

## PolyMag Results

Magnetofection™ is a novel, simple and highly efficient method to transfect cells in culture and in vivo. It exploits magnetic force exerted upon gene vectors associated with magnetic particles to drive the vectors towards, possibly even into, the target cells. In this manner, the complete applied vector dose gets concentrated on the cells within a few minutes so that 100% of the cells get in contact with a significant vector dose.

This has several important consequences:

1. Greatly improved transfection rates in terms of percentage of cells transfected compared to standard transfections.
2. Up to several thousand folds increased levels of transgene expression compared to standard transfections.
3. High transfection rates and transgene expression levels are achievable with extremely low vector doses, which allow saving expensive transfection reagents.
4. Extremely short process time in comparison to standard procedures. A few minutes of incubation of cells with gene vectors are sufficient to generate high transfection efficiency.

OZ Biosciences offers three types of ready-to-use reagents: **SilenceMag** specific to siRNA delivery, **CombiMag** designed to be associated with all vectors: transfection reagents and viruses and **PolyMag**. **PolyMag** has been designed and used successfully to deliver all types of nucleic acids such as plasmid DNA, RNA, Oligonucleotides & siRNA. **PolyMag** is a universally applicable magnetic particle preparation for high transfection efficiency. Nucleic acids to be transfected and the magnetic particles are mixed in a one-step procedure.

## Nucleic Acid Types

Nucleic Acid or Virus Type	PolyMag	SilenceMag	CombiMag
<b>DNA (plasmid)</b>	√	NA	√
<b>Oligonucleotides</b>	√	ND	√
<b>mRNA</b>	√	ND	√
<b>siRNA</b>	√	√	√
<b>dsRNA</b>	ND	√	√
<b>shRNA</b>	ND	√	√
<b>Adenovirus</b>	NA	NA	√
<b>Retrovirus</b>	NA	NA	√

## Cell Types

**PolyMag** is generally applicable on numerous cell types. This technology has been tested successfully on a variety of immortalized and primary cells.

## Transfection of Primary Cells

### Confluent Primary Human Keratinocytes

Transfected with a commercial reagent L or **PolyMag**

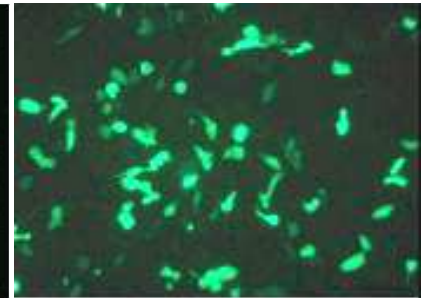
*Reporter Gene: GFP*

We are grateful to the laboratories of Dr. C. Plank (Technical University, Munich) for kindly providing these data.

Standard Transfection



Magnetofection™



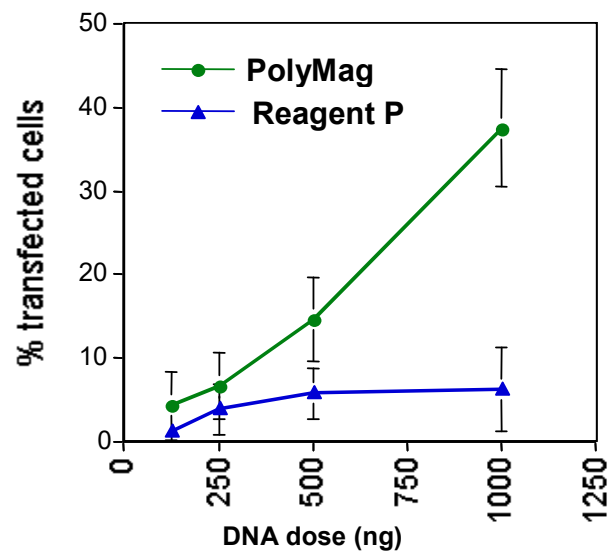
### Primary Porcine Aortic Endothelial Cells

Transfected with a commercial reagent P or **PolyMag**

*Reporter Gene: GFP*

*Culture dish: 96-well plate*

We are thankful to Dr. F. Kroetz (Ludwig-Maximilians University, Munich) for nicely providing these data.



## Transfection of Cell Lines

### CT-26 Colon Carcinoma / Mouse

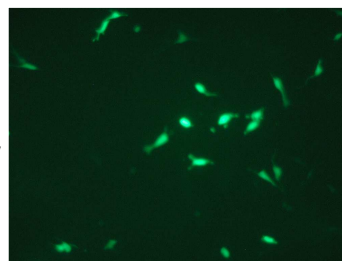
Transfected with a commercial reagent D or **PolyMag**

*Reporter Gene: GFP*

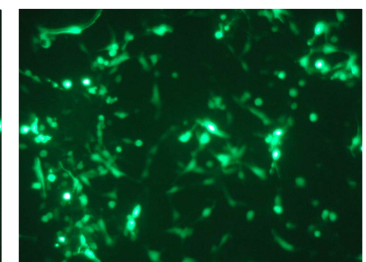
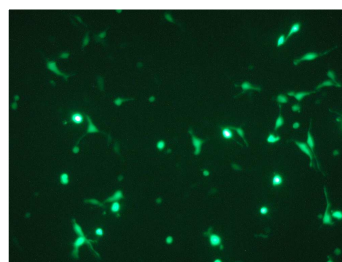
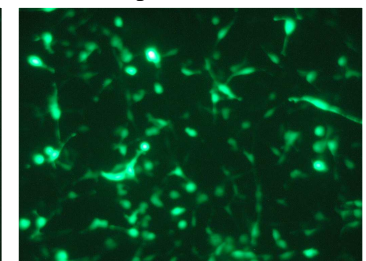
*Culture dish: 24-well plate*

We are grateful to the laboratories of Dr. C. Plank (Technical University, Munich) for kindly providing these data.

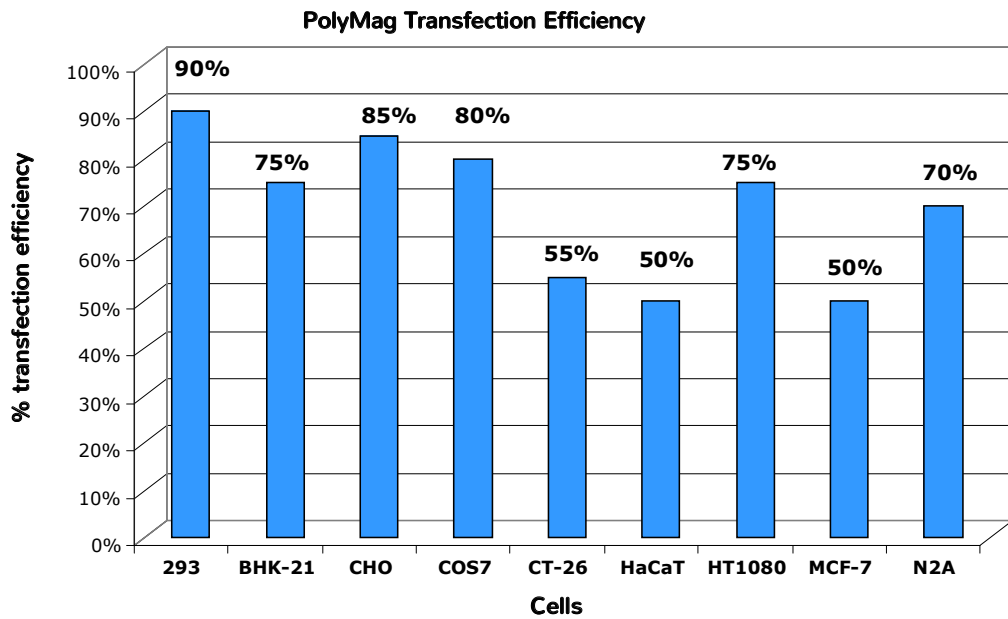
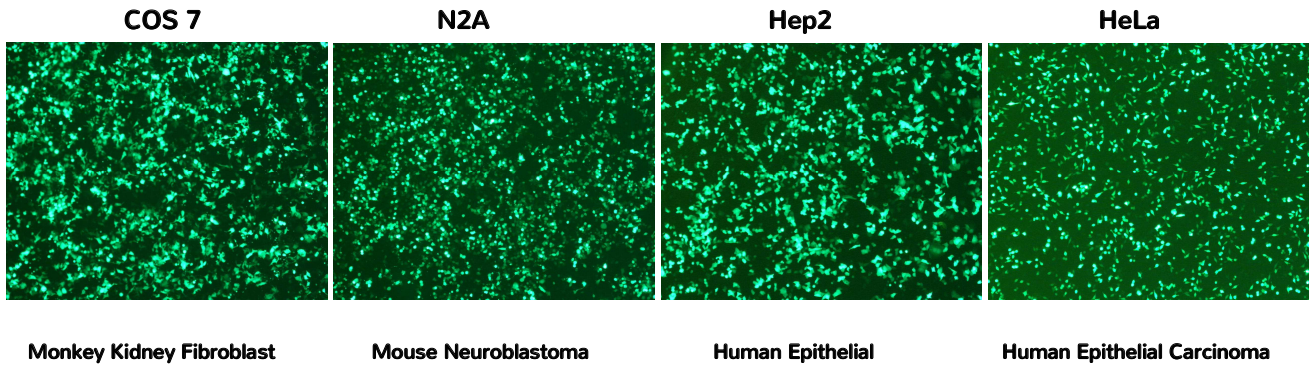
Standard Transfection



Magnetofection™



Various examples of **PolyMag** transfection efficiency (24-well plate; 0.5  $\mu$ l PolyMag & 0.5  $\mu$ g DNA/ well):

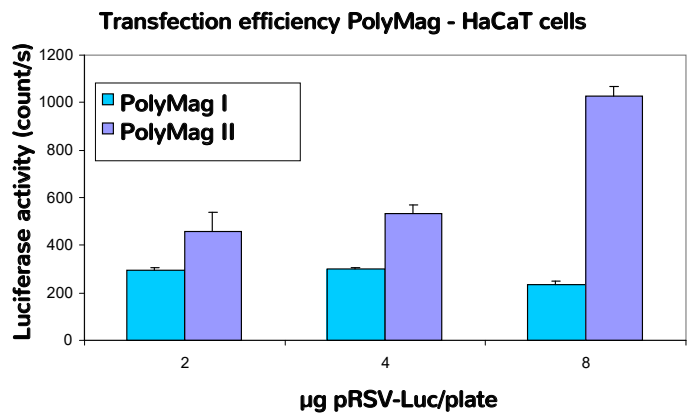


**Improved and Optimized PolyMag Formulation**

OZ Biosciences has developed a new very efficient **PolyMag** formulation for Magnetofection™. This improved PolyMag II reagent gives reliable higher transfection efficiencies in numerous cell types (see figures), transports all type of nucleic acids (DNA, siRNA, ODN). This **PolyMag** optimized formulation is the actual commercialized **PolyMag** reagent.

Most of the herein described results were obtained with the new optimized **PolyMag** formulation.

We are thankful to Dr. L. Journot (CNRS-UPR2580, Montpellier) for nicely providing these data.



## Delivery of Oligonucleotides

### HUVEC-C (Primary) / Human

Transfected with a commercial reagent E or

#### PolyMag

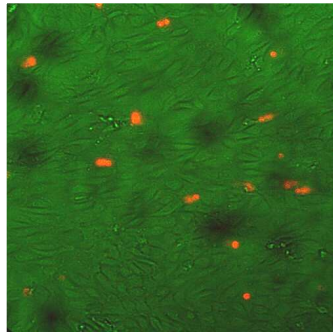
Fluorescent Oligonucleotides

*Culture dish: 24-well plate*

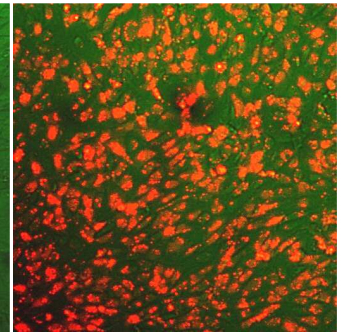
*ODN: 200 ng / well*

We are grateful to Dr. F. Kroetz (Ludwig-Maximilians University, Munich) for kindly providing these data.

### Standard Transfection



### Magnetofection™



## siRNA Delivery Efficiency

### GFP stably transfected HeLa cells.

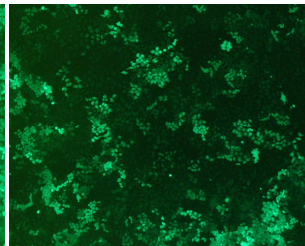
*Target siRNA: GFP*

*Cell culture dish: 96-well plate*

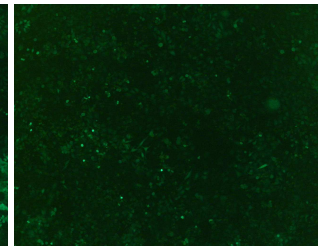
*Transfection volume: 200  $\mu$ l.*



Control



PolyMag + siRNA/10nM



PolyMag + siRNA/40nM

### Primary Human Umbilical Vein Endothelial Cells (HUVEC).

*Target siRNA: Transcription factor # 1*

*Control siRNA: Transcription factor # 2*

(Belongs to TF#1 family)

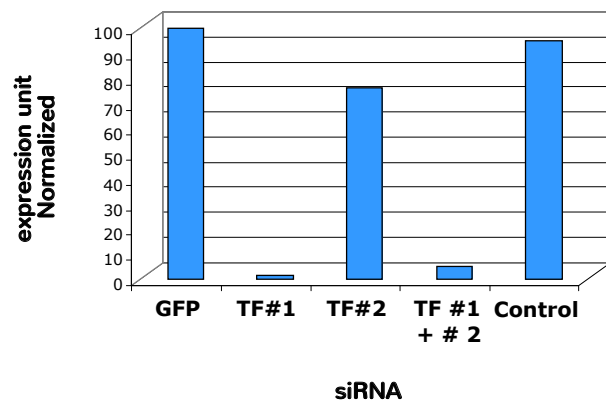
*Control siRNA: GFP*

*Cell culture dish: 60 mm*

*SiRNA final concentration: 25nM (2x treatments)*

We are grateful to Dr. D. Mathieu (CNRS-UMR5535, Montpellier) for kindly providing these data.

### TF # 1 Gene Silencing mediated by PolyMag/siRNA



## Cytotoxicity Electroporation vs. Magnetofection

