

Laboratory report on seed health test

- Amount of seed-borne pathogens
- Amount of disease incidence

Objectives of seed health testing

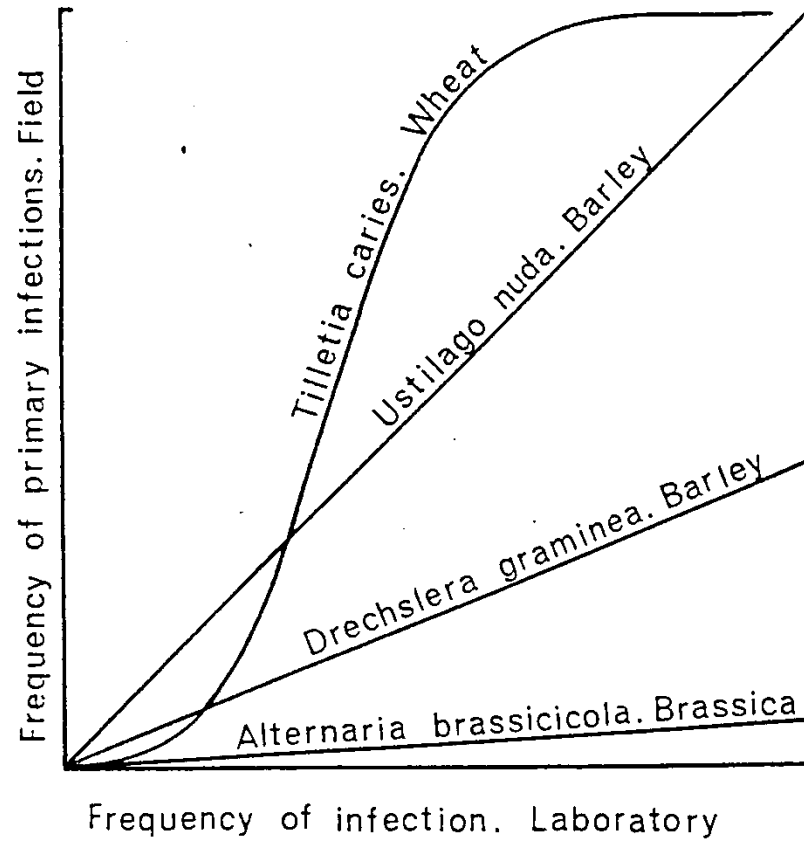
- Evaluation for planting value
- Seed treatment
- Certification
- Quarantine
- Human consumption & animal feed

Tolerance/Threshold level

(Evaluation for planting value)

Importance of primary infections for:

1. Disease development
2. Yield reduction
3. Infection of new seed crop

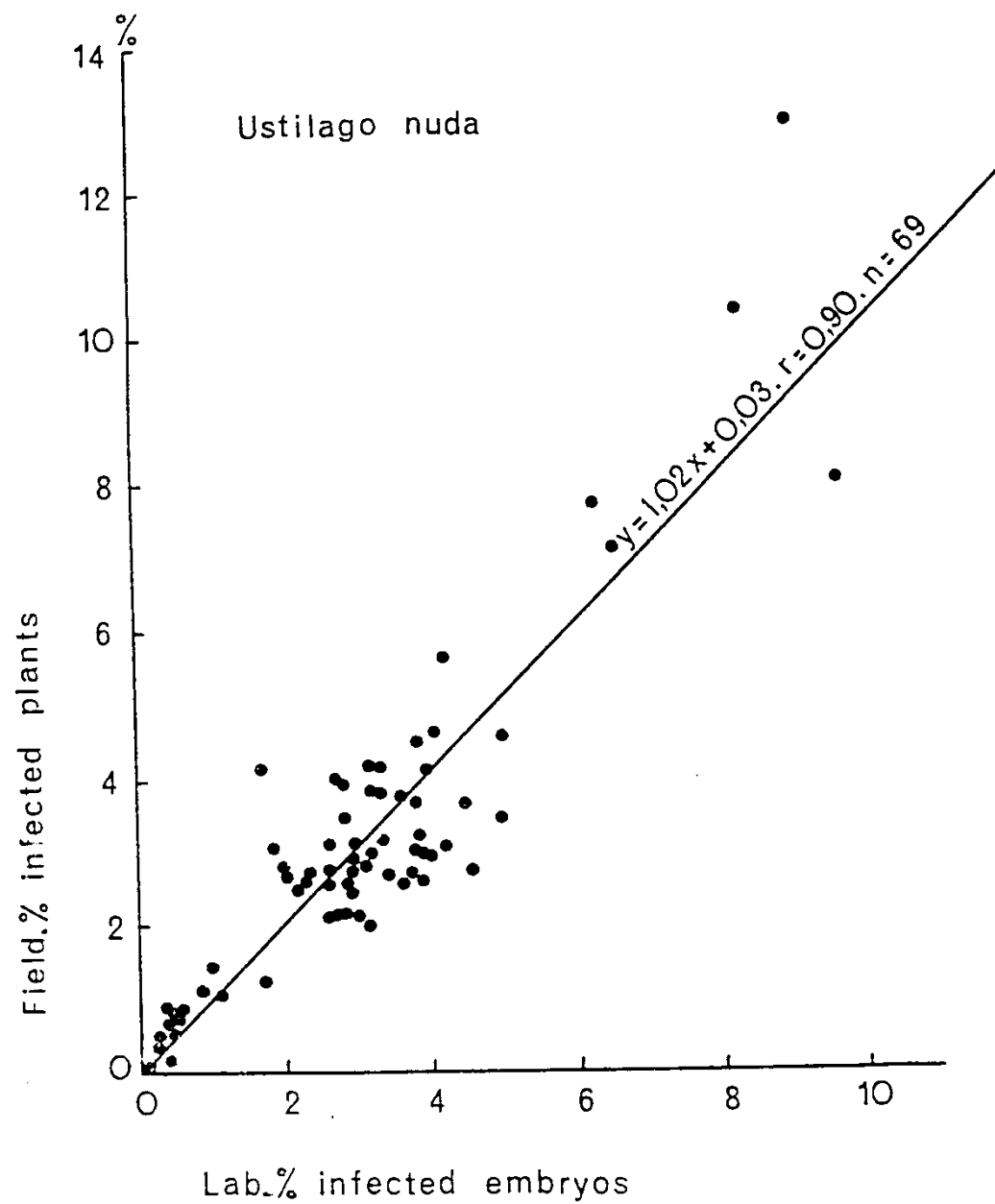


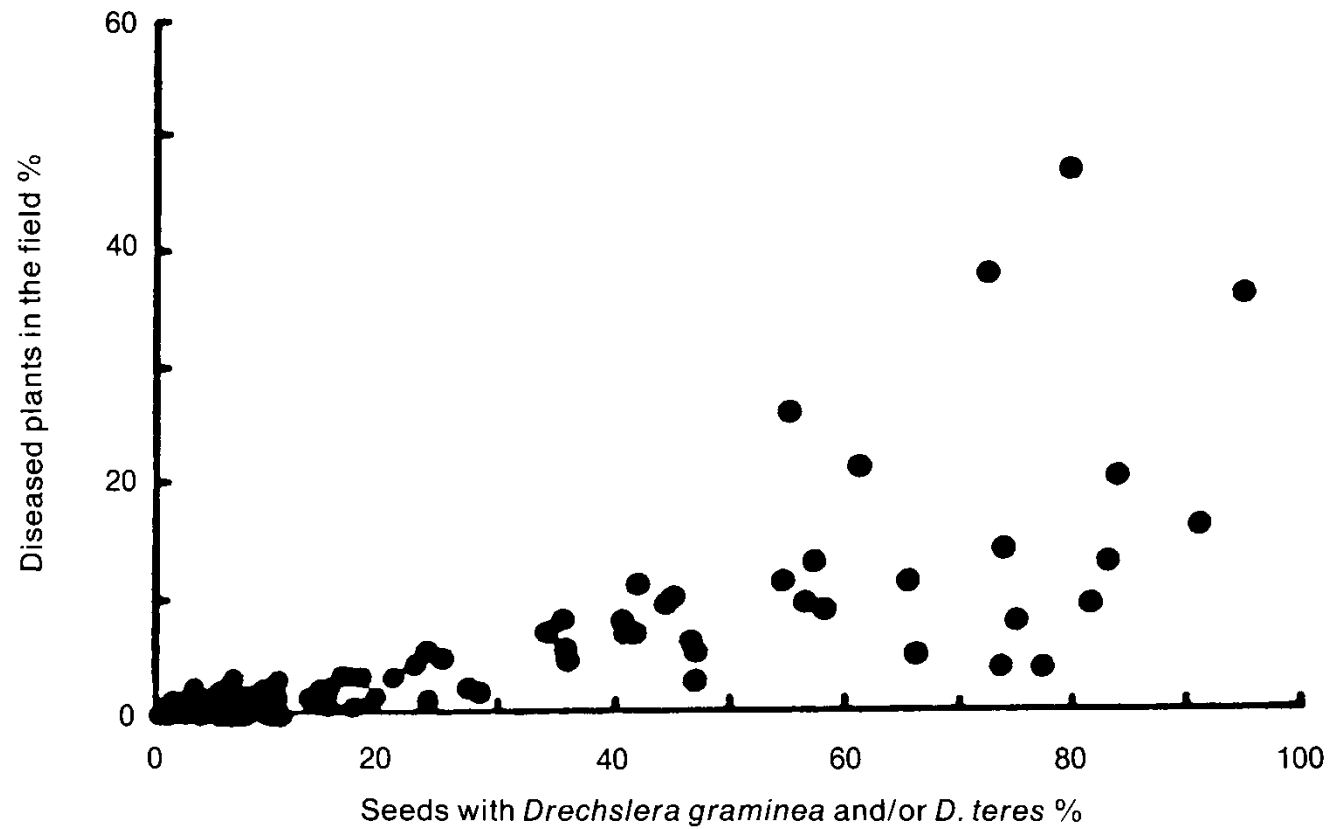
Types of relationships between test results
and primary infections in the field.

Seed transmission relationships.

Pathogen	% infection in		Transmission ratio	Crop	Reference
	Lab/glhse	Field soil			
<i>Polyspora lini</i>	15.0	1.7	9:1	Flax	Henry and Campbell, 1938
<i>Colletotrichum lini</i>	66.3	17.0	4:1	Flax	Henry and Campbell, 1938
<i>Pyrenophora graminea</i> and <i>P. teres</i>	50–75	5–10	10:1 to 7.5:1	Barley	Jorgensen, 1977
	50–65	0–11	0:0 to 6.0:1	Barley	Jorgensen, 1977
<i>Ascochyta pisi</i>	11.2	3.3	4:1	Peas	Maude and Kyle, 1970
	6.3	0.4	16:1	Peas	Maude and Kyle, 1970
	34.0	6.5	5:1	Peas	Maude and Kyle, 1970
<i>Alternaria brassicicola</i>	62.0	11.0	6:1	Cabbage	Maude and Humpherson-Jones, 1980b
	11.5	1.2	10:1	Kale	Maude and Humpherson-Jones, 1980b
	1.5	0.0	0:0	Kale	Maude and Humpherson-Jones, 1980b
<i>Alternaria brassicae</i>	28.0	9.3	3:1	Cabbage	R.B. Maude, pers. comm., 1991
	10.5	1.2	9:1	Oilseed rape	R.B. Maude, pers. comm., 1991
	14.0	1.2	12:1	Oilseed rape	R.B. Maude, pers. comm., 1991
<i>Pseudomonas syringae</i> pv. <i>phaseolicola</i>	9.2	0.84	11.0:1	Phaseolus bean	Taylor, 1970b
	3.5	0.37	9.5:1	Phaseolus bean	Taylor, 1970b
	1.4	0.15	9.3:1	Phaseolus bean	Taylor, 1970b
	0.1	0.0	0:0	Phaseolus bean	Taylor, 1970b
<i>Pseudomonas syringae</i> pv. <i>phaseolicola</i>	16.1	1.80	8.9:1	Phaseolus bean	Taylor <i>et al.</i> , 1979b
	1.1	0.13	8.5:1	Phaseolus bean	Taylor <i>et al.</i> , 1979b
	2.4	0.22	10.9:1	Phaseolus bean	Taylor <i>et al.</i> , 1979b
	2.4	0.42	5.7:1	Phaseolus bean	Taylor <i>et al.</i> , 1979b
	5.4	0.57	9.5:1	Phaseolus bean	Taylor <i>et al.</i> , 1979b

Lab/glhse, laboratory/glasshouse tests

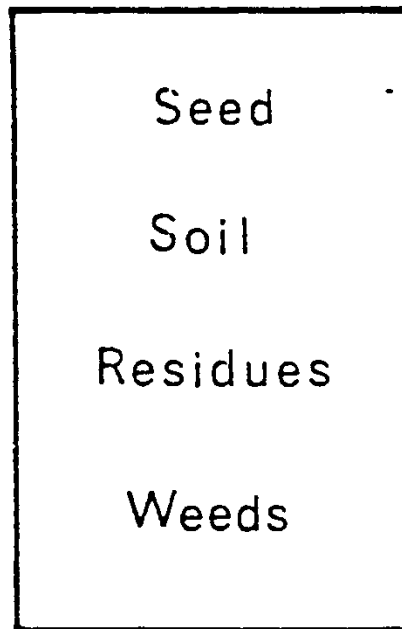




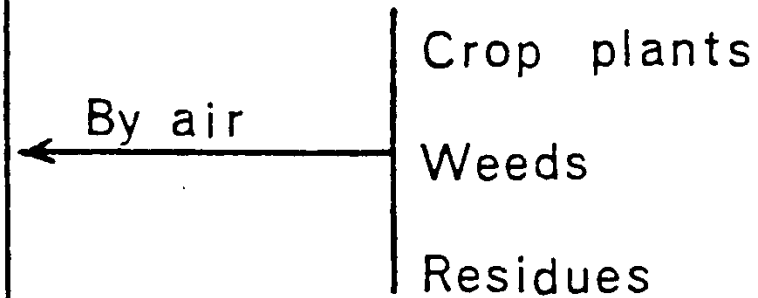
Correlation between percentage seeds infected with *Drechslera graminea* and/or *D. teres* according to the freezing blotter method and percentage diseased plants in the field

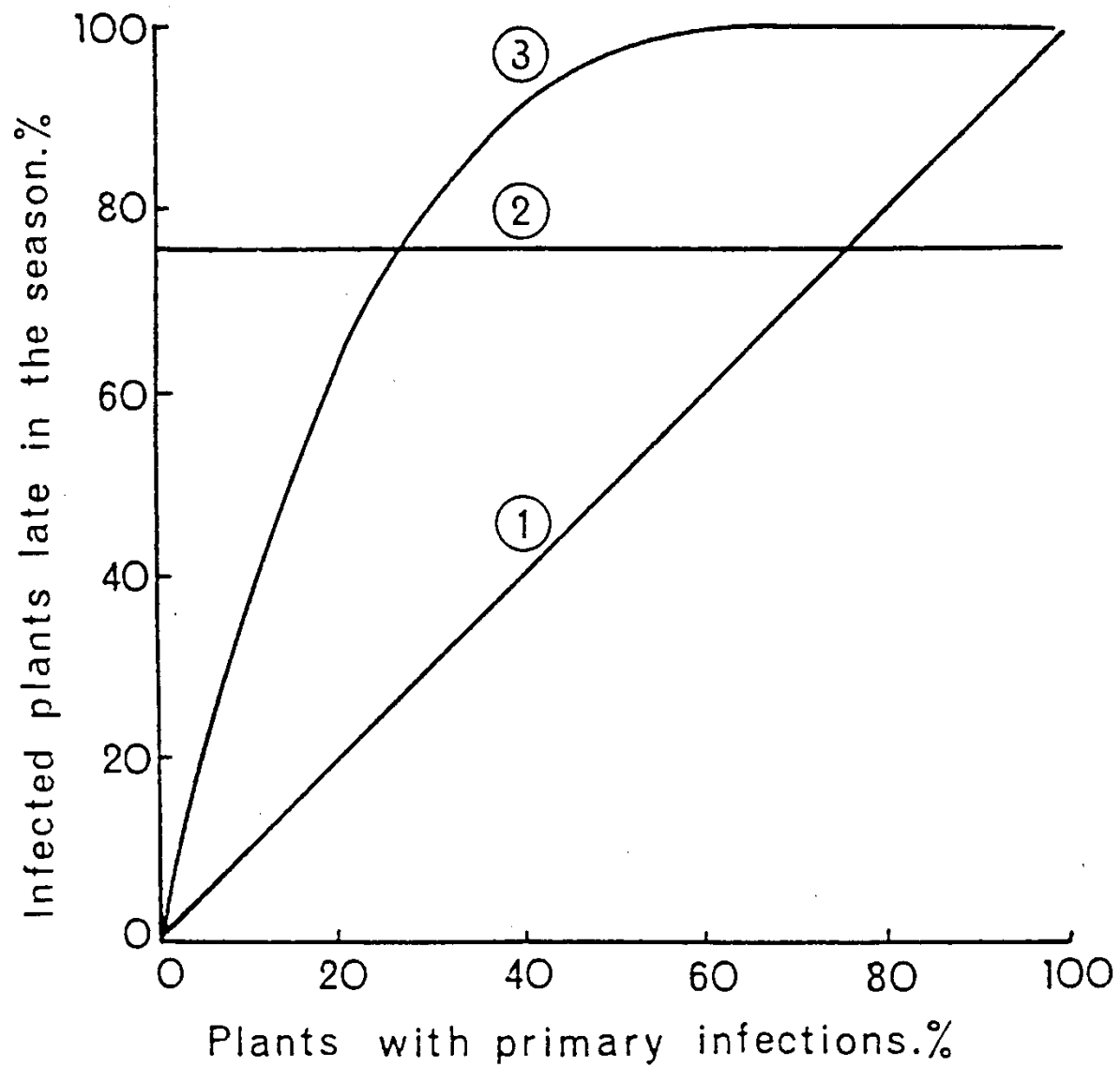
Sources of inoculum

In the field



Outside the field





Average infection of species of *Fusarium* and *Drechslera* in about 100 samples of barley seeds in each of 10 years.

Species	Average percentage of infected seeds										Average
	1965	1966	1967	1968	1969	1970	1971	1972	1973	1974	
<i>Fusarium avenaceum</i>	1.1	0.7	0.6	0.2	0.2	2.1	2.0	1.2	1.0	2.4	1.2
<i>Fusarium culmorum</i>	0.1	0.1	0.1	0.1	0.1	0.3	0.4	0.1	0.1	0.1	0.1
<i>Fusarium graminearum</i>	0.6	0.2	0.1	0.1	0.1	0.4	0.2	0.1	0.1	0.1	0.2
<i>Fusarium nivale</i>	1.3	0.3	0.1	0.5	0.1	0.5	0.9	5.1	0.2	1.4	1.0
<i>Fusarium poae</i>	0.4	0.5	0.5	0.6	0.9	2.9	1.6	1.2	1.8	2.1	1.3
<i>Fusarium</i> spp.	0.1	0.5	0.2	0.5	0.1	1.3	4.7	3.5	2.1	5.1	1.8
All species of <i>Fusarium</i>	3.6	2.3	1.6	2.0	1.5	7.5	9.8	11.0	5.3	11.2	5.6
<i>Drechslera sorokiniana</i>	0.8	4.0	3.5	0.8	0.7	5.8	5.2	4.4	2.0	1.4	2.7
<i>Drechslera gramineum</i> / <i>D. teres</i>	0.2	0.3	0.1	0.2	1.4	0.2	5.1	3.4	14.4	3.1	2.8

Correlation and regression between seed infection and root discoloration.

Pathogen	Year	Number of samples	Average percentage		Regression coefficient		Correlation coefficient r
			infected seed ($\bar{x}$)	discoloured rootlets ($\bar{y}$)	b*	a*	
<i>Fusarium nivale</i>	1972	92	4.74	5.30	0.71	1.96	0.78
<i>Fusarium nivale</i>	1973	23	11.04	14.83	1.10	2.75	0.91
<i>Drechslera sorokiniana</i>	1977	94	20.01	9.66	0.53	-0.88	0.89
<i>Drechslera sorokiniana</i>	1978	109	25.22	15.29	0.50	2.77	0.85
<i>Drechslera sorokiniana</i>	1979	76	19.36	11.11	0.41	3.08	0.82

*Regression: $y = bx + a$

The effect on grain yield of fungicidal treatment of seed with different degrees of infection determined by the Doyer filter paper method (calculated from Lindegaard and Christensen, 1971). Seed treatment: 100 ml organic mercury fungicide containing about 1.2% mercury.

Percentage infection	Number of experiments	Average increase in grain yield due to seed treatment kg/ha
0	148	-19.1 ⁺
1- 5	271	-22.6 ⁺⁺
6-15	176	-10.9 ^{NS}
16-40	48	29.5 ⁺⁺
41-70	18	50.6 ^{NS}

⁺ Significantly different from zero at the 94% level

⁺⁺ Significantly different from zero at the 99% level

^{NS} Not significantly different from zero

Inoculum thresholds and crop losses.

Crop	Pathogen	No. of affected seeds/ seedlings causing economic loss
Lettuce	Lettuce mosaic virus	1/30,000
Bean	<i>Pseudomonas syringae</i> pv. <i>phaseolicola</i>	1/10,000 to 1/16,000
Cabbage	<i>Leptosphaeria maculans</i>	1/10,000
Celery	<i>Septoria apiicola</i>	1/7000
Onion	<i>Botrytis allii</i>	1/100
Peas	<i>Ascochyta pisi</i>	> 5/100
Field bean	<i>Didymella fabae</i>	> 2/100

Standards for seedborne pathogens in UK seeds regulations (from Rennie, 1993).

Regulations	Crop	Pathogen	Seed category	Standard*
Fodder	Field beans	<i>Ascochyta fabae</i>	Pre-basic	1 infected seed in 1000
			Basic	2 infected seeds in 1000
			C1	2 infected seeds in 500
			C2	1% [†]
	Peas	<i>Pseudomonas syringae</i> pv. <i>pisi</i>	Multiplication categories	Nil in 1 kg
Vegetables	Brassicas	<i>Phoma lingam</i>	Basic	Nil in 1000 seeds [†]
	Red beet	<i>Phoma betae</i>	Basic	Nil in 200 seeds [†]
	Celery	<i>Septoria apiicola</i>	Basic and certified	Nil in 400 seeds [†]
	Celery	<i>Phoma apiicola</i>	Basic and certified	Nil in 400 seeds [†]
	Peas	<i>Ascochyta</i> spp.	Basic	Nil in 200 seeds [§]
			Certified	20 seeds max. in 200 [§]
	Lettuce	Lettuce mosaic virus	Basic and certified	Nil in 5000 seeds
	Phaseolus bean	<i>Colletotrichum lindemuthianum</i>	Basic	Nil in 600 seeds
	Phaseolus bean	<i>Pseudomonas syringae</i> pv. <i>phaseolicola</i>	Basic	Nil in 5000 seeds
	Vicia bean	<i>Ascochyta fabae</i>	Basic	Nil in 600 seeds

Plant Quarantine

Phytosanitary Certificate for hybrid seeds produced by contracted must be free from following organisms:

Host	Pathogen						
Tomato	<i>Clavibacter michiganensis</i> sub sp. <i>michiganensis</i>				<i>Phoma destructiva</i>		
	<i>Xanthomonas campestris</i> pv. <i>vesicatoria</i>				<i>Colletotrichum phomoides</i>		
	<i>Pseudomonas syringae</i> pv. <i>tomato</i> (<i>P. punctulans</i>)				Tomato Black Ring Virus		
	<i>Psedomonas corrugata</i>				Tomato Bushy Stunt Virus		
	<i>Pseudomonas solanocearum</i>				Tomato Ring Spot Virus		
	<i>Didymella lycopersici</i>				Tobacco Mosaic Virus		
	<i>Fusarium oxysporum</i>				Potato Spindle Tuber Viriod		
	or <i>F. oxysporum</i> f. sp. <i>lycopersici</i>				(Tomato Bunchy Top Virus)		
	or <i>F. oxysporum</i> f. sp. <i>lycopersici</i> race III						
	<i>Verticilium albo-atrum</i>				Tomato Big Bud (MLO)		
	<i>V. dahliae</i>						

Phytosanitary Certificate for hybrid seeds produced by contracted must be free from following organisms

Host	Pathogen
Watermelon	<i>Pseudomonas syringae</i> pv. <i>lachrymans</i>
	<i>Xanthomonas cucurbitae</i>
	<i>Fusarium oxysporum</i> f. sp. <i>niveum</i>
	<i>Mycosphaerella melonis</i>
	<i>Physalospora rhodina</i>
	<i>Colletotrichum lagenarium</i>
	Musk Melon Necrotic Spot Virus
	Squash Mosaic Virus
	Prunus Necrotic Ring Spot Virus

Chemical seed treatment

Purposes of Seed Treatment

- Eradicate seedborne pathogens



Loose
smut
fungus
in seed
embryo

Purposes of Chemical Seed Treatment

- Protect seeds and seedlings



Soybean
seed
decay and
seedling
blight

Alternatives and Supplements

- Broadcast sprays
- Certified seed
- Crop rotation
- Planting date
- Variety resistance
- Volunteer control

Advantages

- Seedborne pathogens are vulnerable
- Precision targeting
- Optimum timing
- Low dose
- Relatively easy to apply

Disadvantages

- Accidental poisoning
- Cropping restrictions
- Limited dose capacity
- Limited duration of protection
- Limited shelf life of treated seed
- Worker exposure
- Possible phytotoxicity

When to use a seed treatment

- When planting for seed production
- When using low test weight or older seed
- When planting in unfavorable conditions
- In fields with a history of stand problems
- When replanting is not feasible
- When seed-borne pathogens may be present
- When yield potential is high or seed is expensive

Classification of non-systemic and systemic organic fungicides used as seed treatments in the UK (from Hassal, 1990; Soper, 1991; Tomlin, 1994; Whitehead, 1995).

Type	Family/action	Group	Used in seed treatment
Non-systemic	Organosulphurs	Dithiocarbamates	Thiram
		Phthalimides	Captan
	Chlorinated aromatics	Chlorinated nitro compounds	(Quintozene)
		Chlorinated quinones	(Dichlone, chloranil)
Non-systemic or weakly systemic		Benzotriazines	Triazoxide
		Cyanopyrroles	Fenpiclonil
		Guanadines	Guazatine
		Azoles	Imazalil, prochloraz, bitertanol
Systemic		Dicarboximides	Iprodione, vinclozolin
		Benzimidazoles	Benomyl, thiabendazole
			carbendazim, fuberidazole
		Phenylamides	Carboxin
	Steroid inhibitors Reduction Demethylation	Morpholines	Fenpropimorph
			Triadimenol, flutriafol
		Azoles	tebuconazole
		Isoxazoles	Hymexazol
		Pyrimidines	Ethirimol
	Miscellaneous groups	Acylalanines	Metalaxyl
		Oxazole ketones	Oxadixyl
		Ethyl phosphonates	Fosetyl aluminium

Advantages and disadvantages of the major types of seed treatment formulation
(from Godwin *et al.*, 1988).

Formulation	Advantage	Disadvantage
DS (powder for dry seed treatment)	'On farm' application possible	Reduced adhesion Reduced efficacy
WS (water dispersable powder for slurry treatment)	'On farm' application possible	Efficacy not maximized
FS (flowable concentrate)	Accurate loading Good adhesion Excellent efficacy Application machinery can be washed out No organic solvents used	Specialist machinery needed for application
LS (solution)	Accurate loading Excellent adhesion Efficacy maximized	Organic solvents present can be phytotoxic Specialist machinery needed for application
FC (formulated products applied in polymer film coats)	Accurate loading Excellent adhesion Efficacy maximized No phytotoxicity No hazards to health or environment	Specialist machinery needed for application

