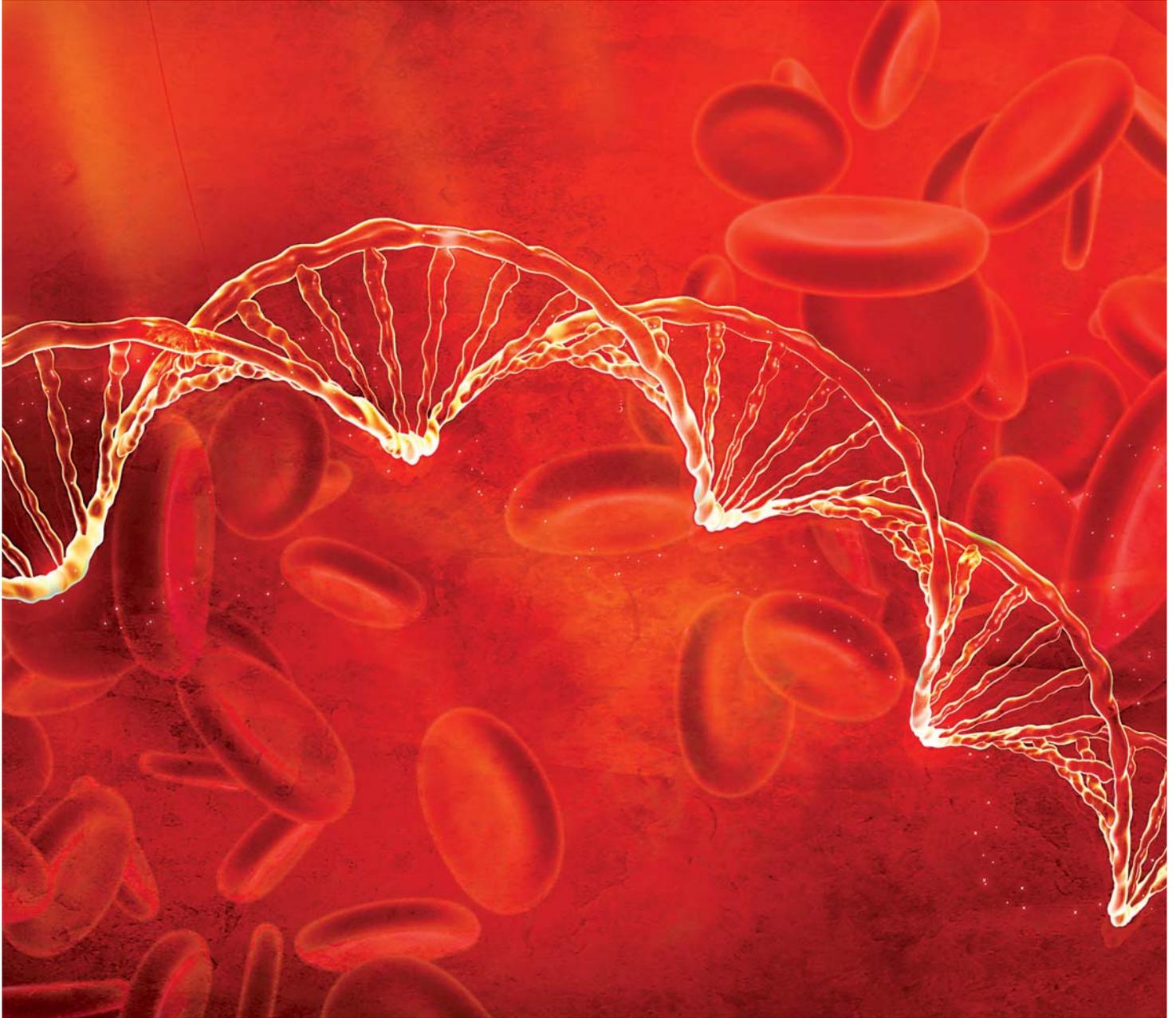


invitrogen™
by *life* technologies™



Gateway® cloning technology

The easy-to-use choice for everyday cloning

life
technologies™

The trusted leader in cloning technology

Gateway® cloning technology has been cited by life science researchers more than 1,500 times. It's no wonder Gateway® cloning has been the go-to choice for years, by researchers with varying experience—from beginners to advanced—for protein expression, functional analysis, and much more.

Circumvent the roadblocks of traditional restriction enzyme cloning—no need for ligase, subcloning steps, or the hours spent to screen countless colonies. Experience Gateway® cloning technology.

- Fast reactions—1-hour room-temperature cloning reactions
- Accurate results—cloning reactions achieve >95% efficiency to deliver the clone you need
- Versatile technology—easily shuttle DNA material/insert from vector to vector
- Streamlined protocol—no need for resequencing; use the same clone from target identification to validation



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Basic cloning protocol: three steps to better efficiency

Entry clones, enzymes, and Destination vectors

Determine the Entry clone

The Entry clone is how and where you start your experiment, as it contains your gene of interest or DNA fragment flanked by *attL* sequences, which are then used to recombine with *attR* sequences to create your desired expression clone. Choose one of our TOPO[®] cloning vectors to create your Entry clone, or purchase a premade clone from our validated Ultimate[™] ORF Clone Collection.

Mediate the reaction with Clonase[®] enzymes

Once the Entry clone is ready, the gene of interest is easily shuttled to a secondary plasmid, the Destination vector. This reaction is mediated by LR Clonase[®] enzyme mix, which contains the protein machinery necessary to excise the gene of interest from the Entry clone and integrate it into the Destination vector, which then becomes your expression clone. Reversing this reaction is simple: it requires a BP reaction (recombination between *attB* and *attP* sites) using BP Clonase[®] enzyme mix.

Both LR Clonase[®] and BP Clonase[®] enzyme mixes are supplied in easy-to-use master mix formats, ensuring consistency and reliability from reaction to reaction.

Select the Destination vector

Once you clone your gene of interest or DNA fragment into a Gateway[®] vector, you can shuttle it to as many expression and functional analysis systems as you need.

The diverse selection of expression vectors available with Gateway[®] cloning technology is vast and broad. From expression proteins in *E. coli*, yeast, insect, or mammalian cells to RNAi studies, from crystallography to protein–protein interaction functional studies, there is a Destination vector for your application. And for those applications that require a specialized or customized vector, the Gateway[®] Vector Conversion System can convert any vector into a Gateway[®] cloning–compatible vector.

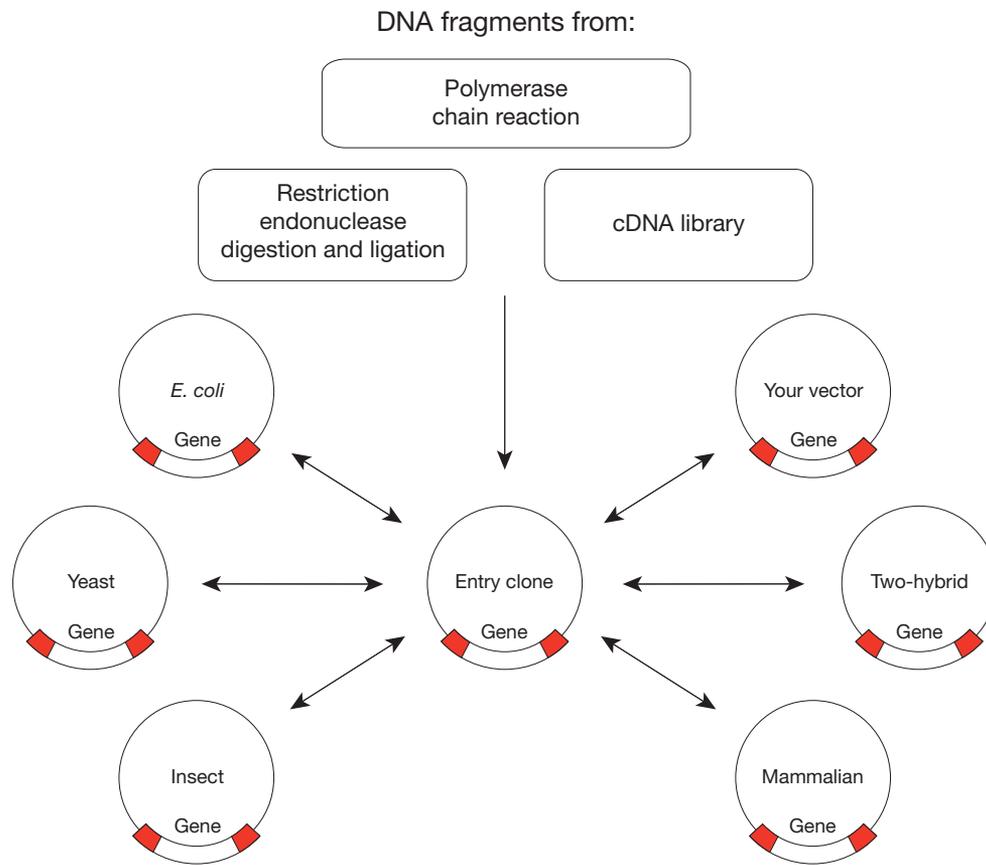


Figure 1. Gateway® technology facilitates cloning of genes into and back out of multiple vectors via site-specific recombination. Once a gene is cloned into an Entry clone you can then move the DNA fragment into one or more Destination vectors simultaneously.



Product selection guide

Learn which products to implement at each stage

Creating an Entry clone

Using TOPO® vectors and PCR amplification/restriction-enzyme vectors are the most common ways to construct your own Entry clone.

TOPO® vectors—both options offer 5-minute cloning and >95% efficiency

pCR®8/GW/TOPO® Vector Kits

- Convenient sequencing
- Robust selection in *E. coli* with spectinomycin resistance
- Easy excision of insert DNA with flanking EcoRI sites

pENTR™/D-TOPO® Vectors

- Fast Directional TOPO® cloning
- Delivers insert in correct orientation
- Contains necessary *attL* sequences for recombination into any Destination vector
- Select versions carry a TEV protease cleavage site for producing native proteins after expression

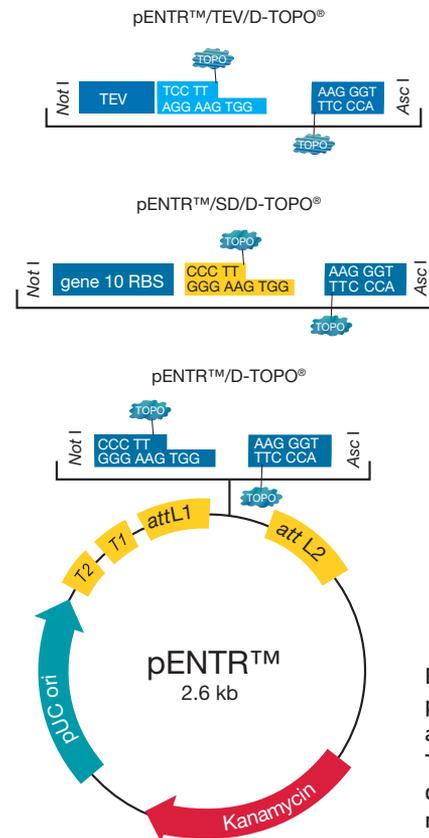


Figure 2. Several pENTR™ vectors are available for directional TOPO® cloning and direct access to the multitude of Gateway® expression vectors.

Purchasing a premade clone

You can also utilize Gateway® technology with a ready-to-use clone from our extensive clone collection. The Ultimate™ ORF Clone Collection consists of high-quality, full-insert sequenced human and mouse open reading frames already cloned into the pENTR™ 221 Gateway® Entry vector for limitless downstream analysis capabilities. Clones contain

DNA- and amino acid sequence-verified, expression-ready cDNAs, including kinases, G-protein-related, phosphatases, ion channels, GPCRs, chemokines, nuclear receptors, and cytokines.

View the Ultimate™ ORF Clone Collection at lifetechnologies.com/orf

PCR amplification or restriction-enzyme cloning vectors

pDONR™ and pENTR™ Vectors

These vectors allow you to clone a PCR product amplified with primers containing *attB* sequences (pDONR™) or specific restriction sites (pENTR™). Using PCR to generate the Entry clone, two short artificial *attB* sequences (*attB1* and *attB2*) must flank your gene of interest and be added to specific primers that are used to amplify the gene of choice. The DNA fragment is combined with a donor vector that contains *attP1* and *attP2* sequences and with BP Clonase® II enzyme.

- >90% of the colonies contain the Entry clone with the gene of interest in the correct orientation
- Final Entry clones are ready for recombination with any Gateway® Destination vector

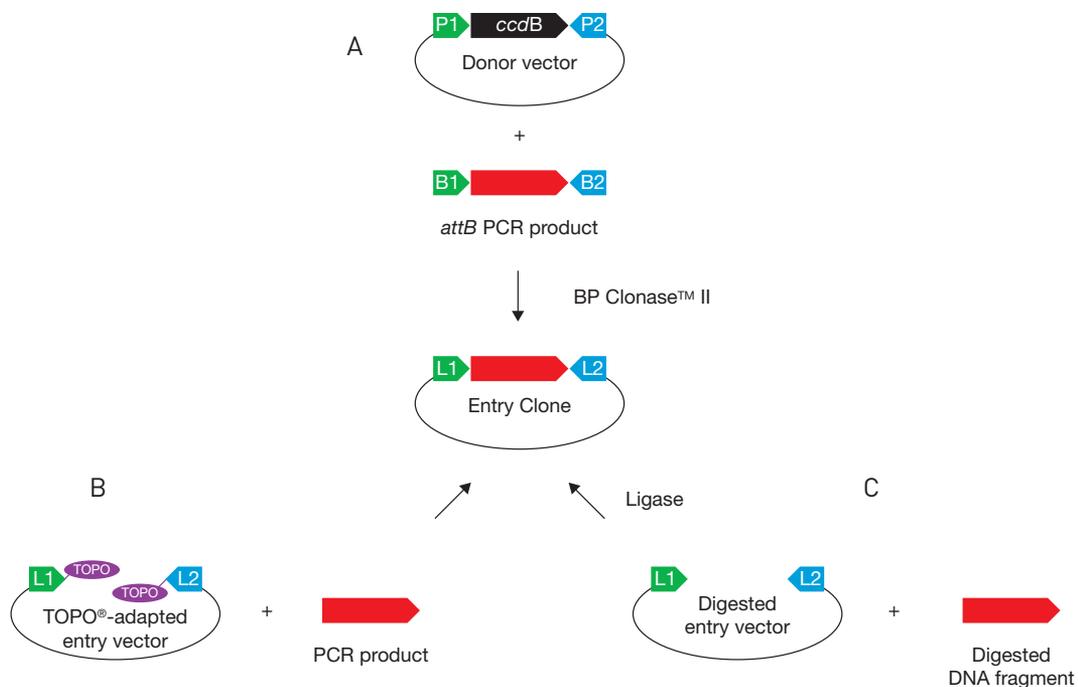


Figure 3. Strategies to build the Entry clone. The three possible methods that lead to the Entry clone are depicted: (A) BP cloning, (B) TOPO® cloning, and (C) restriction enzyme and ligase cloning. Red arrows represent the fragment of interest. Adapted from Katzen F (2007) *Expert Opin Drug Discov* 2(4):571–589.



Clonase[®] enzyme mix selection guide

| | BP Clonase [®] II Enzyme Mix | LR Clonase [®] II Plus Enzyme Mix |
|--|--|--|
| Application | Creating Entry clones | Creating expression clones |
| Proteins involved in site-specific recombination | <ul style="list-style-type: none"> • Int (integrase) • IHF (integration host factor) | <ul style="list-style-type: none"> • Int (integrase) • IHF (integration host factor) • Xis (excisionase) |
| Activity | <ul style="list-style-type: none"> • DNA recombinase • DNA binding protein • High efficiency for Entry clone construction • Single-mix format eliminates pipetting steps and hands-on errors | <ul style="list-style-type: none"> • DNA recombinase • DNA-binding protein • Highest cloning efficiency for single- and multiple-fragment cloning • Optimized for difficult cloning reactions • Works with MultiSite Gateway[®] Pro technology |
| Advantages | <ul style="list-style-type: none"> • Easy-to-use, single-mix format ensures enzyme stability • Convenient 10 μL reaction setup | <ul style="list-style-type: none"> • Easy-to-use, single-mix format ensures enzyme stability • Convenient 10 μL reaction setup |

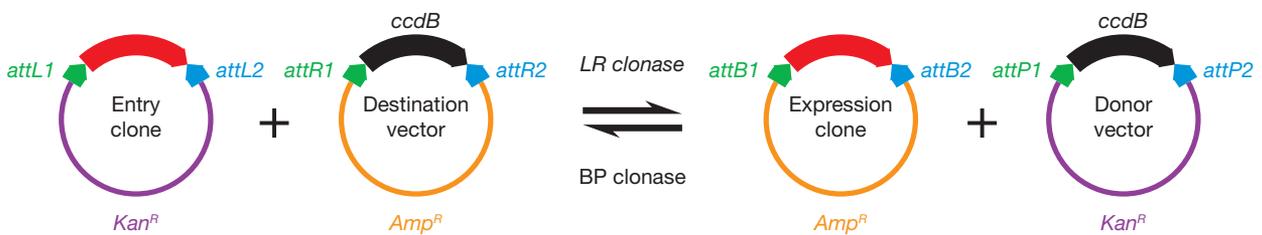


Figure 4. The Gateway[®] reactions. The scheme shows the four types of plasmids and enzyme mixes involved in Gateway[®] cloning reactions. Red arrows represent the fragment of interest. Adapted from Katzen F (2007) *Expert Opin Drug Discov* 2(4):571–589.

Destination vector selection guide

Gateway® cloning technology is especially noted for its utility in protein expression. The flexibility and diverse selection of Destination vectors and host systems is particularly attractive for multidisciplinary protein expression studies.

Destination vectors for protein expression

| Host system for protein expression | Gateway® Destination vector family |
|---|--|
| <i>E. coli</i> | <ul style="list-style-type: none">• pDEST™ 14, 15, 17, and 24• pET160 and pET161 DEST vectors |
| Yeast | pYES2-DEST52 |
| Insect cells | BaculoDirect™ C-Term Expression Kit |
| Mammalian cells (constitutive expression) | pcDNA™ mammalian expression vector family |
| Mammalian cells (regulated expression) | pT-REx-DEST30 and pT-REx-DEST31 vectors |
| Mammalian cells (viral delivery) | ViraPower™ Lentiviral Expression Systems |

Destination vectors for additional application areas

| Application | Gateway® Destination vector family |
|-------------------------------------|---|
| Antibody or antigen production | Champion™ pET Expression systems |
| Localization | VividColors™ pcDNA™ GFP Destination vector family |
| Protein array | Expressway™ Plus Expression System |
| Protein-protein interaction studies | ProQuest™ Two-Hybrid System using Gateway® technology |
| Reporter assay | GeneBLAzer® pcDNA™ vector family |
| RNAi | GeneBLAzer® pcDNA™ vector family |



MultiSite Gateway[®] Pro Kits

Mix and match fragments while maintaining orientation

MultiSite Gateway[®] Pro Kits

What if you could easily and accurately assemble multiple DNA fragments in the order and orientation of your desire? This approach, called MultiSite Gateway[®] Pro technology, allows the mixing and matching of functional fragments in a concerted fashion to generate multi-segment constructs. MultiSite Gateway[®] Pro technology enables you to perform pathway reconstitution, multiple gene expression and regulation, protein interaction studies, and more.

This approach has several applications covering the engineering of proteins, pathways, and cells, and provides a highly flexible platform for functional analysis.

The full power of Gateway[®] cloning technology is realized with MultiSite Gateway[®] Pro kits, which allow for the simultaneous assembly of multiple fragments into a single vector in a predefined order, orientation, and reading frame (Figures 5 and 6).

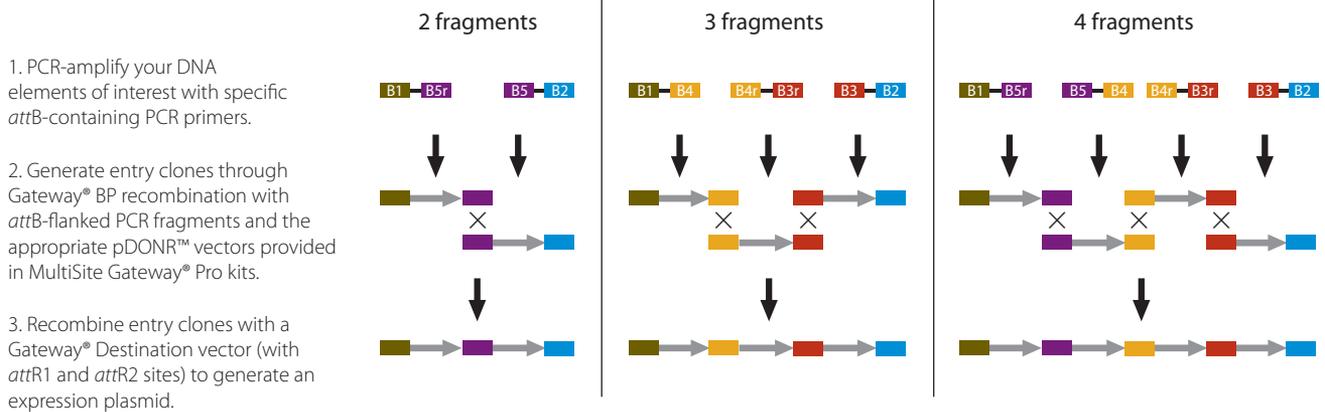


Figure 5. How MultiSite Gateway[®] Pro technology works.

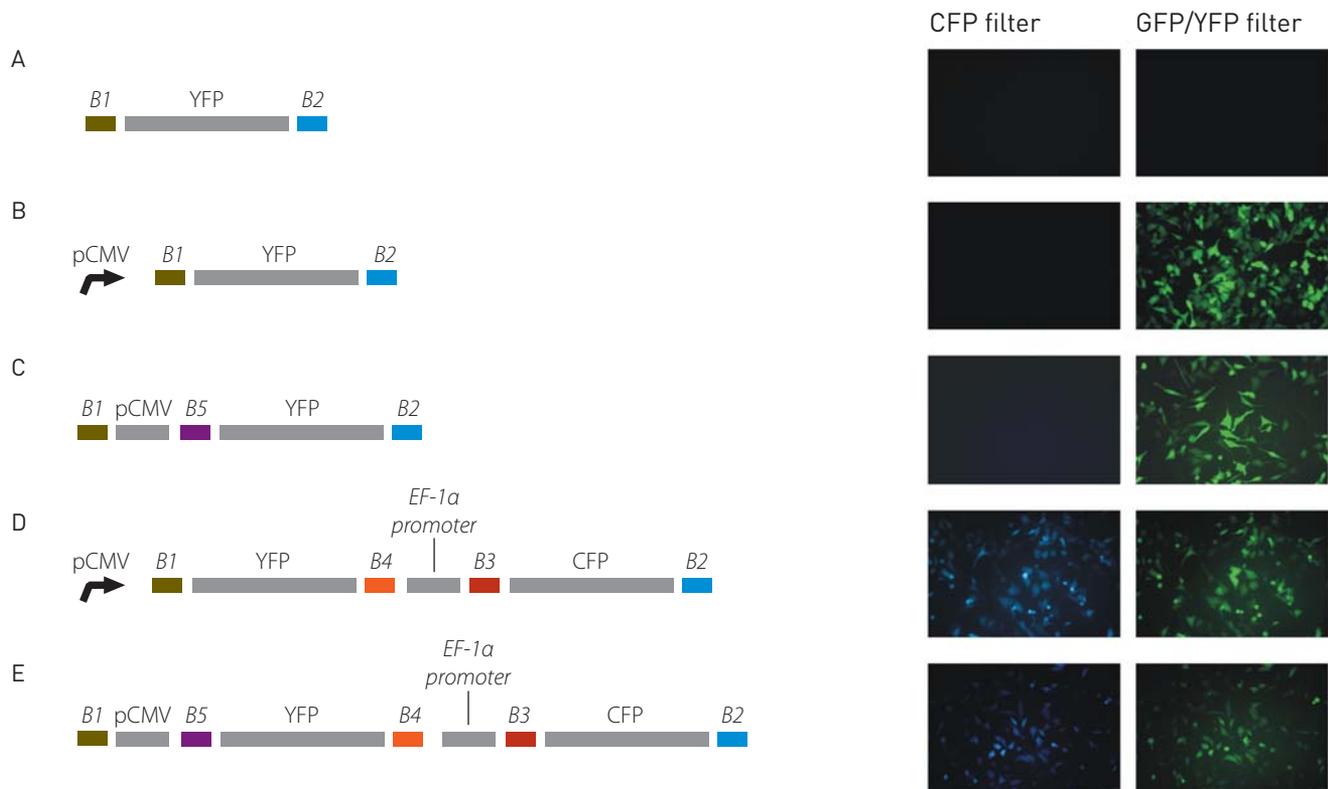


Figure 6. An example of using MultiSite Gateway® Pro technology to study expression of multiple genes in human cells. Entry clones containing genes for YFP and CFP and the CMV and EF-1α promoters were recombined into pcDNA™ 6.2/V5-PL-DEST [A, C, and E] or into pcDNA™ 6.2/V5-DEST [B and D]. The resulting expression clones were used to transfect HeLa cells. Expression was verified under a fluorescence microscope. The plasmid pcDNA™ 6.2/V5-PL-DEST is a promoterless derivative of pcDNA™ 6.2/V5-DEST, which carries the CMV promoter.



Ordering information

TOPO® TA cloning

| Product | Description | Quantity | Cat. No. |
|-------------------------------|---|----------|----------|
| pCR™8/GW/TOPO® TA Cloning Kit | Efficient TOPO® TA cloning kit simplifies Entry clone construction | 20 rxns | K250020 |
| pCR™8/GW/TOPO® TA Cloning Kit | Efficient TOPO® TA cloning with fast-growing competent <i>E. coli</i> that shortens the time for Entry clone construction | 20 rxns | K252020 |
| pCR™8/GW/TOPO® TA Cloning Kit | Efficient TOPO® TA cloning with fast-growing competent <i>E. coli</i> and plasmid purification drastically shortens and simplifies Entry clone construction, saving time and hassle | 20 rxns | K252002 |

Directional TOPO® cloning

| Product | Description | Quantity | Cat. No. |
|--------------------------------|--|----------|----------|
| pENTR™/D-TOPO® Cloning Kit | Directional TOPO® cloning kit that produces expression-ready Entry clones | 20 rxns | K240020 |
| pENTR™/SD/D-TOPO® Cloning Kit | Directional TOPO® cloning kit, including the Shine-Dalgarno sequence that creates an <i>E. coli</i> expression-ready Entry clone | 20 rxns | K242020 |
| pENTR™/TEV/D-TOPO® Cloning Kit | Directional TOPO® cloning kit that creates expression-ready Entry clones with 5' TEV sequence for N-terminal tag removal (creating native proteins) | 20 rxns | K252520 |
| pENTR™/TEV/D-TOPO® Cloning Kit | Directional TOPO® cloning kit with fast-growing competent <i>E. coli</i> that shortens the time for Entry clone construction while creating expression-ready Entry clones with a 5' TEV sequence for N-terminal tag removal creating native proteins | 20 rxns | K253520 |

PCR cloning using BP recombination

| Product | Description | Quantity | Cat. No. |
|---|---|----------|-----------|
| PCR Cloning System with Gateway® technology | Complete kit for directional cloning into a Gateway® vector with pDONR™ 221 vector with kanamycin selection | 20 rxns | 12535029 |
| PCR Cloning System with Gateway® technology | Complete kit for directional cloning into a Gateway® vector with pDONR™/Zeo vector with Zeocin™ selection | 20 rxns | 125350237 |
| pDONR™ 221 Vector | | 6 µg | 12536017 |
| pDONR™ Zeo Vector | | 6 µg | 12535035 |

Restriction enzyme cloning

| Product | Description | Quantity | Cat. No. |
|------------------|---|----------|----------|
| pENTR™ 1A Vector | Restriction enzyme cloning vector that produces in-frame (rf = 0), expression-ready Entry clones, including both Shine-Dalgarno and Kozak sequences | 10 µg | 11813011 |
| pENTR™ 2B Vector | Restriction enzyme cloning vector that produce in-frame (rf = +1), expression-ready Entry clones | 10 µg | 11816014 |
| pENTR™ 3C Vector | Restriction enzyme cloning vector that produce in-frame (rf = +2), expression-ready Entry clones | 10 µg | 11817012 |
| pENTR™ 4 Vector | Same as pENTR™ 1A Vector except with NcoI instead of DraI in MCS that produces in-frame (rf = 0), expression-ready Entry clones | 10 µg | 11818010 |
| pENTR™ 11 Vector | Same as pENTR™ 1A Vector except with NspV instead of DraI in MCS that produces in-frame (rf = 0), expression-ready Entry clones | 10 µg | 11819018 |

Multi-fragment assembly with Gateway® technology

| Product | Description | Quantity | Cat. No. |
|---------------------------------|---|----------|----------|
| MultiSite Gateway® Pro 2.0 Kit | Cloning two fragments into a Gateway® Destination vector | 20 rxns | 12537102 |
| MultiSite Gateway® Pro 3.0 Kit | Cloning three fragments into a Gateway® Destination vector | 20 rxns | 12537103 |
| MultiSite Gateway® Pro 4.0 Kit | Cloning four fragments into a Gateway® Destination vector | 20 rxns | 12537104 |
| MultiSite Gateway® Pro Plus Kit | Allows for flexible cloning of up to four fragments into a Gateway® Destination vector | 20 rxns | 12537100 |
| pcDNA™ 6.2/V5 PL-DEST Vector | A promoterless version of our most popular pcDNA™ vector for use with any of the MultiSite Gateway® Pro Kits Vector has C-terminal V5 and blasticidin selection | 6 µg | 12537162 |



Ordering information

Ultimate™ ORF Clones

| Product | Description | Cat. No. |
|---------------------------|--|----------|
| Ultimate™ Human ORF Clone | The Ultimate™ ORF Clone Collection contains DNA- and amino acid sequence-verified and expression-ready cDNAs, including kinases, G-protein-related, phosphatases, ion channels, GPCRs, chemokines, nuclear receptors, and cytokines; visit lifetechnologies.com/orf to select your clone today | HORF01 |
| Ultimate™ Mouse ORF Clone | | MORF01 |

BP Clonase® enzymes

| Product | Description | Quantity | Cat. No. |
|------------------------------------|---|----------|----------|
| Gateway® BP Clonase® II Enzyme Mix | A proprietary blend of both Int (integrase) and IHF (integration host factor) proteins that catalyze the <i>in vitro</i> recombination of PCR products or DNA segments from clones and a donor vector | 20 rxns | 11789020 |
| | | 100 rxns | 11789100 |
| Gateway® BP Clonase® Enzyme Mix | | 20 rxns | 11789013 |
| | | 100 rxns | 11789021 |

LR Clonase® enzymes

| Product | Description | Quantity | Cat. No. |
|---|--|----------|----------|
| Gateway® LR Clonase® II Plus Enzyme Mix | A proprietary blend of Int (integrase), IHF (integration host factor), and Xis (excisionase) enzymes that catalyze <i>in vitro</i> recombination between an Entry clone and a Destination vector | 20 rxns | 12538120 |
| | | 100 rxns | 12538200 |
| Gateway® LR Clonase® II Enzyme Mix | | 20 rxns | 11791020 |
| | | 100 rxns | 11791100 |
| Gateway® LR Clonase® Enzyme Mix | | 20 rxns | 11791019 |
| | | 100 rxns | 11791043 |

Competent cells

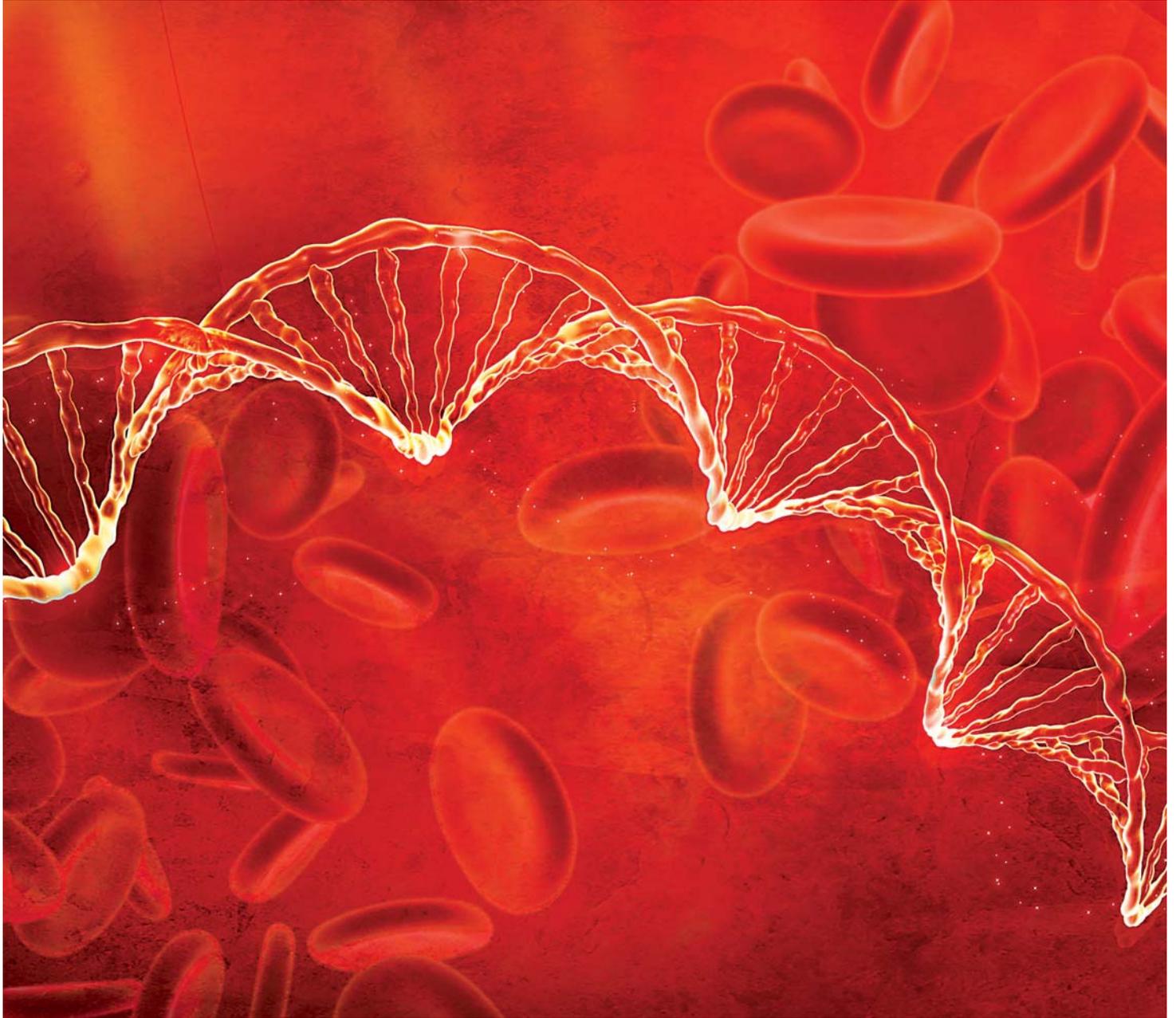
| Product | Description | Quantity | Cat. No. |
|---|--|--------------------|----------|
| One Shot® <i>ccdB</i> Survival™ Competent Cells | Designed for propagation of plasmids containing the <i>ccdB</i> gene | 10 transformations | C751003 |

Converting your proprietary cloning vectors with Gateway® technology

| Product | Description | Quantity | Cat. No. |
|-----------------------------------|--|----------|----------|
| Gateway® Vector Conversion System | Convert any cloning vector into a Gateway® Destination vector using restriction endonucleases and ligase | 20 rxns | 11828029 |

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