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Correction: Targeting Tumor Cell Invasion and Dissemination In Vivo by an Aptamer that Inhibits Urokinase-Type Plasminogen Activator through a Novel Multifunctional Mechanism

The inaugural Rapid Impact article, by Kobayashi and colleagues (beginning on page 828), reveals a crosstalk between two dominant cell cycle tumor suppressor proteins such that p14ARF regulates the stability of p16INK4A through a degradation mechanism involving the REGγ subunit of the 20S proteasome. Accompanying the article online, and presented on the cover, the AACR and Molecular Cancer Research are proud to introduce a new article feature called Visual Overview in which the novel findings of the article are graphically depicted.