Methylene blue improves hemodynamic shock but increases lipoperoxidation in severe acute pancreatitis pig model

Azul de metileno melhora as condições hemodinâmicas do choque circulatório, mas aumenta a peroxidação lipídica em um modelo suino de pancreatite aguda grave

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ABSTRACT

Purpose: Study hemodynamic pattern and lipoperoxidation during methylene blue (MB) treatment on taurocholate-enterokinase induced acute pancreatitis (AP).

Methods: Thirty pigs were equally divided in control group; MB group; AP group; MB previous AP group; and MB after 90 min of induced AP group. MB was given iv in a bolus dose (2mg.kg-1) followed by maintenance dose (2mg.kg-1.h-1). Hemodynamic parameters were recorded continuously during 180 min by Swan-Ganz catheter. Blood samples were taken every 60 min to determine arterial and venous nitrate, malondialdehyde (MDA) and amylase. Pancreatic tissue was removed for histopathologic study.

Results: In AP group MBP and CO decreased over time 33% (p<0.05) and 52% (p<0.05), respectively. In MB previous induced-AP group, there was 70 minutes delay (p<0.05) to decrease MBP and CO. In MB group arterial and venous nitrite decreased (p<0.05) over time. MB infusion increased (p>0.05) serum MDA when associated to AP. After induced AP, MB did not reverse MBP and CO decrease. There was no difference in serum amylase and necro-hemorrhagic findings with MB treatment.

Conclusions: In this taurocholate-induced AP model MB treatment delayed hemodynamic shock and decreases serum nitrate levels but increases serum MDA levels. No volemic replacement was done and it may have been a mitigated factor to a poor tissue perfusion and impairment microcirculation. Further investigations are needed to elucidate MB treatment role during AP treatment.

Key words: Acute Pancreatitis. Methylene Blue. Nitric Oxide. Shock. Lipoperoxidation. Taurocholate.

RESUMO

Objetivo: estudar o perfil hemodinâmico e a lipoperoxidação durante o tratamento com azul de metileno (AM) de pancréatite aguda (PA) induzida por taurocolato-enterokinase. Métodos: Trinta porcos foram igualmente divididos em: grupo controle, grupo AM; grupo PA; grupo AM prévio à PA; grupo AM após 90 minutos após a indução da PA. O AM foi administrado sob a forma de bolus EV (2mg.kg-1) seguido por dose de manutenção (2mg.kg-1.h-1). Os parâmetros hemodinâmicos foram registrados continuamente durante 180 min com auxílio de cateter de Swan-Ganz. Amostras sanguíneas foram colhidas a cada 60 min para a determinação arterial e venosa de nitrato, malondialdeído (MDA) e amilase. Foi feito o corte pancreático para estudo histopatológico. Resultados: No grupo PA a pressão arterial média (PAM) e o débito cardíaco (DC) diminuíram respectivamente 33% (p<0.05) e 52% (p<0.05) no decorrer do tempo. No grupo AM prévio à indução da PA ocorreu 70 minutos de demora (p<0.05) para as diminuições da PAM e DC. No grupo AM houve diminuição temporal do nitrato arterial e venoso (p<0.05). A infusão de AM aumentou os valores de MDA sérica quando associado a PA (p<0.05). Após a indução da PA a infusão de AM não reverteram as quedas da PA e DC. Não houve diferenças nos níveis de amilase sérica e achados histológicos com o tratamento com o azul de metileno. Conclusões: No presente modelo de PA induzida por taurococolato o AM retardou o desenvolvimento do choque circulatório, diminuiu os níveis de nitrito mas aumentou os níveis de MDA. Não se realizou nenhuma reposição volêmica que poderia melhorar a perfusão tecidual e melhora da microcirculação. Investigações adicionais são necessárias para elucidar o papel terapêutico do AM no tratamento da PA aguda.


1. Research performed at Department of Surgery and Anatomy, Ribeirão Preto Faculty of Medicine, University of São Paulo, SP, Brazil.
Introduction

The nitric oxide (NO) role in acute pancreatitis (AP) pathogenesis remains controversial with some studies suggesting that it potentiates the oxidative stress and pancreatic parenchyma injury, while others report that it reduces the pancreatic dysfunction by increasing pancreatic blood flow in response to NO releasing by endothelial cells. Mb blocks GMP-c increase, the ultimate effector pathway of NO production with vasodilatation action. MB also may inhibit the NO synthesis.

MB clinical use has been reported in sepsis, renal and hepatic transplantation, anaphylactic shock and shock associated to extracorporal circulation.

In the present study, MB was employed to evaluated the hemodynamic and biochemical effects in a severe AP model in pigs.

Methods

Surgical preparation of the animals. Thirty healthy male Dalland pigs weighing between 20 to 24 kg were studied at the Laboratory of Experimental Surgery, Department of Surgery and Anatomy, Faculdade de Medicina de Ribeirão Preto, University of São Paulo (FMRP-USP). The experimental model protocol and animal handling procedures had been accomplishment including tracheostomy, venous and arterial access, hemodynamic monitoring, blood samples collection, main pancreatic duct cannulation, the animals were observed by a period of 45 minutes for hemodynamic stability. After this stability period, hemodynamic parameters were collected and registered continuously for 180 min using a MP System 100 (BiO PacSystem, Incorporation, Santa Barbara, CA, U.S.A.). Arterial and venous blood samples were collected to biochemical analyses every 60 min from the beginning of the experiment (instant zero).

Experimental Protocol. Group Control (CTL) (n=6): the animals received in sequence: a) 20 mL i.v. “bolus” of saline followed by saline maintenance (3 mL.kg-1 body weight.h-1 i.v.) b) continuous infusion of saline during the period of 1 min in the main pancreatic duct; Group MB (n=6): the animals received in sequence: a) MB solution i.v.; b) continuous infusion of saline during the period of 1 min in the main pancreatic duct; Group AP (n=6): the animals received in sequence: a) 20 mL i.v. “bolus” of saline followed by saline maintenance (3 mL.kg-1 body weight.h-1 i.v.) b) AP induction; Group MB plus AP (n=6): the animals received in sequence: MB solution i.v.; b) AP induction; Group AP plus MB (n=6): the animals received in sequence: a) 20 mL i.v. “bolus” of saline followed by saline maintenance (3 mL.kg-1 body weight.h-1 i.v.) b) AP induction; c) after 90 minutes of AP induction, MB solution i.v..

AP induction. AP was induced immediately after basal parameters of hemodynamic, arterial and venous blood samples were collected. In brief, a continuous intraductal infusion of a 5% solution of sodium taurocholate (Sigma, St. Louis, Me, USA) in a dose of 1 mL.kg-1 body weight with enterokinase (Sigma, St. Louis, Me, USA) in a dose of 8 U.kg-
1 body weight during a 1 minute period was performed as described also were.\textsuperscript{10,11}

**MB infusion.** MB (Merck Diagnostica, Darmstadt, Germany) infusion was carried through an initial dose of 2mg.kg\(^{-1}\) body weight diluted in 20 ml of 0.9\% sodium chloride solution followed by a maintenance dose of 2 mg.kg\(^{-1}\) body weight.h\(^{-1}\) was done by a continuous intravenous infusion throughout the experiment.

**Nitrate Dosage.** Serum nitrate was indirectly measured by a Sievers NOAnalizer 280i using a NO/ozone chemiluminescence technique.\textsuperscript{12}

**Malondialdehyde Dosage.** MDA dosage was done using a Lipid Peroxidation Assay (Calbiochem San Diego, CA, U.S.A) as described previously.\textsuperscript{13}

**Biochemical Serum Analysis.** Amilase, glicemia, creatinin, hematocrit, blood gas analyze were done using either a Stat Profile 9 Plus Analyzer (New Biomedical Co., ME, USA.) or a RapidLab 860 (Chiron Diagnostics Co. Me, the U.S.A) or COULTER T-540 (Coulter Co., FL, U.S.A) or a Dimension device AIR (Dade Behring Incorporation, Deerfield, IL, U.S.A).

**Pancreas Biopsy and Morphologic Study.** At the end of data collection the previous laparotomy was extended from the xiphoid to the suprapubic region and the animals were sacrificed by exsanguination. The pancreas was identified, isolated and removed in all its extension. The pancreas was immediately fixed in a 10\% formol solution, after which time the slides were prepared from paraffin blocks and stained with hematoxylin-eosin stain. Biopsy samples from all 30 animals were then examined blindly in order to eliminate observer bias. The slides were screened under light microscope for swelling, inflammatory infiltration, necrosis and hemorrhage.

**Statistical Analysis**

The results were expressed in arithmetic mean and standard deviation (sd). Serum amylase, glucose, creatinin and hematocrit were analyzed in 60 minutes intervals from time zero. Collected data were stored in a computer and plot in spread sheets of the Excel program (Microsoft, Seattle, WA, the U.S.A), and statistically analyzed using a Minitab 12.2.2 software (Minitab Incorporation, State College, PA, USA) and a S-Plus 3.3 software (Mathsoft Engineering & Education Incorporation, Cambridge, MA, U.S.A). The longitudinal comparisons were made by analysis of variance for repeated measures, and each instant for analysis of variance (ANOVA) with comparisons of Tukey, or Kruskal-Wallis test, based on rank sum. Fisher’s angular transformation was used to express results in percentage. P <0.05 were considered statistically significant.

**Results**

Hemodynamic results are summarized in Table 1. Figure 1 illustrate the continuously alterations in CO that occurred in all groups. MBP, CVP, PCP, CO, CI and DO\(_2\) decreased (p<0.05) over time in all AP groups. HR increased (p<0.005) in all AP groups over time. SVR and PVR increased (p<0.05) in AP and AP+MB groups. In MB+AP group, there was a 70 minutes delay in the MBP and CO decrease (p<0.05) when compared to AP group. DO\(_2\) diminished in all AP groups although MB immediately before AP induction delayed DO\(_2\) decrease. O\(_2\)EI increased (p<0.05) in all AP groups.

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**TABLE 1 - Hemodynamics results in all groups**

| Group   | Time (min) | MBP (mmHg) | CVP (mmHg) | HR (Bpm) | MPAP (mmHg) | PCP (mmHg) | CO (L/min) | CI (L/min/m²) | SVR (dynas/cm²) | PVR (dynas/cm²) | DO₂ (mlO₂/min) | VO₂ (mlO₂/min) | O₂ El % | SvO₂ % |
|---------|------------|------------|------------|----------|-------------|------------|------------|--------------|----------------|----------------|----------------|----------------|----------|---------|-------|
| **CTL** | 0          | 121(17)    | 9(2)       | 107(30)  | 21(3)       | 10(1)      | 3.0(0.7)   | 4.3(0.9)     | 2929(621)      | 305(127)       | 410(135)       | 91(23)         | 20(3)    | 70(17) |
|         | 60         | 126(10)    | 8(1)       | 104(30)  | 20(3)       | 10(1)      | 3.8(0.8)   | 5.0(1.0)     | 2601(1659)     | 241(118)       | 518(145)*      | 137(50)*       | 25(8)   | 70(18) |
|         | 120        | 141(20)    | 8(1)       | 110(26)  | 19(2)       | 10(1)      | 3.8(1.0)   | 5.0(1.5)     | 2904(680)      | 233(79)        | 537(161)       | 172(82)        | 30(8)*  | 70(15) |
|         | 180        | 142(13)    | 7(1)       | 102(27)  | 16(4)       | 9(1)       | 3.6(0.8)   | 4.7(1.1)     | 3151(835)      | 216(179)       | 515(156)       | 114(37)        | 22(9)   | 70(17) |
| **MB**  | 0          | 112(30)    | 9(1)       | 107(26)  | 21(3)       | 10(1)      | 2.7(0.8)   | 3.5(1.1)     | 3332(1632)     | 367(151)       | 379(145)       | 97(27)         | 25(6)   | 72(18) |
|         | 60         | 133(29)    | 9(2)       | 106(22)  | 22(3)       | 11(2)*     | 2.9(0.7)   | 3.8(0.9)     | 3134(1310)     | 332(123)       | 408(89)        | 113(17)        | 27(8)   | 72(14) |
|         | 120        | 127(22)    | 9(2)       | 111(33)  | 22(3)       | 11(2)     | 2.8(0.5)   | 3.8(0.7)     | 3523(891)      | 336(92)        | 423(84)        | 93(22)         | 21(9)   | 73(12) |
|         | 180        | 127(22)    | 9(2)       | 110(22)  | 23(3)       | 11(2)*     | 2.8(0.5)   | 3.7(0.7)     | 3580(1053)     | 34(104)        | 429(83)        | 124(90)        | 26(15)  | 75(9)  |
| **AP**  | 0          | 120(14)    | 8(1)       | 119(17)  | 16(3)       | 10(2)      | 3.2(0.4)   | 4.2(0.6)     | 2992(652)      | 183(63)        | 503(93)        | 120(88)        | 23(21)  | 75(24) |
|         | 60         | 91(9)*     | 6(2)*      | 142(26)  | 14(3)       | 8(1)*      | 2.4(0.4)   | 3.2(0.6)     | 3000(679)      | 207(72)        | 411(73)        | 131(19)        | 31(9)   | 69(26) |
|         | 120        | 85(12)*    | 4(2)*      | 169(41)  | 16(4)       | 7(2)*      | 1.8(0.2)*  | 2.3(0.4)*    | 3739(860)*     | 411(117)*      | 344(47)*       | 126(22)        | 35(6)   | 59(24)*|
|         | 180        | 80(12)*    | 2(2)*      | 213(51)* | 13(2)       | 6(3)*      | 1.5(0.2)*  | 2.0(0.3)*    | 4229(755)*     | 339(117)       | 320(58)*       | 140(40)        | 43(14)* | 48(18)*|
| **MB+AP** | 0        | 108(22)    | 7(1)       | 97(15)   | 14(1)       | 10(1)      | 2.8(0.7)   | 3.8(0.9)     | 3007(631)      | 128(49)        | 400(116)       | 123(68)        | 30(12)  | 73(17) |
|         | 60         | 92(13)*    | 5(2)       | 119(24)  | 13(3)       | 10(2)      | 2.8(0.4)   | 3.7(0.4)     | 2582(402)      | 93(54)         | 420(57)        | 112(33)        | 26(10)  | 69(20) |
|         | 120        | 81(8)*     | 4(3)       | 139(21)* | 14(3)       | 8(2)       | 2.0(0.3)*  | 2.8(0.5)*    | 3116(555)      | 214(150)       | 348(55)*       | 127(17)†       | 36(10)*  | 61(21)*|
|         | 180        | 75(17)*    | 3(2)*      | 154(43)* | 13(4)*      | 8(1)*      | 1.9(0.4)*  | 2.5(0.5)*    | 3163(733)      | 212(189)       | 340(73)†       | 114(26)†       | 33(8)   | 66(19)*|
| **AP+MB** | 0        | 89(14)     | 7(1)       | 104(18)  | 15(2)       | 10(1)      | 2.7(0.5)   | 3.6(0.7)     | 2619(768)      | 150(56)        | 399(108)       | 55(20)         | 13(8)   | 86(9)  |
|         | 60         | 79(7)      | 6(3)*      | 123(19)  | 15(3)       | 7(2)*      | 2.5(0.5)   | 3.3(0.7)     | 250(1413)      | 276(168)*      | 394(79)        | 98(31)*        | 23(5)   | 84(6)  |
|         | 120        | 71(11)     | 5(2)*      | 136(21)  | 14(3)       | 7(3)       | 1.9(0.4)*  | 2.4(0.6)*    | 2959(421)*     | 294(184)       | 344(93)        | 113(23)        | 32(7)   | 67(16)*|
|         | 180        | 67(12)*    | 3(2)*      | 145(18)* | 17(2)*      | 6(1)*      | 1.5(0.4)*  | 2.1(0.6)*    | 3431(749)*     | 583(133)*      | 293(81)*       | 117(24)*       | 39(7)*  | 64(9)* |

MB (acute pancreatitis plus methylene blue group), MBP, CVP, HR, MPAP, PCP, CO, CI, SVR, PVR, DO₂, VO₂, O₂El, SvO₂. Values are present as mean (standard deviation); * p<0.05 versus t (0); † p<0.05 versus CTL; ‡ p<0.05 versus MB; # p<0.05 versus AP.
Table 2 shows the results of serum nitrate arterial and venous and amilase. Serum arterial nitrate decreased continuously (p<0.05) in MB group over time. Serum arterial nitrate increased (p<0.05) in both AP and MB+AP groups when compared to CTL and MB groups at 180 min, respectively.

Serum venous nitrate decreased (p<0.05) in MB group at 180 min. In contrast, serum venous nitrate increased (p<0.05) in AP group at 180 min. Serum venous nitrate increased (p<0.05) in both AP and MB+AP groups when compared to CTL and MB groups at 180 min, respectively. MDA increased (p<0.05) in MB group at 180 min. In the MB+AP group MDA increased (p<0.05) at 120 and 180 min. MDA in MB group was elevated (p<0.05) when compared to CTL group at 180 min. MDA in MB+AP group were elevated (p<0.05) when compared to MB and AP groups at 60, 120 and 180 min. It were also elevated (p<0.05) in AP+MB group when compared to AP group at 120 and 180 min. (Figure 2).

**TABLE 2** - Serum arterial and venous nitrate and serum amylase results in all groups.

<table>
<thead>
<tr>
<th>Time(min)</th>
<th>Arterial Nitrate (mM)</th>
<th>Venous Nitrate (mM)</th>
<th>Amylase (ULL⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>93(56)</td>
<td>97(59)</td>
<td>1507(656)</td>
</tr>
<tr>
<td>60</td>
<td>94(40)</td>
<td>90(38)</td>
<td>1409(616)</td>
</tr>
<tr>
<td>120</td>
<td>91(41)</td>
<td>83(34)</td>
<td>1388(625)</td>
</tr>
<tr>
<td>180</td>
<td>80(38)</td>
<td>75(40)</td>
<td>1371(612)</td>
</tr>
<tr>
<td>CTL</td>
<td>116(41)</td>
<td>112(38)</td>
<td>1690(436)</td>
</tr>
<tr>
<td>MB</td>
<td>108(40)*</td>
<td>104(39)</td>
<td>1576(419)</td>
</tr>
<tr>
<td>AP</td>
<td>99(22)</td>
<td>97(31)</td>
<td>1575(432)</td>
</tr>
<tr>
<td>MB+AP</td>
<td>169(105)</td>
<td>161(97)</td>
<td>1552(434)</td>
</tr>
<tr>
<td>AP+MB</td>
<td>94(11)</td>
<td>93(7)</td>
<td>124(29)*</td>
</tr>
</tbody>
</table>

**TABLE 3** - Serum glucose, creatinine and hematocrit results in all groups.

<table>
<thead>
<tr>
<th>Time(min)</th>
<th>Glucose (mg.dL⁻¹)</th>
<th>Cr (mg.dL⁻¹)</th>
<th>Ht (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>108(15)</td>
<td>0.9(0.1)</td>
<td>30(2)</td>
</tr>
<tr>
<td>60</td>
<td>100(18)</td>
<td>0.9(0.1)</td>
<td>28(2)</td>
</tr>
<tr>
<td>120</td>
<td>93(13)</td>
<td>0.9(0.1)</td>
<td>28(3)</td>
</tr>
<tr>
<td>180</td>
<td>93(8)</td>
<td>0.9(0.1)</td>
<td>29(3)</td>
</tr>
<tr>
<td>CTL</td>
<td>108(30)</td>
<td>1.1(0.2)</td>
<td>33(2)</td>
</tr>
<tr>
<td>MB</td>
<td>76(22)</td>
<td>1.1(0.2)</td>
<td>32(2)</td>
</tr>
<tr>
<td>AP</td>
<td>70(26)*</td>
<td>1.1(0.2)</td>
<td>32(2)</td>
</tr>
<tr>
<td>MB+AP</td>
<td>58(12)*</td>
<td>1.0(0.1)</td>
<td>32(2)</td>
</tr>
<tr>
<td>AP+MB</td>
<td>55(11)*</td>
<td>0.8(0.1)</td>
<td>34(3)</td>
</tr>
</tbody>
</table>

**FIGURE 2** - Malondialdehyde (MDA) relative variation in all groups. CTL (control group), MB (methylene blue group), AP (acute pancreatitis group), MB+AP (methylene blue plus acute pancreatitis group), AP+MB (acute pancreatitis plus methylene blue group). * p<0.05 versus t (0); † p<0.05 CTL versus AP; ‡ p<0.05 AP versus AP+MB; # p<0.05 MB versus MB+AP
Serum amylase levels were elevated (p<0.05) in AP group at 60 min. Serum amylase levels were elevated (p<0.05) in MB+AP and AP+MB groups at 60 and 180 min. There was increased (p<0.05) serum amylase when compared CTL group to AP group at 60 min.

Serum glucose, creatinin and hematocrit results are showed in Table 3. Serum glucose levels decreased (p<0.05) in MB group at 120 and 180 min; in MB+AP group at 60, 120 and 180 min; in AP and AP+MB groups at 60 and 180 min. No difference (p>0.05) were seen among groups.

Creatinin serum levels increased (p<0.05) in all AP groups AP at 180 min, moreover in AP+MB group that increased (p<0.05) at 120 min. Creatinin serum levels were elevated (p<0.05) in MB+AP group when compared to AP group at 180 min.

Hematocrit increased (p<0.05) in all AP groups. At the end of the experiment, there were a mean increase of 32, 26, and 20 % in AP, MB+AP and AP+MB groups, respectively.

Pancreatic morphological study from CTL and MB groups showed normal gland. However, pancreas from all AP groups presented with bright dark reddish surface with black spots, swelled and jelly consistence. Microscopy study in CTL and MB groups was unremarkable but it revealed necrosis and hemorrhage in all AP groups. None leukocytes infiltration was seen.

**Discussion**

MB has been used as a therapeutic drug for more than 110 years mainly for methemoglobinemia treatment, cyanate antidote and fistula localization. In AP group, venous nitrate levels increased over time and arterial nitrate levels were higher when compared with group CTL. This increase was associated with hypotension, CO and PCP decrease and hemoconcentration. In severe AP, inducible nitric oxide synthase (iNOS) overproduces NO. Excessive NO production cause pathological vasodilatation with capillary permeability increase and third space fluid loss and tissue injury by NO derived free-radicals overproduction. AL MUFTI et al. (1998) demonstrated an increase in iNOS activity associated to systemic NO increase and arterial hypotension in AP experimental models. Thus, inhibition of iNOS may have a potential benefic effect in a variety of pathophysiologic alterations in AP.

The findings of MBP, CVP, CO and CI reduction in AP group associated to HR, SVR and PVR increase with hemoconcentration confirm the method reproducibility regarding precocious hypovolemic shock manifestations in this AP experimental model. CO and CI reduction during AP induction is primarily related to hypovolemia and due to preload reduction. Experimental studies had demonstrated that MBP decrease correspond to 26 to 45% circulating volume decrease. In this study it was 32% in AP group.

Our objective was to evaluate initial AP treatment with MB without volemic replacement. When compared AP groups there was a delay in CO and CI reduction in MB+AP group although there were no difference at the end of the experiment. In AP group it initiated at 70 minutes and only at 140 minutes in MB+AP group. However, MB given 90 minutes after AP induction, did not revert the hemodynamic changes. When there is volemic replacement, CO and CI do not decrease in severe AP initial phase and hemoconcentration precedes hemodinamic alteration. Hemoconcentration was present 10 minutes before CO and CI decrease in AP group. Furthermore, in MB+AP group occurred hemoconcentration at least 50 minutes before CO and CI decrease. We observed that MB (dose of 2mg.kg⁻¹.h⁻¹) increased PVR, that also was reported in shock septic patients treated with MB (4mg.kg⁻¹.h⁻¹) MB inhibition on NO vasodilatation and prostaglandins action may explain PVR increase in septic shock and inflammatory conditions where both action are increased differently from normal physiological conditions as seen in MB group.

DO₂ diminished in AP groups although MB immediately before AP induction delayed DO₂ decrease, most probably related to CO decrease. O₂EL increase observed in AP groups reflects AP hypovolemic condition and it suggests a compensation attempt due to DO₂ decrease and VO₂ increase. However, AP patients that had volemic correction with crystalloid solutions present DO₂ and VO₂ increase suggesting that AP increases metabolic demand.

Serum nitrate levels increase was not observed when MB was used either before or after AP induction. An unexpected reduction of arterial and venous serum nitrate levels were observed suggesting inhibition of NO effect independent of GMPc inhibition. However, this effect seems partial because MB did not decrease serum nitrate levels in the MB+AP or AP+MB groups. This mechanism can be due to the production of superoxide, a NO inactivator. Oxidatio of the reduced form of MB induces the generation of this free radical, justifying the findings of MDA increases associated to MB infusion.

Free radicals (FR) generation seems to be involved in AP experimental models correlated to ischemia, alcohol and biliary etiology. However, MDA serum levels did not increased in AP group. Different experimental models of AP seem to present different degrees and mechanisms of FR formation, with lesser impact in taurocholate-induced AP. Adversely, MB promoted increase of MDA serum levels with a significant impact when it was associated with AP. There are evidence that L-NAME induces the RL production MARCZIN et al. demonstrated that, in physiological conditions, MB inactives NO with consequent increase in superoxide concentration and lipoperoxidation. Therefore, it seems that MDA serum increase in group MB corresponds probably to NO inactivation with consequent NO physiological actions blockade. MDA levels increase, associated with the use of MB during AP, seems not to be associated to initial NO production by iNOS. The NO release, in great amounts, demand necessary time since the initial stimulation to its final production that involves cellular enzymatic mechanisms. The NO increase occurred 180 min after AP induction.

NO may be a cytotoxic mediator during taurocholate-induced AP. In fact, NO cytotoxicity is
attributed to its reaction with ion superoxide forming peroxynitrite.35,36 Although peroxynitrite production precedes iNOS expression, there was no MDA increase in AP group.

Initial studies in sepsis had shown that NO overproduction is responsible by hypotension and that iNOS blockage improves it without improving mortality.37,39 The reason is due to no selective iNOS inhibitors, that NO blocking interferes with perfusion regulation, capillary permeability and leukocytes/platelets interactions. A possible explanation for these findings may be that NO blockage by MB improved systemic hemodynamic parameters during AP, although in the cellular level it promoted a poor tissue perfusion due to its vasoconstrictor action. Moreover, MB increases platelets activity in hypovolemic and homoconcentration conditions propitiating lipoperoxidation and tissue perfusion aggravation. Serum amylase levels increase, MB+AP and AP+MB groups, may correspond to suppression of NO beneficial effect in pancreatic microcirculation. NO is a mediator that increases pancreatic blood flow in response to physiological stimulation of CCK and secretin.40 NO may confer protection against AP development, probably due to pancreatic microcirculation improvement and leukocytes adhesion reduction.41-43

Serum creatinin increase in AP groups is in accordance with LEVY et al.44 that demonstrated acute renal insufficiency consequence to hypovolemia in taurocholate-induced AP model. Serum creatinin worsening in AP+MB group confirms MB vasoconstrictor effect on renal circulation thus aggravating a vasoconstriction due to hypovolemia.

Hypoglycemia occurred in all groups except in group CTL. Hypoglycemia in AP groups may be explained by cells beta destruction and insulin release during taurocholate-induction AP. On the other hand, MB group also presented hypoglycemia over time probably due to a blockage of an insulin inhibitory secretion mechanism NO dependent.45,46 There was no improvement in necrohemorrhagic findings with MB treatment.

In this taurocholate-induced AP model we concluded that MB treatment delayed hemodynamic shock and decreases serum nitrate levels but increases serum MDA levels. In our experiment, no volemic replacement was done and it may have been a mitigated factor to a poor tissue perfusion and impairment microcirculation. Further investigations are needed to elucidate MB treatment role during AP treatment.

References

Liver histological, portal flow and plasmatic nitric oxide alterations caused by biliary obstruction and drainage in rats


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Comments:

Current literature data indicates that methylene blue may reverse or ameliorate hemodynamic alterations caused by the systemic inflammatory syndrome through blockage of nitric oxide activity. Nitric oxide is a vasodilator and regulates the exocrine and endocrine (insulin) pancreatic secretion and the microvascular blood flow. The use of NO in acute experimental pancreatitis shows controversial results. Some studies (Werner et al., 1997; Molero et al., 1995) suggest a protective effect probably due to increased blood flow in the pancreatic micro circulation. Waindebach et al. (1995) did not observe any effect of NO in acute pancreatitis, but others (Dabrowiski et al.,1994; Iomis et al., 1995) had evidence of an injurious effect, thought to be due to oxidative stress, vasodilatation and tissue hypoperfusion. In this study, Meirelles et al. proposed the study of hemodynamic profiles and lipid peroxidation in pigs with acute pancreatitis induced by taurocholate enterokinase, and submitted to treatment with methylene blue, as a NO inhibitor, before and after pancreatitis induction. The dye was able to reduce serum nitrate levels, but lipid peroxidation was increased as shown by serum levels of malon dialdehyde. The results in the study warrant the conclusion that the use of methylene blue before induction of severe acute experimental pancreatitis offered a transitory protection to hemodynamic shock. However, there was a deleterious effect on lipid peroxidation and the claim that there is an improvement in circulatory shock by the use of methylene blue should be avoided.

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