

Review

Not peer-reviewed version

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Posted Date: 28 November 2023

doi: 10.20944/preprints202311.1733.v1

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Review

Omentin: An Atheroprotective Adipokine for Vascular Health

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Abstract: Omentin is an adipokine mainly produced by visceral fat tissue, and it has two isoforms: omentin-1 and omentin-2. Omentin-1 is predominantly secreted by visceral adipose tissue, deriving specially from the stromal vascular fraction cells of white adipose tissue (WAT). Levels of omentin-1 are also expressed in other WAT depots, like epicardial adipose tissue. Omentin-1 exerts several beneficial effects in glucose homeostasis in obesity and diabetes. In addition, research has suggested that omentin may have atheroprotective (protective against the development of atherosclerosis) and anti-inflammatory effects, potentially contributing to cardiovascular health. This review highlights the potential therapeutic targets of omentin-1 in metabolic-related disorders.

Keywords: omentin; adipokines; diabetes mellitus; endothelial dysfunction; inflammation; oxidative stress

1. Introduction

Adipose tissue, in addition to storing energy, secretes a range of cellular components and inflammatory mediators. Tumor necrosis factor- α (TNF α), interleukin-6 (IL-6), leptin, retinol-binding protein, resistin, adiponectin, omentin, apelin, visfatin, and other substances are secreted by different adipose tissue depots with regional heterogeneity (Galic et al., 2010). These secretions have an impact on the metabolism of carbohydrates and lipids, and they also play a significant part in pathological processes that include insulin resistance, type 2 diabetes, atherosclerosis, inflammation, and dysfunction of the vascular endothelium (Galic et al., 2010).

Omentin-1 is a glycoprotein that has emerged as a key player in the complex interplay between adipose tissue and various physiological processes. Omentin-1 is primarily produced by the stromal vascular cells of the visceral adipose tissue (Yang et al., 2006). This adipokine exist in human blood and it is highly expressed in human visceral and epicardial adipose tissues, with lower levels found in other white adipose tissue depots such as subcutaneous WAT (Fain et al., 2005). Omentin-1 is expressed in various cells including endothelial cells, mesothelial cells, vascular cells, intestinal Paneth cells among others; exerting a paracrine, autocrine, and endocrine signaling influence (Watanabe et al., 2017). Importantly and similarly to adiponectin circulating omentin levels are reduced in obese subjects. In fact, individuals with poor glucose regulation have lower serum levels of omentin-1, and this depletion may play a role in the emergence of insulin resistance, type 2 diabetes, obesity, and metabolic syndrome. Indeed, there is a negative correlation between the serum concentration of omentin-1 and the following: body mass index, insulin resistance index, leptin, plasma glucose, fasting insulin, TNF α , and IL-6 (Pan et al., 2010; Herder et al., 2015; Liu et al., 2011; Sperling et al., 2016). This adipokine is downregulated in association with obesity linked metabolic disorders including type 2 diabetes, and insulin resistance (de Souza Batista et al., 2007; Elsaid et al., 2018; Pan et al., 2010; Pan et al., 2019).

Several factors can influence the production and secretion of omentin including obesity, insulin sensitivity, inflammation, genetic factors, and hormones, such as adiponectin and insulin, fibroblast growth factor-21 and dexamethasone may also influence omentin production (Watanabe et al., 2017). Adiponectin, another adipokine, has been linked to omentin, and insulin sensitivity may play a role in omentin regulation. Understanding these factors is essential for unraveling the complex regulatory

mechanisms of omentin and its potential implications for metabolic and cardiovascular health. Ongoing research continues to explore the intricate interplay between omentin, adipose tissue, and systemic physiology.

This review highlights the potential therapeutic targets of omentin-1 in metabolic-related disorders. Relevant pre-clinical and clinical studies were summarized and discussed. Pubmed search was performed between 1990-2023 using the words: omentin-1, endothelial function, obesity, type 2 diabetes, inflammation, and oxidative stress.

2. Vascular Effects of Omentin

Omentin has been implicated in several positive effects on vascular function, making it an area of interest in cardiovascular research. Omentin exerts several positive effects on vascular function, including vasodilation, anti-inflammatory actions, and potential anti-atherosclerotic effects (Hiramatsu-Ito et al., 2016).

2.1. Vasodilation

Omentin has been associated with the promotion of vasodilation, which is the relaxation of blood vessels (Yamawaki et al., 2010). This vasodilatory effect is important for maintaining proper blood flow and reducing the resistance within the vascular system (Kazama et al., 2013). Enhanced vasodilation contributes to optimal cardiovascular function (Sena et al., 2014).

2.2. Anti-Inflammatory Actions

Omentin exhibits anti-inflammatory properties. Inflammation plays a crucial role in the development of vascular diseases, including atherosclerosis (Sena et al., 2014). By exerting anti-inflammatory actions, omentin may help mitigate the inflammatory processes within blood vessels, reducing the risk of vascular damage and atherosclerotic plaque formation (Uemura et al., 2015; Askin et al., 2020). Indeed, omentin-1 inhibits TNF- α , IL-6 and other inflammatory cytokines ultimately impacting vascular and tissue functions (Kazama et al., 2012; Fernández-Trasancos et al., 2017; Wang et al., 2019). Several studies have described the anti-inflammatory and anti-atherosclerotic effects of omentin through intracellular signaling pathways involving Mitogen-activated protein kinase (p38, JNK, ERK), nuclear factor- κ B, and AMP-activated protein kinase/Akt (Watanabe et al., 2017).

2.3. Endothelial Protection

Omentin appears to have protective effects on the endothelium, the inner lining of blood vessels. Endothelial health is essential for maintaining vascular integrity and function (Sena et al., 2014, 2017). In patients with type 2 diabetes, lower levels of omentin-1 have been linked to endothelial dysfunction (Hayashi et al., 2019). Omentin-1 has an endothelial-dependent effect on the vascular reactivity of isolated blood vessels, according to research by Yamawaki and colleagues (2010). Accordingly, we discovered that omentin-1 treatment restored endothelial dysfunction in type 2 diabetes by normalizing ACh-induced relaxation of aortic rings in diabetic Goto-Kakizaki (GK) rats fed a high-fat diet. In arteries without perivascular adipose tissue (PVAT), omentin-1 had no effect on endothelial-independent vasorelaxation. Moreover, the aortas of diabetic GK rats mounted with PVAT showed a reduction in the ET1-induced constrictor response in response to omentin-1 (Leandro et al., 2021). Notably, *ex vivo* omentin-1 vasorelaxation in aortic rings seems to be successful and mostly unrelated to increased peripheral insulin sensitivity (Yang et al., 2006).

Omentin's positive influence on endothelial cells contributes to the prevention of endothelial dysfunction, a key factor in the development of cardiovascular diseases (Dong et al. 2021; Liu et al., 2020; Maruyama et al., 2012).

2.4. Nitric Oxide Production

Omentin has been reported to stimulate the production of NO in endothelial cells. Nitric oxide is a signaling molecule with vasodilatory properties. Increased NO production helps regulate blood vessel tone, ensuring proper blood flow and reducing the risk of vascular constriction (Sena et al., 2014). Omentin-1 has been shown in earlier research to protect endothelial cells by inducing NO production and endothelial NO synthase (eNOS) activation (Yamawaki et al., 2010; Qi et al., 2016). Furthermore, omentin-1 was able to raise NO metabolites in the aortas and considerably raise the ratio of p-eNOS to total eNOS, suggesting that omentin-1's ability to restore endothelial function is caused by an increase in NO bioavailability (Leandro et al., 2021). Vascular oxidative stress was decreased, and nitric oxide (NO) bioavailability was improved by omentin-1 treatment in diabetic GK rats fed (Leandro et al., 2021).

Indeed, omentin-1 can improve endothelial function in normal mice's arteries that have endothelial dysfunction brought on by high glucose concentrations. This improvement is mediated by AMPK and PPAR δ and results in an increase in Akt/eNOS activity and NO production (Liu et al., 2020).

2.5. Anti-Atherogenic Effects

Omentin has an enormous potential against atherosclerotic initiation and progression. Atherosclerosis is a condition characterized by the buildup of fatty deposits (plaque) in the arterial walls, leading to narrowing and hardening of the arteries (Lusis, 2000).

Omentin's ability to promote vasodilation, reduce inflammation, and protect endothelial cells may contribute to its potential anti-atherogenic properties (Leandro et al., 2021; Watanabe et al., 2016). In addition, omentin-1 as anti-inflammatory, antioxidant and anti-apoptotic properties positively impacting endothelial function and preventing atherosclerosis (Gu et al., 2019; Liu et al., 2011; Watanabe et al., 2016).

Recent research has demonstrated the extensive involvement of omentin in numerous pathophysiological processes, including atherogenesis, obesity, insulin resistance, inflammatory response, and regulation of vascular endothelial function (Du et al., 2016; Ge et al., 2021; Harada et al., 2016; Liu et al., 2020; Varona et al., 2019). It improves insulin sensitivity, lowers inflammation, prevents atherosclerosis, controls the activity of vascular endothelial cells (Leandro et al., 2021; Lin et al., 2021), and protects the cardiovascular system (Greulich et al., 2013).

Omentin-1 has gained attention due to its potential significance in vascular function and its role in metabolic regulation.

3. Metabolic Regulation

Omentin has been implicated in metabolic regulation, particularly in influencing insulin sensitivity and glucose metabolism (Yang et al., 2006; Jialal et al., 2013; Koleva et al., 2013). Omentin is a protein that has been studied for its potential roles in metabolic health and regulation. It has been associated with improved insulin sensitivity, glucose and lipid metabolism.

3.1. Insulin Sensitivity

Omentin has been associated with improvements in insulin sensitivity. Insulin sensitivity refers to how effectively cells respond to insulin's signaling to uptake glucose from the bloodstream. Enhanced insulin sensitivity is generally considered beneficial for metabolic health as it helps maintain normal blood glucose levels.

Omentin-1 levels have been demonstrated to be lowered in dysmetabolic conditions, including diabetes mellitus (Eimal Latif et al., 2021; Pan et al., 2010; Pan et al., 2019), obesity (de Souza Batista et al., 2007), and impaired glucose tolerance (Cetin Sanlialp et al., 2022). On the other hand, following aerobic exercise (Saremi et al., 2010), hypocaloric weight loss (Moreno-Navarrete et al., 2010), and metformin therapy (Tan et al., 2010), omentin-1 levels are increased. Higher levels of omentin-1 are advantageous for improving insulin-stimulated glucose transport because they activate Akt signaling, which regulates downstream processes like glucose metabolism (Watanabe et al., 2017).

Like adiponectin, it increases insulin sensitivity through and increment in insulin mediated glucose uptake in adipose tissue. Omentin-1 is downregulated by glucose/insulin levels (Watanabe et al., 2017).

We have previously demonstrated that omentin-1 could lower insulin resistance in GK rats fed with high-fat diet (Leandro et al., 2021). Prior research indicates that this adipokine plays a significant role in regulating insulin sensitivity. Indeed, chronic omentin-1 infusion into ApoE^{-/-} mice may improve insulin resistance via PPAR γ , leading to a decrease in plasma glucose concentration (Yang et al., 2006). The beneficial metabolic effects may indirectly contribute to vascular health by addressing risk factors for cardiovascular diseases.

Omentin-1 has also been associated with beneficial effects against bone metabolic disorders (Xie et al., 2011; Rao et al., 2018). Understanding its role in these processes is essential for exploring its potential therapeutic applications in metabolic disorders.

3.2. Glucose Uptake

Omentin appears to influence glucose uptake in peripheral tissues, such as skeletal muscle and adipose tissue. By promoting glucose uptake, omentin may contribute to the regulation of blood glucose levels. This effect is particularly relevant in the context of insulin resistance, a condition where cells become less responsive to insulin's actions, leading to elevated blood sugar levels.

3.3. Adiponectin Interaction

Omentin and adiponectin, another adipokine, share structural and functional similarities. Omentin may interact with adiponectin receptors and both adipokines have been linked to improvements in insulin sensitivity (Brunetti et al., 2014). The specific mechanisms through which omentin and adiponectin cooperate in metabolic regulation are still an area of active research.

3.4. Anti-Inflammatory Effects

Chronic inflammation is associated with insulin resistance. Omentin's anti-inflammatory properties may contribute to improved insulin sensitivity by reducing inflammation in tissues like adipose tissue and the liver (Herder et al., 2015; Michalczyk et al., 2021; Waluga et al., 2019). Inflammation interferes with insulin signaling pathways, and by mitigating inflammation, omentin may help maintain proper insulin responsiveness. Omentin blunts cytokine expression in different cell types (Fernández-Trasancos et al., 2017; Kazama et al., 2012; Kazama et al., 2015; Wang et al., 2019; Yamawaki et al., 2011) and is negatively associated with systemic inflammatory markers such as TNF α and IL-6 (Zabetian-Targhi et al., 2016). Thus, omentin is a biomarker for metabolic health that may function to dampen obesity-related cytokine effects (Shibata et al., 2012; Zhou et al., 2020). Indeed, it is possible that omentin-1 works by blocking the NF- κ B pathway and triggering the AMPK- and Akt-dependent pathways (Kataoka et al., 2014). Anti-diabetic medications may have an impact on the level of circulating omentin-1, which is inversely linked to the incidence of type 2 diabetes and certain of its complications, such as cardiomyopathy, retinopathy, and diabetic vascular disease (Okamura et al., 2023).

3.5. Lipid Metabolism

Omentin has been suggested to influence lipid metabolism (Herder et al., 2015; Michalczyk et al., 2021). It may play a role in regulating the breakdown of fats (lipolysis) and lipid storage in adipose tissue (Herder et al., 2015; Michalczyk et al., 2021). By modulating lipid metabolism, omentin could impact insulin sensitivity and overall metabolic health.

3.6. Potential Hormonal Interactions

Omentin's effects on metabolic regulation may involve interactions with various hormones, including insulin, adiponectin, and others. These interactions contribute to the complex network of

signaling pathways that regulate glucose and lipid metabolism (Herder et al., 2015; Michalczyk et al., 2021).

While the evidence suggests a link between omentin and metabolic regulation, it's crucial to note that research in this field is ongoing, and the precise mechanisms involved are still being uncovered. The potential therapeutic applications of omentin in addressing metabolic disorders, such as insulin resistance and type 2 diabetes, warrant further exploration and investigation.

4. Clinical Implications

Herein we summarize some clinical implications or potential therapeutic applications of omentin in vascular and metabolic disorders.

4.1. Cardiovascular Diseases

Omentin's positive effects on vascular function, including vasodilation, anti-inflammatory actions, and potential anti-atherosclerotic effects, suggest potential applications in cardiovascular diseases (Biscetti et al., 2019; Biscetti et al., 2020). Therapies aimed at increasing omentin levels or enhancing its activity could be explored for conditions such as atherosclerosis, hypertension, and other cardiovascular disorders (Cetin Sanlialp et al., 2022; Fang et al., 2022; Okamura et al., 2023).

4.2. Metabolic Disorders

Omentin's involvement in metabolic regulation, insulin sensitivity, and glucose metabolism makes it a potential target for metabolic disorders. Strategies to modulate omentin levels or activity may be considered in the management of insulin resistance, type 2 diabetes (Biscetti et al., 2020), and obesity (Weng et al., 2017; Zhou et al., 2020).

4.2.1. Omentin and obesity

In obesity associated with insulin resistance higher circulating levels of retinol-binding protein 4 (Sun et al., 2013), visfatin (Jacques et al., 2012), chemerin (Weng et al., 2017), vaspin (Feng et al., 2014) and resistin (Fontana et al., 2015) and to lower levels of omentin-1 (Narumi et al., 2014) and adiponectin (Wu et al., 2014) have been reported.

However, adipokine expression levels in specific adipose tissue depots might not necessarily correlate with the adipokine levels in the circulation (Margaritis et al., 2013), suggesting the existence of complex mechanisms that regulate the biology and secretome of the adipose tissue. Importantly, in obesity, omentin-1 plays a significant anti-inflammatory role, most likely through upregulating Th-2 cytokines like IL-13 and IL-14. Increased concentrations of omentin are thought to lower the levels of inflammatory cytokines (Zabetian-Targhi et al., 2016).

4.2.2. Insulin Resistance and Type 2 Diabetes

Improving insulin sensitivity is a key goal in managing insulin resistance and type 2 diabetes. Omentin's potential to enhance insulin sensitivity suggests that it could be a therapeutic target for individuals with insulin resistance or those at risk of developing type 2 diabetes (Eimal Latif et al., 2021; Zabetian-Targhi et al., 2016; Zhou et al., 2020).

4.2.3. Obesity and Metabolic Syndrome

Omentin's role in adipose tissue and its potential influence on lipid metabolism may have implications for obesity and metabolic syndrome (Cetin Sanlialp et al., 2022; Liu et al., 2011; Varona et al., 2019; Zhang et al., 2017). Therapeutic approaches that aim to modulate omentin levels or activity could be explored in the context of obesity management and preventing metabolic syndrome-related complications.

4.3. Inflammatory Disorders

Omentin's anti-inflammatory properties make it relevant in conditions associated with chronic inflammation. This includes inflammatory disorders that can impact vascular health, such as rheumatoid arthritis (Robinson et al., 2017). Omentin-based interventions might be investigated as a complementary approach to address inflammation in these conditions.

4.4. Future Therapeutic Developments

Ongoing research may uncover novel therapeutic strategies, such as the development of omentin-based drugs or interventions that target omentin receptors. These advancements could open new avenues for personalized medicine approaches tailored to individuals with specific vascular and metabolic profiles.

Importantly, while the potential therapeutic applications of omentin are promising, further research, including clinical trials, is needed to establish its efficacy, safety, and optimal modes of administration. The field of omentin research is dynamic, and advancements in understanding its role in health and disease may lead to new therapeutic opportunities in the future.

5. Challenges and Future Directions

While omentin holds promise for its potential benefits in vascular function, there are several challenges and gaps in understanding that researchers face.

Receptor Identification and Signaling Pathways

Identifying the specific receptors through which omentin exerts its effects on vascular cells remains a challenge. Omentin receptors and the complete signaling pathways and downstream effects are not fully elucidated. A more comprehensive understanding of the molecular mechanisms involved is needed.

5.1. Tissue-Specific Effects

Omentin is expressed in various tissues, including adipose tissue, but its effects may be tissue specific. Understanding how omentin functions in different tissues and whether its effects on vascular function vary in different vascular beds is a complex aspect that requires further investigation.

5.2. Interactions with Other Adipokines

Omentin shares similarities with other adipokines, such as adiponectin or apelin. The interactions and potential synergies between omentin and other adipokines in modulating vascular function are not fully understood. Disentangling these interactions is crucial for a more comprehensive view of omentin's role.

5.3. Dose-Response Relationships

Determining optimal dosage levels for potential therapeutic interventions is challenging. Omentin's effects may vary based on concentration and understanding the dose-response relationships is essential for designing effective treatments without adverse effects.

5.4. Role in Disease Progression

The context-dependent role of omentin in various disease states is not fully understood. For example, while it may have anti-atherosclerotic effects, its role in advanced stages of vascular diseases or in conditions with chronic inflammation needs further exploration.

5.5. Biomarker Validity

Omentin has been proposed as a potential biomarker for certain metabolic and cardiovascular conditions. However, the validity, specificity, and sensitivity of omentin as a biomarker need to be thoroughly validated in diverse populations and clinical settings.

Initiate longitudinal studies to investigate the association between circulating omentin levels and the development of vascular and metabolic disorders over time. This could provide insights into the predictive value of omentin as a biomarker.

5.6. Limited Clinical Data

While preclinical studies suggest beneficial effects, translating these findings to clinical applications poses challenges. Establishing the safety, efficacy, and feasibility of interventions targeting omentin in humans requires well-designed clinical trials. Conduct well-designed clinical trials to assess the safety and efficacy of interventions targeting omentin in humans. Investigate the potential therapeutic applications of omentin in cardiovascular diseases, metabolic disorders, and other related conditions.

The number of clinical studies investigating omentin's role in vascular function is relatively limited compared to preclinical research. Expanding the clinical evidence base is critical for understanding its relevance in human health and disease.

Addressing these challenges will contribute to a more comprehensive understanding of omentin's role in vascular function and facilitate the development of targeted therapeutic interventions. As research progresses, these challenges are likely to be addressed, leading to a clearer picture of omentin's potential in clinical applications.

Future research on omentin could explore several key areas to deepen our understanding of its role in health and disease.

5.7. Mechanistic Insights

Elucidate the precise molecular mechanisms through which omentin exerts its effects on vascular cells. Identify the specific receptors and downstream signaling pathways involved in omentin-mediated vasodilation, anti-inflammatory actions, and other vascular effects.

5.8. Role in Inflammation

Further investigate omentin's role in modulating inflammation, both locally within adipose tissue and systemically. Explore how omentin's anti-inflammatory actions may impact the progression of inflammatory diseases, including those affecting the vasculature.

5.9. Genetic Variations

Explore the impact of genetic variations in the omentin gene on omentin expression and function (Rathwa et al., 2019). Investigate whether specific genetic polymorphisms are associated with altered susceptibility to vascular and metabolic disorders.

5.10. Omentin as a Therapeutic Target

Evaluate the feasibility and efficacy of developing therapeutic interventions that directly target omentin or its receptors. Investigate the potential use of omentin-based drugs or interventions to improve vascular function and metabolic health.

5.11. Sex Differences

Explore potential sex differences in omentin levels and effects on vascular function. Investigate whether omentin's role varies between males and females, which could have implications for personalized medicine approaches.

As research in the field of omentin continues, addressing these areas of exploration will contribute to a more comprehensive understanding of its physiological functions and potential clinical applications.

6. Conclusion

A schematic diagram summarizing the key features of omentin is presented.



Figure 1. Schematic representation of the beneficial effects of omentin-1.

- Omentin-1 is an adipocytokine widely expressed in a variety of cells, exhibiting microbial defense, antioxidative, anti-inflammatory, and anti-apoptotic properties.
- Omentin-1 exhibits a wide range of therapeutic potential in diabetes by reducing comorbidities linked to type 2 diabetes mellitus, such as vascular diseases and diabetic nephropathy (Senthilkumar et al., 2018; Song et al., 2018). Noteworthy, of the major adipocytokines that inhibit atherosclerosis, omentin-1 has a strong correlation with inflammation, macrophage differentiation, arterial calcification, and plaque formation (Xu et al., 2019).
- Omentin-1 inhibits insulin resistance, atherosclerosis, and inflammation through the intracellular signaling pathways of AMP-activated protein kinase/Akt/nuclear factor- κ B/mitogen-activated protein kinase (ERK, p38, and JNK).
- Omentin-1 may be used as a biomarker for metabolic syndrome, obesity, diabetes, atherosclerosis, ischemic heart disease, and inflammatory diseases.
- This review sheds light on how omentin-1 might be used to treat these diseases and serve as a biomarker.

Funding: This work was supported by the Fundação para a Ciência e Tecnologia, Portugal: Reference number: 2022.04526.PTDC. FCT 2023.

Institutional Review Board Statement:

Informed Consent Statement:

Data Availability Statement:

Conflict of Interest: The author declares no conflicts of interest.

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