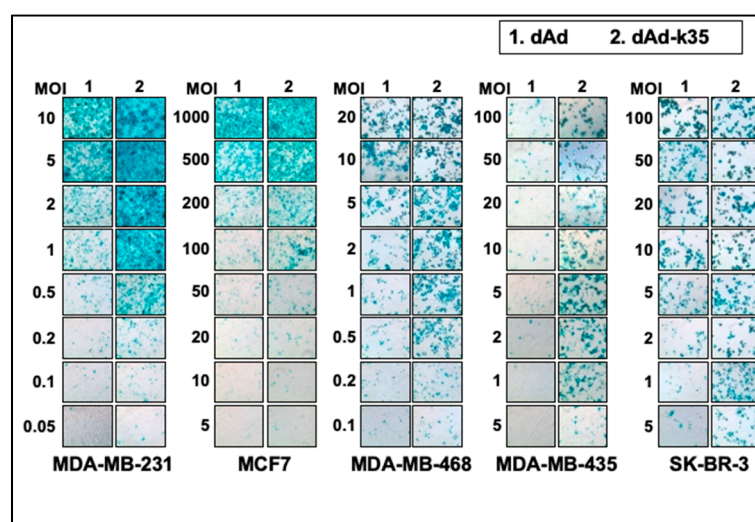


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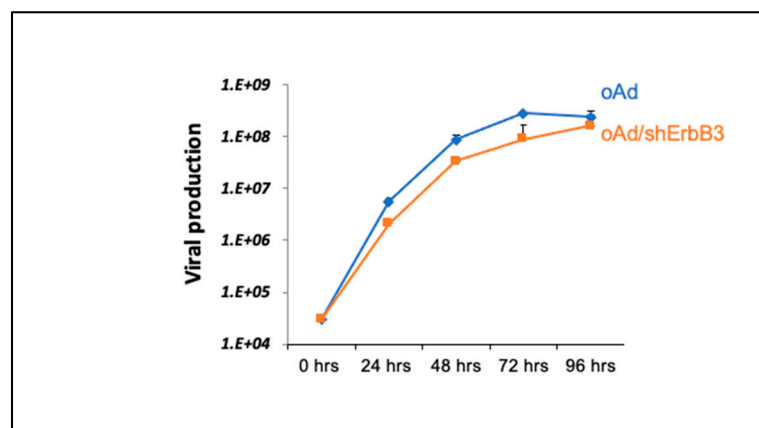
ErB3-targeting oncolytic adenovirus causes potent tumor suppression by induction of apoptosis in cancer cells

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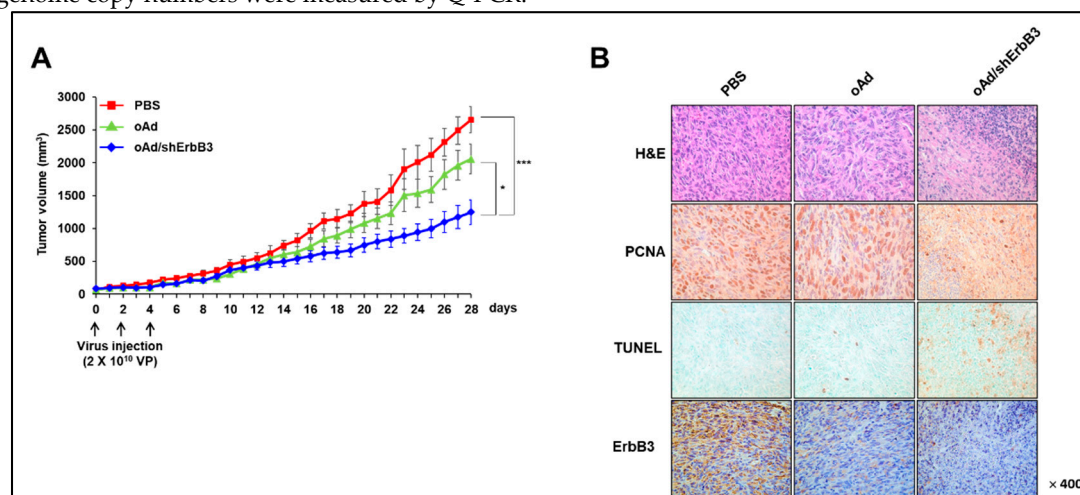
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Supplementary Figure S1. Enhanced transduction efficiency of dAd-k35-LacZ. Human breast cancer cells (MDA-MB-231, MCF7, MDA-MB-468, MDA-MB-435, and SK-BR-3) were transduced with dAd-LacZ (lane 1) or dAd-k35-LacZ (lane 2) with various amount of virus (0.05 - 1000 MOI). At 48 h post-transduction, cells transduced with dAd-LacZ or dAd-k35-dAd-LacZ were stained with X-gal for analyzing the β -galactosidase.



Supplementary Figure S2. Viral replication of oAd and oAd/shErbB3. MDA-MB-231 cells seeded in a 12-well plate were infected at MOI of 0.1 with oAd or oAd/shErbB3. At 0, 24, 48, 72, or 96 h post infection, viral genome copy numbers were measured by Q-PCR.



Supplementary Figure S3. Tumor growth inhibition by shErbB3-expressing oncolytic Ad. MCF-7/Mot tumor-bearing mice were injected with 2×10^{10} VP of either oAd or oAd/shErbB3 along with PBS control (Q2D \times 3). * $p < 0.01$ or *** $p < 0.001$. **(A)** Tumor volume was measured every other day. Data represent as mean \pm SE ($n = 5$ to 8). **(B)** Histological and immunohistochemical analysis. Tumors treated with PBS, oAd, or oAd/shErbB3 were harvested and subjected to H & E staining and immunohistochemical staining for PCNA, TUNEL, and ErbB3, respectively.