

The Effect of Surgery on the Levels of Matrix Metalloproteinases in Patients with Inguinal Hernia

Sabina STROHALMOVÁ^{1,2}, Kateřina LEVOVÁ², Aleš Antonín KUBĚNA², Zdeněk KRŠKA¹, David HOSKOVEC¹, Tomáš ZIMA², Marta KALOUSOVÁ²

¹1st Department of Surgery - Department of Abdominal, Thoracic Surgery and Traumatology, First Faculty of Medicine, Charles University and General University Hospital in Prague, Prague, Czech Republic, ²Institute of Medical Biochemistry and Laboratory Diagnostics, First Faculty of Medicine, Charles University and General University Hospital in Prague, Prague, Czech Republic

Received December 1, 2020

Accepted April 13, 2021

Epub Ahead of Print June 2, 2021

Summary

Matrix metalloproteinases (MMPs) are associated with the alteration of extracellular matrix. The purpose of this study was to investigate how the levels of matrix metalloproteinases and their inhibitors – TIMPs are influenced by the presence of inguinal hernia as well as by its surgical treatment. The studied group consisted of 25 patients with inguinal hernia and 21 healthy controls for comparison. Two blood samples - before and after the treatment were collected from patients. Serum concentrations of MMPs and TIMPs were analysed by multiplex immunoassays. There was a difference in circulating levels of MMPs in patients before the surgery compared to healthy controls – the concentrations of MMP-2 and MMP-9 were significantly lower ($p=0.026$, $p=0.018$, respectively). After the surgery, the levels of MMPs, especially MMP-2 ($p<0.0001$), were significantly decreased in patients compared to the preoperative values, apart from MMP-9. On the contrary, MMP-9 showed significant increase after the surgery ($p<0.0001$). Circulation levels of TIMP-2 in patients were significantly decreased in comparison with controls ($p=0.004$), whereas levels of TIMP-1 were similar to controls. Both tested metalloproteinase inhibitors showed a significant decrease in detected levels (TIMP-1 $p=0.0004$; TIMP-2 $p<0.0001$) after the procedure compared to the preoperative values. The levels of MMPs, especially MMP-2 and MMP-9, and their inhibitors TIMP-1 and TIMP-2 are involved by the presence of inguinal hernia as well as are influenced by the surgery.

Key words

Matrix metalloproteinase • TIMP • Inguinal hernia

Corresponding author

Marta Kalousová, Institute of Medical Biochemistry and

Laboratory Diagnostics, First Faculty of Medicine, Charles University and General University Hospital in Prague, Kateřinská 32, 121 08 Prague 2, Czech Republic. E-mail: marta.kalousova@lf1.cuni.cz

Introduction

Hernia is a pathological process, protrusion of tissues and organs outside their physiological position. There can be several types of hernias, such as umbilical, femoral, inguinal hernia, and the inguinal hernia is considered to be the most common type among the abdominal wall hernias (Dabbas *et al.* 2011, Ramanan *et al.* 2014, Zöller *et al.* 2013).

On the basis of the passage through the abdominal wall we can divide inguinal hernias to direct and indirect. Direct inguinal hernia protrudes through fossa inguinalis medialis to trigonum Hesselbachi and exits in anulus inguinalis superficialis. Indirect inguinal hernia is going through inguinal canal with spermatic cord in men or ligamentum teres uteri in women – it enters in fossa inguinalis lateralis (anulus inguinalis profundus) and exits in anulus inguinalis superficialis (Holzheimer 2005). Indirect inguinal hernia can be both congenital and acquired, direct inguinal hernia is only acquired. Risk factors of developing inguinal hernia are states or activities connected with higher intraabdominal pressure – for example excessive effort, cough, constipation (Jiang *et al.* 2015). The treatment of inguinal hernia is surgical, while elective surgery is preferred.

Matrix metalloproteinases (MMPs) are a family of endopeptidases which contribute to various physiological and pathological processes. They degrade connective tissue and therefore participate in processes related to ontogenetical development, such as morphogenesis, angiogenesis, growth or wound healing (Cauwe *et al.* 2007, Lee *et al.* 2004). MMPs play a role in all the processes connected with increased synthesis, degradation, organisation or maturation of extracellular matrix. Thus, they are key regulators of numerous aspects of tissue remodeling and repair (Gill *et al.* 2008). Furthermore, altered activity of MMPs is linked with degenerative processes, as well as affects inflammatory response or cancer development (Gill *et al.* 2008, Jabłońska-Trypuć *et al.* 2016, Rosenberg 2009, Roy *et al.* 2009).

It has been previously reported that altered activity of metalloproteinases affects the etiopathogenesis of inguinal hernia through the imbalance in the rate of collagen metabolism (Antoniou *et al.* 2009, Henriksen *et al.* 2015, Klinge *et al.* 1999, Peeters *et al.* 2014).

MMPs action is mainly influenced by their endogenous tissue inhibitory proteins (TIMPs). Hence, the offset between TIMPs and MMPs levels is important for the extracellular matrix remodelling in the tissue.

The purpose of the study was to investigate how the circulating levels of matrix metalloproteinases and their inhibitors – TIMPs are influenced by the presence of inguinal hernia as well as by its surgical treatment.

Methods

Study objects

The study group consisted of 25 patients diagnosed with inguinal hernia (22 men and 3 women, median age 72, IQR 55-79 years) from the 1st Department of Surgery - Department of Abdominal, Thoracic Surgery and Traumatology, First Faculty of Medicine, Charles University and General University Hospital in Prague. These patients underwent elective surgery and had no signs of incarcerated hernia. Patients were cardiopulmonary compensated and without symptoms of any infection. Patients with malignancies were not included in the study.

As for the type of hernia, eight patients had direct inguinal hernia, ten patients indirect hernia, three patients had combination of direct and indirect hernia. Four patients suffered from recidive of inguinal hernia. Fourteen patients were treated for arterial hypertension,

ten patients with mild hypertension, four with severe hypertension. One of the patients with severe hypertension suffered from hypertensive kidney disease (serum level of creatinine 140 µmol/l). Two patients were treated for diabetes mellitus on the treatment with GLP1 – agonists, one with complications – diabetic neuropathy and diabetic nephropathy (serum level of creatinine below 150 µmol/l). Another patient had chronic kidney disease, creatinine 109 µmol/l. Eleven patients were diagnosed with heart diseases – five patients with ischemic heart disease, seven patients with atrial fibrillation, one patient with aortic regurgitation. Six patients had no internal comorbidities.

Twenty one healthy controls (eight men and thirteen women, median age 41 (IQR 30-53 years)) were used for comparison baseline levels before the surgery. Controls were volunteers without presence of inguinal hernia, without symptoms of acute infection or disease, without elevation of inflammatory markers in blood (CRP, leucocytes), without hepatic or renal failure, without presence of a tumour, and with blood count in the reference range. They were not taking any chronic medication. Healthy volunteers without internal comorbidities or presence of hernia were chosen to avoid the influence on the levels of MMPs. Results of routine laboratory examinations of the studied groups are presented in Table 1.

This study was performed in adherence to the principles of the Helsinki Declaration and approved by the Ethics Committee of The General University Hospital in Prague. All subjects gave their informed consent with participation in the study.

Laboratory analyses

Blood samples from all individuals were obtained after overnight fasting from the cubital vein into tubes without anticoagulant to obtain serum. From all patients, blood collection occurred preoperatively 1-3 h before the surgery and postoperatively 24 hours after the surgery. Blood collection occurred together with blood collection for standard laboratory analyses. Test tubes with blood were after coagulation centrifuged for 15 min at 900 x g and serum was stored at -80 °C until measurement of metalloproteinases and selected inhibitors.

Multiplex xMAP assay

Serum samples were analyzed with a 5 matrix metalloproteinases MILLIPLEX MAP Human MMP Magnetic Bead Panel 2 kit for simultaneous analysis of

Table 1. Basic laboratory characteristics of the study groups

Parameter	Patients	Healthy Controls	p value
Number	25	21	
Age (years)	72 (55-79)	41 (30-53)	<0.0001
Protein (g/l)	69 (66-71)	72 (67-74)	ns
CRP (mg/l)	1.8 (1.0-4.1)	3.1 (1.5-4.3)	ns
ALT (μ kat/l)	0.34 (0.28-0.49)	0.40 (0.31-0.48)	ns
AST (μ kat/l)	0.42 (0.36-0.48)	0.37 (0.30-0.45)	ns
GGT (μ kat/l)	0.43 (0.3-1.0)	0.36 (0.30-0.56)	ns
Urea (mmol/l)	6.2 (4.9-7.2)	4.9 (3.9-6.4)	ns
Creatinine (μ mol/l)	84 (78-97)	65 (58-77)	ns
Hemoglobin (g/l)	147 (142-154)	135 (131-151)	0.026
Leukocytes ($\times 10^9/l$)	6.5 (4.8-8.2)	6.5 (5.7-8.4)	ns
Thrombocytes ($\times 10^9/l$)	214 (178-252)	243 (204-305)	ns

All data shown as median (interquartile range). Abbreviations: CRP – C-reactive protein, ALT – Alanine aminotransferase, AST – Aspartate aminotransferase, GGT – Gamma-glutamyltransferase. All results were considered statistically significant at $p < 0.05$, ns, not significant.

tested metalloproteinases (MMP-1,-2,-7,-9,-10) and with MILLIPLEX MAP Human TIMP Magnetic Bead Panel 1 kit (Merck KGaA, Darmstadt, Germany) to obtain the concentrations of TIMP-1 and TIMP-2 inhibitors. Analytes were quantified using MAGPIX system, which utilizes xMAP technology and data acquisition xPONENT software (Luminex Corp., Austin, TX, USA). Data were then analyzed by MILLIPLEX[®] Analyst 5.1 software (Merck KGaA, Darmstadt, Germany). Concentrations of analytes were determined using 5PL curve fit. Two quality controls (low, high) were run with each assay. All quality control results were found to fall within the expected ranges provided by the manufacturer. Results of all MMPs and TIMPs are expressed in ng/ml.

Statistical analysis

Statistical tests were used to examine the differences among studied groups (patients before and after treatment, controls). We used the nonparametric Mann Whitney U test for group comparison (age), and the Generalized Linear Model with Gamma Distribution and reciprocal link function for comparisons with adjustment for sex and age (biochemical variables). Differences between patient's serum analytes levels before and after the treatment were analyzed using Wilcoxon signed rank test. Spearman's rank correlation coefficient was used to determine the correlations between studied analytes. Data are expressed as median (IQR, interquartile range). Differences among the subgroups with different types of hernia were evaluated

using the Kruskal–Wallis test. All results were considered statistically significant at $p < 0.05$ and all reported p values were 2-sided. Statistical analysis was performed using GraphPad Prism software 8 (San Diego, CA, USA) and for Generalized linear model the Wolfram Mathematica software, version 11.3 (United Kingdom) was used.

Results

Serum values of tested metalloproteinases and selected main inhibitors were analyzed at two stages - before and after the surgery, and the corresponding analytes' levels were additionally compared to the healthy control group.

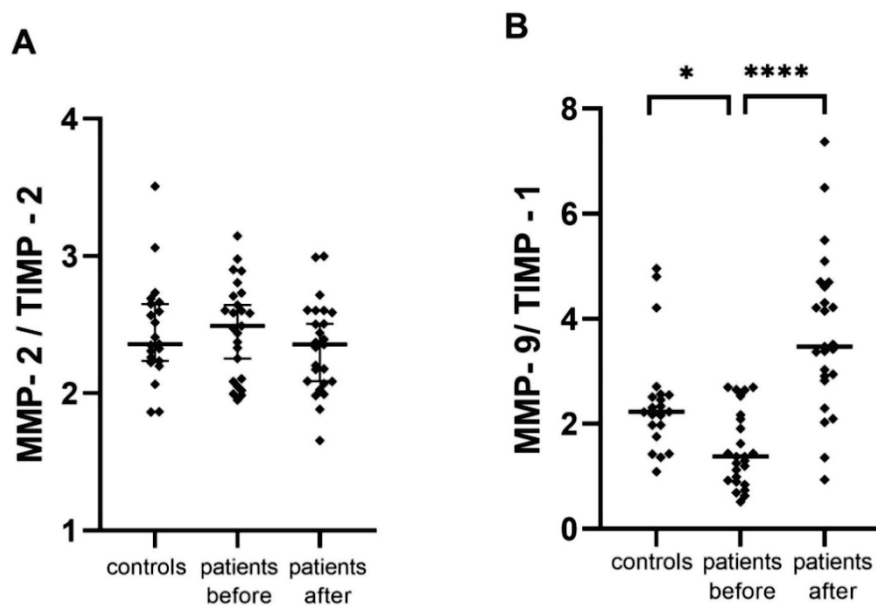
Serum concentrations of metalloproteinases and TIMPs in patients compared to the control group are listed in Table 2. Patients with inguinal hernia had significantly decreased preoperative concentrations of MMP-2 and MMP-9 ($p=0.026$, $p=0.018$, respectively), whereas circulating blood levels of MMP-1,-7,-10 were without change. Different trends were likewise found for tested inhibitors. Preoperative levels of TIMP-2 in patients were significantly decreased in comparison with the control group ($p=0.004$), whereas levels of TIMP-1 were similar to controls.

Patients' serum levels of all tested analytes were significantly different when compared blood draw before and after the surgery (Table 2). After surgery there was a significant decrease in MMPs levels apart from MMP-9, while most significant decrease was observed

Table 2. MMPs and TIMPs serum levels of patients and control group.

Analyte ng/ml	Patients		Controls	p value Before vs After	p value Before vs controls
	Before	After			
<i>MMP-1</i>	9.5 (7.0-12.2)	9.2 (6.6-12.0)	5.5 (3.3-10.7)	0.016	ns
<i>MMP-2</i>	112.3 (92.9-126.0)	91.4 (80.9-103.8)	117.4 (100.7-131.9)	<0.0001	0.026
<i>MMP-7</i>	13.8 (10.7-19.6)	12.5 (9.7-16.3)	11.6 (8.7-12.9)	0.021	ns
<i>MMP-9</i>	133.5 (99.3-210.9)	345.3 (254.5-455.4)	236.4 (177.5-279.0)	<0.0001	0.018
<i>MMP-10</i>	0.44 (0.33-0.72)	0.44 (0.29-0.54)	0.44 (0.35-0.54)	0.014	ns
<i>TIMP-1</i>	103.8 (93.9-115.4)	90.9 (79.1-109.2)	100.4 (91.7-112.5)	0.0004	ns
<i>TIMP-2</i>	45.8 (42.4-47.8)	39.5 (37.0-41.7)	48.2 (44.3-52.3)	<0.0001	0.004

Data are expressed as median (interquartile range). All results were considered statistically significant at $p < 0.05$, ns, not significant.

**Fig. 1.** Ratios of MMPs/TIMPs among the study groups, * $p = 0.026$; **** $p < 0.0001$

for MMP-2 ($p < 0.0001$). In contrast, we observed a significant elevation of MMP-9 levels ($p < 0.0001$) after the surgery. Both tested metalloproteinase inhibitors showed a significant decrease in detected levels (TIMP-1 $p = 0.0004$; TIMP-2 $p < 0.0001$) after the procedure compared to the preoperative values. We did not observe any differences in patients MMPs/TIMPs levels when evaluating subgroups with various types of hernia (direct or indirect, combination of direct/indirect or recidive).

The correlation between tested analytes was also determined. There was a significant correlation between MMP-2 and TIMP-2 in both patients before the surgery and controls ($\rho = 0.66$, $p = 0.0004$; $\rho = 0.64$, $p = 0.002$, respectively) and this correlation was no longer apparent after the procedure. However, after the surgery both inhibitors TIMP-1,2 were significantly correlated ($\rho = 0.64$, $p = 0.001$).

Serum levels of MMP-2 and MMP-9 were also

evaluated as ratios with corresponding dominant inhibitors (Fig. 1). The MMP-9/TIMP-1 preoperative levels ratio was significantly decreased compared to the control group ($p=0.026$), and after the procedure there was a significant increase with respect to the preoperative levels ($p<0.0001$), thereby MMP-9/TIMP-1 ratio copied the same trend as MMP-9 serum levels.

Discussion

In this study, we analyzed the circulating levels of MMPs and TIMPs in patients with inguinal hernia before and after the surgical treatment and compared the preoperative values to healthy controls.

Inguinal hernia formation is connected with tissue alteration. The process of hernia development is associated with the disturbance of regulation of metabolism of collagen (Henriksen *et al.* 2011). MMPs enzymes play a key role in extracellular matrix metabolism during tissue remodeling and repair (Gill *et al.* 2008). They are predominantly extracellularly, however several studies have shown that MMP-1 (Limb *et al.* 2005), MMP-2 (Aldonyte *et al.* 2009, Kwan *et al.* 2004), MMP-11 (Luo *et al.* 2002) and MMP-13 (Cuadrado *et al.* 2009) can also be found intracellularly and may act on intracellular proteins.

In our study, we focused on how are the circulating levels of tested analytes influenced by the presence of inguinal hernia as well as by the surgery. We found significant differences in the preoperative levels of MMP-2, MMP-9 and TIMP-2 compared to the controls. While preoperative serum levels of MMP-2, MMP-9 and TIMP-2 were decreased ($p=0.026$, $p=0.019$, $p=0.004$, respectively), levels of MMP-1, MMP-10 and TIMP-1 were similar to controls.

There are several studies, which were also concentrated on the circulating levels of matrix metalloproteinases and their inhibitors in inguinal hernias. But the levels of MMPs both in serum and plasma, especially for MMP-2 and MMP-9, as well as for TIMPs were reported inconsistently.

Relationship between inguinal hernia and selected preoperative serum levels of MMPs as well as TIMPs in patients with direct, indirect and bilateral type of hernia was investigated by Isik and coworkers. When compared to healthy donors, all three groups of patients revealed significant elevation of all four tested MMPs (MMP-1,2,9,13) whereas levels of TIMPs were decreased. These analytes were also tested in tissue

homogenates taken from transversalis fascia with similar trend results as for serum samples (Isik *et al.* 2017). In contrast, our data indicate that serum preoperative levels of MMP-1 and TIMP-1 were without significant changes and MMP-2,-9 were significantly decreased compared to the controls ($p=0.026$, $p=0.018$, respectively). Significant decrease of TIMP-2 was in line with our results.

Decreased circulating levels of MMP-2,-9 were previously also reported in plasma of patients with inguinal hernia (Antoniou G. A. *et al.* 2011). In this study, 91 patients and 35 controls were compared for both circulating and tissue levels of MMP-2,9 and TIMP-1,2. Whereas their results of tested biomarkers in tissue of the abdominal fascia were consistent with those reported by Isik *et al.* (2017), they differed in circulating preoperative levels. Antoniou *et al.* (2011) found plasma concentrations of MMP-9 and MMP-2 significantly decreased in patients with inguinal hernia in comparison with the control group, which corresponds to our results. On the other hand, several former studies reported significantly higher MMP-2 levels in both serum and tissue samples in patients with direct or indirect inguinal hernia than those in controls (Bellón *et al.* 2001, Li *et al.* 2018, Smigielski *et al.* 2011, Smigielski *et al.* 2009, Wang *et al.* 2020).

Serum MMP-2,-9 were also tested in patients with inguinal hernias along with procollagen type I,III N-terminal propeptides (PINP, PIIINP, respectively) (Li *et al.* 2018). In contrast to our MMP-9 results, as well as Isik *et al.* (2017) and Antoniou *et al.* (2011), they observed no significant changes in serum MMP-9 preoperative levels between the patients and controls (Li *et al.* 2018).

MMPs proteolytic activities influence crucial cellular processes as well as many other physiological events and their activities are mainly regulated by TIMPs. Results regarding the circulating plasma levels of TIMP-1,-2 were also previously reported differently. Significantly higher circulating plasma levels in patients with inguinal hernia compared to controls were previously reported for TIMP-2 and decreased levels of TIMP-1 (Antoniou *et al.* 2011). Furthermore, elevation of TIMP-2, but only in recurrent inguinal hernia patients were noticed by Smigielski *et al.* (2011), while the levels of TIMP-2 in other groups (direct, indirect hernias) were similar to controls. On the other hand significantly decreased serum levels of TIMP-1,2,3 compared to control group were reported (Isik *et al.* 2017). In our study preoperative levels of TIMP-2 were significantly

decreased in comparison with controls ($p=0.004$), whereas levels of TIMP-1 were similar to controls. Thus, the decrease of circulating levels of MMP-9 in patients with inguinal hernia before the surgery was not followed by an increase of the main MMP-9 inhibitor – TIMP-1, because TIMP-1 serum levels were similar to those detected in controls. We found a significant correlation between both inhibitors (TIMP-1, TIMP-2) ($\rho=0.64$, $p=0.001$) after the surgery, where both inhibitors showed a significant decrease in detected levels (TIMP-1 $p=0.0004$; TIMP-2 $p<0.0001$) compared to preoperative values.

Some previous studies were focused on differences in MMPs levels according to the type of hernia. We did not observe any differences in patients MMPs and TIMPs levels both before and after the procedure when they were evaluated according to the type of hernia (direct or indirect). These results are in agreement with those previously reported for MMP-1,-2,-9 (Isik *et al.* 2017, Li *et al.* 2018, Smigielski *et al.* 2009).

By reason of each TIMP displays preferential MMP-binding specificity, the MMP/TIMP ratios can determine the overall proteolytic activity. We observed a higher MMP-9/TIMP-1 ratio of circulating levels in patients after the procedure compared to the preoperative values, consistent with a net MMP-9 increase in patients after surgery. Significant elevation of MMP-2/TIMP-2 mRNA ratios in primary inguinal hernias patients (both direct and indirect) compared to the controls was previously reported, where the MMP-2 and TIMP-2 expression in anterior rectus sheath fascia was done (Wang *et al.* 2020). In contrast, we did not observe a significant change in MMP-2/TIMP-2 circulating levels ratio between patients (blood draw before and after the surgery) and control group.

Most studies concerned with MMPs and TIMPs circulating levels at a single time point – as preoperative values. Our study was aimed to describe the changes in biomarkers levels before and after the surgical procedure. In patients after surgery, there was a significant decrease in MMPs levels, except for MMP-9. MMP-9 showed a significant increase ($p<0.0001$). Within the standardization of procedures, we dealt with the impact of surgical intervention on selected biomarkers. By default, samples are taken into the biobank prior to the planned

surgical intervention and any subsequent sampling can significantly affect the results. The results of the study confirm that the sampling time with respect to the surgical performance can significantly affect the levels of the studied parameters. Standardization of the collection time into the biobank with respect to the surgical performance is therefore a very important element for ensuring the comparability of the collected samples.

We have to admit some limitations of our study. Control group individuals were notably younger than patients with inguinal hernia. Furthermore, the percentage of women in the control group was higher than in the patient's group. Comparisons of MMPs and TIMPs levels between the groups were therefore statistically adjusted for the age and sex differences. Likewise, statistical evaluation of differences in MMPs and TIMPs levels according to the type of hernia has an informative value only between groups with direct and indirect type of hernia, due to the small number of patients in two remaining groups - combination of direct/indirect and recidive inguinal hernia.

The levels of MMPs, especially MMP-2 and MMP-9, and their inhibitors TIMP-1 and TIMP-2 are involved by the presence of inguinal hernia as well as are influenced by the surgery.

Conflict of Interest

There is no conflict of interest.

Acknowledgements

This work was supported by European Regional Development Fund project [EF16_013/0001674], by Ministry of Health, Czech Republic – conceptual development of research organization [VFN-64165], and by research projects Progres Q25, and SVV [260524/2020]. The authors are thankful to the laboratory staff mainly to Mrs. Dita Hudcová and Mrs. Marcela Bartáková from the Institute of Medical Biochemistry and Laboratory Diagnostics, First Faculty of Medicine, Charles University and General University Hospital in Prague for technical assistance. The authors would like to thank also to physicians and nurses from the First Department of Surgery - Department of Abdominal, Thoracic Surgery and Traumatology, First Faculty of Medicine, Charles University and General University Hospital in Prague.

References

- ALDONYTE R, BRANTLY M, BLOCK E, PATEL J, ZHANG J: Nuclear localization of active matrix metalloproteinase-2 in cigarette smoke-exposed apoptotic endothelial cells. *Exp Lung Res* 35: 59-75, 2009. <https://doi.org/10.1080/01902140802406059>
- ANTONIOU GA, TENTES IK, ANTONIOU SA, SIMOPOULOS C, LAZARIDES MK: Matrix metalloproteinase imbalance in inguinal hernia formation. *J Invest Surg* 24: 145-150, 2011. <https://doi.org/10.3109/08941939.2011.558610>
- ANTONIOU SA, ANTONIOU GA, GRANDERATH FA, SIMOPOULOS C: The role of matrix metalloproteinases in the pathogenesis of abdominal wall hernias. *Eur J Clin Invest* 39: 953-959, 2009. <https://doi.org/10.1111/j.1365-2362.2009.02199.x>
- BELLÓN JM, BAJO A, GA-HONDUVILLA N, GIMENO MJ, PASCUAL G, GUERRERO A, BUJÁN J: Fibroblasts from the transversalis fascia of young patients with direct inguinal hernias show constitutive MMP-2 overexpression. *Ann Surg* 233: 287-291, 2001. <https://doi.org/10.1097/00000658-200102000-00020>
- CAUWE B, VAN DEN STEEN PE, OPDENAKKER G: The biochemical, biological, and pathological kaleidoscope of cell surface substrates processed by matrix metalloproteinases. *Crit Rev Biochem Mol Biol* 42: 113-185, 2007. <https://doi.org/10.1080/10409230701340019>
- CUADRADO E, ROSELL A, BORRELL-PAGES M, GARCIA-BONILLA L, HERNANDEZ-GUILLAMON M, ORTEGA-AZNAR A, MONTANER J: Matrix metalloproteinase-13 is activated and is found in the nucleus of neural cells after cerebral ischemia. *J Cereb Blood Flow Metab* 29: 398-410, 2009. <https://doi.org/10.1038/jcbfm.2008.130>
- DABBAS N, ADAMS K, PEARSON K, ROYLE G: Frequency of abdominal wall hernias: is classical teaching out of date? *JRSM Short Reports* 2: 5, 2011. <https://doi.org/10.1258/shorts.2010.010071>
- GILL SE and PARKS WC: Metalloproteinases and their inhibitors: regulators of wound healing. *Int J Biochem Cell Biol* 40: 1334-1347, 2008. <https://doi.org/10.1016/j.biocel.2007.10.024>
- HENRIKSEN NA, MORTENSEN JH, SORENSEN LT, BAY-JENSEN AC, AGREN MS, JORGENSEN LN, KARSDAL MA: The collagen turnover profile is altered in patients with inguinal and incisional hernia. *Surgery* 157: 312-321, 2015. <https://doi.org/10.1016/j.surg.2014.09.006>
- HENRIKSEN NA, YADETE DH, SORENSEN LT, AGREN MS, JORGENSEN LN: Connective tissue alteration in abdominal wall hernia. *Br J Surg* 98: 210-219, 2011. <https://doi.org/10.1002/bjs.7339>
- HOLZHEIMER RG: Inguinal Hernia: classification, diagnosis and treatment--classic, traumatic and Sportsman's hernia. *Eur J Med Res* 10: 121-134, 2005.
- ISIK A, GURSUL C, PEKER K, AYDIN M, FIRAT D, YILMAZ I: Metalloproteinases and their inhibitors in patients with inguinal hernia. *World J Surg* 41: 1259-1266, 2017. <https://doi.org/10.1007/s00268-016-3858-6>
- JABŁOŃSKA-TRYPUĆ A, MATEJCZYK M, ROSOCHACKI S: Matrix metalloproteinases (MMPs), the main extracellular matrix (ECM) enzymes in collagen degradation, as a target for anticancer drugs. *J Enz Inhib Med Chem* 31: 177-183, 2016. <https://doi.org/10.3109/14756366.2016.1161620>
- JIANG ZP, YANG B, WEN LQ, ZHANG YC, LAI DM, LI YR, CHEN S: The etiology of indirect inguinal hernia in adults: congenital or acquired? *Hernia* 19: 697-701, 2015. <https://doi.org/10.1007/s10029-014-1326-5>
- KLINGE U, ZHENG H, SI Z, SCHUMPELICK V, BHARDWAJ RS, MUYS L, KLOSTERHALFEN B: Expression of the extracellular matrix proteins collagen I, collagen III and fibronectin and matrix metalloproteinase-1 and -13 in the skin of patients with inguinal hernia. *Eur Surg Res* 31: 480-490, 1999. <https://doi.org/10.1159/000008728>
- KWAN JA, SCHULZE CJ, WANG W, LEON H, SARIAHMETOGLU M, SUNG M, SAWICKA J, SIMS DE, SAWICKI G, SCHULZ R: Matrix metalloproteinase-2 (MMP-2) is present in the nucleus of cardiac myocytes and is capable of cleaving poly (ADP-ribose) polymerase (PARP) in vitro. *FASEB J* 18: 690-692, 2004. <https://doi.org/10.1096/fj.02-1202fje>
- LEE MH, MURPHY G: Matrix metalloproteinases at a glance. *J Cell Sci* 117: 4015-4016, 2004. <https://doi.org/10.1242/jcs.01223>

-
- LI J, ZHANG X, SUN Q, LI W, YU A, FU H, CHEN K: Circulating matrix metalloproteinases and procollagen propeptides in inguinal hernia. *Hernia* 22: 541-547, 2018. <https://doi.org/10.1007/s10029-018-1751-y>
- LIMB GA, MATTER K, MURPHY G, CAMBREY AD, BISHOP PN, MORRIS GE, KHAW PT: Matrix metalloproteinase-1 associates with intracellular organelles and confers resistance to lamin A/C degradation during apoptosis. *Am J Pathol* 166: 1555-1563, 2005. [https://doi.org/10.1016/S0002-9440\(10\)62371-1](https://doi.org/10.1016/S0002-9440(10)62371-1)
- LUO D, MARI B, STOLL I, ANGLARD P: Alternative splicing and promoter usage generates an intracellular stromelysin 3 isoform directly translated as an active matrix metalloproteinase. *J Biol Chem* 277: 25527-25536, 2002. <https://doi.org/10.1074/jbc.M202494200>
- PEETERS E, DE HERTOOGH G, JUNGE K, KLINGE U, MISEREZ M: Skin as marker for collagen type I/III ratio in abdominal wall fascia. *Hernia* 18: 519-525, 2014. <https://doi.org/10.1007/s10029-013-1128-1>
- RAMANAN B, MALOLEY BJ, FITZGIBBONS RJ Jr.: Inguinal hernia: follow or repair? *Adv Surg* 48: 1-11, 2014. <https://doi.org/10.1016/j.yasu.2014.05.017>
- ROSENBERG GA: Matrix metalloproteinases and their multiple roles in neurodegenerative diseases. *Lancet. Neurol* 8: 205-216, 2009. [https://doi.org/10.1016/S1474-4422\(09\)70016-X](https://doi.org/10.1016/S1474-4422(09)70016-X)
- ROY R, YANG J, MOSES MA: Matrix metalloproteinases as novel biomarkers and potential therapeutic targets in human cancer. *J Clin Oncol* 27: 5287-5297, 2009. <https://doi.org/10.1200/JCO.2009.23.5556>
- SMIGIELSKI J, BROCKI M, KUZDAK K, KOŁOMECKI K: Serum MMP 2 and TIMP 2 in patients with inguinal hernias. *Eur J Clin Invest* 41: 584-588, 2011. <https://doi.org/10.1111/j.1365-2362.2010.02445.x>
- SMIGIELSKI J, KOŁOMECKI K, ZIEMNIAK P, DROZDA R, AMSOLIK M, KUZDAK K: Degradation of collagen by metalloproteinase 2 in patients with abdominal hernias. *Eur Surg Res* 42: 118-121, 2009. <https://doi.org/10.1159/000187643>
- WANG D, HAN Y, XU X, CHEN J, CHEN Y: Matrix Metalloproteinases (MMP-2) and Tissue Inhibitors of Metalloproteinases (TIMP-2) in Patients with Inguinal Hernias. *World J Surg*, 2020. <https://doi.org/10.1007/s00268-020-05674-0>
- ZÖLLER B, JI J, SUNDQUIST J, SUNDQUIST K: Shared and nonshared familial susceptibility to surgically treated inguinal hernia, femoral hernia, incisional hernia, epigastric hernia, and umbilical hernia. *J Am Coll Surg* 217: 289-299.e281, 2013. <https://doi.org/10.1016/j.jamcollsurg.2013.04.020>
-