

Core LIVER: Extend Your Liver's Lifeline

written by Nick Andrews | November 22, 2021

Supplements designed to protect against *liver toxicity* are incredibly important for anyone who cares about their metabolic health. Regardless of the type of athlete or dieter you are, you should know that you can still benefit from taking something to support your liver. That's because *strenuous exercise* has been shown to significantly elevate liver enzymes in healthy adults.[1]



As a part of the incredibly well-received *Core Lifeline Series*, which includes a gut health masterpiece, *Core GUT*, and an inflammation mediator in *Core SOOTHE*, CORE Nutritionals added a potentially *more* important product to the line: a comprehensive liver support and detoxification supplement, **Core LIVER**.

Core LIVER: Comprehensive Liver Support & Detoxification

With just four ingredients and one absorption amplifier, Core LIVER is driven by a *massive* amount of milk thistle extract – at a higher dosage than we've ever seen. In addition, it includes an ingredient that may be new to most of us, in the form of *Katuki* extract, which has been shown to improve liver enzyme levels and reduce liver fat.

All of the details are covered below, but first take a moment to check PricePLOW's coupon-powered prices and sign up for our CORE Nutritionals alerts:

Core Nutritionals Liver – Deals and Price Drop Alerts

Get Price Alerts

Get LIVER Price Alerts
Get Core Nutritionals alerts
Get Liver Cleanse price drops

Also get hot deal alerts

No spam, no scams.

Disclosure: PricePLOW relies on pricing from stores with which we have a business relationship. We work hard to keep pricing current, but you may find a better offer.

Posts are sponsored in part by the retailers and/or brands listed on this page.

Core LIVER Ingredients

In a single *three-capsule* serving of CORE Nutritionals Liver, you get the following:

- **Milk thistle extract (80% silymarin) – 1260 mg**

Supplement Facts		
Serving Size: 3 Capsules Servings Per Container: 30		
	Amount Per Serving	% DV
Milk Thistle (<i>Silybum marianum</i>) (seeds) (80% Silymarin)	1260 mg	**
Katuki Extract (<i>Picrorrhiza kurrad</i>) (root) (std. to 5% kutkin)	400 mg	**
Setria® L-Glutathione	250 mg	**
TUDCA (Tauroursodeoxycholic Acid)	250 mg	**
Bioperine® (Black Pepper Extract) (<i>Piper nigrum</i>) (std. to 95% Piperine)	10 mg	**

**Daily Value not established.

OTHER INGREDIENTS: Cellulose (Veggie Capsule), Rice Flour, Magnesium Stearate, Silicon Dioxide.

Core LIVER Ingredients. Prepare to meet our newcomer, *Katuki* extract!

Any liver-protecting supplement worth its salt is going to contain some **milk thistle**. Also known as *Silybum marianum*, milk thistle is native to the Mediterranean Basin and Asia Minor, and has been recognized for its healing properties since at least 2,000 years ago when it was mentioned in the writings of ancient Greek physician and botanist, Dioscorides.[2]

The “per serving” dose of milk thistle used in CORE Nutritional Liver is *very* high for a comprehensive liver aid that contains additional ingredients. If you haven’t taken milk thistle before, we recommend starting with one capsule (which includes 1260 milligrams of the herb) and *working your way up* to the recommended three doses.

The first thing to notice about this particular milk thistle extract is that it has *80% standardization* for *silymarin* (the bioactive ingredient), which is a very high level of purity for milk thistle extracts. This, combined with the dosage, makes for a *powerful* supplement.

What is *silymarin*?

But what is *silymarin*? It's not actually a single compound – it's a *class* of *three* different “flavonolignans” called *silydianin*, *silibinin*, and *silychristin*.^[3] Although there *are* other bioactive compounds in milk thistle, these are the strongest.

Traditionally, milk thistle was used as a liver tonic^[3] and in the treatment of “melancholy diseases,”^[2] which was a catch-all category for a number of different ailments in pre-modern medicine. In the modern age, we enjoy a *huge* volume of published research attesting to milk thistle's preventative and therapeutic benefits.



Silybum Marianum, better known as *Milk Thistle*, has a tremendous amount of research for liver detoxification

That research includes, but is not limited to:

- Protection and rehabilitation from drug-induced liver disease^[4-6]
- Improvement of liver function post alcohol abuse^[7-12]
- Amelioration of general liver toxicity^[13-15]
- Protection of the liver from environmental toxins^[16]
- Protection against general liver disease^[17,18]

Although milk thistle extract does have an *impressive* ability to protect the liver from damage, the *best* way to prevent liver injury is to *remove or avoid*

whatever toxin you're trying to protect against. Just because you're taking a liver-support supplement does *not* give you license to abuse your liver with reckless abandon. The supplement will only *help* keep your liver healthy – you still need to support it with good lifestyle and dietary choices.

- **Katuki Extract (*Picrorrhiza kurrao*) standardized to 5% kutkin – 400 mg**

Bile is a substance produced by the liver. When it's released, it removes toxins, cholesterol, fat, drugs, and other metabolites from the liver.[19] So when it comes to optimal liver function, *bile secretion* is key] If liver function is impaired and you develop a *bile deficiency*, it can cause a wide range of problems, from impaired hormone synthesis[20] to gallstones[21].



Picrorrhiza Kurroa – the botanical source of Katuki extract

That's where *Picrorrhiza kurrao* comes in. First described in the Ayurvedic system of classical Indian medicine,[22] *Picrorrhiza* has been shown to protect the liver in both humans *and* animals. When given to rats whose livers were deliberately injured by researchers, the animals taking *Picrorrhiza* extracts showed *reductions* in liver enzymes and liver *fat* deposition – both clear indications of *improved* liver function.[22] In humans, a similar effect was observed. Patients who had been diagnosed with acute viral hepatitis saw their *bilirubin* levels return to baseline much *faster* when given a preparation of

Picrorrhiza. [22]

In another study, researchers examined *Picrorrhiza's* ability to reverse the pathology of non-alcoholic fatty liver disease (NAFLD) in rats and found that it *significantly* regressed several measures of NAFLD, as indicated by a histological study of the animals' livers. [23] This is a promising finding since NAFLD is a common disease in humans.

As the most novel ingredient in CORE Liver, Katuki extract introduces a new mechanism for liver health, and could be what sets this supplement apart from its competition. Aside from the rest of the formula, which is already powerful, Katuki makes it worth trying for those who keep an eye on their liver enzymes and are looking for something new.

- **Setria L-Glutathione – 250 mg**

Often referred to as the “master antioxidant,” **glutathione (GSH)** is a non-essential amino acid whose job it is to remove damaging free radical compounds from the body. [24,25] Because the body's rate of glutathione production is limited by the availability of *cysteine*, various supplementation strategies have been developed to overcome this bottleneck and provide the body with *optimal* glutathione levels.



In five incredible capsules, Core GUT brings prebiotics, probiotics, *postbiotics*, and digestive enzymes to help improve your gut health and digestion!

Straight glutathione has limited bioavailability, at least when administered orally. [26] So that's where designer preparations, such as **Setria** glutathione, come in handy. Setria has been shown to be *significantly* more bioavailable

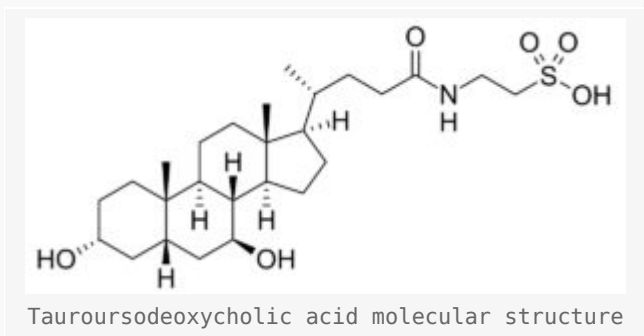
than ordinary glutathione, even in oral preparations. In fact, Setria is so much *more* bioavailable that it has *clinically significant effects*.

In the words of Penn State University College of Medicine researchers who tested an oral preparation of Setria, "*These findings show, for the **first time**, that daily consumption of GSH supplements was effective at increasing body compartment stores of GSH*".[27] In another study, supplementation with Setria and *citrulline* was shown to increase nitric oxide (NO) levels – an indirect way to measure the effectiveness of glutathione supplementation since glutathione protects NO from being reduced by *nitric oxide synthase*. [28]

- **TUDCA (Tauroursodeoxycholic Acid) – 250 mg**

Tauroursodeoxycholic acid is an exciting ingredient to see in a liver supplement. Traditionally, it hasn't been used very often just because it's costly and difficult to obtain. But its *efficacy* in protecting the liver is truly amazing.

TUDCA is *ursodeoxycholic acid* (UDCA) conjugated with the amino acid *taurine*. UDCA is used to treat *cholestasis*, a *bile-deficiency condition* that leads to an accumulation of toxins in the liver. Cholestasis can be caused by a physical blockage in bile ducts, but chronic drug use can also lead to liver toxicity.[29]



The consequences of cholestasis are grim. Usually, fat digestion slows to a halt and can eventually destroy the liver– then cell death often follows.[30] If left untreated, this condition can lead to total organ failure.

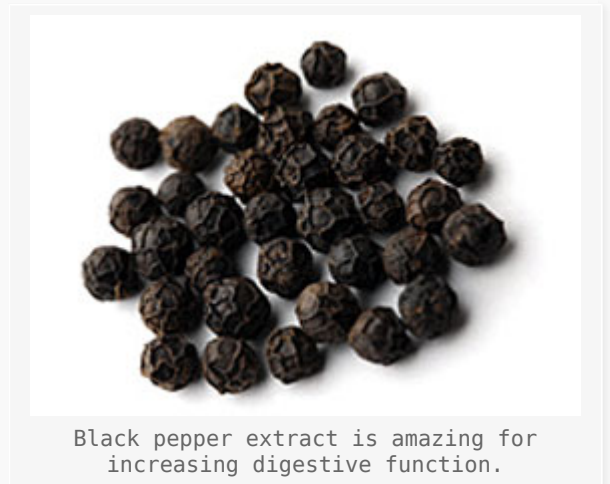
Where TUDCA comes in and helps protect the liver is by occupying certain *receptors* on the surface of the cell that are necessary for cell death to occur. Additionally, it helps stimulate the production of bile and the elimination of toxic metabolites from the liver, thus decreasing the overall burden on the liver.

In a large study directly comparing TUDCA to UDCA, (which is an established treatment for cholestasis), 250 milligrams of TUDCA proved to be, in the words

of the researchers, “safe and as efficacious as UDCA for the treatment of PBC [primary biliary cholangitis].”[31] In another study—a six month trial— TUDCA was found to be safer and more effective than UDCA in the treatment of liver cirrhosis.[32]

So, overall, TUDCA looks to be a lot better than the old workhorse UDCA.

- **BioPerine (Black Pepper (fruit) Extract std. to 95% Piperine) – 5mg**



The reason why **BioPerine** tends to show up in high-quality supplements is because by inhibiting the action of stomach enzymes, it protects bioactive compounds from breaking down in the stomach and, thus, substantially increases their bioavailability.

Capsule supplements are really one of the best places to use BioPerine since the spicy taste doesn't agree with everyone. When it's in a capsule, you can avoid dealing with the flavor or heat, if you so choose.

Dosage

Per the label shown below, take three capsules daily on an empty stomach. This can be all at once, or you can split them a bit into AM/PM if you must.

Dosing milk thistle extract at 1260 mg is appropriate, and in line with the industry standard dose of 1000 mg per day. Research shows that milk thistle extract is tolerated exceptionally well by humans, even at doses of 2 grams per day or more[33] – so there's no need to underdose this supplement. Still, it is, objectively speaking, a large dose, so if you've never taken milk thistle extract before, it would be wise to start with one capsule per day and work your way up to three.

Core LIVER: Another great Core Lifeline Series supplement



Introducing the Core Lifeline series! With Doug Miller at the helm, this is an unsurprising move – but wait until you see the dosages in these supplements!

The health of your liver drives countless other metabolic and organ systems related to your general health and well-being. Those who are consistent with blood work and nutritional monitoring can see how dietary and activity changes affect liver enzyme levels.

Core LIVER was specifically developed to keep the liver working when you're training hardest. Fans of the large, well-standardized milk thistle dose (as we are), are always impressed by the growing amount of research on TUDCA and welcome a new ingredient in the form of Katuki extract.

Although Core LIVER may not be a supplement everyone needs to take, it's one that should significantly help those who do want it to focus on this critical organ. Core GUT and Core SOOTHE may be the supplements in the Lifeline Series that get the most attention due to their general inclusivity, but LIVER may be the most important of the line.

Core Nutritionals Liver – Deals and Price Drop Alerts

Get Price Alerts

Get LIVER Price Alerts
Get Core Nutritionals alerts
Get Liver Cleanse price drops

Also get hot deal alerts

No spam, no scams.

Disclosure: PricePLOW relies on pricing from stores with which we have a

business relationship. We work hard to keep pricing current, but you may find a better offer.

Posts are sponsored in part by the retailers and/or brands listed on this page.

WARNING: This product is only intended for use in healthy adults 18 years of age or older. Pregnant or nursing women should not use this product. Consult your healthcare provider before using this product, especially if you are taking any prescription, over the counter medication, dietary supplement product, or if you have any pre-existing medical condition including, but not limited to: high or low blood pressure, cardiac arrhythmia, stroke, heart, liver, kidney or thyroid disease, seizure disorder, psychiatric disease, diabetes, difficulty urinating due to prostate enlargement, or if you are taking a MAOI (Moclobemide Oxidine Inhibitor) or any other medication. Discontinue use and consult your health care professional if you experience adverse reaction to this product. Discontinue use 2 weeks prior to surgery. Do not exceed recommended serving. Do not use if safety seal is broken or missing. Keep out of reach of children.

DIRECTIONS: Take 3 capsules daily on an empty stomach.

These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

WARNING: Consuming this product can expose you to chemicals including lead, which is known to the State of California to cause cancer. For more information, go to www.P65Warnings.ca.gov.

Supplement Facts
Serving Size: 3 Capsules
Servings Per Container: 30

	Amount Per Serving	% DV
Milk Thistle (Silybum marianum seeds) (90% Silymarin)	1200 mg	**
Kudzu Extract (Pueraria lobata root) (90% to 95% Kudzu)	400 mg	**
Silybin-1-Glucuronate	250 mg	**
T.D.C.A. (Transurodeoxycholic Acid)	250 mg	**
Bioperine® (Black Pepper Extract) (Piper nigrum seed, to 98% Piperin)	10 mg	**

OTHER INGREDIENTS: Cellulose (Veggie Capsule), Rice Flour, Magnesium Stearate, Silicon Dioxide.

BioPerine® is a registered trademark of Selmar Corporation.
Setria® is a registered trademark of KNOX, HANNO BIO CO., LTD.

Manufactured in a U.S. cGMP Facility.
Distributed by Core Nutritionals, LLC, Statesville, NC 28685.
(888) 978-2332, info@corenutritionals.com, www.corenutritionals.com

90 VEGETABLE CAPS
DIETARY SUPPLEMENT

The full Core LIVER Label

References

1. Prabin Khatri, Aryan Neupane, Suman Raj Sapkota, Bibhav Bashyal, Dipesh Sharma, Ashmita Chhetri, K. C. Chirag, Ashish Banjade, Priyanka Sapkota, Subarna. Bhandari, "Strenuous Exercise-Induced Tremendously Elevated Transaminases Levels in a Healthy Adult: A Diagnostic Dilemma", *Case Reports in Hepatology*, vol. 2021, Article ID 6653266, 3 pages, 2021. <https://doi.org/10.1155/2021/6653266>
2. Siegel, Abby B, and Justin Stebbing. "Milk thistle: early seeds of potential." *The Lancet. Oncology* vol. 14,10 (2013): 929-30. doi:10.1016/S1470-2045(13)70414-5 <https://www.ncbi.nlm.nih.gov/labs/pmc/articles/PMC4116427/>
3. Abenavoli, Ludovico, et al; "Milk Thistle in Liver Diseases: Past, Present, Future.;" *Phytotherapy Research : PTR*; U.S. National Library of Medicine; Oct. 2010; <https://www.ncbi.nlm.nih.gov/pubmed/20564545>
4. Saba, P. et al; "Effetti terapeutica della silimarina nelle epatopatie croniche indotte da psicofarmaci"; *Gaz Med Ital*; 135:236251; 1976
5. World Health Organization; "WHO Monographs on Selected Medicinal Plants – Volume 2"; 2004; <https://apps.who.int/medicinedocs/en/d/Js4927e/29.html>
6. Palasciano G et al; "The effect of silymarin on plasma levels of malondialdehyde in patients receiving long-term treatment with psychotropic drugs"; *Current Therapeutic Research*; 55:537-545; 1994; <https://moh-it.pure.elsevier.com/en/publications/the-effect-of-silymarin-on-plasma-levels-of-malon-dialdehyde-in-p>
7. Di Mario FR, Melzer J, Meier R; "Die Wirkung von Silymarin auf Leberfunktionsproben bei Patienten mit alkoholbedingter Lebererkrankung"; *Doppelblindstudie*. In: Di Ritis F, Csomos G, Braatz R, editors. *Der toxisch-metabolische Leberschaden* Lübeck: Hansisches Verlagskontor; p. 54–8; 1981
8. Hellerbrand, C., Schattenberg, J.M., Peterburs, P. et al; "The potential of silymarin for the treatment of hepatic disorders"; *Clinical Phytoscience*; 2, 7; 2017; <https://link.springer.com/article/10.1186/s40816-016-0019-2>
9. Fehér, J et al; "Liver-protective action of silymarin therapy in chronic alcoholic liver diseases"; *Orvosi Hetilap*; 130(51):2723-7; December 17, 1989; <https://www.ncbi.nlm.nih.gov/pubmed/2574842>
10. Ferenci, P et al; "Randomized controlled trial of silymarin treatment in patients with cirrhosis of the liver"; *Journal of Hepatology*; 9(1):105-13; July 1989; <https://www.ncbi.nlm.nih.gov/pubmed/2671116>
11. Velussi, M, et al; "Long-term (12 months) treatment with an anti-oxidant drug (silymarin)

- is effective on hyperinsulinemia, exogenous insulin need and malondialdehyde levels in cirrhotic diabetic patients"; *Journal of Hepatology*; 26(4):871-9; April 1997; <https://www.ncbi.nlm.nih.gov/pubmed/9126802>
12. Parés, A, et al; "Effects of silymarin in alcoholic patients with cirrhosis of the liver: results of a controlled, double-blind, randomized and multicenter trial"; *Journal of Hepatology*; 28(4):615-21; April 1998; <https://www.ncbi.nlm.nih.gov/pubmed/9566830>
 13. Shawn M. Talbott, Kerry Hughes; "The Health Professional's Guide to Dietary Supplements"; Lippincott Williams & Wilkins; 2007; https://books.google.com/books?id=hV2_TdmoDo8C
 14. Schuppan D, Strösser W, Burkard G, et al; "Legalon lessens fibrosing activity in patients with chronic liver diseases" [in German]. *Z Allgemeinmed*; 74:577-84; 1998
 15. Albrecht M, Frerick H, Kuhn U, et al; "Therapy of toxic liver pathologies with Legalon" [in German]; *Z Klin Med*; 47:87-92; 1992
 16. Szilárd S, et al; "Protective effect of Legalon in workers exposed to organic solvents"; *Acta Medica Hungarica*; 45(2):249-56; 1998; <https://www.ncbi.nlm.nih.gov/pubmed/3073356>
 17. Polachi, Navaneethakrishnan, et al; "Modulatory Effects of Silibinin in Various Cell Signaling Pathways against Liver Disorders and Cancer – A Comprehensive Review."; *European Journal of Medicinal Chemistry*; U.S. National Library of Medicine; 10 Nov. 2016; <https://www.ncbi.nlm.nih.gov/pubmed/27517806>
 18. de Avelar, Camila Ribeiro et al; "Effect of silymarin on biochemical indicators in patients with liver disease: Systematic review with meta-analysis."; *World journal of gastroenterology*; vol. 23,27; 2017; 5004-5017; <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5526770/>
 19. "Bile Secretion – an Overview | ScienceDirect Topics." www.sciencedirect.com, www.sciencedirect.com/topics/medicine-and-dentistry/bile-secretion.
 20. Zhou, Huiping, and Phillip B Hylemon. "Bile acids are nutrient signaling hormones." *Steroids* vol. 86 (2014): 62-8. doi:10.1016/j.steroids.2014.04.016 <https://www.ncbi.nlm.nih.gov/labs/pmc/articles/PMC4073476/>
 21. Carey MC. Pathogenesis of gallstones. *Am J Surg*. 1993 Apr;165(4):410-9. doi: 10.1016/s0002-9610(05)80932-8. PMID: 8480873. <https://pubmed.ncbi.nlm.nih.gov/8480873/>
 22. Vaidya, AB, et al. "Picrorhiza Kurroa (Kutki) Royle Ex Benth as a Hepatoprotective Agent—Experimental & Clinical Studies." *Journal of Postgraduate Medicine*, 1996, www.jpgmonline.com/article.asp?issn=0022-3859;year=1996;volume=42;issue=4;spage=105;epage=8;aulast=Vaidya. Accessed 18 Nov. 2021.
 23. Shetty, Sapna N et al. "A study of standardized extracts of Picrorhiza kurroa Royle ex Benth in experimental nonalcoholic fatty liver disease." *Journal of Ayurveda and integrative medicine* vol. 1,3 (2010): 203-10. doi:10.4103/0975-9476.72622 <https://www.ncbi.nlm.nih.gov/labs/pmc/articles/PMC3087357/>
 24. Winterbourn, C. C., and D. Metodiewa. "The Reaction of Superoxide with Reduced Glutathione." *Archives of Biochemistry and Biophysics*, vol. 314, no. 2, 1 Nov. 1994, pp. 284-290, 10.1006/abbi.1994.1444; <https://pubmed.ncbi.nlm.nih.gov/7979367/>
 25. Cardey, Bruno, et al. "Mechanism of Thiol Oxidation by the Superoxide Radical." *The Journal of Physical Chemistry A*, vol. 111, no. 50, Dec. 2007, pp. 13046-13052, 10.1021/jp0731102; <https://pubmed.ncbi.nlm.nih.gov/18044848/>
 26. Witschi, A., et al. "The Systemic Availability of Oral Glutathione." *European Journal of Clinical Pharmacology*, vol. 43, no. 6, 1992, pp. 667-669, 10.1007/BF02284971; <https://pubmed.ncbi.nlm.nih.gov/1362956/>
 27. Richie JP Jr, Nichenametla S, Neidig W, Calcagnotto A, Haley JS, Schell TD, Muscat JE. Randomized controlled trial of oral glutathione supplementation on body stores of glutathione. *Eur J Nutr*. 2015 Mar;54(2):251-63. doi: 10.1007/s00394-014-0706-z. Epub 2014 May 5. PMID: 24791752. <https://pubmed.ncbi.nlm.nih.gov/24791752/>
 28. McKinley-Barnard, Sarah et al. "Combined L-citrulline and glutathione supplementation increases the concentration of markers indicative of nitric oxide synthesis." *Journal of the International Society of Sports Nutrition* vol. 12 27. 10 Jun. 2015, doi:10.1186/s12970-015-0086-7 <https://www.ncbi.nlm.nih.gov/labs/pmc/articles/PMC4472409/>
 29. Ishak, Kamal, and Hyman Zimmerman. "Hepatotoxic Effects of the Anabolic/Androgenic Steroids." *Seminars in Liver Disease*, vol. 7, no. 03, Aug. 1987, pp. 230-236, 10.1055/s-2008-1040579; <https://pubmed.ncbi.nlm.nih.gov/3317860/>
 30. Amaral, Joana D., et al. "Bile Acids: Regulation of Apoptosis by Ursodeoxycholic Acid." *Journal of Lipid Research*, vol. 50, no. 9, Sept. 2009, pp. 1721-1734, 10.1194/jlr.r900011-jlr200; <https://www.ncbi.nlm.nih.gov/labs/pmc/articles/PMC2724780/>
 31. Ma, Hong et al. "A multicenter, randomized, double-blind trial comparing the efficacy and safety of TUDCA and UDCA in Chinese patients with primary biliary cholangitis." *Medicine*

vol. 95,47 (2016): e5391. doi:10.1097/MD.0000000000005391;
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5134868/>

32. Pan XL, Zhao L, Li L, et al. "Efficacy and safety of tauroursodeoxycholic acid in the treatment of liver cirrhosis: a double-blind randomized controlled trial"; *J Huazhong Univ Sci Technolog Med Sci*2013;33:189–94; <https://www.ncbi.nlm.nih.gov/pubmed/23592128>
33. Soleimani V, Delghandi PS, Moallem SA, Karimi G. Safety and toxicity of silymarin, the major constituent of milk thistle extract: An updated review. *Phytother Res*. 2019 Jun;33(6):1627-1638. doi: 10.1002/ptr.6361. Epub 2019 May 8. PMID: 31069872. <https://pubmed.ncbi.nlm.nih.gov/31069872/>