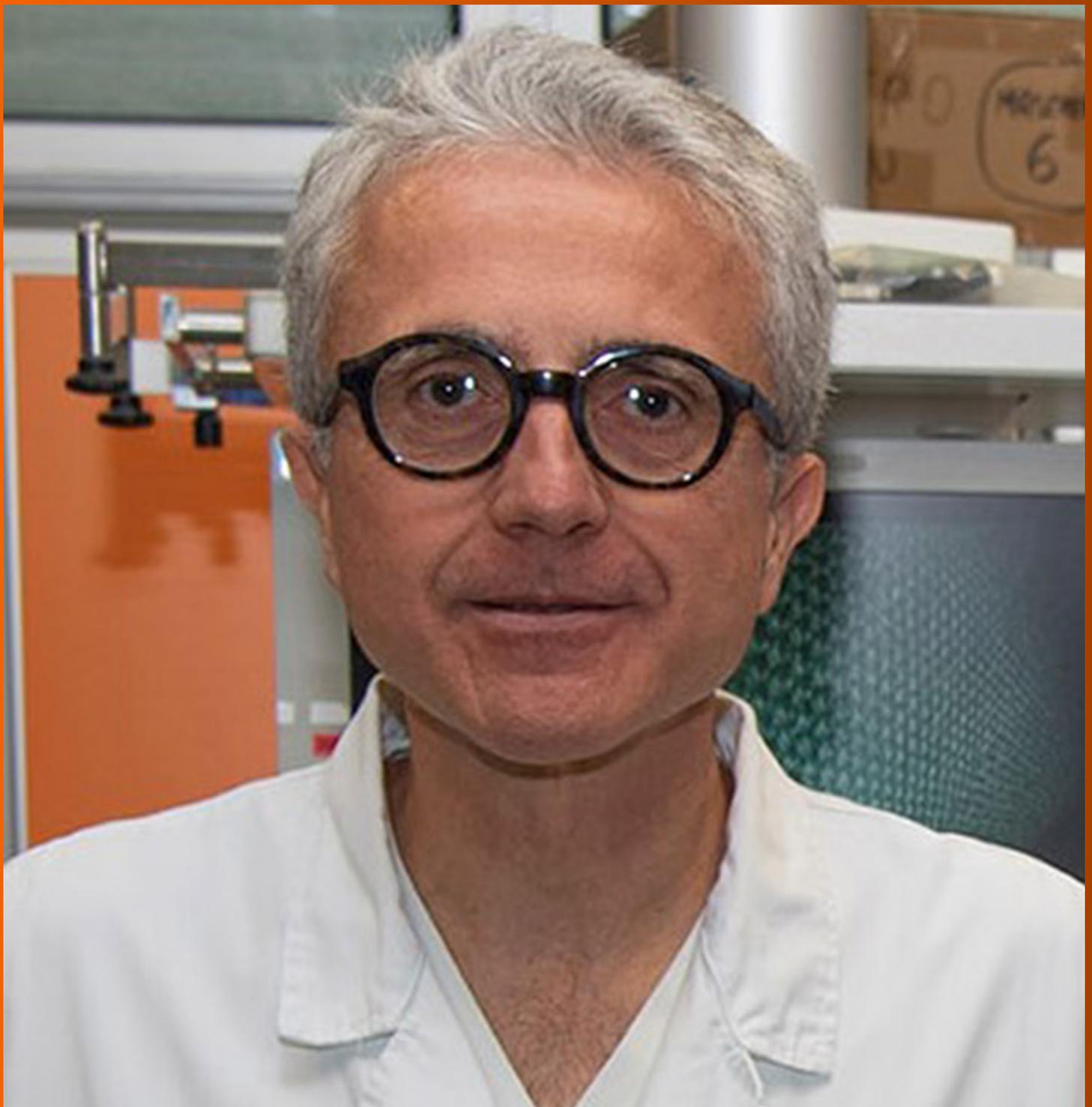


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Intestinal barrier in inflammatory bowel disease: A bibliometric and knowledge-map analysis

Feng Zhou, Nan-Zhen Wu, Yong Xie, Xiao-Jiang Zhou

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Abstract

BACKGROUND

Barrier surfaces composed of specialized epithelial cells separate the host body from the external environment, and are essential for maintaining proper intestinal physiologic and immune homeostasis.

AIM

To explore the development trends and research hotspots of intestinal barrier research in inflammatory bowel disease (IBD).

METHODS

The publications related to the intestinal barrier in IBD were obtained from the Web of Science Core Collection database. Bibliometric analysis and visualization were conducted using VOSviewer, CiteSpace and R software.

RESULTS

A total of 4482 articles published between 2002 and 2022 were identified. The United States is dominant in intestinal barrier research, whereas the University of Chicago is the most active institution. Jerrold from Harvard Medical School was the most productive authors with the most citations. The journals Inflammatory Bowel Disease and Gastroenterology have made significant contributions in this field. The keywords appearing at high frequency related to the intestinal barrier in IBD were detected, including nuclear factor kappa B, tumor necrosis factor- α , apoptosis, oxidative stress and probiotics. Among them, antioxidants, *Akkermansia muciniphila*, nanoparticles, short-chain fatty acids and extracellular vesicles have received growing interest in recent research.

CONCLUSION

The intestinal barrier field is developing rapidly with extensive cooperation. Targeting the gut microbiota and dietary metabolism to regulate the intestinal barrier has shown promising prospective applications and has generated broad interest. The importance of the intestinal barrier in IBD is gradually being fully recognized, providing a new therapeutic perspective for improving inflammation and prognosis.

Key Words: Intestinal barrier; Inflammatory bowel disease; Bibliometrics; Visualization; Hotspots

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Core Tip: The complete composition and function of the intestinal mucosal barrier are essential for maintaining proper intestinal physiological and immune homeostasis. The study evaluated the developmental trends and research hotspots of intestinal barrier research in inflammatory bowel disease (IBD) using bibliometric methods. The results showed that Intestinal barrier field is developing rapidly with extensive cooperation. Targeting the gut microbiota and dietary metabolism to regulate intestinal barrier shown promising prospective applications. This may provide guidance and new insights for further research into improving IBD by modulating the intestinal barrier.

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INTRODUCTION

Specialized epithelial cells constitute barrier surfaces that separate host body from the external environment. The complete composition and function of intestinal mucosal barrier is essential for maintaining proper intestinal physiologic and immune homeostasis[1]. There is growing evidence that a deficit of in the intestinal barrier contributes to inflammation activation, especially in inflammatory bowel disease (IBD).

The intestinal barrier is mainly composed of intestinal epithelial cells, tight junctions and mucous layers. Genetic predisposition and environmental influences generally lead to disruption of the intestinal barrier in IBD, including tight junction changes, mucous layer defects, reduced antimicrobial peptide expression, and autophagy dysregulation[2,3]. This imbalance in the intestinal barrier allows external antigens to invade the body through damaged intestinal barrier, causing inappropriate immune activation. A series of studies have shown that mucosal healing and reestablishment of barrier integrity are significantly associated with clinical remission and improved patient outcomes[4-6]. Knowledge of the research status and focus hotspots of the current field is helpful to further explore the pathogenesis of IBD and search for potential therapeutic targets.

Bibliometrics has become an important methodology for researchers to actively accumulate knowledge and collect evidence. The bibliographic and visual analysis provides new angles to identify development trends and predict research prospects of this field by mathematics and statistics[7-9]. Therefore, this study aimed to explore the knowledge landscape of the intestinal barrier in IBD, hoping to provide guidance and new insights for further research in the field of IBD.

MATERIALS AND METHODS

Search strategies

Bibliometric and visualization analyses were conducted through the Web of Science Core Collection (WoSCC) database, which is one of the most widely used source for academic and bibliometric analysis. The WoS database covers a broad spectrum of academic disciplines and offers precise citation data, facilitating in-depth evaluation of topic development and trend evolution[10-12]. The search strategy was as follows: TS (Topic) = ("inflammatory bowel disease" OR "ulcerative colitis" OR "crohn's disease") AND TS = (intestin* OR gut OR bowel) AND TS = (barrier OR integrity OR permeability). The publication period was between 2001 and 2021 with the publication type limited to original articles written in English (Figure 1).

Data collection

Bibliometric indicators were extracted from selected publications to quantitatively evaluate the characteristics of the literature and its trends, including titles, publication years, authors, affiliations, countries/regions, journals, abstracts, keywords, citations and references. The H-index of scholars, impact factor (IF) and Journal Citation Reports division of journals were also obtained from the Web of Science. Inappropriate formulations of elements were modified and multiple phrases with the same meaning were merged into a single element. The author's significant publications were obtained from the *Reference Citation Analysis* (RCA, <https://referencecitationanalysis.com/>) database. RCA is an open citation

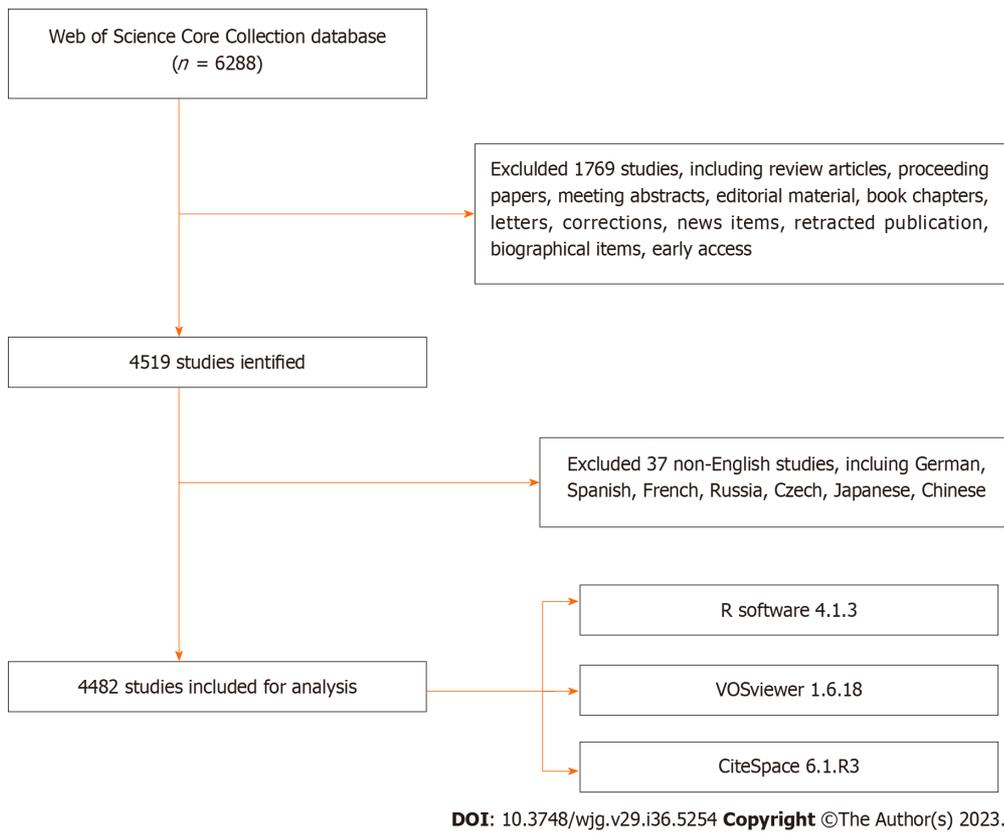


Figure 1 Flowchart of the literature screening.

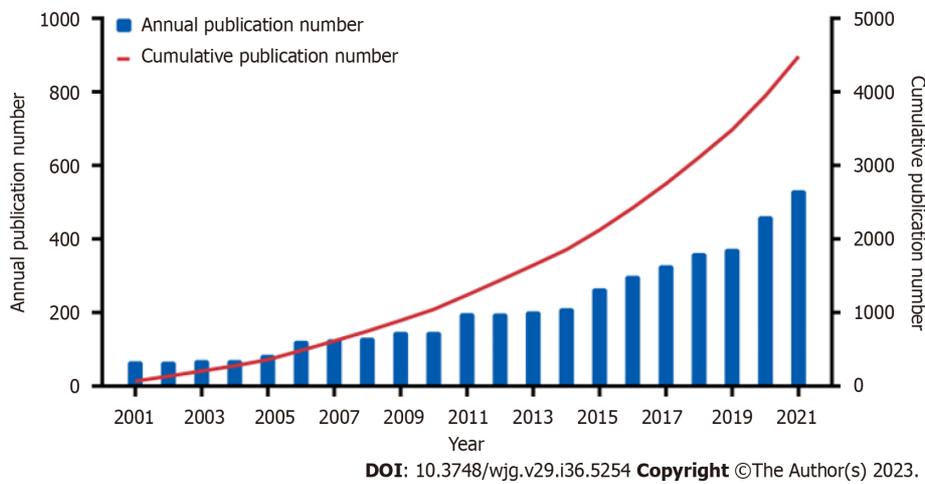


Figure 2 Annual and cumulative growth trends of publications.

analysis database covering various fields and is owned by Baishideng Publishing Group Inc., situated in Pleasanton, CA 94566, United States[13-15]. Scientific data were searched and collected independently by two researchers to ensure the reliability of the results.

Bibliometric analysis

Bibliometric analysis is a quantitative research method that is widely employed to evaluate academic development in a specific field. VOSviewer, CiteSpace and R package were used for constructing the scientometric network and knowledge visualization. VOSviewer was adopted to conduct co-authorship analysis of authors and countries/regions, co-occurrence analysis of keywords and co-citation analysis of journals. Co-authorship and co-occurrence relationships were defined as the simultaneous occurrence of two items, while co-citation relationships referred to two items being cited simultaneously by a third item[16,17]. The network graph generated by VOSviewer displays the size of nodes based on the number of publications, and the connection represents the relationship between nodes. Closely related nodes were

Table 1 The top 10 countries/regions and institutions in the intestinal barrier field relevant to inflammatory bowel disease

Country	Centrality	Count	Citation	CPR	Institution	Centrality	Count	Citation	CPR
United States	0.30	1534	90667	59.1	University of Chicago	0.08	87	8288	95.3
China	0.26	1383	25867	18.7	University of Calgary	0.06	84	4051	48.2
Germany	0.18	533	37129	69.7	Emory University	0.09	83	6183	74.5
Canada	0.09	352	17335	49.2	Shanghai Jiao Tong University	0.07	74	2323	31.4
United Kingdom	0.28	336	23096	68.7	University of Toronto	0.05	68	3125	46.0
Italy	0.12	266	12121	45.6	Harvard Medical School	0.07	62	1883	30.4
Japan	0.10	244	8467	34.7	Chinese Academy of Sciences	0.04	62	1600	25.8
France	0.16	226	11080	49.0	University of California San Diego	0.07	59	6034	102.3
Sweden	0.02	178	11089	62.3	University of Alberta	0.06	58	3310	57.1
Netherlands	0.03	168	7693	45.8	McMaster University	0.05	58	4047	69.8

CPR: Citations per publication.

divided into clusters of the same color.

CiteSpace was used to calculate the node centrality and keyword burst trends. Centrality measures the quantity of all the shortest paths passing through a node in a network, with nodes having centrality greater than 0.1 identified as key nodes of significant importance. Burst detection was employed to identify keywords that experienced a sudden surge in their frequency of appearance during a specific time frame. The green line indicates the occurrence and development of keywords, while the red line represents the duration of the burst.

In addition, R software was used to develop the distribution of scientific productivity among authors and different countries/regions. Lotka's Law demonstrates the relationship between the frequency of scientific paper authorship and the number of publications, revealing an unbalanced distribution of author productivity.

RESULTS

Growth trend of publications

A total of 4482 publications related to the intestinal barrier in IBD were published between 2002-2022. There were only a small number of studies in the early stage, with less than 100 articles each year (Figure 2). During the middle period from 2006 to 2014, the number of publications showed a slow increase. Subsequently, annual publications have grown rapidly, with 53.8% of all publications occurring in the past two decades. Increasing attention to the intestinal barrier in IBD from researchers has driven further development of this field.

Countries/regions and institutions

Intestinal barrier research in IBD has been carried out at 4650 institutions from 89 countries/regions (Figure 3A). The United States is dominant in this field with the highest number of publications ($n = 1426$) and citations ($n = 89615$). The next most productive countries/regions were China and Germany (Table 1). In addition, Germany achieved the highest citations per publication at 73.2. These countries were divided into seven clusters to show the extensive cooperation between nodes (Figure 3B). Multiple nodes played a strong bridging role in the collaboration, most notably for the United States, the United Kingdom and China.

The University of Chicago conducted the largest number of studies ($n = 8236$), followed by Emory University ($n = 6437$) and the University of Calgary ($n = 4064$). Among them, the University of California San Diego exhibited the highest average citations of 102.3. Cooperation between institutions is more evident than that between countries, which enables further exploration and collaboration on the subject (Figure 3C).

Authors and cited authors

A total of 26245 authors have participated in the field of intestinal barrier research in IBD. Seventy-six percent of the authors contributed one article, whereas only 6.0% contributed more than 3 articles (Figure 4A). Turner[18] ($n = 35$), Söderholm *et al*[19-21] ($n = 32$) and Keshavarzian *et al*[22-25] ($n = 28$) were the most productive authors (Table 2). Although Turner[18] from Harvard Medical School also has the most citations ($n = 3399$), the nodes of these scholars were more dispersed across different clusters rather than being aggregated (Figure 4B). Close communication and similar research directions exist in the nodes of the same cluster. In addition, the clusters displaying different colors are not entirely independent and exhibit a relatively low level of connections.

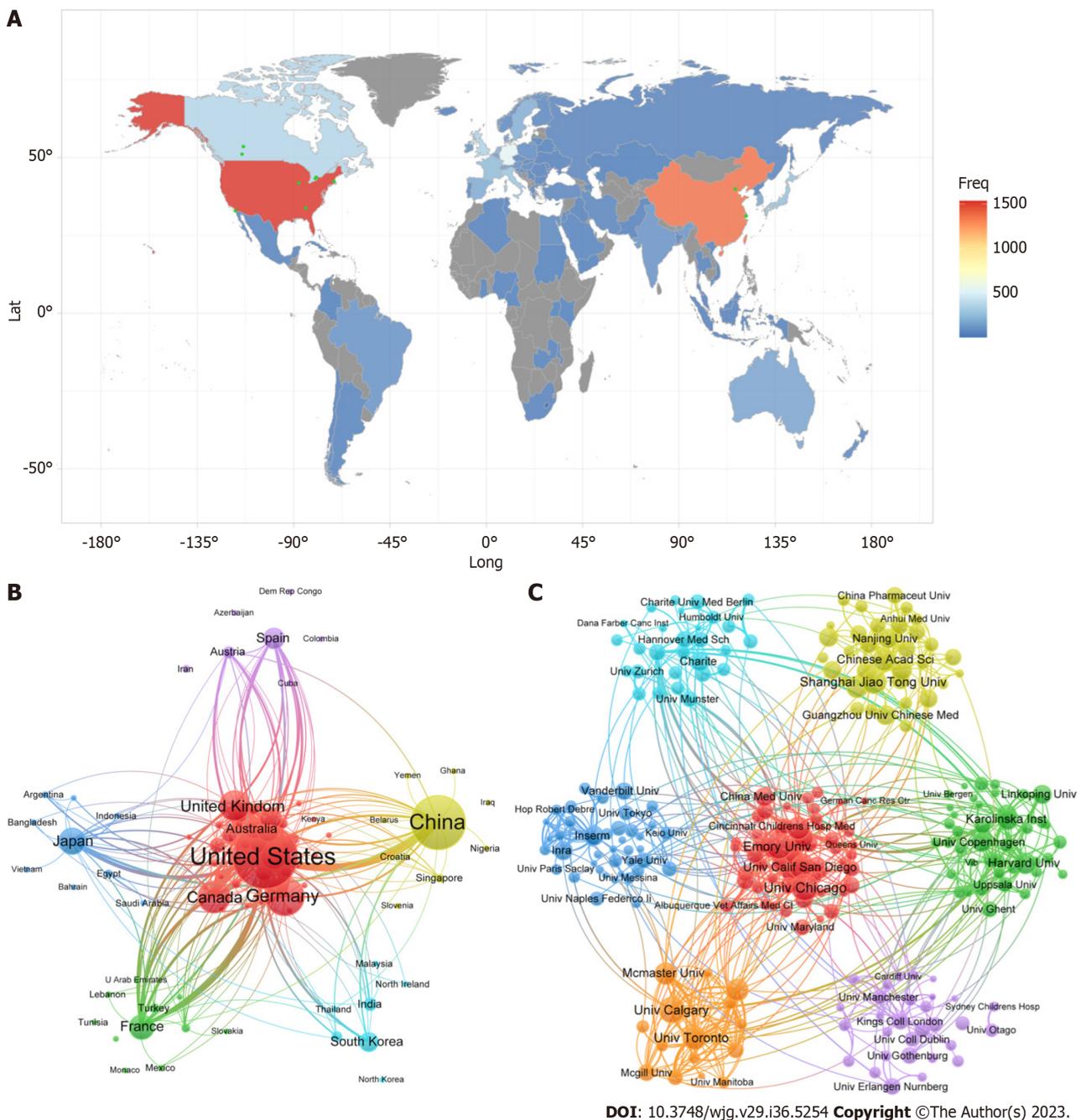


Figure 3 Network visualization map of countries/regions and institutions. A: Geographic distribution map based on the total publications; B: Cooperation map of countries/regions; C: Cooperation map of institutions. The nodes in the network map represent countries/regions, and the node size is proportional to the number of publications. The thickness of the connecting line between nodes indicates the strength of the co-authorship relationships.

Journals and cited academic journals

There were 1045 journals involved in the publication of intestinal barrier research in IBD. IBDs contributed the most articles ($n = 258$), while Gastroenterology received the most citations (Table 3). The next most productive journals were PLoS One ($n = 145$) and the American Journal of Physiology-Gastrointestinal and Liver Physiology ($n = 119$). Articles published in established journals exhibit a higher average citation count, which is consistent with the IF of the journal.

Among them, the majority of journals had an early initiation into this field (Figure 5A). Despite being relatively new to the field, *Frontiers in Immunology*, *Food & Function* and *Scientific Reports* demonstrated rapid development. As shown in Figure 5B, these journals were divided into five clusters based on the similarity of their themes. The higher the number of co-citations between two journals, the stronger is the link between the two nodes.

Keywords co-occurrence, clusters and bursts

Keywords were extracted for multidimensional analysis to determine the research hotspot and exploration direction in the specific field. The keywords appearing at high frequency related with intestinal barrier in IBD were identified, including molecules, pathological processes and treatments (Table 4). Nuclear factor kappa B (NF- κ B) ($n = 418$), tumor

Table 2 The top 10 productive authors and cited authors in the intestinal barrier field relevant to inflammatory bowel disease

Ref.	Count	H-index	Ref.	Count	H-index
Turner[18]	35	78	Turner[18]	3399	78
Söderholm <i>et al</i> [19-21]	32	46	Neurath[35]	3257	87
Keshavarzian <i>et al</i> [22-25]	28	77	Ma <i>et al</i> [40]	2811	26
Rogler[59]	26	66	Hansson[60]	2785	71
Neurath[35]	25	87	Johansson <i>et al</i> [61]	2725	47
Schulzke <i>et al</i> [62]	23	58	Al-Sadi <i>et al</i> [63]	2489	21
Merlin <i>et al</i> [64]	23	62	Danese and Fiocchi[46]	2440	96
Ma <i>et al</i> [40]	22	26	Becker <i>et al</i> [44]	2319	61
Fromm <i>et al</i> [34]	21	66	Schulzke <i>et al</i> [62]	1852	58
Vermeire <i>et al</i> [51]	21	122	Nusrat <i>et al</i> [33]	1838	47

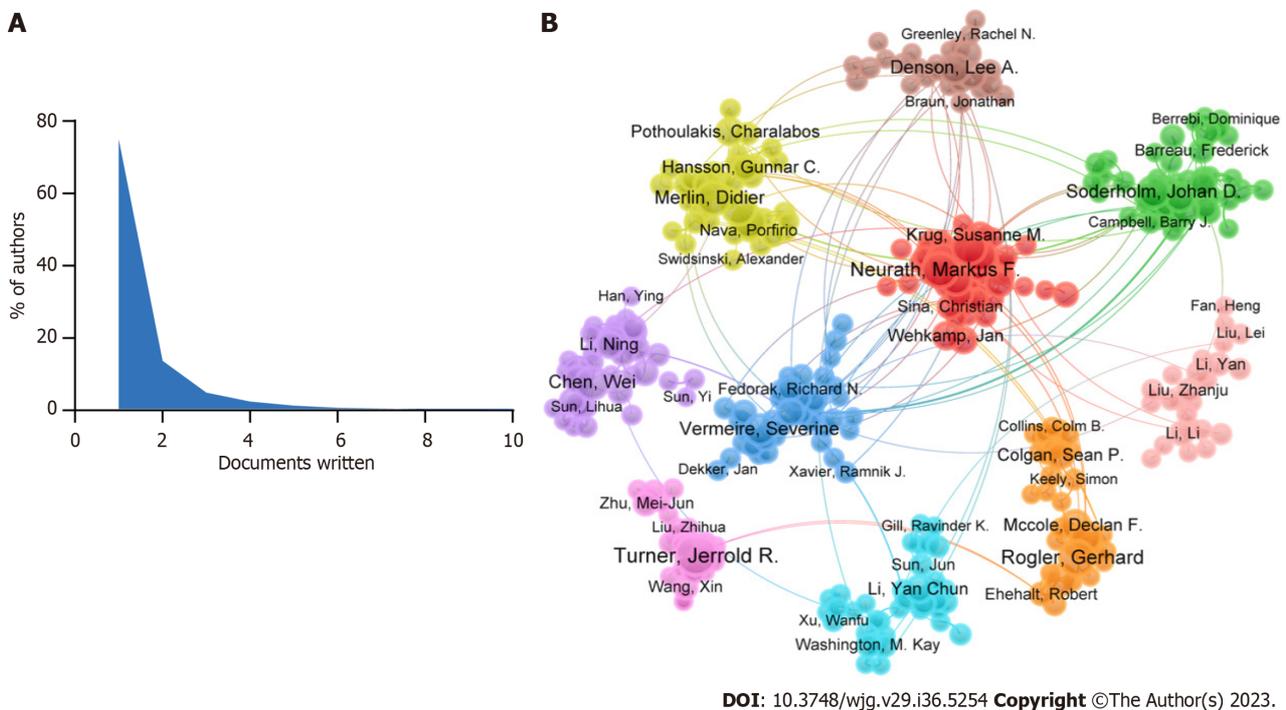


Figure 4 Network visualization map of authors. A: Scientific productivity of authors based on Lotka's Law; B: Cooperation map of authors. Each node represents a different author, and the connections between the nodes reflect the co-authorship relationships.

necrosis factor- α (TNF- α) ($n = 329$) and interferon (IFN)- γ ($n = 173$) were the most commonly implicated molecules. Apoptosis ($n = 292$), oxidative stress ($n = 232$) and metabolism ($n = 88$) were widely involved in research on the intestinal barrier. Regarding treatments, probiotics ($n = 237$), nanoparticles ($n = 77$) and antioxidant ($n = 73$) have attracted extensive attention.

The network map was classified into four clusters by co-occurrence analysis, which portrayed the fundamental knowledge structure of related research fields (Figure 6). Based on the closeness between nodes, the largest blue cluster contains keywords associated with clinical characteristics, such as risk, children and therapy. The red cluster was mainly composed of tight junctions, permeability and transport, which drew attention to barrier function in IBD. The yellow cluster mainly focused on the gut microbiota, including probiotics, metabolism and dysbiosis. The green cluster was involved in the regulation of inflammation, with keywords such as apoptosis, NF- κ B and oxidative stress.

The visual keyword burst trend was constructed to show the major research interests in the intestinal barrier field (Figure 7). Early keyword bursts were mainly concentrated on intestinal barrier function and related inflammatory molecules, where intestinal permeability showed strong bursts. During the middle period, only a limited number of keywords exhibited an outbreak trend, such as autophagy, matrix metalloproteinase, and vitamin D. In contrast, targeted intestinal barrier therapy has been receiving growing interest from researchers, with a particular focus on antioxidants, *Akkermansia muciniphila*, nanoparticles, short-chain fatty acids, and extracellular vesicles as recent areas of research

Table 3 The top 10 journals and co-cited journals in the intestinal barrier field relevant to inflammatory bowel disease

Journal	Count	IF (2022)	JCR (2022)	Cited journal	Citation	IF (2022)	JCR (2022)
<i>Inflammatory Bowel Diseases</i>	258	7.290	Q1	<i>Gastroenterology</i>	14345	33.883	Q1
<i>PLoS One</i>	145	3.752	Q2	<i>Gut</i>	12840	31.793	Q1
<i>American Journal of Physiology-Gastrointestinal and Liver Physiology</i>	119	4.871	Q1/Q2	<i>Inflammatory Bowel Diseases</i>	10188	69.504	Q1
<i>Gastroenterology</i>	118	33.883	Q1	<i>American Journal of Physiology-Gastrointestinal and Liver Physiology</i>	8962	4.871	Q1/Q2
<i>World Journal of Gastroenterology</i>	107	5.374	Q2	<i>Plos One</i>	6557	3.752	Q2
<i>Gut</i>	94	31.793	Q1	<i>Proceedings of the National Academy of Sciences of the United States of America</i>	6250	12.779	Q1
<i>Frontiers in Immunology</i>	86	8.786	Q1	<i>Nature</i>	4978	69.504	Q1
<i>Food & Function</i>	83	6.317	Q1	<i>World Journal of Gastroenterology</i>	4509	5.374	Q2
<i>Digestive Diseases and Sciences</i>	78	3.487	Q3	<i>Journal of Immunology</i>	4142	5.426	Q2
<i>Scientific Reports</i>	74	4.996	Q2	<i>Journal of Clinical Investigation</i>	3390	19.456	Q1

IF: Impact factor; JCR: Journal Citation Reports.

Table 4 The top 10 molecules, pathological processes and disease in the intestinal barrier field relevant to inflammatory bowel disease

Molecule	Count	Pathological process	Count	Treatment	Count
NF-kB	418	Apoptosis	292	Probiotic	237
TNF- α	329	Oxidative stress	232	Nanoparticles	77
IFN-gamma	173	Metabolism	88	Antioxidant	73
Mucin	100	Dysbiosis	83	Butyrate	70
Occludin	90	Autophagy	82	Infliximab	66
Nitric oxide	87	Infection	80	Polysaccharide	48
ZO-1	55	Bacterial translocation	68	Prebiotic	40
Claudin-2	54	Endoplasmic reticulum stress	65	Fecal microbiota transplantation	37
E-cadherin	51	Transport	59	Lactobacillus	32
Antimicrobial peptide	50	Absorption	50	Vitamin D	26

NF-kB: Nuclear factor kappa B; TNF- α : Tumor necrosis factor- α ; IFN: Interferon.

attention.

DISCUSSION

The intestinal barrier refers to the physiological barrier composed of intestinal epithelial cells and mucous layers, which plays a crucial role in the onset and progression of IBD[26-28]. The disruption of the intestinal barrier in IBD patients is closely related to the severity of intestinal inflammation and immune response. The significant increase in the number of annual publications indicates a rapidly growing interest and recognition of the intestinal barrier field.

The United States occupied the dominant places in the intestinal barrier field with the highest number of publications and citations. Although China is developing rapidly in this field, it exhibits lower average citation rates. Other countries such as Germany and the United Kingdom have conducted more in-depth research and have the highest average citation rates. It is necessary to increase the depth and influence of research to address the imbalance between the number of publications and citations. Extensive collaboration is observed in countries such as the United States and the United Kingdom, however, institutional collaboration tends to show a trend of geographic alignment in terms of its distribution.

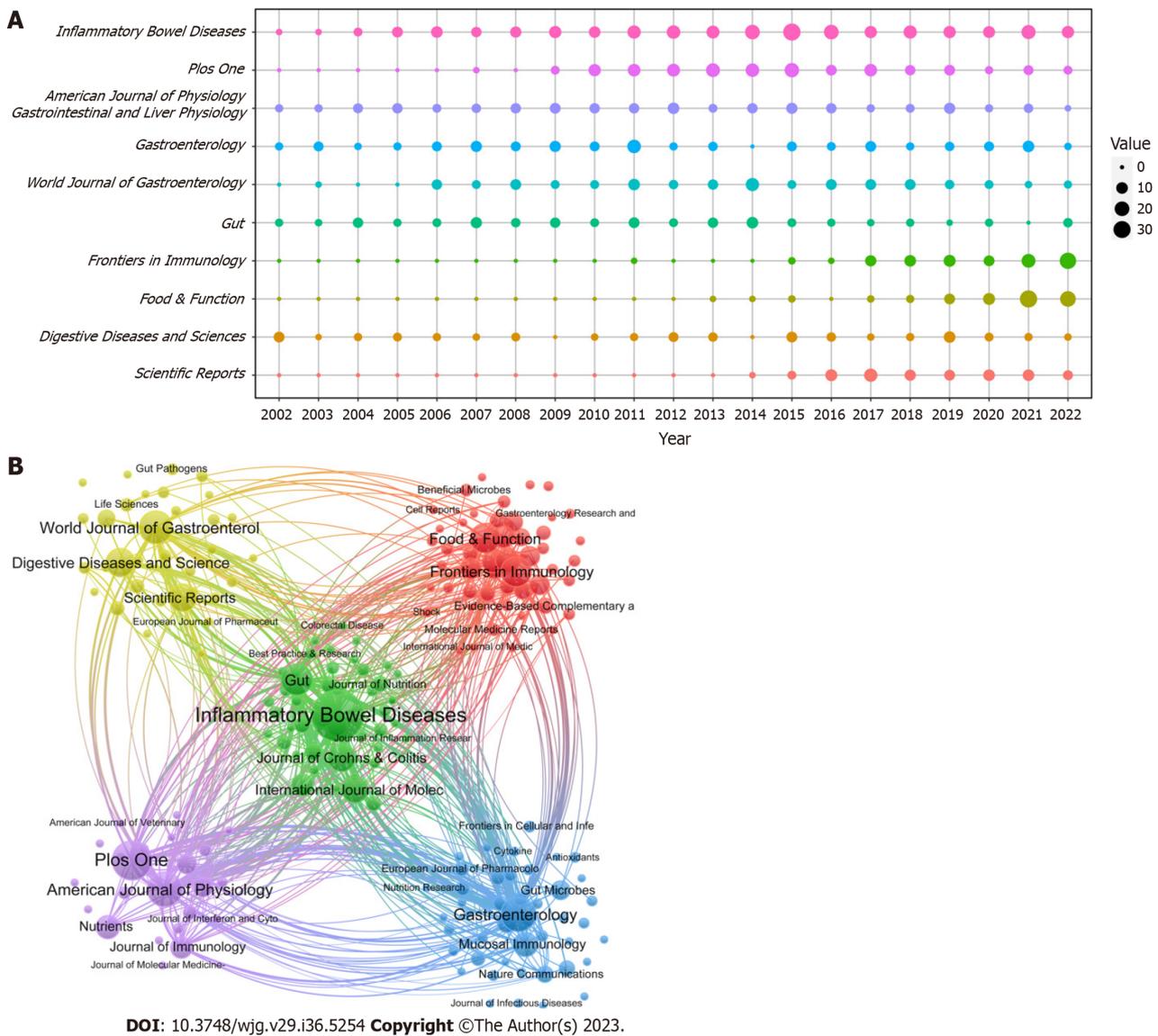
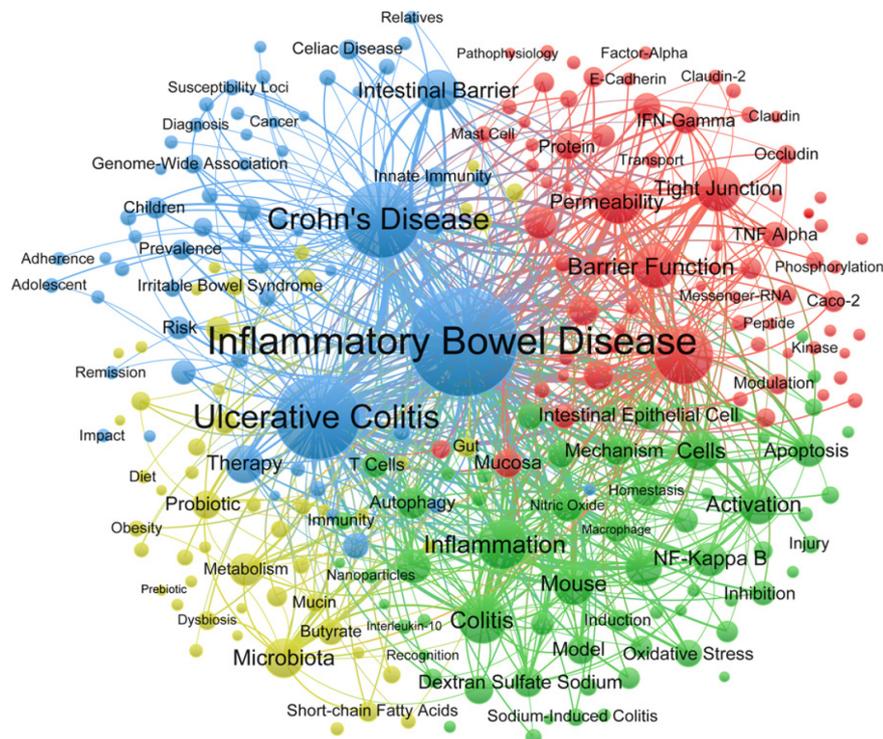


Figure 5 Network visualization map of authors. A: Annual number of publications for the top 10 journals; B: Cooperation map of Journals. The nodes in the graph represented journals, and the connections between the nodes reflect the co-authorship relationships.

The author’s productivity is consistent with Lotka’s Law. Turner[18] from Harvard Medical School has made significant contributions to intestinal barrier research. He and His group found that MLCK-dependent tight junction dysfunction can cause immune system activation and experimental colitis induction[18,29]. Söderholm *et al*[19-21] mainly investigated how chronic stress triggers intestinal inflammation, which is closely associated with mast cells. In contrast, Keshavarzian *et al*[22-25] focused on the connections between colitis and other systemic diseases, such as Alzheimer’s disease. These findings demonstrate the feasibility of improving IBD through barrier repair, which could become a strong candidate for nonimmune suppressive methods to achieve or maintain disease remission.

Intestinal barrier research was initially published in gastroenterology journals, particularly in IBD. The exploration of the intestinal barrier has revealed extensive relationships between this field and several other research areas, such as physiology, biochemistry and immunology. As a result, there has been an increase in the number of interdisciplinary journals to better cater to the needs of this rapidly expanding field. Comprehensive journals have a wide readership and are more conducive to interdisciplinary collaboration.

The integrity of the intestinal barrier has been found to be closely associated with host defense and regulates the progression of IBD. Keywords in the field of the intestinal barrier mainly revolve around four aspects, encompassing clinical characteristics, barrier function, the gut microbiota, and inflammation regulation. Studies have reported that disruptions in the intestinal barrier lead to persistent intestinal symptoms and poor prognosis in IBD patients[30,31]. The increase in intestinal permeability is a marker of intestinal barrier dysfunction. The tight junction between intestinal epithelial cells are dynamic structures capable of swift adjustments, which determine the mucosal permeability[32-34]. The active transcellular transport of ions and water relies on the presence of intact tight junction barriers. Damage to epithelial cells triggers the release of inflammatory cytokines such as TNF and IFN- γ [35-37]. Stimulation of TNF downregulates the expression of tight junction proteins and changes their localization in colonic epithelial cells[38-40]. Insufficiency of interleukin (IL)-10, excessive production of nitric oxide, and impaired autophagy can also induce an



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Figure 6 The network map of keywords in the intestinal barrier field relevant to inflammatory bowel disease. The network map displays the size of nodes based on the number of publications, and the connection represents the co-occurrence relationships between nodes. Closely related nodes were divided into clusters of the same color.

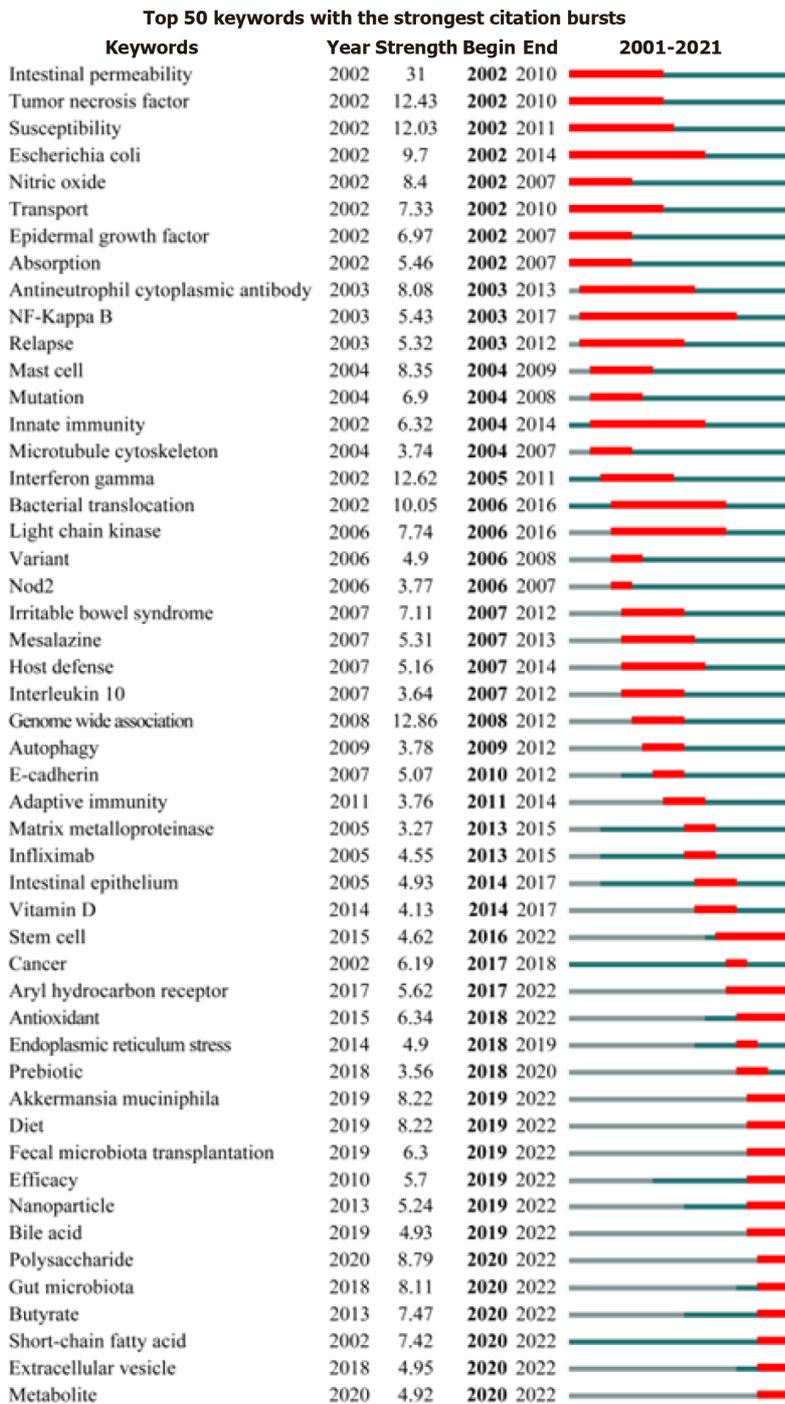
overactive immune system and enhance barrier loss[41-44], which in turn causes further leakage of luminal substances and amplifies local inflammation. The homeostasis of intestinal barrier can be affected either directly by inducing intracellular signaling pathways, or indirectly by the gut microbiota as an interacting partner to a greater extent[45,46]. Reduced intestinal community richness in patients with IBD leads to loss of intestinal homeostasis and inappropriate immune activation, which is characterized by a decrease in Firmicutes and proliferation of facultative anaerobic *Enterobacteriaceae* and adherent invasive *Escherichia coli*[47-49].

Recently, there has been widespread research interest in improving the intestinal barrier by targeting gut microbiota and dietary metabolism. Exogenous administration of probiotics can influence the composition and characteristics of the intestinal microbiota to exert intervention effects, typically in combination with prebiotics. The colonization of the beneficial bacterium *Akkermansia muciniphila* increased the number of goblet cells and upregulated the expression of genes encoding mucin[50]. Although fecal microbiota transplantation (FMT) has been shown to induce clinical remission of ulcerative colitis, the long-term effects of FMT as maintenance therapy have not been validated in IBD[51,52]. Microbial and dietary metabolites also demonstrate a regulatory influence on the intestinal barrier. Short-chain fatty acids produced by dietary fiber fermentation are the main energy source of colon cells, promoting the integrity of epithelial tight junctions and mucus secretion, which may be related to the activation of AMPK and STAT3[53,54]. The observed abnormality in bile acid metabolism may contribute to an enhanced epithelial inflammatory response. The reduction in secondary bile acids weakens the inhibition of IL-8 secretion[55] and the activation of TGR5 receptors in intestinal stem cells[56]. In addition, extracellular vesicles produced by host cells and microorganisms are also involved in cellular communication. Immune cells such as mast cells can secrete extracellular vesicles containing miR-223 to regulate intestinal permeability with downregulation of tight junction proteins[57]. Notably, extracellular vesicles themselves also represent a natural nanomaterial for drug delivery, that can improve the therapeutic effect and reduce systemic exposure in healthy tissues to promote remission of IBD[58].

The bibliometric analysis offers novel and objective insights into evolving intestinal barrier research hotspots and trends across multiple dimensions. Inevitably, there were several limitations to this study. The data were extracted from the WoSCC database, potentially omitting some information from other sources. Furthermore, while bibliometric methods based on natural language processing are susceptible to biases from subject categorization and citation behavior, a sufficient sample size helps ensure the accuracy of research findings to some extent.

CONCLUSION

In conclusion, the intestinal barrier field is developing rapidly with extensive cooperation. Targeting the gut microbiota and dietary metabolism to regulate the intestinal barrier shown promising prospective applications and has generated



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Figure 7 The top 50 keywords with the strongest citation bursts in the intestinal barrier field relevant to inflammatory bowel disease. The green line indicates the occurrence and development of keywords, while the red line represents the duration of the burst.

broad interest. The importance of the intestinal barrier in IBD is gradually being fully recognized, providing a new therapeutic perspective for improving inflammation.

ARTICLE HIGHLIGHTS

Research background

Barrier surfaces composed of specialized epithelial cells separate the host body from the external environment, and are essential for maintaining proper intestinal physiologic and immune homeostasis. There is growing evidence that the mucosal healing and reestablishment of barrier integrity are significantly associated with clinical remission and improved patient outcomes.

Research motivation

With the rapid expansion of research topics, it has become challenging for researchers to accumulate knowledge and actively collect evidence in specific field. The bibliographic and visual analysis provides new angles to identify development trends and predict research prospects of this field by mathematics and statistics.

Research objectives

The purpose of this study was to provide a comprehensive overview of the knowledge structure and research hotspots of intestinal barrier in inflammatory bowel disease (IBD).

Research methods

The publications related to intestinal barrier in IBD were obtained from the Web of Science Core Collection database. Bibliometric analysis and visualization were conducted using VOSviewer, CiteSpace and R software.

Research results

A comprehensive analysis of 4482 articles published between 2002 and 2022 has revealed significant insights in the realm of intestinal barrier research. Intestinal barrier research mainly focuses on four aspects, including clinical characteristics, barrier function, gut microbiota, and inflammation regulation. The keywords appearing at high frequency related with intestinal barrier in IBD were detected, including nuclear factor kappa B, tumor necrosis factor- α , apoptosis, oxidative stress and probiotics. Among them, antioxidants, Akkermansia muciniphila, nanoparticles, short-chain fatty acids and extracellular vesicles have received extensive interest in current research.

Research conclusions

The intestinal barrier field is developing rapidly with extensive cooperation. Targeting the gut microbiota and dietary metabolism to regulate intestinal barrier shown promising applied prospective and has generated broad interest.

Research perspectives

The importance of the intestinal barrier in IBD is gradually being fully recognized, providing a new therapeutic perspective for improving inflammation.

FOOTNOTES

Author contributions: Zhou XJ and Xie Y designed the study; Zhou F and Wu NZ conducted data extraction; Zhou F performed data analysis and drafted the manuscript; Zhou XJ and Xie Y interpreted the data and revised the manuscript; All authors have read and approve the final manuscript.

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REFERENCES

- Villablanca EJ**, Selin K, Hedin CRH. Mechanisms of mucosal healing: treating inflammatory bowel disease without immunosuppression? *Nat Rev Gastroenterol Hepatol* 2022; **19**: 493-507 [PMID: 35440774 DOI: 10.1038/s41575-022-00604-y]
- Ramos GP**, Papadakis KA. Mechanisms of Disease: Inflammatory Bowel Diseases. *Mayo Clin Proc* 2019; **94**: 155-165 [PMID: 30611442 DOI: 10.1016/j.mayocp.2018.09.013]

- 3 **Chen Y**, Cui W, Li X, Yang H. Interaction Between Commensal Bacteria, Immune Response and the Intestinal Barrier in Inflammatory Bowel Disease. *Front Immunol* 2021; **12**: 761981 [PMID: 34858414 DOI: 10.3389/fimmu.2021.761981]
- 4 **Schultz I**, Keita ÁV. Cellular and Molecular Therapeutic Targets in Inflammatory Bowel Disease-Focusing on Intestinal Barrier Function. *Cells* 2019; **8** [PMID: 30813280 DOI: 10.3390/cells8020193]
- 5 **Michaels M**, Madsen KL. Immunometabolism and microbial metabolites at the gut barrier: Lessons for therapeutic intervention in inflammatory bowel disease. *Mucosal Immunol* 2023; **16**: 72-85 [PMID: 36642380 DOI: 10.1016/j.mucimm.2022.11.001]
- 6 **Otte ML**, Lama Tamang R, Papapanagiotou J, Ahmad R, Dhawan P, Singh AB. Mucosal healing and inflammatory bowel disease: Therapeutic implications and new targets. *World J Gastroenterol* 2023; **29**: 1157-1172 [PMID: 36926666 DOI: 10.3748/wjg.v29.i7.1157]
- 7 **Agarwal A**, Durairajanayagam D, Tatagari S, Esteves SC, Harlev A, Henkel R, Roychoudhury S, Homa S, Puchalt NG, Ramasamy R, Majzoub A, Ly KD, Tvrda E, Assidi M, Kesari K, Sharma R, Banihani S, Ko E, Abu-Elmagd M, Gosalvez J, Bashiri A. Bibliometrics: tracking research impact by selecting the appropriate metrics. *Asian J Androl* 2016; **18**: 296-309 [PMID: 26806079 DOI: 10.4103/1008-682X.171582]
- 8 **Chen C**, Song M. Visualizing a field of research: A methodology of systematic scientometric reviews. *PLoS One* 2019; **14**: e0223994 [PMID: 31671124 DOI: 10.1371/journal.pone.0223994]
- 9 **Chen C**. Searching for intellectual turning points: progressive knowledge domain visualization. *Proc Natl Acad Sci U S A* 2004; **101** Suppl 1: 5303-5310 [PMID: 14724295 DOI: 10.1073/pnas.0307513100]
- 10 **Cheng K**, Guo Q, Shen Z, Yang W, Wang Y, Sun Z, Wu H. Bibliometric Analysis of Global Research on Cancer Photodynamic Therapy: Focus on Nano-Related Research. *Front Pharmacol* 2022; **13**: 927219 [PMID: 35784740 DOI: 10.3389/fphar.2022.927219]
- 11 **Cheng K**, Zhang H, Guo Q, Zhai P, Zhou Y, Yang W, Wang Y, Lu Y, Shen Z, Wu H. Emerging trends and research foci of oncolytic virotherapy for central nervous system tumors: A bibliometric study. *Front Immunol* 2022; **13**: 975695 [PMID: 36148235 DOI: 10.3389/fimmu.2022.975695]
- 12 **Yeung AWK**. Web of Science, PubMed and publishers for mislabelled review papers. *Curr Sci* 2019; **116**: 1909-1914 [DOI: 10.18520/cs/v116/i11/1909-1914]
- 13 **Wang JL**, Ma YJ, Ma L, Ma N, Guo DM, Ma LS. Baishideng's Reference Citation Analysis database announces the first Journal Article Influence Index of 101 core journals and a list of high-quality academic journals in gastroenterology and hepatology. *World J Gastroenterol* 2022; **28**: 5383-5394 [PMID: 36312837 DOI: 10.3748/wjg.v28.i37.5383]
- 14 **Wang JL**, Ma YJ, Ma L, Ma N, Guo DM, Ma LS. Baishideng's Reference Citation Analysis database announces the first Journal Article Influence Index of 104 core journals and a list of high-quality academic journals in orthopedics. *World J Orthop* 2022; **13**: 891-902 [PMID: 36312521 DOI: 10.5312/wjo.v13.i10.891]
- 15 **Wang JL**, Ma YJ, Ma L, Ma N, Guo DM, Ma LS. Baishideng's Reference Citation Analysis database announces the first Article Influence Index of multidisciplinary scholars. *World J Clin Cases* 2022; **10**: 10391-10398 [PMID: 36312463 DOI: 10.12998/wjcc.v10.i29.10391]
- 16 **Wu Z**, Cheng K, Shen Z, Lu Y, Wang H, Wang G, Wang Y, Yang W, Sun Z, Guo Q, Wu H. Mapping knowledge landscapes and emerging trends of sonodynamic therapy: A bibliometric and visualized study. *Front Pharmacol* 2022; **13**: 1048211 [PMID: 36699067 DOI: 10.3389/fphar.2022.1048211]
- 17 **Wu H**, Cheng K, Tong L, Wang Y, Yang W, Sun Z. Knowledge structure and emerging trends on osteonecrosis of the femoral head: a bibliometric and visualized study. *J Orthop Surg Res* 2022; **17**: 194 [PMID: 35346273 DOI: 10.1186/s13018-022-03068-7]
- 18 **Turner JR**. Molecular basis of epithelial barrier regulation: from basic mechanisms to clinical application. *Am J Pathol* 2006; **169**: 1901-1909 [PMID: 17148655 DOI: 10.2353/ajpath.2006.060681]
- 19 **Söderholm JD**, Yang PC, Ceponis P, Vohra A, Riddell R, Sherman PM, Perdue MH. Chronic stress induces mast cell-dependent bacterial adherence and initiates mucosal inflammation in rat intestine. *Gastroenterology* 2002; **123**: 1099-1108 [PMID: 12360472 DOI: 10.1053/gast.2002.36019]
- 20 **Söderholm JD**, Yates DA, Gareau MG, Yang PC, MacQueen G, Perdue MH. Neonatal maternal separation predisposes adult rats to colonic barrier dysfunction in response to mild stress. *Am J Physiol Gastrointest Liver Physiol* 2002; **283**: G1257-G1263 [PMID: 12388189 DOI: 10.1152/ajpgi.00314.2002]
- 21 **Söderholm JD**. Mast cells and mastocytosis. *Dig Dis* 2009; **27** Suppl 1: 129-136 [PMID: 20203509 DOI: 10.1159/000268133]
- 22 **Keshavarzian A**, Farhadi A, Forsyth CB, Rangan J, Jakate S, Shaikh M, Banan A, Fields JZ. Evidence that chronic alcohol exposure promotes intestinal oxidative stress, intestinal hyperpermeability and endotoxemia prior to development of alcoholic steatohepatitis in rats. *J Hepatol* 2009; **50**: 538-547 [PMID: 19155080 DOI: 10.1016/j.jhep.2008.10.028]
- 23 **Keshavarzian A**, Green SJ, Engen PA, Voigt RM, Naqib A, Forsyth CB, Mutlu E, Shannon KM. Colonic bacterial composition in Parkinson's disease. *Mov Disord* 2015; **30**: 1351-1360 [PMID: 26179554 DOI: 10.1002/mds.26307]
- 24 **Keshavarzian A**, Mutlu E, Guzman JP, Forsyth C, Banan A. Phosphodiesterase 4 inhibitors and inflammatory bowel disease: emerging therapies in inflammatory bowel disease. *Expert Opin Investig Drugs* 2007; **16**: 1489-1506 [PMID: 17714033 DOI: 10.1517/13543784.16.9.1489]
- 25 **Keshavarzian A**, Engen P, Bonvegna S, Cilia R. The gut microbiome in Parkinson's disease: A culprit or a bystander? *Prog Brain Res* 2020; **252**: 357-450 [PMID: 32247371 DOI: 10.1016/bs.pbr.2020.01.004]
- 26 **Mehandru S**, Colombel JF. The intestinal barrier, an arbitrator turned provocateur in IBD. *Nat Rev Gastroenterol Hepatol* 2021; **18**: 83-84 [PMID: 33318680 DOI: 10.1038/s41575-020-00399-w]
- 27 **Odenwald MA**, Turner JR. The intestinal epithelial barrier: a therapeutic target? *Nat Rev Gastroenterol Hepatol* 2017; **14**: 9-21 [PMID: 27848962 DOI: 10.1038/nrgastro.2016.169]
- 28 **Yu LC**. Microbiota dysbiosis and barrier dysfunction in inflammatory bowel disease and colorectal cancers: exploring a common ground hypothesis. *J Biomed Sci* 2018; **25**: 79 [PMID: 30413188 DOI: 10.1186/s12929-018-0483-8]
- 29 **Graham WV**, He W, Marchiando AM, Zha J, Singh G, Li HS, Biswas A, Ong MLDM, Jiang ZH, Choi W, Zuccola H, Wang Y, Griffith J, Wu J, Rosenberg HJ, Snapper SB, Ostrov D, Meredith SC, Miller LW, Turner JR. Intracellular MLCK1 diversion reverses barrier loss to restore mucosal homeostasis. *Nat Med* 2019; **25**: 690-700 [PMID: 30936544 DOI: 10.1038/s41591-019-0393-7]
- 30 **Chang J**, Leong RW, Wasinger VC, Ip M, Yang M, Phan TG. Impaired Intestinal Permeability Contributes to Ongoing Bowel Symptoms in Patients With Inflammatory Bowel Disease and Mucosal Healing. *Gastroenterology* 2017; **153**: 723-731.e1 [PMID: 28601482 DOI: 10.1053/j.gastro.2017.05.056]
- 31 **Turpin W**, Lee SH, Raygoza Garay JA, Madsen KL, Meddings JB, Bedrani L, Power N, Espin-Garcia O, Xu W, Smith MI, Griffiths AM, Moayyedi P, Turner D, Seidman EG, Steinhart AH, Marshall JK, Jacobson K, Mack D, Huynh H, Bernstein CN, Paterson AD; Crohn's and Colitis Canada Genetic Environmental Microbial Project Research Consortium; CCC GEM Project recruitment site directors include Maria

- Abreu, Croitoru K. Increased Intestinal Permeability Is Associated With Later Development of Crohn's Disease. *Gastroenterology* 2020; **159**: 2092-2100.e5 [PMID: 32791132 DOI: 10.1053/j.gastro.2020.08.005]
- 32 **Slifer ZM**, Blikslager AT. The Integral Role of Tight Junction Proteins in the Repair of Injured Intestinal Epithelium. *Int J Mol Sci* 2020; **21** [PMID: 32024112 DOI: 10.3390/ijms21030972]
- 33 **Nusrat A**, Turner JR, Madara JL. Molecular physiology and pathophysiology of tight junctions. IV. Regulation of tight junctions by extracellular stimuli: nutrients, cytokines, and immune cells. *Am J Physiol Gastrointest Liver Physiol* 2000; **279**: G851-G857 [PMID: 11052980 DOI: 10.1152/ajpgi.2000.279.5.G851]
- 34 **Fromm M**, Krug SM, Zeissig S, Richter JF, Rosenthal R, Schulzke JD, Günzel D. High-resolution analysis of barrier function. *Ann N Y Acad Sci* 2009; **1165**: 74-81 [PMID: 19538291 DOI: 10.1111/j.1749-6632.2009.04047.x]
- 35 **Neurath MF**. Targeting cytokines in inflammatory bowel disease. *Sci Transl Med* 2022; **14**: eabq4473 [PMID: 36516267 DOI: 10.1126/scitranslmed.abq4473]
- 36 **Abraham C**, Abreu MT, Turner JR. Pattern Recognition Receptor Signaling and Cytokine Networks in Microbial Defenses and Regulation of Intestinal Barriers: Implications for Inflammatory Bowel Disease. *Gastroenterology* 2022; **162**: 1602-1616.e6 [PMID: 35149024 DOI: 10.1053/j.gastro.2021.12.288]
- 37 **Luissint AC**, Parkos CA, Nusrat A. Inflammation and the Intestinal Barrier: Leukocyte-Epithelial Cell Interactions, Cell Junction Remodeling, and Mucosal Repair. *Gastroenterology* 2016; **151**: 616-632 [PMID: 27436072 DOI: 10.1053/j.gastro.2016.07.008]
- 38 **Al-Sadi R**, Guo S, Ye D, Rawat M, Ma TY. TNF- α Modulation of Intestinal Tight Junction Permeability Is Mediated by NIK/IKK- α Axis Activation of the Canonical NF- κ B Pathway. *Am J Pathol* 2016; **186**: 1151-1165 [PMID: 26948423 DOI: 10.1016/j.ajpath.2015.12.016]
- 39 **Yu M**, Wang Q, Ma Y, Li L, Yu K, Zhang Z, Chen G, Li X, Xiao W, Xu P, Yang H. Aryl Hydrocarbon Receptor Activation Modulates Intestinal Epithelial Barrier Function by Maintaining Tight Junction Integrity. *Int J Biol Sci* 2018; **14**: 69-77 [PMID: 29483826 DOI: 10.7150/ijbs.22259]
- 40 **Ma TY**, Iwamoto GK, Hoa NT, Akotia V, Pedram A, Boivin MA, Said HM. TNF-alpha-induced increase in intestinal epithelial tight junction permeability requires NF-kappa B activation. *Am J Physiol Gastrointest Liver Physiol* 2004; **286**: G367-G376 [PMID: 14766535 DOI: 10.1152/ajpgi.00173.2003]
- 41 **Koelink PJ**, Bloemendaal FM, Li B, Westera L, Vogels EWM, van Roest M, Gloudemans AK, van 't Wout AB, Korf H, Vermeire S, Te Velde AA, Ponsioen CY, D'Haens GR, Verbeek JS, Geiger TL, Wildenberg ME, van den Brink GR. Anti-TNF therapy in IBD exerts its therapeutic effect through macrophage IL-10 signalling. *Gut* 2020; **69**: 1053-1063 [PMID: 31506328 DOI: 10.1136/gutjnl-2019-318264]
- 42 **Cross RK**, Wilson KT. Nitric oxide in inflammatory bowel disease. *Inflamm Bowel Dis* 2003; **9**: 179-189 [PMID: 12792224 DOI: 10.1097/00054725-200305000-00006]
- 43 **Larabi A**, Barnich N, Nguyen HTT. New insights into the interplay between autophagy, gut microbiota and inflammatory responses in IBD. *Autophagy* 2020; **16**: 38-51 [PMID: 31286804 DOI: 10.1080/15548627.2019.1635384]
- 44 **Becker C**, Dornhoff H, Neufert C, Fantini MC, Wirtz S, Huebner S, Nikolaev A, Lehr HA, Murphy AJ, Valenzuela DM, Yancopoulos GD, Galle PR, Karow M, Neurath MF. Cutting edge: IL-23 cross-regulates IL-12 production in T cell-dependent experimental colitis. *J Immunol* 2006; **177**: 2760-2764 [PMID: 16920909 DOI: 10.4049/jimmunol.177.5.2760]
- 45 **Iyer N**, Corr SC. Gut Microbial Metabolite-Mediated Regulation of the Intestinal Barrier in the Pathogenesis of Inflammatory Bowel Disease. *Nutrients* 2021; **13** [PMID: 34959809 DOI: 10.3390/nu13124259]
- 46 **Danese S**, Fiocchi C. Etiopathogenesis of inflammatory bowel diseases. *World J Gastroenterol* 2006; **12**: 4807-4812 [PMID: 16937461 DOI: 10.3748/wjg.v12.i30.4807]
- 47 **Schirmer M**, Garner A, Vlamakis H, Xavier RJ. Microbial genes and pathways in inflammatory bowel disease. *Nat Rev Microbiol* 2019; **17**: 497-511 [PMID: 31249397 DOI: 10.1038/s41579-019-0213-6]
- 48 **Palmela C**, Chevarin C, Xu Z, Torres J, Sevrin G, Hirten R, Barnich N, Ng SC, Colombel JF. Adherent-invasive Escherichia coli in inflammatory bowel disease. *Gut* 2018; **67**: 574-587 [PMID: 29141957 DOI: 10.1136/gutjnl-2017-314903]
- 49 **Glassner KL**, Abraham BP, Quigley EMM. The microbiome and inflammatory bowel disease. *J Allergy Clin Immunol* 2020; **145**: 16-27 [PMID: 31910984 DOI: 10.1016/j.jaci.2019.11.003]
- 50 **Everard A**, Belzer C, Geurts L, Ouwerkerk JP, Druart C, Bindels LB, Guiot Y, Derrien M, Muccioli GG, Delzenne NM, de Vos WM, Cani PD. Cross-talk between Akkermansia muciniphila and intestinal epithelium controls diet-induced obesity. *Proc Natl Acad Sci U S A* 2013; **110**: 9066-9071 [PMID: 23671105 DOI: 10.1073/pnas.1219451110]
- 51 **Vermeire S**, Joossens M, Verbeke K, Wang J, Machiels K, Sabino J, Ferrante M, Van Assche G, Rutgeerts P, Raes J. Donor Species Richness Determines Faecal Microbiota Transplantation Success in Inflammatory Bowel Disease. *J Crohns Colitis* 2016; **10**: 387-394 [PMID: 26519463 DOI: 10.1093/ecco-jcc/jjv203]
- 52 **Tan P**, Li X, Shen J, Feng Q. Fecal Microbiota Transplantation for the Treatment of Inflammatory Bowel Disease: An Update. *Front Pharmacol* 2020; **11**: 574533 [PMID: 33041818 DOI: 10.3389/fphar.2020.574533]
- 53 **Miao W**, Wu X, Wang K, Wang W, Wang Y, Li Z, Liu J, Li L, Peng L. Sodium Butyrate Promotes Reassembly of Tight Junctions in Caco-2 Monolayers Involving Inhibition of MLCK/MLC2 Pathway and Phosphorylation of PKC β 2. *Int J Mol Sci* 2016; **17** [PMID: 27735862 DOI: 10.3390/ijms17101696]
- 54 **Zheng L**, Kelly CJ, Battista KD, Schaefer R, Lanis JM, Alexeev EE, Wang RX, Onyiah JC, Kominsky DJ, Colgan SP. Microbial-Derived Butyrate Promotes Epithelial Barrier Function through IL-10 Receptor-Dependent Repression of Claudin-2. *J Immunol* 2017; **199**: 2976-2984 [PMID: 28893958 DOI: 10.4049/jimmunol.1700105]
- 55 **Duboc H**, Rajca S, Rainteau D, Benarous D, Maubert MA, Quervain E, Thomas G, Barbu V, Humbert L, Despras G, Bridonneau C, Dumetz F, Grill JP, Masliah J, Beaugerie L, Cosnes J, Chazouillères O, Poupon R, Wolf C, Mallet JM, Langella P, Trugnan G, Sokol H, Seksik P. Connecting dysbiosis, bile-acid dysmetabolism and gut inflammation in inflammatory bowel diseases. *Gut* 2013; **62**: 531-539 [PMID: 22993202 DOI: 10.1136/gutjnl-2012-302578]
- 56 **Sorrentino G**, Perino A, Yildiz E, El Alam G, Bou Sleiman M, Gioiello A, Pellicciari R, Schoonjans K. Bile Acids Signal via TGR5 to Activate Intestinal Stem Cells and Epithelial Regeneration. *Gastroenterology* 2020; **159**: 956-968.e8 [PMID: 32485177 DOI: 10.1053/j.gastro.2020.05.067]
- 57 **Li M**, Zhao J, Cao M, Liu R, Chen G, Li S, Xie Y, Xie J, Cheng Y, Huang L, Su M, Xu Y, Zheng M, Zou K, Geng L, Xu W, Gong S. Mast cells-derived MiR-223 destroys intestinal barrier function by inhibition of CLDN8 expression in intestinal epithelial cells. *Biol Res* 2020; **53**: 12 [PMID: 32209121 DOI: 10.1186/s40659-020-00279-2]
- 58 **Li DF**, Yang MF, Xu J, Xu HM, Zhu MZ, Liang YJ, Zhang Y, Tian CM, Nie YQ, Shi RY, Wang LS, Yao J. Extracellular Vesicles: The Next

- Generation Theranostic Nanomedicine for Inflammatory Bowel Disease. *Int J Nanomedicine* 2022; **17**: 3893-3911 [PMID: 36092245 DOI: 10.2147/IJN.S370784]
- 59 **Rogler G.** The effects of NOD2/CARD15 mutations on the function of the intestinal barrier. *J Crohns Colitis* 2007; **1**: 53-60 [PMID: 21172185 DOI: 10.1016/j.crohns.2007.08.008]
- 60 **Hansson GC.** Mucins and the Microbiome. *Annu Rev Biochem* 2020; **89**: 769-793 [PMID: 32243763 DOI: 10.1146/annurev-biochem-011520-105053]
- 61 **Johansson ME,** Gustafsson JK, Holmén-Larsson J, Jabbar KS, Xia L, Xu H, Ghishan FK, Carvalho FA, Gewirtz AT, Sjövall H, Hansson GC. Bacteria penetrate the normally impenetrable inner colon mucus layer in both murine colitis models and patients with ulcerative colitis. *Gut* 2014; **63**: 281-291 [PMID: 23426893 DOI: 10.1136/gutjnl-2012-303207]
- 62 **Schulzke JD,** Siegmund B, Günzel D. New insights into intestinal secretion. *Gut* 2014; **63**: 1371-1372 [PMID: 23868326 DOI: 10.1136/gutjnl-2013-305214]
- 63 **Al-Sadi R,** Dharmaparakash V, Nighot P, Guo S, Nighot M, Do T, Ma TY. Bifidobacterium bifidum Enhances the Intestinal Epithelial Tight Junction Barrier and Protects against Intestinal Inflammation by Targeting the Toll-like Receptor-2 Pathway in an NF- κ B-Independent Manner. *Int J Mol Sci* 2021; **22** [PMID: 34360835 DOI: 10.3390/ijms22158070]
- 64 **Merlin D,** Sitaraman S, Liu X, Eastburn K, Sun J, Kucharzik T, Lewis B, Madara JL. CD98-mediated links between amino acid transport and beta 1 integrin distribution in polarized columnar epithelia. *J Biol Chem* 2001; **276**: 39282-39289 [PMID: 11507094 DOI: 10.1074/jbc.M105077200]



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