

# Transfection of endometrial cells with siRNA

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## **Abstract**

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#### **Protocol**

## Step 1.

Shortly before transfection, seed  $1x \ 10^5$  cells per well of a 24-wellplate with 0.5 ml appropriate culture medium containing serum.

## Step 2.

For the short time until transfection, incubate the cells under normal growthconditions (typically 37°C and 5% CO2).

## Step 3.

Dilute 37.5 ng siRNA in 100  $\mu$ l culture medium without serum (this will givea final siRNAconcentration of 5 nM after adding complexes to cells instep 5). Add 3  $\mu$ l HiPerFect Transfection Reagent to the diluted siRNA andmix by vortexing.

#### Step 4.

Incubate for 5 min at room temperature (20°C) to allow theformation of transfection complexes.

#### Step 5.

Add the complexes drop-wise onto the cells. Gently swirl the plate to ensureuniform distribution of the transfection complexes.

## Step 6.

Incubate the cells with the transfection complexes under their normalgrowth conditions, and monitor gene silencing after 48h. Changethe medium as required.

## Step 7.