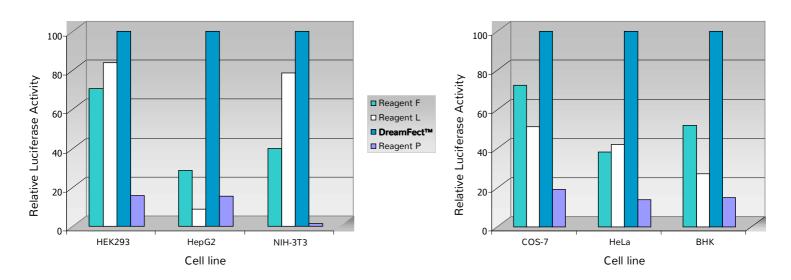


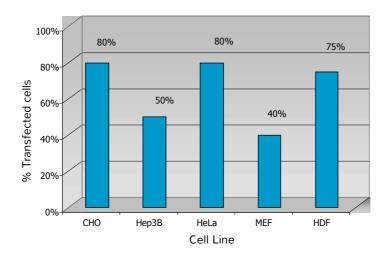
DreamFect [™] complementary results

DreamFect [™] efficiency compared to other transfection reagents



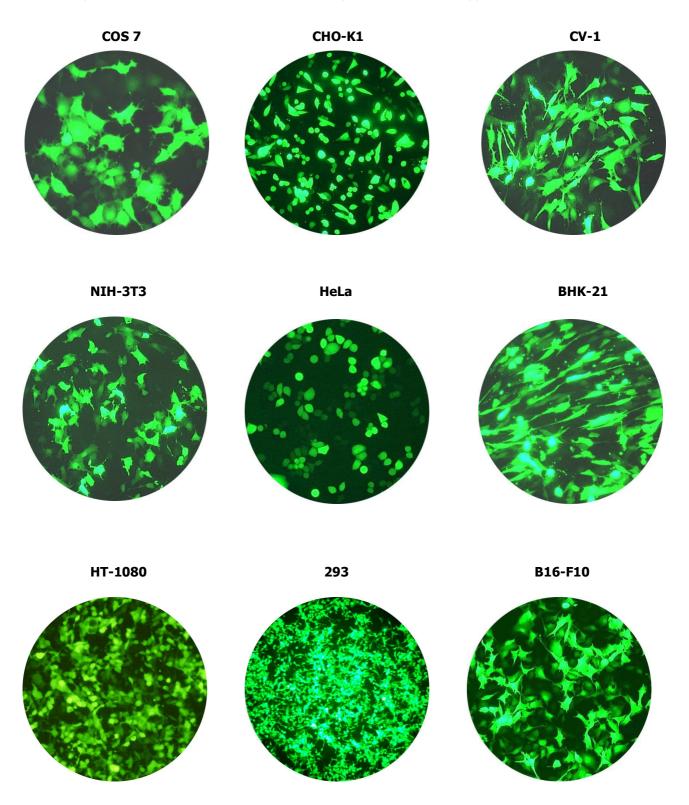
Cells were transfected with 0.1 μ g/well of pND2Luc plasmid DNA in 96-well plates. **DreamFect TM** transfections were performed as described in the instruction manual with 0.4 μ l/well of DreamFectTM reagent. The other transfection reagents were assayed according to the manufacturer's instruction. Luciferase activities were measured with a Luciferase assay kit and results are expressed as relative values.

Percentage of transfected cells achieved with *DreamFect* ™ on several cell lines



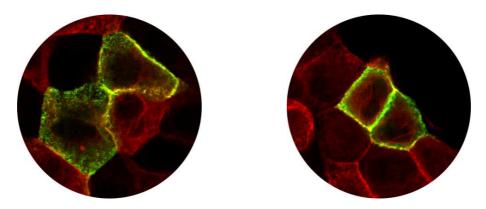
Green Fluorescent Protein expression in different cell lines

Cells (2x10⁵) were transfected with 1 μ g/well of pEGFP plasmid and 4 μ l of **DreamFect** TM reagent in 12-well plates. E-GFP expression was monitored 48 h after transfection by fluorescence microscopy.



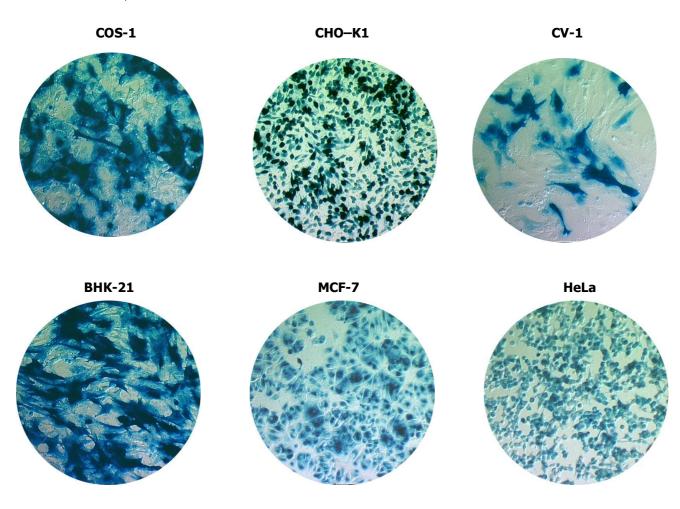
Madin Darby Canine Kidney (MDCK) cells $(4x10^5)$ were transfected with 1 μ g of plasmid DNA, encoding for a chimeric E-GFP protein targeted to the cell membrane (PH Domain), and 4 μ l of **DreamFect** reagent on coverslips in a 6-well

plate. 24 hours post-transfection, cells were fixed with 4% PFA and actin was stained with Texas Red-labeled phalloidin. E-GFP expression and localization was monitored by confocal microscopy.



β-Galactosidase expression in different cell lines transfected with *DreamFect* ™

Cells $(4x10^5)$ were transfected with 2 µg/well of pLacZ plasmid and 8 µl of **DreamFect** TM reagent in 6-well plates. β-Galactosidase expression was revealed 48 h after transfection using OZ Biosciences' X-Gal staining kit (catalog number GX-10003).



Page 3 of 3