

A GATEWAY™ Destination Vector For High-Throughput Construction of *Neurospora crassa* histidine-3 Gene Replacement Plasmids

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A GATEWAY™ Destination Vector For High-Throughput Construction of *Neurospora crassa* histidine-3 Gene Replacement Plasmids

Abstract

We report the construction of a Destination Vector, called pJHAM007, for the targeted integration of DNA sequences at the *histidine-3* (*his-3*) locus of *Neurospora crassa*. pJHAM007 has all the necessary features required to perform a simple, rapid and efficient GATEWAY™ recombinational cloning with an Entry Clone to yield a *his-3*-gene replacement Destination Vector.

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The LR and the BP GATEWAY™ reactions were both performed as recommended by the manufacturer (Invitrogen).

Results and Discussion

Construction of pJHAM007, a GATEWAY™-compatible *his-3*-gene replacement Destination Vector. To demonstrate the feasibility of using GATEWAY™ to direct the integration of constructs at the *his-3* locus of *N. crassa*, we started by converting the *his-3*-gene replacement vector pJHAM003 (Lee, *et al.* 2003 *Curr. Genet.* DOI 10.1007/s00294-002-0366-z) into a Destination Vector called pJHAM007. The *his-3*-gene replacement plasmid, pJHAM007, was constructed by inserting a 2 kb *Bgl*III-*Eco*RV cassette removed from pDEST14 (Invitrogen) and containing the *att*R1, the *cat*⁺ = *chloramphenicol acetyl transferase* gene (chloramphenicol resistance), the *ccd*B gene and the *att*R2, into the *Bam*HI-*Pme*I restriction sites of pJHAM003 (Figure 1). We confirmed the construction of this plasmid by digesting pJHAM007 using several different restriction enzymes. Following a similar strategy, gene-replacement vectors for other loci can be converted into GATEWAY™ Destination Vectors.

RC-mediated Cloning of PCR Products To Generate Entry Clones. Complementation of a mutant phenotype by an ectopically integrated DNA fragment is a common way to demonstrate gene function. Because filamentous fungi can have large genes and complex promoters, it is not uncommon for those DNA fragments to be large, in order for them to contain all the necessary regulatory elements.

To direct the integration of a DNA fragment at the *his-3* locus using the *his-3*-gene replacement Destination Vector pJHAM007, the PCR fragment first needs to be present in an Entry Clone. It is known that the efficiency of the *in vitro* RC reactions decreases with increasing size of the DNAs involved (Hartley, *et al.* 2000 *Genome Res.* **10**: 1788-1795), but given the predicted need to occasionally use large DNA fragments, we first decided to test the limits of the RC-cloning reaction.

We therefore designed PCR primers containing *att*B sites to amplify a 9 kb DNA region from Linkage Group I (LG I) containing the *N. crassa* NCU02764.1 gene. PCR products flanked by *att*B sites can be generated by incorporating *att*B sites (25 base + 4 G residues) at the 5'-end of PCR primers (*att*B1 in the forward primer and *att*B2 in the reverse primer) and cloned by BP Recombination, into *att*P-containing vectors in the presence of Int and IHF (BP Clonase) to generate Entry Clones capable of recombining with Destination Vectors (Hartley, *et al.* 2000 *Genome Res.* **10**: 1788-1795). For this, 500 ng of the 9 kb PCR product were mixed with 300 ng of linearized pDONR201 plasmid (Invitrogen) and BP Clonase following the manufacturer's recommendations (Invitrogen). Following transformation into *E. coli* DH5 α cells and *Kanamycin-Resistance* (Km^R) selection, we analyzed 24 positive clones. Of those, only five were determined to be correct. We selected one correct clone and named it pJHAM009. The resulting plasmid, pJHAM009, is a GATEWAY™-compatible Entry Clone and its insert can therefore be transferred to any other Destination Vector with great versatility and efficiency.

RC-mediated Cloning of the insert from the Entry Clone (pJHAM009), into the *his-3*-Gene Replacement Destination Vector (pJHAM007), to generate the *his-3*-Gene Replacement plasmid (pJHAM010). The next step in the construction of a *his-3*-gene replacement Destination Vector is the transfer of the DNA insert present in the Entry Clone to the Destination Vector. This reaction is catalyzed by the LR Clonase, a mixture of the Int, IHF, and Xis proteins. For this, 140 ng of pJHAM009 Entry Clone were mixed with 300 ng of linearized pJHAM007 plasmid and LR Clonase following the manufacturer's recommendations (Invitrogen). Following transformation into *E. coli* DH5 α cells and *Ampicillin-Resistance* (Ap^R) selection, we analyzed 24 clones, among thousands of transformants. All 24 clones were determined to contain the expected insert size and all had the predicted restriction pattern. We selected one positive clone and named it pJHAM010. These results demonstrate that once in an Entry Clone, inserts as large as 9 kb can be transferred to any other Destination Vector with great versatility and efficiency, especially considering the large size of the resulting plasmid (in this case 21 kb).

Use of plasmid pJHAM007-derivative (pJHAM010) to direct the integration of DNA inserts at the *his-3* chromosomal position. To demonstrate the feasibility of using gene replacement plasmids derived from pJHAM007 to direct the integration of DNA inserts at the *his-3* chromosomal position, we transformed strains DLNCT62A and DLNCR83A (Lee, *et al.* 2003 *Curr. Genet.* DOI 10.1007/s00294-002-0366-z), each using 50 μ g of *Sfi*I-linearized plasmid pJHAM010. We selected 12 colonies from each transformation and tested them for FUDR (2'-deoxy-5-fluorouridine or (+)-5-fluorodeoxyuridine, filter sterilized) and hygromycin-resistance/sensitivity as described in (Lee, *et al.* 2003 *Curr. Genet.* DOI 10.1007/s00294-002-0366-z). Six FUDR-resistant, hygromycin-sensitive transformants obtained from DLNCT62A and seven obtained from DLNCR83A, were selected for further analysis. DNA was extracted from one DLNCT62A- and one DLNCR83A-derived transformant and Southern blot hybridization analysis was used to determine if a true gene replacement event had occurred in these transformants. As predicted, both transformants contained the insert present in plasmid pJHAM010 integrated at the *his-3* locus (data not shown).

In summary, a large PCR fragment obtained from *N. crassa* genomic DNA was cloned into a GATEWAY™-compatible Entry Vector, transferred into a *his-3*-gene replacement Destination Vector and integrated back at the *N. crassa his-3* chromosomal locus with minimal work and in a short period of time. We predict that the use of this technology will accelerate the development of our understanding of *N. crassa* biology considerably.

