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Background {#Sec1} ===== The therapeutic potential of cells administered in vivo is most likely restricted by their low survival rates. While mesenchymal stem cells (MSCs) are highly sensitive to hypoxia, conditioned medium from hypoxic cells increases the survival of MSCs in vitro. We have previously shown that human MSCs (hMSCs) in vitro respond to hypoxia or conditioned media from hypoxic cells by a change in morphology to a spindle-shaped phenotype and an increased expression of HIF1-alpha protein \[[@CR1]\]. This suggests that the hypoxia-associated phenotype is induced in vitro by soluble factors secreted by hypoxic hMSCs. In this study we investigate if the hypoxia-induced phenotype can be induced in vitro in undifferentiated hMSCs by conditioned medium of hypoxic hMSCs.

Methods {#Sec2} ===== Hypoxic hMSCs were established by two-step hypoxic preconditioning of hMSCs. For these experiments, hMSCs were derived from placental tissue of uncomplicated, term pregnancies and expanded in culture. Hypoxic hMSC-conditioned media were obtained from culture medium of hypoxic hMSCs cultivated for 48 hours. Culture medium was carefully removed and replaced with an equal volume of fresh medium. Culture medium from normoxic cells was collected as control. hMSCs were incubated with hypoxic or normoxic conditioned media for 4 hours, 6 hours or 24 hours. hMSCs were then fixed and immunostained for HIF1-alpha, HIF2-alpha and HIF3-alpha. hMSCs were also stained with propidium iodide for analysis of cell death.

Results {#Sec3} ===== The data in Figure [1](#Fig1){ref-type="fig"} show that conditioned media from hypoxic hMSCs are able to induce a hypoxia-like phenotype in undifferentiated hMSCs within 4 hours (Figure [1](#Fig1){ref-type="fig"}) and 6 hours (Figure [2](#Fig2){ref-type="fig"})

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website from his colleague, Barry Seidman,. MPEG-2, DivX, and the Good Old Days:LTE-Advanced (LTE-A) is a network technology that aims to achieve a total of 100 Mbps/50 Mbps on the downlink/uplink, and roughly 100 Mbps/20 Mbps on the uplink through the application of multiple-input multiple-output (MIMO) transmission and a supporting efficient cell structure. In a future 5G system, 5G-dedicated small cell technologies may be introduced to supplement a coverage of macro base stations, and a coverage hole phenomenon caused by a coverage of a macro base station, such as inter-cell interference, etc., may be reduced or avoided. In an aspect of the 5G system, to reduce or avoid the coverage hole phenomenon, a small cell base station may be installed in each coverage hole of a macro base station (that is, a plurality of small cells), and a user

equipment (UE) may receive service through the small cell base station, thereby enhancing a cell capacity. As shown in FIG. 1, in an LTE system, an evolved Node B (eNB) transmits a pilot symbol periodically on the downlink in all occasions, and a User Equipment (UE) calculates a downlink channel estimation value through the received pilot symbol. When the UE does not receive the pilot symbol for a period of time, the UE stops channel estimation. A parameter of the pilot symbol is predefined according to the length of the pilot symbol (for example, 20 symbols). The UE generally performs channel estimation after the whole pilot symbol is received, which causes a long channel estimation time (two seconds or above in an example). This may cause a channel estimation fault, and degrade a channel estimation performance.

Dealing with Aplastic

anemia patients. Aplastic anemia (AA) is a bone marrow failure syndrome characterised by pancytopenia in the absence of congenital or acquired causes of bone marrow failure. The diagnosis of AA is based on three major criteria and two minor criteria. The major criteria include aplastic anaemia-related haemorrh

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