Review Article
Role of matrix metalloproteinase in wound healing

Mimansa Kandhwal1*, Tapan Behl1*, Sukhbir Singh1, Neelam Sharma1, Sandeep Arora1, Saurabh Bhatia2,3, Ahmed Al-Harrasi2, Monika Sachdeva4, Simona Bungau5

1Chitkara College of Pharmacy, Chitkara University, Punjab 140401, India; 2Natural & Medical Sciences Research Centre, University of Nizwa, Nizwa 342001, Oman; 3School of Health Science, University of Petroleum and Energy Studies, Dehradun 248007, Uttarakhand, India; 4Fatima College of Health Sciences, Al Ain 50, United Arab Emirates; 5Department of Pharmacy, Faculty of Medicine and Pharmacy, University of Oradea, Oradea 410028, Romania. *Equal contributors.

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Abstract: Matrix metalloproteinases (MMPs) are a group of endopeptidases that play a vital role in the restoration of damaged skin. Through mediating various cellular events such as angiogenesis and vasodilation, MMPs are very crucial for the mechanism of wound healing. These enzymes are endopeptidases that are reliant on zinc which are concealed through the extracellular matrix (ECM). MMPs have different targets in different phases of wound healing through which they are capable of promoting timely healing in the body. This review discusses all the possible role of MMPs and their inhibitors that are involved during every step of the wound healing process. This review highlights the latest advances in the respective field about the regulation and mediation of MMPs in human skin and how these studies can be applied to other branches of medical sciences as well. Published papers were searched via MEDLINE, PubMed and MDPI from the available peer reviewed journals. Research done in the past suggests that active MMPs are involved in the healing progression of the wounds or they have a positive effect towards healing of wounds. Present studies in the relative field will further enhance the knowledge about enzymes working along with their inhibitors. These studies will help in a way to resolve some of the parameters that are necessary for modulating them either positively or negatively.

Keywords: MMPs, chronic wounds, wound healing, TIMPs, inflammation, chemokines, extracellular matrix

Introduction

Matrix metalloproteins, are endopeptidases that help in regeneration of skin, and these are calcium dependent zinc containing endopeptidases also known as metallopeptidases or matrixins. Skin, the largest organ of human body is responsible for protecting the body from outer infection or damage, apart from that it is also responsible for sustaining the fluid balance and thermo regulation [1]. Primary functions of this protective organ include regulation of the body temperature, which permits sensations like heat, cold and touch and importantly it provides protection from the outer environment. Each layer of skin has its own distinctive feature. The layers have their own separate roles and functions to perform [2-4]. It provides a water proof barrier, sweat glands to maintain body temperature and cushioning effect to the skeleton. The dermis layer is responsible for the tensile potency of the skin along with the ECM [5]. This cushioning effect is more pronounced in the presence of the ECM. It is an absolute 3D structure made up of connective tissue and fat cells including collagen, proteoglycans, and glycoproteins (almost 300 different types of protein) [6]. Proper functioning of the skin is very important for the body as it is the first line defensive organ of the human body. Any damage to any of the layer of human skin may alter the proper functioning of the skin which may result in instability. Breakdown in the protective or defensive function of the skin may be considered as a wound which has to be healed in a well-planned manner. The skin itself is directly involved in the healing of cutaneous wounds while it also plays an indirect role in the healing of internal or chronic wounds as well. The integrity of skin is supremely important while main-
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taining physiological hemostatis and feasibility of the internal tissues as well. Any damage to the epithelial cells which are present in the epidermal layer of the skin may infringe upon the integrity of the skin [7-10]. This may lead to bacterial infiltration into the skin as the mechanical strength of the skin is compromised. These bacterial infringements through the skin from the external environment may reach up to the dermis layer and the ECM and cause further damage. In order to protect the affected part from further infection the wound has to be healed efficiently. Categorically, wounds can be differentiated into various types that are: Superficial wounds (loss of epidermis layer only), partial wound (loss of epidermis and dermis layer) and full thickness wounds which involve the loss of the dermis subcutaneo- us fat and sometimes even bone. Apart from these categories, wounds are to be differentiated on the basis of the time taken to heal properly and there are two sub categories which are acute wounds and chronic wounds [11]. Wounds that heal in a synchronized manner of the healing cascade are generally known as acute wounds. While on the other hand non-compliance of some wounds to the healing pattern may lead to delay in their healing. This may occur because of the prolonged residency of one or the other healing phases [12, 13]. For normal functioning of the skin, the damage has to be repaired immediately in order to prevent the further impairment due to infections. Superficial wounds do not possess any threatening damage, whereas partial and full thickness wounds may involve the loss of dermal layer and subcutaneous layer as well. The normal healing process entails a systematic and organized sequence of four dissimilar yet overlapping stages that are hemostatis, inflammatory phase, proliferation phase and remodeling phase [14, 15]. However, interruption in any phase of this dynamic process may eventually lead to the onset of chronic skin ulcers. A functional co-ordination among all the four partially overlapping phases is mandatory in order to commence the progression of tissue restoration. For the healthy progression of wound healing the ECM has to be remodeled efficiently as it is responsible for the tensile strength of the skin and in this process endopeptidases such as matrix metalloproteinases (MMPs) are thoroughly involved. The prominence of this review article is to emphasize the role of MMPs that are systematically involved the wound healing process. At present there are 24 known MMPs that are likely to be present in the human body [16]. MMPs are generally endopeptidases that are competent at eliminating all the damaged components of ECM. The ECM generally promotes all the cellular responses in the process of wound healing including cell adhesion, cell migration and tissue remodeling. Removal of all the damaged proteins and cells within the ECM is the main line of action of the MMPs. Their engagement is seen in all the four phases of the wound healing. During the inflammatory phase they are capable of removing the damaged proteins and forming a temporary ECM [17-21]. While during the second phase that is the proliferation phase, they help in degrading the capillary basement membrane. MMPs also promote angiogenesis during the wound healing cascade. The respective role of MMPs in wound healing can be studied by carefully evaluating all the events that participate in this sequential process. First, the initial phase (hemostosis) is tissue injury which involves epidermal layer, platelets and TGF-β. Immediately after hemostasis the inflammatory phase begins and this involves neutrophils, macrophages, ROS, TNF, VEGF, and PDGF. Granulation and angiogenesis are also two of the most crucial events in the wound healing process [22]. Involvement of fibroblasts, macrophages, endothelial cells are there in these events. Keratinocytes and collagen fiber crosslinking are typically involved in the process of re-epithelialization and tissue remodeling. All of the stages must be completed in a timely way and in a sequential manner in order for the wound to be fully healed [23]. Here, MMPs play a critical role in each and every phase of wound healing. This review discusses the role of MMPs and their inhibitors in different phases of wound healing. This review discusses the entire possible role of MMPs and their inhibitors that are involved during every step of the wound healing process. This review highlights the latest advances in the respective field about the regulation and mediation of MMPs in the human skin and how these studies can be applied to other branches of medical science as well.

Wound healing cascade

Wound healing undergoes mainly three phases namely the inflammatory phase, the prolifera-
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tive phase and maturation phase. The inflammatory phase forms clots and is followed by inflammation cell build up in the wound or injury site. This phase lasts up to 2-5 days and includes vasoconstriction, platelet aggregation and clot formation followed by vasodilation and phagocytosis. The second phase lasts for about 2-3 weeks which consists of granulation, contraction and epithelialization. Thereafter comes the maturation phase in which new tissue is formed and this phase last for approximately 3 weeks. The superficial wound quickly crosses the threshold of the primary stage of the healing process, which is hemostasis, upon injury to the skin. Excessive bleeding is protected by the onset of clotting in the blood vessels that is further accomplished by accumulation of platelets around the ruptured endothelium region resulting in the formation of the plug. A blood clot is formed upon further cascade of events. The serine proteinase thrombin cleaves fibrinogen into fibrin threads, which bind with platelets to create a clot. Not only does the development of a blood clot halt bleeding, but it also functions as a temporary matrix for cell migration [24, 25]. The cells surrounding the clot release various inflammatory cytokines and growth factors which further signal attraction of various cells such as neutrophils, lymphocytes, monocytes and macrophages and initiate the phase of inflammation. Neutrophils arrive at the site of injury only within a few hours. They are highly responsible for the release of fibronectin which has multiple roles [25, 26]. Fibronectin and fibrin create a provisional matrix which initiates migration of cells and helps them stick together. Their adhesion power is totally dependent on the intensity of the injury that has occurred [23]. Growth factors cause macrophage activation, which results in the production of reactive oxygen species (ROS), multiple growth factors such as platelet-derived growth factor (PDGF), vascular endothelial growth factor (VEGF), MMPs, transforming growth factor-beta (TGF-beta), and fibroblast growth factor (FGF). Angiotensin II is an important factor that can induce the stimulation of macrophages. It is mainly regulated upon reaction of the renin-angiotensin system (RAS) pathway. It generally co exists in the fibroblasts, macrophages and endothelial cells present in human skin. Secretion of MMPs and ROS is usually initiated by the activation of AngII along with the trigger of inflammatory cells. Stimulation of these factors initiates the further intensification of keratinocytes. They are also responsible for their migration through the injured cells as well. TGF-βs plays crucial part in regulation and development of the ECM. Humans have 3 isoforms (TGF-β1-3), each presenting distinct function in regulation of components of the ECM, and proliferation of cells and even cell death [27-29]. The most well-known is TGF-1, which regulates the generation and destruction of numerous elements involved in the healing process of wounds [25]. Upon binding of TGF-1 to its corresponding receptor, synthesis and manufacturing of ECM components such as collagen, fibronectin, and hyaluronic acid are triggered in several cell types, including fibroblasts [30]. Collagen is mainly synthesized by fibroblastic cells and other constituents which are accumulated within the ECM [16]. Fibroblasts are also accountable for the stimulation of ROS which further secrete defensive factors of the body against any kind of microbial attack in the body that are peroxide ions and superoxide ions [31]. The synthesized ROS generate various cytokines which further increase the proteinase production and alter the fragments of the ECM [31]. The main characteristic of ROS is that it has twofold functions that are associated with it. The antimicrobial function is of great advantage to the human body as it protects the body from foreign particles. However, excessive levels of ROS may damage the constituents of ECM [31, 32]. This moderate equilibrium between the secretions of ROS may activate the intricate pathways in the body which activates the MMPs secreted in the wounded area. The elevated levels of ROS could harm the tissues and in turn may alter the wound healing process [33-35]. Dealings that are involved in the next event of the cascade are of great importance and the events are angiogenesis and granulation. In this phase the granulated tissue is supported by the blood supply. Vascular support is increased in the affected tissue and the site surrounding it as well. This process is a reciprocating way that the human body has to deal with excessive blood loss which occurs during wounds. The vascular supply also allows the continuous supply of oxygen to the wounded area. Granulation tissue is generally soft pink to red in appearance which marks the presence continuous vascular supply to the wounds [31]. Around the wounded area the presence of
fibroblasts markedly start increasing through the process of migration [36]. Proliferation of the fibroblasts is also a distinctive feature of this phase in the wound healing as it interacts with the growth factors and the ECM which initiates the process of fibroblasts proliferation. Integrins also play a vital role in the mediation of the whole proliferation process as they are the respective receptors for the fibroblasts. They are also responsible for the biochemical signaling of the earlier mentioned process. The extracellular sphere of influence of integrins bind with the ECM portion and the internal surface of the receptor and are accountable for several signaling procedures [37]. Exposure of the integrins and other fibroblast receptors like discoidin to collagen is responsible for the processing of MMP 2 in the wounded tissue. MMP2 allows the migration of the fibroblast during angiogenesis and ECM remodeling. The interaction of integrins with collagen is responsible for the strength of the ECM and in turn encourages the secretion of supplementary growth factors. In a mouse model, Rossiter et al. confirmed that the absence of keratinocyte-specific VEGF is responsible for aberrant angiogenesis and delayed wound healing [38]. Final support given by the granulation tissue to the epithelial cells is the re-epithelialization phase of the wound healing. Keratinocytes are generally involved during this phase and they are the main mediators that can positively influence the healing phase. Likewise to the other cells in the earlier phases of wound healing keratinocytes also undergo the same process of migration, differentiation and proliferation. Migration of keratinocytes occurs towards the centre of the wound in order to close the wounded site completely. In order to do that keratinocytes must not lose their grip with the adjacent tissue. Keratinocytes also have a distinct feature of changing their shape and size before migration in order to accelerate the process of wound closure [39-41]. Here again integrins are involved as keratinocytes and move towards the newly molded ECM. During this process MMPs are continuously activated through macrophages and keratinocytes in order to degrade the matrix components to achieve healthy healing of the tissue [42-44]. After all the complexity of the events, the wounded tissue enters the last phase which is tissue remodeling [31]. In this phase type 3 collagen is replaced by type 1 collagen in order to give more tensile strength to the tissues. Different phases of wound healing are explained in Figure 1.
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MMPs and its inhibitors

There are currently 28 known MMPs out of which 24 are human enzymes. MMPs generally exist in three different categories that are inactive, active and complex MMPs. These enzymes are endopeptidases which are reliant on zinc and are concealed through the ECM. The primary structure of MMPs comprises a single arrangement of the N-terminus, a pro-domain which is capable of capping the active binding site and a catalytic sphere of influence [45, 46]. The activities of MMPs are synchronized by a group of endogenous tissue inhibitors of metalloproteinase (TIMPs). These inhibitors are further divided into 4 subtypes (TIMPS 1, 2, 3, 4). The inhibitors bind specifically to block the functioning of MMPs. MMP 7 and MMP 26 are also classified as matrix-lysins that exhibit the very basic structure comprising the minimal domain which is discussed above. The catalytic domain is exemplified by the zinc-binding HExxHxxGxxH motif, consisting of three conserved histidines [47, 48]. Numerous MMPs have an additional domain known as the hemopexin-like domain, which is located at the C-terminus of the previously stated basic arrangement and functions in conjunction with it. According to the current hypothesis, the hemopexin-like domain has a role in substrate identification. These domains are shown to be associated with MMP-3 and MMP-10 (also known as stromelysin-1 and-2), MMP-1, 8, and -13 (also known as collagenases), MMP-12 (metalloelastase), MMP-20 (enamelysin), and MMP-22 and -27 (also known as stromelysin) [49].

MMPs and wound healing

Role in inflammation

The inflammatory phase is generally associated with incursion and commencement of the leukocytes. There are several factors which are involved directly or indirectly in order to manipulate this inflammatory stage of wound healing. Many of those factors include chemokines, cytokines, ECM splinters and lipid mediators. Antimicrobial peptides are also among one of those influencing factors as well. Metalloproteinases are a group of proteins which are held responsible for the secretion and pursuit of these factors. Studies from different labs revealed that MMPs that originated or developed from epithelial cells normalize various events in the inflammatory stage which is also the initial stage of the wound healing. The events which are to be regulated through MMPs in this phase are transepithelial migration of leukocytes and partition of signaling proteins such as chemokines. It has been validated through the work done by Parks et al. that MMPs are prominently expressed in the inflammatory cells [50]. While on the other hand stromal and epithelial cells at the wounded site have also corroborated to express multiple MMPs including MMP 1, 2, 3, 7, 9, 10 and 28. All these respective MMPs could possibly mediate the activity of chemokines. This regulation could occur through distressing the production of the chemokine slope or via direct proteolysis [51-54].

These signaling proteins or chemokines are subcategorized into small families like CC chemokines and CXC chemokines. This subdivision occurs on the basis of their discrete function of magnetizing the leukocytes resulting in their influx. Apart from this their N-terminal cysteine residues are also a major factor which is taken into consideration while dividing these subfamilies [55, 56]. The first family which is the CC chemokines plays a considerable role in the chemotaxis of monocytes. However CXC chemokines are generally held responsible for the regulation of neutrophil chemotaxis. If the other dividing factor is also taken into consideration while comparing these two closely related chemokine subfamily varies. The respective function or activity of these signaling proteins is downgraded by MMPs by their division or severance. CC chemokines are responsible for downstream signaling through the formation of receptor antagonist in the presence of MMPs. While on the other hand CXC chemokines variably respond to the MMPs. It has been noted that some of the CXC chemokines are totally impervious to the presence of MMPs while some others are promptly affected by MMP presence. This dissimilarity elaborates on a major operative variation of humans and mice [57, 58]. MMP 8 and 9 are accountable for the processing of Human CXCL8. This process is responsible for the significant amplification in the activity of CXCL8. Functional responses of MMP such as modulations in the migration of inflammatory cells at
the wound site are totally different in humans and mice. This discrepancy is the core reason needed research in this field. There is another CXC chemokine which is found to be handled or processed by several MMPs, which is CXCL5 (LIX). The MMPs which are thoroughly involved in the operation of this chemokine are 1, 2, 8, 9 and 13. These are validated by the studies of Tester et al. as well. This phase of chemokine processing is ultimately responsible for the enhancement of more inflammatory cells to be involved in the wound healing process. Furthermore the processing of CXCL12 and several MMPs such as 1, 2, 3, 9, 13 and 14 are thoroughly involved in the process [58, 59]. Results of CXCL 12 processing via MMPs are totally different than the previous CXCL5 processing which consequentially alleviates the efficiency of chemokines. Out of all the known MMPs, MMP1, 3 and 9 are validated to possess extensive capability of supervising the signaling of chemokines. MMPs are generally involved in absolute chemokine breakdown and generate antagonist for chemokine receptors in order to enhance the performance of chemokines. In addition to MMPs, ADAM 10 is capable of sorting chemokines. ADAM10 is capable of cleaving CXCL16 off the cell surface, therefore allowing it to bind to its particular receptor and influence T-cell initiation at the injured region [60]. This shows that both the metalloproteinases that are MMPs and ADAMs, are thoroughly involved in the mediation of chemokines. Positive and negative regulations are carried out by these metalloproteinases. MMP expression and regulation in the wound healing cascade is diagrammatically represented in Figure 2.

Role in epithelial repair

Re-epithelialization is one of the most important and necessary phases of the sequential process of wound healing. Progression of building healthy tissue all over again at the denuded area is generally categorized as re-epithelializa-
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In this tricky process the wounded cells have to lose their grip on the ECM fragments so that they can freely move across the ECM in order to heal completely. Involvement of numerous MMPs can be seen in this phase of wound healing such as 1, 3, 7, 9, 10, 14 and 28 [61-63]. MMP1 which is also known as collagenase 1 is expressed at the wounded site during this particular phase of wound healing but its activity is valid till the closure of the wound after that it automatically gets turned off [64, 65]. Keratinocytes usually are transferred from the basal lamina at the wounded stage and then they bump into a matrix present in the dermal layer of the skin which is highly rich in type 1 collagen. MMP expression was initiated via ligation of keratinocytes to this particular ECM component with the help of α2β1 integrins. Findings from the labs of Gill et al. suggested that Keratinocyte migration was assisted by MMP1 above the dermal layer matrix. This procedure was carried out through alleviating the effectiveness of collagen and integrin fragments [66-68]. Reduced contact between the two results in the migration of keratinocyte. MMP 10 (stromelysin-2) is confined with MMP1 however MMP 3 (stromelysin-10) is restricted only to the cells following the migrating overlook. The non-overlapping location of MMP3 and 10, two very similar proteinases, suggests that these two MMPs have separate roles in re-epithelialization. In addition to all of these favorable functions, MMPs participate in the development of airway epithelial cells [69, 70]. There is certain evidence from nude mice which uses human tracheal xenografts to confirm the presence of MMP 7 and MMP 9. Their expression and activity were found maximum during the later phases of the wound healing progression. Specific inhibition of these respective MMPs contributes to the impairment in the differentiation of the epithelial cells. This evidence collectively proves that MMPs are necessary in wound healing for proper differentiation of epithelial cells [71]. MMP 10 is also produced by the epithelial layer during the cell migration process revealing that these MMPs are mandatory during cell migration. Tampa et al. corroborated that MMP 10 enhances the process of cell migration. This phase that is the repairing phase of wound healing which includes various cellular responses such as migration, proliferation and differentiation and even cell death [72, 73]. Now, it is quite evident that these events or responses either separately or collectively are able to generate MMPs due to the line of action of these proteinases that have been deciphered up until now. All this evidence corroborates the idea that practically all the MMPs have been confirmed as positive influencers of the wound healing process apart from MMP 2 and MMP 9 that may have the negative or inhibitory effects on the cell proliferation [74].

Role in wound contraction

Specifically, MMP 3 is required for the last step of the wound healing process, which is the resolution phase. It is very evident from the studies of Bullard et al. that the contraction of wound was slowed down in mice lacking MMP3 [75-77]. Slowdown in the pace of contraction of wound healing ultimately increases the size of the wound, which may slow down the whole process of wound healing. In this situation the epithelial cells might also take longer to migrate during the migration phase, which is also a negative sign in the healing process. There is one other role of MMPs which was originally thought of as its main function which is the elimination of the ECM fragments. ECM degradation is a very vital and necessary step in the process of healthy wound healing. Currently, it has been shown that the primary function of most of the MMPs is to activate several cellular factors such as growth factors, cytokines and chemokines. They are certainly capable of activating their respective receptors as well. Reframing of the collagen generally consists of the mortification of the present collagen fibers in order to build the new ones. This is the most important event in the resolution phase of the wound healing. Because MMPs are capable of this process, it seems plausible that they would have a key role in collagen remodeling during wound resolution. Different types of mammalian MMPs and their possible role in wound healing are mentioned in Table 1.

MMP inhibitors during the wound healing cascade

Since the activity of metalloproteinase in wound healing is controlled by endogenous tissue inhibitors of metalloproteinases, it is important to understand how they work. These inhibitors have the ability to attach to particular locations in the body’s immune system, resulting in
## Table 1. Mammalian MMPs and their respective roles in wound healing cascade and several other diseases

<table>
<thead>
<tr>
<th>MMPs (Synonym)</th>
<th>Possible Role in Wound Healing</th>
<th>Favorable Substrates</th>
<th>Possible Role in Other Diseases</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>MMP 1 (Collagenase-1)</td>
<td>It elevates the degree of migration of the keratinocytes resting on fibrillar collagen. Its level is drastically increased in diabetic patients also suffering from diabetic foot ulcer. Excessive production in the keratinocytes may delay the repair process.</td>
<td>Collagen I, II, III, VII and X</td>
<td>Release growth factors and have negative impact in cancer</td>
<td>[68, 78]</td>
</tr>
<tr>
<td>MMP-8 (Collagenase-2)</td>
<td>Produced by neutrophils. Positively stimulates the cutaneous healing. Increased levels can be seen in diabetic patient (Diabetic foot ulcers).</td>
<td>Collagen I, II, III</td>
<td>Act as biological markers in respiratory disorders such as asthma and chronic obstructive pulmonary disease</td>
<td>[79, 80]</td>
</tr>
<tr>
<td>MMP-13 (Collagenase-3)</td>
<td>Initiates Re-epithelialization through contracting the wound.</td>
<td>Collagen I, II, III, IV, IX, X</td>
<td>Involved in the pathogenesis of osteoarthritis</td>
<td>[81, 82]</td>
</tr>
<tr>
<td>MMP-2 (Gelatinase A)</td>
<td>Hasten the process of Wound healing. Stimulate the activation of MMP 9. Keratinocyte immigration or relocation.</td>
<td>Gelatin; collagen I, IV, V, VII, and X</td>
<td>Their levels significantly rise in colorectal cancer tissues</td>
<td>[83, 84]</td>
</tr>
<tr>
<td>MMP-9 (Gelatinase B)</td>
<td>Advances Cell Migration in whole body except Cornea.</td>
<td>Elastin, Fibrillin, Collagen I, III, IV, V</td>
<td>Regulates pathological remodeling processes that involve inflammation and fibrosis in cardiovascular disease</td>
<td>[85, 86]</td>
</tr>
<tr>
<td>MMP-3 (Stromelysin 1)</td>
<td>Stimulates the activation of MMP-9 and positively influence the wound retrenchment and impaired healing.</td>
<td>Collagen IV, V, IX, X; fibronectin; Elastin; gelatin</td>
<td>Involved in the pathology of rheumatoid arthritis (RA) and ankylosing spondylitis (AS)</td>
<td>[74, 87]</td>
</tr>
<tr>
<td>MMP-10 (Stromelysin 2)</td>
<td>Expressed at Keratinocytes at the exposed side of the wound.</td>
<td>Collagen IV, V, IX, X; fibronectin; Elastin</td>
<td>Involved in skeletal development</td>
<td>[88, 89]</td>
</tr>
<tr>
<td>MMP-7 (Matrilysin)</td>
<td>Obligatory role in re epithelialization process.</td>
<td>Elastin; fibronectin; laminin; nidogen; collagen IV; Tumor necrosis factor</td>
<td>Their levels significantly rise in patients suffering from carotid atherosclerosis</td>
<td>[90, 91]</td>
</tr>
<tr>
<td>MMP-12 (Metalloelastase)</td>
<td>Promising monitoring of angiogenesis due its capability of producing angiotatin</td>
<td>Collagen IV; gelatin; fibronectin; laminin</td>
<td>It degrades extracellular matrix elastin and enables infiltration of immune cells responsible for inflammation and granuloma formation</td>
<td>[92, 93]</td>
</tr>
</tbody>
</table>
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inflammation suppression. The variables that are involved in the process of inflammation include several cytokines and chemokines, with TNF being the most prominent. TNF is a cytokine that induces acute inflammation and is activated by ADAM 17 or TACE (TNF converting enzyme). TNF, in a manner similar to that of monocytes, increases the production of MMP9 by activating the transcription factors NFkB and NFxE [94]. MAP kinase pathways in inflammatory cells further regulates or alters many aspects of wound healing. Another inhibitor of MMP in wound healing is TIMP3 and in its absence ADAM17 is enhanced which leads to constitutively increase TNFα release [95-97]. This release further triggers the inflammatory cell infiltration into the liver. Also increased levels of TNFα are responsible for increased IL6 (interleukin 6) release. This behavior causes increased susceptibility to LPS (lipopolysaccharide) induced mortality [98]. TIMP 3 is the sole inhibitor that is capable of functioning during an inflammatory response. During a lung infection or damage, the release of TIMP1 is enhanced [98]. It has been shown that the loss of TIMP1 results in an increase in vascular permeability and neutrophil diapedesis into pulmonary tissue. TIMP1 is also crucial for limiting leukocyte extravasation and vascular permeability via the mechanism of endothelial cell apoptosis, which is a process that occurs in the absence of TIMP1 [97]. TIMP1, on the other hand, has an anti-apoptosis impact on cytokine-simulated endothelial cells via the phosphatidylinositol-3-kinase pathway, which is a novel discovery (Figure 3).

Figure 3. Pictorial illustration of the dermis layer and the role of MMPs and their respective inhibitors in wound healing.

TIMP 3, which functions as a metalloproteinase inhibitor, is required for the regulation of the inflammatory response by modulating cytokine signaling and inflammatory cell adhesion, among other things. On the basis of these findings, it may be inferred that MMP inhibitors play an extremely important role in all areas of wound healing [73, 97]. Apart from that, the TIMP3 and TIMP1 proteins are also capable of controlling cell migration by inhibiting the activity of particular MMPs. The specific function of TIMPs in the control of cell migration has not yet been determined, but some research has shown that TIMP2 increases the migration of keratinocytes into cell culture when in vivo

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Techniques are used. According to the findings of recent research, TIMPs are capable of performing MMP-independent tasks in addition to their capacity to block MMP activity. As evidenced, TIMP3 is a potent inhibitor of the angiogenesis process in wound healing. It also has the capability to directly bind to the vascular endothelial growth factor receptor and prevent the direct binding of VEGF [73]. The angiogenesis phase is of great importance during the cascade of healing wounds as it aids the formation of granulation tissue in order to enhance the vascular supply in and around the affected area. This crucial phase is regulated by TIMP3 and this particular inhibitor of MMPs can dramatically impact the wound healing process. After all the granulation and angiogenic phase, the final stage of the wound healing is the closure of the wound and the contraction of the wound is an important event towards the success of this closure. In the process of re-epithelialization, cutaneous wounds are treated with GM 6001 but it also responsible for the decrease in the pace of wound contraction as well [24, 99, 100]. One more adverse effect of this treatment is the inability to differentiate between the granulation tissue and healthy tissue which leads to impairment in the wound restoration phase. On the other hand the role of the ECM in the wound resolution phase is an important aspect which can be carefully examined and monitored. Collagen is the key requirement for converting the granulation tissue into scar tissue and this process is thoroughly regulated by TIMP3. Fibronectin along with collagen are inhibited and their release is further decreased in the absence of TIMP3.

Conclusion

It has been found that MMPs exist in basically in three distinct forms, which are pro MMPs, active MMPs and TIMPs (complex) MMPs. According to research, it has been confirmed and verified by different scientists that active MMPs are involved in the healing progression of wounds or they have a positive effect towards the healing of wounds. At present there are no such methods or mechanisms which can evenly classify between these three forms of MMPs which confirms that the activity of MMPs in wound healing is still challengeable. However quantitative profiling of a few active MMPs is crucial in examining every possible role of MMPs during wound healing progression, specifically during the ECM reframing phase. Medical research is a tool through which new therapeutic involvement for specific disorders can be identified and examined through various phases. In the case of wound healing, all the respective enzymes and receptors must be carefully examined and studied before introducing any kind of novel drug or molecule. This same principal should be considered in the case of MMPs and wound healing as well. Future experimental work that may include animal models to study MMPs, ADAMs or TIMPs, must be done in order to further explore the field. These studies will further enhance the knowledge on how exactly these enzymes work. Studies of their respective inhibitors are also equally important. These studies will help in a way to resolve all the parameters that are necessary for modulating them either positively or negatively.

Future perspectives

In recent years it has been verified by scientists that MMPs possess several beneficial properties which are the underlying principles of pathologies of different diseases. They have a characteristic role in the degradation of the ECM which ultimately releases numerous biological factors that further act as activators in the wound healing process. They are generally involved in the activation of various growth factors and also in detachment of the membrane. These are the predominant functions of MMPs, and substrates of MMPs are basically non-matrix molecules. Currently many novel techniques are being used in order to study the substrates of MMPs in detail. This could be the ground of future research of meta-analysis which would be able to differentiate downstream effects and profile substrates precise to each protease. For studying or analyzing the enhanced role of MMPs in different physiologies and pathologies, efficient analysis of protease substrate is very important. Almost all the variants of the MMP family are actively involved in several biological events in the body. However, few of them are restricted or specified to certain cells only like MMP 20 (specified for dental tissue). Their constitutive physiological expression is normally low, with transiently higher rates due to homeostatic matrix remodeling or specific developmental
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events. These short expression peaks underline their stringent regulation under physiological conditions and highlight their important role in tissue homeostasis and development. Uneven regulation of the functions of MMP specifically in the pre-chemo phase of cancer is a matter of conflict for MMPs as promising therapeutic targets. Although the first clinical trial failed in which broad spectrum MMP inhibitors were involved. The main ground on which future research will rely is the blueprint or draft of the desired therapeutic agents in inhibition of selective MMP variants. Management of wound healing is still a very challenging issue which has to be supervised with more efficacy and persistence in order to prevent acute wounds from becoming chronic. Thorough experimentation and study in this relative field is very important so that this can be addressed more efficiently. Careful examination of MMPs, their regulation and specific roles with respect to the ECM and its components may provide a new basis for further research in this respective field of wound healing. Research is still underway for framing innovative methods regarding MMPs in treating wounds. This promising field is capable of recognizing novel inhibitors to deal with this problem.

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Disclosure of conflict of interest

None.

Address correspondence to: Dr. Tapan Behl, Chitkara College of Pharmacy, Chitkara University, Punjab 140401, India. Tel: +91-8527517931; E-mail: tapanbehl31@gmail.com

References


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[88] Vaalamo M, Karjalainen-Lindsberg ML, Puolakkainen P, Kere J and Saarialho-Kere U. Distinct expression profiles of stromelysin-2 (MMP-10), collagenase-3 (MMP-13), macrophage metal-
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loelastase (MMP-12), and tissue inhibitor of metalloproteinases-3 (TIMP-3) in intestinal ulcerations. Am J Pathol 1998; 152: 1005-1014.


[97] Boulday G, Fitau J, Coupel S, Soulillou JP and Charreau B. Exogenous tissue inhibitor of metalloproteinase-1 promotes endothelial cell survival through activation of the phosphati-

