

1
Name of Journal: *World Journal of Gastroenterology*

Manuscript NO: 40167

Manuscript Type: ORIGINAL ARTICLE

Basic Study

Mechanism of combined use of vitamin D and puerarin in anti-hepatic fibrosis by regulating the Wnt/ β -Catenin signalling pathway

Huang GR *et al.* Anti-hepatic fibrosis mechanism

Gan-Rong Huang, Si-Jun Wei, Yan-Qiang Huang, Wei Xing, Lu-Yao Wang,
 Ling-Ling Liang

Abstract

AIM

To reveal the protective mechanism of the combined use of vitamin D and puerarin in the progression of hepatic fibrosis induced by carbon tetrachloride (CCl₄).

Match Overview

1	Internet 16 words crawled on 05-Oct-2017 real.mtak.hu	<1%
2	Crossref 14 words Ya Zhang, Hang Zhao, Hua Li, Wei Cao, Fang Wang, Tian Z hang, Si-Wang Wang. "Protective Effects of Amarogentin i...	<1%
3	Internet 13 words crawled on 25-Nov-2017 repositorio.uc.cl	<1%
4	Crossref 12 words Arai, Yohei, Eiichiro Kanda, Soichiro Iimori, Shotaro Naito, Yumi Noda, Tomoki Kawasaki, Hidehiko Sato, Ryoichi And	<1%
5	Internet 12 words crawled on 12-May-2018 www.wjgnet.com	<1%
6	Crossref 12 words Yang Yang, Xiao-xia Chen, Wan-xia Li, Xiao-qin Wu, Chen g Huang, Juan Xie, Yu-xin Zhao, Xiao-ming Meng, Jun Li.	<1%
7	Crossref 12 words "Abstract", Hepatology International, 06/2008	<1%

找到约 9,350 条结果 (用时 0.62 秒)

Wnt/ β -Catenin Signaling Promotes Renal Interstitial Fibrosis

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2663839/> ▼ 翻译此页

作者: W He - 2009 - 被引用次数: 392 - 相关文章

In addition to this canonical pathway, Wnt proteins may exert their activities through ... Wnt/ β -catenin signaling has been shown to play a role in kidney To examine the functional consequence of Wnt regulation in renal fibrosis, we next sought (C and D) ELISA analysis shows an increased DKK1 protein in liver (C) and ...

缺少字词: puerarin

The Wnt/ β -catenin signaling pathway in liver biology and disease

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3298845/> ▼ 翻译此页

作者: J Behari - 2010 - 被引用次数: 93 - 相关文章

A few years later, Nusse and Varmus, while working on the mechanisms of ... The canonical Wnt/ β -catenin pathway is centered on regulating the levels of its major Regulation of hepatic vitamin C biosynthesis by the Wnt/ β -catenin pathway signaling could be a potential therapeutic strategy in preventing liver fibrosis.

缺少字词: puerarin

MicroRNA-17-5p-activated Wnt/ β -catenin pathway contributes to the ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4807984/> ▼ 翻译此页

作者: F Yu - 2016 - 被引用次数: 30 - 相关文章

2015年12月2日 - However, the underlying molecular mechanisms responsible for the ... Aberrant Wnt/ β -catenin signaling has been shown to be involved in ... with anti-hepatic fibrosis activity, is often used as an anti-liver fibrotic product in China [17, 18]. ... of Wnt/ β -catenin pathway and the up-regulation of WIF1 expression.

缺少字词: vitamin puerarin

International Journal of Molecular Medicine - Spandidos Publications

<https://www.spandidos-publications.com/ijmm/articlesbydate.jsp> ▼

09/02/2018, Madecassoside activates anti-neuroinflammatory mechanisms by inhibiting

09/01/2018, Synergistic antitumor effect of brusatol combined with cisplatin on 26/08/2016, Low vitamin D-modulated calcium-regulating proteins in 22/01/2016, Role of the Wnt/ β -catenin signaling pathway in the response of ...

BCL2 | Proteintech Group 12789-1-AP product information - Labome

<https://www.labome.com/product/Proteintech-Group/12789-1-AP.html> ▼ 翻译此页

[全部](#)[图片](#)[新闻](#)[购物](#)[地图](#)[更多](#)[设置](#)[工具](#)

找到约 257 条结果 (用时 0.79 秒)

Mutations in SRD5B1 (AKR1D1), the gene encoding Δ 4-3-oxosteroid ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1773813/> - 翻译此页

作者: HA Lemonde - 2003 - 被引用次数: 73 - 相关文章

Conclusions: **Analysis** of blood samples for SRD5B1 **mutations** can be used to ... **Patients** with **genetic** 5 β -reductase deficiency may respond well to **treatment** with ... Keywords: bile acid synthesis, inborn error, giant cell hepatitis, **cholestasis**, bile ... This **infant** had failed to improve on **ursodeoxycholic acid** therapy but liver ...

Emmanuel Jacquemin's scientific contributions while affiliated with ...

https://www.researchgate.net/scientific-contributions/61814204_Emmanuel_Jacquemin

Mutations in the **novel gene** FOPV are associated with familial autosomal dominant ... Cholic **Acid** to **Treat** HSD3B7 and **AKR1D1** Deficiencies **sufficient** serum vitamin E level in the majority of children with **cholestasis**, single-center, **case-cohort study** including 16 **patients** with PSIS diagnosed before one year of age.

Kenneth DR Setchell's research works | Cincinnati Children's Hospital ...

https://www.researchgate.net/scientific-contributions/8826184_Kenneth_DR_Setchell

Methods: This was a single-center, **case-control** pilot **study** conducted at ... Among SUBA, **chenodeoxycholic acid** (CDCA) was significantly elevated in A **novel** homozygous frameshift germline **mutation** (c.587delG) in the **AKR1D1 gene**; which ... management of the youngest **patient** led to **successful treatment** of the liver ...

Chapter 165. Disorders of Bile Acid Synthesis | Rudolph's Pediatrics ...

<https://mhmedical.com/content.aspx?aid=6727328> - 翻译此页

Book cover The **first patients** were **treated** with **chenodeoxycholic acid** 13,8at a dose of ... proven **mutations** in SRD5B1 (**AKR1D1**, the **gene** encoding the 5 β -reductase ... Some **cases** showing a urine bile acid profile suggestive of 5 β -reductase **Analysis** of urinary bile acids in a **cholestatic infant** revealed major peaks ...

[PDF] Investigations into the genetic causes of liver ... - eTheses Repository

etheses.bham.ac.uk/7101/1/McKayBounford16PhD.pdf

[全部](#)[图片](#)[新闻](#)[购物](#)[地图](#)[更多](#)[设置](#)[工具](#)

找到约 257 条结果 (用时 0.79 秒)

Mutations in SRD5B1 (AKR1D1), the gene encoding Δ 4-3-oxosteroid ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1773813/> - 翻译此页

作者: HA Lemonde - 2003 - 被引用次数: 73 - 相关文章

Conclusions: **Analysis** of blood samples for SRD5B1 **mutations** can be used to ... **Patients** with **genetic** 5 β -reductase deficiency may respond well to **treatment** with ... Keywords: bile acid synthesis, inborn error, giant cell hepatitis, **cholestasis**, bile ... This **infant** had failed to improve on **ursodeoxycholic acid** therapy but liver ...

Emmanuel Jacquemin's scientific contributions while affiliated with ...

https://www.researchgate.net/scientific-contributions/61814204_Emmanuel_Jacquemin

Mutations in the **novel gene** FOPV are associated with familial autosomal dominant ... Cholic **Acid** to **Treat** HSD3B7 and **AKR1D1** Deficiencies **sufficient** serum vitamin E level in the majority of children with **cholestasis**, single-center, **case-cohort study** including 16 **patients** with PSIS diagnosed before one year of age.

Kenneth DR Setchell's research works | Cincinnati Children's Hospital ...

https://www.researchgate.net/scientific-contributions/8826184_Kenneth_DR_Setchell

Methods: This was a single-center, **case-control** pilot **study** conducted at ... Among SUBA, **chenodeoxycholic acid** (CDCA) was significantly elevated in A **novel** homozygous frameshift germline **mutation** (c.587delG) in the **AKR1D1 gene**; which ... management of the youngest **patient** led to **successful treatment** of the liver ...

Chapter 165. Disorders of Bile Acid Synthesis | Rudolph's Pediatrics ...

<https://mhmedical.com/content.aspx?aid=6727328> - 翻译此页

Book cover The **first patients** were **treated** with **chenodeoxycholic acid** 13,8at a dose of ... proven **mutations** in SRD5B1 (**AKR1D1**, the **gene** encoding the 5 β -reductase ... Some **cases** showing a urine bile acid profile suggestive of 5 β -reductase **Analysis** of urinary bile acids in a **cholestatic infant** revealed major peaks ...

[PDF] Investigations into the genetic causes of liver ... - eTheses Repository

etheses.bham.ac.uk/7101/1/McKayBounford16PhD.pdf