

SUMMARY OF AUTHORISATION PATHWAYS

Assessed by Regulator

	Type of dealing	Statutory approval timeframe ¹	Public consultation ²	GTTAC consideration
Licenses	Licenses requiring public consultation on the RARMP	150-200 days	YES	YES
	Environmental release of a novel gene drive organism	400 days	YES	YES
	Inadvertent dealings licences	90 days	NIL	OPTIONAL
	Other than licences mentioned above, including designated dealings.	90-150 days	NIL	AS REQUIRED
Permits	Plant field trials, clinical trials, GMO administered under the TGA Special Access Scheme.	30 days	NIL	NIL
GMO Register	Determination for listing a new dealing on the GMO Register	NIL	OPTIONAL	OPTIONAL

Assessed by Institutional Biosafety Committee (IBC)

	Type of dealing	Statutory approval Timeframe ¹	Public consultation	GTTAC consideration
Notifiable Dealings	Post-notified notifiable dealings	N/A	NIL	NIL

Other

	Type of dealing	Statutory approval Timeframe ¹	Public consultation	GTTAC consideration
Notifiable Dealings³	Pre-notified notifiable dealings	N/A	NIL	NIL
Non-notifiable Dealings⁴	Only dealings to be specified to be non-notifiable dealings (NND) in Regulations	N/A	NIL	NIL
GMO Register⁴	Any dealings currently specified on the GMO Register	N/A	NIL	NIL

¹ Days refers to business days.

² The Regulator may consult the public on any RARMP, if the Regulator considers it in the public interest to do so.

This table only includes circumstances when the Regulator must consult the public.

³ These dealings are authorised in line with requirements set out in the regulations and rules.

⁴ These dealings require self-assessment by the proponent.