

Table S3. Molecular investigation of CRISPR/Cas9-tempted transmutations in *bnarfl11* and it has spread to from T₀ to T₁ generations.

| Examination of T ₀ lines | | | Mandilion Segregation of mutation to T ₁ lines | | | | |
|-------------------------------------|----------------------|-----------------------|---|--------|--------|---------|--------------------------------|
| Plant identification | Genetic constitution | Mutation ^a | plants examined | WT | Hetero | Homo | transmission rate ^b |
| <i>bnarfl11-1</i> | AA09 | +2, +1, -1 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-5</i> | AA09 | +1, -31, -1, +1 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-8</i> | AA09 | -3 | 14 | AA (0) | Aa (7) | aa (7) | 50% |
| <i>bnarfl11-15</i> | AA09 | -9, +1 | 14 | AA (0) | Aa (5) | aa (0) | 35% |
| <i>bnarfl11-19</i> | AA09 | -143, -11 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-23</i> | AA09 | +2, -18 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-27</i> | AA09 | +2, -2 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-34</i> | AA09 | -22, +1 | 14 | AA (0) | Aa (7) | aa (7) | 50% |
| <i>bnarfl11-38</i> | AA09 | -4, +1, -2 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-44</i> | AA09 | +2, -7, +2 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-57</i> | AA09 | +1, -1 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-69</i> | AA09 | +1, -2, -5 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-73</i> | AA09 | 0 | 14 | AA (7) | Aa (7) | aa (0) | 50% |
| <i>bnarfl11-75</i> | AA09 | -6, +3, +1, -2 | 14 | AA (0) | Aa (0) | aa (14) | 100% |
| <i>bnarfl11-79</i> | AA09 | -8, +5 | 14 | AA (0) | Aa (7) | aa (7) | 50% |
| <i>bnarfl11-83</i> | AA09 | +4, -1, -2 | 14 | AA (0) | Aa (7) | aa (7) | 50% |
| <i>bnarfl11-87</i> | AA09 | +2, -11, -4, +2 | 14 | AA (0) | Aa (0) | aa (14) | 100% |

* WT stands for wild type; Hetero stands for heterozygous; Homo stands for homozygous, a "-" and "+" denotes the loss and addition of the specified number bases simultaneously. B was determined based on the number of lines with mutation by the entire line verified for mutation.