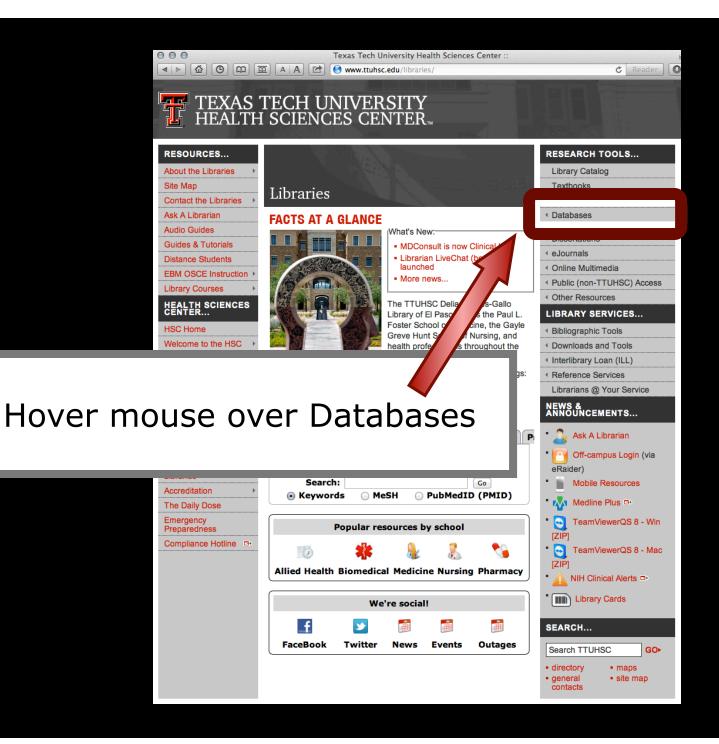
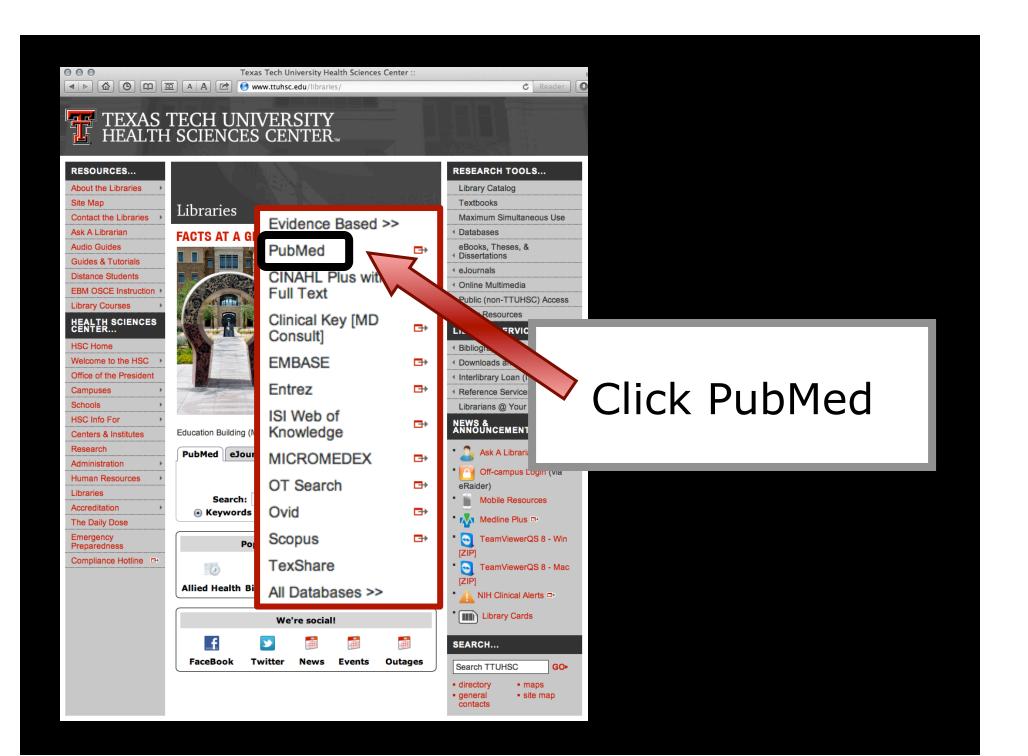


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Click MeSH Database

MeSH Database

Terms in the database are called Medical Subject Headings or MeSH.

is a "controlled vocabulary list" of more than <u>25,000</u> subject headings.

10-15 **MeSH** are assigned to each article in *PubMed*.

The *MeSH* index of 10-15 subject headings is like a keyword abstract of the article.

MeSH are updated annually in December.

Frequently Used Mesh Categories – 2011

A Anatomy

- A1 Body Regions
- A2 Musculoskeletal System
- A3 Digestive System
- A4 Respiratory System
- A5 Urogenital System
- A6 Endocrine System
- A7 Cardiovascular System
- A8 Nervous System
- A9 Sense Organs
- A10 Tissues
- A11 Cells
- A12 Fluids and Secretions
- A13 Animal Structures
- A14 Stomatognathic System
- A15 Hemic and Immune Systems
- A16 Embryonic Structures
- A17 Integumentary System
- A18 Plant Structures
- A19 Fungal Structures
- A20 Bacterial Structures
- A21 Viral Structures

B Organisms

- B1 Eukaryota
- B2 Archaea
- B3 Bacteria
- B4 Viruses
- B5 Organism Forms

C Diseases

- C1 Bacterial Infections & Mycoses
- C2 Virus Diseases
- C3 Parasitic Diseases
- C4 Neoplasms
- C5 Musculoskeletal Diseases
- C6 Digestive System Diseases
- C7 Stomatognathic Diseases
- C8 Respiratory Tract Diseases
- C9 Otorhinolaryngologic Diseases
- C10 Nervous System Diseases
- C11 Eye Diseases
- C12 Male Urogenital Diseases
- C13 Female Urogenital Diseases & Pregnancy Complications
- C14 Cardiovascular Diseases
- C15 Hemic & Lymphatic Diseases
- C16 Congenital, Hereditary, & Neonatal Diseases & Abnormalities
- C17 Skin & Connective Tissue Diseases
- C18 Nutritional & Metabolic Diseases
- C19 Endocrine System Diseases
- C20 Immune System Diseases
- C21 Disorders of Environmental Origin
- C22 Animal Diseases
- C23 Pathologic Conditions,
- Signs, and Symptoms
- C24 Occupational Diseases
- C25 Substance-Related Disorders
- C26 Wounds and Injuries

D Chemicals & Drugs

- D1 Inorganic Chemicals
- D2 Organic Chemicals
- D3 Heterocyclic Compounds
- D4 Polycyclic Compounds
- D5 Macromolecular Substances
- D6 Hormones, Hormone Substitutes, & Hormone Antagonists
- D8 Enzymes, & Coenzymes
- D9 Carbohydrates
- D10 Lipids
- D12 Amino Acids, Peptides, & Proteins
- D13 Nucleic Acids, Nucleotides, & Nucleosides
- D20 Complex Mixtures
- D23 Biologic Factors
- D25 Biomedical and Dental Materials
- D26 Pharmaceutical Preparations
- D27 Chemical Actions and Uses

Other Categories Include:

Analytical, Diagnostic, & Therapeutic Techniques & Equipment

Psychiatry and Psychology

Biological Sciences

Natural Sciences

Anthropology, Education, Sociology & Social Phenomena

Technology, Industry, Agriculture

Information Science

Named Groups

Health Care

Publication Characteristics

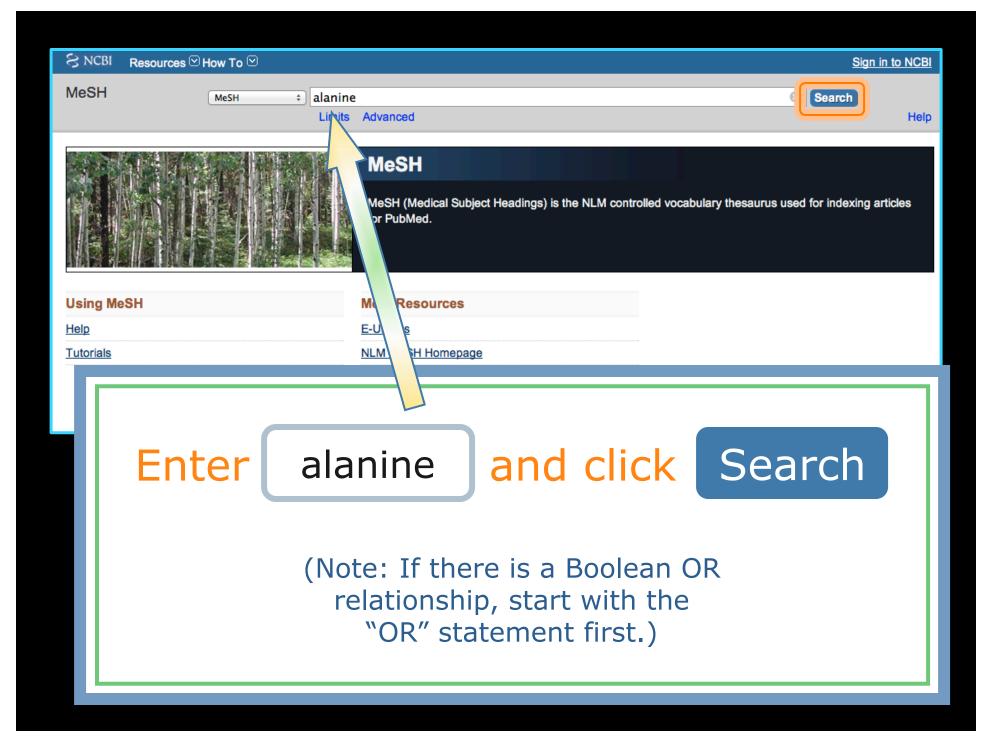
Geographic Locations

Create a Search Strategy Plan

Identify the question and key concepts:

alanine or threonine in the genetics of neoplasms

Write the search program using medical subject headings (MeSH)



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Μ	eSH MeSH alanine Save search Limits Advanced	Search Help
Di	splay Settings: ♡ Summary, 20 per page Send to: ♡	PubMed search builder
1.	Alanine A non-essential amino acid that occurs in high levels in its free state in plasma. It is produced from pyruvate by transamination. It is involved in sugar and acid metabolism, increases IMMUNITY, and provides energy for muscle tissue, BRAIN, and the CENTRAL NERVOUS SYSTEM.	Add to search builder AND +
2.	beta-Alanine An amino acid formed in vivo by the degradation of dihydrouracil and carnosine. Since neuronal uptake and neuronal receptor sensitivity to beta-alanine have been demonstrated, the compound may be a false transmitter replacing GAMMA-AMINOBUTYRIC ACID. A rare genetic disorder, hyper-beta-alaninemia, has been reported. Year introduced: 1992	Search PubMed Find related data Database: Select +
□ 3.	Alanine Transaminase An enzyme that catalyze Nomenclature, 1992) El Year introduced: 1998(*	Find items ails
	Term Definitions	

1) Check the box next to the MeSH heading.

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to 20 of 609

Search

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<< First < Prev Page 1 of 31 Next > Last >> Alanine 1. A non-essential amino acid that occurs in high levels in its free state in plasma. It is produced from pyruvate by transamination. It is involved in sugar and acid metabolism, increases IMMUNITY, and provides energy for muscle tissue, BRAIN, and the CENTRAL NERVOUS SYSTEM.

beta-Alanine

Display Set#

2. An amino acid formed in vivo by the degradation of dihydrouracil and carnosine. Since neuronal uptake and neuronal receptor sensitivity to beta-alanine have been demonstrated, the compound may be a false transmitter replacing GAMMA-AMINOBUTYRIC ACID. A rare genetic disorder, hyper-beta-alaninemia, has been reported. Year introduced: 1992

Alanine Transaminase

3. An enzyme that catalyzes the conversion of L-alanine and 2-oxoglutarate to pyruvate and L-glutamate. (From Enzyme Nomenclature, 1992) EC 2.6.1.2. Year introduced: 1998(1966)

2) Then click

Add to search builder

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Results: 1 to 20 of 228 Selected: 1 << First < Prev Page 1 of 12 Next > Last >>	("Alanine"[Mesh]) OR "Threonine" [Mesh]
 Threonine An essential amino acid occurring naturally in the L-form, which is the active form. It is found in eggs, milk, gelatin, and other proteins. 	
 Protein-Serine-Threonine Kinases A group of enzymes that catalyzes the phosphorylation of serine or threonine residues in proteins, with ATP or other nucleotides as phosphate donors. Year introduced: 1993 	Add to sear uilder OR ‡ Search Pu' d
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alanine OR threonine

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All MeSH Categories Diseases Category Neoplasms

<u>Cysts</u> <u>Arachnoid Cysts</u> <u>Bone Cysts</u> + <u>Branchioma</u> <u>Breast Cyst</u> Bronchogenic Cyst

Neoplasms by Site

Abdominal Neoplasms + Anal Glang Neoplasms Bone Neoplasms + Breast Neoplasms + Digestive System Neoplasms + Endocrine Gland Neoplasms + Eye Neoplasms + Head and Neck Neoplasms + Hematologic Neoplasms + Mammary Neoplasms, Animal + Nervous System Neoplasms + Pelvic Neoplasms Skin Neoplasms + Soft Tissue Neoplasms + Splenic Neoplasms Thoracic Neoplasms + Urogenital Neoplasms +

MeSH terms arranged in tree structure hierarchy

Terms are listed from broadest to narrowest. If Abdominal Neoplasms yields zero results, try using a broader term: Neoplasms by Site. <u>Neoplasms, Vascular Tissue</u> + <u>Nevi and Melanomas</u> + Odoptogenic Tumors +

Neoplasms by Site Abdominal Neoplasms + Anal Gland Neoplasms Bone Neoplasms + Breast Neoplasms + Digestive System Neoplasms + Endocrine Gland Neoplasms + Eye Neoplasms + Head and Neck Neoplasms + Hematologic Neoplasms + Mammary Neoplasms, Animal + Nervous System Neoplasms + Pelvic Neoplasms Skin Neoplasms + Soft Tissue Neoplasms + Splenic Neoplasms Thoracic Neoplasms + Urogenital Neoplasms +

> Carcinoma 256, Walker Carcinoma, Brown-Pearce Carcinoma, Ehrlich Tumor

Note:

PubMed automatically OR's indented terms under a broader term...

... *unless* you select the following command

epidemiology

physiopathology
 prevention and control

Do I

Restrict to MeSH Major Topic.

Do not include MeSH terms found below this term in the MeSH hierarchy.

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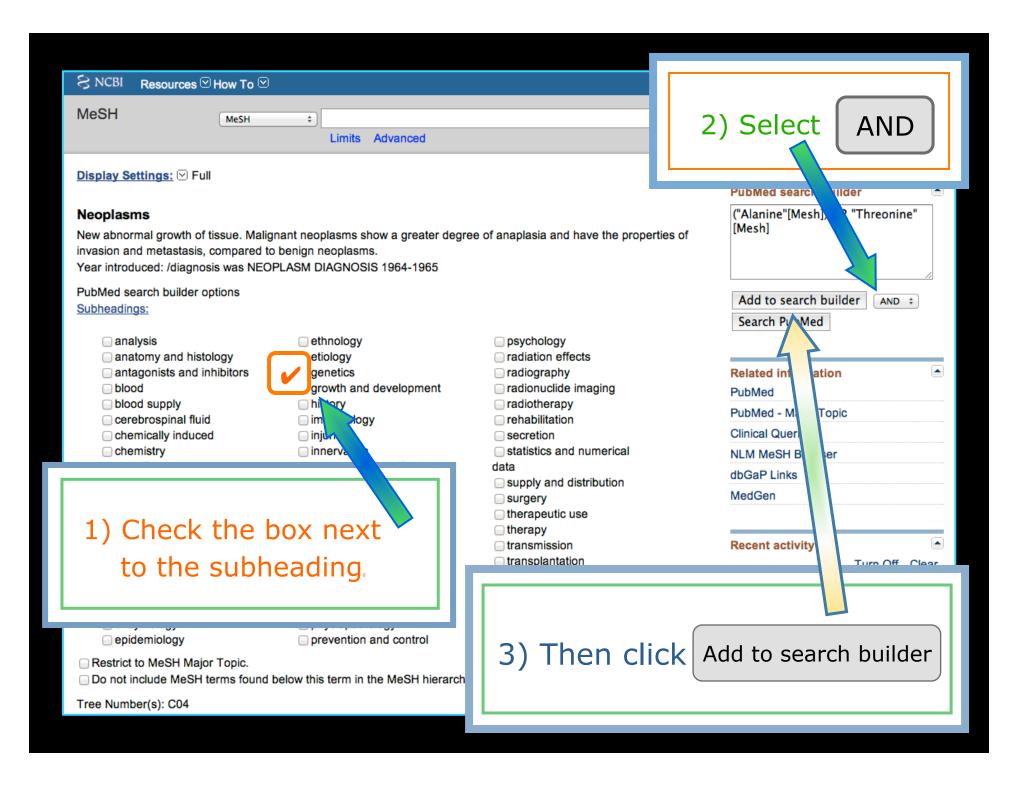
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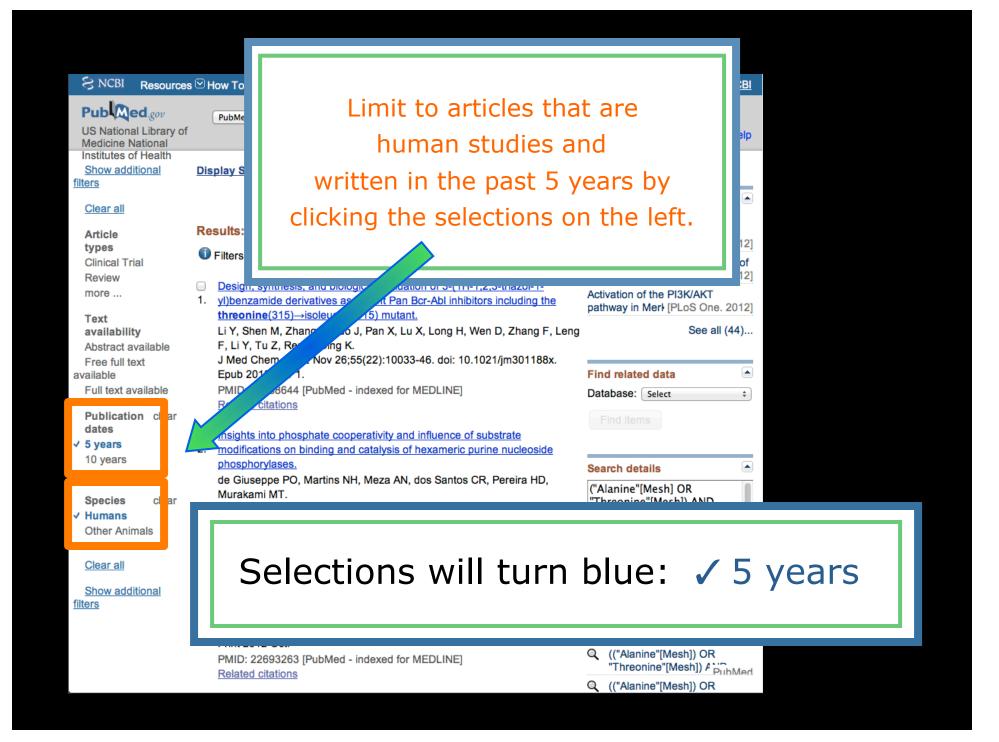
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Article types	Results: 1 to 20 of 64 << First < Prev	Insights into phosphate cooperativity an [PLoS One. 2012]
Clinical Trial Review more Text	 Design, synthesis, and biological evaluation of 3-(1H-1,2,3-triazol-1- <u>yl)benzamide derivatives as Potent Pan Bcr-Abl inhibitors including the</u> <u>threonine(315)</u>→isoleucine(315) mutant. 	The threonine protease activity of testes-specific p [PLoS One. 2012] Activation of the PI3K/AKT pathway in Merl [PLoS One. 2012]
availability Abstract available	Li Y. Shen M. Zhang Z. Luo J. Pan X. Lu X. Long H. Wen D. Zhang F. Leng F	See all (44)
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Clear all	PMID: 220 0 [PubMed - indexed for MEDLINE] Free PMC Article Relate atoms Cue of a pathogenic mutant human glucagon receptor by	AND ("2008/03/14"[PDat] : "2013/03/12"[PDat] AND Search See more

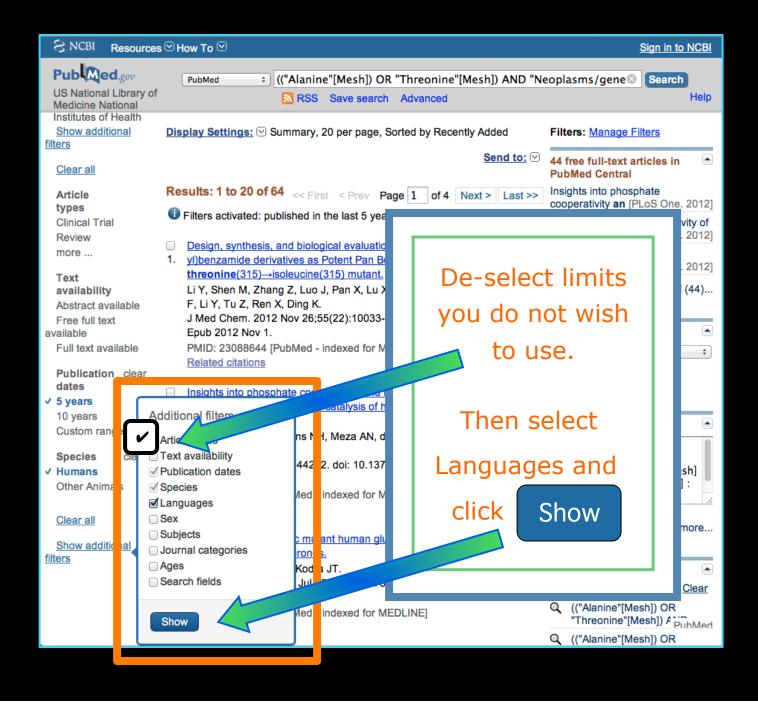
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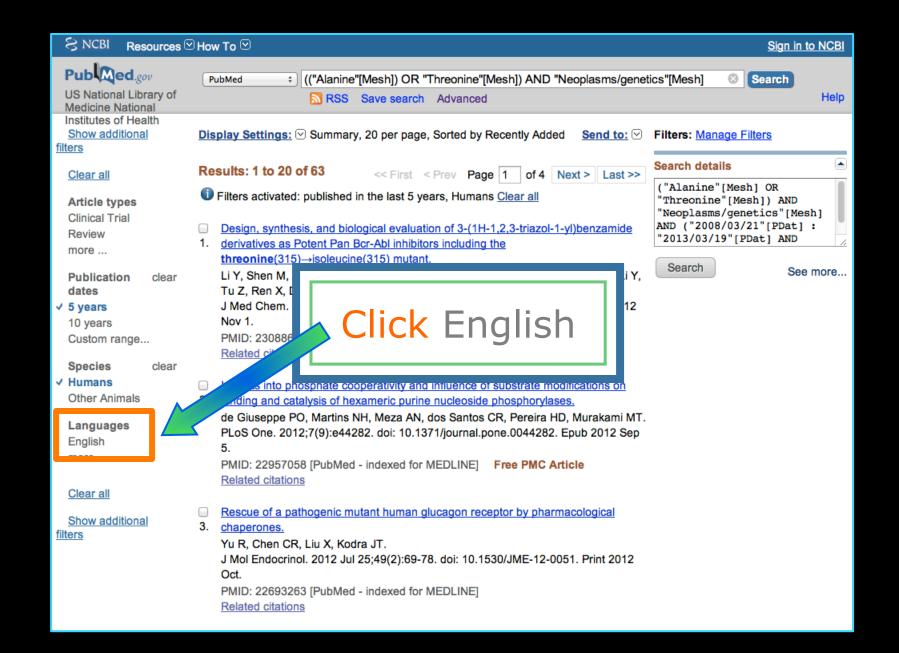
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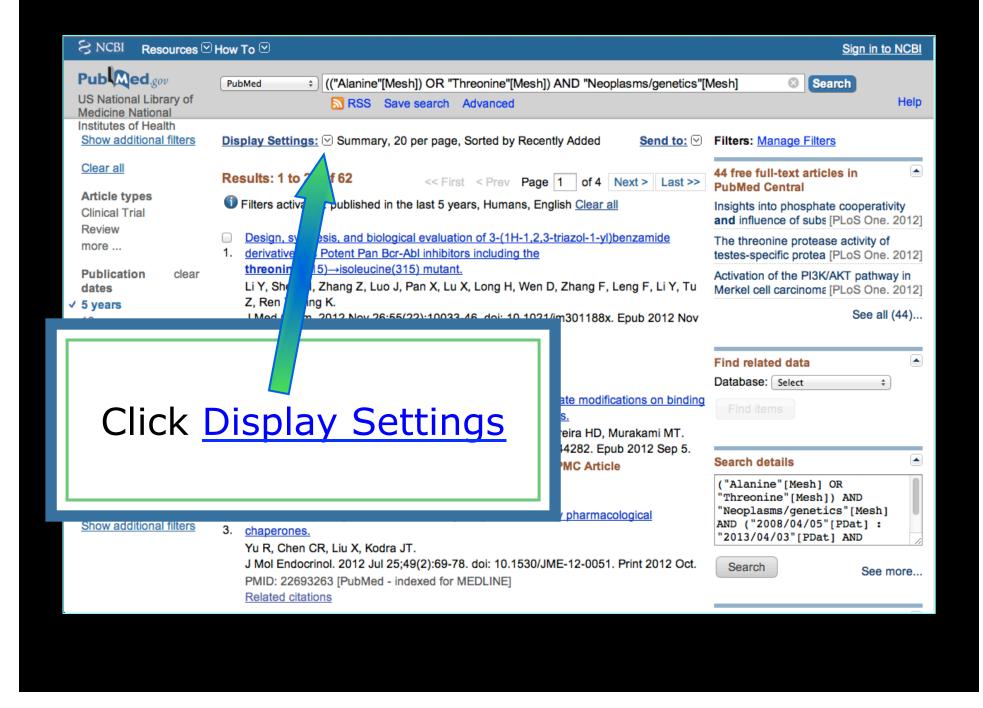


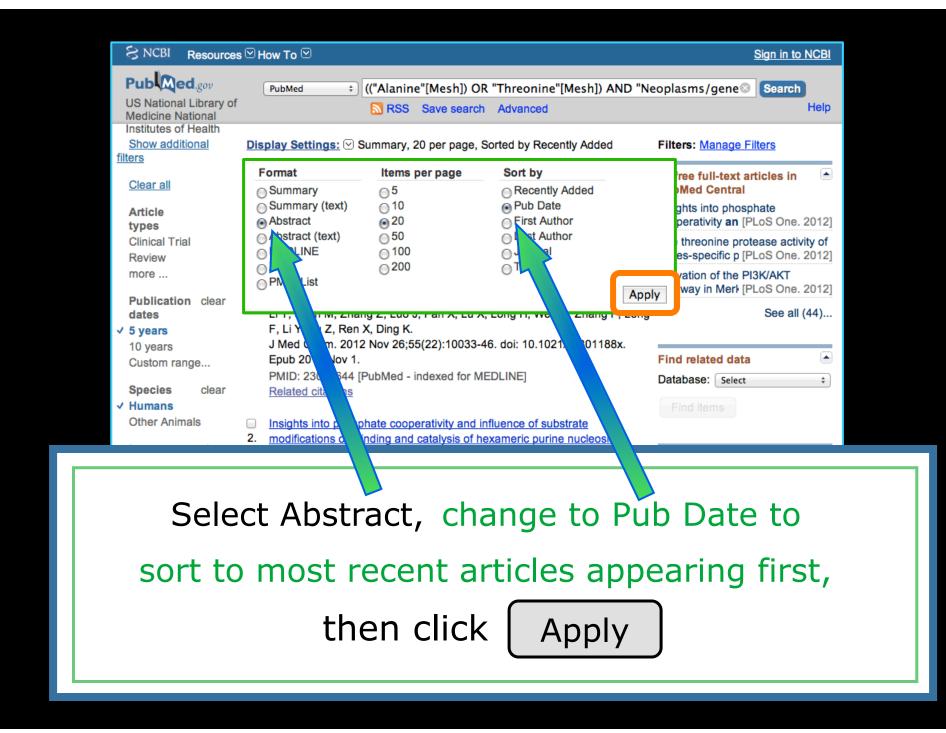


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Insights into phosphate cooperativity and influence of substrate modifications on binding and catalysis of hexameric purine nucleoside phosphorylases.

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Threonine/chemistry

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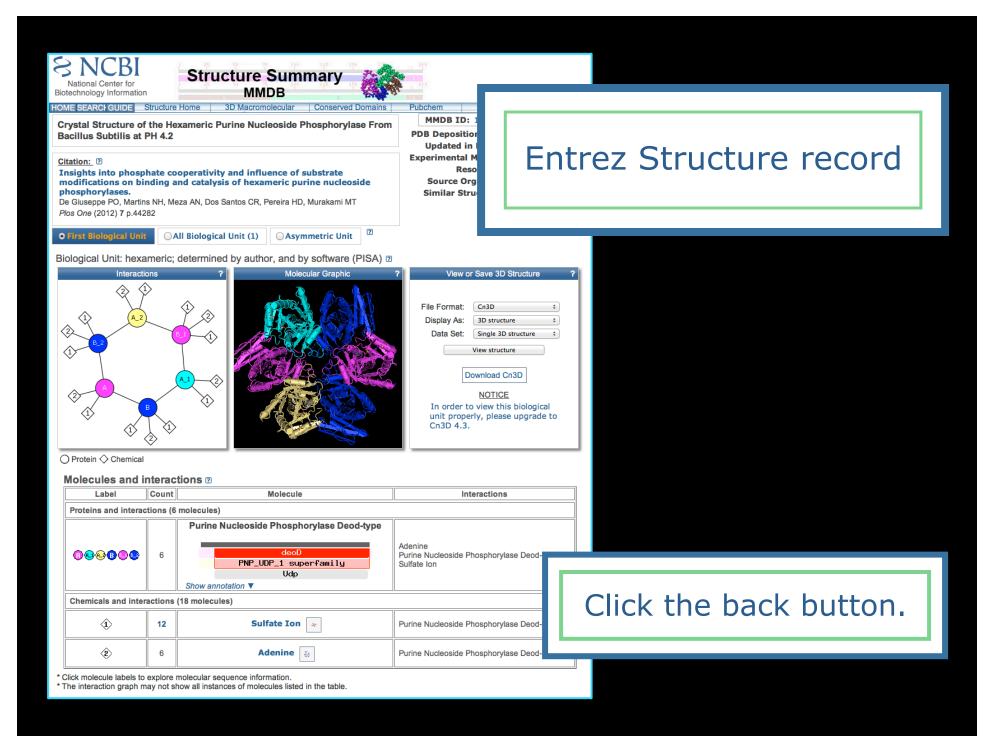
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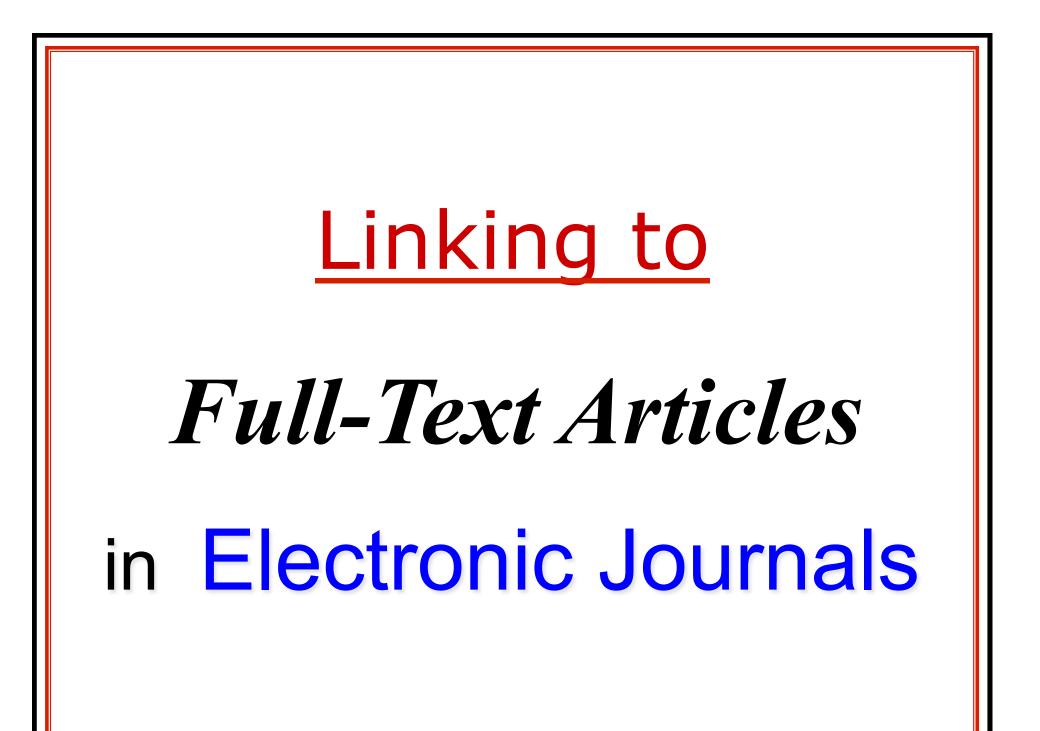
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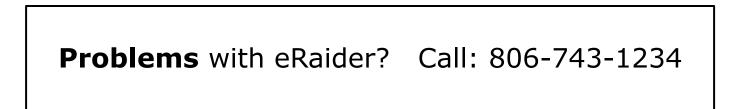
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Priscila O. de Giuseppe, ¹ Nadia H. Martins, ¹ Andreia N. Meza, ¹ Camila R. dos Santos, ¹ Humberto D'Muniz Pereira, ² and Mario T. Murakami ¹ , ²	Crystal structure of calf spleen purine nucleoside phosphorylase with two full trimers in the [J Mol Bi Structural analyses reveal two distinct families of pucleoside phosphorylases [Biochem 102]
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Insights into Phosphate Cooperativity and Influence of Substrate Modifications on Binding and Catalysis of Hexameric Purine Nucleoside Phosphorylases

Priscila O. de Giuseppe¹, Nadia H. Martins¹, Andreia N. Meza¹, Camila R. dos Santos¹, Humber D'Muniz Pereira², Mario T. Murakami¹*

1 Laboratório Nacional de Biociências (LNBio), Centro Nacional de Pesquisa em Energia e Materiais, Campinas, São Paulo, Brazil, 2 Instituto de Física de São de Cristalografia, Universidade de São Paulo, São Carlos, São Paulo, Brazil

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Competing Interests: The authors have declared that no competing interests exist.

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Introduction

Purine nucleoside phosphorylases (PNPs; EC 2.4.2.1) are versatile enzymes that catalyze the reversible phosphorolysis of purine (2'deoxy)ribonucleosides producing bases and (2'deoxy)ribose 1-phosphate [1]. Their key role in the purine salvage pathway made PNPs attractive targets for drug design against several pathogens, such as *Mycobactorium tuberculosis* [2,3], *Plasmolium fakipanm* [4–7], *Trichomasa saginati* [3–10] and *Schistosan annasoni* [11,12], which lacks the *de novo* pathway for purine nucleotides synthesis. Due to their catalytic function, PNPs have also been investigated for the synthesis of nucleoside analogues (NAs) [13] and the activation of prodrugs in anti-cancer gene therapies [14].

NAs can be used in the treatment of a range of human viral infections, such as those caused by HIV, herpesvins and hepatitis B/C virus [15–19]. They are among the first cytotoxic molecules to be used in the treatment of cancer [20] and have been studied as potential drugs against tuberculosis [21,22], makria [7,23], trichomoniasis [24] and schistosomiasis [25]. The chemical synthesis of these compounds is generally a costly process that includes several protection and deprotec [13,26]. This has encouraged the development of new n the synthesis of NAs using PNPs and other enzymes as 1 [13,27,28]. The main advantages of this approach are stereospecificity, regioselectivity and efficiency of enzyn employment usually dispenses group protection and p steps, optimizing the process [13].

The differences in substrate specificity regarding tr hexameric PNPs have allowed the development of su herapies strategies against solid tumosr [14,29]. Trin are mainly found in mammalian species and are s guanine and hypoxanthine (2'-deoxy)ribonuclcoside hexameric PNPs are prevalent in bacteria and accept well as guanine and hypoxanthine (2'-deoxy)ribonuc substrates [1]. Thus, nontoxic adenosine analogues, poor substrates for human PNP, can be cleaved to cyto specifically in tumor cells transfected with the bacterial

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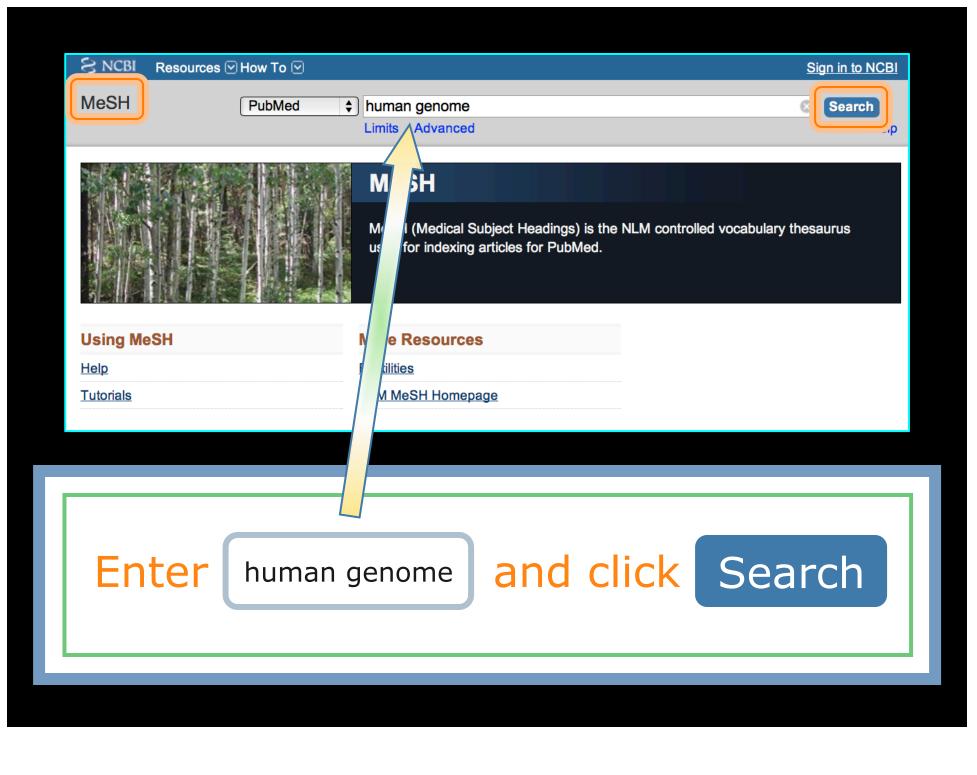
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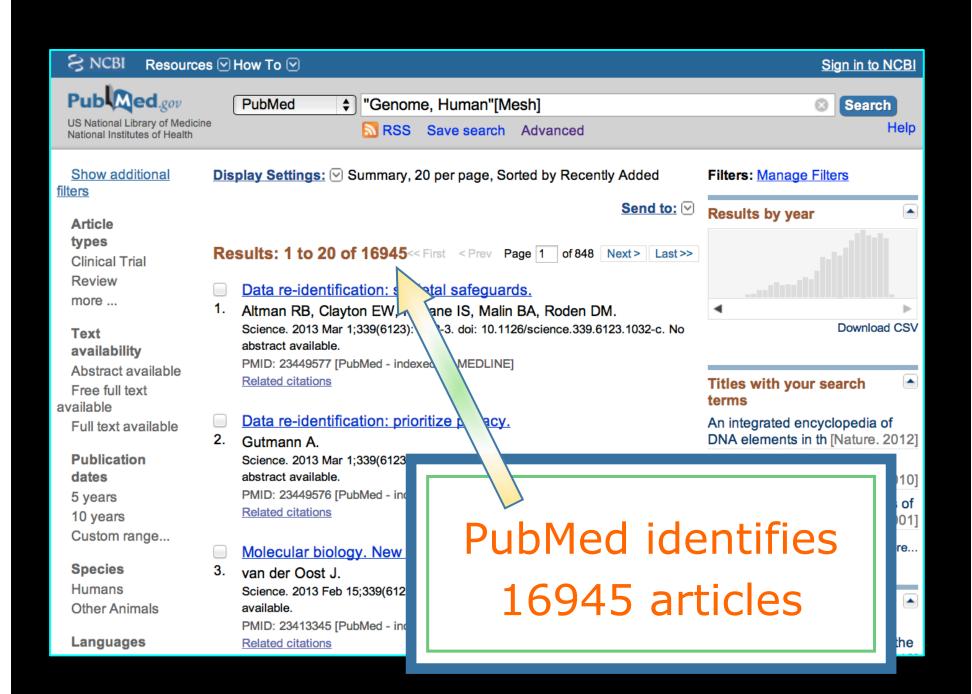
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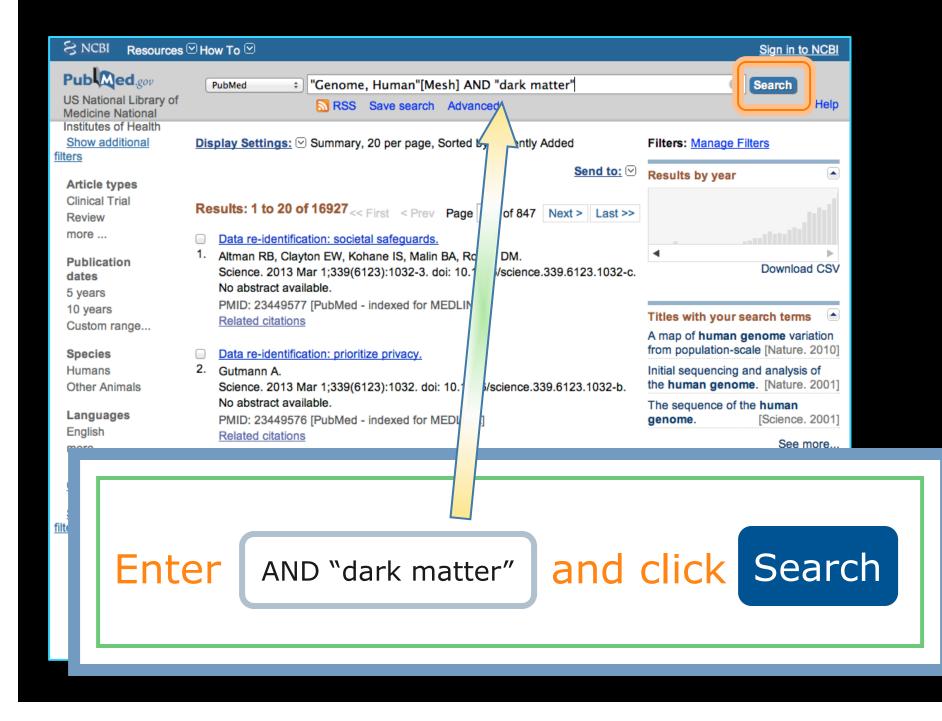
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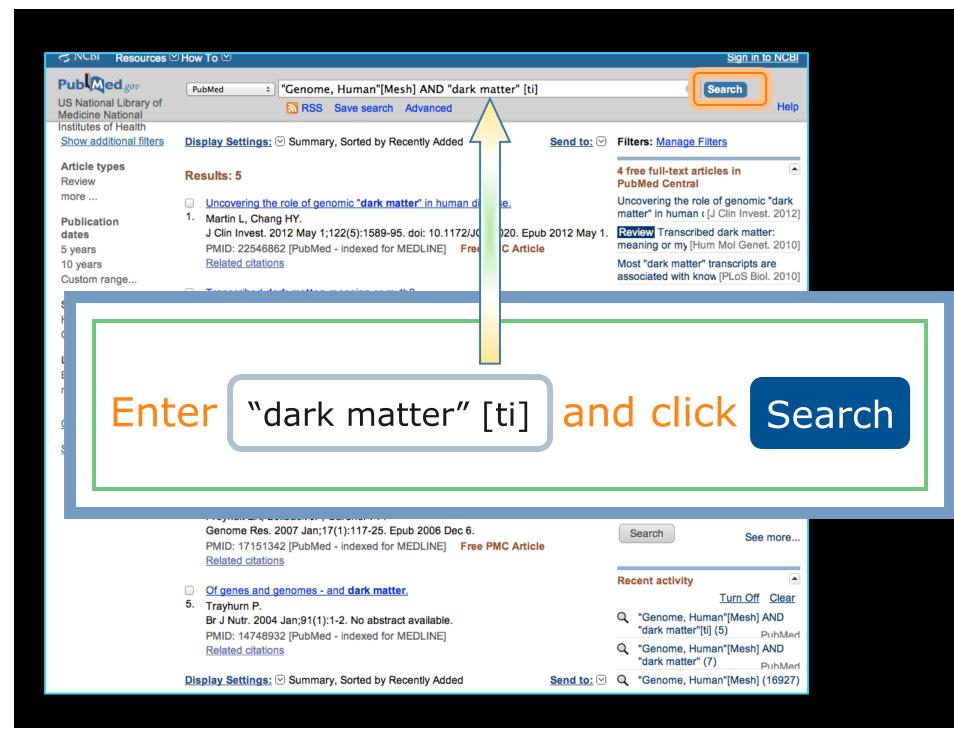


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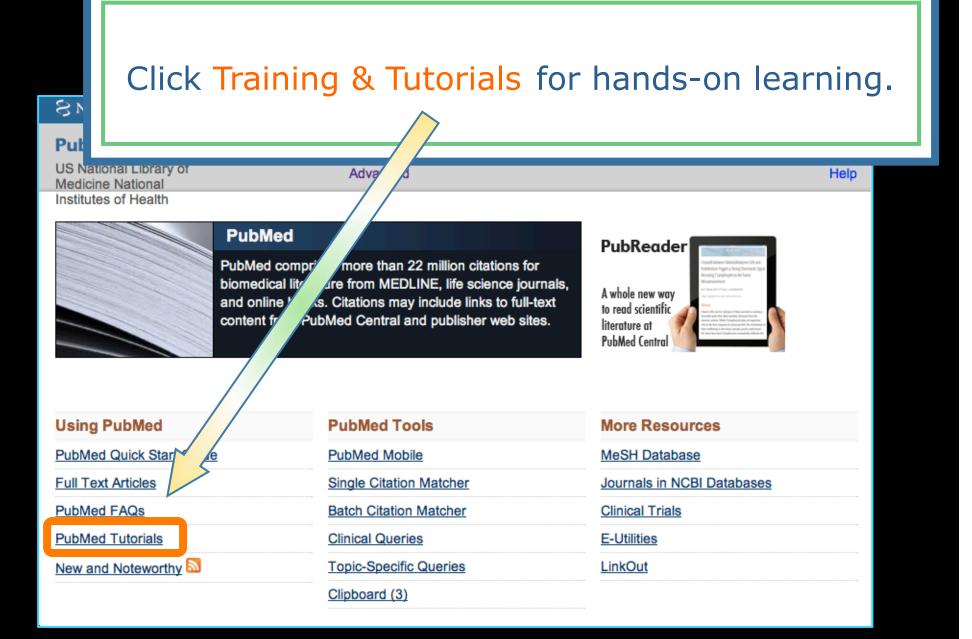
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