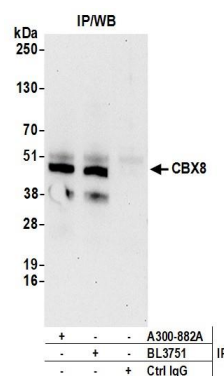
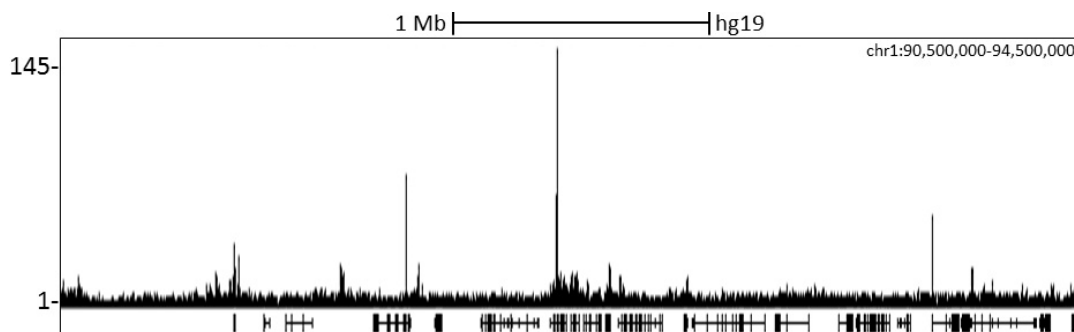


Detection of human CBX8 by western blot. *Samples:* Whole cell lysate (50 μ g) from HeLa and HEK293T cells prepared using NETN lysis buffer. *Antibody:* Affinity purified rabbit anti-CBX8 antibody A300-882A (lot A300-882A-2) used for WB at 0.1 μ g/ml. *Detection:* Chemiluminescence with an exposure time of 30 seconds.



Detection of human CBX8 by western blot of immunoprecipitates. *Samples:* Whole cell lysate (0.5 or 1.0 mg per IP reaction; 20% of IP loaded) from HeLa cells prepared using NETN lysis buffer. *Antibodies:* Affinity purified rabbit anti-CBX8 antibody A300-882A (lot A300-882A-2) used for IP at 6 μ g per reaction. CBX8 was also immunoprecipitated by rabbit anti-CBX8 antibody BL3751. For blotting immunoprecipitated CBX8, A300-882A was used at 0.4 μ g/ml. *Detection:* Chemiluminescence with an exposure time of 30 seconds.



Localization of CBX8 Binding Sites by ChIP–sequencing. Chromatin from K562 cells was immunoprecipitated with anti–CBX8 antibody A300–882A and analyzed by DNA sequencing. The figure illustrates the peak distribution of CBX8 binding within a 4 Mb region of chromosome 1 as detected using anti–CBX8 antibody A300–882A. ChIP–seq validation performed by Diogenode, Denville, NJ.