Bibliographie/Veille

Cas Clinique

- Patiente de 8 ans
- Douleurs musculaires
- Faiblesse musculaire
- Éruption cutané
- Pas de fièvre

- Hypothèse diagnostique ?

PubMed.gov

faiblesse musculaire et eruption cutané

Search

Advanced Create alert Create RSS User Guide

Sorted by: Best match Display options

MY NCBI FILTERS

TEXT AVAILABILITY

- [ ] Abstract
- [ ] Free full text
- [ ] Full text

ARTICLE ATTRIBUTE

- [ ] Associated data

No results were found.

First Prev Page 0 of 0 Next Last

Your search was processed without automatic term mapping because it retrieved zero results.

First Prev Page 0 of 0 Next Last

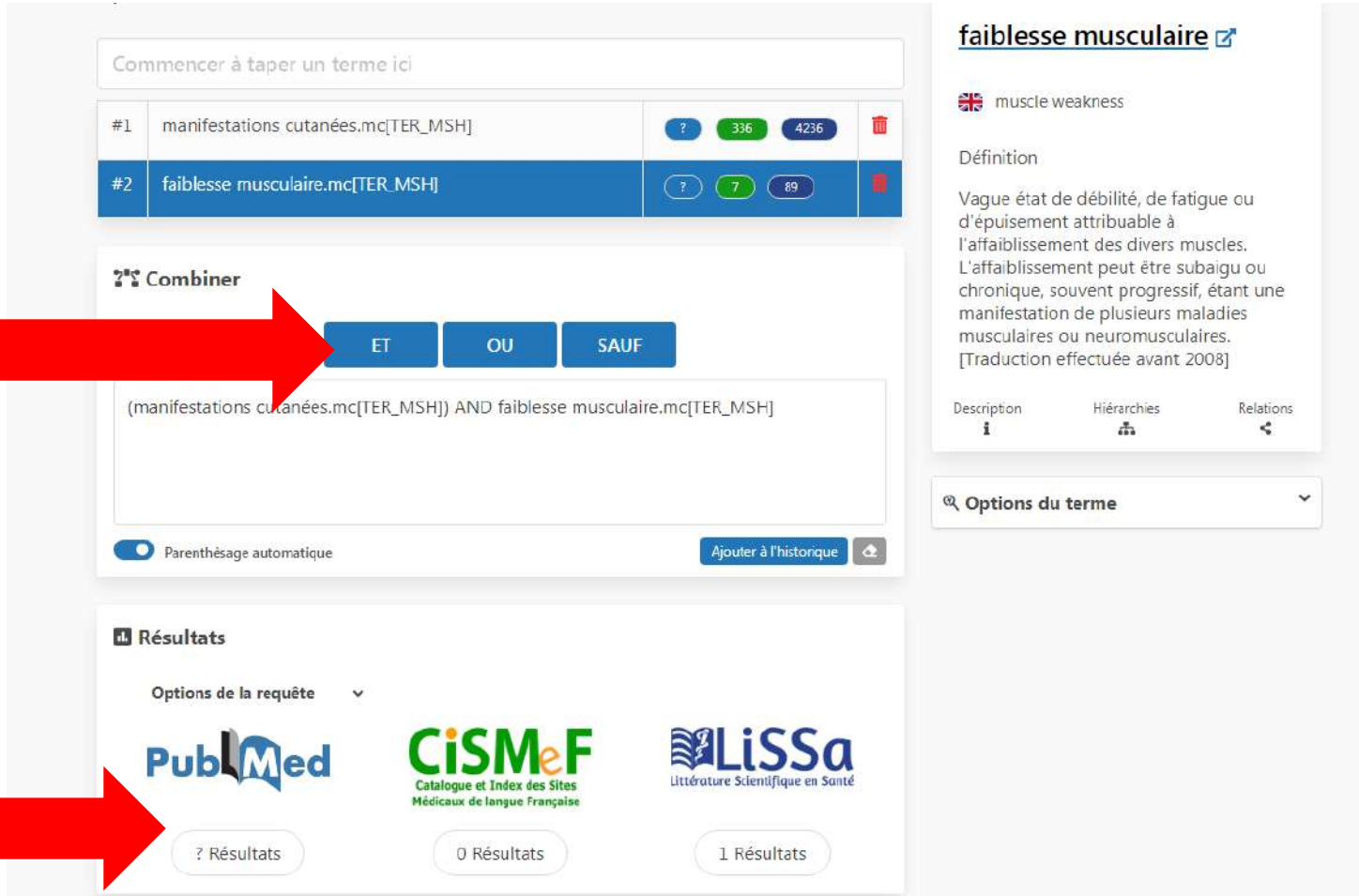
Trouver les bons mots clefs

- https://www.cismef.org/cismef/

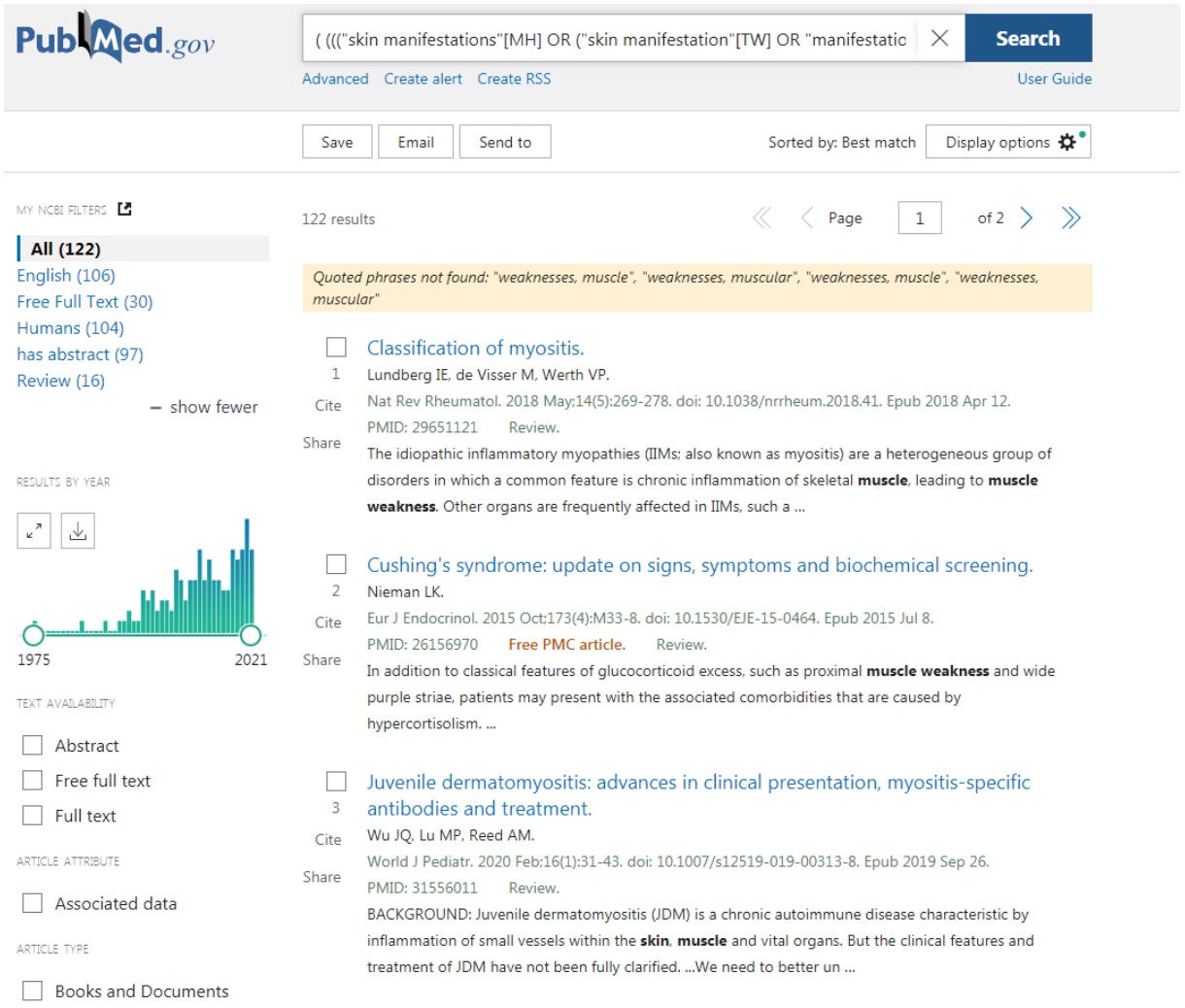
CRBM


Constructeur de Requêtes Bibliographiques Médicales
Créer simplement des requêtes bibliographiques complexes en combinant des termes médicaux de langue française (MeSH)
Chercher un terme
Commencer à taper un terme ici
#1
manifestations cutanées.mc[TER_MSH]
?
336
4236
Combiner
Ajouter
Parenthèsage automatique
Ajouter à l'historique
manifestations cutanées
skin manifestations
Dermatologic disorders attendant upon non-dermatologic disease or injury.
Description
Hiérarchies
Relations
Options du terme

CRBM


Commencer à taper un terme ici
#1 manifestations cutanées.mc[TER_MSH] ? 336 4236
#2 faiblesse musculaire.mc[TER_MSH] ? 7 89
Combiner
ET
OU
SAUF
(manifestations cutanées.mc[TER_MSH]) AND faiblesse musculaire.mc[TER_MSH]
Parenthésage automatique
Ajouter à l'historique
Résultats
Options de la requête
PubMed
CiSMeF
Catalogue et Index des Sites Médicaux de langue Française
LiSSa
Littérature Scientifique en Santé
? Résultats
0 Résultats
1 Résultats
faiblesse musculaire
muscle weakness
Définition
Vague état de débilité, de fatigue ou d'épuisement attribuable à l'affaiblissement des divers muscles. L'affaiblissement peut être subaigu ou chronique, souvent progressif, étant une manifestation de plusieurs maladies musculaires ou neuromusculaires. [Traduction effectuée avant 2008]
Description
Hiérarchies
Relations
Options du terme

pubmed


PubMed.gov
((((("skin manifestations"[MH] OR ("skin manifestation"[TW] OR "manifestatic
Search
Advanced
Create alert
Create RSS
User Guide
Save
Email
Send to
Sorted by: Best match
Display options
MY NCBI FILTERS
All (122)
English (106)
Free Full Text (30)
Humans (104)
has abstract (97)
Review (16)
show fewer
RESULTS BY YEAR
1975
2021
TEXT AVAILABILITY
Abstract
Free full text
Full text
ARTICLE ATTRIBUTE
Associated data
ARTICLE TYPE
Books and Documents
122 results
Page 1 of 2
Quoted phrases not found: "weaknesses, muscle", "weaknesses, muscular", "weaknesses, muscle", "weaknesses, muscular"
Classification of myositis.
1
Lundberg IE, de Visser M, Werth VP.
Cite
Share
Nat Rev Rheumatol. 2018 May;14(5):269-278. doi: 10.1038/nrrheum.2018.41. Epub 2018 Apr 12.
PMID: 29651121 Review.
The idiopathic inflammatory myopathies (IIMs; also known as myositis) are a heterogeneous group of disorders in which a common feature is chronic inflammation of skeletal muscle, leading to muscle weakness. Other organs are frequently affected in IIMs, such a ...
Cushing's syndrome: update on signs, symptoms and biochemical screening.
2
Nieman LK.
Cite
Share
Eur J Endocrinol. 2015 Oct;173(4):M33-8. doi: 10.1530/EJE-15-0464. Epub 2015 Jul 8.
PMID: 26156970 Free PMC article. Review.
In addition to classical features of glucocorticoid excess, such as proximal muscle weakness and wide purple striae, patients may present with the associated comorbidities that are caused by hypercortisolism. ...
Juvenile dermatomyositis: advances in clinical presentation, myositis-specific antibodies and treatment.
3
Wu JQ, Lu MP, Reed AM.
Cite
Share
World J Pediatr. 2020 Feb;16(1):31-43. doi: 10.1007/s12519-019-00313-8. Epub 2019 Sep 26.
PMID: 31556011 Review.
BACKGROUND: Juvenile dermatomyositis (JDM) is a chronic autoimmune disease characteristic by inflammation of small vessels within the skin, muscle and vital organs. But the clinical features and treatment of JDM have not been fully clarified. ...We need to better un ...

Pubmed

Review > Nat Rev Rheumatol. 2018 May;14(5):269-278. doi: 10.1038/nrrheum.2018.41. Epub 2018 Apr 12.

Classification of myositis

Ingrid E Lundberg [1 2], Marianne de Visser [3], Victoria P Werth [4]

Affiliations + expand

PMID: 29651121 DOI: 10.1038/nrrheum.2018.41

Abstract

The idiopathic inflammatory myopathies (IIMs; also known as myositis) are a heterogeneous group of disorders in which a common feature is chronic inflammation of skeletal **muscle**, leading to **muscle weakness**. Other organs are frequently affected in IIMs, such as the **skin**, joints, lungs, gastrointestinal tract and heart, contributing to morbidity and mortality. Currently, IIMs are most often subclassified into polymyositis, dermatomyositis and inclusion body myositis, but this subclassification has limitations as these subgroups often have overlapping clinical and histopathological features, and outcomes vary within the subgroups; additionally, subgroups without considerable myopathy are not included. A new way of subgrouping patients could be on the basis of the presence of myositis-specific autoantibodies. These autoantibodies are associated with distinct clinical features and, moreover, can help to identify subsets of IIMs in which extramuscular symptoms, such as **skin manifestations**, arthritis or interstitial lung disease, might be the presenting or predominant feature when **muscle** symptoms are mild or absent. The recognition that subphenotypes with single-organ involvement other than muscles exist is important for identifying patients with early disease, for clinical care demanding team management and in designing clinical studies to improve our understanding of this heterogeneous disease to develop new therapies.

Accès grâce au CHC

Published: 12 April 2018

Classification of myositis

Ingrid E. Lundberg, Marianne de Visser & Victoria P. Werth

Nature Reviews Rheumatology **14**, 269–278 (2018) | Cite this article

Key Points

- The clinical spectrum of idiopathic inflammatory myopathies (IIMs) has evolved from diseases in which muscle weakness was the main manifestation to systemic inflammatory diseases with multiple organ involvement.

- The EULAR–ACR classification criteria for adult and juvenile IIMs and their major subgroups capture patients with the typical skin rash of dermatomyositis without muscle weakness (amyopathic dermatomyositis).

Associated Content

Focus

Focus on Myositis

Pubmed

Review > Nat Rev Rheumatol. 2018 May;14(5):269-278. doi: 10.1038/nrrheum.2018.41. Epub 2018 Apr 12.

Classification of myositis

Ingrid E Lundberg [1 2], Marianne de Visser [3], Victoria P Werth [4]

Affiliations + expand

PMID: 29651121 DOI: 10.1038/nrrheum.2018.41

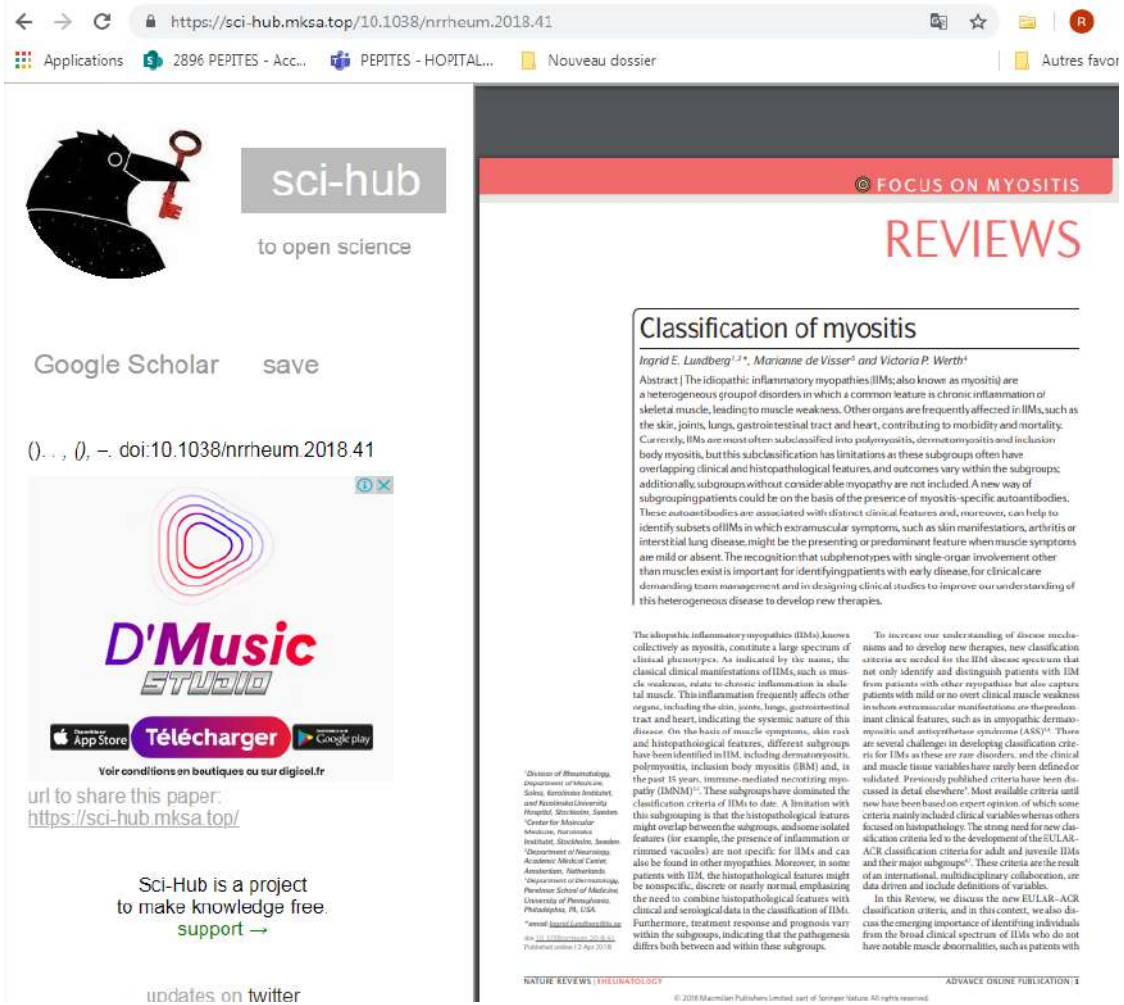
Abstract

The idiopathic inflammatory myopathies (IIMs; also known as myositis) are a heterogeneous group of disorders in which a common feature is chronic inflammation of skeletal **muscle**, leading to **muscle weakness**. Other organs are frequently affected in IIMs, such as the **skin**, joints, lungs, gastrointestinal tract and heart, contributing to morbidity and mortality. Currently, IIMs are most often subclassified into polymyositis, dermatomyositis and inclusion body myositis, but this subclassification has limitations as these subgroups often have overlapping clinical and histopathological features, and outcomes vary within the subgroups; additionally, subgroups without considerable myopathy are not included. A new way of subgrouping patients could be on the basis of the presence of myositis-specific autoantibodies. These autoantibodies are associated with distinct clinical features and, moreover, can help to identify subsets of IIMs in which extramuscular symptoms, such as **skin manifestations**, arthritis or interstitial lung disease, might be the presenting or predominant feature when **muscle** symptoms are mild or absent. The recognition that subphenotypes with single-organ involvement other than muscles exist is important for identifying patients with early disease, for clinical care demanding team management and in designing clinical studies to improve our understanding of this heterogeneous disease to develop new therapies.

Le coté obscure : SCI-HUB

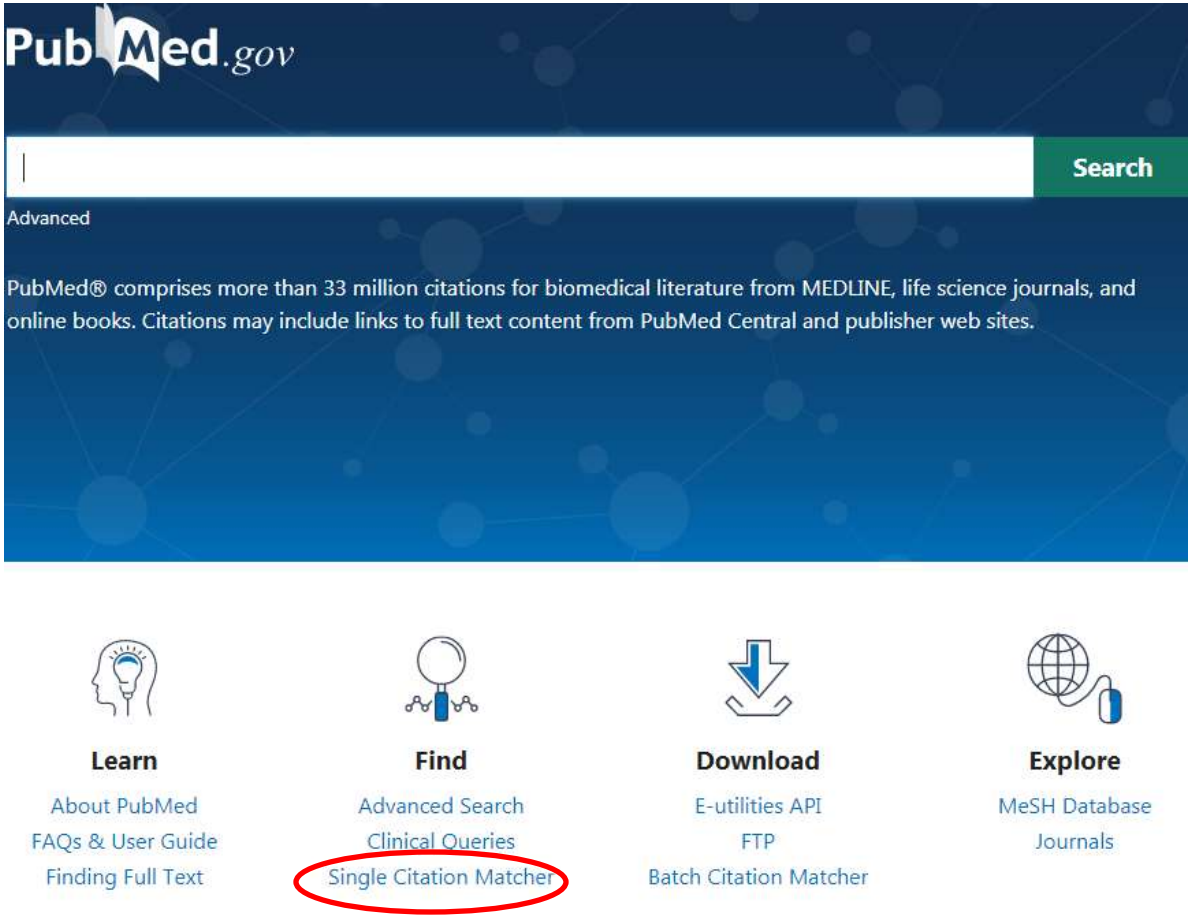
- https://sci-hub.mksa.top/

SCI-HUB

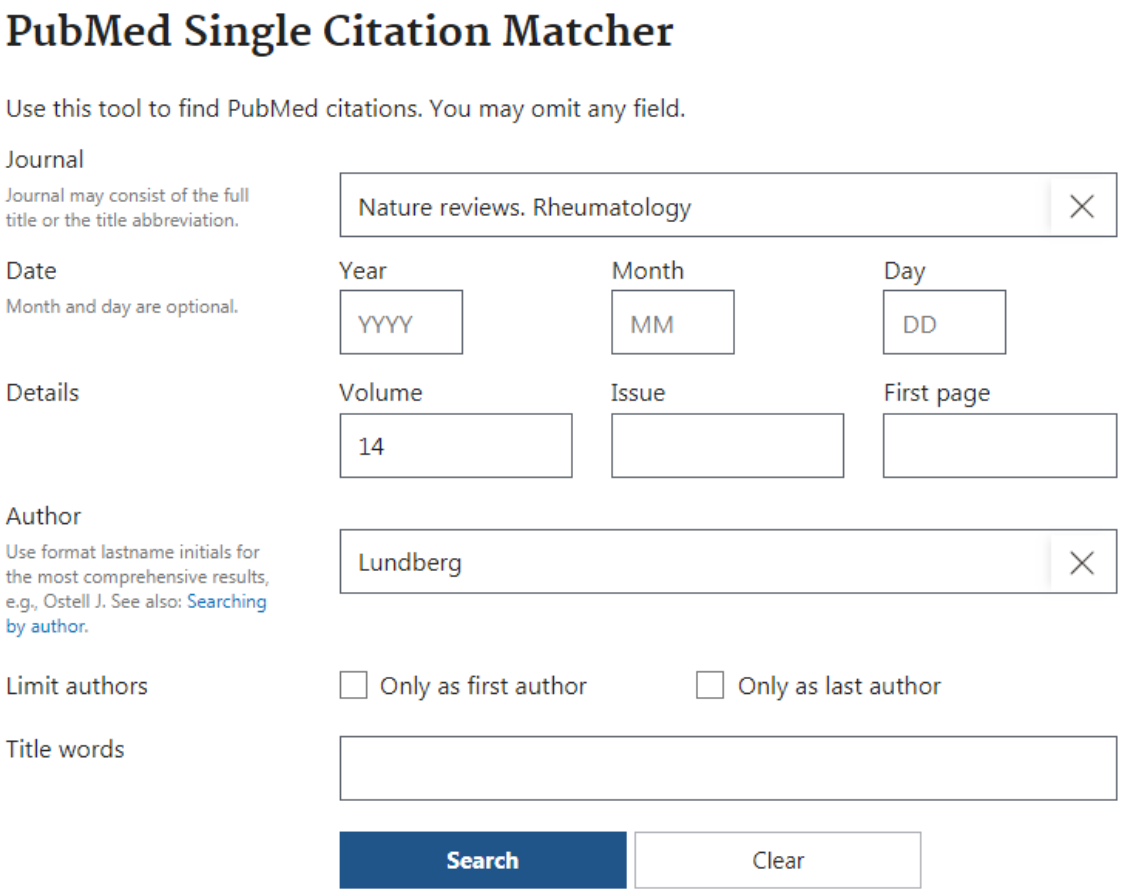

https://sci-hub.mksa.top/10.1038/nrrheum.2018.41
Applications
2896 PEPITES - Acc...
PEPITES - HOPITAL...
Nouveau dossier
Autres favor
sci-hub
to open science
Google Scholar save
(). . , (), –. doi:10.1038/nrrheum.2018.41
D'Music
STUDIO
App Store
Télécharger
Google play
Voir conditions en boutiques ou sur digicel.fr
url to share this paper:
https://sci-hub.mksa.top/
Sci-Hub is a project
to make knowledge free.
support →
updates on twitter
FOCUS ON MYOSITIS
REVIEWS
Classification of myositis
Ingrid E. Lundberg, Marianne de Visser and Victoria P. Werth
Abstract | The idiopathic inflammatory myopathies (IIMs; also known as myositis) are a heterogeneous group of disorders in which a common feature is chronic inflammation of skeletal muscle, leading to muscle weakness. Other organs are frequently affected in IIMs, such as the skin, joints, lungs, gastrointestinal tract and heart, contributing to morbidity and mortality. Currently, IIMs are most often subclassified into polymyositis, dermatomyositis and inclusion body myositis, but this subclassification has limitations as these subgroups often have overlapping clinical and histopathological features, and outcomes vary within the subgroups; additionally, subgroups without considerable myopathy are not included. A new way of subgrouping patients could be on the basis of the presence of myositis-specific autoantibodies. These autoantibodies are associated with distinct clinical features and, moreover, can help to identify subsets of IIMs in which extramuscular symptoms, such as skin manifestations, arthritis or interstitial lung disease, might be the presenting or predominant feature when muscle symptoms are mild or absent. The recognition that subphenotypes with single-organ involvement other than muscles exist is important for identifying patients with early disease, for clinical care demanding team management and in designing clinical studies to improve our understanding of this heterogeneous disease to develop new therapies.
The idiopathic inflammatory myopathies (IIMs), known collectively as myositis, constitute a large spectrum of clinical phenotypes. As indicated by the name, the classical clinical manifestations of IIMs, such as muscle weakness, relate to chronic inflammation in skeletal muscle. This inflammation frequently affects other organs, including the skin, joints, lungs, gastrointestinal tract and heart, indicating the systemic nature of this disease. On the basis of muscle symptoms, skin rash and histopathological features, different subgroups have been identified in IIM, including dermatomyositis, polymyositis, inclusion body myositis (IBM) and, in the past 15 years, immune-mediated necrotizing myopathy (IMNM)[1–3]. These subgroups have dominated the classification criteria of IIMs to date. A limitation with this subgrouping is that the histopathological features might overlap between the subgroups, and some isolated features (for example, the presence of inflammation or rimmed vacuoles) are not specific for IIMs and can also be found in other myopathies. Moreover, in some patients with IIM, the histopathological features might be nonspecific, discrete or nearly normal, emphasizing the need to combine histopathological features with clinical and serological data in the classification of IIMs. Furthermore, treatment response and prognosis vary within the subgroups, indicating that the pathogenesis differs both between and within these subgroups.
To increase our understanding of disease mechanisms and to develop new therapies, new classification criteria are needed for the IIM disease spectrum that not only identify and distinguish patients with IIM from patients with other myopathies but also capture patients with mild or no overt clinical muscle weakness in whom extramuscular manifestations are the predominant clinical features, such as in amyopathic dermatomyositis and antisynthetase syndrome (ASS)[4]. There are several challenges in developing classification criteria for IIMs as these are rare disorders, and the clinical and muscle tissue variables have rarely been defined or validated. Previously published criteria have been discussed in detail elsewhere[5]. Most available criteria until now have been based on expert opinion, of which some criteria mainly included clinical variables whereas others focused on histopathology. The strong need for new classification criteria led to the development of the EULAR–ACR classification criteria for adult and juvenile IIMs and their major subgroups[6]. These criteria are the result of an international, multidisciplinary collaboration, are data driven and include definitions of variables.
In this Review, we discuss the new EULAR–ACR classification criteria, and in this context, we also discuss the emerging importance of identifying individuals from the broad clinical spectrum of IIMs who do not have notable muscle abnormalities, such as patients with
NATURE REVIEWS | RHEUMATOLOGY
ADVANCE ONLINE PUBLICATION | 1

Single citation matcher

- Lundberg, I., de Visser, M. & Werth, V. Classification of myositis. *Nat Rev Rheumatol* **14,**

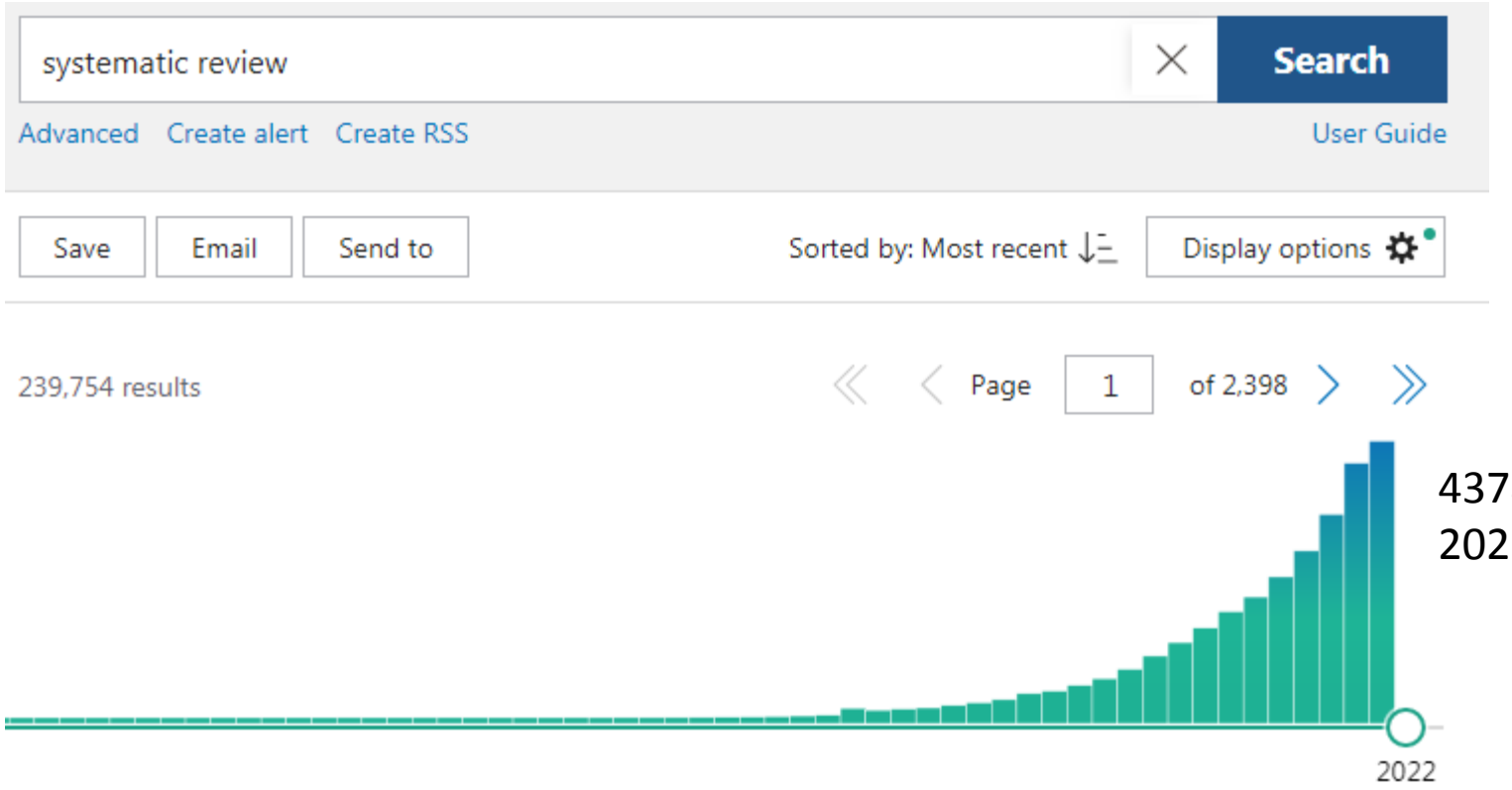

PubMed.gov
Search
Advanced
PubMed® comprises more than 33 million citations for biomedical literature from MEDLINE, life science journals, and online books. Citations may include links to full text content from PubMed Central and publisher web sites.
Learn
About PubMed
FAQs & User Guide
Finding Full Text
Find
Advanced Search
Clinical Queries
Single Citation Matcher
Download
E-utilities API
FTP
Batch Citation Matcher
Explore
MeSH Database
Journals

Single citation match



Pubmed-Medline

- 33 millions de références
- 30 000 revues



43780 en 2021

Pubmed-Medline

- MESH : Medical Subject Headings

Le MeSH comprend les 16 catégories thématiques suivantes.

- Anatomie [A]
- Organismes [B]
- Maladies [C]
- Produits chimiques et pharmaceutiques [D]
- Techniques et équipements analytiques, diagnostiques et thérapeutiques [E]
- Psychiatrie et psychologie [F]
- Phénomènes et processus [G]
- Disciplines et professions [H]
- Anthropologie, enseignement, sociologie et phénomènes sociaux [I]
- Technologie, industrie et agriculture [J]
- Sciences humaines [K]
- Sciences de l'information [L]
- Groupes individualisés [M]
- Soins de santé [N]
- Caractéristiques d'une publication [V]
- Lieux géographiques [Z]

MESH Bilingue

- http://mesh.inserm.fr/FrenchMesh/

Recherche simple

serpent | Rechercher

Recherche avancée

Descripteur français	contient		et
Descripteur français	contient		et
Descripteur français	contient		et
Descripteur français	contient		

Réinitialiser | Rechercher

Descripteurs trouvés

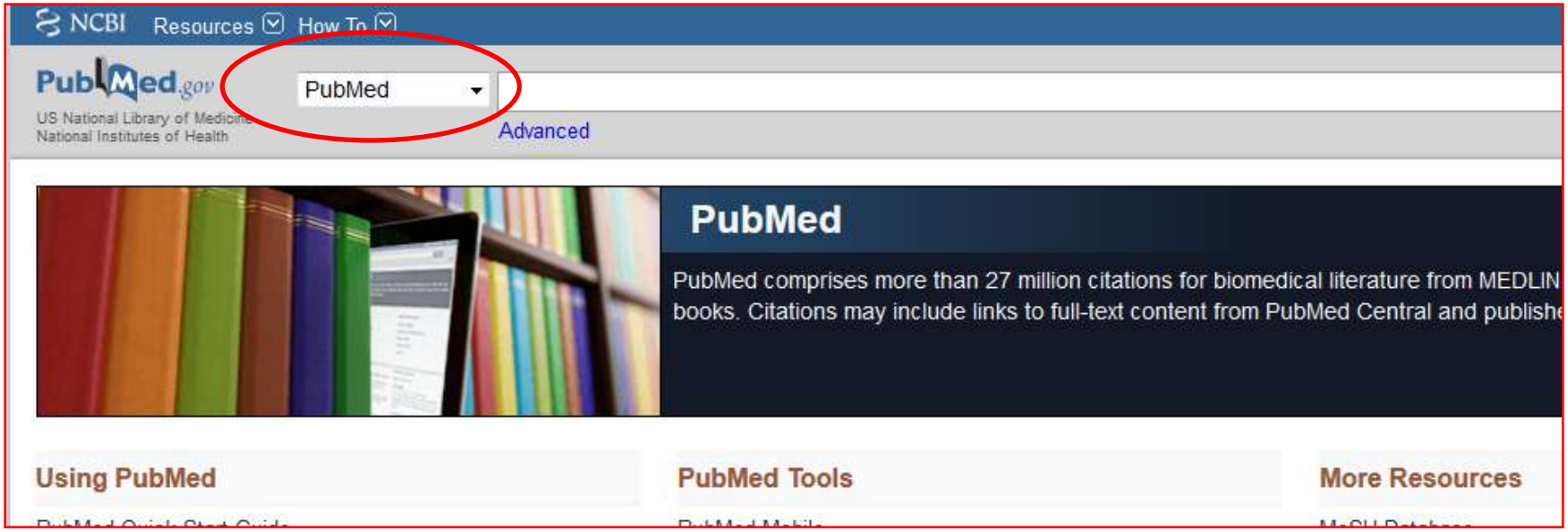
5 document(s) trouvé(s) en 2 ms.

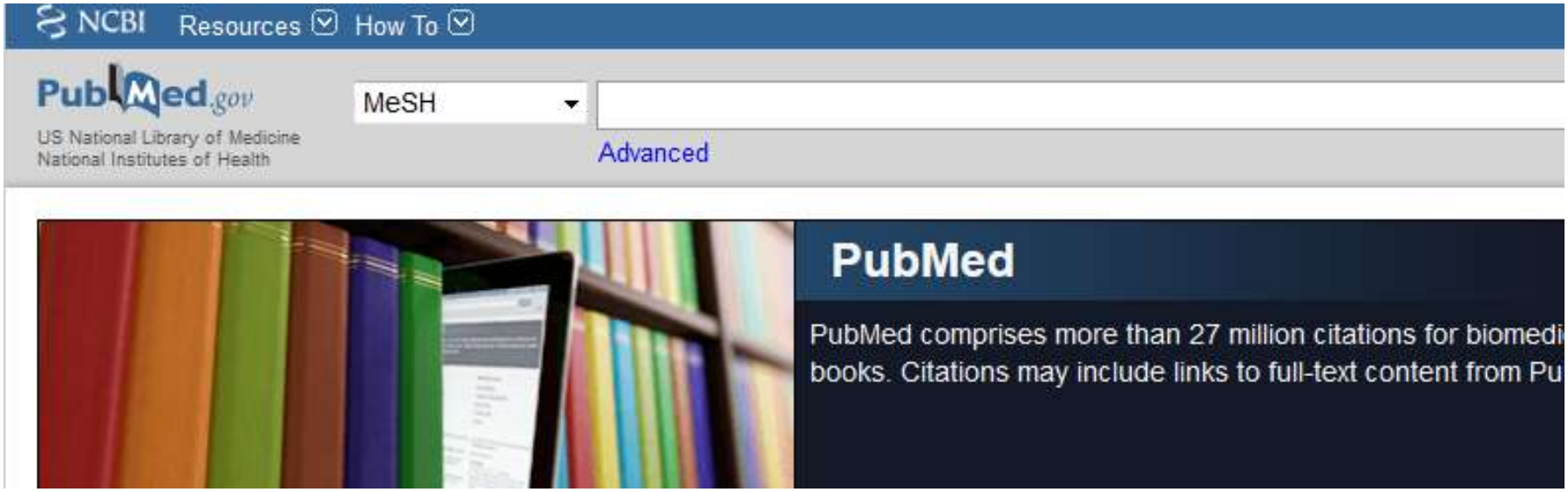
Descripteur français	Terme(s) français	Descripteur américain	Terme(s) américain(s)	Code(s) d'arborescence	Localisation
Morsures de **serpent**	Envenimation par les **serpents** /	Snake Bites	Bite, Snake / Bites, Snake / Snake Bite / Snakebites / Snakebite / Snake Envenomation / Envenomation, Snake / Envenomations, Snake / Snake Envenomations /	C25.723.127.442 C26.176.724	Arborescence
Venins de **serpent**		Snake Venoms	Venoms, Snake / Snake Venom / Venom, Snake /	D20.888.850 D23.946.833.850	Arborescence
Trichosanthes	Trichosanthes kirilowii / **Serpent** végétal /	Trichosanthes	Trichosanthe / Trichosanthes kirilowii / Trichosanthes kirilowius / kirilowius, Trichosanthes / Gourd, Snake / Snake Gourd / Gourds, Snake / Snake Gourds /	B01.650.940.800.575.100.300.827	Arborescence
Venins de crotalidé	Crotaline / Venin de **serpent** à sonnette / Venin de crotale / Crotamine / Crotactine / Crotalotoxine /	Crotalid Venoms	Venoms, Crotalid / Pit Viper Venoms / Venoms, Pit Viper / Pit Viper Venom / Venom, Pit Viper / Viper Venom, Pit / Crotalid Venom / Venom, Crotalid / Crotalin / Crotaline Snake Venom / Snake Venom, Crotaline / Venom, Crotaline Snake / Rattlesnake Venoms / Venoms, Rattlesnake / Rattlesnake Venom / Venom, Rattlesnake / Crotamin / Crotactin / Crotalotoxin /	D20.888.850.960.200 D23.946.833.850.960.200	Arborescence
Elapidae	Élapidés / Hydrophiinae / Lauticaudinae / Lauticaudus / Dendroaspis / Mambas / Micrurus / **Serpent** corail / Hemachatus / Haemachatus / Naja haje / Cobra d'Egypte / Cobra égyptien / Naja naja / Cobra asiatique / Cobra d'Asie / Cobra indien / Boulengerina / Cobras d'eau douce / Cobra / Hydrophiidae / Hydrophiidés / **Serpents** marins / Naja /	Elapidae	Hydrophiinae / Lauticaudinae / Lauticauda / Mamba / Mambas / Dendroaspis / Micrurus / Coral Snake / Coral Snakes / Snake, Coral / Snakes, Coral / Hemachatus / Haemachatus / Ringhals / Ringhal / Naja haje / hajes, Naja / Egyptian Cobra / Cobra, Egyptian / Cobras, Egyptian / Egyptian Cobras / Naja naja / Asiatic Cobra / Asiatic Cobras / Cobra, Asiatic / Cobras, Asiatic / Boulengerina / Water Cobra / Cobra, Water / Cobras, Water / Water Cobras / Cobra / Cobras / Hydrophiidae / Sea Snakes / Sea Snake / Snake, Sea / Snakes, Sea / Naja /	B01.050.150.900.833.672.350	Arborescence

Construire sa recherche : PICO

Type de question clinique	Diagnostic	Étiologie	Thérapeutique /prévention	Pronostic
P	Population, Patient, Problème			
I (élément soumis à évaluation)	Nouveau test	Facteur d'exposition (facteur de risque)	Intervention thérapeutique	Facteur pronostic
C (Comparatuer)	Test de référence	Absence du facteur d'exposition evalué	Ttt ou intervention de référence, placébo, absence du ttt évalué	Absence du facteur pronostique, …
O (critère de jugement)	Identification de la maladie	Survenue de la maladie	Résultat clinique, événement	Résultat clinque

Construire sa recherche




NCBI
Resources
How To
PubMed.gov
US National Library of Medicine
National Institutes of Health
MeSH
Advanced
PubMed
PubMed comprises more than 27 million citations for biomedi
books. Citations may include links to full-text content from Pu

MeSH

MeSH | snake | Search

Create alert Limits Advanced

Summary 20 per page Send to:

Search results

Items: 1 to 20 of 21

<< First < Prev Page 1 of 2 Next > Last >>

1. **Snakes**
   Limbless REPTILES of the suborder Serpentes.

2. **Snake Venoms**
   Solutions or mixtures of toxic and nontoxic substances elaborated by **snake** (Ophidia) salivary glands for the purpose of killing prey or disabling predators and delivered by grooved or hollow fangs. They usually contain enzymes, toxins, and other factors.
   Year introduced: 1976(1975)

3. **Snake Bites**
   Bites by **snakes**. Bite by a venomous **snake** is characterized by stinging pain at the wound puncture. The venom injected at the site of the bite is capable of producing a deleterious effect on the blood or on the nervous system. (Webster's 3d ed; from Dorland, 27th ed, at **snake**, venomous)

4. **Coral Snakes**
   Elapid **snakes** indigenous to the Southern United States, Central, and South America. They are generally less than 1 meter in length and have a brightly-colored ringed pattern.
   Year introduced: 2018

5. **Colubridae**
   The largest family of **snakes** comprising five subfamilies: Colubrinae, Natricinae, Homalopsinae, Lycodontinae, and Xenodontinae. They show a great diversity of eating habits, from eating almost anything to having a specialized diet. They can be oviparous, ovoviviparous, or viviparous. The majority of North American **snakes** are colubrines. Among the colubrids are king **snakes**, water **snakes**, and garter **snakes**. Some genera are poisonous. (Goin, Goin, and Zug, Introduction to Herpetology, 3d ed, pp321-29)
   Year introduced: 1994

6. **Trichosanthes**
   A plant species of the family CUCURBITACEAE that is a source of TRICHOSANTHIN (a ribosomal inhibitory protein).
   Year introduced: 2002

PubMed Search Builder

Add to search builder AND

Search PubMed

YouTube T

Find related data

Database: Select

Find items

Search details

"snakes"[MeSH Terms] OR snake[T
Word]

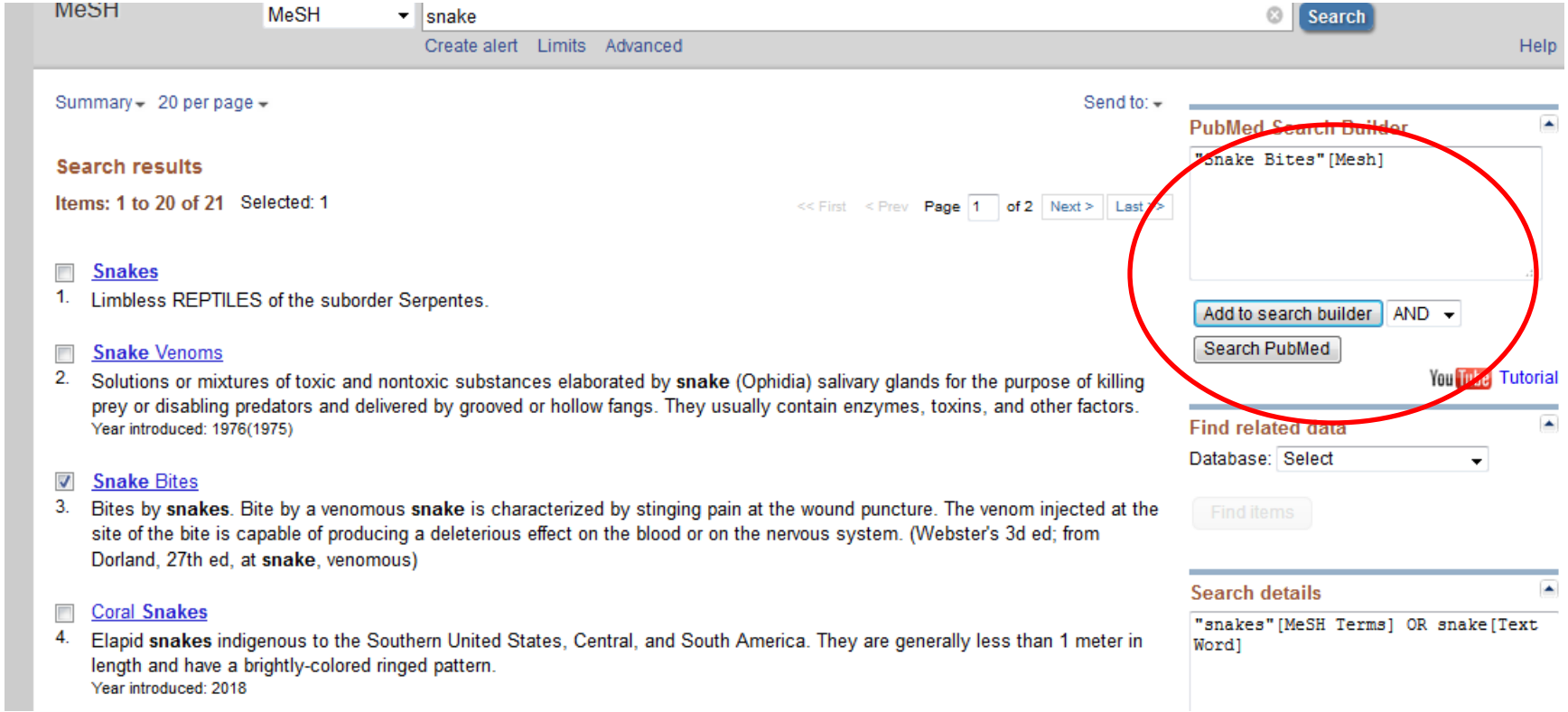
Search See

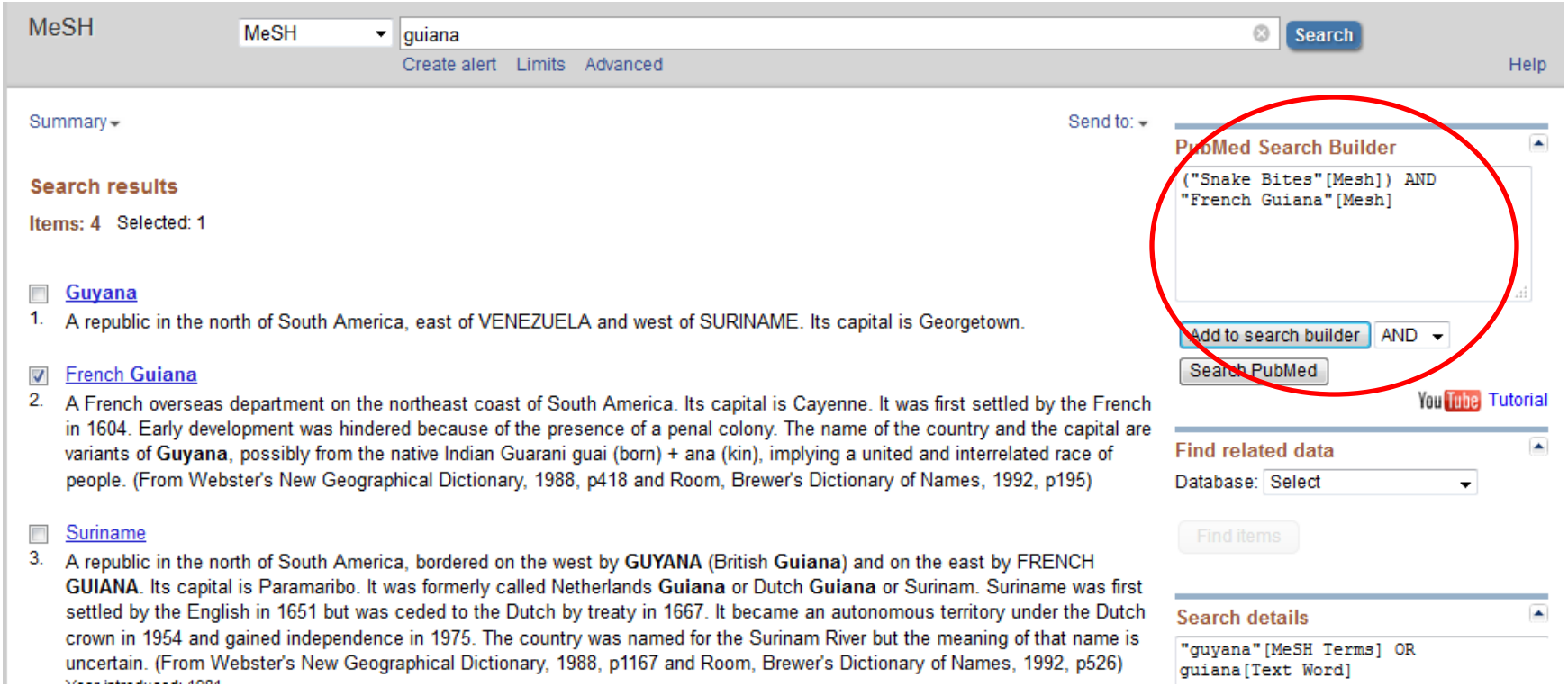
Recent Activity

Turn Off

snake (21)

Snake Bites


MeSH
MeSH
snake
Search
Create alert
Limits
Advanced
Help
Summary
20 per page
Send to:
Search results
Items: 1 to 20 of 21
Selected: 1
<< First
< Prev
Page 1 of 2
Next >
Last >>
Snakes
1. Limbless REPTILES of the suborder Serpentes.
Snake Venoms
2. Solutions or mixtures of toxic and nontoxic substances elaborated by snake (Ophidia) salivary glands for the purpose of killing prey or disabling predators and delivered by grooved or hollow fangs. They usually contain enzymes, toxins, and other factors.
Year introduced: 1976(1975)
Snake Bites
3. Bites by snakes. Bite by a venomous snake is characterized by stinging pain at the wound puncture. The venom injected at the site of the bite is capable of producing a deleterious effect on the blood or on the nervous system. (Webster's 3d ed; from Dorland, 27th ed, at snake, venomous)
Coral Snakes
4. Elapid snakes indigenous to the Southern United States, Central, and South America. They are generally less than 1 meter in length and have a brightly-colored ringed pattern.
Year introduced: 2018
PubMed Search Builder
"Snake Bites"[Mesh]
Add to search builder
AND
Search PubMed
YouTube Tutorial
Find related data
Database: Select
Find items
Search details
"snakes"[MeSH Terms] OR snake[Text Word]


MeSH
MeSH
guiana
Search
Create alert
Limits
Advanced
Help
Summary
Send to:
Search results
Items: 4 Selected: 1
Guyana
1. A republic in the north of South America, east of VENEZUELA and west of SURINAME. Its capital is Georgetown.
French Guiana
2. A French overseas department on the northeast coast of South America. Its capital is Cayenne. It was first settled by the French in 1604. Early development was hindered because of the presence of a penal colony. The name of the country and the capital are variants of Guyana, possibly from the native Indian Guarani guai (born) + ana (kin), implying a united and interrelated race of people. (From Webster's New Geographical Dictionary, 1988, p418 and Room, Brewer's Dictionary of Names, 1992, p195)
Suriname
3. A republic in the north of South America, bordered on the west by GUYANA (British Guiana) and on the east by FRENCH GUIANA. Its capital is Paramaribo. It was formerly called Netherlands Guiana or Dutch Guiana or Surinam. Suriname was first settled by the English in 1651 but was ceded to the Dutch by treaty in 1667. It became an autonomous territory under the Dutch crown in 1954 and gained independence in 1975. The country was named for the Surinam River but the meaning of that name is uncertain. (From Webster's New Geographical Dictionary, 1988, p1167 and Room, Brewer's Dictionary of Names, 1992, p526)
PubMed Search Builder
("Snake Bites"[Mesh]) AND "French Guiana"[Mesh]
Add to search builder
AND
Search PubMed
YouTube Tutorial
Find related data
Database: Select
Find items
Search details
"guyana"[MeSH Terms] OR guiana[Text Word]

PubMed.gov

("Snake Bites"[Mesh]) AND "French Guiana"[Mesh]

Search

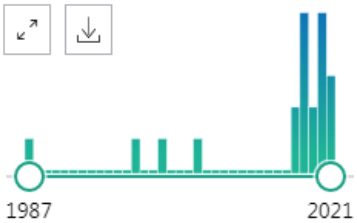
Advanced Create alert Create RSS User Guide

Save Email Send to Sorted by: Most recent Display options

MY NCBI FILTERS

All (15)
English (12)
Free Full Text (3)
Humans (14)
has abstract (10)
Review (2)
– show fewer

RESULTS BY YEAR



TEXT AVAILABILITY

Abstract

15 results

Page 1 of 1

1 Cite Share

Clinical Features and Management of Snakebite Envenoming in French Guiana.

Resiere D, Houcke S, Pujo JM, Mayence C, Mathien C, NkontCho F, Blaise N, Demar MP, Hommel D, Kallel H.

Toxins (Basel). 2020 Oct 19;12(10):662. doi: 10.3390/toxins12100662.

PMID: 33086750 Free PMC article.

2 Cite Share

Reply to Comment on Failure of a Mexican antivenom on recovery from snakebite-related coagulopathy in French Guiana.

Heckmann X, Lambert V, Mion G, Ehrhardt A, Marty C, Perotti F, Carod JF, Jolivet A, Boels D, Andi IL, Larréché S.

Clin Toxicol (Phila). 2021 Mar;59(3):267-268. doi: 10.1080/15563650.2020.1814317. Epub 2020 Sep 1.

PMID: 32870069 No abstract available.

3 Cite Share

Comment on Failure of a Mexican antivenom on recovery from snakebite-related coagulopathy in French Guiana.

Greene SC, Brandehoff NP.

Clin Toxicol (Phila). 2021 Mar;59(3):266. doi: 10.1080/15563650.2020.1800722. Epub 2020 Aug 25.

PMID: 32840395 No abstract available.

Nouveau cas

- Patient drépanocytaire vu aux urgences
- Le chef demande une présentation
- ….

Autre exemple

Recherche simple

Drépanocytose | Rechercher

Recherche avancée

Descripteur français	contient		et
Descripteur français	contient		et
Descripteur français	contient		et
Descripteur français	contient		

Réinitialiser | Rechercher

Descripteurs trouvés

2 document(s) trouvé(s) en 2 ms.

Descripteur français	Terme(s) français	Descripteur américain	Terme(s) américain(s)	Code(s) d'arborescence	Localisation
Drépanocytose	Anémie SS / Anémie drépanocytaire / Anémie falciforme / Anémie à cellules falciformes / Anémie à drépanocytes / Anémie à hématies falciformes / **Drépanocytose** homozygote / Hémoglobinopathie à hématies falciformes / Hémoglobinose S / Hémoglobinose SS / Sicklanémie / Sicklémie /	Anemia, Sickle Cell	Anemias, Sickle Cell / Sickle Cell Anemias / Hemoglobin S Disease / Disease, Hemoglobin S / Hemoglobin S Diseases / Sickle Cell Anemia / Sickle Cell Disorders / Cell Disorder, Sickle / Cell Disorders, Sickle / Sickle Cell Disorder / Sickling Disorder Due to Hemoglobin S / HbS Disease / Sickle Cell Disease / Cell Disease, Sickle / Cell Diseases, Sickle / Sickle Cell Diseases /	C15.378.071.141.150.150 C15.378.420.155 C16.320.070.150 C16.320.365.155	Arborescence
Drépanocytose SC	Hémoglobinopathie SC / Hémoglobinose SC /	Hemoglobin SC Disease	Disease, Hemoglobin SC / Diseases, Hemoglobin SC / Hemoglobin SC Diseases / SC Disease, Hemoglobin / SC Diseases, Hemoglobin / Sickle Cell Hemoglobin C Disease / SC Disease / Disease, SC / Diseases, SC / SC Diseases /	C15.378.071.141.150.150.440 C15.378.420.155.440 C16.320.070.150.440 C16.320.365.155.440	Arborescence

PubMed.gov
US National Library of Medicine
National Institutes of Health

PubMed | Anemia, Sickle Cell | Search

Create RSS | Create alert | Advanced

Article type
Clinical Trial
Review
Customize ...

Text availability
Abstract
Free full text
Full text

PubMed Commons
Reader comments
Trending articles

Publication dates
5 years
10 years
Custom range...

Species
Humans
Other Animals

Clear all

Show additional filters

Format: Summary | Sort by: Most Recent | Per page: 20 | Send to

Best matches for Anemia, Sickle Cell:

Emerging drugs for **sickle cell anemia**.
Singh PC et al. Expert Opin Emerg Drugs. (2015)

[**Sickle cell anemia** causes varied symptoms and high morbidity. Serious prognosis in the most common genetic disease in the world].
Kjellander C et al. Lakartidningen. (2015)

Study of alpha hemoglobin stabilizing protein expression in patients with β thalassemia and **sickle cell anemia** and its impact on clinical severity.
Mahmoud HM et al. Blood Cells Mol Dis. (2015)

Switch to our new best match sort order

Search results

Items: 1 to 20 of 21985

<< First | < Prev | Page 1 of 1100 | Next > | Last >>

1. Early Noninvasive Ventilation and Nonroutine Transfusion for Acute Chest Syndrome in **Sickle Cell** Disease in Children: A Descriptive Study.
Heilbronner C, Merckx A, Brousse V, Allali S, Hubert P, de Montalembert M, Lesage F.
Pediatr Crit Care Med. 2018 Jan 19. doi: 10.1097/PCC.0000000000001468. [Epub ahead of print]
PMID: 29356722

2. The first South American case of pre-implantation genetic diagnosis to select compatible embryo for cord blood transplantation as treatment for **sickle cell anemia**.
Martinhago CD, Endo KR, Oliveira MA, Dias AM, Pereira GS, Azzolini AM, Estrada PR, Bruzaca CG, Martinhago ACN.
JBRA Assist Reprod. 2018 Jan 18. doi: 10.5935/1518-0557.20180017. [Epub ahead of print]
PMID: 29345446 **Free Article**
Similar articles

Filter your results:

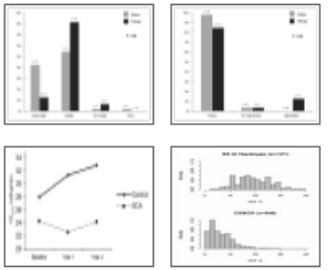
All (21985)
English (19960)
Free Full Text (5735)
Humans (20766)
Items with Abstracts (14176)
Review (2707)

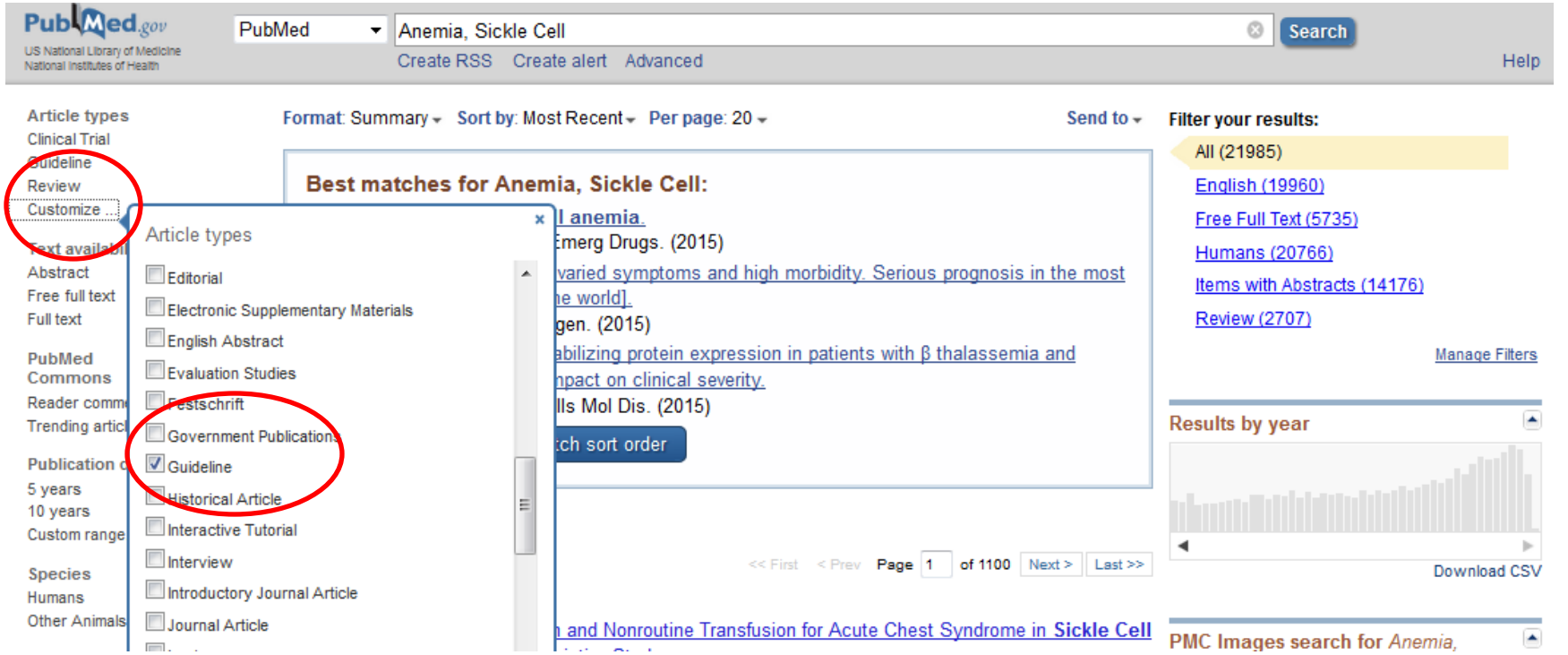
Manage Fil

Results by year

Download

PMC Images search for *Anemia, Sickle Cell*




PubMed.gov
US National Library of Medicine
National Institutes of Health
PubMed
Anemia, Sickle Cell
Search
Create RSS
Create alert
Advanced
Help
Article types
Clinical Trial
Guideline
Review
Customize ...
Format: Summary
Sort by: Most Recent
Per page: 20
Send to
Filter your results:
All (21985)
English (19960)
Free Full Text (5735)
Humans (20766)
Items with Abstracts (14176)
Review (2707)
Manage Filters
Best matches for Anemia, Sickle Cell:
Abstract
Free full text
Full text
PubMed Commons
Publication
5 years
10 years
Custom range
Species
Humans
Other Animals
Article types
Editorial
Electronic Supplementary Materials
English Abstract
Evaluation Studies
Festschrift
Government Publications
Guideline
Historical Article
Interactive Tutorial
Interview
Introductory Journal Article
Journal Article
Results by year
Download CSV
<< First
< Prev
Page 1
of 1100
Next >
Last >>
PMC Images search for Anemia,

PubMed.gov
US National Library of Medicine
National Institutes of Health

PubMed | Anemia, Sickle Cell | Search

Create RSS | Create alert | Advanced | Help

Article types
Clinical Trial
Guideline
Review
Customize ...

Text availability
Abstract
Free full text
Full text

PubMed Commons
Reader comments
Trending articles

Publication dates
5 years
10 years
Custom range...

Species
Humans
Other Animals

Clear all

Show additional filters

Additional filters
- [x] Article types
- [x] Text availability
- [x] PubMed Commons
- [x] Publication dates
- [x] Species
- [ ] Languages

Format: Summary | **Sort by**: Most Recent | **Per page**: 20 | **Send to**

Best matches for Anemia, Sickle Cell:

Emerging drugs for **sickle cell anemia**.
Singh PC et al. Expert Opin Emerg Drugs. (2015)

[**Sickle cell anemia** causes varied symptoms and high morbidity. Serious prognosis in the most common genetic disease in the world].
Kjellander C et al. Lakartidningen. (2015)

Study of alpha hemoglobin stabilizing protein expression in patients with β thalassemia and **sickle cell anemia** and its impact on clinical severity.
Mahmoud HM et al. Blood Cells Mol Dis. (2015)

Switch to our new best match sort order

Search results

Items: 1 to 20 of 21985

<< First | < Prev | Page 1 of 1100 | Next > | Last >>

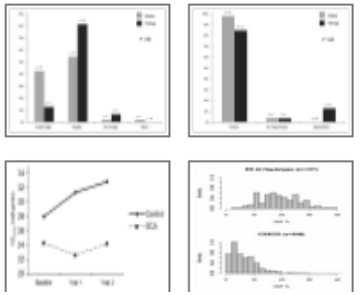
1. Early Noninvasive Ventilation and Nonroutine Transfusion for Acute Chest Syndrome in **Sickle Cell** Disease in Children: A Descriptive Study.
Heilbronner C, Merckx A, Brousse V, Allali S, Hubert P, de Montalembert M, Lesage F.
. 2018 Jan 19. doi: 10.1097/PCC.0000000000001468. [Epub ahead of print]

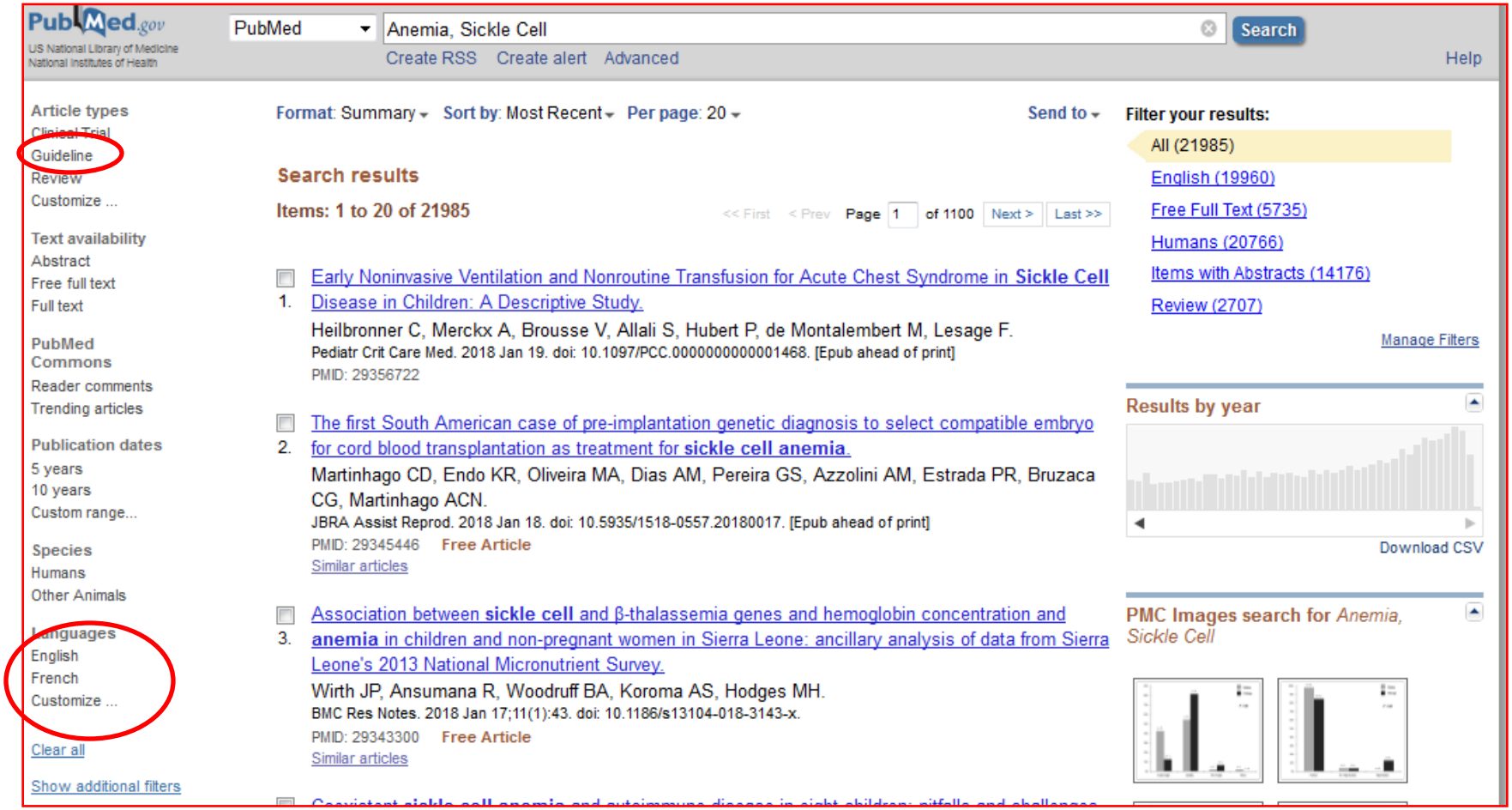
merican case of pre-implantation genetic diagnosis to select compatible embryo
nsplantation as treatment for **sickle cell anemia**.
Endo KR, Oliveira MA, Dias AM, Pereira GS, Azzolini AM, Estrada PR, Bruzaca ACN.
. 2018 Jan 18. doi: 10.5935/1518-0557.20180017. [Epub ahead of print]
ree Article

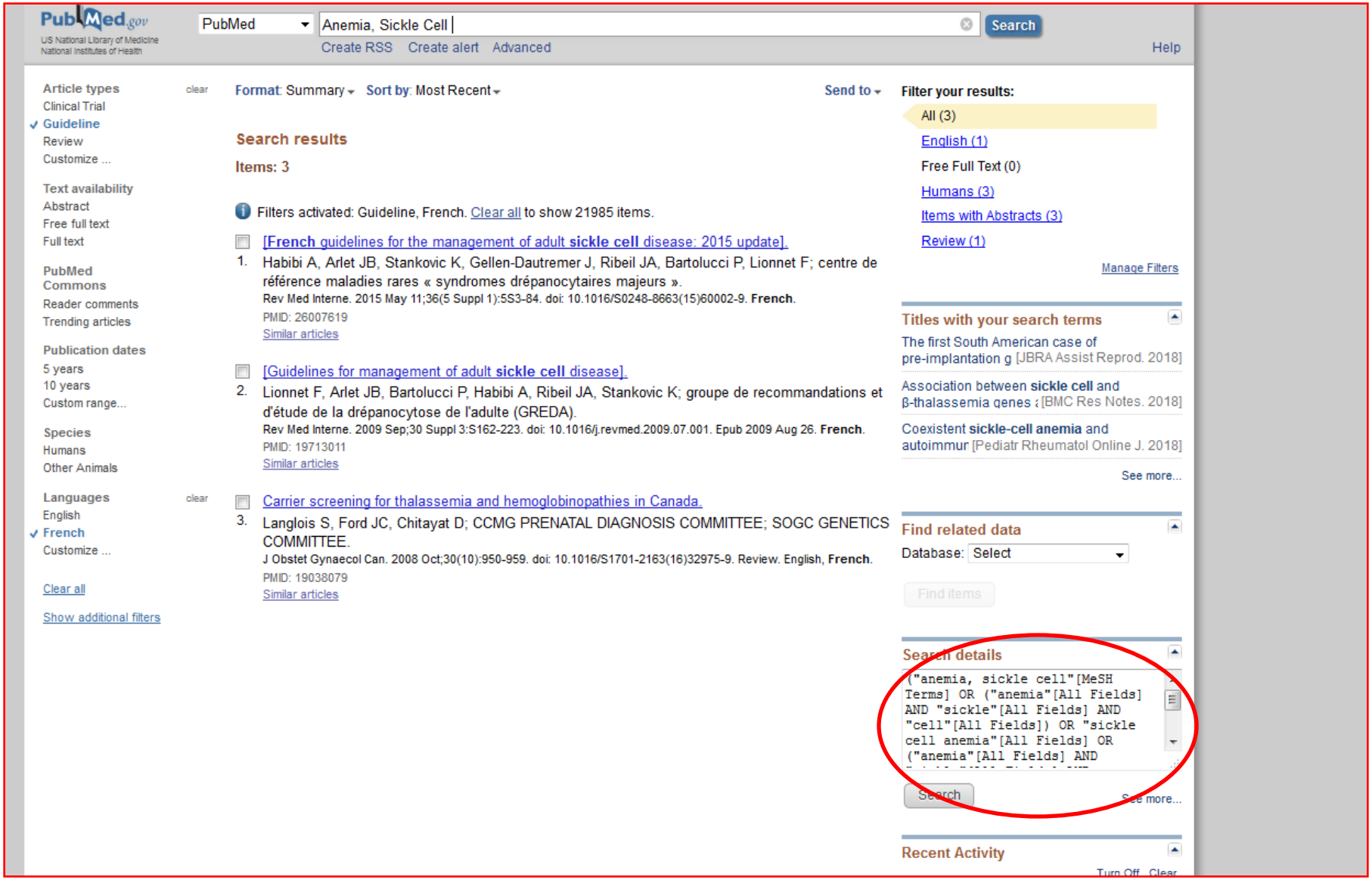
Filter your results:
All (21985)
English (19960)
Free Full Text (5735)
Humans (20766)
Items with Abstracts (14176)
Review (2707)
Manage Filters

Results by year
Download CSV

PMC Images search for *Anemia, Sickle Cell*




PubMed.gov
US National Library of Medicine
National Institutes of Health
PubMed
Anemia, Sickle Cell
Search
Create RSS
Create alert
Advanced
Help
Article types
Clinical Trial
Guideline
Review
Customize ...
Text availability
Abstract
Free full text
Full text
PubMed Commons
Reader comments
Trending articles
Publication dates
5 years
10 years
Custom range...
Species
Humans
Other Animals
Languages
English
French
Customize ...
Clear all
Show additional filters
Format: Summary
Sort by: Most Recent
Per page: 20
Send to
Search results
Items: 1 to 20 of 21985
<< First
< Prev
Page 1 of 1100
Next >
Last >>
1. Early Noninvasive Ventilation and Nonroutine Transfusion for Acute Chest Syndrome in Sickle Cell Disease in Children: A Descriptive Study.
Heilbronner C, Merckx A, Brousse V, Allali S, Hubert P, de Montalembert M, Lesage F.
Pediatr Crit Care Med. 2018 Jan 19. doi: 10.1097/PCC.0000000000001468. [Epub ahead of print]
PMID: 29356722
2. The first South American case of pre-implantation genetic diagnosis to select compatible embryo for cord blood transplantation as treatment for sickle cell anemia.
Martinhago CD, Endo KR, Oliveira MA, Dias AM, Pereira GS, Azzolini AM, Estrada PR, Bruzaca CG, Martinhago ACN.
JBRA Assist Reprod. 2018 Jan 18. doi: 10.5935/1518-0557.20180017. [Epub ahead of print]
PMID: 29345446 Free Article
Similar articles
3. Association between sickle cell and β-thalassemia genes and hemoglobin concentration and anemia in children and non-pregnant women in Sierra Leone: ancillary analysis of data from Sierra Leone's 2013 National Micronutrient Survey.
Wirth JP, Ansumana R, Woodruff BA, Koroma AS, Hodges MH.
BMC Res Notes. 2018 Jan 17;11(1):43. doi: 10.1186/s13104-018-3143-x.
PMID: 29343300 Free Article
Similar articles
Filter your results:
All (21985)
English (19960)
Free Full Text (5735)
Humans (20766)
Items with Abstracts (14176)
Review (2707)
Manage Filters
Results by year
Download CSV
PMC Images search for Anemia, Sickle Cell


PubMed.gov
US National Library of Medicine
National Institutes of Health
PubMed
Anemia, Sickle Cell
Search
Create RSS
Create alert
Advanced
Help
Article types
clear
Clinical Trial
Guideline
Review
Customize ...
Text availability
Abstract
Free full text
Full text
PubMed Commons
Reader comments
Trending articles
Publication dates
5 years
10 years
Custom range...
Species
Humans
Other Animals
Languages
clear
English
French
Customize ...
Clear all
Show additional filters
Format: Summary
Sort by: Most Recent
Send to
Search results
Items: 3
Filters activated: Guideline, French. Clear all to show 21985 items.
[French guidelines for the management of adult sickle cell disease: 2015 update].
1. Habibi A, Arlet JB, Stankovic K, Gellen-Dautremer J, Ribeil JA, Bartolucci P, Lionnet F; centre de référence maladies rares « syndromes drépanocytaires majeurs ».
Rev Med Interne. 2015 May 11;36(5 Suppl 1):5S3-84. doi: 10.1016/S0248-8663(15)60002-9. French.
PMID: 26007619
Similar articles
[Guidelines for management of adult sickle cell disease].
2. Lionnet F, Arlet JB, Bartolucci P, Habibi A, Ribeil JA, Stankovic K; groupe de recommandations et d'étude de la drépanocytose de l'adulte (GREDA).
Rev Med Interne. 2009 Sep;30 Suppl 3:S162-223. doi: 10.1016/j.revmed.2009.07.001. Epub 2009 Aug 26. French.
PMID: 19713011
Similar articles
Carrier screening for thalassemia and hemoglobinopathies in Canada.
3. Langlois S, Ford JC, Chitayat D; CCMG PRENATAL DIAGNOSIS COMMITTEE; SOGC GENETICS COMMITTEE.
J Obstet Gynaecol Can. 2008 Oct;30(10):950-959. doi: 10.1016/S1701-2163(16)32975-9. Review. English, French.
PMID: 19038079
Similar articles
Filter your results:
All (3)
English (1)
Free Full Text (0)
Humans (3)
Items with Abstracts (3)
Review (1)
Manage Filters
Titles with your search terms
The first South American case of pre-implantation g [JBRA Assist Reprod. 2018]
Association between sickle cell and β-thalassemia genes : [BMC Res Notes. 2018]
Coexistent sickle-cell anemia and autoimmur [Pediatr Rheumatol Online J. 2018]
See more...
Find related data
Database: Select
Find items
Search details
("anemia, sickle cell"[MeSH Terms] OR ("anemia"[All Fields] AND "sickle"[All Fields] AND "cell"[All Fields]) OR "sickle cell anemia"[All Fields] OR ("anemia"[All Fields] AND
Search
See more...
Recent Activity
Turn Off Clear

Format: Abstract Send to

Rev Med Interne. 2015 May 11;36(5 Suppl 1):5S3-84. doi: 10.1016/S0248-8663(15)60002-9.

[French guidelines for the management of adult sickle cell disease: 2015 update].

[Article in French]
Habibi A[1], Arlet JB[2], Stankovic K[3], Gellen-Dautremer J[4], Ribeil JA[5], Bartolucci P[6], Lionnet F[7]; centre de référence maladies rares « syndromes drépanocytaires majeurs ».

Author information

Abstract

Sickle cell disease is a systemic genetic disorder, causing many functional and tissular modifications. As the prevalence of patients with sickle cell disease increases gradually in France, every physician can be potentially involved in the care of these patients. Complications of sickle cell disease can be acute and chronic. Pain is the main symptom and should be treated quickly and aggressively. In order to reduce the fatality rate associated with acute chest syndrome, it must be detected and treated early. Chronic complications are one of the main concerns in adults and should be identified as early as possible in order to prevent end organ damage. Many organs can be involved, including bones, kidneys, eyes, lungs, etc. The indications for a specific treatment (blood transfusion or hydroxyurea) should be regularly discussed. Coordinated health care should be carefully organized to allow a regular follow-up near the living place and access to specialized departments. We present in this article the French guidelines for the sickle cell disease management in adulthood.

KEYWORDS: Acute chest syndrome; Blood transfusion; Crise vaso-occlusive; Drépanocytose; Hydroxyurea; Hydroxyurée; Sickle cell disease; Syndrome thoracique aigu; Transfusion sanguine; Vaso-occlusive crisis

PMID: 26007619 DOI: 10.1016/S0248-8663(15)60002-9

[Indexed for MEDLINE]

ELSEVIER

La Revue de Médecine Interne

Volume 36, Issue 5, Supplement 1, 11 May 2015, Pages 5S3-5S84

Recommandations

Recommandations françaises de prise en charge de la drépanocytose de l'adulte : actualisation 2015

French guidelines for the management of adult sickle cell disease: 2015 update

A. Habibi [a, b, c], J.-B. Arlet [d, e], K. Stankovic [f], J. Gellen-Dautremer [a], J.-A. Ribeil [e, g], P. Bartolucci [a, b], F. Lionnet [f, h], centre de référence maladies rares « syndromes drépanocytaires majeurs »

Show more

https://doi.org/10.1016/S0248-8663(15)60002-9

Resume

La drépanocytose est une pathologie génétique du globule rouge, qui provoque des atteintes fonctionnelles et tissulaires de façon systémique. La prévalence de la drépanocytose de l'adulte est en augmentation continue en France. Tout médecin peut être conduit à prendre en charge un patient drépanocytaire. Les complications s'expriment de façon aiguë ou chronique. La douleur représente le principal symptôme qui doit être traité

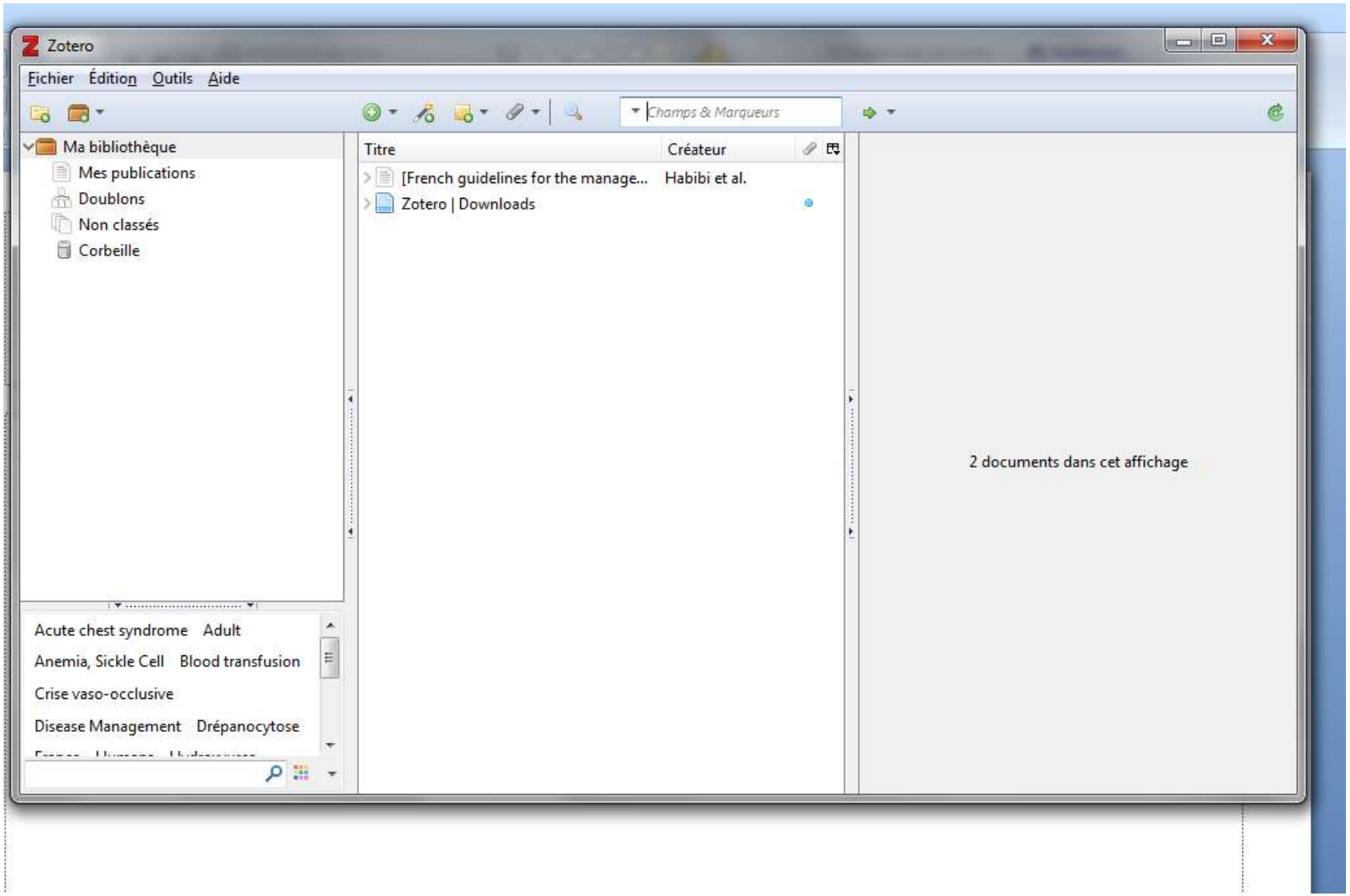
Logiciel de gestion de biblio

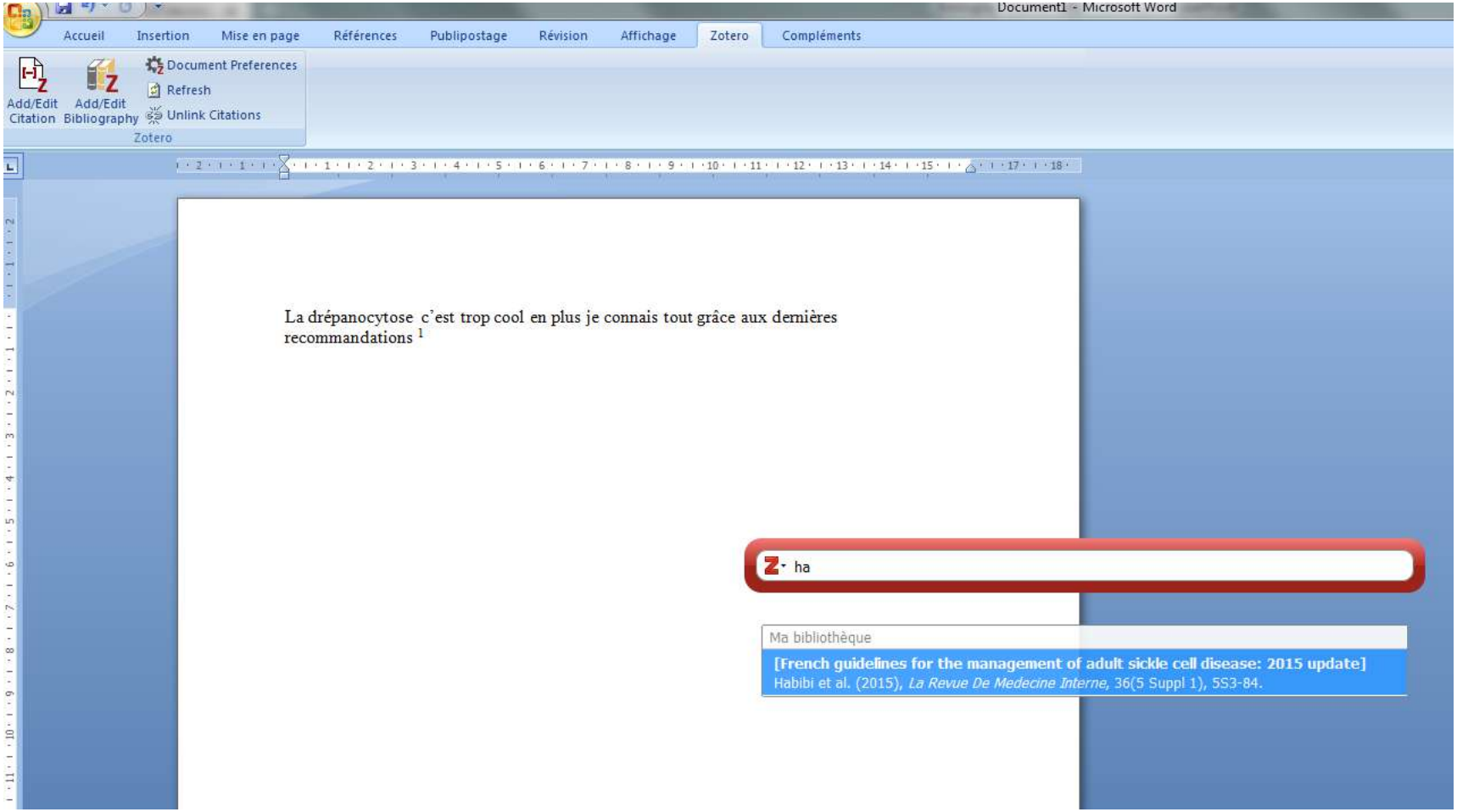
- Zotero
- Endnote
- Bibtex
- RefWorks

zotero

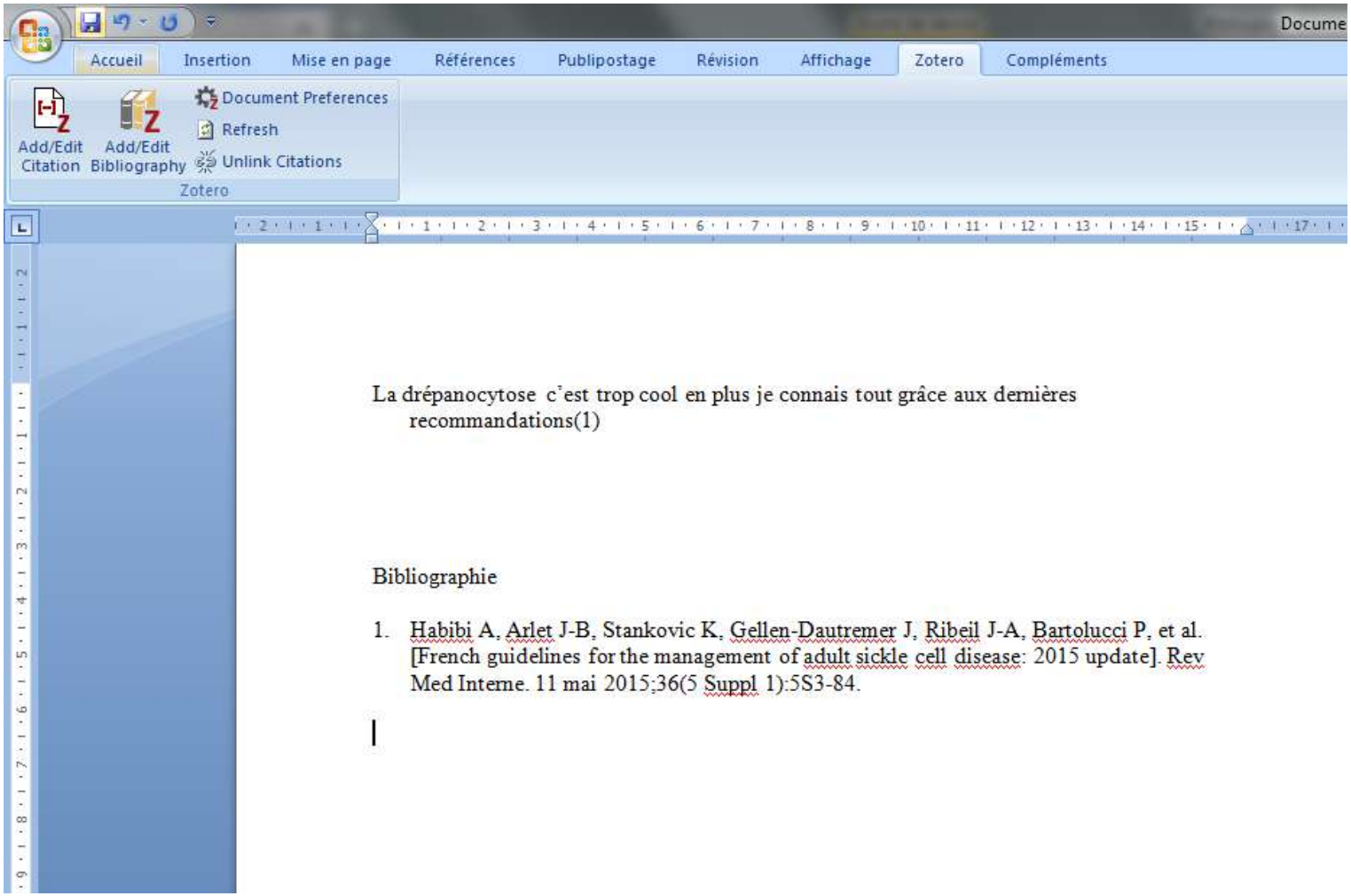

zotero
Log In · Register
Upgrade Storage
Home
Groups
People
Documentation
Forums
Get Involved
Search support
Search
Zotero [zoh-TAIR-oh] is a free, easy-to-use tool to help you collect, organize, cite, and share your research sources.
Download Now
Available for Mac, Windows, and Linux
Collect
Organize
Cite
Sync
Collaborate
Grab your research with a single click.
A personal research assistant.
Zotero is the only research tool that automatically senses content in your web browser, allowing you to add it to your personal library with a single click. Whether you're searching for a preprint on arXiv.org, a journal article from JSTOR, a news story from the New York Times, or a book from your university library catalog, Zotero has you covered with support for thousands of sites.
Save to Zotero
Title
bookreviews.pdf
in progress.doc
Outline for Hamlet
research data.xls
Store anything.
Zotero collects all your research in a single, searchable interface. You can add PDFs, images, audio and video files, snapshots of web pages, and really anything else. Zotero automatically indexes the full-text content of your library, enabling you to find exactly what you're looking for with just a few keystrokes.
Follow Zotero on Twitter
Zotero News
Zotero 5.0 and Firefox: Frequently
Get Involved
Zotero wouldn't be the powerful,


Zotero
Upgrade Storage
Home
Groups
People
Documentation
Forums
Get Involved
Search
Home > Downloads
Using Zotero with Firefox? We've made some important changes to the way Zotero works.
Zotero 5.0 for Windows
Your personal research assistant
Download
Other platforms
macOS · Linux 32-bit · Linux 64-bit
Zotero Connector
Save to Zotero from your browser
Install Firefox Connector
Zotero Connectors automatically sense content as you browse the web and allow you to save it to Zotero with a single click.


Zotero
Fichier Édition Outils Aide
Champs & Marqueurs
Ma bibliothèque
Mes publications
Doublons
Non classés
Corbeille
Titre
Créateur
[French guidelines for the manage...
Habibi et al.
Zotero | Downloads
2 documents dans cet affichage
Acute chest syndrome Adult
Anemia, Sickle Cell Blood transfusion
Crise vaso-occlusive
Disease Management Drépanocytose


Document1 - Microsoft Word
Accueil
Insertion
Mise en page
Références
Publipostage
Révision
Affichage
Zotero
Compléments
Add/Edit Citation
Add/Edit Bibliography
Document Preferences
Refresh
Unlink Citations
Zotero
La drépanocytose c'est trop cool en plus je connais tout grâce aux dernières recommandations [1]
ha
Ma bibliothèque
[French guidelines for the management of adult sickle cell disease: 2015 update]
Habibi et al. (2015), La Revue De Medecine Interne, 36(5 Suppl 1), 5S3-84.

Choisir style Vancouver



Google Scholar

- https://scholar.google.com/
- Cherche dans toutes la documentation scientifique
- Mais possible biais de recherche
- Et pas de sélection des revues

Autre outils

- https://www.uptodate.com/contents/search
- https://www.dynamed.com/
- https://www.tripdatabase.com/

Endometriosis

Overview and Recommendations

Background

- Endometriosis results from the presence of endometrial tissue (glands or stroma) implanted outside of the uterus which may result in pelvic pain or infertility.
- The most common location of endometriosis is the ovaries. Other typical locations include the peritoneum, uterosacral ligaments, and retrouterine pouch.
- Associated symptoms , including chronic pelvic pain and subfertility, may be due to the estrogen-stimulated inflammatory response of the endometriotic lesions.
- Women with history of endometriosis may have slight increased risk of ovarian cancer but this risk may be reduced with > 10 years of oral contraceptive use.

Evaluation

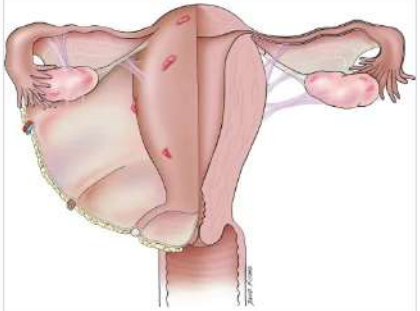
- Pelvic pain is the most common symptom associated with endometriosis and is usually chronic.
- Pain is most intense during the late luteal phase and at the beginning of menses.
- Suspect the diagnosis of endometriosis in women with signs and symptoms on history and physical exam, including (Strong recommendation):
  - severe dysmenorrhea unresponsive to nonsteroidal anti-inflammatory drugs
  - pelvic tenderness and nodularity on palpitation of uterosacral ligament and rectovaginal fascia
- Ultrasound is the first-line imaging test to evaluate for suspected endometriosis (Strong recommendation).
- Definitive diagnosis made by laparoscopic visualization of endometrial lesions with histologic confirmation (Strong recommendation)

TOPIC EDITOR
Elvia Greathouse MD

RECOMMENDATIONS EDITOR
Zbys Fedorowicz MSc, DPH, BDS, LDSRCS

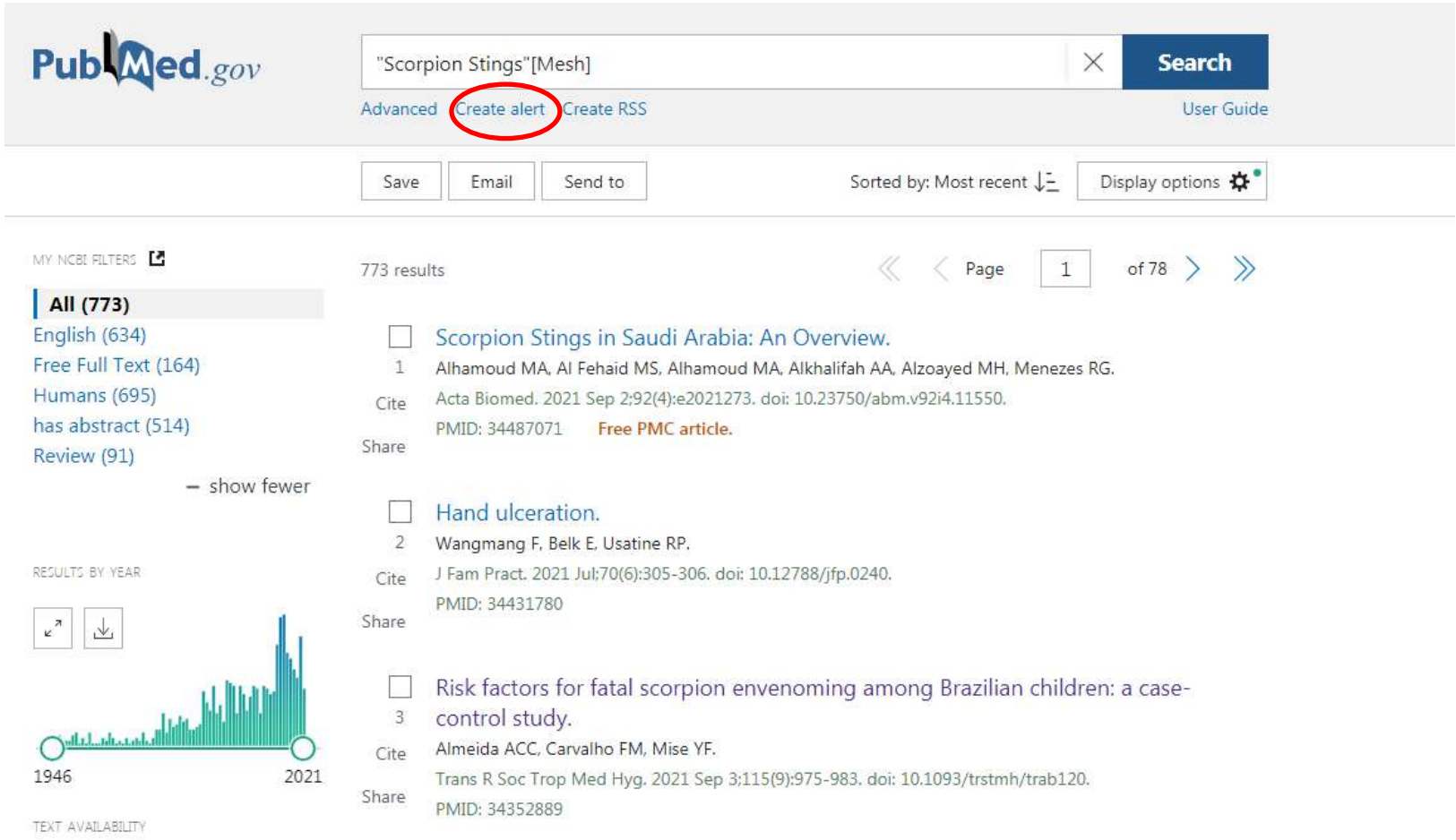
DEPUTY EDITOR
Alan Ehrlich MD, FAAFP

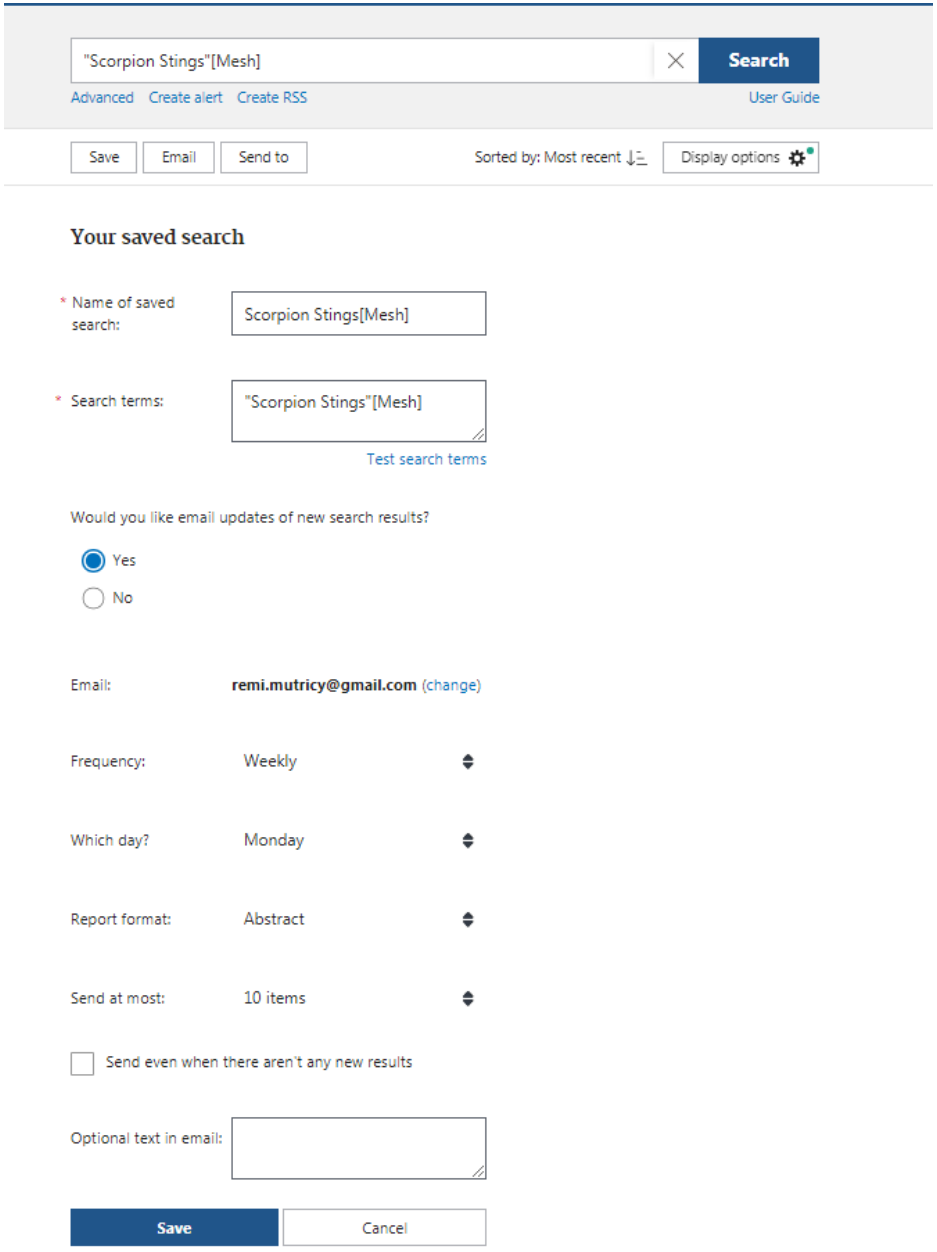
Images All (5)



Endometriosis

VEILLE : pubmed




"Scorpion Stings"[Mesh]
Search
Advanced
Create alert
Create RSS
User Guide
Save
Email
Send to
Sorted by: Most recent
Display options
Your saved search
* Name of saved search:
Scorpion Stings[Mesh]
* Search terms:
"Scorpion Stings"[Mesh]
Test search terms
Would you like email updates of new search results?
Yes
No
Email:
remi.mutricy@gmail.com (change)
Frequency:
Weekly
Which day?
Monday
Report format:
Abstract
Send at most:
10 items
Send even when there aren't any new results
Optional text in email:
Save
Cancel

What's new for 'snake bites' in PubMed

My NCBI à moi 22 janv.

This message contains My NCBI what's new results from the National Center for Biotechnology Information (NCBI) at the U.S. National Library of Medicine (NLM).
Do not reply directly to this message.

Sender's message:

Sent on Monday, 2018 January 22
Search: **snake bites**

View complete results in PubMed (results may change over time).

Edit saved search settings, or unsubscribe from these e-mail updates.

PubMed Results

Item 1 of 1

1. A Tragedy of Errors.
   Srikanth L.
   J Pain Palliat Care Pharmacother. 2016 Jun;30(2):131-3. doi: 10.3109/15360288.2016.1173757. Epub 2016 May 12.
   PMID: 27171792 [PubMed - indexed for MEDLINE]
   Similar articles

R Répondre

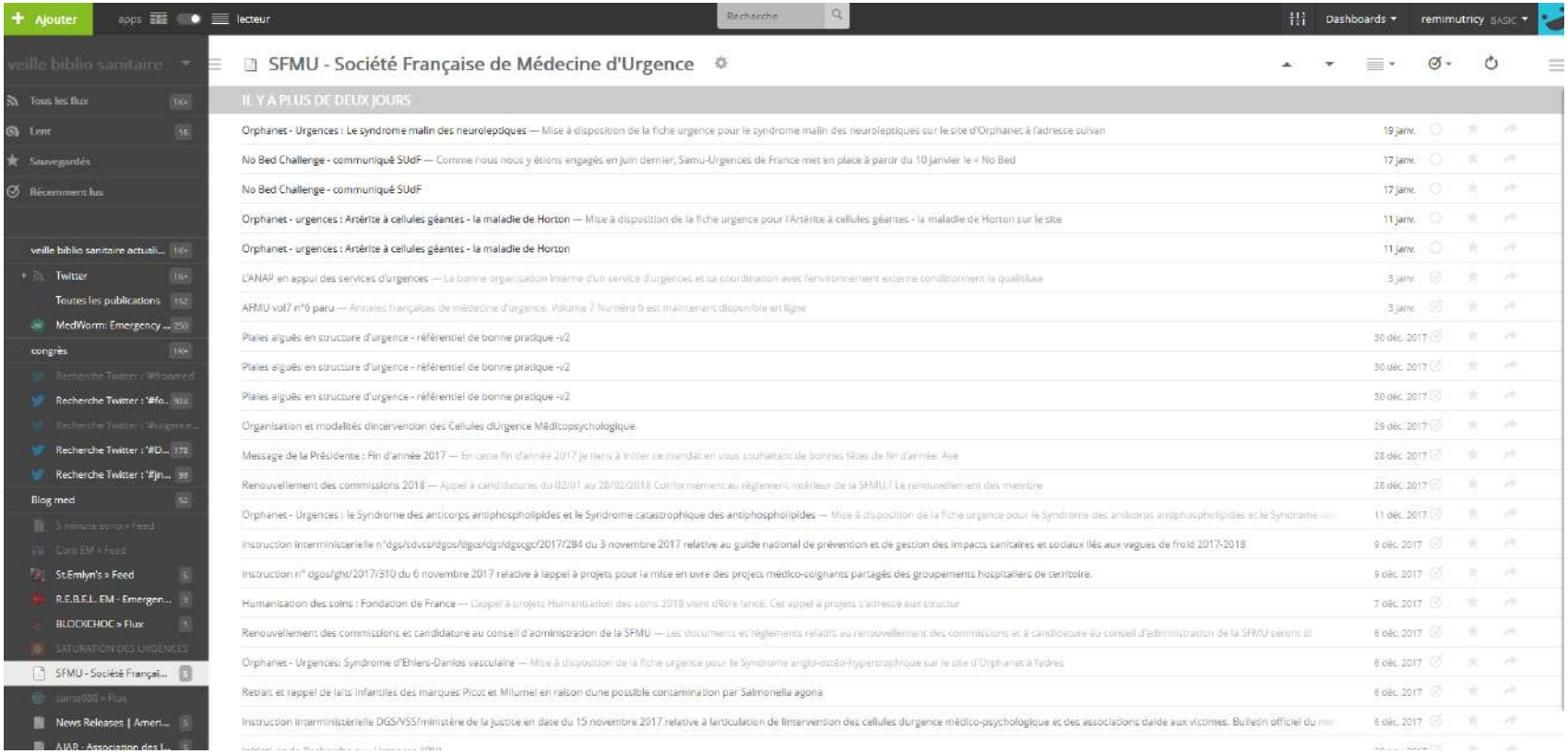
Veille flux RSS

- RSS : *Rich Site Summary* ou *Really Simple Syndication*

L'icône des flux RSS.

Veille flux RSS

- Agregateur de Flux : outlok, Thunderbird, netvibes.
- https://www.netvibes.com/fr



Trucs et BLOG à suivre

- Médecine d'urgences :
  - Santé publique France
  - http://blog.5minsono.com/
  - https://coreem.net/
  - http://stemlynsblog.org/
  - http://rebelem.com/
  - http://blockchoc.org/
  - http://www.sfmu.org/fr/
  - http://www.ajar-online.fr/
  - https://echo-urgences.com/
  - http://la-mine.net/

- Médecine générale
  - https://medicalement-geek.blogspot.com/
  - Jaddo.fr
  - …

Google alerte

- https://www.google.fr/alerts