

Strategy for monitoring and sustainable integrated control or eradication of *Glossina brevipalpis* and *G. austeni* (Diptera: Glossinidae) in South Africa

by

Karin Kappmeier Green

Submitted in partial fulfillment of the requirements for the degree Philosophiae Doctor in the Faculty of Natural and Agricultural Science, University of Pretoria

> Supervisor: Prof. C.H. Scholtz Co-supervisor: Prof. B.G. Williams

> > March 2002



SUMMARY

Glossina brevipalpis Newstead and G. austeni Newstead (Diptera: Glossinidae) are the vectors of trypanosomosis or nagana in cattle in N.E. KwaZulu-Natal, South Africa. Before intervention by means of target technology could be applied successfully to control these species, studies were still needed on the two species' attraction to natural host odours, trapping, their movement and dispersal, feeding responses towards hosts and their geographical distribution. Studies that were conducted with host odours to find an attractive odour for G. austeni, proved that CO₂ was seemingly the main attractive component of host odour for this species. The existing chemicals of the best SA odour, comprising of octenol released at c. 9,1 mg/h, 4-methylphenol released at c.15,5 mg/h and acetone released at c. 350 mg/h, still remained to be the main attractive components for G. brevipalpis. A sticky trap, namely a bicoloured electric blue/black XT, was refined to use in tsetse distribution surveys. A new trap, the H trap, was developed and proved to be effective in catching relatively high numbers of both species. This trap was used to capture live tsetse for mark-release-recapture studies to assess the population size and movement of a tsetse population. These studies revealed that target densities of about 4 targets/km² for G. brevipalpis and 7 targets/km² for G. austeni should be effective to control these species successfully with odour-baited insecticide-impregnated targets. G. austeni was confined to densely shaded areas but it still traversed short distances of up to 345 m of "unsuitable" habitat between pockets of vegetation. G. brevipalpis was considered a much more mobile fly and traversed wide areas of 1,345 m. Both species were readily attracted to cattle, but not to goats nor bushpig. They also fed more readily on cattle. Both species would also feed at night. It was recommended that insecticide-treated cattle could be used as mobile targets to control both G. brevipalpis and G. austeni in areas where cattle predominate.

Tsetse surveys through the northeastern parts of KwaZulu-Natal showed that there were two distinct bands of distribution for *G. brevipalpis*. The main sources of this species seemed to be the game reserves and other natural areas. *G. austeni* was more widespread with a continuous north to south distribution. A Geographic Information System was used to map tsetse distribution and their apparent densities. This was collated with trypanosomosis incidence and prevalence, diptank (cattle) distribution,



land tenure/designation, landcover and vegetation types, which were also mapped. Finally, a strategy was proposed for the monitoring and sustainable integrated control and eventual eradication of both *G. brevipalpis* and *G. austeni* throughout N.E. KwaZulu-Natal. This involves the subdivision of the area into five manageable zones with successive pre-suppression, suppression and eradication operations following in each of the zones. With this proposed strategy eradication of both species could be achieved within 8 - 12 years after initiation.



SAMEVATTING

Glossina brevipalpis Newstead en G. austeni Newstead (Diptera: Glossinidae) is vektore van trypasosomose (nagana) in beeste in noordoos KwaZulu-Natal, Suid-Afrika. Voordat ingryping deur middel van teiken-tegnologie suksesvol toegepas kan word om hierdie spesies te beheer, was dit nodig om die spesies se aanlokking na natuurlike gasheerreuke, vangmetodes, hulle beweging en verspreiding, voergedrag ten opsigte van gashere en hulle geografiese verspreiding, te bestudeer. Studies wat met gasheerreuke uitgevoer was om 'n aanloklike geur vir G. austeni te vind, het bewys dat CO₂ bleikbaar die hoof aanlokkingskomponent vir hierdie spesie is. Dit is bevestig dat die bestaande chemikalieë van die beste SA geur, wat bestaan uit oktenol vrygelaat teen c. 9,1 mg/h, 4-metielfenol vrygelaat teen c.15,5 mg/h en asetoon vrygelaat teen c. 350 mg/h, die hoof aanloklikkingskomponente vir G. brevipalpis is. 'n Gomval ("sticky trap"), naamlik 'n tweekleurige blou/swart XT, was aangepas om in tsetse verspreidingsopnames te gebruik. Die nuutontwikkelde H-val was bewys om doeltreffend te wees om relatiewe hoë getalle van beide tsetse spesies te kan versamel. Hierdie val was gebruik om lewende vlieë vir vang-merk-vrylaat studies te versamel om die populasie grootte en beweging van 'n tsetse populasie te bepaal. Laasgenoemde studies het getoon dat teikendigthede van omtrent 4 en 7 teikens/km² vir G. brevipalpis en G. austeni ondeskeidelik voldoende behoort te wees om hierdie spesies suksesvol met geurlokaas en insekmiddel-geïmpregneerde teikens te beheer. G. austeni is beperk tot digte skaduryke areas maar kan kort afstande, tot 345 m, van ongunstige habitat, tussen plate van digte plantegroei oorbrug. G. brevipalpis was beskou as 'n baie meer mobiele vlieg en het wye areas van 1,345 m oorbrug. Albei spesies word geredelik aangelok na beeste, maar nie na boerbokke of bosvarke nie. Hulle het ook meer geredelik op bees gevoed. Beide spesies kan ook in die nag voed. Dit word aanbeveel dat beeste wat met 'n insekmiddel behandel is as mobiele teikens gebruik word om beide G. brevipalpis en G. austeni te beheer, in areas waar beeste die oorheersende gasheer is.

Tsetse verspreidingopnames in die noordoostelike KwaZulu-Natal het gewys dat G. brevipalpis in twee hoofverspreidings-bande voorkom. Die belangrikste bron van hierdie spesie skyn natuur-reservate asook ander natuurlike gebiede te wees. G. austeni is meer wydverspreid met 'n aaneenlopende noord tot suid verspreiding. 'n



Geografiese Inligtingstelsel was gebruik om tsetse verspreiding en oënskynlike digthede te karteer. Trypanosomose-gevalle en -voorkoms, diptenkverspreiding, grondgebruik, gronbedekking en plantegroeitipes is ook gekarteer en met tsetse verspreiding en digthede vergelyk. Laastens, was 'n strategie vir die monitering en onderhoubare geïntegreerde beheer en uiteindelike totale uitwissing van beide G. brevipalpis en G. austeni in die hele N.O. KwaZulu-Natal voorgestel. Dit behels die onderverdeling van die gebied in vyf bestuurbare zones met opeenvolgende presuppressie, suppressie and uitroei operasies wat in elk van die zones volg. Met hierdie voorgestelde strategie kan beide tsetse spesies binne 8 - 12 jaar uitgeroei word.



ACKNOWLEDGEMENTS

I am very grateful to Errol Nevill, my supervisor at ARC-Onderstepoort Veterinary Institute (ARC-OVI), for valuable discussions and encouragement throughout the years. I also greatly appreciate input from my promoters: Clarke Scholtz, University of Pretoria, for his advice and support, and Brian Williams, World Health Organization, Switzerland, for his continued support, guidance and encouragement.

During the several years of field work at Hellsgate Tsetse Research Station, this work would not have been possible without the help of one or more of the following persons: Pieter Trollip, S'Busiso Ngwane, Petros Gazu, Jerome Ntshangase, Johan Esterhuizen, Daniël de Klerk, Gert Venter and Justin Greger. I also thank the Commanding Officers of Hellsgate Military Base for permitting research to be conducted on their premises and the Regimental Sergeant Majors for continued support and provision of infrastructure.

Stephen Torr, Natural Resources Institute (NRI), United Kingdom, was invaluable in teaching me odour sampling techniques in the field and also measuring carbon dioxide of the experimental cow with an infrared gas analyzer. David Hall and his team of chemists at NRI who performed the chemical analysis of odour samples are also acknowledged. I also thank Stephen for his comments and interpretation of the results of odour studies.

I am grateful to John Hargrove, Regional Tsetse and Trypanosomosis Control Programme (RTTCP), Harare, Zimbabwe, for helping with the analysis of the population dispersal data and the invaluable advice, suggestions and comments he made. I particularly also appreciate the input, criticism and guidance from Glyn Vale, RTTCP, whom I regard as my mentor in the field of tsetse.

I thank Tim Robinson, for consulting on the undertaking of tsetse distribution surveys and providing means of analysis of tsetse distribution data and the DAVID database. I am especially indebted to personnel of the Entomology Division, ARC-OVI, who conducted several of the surveys: Errol Nevill and Daniël de Klerk, who were also assisted by Lia van der Walt, Gert Venter, Hilda Nevill and Fred Potgieter. A special



thanks to Hluhluwe and Jozini State Veterinary Animal Health Technicians for assistance and conducting many of the surveys in communal farming areas. Thanks also to KwaZulu-Natal Wildlife, SAFCOL, MONDI, commercial farmers and communal farmers to allow us to survey their premises. I thank Hilda Nevill and Lia van der Walt for entering survey data into the DAVID database. I am especially indebted to Alta Stenson and Roy Williams for producing distribution maps with Arc-View GIS.

A word of thanks to the owners who lend the animals to use in various experiments: Ralph Dobeyn for the cow (Oublom), Petrus Gazu for the young calf (Snoet) and three goats and Rob Bagnall for obtaining the bushpigs (Pumba and Varkie). A special word of thanks for the patience of these animals when spending many hours in the ventilated pit during odour experiments or when tethered to a pole in the bush.

Lastly, I thank my husband, Bernard, and some close friends and family for regularly drawing my attention away from tsetse flies.

Funding received by the ARC-OVI and National Department of Agriculture, Directorate of Veterinary Services is gratefully acknowledged.



CONTENTS

Summary .		
	ıgiii	
Acknowled	lgements v	
Contents	vii	
List of Tab	ix	
List of Figu	ures xii	
List of acro	onyms and abbreviations xv	
CHAPTER	R 1:INTRODUCTION	
	1.1 Literature review 1	
	1.2 Justification	
	1.3 Problem and hypothesis 12	
	1.4 Objectives	
	1.5 Expected benefits arising from this study	
CHAPTER	R 2:GENERAL MATERIALS AND METHODS	
	2.1 Study Area	
	2.2 General techniques and equipment 16	Y .
	2.3 Experimental design and analysis	
CHAPTER	R 3:STUDIES TO FIND AN ATTRACTIVE ODOUR BAIT	
	3.1 Abstract	
	3.2 Introduction	
	3.3 Materials and Methods	
	3.4 Experiments and Results	
	3.5 Discussion 55	
CHAPTER	R 4:DEVELOPMENT OF SUITABLE TRAPS	
	4.1 Abstract	
	4.2 Introduction	
	4.3 Materials and Methods	
	4.4 Experiments and results72	
	4.5 Discussion	
СНАРТЕ	R 5:POPULATION DISPERSAL AND MOVEMENT	
	5.1 Abstract	
	5.2 Introduction	



	5.3 Materials and Methods	97
	5.4 Results	104
	5.5 Discussion	120
CHAPTE	ER 6: FEEDING RESPONSES	
	6.1 Abstract	125
	6.2 Introduction	125
	6.3 Materials and Methods	127
	6.4 Experiments and Results	129
	6.5 Discussion	135
CHAPTE	ER 7:TSETSE DISTRIBUTION AND ABUNDANCE	
	7.1 Abstract	139
	7.2 Introduction	139
	7.3 Materials and Methods	143
	7.4 Results	148
	7.5 Discussion	170
CHAPTE	ER 8: DISCUSSION AND STRATEGY FORMULATION	ON
	8.1 Tsetse monitoring and control options	186
	8.2 Research needs addressed	188
	8.3 Strategy formulation	189
CHAPTE	ER 9: CONCLUSIONS, CONSEQUENCES AND FUT	URE PRIORITIES
	9.1 Conclusions	204
	9.2 Consequences	206
	9.3 Future priorities	207
REFERE	ENCES	208
APPEND	OIX I	220



LIST OF TABLES

Chapter 3
3.1 Attractiveness of targets baited with various release rates of CO ₂ and the
best SA blend
3.2 Indices of catches of targets baited with CO2, phenols and acetone relative
to the control treatment
3.3 Indices of catches of targets baited with CO2, the SA blend and a combination
of CO ₂ added to the blends relative to the control treatment
3.4 Summary of the results to evaluate the importance of natural cow odour vs.
visual stimuli
3.5 Summary of CO ₂ measurements taken during October 1997
3.6 Mean catches of targets baited with natural cow odour and with CO2 released
at the same rate as produced by the cow
3.7 Summary of the results showing the importance of the remaining odour
components other than octenol, 4-methylphenol, acetone and carbon
dioxide
3.8a Rates of production of acetone and butanone from cow and synthetic source
(AOP)
3.8b Analyses of volatiles collected on Porapak (ratio relative to 4-methylphenol
= 100)
3.8c Rates of production of carboxylic acids in cattle odour
3.8d Estimates of the mean rates of production of various tsetse attractants as
obtained for the second run of chemical absorbtion (1998) and analyses 47
3.9 List of synthetic cow (SC) components used to simulate the natural cow
and the recommended release rates. The sachet sizes, which gave more
or less the correct dosages are indicated
3.10 Indices of mean catches of flies attracted to natural and synthetic cow (SC)
odour relative to the control
3.11 Indices of mean catches attracted to cow, bushpig and goat odours relative
to 'no odour'
3.12 Indices of mean catches obtained with odours released from cow, man and
a combination of cow and man relative to the control
3.13 Summary of indices of the attractiveness of natural cow odour vs. 'no odour'
for five experiments (A-E)



Chapter 4	
4.1 Comparisons of various shapes and colours of sticky traps in four	
experiments	14
4.2 Comparisons of e.blue/black 3-dimensional XTs with 2-dimensional	
Monopanels of various sizes	6
4.3 Behavioural responses of a) G. brevipalpis and b) G. austeni in and around the	
H3, H4 and H5 trap modifications as determined with electric nets	19
4.4 Final comparisons of the H4 and H5 modifications with the B4, B5 and Nzi	
traps 8	2
Chapter 5	
5.1 Indices of increase of the recommended target relative to the H trap	14
5.2 Summary of details on the number of flies released and recaptured at the	
various trap sites - 13 January to 5 March 1999 10	16
5.3 Summary of estimates on population density and expected target densities	
needed for various options of killing percentages	6
5.4 Summary of mark-release-recapture results for Blocks B, C, D and E to	
investigate the use of open areas as natural barriers to the movement of	
G. brevipalpis and G. austeni - 3 September to 17 December 1998 11	8
Chapter 6	
6.1 Relative attraction of G. brevipalpis and G. austeni males and females to cow,	
bushpig and goats (in two experiments within sand forest)	1
6.2 Feeding percentages of G. brevipalpis and G. austeni males and females on	
cow, bushpig and goats (in two experiments within sand forest)	1
6.3 Relative attraction of G. brevipalpis and G. austeni males and females at	
various times of day inside sand forest (Site 1) and in the adjacent open	
grassland area (Site 2)	4
6.4 Feeding percentages of G. brevipalpis and G. austeni males and females at	
various times of day inside sand forest (Site 1) and in the adjacent open	
grassland area (Site 2)	4
Chapter 7	
7.1 Summary of survey units surveyed in natural and commercial areas	:0



7.25	Summary of survey units (diptank localities) surveyed in communal farming	
	areas	
Cha	apter 8	
8.1	Detailed information of Zones I - V listed according to natural and commercial	
	areas and communal areas. The species present and the approximate size of	
	each zone is given	
8.2	Technical work plan of project phases to be applied in each of the zones (I - V)	
	as projected during indicated timeframe (1 - 8)	
Арр	pendix 1	
A.1	Details of survey sampling site coordinates and trap catches	



LIST OF FIGURES

Chapter 2	
2.1 Visual (1 x 1 m phthalogen blue) and non-visual (0,5 x 1 m net) electric grids	
incorporated to form a flanked target (i.e. p.blue/net)	7
Chapter 3	
3.1 Schematic representation of cow in underground ventilated pit 2	6
3.2 Extractor fan outlet placed c . 50 cm downwind of a flanked p/blue electric	
target2	7
3.3a Odour extraction and sampling setup. Extracted air from the pit (housing an	
animal) was sampled via tubing with air pumps (shown right). Filters	
were inserted through the sampling ports in the ventilation shaft of the	
pit (center). Carbon dioxide was measured similarly by means of an	
infrared gas analyzer (shown left)	9
3.3b Collections of carboxylic acids were made by sampling through filters	
containing Chromasorb P AW filters (left), volatiles (i.e. phenols and octenol)
were collected on Porapak® filters (centre) and ketones and aldehydes	
(carbonyl compounds) were trapped with silica SepPak® cartridges (right)3	0
3.4 Front view of ramp of pit. Setup shows 1,5 x 1 m electric net at far side of	
pit where ventilation shaft exits. Note fibre-glass sheet in roof of pit allow-	
ing light into pit	7
3.5 CO ₂ release rates of cow during the mornings and afternoons as determined	
by means of the regression between measured concentration (ppm minus	
background) against nominal CO ₂ rates (1/min) obtained in Table 3.5 4	0
Chapter 4	
4.1 Sticky traps for G. austeni: (a) Rectangular sticky screen; (b) 3-DT; (c) cross-	
shaped target (XT)6	5
4.2 Diagrammatic representations of the prototype H trap (a) with its H1,	
H2 and H3 modifications (b-d) and details of collecting device (e)	0
4.3 Photograph of the final H trap design for the capture of G. brevipalpis and	
G. austeni (the trap is held upright by fastening the corners to four rigid	
metal poles (1,2 m long) and the cones are suspended from two flexible	
steel rods (1,4 m long)	34



4.4 Diagrammatic representation of the final H trap with details of materials
and measurements for trap construction
Chapter 5
5.1 Copy of airphoto of Ndlozi peninsula, Lake St. Lucia, showing the vegetation of
the Hellsgate study area. The positions of various Blocks (A-E) used in mark-
release-recapture trials are shown
5.2a Yellow artists' oilpaint was used to colour-code flies on positions of thorax
as also indicated in Fig 5.2 b
5.2b Positions on thorax used for marking (e.g. for position 18, positions 10 + 8
are marked)
5.3a Summary of the dispersal rates for G. brevipalpis males
5.3b Summary of the dispersal rates for G. brevipalpis females
5.3c Summary of the dispersal rates for G. austeni males
5.3d Summary of the dispersal rates for G. austeni females
5.4 Daily recapture rate at various days after release for G. brevipalpis a) males
and b) females114
5.5 Daily recapture rate at various days after release for G. austeni a) males
and b) females
Chapter 6
6.1 Cow in the centre of an incomplete ring of electric nets (8 m diam.) covering
35 % of the circumference of the ring
Chapter 7
7.1 Historical distribution of the tsetse flies Glossina pallidipes, G. austeni and G.
brevipalpis (after Du Toit 1954)
7.2 Reference map to indicate localities of magisterial districts, major game reserves
and conservation areas, lakes and major rivers
7.3 Distribution of Glossina brevipalpis and G. austeni expressed as positive
and negative trap catches
7.4 Apparent density of Glossina brevipalpis expressed as the number of
flies/trap/day



7.5 Apparent density of Glossina austeni expressed as the number of	
flies/trap/day	1
7.6 Diptank positions in magisterial districts of Ingwavuma, Ubombo, Hlabisa,	
Nongoma and Mhlabathini indicating the distribution of cattle of	
communal farmers (diptank areas positive for tsetse during surveys are	
numbered)	3
7.7 Approximate distribution of cattle affected by trypanosomosis during 1990-	
1992 in N.E. KwaZulu-Natal	5
7.8 Prevalence of trypanosomosis in N.E. KwaZulu-Natal as determined by BCT	
and Ag-ELISA	6
7.9 Landcover map	8
7.10 Vegetation type map	9
Chapter 8	
8.1 Summary of research needs addressed during this study necessary to develop	
a strategy for the monitoring and control of Glossina brevipalpis and G.	
austeni in N.E. KwaZulu-Natal, linked to tsetse monitoring and control	
options	5
8.2 Distribution of G. brevipalpis and G. austeni according to positive trap catches	
of the distribution survey. Zones $I-V$ are indicated as part of a strategy to	
eradicate the two tsetse species from N.E. KwaZulu-Natal. Positions of	
temporary target barriers are indicated. The remaining boundaries of zones	
are natural barriers of fly-free areas	7



LIST OF ACRONYMS AND ABBREVIATIONS

3DT 3-Dimensional trap

Ag-ELISA Antigen – Enzyme Linked Immuno-Sorbent Assay

ANOVA Analysis of variance

AOP Acetone, octenol and phenols mixture

ARC-ISCW Agricultural Research Council-Institute for Soil, Climate and Water

ARC-ITSC Agricultural Research Council-Institute for Tropical and Subtropical

Crops

ARC-OVI Agricultural Research Council-Onderstepoort Veterinary Institute

AVHRR Advanced Very High Resolution Radiometer

BCT Buffy Coat Technique

CCD Cold Cloud Duration

CSIR Centre/Council for Scientific and Industrial Research

DAVID Disease and Vector Integrated Database

FAO Food and Agriculture Organization

GIS Geographic Information System

GPS Global Positioning System

IAEA International Atomic Energy Agency

IGR Insect Growth Regulator

IPAR Intercepted Photosynthetically Active Radiation

KZN KwaZulu-Natal

KZNNCS KwaZulu-Natal Nature Conservation Services

LIT Lethal Insect Technique

LP Legpanel

MP Monopanel

NDVI Normalized Difference Vegetation Index

NOAA National Oceanic and Atmospheric Administration

NRI Natural Resources Institute

PAAT Programme Against African Trypanosomiasis

PATTEC Pan African Tsetse and Trypanosomiasis Eradication Campaign

RT Rectangular sticky screen

RTTCP Regional Tsetse and Trypanosomiasis Control Programme

SAFCOL South African Forestry Company Limited

SANDF South African National Defense Force



SAT	Sequential Aerosol Technique
SC	Synthetic Cow
SIT	Sterile Insect Technique
XLP	Cross-shaped legpanel
XT	Cross-shaped target (sticky trap)