

Supplementary Material

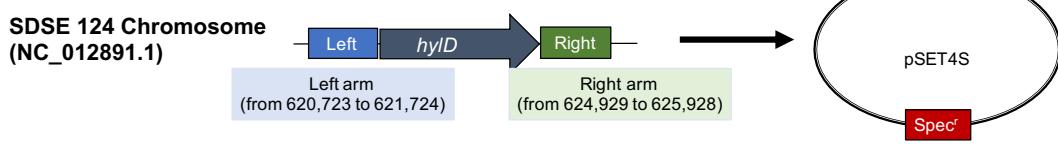
Novel Hyaluronate Lyase Involved in Pathogenicity of *Streptococcus*

dysgalactiae* subsp. *equisimilis

Van An Nguyen, Kohei Ogura, Miki Matsue, Norihiko Takemoto, Kanae Mukai, Yukari Nakajima,
Thuy Linh Hoang, Yasunori Iwata, Norihiko Sakai, Takashi, Wada, Wataru Hashimoto, Shigefumi
Okamoto, Hiroshi Ichimura

* Correspondence: Kohei Ogura: ogura@staff.kanazawa-u.ac.jp

Cloning



Gene recombination (When the left arm firstly crossovers)

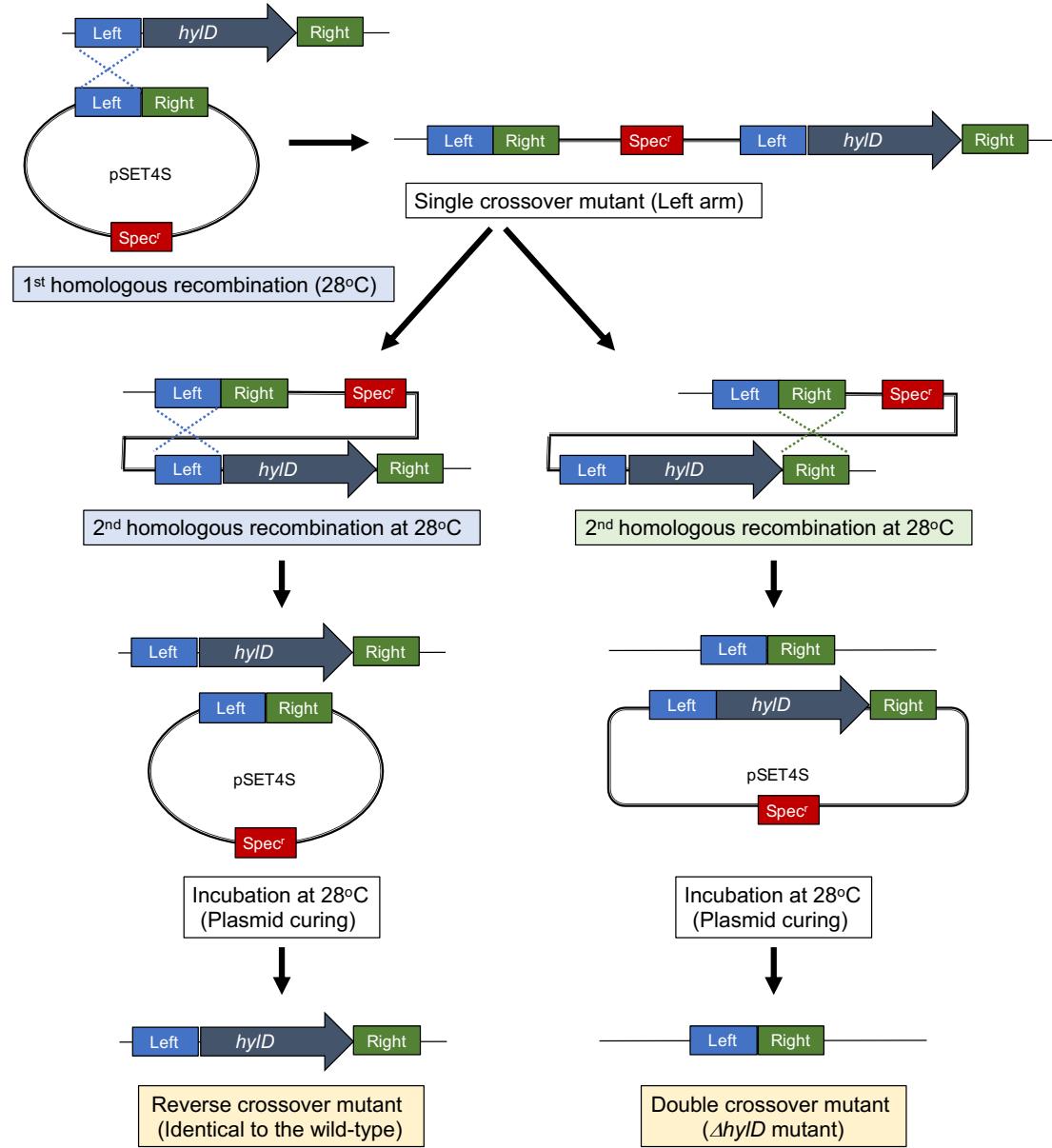


Figure S1. Diagram of procedure for prepare Δ hyd mutant strain. While this picture only shows a case when the first crossover mutant is obtained by homologous recombination of left arm (1 kb upstream sequences of the *hyd* gene in the SDSE chromosome), recombination of right arm (1 kb downstream sequences) also occurred. After transformation of the SDSE-124 by electroporation (1.75 kV pulse), All of the procedures were conducted at room temperature or 28°C.

Figure S2. Alignments of HylB (from GBS strain COH1) and HyID (from SDSE strain 124) proteins using Clustal W. Catalytic residues were colored by yellow. The loop region ($^{804}\text{KKLTID}^{809}$) was indicated by blue.

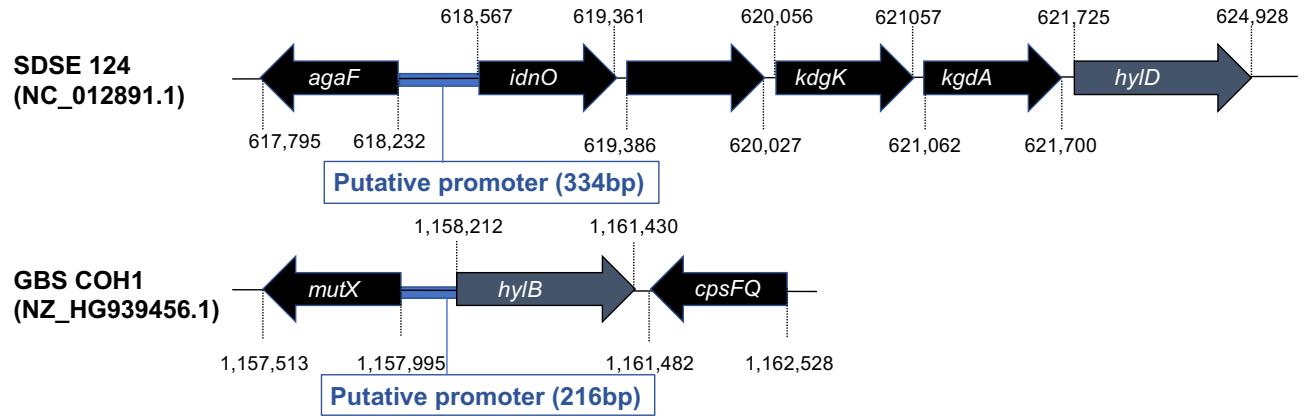


Figure S3. Genetic arrangements of *hylD* and *hylB* genes. Putative promoter regions were indicated in blue.