

**Effect of Nutrition on Postharvest Quality and Grey Mould
Development in Strawberries**

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Table of Contents

Table of Figures	vii
Table of Tables	xii
Abstract	xv
Declaration	xviii
Acknowledgements	xix
Abbreviations	xxi

Chapter One

General Literature Review

1.1 Introduction	1
1.2 Strawberry	1
1.3 Grey mould.....	4
1.3.1 Symptoms of grey mould.....	4
1.3.2 Causal agent of grey mould	5
1.3.3 Infection pathways of <i>B. cinerea</i>	7
1.3.3.1 Infection of flowers	7
1.3.3.2 Infection of ripening fruit	8
1.3.3.3 Infection and extensive colonisation of leaf residues.....	8
1.3.4 Disease cycle.....	9
1.3.5 Control of grey mould.....	11
1.4 Effect of calcium on fruit quality and postharvest diseases	11
1.4.1 Preharvest calcium application	14
1.4.2 Postharvest calcium application.....	16
1.5 Effect of boron on fruit quality and postharvest diseases	19
1.6 Justification and aims.....	22

Chapter Two

General Materials and Methods

2.1	Introduction	24
2.2	Plant materials	24
2.3	Standard nutrient solutions.....	25
2.4	Assessment of postharvest fruit quality	27
2.4.1	Shelf life.....	27
2.4.2	External appearance.....	27
2.4.3	Fruit firmness	27
2.4.4	pH.....	27
2.4.5	Soluble solids content (SSC)	27
2.4.6	Titrateable acidity (TA).....	28
2.5	Nutrient analysis.....	28
2.6	<i>B. cinerea</i> isolation, maintenance and inoculation.....	29
2.6.1	Isolation and maintenance.....	29
2.6.2	Preparation of conidia suspension	30
2.6.3	Inoculation of flowers	30
2.7	Assessment of grey mould development.....	30
2.8	Statistical analysis	31

Chapter Three

Effect of Preharvest Calcium Application

3.1	Introduction	32
3.2	Materials and methods	33
3.2.1	Plant materials and growth conditions.....	33
3.2.2	Treatments.....	33
3.2.2.1	Using a closed pot system	33
3.2.2.2	Using an automatic fertigation system	34
3.2.2.3	Using a manual fertigation system	37
3.2.3	Inoculation of flowers with <i>B. cinerea</i>	38

3.2.4	Storage conditions and postharvest assessment.....	39
3.2.5	Assessment of grey mould development	39
3.2.6	Statistical analysis.....	39
3.3	Results	40
3.3.1	Using a closed pot system.....	40
3.3.2	Inoculation of flowers with <i>B. cinerea</i>	40
3.3.3	Calcium treatments through automatic fertigation system	40
3.3.3.1	Effect on grey mould development	40
3.3.3.2	Effect on shelf life	44
3.3.3.3	Effect on fruit firmness.....	44
3.3.3.4	Effect on postharvest quality	47
3.3.3.4.1	External appearance.....	47
3.3.3.4.2	Soluble solids content	47
3.3.3.4.3	pH.....	47
3.3.3.4.4	Titrateable acidity	48
3.3.3.4.5	Effect on calcium content in fruit and leaf tissues.....	48
3.3.4	Calcium treatments through a manual fertigation system.....	58
3.3.4.1	Effect on grey mould development	58
3.3.4.2	Effect on shelf life	58
3.3.4.3	Effect on fruit firmness.....	59
3.3.4.4	Effect on postharvest quality	64
3.3.4.4.1	External appearance.....	64
3.3.4.4.2	Soluble solids content	64
3.3.4.4.3	pH.....	65
3.3.4.4.4	Titrateable acidity	65
3.3.4.4.5	Effect on calcium content in fruit and leaf tissues.....	65
3.4	Discussion	76
3.5	Conclusions	80

Chapter Four

Effect of Preharvest Boron Application

4.1	Introduction	81
4.2	Materials and methods	82
4.2.1	Plant materials and growth conditions	82
4.2.2	Treatments.....	82
4.2.3	Inoculation of flowers with <i>B. cinerea</i>	83
4.2.4	Storage conditions and postharvest assessment.....	84
4.2.5	Assessment of grey mould development	84
4.2.6	Statistical analysis.....	84
4.3	Results	85
4.3.1	Effect on grey mould development.....	85
4.3.2	Effect on fruit firmness	90
4.3.3	Effect on postharvest quality	90
4.3.3.1	External appearance and shelf life.....	90
4.3.3.2	Soluble solids content.....	91
4.3.3.3	pH	92
4.3.3.4	Titrateable acidity.....	92
4.3.4	Effect on boron content in leaf tissues	100
4.4	Discussion	104
4.5	Conclusions	107

Chapter Five

Effect of Postharvest Calcium Treatment

5.1	Introduction	109
5.2	Materials and methods	110
5.2.1	Fruit materials	110
5.2.2	Postharvest treatments	111
5.2.3	Inoculation of fruit.....	111
5.2.4	Storage conditions.....	111

5.2.5	Effect of calcium lactate on <i>B. cinerea</i> development	111
5.2.6	Effect of calcium chloride on <i>B. cinerea</i> development.....	112
5.2.7	Direct comparison of calcium lactate and calcium chloride in delaying <i>B. cinerea</i> development	113
5.2.8	Further evaluation of the most effective treatments from prior experiments	113
5.2.9	Botrytis fruit rot evaluation.....	114
5.2.10	Statistical analysis.....	114
5.3	Results	117
5.3.1	Effect of calcium lactate on <i>B. cinerea</i> development <i>in vitro</i>	117
5.3.2	Effect of calcium lactate on <i>B. cinerea</i> development in fruit	117
5.3.3	Effect of calcium chloride on <i>B. cinerea</i> development <i>in vitro</i>	123
5.3.4	Effect of calcium chloride on <i>B. cinerea</i> development.....	123
5.3.5	Direct comparison of calcium lactate and calcium chloride in delaying <i>B. cinerea</i> development	126
5.3.6	Further evaluation of the most effective treatments from prior experiments	130
5.4	Discussion	132
5.5	Conclusion.....	136

Chapter Six

Effect of Calcium and Boron on Botrytis Leaf Blight

6.1	Introduction	137
6.2	Materials and methods	138
6.2.1	Plant materials.....	138
6.2.2	Conidia suspension and inoculation of leaf	139
6.2.3	Evaluation of Botrytis leaf blight.....	140
6.2.4	Nutrient analysis	140
6.2.5	Statistical analysis.....	140
6.3	Results	142

6.3.1	Effect of calcium on the development of Botrytis leaf blight.....	142
6.3.2	Correlation between calcium concentration in nutrient solution, calcium content in leaf and blight lesion size.....	142
6.3.3	Effect of boron on the development of Botrytis leaf blight.....	149
6.3.4	Correlation between boron concentration in nutrient solution, boron content in leaf and blight lesion size.....	150
6.4	Discussion.....	156
6.5	Conclusion.....	159

Chapter Seven

General Discussion

7.1	Calcium and boron mobility may affect postharvest quality.....	161
7.2	Impact of calcium and boron on grey mould.....	162
7.3	Factors influence the amount of calcium taken into fruit during postharvest application.....	164
7.4	Cultivar comparison.....	165
7.5	Further research.....	166
7.6	Conclusions.....	167

References	168
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Appendix 1	182
-------------------------	------------

Appendix 2	183
-------------------------	------------

Appendix 3	184
-------------------------	------------

Appendix 4	185
-------------------------	------------

Table of Figures

Figure 1.1	A strawberry fruit, illustrating calyx, enlarged receptacle (edible portion) and achenes.....	3
Figure 1.2	Grey mould on strawberry.....	6
Figure 1.3	Longitudinal section of strawberry flower, illustrating pathways of flower infection: petals, pistils and stamens.....	8
Figure 1.4	Infection of ripening fruit. <i>Botrytis cinerea</i> spreads by fruit-to-fruit contact such as this.....	9
Figure 1.5	Disease cycle of grey mould on strawberry (modified from http://ohioline.osu.edu/hyg-fact/3000/3017.html).....	10
Figure 1.6	Schematic representation of cell walls, illustrating the Ca ⁺ bridges between pectin molecules in the cell walls (inset).....	13
Figure 2.1	Inoculated plants, showing how flowers were covered with a plastic bag to maintain moisture after inoculating with <i>Botrytis cinerea</i> or sterile water (as a control).	31
Figure 3.1	The closed pot system used to apply calcium.....	33
Figure 3.2	The automatic fertigation system in the glasshouse.....	36
Figure 3.3	Effect of preharvest calcium application through an automatic fertigation system on the percentage of fruit displaying rot for ‘Aromas’ fruit from <i>Botrytis cinerea</i> -inoculated flowers (solid line) and water-treated flowers as control (dashed line).....	42
Figure 3.4	Effect of preharvest calcium application through an automatic fertigation system on the percentage of fruit displaying rot for ‘Selva’ fruit developed from <i>Botrytis cinerea</i> -inoculated flowers (solid line) and water-treated flowers as control (dashed line).....	43
Figure 3.5	Effect of preharvest calcium application through the automatic fertigation system on shelf life of ‘Aromas’ (a) and ‘Selva’ (b) during storage at 10°C, 90±5% RH for 10 days.....	45

Figure 3.6	Effect of preharvest calcium application through the automatic fertigation system on fruit firmness of ‘Aromas’ (a) and ‘Selva’ (b) during storage at 10°C, 90±5% RH for 10 days.	46
Figure 3.7	Effect of preharvest calcium application through the automatic fertigation system on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Aromas’ during storage at 10°C, 90±5% RH for 0 to 10 days.	52
Figure 3.8	Effect of preharvest calcium application through the automatic fertigation system on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Selva’ during storage at 10°C, 90±5% RH for 0 to 10 days.	53
Figure 3.9	Effect of preharvest calcium application through the manual fertigation system on the percentage of fruit displaying rot for ‘Aromas’ fruit developed from <i>Botrytis cinerea</i> -inoculated flowers (solid line) and water-treated flowers as control (dashed line).	60
Figure 3.10	Effect of preharvest calcium application through the manual fertigation system on the percentage of fruit displaying rot for ‘Selva’ fruit developed from <i>Botrytis cinerea</i> -inoculated flowers (solid line) and water-treated flowers as control (dashed line).	61
Figure 3.11	Effect of preharvest calcium application through the manual fertigation system on shelf life of ‘Aromas’ (a) and ‘Selva’ (b) during storage at 10°C, 90±5% RH for 10 days.	62
Figure 3.12	Effect of preharvest calcium application through the manual fertigation system on fruit firmness of ‘Aromas’ (a) and ‘Selva’ (b) during storage at 10°C, 90±5% RH for 10 days.	63
Figure 3.13	Effect of preharvest calcium application through the manual fertigation system on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Aromas’ during storage at 10°C, 90±5% RH for 0 to 10 days.	69

Figure 3.14	Effect of preharvest calcium application through the manual fertigation system on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Selva’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	70
Figure 3.15	Appearance of plants grown in soil and manually fertigated with various concentrations of calcium.	71
Figure 4.1	Appearance of plants grown in soil and fertigated with various concentrations of boron.	87
Figure 4.2	Effect of preharvest boron application and inoculation with <i>Botrytis cinerea</i> on the number of ‘Aromas’ flowers that died (a), aborted (b) and developed into fruit (c).	88
Figure 4.3	Effect of preharvest boron application and inoculation with <i>Botrytis cinerea</i> on the number of ‘Selva’ flowers that died (a), aborted (b) and developed into fruit (c).	89
Figure 4.4	Effect of preharvest boron application on fruit firmness of ‘Aromas’ (a) and ‘Selva’ (b) during storage at 10°C, 90±5% RH for 10 days....	93
Figure 4.5	Effect of preharvest boron application on general appearance of fruit from ‘Aromas’ (a) and ‘Selva’ (b) at harvest then after 4, 8 and 10 days of storage at 10°C, 90±5% RH.	96
Figure 4.6	Effect of preharvest boron application on the expected duration of shelf life for ‘Aromas’ and ‘Selva’.....	97
Figure 4.7	Effect of preharvest boron application on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Aromas’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	98
Figure 4.8	Effect of preharvest boron application on soluble solids content (a), pH (b) and titratable acidity (c) of ‘Selva’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	99
Figure 5.1	Diagram of general methodology of postharvest calcium treatment.	115
Figure 5.2	Diagram of modified methodology used to further evaluate the most effective treatments: calcium lactate at 3000 ppm Ca and calcium chloride at 4500 ppm Ca as detailed in Section 5.2.8.....	116

Figure 5.3	Colony diameter of <i>Botrytis cinerea</i> on potato dextrose agar amended with calcium lactate at 0, 1500, 3000 and 4500 ppm Ca.....	117
Figure 5.4	Development of rot lesion area on fruit treated with calcium lactate and storage at 10°C, 90±5% RH for 7 days after inoculation with <i>Botrytis cinerea</i>	119
Figure 5.5	Effect of calcium lactate dips on the diameter of rot lesions after storage at 10°C, 90±5% RH for up to 7 days.....	120
Figure 5.6	Effect of calcium lactate and storage period after calcium treatment prior to inoculation on rot lesion development, 7 days after inoculation.	121
Figure 5.7	Effect of calcium lactate dips on rot lesion development after storage at 10°C, 90±5% RH for up to 7 days.....	122
Figure 5.8	Colony diameter of <i>Botrytis cinerea</i> on potato dextrose agar amended with calcium chloride at 0, 1500, 3000 and 4500 ppm Ca.....	123
Figure 5.9	Effect of calcium chloride and storage period after calcium treatment prior to inoculation on rot lesion development, 7 days after of inoculation.....	124
Figure 5.10	Effect of calcium chloride dips on rot lesion development after storage at 10°C for up to 7 days..	125
Figure 5.11	Comparison of the effect on Botrytis rot lesion area of treatment with calcium lactate (blue bars) and calcium chloride (pink bars) after 7 days of storage.....	127
Figure 5.12	Botrytis rot near the calyx (A) on fruit mock-inoculated with sterile nanopure water (SNW) by wounding at site B:.....	128
Figure 5.13	The percentage of fruit which presented visible rot near the calyx after 7 days of storage at 10°C, 90±5% RH.....	129
Figure 5.14	Effect of calcium lactate at 3000 ppm Ca (CL3000Ca) and calcium chloride at 4500 ppm Ca (CC4500Ca) on the development of Botrytis rot on late-season strawberries (May, 2006) (a) and on early-season strawberries (December, 2006) (b).....	131

Figure 6.1	<i>Botrytis cinerea</i> -inoculated leaves in a 15-cm diameter Petri dish....	141
Figure 6.2	Leaves of cultivar ‘Aromas’ 3 and 7 days after inoculation with <i>Botrytis cinerea</i> , illustrating the effect of calcium on the severity of Botrytis leaf blight on leaves detached from plants that received 0, 100, 300 and 500 ppm Ca.....	144
Figure 6.3	Leaves of cultivar ‘Selva’ 3 and 7 days after inoculation with <i>Botrytis cinerea</i> , illustrating the effect of calcium on the severity of Botrytis leaf blight on leaves detached from plants that received 0, 100, 300 and 500 ppm Ca.....	145
Figure 6.4	Effect of calcium, incorporated in fertigation nutrient solution, on severity of Botrytis leaf blight in ‘Aromas’ during 7 days after inoculation.....	146
Figure 6.5	Effect of calcium, incorporated in fertigation nutrient solution, on severity of Botrytis leaf blight in ‘Selva’ during 7 days after inoculation.....	147
Figure 6.6	Leaves of cultivar ‘Aromas’ 3 to 7 days after inoculation with <i>Botrytis cinerea</i> , illustrating the effect of boron on the severity of Botrytis leaf blight on leaves detached from plants that received 0, 0.25, 0.5 and 1.0 ppm B.	151
Figure 6.7	Leaves of cultivar ‘Selva’ 3 to 7 days after inoculation with <i>Botrytis cinerea</i> , illustrating the effect of boron on the severity of Botrytis leaf blight on leaves detached from plants that received 0, 0.25, 0.5 and 1.0 ppm B.	152
Figure 6.8	Effect of boron, incorporated in fertigation nutrient solution, on severity of Botrytis leaf blight in ‘Aromas’ during 7 days after inoculation.....	153
Figure 6.9	Effect of boron, incorporated in fertigation nutrient solution, on severity of Botrytis leaf blight in ‘Selva’ during 7 days after inoculation.....	154

Table of Tables

Table 2.1	Composition of 10X full strength Hoagland's solution [modified from Hoagland and Arnon (1938)]. The volume was made up to 20 L with reverse osmosis water.....	26
Table 3.1	The composition of standard nutrient solutions applied to Mount Compass sand prior to planting in the closed pot system [modified from Hoagland and Arnon (1938)]......	35
Table 3.2	Experimental calcium treatments applied to Mount Compass sand immediately prior to planting in the closed pot system.....	35
Table 3.3	Calcium content in 20 L fertigation tank used in the automatic fertigation system.	37
Table 3.4	The amount of calcium sulphate added into a 20 L fertigation tank for use in manual fertigation.	38
Table 3.5	The percentage of flowers that died resulting from different methods of inoculation with <i>Botrytis cinerea</i> conidia suspension.	41
Table 3.6	The percentage of flower death resulting from inoculation with <i>Botrytis cinerea</i> conidia suspension at different concentrations.	41
Table 3.7	Effect of preharvest calcium application through the automatic fertigation system on external appearance of 'Aromas' during storage at 10°C, 90±5% RH for 0 to 10 days.	49
Table 3.8	Effect of preharvest calcium application through the automatic fertigation system on external appearance of 'Selva' during storage at 10°C, 90±5% RH for 0 to 10 days.	50
Table 3.9	Effect of calcium treatment through the automatic fertigation on foliar concentration of nutrients in 'Aromas'.....	54
Table 3.10	Effect of calcium treatment through the automatic fertigation on nutrient concentrations in 'Aromas' fruit.	55
Table 3.11	Effect of calcium treatment through the automatic fertigation on foliar concentration of nutrients in 'Selva'.	56

Table 3.12	Effect of calcium treatment through the automatic fertigation on nutrient concentrations in ‘Selva’ fruit.....	57
Table 3.13	Effect of preharvest calcium application through the manual fertigation system on external appearance of ‘Aromas’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	67
Table 3.14	Effect of preharvest calcium application through the manual fertigation system on external appearance of ‘Selva’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	68
Table 3.15	Effect of calcium treatment through the manual fertigation on foliar concentration of nutrients in ‘Aromas’.....	72
Table 3.16	Effect of calcium treatment through the manual fertigation on nutrients concentration in ‘Aromas’ fruit.....	73
Table 3.17	Effect of calcium treatment through the manual fertigation on foliar concentration of nutrients in ‘Selva’.....	74
Table 3.18	Effect of calcium treatment through the manual fertigation on nutrient concentrations in ‘Selva’ fruit.....	75
Table 4.1	The amount of boric acid and potassium hydroxide added into a 20 L fertigation tank.....	83
Table 4.2	Effect of preharvest boron application on external appearance of ‘Aromas’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	94
Table 4.3	Effect of preharvest boron application on external appearance of ‘Selva’ during storage at 10°C, 90±5% RH for 0 to 10 days.....	95
Table 4.4	Effect of preharvest boron application on foliar concentration of nutrients in ‘Aromas’.....	102
Table 4.5	Effect of preharvest boron application on foliar concentration of nutrients in ‘Selva’.....	103
Table 6.1	Calcium content in ‘Aromas’ and ‘Selva’ leaves.....	148

Table 6.2	The linear correlation coefficients between calcium concentration in the fertigation nutrient solution, calcium content in leaf and blight lesion size (7 days after inoculation) in cultivar ‘Aromas’ in September 2006.	148
Table 6.3	The linear correlation coefficients between calcium concentration in the fertigation nutrient solution, calcium content in leaf and blight lesion size (7 days after inoculation) in cultivar ‘Selva’ in September 2006.	148
Table 6.4	Boron content in ‘Aromas’ and ‘Selva’ leaves.....	155
Table 6.5	The linear correlation coefficients between boron concentration in the fertigation nutrient solution, boron content in leaf and blight lesion size (7 days after inoculation) in cultivar ‘Aromas’ in July 2007.	155
Table 6.6	The linear correlation coefficients between boron concentration in the fertigation nutrient solution, boron content in leaf and blight lesion size (7 days after inoculation) in cultivar ‘Selva’ in July 2007.	155

Abstract

Strawberries are an extremely perishable fruit mainly due to their soft texture and sensitivity to fungal infection. The fungal pathogen *Botrytis cinerea* is responsible for grey mould on strawberries and is the main causal agent of postharvest decay and subsequent economic loss. As an alternative to fungicides, manipulation of plant nutrition, such as calcium and boron, has been suggested as a means of disease management. This project investigated the effects of calcium and boron application on fruit quality and grey mould development in strawberry.

The effect of calcium on fruit quality, grey mould development and leaf blight in strawberry cultivars ‘Aromas’ and ‘Selva’ was investigated through preharvest and postharvest applications. To determine the effect of preharvest application, calcium sulphate in 0.25X strength Hoagland’s solution was applied at 0, 100, 300 and 500 ppm Ca through fertigation. Fully-ripened fruit were harvested and evaluated for postharvest quality at harvest and then after storage at 10°C, 90±5% RH for 2 to 10 days. Although fruit firmness of both cultivars declined slightly during storage, this was not affected by preharvest calcium application. Similarly, preharvest calcium treatment had no effect on the external appearance, pH, soluble solids content (SSC) or titratable acidity (TA).

No grey mould development was observed on fruit at harvest when flowers were inoculated with a conidia suspension of *B. cinerea* (10^4 conidia per mL). However, fruit harvested from plants that received calcium at any concentration had less incidence of grey mould during storage at 10°C, 90±5% RH for 14 days than fruit harvested from plants that received no calcium for both cultivars. For ‘Aromas’, 79% and 51% of fruit, and for ‘Selva’, 69% and 43% of fruit, showed rot when treated with 0 and 500 ppm Ca, respectively. The shelf life of ‘Aromas’ and ‘Selva’ increased by about 8% when plants received 500 ppm Ca in comparison with plants that received 0 ppm Ca.

After 7 days of incubation at 22 to 24°C, there was no difference between blight lesions on wound-inoculated detached leaves from different calcium treatments for either cultivar. However, the lesions on ‘Selva’ were smaller than on ‘Aromas’. The calcium levels in leaves from plants that received calcium at any concentration were adequate for strawberry growing and significantly higher ($P < 0.05$) than in leaves from plants that received 0 ppm Ca. However, calcium treatment did not ensure transfer of calcium to fruit tissues.

Calcium lactate and calcium chloride were used as postharvest calcium treatments at 1500, 3000 and 4500 ppm Ca. Fruit of ‘Selva’ were dipped in calcium solution for 5 min and wound-inoculated with *B. cinerea* (10^6 conidia per mL). Calcium lactate and calcium chloride at 3000 and 4500 ppm Ca, respectively, were most effective in delaying Botrytis rot development on ‘Selva’ after 7 days of storage at 10°C, 90±5% RH. Storage for least 24 h after calcium dips prior to inoculation was required to delay the development of fruit rot. Fruit harvested early in the season seemed to be less susceptible to grey mould than those harvested later. However, calcium treatment tended to be more effective when applied to late-season fruit.

Preharvest boron treatment, applied as for calcium but at 0, 0.25, 0.5 and 1.0 ppm B, had no effect on fruit firmness of either cultivar. However, firmness of ‘Aromas’ fruit was slightly greater than ‘Selva’ fruit for all treatments. The amount of boron applied had no effect on the external appearance, pH, SSC or TA for either cultivar after storage of fruit for up to 10 days.

Application of boron had no effect on fruit grey mould development in either cultivar. Furthermore, boron had minimal effect on the incidence of blight on wound-inoculated detached leaves of ‘Aromas’ 7 days after inoculation. However, blight lesion diameters on ‘Selva’ leaves in the 1.0 ppm B treatment (8.0 mm) were significantly smaller ($P < 0.001$) than in the 0 ppm B treatment (13.0 mm).

Phytotoxicity was observed in boron treatments even at the level considered optimum for strawberry growing. Severity increased with increasing boron concentration but no consistent effect on flower death or flower abortion was observed.

In conclusion, strawberry is sensitive to boron toxicity. Calcium may enhance fruit firmness and, consequently, delay grey mould development if calcium penetrates the fruit. Postharvest calcium treatment tended to be more effective in delaying development of grey mould when applied to late-season fruit. Calcium lactate is a potential alternative to calcium chloride for reducing decay caused by *B. cinerea* in strawberry without providing undesirable bitterness. This finding may provide a basis for application in industry.

Declaration

I declare that the work presented in this thesis contains no material which has been accepted for the award of any other degree or diploma in any university or other tertiary institution. To the best of my knowledge and belief, this thesis contains no material previously published or written by another person, except where due reference has been made in the text.

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Abbreviations

μL	microlitre
ANOVA	Analysis of Variance
ARC	Australian Research Council
B	boron
Ca	calcium
cm	centimetre
DN	day-neutral
e.g.	for example
<i>et al.</i>	and others
FAO	Food and Agricultural Organisation of the United Nations
g	gram
GA	General Appearance
h	hour
i.e.	that is
ICP-OES	Inductively Coupled Plasma Optical Emission Spectrometer (ARL model 3580 B)
kg	kilogram
kgf	kilogram-force
L	litre
LD	long-day
LSD	Least Significant Difference
mg	milligram
min	minute
mL	millilitre
mm	millimetre
mm^2	square millimetre
$^{\circ}\text{C}$	degree celsius

pH	potential of hydrogen
ppm	parts per million
RH	relative humidity
RO	reverse osmosis
SARDI	South Australia Research and Development Institute
SD	short-day
SE	standard error
SNW	sterile nanopure water
SSC	soluble solids content
TA	titratable acidity
TSS	total soluble solids
UC	University of California
v/v	volume by volume