

Supporting Information

PEGylated graphene oxide-mediated quercetin modified collagen hybrid scaffold for enhancement of MSCs differentiation potential and diabetic wound healing

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Table S1 Primers for generating PCR fragments

Name of primer	Forward primer (5'-3')	Reverse primer (5'-3')
β-actin	CGTTGACATCCGTAAAGACC	TAGGAGCCAGAGCAGTAATC
LPL	AGTTTGACCGCCTCCGG	TCCTGTCACCGTCCATCCATGGA
PPAR γ	ACTGCCGGATCCACAAAAA	TCTCCTTCTCGGCCTGTG
ALP	AACCCAGACACAAGCATTCC	CCAGCAAGAAGAACCTTTG
Runx-2	TGCCACCTCTGACTTCTGCC	CGCTCCGGCCCACAATCTC
Col I	TGACTGGAAGAGCGGAGAGT	GACGGCTGAGTAGGAAACAC
Col III	AGGTTCTCCTGGTGCTGCT	GGATGCCCACTTGTCCAT
a-SMA	AATGGCTCTGGGCTCTGTAA	CTCTTGCTCTGGGCTTCATC

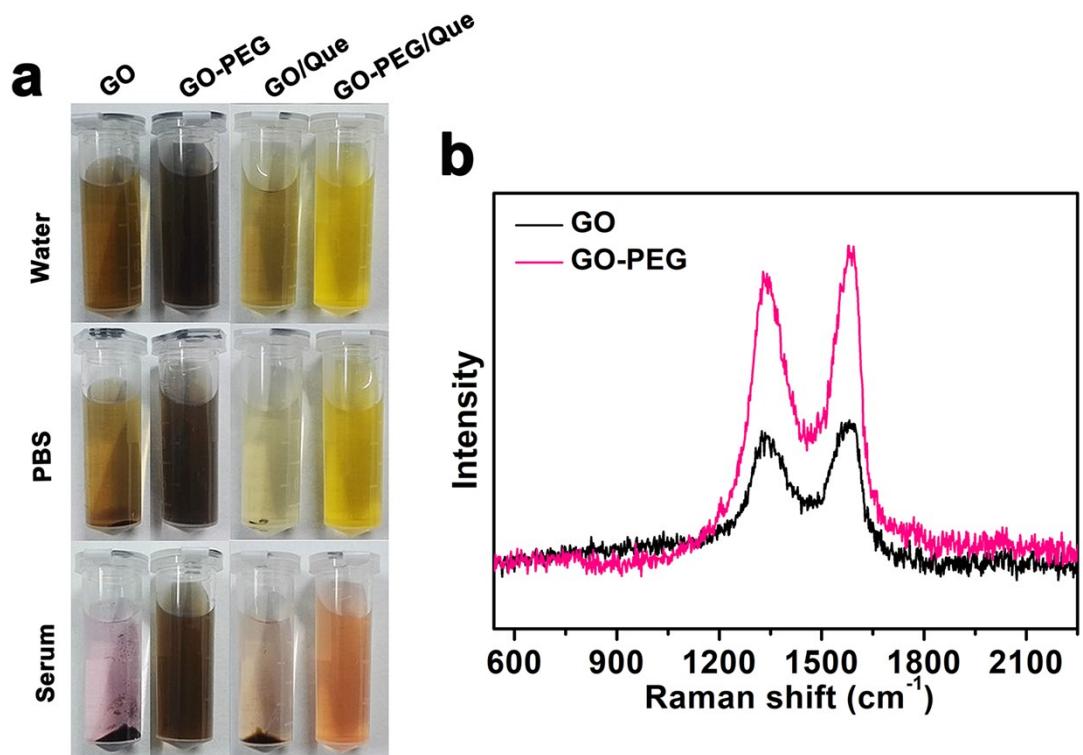


Fig. S1 (a) Photos of the stability of GO, GO-PEG, GO/Que and GO-PEG/Que in Water, PBS and Serum, recorded after centrifugation at 10,000 g for 5 min. (b) Raman spectra of GO and GO-PEG, recorded at 514.5 nm.

Table S2 Wavenumbers (cm^{-1}) and assignments of the main infrared bands of GO and GO-derivatives^a

Compounds	$\nu\text{-OH}$	$\nu\text{-CH}_2\text{-}$	$\nu\text{C=O}$	$\nu\text{-NH-CO-}$	$\nu\text{C}_2\text{=C}_3$	$\nu\text{C=C(A)}$	$\delta\text{C}_3\text{-OH}$	$\nu\text{C-O}$	$\nu\text{C-O-C}$	$\nu\text{benzene}$
GO	3342s	—	1722m	—	1621m	—	1374b.w	1250	1043m	—
GO-COOH	3136b.s	—	1706m	—	1581s	—	1363b.s	—	1043s	—
GO-PEG	3200b.s	2850w	—	1698w	1616b.s	—	1347m	—	1060s	—
Que	3239b.s	—	1660m	—	1604s	1519s	1378m	1315m	1091w	817m
GO/Que	3255b.s	—	1635vw	—	1606m	1520w	1380w	1320vw	1058s	812w
GO-PEG/Que	3254b.s	2849w	1637vw	1660m	1606s	1518s	1378m	1316m	1090w	816m

a Abbreviations: b., broad; m., medium; s., strong; w., weak; vw., very weak

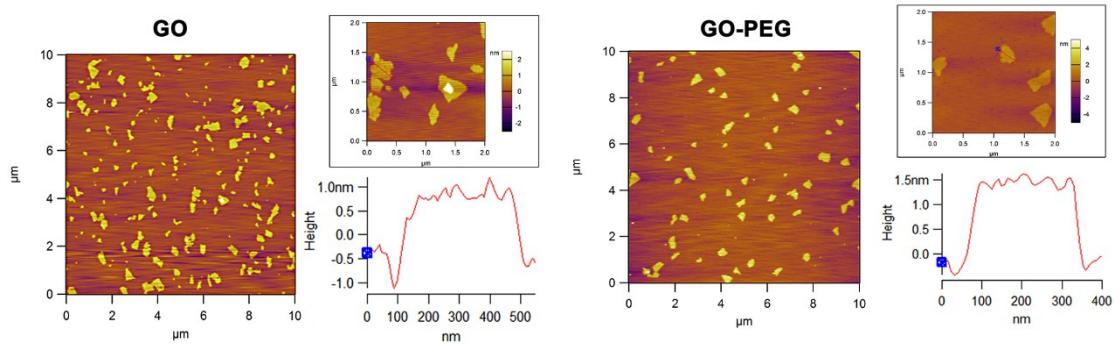


Fig. S2 AFM images and representative height profiles of GO and GO-PEG.

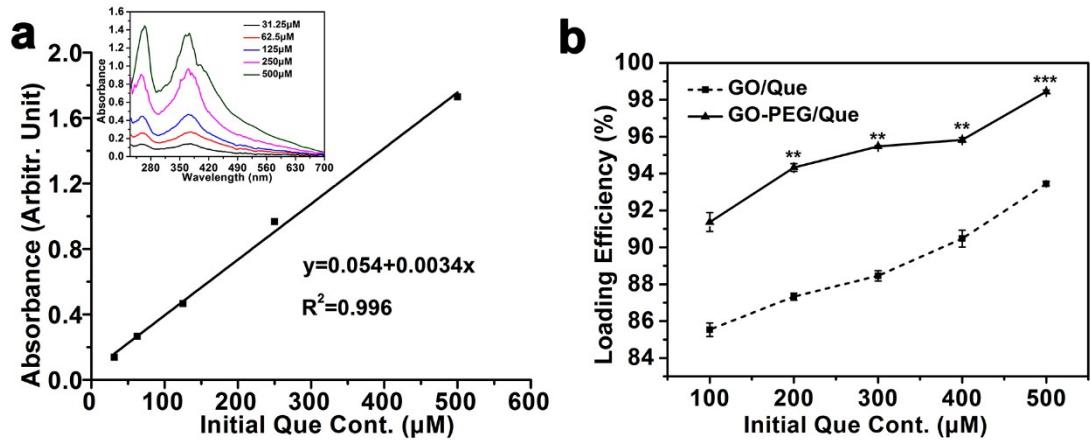


Fig. S3 Loading efficiency of GO-PEG/Que. (a) UV-Vis absorbance and standard curve of different initial Que concentrations at a wavelength of 370 nm. (b) Loading efficiency of different initial amount of Que loaded onto 0.1 mg/mL GO and GO-PEG (* $p \leq 0.05$, ** $p \leq 0.01$, $n = 3$).

Table S3. Raman-selected bands (cm^{-1}) and assignments in the range of 700-1700 cm^{-1} region of ADM scaffold ^a

Tentative assignments	Raman shift/ cm^{-1}
γ -(C-C)-Pro	857 923
γ -(C-C)-HyPro	880
γ -(C-C)-backbone	939
Phe	1004 1207
Pro	1034
γ -(C-C)	1167
HyPro	1178
Amide III	1248 1280
δ -(CH)	1322 1444
CH_2 -Pro	1346
δ -(COO-)	1428
C-H	1458
Amide I	1640 1673

^a Abbreviations: γ , stretching vibration; δ , deformation vibration; Phe, phenylalanine; Pro, proline; Hypro, hydroxyproline.

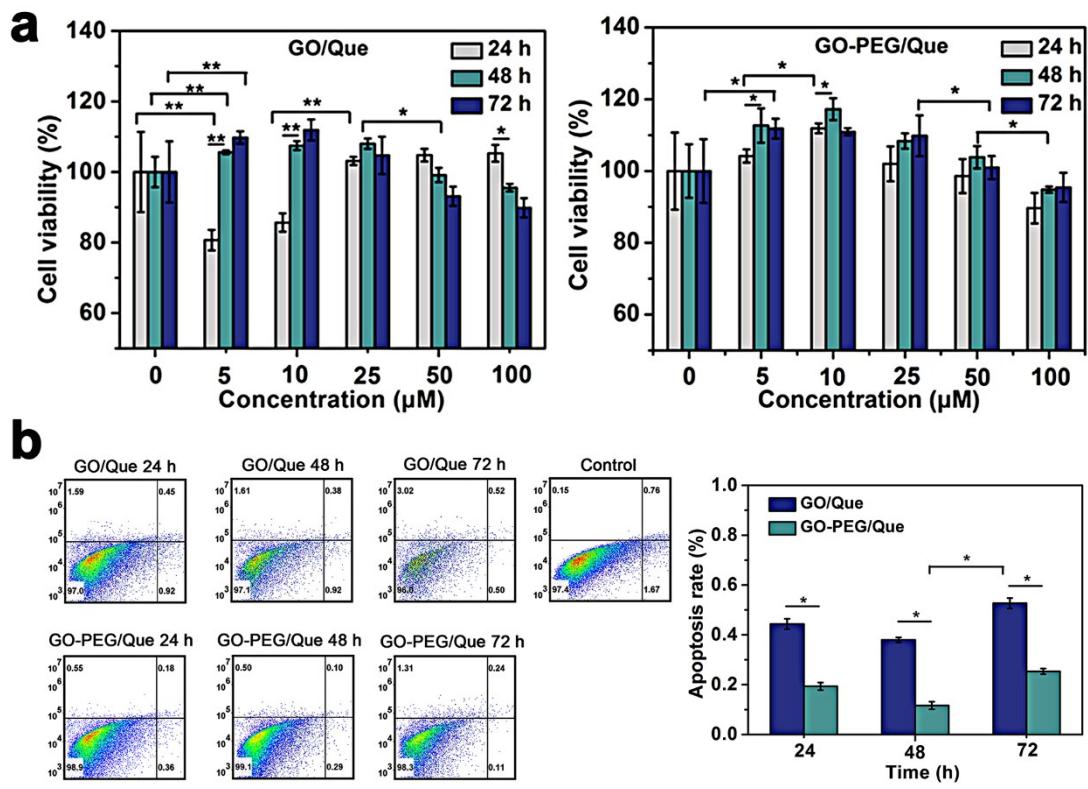


Fig. S4 *In vitro* cell viability and toxicity. (a) MTT assay to assess the viability of MSCs incubated with various GO/Que and GO-PEG/Que concentrations (0–100 μ M) for 24 h, 48 h and 72 h, respectively (* $p \leq 0.05$, ** $p \leq 0.01$, $n = 6$). (b) MSCs after 24 h, 48 h and 72 h of incubation with GO/Que and GO-PEG/Que were subjected to flow cytometry analysis (* $p \leq 0.05$, $n = 3$).

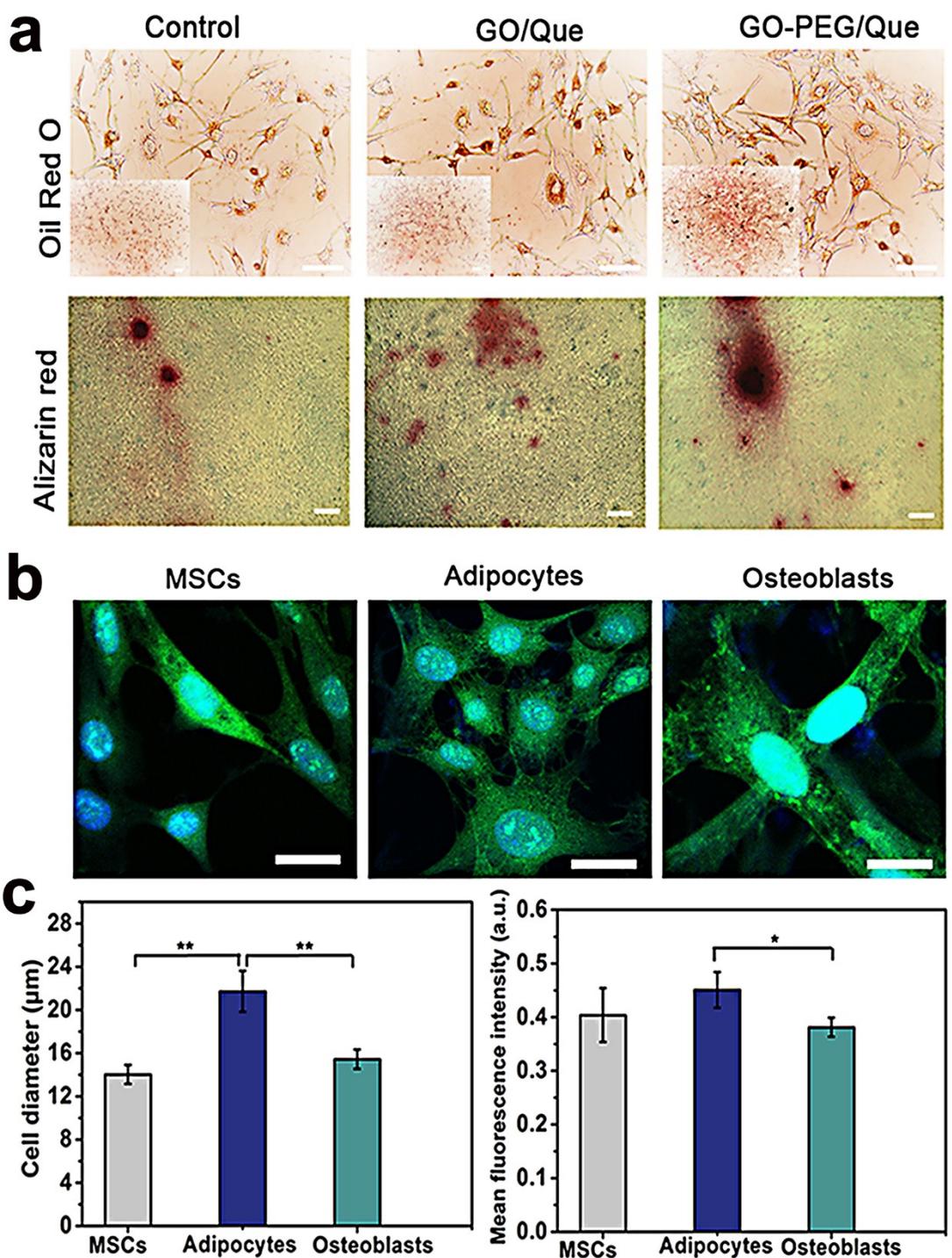


Fig. S5 Adipogenic and osteogenic differentiation of MSCs *in vitro*. (a) Oil red O and Alizarin red staining of MSCs after incubation with induction medium (control), GO/Que and GO-PEG/Que ($n = 6$). Scale bar: 100 μm . (b) Morphology of the MSCs, adipocytes and osteoblasts. Scale bar: 10 μm . Cell diameter (c) and mean fluorescence intensity (d) of MSCs, adipocytes and osteoblasts (* $p \leq 0.05$, ** $p \leq 0.01$, $n = 6$).