Society for Applied Bacteriology Symposium Series No. 2

ACTINOMYCETALE

Characteristics and Practical Importance

edited by G. Sykes and F. A. Skinner

THE SOCIETY FOR APPLIED BACTERIOLOGY SYMPOSIUM SERIES NO. 2

ACTINOMYCETALES: CHARACTERISTICS AND PRACTICAL IMPORTANCE

Edited by

G. SYKES

AND

F. A. SKINNER



ACADEMIC PRESS INC. (LONDON) LTD 24-28 OVAL ROAD LONDON N.W.1

U.S. Edition published by
ACADEMIC PRESS INC.
111 FIFTH AVENUE
NEW YORK, NEW YORK 10003

Copyright © 1973 By the Society for Applied Bacteriology

ALL RIGHTS RESERVED

NO PART OF THIS BOOK MAY BE REPRODUCED IN ANY FORM BY PHOTOSTAT, MICROFILM, OR ANY OTHER MEANS, WITHOUT WRITTEN PERMISSION FROM THE PUBLISHERS

Library of Congress Catalog Card Number: LCCCN 72-9777 ISBN: 0-12-679950-4

Printed in Great Britain by The Whitefriars Press Ltd., London and Tonbridge, England

Contributors

- NINA S. AGRE, Laboratory for Physiology of Actinomycetes, Institute of Microbiology, USSR Academy of Sciences, Moscow, U.S.S.R.
- R. W. ATTWELL, Postgraduate School of Studies in Biological Sciences, University of Bradford, Bradford BD7 1DP, England
- H. BEERENS, Institut Pasteur, 20, Boulevard Louis XIV, 59000 Lille, France
- W. BLYTH, Department of Botany, University of Edinburgh, Edinburgh EH9 3JH Scotland
- G. H. BOWDEN, The London Hospital Medical College. Dental School, Turner Street, London, England
- R. M. BRADSHAW, Hartley Botanical Laboratories, University of Liverpool, Liverpool, England
- CATHERINE L. BULLEN, Public Health Laboratory, Lewsey Road, Luton, England
- N. P. BURMAN, Metropolitan Water Board, Rosebery Avenue, London EC1R 4TP, England
- M. CATTEAU, C.E.R.T.I.A., 369, Rue Jules Guesde, Flers Bourg, 59650 Villeneuve d'Ascq, France
- T. CROSS, Postgraduate School of Studies in Biological Sciences. University of Bradford, Bradford BD7 1DP, England
- M. GOODFELLOW, Department of Microbiology Medical School, University of Newcastle-upon-Tyne, Newcastle-upon-Tyne NE1 7RU, England
- D. GOTTLIEB, Department of Plant Pathology, University of Illinois, Urbana, Illinois 61801, U.S.A.
- J. M. HARDIE, The London Hospital Medical College, Dental School, Turner Street, London, England
- D. A. HOPWOOD, John Innes Institute, Colney Lane, Norwich NOR 70F, England
- L. V. KALAKOUTSKII, Laboratory for Physiology of Actinomycetes, Institute of Microbiology, USSR Academy of Sciences, Moscow, U.S.S.R.
- J. LACEY, Rothamsted Experimental Station, Harpenden, Herts, England
- D. H. LAPWOOD, Rothamsted Experimental Station, Harpenden, Herts, England
- J. LOSFELD, U.E.R. Mathématiques, Université des Sciences, 59000 Lille, France
- F. PONCELET, C.E.R.T.I.A., 369, Rue Jules Guesde, Flers Bourg, 59650 Villeneuve d'Ascq, France
- L. M. POUZHARITSKAJA, Laboratory for Physiology of Actinomycetes Institute of Microbiology, USSR Academy of Sciences, Moscow, U.S.S.R.
- G. P. SHARPLES, Hartley Botanical Laboratories, University of Liverpool, Liverpool, England

- W. B. TURNER, I.C.I. Ltd., Pharmaceuticals Division, Alderley Park, Macclesfield, Cheshire, England
- KATHLEEN WILLIAMS, Public Health Laboratory, Lewsey Road, Luton, England
- S. T. WILLIAMS, Hartley Botanical Laboratories, University of Liverpool, Liverpool, England
- A. T. WILLIS, Public Health Laboratory, Lewsey Road, Luton, England

Preface

THIS second volume in the Symposium series of the Society for Applied Bacteriology publications is concerned with a review of the characteristics and significance of the Actinomycetales. The Symposium was held at the Loughborough University of Technology, England in July 1972. Each chapter has been contributed by a specialist in his field and so the book as a whole constitutes a unique collection of information on this group of micro-organisms which has long been recognized as having many unusual and interesting features.

As might be expected of such a diverse group of organisms there is much to be learned of their physiological characteristics and several chapters are devoted to such studies. Under this heading there are valuable and comprehensive contributions on the taxonomy and genetics of the group and on the production and germination of their spores. In other chapters the occurrence and significance of the actinomycetes in water, soil and the human digestive tract, and their role in causing potato scab disease is discussed.

It is relevant to stress the medical importance of the actinomycetes both in causing and curing disease. The realization that some streptomycetes could produce antibacterial substances stimulated great interest in the group, especially in the years immediately following the Second World War, and led to the development of the present impressive array of antibiotics of clinical value. The pathogenic properties of the anaerobic actinomycetes have been known for a long time but more recently attention has been directed to the activities of the aerobic types as allergens in pulmonary conditions such as Farmer's Lung disease.

G. SYKES
3 Grosvenor Court
Egerton Road
Weybridge
Surrey, England

F. A. SKINNER Rothamsted Experimental Station Harpenden Hertfordshire England

Contents

LIST	OF CONTRIBU	TORS										
PRE	FACE .			~					1			vi
Gene	eral Consideration	and Imp	olicat	ions of	the A	Actino	mycet	tales				
D	. GOTTLEIB											
	Introduction											1
	Habitat .											2
	Soil competition										,	3
	Antibiotics			× 100				٠.				4
	Species pathoge											5
	Plant diseases											6
	Animal nutritio											6
	Food preservati	on										7
	Co-operative tax	konomy						.,				7
	References											9
Taxo	nomy and Classif	fication o	f the	Actino	myce	etes						
T	. CROSS and M. (GOODFE	LLO	W								
	Introduction								, .			11
	Family Actinom											13
	Family Actinop	lanaceae										30
	Family Dermato						. 3	4				38
	Family Frankiac	ceae			· ,					2.1		40
	Family Micromo											41
	Family Mycobac	cteriaceae										43
	Family Nocardia	aceae										57
	Family Strepton	nycetacea	le			. ,						64
	Family Thermoa	actinomy	cetac	eae					1.0			76
	Family Thermor	monospoi	acea	e								78
	Genera in search											88
	References								٠.			91
The I	Fine Structure of	the Actin	omy	cetales								
	T. WILLIAMS, G					. BRA	DSHA	W				
	Introduction										1	13
	Methods for exam											13
	(a) Ultrathin											
	(b) Whole cel											18

X CONTENTS

	(c)	Carbon repl	icas of	whol	e cells							114
	(d)	Carbon repl	icas of	freez	e etch	ed cell	S					114
	(e)	Negative sta	ining	of disr	upted	cells						114
	(f)	Scanning ele	ectron	micro	scopy							114
		al cell conten										114
	Mesos	omes										115
	Cross	walls										117
	Modif	ications to ba	asic ce	ll stru	cture						;	119
	Flagel	la .										120
	Forma	tion of spore	es								٠.	121
	(a)	Fragmentati	ion of	a shea	thless	hypha	(grou	ip 1a)				123
	(b)	Fragmentati	ion of	a shea	thed l	hypha	(group	1b)				123
	(c)	Endogenous	spore	form	ation	(group	2)					125
	Ackno	wledgements	S								2	126
	Refere	ences										127
Gene	tics of	the Actinomy	ycetale	es								
D	А. Но	PWOOD										
	Introd	uction										131
	An ou	tline of the	conj	ugatio	n syst	em in	Strep	tomy	ces co	elicolo	or	
	A3((2)						1				132
	(a)	The fertility	types									132
	(b)	Zygote geno	mes									135
	(c)	The conjuga	tion p	rocess	3							136
	(d)	Conclusions										138
	Other	conjugationa	l syste	ms							٠.	138
	(a)	Other strept	omyce	etes .						. 4		138
	(b)	Nocardia										140
	(c)	Micromonos	spora									141
	Transf	ormation sys Thermoactin	tems									141
	(a)	Thermoactin	nomyc	es vul	garis						٠.	141
	(b)	Streptomyc	etes				. 1					142
	Actino	mycete gene	tics ar	nd ind	ustrial	proble	ems					143
	(a)	Mutagenesis										143
	(b)	The predicti Ultrafertility	ve valu	ue of 1	the lin	kage n	nap					145
	(c)	Ultrafertility	y									146
	(d)	Interspecific	and i	nter-st	train r	ecomb	inatio	n				147
	(e)	The desirab	ility o	f harn	essing	transc	luctio	n or tr				
		in strain imp	roven	nent								149
	Actino	mycete gene	tics an	d the	study	of mo	rphog	enesis				
	Ackno	wledgements	S									150
		nces										150

	Streptomyc								l Beha	viour		
L	. V. KALAI	KOUTSK	II and	L. M	. Pot	ZHAI	RITSK	AJA				
	Introducti											155
	Spore viab											156
	Resistance	e to delete	erious	agent	ts							157
	Spore stru	cture and	d comp	positi	on							158
	(a) Str	ucture mposition									979	158
	(b) Co	mposition	n									160
	(c) Me	tabolic ac	ctivity									164
	Germinati											165
	(a) Eff	ect of en	vironn	nenta	l facto	ors				,.		165
		rmination		ts								168
	Conclusion	n										170
	Reference	S										171
Endo	spores of A	ctinomy	cetes:	Dorn	nancy	and C	Germin	ation				
	V. KALAK											
	Introducti	on										179
	The spore											181
	Germinati	on stages										182
	(a) Eff	ects of lo	w tem	perat	ures	on acti	ivation	1				182
	(b) Oth	ner activa	tion to	reatm	ents							184
	(c) Wh	at change	s occi	ır dur	ing ac	ctivatio	on?					186
	(d) Ger	rmination	1						1.			189
	(e) Het	terogenei	ty of s	pore	popul	lations	3					190
	Effect of	repeated	transf	er of	cultu	res on	the a	bility	of the	ir spo	res	
	tog	germinate										191
	Concludin	g remarks	S									192
	References											
Gern	ination of	Actinomy	cete S	Spore	S							
R	W. ATTWI	ELL and	T. CR	OSS								
	Introducti	on										197
	Actinomy	cetes stud	lied									198
	Spore p	roduction	n									198
	Spore p Appearance	e in micr	ocultu	ire								199
	Changes in											
		uence of										200
	(b) Age											200
			ent									201
	(d) Effe	t treatme	tibiot	ics								202
	Discussion											

xii CONTENTS

Summary							205
Acknowledgements							206
References							
Secondary Metabolism with Special Refer	ence to	Acti	nomy	cetale	2		
W. B. TURNER	chec te	Acti	noniy	cctares	3		
Introduction							209
Relationship of primary and second				•			210
Biosynthetic origins of the secondar	w met	holit	13111			•	210
(a) Incorporation of the intact g					,		
(b) Compounds related to the nu			ton				211
(c) The shikimic acid pathway		ucs					
(d) The polyketide and polyprop							01/
							010
(f) Compounds derived from am	ino ac	ide.					216
Conclusions							216
References							217
References							41 /
The Occurrence and Significance of Acting	omy ce	tes					
N. P. BURMAN							
Introduction							219
Methods							219
Occurrence and significance						٠.	221
(a) In river water							221
(b) In stored water							
(c) In slow sand filters .							223
(d) In biological sludge blankets				4.			
(d) In biological sludge blankets(e) In chlorinated water							
(f) In the distribution system	•	•					227
References							
						•	22)
Actinomycetes in Soils, Composts and Foo	dders						
J. LACEY							
Introduction							231
Methods of assessment and their into							
Soils							233
(a) Numbers and types of actino	mycete	es					000
(b) State of actinomycetes in soil	l						235
(c) Factors influencing actinomy							235
Manure and composts .							237
(a) Animal manners: mushroom	compo	sts					
(b) Vegetable composts .							240
(c) Town wastes			14				240
(0) 10 1111 1140000				-			0

CONTENTS													
Fodders													
											240		
	(b) Cereal grains										242		
	(c) Straw										242		
	Other materials										242		
	, ,										242		
	(b) Miscellaneou										243		
	The role of actinon	nycete	S .						· 10.		243		
	(a) Interactions	with o	other	organ	isms						244		
	(b) Antibiotic p												
	(c) Degradation	of org	ganic	matte	r						245		
	Conclusion										246		
	References												
Ctuan	tomy ces scabies and	Dotat	0 500	h Dicc	2000								
	H. LAPWOOD	rotat	o sca	U DISC	ease								
D.											252		
	The disease				*								
	(a) General		*										
	(b) Symptoms										253		
	The pathogen										254		
	(a) Identity										254		
	(b) Isolation and	d cultu	ire								255		
	(a) Identity (b) Isolation and Infection and disea	se deve	elopm	ent							255		
	Factors affecting th	ne path	nogen	and d	lisease	incid	lence		• "		256		
	Control measures										257		
	(a) Seedborne i	noculu	ım								258		
	(b) Soilborne in	ocului	m								258		
	(c) Prevention (of infe	ction								258		
	References								9				
Farm	er's Lung Disease												
W	BLYTH												
	Introduction	,					8 4				261		
	Incidence										267		
	Organisms known t						eir occu	irrence	e		263		
	Inhalation, retention										263		
	Sputum culture	4114	J		. 0. 10	Pilu	ory unit	- 20113			265		
	Common types of	resnira	tory l	wner	enciti	vity.					265		
	Clinical features	сърпа	cory I	ry pers	0113111	vity					266		
		ontino							•				
	Histopathological f	eature	2								267		

xiv CONTENTS

	Serological	evidence									268
	(a) Mou	ldy hay an	d therm	ophili	ic actir	nomyc	etes				268
	(b) Funs	ldy hay an gal antigen	S								
	Relationship	ns hetweer	serolog	v and	natho	ogenesi	2				
	Treatment a										273
	References		1						- 7		273
	References										213
Comr	nensal and P	athogenic	Actinor	nvces	Specie	e in M	an				
	H. BOWDE				Specie	22 111 141	an				
	Introductio										277
	Early studie										277
	Actinomyce										279
	Actinomyce	es as natho	gens	Sairisi	113 111 1	11411					281
	(a) In m	an .	gens								281
		erimental i								٠	
											283
	Some chara				_						284
		ohology									285
	(b) Colir	nial . ılar .									286
		iological									
	Serology of		-						7		290
	References				*		1.		*		295
	pid Method	for the le	dentifica	ation	of Bifi	idobacı	teriun	2 Spec	ies Us	ing	
	naracters	D D		Y D							
Μ.	CATTEAU		ELET, I	H. BEI	EREN	S and J	. LOS	SFELI)		
	Introductio										301
	Matériel et										301
	(a) Soud										301
	(b) Prép	aration de	s culture	es	,						302
		emensemer									303
	(d) Exp	loitation d	es résult	tats							303
	Résultats										303
	Discussion						· ·				309
	Références										309
	References									1	507
	Significance										
C	ATHERINE	L. BULLI	EN, A. 7	T. WIL	LIS at	nd KA	THLE	EN W	ILLIA	MS	
	Introduction	on .	11.						1	7	311
	The compo	sition of b	reast m	ilk an	d cows	' milk					313
	(a) Imp	ortant cor	stituent	s .							313

CONTENTS XV

	Some properties of	breast	milk	and co	ows' r	nilk					314
	(a) Coagulability	/									314
	(b) Buffering car	pacity	of br	east n	nilk ar	nd cow	s' milk				314
	Growth of Escheric	hia co	oli and	d anae	robic	Lactol	oacilli	in bre	ast mi	lk	
	and cows' milk										314
	Growth of anaerobi	c lacto	bacil	li and	E. co.	li in lac	ctose b	roth			316
	Continuous culture	of a	naero	bic la	ctoba	cilli ar	E.	coli ii	n brea	st	
	milk and cows' n	nilk									316
	Decrease of buffering										
	Growth in breast mi	lk and	dcow	s' milk	prep	aration	1S				319
											319
	Anaerobic lactobac	illi in	the	vagina	e of	pregna	nt wo	men	and th	1e	
	mouths of newbo	orn inf	ants								320
											321
	Acknowledgements									į.	324
	References										324
Techr	iques for the Isola	tion a	and C	harac	teriza	tion of	f Acti	nomy	ces an	d	
(b) Buffering capacity of breast milk and cows' milk Growth of Escherichia coli and anaerobic Lactobacilli in breast milk and cows' milk Growth of anaerobic lactobacilli and E. coli in lactose broth Continuous culture of anaerobic lactobacilli and E. coli in breast milk and cows' milk Decrease of buffering capacity of cows' milk Growth in breast milk and cows' milk preparations The origin of anaerobic lactobacilli in the infant intestine Anaerobic lactobacilli in the vaginae of pregnant women and the mouths of newborn infants Discussion Acknowledgements					327						
	•										
Techr	iques for the Isol	ation	and	Ident	ificat	ion of	Aer	hic	Actino)-	
											335
ing co	ares. Dummary Of	tile 1	Discus	331011	DC2210	PAR.					222

General Consideration and Implications of the Actinomycetales

D. GOTTLIEB

Department of Plant Pathology, University of Illinois, Urbana, Illinois 61801, U.S.A.

CONTENTS

1.	Introduction				v.													1
2.	Habitat																	2
3.	Soil competition				Ţ		1.								ū.		1000	3
4.	Antibiotics .						**							*				4
5.	Species pathogenic	to	ma	n							ž						1	5
6.	Plant diseases						25											6
	Animal nutrition																	
8.	Food preservation							¥					٠				×	7
9.	Co-operative taxon	on	ıy						*			**						. 7
10	References							9		4	5		(4)					

1. Introduction

THE ACTINOMYCETES have always been a strange group of organisms to the bacterial taxonomists. The number of investigators studying this group of microbes has always been small and for the most part these organisms have been neglected by medical bacteriologists, physiologists and biochemists. Such affairs were to be expected when one recalls that until recently not even their taxonomic position was certain (Breed, Murray & Smith, 1957). Bacteriologists considered them as bacteria and mycologists generally considered them as fungi. The diseases caused by them have been described in books on medical mycology (e.g. Conant et al., 1954). This era is now over and the actinomycetes are generally accepted as bacteria. They have no nuclear membrane, are sensitive to lysozyme for the most part and to the common antibacterial agents; there is a similarity to the type of bacterial flagella when these organelles are present and the types of cell wall resemble those of bacteria. The hyphal diameters are much smaller than those of fungi and are close to those of the bacteria.

Genera in actinomycetes consist of varied groups of bacteria whose common feature is the formation of hyphae at some stage of development. This tendency is tenuous and often requires imagination to believe in it. In some species of the genus *Mycobacterium* hyphae are never seen, whereas other species have incipient hyphae only in young cultures. An even worse situation exists in the various genera of the Actinomycetaceae; these form microcolonies which have very transitory filaments that are extremely difficult to see and the dominant forms are diphtheroid, bifid or even rod shaped. Hyphal development is more

pronounced in the nocardias but they, too, fragment. The time at which this occurs varies with the species and it determines the types of colony which form. Colonies in which the filaments break up early tend to be soft or mucoid whereas those whose filaments break up after a few days have time for a branched mycelium to form and develop a leathery texture: they later fragment to bacillary or coccoid elements. In the Streptomycetaceae, filament development is strong and fragmentation is rarely found. Examples are in the Streptomyces, Micromonospora and Actinoplanes spp.

Reproduction is usually asexual, though sexual processes have been shown to occur by genetic analyses. In the nonhyphal forms, asexual reproduction is by fragmentation or perhaps even by the usual fission of single cells. Where stable hyphae are produced, vegetative reproduction is by well formed spores resembling fungal arthrospores, borne either free or in sporangia, as in the Actinoplanaceae. The free spores are in the form of sporophores and may consist of 1, 2 or many spored chains arising singly or verticillate from primary hyphae. The long chains are straight, looped or spiral. Except for the Actinomycetaceae, which contain aerobic or micro-aerophilic genera, the actinomycetes are generally aerobic.

2. Habitat

Members of the actinomycetes which live a saprophytic existence can be found with greater or less frequency in most ecological settings, soil having the greatest population density. Numerically they are less dominant than the other bacteria and more prominent than the fungi. The numbers vary a great deal but it is not uncommon in fertile soils from many different parts of the world to find -5×10^6 propagules/g of soil (Taber, 1960; Gottlieb, unpublished). They occur in greatest numbers in the top few inches of the soil and decrease with depth (Waksman & Purvis, 1932). The sensitivity of Streptomyces spp. to acids at pH 5.0 or less precludes high populations in such soils (Taber, 1960). Similarly, the writer has not found them in soils from the Arctic tundra of northern Sweden or in soils of the Great Salt Desert of the U.S.A. High densities of spores can also be found on potato or sweet potato that have been infected by pathogenic Streptomyces spp. Streptomyces spp. are also common on most plant parts and, being dustborne, can be isolated from almost all artifacts and natural materials (Grein & Meyer, 1958). They are found in marine littoral and associated with Micromonospora spp., especially in fresh water lakes and in bottom muds. By their nature thermophilic forms, are found commonly in composts of manures. Other types are favoured by particular oxygen environments; Actinomyces israeli, for example, is anaerobic, whilst a number of other species are micro-aerophilic.

The streptomycetes have 3 important roles in the soil. One is their function in decomposing the organic matter of the soil (Waksman & Lomanitz, 1925;

Reynolds, 1954; Williams, 1966). Another is their effect on soil structure in binding clay particles by the hyphal threads to impart a granular viable structure that is conducive to crop production. The actinomycetes are undoubtedly responsible in part for the earthy odour of soil (Gerber & Lechevalier, 1965).

3. Soil Competition

The third interesting role of actinomycetes in soil is their ability to produce antibiotics which are inimical to other soil microbes. This is on the assumption that antibiotics are produced in normal untreated soils in the same way as they are produced in special laboratory media. This results in an intriguing hypothesis to explain some of the competitive effects in soil, resulting in the increase of some species and loss of other species among the microbial components of the soil. Competition certainly occurs between organisms in soil, but it is difficult to demonstrate that the actinomycetes produce in untreated soils antibiotics at levels high enough to inhibit other soil organisms. There are data to show that actinomycetes, especially members of the genus Streptomyces, can produce antibiotics in some sterilized soils, with or without added metabolites, and even in nonsterile soils, but the question remains: which of these conditions represents the natural state of the soil? Two main considerations are (1) natural soils do not have the necessary metabolites in concentrations necessary to stimulate and allow antibiotic production and (2) the necessary metabolites are normal in soil and are part of the natural life cycles in soils. An added difficulty is that of assessing the 'clinical' level of antibiotic concentration in soil.

A few examples clarify the situation. The addition of S. griseus and Bacillus subtilis to normal or treated sterile soil resulted in a decrease in the population of the Bacillus but there was no detectable amount of streptomycin. Moreover if streptomycin had been produced it would have been adsorbed by the negatively charged clay component of the soil to render it inactive (Siminoff & Gottlieb, 1951). On the other hand S. venezuelae produces chloromycetin, a neutral molecule. This actinomycete produced the antibiotic in trace amounts in sterile, but not in nonsterile soil, and the amounts were increased by adding green alfalfa hay or tryptone to the soil (Gottlieb & Siminoff, 1952). The antibiotic was not found in field plots that had been inoculated with the streptomycete nor in any of the 91 soils that were collected in field surveys, including areas from which the organism had originally been isolated (Ehrlich, Anderson, Coffey & Gottlieb, 1952, 1953). Actidione, another neutral antibiotic produced by S. griseus gave similar results (Gottlieb, Siminoff & Martin, 1952). The results of studies with S. rimosus and S. aureofaciens, which produce the amphoteric compounds terramycin and aureomycin, respectively, are different (Martin & Gottlieb, 1952). The actinomycetes inhibited B. polymyxa in soil even though relatively high concentrations of the antibiotics did not. In this case the

inactivation or removal from soil of the antibiotics was presumably due to the pH value of the soil.

There appears to be no correlation between the ability of an organism to inhibit the growth of other organisms in soil and its ability to produce an antibiotic that inhibits them. As a part of one of our studies, 15 actinomycetes were isolated that produced an antibiotic which inhibited *Rhizoctonia solani* and 15 others that did not. The actinomycetes were separately added to sterile soil 2 weeks before, at the same time as, and 2 weeks after, the *Rhizoctonia*. After 1 month in pots the relative populations of *Rhizoctonia* in the soils were determined by the numbers of radish and pea seedlings that did not emerge or were damped off, and there were no significant differences between groups. Some isolates that produced antibiotics antagonized the fungus, whereas others did not. Similarly some that did not produce antibiotics inhibited the test fungus whereas others did not.

4. Antibiotics

It is because of their ability to produce antibiotics that the actinomycetes are perhaps best known. Except for a few that are produced by fungi, e.g. penicillin and cephalosporin, and a few that are produced by bacteria, e.g. bacitracin and polymyxin, all other antibiotics that are medically useful and have a wide application are synthesized by actinomycetes. A compilation of the microbial sources of antibiotics discovered in the U.S.A. and Japan between 1953 and 1970 reveal that c. 85% are produced by actinomycetes, 11% by fungi and 4.5% by bacteria. These differences may not reflect the actual proclivities of the different groups to produce antibiotics for similar ratios might be obtained if more effort was spent on the last groups. Yet the decision, if it were so made, to spend most time on the actinomycetes is in my experience the result of the fact that the actinomycetes have shown the greatest potential for producing new and useful antibiotics. The exact number of antibiotics that are known is difficult to even estimate. About 2700 substances have been reported but I would estimate that c. 50% of them are so poorly characterized that their uniqueness is suspect. The most commonly reported new antibiotics are those with antibacterial activity, the proportion in Japan being 48% and in the U.S.A. 56%; next are antifungal agents, in Japan, 32% and in the U.S.A. 24%. Antitumour antiviral and antiprotozoal antibiotics comprise the remainder. These figures probably reflect not so much the innate ability of microbes to produce such agents but the amount of research effort that has been placed in the search for antibiotics of different therapeutic abilities.

Another means of determining the importance of antibiotic production is by the amount and value of the product that is produced. With these data one can get into ethical difficulties such as the value of saving one life against the value