

Table 1. Examples of studies that used assessment of seed viability and germination at the end of the experiment to estimate initial viability.

Species tested	Method of determining initial viability from end-of-experiment data	Experimental conditions	Differing initial viability between treatments	Reference
21 woody native species of Chilean matorral	Viability calculated as: those germinated during the monitoring period plus non-germinated seeds identified as viable by tetrazolium test	Moist absorbent paper in petri dishes at 20/10°C with 12/12 h light/dark for 36 d	Yes (3 species), but data not adjusted for viability	Gómez-González et al. (2017)
13 native and one introduced species (<i>Acacia saligna</i>) of South African fynbos	Viability determined as the sum of germinated seeds and seeds appearing fresh on dissection of ungerminated seeds	1% water agar at 10/20°C with a 12 h light and dark cycle for 91 d	Yes (at least 2 species)	Hall et al. (2017)
65 species commonly occurring on New England tableland (NSW Australia)	Seeds that did not germinate, but looked viable, were analysed using tetrazolium test. Viability based on treatment with highest germination plus any seeds that remained dormant but viable	Moist pad in dish at 25/15°C with a 12/12 h light/dark for 28 or 56 d	Probably (as used highest viability levels between treatments)	Clarke et al. (2000)
<i>Asterolasia buxifolia</i> , riparian habitat of SE Australia	Embryo dissected from 20 seeds that did not germinate and viability confirmed if embryo and endosperm intact	Moist filter paper in petri dishes at 11/3°C with 12/12 h in light/dark for 77 d	Data not adjusted for viability*	Collette and Ooi (2017)
33 herb and small shrub species in fire-prone Turkey	Embryo of seeds that did not germinate examined and viability confirmed if embryo intact	0.8% agar in Petri dishes at 20°C in dark for 35 d	Data adjusted for viability	Serter Çatav et al. (2017)
46 legumes species of tropical savanna, Brazil	Initial viability equals the sum of germinated and dormant seeds in control	Moist filter paper in petri dishes at 27°C with 12/12 h light/dark for 28 d	No data given at treatment level	Daibes et al. (2019)
13 species of West African	Cut test – condition of embryo,	Moist filter paper in bell jars at 25°C light	No data given at	Dayamba et

of grasslands and woodlands of SE Australia	seeds post-trial	light/dark for 56 d	treatment level	al. (2019)
2 alien and 2 indigenous legume species in S African fynbos	Germination level of scarified seeds conducted at same time as other treatments	0.02% benomyl solution in petri dishes at 20°C with 12/12 h light/dark for 30 or 60 d	No data given at treatment level	Jeffery et al. (1988)
3 species of <i>Acronychia</i> in E Australian rainforest	Ungerminated seeds checked for firmness by pressing seed with forceps. Then firm seeds checked for viability via cut test	0.8% water agar in petri dishes at 25/10°C with a 12/12 h light/dark for 28 d	No data given at treatment level	Liyange et al. (2020)
9 herbaceous species in Brazilian grassland	Tetrazolium test on ungerminated seeds	Moist filter paper in petri dishes at 20°C or 25°C at 16/8 h in light/dark for 21 d	No data given at treatment level	Overbeck et al. (2005)
<i>Brassica napus</i>	Cotyledon condition of ungerminated seeds post-trial. Necrotic cotyledons = nonviable, yellow-milky cotyledons = viable	Moist filter paper in petri dishes at 20°C in dark for 35 d	No	Shayanfar et al. (2020)

*Viability 80% at 100°C but only 65% at the lower temperature of 80°C appears anomalous and might indicate an unexpected treatment effect on viability (they should have been the same or the reverse if there was a heat effect on viability) but no statistical analyses were undertaken