Original Article

Efficacy of glucocorticoids in rodents of severe acute pancreatitis: a meta-analysis

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Abstract: Background: The use of corticosteroid in the management of severe acute pancreatitis (SAP) remains contentious and is still being debated despite many pre-clinical studies demonstrating benefits. The limitations of clinical research on corticosteroid in SAP are disparities with regard to benefit, a lack of adequate safety data and insufficient understanding of its mechanisms of action. Thus, we performed a meta-analysis to assess the effectiveness of corticosteroid in experimental SAP and take a closer look at the relation between the animal studies and prospective trials. Methods: Studies investigating corticosteroid use in rodent animal models of SAP were identified by searching multiple three electronic databases through October 2013, and by reviewing references lists of obtained articles. Data on mortality, changes of ascitic fluid and histopathology of pancreas were extracted. A random-effects model was used to compute the pooled efficacy. Publication bias and sensitivity analysis were also performed. Results: We identified 15 published papers which met our inclusion criteria. Corticosteroid prolonged survival by a factor of 0.35 (95% CI 0.21-0.59). Prophylactic use of corticosteroid showed efficacy with regards to ascitic fluid and histopathology of pancreas, whereas therapeutic use did not. Efficacy was higher in large dose and dexamethasone groups. Study characteristics, namely type of steroids, rout of delivery, genders and strains of animal, accounted for a significant proportion of between-study heterogeneity. No significant publication bias was observed. Conclusions: On the whole, corticosteroids have showed beneficial effects in rodent animal models of SAP. Prophylactic use of corticosteroid has failed to validate usefulness in prophylaxis of postendoscopic retrogradcholangiopancreatography pancreatitis. Further appropriate and informative animal experiments should be performed before conducting clinical trials investigating therapeutic use in SAP.

Keywords: Glucocorticoids, acute pancreatitis, animals studies, meta-analysis

Introduction

Acute pancreatitis (AP) is a common clinical condition with variable severity in which some patients experience mild, self-limited attacks while others manifest a severe, highly morbid, and frequently lethal attack [1, 2]. The exact mechanisms by which diverse etiological factors induce an attack are still unclear, while it is generally believed that the release of a variety of inflammatory mediators causes a cascade-like reaction and leads to systemic inflammatory response syndrome (SIRS). An excessive SIRS leads to distant organ damage and multiple organ dysfunction syndrome (MODS), which established the role played by inflammatory mediators in the aggravation of AP and the resultant fatal condition [3-5].

Glucocorticoids have well-known immunosuppressive effects and were commonly believed to be involved in the regulation of cytokine production and the inflammatory processes [6]. Furthermore, glucocorticoids protected the acinar cells, stabilized the cell membrane, and directly regulated amylase synthesis [7, 8]. Some earlier studies supported the beneficial effects of the prophylactic or therapeutic use of glucocorticoids with regard to hemodynamic changes, pancreatic edema formation, histological changes in the pancreas, and even survival [9-11], but others cast doubt on the favorable effects and by contrast, showed the exacerbation of pancreatitis by the administration of glucocorticoids [12-14]. In addition, the possible involvement of steroids in the pathogenesis of acute pancreatitis and considerable side-effects have long been a drawback in the use of this agent in pancreatitis [15, 16]. Therefore, whether or not glucocorticoids are an effective medicine for pancreatitis is a matter of much dispute. Clinically, well-designed
controlled prospective studies have been performed to evaluate the effects of corticosteroid use in prevention of AP after diagnostic or therapeutic endoscopic retrograde cholangiopancreatography (ERCP), by which the designs were compatible to the idea of prophylactic use in animal studies [17-19]. However, the clinical utilization of corticosteroid in pancreatitis has always been controversial and clinical trial for therapeutic purposes in severe acute pancreatitis (SAP) patients with is still scarce due to disparities with regard to beneficial effects and risks, a lack of adequate safety data and insufficient understanding of its mechanisms of action.

Meta-analyses of data from human studies are invaluable resources in the life sciences. Similarly there are a number of benefits in conducting systematic reviews, especially meta-analysis on data from animal studies. They can be used to inform clinical trial design, highlight areas which may benefit from further preclinical research, judge the safety and efficacy of drugs/treatments or provide insights into discrepancies between preclinical and clinical trial results [20, 21]. For discovering the molecular mechanisms underlying AP, rodent models essentially contribute to the understanding of the pathological reaction and are thus often used in assessment of drug efficiency in preclinical research of SAP [22]. Therefore, in this paper, we performed a meta-analysis of the use of corticosteroid in rodent animal models of experimental SAP. The objectives of the present study were to: (1) collate the experimental evidence for glucocorticoids administered before or after on set SAP in rodent animal models and explore whether the contradictory results in animal studies arose from differences in the dose and type of steroids and experimental models et al. (2) assess whether combined results of prophylactic use of corticosteroids in animal studies could revealed indications found in clinical studies; (3) propose the development of further preclinical hypotheses to test in animals and ultimately aid in the design of future clinical trials.

Materials and methods

Search strategy

The following electronic databases were searched for original articles concerning the effects of glucocorticoids on experimental SAP published up to October 2013: PubMed, Embase and Web of Knowledge. The following key words were used to identify possible publications: “glucocorticoids” or “glucocorticoid” or “corticosteroids” or “corticosteroid” or “steroids” or “steroid” or “hydrocortisone” or “hydrocortisone sodium succinate” or “cortisone” or “prednisone” or “prednisone acetate” or “prednisolone” or “prednisolone acetate” or “methylprednisone” or “6alpha-methylprednisolone sodium succinate” or “dexamethasone” or “dexamethasone sodium phosphate” or “betamethasone” or “betamethasone sodium phosphate” and “pancreatitis” or “pancreatitis” or “pancreas” or “pancreatic inflammation”. Search results were limited to animal subjects. No language restriction was used. In addition, we searched for possible eligible studies in the references within the retrieved articles, as well as in review articles.

Criteria for inclusion and exclusion

The selection of studies was performed on the basis of the title and abstract. Two well trained investigators (M.Y. and Z.Y.) independently screened all the abstracts for the inclusion criteria. Differences were resolved by a third investigator (Y.Z.). Studies were included if they fulfilled the following criteria: (1) reported quantitative estimates of the effects of glucocorticoids on mortality, change of volume of ascites and histopathology of pancreas in experimental SAP; (2) number of animals per group was given. Papers were excluded if they fulfilled one of the following criteria: (1) animal studies performed in non-rodent species such as dogs, pigs; (2) Glucocorticoid treatment was combined with co-treatments; (3) glucocorticoid treatment was longer than 5days; (4) studies were specially excluded where in vitro or edematous pancreatitis models were used; (5) double publication or data republication. If duplicate article was published using the same case series, the data from the most informative manuscript was included.

Assessment of risk of bias in included studies

Two reviewers (M.Y. and Z.Y.) independently assessed the associated risk of bias using the 10 criteria (rating: yes, no, unclear) adjusted from recommendation by the Cochrane Back Review Group [23]. When necessary, discrep-
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Articles identified through database searching (n=7145)

Excluded based on the titles and abstracts (n=7083)

Potentially relevant articles screened (n=62)

Excluded articles: Non-roden animals (n=16)
Data republication (n=14)
Steroids treatment longer than 5 days (n=8)
Mild or in vitro acute pancreatitis (n=5)
Combined treatment (n=4)
Insufficient data for meta-analysis (n=2)

Studies included:
Mortality (n=7)
Ascites (n=4)
Histopathology (n=9)

Figure 1. Identification of eligible studies from different databases.

ancies were rechecked with a third reviewer (Y.Z.) and consensus achieved by discussion. The items in the criteria were described in Figure 2, basing on the possible presence of selection bias (items 1, 2 and 3), performance bias (items 5 and 6), detection bias (items 4, 7 and 8) and attrition bias (items 9 and 10) [24]. The “green” indicated low risk of bias, the “red” indicated high risk of bias, “yellow” indicated unknown risk of bias. Studies that met 5 of the 10 criteria and had no serious flaw (such as drop-out rate higher than 30% due to inappropriate manipulation) were rated as having low risk of bias.

Data extraction and data synthesis

From the studies included, the following data were extracted: first author’s last name, year, animals' genders, species and strains, number of animals per group, method of AP induction, type and dose of glucocorticoids, timing of glucocorticoids administration relative to AP induction, route of administration, timing data collection and outcome measures. The patient-important mortality (primary) and surrogate parameters including changes in the volume of ascites and histopathology of pancreas (secondary) were used to measure the efficacy of glucocorticoids. Data were processed as following: for mortality, the Risk Ratio was determined; for the outcome measure “volume of ascites” and “histopathology of the pancreas”, the standardized mean difference (SMD) was calculated. Where a publication reported more than one experiment, we considered these to be independent experiments and extracted data for each of these [25]. When a single control group was used for multiple treatment groups this was adjusted by dividing by the number of treatment groups served (surrogate parameters) [24]. In case histopathological data was not presented in an overall score, we calculated an overall score by uniformly weighing the separate means and SE’s of fibrosis, acinar cell loss etc. For mortality, we merge the events and sample sizes of the treatment group into one group, the same for the control group [24]. If data were not reported, attempts were made to contact the authors for additional information. Where data were only presented graphically, we used digital ruler software (GetData software) to extract.

Statistical analysis

Taking into account both within-study and between-study variabilities, we used the random effects model of inverse variance method which is more conservative than fixed-effects to aggregate data (either OR or SMD) and obtain an overall effect size and 95% confidence interval [26]. Heterogeneity across studies was assessed by performing X² tests (assessing the P value) and calculating I² statistic, a quantitative measure of inconsistency across studies. Studies with an I² of 25% to 50% were considered to have low heterogeneity, I² of 50% to 75% was considered moderate heterogeneity, and I² > 75% was considered high heterogeneity. If I² > 50%, potential sources of heterogeneity were identified by sensitivity analyses conducted by omitting one study in each turn and investigating the influence of a single study on the overall pooled estimate [27].

To explore the impact of study characteristics on estimates of effect size, we performed a stratified meta-analysis with experiments grouped according to the following: timing of corticosteroid treatment (prophylactic or therapeutic), equivalent dosage of dexamethasone. Given consideration of different type of steroids, we used equivalent dosage of dexamethasone to evaluate the impact of dose, using a conversion rate as following: hydrocortisone 20 mg = cortisol 25 mg = prednisone or pred-
# Table 1. Main characteristics of included studies

<table>
<thead>
<tr>
<th>Study</th>
<th>Species</th>
<th>N (c)/n (exp)</th>
<th>Method AP induction</th>
<th>Type and timing of GC</th>
<th>Duration and dose of GC</th>
<th>Admin rout</th>
<th>Timing data collection</th>
<th>Outcome measures</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thomas et al. 1998</td>
<td>male; Rats/sprague-dawley</td>
<td>10/32</td>
<td>Cerulean (30 µg/kg iv) superimposed on 10 mM glycodeloxy-cholic acid (id) for 12 h</td>
<td>Pred; 6 h after AP induction</td>
<td>2 mg/kg; 10 mg/kg; 50 mg/kg/d</td>
<td>Iv</td>
<td>36 h</td>
<td>Mortality; ascites; histological score</td>
</tr>
<tr>
<td>Ramudo et al. 2010</td>
<td>Male; rats/wistar</td>
<td>4/8</td>
<td>Bile-pancreatic duct obstruction (BPDO) for 12 h</td>
<td>Dx, 30 min before and 1 h after AP induction</td>
<td>1 mg/kg</td>
<td>lm</td>
<td>12 h</td>
<td>Histological score</td>
</tr>
<tr>
<td>Ramudo et al. 2010</td>
<td>Male; rats/wistar</td>
<td>8/16</td>
<td>Retrograde infusion of 3.5% sodium taurocholate (id) for 12 h</td>
<td>Dx, 30 min before and 1 h after AP induction</td>
<td>1 mg/kg</td>
<td>lm</td>
<td>3 h, 6 h</td>
<td>Histological score</td>
</tr>
<tr>
<td>Gloor et al. 2001</td>
<td>Female; rats/wistar</td>
<td>8/21</td>
<td>Retrograde infusion of 5% sodium taurocholate (id)</td>
<td>Hc; 10 min after AP induction</td>
<td>10 mg/kg</td>
<td>Iv</td>
<td>1.5 h, 3 h, 6 h, 12 h, 24 h, 72 h</td>
<td>Mortality</td>
</tr>
<tr>
<td>Zhang et al. 2007</td>
<td>Male; rats/SD</td>
<td>45/45</td>
<td>5% sodium taurocholate (1 mg/kg id)</td>
<td>Dx; 15 min after AP induction</td>
<td>5 mg/kg</td>
<td>Iv</td>
<td>3 h, 6 h, 12 h</td>
<td>Mortality; histological score</td>
</tr>
<tr>
<td>Muller et al. 2008</td>
<td>Female; rats/Wistar</td>
<td>6/6</td>
<td>Retrograde infusion of 5% sodium taurocholate (1 mg/kg id)</td>
<td>Dx; 1 h prior to AP induction</td>
<td>1 mg/kg</td>
<td>lm</td>
<td>6 h</td>
<td>Mortality</td>
</tr>
<tr>
<td>Wang et al. 2003</td>
<td>Male; rats/wistar</td>
<td>18/18</td>
<td>Retrograde infusion of 5% sodium taurocholate (id)</td>
<td>Dx; 5 min after AP induction</td>
<td>0.5 mg/kg</td>
<td>Iv</td>
<td>12 h</td>
<td>Mortality; histological score</td>
</tr>
<tr>
<td>Ou et al. 2012</td>
<td>male; rats/SD</td>
<td>36/36</td>
<td>Retrograde infusion of 3.5% sodium taurocholate (id)</td>
<td>Dx; 15 min after AP</td>
<td>5 mg/kg</td>
<td>Iv</td>
<td>3 h; 6 h; 12 h</td>
<td>Mortality</td>
</tr>
<tr>
<td>Kandil et al. 2006</td>
<td>unknown; rats/SD</td>
<td>8/8</td>
<td>Retrograde infusion of 40 g/L sodium taurocholate (id)</td>
<td>Dx; 4 d before AP induction</td>
<td>2 mg/kg (per day), continued for 4 days</td>
<td>Ip</td>
<td>24 h</td>
<td>Histological score</td>
</tr>
<tr>
<td>Paszt et al. 2004</td>
<td>Male; rats/Wistar</td>
<td>12/24</td>
<td>Retrograde infusion of sodium taurocholate (id)</td>
<td>Dx or hydrocortisone; just prior to AP induction</td>
<td>4 mg/kg; 20 mg/kg</td>
<td>Sc</td>
<td>24 h</td>
<td>Mortality</td>
</tr>
<tr>
<td>Richter et al. 1994</td>
<td>Female; rats/Wistar</td>
<td>6/6</td>
<td>Retrograde infusion of trypsin (id)</td>
<td>Pre; 1 h before AP induction</td>
<td>3 mg/kg</td>
<td>Iv</td>
<td>6 h, 24 h, 48 h</td>
<td>Histological score</td>
</tr>
<tr>
<td>Takaoka et al. 2002</td>
<td>Male; rats/Wistar</td>
<td>3/27</td>
<td>Closed duodenal loop induced pancreatitis group (CDL)</td>
<td>Mp; 5 min just before the preparation of CDL</td>
<td>30 mg/kg</td>
<td>Iv</td>
<td>6 h</td>
<td>Aspects; histological score</td>
</tr>
<tr>
<td>Paszt et al. 2008</td>
<td>Male; rats/Wistar</td>
<td>6/6</td>
<td>Administration of L-arginine (ip)</td>
<td>Mp; Just before the induction of AP</td>
<td>30 mg/kg</td>
<td>Sc</td>
<td>24 h</td>
<td>Histological score</td>
</tr>
<tr>
<td>Osman et al. 1999</td>
<td>Mixed; rabbits/New Zealand White</td>
<td>7/10</td>
<td>Chenodeoxycholic bile acid (id)</td>
<td>Hc; 30 min before AP induction</td>
<td>50 mg/kg</td>
<td>Sc+iv</td>
<td>12 h</td>
<td>Mortality; ascites</td>
</tr>
<tr>
<td>Sun et al. 2007</td>
<td>unknown; rats/SD</td>
<td>20/20</td>
<td>5% Chenodeoxycholic bile acid (2 mg/kg id) and literature of pancreatic duct</td>
<td>Hc; 30 min before AP induction</td>
<td>10 mg/kg</td>
<td>Sc+iv</td>
<td>1 h, 3 h, 6 h, 12 h</td>
<td>Aspects; histological score</td>
</tr>
</tbody>
</table>

Abbreviation: SD: sprague-dawley; id: intraductal; ip: intraperitoneal; iv: intravenous; Sc: subcutaneous; Dx: dexamethasone; Hc: hydrocortisone; Pre: prednisone; Pred: prednisolone; Mg: methylprednisone; AP: acute pancreatitis.
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nisolone 5 mg = methylprednisolone 4 mg = dexamethasone 0.75 mg; subsequently we converted these continuous variable into semi-quantitative data: low dose (< 0.5 mg/kg), moderate dose (0.5-1 mg/kg) and large dose (> 1 mg/kg). Additionally, for outcomes with 10 or more experiments included, we further find the source of heterogeneity according to the following: method of AP induction, type of steroids, route of drug delivery, time for outcome measurement, strain and gender of animals used. Publication bias was examined by Egger’s test [28]. Trim and fill analysis was performed to yield an effect of adjusted for funnel plot asymmetry. All of the calculations were conducted by STATA version 12.0 (Stata Corporation, College Station, TX, USA). P values were 2-sided and P < 0.05 was considered statistically significant.

Results

Description of the included studies

Based on our search criteria, we identified 7145 articles from PubMed, EMBASE and Web of Knowledge or by hand-searching. According to the search strategy described above mentioned, we retrieved 62 papers that seemed to meet our selection criteria. After studying the full-text articles, 47 were excluded, among which 16 were excluded for non-rodent animals used, 14 for data republication, 6 for long-term corticosteroid treatment (> 5 days), 5 for mild or in-vitro AP models, 4 for co-treatments, 2 for unavailable quantitative data. Finally, a total of fifteen articles with approximately 500 animals were included in this meta-analysis [29-43] (Figure 1). The study characteristics varied considerably between the included papers (Table 1). Male rat models (Wistar, Sprague-Dawley) [29-31, 33, 35, 36, 38-40] were used in most of the 15 studies, and female gender rats were used in 3 studies [32, 34, 43], while the gender of remaining studies were not mentioned [37, 41, 42]. Most studies used rats [29-40, 42, 43] and one performed in rabbits [41]. Nine different techniques were used to induce AP. Within these, most studies induce AP through retrograde infusion of sodium taurocholate into bile pancreatic duct [31-38]; 2 studies by retrograde infusion of chenodeoxycholic superimposed on cerulean (iv) or litigation of pancreatic duct [29, 42]; 2 by bile-pancreatic duct obstruction (12 h) or closed duodenal loop [30, 39]; 1 study via purely infusion of glycodeoxycholic acid [41]; 1 by retrograde infusion of trypsin (id) [43], and the remaining one used L-arginine (ip) [40]. Also the administration time and delivery routes, the dosage and the type of steroids varied greatly between the studies. Seven studies reported mortality, four including 9 experiments presented volume of ascitic fluids and nine studied histopathology of pancreas could be pooled in the final analysis (Table 1).

Quality of reporting and risk of bias

Figure 2 showed the overall results of the risk of bias assessment of the 15 studies included in this meta-analysis. 73% of the studies stated...
Figure 3. Forest plot of effects of glucocorticoids supplementation on mortality in experimental acute pancreatitis. Forest plot of the data of seven included studies. The forest plot displays the OR, 95% confidence interval and relative weight of the individual studies. The diamond indicates the global estimate and its 95% confidence interval.
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Figure 4. Forest plot of the effects of glucocorticoids supplementation on volume of ascites in experimental acute pancreatitis. The forest plot displays the SMD, 95% confidence interval and relative weight of the individual studies. The diamond indicates the global estimate and its 95% confidence interval.
that the allocation of the experimental units to the treatment groups was randomized. However, none of these studies mentioned the method of randomization used and only one provided sufficient details so that the adequacy of the method could be judged. None of the papers described whether or not the allocation to the different groups during the randomization process was concealed. 53% of the studies reported that they blinded the outcome assessment. Figure 2 showed that only four out of the 15 studies scored 5 out of the 10 items as low risk of bias. Many items were scored as “unclear risk of bias”, but few studies were deemed fatally flawed, indicating reliability of our results to some extent.

Effects of glucocorticoids

Mortality: Seven publications studied the effect of glucocorticoids administration on mortality in experimental SAP. A combination of all the animal studies included showed a significant effect on reduced mortality (RR 0.35 [0.21, 0.59]; n = 7). No significant heterogeneity was found (P = 0.71; I² = 0.0%) even though a study performed in rabbits was included in this meta-analysis. Comparison of the effects of glucocorticoids administration before or after inducing SAP on the risk of mortality revealed reduced mortality in both groups (before; RR 0.46 [0.24, 0.88]; n = 3; after; RR 0.24 [0.1, 0.54]; n = 4). Subgroup analyses on the study characteristic “equivalent dosage of dexamethasone” also showed improvement in mortality regardless of the dosage (large RR 0.38 [0.22, 0.65]; n = 6; low RR 0.15 [0.02, 0.92]; n = 1) (Figure 3).

Ascites: Four papers including 9 experiments reported change of pancreatic ascites production due to corticosteroid treatment. Overall analysis showed that corticosteroid significantly reduced the volume of ascites compared to controls (SMD = -2.53 [-3.76, -1.29]; n = 9; P = 0.000). Heterogeneity was high (P = 0.000; I² = 76.9%) and remained moderate after eliminating the study performed in rabbits (P = 0.04; I² = 52.3%) (Figure 4).

Histopathology of the pancreas

Nine out of 17 papers could be included in quantitative analysis, which covered 18 comparisons in total. Overall analysis showed that glucocorticoids can reduced/improved the histopathology of the pancreas (SMD = -0.86 [-1.40, -0.31]; n = 18; P = 0.002) (Figure 5). Heterogeneity was high (P = 0.000, I² = 62.4%). Subgroup analysis revealed that prophylactic corticosteroid use significantly decrease the pathological score, whereas no significant change was observed in therapeutic use group (before; SMD = -1.23 [-2.12 -0.34]; n = 9; P = 0.007; after; SMD = -0.54 [-1.28, 0.20]; n = 9; P = 0.152). Taking into account of dosage, large dose group significantly ameliorated histopathology, both low and moderate groups did not (low; SMD = -0.39 [-0.91, 0.13]; n = 6; P = 0.14; moderate; SMD = -1.48 [-3.13, 0.17]; n = 6; P = 0.078; large; SMD = -0.94 [-1.61, -0.27]; n = 6; P = 0.021).

Subgroup analyses also revealed that the significant efficacy was only observed in dexamethasone group (dexamethasone; SMD = -2.01 [-3.42, 0.6]; n = 8; P = 0.005; hydrocortisone; SMD = -0.36 [-0.99, 0.27]; n = 4; P =

<table>
<thead>
<tr>
<th>Volume of ascites</th>
<th>SMD</th>
<th>LL</th>
<th>UL</th>
<th>n</th>
<th>P</th>
<th>p of heterogeneity</th>
<th>I²</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>-2.53</td>
<td>-3.76</td>
<td>-1.29</td>
<td>9</td>
<td>0.000</td>
<td>0.000</td>
<td>76.90%</td>
</tr>
<tr>
<td>Timing of treatment</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prophylactic (before)</td>
<td>-3.29</td>
<td>-5.03</td>
<td>-1.54</td>
<td>6</td>
<td>0.000</td>
<td>0.000</td>
<td>81.30%</td>
</tr>
<tr>
<td>Therapeutic (after)</td>
<td>-1.32</td>
<td>-2.72</td>
<td>0.08</td>
<td>3</td>
<td>0.065</td>
<td>0.124</td>
<td>52%</td>
</tr>
<tr>
<td>Equivalent dosage of Dx</td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Large</td>
<td>-4.35</td>
<td>-7.11</td>
<td>-1.59</td>
<td>4</td>
<td>0.002</td>
<td>0.000</td>
<td>84.90%</td>
</tr>
<tr>
<td>low</td>
<td>-1.53</td>
<td>-2.5</td>
<td>-0.56</td>
<td>5</td>
<td>0.002</td>
<td>0.113</td>
<td>46.50%</td>
</tr>
</tbody>
</table>

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Figure 3

Figure 4

Figure 5
Glucocorticoids and severe acute pancreatitis

<table>
<thead>
<tr>
<th>Study</th>
<th>SMD (95% CI)</th>
<th>Weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zhang et al, 2007</td>
<td>-1.05 (-1.49, -0.61)</td>
<td>9.89</td>
</tr>
<tr>
<td>Thomas et al, 1998</td>
<td>0.00 (-1.55, 1.55)</td>
<td>5.68</td>
</tr>
<tr>
<td></td>
<td>0.19 (-1.14, 1.52)</td>
<td>6.46</td>
</tr>
<tr>
<td></td>
<td>0.00 (-1.57, 1.57)</td>
<td>5.60</td>
</tr>
<tr>
<td>Sun W, 2007</td>
<td>-0.57 (-1.84, 0.70)</td>
<td>6.69</td>
</tr>
<tr>
<td></td>
<td>-0.21 (-1.46, 1.03)</td>
<td>6.80</td>
</tr>
<tr>
<td></td>
<td>-0.47 (-1.74, 0.79)</td>
<td>6.73</td>
</tr>
<tr>
<td></td>
<td>-0.19 (-1.43, 1.05)</td>
<td>6.80</td>
</tr>
<tr>
<td>Ramudo et al, 2010 (NaTo)</td>
<td>-0.87 (-2.67, 0.93)</td>
<td>4.88</td>
</tr>
<tr>
<td></td>
<td>1.16 (-0.72, 3.04)</td>
<td>4.66</td>
</tr>
<tr>
<td></td>
<td>-4.07 (-7.37, -0.78)</td>
<td>2.16</td>
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<td></td>
<td>-0.07 (-1.77, 1.63)</td>
<td>5.19</td>
</tr>
<tr>
<td>Ramudo et al, 2010 (BPDO)</td>
<td>-3.85 (-7.02, -0.69)</td>
<td>2.30</td>
</tr>
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<td></td>
<td>-3.40 (-6.31, -0.50)</td>
<td>2.62</td>
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<tr>
<td>Kandil et al, 2006</td>
<td>-7.13 (-9.94, -4.31)</td>
<td>2.75</td>
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<tr>
<td>Richter et al, 1995</td>
<td>-0.74 (-1.92, 0.44)</td>
<td>7.06</td>
</tr>
<tr>
<td>Tokaoka et al, 2002</td>
<td>-0.54 (-1.57, 0.50)</td>
<td>7.64</td>
</tr>
<tr>
<td>Paszt et al, 2008</td>
<td>-1.99 (-3.42, -0.56)</td>
<td>6.10</td>
</tr>
<tr>
<td>Overall</td>
<td>(I-squared = 62.4%, p = 0.000)</td>
<td>-0.86 (-1.40, -0.31)</td>
</tr>
</tbody>
</table>

NOTE: Weights are from random effects analysis
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When a single study involved in the meta-analysis was deleted each time, the results of meta-analysis remained unchanged, indicating that the results of the present meta-analysis were stable (data not shown). There was no statistical evidence of publication bias among studies for both mortality and histopathology of pancreas by using Egger’s test ($P = 0.17$; $P = 0.44$, respectively). For volume of ascites, Egger’s test revealed significant publication bias ($P = 0.001$). The overall conclusion indicated no overestimation after Trim and Fill analysis filled with two studies (SMD $= -3.29 [-4.7, -1.89]$; $n = 11$; $P = 0.000$).

Discussion

The results of this meta-analysis showed the beneficial effects of glucocorticoids with regard to the primary outcome measure survival in animal studies of SAP whenever these agents were employed prophylactically or therapeutically. When it came to other outcome measures (changes of pancreatic ascites and histopathology of pancreas), overall analysis also showed a positive effect, however, subgroup analyses indicated that the improvement was not statistically significant in therapeutic treatment group. Furthermore, efficacy seemed to be influenced by dosage and type of steroids, strains or genders, drug delivery routes et al.
and part of those characteristics accounted for a quite considerable proportion of between-study heterogeneity.

Despite the improvements in intensive care and surgical therapy, the mortality rate for acute necrotizing pancreatitis remains high [44]. The clinical presentation may range from a local inflammatory process to a severe pancreas injury associated with extra-pancreatic manifestations such as circulatory, renal or pulmonary complications [45, 46]. Activation of the inflammatory cascade mediated by various inflammatory mediators is the key causes of acute pancreatitis aggravation, eventually results in microcirculatory disturbances, multiple organ failure and even death [46]. Glucocorticoids are potent anti-inflammatory drugs, both directly and indirectly attenuating the synthesis, release, and action of inflammatory cytokines and certain other mediators [6]. It is therefore assumed that glucocorticoids could alleviate the severity of AP in terms of its inhibitory role in excessive inflammatory reactions. Among our included studies, many indicated that exogenous glucocorticoids significantly improved both the local pancreatic inflammatory response as well as systemic inflammatory parameters including the serum concentrations of tumor necrosis factor-alpha (TNF-alpha), interleukins (ILs), platelet-activating factor and C-reactive protein et al. [30, 32, 35, 38, 41]. Furthermore, glucocorticoids were found to attenuate the pancreas damage by protecting acinar cells during cerulein-induced AP [7]. In the Journal of Gastroenterology, Takaoka et al. [39] have very clearly shown that in rats with SAP induced by a closed duodenal loop, the intravenous administration of methylprednisolone (30 mg/kg) alleviated the severity of the acute pancreatitis in terms of pancreatic edema formation and elevation of serum amylase, as well as alleviating its complications the production of ascites through vascular permeability.

Our results showed that corticosteroid is effective at reducing pancreatic ascites production and histopathological score of pancreas and improving survival in experimental models of SAP. It is noteworthy in our study that the short-term pre-treatment with glucocorticoids resulted in both reducing pancreatic ascites and ameliorating histopathology of pancreas, however, post-treatment proved to be ineffective to hinder pancreatic damage and pancreatic ascites production. A plausible mechanism by which may explain the observation was the tight association between inflammatory response and severity of AP. Prophylactic treatment can limit both the local and systemic inflammatory response at the very earliest phase, which resulted in a beneficial effect on the progression of the disease in all directions. Therapeutic treatment just partly blocked inflammatory responses, so the improvement were not statistical significant. However, longer survival has been achieved, demonstrating its beneficial potential in overall prognosis. Taking the dose and type of steroids into consideration, less ascites and improved histopathology of pancreas was found in large dose and dexamethasone groups. The anti-inflammatory potency of glucocorticoid compound was different from each other, it was quite reasonable to find that dexamethasone (with stronger anti-inflammatory potency) showed a positive effectiveness whereas cortisolone, hydrocortisone or prednisone (with weaker anti-inflammatory potency) did not in this setting, so did the large dose of steroids.

The design of prophylactic use in an animal study could be clinically relevant with studies which evaluated the preventive role of corticosteroids in post-ERCP pancreatitis. In light of the results presented above-mentioned strong evidence for efficacy of prophylactic use, it was defensible and understandable that the clinical trials were started [17, 18]. However, Giovanni et al. [47] showed in a controlled prospective study involving 529 patients, that 100 mg of hydrocortisone did not significantly affect acute pancreatitis after diagnostic or therapeutic-ERCP, and the latest meta-analysis suggested that prophylactic corticosteroid use cannot prevent pancreatic injury after ERCP and their use in the prophylaxis of post-ERCP pancreatitis is not recommended [19]. Nearly all interventions which have shown promise in preclinical studies have failed to translate successfully to the clinical setting. In our study, lack of correspondence between animal data and results from clinical trials might be explained below: first of all, glucocorticoids are potent anti-inflammatory drugs and can improved inflammatory mediated pancreas damage, but the beneficial effect itself is not strong enough to improve the
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final clinical results. More importantly, clinical trials mostly used hydrocortisone and sex in their groups was balanced, whereas most of our studies used dexamethasone and male Wistar rat models were more common. When the study was subdivided by type of steroids and sex of rodents, we found no significant relationship between corticosteroid use and improved histopathology of pancreas in hydrocortisone and female or sex-mixed groups. From the point of view, the results in animal studies might correspond well to clinical studies. Additionally, animal studies are prone to bias, mostly overestimation of the treatment [48]. Therefore, contradictory results might arise from lack of control of bias and more appropriate animal experiments in relation to the design of human studies.

The utilization of steroid for therapeutic purposes in the patients with acute pancreatitis is still controversial, and the effectiveness has been suspected for many years. Case reports from several decades ago have suggested a potential clinical benefit of hydrocortisone in acute hemorrhagic pancreatitis [49]. However, the available data with positive results about steroid in acute pancreatitis up to now were mostly experimental studies [33]. In our meta-analysis, the efficacy was not statistically significant in therapeutic treatment group with regard to ascitic fluid production and histopathology of pancreas, whereas better survival was observed, indicating the possible usefulness of the steroid therapy for SAP. It might be worthwhile to investigate this point further clinically. However, the beneficial findings of prophylactic steroids use obtained in rodent animal studies were not validated in interventional clinical studies. Therefore, we assumed that therapeutic corticosteroid use in SAP could also lead to no benefit in clinical application, not only because of the previous inconsistency but also for its relatively lower efficacy even in animal studies. Even for prolonged survival, we should interpret with caution because of data collection time in animal studies. Corticosteroid significantly improved recent mortality in animal studies, within these, the longest observation time was 72 hours, in other word, effects of corticosteroid on late mortality was not investigated at all. However, SAP is characterized by completely different two phases. The early phase is associated with SIRS, multiple organ damage and early mortality, the late phase is characterized by infections; so whether corticosteroid improve overall survival need to be elucidated by further experimental studies. Furthermore, as known to all of us, corticosteroid has unpleasant side effects, even induction of AP, whereas no safety data in these animal studies has been discussed, so further appropriate and informative animal experiments should be performed before conducting clinical trials.

Taking consideration of the meta-analysis itself, there are limitations to our approach. First, although our search strategy was designed to be robust, we cannot rule out the possibility of missing studies. Second, the heterogeneity among the various animal studies was quite considerable. We performed a random effect meta-analysis in order to minimize the risk of finding erroneous estimates offering more consistent results. Our analysis was based on study mean effects, as we did not have access to individual data. Study characteristics actually accounted for a quite considerable proportion of between-study heterogeneity, but the results of these sensitivity and stratified analyses should be interpreted with caution for small number of experiments in some subgroups interactions. Furthermore, only half of the studies reported that they blinded the outcome assessment and only one study provided sufficient details to judge the adequacy of the method of randomization, so the methodological quality of the individual animal studies urgently need to be improved in order to increase the potential value of animal studies as a preparation for clinical applications.

To the best of our knowledge, this is the first meta-analysis to evaluate the effect of glucocorticoids in rodent animal models. This meta-analysis has demonstrated the beneficial effects of glucocorticoids with regard to the primary outcome measure survival irrespective of administration before or after AP induction. In addition, significant effectiveness has been obtained on both reduced histopathology of pancreas and ascitic fluid production for prophylactic use before AP induction, whereas therapeutic use after AP induction did not. For prophylactic use, contradictory results might arise from lack of more appropriate animal experiments in relation to the design of human studies. Further appropriate and informative
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animal experiments should be performed before conducting clinical trials.

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Disclosure of conflict of interest

None.

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