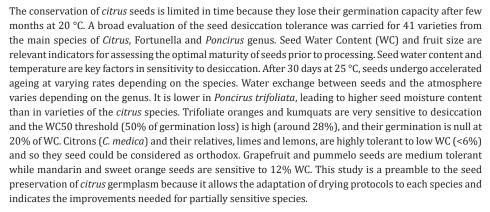


The Tolerance of *Citrus* Seeds to Desiccation Reveals a Range of Behaviors from Orthodox to Recalcitrant

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Abstract



Keywords: Water content; Relative humidity; Germination; Seed maturity; *Citrus* diversity; Seed ageing; Orthodox; Recalcitrant

Introduction

According to the recent genomic studies, allopatric evolution, hybridization and migration can explain the citrus evolution and diversification [1-3]. The main citrus crops, sweet orange (C. sinensis (L.) Osb.), lemon (C. limon (L.) Burm.), sour orange (C. aurantium L.), lime (C. aurantifolia (Christm.) Swing.), yuzu (C. junos Sieb. ex-Tan.), belonged from natural interspecific hybridizations between five ancestral taxa, citron (C. medica L.), pummelo (C. maxima Burm.) Merr.), mandarin (C. reticulata Blanco), and Papeda species such as C. micrantha Wester and C. ichangensis Swing. Trifoliate oranges (Poncirus trifoliata (L.) Raf.) and kumquats (Fortunella sp.) were also originated from Asia but due to the biology of reproduction (shift of blooming) limited the intergeneric natural hybridizations. However, the trifoliate orange is a genitor of breeding programs for production of intergeneric hybrids used as rootstocks because it remains a source of resistances to diseases and environmental stresses. The citrus rootstocks are propagated by seedling. Maintaining the viability of citrus seed over time is essential to meet the short-term (few months) conservation needs of commercial seed producers and breeders. Storage of the seeds at a relatively low temperature (5 °C) with appropriate fungicide treatment and relatively moderate drying proved adequate for short-term maintenance [4-6]. This short-term maintenance is adapted for the production of citrus rootstock seed, to bridge the gap between harvest time and spring sowing time. During the formation of seeds after fertilization, many citrus species or genotypes have the particularity of producing somatic embryos, which allow clonal propagation. Citrus





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rootstocks are clonally amplified by seedling of polyembryonic seeds [7]. Maintaining long-term seed viability is a necessity for ex situ conservation of *citrus* genetic resources, for which an effective and low-cost method of long-term conservation is needed. Many research and storage advances have been achieved predominantly in cultivated *citrus* species. The most promising potential long term ex situ technique appears to be cryopreservation of seeds and it is considered as the safest option to prevent seed viability loss [8,9].

Understanding behavior of seed storage depends on a knowledge of a species taxonomy, plant ecology, and seed characteristics [10]. Citrus are classified as intermediate seeded species because can withstand partial dehydration, but they cannot be stored under conventional genebank conditions because they are cold sensitive and desiccation does not increase their longevity [11,12]. Tolerance to seed desiccation varies within and between citrus species from orthodox to recalcitrant [10]. The moisture control is a key element for freezing of citrus seed [9]. The limit of citrus seeds dehydration for liquid nitrogen freezing treatment corresponds to the unfrozen Water Content (WC) in the tissue, confirming that seed survival strictly depends on avoidance of intracellular ice formation. Moisture contents between 10 to 20% (fresh weight basis) are often optimal for survival of freezing [13]. A fixed seed moisture content can be achieved by incubation of seeds in different atmospheric relative humidity controlled by salt solutions [9]. The WC equilibrium was achieved after few days or weeks of incubation, depending on species and Relative Humidity (RH). Other drying method are frequently used as laminar airflow and silica. The removal of the seed coat could permit lemon seeds to be dried to a low moisture content without any damage [14]. However, the removal of the seed coat is not necessary, if provided sufficient time is allowed for intact seeds to germinate [15]. Damage caused by low temperatures and dehydration in plant cells may be irreversible and cause cell death depending on the intensity of the stress [16]. Water is an essential element in maintaining the integrity of the cell membrane. Indeed, a water deficiency leads to a modification of the phospholipid structure of the lipid bilayer, which leads to leakage of electrolytes to the intercellular medium [17]. Many studies on seed desiccation tolerance have been down on citrus cultivars but there is considerable variability among seed lowest safe moisture content (WC) when comparing different studies [18]. In sour orange (C. aurantium), for example, viability is 25% for a WC of 13.8% [19], 40% for a WC of 14.4% [20] and 80% for a WC of 10% [8]. Similarly high discrepancies are observed in lemon (C. limon): 1% for a WC of 10% [21], 85% for a WC of 7.5% [22] and 90% for a WC of 10.3% [23]. These differences in viability between studies cannot be fully explained by the varietal diversity of these 2 species, as it has been shown that their diversification has mainly been through mutations and that human selection has mainly retained new phenotypes on fruit traits such as color, shape, size juice composition and seed lessness [24,25]. In ancestral species, on the other hand, sexual crossing is the source of varietal diversification, generating more genetic diversity than mutation [26]. It is therefore normal to observe diversity in dried seed viability between varieties of pumelo (C. maxima) [27,28]. Various categorization of the lemon seeds was proposed by different authors

from orthodox [20] and intermediate to recalcitrant [21,22]. The definition of desiccation tolerance is based on the ability of seeds not to lose viability below 10% moisture content [29]. The lower limit of the threshold water content before injuries for intermediate seeds is around 15% [30]. Most studies of desiccation tolerance were done on very few varieties at the same time and with the same conditions of seed drying, making it impossible to apply a suitable method for short or long-time conservation adapted to each citrus species. Since the control of WC is crucial for the preservation of germination capacity and viability of citrus seeds, the main objective of this study was to evaluate, the seed desiccation tolerance in a wide range of desiccation levels for the citrus species. This multi-species study has other objectives. The evaluation of some factors that can modify seed quality (maturity and temperature during drying), and the inheritance of desiccation tolerance across interspecific and intergeneric hybrids. The list of investigated citrus was defined to represent the citrus genetic diversity including ancestral species and cultivated groups.

Materials and Methods

Plant material

Fruits were picked on the trees of the INRAE-Cirad *citrus* biological resource center (San Giuliano, Corsica, France; 42° 18′ 55′ N′ and 9° 29′ 29′ E′) [31]. The seeds were extracted from maturate fruit at the ripening time of each variety (Full fruit size), washed with water and few drops of detergent, partially dried at room temperature (22-25 °C) on the open-air desk for 20h, before to be used or stored at 4 °C. Thirty-four *citrus* cultivars were selected to represent the four *Citrus* ancestral species of *Citrus* and their interspecific hybrids such as sweet oranges, sour oranges, lemons and limes. In addition, Fortunella and *Poncirus* genus were represented each one by three and two cultivars respectively, completed by 2 *Citrus* x *Poncirus* hybrids usually used as rootstocks in the *citrus* industry, citranges (*C. sinensis* x *P. trifoliata*).

Seed desiccation

Seeds were incubated in a hermetically sealed jar containing a saturated solution of different salts in order to control the relative humidity of the atmosphere. The seeds were kept above this solution, without contact with it, on a support allowing the exchange of water molecules between the atmosphere of the jar and the seeds. The relative humidity (RH) of the atmosphere of the jar were controlled by different saturated salt solutions, NaOH (6%), KOH (9%), MgCl₂ (34%), K₂CO₃ (45%), NaCl (75%), NH₄Cl (78%), (NH₄)2SO₄ (81%), KCl (85%) and Na₂CO₃ (87%) [32,33]. Seeds loose water molecules by exchange with the atmosphere until they reach equilibrium [33]. The time required for seed moisture content to stabilize depends on RH and variety [34]. To ensure MC stability, seeds were incubated for 12 days at RH levels of 6 to 45%, 20 days at RH levels of 75 and 78%, and 30 days at RH levels of 81 and 85%.

Measurement of the seed moisture (Water Content: WC)

Ten normal developed seeds were weighed (P1) and then completely dried by incubation for $16\,h$ at $104\,^{\circ}C$ and then weighed

again (P2). The water content of the seed is WC = ((P1-P2)/P1) \times 100.

Germination test

Seeds were sowed on fine water saturated vermiculite in petri dishes (15x15cm), placed at 25 °C, in the dark. After 4 weeks, the number of germinated seeds with at minimum a 20mm-long emerging root axis, was scored. The germination rate (%) was then calculated.

Seed maturity

The seed weight, the fruit weight, the germination capacity and the seed moisture content were measured during 7 months from 10^{th} of June (40-60 days after anthesis) to 10^{th} of January (when the fruit was totally developed), for 3 *citrus* genotypes usually used as rootstock, 'Volkamer' lemon, 'Maroc' sour orange, and 'Carrizo' citrange. The flowering time diverges between the 3 genotypes it extended from mid-April to mid- May for the 'Volkamer' lemon and the sour orange, but it was 2-3 weeks earlier for 'Carrizo' citrange. For each parameter, during the period of fruit development nine dates were chosen for sampling 8 fruits for the weight measurement, 10 seeds for weight and WC assessment, and 3x30 seeds for germination.

Effect of temperature and moisture content on seed germination during drying process and storage

Six *citrus* rootstocks were selected for this experiment, three *Citrus* species ('Volkamer' lemon, 'Rangpur' lime and sour orange), 'Pomeroy' trifoliate orange (*P. trifoliata*) and two intergeneric hybrids, 'Carrizo' and 'C35' citranges (*C. sinensis* x *P. trifoliata*). The jars containing seeds and salt solutions for RH of 45% ($\rm K_2CO_3$), 75% (NaCl) and 85% (KCl), were disposed into an incubator (25 °C) or into a cold chamber (4 °C). The experiment was conducted over 270 days with samples of seeds were kept at 30, 60, 90, 120, 150, 180 and 270 days of incubation time to evaluate the WC and GR parameters.

Evaluation of the seed tolerance of *citrus* species against a large range of desiccation levels

The proportion of germinated seeds of 39 genotypes from different *citrus* varieties, cultivars and species, was measured after drying the seeds under different RHs (85%, 81%, 75%, 62%, 45%, 23%, 6%) by using the salt solutions of KCl, $(NH_4)2SO_4$, NaCl, NH_4NO_3 , K_2CO_3 , K acetate, NaOH, respectively, at 25 °C and until the WC has reached the equilibrium. The tolerance to desiccation was evaluated by comparing the germination rate of dried seeds with the germination of untreated seeds directly sowed after extraction. Fifty seeds were used for the evaluation of germination rate for each *citrus* accession.

Statistical analysis

Data are expressed as mean values \pm SD. Differences between varieties were assessed by Duncan test and one-way analysis of variance to determine significant differences for different RH applied for seed desiccation. Statistical significance was determined with p<0.05.

Results

Evolution of germination capacity according to the seed and fruit development

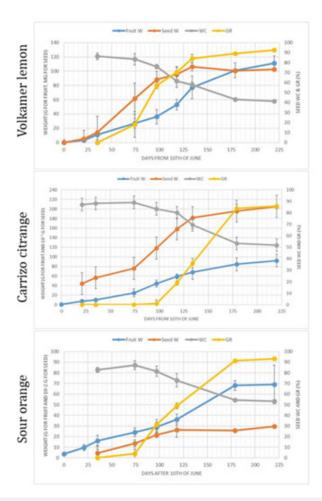


Figure 1: Evolution of fruit weigh (W), seed weight and moisture (WC) and of the Germination Rate (GR) during fruit development (from 10th of June/0 to 20th of January/220 days) for 'Volkamer' lemon, Sour orange and 'Carrizo' citrange. Vertical bars represent ± SD.

The kinetics of the seed weight, the fruit weight as well as the germination rate was similar between 'Volkamer' lemon, 'Carrizo' citrange and Sour orange (Figure 1). The kinetics of orange trifoliate seed parameters have not been plotted because they are lagged by almost 2 months due to its early flowering. In addition, no data were available from October onwards. The seed moisture content (WC) was still high (around 90%) at the beginning of the experiment (in June) and then decreased slowly from mid-September (100 days after the first measurement) until stabilization approximatively to 50% for 'Carrizo' citrange and sour orange and to 40% for 'Volkamer' lemon. During the phase of seed water loss, the seed weight always increased. Germination Rate (GR) correlated negatively with seed WC. Regardless of the variety, GR is maximum when seed WC was below 55% (it is the same with trifoliate orange). The higher germination rates were observed when the fruit and the seed weights reached their maximum. The seeds of the 'Volkamer' lemon had an earlier maturity of approximately 30 days

compared to the two other *citrus* fruits (end of October versus end of November). Trifoliate orange seeds reach maturity in September. Taking into account these results on the seed maturity, the fruits of all the studied *citrus* fruits were harvested between December and February when they had reached their maximum size (or weight). The fruits of *Poncirus trifoliata* were kept in October and seeds of kumquats in March to ensure optimal seed maturity because their bloomed in July.

The seed WC according to the species and the degree of dehydration

At the end of desiccation process, the equilibrated seed WC (when it reached a stable value) differed between *citrus* varieties

(Table 1). It was always significantly higher in *Poncirus trifoliata* than in other genotypes, whatever the RH. However, this difference in WC compared to other varieties decreases as RH decreases. The WC value for trifoliate orange seeds, observed at 87% RH, has not yet reached equilibrium after 47 days of desiccation, as the standard deviation is still very high (4.7). The WC values for the other varieties are stable with a standard deviation close to 1.0, they have therefore reached equilibrium. In 'Carrizo' citrange the equilibrated WC are significantly higher than those of *Citrus* species and trifoliate orange only at RH of 75% and 87%. This intermediate behavior is in agreement with its intergeneric hybrid origin (*CitrusxPoncirus*). The behaviors of the five *Citrus* species and Nagami kumquat are similar.

Table 1: The average of final seed WC (g H_2 Og-1dw) %) and standard deviation of 8 citrus species desiccated with four relative humidity (RH in %) and just after extraction (T_0 fruit). Values followed by the same letter did not differ significantly at P<0.05 using Duncan's multiple range test.

Citmus on	T _o	RH %						
Citrus sp.	Fruit	87	75	35	9			
Trifoliate orange	43.3±4.4	33.8±4.7b	17.6±2.3b	12.6±1.7e	8.6±1.2a			
Carrizo citrange	39.5±4.4	24.1±0.1ab	12.5±0.6ab	7.8±0.4c	3.9±0.1c			
Nagami kumquat	49.9±2.2	15.9±1.0d	9.1±0.6c	4.3±0.2d	2.7±0.3e			
Corsican citron	53.4±0.5	18.0±0.2cd	11.4±0.9ac	5.0±0.4bd	1.8±0.6e			
Chandler pummelo	49±2.5	17.6±1.3cd	9.3±0.2c	5.8±0.1ab	3.8±0.1c			
Cleopâtra mandarin	58.4±2.9	21.6±0.4abc	10.4±0.3ac	7.2±0.4ac	4.2±0.2b			
Maroc sour orange	57.1±4.1	18.0±1.1cd	10.5±1.5ac	7.9±0.3c	3.5±0.4cd			
Rangpur lime	42.5±1.1	17.2±1.1cd	9.4±0.7c	6.7±0.8ac	2.9±0.2e			

The germination varies with the temperature, the desiccation intensity and the genotype

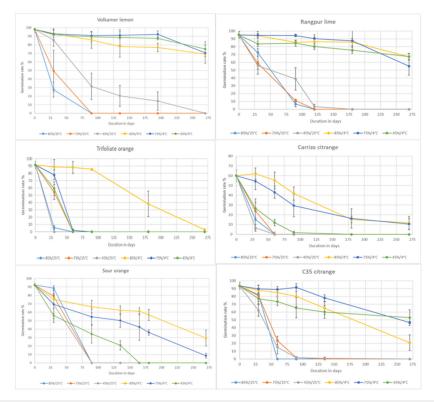


Figure 2: Evolution of seed germination of 6 citrus rootstocks during the desiccation period with different relative humidity (45, 75 85% RH) and two temperatures of incubation (25 °C and 4 °C). Vertical bars represent ± SD.

In general, and for all varieties, the highest germination rates were observed when seeds were dried at 4 °C (Figure 2). The temperature has a strong effect on the preservation of the seed germination capacity. After 100 days of incubation at 25 °C, the seeds of all the varieties do not germinate any more with the exception of the seeds of 'Volkamer' lemon and 'Rangpur' lime incubated at 45% RH their germinations are close to 30%. The germination capacity is longer preserved when the seeds were maintained at 4 °C. After 100 days of incubation in 85% RH at 4 °C, the germination was higher than 80% for 'Volkamer' lemon, 'Rangpur' lime, 'C35' citrange and trifoliate orange, and higher than 60% for sour orange. The germination kinetics of 'Rangpur' lime and 'Volkamer' lemon seeds during drying are very similar whatever the conditions. For 'C35' citrange, unlike the temperature the relative humidity does not seem to have an effect on the germination capacity. Its germination decreased and at twice as fast at 25 °C as for the previous two rootstocks and at 4 °C it was reduced to around 20-50%. For the three other rootstocks, the relative humidity has an incidence on the germination rate. At 45% RH and 4 °C, the germination capacity was null after 60 days for the trifoliate orange, 90 days for the 'Carrizo' citrange and 150 days for the sour orange. The seeds of these three rootstocks are not resistant to excessive drying. The seeds of trifoliate orange were even very sensitive to a RH of 75% since the germination rate is null after 60 days of incubation. After 100 days of incubation, the germination decreased by 30% for sour orange and 50% for 'Carrizo' citrange, with seeds incubated at 75% or 85% of RH. At the end of 9 months, the seeds of these two cultivars have a germination rate ranged between 10 and 20%. There is a difference in final water content (at equilibrium) and in the rate of decay depending on the drying temperature. At 4 °C, water loss from the seed is slowed down and the equilibrium WC value is higher than that of seeds incubated at 25 °C, regardless of relative humidity. WC values observed after 30 days at 25 °C are only reached after 90 days at 4 °C for all rootstocks. For example, in 'Volkamer' lemons, at 85% RH, the WC stabilizes at 11.6% after 60 days at 25 °C, whereas at 4 °C stability is achieved at 13.5% after 135 days of incubation. At 75% RH and 25 °C, the WC is already stabilized at 8.9%, while at 4 °C, the WC stability is around 10.1% and only after 90 days of incubation. Similar differences are also

observed for incubation at 45% RH.

Seed tolerance against water loss during drying process performed at $4\,^\circ\text{C}$

The germination rate according to the water content makes it possible to evaluate the sensitivity or the tolerance of seeds to desiccation and to measure the water content corresponding to a reduction of 50% of the germination rate (WC50). A previous experiment had shown that the time required to achieve WC stability during desiccation was long at high relative humidity (>20 days) and varied depending on the species [34]. Therefore, only the data obtained from incubation made at 4 °C and during the first 160 days were used to avoid or limit the effect of the seed ageing (Figure 3). The evolution of germination rater as a function of WC of 'Volkamer' lemon and 'Rangpur' lime was almost linear with quite no slope. The decrease in germination rate between the highest (42%) and lowest (7%) WC was only about 10% of the initial value for these two citrus species. The germination rate of the 'C35' citrange decreased steadily but gently during the decrease of WC and the total reduction in water content was only 25% at the lower seed WC. The curves of sour orange and 'Carrizo' citrange evolved similarly with a drastic slowdown of the germination rate. The curve corresponding to the trifoliate orange presented a very inclined slope with a very pronounced decrease of the germination rate when the WC was below 30%. The graphical estimation of the WC50 from these curves could only be evaluated for these last three citrus genotypes because only they showed a significant decrease in germination rate. The WC50 was graphically estimated around 13% for sour orange, 24% for 'Carrizo' citrange and 27% for trifoliate orange. It should be noted that the initial germination rate of 'Carrizo' citrange seeds was low (60%) in contrast to other citrus fruits which were at about 90%. It is thus preferable to evaluate the level of sensitivity of seeds to desiccation in relative proportions of decrease of the germination rate rather than on absolute values. In agreement with the WC50 values and the curves of germination rate versus water content, it was concluded that trifoliate orange and 'Carrizo' citrange are the most sensitive varieties to desiccation, sour orange less sensitive, 'C35' citrange tolerant, 'Rangpur' lime and 'Volkamer' lemon highly tolerant.

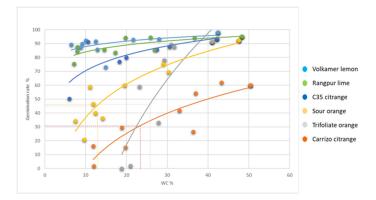


Figure 3: Kinetics of the Germination Rate (GR) and the seed WC (% of fresh matter weight) for six citrus rootstocks, made only with values obtained at 4 °C and the period of the first 160 days of incubation. The dotted lines in tendency curves indicate the water content (WC50) corresponding to half reduction of the initial germination rate (50% loss of GR) for the citrus genotypes presenting a sufficient GR decrease.

Evaluation of the diversity of seed desiccation tolerance of *citrus* species

Each species was represented by 2-4 cultivars to assess the seed behavior against a range of desiccation degrees for each taxonomic group (Table 2). The initial germination rate when the seeds were sawed just after extraction, was high (>72%) for all studied cultivars. In general, there was little or no variation between cultivars of the same species. Given the absence of repetition (only one measurement carried out on 50 seeds), the variation between GR lower than 15% are not taken into account and only relative decreases higher than 30% are considered as informative for the qualification of the tolerance status to desiccation conditions of each citrus cultivar. The four ancestral species of the genus Citrus showed variable behavior with notably tolerance to low RH (6% and 23% corresponding to a range of 3.5-7.5% WC) for all citrons (C. medica) and Papedas that maintained a viability higher than 80% (excepted for Combava). Pummelos (C. maxima) were also tolerant (more than 50% of GR decreases were observed only from 23% and lower RH). Among ancestral species, mandarins (C. reticulata) were the most sensitive with a loss close to 50% of germination at 62% RH (the WC is close to 10% for 62% RH). Kumquats (Fortunella sp.) lost more than 50% of germination at 75% RH while trifoliate oranges (P. trifoliata) lost the same proportion at 81% RH. Limes and lemons displayed generally the same tolerance level as their citron parent except for 'Alemow' (C. macrophylla) which lost more than 50% of germination between 23% and 6% RH. The germination capacity of sour oranges was maintained high at 45% RH while grapefruits have a behavior comparable to their pummelos parent, as well as sweet oranges had a behavior close to that of mandarins. Intervarietal diversity can be observed in few species based on seed behavior at low RH (or high drying conditions). This is the case for pummelos, mandarins, sour oranges and citranges. 'Cleopatra' and 'Changsa' mandarins are more sensitive than 'Sunki' with a decrease of the germination at 45% RH, representing 79, 59 and 26% of the initial rate, respectively. The germination of 'Sans Pepin' pummelo decreases drastically at 45% RH (53% of loss), while it is still very high for the other two cultivars (>80%). The behavior of C35 and Carrizo citranges was opposite with a high sensitivity of Carrizo. Despite this intra specific variability, a classification of species from the most sensitive to the most tolerant to seed desiccation can be proposed: trifoliate oranges<kumquats<'Carrizo' citrange<sweet oranges<mandarins<'C35'citrange<grapefruits< pummelos<sour oranges<'Alemow'<lemons<papedas-limes-'Nasnaran'<citrons.

Table 2: Percentage of germination (GR) of citrus cultivars and species with seeds incubated in a range of RH. To represents the germination rate of untreated seeds directly sowed after extraction from fruit, the grey color and its intensity represents the diminution of GR relatively to the T_0 : light grey for decreases between 30% - 50%, medium grey 50%-70% and deep grey for higher than 70%.

Group	Cultivar	At T ₀	RH							
			85%	81%	75%	62%	45%	23%	6%	
Citron	Diamante	96	98	98	96	96	94	100	96	
	Ethrog	92	92	98	96	ND	94	54	50	
	Poncire commun	94	92	94	98	92	85	86	82	
	Corsican	98	92	94	88	96	96	88	90	
Citron	Alemow	86	94	90	88	ND	88	34	24	
hybrid	Femminello	96	100	98	98	96	90	92	92	
Lemon	Eureka	92	92	92	88	82	80	70	76	
	Volkamer	100	98	100	92	100	96	100	92	
	Rough	92	94	96	96	84	98	72	56	
	Mexican	94	94	94	96	92	94	76	70	
	Pursha	96	98	94	98	90	52	ND	ND	
Lime	Nestour	88	90	94	96	88	84	78	78	
	Rangpur	96	96	94	94	94	100	98	96	
Mandarin	Nasnaran	94	94	98	94	94	92	90	80	
	Sunki	98	90	92	76	68	48	32	10	
hybrid	Cleopatra	96	94	94	58	44	20	12	0	
	Willow leaf	96	98	90	66	40	32	12	4	
Mandada	Carvalhal	88	74	62	56	ND	28	ND	ND	
Mandarin	Changsa	88	90	78	60	54	40	6	0	
	Chandler	95	94	88	90	70	94	76	60	
	Reinking	92	84	94	92	88	78	ND	ND	
Pummelo	Sans Pépin	87	96	88	96	98	40	8	10	
	Eingedi	98	82	88	94	84	82	46	30	

Grapefruit	Marsh	96	92	96	88	73	64	56	36
	Duncan	90	90	86	84	82	72	20	18
	Parson Brown	82	72	72	46	28	26	18	12
Sweet orange	Pineapple	86	88	84	52	ND	40	4	0
	Mme Vinous	84	88	80	58	40	26	16	10
orange	Maroc	92	92	96	92	84	76	60	50
	Australian	88	92	82	80	82	80	48	48
Sour orange	Gou Tou	90	92	88	90	80	54	34	34
	Combava	94	88	82	76	82	76	88	72
Papeda	Kalpi papeda	98	98	92	92	94	88	88	80
	Melanesian	90	88	82	80	80	92	82	88
0	Carrizo	90	90	68	10	2	0	0	0
Citrange	C35	92	80	ND	82	ND	60	ND	ND
Trifoliate orange	Flying Dragon	82	28	24	40	0	0	0	0
	Rubidoux	74	24	28	58	0	0	0	0
	Marumi	80	42	36	24	0	0	0	0
V	Meiwa	88	40	10	4	0	0	0	0
Kumquat	Nagami	72	60	50	20	16	0	0	0

Discussion

Seed quality and maturity

Three main factors important to seed storage are moisture content, temperature and seed quality. Among these factors, seed quality is an ambivalent and complex factor because it is subject to environmental and genetic controls [35]. Most often, seed lots are of heterogeneous quality, often due to differences in flowering period, dry matter accumulation, dormancy, and maturation date, all of which are genetically regulated characteristics that affect seed longevity [36]. Seed quality, in terms of sensitivity to desiccation, has been suggested as the reason for the variability observed in wild Coffea arabica species [37]. Seed development initiates by embryogenesis followed by an embryogenic morphogenesis. After that, developing seeds enter in a maturation stage, corresponding to reserve accumulation, by reorganization of metabolism and synthesis of storage compounds (starch, storage protein and oil). Finally, the desiccation stage is the last step of seed maturation [38]. Seed desiccation plays a role in redirecting metabolism from developmental to a germinative mode by inducing hydrolytic enzymes essential to the growth of seedling [39]. Seed desiccation is an active stage in terms of gene expression and metabolism [40]. The transition of developing seeds from the phase of reserve accumulation to desiccation is associated with distinct gene expression and metabolic switches. Maximum desiccation tolerance is often acquired at mass maturity (max. dry weight accumulation) [12,39]. The use of seeds that have not reached their maximum maturity and therefore tolerance to desiccation could explain the contradictory results in the literature on seed responses to desiccation [41]. Our results are in agreement with this description of seed maturation by the increase of germination capacity related to the water loss. This desiccation phase is therefore necessary

for the seed to acquire maximum germination capacity. The time required for the seeds to reach full maturity (>80%) depends on the variety, with nearly 45 days separating the earliest ('Volkamer' lemon) from the latest (sour orange) even though flowering occurs at the same time for both species. Once the fruits have reached maturity, the water content of the seeds and the germination rate no longer vary during the fruit harvesting period [42]. These stages of seed development are often identified using non-destructive visual maturity indicators such as the size, color, and hardness of the fruit or seeds [43,44]. To obtain seeds with uniform maturity, which is essential for comparing and interpreting desiccation tolerance capacities, it is useful to identify maturity markers. Because the fruits of citrus trees used as rootstocks are not consumed, it is difficult to determine indicators of fruit ripeness. In general, there is a good correlation with abscission [44]. The easiest indicator to observe is the size (or mass) of the fruit. When it exceeds 85% of its final weight, the seed germination reaches its maximum (> 90%) regardless of the variety. The water content of the seeds in the fruit is also a good indicator of seed maturity. The germination is highest when it is close to or below 55%. Water content was the indicator used in subsequent experiments to ensure optimal seed maturity before treatment.

Fluctuation of water exchanges between seeds and the atmosphere

The deadline to reach a stable WC (cessation of water exchange between seeds-atmosphere-salt solution) varies according to the *citrus* species, suggesting different water flow regulation through the membrane of seed teguments. Seed coats are often suggested as a contributing factor (chemical or physical inhibitors) in the erratic, slow and reduced germination in response to desiccation and storage that has been observed in *citrus* seeds [14,45,46]. The

development of physical dormancy via a seed coat barrier to water uptake and gas exchange [47]. Crane [48] suggested such interactions may account for the damage observed in intermediate species on long-term storage [48]. *Citrus* seeds have characteristically high oil content, between 37% to 52% lipid content [9]. Therefore, this phenomenon could contribute to the variable and unusual seed physiology observed in *citrus*. Thus, analysis of seed oils may allow more accurate interpretation of survival responses in *citrus* species after drying.

Seed ageing

Seed aging is a natural process that causes a decrease in seed viability and vigor over time, reducing their ability to germinate. This phenomenon is of great concern to plant breeders and has implications for the preservation of plant genetic resources. Physiologically, aging seeds undergo a series of complex biochemical changes, including lipid peroxidation, protein oxidation, DNA damage, and changes in the antioxidant system [49]. This decline is caused by various factors, including the accumulation of Reactive Oxygen Species (ROS), changes in the composition and integrity of membrane lipids, and alterations in gene expression and protein synthesis [50]. The accumulation of ROS causes them to react with unsaturated fatty acids and causes changes in cell membranes, such as lipid peroxidation and, ultimately, their destruction [51]. Enzymatic detoxification and cell membrane repair are the main ways to delay ageing [49]. The activity of ROS-trapping enzymes and non-enzymatic antioxidant enzymes gives seeds high resistance to oxidative damage and minimizes damage to cells [52]. No studies have yet been conducted on the molecular changes that occur during the ageing of citrus seeds that could explain the rapid loss of germination when seeds are stored at room temperature. In our study, seed ageing is particularly accelerated in trifoliate oranges (P. trifoliata). This is evident early on, as germination rates are low for high water contents (81 and 85%) because the time taken to reach equilibrium for WC is close to 50 days at 25 °C (data not shown), whereas at 75% RH, WC equilibrium is reached after only 25 days. Germination is therefore higher at this RH. Low temperatures (4 °C), even if they slow down the flow of water between seeds and the atmosphere, help to limit the ageing phenomenon.

Diversity of seed response to desiccation in the citrus family

It is well known that seeds of most *Citrus* species are not orthodox and that their long-term storage under conventional conditions (low seed water content and at freezing temperatures) is not possible [10]. Nevertheless, *citrus* species show high variability in storage responses, some are desiccation tolerant and others have complex intermediate seed storage behavior, with some loss of viability at lower moisture contents [10]. Lime (*Citrus aurantifolia*), lemon (*C. limon*), Rough lemon (*C. jambhiri*) and grapefruit (*C. paradisi*) are the most tolerant species; at low water contents (<10%), they maintain germination rates above 80% [9,20,53]. In contrast, 'Calamondsi' (*C. madurensis*), 'Savage' citrange, 'Sacaton' citrumelo, trifoliate orange (*P. trifoliata*) and mandarin (*C. reticulata*) are known to be *citrus* fruits that are sensitive to seed desiccation [9,27,54]. However, there are also

many discrepancies between different studies. For example, for sour orange (C. aurantifolia) seeds with water contents between 10 and 14%, the germination fluctuates between 40 and 80% [8,20]. Even greater variations are observed in lemon (C. limon), where the germination varies from 1 to 85% for water contents of 10 and 7.5% respectively [21,22]. It is therefore difficult to make comparisons between different studies because drying conditions vary between studies (coated or uncoated seeds; drying using air flow, silica, atmosphere with controlled relative humidity, temperature). Germination conditions could also be a cause of variation. For example, lemon (C. limon) seeds needed a higher germination temperature (i.e. 30 °C) and a longer germination time (5 weeks) post drying [15]. Many germination tests in citrus had previously been conducted over shorter time periods and at a lower temperature (e.g. 20 °C) [15]. This may be a factor involved (i.e. insufficient time for germination) in some of the variable reports of desiccation sensitivity in citrus. A longer time for water uptake is suggested as the reason for the resultant delay in germination on drying of citrus seeds [46]. In our study, where seed maturity is controlled and drying and germination conditions are identical for all varieties and species, we do not observe any fluctuations between clonal varieties of the same species. The varietal diversity of most secondary species cultivated for fruit consumption is based solely on the clonal selection of somatic bud mutations identified in orchards. This applies, among others, to lemon trees (C. limon, [24]), orange trees (C. sinensis, [55]), and grapefruit trees (C. paradisi, [56]). For each species, cultivars have the same level of tolerance to desiccation. Another critical point could be the classification of citrus fruits and the use of a variety as a reference for a species. For example, in sour oranges (C. aurantifolia), there is genetic variability linked to hybridization involving different parents [25]. The 'Gou Tou' and 'Australian' varieties do not have the same genetic origin and therefore do not have the same genotype as the 'Maroc' variety, for example. The tolerance of their seeds to desiccation may therefore vary. Lemons are another example of genetic diversity, with varieties of diverse genetic origins. The 'Femminello' and 'Eureka' varieties are cultivars for consumption. They have the same genetic origin, a hybridization between a sour orange and a citron [1]. They differ only in mutations identified by clonal selection. The 'Volkamer' and 'Rough' lemons, used as rootstocks, are the result of crosses between citrons and mandarins [24]. Although genetically different, these lemon varieties have the same high level of drought tolerance.

The inheritance of tolerance to low water content

Species, *C. aurantifolia, C. macrophylla, C. limon, C. jambhiri, C. karna*, and *C. limonia* share citron as a common ancestor [24]. Although some of them have a parent that is sensitive to low water content (mandarin and sour orange), they have inherited the drought tolerance of *C. medica*. This trait may be dominant in this ancestral species. The example of citranges is slightly different. 'Carrizo' is thought to have inherited the sensitivity of its pollinating parent, the trifoliate orange (*P. trifoliata*) while, 'C35' citrange is more tolerant to desiccation than its other parent, the sweet orange. This is therefore a case of transgressive inheritance. 'Nasnaran' mandarin is very tolerant to desiccation. But in reality, this is a

hybrid between mandarin and a Papeda (*C. micrantha*, not present in this study) [57]. His tolerance is therefore a legacy from his Papeda progenitor. Other secondary species related to *C. reticulata* differ in their level of tolerance. Sour oranges (*C. aurantium*) have a tolerance equivalent to that of pummelo, while oranges (*C. sinensis*) inherited the sensitivity of mandarins. The pummelo (*C. maxima*) has transmitted its tolerance to the grapefruit (*C. paradisi*), which is a cross between *C. sinensis* and *C. maxima* [1].

Conclusion

To impartially assess the tolerance of seeds to different degrees of desiccation, it is first necessary to ensure the quality of the seeds, i.e., their optimum germination capacity. The time required to reach this maturity varies depending on the species and can be assessed by the water content of the seeds, which must be less than 55%. To prevent accelerated aging, seeds must be dried at 4 °C or 25 °C, but in the latter case for a short period (21 days). Seed tolerance to desiccation varies considerably depending on the species. All varieties of citrons and their hybrids (lemons and limes) have a high tolerance to low water content (\sim 5%) and their seeds can be considered as orthodox. Germination capacity decreases with seed water content in grapefruit, pummelos, and sour oranges, so their tolerance is moderate, while seeds of sweet oranges and mandarins are recalcitrant (sensitive to water content close to 15%). The most sensitive species are kumquats (Fortunella sp.) and trifoliate oranges (P. trifoliata), which lose more than 50% of their germination capacity at water contents close to 25%. Tolerant species can be included in the seed cryopreservation program using the desiccation methods developed in this study.

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Conflicts of Interest

The authors declare no conflict of interest.

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