


Stable Cells with NF- κ B-ZsGreen Fused Genes Created by TALEN Editing and Homology Directed Repair for Screening Anti-inflammation Drugs

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Background: NF- κ B is a sequence-specific DNA-binding transcription factor that plays key roles in inflammation and cancer. It is well known that NF- κ B is over-activated in these diseases. NF- κ B inhibitors are therefore developed as promising drugs for these diseases. However, finding NF- κ B inhibitors is dependent on effective screening platforms.

Methods: For providing an easy and visualizable tool for screening NF- κ B inhibitors, and other NF- κ B-related studies, this study edited all five genes of NF- κ B family (RELA, RELB, CREL, NF- κ B1, NF- κ B2) in three different cell lines (293T, HepG2, and PANC1) with both TALEN and CRISPR. The edited NF- κ B genes were repaired by homology-dependent repair using a linear homologous donor containing ZsGreen coding sequence. The edit efficiency was thus directly evaluated by detecting cellular fluorescence. The editing efficiency was also confirmed by PCR detection of NF- κ B-ZsGreen fused genes.

Results: It was found that all genes were more efficiently edited by TALEN in all cells than CRISPR. The positive cells were then isolated from the TALEN-edited cell pool by flow cytometry. The purified positive cells were finally evaluated by regulating NF- κ B activity with a known NF- κ B inhibitor, BAY 11-7082, and an NF- κ B-targeting artificial microRNA, miR533. The results revealed that all the labeled NF- κ B genes responded well to the two kinds of NF- κ B activity regulators in all cell lines.

Conclusion: This study thus obtained 15 cell lines with NF- κ B-ZsGreen fused genes, which provide an easy and visualizable tool for screening NF- κ B inhibitors and other NF- κ B-related studies.

Keywords: NF- κ B, TALEN, edit, HDR, ZsGreen

Introduction

Targeted integration of transgenes is usually achieved by a homologous recombination (HR)-mediated method.^{1,2} It requires a repair template that harbors left and right homology arms (HAs) (500–3000 bp), thus allowing precise insertion of large DNA fragments. To facilitate such targeted integration of transgenes, a targeted DNA double-strand break (DSB) has to be produced in the genome by using custom-designed nucleases, such as zinc-finger nuclease (ZFN),^{3–6} transcription activator-like effector nucleases (TALEN),^{2,7} and the clustered regularly interspaced short palindromic repeats (CRISPR)/CRISPR-associated protein-9 nuclease (Cas9).^{1,8} However, DSBs are lethal to cells so that it must be repaired by the cell's

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DNA repair machinery. DSBs can be repaired via imprecise nonhomology-based repair mechanisms, such as non-homologous end-joining (NHEJ), or by precise homology-dependent repair (HDR).⁹ HDR utilizes a template DNA that contains sequences homologous to the flanking sequences of the DSB.¹⁰ The homologous sequences are termed as homology arms. The HDR can use donors of variant forms including single-stranded DNA oligonucleotides (ssODNs) and double-stranded DNA (PCR fragments).¹¹ The PCR fragments with 35-bp homology arms can function as efficient donors for genome editing in mouse embryos and human cells.¹² The genome editing and HDR are now widely used to produce various chimeric antigen receptor (CAR) immunotherapeutic cells such as CAR-T. Additionally, other cell therapies also rely on gene modification to provide cells with particular functions.

Nuclear factor κ B (NF- κ B) is a family of DNA-binding transcription factors. The NF- κ B family consists of five members, RelA/p65, RelB, c-Rel, p50, and p52, which are coded by the genes of RELA, RELB, CREL, NF- κ B1, and NF- κ B2, respectively. These members are classified into two groups: a group of p50 and p52, and the other group of RelA, RelB, and c-Rel.¹³ The former has the DNA-binding domain (DBD) but no transcription activation domain (TAD); however, the latter has both DBD and TAD. The main functional and canonical dimer of NF- κ B family is p50-p65 heterodimer.¹³ NF- κ B plays various key roles in many physiological and pathological processes, such as immunity,^{13–15} cell proliferation,¹⁶ apoptosis, inflammation,^{17,18} and especially oncogenesis.¹⁹ Some studies indicate that many diseases are associated with the aberrant activation of NF- κ B and its signaling pathway.^{18,20} For instance, NF- κ B is abnormally activated in many human cancers to promote survival and malignancy by upregulating antiapoptotic genes.²¹ Therefore, many pharmaceutical companies and scientists have been committing to develop NF- κ B inhibitors for inflammation and cancer therapy.²²

Chemicals are important fields to develop inhibitors of NF- κ B activity. For instance, BAY 11–7082 is a widely used NF- κ B inhibitor that inhibits NF- κ B activity by inhibiting the phosphorylation of I κ B α .²³ In addition, microRNA (miRNA) is a kind of natural small RNA (~22 nt), which can regulate gene expression by blocking target mRNA translation or inducing target mRNA degradation.^{24–26} For finding the NF- κ B inhibitors, an effective cellular platform is essential, in which the change

of NF- κ B activity under the treatment of various potential drugs should be better facilely and quantitative monitored. For this purpose, various NF- κ B reporter stable cell lines with NF- κ B reporter construct that contains several repeats of NF- κ B binding sites, a minimal promoter upstream of the green fluorescent protein (GFP) and/or luciferase coding region were developed,^{27–33} which allows an assessment of NF- κ B activity via fluorescence microscopy, flow cytometry, and/or luminometry. However, these constructs are independent on the natural promoters of NF- κ B genes. They report a signal resulted from NF- κ B activity but not NF- κ B itself. In addition, the current platforms are only developed for NF- κ B RelA; however, NF- κ B family consists of five genes. RelA just forms the canonical NF- κ B signaling pathway, while others such as p52-RelB consist of a noncanonical NF- κ B signaling pathway.^{15,34} These different pathways were activated by different sets of stimulators and activate different sets of target genes.^{15,34} Therefore, more simple and comprehensive cell platforms are still in demand in the industry.

This study systematically edited all five genes of NF- κ B family in three different cell lines (293T, HepG2, PANC1) with both TALEN and CRISPR. The linear HDR donor containing ZsGreen coding sequence was used to repair the TALEN/CRISPR-produced DBSs. In this way, a ZsGreen coding sequence was fused to the ends of NF- κ B genes, which would produce ZsGreen-labeled NF- κ B proteins under the control of natural promoters of NF- κ B genes. The editing results were characterized by the fluorescent imaging of cells, quantitative analysis of cell fluorescence with flow cytometry, and quantitative PCR (qPCR) detection of NF- κ B-ZsGreen fused genes. The editing results were also evaluated by treating cells with an NF- κ B stimulator, TNF α . Finally, the positive cells were isolated from the edited cells with flow cytometry according to cellular fluorescence and further validated by treating cells with two kinds of NF- κ B inhibitors, chemical and microRNA inhibitors. This study thus obtained 15 cell lines with NF- κ B-ZsGreen fused genes, which provide an easy and visualizable tool for screening NF- κ B inhibitors and other NF- κ B-related studies.

Methods

TALEN and CRISPR Vectors

The TALEN and CRISPR expression plasmids that express TALEN and CRISPR/Cas9-sgRNA targeting five genes of NF- κ B family came from our recent study.³⁵ All

vectors used for cell transfection were purified from DH5 α using EndoFree Plasmid kits (QIAGEN, Germany) and quantified with a NanoDrop 2000c Spectrophotometer (Thermo Fisher Scientific, USA).

HDR Donor Preparation

The linear HDR donors that were used to fusing ZsGreen coding sequences to NF- κ B family genes were prepared by PCR amplification. Oligonucleotide ending with 35-nt sequences that were used as homology arms targeting NF- κ B family genes were used as primers (Table S1). A linear DNA fragment containing the Gslink-ZsGreen coding sequences was used as a template. The PCR reaction (30 μ L) contained 1 ng Gslink-ZsGreen DNA fragment (template), 10 μ M hom-F (Table S1), 10 μ M hom-R (Table S1), and 1 \times PrimeSTAR[®] HS DNA Polymerase (Takara, Japan). PCR program was 96°C 3 min, 28 cycles of 96°C 20 s, 58°C 20 s, and 72°C 1 min, and 72°C 5 min. The PCR products were purified with a Gel Extraction Kit (QIAGEN).

Cell Culture and Transfection

293T, HepG2, and PANC1 cells were obtained from the cell resource center of Shanghai institute for biological sciences, Chinese Academy of Sciences; China. 293T is a human embryonic kidney cell with stable gene sequence coding SV40 large T antigen, HepG2 is a human hepatoma cell, and PANC1 is a human pancreatic carcinoma cell. 293T is widely used in gene editing studies due to its high transfection efficiency. Hepatoma is a cancer that most seriously threatens the health and life of the Chinese. The pancreatic carcinoma is recognized as the king of carcinoma due to very limited therapy and high mortality. Therefore, the three cell lines were selected to use in this study. Cells were cultured with DMEM (HyClone, USA) containing 10% fetal bovine serum (FBS) (HyClone), 100 U/mL penicillin, and 100 μ g/mL streptomycin in a humidified incubator with 5% CO₂ for 37°C. Cells were seeded in a 24-well plate at a density of 5 \times 10⁴ cells/well and incubated for 24 h. The cells were co-transfected with 400 ng of TALEN vector (200 ng left TALEN and 200 ng right TALEN) or CRISPR/Cas9-sgRNA vector and 400 ng of HDR donor. The cell transfection was performed with lipofectamine 2000 (Invitrogen, USA) according to the manufacturer's instruction. The transfected cells were cultured for an additional 72 h. The cells were rinsed with PBS and imaged with a fluorescent microscope, IX51 with DP71 (Olympus, Japan), and quantitatively analyzed with a flow cytometer, BD Calibur (BD, USA). Ten thousand cells were analyzed in flow cytometry.

PCR Detection of Fused Genes

The genomic DNA (gDNA) was extracted from the edited cells by using a gDNA extraction kit (Tiangen, China). The gDNA was detected by quantitative PCR (qPCR) using a StepOnePlus[™] Real-Time PCR system (Applied Biosystems, USA). The fused NF- κ B genes were detected with forward primers (RELA-F, RELB-F, CREL-F, NFKB1-F, or NFKB2-F; Table S2) and a reverse primer (ZsGreen-R; Table S2). GAPDH gene was also detected as an internal control with primers GAPDH-F and GAPDH-R (Table S2). The PCR reaction (20 μ L) consisted of 100 ng of gDNA, 10 μ M forward primer, 10 μ M reverse primer, and 2 \times SYBR Mix (Roche, Switzerland). PCR program was 95°C 3 min, 40 cycles of 95°C 15 s, and 60°C 1 min. The PCR products were detected by agarose gel electrophoresis.

TNF α Stimulation of NF- κ B Fused Gene

The TALEN-edited cells were seeded in 24-well plates and incubated for 24 h at 37°C in 5% CO₂. The cells were then stimulated with TNF α (Sigma, USA) at a final concentration of 10 ng/mL for 1 h. The control cells were not treated by TNF α . The cells were then washed with cold PBS and collected by trypsinization. The collected cells were washed with cold PBS. The cells were fixed in formaldehyde (Sigma) at a final concentration of 1% (w/v) for 10 min at room temperature. The cross-linking was terminated by adding glycine at the final concentration of 0.125 M. The cells were precipitated by centrifugation at 1500 rpm for 10 min at 4°C. The cells were lysed with 500 μ L of lysis buffer (10 mM Tris-HCl, pH 7.4, 10 mM NaCl, 3 mM MgCl₂, and 0.1% IGEPAL CA-630). The nuclei were collected by centrifugation at 1500 rpm for 10 min at 4°C. The nuclei were washed twice with PBS. Finally, the nuclei were imaged with fluorescent microscope IX51 with DP71 (Olympus) and analyzed with flow cytometry (Calibur).

Screening of Positive Cells

The TALEN-edited cells were collected by trypsinization. The cells were resuspended in 200 μ L DMEM (HyClone) containing 2% fetal bovine serum (FBS) (HyClone), 100 U/mL penicillin, and 100 μ g/mL streptomycin. The cells were used to screen positive cells with green fluorescence through a sorter of flow cytometry (BD FACSAria[™] III, USA). The fluorescent cells were purified from one million edited cells. The screened cells (about

100–200 thousand) were cultured at 37°C for 48 h in a 24-well plate.

Validation of Screened Cells by NF- κ B Regulators

The positive cells cultivated in 24-well plates were treated with 50 μ M BAY 11–7082 (Abmole, USA) for 1 h. The cells were imaged with fluorescent microscope (IX51 with DP71) and analyzed with flow cytometry (Calibur). The positive 293T, HepG2, and PANC1 cells in 24-well plates were transfected with 800 ng of pDMP-miR533, respectively. The cells were cultured for an additional 72 h. The cells were imaged with a fluorescent microscope (IX51 with DP71) and analyzed with flow cytometry (Calibur).

PCR Detection of NF- κ B Target Gene Expression

The total RNA was purified from cells using the Trizol[®] reagent (Invitrogen). Then, 500 ng of total RNA was reversely transcribed into cDNA using PrimeScript[™] RT

Master Mix (Takara) according to the manufacturer's instructions in a total volume of 10 μ L. Triplicate samples per treatment were evaluated on a ABI Step One Plus (Applied Biosystems). The reaction mixture contained 2 μ L cDNA, 0.25 μ M of primers, and 10 μ L of Hieff qPCR SYBR Green Master Mix (Yeasen, China) in a total volume of 20 μ L. The primers used in qPCR were list in Table S3. PCR program was 95°C 5 min, 40 cycles of 95°C 10 s, and 60°C 30 s. The melting curve analysis was performed. The threshold value Ct values were normalized by subtracting the Ct values obtained for GAPDH. The Δ Ct values were then used to calculate relative changes of mRNA expression.

Results

Editing NF- κ B Family Genes in Three Cell Lines with Both TALEN and CRISPR

Five pairs of TALEN and five CRISPR expression plasmids were prepared by our recent study that developed an easy ready-to-use TALEN plasmid preparation pipeline.³⁵

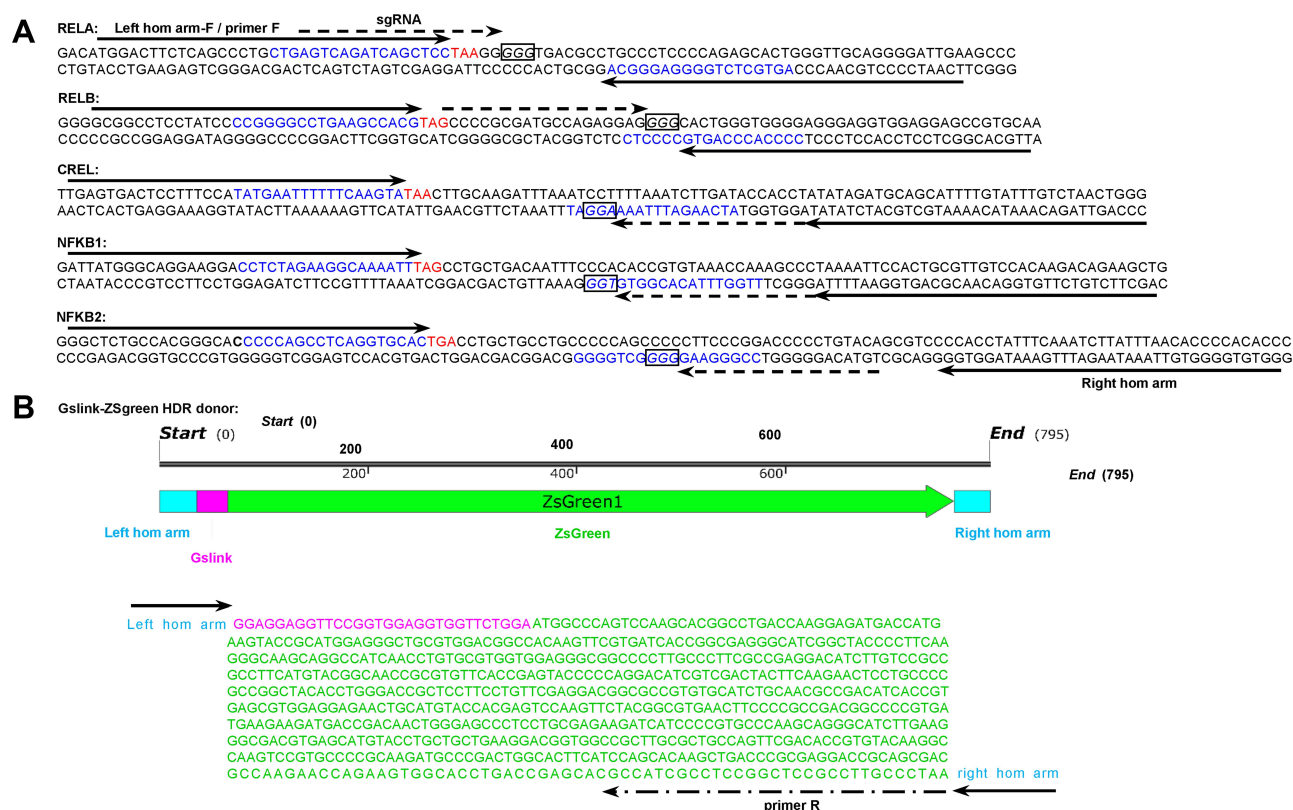


Figure 1 TALEN and CRISPR targets in NF- κ B genes and HDR donor. **(A)** The selected left and right homologous (hom) arms and TALEN and CRISPR targets in the NF- κ B family genes. Up strand: sense strand (from left to right: 5' to 3'); down strand: anti-sense strand (from left to right: 3' to 5'); bases labeled with a solid arrow: left hom arm/primer F and right hom arm; bases in blue: TALEN target sites; bases labeled with dotted arrow: sgRNA target sites; bases in red: stop codon; bases in square: PAM. **(B)** The HDR donor containing Gslink and ZsGreen coding sequences and ending with homology arms targeting to five NF- κ B family genes. The length of left and right hom arms is 35 nt.

Five pairs of TALEN expression plasmids were constructed to express the TALEN proteins that can produce double-strand breaks (DSBs) at the end of coding sequences of five NF- κ B family genes (Figure 1A). Five CRISPR expression plasmids were constructed to express Cas9 protein and single-guide RNA (sgRNA) that can produce DSBs at the same sites as TALENs (Figure 1A).

A linear HDR donor that contains a Gslink and ZsGreen coding sequences and ends with 35-nt homologous (hom) arms was used as a template for repairing the DSBs (Figure 1B).

Five NF- κ B family genes (RELA, RELB, CREL, NF- κ B1, NF- κ B2) were systematically edited in three different cell lines (293T, HepG2 and PANC1) by co-transfecting cells

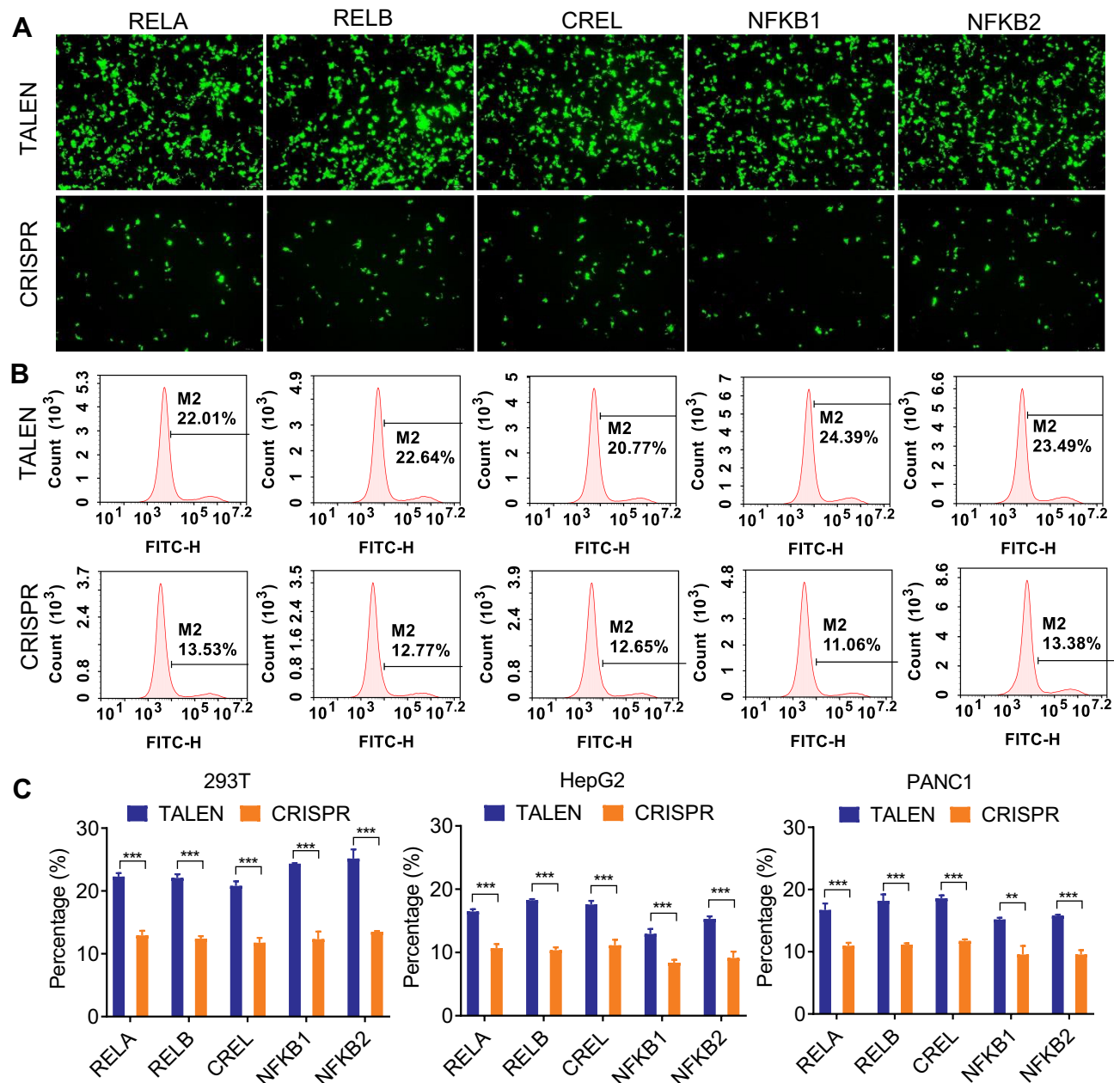


Figure 2 Editing NF- κ B genes in three cell lines with TALEN and CRISPR. Linear dsDNA fragments containing a Gslink-ZsGreen-coding sequence and ending with homology arms homologous to NF- κ B family genes were used as HDR donors. (A) Representative fluorescent images of 293T cells after the TALEN and CRISPR editing and HDR. The representative fluorescent images of HepG2 and PANC1 cells after the TALEN and CRISPR editing and HDR are shown in Figure S1A and S2A. The scale bar is 100 μ m. (B) Flow cytometry assay of 293T cells after the TALEN and CRISPR editing and HDR. The flow cytometry assays of HepG2 and PANC1 cells after the TALEN and CRISPR editing and HDR are shown in Figure S1B and S2B. (C) The statistical analysis of flow cytometry assays of three biological replicates. The results are shown as mean values \pm standard deviation (SD). The Student t test was performed to assess the statistical significance. *** $p < 0.001$; ** $p < 0.01$.

with TALEN/CRISPR and HDR donor. The successful editing and HDR would produce an NF- κ B-ZsGreen fused gene in cells, which allows cells to express an NF- κ B-ZsGreen fused protein under the control of natural promoter of NF- κ B gene. By directly imaging cells with fluorescent microscope and quantitatively analyzing cells with flow cytometry, the editing efficiency can be evaluated. The results indicated that all five genes were successfully edited by both TALEN and CRISPR in all cells (Figure 2, Figure S1, and Figure S2).

However, both fluorescent imaging (Figure 2A, Figure S1A, and Figure S2A) and flow cytometry quantitative assay (Figure 2B, Figure S1B, and Figure S2B) revealed that TALEN had higher editing efficiency than CRISPR. On average, TALEN edited 22.2%, 15.5%, and 17% cells in 293T, HepG2, and PANC1 cells, respectively, while CRISPR edited 13.2%, 9.7%, and 10.3% cells in 293T, HepG2, and PANC1 cells, respectively (Figure 2C). The editing success and efficiency were also confirmed by the

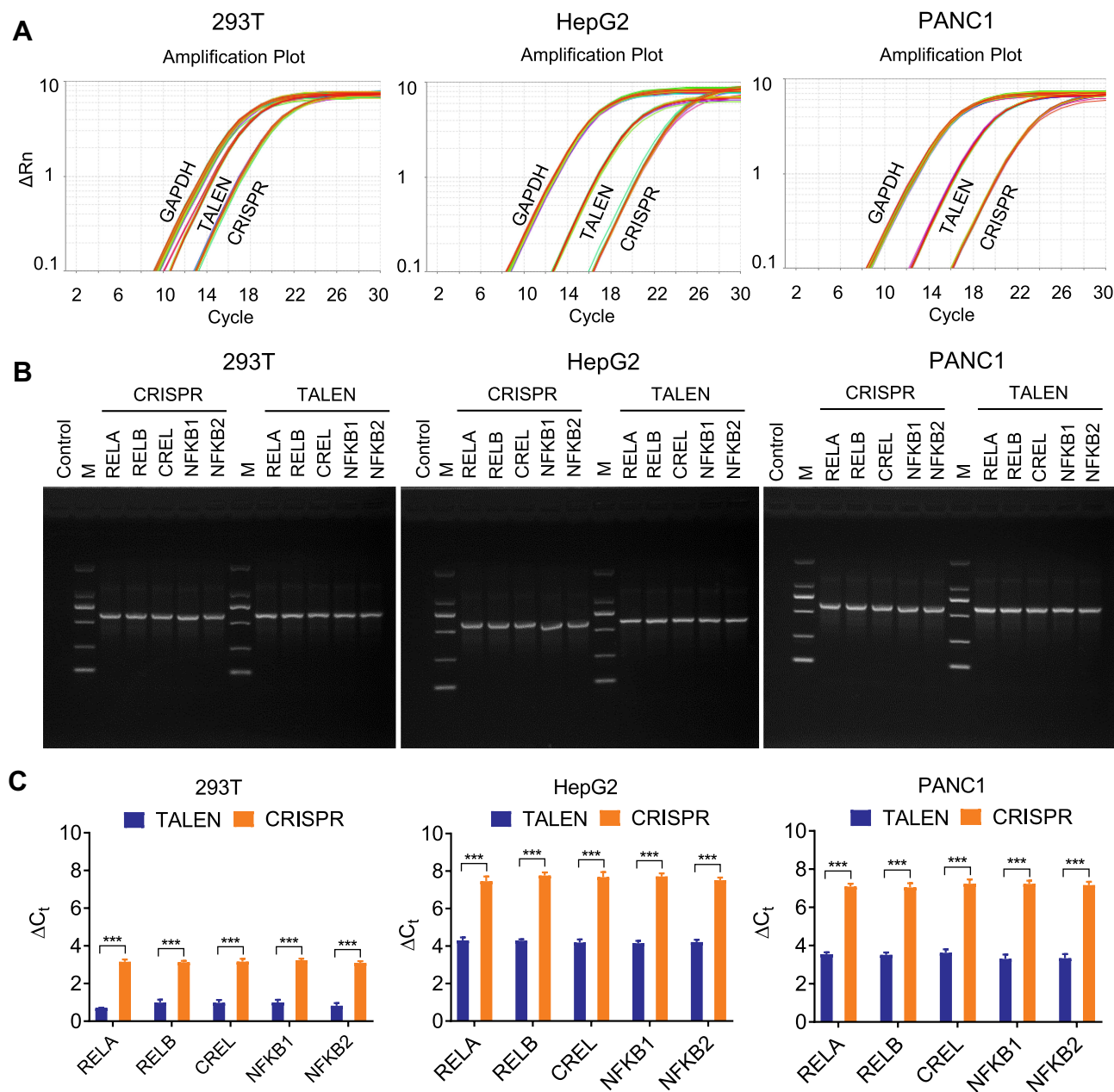


Figure 3 Detection of editing efficiency of TALEN and CRISPR with qPCR. A same amount of gDNA (100 ng) from all cells after the TALEN and CRISPR editing and HDR was used to detect the fused NF- κ B-ZsGreen genes by qPCR. (A) The amplification plots of fused NF- κ B-ZsGreen genes. (B) Visualization of qPCR products with agarose gel electrophoresis. (C) The statistical analysis of qPCR detection. ΔC_t was calculated by $C_{t(NF-\kappa B-ZsGreen)} - C_{t(GAPDH)}$. The results are shown as mean values \pm standard deviation (SD). The Student *t* test was performed to assess the statistical significance. ****p* < 0.001.

subsequent qPCR detection of NF- κ B-ZsGreen fused genes in gDNA (Figure 3).

To further validate the edited NF- κ B genes, the TALEN-edited cells were treated with an NF- κ B stimulator, TNF α . Because the activated NF- κ B proteins will translocate into nuclear, the nuclei were isolated from the TNF α -treated and -untreated cells. The isolated nuclei were analyzed by both fluorescent imaging and flow cytometry. The results indicated that the TNF α treatment

induced a significant increase of fluorescent nuclei in all cells (Figure 4, Figure S3, and Figure S4).

Screening of Positive Cells and Treatment

To obtain the pure positive cells, the TALEN-edited cells were sorted with flow cytometry according to ZsGreen fluorescence. The results indicated that the pure positive cells could be efficiently obtained from TALEN-edited cell lines (Figure S5 and Figure 5). Therefore, the pure positive

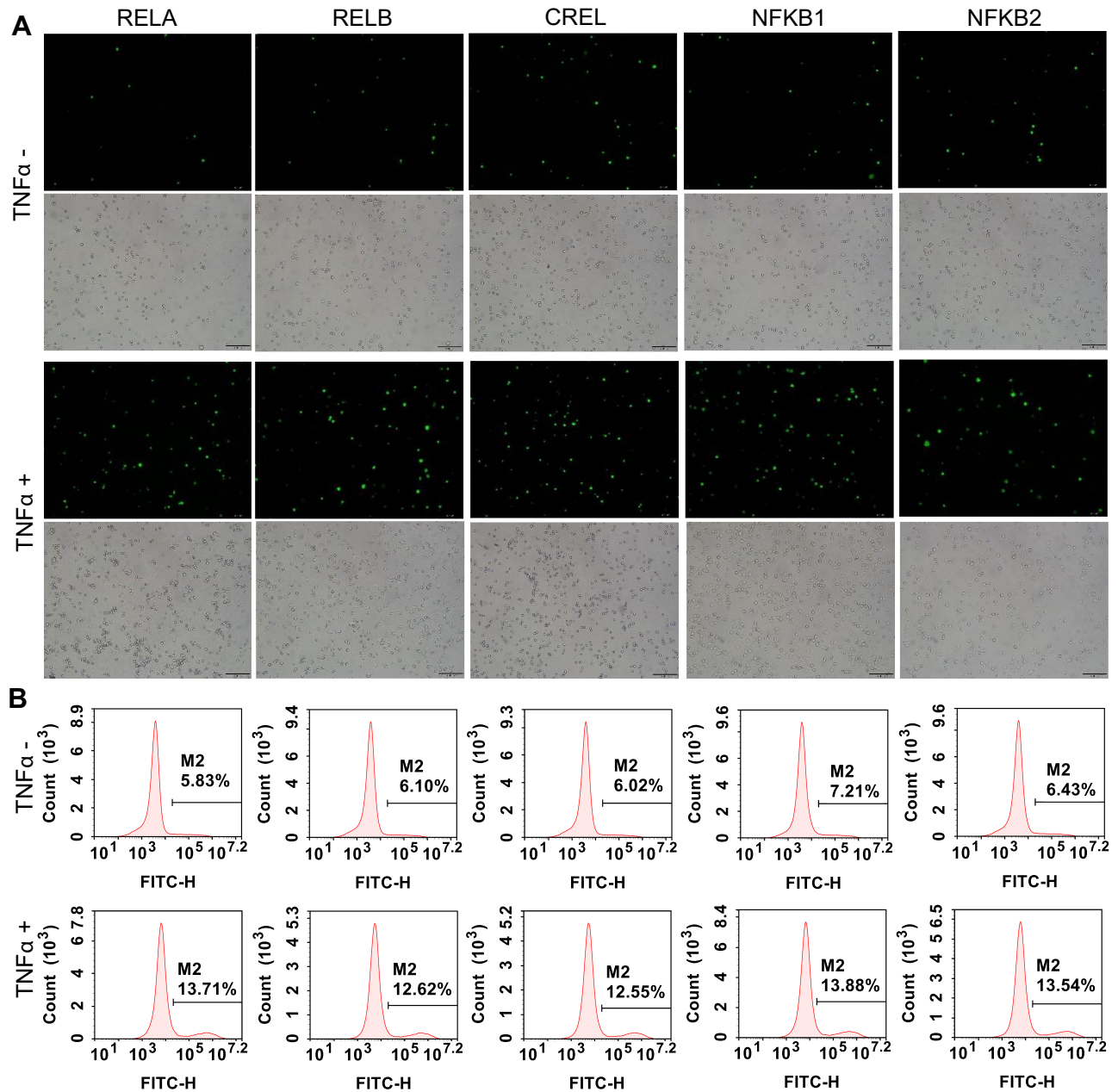


Figure 4 TNF α stimulation to the TALEN-edited cells. The TALEN-edited cells were treated by TNF α , an NF- κ B stimulator, and the nuclei were isolated from the treated cells. **(A)** Representative fluorescent images of the isolated nuclei of 293T cells. The representative fluorescent images of the isolated nuclei of HepG2 and PANC1 cells are shown in Figure S3A and S4A. The scale bar is 100 μ m. **(B)** Flow cytometry assay of the isolated nuclei of 293T cells. The flow cytometry assays of the isolated nuclei of HepG2 and PANC1 cells are shown in Figure S3B and S4B.

cells were obtained for five NF- κ B family genes from three cell lines (15 cell lines in total) (Figure 5).

To validate the potential application of these purified cell lines in screening of NF- κ B inhibitors, all cell lines were treated with two kinds of representative NF- κ B inhibitors, a small molecular chemical, BAY 11-7082, and a microRNA, miR533. The results indicated that both BAY 11-7082 and miR533 significantly decreased the number of fluorescent cells in all 15 cell lines (Figure 6, Figure 7, and Figure S6–S9).

To further validate the response of stable cells to the treatment of NF- κ B inhibitors, the expression of some typical NF- κ B target genes in the BAY 11-7082-treated and -untreated stable HepG2 and PANC1 cells with RELA-ZsGreen gene was detected by qPCR. The detected NF- κ B target genes included CCL2, IL1A, IL1B, NFKB1, NFKBIA, and PTX3. These genes are all NF- κ B direct target genes (NF- κ B Target Genes, Boston University Biology, <https://www.bu.edu/nf-kb/gene-resources/target-genes/>). The results indicated

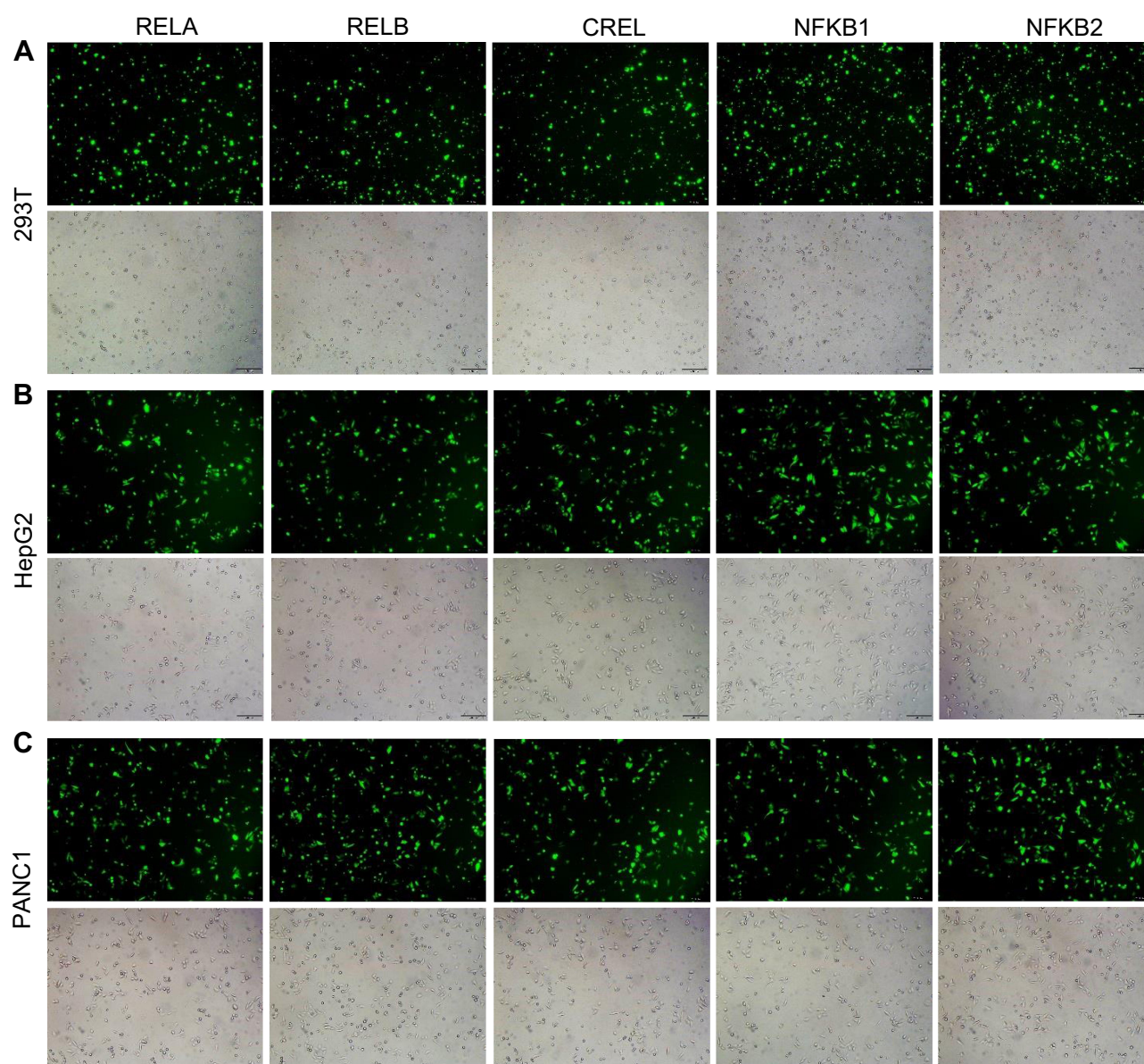


Figure 5 Fluorescent imaging of flow cytometry-sorted positive cells. (A) 293T cells. (B) HepG2 cells. (C) PANC1 cells. The scale bar is 100 μ m. The flow cytometry assays of the sorted cells are shown in Figure S5.

that the BAY 11-7082 treatment to the stable cells leads to the significant downregulation of these genes ([Figure S10](#)).

Discussion

NF- κ B is a key transcription factor involved in inflammation and cancer. To provide an easy and visualizable tool

for screening NF- κ B inhibitors and other NF- κ B-related studies, this study edited all five genes of NF- κ B family in three different cell lines with both TALEN and CRISPR. The edited NF- κ B genes were repaired by HDR using a linear homologous donor containing ZsGreen coding sequence. In this way, the NF- κ B proteins were labeled by fluorescent ZsGreen in cells. By detecting the edited

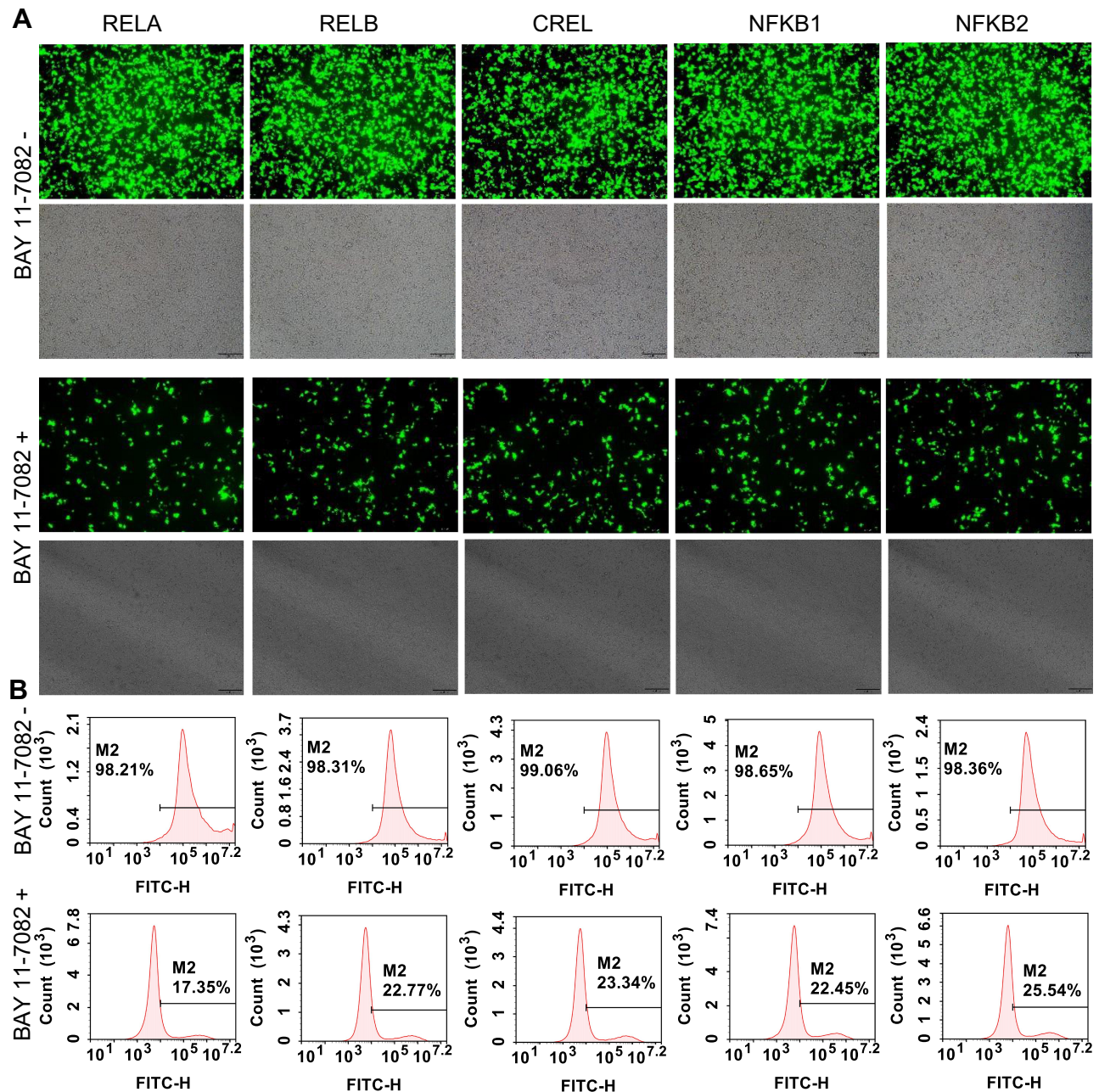


Figure 6 BAY 11-7082 treatment to the flow cytometry-sorted positive cells. **(A)** Representative fluorescent images of BAY 11-7082-treated and -untreated 293T cells. The representative fluorescent images of the BAY 11-7082-treated and -untreated HepG2 and PANC1 cells are shown in Figure S6A and S7A. The scale bar is 100 μ m. **(B)** Flow cytometry assay of BAY 11-7082-treated and -untreated 293T cells. The flow cytometry assay of BAY 11-7082-treated and -untreated HepG2 and PANC1 cells are shown in Figure S6B and S7B.

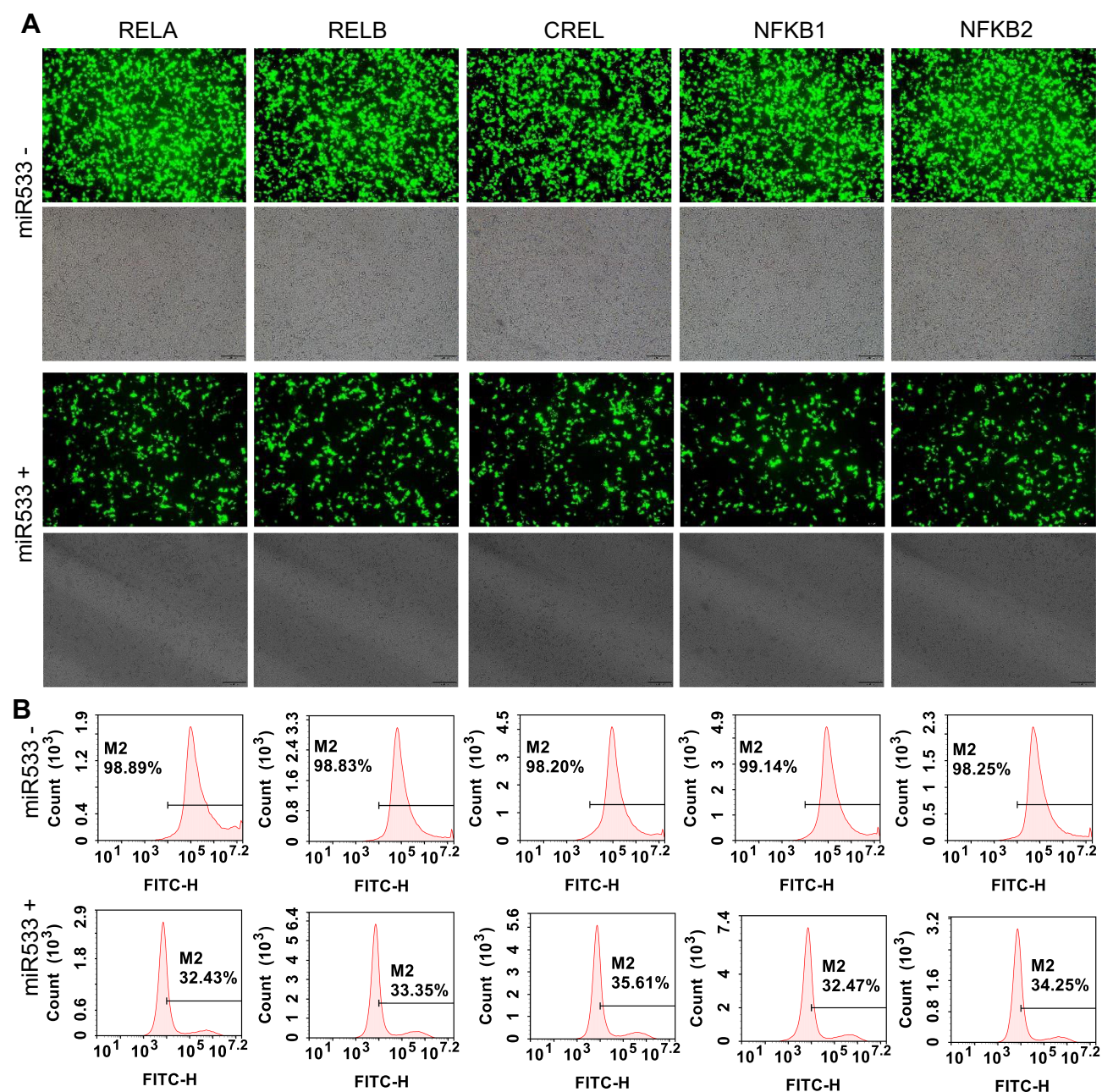


Figure 7 MiR533 treatment to the flow cytometry-sorted positive cells. **(A)** Representative fluorescent images of miR533-treated and -untreated 293T cells. The representative fluorescent images of the miR533-treated and -untreated HepG2 and PANC1 cells are shown in Figure S8A and S9A. The scale bar is 100 μ m. **(B)** Flow cytometry assay of miR533-treated and -untreated 293T cells. The flow cytometry assay of miR533-treated and -untreated HepG2 and PANC1 cells are shown in Figure S8B and S9B.

cells with fluorescent imaging, flow cytometry, and qPCR, it was demonstrated that TALEN had higher editing efficiency than CRISPR in editing all genes in all cells. The positive cells were easily isolated from the TALEN-edited cells by flow cytometry. The purified positive cells responded well to two kinds of known NF- κ B activity regulators in all cell lines. As a result, this study obtained 15 cell lines with five NF- κ B-ZsGreen fusing genes,

providing useful cellular platforms for screening potential NF- κ B-target drugs.

In our recent study, we developed an easy and rapid pipeline for preparing ready-to-use TALEN expression plasmids.³⁵ Using the pipeline, this study constructed the TALEN expression plasmids that can express the TALEN proteins targeting five genes of NF- κ B family. By the quantitative analysis of cell fluorescence with flow

cytometry and the edited genes with qPCR, this study demonstrated that all genes were more effectively edited by TALEN than CRISPR in all cells, indicating that TALEN is still a valuable genome editing tool with better performance than CRISPR. The high editing efficiency is very important for a gene editing tool. Additionally, TALEN can be used to edit any sequence in genome, whereas CRISPR is limited by the requirement of protospacer adjacent motif (PAM). Moreover, TALEN has high targeting performance due to cutting a target by a pair of TALEN; however, CRISPR is still challenged by its potential off-target in application. As previously reported, this study also demonstrated that a linear DNA fragment ended with short homology arm (35 nt) can be used as an efficient HDR donor in genome editing. This kind of linear HDR donor can be facilely prepared by PCR amplification using primers ended with 35-nt homology arms.

This study used miR533 to represent an NF- κ B inhibitor of microRNA. This microRNA is an artificial microRNA, which was designed and verified by our previous work.³⁶ In this study, this microRNA was expressed by a plasmid that contains an NF- κ B-specific promoter and a downstream microRNA expression sequence. The NF- κ B-specific promoter consists of an NF- κ B decoy sequence and minimal promoter. The microRNA expression sequence can transcribe a microRNA targeting RELA coding sequence. This study showed that the expression of other four NF- κ B genes other than RELA could be also inhibited by this microRNA. The reason is that the NF- κ B genes RELB,³⁷ CREL,^{38,39} NF- κ B1,⁴⁰ and NF- κ B2⁴¹ are the direct target genes of RELA (NF- κ B Target Genes, Boston University Biology, <https://www.bu.edu/nf-kb/gene-resources/target-genes/>).

Conclusions

This study edited five NF- κ B family genes in three cell lines with TALEN and CRISPR. The edited genes were fused to the ZsGreen coding sequences by HDR. The results indicated that TALEN had higher editing efficiency than CRISPR in editing all genes in all cells. The screened positive cells formed 15 cell lines with five NF- κ B-ZsGreen fusing genes. These cell lines could respond to variant NF- κ B inhibitors and thus become useful platforms for screening potential NF- κ B-target drugs.

Abbreviations

TALE, transcription activator-like effector; TALEN, TALE nuclease; DBD, DNA-binding domain; TAD, transcription activation domain; HR, homologous recombination; HAS,

homology arms; ZFN, zinc-finger nuclease; CRISPR, clustered regularly interspaced short palindromic repeats; sgRNA, single-guided RNA; ssODNs, single-stranded DNA oligonucleotides; CAR, chimeric antigen receptor; PCR, polymerase chain reaction; qPCR, quantitative PCR; gDNA, genomic DNA; NF- κ B, nuclear factor kappa B; NHEJ, non-homologous end joining; HDR, homology-dependent repair; DSB, double-strand break; TF, transcription factor; FBS, fetal bovine serum.

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Disclosure

The authors report no conflicts of interest in this work.

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