

Differing responses of osteogenic cell lines to glycerophosphate.

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Supplementary materials

Gene	Forward primer	Reverse primer
Tyrosine 3-monooxygenase, <i>YWHAZ</i>	ACTTTTGGTACATTGCTTCAA	CCGCCAGGACAAACCAGTAT
Osteocalcin, <i>OCN</i>	GGCAGCGAGGTAGTGAAGAG	CTGGAGAGGAGCAGAACTGG
Alkaline phosphatase, <i>ALP</i>	TGCTCTGCGCAGGATTG	GGAGACACCCATCCCATCTC
Runt-related transcription factor 2, <i>RUNX2</i>	CGCCTCACAAACAACCACAG	TCACTGTGCTGAAGAGGCTG
Phosphate regulating endopeptidase homolog X-linked, <i>PHEX</i>	ACTTTGCACTGCACTGGACT	TCCATCAGAAGGGCCGTAGA
Marker of proliferation Ki-67, <i>MKI67</i>	GCCCGGGGACGTAGCCTGTA	ACCGTCGACCCCGCTCCTTT
Proliferating cell nuclear antigen, <i>PCNA</i>	CCACGTCTCTTTGGTGCAG	CCGGCGCATTTTATGATTTTGG

Table 1. Primers were designed to assess the expression of osteogenic differentiation and cell proliferation using qPCR.

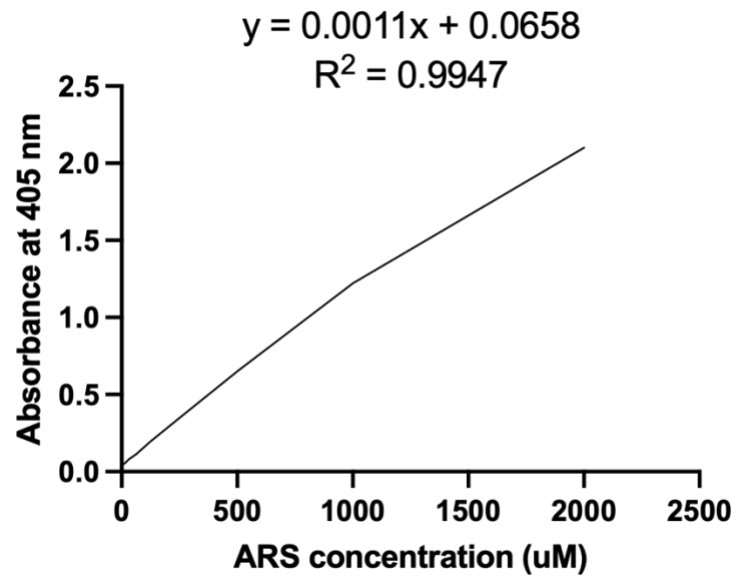


Figure 1. Standard curve of serial dilutions of alizarin red stain used to quantify optical density of mineralising hMSCs and Saos-2 cell cultures supplemented with combinations of Asc, Dex and β -Gly.