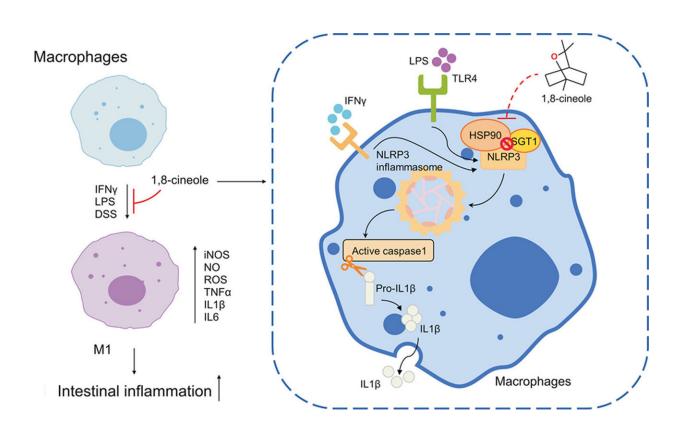


Research demonstrates the potential of natural compounds to treat gastrointestinal disorders

November 27 2023



Credit: *Journal of Pharmaceutical Analysis* (2023). DOI: 10.1016/j.jpha.2023.07.001

Chronic gastrointestinal (GI) disorders are becoming increasingly common throughout the world, but many of them still lack effective



treatment. Researchers have now turned to natural compounds, such as those present in traditional medicines, to search for potential drug candidates for difficult-to-treat diseases. The latest issue of the *Journal of Pharmaceutical Analysis (JPA)* features three articles that report promising findings and could pave the way to treatments for complex diseases affecting the liver and intestine.

The first study focuses on 1,8-cineole, a compound found in volatile oil extracts of the Amomum compactum plant, for treating <u>ulcerative colitis</u> (UC). Motivated by its established use for the treatment of pneumonia and preliminary data on its beneficial effects on the gut, a group of researchers investigated the anti-inflammatory mechanisms of this compound in the context of UC. To this end, they employed dextran sulfate sodium to induce colitis in C57BL/6 mice and conducted several assays on <u>cell cultures</u> to determine 1,8-cineole's targets and its effects.

The article was <u>published</u> in *JPA* in September 2023. The findings revealed that this molecule inhibits the binding between <u>heat shock</u> <u>protein</u> 90 (HSP90) and nucleotide-binding and oligomerization domain-, leucine rich repeat-, and pyrin domain-containing 3 (NLRP3), thereby inhibiting the activation of the NLPR3 inflammasome and the M1 polarization of macrophages. This prevents the activation of inflammatory pathways and promotes the recovery of the intestinal wall.

"Our findings provide convincing evidence that 1,8-cineole is a promising candidate in the search for a novel therapeutic strategy for UC treatment," comments corresponding author Dr. Xiaoyin Chen.

In the <u>second study</u>, scientists turned to ginsenoside Rk2, a bioactive component found in steamed ginseng (Panax notoginseng), and investigated its potential as a treatment for alcoholic lived disease (ALD). They used an ethanol feeding regime to induce ALD-like liver injury in mice and then showed that Rk2 could alleviate the observed



deleterious effects.

Simply put, as an NLPR3 inhibitor, Rk2 could ameliorate liver inflammation. Moreover, Rk2 also restored intestinal barrier function by enhancing NLPR6 signaling in the intestine, demonstrating the multifaceted nature of its therapeutic potential.

"Our findings indicate that Rk2 is a promising agent for the prevention and treatment of ALD and other NLPR3-driven diseases," concludes corresponding author Dr. Jian-Bo Wan.

Finally, in the <u>third study</u>, researchers explored the relationship between gut microbiota and specnuezhenide (SNZ), an active component of the traditional Chinese medicine Fructus Ligustri Lucidi. Their findings revealed that SNZ is metabolized by the enzyme carboxylesterase produced by intestinal bacteria, yielding the metabolites salidroside and tyrosol.

In mice, orally administered SNZ could modify the makeup of the gut microbiota and fungal groups, and also had the potential to inhibit colorectal tumor growth. Taken together, these results suggest the importance of the metabolism of gut microbiota.

"Intestinal bacterial metabolic enzymes are an important supplement to human liver and kidney metabolism. Thus, in clinical drug use, the characteristics of the <u>gut microbiota</u> in individuals must be taken into account for personalized treatment," remarks corresponding author Dr. Yan Wang.

These studies will hopefully pave the way to naturally effective treatments for challenging GI disorders.

More information: Shengsuo Ma et al, 1,8-cineole ameliorates colon



injury by downregulating macrophage M1 polarization via inhibiting the HSP90-NLRP3-SGT1 complex, *Journal of Pharmaceutical Analysis* (2023). DOI: 10.1016/j.jpha.2023.07.001

Jian Zou et al, Ginsenoside Rk2, a dehydroprotopanaxadiol saponin, alleviates alcoholic liver disease via regulating NLRP3 and NLRP6 inflammasome signaling pathways in mice, *Journal of Pharmaceutical Analysis* (2023). DOI: 10.1016/j.jpha.2023.05.005

Hang Yu et al, Gut microbiota-based pharmacokineticpharmacodynamic study and molecular mechanism of specnuezhenide in the treatment of colorectal cancer targeting carboxylesterase, *Journal of Pharmaceutical Analysis* (2023). DOI: 10.1016/j.jpha.2023.06.012

Provided by Cactus Communications

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