# Compilation of tRNA sequences and sequences of tRNA genes

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## **INTRODUCTION**

The 1995 compilation contains 2700 sequences of tRNAs and tRNA genes. The last edition which appeared two years ago (1) was supplemented by 694 new sequences. The sequences of tRNA mutants and of tRNAs originating from transformed or differentiated cells were not considered.

The tRNAs included in the compilation are listed in Table 1. Each tRNA or tRNA gene is specified by the (abbreviated) name of the organism from which it was isolated and a four digit code: the first three digits identify the organism, the last digit specifies the particular isoacceptor. The amino acid specificity of the tRNA is indicated by a one-letter amino acid code. The tRNAs coding for selenocysteine were annotated with the letter Z. Initiator tRNAs are annotated with the letter X.

The sequences, references and footnotes of tRNAs and tRNA genes listed in Table 1 are deposited in the European Bioinformatics Institute (EBI) Data Library. The references are restricted to the first complete publication of the sequence unless additional information (e.g. base modification, corrections, etc.) was later obtained. In such cases additional references were added.

In order to facilitate a computer analysis an alignment is used which is most compatible with the tRNA phylogeny and known three-dimensional structures of tRNA. The corresponding numbering system is shown in Figure 1.

As was the case in the previous edition (1), this publication does not contain a sequence printout. Instead, the sequences have been deposited in the EBI Data Library. This publication should be therefore quoted as a reference for data obtained from the electronically accessible database. Information on how to access the sequence files can be obtained by electronic mail: send email to Netserv@ebi.ac.uk containing the command 'HELP TRNA.' The help file will contain all the information needed to obtain the requested sequence. The tRNA database is also available via anonymous FTP from ftp.ebi.ac.uk in the directory pub/databases/tRNA. It is also distributed on the EBI CD-ROM. Contact the EBI Data Library, Hinxton Hall, Hinxton, Cambridge CB10 1RQ, UK (Fax: +44 1223 494468, email: DataLib@EBI.AC.UK).

Researchers who wish to obtain the sequence information on a floppy disk, or as a hardcopy should contact M. Sprinzl, Laboratorium für Biochemie, Universität Bayreuth, D-95440 Bayreuth, Germany, Fax: +49 921 552432, email: Ma-

thias. Sprinzl@uni-bayreuth.de. Software allowing search for tRNA sequences according to several criteria, e.g. source, partial sequence, modified nucleoside, anticodon, amino acid specificity and printout of sequences in a form of tables or cloverleaves, is also available.

#### **Presentation of sequences**

The sequences in the database are divided into three parts. The first two parts contain the sequences of the tRNA genes and tRNAs, respectively, which can be fitted into the canonical tRNA alignment. The third part contains tRNA and tRNA gene sequences, mainly of animal mitochondria, whose secondary structures differ from most tRNAs and could not be aligned according to Figure 1.

An example for sequence presentation in the database is given in Table 2. Each sequence in the compilation occupies two consecutive lines. The first line begins with the letter 'D' or 'R' and contains the six-position identification code of the sequence ('D' or 'R' for DNA or RNA, respectively; a one-letter code for the amino acid, X for methionine-initiator, Z for selenocysteine; and the four-digit code specifying the organism and isoacceptor. After this, the sequence of the anticodon (in the case of tRNA sequences in its modified form) is given, followed by the name and the kingdom of organism (Table 1), and the sequence (99 standard positions). The second line begins with the sign '+' and contains the information about base-pairing (double helical regions only, tertiary interactions are not annotated). All other lines in the compilation begin with signs other than 'D,' 'R' or '+' (usually '\*') and contain comments.

Nucleotides involved in Watson-Crick pairs are marked with '=', the G·U pairs are indicated with the sign '\*'. Nucleotides 26 and 44 are considered to form a base pair included in the anticodon stem (Fig. 1).

The sequences in original publications denoted as 'yeast' are assigned to *Saccharomyces cerevisiae*. The user should be aware, however, that some of these organisms have possibly been misclassified and that the original literature should be consulted.

This compilation uses a one-letter code for all nucleotides including modified ones. For standard nucleotides, adenosine, cytidine, guanosine, thymidine and uridine the usual abbreviations, A, C, G, T and U, respectively, are used. To designate

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Table 1. List of tRNA sequences and sequences of tRNA genes included in the compilation

PART ONE: Sequences of	LPRA gene	h
Source	Code	ITNA genes
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PHAGE TA PHAGE TO	C23	GUIGRST ACCCPGHICLMAPGGGTVWXY
ANCHAEBACTERIA	600-10	9
ANCIUS GLOSUS FULG.	G34	A
HALOBACTERUM OUT.	938	AG .
HALOBACTERI JM WATL	045	ie W
HALOFERAX VOLCANII METHANOBAC FORMIL	050	CW
METHUNORAC THERM.	062	A .
METHANOTHRY SCEN	000	ADEPHEADPOPERVY A
METHANOTHERM REA PLIMINOSACTER AMYLO	057 046 070	ADDERMANDET
METHANGCOC VOLTAR METHANGETHUS KAND.	074	DKPTV
METHANOSPIR, HUNG,	C78	NLOS A
THERMORASMA ACID.	036	rólivx M
THERMOCOCCUS CILER	694	APT CNA
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EURACTENA	110-23	B
MYTOPLASMA CAPRIC. MYTOPLASMA MYCOID.	118	ACDITCHIK (LLMNPORKS) TTVWWXY ACCICAMAPPRISTVX
MYTORLASMA PNEU. MYTORLASMA PODD	120	ACDEBOG-WICCLIAN/PORFRSSSSTTTVWWXYY
achorbiplasava laid.	123	IA AACDEFG-BIKKLILIMMAGRESTVW
SPIPOPLASMA MELF.	120	OWW ACCRIMPISK
NORMELIA BLIRGDORF. MATONELIA BACIL	128	AI K
STREET DUTY CES GRES.	130	î
STREPTORIYEES FOIL	131	DORK
STREPTORYCES AMBO.	136	OOESSOCIANYOSPOWY
YYOOBACT, TUBERS	139	90
OLEUSHILLA, ASROOM.	148	
CLOSTRIDIUM THERM	142	
SEELE FORMER DACK	144	
LOSTRIBUM ACETO.	105	
NTIROCOCCUS HIFAI STAPHYLOCOC AURS.		CTOGGGHKILLIMPORSSTTVVWXY
ACTOBAC BULG. ACTOBAC DELBRUEC.	150 0	FONFRSY
ACTOCOCCUR LACTIS	153 S	ABPOINSX
ACILLUS SUSTILIS ACILLUS CIRCULANS ACILLUS SP. PS3		AAACDEFFGGGHEIKKULLULMWNWPQRSSSTTTVWXXY
ACILUS SP. PS3	157 D	DVSV
HERMUS THERMOPHS. HODOTHERMUS MAR.	160 A	GTTY
HIDBACILLUS FERRO	162 A	i ACDEFGGGHIKULLUMNPPPOOMHRHISSESTTTTTVVVWXXXYV
ALMONEULA TYPHI.	170 H	LPRR
FECHODESMILM SPEC	173 A	
HOTOBACT, PHOSPH, HOTOBAC, LEIDONA.	174 H	
EROMONAS HYDROPH.	178 A	EHLPN
SEUDOMONAS AER SEUDOMONAS FLUOR.	184 A	Fig. (1)
AMPYLOGAC JEJUNI AULOBACTER CRES.	195 A	
HISOBIUM MELILOTI ZOANCUS SP.BH72	194 L 195 L	
OFFICE PROTEINS.	198 L	
NACYSTIS MIDULANG	210 A	
YMECHOCYSTIS SP. YANOPHORA PARAD.		ACC PORLS
TANELLA LITTORA.	222 A 224 A	
MGANELLES		
HOROPLASTS	440.000	
	240-358	
YANDPHORA FARAD. 'LAIELLA LITTORA.	240 Al	
LAMYDOMONAS REIN LAMYDOMO, MORWU,		COECHITW
CORRESCON ESCU.	245 A	RS
JOUMIS SATIVUS	249 DI 250 E	
RTARIA LONGA KREMA GRACILIS	251 At	CODERGINIOLLANPORRESTYWXXXY
TYPTOMONAS SPEC.	204 At	R
ITTHAMMON SP.	257 AL	
ANIONIM CALDAR. JETHODISCUS LUT.	268 AJ 268 AJ	
ARCHANTIA POLYM. IBOUTA REFLEXA	260 A	DEFGGH KULLIMNPPORREGESTTVVWXY
RESOCHAETE ORDIC.	262 Al	ILMY
PIDGUM YUUGARE TII CUM AESTIVUM	268 Ct	MESTVX MODIMPRISTWXY
NZA SATIVA A MAYS	230 AL	XTERGHELLMANPORRSSETTVVWY PGHLLMNPRSSSTVVWX
FAGUS VIRGINIA.	274 LN	R
MARDOPSIS THAL.	276 IM 278 R	μ
ASSICA GLERACEA YORK MAX	280 L	wv
THE PER PERSON.		
DICAGO BATTVA	299 H	
ENCAGO BATTVA COTIANA TABACUM COTIANA DEBNEYI	292 AC	DEPGGHBIO.LLMNPGBR888TTVVWXY
ENCAGO BATTIVA COTIANA TABACUM	292 AC	

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ORSEN MONEY
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CEPHALOPRUS MAXIN.
DAMALISCUS DORCAS
GAZELLA THORSON
KOSIUS FLUPRIPRIM.
MADOGUA MIRID
ORYX GAZELLA
TRAGELAPHUS IMBER.
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TETHANYMERA PYRIC
LIDSHMANA TARRHY!
BICTYOSTELUM DIS.
HYPPANIM POLYCEPH.
MUJROSPORA CRASSA
CANCIDA ARCCAMS
RHYTOPHTHURA PAR.
PODOSPURA ANGERNA
SACDIARDIMYCES CRI.
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CAMDIDA CYLINDRA
          PLANTS.
       CHLAWYOR TRACHOM.
ARABOOPSIS THALL
OLYCINE MAX.
HUSBOLLIS YULGARIS
NICOTIMAN RUSTICA
PETUMA BY
HELIANTHUS ANNULS
SURCHUM RICOLOR
GRYZA SATIVA
THITIQUM AESTIVUM
TRITIQUM YULGARE
SOYBEAN
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       PART TWO: 18MA Sequences
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614 W
618 PP
622 GILPORST
626 DLMPQ
     AVIAN ONCO. VIRUS
CHICKEN ASVOAMMES
MOUSE NAMULY
PHAGE TO
PHAGE TO
     AMEHADBACTERIA.
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  HALORACTERUM CUT.
HALOPERAX VELICAMI
HALOCOCOUR MORPHUM
METHANORAC, THERM
SULFRICIBUS ACEDO,
THERMORIASMA AGIDO
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AAACDBERODGCHSEXLULLUMPPPQRERSSSTTVVWXV
SI
   DURACTERIA
                                                                                                                         110-236
OUNCERNA

NYCOPLASMA CAPRIC.

NYCOPLASMA MYCOPLASMA MYCOPLASMA MYCOPLASMA CITRI

STREFTOWNCES CHE.

STREFTOWNCES COEL.

STAPHYLOCOL EPIG.

NYCOPAC. SMFG.

PACELLUS SUBTILES

THERMUS THERMOPHIL

ECIDI

SALADOMELLA TYPH

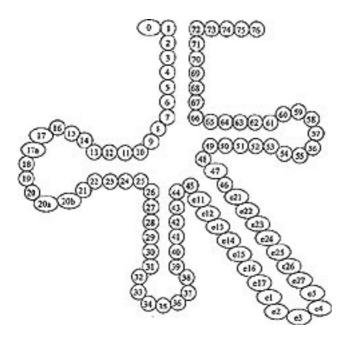
PHOCOSTRIL, TUR.

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SYNECHOCYSTIS SP.
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  ORGANBLES
 CHLOROPLASTS
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CHAMPOONOMAS REM
BUELRIA GRACIE
CODILIN FRAGE E
SCRIGGERIA CRIZA
LIPTUS ALBUS
HORDOM VILGARE
TRITCUM MASTIVUM
ZEA MAYS
GLYCINE MAX
NICOTIAMA TABACUM
PHASICILUS VILGARES
SPIRACIA GLERACEA
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SINGLE CRLL ORGANISMS 260-419
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TETRAHYMENA THERM.
MEUROSPORA CRASSA
SACOHAROMYCES CER.
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PGHKLMPRESSSTWXY
PLANTS.
SOLANUM TUBEROSUM
OGNOTHERA SP.
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PHASEOLUS VULGARIS	440	RULMPWOLY
AMMALS	460	689
ASCARIS SUUM	454	FMS
ARDES ALBOPICTUS HAMETER	480	DEGINORSVX
BAT LIVER	528	DOWNTHWAM DRUKE
BOYINE UVER	538	DGHULRSSSTVWXX
HUMAR	558	S
BUKAMYOTIC CYTOPLASM	9004	999
SINGLE CELL ORGANISMS AND FUND!	600-	009
BUGLENA GRACEUS TETRAHYMENA THERM.	604	DF
SCENEDESMUS DELIC.	608	DDDX FXY
REUPPORPORA CRASSA	620	FX
SACCHAROMYCES CER.	628	ACDEFFGGIHHIKELLLMNPPRIRSSSTTVVVWXY
SCHZDSACOHA, POM. TORULOPSIS UTILIS	632	EFXY
CANDIDA CYLINDRA.	636	ALPYXY ULBSSSS
PLANTS	670-	749
HÖRDEUM VULGARE	678	eer
WHICAT GERM	682	PORMEWXYY
BRASSICA NAPUS	696	F
LUPWUS LUTEUS	694	EFGH WINPEVXY
PHASEOLUS VULGARIS	998 702	tutx
PISUM SATIVUM SPINACIA DLERAÇIA	704	5
NICUTIANA RUSTICA	705	SSSSSYY
SOLAMUM TUBEROSUM CUCUMIS SATIVUS	707	rw.
ANMALS.	750-6	
CAENORHABOL BLEG.	266	
ASTEPINA AVURENSIS		×
BOMET'X MORI	758	AAFFGGI
DROSOPHILA MELANO.	7.79	BPHEX368VVVXY
EUPHAUSIA GROTEA XENOPUS LAEVIS	786	X
SALMON LIVER	792 796	DFX
CHICKEN	904	ŵ
MOUSE	810	EFFRIKKMOORITYXZ
RAT	916	DDEKNOLLNINGSSSVVX
RABBIT LIVER BOVINE LIVER	922	DEKIOUMY
CALF UVER	934	DER.NORRRTWYZ
COW MANMARY GLAND	9.40	ii.
SHEEP LIVER HUMAN	945	HX
- Constant	20.00	AABFOOHLMMNDQSVXYV2
PART THREE IPSIA and IRW	A gene	sequotees that differ from the conversional alignmen
Source		15NA/15NA gano
MTOC-IONDINA	360-6	***************************************
SINGLE COLL ORGANISMS	35D-4	
AND FUNGI		
TRICHOPHYTON MENT.	401	E
ANMALS	790-9	
OPES MELLIFERAL	482	7
MPHNIA PULEX	483	Č.
MALACHOPTERA PHYS.	634	MSS
MALAENDIFTERA MUSIC.	534 535	8
(ALICHOERUS ORYPUS	537	Ē.
HOCA VITUUNA SAMANG	538	8
	842	.5
CHIMPANGSE	572	5
	572 572 580	5 9



**Figure 1.** Numbering of nucleotides in tRNAs. Circles represent nucleotides which are always present; the ovals, nucleotides which are not present in each structure: these are nucleotides before the position 1 on the 5'-end, before and after the two invariant GMP residues 18 and 19 in the D-loop, and the nucleotides in the variable loop. The nucleotide to be added at a given site is indicated by the number of the preceding nucleotide followed by a colon and a letter in alphabetical order. The nucleotides in the variable stem have the prefix 'e' and are located between positions 45 and 46 obeying the base-pairing rules. The nucleotides in the 5'-strand and the 3'-strand are numbered by e11, e12, e13, ... and e21, e22, e23, ... respectively; and the second digit identifies the base-pair. In the case of a long variable region, the loop can be formed by up to five nucleotides: e1, e2, e3, e4 and e5.

modified nucleotides, the other ASCII signs are employed as defined in Table 3. Terminology and structure of the modified nucleosides occurring in tRNAs were used according to (2). Positions in particular sequence which are not filled (gaps in the generalised structure, Fig. 1) are indicated by a dash. All nucleotide insertions are denoted by underlining at the place of insertion.

## Numbering and alignment of the variable region

The alignment of the variable region has been done in accordance with Steinberg and Kisselev (3). The extra arm is placed between nucleotides 45 and 46. The extra arm includes two double helical strands forming a stem and a loop. The annotations of the nucleotides in the extra arm positions begin with the letter 'e' (extra) followed by a one- or two-digit number. We have reserved a space for 7 base pairs in the stem and 5 nucleotides in the loop. The nucleotides in the loop are numbered from 1 to 5, whereas the

nucleotides in the stem are numbered from 11 to 17 (5'-branch) and from 27 to 21, in reverse order, (3'-branch), to indicate base pair formation between nucleotides 11–21, 12–22, etc. (Fig. 1). In the tRNAs where the extra arm position 45 is empty but where the nucleotides 46–48 between the extra arm and T-domain are present, the positions will be filled in the order 48, 46, 47, i.e. tRNAs use position 48, 46 and 47 for the first, second and third nucleotide, respectively, depending on the length of the sequence in this region. A similar situation occurs in tRNAs without a long extra arm, where the most variable position 47 is deleted in many sequences.

## Alignment of animal mitochondrial tRNAs

In properly aligned tRNA sequences, nucleotides occupying the same position in different tRNA sequences should play a comparable structural or functional role. Most animal mitochondrial tRNAs cannot be easily aligned with other tRNAs mainly because of the absence of information about their three-dimensional structure. Experimental data, however, point to the existence of tertiary interactions in these tRNAs. In this compilation, we use an alignment which accounts for these interactions as much as possible. Where we could do so, the animal mitochondrial tRNAs were included in Parts I and II. The alignment of animal mitochondrial tRNA is, however, not yet unambiguous.

Some animal mitochondrial tRNAs have completely unusual secondary structure and cannot be fitted in the tRNA alignment used here (Parts I and II). We treated these sequences separately including them in Part III. Here, each particular sequence has its own alignment. To this group belong the tRNAs from: (i) mitochondria of a parasitic worm lacking the T-, or D-domain; (ii) mitochondria of molluscs, insects and echinoderm, with extended anticodon and T-stems; (iii) mammalian mitochondria, lacking the D-domain.

For some tRNA genes the secondary structure pattern cannot be clearly established. We have also included these sequences in Part III. It is possible that post-transcriptional modifications of these tRNAs will result in improvement of the secondary structure.

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Table 2. Format of tRNA sequences in EBI databank

				accept	D-domain		anticodor	domain	variable red	ion	T-domain		accep
mber				stem					extra loop	1			eten
	Amt i	Lodon	10	012345678	91111111111	1122222222	2222333333		4-ccccccccccc		4555555555	5666666	6666777
1		Organism	Kingdo	0	0123456778	900012345	6789012349	678901234	5111111112345		9012345678	9012345	6789012
	- 1	or gament	[ Kingdo	-1	2	ab			1234567	7654321			
260	TGC	PHAGE T5	VIRUS	-GGGCGAATAGT	arcage-gog-	- AGCACACO	ACCRUMINACE E	mornocous.	**				
				*-*	****	*		ICIOGIA	***************************************	G-GGAG			
340	TGC	ARCHAEGLOBUS FUI	G. ARCHAE				GCCTTTGCGA			accaca	GGTTCAAATC		
				*******						accucu	GUITCHANTC		STUCA
380	TGC	HALOBACTERIUM CO	T. ARCHAE	-GGGCCCATAGC	PCAGT GGT-	-AGAGTGCC	TCCTTTGCAA	GGAGGAT		GCCCTG	GGTTCGAATC		
					****					900010			arcon-
420 7	rgc	HALOBACTERIUM HA	L. ARCHAE	-GGGCCCATAGC	CAGTGGT-	-AGAGTGCC	TCCTTTGCAA	GGAGGAT		GCCCTG	OTTGGAATO		
				******									*****
580	IGC	METHANOBAC.FORM	. ARCHAE	-GGGCCCGTAGC				GGCGGAG		GCCCCG	GGTTCAAATC	CCGGTGG	TCCA-
	man.	METHANOBAC . THERM						****					*===*=
520 .	LOC.	METHANOBAC.THER	. ARCHAE	-GGGCCCGTAGC						GCCCCG	GGTTCAAATC	CCGGTGG	STOCA-
560 7	nac:	METHANOCOC, VANI	ABOTTAD	-GGGCCCGTAGC	BEEF		*****				*****		**-
	L-Gro-	HEISENCOC. VALLE.	ARCHAE	-GGGCCCGTAGC	CAGTT-GGG-	- AGAGCGCT					GGTTCAAATC		
70 9	rac	METHANOTHRIX SOF	H ADCHAR	-GGGCTTGTAGC		30300000		*****			-*		*-
		THE PERSON NAMED IN	ar mounts	##*####	CAGCI-GGI-		SCCTTTGCAA	GGCGGAG			GGTCCGAATC		
80 1	rgc :	METHANOTHERM, PE	R. ARCHAE	-GGGCCCATAGC				aacaaaa					
				*******			secci i despe	oocuumu			GUTTCAAATC		
780 1	rgc .	METHANOSPIR. HUN	G. ARCHAE	-GOGCTCGTAGCT	CAGCT-GGA-	- AGAGCGCG	COTTTGCAA	OGCOGAG			GTTCAAATC		
				*-*							MITTEROMETE:		STCCA-
940 T	'GC	THERMOCOCCUS CEL	ER ARCHAE	-GGGCCGGTAGCT	CAGCCTGGG-	-AGAGOGTO	GCTTTGCAA	GCCGAAG			BUTTOGAATO		
													******
80 7	,GC	THERMOPROT. TENA	X ARCHAE	-GGGCCGGTAGT(		-AGGACGCC	CCTTGCGC	GCGGGGAG			GTTCGAATC		
	-	THE PROPERTY OF THE PARTY OF TH		******	*	****	*****				*****		
ST C	Jac.	THERMOPROT. TENA	X ARCHAE	-GGGCCGGTAGT(	TAGCGGA-				***********	ATCCCG	SCITTOGAATOC	ccggccgc	TCCA
40 7	me :	MYCOPLASMA CAPRI	a propose			*		****					*****
40 1		MICOPLASKA CAPRI	C. BUBACT	-GGGCCCTTAGCT	CAGCT-GGG-						GTTCGATCC		
80 T	GC I	MYCOPLASMA MYCOI	D. FURACT	-GGGCCCTTAGCT			*****				*====		
		THE PROPERTY MICOI	D. BUBACI	-doccerrage;	CNGCI-GGG-		GCCTTGCAC	acagggg			GTTCGATCCC		
00 T	GC I	MYCOPLASMA PNEU.	KURACT	-GGGGATGTAGCT							*		
							antituche.				KITTTGAGACO		
30 T	GC I	ACHOLEPLASMA LAI	D. EUBACT	-GGGGCTTTAGCT	CAGCT-GGG-			acaggag			GTTCGAT-CC		*****
1				*******					COLORADO DE NE		MITCORI-CC	ACT MAGE	TCCACC
31 T	GC /	ACHOLEPLASMA LAI	D. EUBACT	-GGGGCCTTAGCT	CAGCT-GGG-	-AGAGCGCCT	GCCTTGCAC	GCAGGAG			KITTOGAT-CO	VacTtacaca	20001.00
						****	****						and *
60 T	ac :	SPIROPLASMA MELI	F. EUBACT	-GGGCCCGTAGCT	CAGCT-GGG-	-AGAGCACCT	GCCTTGCAC	CAGGGG			GTTCGATCC		
				******			*****				*====		
80 T	GC I	BORRELIA BURGDOR	F. EUBACT			AGAGCATCO	GCTTTGCAAG	CCGAGG			GTTCGAGTCC		
				*	****								*****
70 T	ac I	ENTEROCOCCUS HIR	AE EUBACT	-GOOGCCTTAGCT		AGAGCGCCT	GCTTTGCAC			GTCAGCC	GTTCGATCCC	COCTAGGO	TCCA
	00 .	CONDUCT OCCUPANT		*	***			*****					
90 T	OF 3	STAPHYLOCOC, AUR	B. EUBACT	OGGGGCTTAGCT		AGAGCGCCT	GCTTTGCACC			GTCAGCG	GTTCGATCCC	CCTAGTO	TOCACO

Table 3. Modified nucleosides in tRNA and their abbreviations

one	letter code of no	cleatides			
V.			1	G	unknown modified guanosine
	symbol (2)		K.	m1G	1-methylguanosine
	V		L	m2G	N2-methylguancsine
		name [2]		Gm	2"-O-methylguanosina
		V	B	m22G	N2, N2-dimethylguanosine
			1	m22Gm	N2,N2,2'-0-trimethylguanosine
U	U	uridine	7	m7G	7-methylguanosine
C	C	cytidine	T.	fa7d7G	archaepsine
A	A	adenosine	0	Q	queuosine
G	G	guanosine	8	manΩ	mannosyl-queuosine
Т	T	thymine (for sequences of tRNA genes only)	9	galQ	galactosyl-queuosine
-		empty position	Y	yW	wybutosine
	(underline)	insertion (see footnote for further information)	W	o2yW	peroxywybutosine
		unknown nucleotide		11 10 200 0 0000	beautiful antenne
		and the factoring	N	20	unknown modified uriding
н	PA.	unknown modified adenosine	1	mnm5U	5-methylaminomethyluridine
-	m1A	1-methyladenosine	2	s2U	2-thiouridine
1	m2A	2-methyladenosine	J	Um	2'-O-methyluridine
+	i6A	N <sup>6</sup> -isopentenyladenosine	4	94U	4-thiouridine
	ms216A	2-methylthio-N <sup>6</sup> -isopentenyladenosine	8.	ncm5U	5-carbamoylmethyluridine
-	m6A	N <sup>6</sup> -methyladenosine	1	mem5U	5-methoxycarbonylmethyluridine
6	t6A	N <sup>6</sup> -threonylcarbamoyladenosine	S	mnm5s2U	5-methylaminomethyl-2-thiouridine
E	m6t6A	N <sup>6</sup> -methyl-N <sup>6</sup> -threonylcarbamoyladenosine	3	mom5s2U	5-methoxycarbonylmethyl-2-thiouridine
1	ms2t6A	2-methylthio-N <sup>8</sup> -threonylcarbemoyladenosine	v	pmo5U	uridine 5-oxyacetic acid
	Am	2'-O-methyladenosine	5	moBU	5-methoxyuridine
î.	1	Ingsine	1	cmnm5U	5-carboxymethylaminomathyluridine
0	m1I	1-methylinosine	4	cmnm6s2U	5-carboxymathylaminomethyl-2-thiouridine
-	Ar(p)	2'-0-ribosyladenosine (phosphat)	×	acp3U	3-(3-emino-3-carboxypropy)uridine
	lo6A	N <sup>6</sup> -(cis-hydroxylsopentenyl)adenosine	90	mohm5U	5-(carboxyhydroxymethyljuridinemethyl ester
		in -test-type oxyrisoperaterryttadendsine	i	cmnm5Um	5-carboxymethylaminomethyl-2"-O-methyluridine
<	7C	unknown modified cytiding	-	nom5Um.	5-carbamoyimethyl-2'-O-methyluridine
96	62C	2-thiocytidine	D	D	dihydrouridine
В	Cm	2'-O-methyloytiding	P	Y	pseudouridine
M	ao4C	N <sup>4</sup> -acetylcytidine	1	m1Y	1-methylpseudouridine
2	m5C	5-methylcytidine	z	Ym	2"-O-methylpseudouridine
	m3C	3-methylcytidine	T	mbU	ribosylthymine
	k2C	Ivsidine	F	m5s2U	5-methyl-2-thiouridine
5	15C	5-formylcytidin		m5Um	5, 2'-0-dimethyluriding