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Plasmid construction and viral infection

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Yuejun Wang[1]

[1]Department of Prosthodontics, Peking University School and Hospital of Stomatology

Yuejun Wang

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Abstract

The plasmids were constructed using standard methods; all sequences were verified by appropriate restriction digestion and/or sequencing. Human full-length *IGFBP2* cDNA from ASCs fused to a M2-Flag tag was produced with a standard PCR protocol. This sequence (Flag-*IGFBP2*) was subcloned into the pQCXIN retroviral vector with AgeI and BglII restriction sites. For viral infections, MSCs were plated overnight, then infected with retroviruses in the presence of polybrene (6 µg/mL, Sigma-Aldrich, St. Louis, MO, USA) for 12 h. After 48 h, infected cells were selected with 600 µg/mL G418 for 10 days.

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