

Dedication

To:

My fatherAbusara,

My motherNayla with her love and continuous

Support remained for me as unlimited

Source of encouragement.

My husbandAwadalla

For his understanding

during this investigation.

My daughter Hatton and ***My son***Ahamed

For their love

ACKNOWLEDGEMENTS

I thank God Almighty for giving me the health, strength and courage to accomplish this work.

My thanks are due to German Embassy in Khartoum, through German Academic Exchange Service DAAD for giving me in country scholarship opportunity. Also my thanks to Elneelain University for giving me study leave to study my PhD.

My deep appreciation and gratitude to my supervision Prof. Mohammed KhairAbdalla who really encouraged me and made me go deeper and learn more about this field of science, either from direct discussion or thought attend workshop which given by him.

My gratitude and appreciation is extended to my co-supervision Dr. SihamRahamatalla for her guidance, fruitful advice, help and suggestions. Also for giving me opportunity to be one of student group for Prof. Dr. Brockman for six months.

My profound respect and gratitude goes to Prof. Dr. Brockman for his scientific guidance and attend some lectures .My gratitude also extends to Prof. Dr. Reissmann and Mrs. D. Schemmel who have particularly very keen and have provided unreserved assistance throughout the part of the molecular biology period.

Sincere thanks and indebtedness are due to Dr. Ammar Said Ahamed for his generous stand by, suggestions, advice and un limiting help during and after my stood in Berlin.

I am most indebted to Prof. Amir Mohammed Salih faculty of Animal production , University of Khartoum ,who kindly supported me when this research initiated.

I am indebted to all staff member of animal production epically Dr. YaggobMagbol and Dr. HafzAbdrahman, department of agricultural technology and fish science, Elneelan University, for their excellent cooperation during my study.

My deep sincere, grateful thanks, gratitude, appreciation and honour to my family , epically my mother, husband , sisters ,brother , daughter , son and my lovely friend SariaAlmarzogi, for their generous support and patience during the whole period of my study including my absent for six month.

Table of contents

	Page
Dedication	i
ACKNOWLEDGEMENTS	ii
Table of contents	iv
Abstract	vii
Arabic Abstract	ix
Chapter 1 General Introduction	1
Chapter 2 Overview of dairy production system and genetic conservation for Cattle Breeds in tropics	8
2.1 Introduction.....	9
2.2 Dairy production systems in the tropics.....	12
2.2.1 Pastoral systems.....	12
2.2.2 Semi- intensive systems.....	14
2.2.3 Intensive systems.....	14
2.3 Conservation of Local cattle Breeds.....	16
2.3.1 Reasons for conservation of local cattle breeds.....	17
2.3.1.1 Economic-biological reasons.....	17
2.3.1.2 Scientific reasons.....	18
2.3.1.3 Cultural-historical reasons.....	19

2.3.2 Steps for conservation.....	19
2.3.2.1 Inventory.....	19
2.3.2.2 Evaluation.....	20
2.3.2.3 Choice.....	20
2.3.2.4 Preservation.....	20
2.3.2.4.1 Conservation in situ (In situ conservation).....	21
2.3.2.4.2 Ex-situ Conservation.....	22
2.4. Improvement of indigenous breeds in tropical areas by selection..	23
2.4.1 Selection procedures in developed countries.....	26
2.4.1.1 Testing stations.....	26
2.4. 2.2 Progeny testing schemes.....	26
2. 4.2.3 Nucleus or group breeding schemes.....	28
2.5 Molecular techniques.....	30
2.5.1 Marker assisted selection (MAS).....	30
2.5. 2 Positive and negative selectable marker.....	31
2.5.3 There are many type of markers.....	31
2.5.4 Genotyping methods and MAS.....	32
2.5.5 Steps for MAS.....	32
2.5. 6 Problems with MAS/QTL.....	32

2.7 Reference

Chapter 3	<i>DGAT1</i> gene in dairy cattle –Review.....	38
	Abstract, Introduction, Discovery of <i>DGAT1</i> gene , Roles of <i>DGAT1</i> in Triacylglycerol Metabolism, <i>DGAT1</i> Enzymes and Triglyceride Synthesis, <i>DGAT1</i> and milk fat association, The effect of <i>DGAT1</i> gene polymorphisms on milk production, conclusion , Reference.	
Chapter 4	Smallholder dairy production system of Kenana and Butana cattle in Sudan.....	57
	Abstract, Introduction, Material and methods, Results and Discussion, Conclusions, Reference.	
Chapter 5	Variants of diacylglycerol acyltransferase1 (DGAT1) gene in Sudanese dairy cattle (Kenana and Butana).....	73
	Abstract, Introduction, Material and methods, Results and Discussion, Conclusions, Reference.	
Chapter 6	General Discussion	90
Chapter 7	Summary and conclusions.....	101
Appendix	106

ABSTRACT OF THESIS

The objectives of the current study were the characterization of some Sudanese dairy cattle types mainly Kenana and Butana (on farm and molecular characterization) and their dairy production systems, adopted management practices, breeding objectives and constraints of dairy development in Sudan. A structured questionnaire was administered to 101 Kenana and Butana owners randomly selected from the homelands of both breeds. Personal interviews and repeated field visits were conducted. The average herd size of Kenana and Butana cattle was 10 and 6 animals, respectively. Kenana and Butana farmers adopted different management systems. The traditional nomadic system was more prevalent in Kenana area (98%), while all Butana owners used a transhumant system (100%). Kenana and Butana cattle are kept in a mixed crop- livestock production system and livestock species kept by farmers comprise cattle, sheep and goats. Cattle are the dominant species, mainly used for draught power followed by milk production. The purpose of keeping cattle in the study area was to generate income from the sale of milk which was the main source of income for herders in the Kenana area (100%) and up to 50% of Butana owners. However, surplus milk is sold at farm gate to middlemen at low prices, and animals are sold live in the village or nearest markets. The general appearance and body conformation were the criteria for selection of

breeding bulls used by both Kenana and Butana owners (72% and 80.4%, respectively). Disease prevalence was important in both production systems and almost all farmers in both areas reported incidences of diseases. Trypanosomosis were the main problems reported by Kenana herders (61.8%) while Butanan owners complained mainly of ticks. Veterinary services in the country at large have declined in recent years and in some areas have witnessed a degree of collapse. The two breeds were also characterised by genotyping the *diacylglycerol acyltransferase1* (*DGAT1*) gene. In both breeds, the *DGAT1* Lysine variant (232K) that is associated with high fat and protein content as well as high fat yield in other breeds was the most frequent allele. The frequencies of the 232K allele were 96.3% and 84.6% in Kenana and Butana breeds, respectively. At the *DGAT1* promoter VNTR locus, four alleles containing four to seven repeats of the 18 bp motif were found in both breeds. The most frequent allele was the VNTR allele 3 containing five repeats with a frequency of 60.4 % and 57.5 % in Kenana and Butana breeds, respectively. In conclusion, the two examined Sudanese dairy cattle breeds do not differ in allele frequencies at the *DGAT1* locus.

ملخص البحث

الهدف من الدراسة الحالية ، التوصيف (داخل الحقل والمعمل _التوصيف الجزيئي) لبعض أنواع الابقار الحلوب السودانية (كنانة وبطانة) ، بالإضافة لمعرفة أنظمة إنتاج الالبان والممارسات الادارية التي تم اتباعها ، و أهداف التربية والمعوقات لنظم انتاج الالبان في السودان . تم عمل (101) استبيان منظم لاصحاب ماشية (كنانة وبطانة) تم إختيارهم عشوائياً من مناطقهم الاصلية . اوضحت الدراسة ان متوسط حجم القطيع من ابقار (كنانة وبطانة) 6-10 حيوان في المتوسط على التوالي . كما ان نظام التربية المتبع مختلف في كل من المنطقتين (كنانة وبطانة) حيث يتم اتباع النظام الرعوي التقليدي (98%) في كنانة في حين يتم ممارسة النظام شبه الحديث في أبقار بطانة بنسبة (100%) كما يتم اتباع نظام الانتاج الحيواني المختلط في كل من المنطقتين في تربية ورعاية الابقار والاعنام والماعز. الابقار هي اكثر انواع الماشية السائدة وتستخدم أساساً لإنتاج الالبان ، والغرض الاساسي من حفظ الابقار في منطقة الدراسة هو العائد المادي من بيع اللبن الذي هو المصدر الاساسي للدخل عند مربي الابقار في (كنانة وبطانة) 100% الى 50% على التوالي بالإضافة الى ان الفائض من اللبن يتم بيعه للوسطاء بأسعار منخفضة كما تباع الحيوانات الحية في القرية أو الاسواق القريبة . يتم إختيار ثيران التربية التي يتم إستخدامها من قبل المربين (كنانة وبطانة) على أساس المظهر الخارجي العام والشكل بنسبة 72% - 80.4% على التوالي . انتشار الامراض هو احد المشاكل الهامة المؤثرة على نظام الانتاج للمزارعين في (كنانة وبطانة) . حيث تمثل التريانسوما المرض الرئيسي المؤثر على الانتاج عند مربي كنانة (61.8%) بينما يعتبر القراد هو المؤثر الرئيسي للإنتاج عند مربي بطانة . وعموماً تراجعت الخدمات البيطرية في السنوات الاخيرة وفي بعض المناطق شهدت درجة من الانهيار . كما تم النوصيف الجزيئي للجين DGAT1 للسلاطين. اوضحت الدراسة أن الطفرة (232K لايسين ذات علاقة بزيادة الدهن ومحتوى البروتين وكذلك انتاجية الدهون ، وهي تتكرر بنسبة 96.3% - 84.6% في كل من ابقار (كنانة وبطانة) على التوالي. أما الطفرة الثانية (VNTR) الموجودة في الجزء المحفز من الجين وعند التحليل المعلمي تم العثور على أربعة أليلات تحتوي على 4-7 تكرارات ابتداءً من 18 زوج قاعدي موجودة في كل من السلاطين (كنانة وبطانة) والأليل الاكثر تكراراً هو أليل 3 الذي يحتوي على 5 تكرارات فهو موجود بنسبة 60.4% - 57.5% في (كنانة وبطانة) على التوالي. وختاماً نجد ان سلالات الابقار

الطوب السودانية (كنانة وبطانة) لا تختلف عن بعضها فى تكرارات الأليل بالنسبة للجين

. DGAT1