# Summary of the Risk Assessment and Risk Management Plan

**for**

**Licence Application No. DIR 156**

## Introduction

The Gene Technology Regulator (the Regulator) has decided to issue a licence for this application for the intentional release of a genetically modified organism (GMO) into the environment. A Risk Assessment and Risk Management Plan (RARMP) for this application was prepared by the Regulator in accordance with the requirements of the *Gene Technology Act 2000* (the Act) and corresponding state and territory legislation, and finalised following consultation with a wide range of experts, agencies and authorities, and the public. The RARMP concluded that the field trial poses negligible risks to human health and safety and the environment and that any risks posed by the dealings can be managed by imposing conditions on the release.

## The application

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| Application number | DIR 156 |
| Applicant | The Royal Melbourne Institute of Technology (RMIT) University  |
| Project title | Limited and controlled release of buffalograss genetically modified for herbicide tolerance and dwarf phenotype |
| Parent organism | *Stenotaphrum secundatum* (buffalo grass) |
| Introduced genes and modified traits | * Gene encoding the enzyme 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS) from *Arabidopsis thaliana* (thale cress) for tolerance to the herbicide glyphosate
* Gene encoding the enzyme gibberellic acid 2-oxidase 3 from *Spinacia oleracea* (spinach) for shorter stature and slowed growth
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| Proposed location | One site in Victoria |
| Proposed release size | Up to 200 m2  |
| Proposed release dates | April 2018 – April 2019 |
| Primary purpose | To assess agronomic characteristics of the GM buffalo grass plants  |

## Risk assessment

The risk assessment concludes that risks to the health and safety of people, or the environment, from the proposed release are negligible. No specific risk treatment measures are required to manage these negligible risks.

The risk assessment process considers how the genetic modification and proposed activities conducted with the GMOs might lead to harm to people or the environment. Risks are characterised in relation to both the seriousness and likelihood of harm, taking into account current scientific/technical knowledge, information in the application (including proposed limits and controls) and relevant previous approvals. Both the short and long term are taken into account in the risk assessment.

Pathways which may result in harm that were considered included exposure of people or animals to the GM plant material, likelihood of persistence or dispersal of the GMOs, and transfer of the introduced genetic material to other buffalo grass plants. Potential harms associated with these pathways included toxicity or allergenicity to people, toxicity to desirable animals, and environmental harms due to weediness.

The principal reasons for the conclusion of negligible risks are that buffalo grass is not a food crop which reduces human exposure, the GM plant material will not be used for animal feed and that the imposed limits and controls effectively contain the GMOs and minimise exposure to the GMOs and their genetic material.

## Risk management plan

The risk management plan describes measures to protect the health and safety of people and to protect the environment by controlling or mitigating risk. The risk management plan is given effect through the conditions of the licence.

As the level of risk is considered negligible, specific risk treatment is not required. However, since this is a limited and controlled release, the licence includes limits on the size, location and duration of the release, as well as controls to prohibit the use of GM plant material in animal feed, to minimise dispersal of the GMOs or GM pollen from the trial site, to transport GMOs in accordance with the Regulator’s guidelines, to destroy GMOs at the end of the trial, and to conduct post-harvest monitoring at the trial site to ensure all GMOs are destroyed.

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