

Pituitary NLP Search

文献由来知識情報から自然言語処理で抽出した遺伝子・現象の
関係性検索データベース

http://bioinf.mind.meiji.ac.jp/KPJ_DB/Ref_Search_ver2a.html



Account : blog_kato Password : ochid44ckv34

Pituitary NLP Search

Reference Search
Help

Entity

To search with a multi-word entity, connect words with underscore.

Subjective Entity

Objective Entity

Relation Type

Regulation

ChemicalReaction

PromoterBinding

MolSynthesis

MolTransport

DirectRegulation

Expression

Binding

ProtModification

miRNAEffect

Species

Homo sapiens (Human)

Rattus norvegicus (Rat)

PMID

Only number part of PubMed ID is accepted. Separate with a single space when submitting multiple queries.

To be shown

To be excluded

Publication Year

The dominical year in four digits is accepted. Separate with a single space when submitting multiple queries.

Submit

- ・ 遺伝子と遺伝子の関係性が論文でどう書かれているか知りたい
- ・ 遺伝子と現象の関係性を調べたい



例) PROP1で検索

Subjective Entity

Submit

Search Result in Human

Relation	Type	Sentence	PMID	Journal	Year
PROP1 --> ACTH deficiency	Regulation	PROP1 (Prophet of Pit-1) gene mutations also cause gonadotrophin deficiencies and in some cases partial ACTH deficiency.	12780757	Clin Endocrinol (Oxf)	2003
PROP1 --> ACTH deficiency	Regulation	"Mutations in the PROP-1 gene, which are a more common cause of hypopituitarism, lead to a clinical phenotype characterized by GH, PRL, TSH, LH and FSH deficiency, and sometimes ACTH deficiency as well."	12717343	Minerva Endocrinol	2003
PROP1 --> ACTH deficiency	Regulation	"In humans, PROP1 mutations are the most common cause of combined pituitary hormone deficiency, including GH, TSH, gonadotropin, and evolving ACTH deficiencies."	19837867	Endocr Rev	2009
PROP1 --> ACTH deficiency	Regulation	"Occasionally lack of PROP1 causes acquired ACTH deficiency, suggesting that Prop1 is not strictly required for corticotrope differentiation but is essential for differentiation of corticotrope."	11371507	Hum Mol Genet	2001

遺伝子と遺伝子, 遺伝子と現象の関係性がある文章だけを論文から抽出

PROP1 --> ACTH deficiency	Regulation	of the PROP1 transcription factor causes ACTH deficiency directly, the clinical finding of an impaired pituitary adrenal axis suggests that PROP1 is not strictly required for corticotrope differentiation but is essential for differentiation of corticotrope.	15472232	J Clin Endocrinol Metab	2004
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162480報の論文由来 57880 Relation(関係性)搭載

Human, Rat対応

Search Result Example

PITX2とSitus Inversusの間に関係性があることを表す

Regulationの関係

Relation	Type	Sentence	PMID	Journal	Year
PITX2 ----> Situs Inversus	Regulation	"Thus, the findings in this patient failed to indicate that mutation of the PITX2 gene is involved in the pathomechanism of situs inversus totalis associated with inherited form of multiple pituitary hormone deficiency."	21278027	Orv Hetil	2011

抽出元の文章

論文のPubMed ID

論文の掲載誌

論文の出版年

Function

POMC TSH
dwarfism ...

タンパク質名, 化合物名, 病名などからの検索

2001 2002
2003 ...

論文の発行年, PMIDからの検索



Binding, Regulation, Expressionなど関係性の種類別の検索



検索結果の文章の引用元論文Abstのリンク

Method

Pathway Studio®

下垂体に関連するキーワード23個を用いてPubMedから論文を収集し, 自然言語処理ツールMedScan & PathwayStudio®を使用してRelationを抽出



使用キーワード

ACTH / adrenocorticotrophic hormone, adenohipophysis, anterior lobe, dwarf, FSH / follicle-stimulating hormone, GH / growth hormone, hesx1, hypergonadism, hypogonadism, intermediate lobe, LH / luteinizing hormone, neurohypophysis, pit1 / pit-1, pituitary, pituitary adenoma, pitx1, pitx2, POMC / pro-opiomelanocortin, posterior lobe, prolactin / PRL, prop1 / prop-1, Rathke's pouch, TSH / thyroid-stimulating hormone