

# **ESBIOBASE** (Elsevier BIOBASE)

# Subject Coverage

- · Applied microbiology and biotechnology
- Cancer research
- · Cell and developmental biology
- · Clinical chemistry
- Ecological and environmental sciences
- Endocrinology and metabolism
- Genetics and molecular biology
- Neuroscience
- Plant science
- Protein biochemistry
- Toxicology

File Type	Bibliographic					
Features	Alerts (SDIs)	Not available				
	CAS Registry Number® Identifiers		SLART			
	Keep & Share		Structures			
Record Content	<ul> <li>Bibliographic information, abstracts (80%), and indexing</li> <li>Cited references from over 1,900 source journals</li> </ul>					
File Size	9.063.758 records					
Coverage	1994-2021					

## Updates

Static file

## Language

English

## Database Producer

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Email: <a href="mailto:ebd-marketing@elsevier.com">ebd-marketing@elsevier.com</a>

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## Sources

• Over 1,900 primary research journals are monitored for coverage

## **User Aids**

- Online Helps (HELP DIRECTORY lists all help messages available)
- STNGUIDE

#### Cluster

- AGRICULTURE
- ALLBIB
- AUTHORS
- BIOSCIENCE
- CORPSOURCE
- ENVIRONMENT
- HEALTH
- MEDICINE
- PHARMACOLOGY
- TOXICOLOGY

**STN Database Cluster information** 

# **Search and Display Field Codes**

Fields that allow left truncation are indicated by an asterisk (\*)

## **General Search Fields**

Search Field Name	Search Code	Search Examples	Display Codes
Basic Index* (contains single words from the title (TI), abstract (AB), classification code (text) (CC), gene number (GEN), organism (ORGN) and supplementary term (ST) fields)	None or /BI	S SEED GERMINATION S HIGH TEMPERATURE S ?BACTERIUM	AB, CC, GEN, ORGN, TI, ST
Abstract Accession Number Author	/AB /AN /AU	S ?ORGANISM/AB S 1999130558/AN S BANNWARTH B/AU S MANABE, H./AU	AB AN AU
Classification Code *(1) (code and text)	/CC	S 86.3.4/CC S INFECTIOUS DISEASE/CC S ?CYTE?/CC	CC
Corporate Source (1) (authors, affiliations, and e-mail addresses)	/CS	S UNIVERSITY LISBON/CS	CS, EML
Country (of Publication) (ISO code and text)	/CY	S DE/CY S UNITED KINGDOM/CY	CY
Document Type (STN code and text)	/DT (or /TC)	S BA/DT S BOOK ARTICLE/DT	DT
E-mail <b>(1)</b>	ÈML ´	S LOUIS BIO/EML	CS, EML
Entry Date <b>(2)</b> Field Availability	/ED /FA	S ED=FEB 2009 S ORGN/FA	ED FA
Gene Number * International Standard (Document) Number (contains CODEN, ISBN and ISSN)	/GEN /ISN	S L26294/GEN S 1017-7825/ISN S ANEMDG/ISN	GEN ISN, SO
Journal Title	/JT	S ACTA CYTOLOGICA/JT AND PY=1999	JT, JTA, JTF, SO
Language (ISO code and text)	/LA	S FR/LA S FRENCH/LA	LA
Organism * Publication Date (2) Publication Year (2) Publisher (2) Publisher Item Identifier Reference Count (2)	/ORGN /PD /PY /PB /PUI /REC (or /RE.CNT)	S ABIES ALBA/ORGN S JAN 1999/PD S 1997-1998/PY S SPRINGER HEIDELBERG/PB S S0001706X99000030/PUI S 19/REC	ORGN PD, SO PY, SO PB, SO PUI REC, SO
Source (contains journal titles collation, CODEN, ISSN, publication year, and ISBN)	/SO	S BIOCONTROL SCIENCE/SO S NGENEC/SO AND 22/SO	SO
Summary Language (ISO code and text)	/SL	S DE/SL	SL
Supplementary Term * (1)	/ST	S ?FERTILITY/ST S DEATH/ST	ST
Title*	/TI	S PLANT SCIENCE/TI	TI
Update Date (2) Word Count, Title (2)	/UP /WC.T	S UP>MAR 2009 S WC.T<=8	ED WC.T

<sup>(1)</sup> Search with implied (S) proximity is available in this field.(2) Numeric search field that may be searched with numeric operators or ranges.

# **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 TI AU. The fields are displayed or printed in the order requested.

Hit-term highlighting is available for most fields. Highlighting must be ON during SEARCH in order to use the HIT, KWIC, and OCC formats.

Format	Content	Examples
AB AN AU CC CS CY DT (TC) ED (UP) EML (1) GEN ISN (1) JT (1) JTA (1) JTF (1) LA ORGN PB (1) PD (1) PU (1) REC (RE.CNT) (1) SL SO ST TI WC.T (1)	Abstract Accession Number Author Classification Code Corporate Source (authors, affiliations, and e-mail addresses) Country (of Publication) Document Type Entry Date (contains Update Date) E-mail Address Gene Number International Standard (Document) Number Journal Title Journal Title, Abbreviated Journal Title, Full Language Organism Publisher Publication Date Publisher Item Identifier Publication Year Reference Count Summary Language Source Supplementary Term Title Word Count, Title	D AB 1-5 D AN D AU D CC D CS D CY D DT D ED D EML D GEN D ISN D JT D JTA D JTF D LA, SL 1-3 D ORGN D PB D PUI SO D PY D REC D SL D SO D ST D TI D WC.T
ABS ALL DALL IALL BIB IBIB IND SCAN (2) TRIAL (TRI, SAMPLE, SAM, FREE)	AN, AB AN, TI, AU, CS, SO, PUI, DT, CY, LA, SL, ED, AB, CC, ST, ORGN, GEN ALL, with delimiter for post-processing ALL, indented with text labels AN, TI, AU, CS, SO, PUI, DT, CY, LA, SL, ED (BIB is the default) BIB, indented with text labels AN, CC, ST, ORGN, GEN TI, ST (random display without answer numbers) TI, CC, ST, ORGN	D ABS D ALL D DALL D IALL D BIB D IBIB D IND D SCAN D TRI
HIT KWIC OCC	Hit term(s) and field(s) Up to 50 words before and after hit term(s) (KeyWord-In-Context) Number of occurrences of hit term(s) and field(s) in which they occur	D HIT D KWIC D OCC

<sup>(1)</sup> Custom display only.

<sup>(2)</sup> SCAN must be specified on the command line, i.e., D SCAN or DISPLAY SCAN.

# 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	AB	Υ	N
Accession Number	AN	Υ	N
Author	AU	Y (2)	Υ
Citation	CIT (RE)	Y (2,3)	N
Classification Code	CC	Y (2)	Υ
CODEN	CODEN	N	Υ
Corporate Source	CS	Y (2)	Υ
Country of Publication	CY	Υ	Υ
Document Type	DT (TC)	Υ	Υ
E-mail Address	EML	Υ	Y
Entry Date	ED	Υ	Υ
Genbank Number	GENBANK	Y (2)	N
Gene Number	GEN	Y (2)	Υ
International Standard (Document) Number	ISN	Y (4)	Υ
International Standard Book Number	ISBN	N	Y
International Standard Serial Number	ISSN	N	Y
Journal Title	JT	Υ	Y
Journal Title, Abbreviated	JTA	Υ	Y
Journal Title, Full	JTF	Υ	Y
Language	LA	Υ	Y
Occurrence Count of Hit Terms	OCC	N	Y
Organism	ORGN	Υ	Y
Publisher	PB	Υ	Y
Publication Date	PD	Υ	Y
Publication Year	PY	Υ	Y
Publisher Item Identifier	PUI	Υ	Y
Reference Count	REC (RE.CNT)	Υ	Y
Source	SO	Y <b>(5)</b>	Y
Summary Language	SL	Υ	Y
Supplementary Term	ST	Y (2)	N
Title	TI	Y (default)	Y
Update Date	UP	Υ	Υ
Word Count, Title	WC.T	Y	Y

<sup>(1)</sup> 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.

<sup>(2)</sup> SELECT HIT or ANALYZE HIT are not valid with this field.

<sup>(3)</sup> SELECT or ANALYZE CIT allows you to extract the reference from the source documents in this file and have them automatically converted to a citation format for searching in the SCISEARCH file. SEL or ANALYZE CIT extracts first author, publication year, volume, first page, with a truncation symbol and with /RE appended to the terms created by SELECT.

<sup>(4)</sup> Selects or analyzes CODEN and ISSN with /ISN appended to the terms created by SELECT.

<sup>(5)</sup> Selects or analyzes CODEN and ISSN with /SO appended to the terms created by SELECT.

# Sample Record

#### **DISPLAY ALL**

- AN 2009032890 ESBIOBASE
- TI Plasma 25-hydroxyvitamin D, 1,25-dihydroxyvitamin D, and parathyroid hormone in familial hypocalciuric hypercalcemia and primary hyperparathyroidism
- AU Christensen, Signe Engkjaer; Vestergaard, Peter; Rejnmark, Lars; Mosekilde, Leil; Nissen, Peter H.; Heickendorff, Lene; Brixen, Kim
- CS Christensen, Signe Engkjaer; Vestergaard, Peter; Rejnmark, Lars; Mosekilde, Leil (Department of Endocrinology and Metabolism C, Aarhus University Hospital, Aarhus Sygehus, Tage-Hansens gade 2, DK 8000 Arhus C (DK)); Nissen, Peter H.; Heickendorff, Lene (Department of Clinical Biochemistry, Aarhus University Hospital, Aarhus Sygehus, Tage-Hansens gade 2, DK 8000 Arhus C (DK)); Brixen, Kim (Department of Endocrinology, Odense University Hospital, Odense (DK))
  EMAIL: rlq06sec@as.aaa.dk
- SO European Journal of Endocrinology (Dec 2008) Volume 159, Number 6, pp. 719-727, 47 refs.

CODEN: EJOEEP ISSN: 0804-4643 E-ISSN: 1479-683X

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- DT Journal; Article
- LA English
- SL English
- ED Entered STN: 27 Feb 2009
  - Last updated on STN: 27 Feb 2009
- Introduction: Familial hypocalciuric hypercalcemia (FHH) is a lifelong, AΒ benign, inherited condition caused by inactivating mutations in the calcium-sensing receptor (CASR) gene. Both FHH and primary hyperparathyroidism (PHPT) are characterized by elevated P-calcium, normal or elevated plasmaparathyroid hormone (P-PTH), and typically normal renal function. In PHPT, vitamin D metabolism is typically characterized by low plasma levels of 25-hydroxyvitamin D (250HD), and high plasma levels of 1,25-dihydroxyvitamin D (1,25(HO) 2 D). In FHH, the vitamin D metabolism is not very well known. Objective: To compare and evaluate plasma 250HD, 1,25(OH) 2 D, and PTH in FHH and PHPT. Design: Cross-sectional study. Materials: About 66 FHH patients with mutations in the CASR gene, 147 patients with surgically verified PHPT, and 46 controls matched to FHH patients according to age (± 5 years), sex, and season. All patients had a P-creatinine < 140 μmol/l. Methods: We measured P-calcium, P-Ca 2+ , P-albumin, P-creatinine, P-phosphate, Pmagnesium, and P-PTH by standard laboratory methods. P-250HD and P-1,25(OH) 2 D were measured by RIA or enzyme immunoassay, In FHH, all protein-coding exons in the CASR gene were sequenced and aligned to GenBank reference sequence NM\_000388.2. Results: PHPT patients had higher body mass index (2p<0.01), together with higher P-PTH (2p< 0.01) and P-1,25(OH) 2 )D (2p<0.01) compared with FHH patients. The groups had similar levels of P-Ca 2+ and of P-25OHD. The phenotypic expression of the CASR mutations (as determined by the degree of hypercalcemia) did not influence the levels of P-1,25(OH) 2 D. Conclusion: Even though Pcalcium, and P-250HD were comparable, P-1,25(OH) 2 D and P-PTH differed between FHH and PHPT. . COPYRGT. 2008 European Society of Endocrinology. CC 81.3.7.5 CLINICAL CHEMISTRY, PHYSIOLOGY, Endocrinology, Parathyroid pathophysiology; 81.4.7.8 CLINICAL CHEMISTRY, BIOCHEMISTRY, Vitamins,
  - 81.3.7.5 CLINICAL CHEMISTRY, PHYSIOLOGY, Endocrinology, Parathyroid pathophysiology; 81.4.7.8 CLINICAL CHEMISTRY, BIOCHEMISTRY, Vitamins, Vitamin D; 83.9.6.1 ENDOCRINOLOGY AND METABOLISM, ENERGY BALANCE AND NUTRITION, Hormone Control of Calcium Metabolism and Bone Physiology, Parathyroid hormone; 83.9.6.2 ENDOCRINOLOGY AND METABOLISM, ENERGY

BALANCE AND NUTRITION, Hormone Control of Calcium Metabolism and Bone Physiology, Vitamin D and derivatives GENBANK NM\_000388 referred number

GEN

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