## Development of Visible Markers for Transgenic Plants and their Availability for Environmental Risk Assessment

Masanori Tamaoki<sup>a,\*</sup>, Hiroe Imai<sup>b</sup>, Hayato Takahashi<sup>a</sup>, Yumio Toda<sup>c</sup>, Yasuo Niwa<sup>d</sup>, Nobuyoshi Nakajima<sup>a</sup>, Mitsuko Aono<sup>c</sup>, Akihiro Kubo<sup>c</sup>, and Hikaru Saji<sup>c</sup>

- <sup>a</sup> Biodiversity Conservation Research Project, National Institute for Environmental Studies, 16-2 Onogawa, Tsukuba, Ibaraki 305-8506, Japan. Fax: +81-29-850-2490.
  E-mail: mtamaoki@nies.go.jp
- <sup>b</sup> Endocrine Disrupters Research Project, National Institute for Environmental Studies, 16-2 Onogawa, Tsukuba, Ibaraki 305-8506, Japan
- <sup>c</sup> Environmental Biology Division, National Institute for Environmental Studies, 16-2 Onogawa, Tsukuba, Ibaraki 305-8506, Japan
- <sup>d</sup> Graduate School of Nutritional and Environmental Sciences, University of Shizuoka, 52-1 Yada, Shizuoka 422-8526, Japan
- \* Author for correspondence and reprint requests

Z. Naturforsch. 61c, 377-386 (2006); received December 8, 2005/January 11, 2006

Monitoring of transgenic plants in the field is important, but risk assessment has entailed laborious use of invisible marker genes. Here, we assessed three easily visible marker transgenes - green fluorescent protein (GFP), R, and Nicotiana tabacum homeobox (NTH) 15 genes – for their potential use as marker genes for monitoring genetically modified plants. Transgenic Arabidopsis thaliana plants for each of these genes were visibly distinguished from wild-type plants. We determined the germination rate, 3-week fresh weight, time to first flowering, and seed weight of the transgenic plants to evaluate whether the expression of these marker genes affected the growth of the host. Introduction of GFP gene had no effect on the evaluated parameters, and we then used the GFP gene as a marker to assess the outcrossing frequency between transgenic and two Arabidopsis species. Our results showed that the hybridization frequency between transgenic plants and Arabidopsis thaliana was 0.24%, and between transformants and *Arabidopsis lyrata* it was 2.6% under experimental condition. Out-crossing frequency was decreased by extending the distance between two kinds of plants. Thus, the *GFP* gene is a useful marker for assessing the whereabouts of transgenes/transformants in the field. We also demonstrated that the GFP gene is possibly applicable as a selection marker in the process of generation of transgenic plants.

Key words: Green Fluorescent Protein, Risk Assessment, Transgenic Plant