

# TOXCENTER (Toxicology Center Database)



## Subject Coverage

- Adverse Drug Reactions
- Air Pollution
- Animal Venom
- Antidotes
- Carcinogenesis via Chemicals
- Chemically Induced Diseases
- Drug Evaluations
- Environmental Pollution
- Food Contamination
- Mutagenesis
- Occupational Hazards
- Pesticides and Herbicides
- Radiation Teratology
- Toxicological Analysis
- Waste Disposal

## File Type

Bibliographic

## Features

Thesauri      Biosystematic Code (/BC), MeSH Tree Number (/MN), Chemical Name (/CN), Organism (/ORGN), Controlled Term (/CT), Supplementary Term (/ST), Geographic Term (/GT)

[Alerts \(SDIs\)](#)      Weekly or monthly (Weekly is the default)

[CAS Registry Numbers®](#)            Page Images            STN AnaVist     

[Keep & Share](#)            [SLART](#)            [STN Easy](#)     

Learning Database            Structures            STN Viewer     

## Record Content

- Bibliographic data
- Abstracts
- Indexing terms
- CAS Registry Numbers

## File Size

More than 10 million records (1/12)

## Coverage

1907-present

## Updates

Weekly

## Language

English

## Database Producer

Chemical Abstracts Service  
2540 Olentangy River Road  
P.O. Box 3012  
Columbus, Ohio 43210-0012 USA  
Phone: 800-753-4227 (North America)  
Phone: 614-447-3700 (worldwide)  
Fax: 614-447-3751  
E-mail: [help@cas.org](mailto:help@cas.org)

<b>Sources</b>	<ul style="list-style-type: none"><li>• ANEUPL Aneuploidy File</li><li>• BIOSIS® 1946 to the present</li><li>• CAplus<sup>SM</sup> 1907 to the present</li><li>• CIS CIS Abstracts</li><li>• CLINICALTRIALS.GOV ClinicalTrials.gov</li><li>• CRISP Toxicology Research Projects</li><li>• DART Development and Reproductive Toxicology File</li><li>• EMIC Environmental Mutagen Information Center File</li><li>• EPIDEM Epidemiology Information System</li><li>• ETIC Environmental Teratology Information Center File</li><li>• FEDRIP Federal Research in Progress</li><li>• HAPAB Health Aspects of Pesticides Abstract Bulletin</li><li>• HMTC Hazardous Materials Technical Center File</li><li>• IPA 1970 to the present</li><li>• ISRCTN ISRCTN Register of Clinical Trial Numbers</li><li>• MEDLINE® 1946 to the present</li><li>• PESTAB Pesticides Abstracts</li><li>• PPBIB Poisonous Plants Bibliography</li><li>• RISKLINE Swedish National Chemicals Inspectorate</li><li>• TSCATS Toxic Substances Control Act Test Submissions</li></ul>
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<b>User Aids</b>	<ul style="list-style-type: none"><li>• Online Helps (HELP DIRECTORY lists all help messages available)</li><li>• STNGUIDE</li></ul>
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<b>Clusters</b>	<ul style="list-style-type: none"><li>• ALLBIB</li><li>• AUTHORS</li><li>• BIOSCIENCE</li><li>• CASRNS</li><li>• COMPANIES</li><li>• CORPSOURCE</li><li>• ENVIRONMENT</li><li>• FOOD</li><li>• FORMULATIONS</li><li>• HEALTH</li><li>• MEDICINE</li><li>• PHARMACOLOGY</li><li>• TOXICOLOGY</li></ul>
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[STN Database Clusters](#) information (PDF)

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**Pricing** See the [STN Price List](#) or enter HELP COST at an arrow prompt (=>).

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## Search and Display Field Codes

The fields that allow left truncation are marked with an asterisk (\*).

Search Field Name	Search Code	Search Examples	Display Codes
Basic Index * (contains single words from the title (TI), abstract (AB), named person (NA), chemical name (CN), supplementary term (ST), controlled term (CT), geographic term (GT), organism (ORGN), biosystematic codes (BC), and gene name (GEN) fields, as well as CAS Registry Numbers (RN))	None (or /BI)	S GROWTH FACTOR S HEPATITIS S ?ALLEL? S 79-07-2 S MEDICAL DEVICE# (L) MANUFACTURER# S HOME (S) INTERVENTION	AB, BC, CN, CT, GEN, GT, NA, ORGN, RN, ST, TI
Abstract * <b>(1)</b>	/AB	S ANTIPSYCHOT?/AB S INDUCED-ARRHYTHMIA/AB S ?TOXIC?/AB S (HAND (S) TREMOR#)/AB	AB
Accession Number	/AN	S 2001:3269/AN	AN
Author (includes inventor and editor) <b>(2)</b>	/AU	S SCOTT, M?/AU	AU, SO
Author Group	/AUTH	S BROOKS A?/AUTH	AUTH
Biosystematic Code <b>(3,4,5)</b> Superterm <b>(6)</b>	/BC	S PURPLE BACTERIA/BC S 85306/BC S HOMINIDAE/BC AND 57-88-5	BC
Chemical Name <b>(5,7)</b> (contains names from the chemical name (CN) and CAS Registry Number (RN) fields)	/CN	S INTERLEUKIN?/CN S SAL-I/CN	CN, RN
Chemical Name Segment *	/CNS	S ?FLUOR?/CNS	CN, RN
Classification Code <b>(8)</b> (Concept Code)	/CC	S GENERAL BIOLOGY?/CC S "FORESTRY AND FOREST PRODUCTS"/CC S 28:08.04/CC S 60-2/CC S 10006/CC	CC
Clinical Trial Numbers	/NCT	S ISRCTN02140505/NCT S NCT00005487/NCT	NCT
Collaborator	/AUCL	S BEGAY JACK/AUCL	AUCL
Comment	/CM	S TOXICOL?/CM	CM
Controlled Term <b>(5,10,11)</b> (includes main terms)	/CT	S SUNSCREENING AGENTS/CT S (HYPERTENSION (L) BL)/CT S *MEDICARE/CT S F3.375.100./CT	CT
Corporate Source <b>(9,12)</b> (Patent Assignee)	/CS	S FORD/CS S SURGERY DEPARTMENT/CS S ROSIGLITAZONE STUDY GROUP/CS	CS
Country of Publication <b>(13)</b> (code and text)	/CY	S L1 AND US/CY	CY
Digital Object Identifier <b>(32)</b>	/FTDOI	S 10.1111?/FTDOI	FTDOI, DOI
Document Number	/DN	S 2072020/DN	DN
Document Type (code and text) <b>(14)</b>	/DT (or /TC)	S L1 AND J/DT	DT
Duration Begin, Date (Initial Project Date) <b>(15,16)</b>	/DB	S DB>=2000	DB
Duration End, Date (Final Project Date) <b>(15,16)</b>	/DE	S DE>=2000	DE
E-mail Address <b>(17)</b>	/EML	S A.DAY@UTORONTO.CA/EML	CS, EML
Entry Date <b>(15)</b>	/ED	S L8 AND ED>=20020600	ED, UP
Field Availability (code and text)	/FA	S L7 AND RN/FA	Not displayed
File Segment	/FS	S CAPLUS/FS AND MODULATION S CLINICAL TRIALS.GOV/FS	FS

## Search and Display Field Codes (cont'd)

Search Field Name	Search Code	Search Examples	Display Codes
Gene Name (13)	/GEN	S C-JUN/GEN S HUMAN APOE GENE/GEN	GEN
Group Author	/AUGR	S DANACOL GROUP/AUGR	AUGR
Geographic Term (3,5)	/GT	S (PALEARCTIC REGION (S) AFRICA)/GT S UK/GT	GT
Index Term (contains single words from the named person (NA), chemical name (CN), supplementary term (ST), controlled term (CT), geographic term (GT), organism (ORGN), and biosystematic codes (BC) fields, as well as CAS Registry Numbers (RN))	/IT	S GENETIC ENGINEERING/IT S (BACTERIA (L) VIRUS)/IT S (CYTOTOXIN (S) PROTEIN)/IT S 64-17-5/IT	BC, CN, CT, GT, NA, ORGN, RN, ST
International Standard (Document) Number (contains CODEN, ISBN, and ISSN) (17)	/ISN	S LANCAO/ISN S 0022-3263/ISN S 9971-62-253-X/ISN	ISN, SO
Inventor (18)	/IN	S BUSH R?/IN	AU
Journal Title Code (10)	/JTC	S 9892302/JTC S AN7/JTC	SO
Journal Title, Full and Abbreviated (19)	/JT	S DRUG SAF./JT	JT, SO
Language (code and text) (20)	/LA	S L8 AND EN/LA	LA
Meeting Date (3,15,21)	/MD	S MD>20010600	MD, SO
Meeting Location (3)	/ML	S ORLANDO/ML	ML, SO
Meeting Organizer (3,9)	/MO	S ONCOLOGISTS/MO	MO, SO
Meeting Title (3)	/MT	S (ANNUAL AND SCIENTI?)/MT	MT, SO
Meeting Year (3,14)	/MY	S MY>=2001	MD, MY, SO
Named Person or Institution (22)	/NA	S MOZART?/NA	NA
Number of Contract (10)	/NC	S DE07085/NC S NCI/NC	NC
Number of Report (22)	/NR	S NASA-00001502/NR S NASA/NR	NR, SO
Order Number (23)	/ON	S NTIS-OTS0590168/ON	ON
Organism Name (3,5) (includes Superterms) (6)	/ORGN	S RODENTIA/ORGN AND L1 S HUMANS+ALL/ORGN AND L1	ORGN
Other Source (24)	/OS	S (2001:9177 AND IPA)/OS	OS
Patent Country (18)	/PC (or /PCS)	S US/PC	PC, PI
Patent Number (18,31)	/PN (or /PATS)	S US4561876/PN	PI
Patent Number/Kind Code	/PNK	S US2005233010/PNK	PNK
Publication Date (15,19)	/PD	S PD>=20020100	PD, PI, SO
Publication Year (15,19)	/PY	S 2001-2002/PY	PI, PY, SO
Section (25) (code and text)	/SC	S DRUG ANALYSIS/SC S 8/SC	SC
Source (contains publication title, publication date, collation information (volume, issue, pagination), meeting information, patent information, editor, publisher, number of report, space flight mission, investigator, and affiliation, CODEN, ISSN, and ISBN)	/SO	S EPA/OTS/SO S (NEW DRUGS AND 19 AND 1)/SO S (UK AND PAT)/SO S FLIGHT EXPERIMENT/SO S SCIENCE PUBLISHER/SO S (JPCHAX AND ASAP)/SO S NASA00027887/SO	SO
Summary Language (26) (code and text)	/SL	S DE/SL	SL
Supplementary Term (5,27,28)	/ST	S ANTIDOTES/ST S CARDIAC DRUGS/ST S CHEMISTRY/ST (L) MAJOR CONCEPTS/FA	ST

## Search and Display Field Codes (cont'd)

Search Field Name	Search Code	Search Examples	Display Codes
Supporting Organization (Sponsoring Agency) <b>(9,29)</b>	/CSS	S "U.S. DEPT. OF HEALTH AND HUMAN SERVICES"/CSS	CSS
Title *	/TI	S ULCER/TI	TI
Update Date <b>(15)</b>	/UP	S L1 AND UP>=20020600	ED, UP
Uniform Resource Locator <b>(3)</b>	/URL	S HTTP://WWW.BIOSCIENCE?/URL	SO, URL
Zip Code <b>(30)</b>	/ZP	S 02139/ZP	CS, ZP

- (1) Available for all file segments except ANEUPL and ETIC.
- (2) Available for all file segments except TSCATS.
- (3) Available for BIOSIS file segment only.
- (4) The /BC field is a subset of the /ORGN field.
- (5) There is an online thesaurus associated with this field.
- (6) Enter HELP STERMS at an arrow prompt in the file for a list of superterms.
- (7) Available for BIOSIS, CAPLUS, DART, EMIC, ETIC, IPA, and MEDLINE file segments.
- (8) Contains code, text, and major terms for BIOSIS file segment; contains CA section and subsection for CAPLUS file segment; contains code and text for IPA file segment; contains the code for the CIS file segment; contains the TSCA Section for the TSCATS file segment.
- (9) Search with implied (S) proximity is available in this field.
- (10) Available for DART, EMIC, and MEDLINE file segment only.
- (11) MeSH Tree Numbers are also searched in this field. (L) proximity is available with Qualifiers. Postings for MeSH headings do not include narrower terms, while MeSH Tree Numbers do include all narrower levels.
- (12) Available for BIOSIS, CAPLUS, CIS, CRISP, DART, EMIC, FEDRIP, IPA, MEDLINE, PESTAB, AND TSCATS file segments.
- (13) Available for CAPLUS and MEDLINE file segments only.
- (14) Available for ANEUPL, BIOSIS, CAPLUS, CIS, DART, EMIC, ETIC, IPA, MEDLINE and PESTAB file segments.
- (15) Numeric search field that may be searched using numeric operators or ranges.
- (16) Available for FEDRIP file segment only.
- (17) Available for BIOSIS, DART, EMIC, and MEDLINE file segment only.
- (18) Available for BIOSIS and CAPLUS file segments only.
- (19) Available for all file segments except CRISP, FEDRIP, and TSCATS.
- (20) Available for all file segments except EPIDEM, PPBIB, and TSCATS.
- (21) When the meeting date is multiple days, e.g., 5-10 FEB 2001, only the first and last days are searchable.
- (22) Available for MEDLINE file segment only.
- (23) Available for TSCATS file segment only.
- (24) Available for BIOSIS, CAPLUS, EMIC, FEDRIP, IPA, and MEDLINE file segments only.
- (25) Available for IPA file segment only.
- (26) Available for IPA file segment only.
- (27) Available for ANEUPL, BIOSIS, CIS, CRISP, EMIC, EPIDEM, ETIC, FEDRIP, HMTIC, IPA, MEDLINE, PPBIB, RISKLINE, and TSCATS file segments.
- (28) Search single words from the CAPLUS ST field in /BI or /IT.
- (29) Available for CRISP, EPIDEM, FEDRIP file segments only
- (30) Available for CRISP and FEDRIP file segments only.
- (31) Either STN or Derwent format may be used.
- (32) Available in BIOSIS and MEDLINE file segments only. Unable to EXPAND on this field.

## Limiting Search Codes

Only an L-number for an answer set created in TOXCENTER may be limited.

Search Field Name	Search Code	Search Examples
Animal subject	/ANIMAL	S L3/ANI <b>(1)</b>
English language records	/ENGLISH	S L1/ENG,HUM <b>(1,2)</b>
Female subject	/FEMALE	S L2/FEM <b>(1)</b>
Human subject	/HUMAN	S L1/HUM <b>(1)</b>
Male subject	/MALE	S L2/MALE <b>(1)</b>

- (1) The code may be abbreviated to the first three letters.
- (2) An answer set may be limited to more than one subject area.

## Chemical Name (/CN) Thesaurus

All Relationship Codes can be used with both the SEARCH and EXPAND command in the Chemical Name (/CN) thesaurus.

Code	Content	Example
ALL	All associated terms (SELF, CN, RN, RR, EC, UF, USE, HM, PA, INDX, NOTE, PNTE, RE)	E CHAETOGLOBOSINS+ALL/CN E 8057-37-2+ALL/CN
AUTO (1)	Automatic Relationship Code (SELF, USE)	E BROMOACETIC ACID+AUTO/CN
HM	Heading Mapped to (SELF, RN, RR, CN, EC, HM)	E NEOSPORIN+HM/CN
NOTE	Notes associated with the term (SELF, CN, RN, EC, RR, INDX, PA, NOTE, PNTE, RE)	E SERICYSTATIN+NOTE/CN E EC 2.4.1.119+NOTE/CN
PFT	Preferred and Forbidden Terms (SELF, CN, RN, EC, RR, UF, USE)	E COMBRETASTATIN+PFT/CN
RN	CAS Registry Number associated with the name or name associated with CAS Registry Number (SELF, CN, RN, EC)	S MANGANESE CHLORIDE+RN/CN E 20603-88-7+RN/CN
RR	Associated CAS Registry Numbers (SELF, CN, RN, EC, RR)	E FLUVALINATE+RR/CN E 7773-01-5+RR/CN
XUSE	USE and UF terms from the current MeSH (SELF, USE, UF)	E 6-CHRYSENYLAMINE+XUSE/CN S 6-CHRYSENYLAMINE+XUSE/CN

(1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /CN Thesaurus

Code	Description
→	Self
CN	Chemical Name and Enzyme Name
EC	Enzyme Commission Numbers
HM	Heading Mapped To
INDX	Indexer Note
NOTE	Scope Note
PA	Pharmacological Action
PNTE	Previous Indexing Note
RE	Reference
RN	CAS Registry Number
RR	Related Registry Numbers
UF	Used For Term
USE	Used Term

## Controlled Term (/CT) Thesaurus

All Relationship Codes can be used with both the SEARCH and EXPAND command in the Controlled Term (/CT) thesaurus.

The /CT thesaurus contains the current Controlled Terms. MeSH Tree Numbers are searchable terms in the /CT thesaurus.

The /CT and /MN Thesaurus have the same EXPAND capabilities except when expanding MeSH Tree Numbers. The /CT Thesaurus will expand the same Tree Number hierarchy, while the /MN Thesaurus will expand the MeSH terms corresponding to the various MeSH Tree Numbers.

Code	Content	Example
ALL	All associated terms (BT, SELF, MN, DC, NOTE, INDX, ENTC, AQ, PNTE, HNTE, ONTE, MHTH, BXTH, PA, UF, USE, QUSE, NT, QLF, QA, QCAT, QNOTE, QINDX, QHNTE, QONTE, QUF, RT)	E PEPTIC ULCER+ALL/CT E C6.405.748+ALL/CT
AUTO (1)	Automatic Relationship Code (Preferred Terms and Qualifiers) (SELF, USE, QUSE)	E NASAL SINUSES+AUTO/CT
BT	Broader Terms (BT, SELF, MN)	E ADV EFF+AUTO/CT E PREGNANCY TESTS+BT/CT
HIE	Hierarchy (Broader and Narrower Terms) (BT, SELF, MN, NT)	E RECEPTORS, DRUG+HIE/CT
KT	Keyword Terms (multiword phrases containing the term) (SELF, KT)	S SHOCK+KT/CT
MN	Tree Number and descriptor class (SELF, MN, DC)	E PROSTHESIS FAILURE+MN/CT
NOTE	Notes associated with the term (SELF, MN, NOTE, INDX, ENTC, AQ, PNTE, HNTE, ONTE, MHTH, BXTH, PA)	E RHINOVIRUS+NOTE/CT
NT	Narrower Terms (SELF, MN, NT)	S NEURONS+NT/CT
PFT	All Preferred and Forbidden Terms  (SELF, MN, ENTC, AQ, UF, USE)	E FIBRIN TISSUE ADHESIVE+PFT/CT
QLF	Qualifier and associated terms (SELF, AQ, QUSE, QLF, QA, QCAT, QNOTE, QINDX, QHNTE, QONTE, QUF)	S ADVERSE EFFECTS+QLF/CT
QPFT	Qualifier Preferred (SELF, QUSE, QLF, QUF)	E PSYCHOLOGY+QPFT/CT
RT	Related Terms (SELF, MN, RT)	E ETHICS+RT/CT
STD	Standard (Broader, Narrower, and Related Terms) (BT, SELF, MN, NT, RT)	S SPINAL CORD+STD/CT E PNEUMONIA+STD/CT
UF	Used For Terms (Forbidden Terms) (SELF, MN, UF)	E SEX BEHAVIOR+UF/CT
USE	Used Terms (Preferred Terms) (SELF, MN, USE)	E JOINT TUBERCULOSIS+USE/CT
XUSE	USE and UF terms from the current MeSH (SELF, USE, UF)	E RADICULITIS+XUSE/CT

- (1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /CT Thesaurus

Code	Description
→	Self
AQ	Allowable Qualifier
BT	Broader Term
BXTH	Backwards Cross Reference Thesaurus
DC	Descriptor Class
ENTC	Entry Combination
HNTE	History Note
INDX	Indexer Note
KT	Keyword Terms
MH	MeSH Heading
MHTH	MH Thesaurus
MN	MeSH Tree Number
NOTE	Scope Note, Consider Also Terms
NT	Narrower Term
ONTE	Online Note
PA	Pharmacological Action
PNTE	Previous Indexing Note
QA	Qualifier Abbreviation
QCAT	Allowable Categories
QHNTC	Qualifier History Note
QINDX	Qualifier Indexer Note
QLF	MeSH Qualifier (subheading)
QNOTE	Qualifier Scope Note
QONTE	Qualifier Online Note
QUF	Qualifier Use For
QUSE	Qualifier Use
RT	Related Term
UF	Used For Term
USE	Used Term

## Geographic Term (/GT) Thesaurus

All Relationship Codes can be used with both the SEARCH and EXPAND command in the Geographic Term (/GT) thesaurus.

Code	Content	Example
ALL	All associated terms (BT, SELF, UF, USE, NT)	E TANZANIA+ALL/GT
AUTO (1)	Automatic Relationship Code (SELF, USE)	E GOLD COAST+AUTO/GT
BT	Broader Terms (BT, SELF)	E POLAND+BT/GT
KT	Keyword Terms (multiword phrases containing the term) (SELF, KT)	E GERMANY+KT/GT
NT	Narrower Terms (SELF, NT)	E AFRICA+NT/GT
PFT	Preferred and Forbidden Terms (SELF, UF, USE)	E DDR+PFT/GT
STD	Standard (Broader and Narrower Terms) (BT, SELF, NT)	E PALEARCTIC REGION+STD/GT
UF	Used For Terms (Forbidden Terms) (SELF, UF)	E IVORY COAST+UF/GT
USE	Used Terms (Preferred Terms) (SELF, USE)	E ABYSSINIA+USE/GT

(1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /GT Thesaurus

Code	Description
→	Self
BT	Broader Term
KT	Keyword Term
NT	Narrower Term
UF	Used For Term
USE	Used Term

## MeSH Tree Number (/MN) Thesaurus

In the MeSH Tree Number (/MN) Thesaurus, all Relationship Codes can only be used with the EXPAND command.

The /CT and /MN Thesaurus have the same EXPAND abilities except when expanding MeSH Tree Numbers. The /CT Thesaurus will expand the same Tree Number hierarchy, while the /MN Thesaurus will expand the MeSH terms corresponding to the various MeSH Tree Numbers.

The /MN thesaurus does not have any postings. When searching, it is necessary to edit the field code to /CT.

Code	Content	Example
ALL	All associated terms (BT, SELF, MN, MH, DC, NOTE, INDX, ENTC, AQ, PNTE, HNTE, ONTE, MHTH, BXTH, PA, UF, USE, QUSE, NT, QLF, QA, QCAT, QNOTE, QINDX, QHNTE, QONTE, QUF, RT)	E GRANULOMA+ALL/MN E C23.550.+ALL/MN
AUTO (1)	Automatic Relationship Code (Preferred Terms and Qualifiers) (SELF, USE, QUSE)	E PANCREATIC CHOLERA+AUTO/MN
BT	Broader Terms (BT, SELF, MN, MH)	E ILLUSIONS+BT/MN
HIE	Hierarchy (Broader and Narrower Terms) (BT, SELF, MN, MH, NT)	E CHLAMYDIA+HIE/MN
KT	Keyword Terms (multiword phrases containing the term) (SELF, KT)	E SHOCK+KT/MN
MN	Tree Number and descriptor class (SELF, MN, MH, DC)	E ABSCESS+MN/MN
NOTE	Notes associated with the term (SELF, MN, MH, NOTE, INDX, ENTC, AQ, PNTE, HNTE, ONTE, MHTH, BXTH, PA)	E SPINAL NERVES+NOTE/MN
NT	Narrower Terms (SELF, MN, MH, NT)	E TOOTH+NT/MN
PFT	Preferred and Forbidden Terms (SELF, MN, MH, ENTC, AQ, USE, UF)	E PROSTHESIS FAILURE+PFT/MN
QLF	Qualifier and associated terms (SELF, QUSE, QLF, QA, QCAT, QNOTE, QINDX, AQ, QHNTE, QONTE, QUF)	E AE+QLF/MN
QPFT	Qualifier Preferred (SELF, QUSE, QLF, QUF)	E METABOLISM+QPFT/MN
RT	Related Terms (SELF, MN, MH, RT)	E AIR BAGS+RT/MN
STD	Standard (Broader, Narrower, and Related Terms) (BT, SELF, MN, MH, NT, RT)	E ALCOHOLISM+STD/MN E C21.613.455.571.+STD/MN
UF	Used For Terms (Forbidden Terms) (SELF, MN, MH, UF)	E IODIDE PEROXIDASE+UF/MN
USE	Used Terms (Preferred Terms) (SELF, MN, MH, USE)	E OPHTHALMIA+USE/MN
XUSE	USE and UF Terms from the current MeSH (SELF, USE, UF)	E ARSENIC POISONING+XUSE/MN

(1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /MN Thesaurus

Code	Description
→	Self
AQ	Allowable Qualifier
BT	Broader Term
BXTH	Backwards Cross Reference Thesaurus
DC	Descriptor Class
ENTC	Entry Combination
HNTE	History Note
INDX	Indexer Note
KT	Keyword Terms
MH	MeSH Heading
MHTH	MH Thesaurus
MN	MeSH Tree Number
NOTE	Scope Note, Consider Also Terms
NT	Narrower Term
ONTE	Online Note
PA	Pharmacological Action
PNTE	Previous Indexing Note
QA	Qualifier Abbreviation
QCAT	Allowable Categories
QHNT	Qualifier History Note
QINDX	Qualifier Indexer Note
QLF	MeSH Qualifier (subheading)
QNOTE	Qualifier Scope Note
QONTE	Qualifier Online Note
QUF	Qualifier Use For
QUSE	Qualifier Use
RT	Related Term
UF	Used For Term
USE	Used Term

## Organism (/ORGN) Thesaurus

All Relationship Codes can be used with both the SEARCH and EXPAND command in the Organism (/ORGN, /BC) thesaurus.

Code	Content	Example
ALL	All associated terms (BT, SELF, NOTE, UF, USE, NT, RT, BC)	E RODENTIA+ALL/ORGN
AUTO (1)	Automatic Relationship Code (SELF, USE)	E PISCES+AUTO/ORGN
BT	Broader Terms (BT, SELF)	E BOVIDAE+BT/ORGN
HIE	Hierarchy (BT, SELF, NT)	E PISCES+HIE/ORGN
KT	Keyword Terms (multiword phrases containing the term) (SELF, KT)	E VIRUSES+KT/ORGN
NOTE	Notes associated with the term (SELF, NOTE)	E PROTOZOA+NOTE/ORGN
NT	Narrower Terms (SELF, NT)	E AMPHIBIA+NT/ORGN
PFT	Preferred and Forbidden Terms (SELF, UF, USE)	E PISCES+PFT/ORGN
RT	Related Terms (SELF, RT)	E RODENTS+RT/ORGN
STD	Standard (Broader, Narrower, and Related Terms) (BT, SELF, NT, RT)	E AVES+STD/ORGN
UF	Used For Terms (Forbidden Terms) (SELF, UF)	E SALIENTIA+UF/ORGN
USE	Used Terms (Preferred Terms) (SELF, USE)	E BC25990+USE/ORGN

- (1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /ORGN Thesaurus

Code	Description
→	Self
BC	Biosystematic Codes
BT	Broader Term
KT	Keyword Term
NOTE	Scope Note
NT	Narrower Term
RT	Related Term
UF	Used For Term
USE	Used Term

## Supplementary Term (/ST) Thesaurus

All Relationship Codes can be used with both the SEARCH and EXPAND command in the Controlled Term (/ST) thesaurus.

Code	Content	Example
ALL	All associated terms (BT, SELF, NOTE, UF, USE, NT, RT)	E ANIMAL HUSBANDRY+ALL/ST
AUTO (1)	Automatic Relationship Code (SELF, USE)	E PEDIATRICS+AUTO/ST
BT	Broader Terms (BT, SELF)	E ALLERGY+BT/ST
HIE	Hierarchy (Broader and Narrower Terms) (BT, SELF, NT)	E HUMAN MEDICINE+HIE/ST
KT	Keyword Terms (multiword phrases containing the term) (SELF, KT)	E DENTAL+KT/ST
NOTE	Scope Notes (SELF, NOTE)	E DENTAL MEDICINE+NOTE/ST
NT	Narrower Terms (SELF, NT)	E AGRICULTURE+NT/ST
PFT	Preferred and Forbidden Terms (SELF, UF, USE)	E ANTIVIRAL – DRUG+PFT/ST
RT	Related Terms (SELF, RT)	E TOXICOLOGY+RT/ST
STD	Standard (Broader, Narrower, and Related Terms) (BT, SELF, NT, RT)	E CLINICAL IMMUNOLOGY+STD/ST
UF	Used For Terms (Forbidden Terms) (SELF, UF)	E ANTIVIRAL – DRUG+UF/ST
USE	Used Terms (Preferred Terms) (SELF, USE)	E ANTIVIRAL+USE/ST

- (1) Automatic relationship is SET OFF. When SET REL is ON, the result of EXPAND without any relationship code is the same as described for AUTO.

## Field Descriptors for the /ST Thesaurus

Code	Description
→	Self
BT	Broader Term
KT	Keyword Term
NOTE	Scope Note
NT	Narrower Term
RT	Related Term
UF	Used For Term
USE	Used Term

## DISPLAY and PRINT Formats

Any combination of formats may be used to display or print answers. Multiple codes must be separated by spaces or commas, e.g., D L1 1-5 CN CO. The fields are displayed or printed in the order requested.

Hit-term highlighting is available in all fields except CM, CP, ED, MY, PC, and PY. Highlighting must be ON during SEARCH to use the HIT, KWIC, and OCC display formats.

Format	Content	Examples
AB (1)	Abstract	D AB
AN	Accession Number	D AN FS
AU (IN) (2)	Author or Inventor	D AU 1-5
AUCL	Collaborator	D AUCL
AUGR	Group Author	D AUGR
BC (3)	Biosystematic Code	D BC
CC (4)	Classification Code (Concept Code)	D CC 1 8
CM (5)	Comment	D CM SO
CN (6)	Chemical Name	D RN CN
CP (4)	Copyright Notice	D CP
CS (7)	Corporate Source (includes ZP)	D CS
CSS (8)	Supporting Organization (Sponsoring Agency)	D CSS
CT (9)	Controlled Term	D CT
CY (10)	Country of Publication	D CY
DB (11)	Duration Begin, Date (Initial Project Date)	D DB DE
DE (11)	Duration End, Date (Final Project Date)	D DB DE
DN	Document Number	D DN
DOI (FTDOI) (29)	Digital Object Identifier	D DOI or D FTDOI
DT (TC) (12)	Document Type	D DT
ED (UP)	Entry and Last Updated Dates	D 1 3 5 ED
EML (9,13)	E-mail Address	D EML AU
FS	File Segment	D FS 2,7
GEN (10)	Gene Name	D GEN
GT (3)	Geographic Term	D GT
ISN (13,14)	International Standard (Document) Number (CODEN, ISBN, ISSN)	D ISN
JT (13,14)	Journal Title	D JT
JTA (13,15)	Journal Title, Abbreviated	D JTA
JTF (13,16)	Journal Title, Full	D JTF
LA (17)	Language	D LA
MD (3,13)	Meeting Date	D MD
ML (3,13)	Meeting Location	D ML
MO (3,13)	Meeting Organizer	D MO
MT (3,13)	Meeting Title	D MT
MY (3,13)	Meeting Year	D MY
NA (5)	Named Person or Institution	D NA
NC (8)	Number of Contract	D NC
NCT	Clinical Trial Numbers	D NCT
NR (5,13)	Number of Report	D NR
ON (18)	Order Number	D ON
ORGN (3)	Organism	D ORGN
OS (19)	Other Source	D OS
PC (PCS) (13,20)	Patent Country	D PC
PD (13,14)	Publication Date	D PD
PI (20, 28) (PN, PATS)	Patent Information	D PI
PNK	Patent Number/Kind Code	D PNK
PY (13,14)	Publication Year	D PY
RN (21)	CAS Registry Number and Chemical Name	D RN
SC (22)	Section	D SC
SL (23)	Summary Language	D SL
SO	Source	D TI SO
ST (24)	Supplementary Term	D ST
TI (25)	Title	D TI
URL (3,13)	Uniform Resource Locator	D URL
ZP (26)	Zip Code	D ZP

**DISPLAY and PRINT Formats (cont'd)**

Format	Content	Examples
ABS ALL	AN, CP, AB AN, CP, DN, TI, AU, CS, CSS, ON, NC, PI, SO, CM, CY, DT, FS, NCT, OS, LA, SL, ED, DB, DE, AB, SC, CC, BC, CT, ST, NA, GT, ORGN, RN, CN, GEN, DOI	D ABS D ALL
AUTH BIB	AUY, AUCL, AUGR AN, CP, DN, TI, AU, CS, CSS, ON, NC, PI, SO, CY, DT, FS, NCT, OS, LA, SL, ED, DB, DE, DOI (BIB is the default)	D AUTH D L2 3 BIB
CBIB DALL IABS IALL IBIB IND SCAN (25,27)	AN, DN, compressed bibliographic information ALL, delimited for post processing ABS, indented with text labels ALL, indented with text labels BIB, indented with text labels AN, CP, SC, CC, BC, CT, ST, NA, GT, ORGN, RN, CN, GEN TI, CM, CN (random display without answer numbers)	D CBIB D DALL D IABS D IALL D IBIB D IND D SCAN
HIT HITIND KWIC OCC (25)	Fields containing hit terms IND Hit terms with 20 words on either side (Keyword in Context) Number of occurrences of hit terms and fields in which they occur	D HIT D HITIND D KWIC NOH D OCC 1-6

- (1) Available in all file segments except ANEUPL and ETIC.
- (2) Available in all file segments except TSCATS.
- (3) Available in BIOSIS file segment only.
- (4) Available in BIOSIS, CAPLUS, CIS, IPA, and TSCATS file segments only.
- (5) Available in MEDLINE file segment only.
- (6) Available in BIOSIS, CAPLUS, DART, EMIC, ETIC, IPA, and MEDLINE file segments.
- (7) Available in BIOSIS, CAPLUS, CIS, CRISP, DART, EMIC, FEDRIP, IPA, MEDLINE, PESTAB, AND TSCATS file segments.
- (8) Available for CRISP, EPIDEM, FEDRIP file segments only.
- (9) Available in DART, EMIC, and MEDLINE file segment only.
- (10) Available in CAPLUS and MEDLINE file segments only.
- (11) Available in FEDRIP file segment only.
- (12) Available in ANEUPL, BIOSIS, CAPLUS, CIS, DART, EMIC, ETIC, IPA, MEDLINE and PESTAB file segments.
- (13) Custom display only.
- (14) Available in all file segments except CRISP, FEDRIP, and TSCATS.
- (15) Available in ANEUPL, CAPLUS, DART, EMIC, EPIDEM, ETIC, HAPAB, HMTTC, IPA, MEDLINE, PESTAB, PPBIB, and RISKLINE file segments.
- (16) Available in BIOSIS, CAPLUS, DART, ETIC, IPA, MEDLINE, and PESTAB file segments.
- (17) Available in all file segments except EPIDEM, PPBIB, and TSCATS.
- (18) Available in TSCATS file segment only.
- (19) Available in BIOSIS, CAPLUS, EMIC, FEDRIP, IPA, and MEDLINE file segments only.
- (20) Available in BIOSIS and CAPLUS file segments only.
- (21) Available in all file segments except CRISP and FEDRIP.
- (22) Available in IPA file segment only.
- (23) Available in BIOSIS and IPA file segments only.
- (24) Available in ANEUPL, BIOSIS, CAPLUS, CIS, CRISP, EMIC, EPIDEM, ETIC, FEDRIP, HMTTC, IPA, MEDLINE, PPBIB, RISKLINE, and TSCATS file segments.
- (25) No online display fee for this format.
- (26) Available for CRISP and FEDRIP file segments only.
- (27) SCAN must be specified on the command line, i.e., D SCAN or DISPLAY SCAN.
- (28) Patent numbers are available in STN and Derwent format. The format for DISPLAY, PRINT, SELECT, and SORT is set using the SET PATENT command. STN is the default format. Enter SET PAT DERWENT to change to the Derwent format. To reset to the STN format, enter SET PAT STN.
- (29) Available in BIOSIS and MEDLINE file segments only.

**SELECT, ANALYZE, and SORT Fields**

The SELECT command is used to create E-numbers containing terms taken from the specified field in an answer set.

The ANALYZE command is used to create an L-number containing terms taken from the specified field in an answer set.

The SORT command is used to rearrange the search results in either alphabetic or numeric order of the specified field(s).

Field Name	Field Code	ANALYZE/ SELECT (1)	SORT
Abstract (2)	AB	Y (3)	N
Accession Number	AN	Y	N
Author (Inventor) (4)	AU	Y	Y
Author Group	AUTH	Y	Y
Biosystematic Code (5)	BC	Y	N
CAS Registry Number (6)	RN	Y (3)	N
Chemical Name (7)	CN	Y (8)	N
	NAME	Y (3,8)	N
Chemical Name and CAS Registry Number	CHEM	Y (3)	N
Citation	CIT	Y (8,9)	N
Classification Code (10) (Concept Code)	CC	Y	N
Clinical Trial Numbers	NCT	Y	Y
CODEN	CODEN	N	Y
Collaborator	AUCL	Y	Y
Comment	CM	Y (35)	N
Controlled Term (10)	CT	Y	N
Corporate Source (12) (Patent Assignee)	CS	Y	Y
Country Of Publication (13)	CY	Y	Y
Document Number	DN	Y	Y
Document Type (14)	DT	Y	Y
Duration Begin, Date (15) (Initial Project Date)	DB	Y	Y
Duration End, Date (15) (Final Project Date)	DE	Y	Y
E-mail Address (11)	EML	Y	Y
File Segment	FS	Y	Y
GenBank Number (16)	GENBANK	Y (3)	N
	GBN	Y (3)	N
Gene Name	GEN	Y	Y
Group Author	AUGR	Y	Y
Geographic Term (5)	GT	Y (8)	Y
International Standard Book Number	ISBN	N	Y
International Standard (Document) Number (17)	ISN	Y (18)	N
International Standard Serial Number	ISSN	N	Y
Inventor (19)	IN	Y	Y
Journal Title, Abbreviated and Full (17)	JT	Y	Y
Journal Title, Abbreviated (20)	JTA	Y (21)	Y
Journal Title, Full (22)	JTF	Y (21)	Y
Journal Title Code (17)	JTC	N	Y
Language (23)	LA	Y	Y
Meeting Date (5)	MD	Y	Y
Meeting Location (5)	ML	Y	Y
Meeting Organizer (5)	MO	Y	Y
Meeting Title (5)	MT	Y (8)	Y
Meeting Year (5)	MY	Y (8)	Y
Named Person or Institution (24)	NA	Y	N
Number of Contract (25)	NC	Y	Y
Number of Report (26)	NR	Y	Y
Occurrence Count of Hit Terms	OCC	N	Y
Organism (5)	ORGN	Y	N
Order Number (27)	ON	Y	Y
Other Source (28)	OS	Y	Y

**SELECT, ANALYZE, and SORT Fields (cont'd)**

Field Name	Field Code	ANALYZE/ SELECT (1)	SORT
Patent Country (19)	PC	Y (8)	Y
	PCS	Y (8)	Y
Patent Information (19)	PI	Y (29)	Y
Patent Number (19)	PN	Y	Y
	PATS	Y	Y
Patent Number/Kind Code	PNK	Y	N
Publication Date (18)	PD	Y (8)	Y
Publication Year (18)	PY	Y (8)	Y
Section (20)	SC	Y	Y
Source	SO	Y (31)	N
Summary Language (32)	SL	Y	Y
Supplementary Term (33)	ST	Y	N
Supporting Organization (25) (Sponsoring Agency)	CSS	Y	Y
Title	TI	Y (default)	Y
Treatment Code (14)	TC	Y (34)	Y
Uniform Resource Locator (5)	URL	Y	Y
Zip Code (6)	ZP	Y	Y

- (1) HIT may be used to restrict terms extracted to terms that match the search expression used to create the answer set, e.g., SEL HIT TI.
- (2) Available in all file segments except ANEUPL and ETIC.
- (3) Appends /BI to the terms created by SELECT.
- (4) Available in all file segments except TSCATS.
- (5) Available for the BIOSIS file segment only.
- (6) Available in all file segments except CRISP and FEDRIP.
- (7) Available in BIOSIS, CAPLUS, DART, EMIC, ETIC, IPA, and MEDLINE file segments.
- (8) SELECT HIT and ANALYZE HIT are not valid with this field.
- (9) Extracts first author, publication year, volume, and first page with a truncation symbol appended and with /RE appended to the terms created by SELECT.
- (10) Available in BIOSIS, CAPLUS, CIS, IPA and TSCATS file segments only.
- (11) Available in DART, EMIC, and MEDLINE file segment only.
- (12) Available in BIOSIS, CAPLUS, CIS, CRISP, DART, EMIC, FEDRIP, IPA, MEDLINE, PESTAB, AND TSCATS file segments.
- (13) Available in CAPLUS and MEDLINE file segments only.
- (14) Available in ANEUPL, BIOSIS, CAPLUS, CIS, DART, EMIC, ETIC, IPA, MEDLINE and PESTAB file segments.
- (15) Available in FEDRIP file segment only.
- (16) Available in BIOSIS and MEDLINE file segments only.
- (17) Available in all file segments except CRISP, FEDRIP, and TSCATS.
- (18) Selects or analyzes CODEN, ISSN, and ISBN with /ISN appended to the terms created by SELECT.
- (19) Available for BIOSIS and CAPLUS file segments only.
- (20) Available in ANEUPL, CAPLUS, DART, EMIC, EPIDEM, ETIC, HAPAB, HMTTC, IPA, MEDLINE, PESTAB, PPBIB, and RISKLINE file segments.
- (21) Appends /JT to the terms created by SELECT.
- (22) Available in BIOSIS, CAPLUS, DART, ETIC, IPA, MEDLINE, and PESTAB file segments.
- (23) Available in all file segments except EPIDEM, PPBIB, and TSCATS.
- (24) Available in MEDLINE file segment only.
- (25) Available for CRISP, EPIDEM, FEDRIP file segments only
- (26) Available in MEDLINE file segment only.
- (27) Available in TSCATS file segment only.
- (28) Available in BIOSIS, CAPLUS, EMIC, FEDRIP, IPA, and MEDLINE file segments only.
- (29) Selects or analyzes the patent number and appends /PN to the terms created by SELECT.
- (30) Available in IPA file segment only.
- (31) Selects or analyzes CODEN, ISSN, and ISBN with /SO appended to the terms created by SELECT.
- (32) Available in BIOSIS and IPA file segments only.
- (33) Available in ANEUPL, BIOSIS, CAPLUS, CIS, CRISP, EMIC, EPIDEM, ETIC, FEDRIP, HMTTC, IPA, MEDLINE, PPBIB, RISKLINE, and TSCATS file segments.
- (34) Appends /DT to the terms created by SELECT.
- (35) Selects or analyzes the PMID values with /DN appended.

## Sample Records

### DISPLAY IALL (BIOSIS File Segment)

ACCESSION NUMBER: 2011:95669 TOXCENTER  
COPYRIGHT: Copyright (c) 2011 The Thomson Corporation  
DOCUMENT NUMBER: PREV201100132307  
TITLE: Methods for Genotyping Verotoxin-Producing Escherichia coli  
AUTHOR(S): Karama, M.; Gyles, C. L. [Reprint Author]  
CORPORATE SOURCE: Univ Guelph, Dept Pathobiol, Guelph, ON N1G 2W1, Canada  
cgyles@uoguelph.ca  
SOURCE: Zoonoses and Public Health, (OCT 2010) Vol. 57, No. 7-8,  
pp. 447-462.  
ISSN: 1863-1959.  
DIGITAL OBJECT IDENTIFIER: 10.1111/j.1863-2378.2009.01259.x  
DOCUMENT TYPE: Article  
General Review; (Literature Review)  
FILE SEGMENT: BIOSIS  
OTHER SOURCE: BIOSIS 2011:132307  
LANGUAGE: English  
ENTRY DATE: Entered STN: 15 Mar 2011  
Last Updated on STN: 15 Mar 2011

#### ABSTRACT:

Verotoxin-producing Escherichia coli (VTEC) is annually incriminated in more than 100 000 cases of enteric foodborne human disease and in losses amounting to \$US 2.5 billion every year. A number of genotyping methods have been developed to track VTEC infections and determine diversity and evolutionary relationships among these microorganisms. These methods have facilitated monitoring and surveillance of foodborne VTEC outbreaks and early identification of outbreaks or clusters of outbreaks. Pulsed-field gel electrophoresis (PFGE) has been used extensively to track and differentiate VTEC because of its high discriminatory power, reproducibility and ease of standardization. Multiple-locus variable-number tandem-repeats analysis (MLVA) and microarrays are the latest genotyping methods that have been applied to discriminate VTEC. MLVA, a simpler and less expensive method, is proving to have a discriminatory power comparable to that of PFGE. Microarrays are successfully being applied to differentiate VTEC and make inferences on genome diversification. Novel methods that are being evaluated for subtyping VTEC include the detection of single nucleotide polymorphisms and optical mapping. This review discusses the principles, applications, advantages and disadvantages of genotyping methods that have been used to differentiate VTEC strains. These methods have been mainly used to differentiate strains of O157:H7 VTEC and to a lesser extent non-O157 VTEC.

CLASSIFICATION CODE: Pathology - General 12502  
Toxicology - General and methods 22501  
Physiology and biochemistry of bacteria 31000  
Medical and clinical microbiology - Bacteriology 36002

SUPPLEMENTARY TERMS: Major Concepts  
Toxicology; Infection; Methods and Techniques; Human  
Medicine (Medical Sciences)  
SUPPLEMENTARY TERMS: Diseases  
foodborne infection: bacterial disease, toxicity  
SUPPLEMENTARY TERMS: Chemicals & Biochemicals  
verotoxin: toxin  
SUPPLEMENTARY TERMS: Methods & Equipment  
genotyping: laboratory techniques, genetic techniques;  
microarray: laboratory techniques, genetic techniques;  
optical mapping: laboratory techniques, genetic  
techniques; pulsed-field gel electrophoresis:  
electrophoretic techniques, laboratory techniques;  
Multiple-locus variable-number tandem-repeats analysis:  
laboratory techniques, genetic techniques  
SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
single nucleotide polymorphism

**TOXCENTER****DISPLAY IALL (BIOSIS File Segment) (cont'd)**

ORGANISM: Classifier  
 Enterobacteriaceae 06702  
 Super Taxa  
 Facultatively Anaerobic Gram-Negative Rods; Eubacteria;  
 Bacteria; Microorganisms  
 Organism Name  
 Escherichia coli (species): pathogen, serovar-0157:H7  
 Taxa Notes  
 Bacteria, Eubacteria, Microorganisms

ORGANISM: Classifier  
 Hominidae 86215  
 Super Taxa  
 Primates; Mammalia; Vertebrata; Chordata; Animalia  
 Organism Name  
 human (common): host  
 Taxa Notes  
 Animals, Chordates, Humans, Mammals, Primates,  
 Vertebrates

**DISPLAY IALL (CAPLUS File Segment)**

ACCESSION NUMBER: 2001:187664 TOXCENTER  
 COPYRIGHT: Copyright 2011 ACS  
 DOCUMENT NUMBER: CA13519271902Z  
 TITLE: Methods for assessing complement activation  
 AUTHOR(S): Hugli, Tony E.; Stoughton, Roland B.  
 CORPORATE SOURCE: ASSIGNEE: The Scripps Research Institute  
 PATENT INFORMATION: US 6297024 B1 2 Oct 2001  
 SOURCE: (2001) U.S., 28 pp., Cont.-in-part of U.S. Ser. No.  
 173,579.  
 CODEN: USXXAM.  
 COUNTRY: UNITED STATES  
 DOCUMENT TYPE: Patent  
 FILE SEGMENT: CAPLUS  
 OTHER SOURCE: CAPLUS 2001:719020  
 LANGUAGE: English  
 ENTRY DATE: Entered STN: 16 Nov 2001  
 Last Updated on STN: 19 Mar 2002

**ABSTRACT:**

Methods for measuring in vivo activation of the lectin pathway by measuring mannan-binding serine protease activity (MASP) are provided. The methods are accomplished by C3a and C4a levels in in vitro activated EDTA plasma. In particular, the increase in C3a and/or C4a as a function of time is an indicator of the amount of activated MASP in EDTA plasma. Methods are also provided for measuring the alternate and classical pathways of complement activation, exclusive of the lectin pathway, and thereby disorders associated therewith. To perform such measurements, Futhan or other serine protease inhibitor is added to blood or plasma, containing a divalent metal ion chelator, and C3a and C4a are measured.

CLASSIFICATION CODE: 15-4

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
 complement activation MASP1 MASP2 lectin pathway; C3a C4a  
 C5a chelator drug screening; transplant rejection  
 autoimmune disease complement activation

REGISTRY NUMBER: 80295-42-7 (complement C3a)  
 80295-49-4 (complement C4a)  
 80295-54-1 (complement C5a)  
 172306-56-8 (MASP serine protease)  
 214915-11-4 (MASP-1 serine protease)  
 214915-16-9 (MASP-2 serine protease)  
 60-00-4 (EDTA)  
 37259-58-8 (Serine protease)  
 14127-61-8 (Calcium ion)

REGISTRY NUMBER: 82956-11-4; 77-92-9

**DISPLAY IALL (IPA File Segment)**

ACCESSION NUMBER: 2001:153071 TOXCENTER  
COPYRIGHT: Copyright 2001 ASHP  
DOCUMENT NUMBER: 38-10342  
TITLE: Gender gap narrows in smoking prevalence, deaths  
AUTHOR(S): anon  
SOURCE: American Journal of Health-System Pharmacy, (May 15  
2001) Vol. 58, pp. 852, 854.  
CODEN: AHSPEK. ISSN: 1079-2082.  
DOCUMENT TYPE: Journal  
FILE SEGMENT: IPA  
OTHER SOURCE: IPA 2001:10342  
LANGUAGE: English  
ENTRY DATE: Entered STN: 16 Nov 2001  
Last Updated on STN: 16 Nov 2001

**ABSTRACT:**

It was announced that the gap in smoking prevalence between men and women has closed to less than 5 percentage points; women now account for 39% of all smoking related deaths, a proportion that has more than doubled since 1965, according to the latest U.S. Surgeon General's report on smoking. Women who smoke not only share the same health risk as men but are also faced with health consequences that are unique to women, including pregnancy complications, problems with menstrual function, and cervical cancer. The report also states that nicotine addiction appears generally similar between the sexes, however, women's regulation of nicotine intake may be less precise than men's.

Elvira deC. Weiss

SECTION CODE: 22 Sociology, Economics and Ethics; 4 Toxicity  
CLASSIFICATION CODE: 12:00 Autonomic drugs  
SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
Nicotine; smoking; women  
Autonomic drugs; nicotine; women  
Smoking; cigarettes; women  
Cigarettes; smoking; women  
Women; smoking; cigarettes  
Toxicity; nicotine; women  
Sex; consumers; smoking  
Death; cigarettes; smoking  
Pregnancy; cigarettes; smoking  
Cervix neoplasms; cigarettes; smoking  
REGISTRY NUMBER: 54-11-5 (Nicotine)

**DISPLAY IALL (MEDLINE File Segment)**

ACCESSION NUMBER: 2011:94562 TOXCENTER [Full-text](#)  
COPYRIGHT: .COPYRGT. 2010 Japanese Cancer Association.  
DOCUMENT NUMBER: PubMed ID: 21175992  
TITLE: Phase II study of erlotinib plus gemcitabine in Japanese patients with unresectable pancreatic cancer  
AUTHOR(S): Okusaka Takuji; Furuse Junji; Funakoshi Akihiro; Ioka Tatsuya; Yamao Kenji; Ohkawa Shinichi; Boku Narikazu; Komatsu Yoshito; Nakamori Shoji; Iguchi Haruo; Ito Tetsuhide; Nakagawa Kazuhiko; Nakachi Kohei  
CORPORATE SOURCE: National Cancer Center Hospital, Tokyo, Japan  
tokusaka@ncc.go.jp  
SOURCE: Cancer science, (2011 Feb) Vol. 102, No. 2, pp. 425-31.  
Electronic Publication: 2010-12-22.  
Journal code: 101168776. E-ISSN: 1349-7006. L-ISSN: 1347-9032.  
DIGITAL OBJECT IDENTIFIER: 10.1111/j.1349-7006.2010.01810.x  
COUNTRY: England: United Kingdom

## TOXCENTER

## DISPLAY IALL (MEDLINE File Segment) (cont'd)

DOCUMENT TYPE: (CLINICAL TRIAL, PHASE II)  
Journal; Article; (JOURNAL ARTICLE)  
(MULTICENTER STUDY)  
(RESEARCH SUPPORT, NON-U.S. GOV'T)  
(CLINICAL TRIAL)

FILE SEGMENT: MEDLINE  
OTHER SOURCE: MEDLINE 2011195265  
LANGUAGE: English  
ENTRY DATE: Entered STN: 15 Mar 2011  
Last Updated on STN: 15 Mar 2011

## ABSTRACT:

Erlotinib combined with gemcitabine has not been evaluated in Japanese patients with unresectable pancreatic cancer. This two-step phase II study assessed the safety and pharmacokinetics of erlotinib 100 mg/day (oral) plus gemcitabine 1000 mg/m<sup>2</sup> (i.v. days 1, 8, 15) in a 28-day cycle in the first step, and efficacy and safety in the second step. The primary end-point was safety. One hundred and seven patients were enrolled (first step, n = 6; second step, n = 101). The most common adverse event was RASH (compiled using the preferred terms rash, acne, exfoliative rash, dermatitis acneiform, erythema, eczema, dermatitis and pustular rash) in 93.4% of patients. One treatment-related death occurred. While interstitial lung disease-like events were reported in nine patients (8.5%; grade 1/2/3, 3.8/2.8/1.9%), all patients recovered or improved. The median overall survival, the 1-year survival rate and median progression-free survival were 9.23 months, 33.0% and 3.48 months, respectively. The overall response and disease control rates were 20.3% and 50.0%, respectively. In Japanese patients with unresectable pancreatic cancer, erlotinib plus gemcitabine had acceptable toxicity and efficacy that was not inferior to that seen in Western patients.

CONTROLLED TERM: Check Tags: Female; Male  
\*Adenocarcinoma: DT, drug therapy  
Adult  
Aged  
Antineoplastic Combined Chemotherapy Protocols: PK,  
pharmacokinetics  
\*Antineoplastic Combined Chemotherapy Protocols: TU,  
therapeutic use  
Asian Continental Ancestry Group  
\*Carcinoma, Adenosquamous: DT, drug therapy  
Deoxycytidine: AD, administration & dosage  
Deoxycytidine: AE, adverse effects  
Deoxycytidine: AA, analogs & derivatives  
Deoxycytidine: PK, pharmacokinetics  
Humans  
Kaplan-Meier Estimate  
Middle Aged  
\*Pancreatic Neoplasms: DT, drug therapy  
Quinazolines: AD, administration & dosage  
Quinazolines: AE, adverse effects  
Quinazolines: PK, pharmacokinetics  
REGISTRY NUMBER: 103882-84-4 (gemcitabine)  
951-77-9 (Deoxycytidine)  
CHEMICAL NAME: Quinazolines; erlotinib

**DISPLAY IALL (ANEUPL File Segment)**

ACCESSION NUMBER: 2002:348890 TOXCENTER  
DOCUMENT NUMBER: ANEUPL-83-001787  
TITLE: MICROTUBULE ORGANIZING CENTERS DURING THE CELL CYCLE OF  
3T3 CELLS.  
AUTHOR(S): BROOKS R F; RICHMOND F N  
SOURCE: Journal of Cell Science, (1983) 61 231.  
CODEN: JNCSA.  
FILE SEGMENT: ANEUPL  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002  
SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
AGENT TEST RESULTS; MAMMAL MOUSE CELL CULTURE; COLCEMID  
REGISTRY NUMBER: 477-30-5

**DISPLAY IALL (CIS File Segment)**

ACCESSION NUMBER: 2002:523046 TOXCENTER  
DOCUMENT NUMBER: CIS-01-01633  
TITLE: Toxicological profile for alpha-, beta-, gamma- and  
delta-hexachlorocyclohexane (Update)  
AUTHOR(S): Anonymus  
CORPORATE SOURCE: Agency for Toxic Substances and Diseases Registry (ATSDR)  
SOURCE: U.S. Department of Health and Human Services, Public  
Health Service, Agency for Toxic Substances and Disease  
Registry, Division of Toxicology/Toxicology Information  
Branch, 1600 Clifton Road NE, E-29, Atlanta, GA 30333,  
USA, July 1999. xix, 273p. Illus. Approx. 860 ref..  
DOCUMENT TYPE: Book; (Monograph)  
FILE SEGMENT: CIS  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002  
ABSTRACT:  
This profile was prepared in accordance with guidelines set by the US Agency  
for Toxic Substances and Disease Registry and the EPA. The key literature  
related to the toxic effects of hexachlorocyclohexane is identified and  
reviewed. Contents: public health statement; health effects; chemical and  
physical information; production, import, use and disposal; potential for human  
exposure; analytical methods; regulations and advisories; glossary. Health  
hazards include: neurologic effects (dizziness, seizures); haematological  
changes; effects on the immune system; liver and kidney damage; changes in  
reproductive hormone levels in the blood; animal studies show reproductive  
effects, genotoxicity and cancerogenicity (liver cancer). (Update of CIS  
99-234).  
CLASSIFICATION CODE: 120  
SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
HEXACHLOROCYCLOHEXANE; TOXICOLOGY; TOXICITY EVALUATION;  
HEALTH HAZARDS; USA; CRITERIA DOCUMENT; TOXIC EFFECTS;  
LITERATURE SURVEY; EXPOSURE EVALUATION; LIMITATION OF  
EXPOSURE; GLOSSARY; SKIN ABSORPTION; NEUROLOGICAL EFFECTS;  
CARCINOGENIC EFFECTS; HAEMATOLOGICAL EFFECTS; IMMUNOTOXIC  
EFFECTS; MUTAGENIC EFFECTS; ANTIFERTILITY EFFECTS; HEPATIC  
DAMAGE; RENAL DAMAGE; DETERMINATION IN BIOLOGICAL MATTER;  
LEGISLATION  
REGISTRY NUMBER: 319-84-6; 319-85-7; 319-86-8; 58-89-9; 608-73-1

## TOXCENTER

## DISPLAY IALL (CRISP File Segment)

ACCESSION NUMBER: 2002:562111 TOXCENTER  
DOCUMENT NUMBER: CRISP-2000-TW00927-03  
TITLE: TAXOL EFFECTS ON SIGNALING PATHWAYS IN OVARIAN CELLS  
AUTHOR(S): HORWITZ S B  
CORPORATE SOURCE: ALBERT EINSTEIN COLL OF MED, 1300 MORRIS PARK AVENUE,  
BRONX, NY 10461:NEW YORK  
SUPPORTING ORGANIZATION (SPONSORING AGENCY): U.S. DEPT. OF HEALTH AND HUMAN  
SERVICES; PUBLIC HEALTH SERVICE; NATIONAL INSTITUTES OF  
HEALTH, FOGARTY INTERNATIONAL CENTER  
SOURCE: Crisp Data Base National Institutes of Health.  
DOCUMENT TYPE: (Research)  
FILE SEGMENT: CRISP  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

## ABSTRACT:

DESCRIPTION Taxol is a natural product which has significant antitumor activity in a large number of human tumors. It has recently become appreciated that in addition to stabilizing microtubules and blocking cell cycle progress, taxol can alter gene expression by interacting with the MAP-kinase signal transduction pathway. Previous studies from the foreign and US PIs had demonstrated that tyrosine phosphorylation of the adaptor protein, Shc, and formation of Shc/grb2/SOS complexes occurs in a taxol-treated murine macrophage cell line. The investigators propose to extend these observations to examine human ovarian carcinoma cells. The specific aims are to: 1. Study the effect of taxol on activation of the MAP-kinase pathway in human ovarian carcinoma cell lines; 2. Study the effect of taxotere and other taxol analogs compared to that of taxol on the activation of the Ras/MAPK signaling pathway and the relationship of this activity to effects on microtubule stabilization and cytotoxicity; 3. Compare taxol and taxotere in MAPK signaling pathway in taxol-sensitive and taxol-resistant human ovarian carcinoma cells.

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
paclitaxel; antineoplastic; biological signal  
transduction; microtubule; chemical structure function;  
pharmacokinetics; drug adverse effect; drug resistance;  
drug screening ,evaluation; gene expression; human tissue;  
immunoprecipitation; phosphorylation; electrophoresis;  
tissue ,cell culture; cytotoxicity; analog; apoptosis;  
enzyme activity; mitogen activated protein kinase; cell  
proliferation

## DISPLAY IALL (DART File Segment)

ACCESSION NUMBER: 2002:582299 TOXCENTER  
DOCUMENT NUMBER: DART-TER-2001532  
TITLE: Basic fibroblast growth factor (bFGF) induces branching  
and dilatation of developing renal tubules leading to cyst  
formation.  
AUTHOR(S): Li Z; Liu X H; Xu L; Jerebstova M; Ye X; Ray P E  
CORPORATE SOURCE: Centers for Molecular Physiology and Genetic Medicine,  
Children's Research Institute, Children's National Medical  
Center, Washington, DC.  
SOURCE: PEDIATR Res 2002 Apr;51(4 Pt 2):435A. PEDIATRIC RESEARCH,  
ISSN: 0031-3998.  
DOCUMENT TYPE: Abstract; (MEETING ABSTRACT)  
FILE SEGMENT: DART  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

**DISPLAY IALL (DART File Segment) (cont'd)****ABSTRACT:**

**BACKGROUND:** Increased expression of bFGF has been detected in dysplastic kidneys. However, the role of bFGF and the signaling pathways modulating its activity in developing kidneys are not completely understood. **OBJECTIVE:** To determine whether bFGF modulates the growth of rat metanephric kidneys (MK) and define the signaling pathways involved in this process. **DESIGN/METHODS:** MK (n = 80) from 15 day old rat embryos were grown in serum free DMEM-F12 media in the presence or absence of bFGF (5-20 ng/mL), bFGF neutralizing antibodies, 8-Bromo-cAMP (10(-3) M) and Isoproterenol (10(-5) M). Immunohistochemistry and in situ hybridization studies were done using the following markers of cell differentiation: WT-1, Pax, Factor VIII, BF-2; alpha-smooth muscle action, vimentin, Dolichos biflorus agglutinin. Cell apoptosis and proliferation were detected with the TUNEL assay, PCNA, (3H)-thymidine, and cell counts. The number and length of branching developing tubules were measured by morphometric analysis using a computerized program. Cultured primary stromal mesenchymal cells (SMC) were grown and treated as described above for the MK. MAPK kinase (MEK 1-2) and cAMP activity were determined by Western blots and ELISA respectively. To determine the effects of bFGF in vivo, MK were infected with rAd vectors carrying either a secreted 18 kD form of bFGF or the beta-galactosidase gene, and transplanted under the kidney capsule of adult rats (n = 6 each group). Results were evaluated by one-way analysis of variance using the Newman-Keuls test. P values of less than 0.05 were considered significant. **RESULTS:** Basic FGF increased the branching of developing tubules and induced the growth of peritubular mesenchymal stromal and endothelial cells. These changes were partially inhibited by 8-Bromo-cAMP and Isoproterenol. Basic FGF increased (5 folds) the number of SMC through a MEK 1-2-dependent signaling pathway. cAMP blocked these effects. Transplanted MK infected with rAd-bFGF, but not with rAd-beta-gal, showed increased renal growth, angiogenesis, tubulogenesis, and formation of cystic structures. **CONCLUSIONS:** Basic FGF induces renal growth and facilitates the formation of tubular cystic structures at least partially through a MEK 1-2-dependent signaling pathway that could be inhibited by cAMP, both in vitro and in vivo.

**CONTROLLED TERM:** Check Tags: Animal  
Rats  
\*Fibroblast Growth Factor 2: PH, PHYSIOLOGY  
\*Kidney Tubules: EM, EMBRYOLOGY  
\*Kidney Tubules: CY, CYTOLOGY  
\*Kidney Tubules: EN, ENZYMOLOGY  
Tissue Culture  
Cell Division  
Apoptosis  
Mitogen-Activated Protein Kinases: ME, METABOLISM  
Immunohistochemistry  
Enzyme-Linked Immunosorbent Assay  
Blotting, Western  
Signal Transduction

**REGISTRY NUMBER:** 103107-01-3 (Fibroblast growth factor 2)  
**CHEMICAL NAME:** EC 2.7.1.- (Mitogen-activated protein kinases)

## TOXCENTER

## DISPLAY IALL (EMIC File Segment)

ACCESSION NUMBER: 2002:453936 TOXCENTER  
DOCUMENT NUMBER: EMIC-110877  
TITLE: Impact of dimethyl sulfoxide and examples of combined genotoxicity in the SOS chromotest.  
AUTHOR(S): Gebel T; Koenig A  
CORPORATE SOURCE: Medical Institute of General Hygiene and Environmental Health, University of Goettingen, Windausweg 2, D-37073, Goettingen, Germany. tgebel@gwdg.de  
SOURCE: Mutation Research, (1999 Aug 18) 444 (2) 405-11.  
Journal Code: NNA. ISSN: 0027-5107.  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
FILE SEGMENT: EMIC  
OTHER SOURCE: EMIC MED-99453366  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

## ABSTRACT:

The bacterial SOS chromotest with Escherichia coli PQ37 was used for the assessment of genotoxicity of combined xenobiotic treatments. The modulation of test compound genotoxicity by dimethyl sulfoxide (DMSO), a common solvent for test compounds, was assessed as well. It was shown that DMSO modulated SOS chromotest genotoxicity of several xenobiotics: in comparison to test compound dissolution in water, the commonly used addition of 3.2% (v/v) DMSO as solvent lead to a significant increase in the genotoxicity of K(2)RhCl(5) and beta-propiolactone (BPL). However, the effects of cisplatin decreased significantly when DMSO was added. Thus, albeit DMSO is not genotoxic in this test itself, it can interfere with SOS chromotest responses. Further experiments were performed in the absence of DMSO. BPL and cisplatin in combination showed an over-additive synergism in SOS genotoxicity as well as K(2)RhCl(5) and cisplatin did. Addition of Pd(NH(3))(4)Cl(2) and NaAsO(2), which are non-genotoxic in the SOS chromotest, did not enhance the K(2)RhCl(5)- or BPL-mediated SOS sfiA induction. Nevertheless, at the highest subcytotoxic dose of NaAsO(2) tested (200 microm), a slight yet significant suppression of BPL-mediated SOS genotoxicity was observed. These results confirm that the SOS chromotest is a useful tool for the rapid evaluation of the combined genotoxicity of compound mixtures. However, the use of DMSO as test solvent has to be taken with caution.

CONTROLLED TERM: Cisplatin: TO, TOXICITY  
Colorimetry  
\*Dimethyl Sulfoxide: PD, PHARMACOLOGY  
\*Escherichia coli: DE, DRUG EFFECTS  
\*Escherichia coli: GE, GENETICS  
\*Mutagenicity Tests  
\*SOS Response (Genetics): DE, DRUG EFFECTS

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
Taxonomic Name: ESCHERICHIA COLI,PQ37  
Test Object: BACTERIA  
EFFECTS ON NUCLEIC ACIDS: DNA DAMAGE-SOS INDUCTION-SFIA  
MISCELLANEOUS CATEGORY: AGENT INTERACTIONS

REGISTRY NUMBER: 67-68-5 (DMSO)  
57-57-8 (BETA-PROPIOLACTONE)  
15663-27-1 (CISPLATIN)  
65980-75-8 (DIPOTASSIUM PENTACHLORORHODATE)  
13815-17-3 (TETRAAMMINEPALLADIUM(2+) DICHLORIDE)  
7784-46-5 (SODIUM ARSENITE)

**DISPLAY IALL (EPIDEM File Segment)**

ACCESSION NUMBER: 2002:352326 TOXCENTER  
DOCUMENT NUMBER: EPIDEM-008891  
TITLE: THE RELATION OF DIET, CIGARETTE SMOKING, AND ALCOHOL  
CONSUMPTION TO PLASMA BETA-CAROTENE AND ALPHA-TOCOPHEROL  
LEVELS.  
AUTHOR(S): STRYKER W S; KAPLAN L A; STEIN E A; STAMPFER M J; SOBER A;  
WILLETT W C  
SOURCE: AM. J. EPIDEMIOLOG., (1988) 127 (2) 283-296.  
DOCUMENT TYPE: Journal  
FILE SEGMENT: EPIDEM  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

**ABSTRACT:**

THE RELATION OF DIET AND NUTRITIONAL SUPPLEMENTS, CIGARETTE USE, ALCOHOL CONSUMPTION, AND BLOOD LIPIDS TO PLASMA LEVELS OF BETA-CAROTENE AND ALPHA-TOCOPHEROL WAS STUDIED AMONG 330 MEN AND WOMEN AGED 18-79 YEARS. DIETARY CAROTENE, PREFORMED VITAMIN A, AND VITAMIN E INTAKE WERE ESTIMATED BY A SELF-ADMINISTERED SEMIQUANTITATIVE FOOD FREQUENCY QUESTIONNAIRE. THE CORRELATION OF DIETARY CAROTENE WITH PLASMA BETA-CAROTENE WAS REDUCED IN SMOKERS COMPARED WITH NONSMOKERS (R = 0.02 VS. 0.44 AMONG MEN; R = 0.19 VS. 0.45 AMONG WOMEN). SMOKERS HAD MUCH LOWER PLASMA LEVELS OF BETA-CAROTENE THAN DID NONSMOKERS (GEOMETRIC MEAN 8.5 VS. 15.3 MICROGRAMS FOR MEN; 17.3 VS. 26.3 MICROGRAMS/DL FOR WOMEN) DESPITE ONLY SLIGHTLY LOWER INTAKES OF CAROTENOIDS. IN MULTIPLE REGRESSION ANALYSES, MEN WHO SMOKED ONE PACK PER DAY HAD 72% (95% CONFIDENCE INTERVAL (CI) 58-89) OF THE PLASMA BETA-CAROTENE LEVELS OF NONSMOKERS AFTER ACCOUNTING FOR DIETARY CAROTENE AND OTHER VARIABLES; FOR WOMEN, THE CORRESPONDING PERCENTAGE WAS 79% (CI 64-99). IN SIMILAR MODELS, MEN DRINKING 20 G OF ALCOHOL PER DAY HAD 76% (CI 65-88) OF THE BETA-CAROTENE LEVELS OF NONDRINKERS; WOMEN HAD 89% (CI 73-108) OF THE LEVELS OF NONDRINKERS. AN INTERACTION TERM FOR CAROTENE INTAKE AND SMOKING WAS STATISTICALLY SIGNIFICANT IN A MODEL COMBINING BOTH SEXES. THESE RESULTS SUGGEST THAT PLASMA LEVELS OF BETA-CAROTENE AMONG SMOKERS AND, PERHAPS, HEAVY CONSUMERS OF ALCOHOL MAY BE REDUCED SUBSTANTIALLY BELOW LEVELS DUE TO DIFFERENCES IN DIET. THE CORRELATION OF CALORIE-ADJUSTED INTAKE OF VITAMIN E WITH LIPID-ADJUSTED PLASMA LEVELS OF VITAMIN ALPHA-TOCOPHEROL WAS 0.53 FOR MEN (N = 137) AND 0.51 FOR WOMEN (N = 193) AND DID NOT DIFFER BY ALCOHOL CONSUMPTION AND CIGARETTE USE; THESE CORRELATIONS WERE LARGELY ACCOUNTED FOR BY USE OF VITAMIN SUPPLEMENTS. IN LINEAR REGRESSION MODELS, VITAMIN E INTAKE AND PLASMA LIPIDS WERE SIGNIFICANT PREDICTORS OF PLASMA ALPHA-TOCOPHEROL LEVELS./DRUG THERAPY

**SUPPLEMENTARY TERMS: Miscellaneous Descriptors**

VITAMIN E; CAROTENE; LYCOPENE; BETA-CAROTENE; VITAMIN A;  
ADOLESCENCE; ADULT; AGED; ALCOHOL DRINKING; ADMINISTRATION  
AND DOSAGE; BLOOD; CROSS SECTIONAL STUDIES; DIET; FEMALE;  
HUMAN; MALE; MIDDLE AGE; SMOKING; ANALYSIS; PLASMA;  
NUTRITION; EPIDEMIOLOGIC METHODS; LIPIDS; BODY WEIGHT;  
STATISTICS; CALORIC INTAKE

REGISTRY NUMBER: 59-02-9; 36-88-4; 502-65-8; 7235-40-7; 68-26-8

**TOXCENTER****DISPLAY IALL (ETIC File Segment)**

ACCESSION NUMBER: 2002:503699 TOXCENTER  
 DOCUMENT NUMBER: ETICBACK-49739  
 TITLE: EFFECTS OF TERBUTALINE SULFATE ON FETAL CARDIAC FUNCTION  
 AUTHOR(S): PETERSEN R; CARTER L S; CHESCHEIR N C; KATZ V L; CEFALO R  
 C  
 SOURCE: AM J OBSTET GYNECOL, (1989) (161) 509-512.  
 CODEN: AJOGA.  
 DOCUMENT TYPE: Journal  
 FILE SEGMENT: ETIC  
 LANGUAGE: English  
 ENTRY DATE: Entered STN: Dec 2002  
 Last Updated on STN: Dec 2002  
 SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
 Taxonomic Name: CAVIA PORCELLUS, HARTLEY  
 Test Object: MAMMAL, GUINEA PIG  
 Assay Method: VIABILITY, FERTILITY AND MORTALITY;  
 CARDIOVASCULAR SYSTEM; GROWTH  
 Maternal Effects: MATERNAL WEIGHT CHANGES; MATERNAL  
 CARDIOVASCULAR SYSTEM  
 Route Of Administration: SUBCUTANEOUS; INTRAPERITONEAL  
 REGISTRY NUMBER: 23031-32-5 (TERBUTALINE SULFATE)  
 9005-49-6 (HEPARIN)

**DISPLAY IALL (FEDRIP File Segment)**

ACCESSION NUMBER: 2003:105024 TOXCENTER  
 DOCUMENT NUMBER: FEDRIP-00199078  
 TITLE: Neuropsychological Functioning in Persian Gulf War Era  
 Veterans  
 AUTHOR(S): White Roberta F., Ph.D.  
 CORPORATE SOURCE: 110326; Department of Veterans Affairs Research and  
 Development (15), 810 Vermont Ave. N.W., Washington, D.C.  
 20420 United States of America  
 FILE SEGMENT: FEDRIP  
 OTHER SOURCE: FEDRIP 200301-000167  
 LANGUAGE: English  
 ENTRY DATE: Entered STN: 6 May 2003  
 Last Updated on STN: 6 May 2003  
 BEGIN DATE (INITIAL PROJECT YEAR): 19960501  
 ABSTRACT:  
 PERSIAN GULF SYNDROME; INTERVIEWS; ENVIRONMENTAL EXPOSURE; AFFECTIVE SYMPTOMS  
 Abstract Many Persian Gulf War (PGW) veterans have reported a significant  
 decline in their health since returning from the War. These symptoms have  
 included excessive fatigue skin rash, Joint pain, headaches, and disturbances  
 of concentration and memory according to the Persian Gulf Registry  
 questionnaires completed by 12,774 veterans. These veterans may have been  
 exposed to several neurotoxicants ranging from diesel fuels, oil fire smoke,  
 pesticides, and biological or chemical weapons. In addition, some of the  
 reported health symptoms overlap with symptoms of post-traumatic stress  
 disorder (PTSD), multiple chemical sensitivity (MCS), and chronic fatigue  
 syndrome (CFS). Cognitive impairments have been noted in exposures to the  
 previously mentioned neurotoxicants and in the three disorders noted above. In  
 order to document these impairments in PGW veterans, neuropsychological tests  
 of known validity and sensitivity have been used. OBJECTIVES: This study  
 evaluates the neuropsychological functioning of PGW-era veterans who are  
 seeking treatment for any type of health or adjustment complaint. PGW deployed  
 veterans will be compared with non-deployed veterans. The purpose of the group  
 comparisons is to isolate factors that may differentiate Persian Gulf War  
 veterans from non-Gulf War veterans who served during the same time period. In  
 this way, we hope to better understand the constellation of health complaints  
 popularly referred to as Gulf War Syndrome." RESEARCH PLAN/ METHODS: All

**DISPLAY IALL (FEDRIP File Segment) (cont'd)**

research subjects will be administered the previously mentioned neuropsychological battery. In addition, a standardized set of questionnaires and interviews will also be administered to identify health symptoms including physical and mental health conditions. These instruments permit diagnosis PTSD, MCS, CFS and other psychiatric disorders. The PGW veterans will be compared with non deployed veterans. Data from these groups will also be compared with those from a non-treatment seeking research sample of PGW veterans who are being studied using the same neuropsychological instruments. FINDINGS: The project's funding has ceased however, we are currently in a non-funded extension period in order to complete the data collection phase. To date, there are 137 subjects enrolled in the studs including 121 men and 16 women. As per Pi this project has ended as of 12/99.

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
FEDRIP; RPROJ; va; U.S. DEPT. OF VETERANS AFFAIRS; PERSIAN  
GULF SYNDROME INTERVIEWS ENVIRONMENTAL EXPOSURE AFFECTIVE  
SYMPTOMS

**DISPLAY IALL (HAPAB File Segment)**

L11 ANSWER 2 OF 5987 TOXCENTER COPYRIGHT 2002 ACS  
ACCESSION NUMBER: 2002:311239 TOXCENTER  
DOCUMENT NUMBER: HAPAB-72-02714  
TITLE: Determination of total bromine and of bromate in barley  
and malt.  
AUTHOR(S): Brookes P A; Martin P A  
SOURCE: J. Inst. Brewing London, (1972) 78 (2) 165-169. Ref: 17.  
FILE SEGMENT: HAPAB  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

ABSTRACT:  
Improved techniques for the determination of bromine and bromate in barley and malt were devised since published methods were found unsatisfactory. Bromate was determined directly on an aqueous extract of barley using an iodometric amplification reaction. Sodium molybdate/potassium fluoride reagent, potassium iodide solution, sulfuric acid, and starch solution were added to the extract and a titration was carried out with 0.02N sodium thiosulfate. Recoveries ranged from over 100% in wheat flour pellets and barley to as low as 85% in malt flour. Total bromine was determined using the same reaction after ashing the sample in an oxygen combustion flask and oxidizing the resulting bromide to bromate. It was necessary to dry steeped and germinating barleys for four hr at 105 DEG in order to grind samples and obtain satisfactory combustion. 1972  
REGISTRY NUMBER: 7726-95-6

## TOXCENTER

## DISPLAY IALL (HMTC File Segment)

ACCESSION NUMBER: 2002:353910 TOXCENTER  
DOCUMENT NUMBER: HMTC-94-0001885  
TITLE: EVALUATION OF XE-340 AS A TRAPPING MEDIUM FOR AIRBORNE  
ORGANOCHLORINE PESTICIDES  
AUTHOR(S): Yeboah P O; Kilgore W W  
SOURCE: Bulletin of Environmental Contamination and Toxicology  
33(1):13-19, (1984)  
FILE SEGMENT: \*\*\*HMTC\*\*\*  
LANGUAGE: English  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

## ABSTRACT:

Amberlite XAD, a porous polymer sorbent, is the most popular and least expensive commercial sampling system available to evaluate airborne organochlorine pesticides. However, its presampling cleanup procedure is lengthy. An alternative system which is just as effective but less time-consuming than XAD is the XE-340 system. XE-340 is a solid sorbent that incorporates activated carbon and polymeric principles. This paper presents the results of XE-340 sampling efficiency tests using seven organochlorine pesticides: alphachlordane, gamma-chlordane, lindane, heptachlor, heptachlor epoxide, aldrin, and dieldrin. Vapor pressures, fortification levels, and percent recoveries are presented.

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
alpha-chlordane; gamma-chlordane; U036; lindane; U129;  
heptachlor; U129; dieldrin; P037; Characteristics;  
Monitoring; Substance Identification; Chemical Industry;  
Agricultural Chemicals; Pesticides; Monitoring Methods;  
Air Analysis; Halogenated Organics

REGISTRY NUMBER: 319-84-6

**DISPLAY IALL (PESTAB File Segment)**

ACCESSION NUMBER: 2002:346703 TOXCENTER  
DOCUMENT NUMBER: PESTAB-81-2783  
TITLE: Hydrogen peroxide and hydroxyl radical: intermediates in indirect photolysis reactions in water.  
AUTHOR(S): Draper W M; Crosby D G  
CORPORATE SOURCE: Dep. Anim. Dairy & Vet. Sci., Utah State Univ., Logan, UT 84322  
SOURCE: Journal of Agricultural and Food Chemistry, (1981) 29 (4) 699-702. Ref: 22.  
CODEN: JAFCA.  
DOCUMENT TYPE: (Periodical)  
FILE SEGMENT: PESTAB  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

**ABSTRACT:**

Thiobencarb (benthocarb; I) photodecomposes slowly to thiobencarb S-oxide (II) in aqueous solutions exposed to sunlight. The rate of photodecomposition of I in water was greatly increased by addition of small amounts of hydrogen peroxide, tryptophan, or methylene blue. In these solutions, 2-hydroxythiobencarb (III), 3-hydroxythiobencarb (V), II, and N-monoethylthiobencarb (IV) were isolated as photooxidation products of I; the distributions of these products were similar in each case. Thiobencarb did not undergo reaction with photochemically generated, singlet molecular oxygen but was oxidized by hydroxyl radical (Fenton's reagent) to III, V, and II. The free radical photooxidation of I in aqueous acetone was unique; II and IV were among the photoproducts but the phenols, III and V, were absent. These results demonstrate the involvement and common intermediacy of hydrogen peroxide and hydroxyl radical in indirect photolysis reactions in aqueous solutions of hydrogen peroxide, tryptophan, and methylene blue. (Author abstract reprinted by permission of the American Chemical Society.)

REGISTRY NUMBER: 28249-77-6

**DISPLAY IALL (PPBIB File Segment)**

ACCESSION NUMBER: 2002:351132 TOXCENTER  
DOCUMENT NUMBER: PPBIB-03654  
TITLE: Trans-4-hydroxy-2-hexenal: A reactive metabolite from the macrocyclic pyrrolizidine alkaloid senecionine.  
AUTHOR(S): Segall H J; Wilson D W; Dallas J L; Haddon W F  
SOURCE: Science, (1985) 229 (Aug 2) 472-475.  
FILE SEGMENT: PPBIB  
ENTRY DATE: Entered STN: Dec 2002  
Last Updated on STN: Dec 2002

**ABSTRACT:**

The toxicity of macrocyclic pyrrolizidine alkaloids in the livers of man and animals has been attributed to the formation of reactive pyrroles from dihydropyrrolizines. Now a novel metabolite, trans-4-hydroxy-2-hexenal, has been isolated from the macrocyclic pyrrolizidine alkaloid senecionine, in an in vitro hepatic microsomal system. Other alkenals such as trans-4-hydroxy-2-nonenal have previously been isolated from microsomal systems when treated with halogenated hydrocarbons or subjected to lipid peroxidation. The in vivo pathology caused by trans-4-hydroxy-2-hexenal appears to be identical to that previously attributed to reactive pyrroles. There are similarities between the toxic effects of this alkenal and those of centrilobular hepatotoxins such as CCl<sub>4</sub> and other alkenals formed during lipid peroxidation.

SUPPLEMENTARY TERMS: Miscellaneous Descriptors

Chemical analysis; Senecio; Compositae; Liver; Hepatic microsomal enzyme; Centrilobular necrosis; Hepatotoxic

REGISTRY NUMBER: 130-01-8

## TOXCENTER

## DISPLAY IALL (RISKLINE File Segment)

ACCESSION NUMBER: 2002:624842 TOXCENTER [Full-text](#)  
DOCUMENT NUMBER: RISKLINE-2001120009  
TITLE: Toxicology and carcinogenesis studies of  
p,p'-Dichlorodiphenyl sulfone in F344/N rats and B6C3F1  
mice (feed studies)  
AUTHOR(S): Anonymous  
SOURCE: National Toxicology Program Technical Report Series,  
(2001) 501 262 p.  
FILE SEGMENT: RISKLINE  
LANGUAGE: English  
ENTRY DATE: Entered STN: 13 Jul 2004  
Last Updated on STN: 29 Nov 2005

## ABSTRACT:

p,p'-Dichlorodiphenyl sulfone is used as a starting material in the production of polysulfones and polyethersulfones and as a component in reactive dyes in the textile industry; it is also a by-product of pesticide production. p,p'-Dichlorodiphenyl sulfone was nominated for study by the National Cancer Institute because of its history of high production and use, the prospect of increased production and use, and the absence of adequate toxicity testing. Male and female F344/N rats and B6C3F, mice were exposed to p,p'-dichlorodiphenyl sulfone (greater than 99% pure) in feed for 14 weeks or 2 years. Genetic toxicology studies were conducted in Salmonella typhimurium, cultured Chinese hamster ovary cells, and mouse bone marrow. 14-week study in rats. Groups of 10 male and 10 female F344/N rats were fed diets containing 0, 30, 100, 300, 1,000, or 3,000 ppm p,p'-dichlorodiphenyl sulfone (equivalent to average daily doses of approximately 2, 6, 19, 65, or 200 mg p,p'-dichlorodiphenyl sulfone/kg body weight) for 14 weeks. All rats survived until the end of the study. Mean body weights of groups exposed to 300 ppm or greater were significantly less than those of the controls. Liver weights of groups exposed to 100 ppm or greater and kidney weights of 1,000 and 3,000 ppm male rats were significantly greater than those of the controls. Centrilobular hepatocyte hypertrophy of the liver was observed in most male rats exposed to 100 ppm or greater and in all female rats exposed to 300 ppm or greater, and the severities were increased in 300 ppm males and 1,000 and 3,000 ppm males and females. The incidences of nephropathy in 1,000 and 3,000 ppm female rats were significantly increased. Dose-related increases in severity of nephropathy were observed in male rats. 14-week study in mice. Groups of 10 male and 10 female B6C3F1 mice were fed diets containing 0, 30, 100, 300, 1,000, or 3,000 ppm p,p'-dichlorodiphenyl sulfone (equivalent to average daily doses of approximately 3.5, 15, 50, 165, or 480 mg/kg) for 14 weeks. All mice survived until the end of the study. Mean body weights of groups exposed to 300 ppm or greater were significantly less than those of the controls. Liver weights of groups exposed to 300 ppm or greater were significantly increased. Centrilobular hypertrophy of the liver was observed in most males exposed to 100 ppm or greater and in all females exposed to 1,000 or 3,000 ppm, and the severities generally increased with increasing exposure concentration. 2-year study in rats. Groups of 50 male and 50 female rats were fed diets containing 0,10 (males), 30,100, or 300 (females) ppm p,p'-dichlorodiphenyl sulfone for 105 weeks. Dietary concentrations of 10, 30, and 100 ppm resulted in average daily doses of approximately 0.5, 1.5, and 5.0 mg/kg to males. Dietary concentrations of 30, 100, and 300 ppm resulted in average daily doses of approximately 1.6, 5.4, and 17 mg/kg to females. Additional groups of 10 male and 10 female rats were fed the same p,p'-dichlorodiphenyl sulfone-containing diets for 18 months and bled for plasma determinations of p,p'-dichlorodiphenyl sulfone at approximately 2 weeks and 3, 12, and 18 months. Survival of all exposed groups of male and female rats was similar to that of the control groups. Mean body weights of 30 and 100 ppm males were generally less than those of the controls during the latter part of the study, and mean body weights of 100 and 300 ppm female rats were less from weeks 30 and 18, respectively. Feed consumption by the exposed groups was similar to that by the controls throughout the study. The incidences of centrilobular hepatocyte hypertrophy in 100 ppm male and 100 and 300 ppm female rats were significantly greater than those in the controls. The incidences of bile duct hyperplasia and centrilobular degeneration were also significantly increased in 100 and 300

**DISPLAY IALL (RISKLINE File Segment) (cont'd)**

ppm females. No neoplasms were related to chemical exposure. 2-year study in mice. Groups of 50 male and 50 female mice were fed diets containing 0, 30, 100, or 300 ppm p,p'-dichlorodiphenyl sulfone for 105 to 106 weeks. Dietary concentrations of 30, 100, and 300 ppm delivered average daily doses of approximately 4, 13, and 40 mg/kg to males and approximately 3, 10, and 33 mg/kg to females. Additional groups of 10 male and 10 female mice were fed the same p,p'-dichlorodiphenyl sulfone-containing diets for up to 12 months; three mice in each group were bled for plasma determinations of p,p'-dichloro-diphenyl sulfone at approximately 2 weeks or 3 or 12 months. Survival of all exposed groups of male and female mice was similar to that of the control groups. Mean body weights of 300 ppm mice were less than those of the controls throughout most of the study. Feed consumption by the exposed groups was similar to that by the controls throughout the study. The incidences of centlobular hepatocyte hypertrophy in all exposed groups of male mice and in 100 and 300 ppm females were significantly greater than those in the controls. The incidence of eosinophilic foci in 300 ppm females was significantly increased. No neoplasms were related to chemical exposure. Pharmacokinetics. Of p,p'-Dichlorodiphenyl sulfone. p,p'-Dichlorodiphenyl sulfone is rapidly absorbed from the gut and metabolized by a saturable process. Although some p,p'-dichlorodiphenyl sulfone is eliminated unchanged in feces and urine, most of the elimination is via metabolism. Mathematical modeling of the toxicokinetics supports the view that p,p'-dichlorodiphenyl sulfone induces enzymes involved in its metabolism. Genetic toxicology. p,p'-Dichlorodiphenyl sulfone was not mutagenic in any of several strains of *Salmonella typhimurium*, with or without metabolic activation enzymes (S9). Results of the sister chromatid exchange test in cultured Chinese hamster ovary cells were judged to be negative in the presence of S9 and equivocal in the absence of S9, but no induction of chromosomal aberrations was noted, with or without S9. In contrast to the in vitro results, positive results were obtained in an acute in vivo mouse bone marrow micronucleus assay with p,p'-dichlorodiphenyl sulfone administered by intraperitoneal injection three times over a dose range of 200 to 800 mg/kg. Conclusions. Under the conditions of these 2-year feed studies, there was no evidence of carcinogenic activity of p,p'-dichlorodiphenyl sulfone in male F344/N rats exposed to 10, 30, or 100 ppm or in female F344/N rats exposed to 30, 100, or 300 ppm. There was no evidence of carcinogenic activity of p,p'-dichlorodiphenyl sulfone in male or female B6C3F1 mice exposed to 30, 100, or 300 ppm. Exposure to p,p'-dichlorodiphenyl sulfone for 2 years caused increased incidences of nonneoplastic lesions of the liver in male and female rats and mice.

**SUPPLEMENTARY TERMS: Miscellaneous Descriptors**

ANIMAL; acute toxicity; subacute toxicity; chronic toxicity; carcinogenicity; genetic toxicity; toxicokinetics; liver; dose response; textile industry; pesticides

**REGISTRY NUMBER:**

80-07-9

**TOXCENTER****DISPLAY IALL (TSCATS File Segment)**

ACCESSION NUMBER: 2002:612111 TOXCENTER  
 DOCUMENT NUMBER: TSCATS-452426  
 TITLE: MUTAGENICITY EVALUATION OF HEXAMETHYLDISILOXANE. FINAL REPORT.  
 CORPORATE SOURCE: LITTON BIONETICS INC  
 ORDER NUMBER: NTIS/OTS0572722  
 SOURCE: EPA/OTS; Doc #86940001687.  
 FILE SEGMENT: TSCATS  
 ENTRY DATE: Entered STN: Dec 2002  
 Last Updated on STN: Dec 2002

**ABSTRACT:**

Hexamethyldisiloxane did not produce responses in the L5178Y mouse lymphoma cells which indicated consistent genetic activity. Several scattered increases appeared to be the result of spurious fluctuations and cytotoxicity. The only evidence of a genotoxic response was the induction of chromosome aberrations under non-activation test conditions. These data can be considered evidence for weak clastogenic activity.

CLASSIFICATION CODE: TSCA Sect. 8D Rec 04/28/94

SUPPLEMENTARY TERMS: Miscellaneous Descriptors  
 INDUS HLTH FOUNDATION INC; HEXAMETHYLDISILOXANE (CAS# 107-46-0); HEALTH EFFECTS; GENOTOXICITY; GENE MUTATIONS; MAMMALS; MICE; IN VITRO

REGISTRY NUMBER: 107-46-0

**EXPAND in /BC Thesaurus**

=> E FUNGI+ALL/BC

E1 0 BT3 Super Taxa/BC  
 E2 0 BT2 Super Taxa Terms/BC  
 E3 27236 BT1 Plantae/BC  
 E4 11787 --> Fungi/BC  
 E5 11785 UF 15000/BC  
 E6 11787 UF 15000 Fungi/BC  
 E7 8930 NT1 Ascomycetes/BC  
 E8 5155 NT1 Basidiomycetes/BC  
 E9 30548 NT1 Fungi Imperfecti or Deuteromycetes/BC  
 E10 197 NT1 Myxophyta/BC  
 E11 2547 NT1 Phycomycetes/BC  
 \*\*\*\*\* END \*\*\*\*\*

**EXPAND in /CN Thesaurus**

=> E SERICYSTATIN+ALL/CN

E1 0 --> Sericystatin/CN  
 E2 5 USE CSTA protein, human/CN  
 \*\*\*\*\* END \*\*\*\*\*

## EXPAND in /CT Thesaurus

## =&gt; E SUNSCREENING AGENTS+ALL/CT

```
E1          0      BT4   J Technology, Industry, Agriculture/CT
E2          0      BT3   Technology, Industry, and Agriculture/CT
E3         750     BT2   Household Products/CT
E4          0      BT4   D Chemicals and Drugs/CT
E5          0      BT3   Chemical Actions and Uses/CT
E6          3      BT2   Specialty Uses of Chemicals/CT
E7        2139     BT1   Cosmetics/CT
E8          0     BT5   D Chemicals and Drugs/CT
E9          0     BT4   Chemical Actions and Uses/CT
E10         0     BT3   Pharmacologic Actions/CT
E11         0     BT2   Therapeutic Uses/CT
E12       2977     BT1   Dermatologic Agents/CT
E13         0     BT6   D Chemicals and Drugs/CT
E14         0     BT5   Chemical Actions and Uses/CT
E15         0     BT4   Pharmacologic Actions/CT
E16         0     BT3   Physiological Effects of Drugs/CT
E17         0     BT5   D Chemicals and Drugs/CT
E18         0     BT4   Chemical Actions and Uses/CT
E19         3     BT3   Specialty Uses of Chemicals/CT
E20       1404     BT2   Protective Agents/CT
E21       5876     BT1   Radiation-Protective Agents/CT
E22       1677     -->   Sunscreening Agents/CT
E23       1677     MN    D27.505.696.706.776.800./CT
E24       1677     MN    D27.505.954.444.695./CT
E25       1677     MN    D27.720.269.800./CT
E26       1677     MN    D27.720.799.763.764./CT
                DC    an INDEX MEDICUS major descriptor
                NOTE   Chemical or physical agents that protect the
                        skin from sunburn and erythema by absorbing or
                        blocking ultraviolet radiation.
                INDX   D25-26 qualif
                AQ     AD AE AN CH CL CS CT DU EC HI IP ME PD PK PO RE
                        SD ST TO TU
                HNTE   72(66)
                MHTH   NLM (1966)
E27         0     UF    Agents, Sunscreening/CT
E28         0     UF    Sunscreens/CT
E29        643     NT1   Zinc Oxide/CT
***** END *****
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## TOXCENTER

## EXPAND in /GT Thesaurus

=&gt; E AFRICA+ALL/GT

E1	0	BT3 Geopolitical Locations/GT
E2	0	BT2 Geographic Classifiers/GT
E3	2827	BT1 Ethiopian region/GT
E4	0	BT3 Geopolitical Locations/GT
E5	0	BT2 Geographic Classifiers/GT
E6	38036	BT1 Palearctic region/GT
E7	3188	--> Africa/GT
E8	9	NT1 Angola/GT
E9	52	NT1 Benin/GT
E10	15	NT1 Botswana/GT
E11	1	NT1 Burkina-Faso/GT
E12	5	NT1 Burundi/GT
E13	57	NT1 Cameroon/GT
E14	8	NT1 Central African Republic/GT
E15	7	NT1 Chad/GT
E16	19	NT1 Congo/GT
E17	0	NT1 Congo River/GT
E18	5	NT1 Djibouti/GT
E19	3	NT1 Equatorial Guinea/GT
E20	4	NT1 Eritrea/GT
E21	67	NT1 Ethiopia/GT
E22	11	NT1 Gabon/GT
E23	26	NT1 Gambia/GT
E24	105	NT1 Ghana/GT
E25	10	NT1 Guinea/GT
E26	11	NT1 Guinea-Bissau/GT
E27	26	NT1 Ivory Coast/GT
E28	162	NT1 Kenya/GT
E29	6	NT1 Lesotho/GT
E30	5	NT1 Liberia/GT
E31	36	NT1 Madagascar/GT
E32	32	NT1 Malawi/GT
E33	24	NT1 Mali/GT
E34	9	NT1 Mauritania/GT
E35	28	NT1 Mozambique/GT
E36	12	NT1 Namibia/GT
E37	15	NT1 Niger/GT
E38	432	NT1 Nigeria/GT
E39	9	NT1 Rwanda/GT
E40	0	NT1 Sao Tome and Principe/GT
E41	45	NT1 Senegal/GT
E42	8	NT1 Sierra Leone/GT
E43	13	NT1 Somalia/GT
E44	842	NT1 South Africa/GT
E45	55	NT1 Sudan/GT
E46	7	NT1 Swaziland/GT
E47	102	NT1 Tanzania/GT
E48	15	NT1 Togo/GT
E49	86	NT1 Uganda/GT
E50	9	NT1 Zaire/GT
E51	39	NT1 Zambia/GT
E52	71	NT1 Zimbabwe/GT
E53	44	NT1 Algeria/GT
E54	411	NT1 Egypt/GT
E55	13	NT1 Libya/GT
E56	162	NT1 Morocco/GT
E57	125	NT1 Tunisia/GT
*****	END	*****

## EXPAND in /MN Thesaurus

=&gt; E D27.720.269.800.+ALL/MN

E1 BT4 J Technology, Industry, Agriculture/MN  
E2 BT3 Technology, Industry, and Agriculture/MN  
E3 BT2 Household Products/MN  
E4 BT4 D Chemicals and Drugs/MN  
E5 BT3 Chemical Actions and Uses/MN  
E6 BT2 Specialty Uses of Chemicals/MN  
E7 BT1 Cosmetics/MN  
E8 BT5 D Chemicals and Drugs/MN  
E9 BT4 Chemical Actions and Uses/MN  
E10 BT3 Pharmacologic Actions/MN  
E11 BT2 Therapeutic Uses/MN  
E12 BT1 Dermatologic Agents/MN  
E13 BT6 D Chemicals and Drugs/MN  
E14 BT5 Chemical Actions and Uses/MN  
E15 BT4 Pharmacologic Actions/MN  
E16 BT3 Physiological Effects of Drugs/MN  
E17 BT5 D Chemicals and Drugs/MN  
E18 BT4 Chemical Actions and Uses/MN  
E19 BT3 Specialty Uses of Chemicals/MN  
E20 BT2 Protective Agents/MN  
E21 BT1 Radiation-Protective Agents/MN  
E22 --> D27.720.269.800./MN  
E23 MH Sunscreening Agents/MN  
DC an INDEX MEDICUS major descriptor  
NOTE Chemical or physical agents that protect the  
skin from sunburn and erythema by absorbing or  
blocking ultraviolet radiation.  
INDX D25-26 qualif  
AQ AD AE AN CH CL CS CT DU EC HI IP ME PD PK PO RE  
SD ST TO TU  
HNTE 72(66)  
MHTH NLM (1966)  
E24 UF Agents, Sunscreening/MN  
E25 UF Sunscreens/MN  
\*\*\*\*\* END \*\*\*\*\*

## EXPAND in /ORGN Thesaurus

=&gt; E RODENTIA+ALL/ORGN

E1 0 BT6 Super Taxa/ORGN  
E2 0 BT5 Super Taxa Terms/ORGN  
E3 1369945 BT4 Animalia/ORGN  
E4 1293434 BT3 Chordata/ORGN  
E5 1291289 BT2 Vertebrata/ORGN  
E6 1219228 BT1 Mammalia/ORGN  
E7 449350 --> Rodentia/ORGN  
E8 2699 UF 86265/ORGN  
E9 0 UF BC86265/ORGN  
E10 4 NT1 Abrocomidae/ORGN  
E11 39 NT1 Anomaluridae/ORGN  
E12 5 NT1 Aplodontiidae/ORGN  
E13 9 NT1 Bathyergidae/ORGN  
E14 24 NT1 Capromyidae/ORGN  
E15 35 NT1 Castoridae/ORGN  
E16 21967 NT1 Caviidae/ORGN  
E17 208 NT1 Chinchillidae/ORGN  
E18 19987 NT1 Cricetidae/ORGN  
E19 37 NT1 Ctenodactylidae/ORGN  
E20 5 NT1 Ctenomyidae/ORGN  
E21 20 NT1 Dasyproctidae/ORGN  
E22 6 NT1 Dinomyidae/ORGN

## TOXCENTER

=&gt; E RODENTIA+ALL/ORGN (cont'd)

E23	7	NT1	Dipodidae/ORGN
E24	14	NT1	Echimyidae/ORGN
E25	22	NT1	Erethizontidae/ORGN
E26	32	NT1	Geomyidae/ORGN
E27	7	NT1	Gliridae/ORGN
E28	42	NT1	Heteromyidae/ORGN
E29	4	NT1	Hydrochoeridae/ORGN
E30	29	NT1	Hystriidae/ORGN
E31	417448	NT1	Muridae/ORGN
E32	14	NT1	Octodontidae/ORGN
E33	5	NT1	Pedetidae/ORGN
E34	0	NT1	Petromyidae/ORGN
E35	6	NT1	Platacanthomyidae/ORGN
E36	3	NT1	Rhizomyidae/ORGN
E37	367	NT1	Sciuridae/ORGN
E38	13	NT1	Seleviniidae/ORGN
E39	7	NT1	Spalacidae/ORGN
E40	2	NT1	Thryonomyidae/ORGN
E41	9	NT1	Zapodidae/ORGN
E42	449350	RT	Rodents/ORGN

\*\*\*\*\* END \*\*\*\*\*

## EXPAND in /ST Thesaurus

=&gt; E ANIMAL HUSBANDRY+ALL/ST

E1	0	BT3	Major Concepts/ST
E2	0	BT2	Major Concept Terms/ST
E3	56097	BT1	Agriculture/ST
E4	16871	-->	Animal Husbandry/ST

NOTE Studies of the breeding, feeding, housing, and management of animals used to produce consumer goods.

NOTE For studies of disease and health care of domesticated or zoo animals, see Veterinary Medicine. For studies of fish rearing, see Aquaculture. For studies of the care and breeding of laboratory or zoo animals in artificial environments, see Animal Care.

E5	1462	RT	Aquaculture/ST
E6	4355	RT	Animal Care/ST
E7	22378	RT	Veterinary Medicine/ST

\*\*\*\*\* END \*\*\*\*\*

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