Copyright is owned by the Author of the thesis. Permission is given for a copy to be downloaded by an individual for the purpose of research and private study only. The thesis may not be reproduced elsewhere without the permission of the Author.

In vitro and in vivo studies on treatment and prevention of bovine mastitis

A thesis presented in partial fulfilment of the requirements for the degree of Philosophy Doctor

in

Veterinary Science

at Massey University, Palmerston North, New Zealand.

Kiro R Petrovski

© Kiro R Petrovski, 2011 Except as provided by the Copyright Act 1994, no part of this publication may be reproduced or stored in a retrieval system in any form or by any means without the prior written permission of the copyright owner

Abstract

Mastitis prevalence on dairy farms depends on the number of infected cows and the duration of each intramammary infection. Strategies aiming to influence these factors are the subject of research presented in this thesis.

Decreasing the duration of infection can be achieved by successfully treating infected quarters. Treatment of mastitis can occur during lactation or in the dry period. Treatment success is influenced by the concentration of antimicrobial achieved at the site of infection and the length of time it is present. The concentration of antimicrobial should exceed the relevant minimal inhibitory concentration. The susceptibility of mastitiscausing organisms varies among geographical areas and over time. New Zealand's susceptibility data demonstrated a high susceptibility to penicillin. A formulation containing this antimicrobial was administered to healthy lactating cows milked once or twice daily. The concentrations of penicillin in milk were above the minimal inhibitory concentrations for the entire inter-dosing interval. Doubling the number of treatments or milking once-a-day resulted in a significantly increased time above the minimal inhibitory concentrations.

The number of new infections is greatest during the early dry period in mature cows and in the pre-calving period in both heifers and mature cows. Pre-partum administration of delayed release antimicrobial formulations in heifers decreased the incidence of clinical mastitis and resulted in better reproductive performance, but not in increased milk production, when compared to control heifers. More effective prevention of new infections within the dry period was achieved by administering a novel teat sealant to mature cows when compared to a commercial teat sealant and untreated controls.

Strategies for shortening the duration of intramammary infections and decreasing the number of affected cows at the start of lactation investigated in this thesis should reduce the prevalence of mastitis on dairy farms in New Zealand.

KEY WORDS: Aetiology, Antibiotic, Antimicrobial, Challenge, Dry period, Experimental challenge, Heifers, Individual Cow Somatic Cell Count, Internal teat sealant, Mastitis, Milking frequency, Penicillin G, Reproductive performance, *Staphylococcus aureus, Streptococcus uberis, Streptococcus dysgalactiae*, Susceptibility, Time Above the Minimal Inhibitory Concentrations, Udder, Withholding Period.

Acknowledgments

The PhD journey is long and extremely demanding. This thesis represents a multidisciplinary research effort and many people have contributed to its realisation. It is a very difficult task to remember everyone who took a part in helping me through this journey of a PhD candidacy.

I would like to say thanks to my Supervisors: Prof Norman B Williamson (Chief Supervisor), Prof Timothy J Parkinson (Internal Supervisor), Prof Ian J Tucker (External Supervisor) and Assoc Prof Nicolas Lopes-Villalobos (Internal Supervisor) for their guidance and support in this journey.

Thanks also to the co-authors in the various papers: Mohamed Abdalla, Alfredo Caicedo-Caldas, Alejandro (Alex) Grinberg, Richard Laven, Scott McDougall and Paul Rapnicki.

Massey University and particularly the Institute of Veterinary, Animal and Biomedical Sciences are thanked for employing me with financial support from *Bomac a company of Bayer Ltd*.

The experiments reported in this thesis would not have happened without financial support from *Bomac a company of Bayer Ltd* (formerly *Bomac Laboratories Ltd*) and I would like to express my sincere gratitude for their support.

I am indebted to all farm staff in the involved farms, staff at *Gribbles Veterinary Laboratories* Auckland, Christchurch, Dunedin, Hamilton, and Palmerston North, New Zealand and the *Veterinary Diagnostic Laboratory, University of Minnesota*, Minnesota, USA and *MilkTest NZ* (formerly *SAITL Dairy Laboratories*), Hamilton, New Zealand for their friendship, cooperation and interest in my research. Without their full support and cooperation the work reported in this thesis could not have been carried out.

In this thesis five experiments involved the use of dairy cows. For each individual experiment appropriate Animal Ethics Committee was granted by various committees. I would like to say thanks to the committee members for their understanding and approvals of the experiment designs.

It is not possible to list the names of everyone who contributed either directly or indirectly towards this work. Hence, I wish to express my sincere gratitude to all those who discussed matters related to this work and are not mentioned individually. The

anonymous reviewers of the manuscripts are thanked for their constructive criticism of the submitted work.

Some people who assisted me in the preparation of this thesis by taking part in some of the work, helping with agreements for funding, discussion, advice, statistical analysis or just being there when needed:

- 1. Past and present staff at *Bomac a company of Bayer Ltd*, Auckland, New Zealand: Fadil Al Alawi, Danielle Baxter, Ray Castle, Richard Emslie, Renee Hetherington, Wayne Leech, Connell McLaren, Don McLaren, Lina Ma, Rob Nottingham, Michael Syme, Warren Tully and Bruce Vautier.
- 2. Past and present staff at *Estendart Ltd*, Palmerston North, New Zealand: Alan Alexander, Hailey Baird, Rene Corner, Kara Eaton, Gilly Evans, Kathryn Hutchinson, Lina Yang, Jude Vautier and many casual staff.
- 3. Past staff at *Invoco AgResearch*, Palmerston North: Jeremy Lind, Leonora Pearson and Brian Timms.
- 4. Past and present staff at *JL Vet Services*, Palmerston North: Jeremy Lind and many casual staff.
- 5. Past and present farm staff John Allen, Wendy Allen, Hamish Doohan, Christine Finnegan, Conrad Maeke, Phil Martin and Grant Rudman.
- 6. Massey University, past and present staff at the *Agricultural Farm Services* Louise Beazley, Natalie Butcher, Martin Chesterfield, Gareth Evans, Erin Hutchinson, Mark Lawrence, Natalia Martin and Byron Taylor.
- 7. Massey University, past and present staff at the Institute of Veterinary, Animal and Biomedical Sciences Frazer Allen, Rukhshana Akhter, Hugh Blair, Andrea Coleman, Georgie Cowley, Gina de Nicolo, Sharron Hawira-Seanoa, Debbie Hill, Litty Kurian, Sue Leathwick, Gayle McKenna, Hamish Mack, Carol Orr, Rebecca Patisson, Quentin Roper, Kevin Stafford, Peter Wildbore and Dianna Willson.
- 8. Massey University, past and present staff at the *Research Management Services*: Nicola Carse, Mark Cleaver, Don Brown, Leith Hutton and Carolina Tate.
- 9. Massey University, Turitea campus, Library staff Chris Good and Bruce White.
- 10. Massey University, Large Animal Teaching Unit Liz Gillespie and Robin Whitson.
- 11. Massey University, past and present staff at the Veterinary Teaching hospital Lesley England, Kevin Lawrence, Jenny Nixey, Alan Thatcher and Jenny Weston.

- 12. *University of Otago, School of Pharmacy*, past and present staff: Olaf Bork and Zimei Wu.
- 13. Duncan Hedderley from *The New Zealand Institute for Plant and Food Research Ltd*, Palmerston North.
- 14. Hassan Hussein from Cognosco, Morinsville, New Zealand.
- 15. Yuanxiang Shi, a visiting scholar from China.

Finally, my family and my dear wife, Ljubica (Bube) for putting up with me during this project.

Table of Contents

Abstra	ıct	I
Ackno	wledgments	III
List of	Equations	XIII
List of	Figures	XV
List of	Tables	XIX
List of	Abbreviations	XXV
1.	General introduction to the thesis	3
1.1	Background and areas of research of the thesis	3
1.2	Aims of the thesis	10
1.3	Research objectives of the thesis	10
	1.3.1 Part One	10
	1.3.2 Part Two	11
	1.3.3 Part Three	11
	1.3.4 Part Four	11
1.4	References	12
Part c	one	15
Part c		
	Introduction to part one: Antimicrobial susceptibility References	19
2.	Introduction to part one: Antimicrobial susceptibility	19 26
 2.1 3. 	Introduction to part one: Antimicrobial susceptibility References	19 26 gnostic
 2.1 3. 	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia	1926 gnostic33
2.2.13.labora	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand	1926 gnostic33
2.2.13.labora3.1	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract	1926 gnostic3334
2. 2.1 3. labora 3.1 3.2	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction	1926 gnostic3334
2. 2.1 3. labora 3.1 3.2	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction Materials and methods	1926 gnostic333435
2. 2.1 3. labora 3.1 3.2	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction Materials and methods 3.3.1 Microbiological methods	1926 gnostic33343535
2. 2.1 3. labora 3.1 3.2 3.3	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction Materials and methods 3.3.1 Microbiological methods 3.3.2 Statistical analysis	1926 gnostic33343535
2. 2.1 3. labora 3.1 3.2 3.3	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction Materials and methods 3.3.1 Microbiological methods 3.3.2 Statistical analysis Results	1926 gnostic333435353639
2. 2.1 3. labora 3.1 3.2 3.3 3.4 3.5	Introduction to part one: Antimicrobial susceptibility References Culture results from 25,288 milk samples submitted to veterinary dia stories from August 2003 to December 2006 in New Zealand Abstract Introduction Materials and methods 3.3.1 Microbiological methods 3.3.2 Statistical analysis Results Discussion	1926 gnostic333435363943

4.	A descriptive analysis of the antimicrobial susceptibility of m	astitis~causing
bacter	ria isolated from samples submitted to commercial diagnostic labora	atories in New
Zealaı	nd (2003–2006)	55
4.1	Abstract	55
4.2	Introduction	56
4.3	Materials and methods	57
	4.3.1 Criteria for selection of cases	57
	4.3.2 Microbiological methods	58
	4.3.3 Other records	59
	4.3.4 Statistical analysis	59
4.4	Results	60
	4.4.1 Antimicrobial susceptibility of streptococci	61
	4.4.2 Antimicrobial susceptibility of staphylococci	64
	4.4.3 Antimicrobial susceptibility of other bacterial species	66
4.5	Discussion	66
4.6	Acknowledgements	77
4.7	References	77
5.	Susceptibility to antimicrobials of mastitis-causing Staphylo	coccus aureus,
Strept	tococcus uberis and Strep. dysgalactiae from New Zealand and the U	SA as assessed
by the	e disk diffusion test	83
5.1	Abstract	83
5.2	Introduction	84
5.3	Materials and methods	85
	5.3.1 Statistical analysis	87
5.4	Results	89
	5.4.1 Level of susceptibility	94
	Susceptibility of Staphylococcus aureus	96
	Susceptibility of Streptococcus spp	96
	5.4.2 Zones of inhibition	99
	5.4.3 Discordant isolates	104
5.5	Discussion	104
5.6		
0.0	Conclusion	
5.7	Conclusion Acknowledgments	108

6.	Correlation of the antimicrobial susceptibility of Staphylococcu	<i>is aureus</i> and
strepto	ococci isolated from bovine milk samples collected in New Zealand v	when tested by
the ag	gar disk diffusion and broth microdilution methods	115
6.1	Abstract	115
6.2	Introduction	116
6.3	Materials and methods	117
6.4	Results	120
	6.4.1 Overall	120
	6.4.2 Ampicillin	123
	6.4.3 Cloxacillin	126
	6.4.4 Enrofloxacin	128
	6.4.5 Neomycin	130
	6.4.6 Oxytetracycline	132
	6.4.7 Penicillin	134
6.5	Discussion	136
6.6	Conclusion	139
6.7	Acknowledgments	139
6.8	References	139
Part	two	143
7.	Introduction to Part Two: Effects of milking frequency on pharm	acokinetics of
penici	illin G administered by the intramammary route	
7.1	References	153
8.	Milking frequency affects the penicillin G elimination time	s from milk.
conce	ntrations and recovery rate following intramammary administration	<i>'</i>
8.1	Abstract	159
8.2	Introduction	160
8.3	Materials and methods	161
	8.3.1 Experimental animals	161
	8.3.2 Treatment and procedures	162
	8.3.3 Statistical analysis	
8.4	Results	
	8.4.1 Elimination times	166
	8.4.2 Time above MIC	166

	8.4.3 Amount of drug recovered	167
8.5	Discussion	168
8.6	Conclusion	172
8.7	Acknowledgments	172
8.8	References	172
Part :	three	177
9.	Introduction to Part Three: treatment of heifers for mastitis pre-calving	181
9.1	References	187
10.	Treatment before calving of heifers for mastitis improves their rep	roductive
perfor	mance, but not their milk production	195
10.1	Abstract	195
10.2	Introduction	196
10.3	Materials and methods	197
	10.3.1 Procedures	197
	10.3.2 Statistical Analysis	198
10.4	Results	199
	10.4.1 Incidence of clinical mastitis	199
	10.4.2 Prevalence of subclinical mastitis	199
	10.4.3 Days-in-milk	203
	10.4.4 Milk production	203
	10.4.5 Reproductive performance	204
10.5	Discussion	205
10.6	Conclusions	208
10.7	Acknowledgments	208
10.8	References	208
Part :	four21	5
11.	Introduction to Part Four: antimicrobial teat sealant for use at drying off	219
11.1	References	224
12.	A preliminary evaluation of the efficacy of two novel internal tea	at sealant
formu	ations against bacterial challenge in the early dry period	229
	Abstract	
12.2	Introduction	230
122	Materials and methods	230

	12.3.1 Cows and treatments administered	231
	12.3.2 Procedures	231
	12.3.3 Statistical analysis	239
12.4	Results	240
	12.4.1 Length of dry period	240
	12.4.2 Palpation scores	240
	12.4.3 Clinical mastitis	242
	12.4.4 Genotyping isolates of Streptococcus uberis from clinical cases	242
	12.4.5 Intramammary infection	242
	12.4.6 Somatic cells	244
12.5	Discussion	244
12.6	Acknowledgments	246
12.7	References	246
13.	Efficacy of a novel internal dry period teat sealant containing 0.5% chl	orhexidine
agains	t experimental challenge with Streptococcus uberis in dairy cattle	251
13.1	Abstract	251
13.2	Introduction	252
13.3	Materials and methods	253
	13.3.1 Animals	253
	13.3.2 Treatment products and treatment administration	254
	13.3.3 Procedures	254
	13.3.4 Statistical analysis	257
13.4	Results	260
	13.4.1 Dry period	260
	13.4.2 Udder palpation scores	260
	13.4.3 Clinical mastitis during the palpation period	261
	13.4.4 Milk culture results	262
13.5	Discussion	264
13.6	Conclusion	
13.7	Acknowledgments	268

14.	General discu	assion	275
14.1	Part One		275
14.2	Part Two		279
14.3	Part Three		280
14.4	Part Four		281
14.5	Further resear	rch needs identified	283
14.6	References		284
15.	List of referen	nces	289
16.	Bibliography		315

List of Equations

Equation 1-1 Calculations of the prevalence of mastitis on	a dairy farm	3
Equation 4.1 Back-transforming of the model outputs		60

List of Figures

Figure 1.1. Incidence and prevalence of intramammary infections in a herd throughout the season when each infection is of a long duration. The prevalence at the moment of observation is high despite the low incidence
Figure 1.2. Incidence and prevalence of intramammary infections in a herd throughout the season when each infection is of a short duration. The prevalence at the moment of observation is low despite the high incidence
Figure 1.3. Incidence of intramammary infections through the lactational cycle provided no dry cow therapy is used
Figure 2.1.Trend in the number of herds and average herd size from 1974/75 to 2009/10
Figure 3.1. Origin of the milk samples submitted for culturing to five commercial laboratories in New Zealand from August 2003 to December 2006
Figure 3.2. Monthly isolates of <i>Staphylococcus aureus</i> and <i>Streptococcus uberis</i> as percentage of all samples submitted from August 2003 to December 2006 from winter to autumn
Figure 3.3. Monthly isolates of other commonly-isolated mastitis-causing organisms Zealand as percentage of all samples submitted from August 2003 to December 2006 from winter to autumn.
Figure 3.4. Percentage of isolates of <i>Streptococcus uberis</i> , <i>Staphylococcus aureus</i> by seasons
Figure 5.1 Agar disk diffusion and E-test of <i>Staphylococcus</i> isolate
Figure 5.2 Agar disk diffusion and E-test of streptococcal isolate
Figure 6.1. Schema of the graphic presentation of each antimicrobial/isolates of causative organism susceptibility testing outcome
Figure 6.2. Distribution of susceptibility results of isolates of <i>Staphylococcus aureus</i> or streptococci to ampicillin

Figure 6.3. Distribution of susceptibility results of isolates of Staphylococcus aureus or
streptococci to cloxacillin
Figure 6.4. Distribution of susceptibility results of isolates of Staphylococcus aureus on
streptococci to enrofloxacin
Figure 6.5. Distribution of susceptibility results of isolates of Staphylococcus aureus on
streptococci to neomycin
Figure 6.6. Distribution of susceptibility results of isolates of Staphylococcus aureus on
streptococci to oxytetracycline
Figure 6.7. Distribution of susceptibility results of isolates of Staphylococcus aureus or
streptococci to penicillin
Figure 8.1. Treatment by the intramammary route using partial insertion technique
Figure 8.2. Milk samples for various analysis and reserves
Figure 8.3. <i>DeLaval</i> in-line samplers
Figure 8.4. Concentrations of procaine penicillin G in milk (mg/kg) in cows treated 3
times and milked once-a-day or twice daily and treated 6 times and milked twice daily
167
Figure 10.1. Moving average (10-daily) of the predicted geometric mean of individual
cow somatic cell count in treated and control heifers in their first lactation201
Figure 10.2. Percentage of treated heifers with high individual cow test-day somatic cell
count (ICSCC≥200,000/mL) percentage of new infections (change of ICSCC from low to
high) and percentage of cured cases (change of ICSCC form high to low) through their
first lactation
Figure 10.3. Percentage of control heifers with high individual cow test-day somatic cell
count (ICSCC≥200,000/mL) percentage of new infections (change of ICSCC from low to
high) and percentage of cured cases (change of ICSCC form high to low) through their
first lactation

Figure 10.4. Predicted milk volume using the method of Ali and Schaeffer (1987) in
treated and control heifers during their first lactation
Figure 11.1. Treatment design for the second challenge study
Figure 12.1 X-ray picture of the location of the internal teat sealant post treatment with
the sealant showing as a bright white area within the teat cavity
Figure 12.2 X-raying of the teats after treatment in order to evaluate the position of the
teat sealant as shown in Figure 12.1
Figure 12.3. Average daily udder palpation score
Figure 13.1. Survival analysis from treatment to incidence of clinical mastitis during the
first 34 days after drying~off262

List of Tables

Table 4.3. Number of tests and estimated percentage susceptibility, with 95% CI, to
antimicrobials for isolates of Streptococcus uberis from milk samples submitted to five
commercial laboratories in New Zealand over a 40-month period (2003-2006), adjusted
for effect of year of testing, island of origin of sample, and the interaction of year and
antimicrobial63
Table 4.4 Number of tools and estimated parameters are estimated to a superintibility with QEO/ CI to
Table 4.4. Number of tests and estimated percentage susceptibility, with 95% CI, to
antimicrobials for isolates of <i>Staphylococcus aureus</i> from milk samples submitted to five
commercial laboratories in New Zealand over a 40-month period (2003–2006), adjusted
for effect of year of testing, island of origin of sample, and the interaction of year and antimicrobial
antimicropiai
Table 4.5. List of available pharmaceutical products containing a minimum of one
antimicrobial authorised for the treatment of bovine mastitis in New Zealand (2009), as
indicated on the label's recommendations
Table 5.1. Disk potency of antimicrobials used in the study (µg~ micrograms)87
Table 5.1. Plak potency of untilineroplate used in the study (µg interograms)
Table 5.2. Susceptibility of isolates of Staphylococcus aureus and streptococci by country
90
Table 5.3. Susceptibility of streptococcal isolates by country
Table 5.4. Prevalence of susceptibility (mean ± SE) of Staphylococcus aureus and
streptococci isolated from milk samples collected in New Zealand and the USA to a range
of antimicrobials95
Table 5.5. Prevalence of susceptibility (mean \pm SE) of <i>Streptococcus uberis</i> and <i>Streptococcus</i>
dysgalactiae isolated from milk samples collected in New Zealand and the USA to a range
of antimicrobials98
Table 5.6. Diameters of zones of inhibition (mean \pm SE) for susceptible and resistant
isolates of Staphylococcus aureus and streptococci isolated from milk samples collected ir
New Zealand and the USA
Table 5.7. Diameters of zones of inhibition (mean \pm SE) for susceptible and resistant
isolates of Streptococcus uberis and Strep. dysgalactiae isolated from milk samples
collected in New Zealand and the USA

Table 6.1. Interpretive criteria for bacteria isolated from animals (if not stated otherwise based on the Clinical and Laboratory Standards Institute, 2008)
Table 6.2. Average sizes of inhibition at agar disk diffusion (mm \pm SE) compared to microdilution test results (μ g/mL) for various antimicrobial and causative organism combinations
Table 6.3. Correlation parameters for the zones of inhibition measured using the again disk diffusion method with the MIC obtained by the broth microdilution method for various antimicrobial/causative organism combinations
Table 8.1. Elimination times of penicillin G from milk (means \pm SE) in cows under different milking frequency and treatment regime treated with Lactapen G by the intramammary route
Table 8.2. Amount of penicillin G recovered from milk of cows under different milking frequency and treatment regime treated with Lactapen G by the intramammary route
Table 9.1. Reported prevalence of intramammary infections in heifers before the first calving based on culture
Table 9.2. Prevalence of intramammary infections in heifers around the first calving
Table 9.3. Rate of clinical mastitis in heifers around calving or during the first lactation
Table 10.1. Changes in the percentage of heifers with high individual somatic cell counts High ICSCC; SCC≥200,000/mL) from low to high (New infections) and high to low (Cured infections) approximating rates of new infections and cures from subclinical mastitis and the percentage of high ICSCC on test-day in treated and control heifers during their first lactation
Table 10.2. Means ± SE and differences of three reproductive parameters in treated and control heifers
Table 11.1. Summary of studies on the efficacy of teat sealants alone and their use ir combination with other products

Table 12.1.Quarter and teat examination and palpation scores and description (criteria developed by KRP)
Table 12.2. Means and standard errors of the lengths of the dry period in days per group
Table 12.3. Palpation scores in the first 34 days after drying-off adjusted for the random effect of an individual cow
Table 12.4. Effect of treatment on palpation scores in the first 34 days after drying-off
Table 12.5. Incidence of clinical mastitis during the palpation period by treatment
Table 12.6. Prevalence of quarters with intramammary infection after calving243
Table 12.7. Summary of the culture results in per cent (and numbers) after calving (D0 ~ day of calving; D4 ~ day 4 after calving) for all sampled quarters (including those treated for clinical mastitis during the palpation period)
Table 12.8. Means and their standard errors of the somatic cell scores (log of the somatic cell count divided by 1,000) among groups after calving
Table 13.1. The concentration of colony-forming units of a <i>Streptococcus uberis S210</i> strain per millilitre in the challenge broth at different challenge days
Table 13.2. Quarter and teat examination and palpation scores and description255
Table 13.3. Prevalence of infected or non-infected quarters in percent ± standard errors among groups
Table 13.4. Effect of treatment on palpation scores in the first 34 days after drying-off
Table 13.5. Distribution of cases of clinical mastitis, the probability of a quarter being affected by clinical mastitis and probability of a quarter of being affected with clinical mastitis caused by the challenge organism in the first 34 days after drying-off261

Table	13.6.	Summary	of t	he cult	ure re	sults	ın p	ercent	(and	numbers)	among	groups
												263
Table	13.7.	Means and	1 959	% confi	dence	interv	als c	of the o	uarte	r level infe	ection ra	te after
calvin	g											264

List of abbreviations

Abbreviation	Meaning
ACVM	Agricultural Compounds and Veterinary Medicines Group (part of MAF New Zealand)
ATS	Anti-Infective-Containing Internal Teat Sealant
BAGG	Buffered Azide Glucose Glycerol broth
BMSCC	Bulk Milk Somatic Cell Count
CAMP	Christie-Atkins-Munch-Petersen test
CI	Confidence Interval
CLSI	Clinical and Laboratory Standards Institute
CMT	California Mastitis Test
CNS	Coagulase-Negative Staphylococci
DCT	Dry Cow Therapy
EMEA	European Medicines Agency
EUCAST	European Committee on Antimicrobial Susceptibility Testing
I	Intermediate Susceptibility
ICSCC	Individual Cow Somatic Cell Count
IU	International Units
kg	Kilogramme
L	Litre
LF	Left Front
Ltd	Limited
μg	Microgram
mg	Milligram
mL	Millilitre
MIC	Minimal Inhibitory Concentration

List of Abbreviations continued

MRL	Maximum Residue Levels
MRSA	Methicillin-Resistant Staphylococcus aureus
NIRD	National Institute for Research in Dairying
OAD	Once-a-Day
PBP	Penicillin Binding Protein
PFGE	Pulse Field Gel Electrophoresis
R	Resistant
RR	Rear Right
S	Susceptible
SAMM	Seasonal Approach to Managing Mastitis
SAS	Statistical Analysis System
SCC	Somatic Cell Count
SCS	Somatic Cell Score
T>MIC	Time above the Minimal Inhibitory Concentrations
TD	Twice daily
WHP	Withholding period