB1^{VprBP}

(A) HEK293 cells were transfected with NS siRNA or Cep78 siRNA. Lysates were Western blotted with the indicated antibodies. β -actin was used as a loading control. (B) Flag or Flag-Cep78 was expressed in HEK293 cells. Lysates were Western blotted with the indicated antibodies. α -tubulin was used as a loading control. (C) Flag-Cep78 wild type (1-722) or mutant (1-722D290A) was co-expressed with HA-Ub and Myc-VprBP in HEK293 cells. Lysates were immunoprecipitated with an anti-Myc antibody without SDS and Western blotted with the indicated antibodies. IN, input. (D) HEK293 cells were transfected with NS siRNA or Cep78 siRNA (oligo 2 or oligo 6) and constructs expressing Flag-VprBP and HA-Ub. Lysates were immunoprecipitated with an anti-Flag antibody without SDS and Western blotted with the indicated antibodies. IN, input. α -tubulin was used as a loading control.

Appendix Figure S2: Cep78 modulates ubiquitination of two EDD-DYRK2-DDB1^{VprBP} substrates katanin p60 and TERT but not a CRL4^{VprBP} substrate MCM10

(A, B) HEK293 cells were transfected with NS siRNA or Cep78 siRNA and constructs expressing HA-Ub and Flag-*katanin p60* or Flag-*TERT*. Lysates were immunoprecipitated with an anti-Flag antibody in 1% SDS and Western blotted with the indicated antibodies. IN, input. β -actin was used as a loading control. (C, D) HEK293 cells were transfected with constructs expressing HA-Ub, Flag-*katanin p60* or Flag-*TERT*, and GFP or GFP-Cep78. Lysates were immunoprecipitated with an anti-Flag antibody in 1% SDS and Western blotted with the indicated antibodies. IN, input. (E) HEK293 cells were transfected with NS siRNA or Cep78 siRNA and constructs expressing HA-Ub and Myc-*MCM10*. Lysates were immunoprecipitated with an anti-Myc antibody in 1% SDS and Western blotted with the indicated antibodies. IN,

input. β -actin was used as a loading control. **(F)** HEK293 cells were transfected with constructs expressing HA-Ub, Myc-MCM10, and Flag, Flag-Cep78 or Flag-VprBP. Lysates were immunoprecipitated with an anti-Myc antibody in 1% SDS and Western blotted with the indicated antibodies. IN, input. * denotes bands corresponding to the expected proteins.

Appendix Figure S3: CP110 interacts with VprBP and its protein levels are dependent on EDD-DYRK2-DDB1^{VprBP} and Cep78

(A) Flag or Flag-VprBP was expressed in HEK293 cells. Lysates were immunoprecipitated with an anti-Flag antibody and Western blotted with the indicated antibodies. IN, input. **(B)** RPE-1 or HeLa cells were transfected with construct expressing Myc, Myc-VprBP, Flag, Flag-EDD, GFP or GFP-DDB1. Lysates were Western blotted with the indicated antibodies. β -actin was used as a loading control. **(C)** RPE-1 or HeLa cells were transfected with NS siRNA or Cep78 siRNA (oligo 2 or 6). α -tubulin was used as a loading control. **(D)** RPE-1 or HeLa cells were transfected with NS siRNA or Cep78 siRNA that targets the 3'UTR of Cep78 mRNA (oligo 7) and construct expressing Flag or Flag-Cep78. Lysates were Western blotted with the indicated antibodies. α -tubulin was used as a loading control.

Appendix Figure S4: Cep76 protein levels are not affected by EDD-DYRK2-DDB1^{VprBP} or Cep78

(A) HEK293 and HeLa cells were transfected with NS siRNA or Cep78 siRNA. Lysates were Western blotted with the indicated antibodies. β -actin was used as a loading control. **(B)** Flag, Flag-Cep78 or Flag-VprBP was expressed in HEK293 cells. Lysates were Western blotted with the indicated antibodies. β -actin was used as a loading control.

Appendix Figure S5: Cep78 regulates CP110 ubiquitination and protein levels through EDD-DYRK2-DDB1^{VprBP}

(A) HEK293 cells were transfected with constructs expressing Flag-Cyclin F, Flag-Neurl4 or Flag-VprBP, GFP or GFP-Cep78, and HA-CP110. Lysates were immunoprecipitated with an anti-HA antibody in 1% SDS and Western blotted with the indicated antibodies. IN, input. (B) Flag-Cyclin F, Flag-Neurl4 and Myc-VprBP were expressed singly or in combination in HeLa cells. Lysates were Western blotted with the indicated antibodies. α -tubulin was used as a loading control.* denotes bands corresponding to the expected proteins.

Appendix Figure S6: Cep78 controls CP110-dependent centriole elongation in non-ciliated cells

(A) HeLa cells transfected with NS siRNA and Cep78 siRNA (oligo 2 or oligo 6) were stained with DAPI (blue) and antibodies against Cep78 (red) and centrin (green). Scale bar, 1 μ m. (C) HeLa cells were transfected with NS siRNA or Cep78 siRNA that targets the 3'UTR of Cep78 mRNA (oligo 7) and construct expressing an irrelevant Flag-tagged protein (control) or Flag-Cep78. Cells were stained with DAPI (blue) and antibodies against Flag (red) and centrin (green). Scale bar, 1 μ m. (B and D) The percentage of cells with elongated centrioles (centrin filaments) was determined. At least 100 cells were scored in each condition and two independent experiments were performed.