

# TransforMax EC100 elektrokompetente E. coli

Lucigen



epicentre  
an illumina company

## TransforMAX™

10 x 100 µl

Artikel-Nr.: 190055 | Lucigen | Hersteller-Nr.: EC10010

**370,80 €\***

\*zzgl. MwSt. zzgl. Versandkosten

## Beschreibung

**Produkttyp:** Kompetente Zellen

**Verpackung:** 10 x 100 µl

### Applications

- Routine cloning of DNA up to 200 kb.

The highly versatile TransforMax™ EC100™ *E. coli* competent cells are ideal for most cloning applications. The cells provide very high transformation efficiency when tested against a wide range of supercoiled DNAs as well as DNA directly from a ligation reaction (Table 1).

### Benefits

- High transformation efficiency with clones of all sizes, including BAC clones (Table 1).
- *lacZ*?*M15* for blue/white screening of recombinants.
- Restriction minus [*mcrA*, ?(*mrr-hsdRMS-mcrBC*)] enables efficient cloning of methylated DNA.
- Endonuclease minus (*endA1*) to ensure high yields of DNA.
- Recombination minus (*recA1*) for greater stability of large cloned inserts.

### Genotype

*F*<sup>-</sup> *mcrA* ?(*mrr-hsdRMS-mcrBC*) ?80*dlacZ*?*M15* ?*lacX74* *recA1* *endA1* *araD139* ?(*ara*, *leu*)7697 *galU* *galK* ? *rpsL* (*Str*<sup>R</sup>) *nupG*

### TransforMax EC100 Electrocompetent *E. coli*

- Transformation efficiency of >1 x 10<sup>10</sup> cfu/µg of pUC19.

### TransforMax EC100 Chemically Competent *E. coli*

- Transformation efficiency of >5 x 10<sup>8</sup> cfu/µg of pUC19.

DNA	TransforMax™ EC100™ Chemically Competent <i>E. coli</i>	TransforMax™ EC100™ Electrocompetent <i>E. coli</i>
pUC19	$1.4 \times 10^8$	$1.4 \times 10^{10}$
8.1-kb Clone	$1.3 \times 10^7$	Not tested
13.1-kb Clone	$4.3 \times 10^6$	$1.3 \times 10^9$
23.1-kb Clone	$9.2 \times 10^5$	$3.0 \times 10^8$
145-kb BAC Clone	Not tested	$7 \times 10^7$
13.1-kb clone directly from a ligation reaction	$2.2 \times 10^5$	$2.1 \times 10^7$

**Table 1. Comparison of the transformation efficiencies of TransforMax™ EC100™ *E. coli* with a variety of DNAs.**

Transformations were performed using 50 µl of competent cells and either supercoiled DNAs of the indicated sizes or a 1-µl aliquot from a standard 10-µl ligation reaction. Results shown are in cfu/µg of DNA and are the average transformation efficiencies obtained from several trials.