03=
04 <u>Cardiovascular Droiase</u>
05 Arterosclerosis
06 Myocardial Infrarction
07 Strokes—

deleté heart disease" co l'appres n/03

16 <u>Infections Diseise</u> 17 Ural Infections

11/13 Alsheimins + Dementia

add "dementia) + collapse

10 <u>Mental Disease</u> 12 Depression 14 Sch 120phenia

21 immunological disease should read "auto-immune disease"

24 genetrally engineered diseases delete from table

Number 1 problem in developing conntries in year 2000?
01- Infections disease
02- Parasitu disease
03 - Malarin
04 Aids
05 Filarrasis
06- Hepatitis
07- Malnutrition
08. Food supply: feeding the hungry
09- Sanitation proflems
10- Population growth/control
11- Cardio vascular problems
12- Cancer 13- Aging/problems
14- DIARRHEA
15- PNEUMONIA
16- HEART DISEASE
17- HYPERTENSION
18- Todine deficiency
19- IRON DEFICIENCY
20- INFANT MORFAUTY
21- DIABETES
22- HEALTH CARE DELIVERY
23- MENTAL CUNESS
24- HARD DRUGS
25- VIRAL DISEASES
26- EXPLOMATION BY DRUG COMPANIES
27- ALCOHOUSM

98- Don't Know 99- None

9-130) 1-132)

10f2 What should be nations #1 3-134) 5-136) 01-Molecular/Cellular brology 02-Cardiovascular 03-Cardiovascular causes 04 Cardiovascular prevention 05-Cardivascular treatment 06-Infections disease 07 aids 08-Viral diseases 09-Genetic diseases 10-Neurosuence 11-Brain disease 12aging /age related problems (n/s) 13alzheimen 14-Preventive medicin 15-Busic / fundamental research 16-Mental Illness 17-Heart disease / causes 18atherosclerosis Heart disease prevention Heart disease treatment 21-Etrlogy of disease 22-Identifying population at risk 23-Epidem wlogy Bromedisal research all areas of research are important from hove provity over the other 97-Don't Know /NA None

e Veron

#861018

]. 3

26-	Endocrinology
	Immunology
	Vacunation / vacune development
	anti viral agents
	Population control
	Cancer: causes
	Cancer: cures
	Cancer: prevention
	Cancer: Freatment
35-	Degenerative desiases
	Did
37-	Resear her automony / freedom
38-	Life styk related problems : obesity / smiking /alcohol
39-	Increased funding
	Pain research
	Prevention of disc degeneration
	Emphasis on the productive - less on aging /terminally ill
43 -	Inadequate # of vessarch personnel
44-	availability of health care
45 -	
46-	artificial contraception fatortion
47-	Identifying ineffective treatment
48-	Esearch ment human Emphasis on Macht societal needs

TO SANGE OF THE SECRETARY OF

Ê

	PG. 10F2
the diseases frontations you th	ink may be climinated by year 2000
(139-40) (141-42)	
01- Measles (16)	26- KAWASAKI
02. Aids (14)	27 - MALNUTRITION
03- Paliomyelitis (12)	28 - VITAMIN DEFICIENCY
04- Rubella (6)	29- GOITER
05. Malania (6)	30- SCHISTOSOMIAGIS
06- Hepatitis (GEN) (6)	31- HEPATITIS A
07- Chicken pox 5	32- HEPATITIS B
08- Huntingtons (4)	33- PNEUMONIA
09- Lenkemia (3)	34- LEAD, POISONING
10- Diptheria (3)	35 - ENVIRONMENTAL HEALTH PROBLEMS
11- Whooping Longh (3)	36- ENPHYSEMA
12- CHOLERA ()	37- LUNG CANCER
13- Tuberculosis (2)	38- PRENATURITY
14- Mumps (2)	39- HYALINE MEMBRANE DISEASE
15- Dishites (2)	40- RH ERYTHROBLASTOSIS
16- Cystic Febrosis (2)	41- RICKETS
17- Cardiovascular (2)	42- YAWS
y 18- Infection disease	43- CANCER/CANCERCEUS
19 - Uvial disease	44- IMMUNE DISEASES
20- HEREDHARY DISEASES	45- Dysentery
21- TYPHOID FEVER	46-INAUENZA
22- 6CARLET FEVER	47- BREAST CANCER
23- SICKLE CELL ANEMIA	48- HEPATOCELLULAR CARCINOMA
24- THALASSEMIA	49-TRYPANOSOMA
25- HEMOPHILIA	50- RABIES
98 - DON'T KNOW/NOANSWER	51- HEART DISEASE
99- None	

A SECTION OF THE CONTROL OF THE CONT

a4	P6-20=2
CONTID.	
(137-38) (139-40) (141-42)	
2- GAUCHER	
63- 1AY-5ACH5	
54- ARTEROSCIEROSIS	
65- OSTEOARTHRITIS	·
66- LOW BACK PAIN	
59- MULTIPLE SCLEROSIS	
58- Alzheimers	
59- ARTHRITIS	
60- LEPROBY	
61 - PSYCHIATRIC DISEASES	
62- HODEKINS	
63- IATROGENIC BIRTH DEFECTS	
64- SCHIZOPHRENIA	
65 - BILARETTE SMOKING	
66- DEATH FROM ORG AN DISEASE	
67- GALL BLADDAR SURGETLY	
68- DIAMPHERS	
69- ROTA	
70 - OSTEOPOROSIS	
71- CYSTOMYOSIS	•
72- TROPANSTIOMASIS	
73- Herpes	
74- Vitamin A deficiency	

Base: 149 P6. 10F3 Disease or condition which genetic g will have to year 2000? 14)(145-46) Cancer (1/s) (30) 02- Drabetes (16) 03 - Sickle Cell anema (12) (8)Cystic Febrosis (7)(7) 08 alzhuners (5)(4) (4) (4) (4) 12-Gowth hosmone deficiencies (4) 14-(3) Infection disease (n/s) (3) Genetic diseases (1/5) 16-Cardio vascular disease 17-18-(2) Genetically engineered bacteria Hyportensio 20-

(2)

Malnutrition

22-

49- REVALKIONEY FAILURE

PG. 30F3

## Q8 BIGGEST FRUSTRATION FOR GLIENTISTS OVER NEXT 14 YEARS?

	to the same time to the same that the same time to the same time time to the same time time to the same time time time time time time time ti
01	LACK OF APEQUATE FUNDS/SUPPORT
02	GOVERNMENT/BUREAUCRATIC INTERFERENCE/REGULATIONS (N/6)
03	GETTING APPROVAL FOR EXPERIMENTS
04	REGULATIONS REGARDING ANIMAL RESEARCH/ENTERFERENCE WITH ANIMAL RESEARCH
.05	PRUS REGULATORY GROUPS/GETTING DRUSS ACCEPTED
06	LACK OF FREEDOM FOR SCIENTISTS/REGEARCHERS
07	REGULATIONS REGARDING STUDIES ON HUMANS
.08	UNDERSTANDING MOLECULAR/CELL BIOLOGY
. 09	UNDERSTANDING BASIC MECHANISM/ETIOLOGIES
10	COMPLEXITY OF CENTRAL NERVOUS 646TEM FUNCTION
ji	FMMUNOLOGY REJECTION/IMMUNE RESPONSE
12	LACK OF ANIMALS FOR REGEARCH
13	NEED NEW ANTIBIOTICS / DRUGS
14	PREVENTION: MOTIVATING PUBLIC TO PREVENT/CHANGE HABITS /STOP SMOKING
_15	LACK OF GUPPORT FOR LONG-TERIU RESEARCH
16	DRUG REGISTANCE
¡커.	INCREASING COLLABORATIVE ASPECTS/PIFFICLUTY GETTING COLLABORATION FROM EXPERTS
18	SLOWNESS OF PROGRESS
19	LAG BETWEEN ADVANCES & APPLICATION
20	APPEARANCE OF NEW DISEAGE/VIRUS
21	ETIOLOGIES OF MENTAL DISORDERS
22	LACK OF RECOGNITION/INTEREST IN THE FIELD
ત્રે3	INCREASING CURE RATE
24	LACK OF RESEARCH PERSONNEL
25	COMPLEXITY OF THE PROBLEM
	QUALITY OF RESEARCHERS
27	REGULTS/DRUGG PROVEN INEFFECTIVE IN HUMAN TRIALS (WHENEFFECTIVE IN ANIMALS/IN-VITRO)
28	DIFFERENCES BETWEEN NORMAL 4 CANCER CELL
29.	POPULATION'S ACCESS TO TECHNOLOGY/KNOWLEDGE

57 TRANSITION FROM SIMPLE LINKS TO HIGH LEVEL ORGANIZATION OF CENTRAL NERVOUS SYSTEM

αÞ	- CONTIN-
58	NEW SURGERIES WHICH INCREASE SUSCEPTIBILITY OF PATIENT 10 MICRO ORGANISMS
59	SOCIETAL RESISTANCE TO USE HIGH TECHNOLOGY
60	POPULATION UNRESPONSIVE TO FINDINGS
61	UNDERSTANDING PRECISE ROLE OF DIETARY FACTORS IN DISEAGE
62	INABILITY TO REPAIR TRREVERSIBLE CHANGES
63	GROWING INCIDENCE OF CARDIO MYOPATHY
64	SIDE EFFECTS OF DRUGG
65	TO DETERMINE RELEVANT OUTCOME MEASUREMENTS IN INFANT NUTRITION
66	AGE-RELATED PROBLEMS
67	ANTI- GENETIC VIRUSES .
	ABILITY TO THINK CREATIVELY
69-	unperstanding mechanics of tissue + interaction betw. mechanism
70	developing methodologies to isolate neural systems
71-	how animal systems fit into human brain
72	Multiplicity of types of tumors
	The state of the s
:	

98 PON'T KNOW/NO ANGWER

 Oak	Other area of brotochnology research which is more promising that those mentioned?
Brotech	nology neep Input
(an	·-17) (218-19) (220-2)
01	. New methods for protein synthesis
02	· new methods for protein synthesis · automation of DNA sequencing procedures
03	· transgenic mice - taking human gene + put into mice
04	· preventne medicine
.05	· molecular basis of memory + behavior
0lo 11	1. choning of genes (for molecules)
07	. work in yeast - its a enkaryote can excrete proteins
08	· use of bromotecules in electronic Switching mechanisms
09	· safé reproductive pill
10	development of delivery methods for getting proteins into people
11	Selective herbrade engineering pestresistant plants salt to brant weather tobrant genetic engineering to improve plants animals
	genetic engineering to improve plants animals
12	inderstond motecular tevel what is going on in any disease
	-> forming therapeutic proteins - for I a month use -slowly released
98 DK/NA	HEAR HOW CNSWOTKS  14 RECEPTOR MEDIATED SIGNAL TRANSDUCTION  15 REGEARCH INTO LINKING BEHAVIOR TO GENETICS OTHE MOLECULAR LEVEL

Q7a	fundamental question in order to achewe breakthrough in bestechnology
(	222-23) (224-25)
01 1	Understanding how gene is regulated / > (9) Mechanism of gene control
	700000000000000000000000000000000000000
02	Molecular basis of cell function (3)
03	Understanding molecular basis of neuronal patterning (2)
04	Better nethods for westing new genes into genome
05	E Molecular mechanisms of growth + cell differentiation
06	F How genetic sequence determines the 3-D folding /structure
67	Etiology & Pathogenisis of dueases
8	Interrelationship between hormones of the immune pystem
S	Studies on mechanism of action of natural mediators of immune inflammatory of proliferative response
THE RESERVE OF THE PROPERTY OF	
	Kaulation; frontrol of gene
VV.	
	→K/NA
99	DONE

Q. 10a	Biggest advance, woulding from bio-engineering by year 2000
,	14-15) (316-17)
	[ Understanding / controlling rejection / compatitulity (11)
	Controlling immune system / response (3)
02	New drugs (1)
	Identifying genes involved in necognition of pelf & mon-self
	Bone Marrow-meeting gener with bone barrow a getting them expressed
	by recipients
<u>15</u>	Development of ne agents
	Computer assisted muscle movements generated by computer
······································	

98 DK/NA

99 NONE

106	Biggist advance (forcemen) resulting from bio engineering by year 200?
	,
01	Molecular etrology  (3)  Understanding mechanism of cell growth (3)
	Understanding oncogenes (2)
04	D Detection (2)
	B Stemulation of immune system to combat/ (2)
	B Stromelation of immune system to combat/ (2)  identify cancer
03(	new drugs spee a Hack one ogenes on their products
05	More selective delivery of drugs using monoclonal autibody conjugates
06	Improved Treatment
	Understanding the events in the environment that cause mutations
8	Hybrid Teating.
	Discovery of growth & autigrowth factors in cancer cells

98 DON'T KNOW /NO ANSWER

3401 PP

100	Biggist advance for heart disease resulting from bis-engineering	ig by
(322-23	) (324-25)	
Ol	Control of lipid levels in blood	(3.)
0,6	Understanding of lipid metabolism	(2)
<u></u> 0€	Control of lipid levels in blood  Understornding of lipid metabolism  Use of TPA/ tissue plasminogen activation	(2.22)
04	Understanding mechanisms of atherosclerosis	(3)
٥٤	Regulation of chalesteral	(3,)
00	Control/regulation of LDL (bow density lips protein)	(2)
٥٢	1 Early diagnosis of predisposition	
	Understanding of hourt hormones	
	Developing techniques for preventing the occlusion of the coronary blood is	ressels
	10 neverse atherosclerosis specifically plaque formation	
	1 Effective control of hyperteusion	
	•	

98 DK/NA

340U PP

BIOTECHNOLOGY

Q18B

one Application is most promising

(364-65) (366-67)

Manipulation of the host/immune system to prevent rejection 01

02

Genetic manipulation of the graft either to improve function or prevent rejection Introducing new gener into an organism to replace missing or defective ones 13

- Q19 A
- Of Creation of food lenriched food products can be made by biotechnology concentrates
- 02 Gene therapy (insertion of genes to treat gene disorders)
- 03 Development of an effective individualized anticancer treatment for each patient
- 04 Antisense RNA. away you can turn off the expression of certain genes
- O5 Treatment of cancer-some of the growth factors and the drugs predicted on them will have major effects against tumors as new chemothera putic agents
- 06 Limb regeneration by understanding differentiation and growth control
- 07 Regenerating tissues and whole organs
- 08 Concer vaccine
- 09 Correcting inherited diseases by bioengineering i.e. in conjunction with bone marrow transplantation
- 10 Synthetic chemistry
- 11 New pharmacology-effects gene regulation
- 12. If we knew the rules of protein folding we could make specific re-agents and drugs
- 13 New advances on hormonal requirements of cells from the nervous system. There are many compounds needed by braincells which have not been identified
- 14 Design and make small molecules that will do the same thing as large, naturally assisting macromolecules such as proteins, enzymes or peptide hormones
- 15 Early diagnosis and Prevention

- OI Synthetic molecules will replace insulin, growth hormone, TPA, urokinase, streptokinace.
- 02 Antibodies directed against neoplastic cells would revolutionize anticancer therapy
- 03 Limb regeneration-There are really notherapies for severed limbs
- 04 Growth inhibitory factors would replace cytoloxic drugs
- 05 Treating tumors by identifying the specific gene that was mutated. Then you could turn off the mutant gene. That would replace chemotherapy Iradiation therapy
- Ob Development of an effective individualized anti-cancer treatment will replace all chemotherapy
- 07 Hemophilia by Factor III and diabetes by insulin
- 08 Cancer vaccine would replace radiation and chemotherapy
- 09 Regenerating tissues and whole organs will replace transplants
- 10 Prevention will replace all treatment
- 11 Thalasemia treated by transfusion will be condidate for gene therapy

Q6b Other area of rescench in infections disease nurse promised that what y (216-17)(218-19)(220-21)

O1- Understanding of basic mechanisms of immunity

O2- Pathogenisis

O3- ONA/miclicated study

O4- Understanding of molecular mechanisms of parasites

O5- of what diseases

O6- Ding delivery systems

O7- ROLES OF INFECTIONS IN DISEASES THAT ARE NOT THOUGHT 10 BE INFECTIOUS

O8 USE OF MONO CLONAL ANTI-BODIES AS IMMUN-THERAPUTIC ASENTS

O9 IMMUNIOLOGIC 4CHEMOTHERAPUTIC CONTROL OF FILARIAL DISEASES

98 DK/NA 99 None / no others

Infattons Osiane

#861018

Q. 11a Biggest advance in prevention/treatment of MALARIA by

Infections Disease

Fundamental question reeds answer in order to achoese major break through in treatment of infections disease? (222-223)(224-225)

Anderstanding of basic mechanism of cumunity

02- Basis of pathogenuity 03- Mechanisms of drug action + resistance 04- Better understanding of viruses

05. Development of preventire measures

06- Understanding brochemistry + evolution of intermediary metabolism

Identifying patterns of protein

BETTER DRUG TARGETING

09- BETTER DELIVERY METHODS

ioentification of latent viruses + bacterial L' forms

98 DK/NA 99 None

98 DON'T KNOW/ NO ANSWER

99 NONE

DON'T KNOW/NO ANSWER

Q. 11a Brygest advance in prevention/treatment of MALARIA by year 2000? (320-321) (322-323)

01- Develop effective/new vacuine 02- Mos gni to control 03- Anti-malaria drugs

04 USE OF DOT

DON'T KNOW/ NO ANSWER

99 WNE

## Infections Disease

Q. 116 Biggest advance in prevention/heatment of common COLD by year 2000 34
(324-25)(326-27)

01- Interferon

02- Therapeutic drugs; anti-viral / Chemotheapy

03- Immunization

04- Better understanding of weld visuses

05 ACHIEVING THE CONTROL OF THE MATRIATION PROCESS OF ENZYMES

06 UNDERSTANDING HOW INTERFERON WORKS

98 DONT KNOW/NOANEWER 99 No major advance

## (314c L357-58] [359-60] [361-62]

Ol New antiviral lantifungal drugs will replace etisting compounds Itherapies
Ol Pseudomones will replace 8-tartans antibiotics
Ol Vaccines will be replaced by prevention
Old Nucleoside analogs now used for therpes Virus will be replaced by new agents
Old Treatment for measles and hepatitis will be replaced by a vaccine
Old Use of combination therapy using two antibodies
Old Displace the poor compounds we have now
Old Common penicillin will be replaced by new drugs
Old Toxic drugs will be replaced by polynicleotides
In Antiviral drugs lantibiotics will be replaced by Immunization

98 DLINA 99 None Bio Medical Infections Diseases

#861018

Diseases in developing countries which should be top research priority. 366/61) (368-69) 01- Malaria (16) 02- Schistosomus (8) (5) 04-(3)05-Typhond (3)06-Taber culosis (3)Viral discise 07-(3)(3) 08-Parasitii disease ACUTE RESPIRATORY INFECTIONS MEASLES POLIO 98- DK/NA 99- NONE 8 18th Diseases in 4.5. + developed world which should be top reasearch priority 70-71 (372-73) Ards 01-(16) Influenza (4) Pneumoma (2) Cyto megaloverus (3) Pulmonary disease (u/s) (2) 06 -RESPIRATORY DISEASES 07 -INCOGENIC VIRUSES 08-VIRAL INECTIONS PROTOZOAL DISEASES (continued) P:2 PULMONARY BRONCHITIS

98- DON'T KNOW/NOANSWER

7a (	ardio vascular - fundamental question which needs to be answered to achieve break through in treatment of central hervons disorders
223)	(224-225)
01	- Molecular mechanism of heart disease (3)
02	- Molecular mechanism of heart disease (3) - Molecular mechanism of atheroschlerosis (5)
03	- Causes / etiology of heart disease (2)
04	- Causes/etiology of heart disease (2) - Causes/etrology of atheroschlerosis (5)
05-	Development of methodo to affect lifestyle al terations
06	Pathogenesis of hypertension
27-	What leads to progression of heart muscle disease
08-	Michanism of conversion from chronic to acute discise in ischemia
09-	Finding best therapy which makes atherosclerosis disappear
10 -	Understanding factors which make individual susceptible
11-	How type A behavior accelerates heart disease
12-	Understanding what controls levels of high density 11 po proteins
13-	Understanding what controls levels of high density lipo proteins Relationship between at a thunganesis + throm bogenesis
98	DOD'T KKKM/ NO ANSWER
	NONE

Fio Medie ardervas a R.11a		#861018 Base: 18 by year 2000;
(316-17)(3	318-19)	
01-	Prevention of / reduction of smoking. Change in dictary habits	(3)
	Reduction/prevention of atherosclerosis	(3)
	Thrombolytic therapy	(4)
86 -	REDUCTION OF BLOOD LOW DENSITY UPO PROTEIN	
06 -	REDUCTION OF CHOLESTEROL LEVELS	
	LOENTIFICATION OF SUSCEPTIBLE INDIVIDUALS BY MEANS OF GENETIC MARKERS	
08 -	DRUGG TO STOP INFARCTION PROCESS	
09 -	UNDERSTANDING WHAT PRECIPITATES CLOT	
	NEW AGENTS TO PREVENT PLATELET AGGREGATION PRODUCED USING MONOCUNAL ANTI BODY TECHNIQUE	
	MODIFICATION OF TYPE A BEHAVIOR	
12-	BETTER UNDERSTANDING OF THE MOLECULAR MECHANISMS OF ARTERIOSCLEROSIS	

PARTING AND THE SECRETARY OF THE SECRETA

MEDICAL	# 861018
OVASCULAR IIA_CONTO	PG. 20F2
16-11) (318-19)	
13- PROGRESS IN CORONARY BY PASS	
	,
98- DON'T KNOW/NO ANSWER	

DK/NA

None /no major advance

98 DONTKNOW/DOANSWER

99 NONE

	26 What changes in lifestyte would do most to reduce incidence / seventy of cardio vascular disease?	
38)(33	1-40)(341-42)	
01-	Reduction of smoking (9)	
02 -	Dietarry changes (8)	
<i>6</i> 3-	Dietary changes (8) Reduced fat intake (3)	
04-	Better weight control: less calories/food (6)	
65-	Control of hypertension (4)	
06-	Increased exercise /physical activity (6) Modification of Type A behavior Lower cholesterol	
07-	Modification of Type A behavior	
08-	Lower cholesterol	
09-	Less sodium	
	Coll Commercial Coll	
	to the late of the second state of the second state of the second	
F S AMERICA CONT.		
TE TO - Different 121 - Marion - Constitution -		

DON'T KNOW/NO ANSWER

qq

qq

NONE

Ol	(356-57) (358-59) - BETTER / GAFER ANTI - ARRHYTHMIC DRUGG	
02		
03	,	(3)
٥٠		ALINFARCTION (3)
05	- EMPROVED THROMBOLYTIC AGENT/ANTI CLOTTING DRUGS	(3)
06	- AGENTS TO CONTROL/PREVENT ATHEROSCLEROSIS	<u> </u>
07	- SUPROVED TREATMENT/CONTROL OF REJECTION IN TRANSPLANTS	
08	- LAGER TECHNOLOGY/TECHNIQUES	(£)
00	- CATHETER BASED TREATMENT/TECHNIQUES	(2)
19	- DRUG MODIFICATION OF ATHEROMOUS PLAQUE/DRUGGTHAT REDUCE PLAQUES	)
<u>ii</u>		
12	- DRUGG TO PREVENT ARTERIOSCIEROSIG	
13	- DRUGS THAT ALLOW NUTRIENTS INTO THE HEART MUSCLE	
!	- USE OF UPID CONTROL DRUGS	
	5- GENETIC THERAPIES	
10	- NEW BURGICAL ENTERVENTIONS	
	1- MODIFICATION OF TYPE A BEHAVIOR	
](	- BETTER TREATMENT OF HEART PAILURE	
1	- CHANGES IN LIFESTYLE	

98 DK/NA

99- NOTHING

KDiorascular

99 NONE

0/186 What pr 0-71) (372-	rogress well be in our ability to tell who will	suffer from	a (heart attack)
7G) (317- OI-	18) Identification of /knowledge of Genetic markers / DNA mankers	(6)	(4)
and the state of t	Identification of Risk factors	(4)	<u>(</u> †)
03-	Non in vaoire testing/techniques.	.(4)	(3)
64-	Non in vaoire testing/techniques. Imaging techniques	(3)	(5)
05-	Better study / control of hypertonsion ,	/ -	(3)
06-	Screening population for blood cholesteral	levels	
07-	Stress tests		
68-	Better disanstic techniques		
	Better diagnostic techniques Detecting plasma lysid moiety Undustanding lipoproteins		
09-	// di to		
10-	white anding apoproteins		
And the second s		MARIE MARIE MARIE MARIE E ESTA STATE E MARTE	
		THE STREET STREET STREET STREET STREET	
and a company cold and a second		**	
00 0	DON'T KNOW NA		

## GOLD-MEDICAL IMPLANTS

Q6B 16 THERE ANY OTHER MAJOR A	REA OF EMPLANT RESEARCH	WHICH IS MORE	PROMISING
THAN THOSE MENTIONED?			
(216-17) (218-19) (220-21)			

OI ORTHOPEDIC EMPLANTS/REPLACEMENT	(4)
(JOINT REPLACEMENT, SPINAL REPLACEMENT)	
02 MUSCULAR-SKELETAL IMPLANTS	(\$)
03 MUSCULAR ALLO-GRAFTS	(1)
04 PANCREAS ALLO GRAFTS	(i)
05 THYROID PARATHYROID ALLUGRAFTS	(i)
Ob ENDOCRINE IMPLANTS	(i)
07 OSTEOCHODROSKELETAL 1156UE	(1)
08 IMPLANTABLE DEFIBRILLATORS	(1)
09 IMPLANTABLE INFLIGION PLIMP	()
10 PROSTHETIC DEVICES	(1)
IL ELECTRICAL TREATMENT OF OSTEOPOROSIS OF THE SPINE	(i)
12 ELECTRICAL TREATMENT OF OSTEOARTHRITIS	(V)
13 BIOWGICAL FIXATION- OSTEO INDUCTIVE METHODS	
14 BIO ARTIFICIAL ORGANG	
IS REPLACEMENT OF SOFT TISSUEIN MUSCULARSKELETAL SYSTEM	
16 HYBRID ARTIFICIAL ORGANS	
17 IMPLANTING A MATERIAL THAT BEGING AS A MECHANICAL DEVICE BUT THROUGH	
BIODEGRAPATION & REPLACEMENT ENDS AS A BIOLOGICAL DEVICE	
18 FETAL GRAFTING OF TISSUE FOR LIGAMENT/CARTILLE DEFECT	Replacement

98 DON'T KNOW/NO ANSWER

99 NO/NONE

(7)

## GOLD - MEDICAL IMPLANTS

Q12 IN WHAT ONE AREA DO YOU SEE IMPLANTS AS PLAYING THE GREATEST ROLE IN THE YEAR 2000?
(316-17) (318-19)

01	CARPIOVAGCILLAR	(F)
	(HEART VALUE IMPLANTS, ARTIFICIAL HEART REPLACEMENTS)	
02	ORTHOPEDICS	(16)
	CORTHOPEDIC IMPLANTS, HIP IMPLANTS, KNEES, JOINT REPLACEMENTS/	
	IMPLANTS/RECONSTRUCTION, ARTHRITIE, EPINE) MUSCULAR SKELETAL (ne-read: Stim. Elean muscles - ms.)	
13	MUSCULAR SKELETAL (ne-read: Stim. clear muscles -ms.)	(3)
04	· · · · · · · · · · · · · · · · · · ·	(2)
05	IMPLANTABLE DRUG DELIVERY PLIMPS	(D)
06		(D)
Oη	HEARING AIDS	(()
80		
09	AS PREJENTION OF SUDDEN DEATH	
10	MECHANICAL REPLACEMENT FOR DAMAGED ORGANS of hugs that have	
	electrical uiteractions w/ tissue & implant	

<sup>98</sup> DON'TKNOW/NO ANGWER

<sup>99</sup> NOWE

(4)

## GOLD-MEDICAL EMPLANTS

03 HEART

Q14 CAN YOU THINK OF A SPECIFIC INSTANCE WHERE ANIMATE IMPLANTS ARE LIKELY TO

HAUE REPLACED IMPLANTS?

(321-22) (323-24)

OI JOINTO

(JOINT TRANSPLANTS/REPLACEMENT/RESURFACING/ALLOGRAFTS, HIPS,

ALLOGRAFT OF THE KNEE)

(BONE TRANSPLANTS/REPLACEMENT, SEGMENTS OF BONE/SKELETON)

(ARTIFICIAL HEARTS/VALUES, HEART IMPLANTS)

OF LUNGS

OF LIVER

OT PANCREAS

ON KIPNEY

(ARTIFICIAL HEARTS/VALUES, HEART IMPLANTS)

(I)

(I)

(I)

09 LIGAMENTS

10 CONNECTIVE TIGGUE

11 INGULIN GECRETING TIGGUE

98 DK/NA

Q.65	Other areas of CNS research which is more promising than these mentione
(216-	-17), (218-19) (220-21)
	alzheimers (2)
	Understanding multiple oclerosis (2)
	Biological shythms both in clinical & basic research
	NEW PSYCHOTHERAPIES
	NEW BEHAVIORAL TREATMENTS
	Treatments for memory disorders
	Understanding the plastic mechanisms of the brain
08	Prigs for memory
	Drugs for learning
10	Development in infancy + adolescence
1	Understanding embryological, fetal & meonatal development of the CNS
	Understanding virally triggered disorders
	Stabalizing systems of the brain
<u> </u>	Genetics of depression
15	Understanding addictive disorders
ile	Neural death as a part of the pathology of schizophrenia of depression
18 (11) 1 STATE MILLER AND S. BELL MARK THOMAS AND	
en proposati formation nag a a a como e el del de del transformación de trans	,
anne d'andre del manutes resident en recesa à la 1141 filia - 144 de des penerses re	
California Managas de distre de la	
0¢	D. 1412 121 0
	Don't Know/No Ainswer No/None
	1007 None
Programme Community (Community Community Commu	
Kanta araga a remakan	

putral N	knows System
	(2a2-a3) (a24-a5)
Q.7a	Fundamental question needs answer to order to achieve break through
01- A	Understanding motecular/cellular biology of CNS (5)
02-0	Understanding the organization of the brain (2)
03 - C	Under standing the ctrology of CNS diseases (2)
04-B	Understanding dev. of CNS
05-	Discovering new neuro transmitters like peptides +
	Discovering new neuro transmitters like peptides + neuro ammes
06-	Complete genetic library of normal human tabnormal human
07-	Understanding changes in synophi efficacy/synoptic mechanisms
08-	Delermine forctors that bring release of neuro transmitters
	Factors that affect receptor synthesis in healthy of diseased states
	What will allow Cass time to receive to
	What will allow Cors tissue to regenerate  Froctors which govern neuronal degeneration + regenerations
13-	Why brain cells don't replicate themselves
14-	Interelations between different reunal systems
98	DK/NA
99	None

ŕ

011.	PREVENTION/	
QIIa	BIGGEST AD VANCE IN TREATMENT OF ALZHEIMERS BY YE	AR 2000
	(324-25) (326-27)	
	OI UNDER STANDING OF ETIOLOGY/PATHO GENESIS	(13)
	02 UNDERSTANDING OF GENETIC CAUSES/BASIS FOR DISEASE	(2)
	13 FINDING ATREATMENT	(3)
	04 DEVELOPMENT OF DRUGS	(6)
	05 ENVIRONMENTAL FACTORS FOUND TO CAUSE THE DISEASE	(3)
	06 DIET / NUTRIENT TREATMENT	(2)
	07 Demonstration of its autommune origin	
	08 - 4 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	
	og Prevention of the degeneration of specific types of memors	
	jo Discordening endogenous meurotoxins & how to block the action of those neurotor	aris
	il Understanding the brokenical defect	
	12 SURGICAL ALTERATIONS	
	`	,
WE & 0		
	98 DON'T KNOW/NO ANSWER	
	99 NONE	

11c	Biggest advance in prevention/treatment of epilepsy (332-33) (334-35)
,	(332-33) (334-35)
D	1- Better Drus Treatments
	(with ho side effects)
0	2- Understanding heuro transmitters
03	- Hereditary forms will be diagnosable; caught +
er r a a susan recommendad rela	Better obsteties: treated early
04	1- Dangerons use of oxytoxics in labor; artificial
	induction of births
OS	- Hereditary forms will be diagnosable; caught +  Better obsteties: treated early  Dangerons use of oxytoxics in labor; artificial  moduction of births  - Understanding of the pathophyprology
	- Better ways of controlling seizures
0	7- Inhibiting the formation of epileptic foci
	Ludersfanding of brain systems involved in seizures
	- Relationship between brain structure + function
	Bitter suguial elemination of the foir of seizures in these not treatable w/duys
	not treatable w/ duys
	`
aa	8 DK/NA
	9 NONE

2126	Change in lifestyle which would reduce incidence /severity of	:NS.,
(સપધ-	45) (346-47)	is orders;
01	Reduction in also hol use /abuse	(9)
02	Less drug use / abuse	(5)
03	Less smoking	(2)
04	Less lipids	(2)
05	Control of diet /metritional changes	(5)
ما٥	Exercise	(2)
	Reduction of stress	(2)
80	Control of hypertension	(2)
09	Change in sexual habits /promisanty	(2)
	,	()
	autronobile safetye	(2)
11	More attention paid to environal and /dagma understanding afores rolf	
	More attention paid to emotional meeds / deeper understanding of ones self Restriction of family life	
:		
D	Taking responsibility for ones illnesses. People should not entrust doctors with po much of their care	
111	Move away from toxic waste sites	
	Trive may from 10 Howard Ambo	

98 DON'T KNOW/NO ANSWER

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Base	′	2	4
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P6.10F2

2. 14a 2003 important new this april for CNS disorders by year 2000? (363-64) (365-66)

01- Genetic engineering

(8)

02- New drugs

03- New Specific anti-psychotics

04 New/specific anti-depressants

05- Neuropeptides

06-1 Mono clonal anti bodies

67- Treatment for althumers

08- Tissue transplants auto brain

09 - Use of mini formatic Pumps (2)

10 - Treatments for disorders of learning + memory

11- Behovioral Therapies

12- Profective therapies for Huntingtons

13-loe of various new growth factors for neuro regeneration

14- Replication of long acting drugs

16- Combination of behavioral, cognitive + drug treatments

17- Targeted drugs

18- Irumuno suppressive therapy

19- Unti wiral drugs

20 - Prenatal interventions

21- Treatments that exploit adaptive compensatory functions

22- Treatment of endogenous neuro toxens through blochade or through prevention of their manufacture

23 Drugthat prevents demyclimation

24- administration of trophic factors to prevent neuronal death 25- Transplantation of transfected cells to provide definint

98 OK/NA 99 NONE

c Tre	atment which will be replaced a if so by what
(368-	-69) (370-71) (372-73)
02 -	Transplants may replace L-DOPA
03,-,,	Drugs will replace stay in hospitals— Transplants may replace L-Dopa  Origs!  Out; psychotris be replaced by drugs w/ no-
04-	Genetic theapy will completely replace any
05-	Antipsychotic treatments will be replaced by
06-	Sugrial reatment (for brain tumors) replaced by Chemotherapy
07-	(antianxiety drugs) replaced by drugs without abuse potential
	Electro shock eliminated - replaced by drugs + dragnosis
09-1	(antipsychotics) + tranginlizers) replaced by new drugs
10-	New growth fretors would replace rehabitation treatments
11-	Better drugs replace le
11	Trangulizer a W be replaced by new treatment for Alzhemeurs
	Convention
	Dat prog
12-	Anti psychotic fanti depress. replaced by treatments that exploit adaptive compensatory functions of brain Tropic fontors for alz. could replace hydrogene + aut, costorases
	administrating of
/3-	Tropic factors for alz. could replace hypergene + aut, estales
14-	Targeted drugs w/ replace amost drugs

3000-199

ر ( الا	Role of traditional psychoanolytic therapy in year 2000?	
		( )
	Used by affluent people/reserved for those who can afford it	(4
62	an untellectual tool: to analize this personalties slf-improvement/self- knowledge	(5
03	To treat personality/characterological disorders	(2,
04	Educational / historical interest	(2)
		Plant de la Proposition
		er a sikilinganis i
		er - t could also serve a
98	DONTKNOW/NO ANSWER	
99	11.	5)
		****

(	other area of cancer pusearch which is more promising than those mentioned
(216	-In) (218·19) (220-21)
	11
	How to combine drugs + chemotherapy
02-	Brug resistance fundustanding dung resistance
63-	Drug resistance / understanding drug resistance Segnencing of total human genome
	Cellular communication factors
	Basic research
06-	Growth factors / growth factor research
07-	Deorgn of strategies to attack this apentically solid tumo
08-	Epidemiology
	Behavioral aspects of smoking cessation
	Immure modulation research
	Understanding of network systems
	Containment of nuclear accident + industrial pollutants

Steeling Steel steel stage person is

98 DK/NA

Ta	Martin	portant Fundamental Question which much to be answered in order to achieve a major
	Lieuxi	wough in the treatment of cancer
	U	1/2 de to de a contra to
	01-	underganding mechanism of celigioner.
		normal all to cancer cell
,	02-	22-23) (224-25) Understanding mechanism of cell growth +  differentiation / what transform  normal all to cancer cell  Over coming drug resistance.
	03-	Understanding mechanisms of climinating alls that are
	AND THE RESERVE TO SERVE THE PROPERTY OF THE P	Understanding mechanisms of climinating alls that are abnormal.
	04-	How to control it/control of metasticis
	05-	Something ampletely unknown will answer the question
	06-	Detection of some from abrorma leters
	V	Detection of geno typic abnormalities
	07	Identifying viruses which cause cancer
· · · · · · · · · · · · · · · · · · ·	AND THE REAL PROPERTY AND ADDRESS OF THE PARTY	
	er e en e	
* ***		
		The state of the s
		The state of the s
	60	+1./110
	i	JK/NA .
	44	None

w Medine #861018 11A (324-25) (326-27) nun Jab (344-45) (346-47) Base: 26 For lung cancer: Biggest advance in pres Change in lifesty & reduce prevention / freating 126 Eliminating smoke Improved delivery of combination drugs Tentifying ush groups affected by carcing Reduction of fat detake Lus salt 06-Less promiscuity EARLY DIAGNOSIS more maderatchy 98 DK/NA 99 NONE Learning how not to age for breast cancer Bygest advance in prevention /treatment by year 2000? (328-29) (330-31) Early detection Nutritional Changes/ Identification of dietary 03how fut diet Chemotherapy 05 06 Lactation early in ly Markers for patients at high wok Effective / Cers toxic therapeutic a antibody - linked RICIN Better understanding of hormonal influence 12anti-promoters Bitty treatment of carly disea 13-Hormonal Treatment Use of monoclonal antibodies fordiagnoss Comprehensive Screening programs 98 DR/NA 99 NONE

a Bu	the year 2000, 2 or 3 most important completely new types of therapy for caucar
(35	nich are motavailable now
01-	Develop mono donal antibodies
02-	Differentiation agents / therapy
03-	
04-	Improved biologie response modifiers
05-	Ingrovements is organ transplants
06 -	Laser surgery
07-	Modification of unmine system/response
08-	Manyoulation of gene expression
09-	New method for ugulating oneogenes
10-	Interferon
/1-	Different combinations.
12	IDENTIFYING GENETIC SUSCEPTIBILITY
13	MEDICATION THAT DELAYS CELL DIVISION
14	THERAPY DIRECTED AGAINST GENETIC CHANGES OCCUR IN CANCERCELL
15	NEW CHEMÓTHERA PENTIC AGENTS
ال	Hormonal antagoments
TO THE POST OF THE	
	The state of the s

14c		
6	Ópecifi	treatment which will be replaced 4 if so by what
	(3	64-65) (366-67) (368-69)
	01-	Total breast removal will be replaced by early
		Total breast removal will be replaced by early diagnosis
	02	Surgery rodiotherapy + chimotherapy will be climinated by better therapies
	03-	Breast cancer suggery will be replaced by bropsy, radiotherapy + Chemotherapy
	04-	Spleenectomies replaced by interferon + deoxyro formycin
	05-	Chemo replaced by New drugs / agents : monoclonal, cytotoxic
	06-	Present ay to toxic therapy up laced by w/better specifically
	07-	Radiation therapy more localized + specific
	58-	Laser surgery well replace other forms of surgery
	09-	Radial mastertomy uplaced by lump-ectomy/localize
	10 -	Existing dugs replaced by amproved drugs
	//-	Defferentiation agents will replace some the agris
,	12-	Bulogical dings will replace conventional cytotoxic drugs
4	/3-	Therapy/ Drug combinations well replace conf for breast cancer Certain forms of X-RAY replaced by new agents
1	14 -	Certain forms of X-RAY replaced by new agents

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Other area of nutritional research is more promising

(216-17) (218-19) (220-21) 01- Ostroporosis 02- Obesity research 03 - Dyestun 04- Chemoprevention 05 - Regulation of mutition on gene level 06 - Methods of determining body composition ULTRIENT INTERACTION-RELATIONS HIPOF ONE NUTRIENT TO ANOTHER CHRONIC RENAL DISEAGE 69 SOLID TUMOR CANCER 10 ADAPTING SPECIAL DIETS TO ACUTE & CHRONIC ORGAN FAILURE 11 OTHER AGPECTS OF DIET & DEGENERATIVE DESEASE 12 NUTRITIONAL NEEDS OF SMALL PRE-TERM INFANTS 3 NUTRIENT - NON-NUTRIENT INTERACTIONS 14 EPIDEMIOLOGICAL CORRELATIONS BETWEEN CERTAIN DIETARY CONSTITUENTS & CHRONIC DISEASE 15 LINKS BETWEEN DIET CANCER & DRUGS & CANCER 16 UNDERSTANDING NUTRIENT BIOANALABILITY & THE EFFICIENCY BY WHICH WE ABSORB NUTRIENTS FROM FOOD 17 SIGNIFICANCE OF NUTRIENT IMBALANCES & INDIVIDUAL NUTRIENT TOXICITY 18 BIOLOGICAL SIGNIFICANCE OF PROTEINAGE THABITORS 19 FATTY AUD METABOLISM 20 DETERMINING NUTRIET REQUIREMENTS FOR SPECIAL POPULATIONS/INDIVIDUALS 21 GENETIC MATA BOLIC & POYCHOLOGICAL FACTORS THAT EFFECT FOOD INTAKE 22 EFFECTS OF DIETS ON PSCHOLOGICAL & BEHAVIORAL FUNCTIONS 23 EFFELT OF DIET ON GROWTH & MENTAL DEVELOPMENT 24 EDUCATING THE POPULATION TO CHANGE DIETARY HABITS 25 STUDIES IN TASTE OF FOOD SELECTION 26 ENERGY METABOLISM 27 CONVINCING FOOD INDUSTRY OF THEIR BE NUTRITIONAL RESPONSIBILITY

98 DONTKNOW/ NO ANSWER 99- None/no others

:	nm Killion	١
aub		

	<b>U</b>
	Regulation of nervous system by nutrition
35	Specialized use of nutrients subtrates treating specific disease
34	Thermogenesis in human cells
33	Alcoholism
32	Nutrition & Canar
31	Interaction of drugs 9 mutrients
	Identification of trace elements
	Nutrition and other mon-immune defense systems
	Modification of metabolic + nutritive response to unjury to and seps

utrition

Ta	Fundamental question which needs answer to acheeve breakthrough
23)(22	4-25)
01-	Better under standing of details / metabolism
	Better understanding of details/metabolism at molecular/cellular level
02-	Role of specific untrients/nutritional patterns in cancer/hypertension/ba
	cancer/hypertension/ba
13	ROLE OF NUTRITION IN THIMUNO-MODULATION
04	MECHANISM OF ACTION OF VITAMINS & MINERALS
15	BIOCHEMISTRY BEHIND THE EMMUNOLOGY OF CANCER
	TO DETERMINE RELEVANT OUTCOME MEAGUREM ENTS IN INFANT NUTRITION
67	RESEARCH INTO ENERGY INETABOLISM
80	NUTRIENTS & THEIR FUFULENCES ON GENETIC EXPRESSION
69	HOW TO MEASURE DIETARY TUTAKE ON LARGE GROUPS OF PEOPLE/DIFF. PEOPLE
	LINDERSTANDING WHAT CONTROLS OBESITY/FOOD ENTAKE/SELECTION
	MEAGURING INDIVIDUAL DIFFERENCES IN MONITORING TUDIVIDUAL REACTIONS TO STRESS
	Understanding metabolism of brain
	Genetic influences on the metabolism
	REGILLATORS OF BASIC METABOLIC PROCESSES
	INTERACTION OF NUTRIENTS. HOW WHAT PEOPLE EAT AFFECTS THEM
•.	WOERSTANDING METABOLIC PROCESSES & ENDIVIDUAL DIFFERENCES
	EHRLY LIFE NUTRITION INFLUENCE ON LONGTERM RESISTANCE
18-	Notare of metabolic reg. of whole what benome
19	Function of the entero cytes
20	correlation of intracellular por ticles w/intermediary metabolisin of food 5 hoffs
21	control of appetite + body wight food 5 high
	system for RAPID ASSESSMENT of nutritional status
23-	BIOCHEMICAL SUBSTATE TO FUEL CELLS TO REACH ITS MASSIMUM POTENTIAL

98- DK/NA 99 None Bro Medical Nutrition

99- None

7-29)	Regest metritional advance in prevention of cancer ()
01-	High fibri dut (2)
	Reduction in calonus (2)
03-	Dietary prevention of cancer (3.)
04-	Dictary prevention of cancer (3.)  BELATIONSHIP of nutrients to cancer foods (2)  Discovery of promoters of cancer infoods (2)
05-	Low fat / defining roles of fat (2)
06.	
06 -	Arolecular basis of varions mutort wonat problems  Accessing vit. A as inp. / viramins  Ose of wat vitamin A analog/roles of duetary betacaro tene, vit. A
67-	Nutritional support for therapies: radiation, chemo, surgery
08-	Modifications of current the apentic measures of drugs + radio therap
09-	Manypulations of minunological functions
0-	Mand + mental acquilation
11-	Factors which prevent normal cells from behaving properly in
	factors which prevent normal cells from behaving properly in presence of malignant cells
12-	Great fibers which will bind fatty acids  Stop growing tobacco  Guater change in fast food preparation
/3 -	Stop growing tobacco.
14-	Guater change in fast food preparation
15 -	Use of vitamin A, B12, FOLICACID TO PREVENT CHROMOSOME DAMISE
16 -	Stop contaminating air /water/food
17-	Duit concentrates on available controlydiates
	V
<b>.</b>	

Biv-Medical Natrition Olob Biggist advance in 332-333 (334-35) 01- Dictary control of blood cholesterol. 02 - Reduce fat intake 03 - Public education Role of 3/fatty acids (omega) Control of calone intake 06- Physical activity.
07- HAPERTENSIVE DIDETISE AS IT relates to Sodium / potrass Dutory modification (controlis Putrition Dietary prev. of atherosclerosis Use of specifized formulations of non-metabolizable lysid analogs Reduced arterioselerosis disease Control via nutrition of obesity / hupper 14 identify susceptibles + treat them 15 Better dicts Highfiber combin. det , drugs, physical activity ability to reduce atherosclusis ksions martenes 20 --resuis to bind beleaceds 21understooned role of forthy acid in regulation of choleste

98 DK/NA

## Bio Medial Nutrition

01-	Greater emphasis on physical activity
02-	Public education
03-	Calonic control / Eating less
04-	Understanding the genetic factors that virtuence obesity
05-	Clarification of the role of brown fat
06-	The use of lipid analogs
07	Development of appetite suppressant drugs
08-	Oberity Assessment
09-	Development of peptides that control satiety maturally
10-	Development of effective drug therapy that is page
	Understanding energy metabolism + regulation of energy storage
12-	Dealing with patiety / Diets that will mariave patiety
13-	low fat diet
14-	Change in lifestyle
16-	Greater intake of vegetables & fruit
16-	Total dit Nather than isolated elements
<u>  [-</u>	Behavior modification toward certain foods
18-	appetite controlling mechanism
i9 -	Understanding how the body penses its energy balance of the public queut effect on food witake
20-	CAN BE PREVENTED IF CONTROLLED EARLY IN LIFE
21-	BAFE METHODS OF CONTROLLING FAT STORAGE TO ALLOW THE FAT TO BE BURNED OFF AS HEAT
22-	ALTERNATIVELY SAFE REGULATORS OF FAT METABOLISM TO INTERFERE W THE LAYING DOWN OF FAT
23-	PSYCHOLOGICAL AS WELL AS NUTUTIVE CHANGES PSYCHIATRIE SOCUTION
98-	Don't Know/ No Answer
99-	NONE

01-	Incorporating beneficial fatty acids in fords/ use of 3 fatty acids
	use of 3 fatty ands
02-	Duts fails made to modify psysiological function
	life long in people
03-	Dry-rutrient interrelationships / freatments
04-	Identify compounds that modify feeding behavior
05-	Knowledge of nutrotional interactions
06-	· Knowledge of nutritional interactions Developing of diets for all nutrients + all diseases
07-	Use of purified parcreatic enzyme preparation
08-	Insulin pump
09-	Mono clonal antibodies + targeting
10-	Prinatal nutritional symplements
11-	Dut suprovement for school children
12-	Dut suprovement for school children Duts which moderate psychiatric illners fase in the early
/3-	Nutruuts for prematures
14-	agents absorbed + affecting cellular receptors not enter intermediary metabolism
	noi eurs mediang mil -
./5-	Dingsthat prevent catabolic processes
16.	Lipo some delivered medicines
/7	artifical feeding devues
18-	Over the counter foods designed specifically for specific diseases
19-	Developments in TPN
20-	Mutrition of all patients / dentifying nut- risk for selective surgery
21-	The of des fais
22-	Understanding of correct calcium intoke
23-	ability to monitor glucose-ensuling ability to access mairidual mitritant requirements
24-	abouty to access individual metritent requirements
	·

23	Total parenteral nutrition
	Mutritional therapy for prevention/treatment of osteoporosis
	Mutrient combinations which affect prople differently
28-	Genetic corrections of inform errors of metabolism
	During metricuts across cellulas membrane
	Induction of enzyme deficiency
	Nutritional therapy that improves muscle performance
32-	Mutritional therapy beneficial to body but harmful to viral
33-	Mutritional therapy beneficial to body but haimful to viral infections!
34-	Mutritional therapy for chronic reval disease
35-	Meeting trace mineral regionements for patients on prolonged parenteral nutrition
	prolonged parenteral nutrition
36-	aftering metrotion w/out complications of lives in newborn
37-	Briding file acids which lowers cholesteral
	Intravenous amino acid mixtures which modify near tralismitter behavior
	Meanstralls milly behavior
39-	New sources of protein
	Understanding of maryinal defociences + able to treat them
41-	Mutritional therapy for disorders of central nervous system
42-	amino and analogs iseful as substitution for naturally
43-	Broady available ify of microputrients
44.	Recognition of nutritional factors that delay aging
	PRODUCE ON FARMS BETTER MEATS/EGGS
46-	VITAMINS (C, E, BATACARO TENE) to prevent cancer
43-	Broady available lifty of microputrients

99 NOWE

01-	Present fad diets replaced by new fad diets
02-	High potency vitamins will be replaced by carotenoids
53-	Intravascular muroscopii techniques w/dietorytherepy well replace heart surgery
04-	Polynmaturated + low cholesterol diets will be replaced
65-	BYPASS SURGERY REPLACED BY NUTRITIONAL THERAPY
06-	LOW CHOLESTEROL DIETS WILL BE REPLACED BY OTHER APPROACHES
07-	SURGICAL APPROACHES TO OBESITY REPLACED BY PREVENTION / DIET
08	Precise measurements of energy consumption by individual not populations

01-	Improved methods to reduce complications/ . 3
	Improved methods to reduce complications/ adverse reaction/infection
62-	Finding ideal combinations solutions for specific conditions (2
	Defining the requirements for trace elements (2)
03-	Improving combinatori/solutions generally
04-	Simplification to require less supervision. Better preservatives pumps, live technology.
05-	Capitalization on TPN as an experimental tool for better understanding of human mutrition
.06-	Being able to treat outside the hospital
07-	More Knowledge about nutrient requirements 4 interactions
08-	Improved high everyy pourtes
09-	Advances in equipment the logistics of administration
_10	Attempts to make it more balanced generally
_ 11	longer activity
12-	DEFINING REQUIREMENTS FOR TRACE ELEMENTS
13-	PREDICTION OF ITSEFFECT ON DRUG METABOLISM EFFICIENCY & TOXICITY
14-	Clearer indications of complicated cases of the ability to handle them
i5-	POLYNEURIC NUTRIENTS
16-	learning how to monitor patients so we tailor the amounts to their daily needs
17-	SYSTEM WHICH INTRODUCES NUTRIENTS TO LIVER FIRST
	for low workers infrants

Qai

MOST IMPORTANT ADVANCE YOU FORESEE IN ENTERAL NUTRITION

(370-7D (372-73)

- 01 Total replacement of parenteral nutrition
- 02 Development of more efficiently absorbable diets
- 03 Use of more elemental farmulas a management of intestuial diseases
- 04 Furthering the use of enteral nutrition in the developing countries
- 05 Better use of the proper formulas for malnurished children
- 06 Understanding the bio-availability of different nutrients in the diet
- 07 De Understanding dictary component interactions
- 08 Returnts simpler less expensive formulas
- 69 Obesity Control
- 10 Pharmacologic treatment of gastronitestinal dysfunction to endotorin
- Il Understanding the role of early nutrition in adult obesity
- 12 Products for specific conditions
- 13 Defining + incorporating trace elements into enteral motriture
- 14 Development of guidelines for use with drugs
- 15 Shiftin populations diet away from saturated a high fat diet
- 16 A loss of fear of eating
- 17 Providuig mecessary fatty acids 4 minerals
- 18 Better toterance tolerated formulas
- 19 & Understanding a alteration of digestion
- 20 Knowledge of absorption patterns . metabolism of different mutrient mixtures
- 21 Determining digestive rabsorptive capacities, tolerances
- 22 Side effects could be lowered
- 23 Risk identification
- 24 Tubing the stomach of a person with little mental function
- 25 Making them more acceptable
- 26 availability of Jords that have optimal mix of appropriate Yetty acids dother food stuffs in a form people will eat nother than taking additives of supplements
- 27 Education of public to eat variety of Joods withrus the four Basic food Groups a adjusting portion size to every expenditure

33 Better definition of specific mutrient reguments

34 Making their more palatable

35- FORMULATE SAE SUBSTRATES SO CAN TRACE PATH of ABSORPTION + NUTRITION

THRONGH BODY + ICHM WHICH CELLS BENIFIT

36. USE of DIETARY MODIFICATION to MINIMIZE GENETIC INSULTS AS
RESULT of RADIATION, DRUGS, IS MOKING

37- Redefinition of balance of enteral solutions

38- understand specifie tissue requirement

39 - Develop opecialized diet for hospital patient to

98 DON'T KNOW NO ANOWER