VITAMINS AND NITROGEN REQUIREMENTS OF **ARTHROBACTER SPECIES**

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ABSTRACT

Fifty-two strains of Arthrobacter, picked from a collection of 170 strains isolated from glycine enrichments inoculated with soil, sewage and activated sludge were found to be exacting to vitamins or aminoacids. According to their requirements they could be divided into five sub-groups. The I subgroup demanded thiamine alone, the II thiamine and pantothenate, the III thiamine, pyridoxine and glutamic acid, the IV thiamine, pantothenate and glutamic acid or sulphur containing amino acids as nitrogen source. The V subgroup comprised those which demanded organic form of nitrogen, glutamic acid serving the purpose most effectively. Biotin was found stimulatory but not essential for the II and III Subgroups. Some of the strains in subgroup II exhibited an interesting cycle of morphogenesis in media with and without biotin. A role for biotin in the normal life cycle of Arthrobacter has been postulated.

INTRODUCTION

According to Conn and Dimmick (1947), the soil Arthrobacter represent those which can utilize inorganic nitrogen in the absence of vitamins but this observation has not been substantiated by other investigators. Taylor (1938), for example, reported that only 17 out of 106 soil Arthrobacter were able to grow on nitrate nitrogen in the absence of yeast extract. Jensen (1952) revealed that only certain strains of soil coryneform bacteria were able to utilize inorganic nitrogen compounds while others demanded organic form of nitrogen. Morris (1960), and Chan and Stevenson (1962), also found that a good growth of A.globiformis can be obtained with inorganic nitrogen only when biotin was incorporated in the medium and this has subsequently been confirmed by Veldkamp et al. (1963). Mulder (1963) pointed out that almost all of the soil Arthrobacter strains he studied were able to utilize inorganic nitrogen though some demanded biotin for the purpose and others depended on a vitamin mixture or even on a vitamin mixture and casamino acids for growth to occur.

In general, it may be stated that the vitamin requirements of Arthrobacter have not adequately been studied. Of the few available reports, that of Lochhead and Thexton (1952) and Lochhead and Burton (1955) reveal that some of the species required thiamine alone or thiamine and biotin in addition 142

to vitamin B_{12} . Subsequently, Lochhead (1958) described two new species of *Arthrobacter* demanding respectively vitamin B_{12} and terregens factor and Chan (1964) reported on the biotin requirement of *A.globiformis*.

In the present paper are described details of the experiments carried out on the nutritional requirements of 52 Arthrobacter strains picked at random from fresh isolates made from various samples of soil, sewage and activated sludge. It would be clear from the results that organisms of this genus can conveniently be grouped into 5 distinct nutritional types, thus lending support to the conclusion of Knight and Proom (1950) and Dias and Bhat (1963, 1964) that nutritional survey could be a valuable aid in the identification of the sporeforming and diphtheroid bacteria.

MATERIALS AND METHODS

Organisms: Arthrobacter cultures employed in this study were isolated from glycine enrichments inoculated with soils, sewage and activated sludge by the method described previously by Mullakhanbhai and Bhat (1966 b). The pure cultures were maintained on nutrient agar and glycine agar slants. Their characterization was achieved by following the techniques described in 'Manual of Microbiological Methods' Conn et al. (1957). Morphological and Gram staining properties were observed at intervals of 8, 18, 24 and 36 hours of growth on n. agar slants and basal salt liquid medium containing appropriate carbon and nitrogen source plus the growth factors required. Strains mentioned herein, as belonging to Group I, II and IV, were derived from soils whereas those belonging to group III were those exclusively derived from sewage and activated sludge. The group V strains again were from soil and sewage.

Media for nutritional studies: The techniques used in the elucidation of the nutritional requirements were essentially those described by Dias and Bhat (1964) except that the basal mineral base employed was that of Khambata et al. (1960). All tests were done in test tubes $(1.8' \times 15 \text{ cm.})$ each containing 10 ml of the medium. Inoculum in each case was made from a 24 hour old culture (growing on n. agar slant) suspended in normal saline centrifuged at 6000 r.p.m. and washed twice and resuspended in distilled water. Aliquots of 0.05 ml of a thin suspension were inoculated and the tubes incubated at room temp. $(20 - 26^{\circ}C)$. For elucidating the vitamin nutrition, cultures were serially transferred twice and the growth response was measured on a Bausch and Lomb 'Spectornic 20' colorimeter at 540 m μ .

RESULTS

Nutritional properties of sub-group I: The minimal requirements for the growth of sub-group 1 Arthrobacter strains in the basal salt solution containing 1% glucose and 0.05% ammonium sulphate are shown in Table I. Thiamine

					Arthr	obac	ter st	rains	i i	
Vitamins	s excluded		83	84	85	86	87	88	89	90
from th	e mixture			(1	Gro 00 –		respo		ce)	
None			40	45	50	50	40	40	45	50
Thiamine	******		0	0	0	0	0	0	0	0
Nicotinic acid		******	39	48	48	51	39	37	47	49
Calcium pantothenate	e		41	50	46	50	45	39	45	50
Pyridoxine	ana ++18		38	45	49	50	43	40	49	51
p.aminobenzoic acid		******	40	45	50	52	41	38	46	50
Folic acid			41	50	50	49	40	40	43	49
p.aminobenzoic acid	and Folic	acid	40	50	46	50	39	40	45	49
Biotin			41	48	50	50	40	40	47	50
Riboflavin	*****		40	48	49	51	40	39	45	50
Vitamin B ₁₂			40	50	50	50	40	36	43	49
All	****		0	0	0	0	0	0	0	0

TABLE I

Growth of Arthrobacter (Sub-group I) as affected by different vitamins

TABLE II

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				Ar	throt	bacter	stra	ins	
Vitamins exc	luded		75	76	77	78	79	80	81
from the mix	ture					h re tran	-)
None .			50	40	38	40	50	41	39
Thiamine .			0	0	0	0	0	0	0
Nicotinic acid		******	50	40	40	41	49	40	43
Calcium panthothenate _			0	0	0	0	0	0	0
Pyridoxine .		******	51	42	35	42	47	41	40
p.aminobenzoic acid			50	40	37	41	50	42	40
Folic acid .			50	42	40	40	50	39	39
p.aminobenzoic acid and Fo	olio acid		50	42	39	40	48	40	41
Biotin			30	28	30	32	30	40	40
Riboflavin .	*****		51	40	38	42	48	42	40
Vitamin B ₁₂	• 4 100 4 4 4		51	41	40	40	48	42	40
A11		******	0	0	0	0	0	0	0

Growth of Arthrobacter (Sub-group II) as affected by different vitamins

				A	throl	bacter	stra	ins	
Vitamin	s excluded		49	50	51	52	53	54	55
from th	e mixture					th re trans		se ince)	
None	******	*****	45	55	52	55	48	50	47
Thiamine	*****		0	0	0	0	0	0	0
Nicotinic acid	*****	******	43	57	50	55	47	51	49
Calcium panthothenate	******	******	43	55	50	53	49	50	46
Pyridoxine	*****	*****	0	0	0	0	0	0	• 0
p.aminobenzoic acid		6	45	50	52	55	50	49	46
Folic acid	-	******	45	58	50	52	48	49	49
p.aminobenzoic acid an	d Folic acid	******	42	55	51	51	48	50	46
Biotin	******		45	55	52	53	50	50	45
Riboflavin	*****	*****	46	58	50	55	49	51	48
Vitamin B ₁₂	*****	******	44	55	50	55	47	50	48
All			0	0	0	0	0	0	0

TABLE III

Growth of Arthrobacter (Sub-group III) as affected by different vitamins *

 Basal salt solution containing 0.1% Na-glutamate as nitrogen source and 1% glucose as carbon source.

was indispensable for growth for all the strains. The suitability or otherwise of various compounds as substitutes for the essential vitamin is indicated in Table V. It is clear that the requirement for thiamine could not be met with other substitutes. Sorbitol, known for its thiamine sparing action in animal systems, could not substitute the vitamin in the bacterial system. None of the inorganic and organic nitrogenous compounds tested supported good growth (Table IX). Likewise, glutamate only was observed to be suitable as a combined source of of carbon and nitrogen to all the strains and rest of the compounds, served only to a limited extent for strains 87, 88, 89 and 90. It is interesting to note that glutamate was the only compound that supported growth of the first four strains, viz., 83, 84, 85 and 86 (Table XI) and this provided a means for the demarcation of one from the other group.

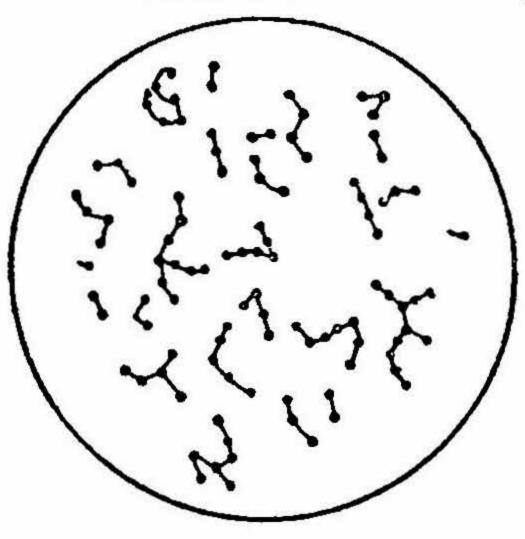
Sub group II: The minimal requirements for growth of this sub-group Arthrobacter strains in basal salt – glucose – ammonium sulphate medium are shown in Table II. All the strains demanded thiamine and pantothenate for growth. Other compounds when tested for their suitability as substitute for the essential vitamins (Table VI) revealed that none could replace either

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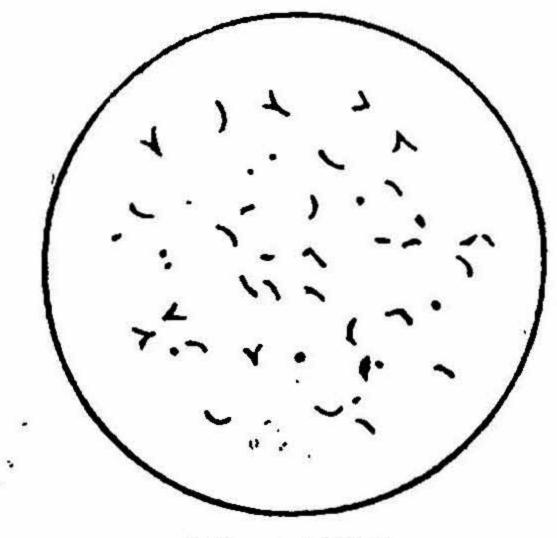
thiamine or pantothenate. Of interest is the finding that β -alanine could not replace pantothenate and that biotin was growth stimulatory but not essential. However, biotin deficient medium promoted the formation of abnormal forms (see Fig. 1) and led to the accumulation of intracellular inclusions. In fact, in the absence of biotin the cell division was interrupted and growth was retarded whereas in the presence of biotin the normal life cycle was completed. The ability of this subgroup strains to utilize various nitrogen compounds is shown in Table X. Ammonium sulphate proved to be a good source of nitrogen. All the amino-acids tested supported growth, asparagine and cysteine proving themselves to be exceptionally good sources. Of the various compounds tested as combined sources of nitrogen and carbon (Table XII), aspartic acid and asparagine supported fairly good growth whereas alanine and glutamate were excellent. Of interest is the observation that sulphur containing amino acids served as satisfactory nitrogen sources but not as sources of combined nitrogen and carbon.

Sub-group III: The minimal requirements for growth of subgroup III Arthrobacter strains in basal salt solution + 1% glucose and 0.1% glutamate are presented in Table III. Pyridoxine and thiamine were indispensable as growth factors to all the strains. In the absence of vitamins even a trace of growth was not observed in first transfer indicating thereby that they had an absolute demand for these growth factors. Table VII represents the growth response of the cultures in media supplemented with various substances. None was found suitable to replace thimaine and pyridoxine. Biotin was found stimulatory but not essential for growth and the demand seemed more



Effect of Biotin on the morphogenesis of Arthrobacter Sp. Sub-group II

4 Days old Cells from medium without Biotin showing a stressed condition with granules, deeply stained



4 Days old Cells from medium containing Biotin showing pleomorphic rods and Cocci (Normal Life Cycle)

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conspicuous when the organisms were cultivated in 05% glutamate containing medium. Glutamate was the only compound that served well as a source of nitrogen, as well as a combined source of carbon and nitrogen, but even in its presence the damand for thiamine and pyridoxine was very sharp and unequivocal suggesting thereby the suitability of the strains for the detection and assay of such factors (Mullakhanbhai and Bhat, 1966 a).

Sub-group IV: The need for growth factors by these Arthrobater strains in basal salt solution +1% glucose and 0.1% glutamate is shown in Table IV. With all strains it was observed that thiamine and pantothenate were indispensable for growth. Like those in the previous subgroups the strains in this subgroup also could not utilize substitutes of the two vitamins (Table VIII). Glutamate and two sulphur containing aminoacids however were suitable as N. sources. Glutamate was also suitable as a combined source of nitrogen and carbon.

Sub-group V: All the strains in this subgroup demanded organic form of nitrogen but could do without any growth factor. Glutamate was the only compound found to be the most suitable substrate for growth, though complex substrates like peptone and casein-hydrolysate also supported their good growth.

Arthrobacter strains									
67	68					73			
	67	67 68	67 68 69	67 68 69 70	67 68 69 70 71	The second se			

TABLE IV

nom the r) – %	tran)
None		 55	48	50	45	52	50	40
Thiamine	*	 0	0	0	0	0	0	0
Nicotinic acid	·······	 55	48	50	47	50	50	42
Calciam pantothenate		 0	0	0	0	0	0	0
Pyridoxine	*****	 53	47	51	47	52	51	40
p aminobenzoic acid		 53	49	50	44	52	51	41
Folic acid	*****	 55	48	50	45	51	50	40
p.aminobenzoic acid and	Folic acid	 55	48	52	47	50	50	42
Biotin		 54	49	50	45	50	52	40
Riboflavin	******	 55	47	51	45	52	50	41
Vitamin B ₁₂		55	47	50	45	50	50	41
All		 0	0	0	0	0	0	C

Basal salt solution containing 0.1% Na-glutamate as nitrogen source and 1% glucose as carbon source.

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TABLE V

Growth response of Arthrobacter (Sub-group I) in various media

				Arth	robac	ter s	strain	S	
Supplements added to mediu	m	83	84	85	86	87	88	89	90
Supplements added to moura						resp			
			(100 -	- % ti	ransn	nittan	ice)	
Vitamin mixture		40	47	50	50	42	40	46	50
Thiamine		35	40	46	46	37	35	40	39
Thiamine + Nicotinic acid	******	37	40	44	47	38	32	40	38
Thiamine + Calcium pantothenate		35	39	45	47	36	34	41	38
Thiamine + pyridoxine	*****	38	40	45	45	37	34	40	39
Thiamine + Folic acid	011 FWE	40	41	43	47	36	36	40	40
Thiamine + p.aminobenzoic acid		40	40	45	45	36	35	38	42
Thiamine + Biotin	*****	39	39	47	45	35	38	38	49
Thiamine + Ribotlavin		35	39	46	47	37	34	37	40
Thiamine + Vitamin B ₁₂		36	40	48	46	37	35	38	41
Thiamine + Casein hydrolysate	*****	42	41	50	46	40	40	42	50
Thiamine + Asparagine		36	38	46	44	38	38	37	41
Thiamine + Glutamate	***-	38	38	48	47	38	38	38	43
Thiamine + Aspartic acid	*****	35	38	48	47	63	37	36	42
Methionine + Sorbitol	******	0	0	0	0	0	0	0	0
Sorbitol + Casein-hydrolysate		0	0	0	0	0	0	0	0
Biotin + Casein-hydrolysate		0	0	0	0	0	0	0	0
Biotin + Sorbitol		0	0	0	0	Û	0	0	0
None	*****	0	0	0	0	0	0	0	0

DISCUSSION

The observation that species in the genus Arthrobacter occur in a wide variety of material and are almost ubiquitous is suggestive of their importance in nature. Though the type species Arthrobacter globiformis is simple in its nutritional requirement other species studied here (as well as by others elsewhere) seem to demand specific growth factors and that this demand is so absolute and specific as to suggest the exploitation of the species for the detection and assay of growth factors present in biological materials. It is also apparent from the above results that the requirements for essential metabolites or growth factors are characteristics for most of the species within a genus eventhough certain strains within a species may behave differently. The morphology of some bacterial cultures grown under laboratory cultural conditions is greatly influenced by the components of the medium in which they are grown. Cultivation of an organism in a medium limiting in one or more components may likewise induce an 'abnormal' morphogenesis. This was clearly observed in some strains of *Arthrobacter* subgroup II where biotin deficiency led to a condition in which cells failed to fragment thereby retarding growth. Bitoin deficiency also caused the accumulation of intracellular inclusions (deeply stained granules) whereas biotin sufficiency restored normalcy in their life cycle. Such deformities in the morphogenesis under deficient conditions have previously been observed by Chan (1964) in *A.globiformis*.

				Ar	throb	acter	strai	ins	
Supplements added	d to med	ium	75	76	77	78	79	80	81
Supplements added				44	Grow				
				(100) – %	trans	smitta	ance)	
Vitamin mixture	*****	******	50	40	35	35	47	45	42
Thiamine + Calcium pantot	henate	******	50	39	35	35	45	45	4(
Thiamine		*** ***	0	0	0	0	0	0	
Calcium pantothenate			0	0	0	0	0	0	ľ
Thiamine + Biotin	*****		0	0	0	0	0	0	ļ
Thiamine + Pyridoxine	,		0	0	0	0	0	0	
Thiamine + Vitamin B_{12}		*****	0	0	0	0	0	0	
Calcium pantothenate + Bio	otin	844 - 1 5 7	0	0	0	0	0	0	
Thiamine + β -alanine	••••		0	0	0	0	0	0	
Thiamine + Asparagine		******	0	0	0	0	0	0	
Thiamine + Glutamate		******	0	0	0	0	0	0	
Thiamine + Casein-hydroly	sate		0	0	0	0	0	ΰ	
Thiamine + Valine + Leucin	e		0	0	0	0	0	0	
Sorbitol + Calcium pantoth	enate	441497	0	0	0	0	0	0	
Thiamine + Sorbitol			0	0	0	0	0	0	
None			0	0	0	0	0	0	. `

TABLE VI

Growth response of Arthrobacter (Sub-group II) in various media

TABLE VII

Growth response of Arthrobacter (Sub-group III) in various media

				AI	throl	bacter	• stra	ins	
Supplements adde	to mediur	n	49	50	51	52	53	54	55
Supplements adde					Grow - %		-)
Vitamin mixture			45	50	52	50	45	50	50
Thiamine			0	0	0	0	0	0	0
Pyridoxine		-	0	0	0	0	0	0	0
Thiamine + Pyridoxine	******		40	46	45	44	40	45	45
Thiamine + Pyriodine + Bi	otin	*****	46	50	50	49	46	48	48
Thiamine + Asparatic acid		6 .	0	0	0	0	0	0	0
Thiamine + y.aminobutyri	c acid	*****	0	0	0	0	0	0	C
Thiamine + Asparagine	******	******	0	0	0	0	0	0	0
Thiamine + Casein-hydrol	ysate	*****	0	0	0	0	0	0	0
Sorbitol + Pyridoxine	*****	******	0	0	0	0	0	0	0
Thiamine + Sorbitol		*****	0	0	0	0	0	0	0
Pyridoxine + Methionine			0	0	0	0	0	0	0
None		*****	0	0	0	0	0	0	0

Taxonomical consideration. The genus Arthrobacter, Conn and Dimmick (1947) was created with a view to segregate the soil "diphtheroids" from the true corynebacteria comprising primarily plant or animal pathogens. The Arthrobacter, in true sense, are saprophytes. They differ from Corynebacterium species in possessing a more complicated life cycle, in the course of which often short branched filamentous rods appear. The latter characteristic renders the boundary between Arthrobacter and Nocardia less distinct. However, in Nocardia branching is more persistent whereas in Arthrobacter true branching does not occur (Skerman, 1959) though occasionally rudimentary budding might be observed. A detailed study of cytology cellular morphology and cell-wall composition of the three genera may reveal some clear-cut demarcations between them and this aspect is under investigation in this laboratory.

In order not to complicate the taxonomy of *Arthrobacter* the authors do not intend to propose new names for the species worked out here. Detailed taxonomic studies have indeed been conducted in this laboratory and a simplified scheme will be proposed in the near future to make the taxonomic position of *Arthrobacter* more clear.

				Ar	throb	acter	strai	ins	
Supplements added	to mediu	m	67	68	69	70	71	72	73
Supplements added				(100			spons mitta		
Vitamin mixture		******	55	48	50	43	50	50	4
Thiamine			0	0	0	0	0	0	
Calcium pantothenate		400 e-1-100	0	0	0	0	0	0	
Thiamine + Calcium panto	thenate		51	44	49	40	48	49	4
Thiamine + Calcium panto	thenate +	Biotin	53	47	49	32	50	48	4
Thiamine + Biotin			0	0	0	0	0	0	
Thiamine + Pyridoxine			0	0	0	0	0	0	
Thiamine + Vitamin B_{12}			0	0	0	0	0	0	
Calcium pantothenate + Bi	otin	*** ***	0	0	0	0	0	0	
Thiamine + β -alanine			0	0	0	0	0	0	
Thiamine + Asparatic acid		******	0	0	0	0	0	0	
Thiamine + Asparagine	*****	******	0	0	0	0	0	0	
Thiamine + Casein-hydroly	sate		0	0	0	0	0	0	
Thiamine + Valine + Leucin	ne		0	0	0	0	0	0	
Thiamine + Sorbitol			0	0	0	0	0	0	
Sorbitol + Calcium pantoth	nenate		0	0	0	0	0	0	
None			0	0	0	0	0	0	

TABLE VIII

Growth response of Arthrobacter (Sub-group IV) in various media

TABLE IX

Growth response of Arthrobacter (Sub-group I) in media containing different nitrogen sources

				2	Arthi	obac	ter s	train	S	
Nitroge	en source		83	84	85	86	87	88	89	90
						wth				1/
				(1	00 -	% tra	insmi	ittand	:e)	
None	b		0	0	0	0	0	0	0	0
Sodium nitrite		B erre	0	0	0	0	0	0	0	0
Sodium nitrate	*****		41	34	15	0	0	0	0	44
Ammonium sulphat	e		35	38	36	39	35	35	37	35
Ammonium nitrate			20	18	22	20	15	15	21	20
Ammonium phosph	ate (Dibasic)		39	30	48	38	20	15	15	42
Aspartic acid			41	35	41	43	23	22	16	34
Asparagine	*****		25	18	32	30	15	16	18	46
Arginine			30	30	41	36	18	18	29	51
Alanine		******	36	25	44	45	28	25	25	57
β-alanine			21	20	24	27	17	50	20	17
Cysteine			42	40	39	41	38	32	30	33
Glycine	6	*****	20	20	26	26	18	15	18	15
Glutamate			41	45	58	59	27	22	30	48
Histidine			45	23	62	56	15	15	15	50
Lysine		****	32	30	41	35	[†] 0	0	0	37
Leucine	50000 ¹⁰		30	28	40	46	18	67	20	30
Methionine		******	22	20	24	28	18	25	18	20
Proline	******	*****	0	15	15	18	0	0	0	29
Phenylalanine		*****	28	20	35	29	15	15	15	20
Serine			25	21	30	27	0	0	0	0
Threonine		***	26	20	27	29	37	50	30	20
Valine		*****	26	25	35	34	0	0	0	17
Urea	*****		0	0	0	0	0	0	0	0
Urate			0	0	0	0	0	0	0	0
Creatinine		*****	0	0	0	0	0	0	0	0
Peptone			31	30	35	27	38	31	30	60
Casein-hydrolysate	Barrent		24	30	35	30	22	20	20	56

TABLE X

Growth response of Arthrobacter (Sub-group II) in media containing different nitrogen sources

		Growth response $(100 - \% \text{ transmittand})$							
Nitrogen	Source		15	South State	- 12 - C			80	81
			0	0					
None	*****				N75 336			0	0
Sodium nitrite	P1 1111	6+++++		2064	9.71	2.84	22 17 - 19 - 19 - 19 - 19 - 19 - 19 - 19 -	0	0
Sodium nitrate								15	19
Ammonium sulphate								49	51
Ammonium nitrate	Ditesta)	*****	555689970. - 43		-FAUTE	172223334	2011/03/03/03	25	27
Ammonium phosphate (Dibasic				0.000		576 - 676	36	35
Aspartic acid		*****				10.000		20	21
Asparagine	*****				2020223			44	47
Arginine		******						15	15
Alanine								18	20
β-alanine			30	26				30	28
Cysteine	******		50	15	40	42	47	44	40
Glycine	0 11114		18	15	16	15	18	20	20
Glutamate			26	33	25	20	25	25	25
Histidine	*****	60 9444	0	50	20	15	0	0	0
Lysine		pp a nort	0	38	15	0	0	0	0
Leucine			15	51	20	26	15	15	15
Methionine	******		36	15	20	23	30	29	27
Proline	*****		0	0	16	15	0	0	0
Phenylalanine	******		26	15	0	0	20	18	15
Serine	*****		0	25	0	0	0	0	0
Threonine			34	21	40	43	30	32	30
Valine			15	0	0	0	15	15	15
Urea			0	υ	0	0	0	0	0
Urate			0	0	0	0	0	0	0
Creatinine			0 0	0	0	0	0	0	0
Peptone	*****	B ***#	52	64	36	30	50	52	54
Casein-hydrolysate	*****		44	60	30	20	-45	45	46

TABLE XI

Suitability of compounds to serve as combined sources of carbon and nitrogen for Arthrobacter (Sub-group I)

					Arthr	obac	ter s	train	S	
2.1	Substrates		83	84	85	86	87	88	89	9(
÷				(1			re s pe ansm		ce)	
None			0	0	0	0	0	0	0	 0
Aspartic acid			0	0	0	0	65	43	50	50
Asparagine		*****	0	0	0	0	22	31	25	68
Arginine		•••••	0	0	0	0	15	24	20	80
Alanine			0	0	0	0	36	26	32	41
β-alanine			0	0	0	0	15	0	15	39
Cysteine			0	0	0	0	0	0	0	0
Glycine			0	0	0	0	0	0	0	0
Glutamate			60	60	64	62	66	59	60	75
Histidine			0	0	0	0	58	62	55	70
Lysine			0	0	0	0	26	27	24	78
Leucine		*****	0	0	0	0	35	32	30	0
Methionine			0	0	0	0	0	0	0	0
Proline			0	0	0	0	35	55	30	0
Phenylalanine			0	0	0	0	0	0	0	0
Serine			0	0	0	0	24	21	20	0
Threonine		******	0	0	0	0	0	0	0	0
Tryptophane			0	0	0	0	0	0	0	0
Valine			0	0	0	0	0	0	0	0
Urea			0	0	0	0	0	0	0	0
Urate			0	0	0	0	0	0	0	0
Creatinine			0	0	0	0	0	0	0	0
Hippurate			0	0	0	0	0	0	0	0
Peptone			56	45	46	58	61	43	60	64
Casein-hydrolys	sate	a a contra	45	45	48	60	21	40	40	61

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TABLE XII

Suitability of compounds to serve as combined sources of carbon and nitrogen for Arthrobacter (Sub-group II)

		Arthrobacter strains								
	Substrates		75	76	77	78	79	80	81	
				Growth response (100 – % transmittance)						
None			0	0	0	0	0	0	0	
Aspartic acid			40	50	56	46	42	38	44	
Asparagine	*****		55	72	33	17	50	46	49	
Arginine	\$=1.4**		20	88	20	0	18	20	20	
Alanine			72	62	23	16	76	70	73	
β-alanine		******	0	56	0	0	0	0	0	
Cysteine	6		0	0	0	0	0	0	0	
Glycine	e o stadad	••	0	0	0	0	0	0	(
Glutamate			80	82	32	15	78	75	8	
Histidine			0	62	54	42	0	0		
Lysine		*****	0	75	28	16	0	0	1	
Leucine			0	0	24	26	0	0		
Methionine		*****	0	0	0	0	0	0	1	
Proline	6		26	32	49	46	27	25	2	
Phenylalanine			0	0	0	0	0	0		
Serine			0	0	27	0	0	0	3	
Threonine		******	0	0	0	0	0	0		
Tryptophane	******		0	0	0	0	0	0		
Valine			0	0	0	0	0	0		
Urea			0	0	0	0	0	0		
Urate		******	0	0	0	0	0	0		
Creatinine	******		0	0	0	0	0	0		
Hippurate	*****		0	0	0	0	0	0		
Peptone			77	75	44	37	70	69	6	
Casein-hydrolysa	te		68	75	29	32	66	65	5	

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