



GENOME EDITING KIT - SIZE: 6 / 35 EDITS\*

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## Super UniGuide

Achieves higher cleavage efficiencies and allows editing of difficult targets.

\* For use with S.p. Cas9 and derived variants. Suitable for 6 edits using electroporation or 35 edits using lipofection. Store at -20°C. Avoid freeze thaw cycles. Vortex and centrifuge briefly after thawing.

For research use only. UBMTA applies.

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## Contents

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- **tracrRNA component of gRNA**

2,000 pmol lyophilized modified RNA

- **Buffer A**

20  $\mu$ l aqueous solution

## Quick protocol

1. Centrifuge all tubes briefly before opening to ensure the contents are collected at the bottom.
2. Prepare a 200  $\mu$ M stock solution by resuspending the ● **tracrRNA** in 10  $\mu$ l of ● **Buffer A**.
3. Mix with an equal amount of any crRNA (not included in this kit; e.g. 2  $\mu$ l of 200  $\mu$ M tracrRNA + 2  $\mu$ l of 200  $\mu$ M crRNA).
4. Incubate the resulting mix at 95°C for 2 minutes, then allow to cool at room temperature for 10 minutes to form a functional gRNA.
5. Add gRNA to the tube containing the other genome editing components before delivery.

Use case suggestion: Use 3  $\mu$ l gRNA per electroporation (per 1 million cells) or 0.5  $\mu$ l gRNA per lipofection (per 6-well). Titration in your system of choice might increase efficiency further.



For more information visit:

[www.clearcutbiolabs.com/solutions/super-uniguide/](http://www.clearcutbiolabs.com/solutions/super-uniguide/)