



## ORIGINAL

# Levosimendan in patients with cardiogenic shock complicating myocardial infarction: A meta-analysis



M. Fang<sup>a,b</sup>, H. Cao<sup>a</sup>, Z. Wang<sup>a,\*</sup>

<sup>a</sup> Department of Intensive Care Medicine, 3rd Hospital of HeBei Medical University, Shi Jiazhuang, China

<sup>b</sup> Department of Anaesthesia and Intensive Care, IRCCS San Raffaele Scientific Institute, Milan, Italy

Received 7 March 2017; accepted 30 August 2017

Available online 7 November 2017

### KEYWORDS

Levosimendan;  
Mortality;  
Myocardial infarction;  
Cardiogenic shock

### Abstract

**Purpose:** Cardiac shock is the leading cause of death in patients with acute myocardial infarction. The objective of this systematic review and meta-analysis was to evaluate whether levosimendan, compared to any type of control, is associated with improved clinical outcomes in patients with cardiogenic shock complicating myocardial infarction.

**Materials and methods:** The PubMed, EMBASE, Cochrane Central Register, and China National Knowledge Information databases were searched for pertinent studies published up until 1 May 2016. Randomized and non-randomized clinical trials comparing levosimendan to standard therapy or placebo, in adult patients with cardiogenic shock complicating myocardial infarction, and reporting at least one outcome of interest were included. The primary outcome was mortality, whereas secondary outcomes were length of ICU stay, SOFA score, cardiac index (CI), cardiac power index (CPI), ejection fraction (EF), end-systolic volume (ESV), mean blood pressure (MBP), pulmonary arterial pressure (PAP), mixed venous oxygen saturation (SvO<sub>2</sub>), pulmonary artery occlusion pressure (PAOP) and glomerular filtration rate (GFR). We pooled risk ratio (RR) and 95% confidence interval (CI) using fixed and random effects models.

**Results:** Thirteen studies comprising a total of 648 patients were included in the analysis. There was a nonsignificant reduction in mortality with levosimendan compared to the controls (RR=0.82 [0.65–1.01], *P* for effect=0.07, *I*<sup>2</sup>=0%). In the levosimendan group PAP and ESV were significantly reduced, while CI, CPI, EF, MBP and SvO<sub>2</sub> were significantly increased. No differences in SOFA score, ICU days, PAOP or GFR were noted.

**Conclusions:** Levosimendan can improve hemodynamic parameters and cardiac function when compared with a control group, with no evidence of benefit in terms of survival.

© 2018 Elsevier España, S.L.U. y SEMICYUC. All rights reserved.

\* Corresponding author.

E-mail address: [18533112993@126.com](mailto:18533112993@126.com) (Z. Wang).

<https://doi.org/10.1016/j.medin.2017.08.009>

0210-5691/© 2018 Elsevier España, S.L.U. y SEMICYUC. All rights reserved.

**PALABRAS CLAVE**

Levosimendán;  
Mortalidad;  
Infarto de miocardio;  
Choque cardiogénico

**Levosimendán en pacientes con choque cardiogénico que complique un infarto de miocardio: metaanálisis****Resumen**

**Objetivo:** El choque cardiogénico es la principal causa de muerte en pacientes con infarto agudo de miocardio. El objetivo de este metaanálisis y revisión sistemática fue evaluar si, en comparación con cualquier tipo de control, el levosimendán se asocia a mejores efectos clínicos en pacientes con choque cardiogénico que complique un infarto de miocardio.

**Materiales y métodos:** Se realizaron búsquedas en las bases de datos PubMed, EMBASE, Cochrane Central Register y China National Knowledge Information para encontrar estudios pertinentes publicados hasta el 1 de mayo de 2016. Se incluyeron ensayos clínicos aleatorizados y no aleatorizados en los que se comparase el levosimendán con el tratamiento estándar o con un placebo en pacientes adultos con choque cardiogénico que complicase un infarto de miocardio, y que informasen sobre al menos una variable de interés. La variable principal fue la mortalidad, mientras que las variables secundarias fueron la duración del ingreso en la UCI, la puntuación SOFA, el índice cardíaco (IC), el índice de potencia cardíaca (IPC), la fracción de eyección (FE), el volumen sistólico final (VSF), la presión arterial media (PAM), la presión arterial pulmonar (PAP), la saturación venosa mixta de oxígeno (SvO<sub>2</sub>), la presión de oclusión de la arteria pulmonar (POAP) y la tasa de filtración glomerular (TFG). Agrupamos la razón de riesgos (RR) y el intervalo de confianza (IC) del 95% por medio de modelos de efectos fijos y aleatorios.

**Resultados:** Se incluyeron en el análisis 13 estudios que incluyeron un total de 648 pacientes. Se observó una reducción no significativa de la mortalidad con levosimendán en comparación con los controles (RR = 0,82 [0,65-1,01], p para efecto = 0,07, I<sup>2</sup> = 0%). En el grupo de tratamiento con levosimendán, la PAP y el VSF se vieron reducidos de forma significativa, mientras que el IC, el IPC, la FE, la PAM y la SvO<sub>2</sub> aumentaron de forma significativa. No se observaron diferencias en la puntuación SOFA, los días de ingreso en la UCI, la POAP ni la TFG.

**Conclusiones:** El levosimendán puede mejorar los parámetros hemodinámicos y la función cardíaca en comparación con un grupo de control, si bien no existen evidencias de beneficios en términos de supervivencia.

© 2018 Elsevier España, S.L.U. y SEMICYUC. Todos los derechos reservados.

**Introduction**

Cardiogenic shock complicates approximately 5% of myocardial infarctions. Consequent marked hypotension, reduced oxygen supply, and inadequate perfusion of various organs can result in multiple organ dysfunction. Despite utilization of an early revascularization strategy and advancing patients' care, cardiogenic shock remains the leading cause of death in this population with high hospital mortality rate, approaching 50%.<sup>1,2</sup>

In cardiogenic shock complicating myocardial infarction early revascularization of the occluded vessel with a restoration of coronary cardiac blood flow preferably by means of PCI is the first line strategy. A supportive approach is to give mechanical support, such as Extracorporeal Membrane Oxygenation (ECMO).<sup>3</sup> For inotropic support in patients with cardiogenic shock the drugs of choice is dobutamine. However, mortality of such patients in cardiogenic shock remains high. Further studies are needed to evaluate new therapeutic approaches to decrease mortality and morbidity of these patients.

Levosimendan is a relatively novel inotropic agent, which acts on cardiac troponin C, stabilizing the bound Ca<sup>2+</sup>, prolonging the interaction between actin and myosin, and thus

enhances cardiac contractility.<sup>4</sup> It is a calcium-sensitizer agent<sup>5</sup> with vasodilatory properties,<sup>6</sup> exerting beneficial effects particularly in cardiac surgery, a setting where it recently showed a survival benefit when compared with dobutamine.<sup>7</sup>

More recently, a number of clinical trials have now been completed. Therefore, the principal objective of this study was to critically review the literature to evaluate whether levosimendan compared to standard therapy, in patients with cardiogenic shock complicating myocardial infarction, is associated with improved clinical outcomes, in particular survival, and hemodynamics.

**Materials and methods****Search strategy**

Appropriate studies were independently searched in BioMed Central, PubMed, EMBASE, Cochrane Central Register of clinical trials, and Chinese database (CNKI, WANGFANG DATA, and CQVIP) by 2 investigators. The full PubMed search strategy is available in the Appendix A. Supplementary data. No language restriction was enforced. The search was final-

ized on 1st May 2016. We decided to use a basic search strategy in order to make the strategy as sensitive as possible.

Abstracts from recent international conferences were searched for additional relevant studies. In addition, we use backward snowballing (i.e. scanning of references of retrieved articles and pertinent reviews). The search strategy aimed to include any controlled study with levosimendan administration in cardiogenic shock complicating myocardial infarction in adult humans.

## Study selection

Two authors reviewed all abstracts to identify potentially eligible controlled trials. If it was possible, full text articles were retrieved and reviewed to determine whether they met the eligibility criteria. If the complete paper was not available in the database, the corresponding author was contacted to get further material. Disagreements between reviewers were resolved by consensus.

The inclusion criteria was reports of controlled trials, comparing levosimendan to any other therapy for cardiogenic shock in adult human and reported at least one outcome of interest. The primary outcome was mortality, whereas secondary outcomes were length of ICU stay, Sequential Organ Failure Assessment (SOFA) score, cardiac index (CI), cardiac powder index (CPI), ejection fraction (EF), end systolic volume (ESV), mean blood pressure (MBP), pulmonary atrial pressure (PAP), mixed venous oxygen saturation (SvO<sub>2</sub>), glomerular filtration rate (GFR), and pulmonary artery occlusion pressure (PAOP). The time points of the collection of these variables should follow what reported by authors. There were no restrictions on time or dose of administration. The exclusion criteria were: duplicate publications, pediatric studies, and non-intravenous administration of levosimendan.

## Data abstraction and study characteristics

Two investigators abstracted baseline, procedural, and outcome data by using a data-recording table developed for this purpose. Data collected included: patient baseline characteristics, study design, sample size, clinical setting, study definition of cardiogenic shock, details of levosimendan and control regimens, and clinical outcomes.

## Internal validity and risk of bias assessment

The internal validity of each randomized controlled trial (RCT) was critically assessed for bias as reported by the Cochrane Collaboration methods.<sup>8</sup> Each report was evaluated for risk of bias associated with the random sequence generation method, allocation concealment, blinding of participants and personnel, completeness of outcome data, free of selective reporting, and other bias. The overall risk of bias was presented as low, unclear, or high.

The internal validity of each non-randomized controlled trial (nRCT) was critically assessed for bias as reported by the Newcastle–Ottawa Scale Risk assessment for case-control studies.<sup>9</sup> Each article was evaluated for risk of bias

associated with selection, comparability, and exposure. We rated the risk of bias by applying a rating of star number to determine whether adequate measures were taken to protect against each potential source of bias in each study. The overall risk of bias was presented as star number.

## Data analysis and synthesis

To analyze the binary outcome, we calculated risk ratio (RR) and 95% confidence interval. Mean difference (MD) and 95% confidence interval were computed for continuous variables. Furthermore, we compute numbers needed to be treated. Heterogeneity was measured using the Cochrane *Q* test, quantified with *I*<sup>2</sup> statistic (*I*<sup>2</sup> > 25% was defined as threshold indicating significant heterogeneity), and Tau-square (Tau<sup>2</sup>). The primary analysis was conducted by means of the Peto fixed effects method when *I*<sup>2</sup> < 25% and with the random effects model when *I*<sup>2</sup> > 25%. Publication bias was evaluated by visually inspecting funnel plot of the primary outcome.

Statistical significance was set at the 2-tailed 0.05 levels for hypothesis testing. Data analysis was performed using Review Manager (RevMan, version 5.3. Copenhagen: The Nordic Cochrane Centre, The Cochrane Collaboration, 2014).

## Results

### Literature search

There were 1137 reports identified by the search, 43 full or abstract articles were retrieved for in depth review (Appendix A. Supplementary data). Finally, 13 studies enrolling 648 participants fulfilled all eligibility criteria. There are 5 RCTs with 254 patients<sup>10–14</sup> and 8 nRCTs with 394 patients.<sup>15–22</sup> The trial characteristics are shown in Table 1 and in Appendix A. Supplementary data. There were 11 studies in English,<sup>10–12,15–22</sup> two in Chinese.<sup>13,14</sup>

Study quality and risk of bias are reported in the Appendix A. Supplementary data. Two RCTs<sup>10,12</sup> were rated as high risk of bias, 2 RCTs<sup>13,14</sup> at unclear risk, and 1 RCT<sup>11</sup> at low risk of bias, according to Cochrane Collaboration methods. Six nRCTs were rated as medium risk of bias,<sup>16–20,22</sup> 2 nRCTs<sup>15,21</sup> were rated as high risk bias, according to Newcastle–Ottawa Scale.

### Levosimendan in patients with cardiogenic shock: mortality

The use of levosimendan in patients with cardiogenic shock was associated with a non significant reduction in mortality at the longest follow-up available (68/187 [36%] in the levosimendan group and 121/270 [53%] in the control group, RR 0.82[0.65,1.01], *P* for effect=0.07, *P* for heterogeneity=0.88, *I*<sup>2</sup>=0%, Tau<sup>2</sup> 0.00, numbers needed to treat = 11; Appendix A. Supplementary data), with 11 studies included. No publication bias was present (Appendix A. Supplementary data).

When including only RCTs, there was no statistically significant reduction in mortality at the longest follow-up

**Table 1** Characteristics of the studies included in the meta-analysis.

Study	Year	Population	N	RCT (Y/N)	Levo bolus (µg/kg)	Levo infusion (µg/kg/min)	Levo duration (h)	Control group	Duration of follow-up (days)
Christoph	2008	CS with STEMI and PCI	22	N	12	0.1	24	IABP	In-hospital
Fuhrmann	2008	CS with AMI	32	Y	12	0.2	24	Enoximone	30
Samimi-Fard	2008	CS with STEMI and PCI	22	Y	24	0.1	24	Dobutamine	360
Soos	2009	CS with STEMI and PCI	106	N	NA	0.1	6	Controlled therapy	200
Omerovic	2010	CS with STEMI	94	N	12	0.1	24–48	Levosimendan contraindicated cohort	30
Poli	2011	CS with STEMI	43	N			24	Dobutamine	In-hospital
Caetano	2012	CS with AMI	37	N				Controlled therapy	In-hospital
Husebye	2013	Subgroup:CS with STEMI	9	Y	NA	0.1	25	Placebo	180
Affronti	2013	CS with AMI and ECMO	17	N	NA	0.1–0.2	24–48	Catecholamine	In-hospital
Katsytadze	2013	CS with AMI	27	N				Necessary conservative treatment	360
Mancone	2013	CS with STEMI and PCI and IABP	48	N	NA	0.05–0.2	24	Controlled therapy	30
Luo	2013	CS with STEMI	83	Y	12	0.1–0.2		Dobutamine	In-hospital
Li	2015	CS with AMI	108	Y	12	0.1–0.2	24	Dobutamine	14

CS, cardiogenic shock; STEMI, ST elevation myocardial infarction; PCI, percutaneous coronary intervention; AMI, acute myocardial infarction; ECMO, extracorporeal membrane oxygenation; IABP, intra-aortic balloon pump; Levo, levosimendan.

available in patients with cardiogenic shock (11/37 [29.7%] in the levosimendan group and 14/39[35.9%] in the control group, RR 0.82 [0.43,1.56], *P* for effect=0.55, *P* for heterogeneity=0.21, *I*<sup>2</sup>=37%, Tau<sup>2</sup> 0.00, numbers needed to treat=3 Appendix A. Supplementary data); with 3 studies included.

Even when including only nRCTs, there was no statistically significant reduction in mortality at the longest follow-up available in patients with cardiogenic shock (57/150 [38%] in the levosimendan group and 107/231[45.0%] in the control group, RR 0.81 [0.65,1.03], *P* for effect=0.08, *P* for heterogeneity=0.96, *I*<sup>2</sup>=0%, Tau<sup>2</sup> 0.00, numbers needed to treat=8, Appendix A. Supplementary data); with 8 studies included (Fig. 1).

### Levosimendan in patients with cardiogenic shock: SOFA

Two studies<sup>10,17</sup> reported SOFA that was not significant difference between levosimendan group and control group

(MD −1.87[−3.92, 0.18], *P* for effect=0.07, *P* for heterogeneity=0.03, *I*<sup>2</sup>=72%, Tau<sup>2</sup> 1.74; Table 2 and Appendix A. Supplementary data).

### Levosimendan in patients with cardiogenic shock: cardiac function

Four studies<sup>10,12,17,21</sup> reported Cardiac Index that was significantly higher in the levosimendan group when compared with the control group (MD 0.17[0.06, 0.29], *P* for effect=0.003, *P* for heterogeneity=0.38, *I*<sup>2</sup>=3%, Tau<sup>2</sup> 0.00; Appendix A. Supplementary data).

Three studies<sup>10,12,17</sup> reported Cardiac Power Index (CI\*MAP\*0.0022) that was significantly higher in the levosimendan group when compared with the control group (MD 0.08[0.03, 0.13], *P* for effect=0.003, *P* for heterogeneity=0.08, *I*<sup>2</sup>=60%, Tau<sup>2</sup> 0.00; Appendix A. Supplementary data).

Four studies<sup>12,14,21,22</sup> reported Ejection Fraction that was significantly higher in the levosimendan group, when com-

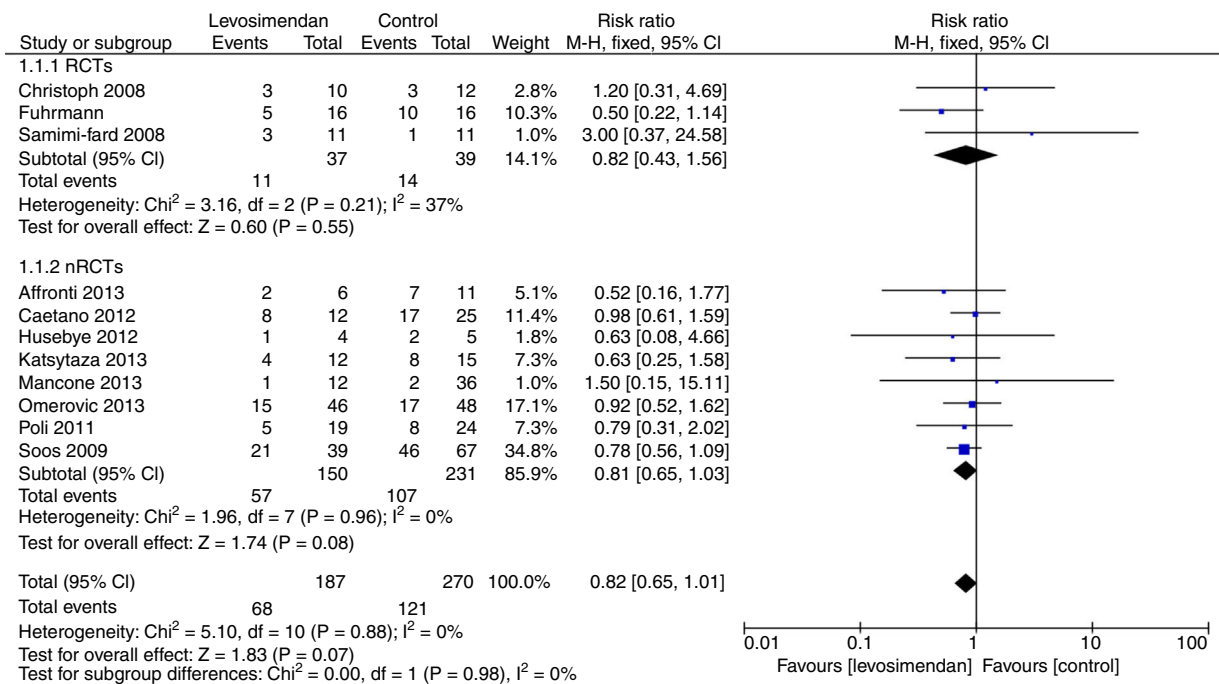


Figure 1 Forest plot for risk of mortality in RCT and nRCT.

Table 2 Secondary end points.

Outcome of interest	No. of studies	MD (95% CI)	P	I <sup>2</sup> (%)	P (Q test)	Tau <sup>2</sup>
SOFA	2	-1.87 [-3.92, 0.18]	0.07	72	0.03	1.74
Cardiac index (L min <sup>-1</sup> m <sup>-2</sup> )	4	0.17 [0.06, 0.29]	0.003	3	0.38	0.00
Cardiac power index (W m <sup>-2</sup> )	3	0.08 [0.03, 0.13]	0.003	60	0.08	0.00
Ejection fraction (%)	4	1.43 [1.03, 1.83]	<0.0001	20	0.29	0.04
End-systolic volume (ml)	3	-6.18 [-9.94, -2.42]	0.001	76	0.02	8.23
Mean blood pressure (mmHg)	2	3.49 [1.05, 5.94]	0.005	0	0.48	0.00
Pulmonary artery pressure (mmHg)	2	-4.00 [-7.41, -0.58]	0.02	24	0.25	1.92
ScvO <sub>2</sub> (%)	2	11.40 [7.35, 15.44]	<0.00001	0	0.71	0.00
ICU days	3	-0.16 [-3.14, 2.83]	0.92	0	0.59	0.00
GFR	2	3.54 [-7.04, 14.13]	0.51	70	0.07	42.31
PAOP	2	1.25 [-0.12, 2.62]	0.07	0	0.88	0.00

pared with the control group (MD 1.43[1.03, 1.83], *P* for effect < 0.00001, *P* for heterogeneity = 0.29, *I*<sup>2</sup> = 20%, Tau<sup>2</sup> 1.33; Appendix A. Supplementary data).

Three studies<sup>13,21,22</sup> reported End-Systolic Volume that was significantly lower in the levosimendan group when compared with the control group (MD -6.18[-9.94, -2.42], *P* for effect = 0.001, *P* for heterogeneity = 0.02, *I*<sup>2</sup> = 76%, Tau<sup>2</sup> 8.23; Appendix A. Supplementary data).

### Levosimendan in patients with cardiogenic shock: hemodynamics

Two studies<sup>10,17</sup> reported Mean Blood Pressure that was significantly higher in the levosimendan group when compared with the control group (MD 3.49[1.05, 5.94], *P* for

effect = 0.005, *P* for heterogeneity = 0.60, *I*<sup>2</sup> = 0%, Tau<sup>2</sup> 0.00; Appendix A. Supplementary data).

Two studies<sup>10,22</sup> reported Pulmonary Atrial Pressure that was significantly lower in the levosimendan group when compared with the control group (MD -4.00[-7.41, -0.58], *P* for effect = 0.02, *P* for heterogeneity = 0.25, *I*<sup>2</sup> = 24%, Tau<sup>2</sup> 1.92; Appendix A. Supplementary data).

Two studies<sup>10,19</sup> reported SvO<sub>2</sub> that was significantly higher in the levosimendan group when compared with the control group (MD 11.40[7.35, 15.44], *P* for effect < 0.00001, *P* for heterogeneity = 0.71, *I*<sup>2</sup> = 0%, Tau<sup>2</sup> 0.00; Appendix A. Supplementary data).

Two studies<sup>10,17</sup> reported pulmonary artery occlusion pressure (PAOP) was not significant difference between levosimendan group and control group (MD 1.25[-0.12, 2.62], *P* for effect = 0.07, *P* for heterogeneity = 0.88, *I*<sup>2</sup> = 0%, Tau<sup>2</sup> 0.00; Appendix A. Supplementary data).

## Discussion

We performed a systematic review and meta-analysis of controlled trials and nRCTs to evaluate the effect of levosimendan compared with standard therapies or placebo on survival and hemodynamic parameters in patients presenting cardiogenic shock complicating myocardial infarction. It revealed that levosimendan is associated with improved cardiac function and many hemodynamic parameters, but it was not associated with a significant reduction in mortality, even if a non-significant trend was present. Notably, this is an important meta-analysis performed on levosimendan administration in patients with cardiogenic shock complicating myocardial infarction.

Cardiogenic shock is very different from the Low Cardiac Output Syndrome (LCOS). In many studies and meta-analyses, levosimendan was proved more effective than standard therapies in the patients with LCOS.<sup>23–25</sup> The most important difference between cardiogenic shock and LCOS is the reduction in cardiac pump performance with hypoperfusion to vital organs. Maybe the effect of Levosimendan could benefit more the population of LCOS than cardiogenic shock.

Levosimendan is a calcium enhancer with calcium-sensitizing activity. C-AMP independent and ATP neutral induce the improvement in calcium sensitivity of cardiac muscle cell.<sup>25</sup> In 2003, Delle-Karth and co-worker<sup>26</sup> published the successful use of levosimendan in patients with cardiogenic shock for the first time. In our meta-analysis, Levosimendan may eliminate the SOFA scores, enhance the cardiac functions, improve cardiac ejection fraction and cardiac volume of systolic and diastolic stage, and improve the other hemodynamic parameters. The effects could come from the levosimendan mechanism. Levosimendan exhibits calcium-dependent binding to the N-terminal domain of cardiac troponin C (TnC) with a higher affinity at high calcium concentrations and lower affinity at low calcium concentrations.<sup>25</sup> The positive inotropic effect is obtained without increasing intracellular calcium concentration or without a significant increase in myocardial oxygen demand, usually seen with other inotropes.<sup>27,28</sup> This implies that less energy is utilized by the cardiomyocytes, because re-internalization of calcium increases ATP expenditure and accounts for 30% of energy consumed by the cardiomyocyte during the contraction-relaxation cycle. A comparison of levosimendan and milrinone<sup>29</sup> showed that both intensify cardiac contraction, but milrinone increased oxygen consumption, levosimendan did not. In other studies, levosimendan was also shown to be superior to dobutamine in term of myocardial efficiency.<sup>30,31</sup>

Again, we found that Levosimendan may improve the hemodynamic parameters. Several clinical observations reveal that levosimendan improve hemodynamics even in patients with cardiogenic shock if it is combined with catecholamines to maintain adequate perfusion pressure.<sup>32,33</sup>

Levosimendan was associated with a non-significant improve in survival. This may suggest that use of levosimendan in patients with cardiogenic shock complicating myocardial infarction does not offer a mortality benefit or, more probably, that too few patients have been randomized

so far to reach powered conclusions on mortality. On the other hand, we also found that levosimendan can improve the hemodynamic parameters. Maybe the pooled analysis could have been insufficiently powered to detect a clinically relevant reduction in mortality in a general population of patients with cardiogenic shock. Perhaps, further larger high-quality RCTs are warranted to reach conclusions on the topic.

This meta-analysis has strengths and limitations. The methodological quality of the studies included in this meta-analysis was not optimal, with only one trial presenting low risk of bias.<sup>11</sup> Not all trials reported all hemodynamic parameters, so these estimations are drawn from small numbers of measurements and should therefore be interpreted with caution. Unfortunately, there are only 5 randomized trials analyzing the effect of levosimendan in patients with cardiogenic shock. Ideally, a large randomized controlled trial on the lookout for the eventual beneficial effects of levosimendan in cardiogenic shock setting should be performed.

## Conclusion

In summary, this systematic review of 13 clinical trials found that there was no evidence of survival benefit when levosimendan was compared to control therapies. On the other hand, levosimendan was associated with an improvement in hemodynamics and cardiac function. Further high quality studies on levosimendan for patients with cardiogenic shock complicating myocardial infarction are needed.

## Authors' contributions

Study design: FM; study conduct: FM, CH; data analysis: FM, CH; data interpretation: FM, CH, WZ; writing and revising paper: FM, CH, WZ. All authors read and approved the final manuscript.

## Funding

There was no funding source for this study.

## Conflict of interests

The authors did not have any conflict of interests.

## Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at [doi:10.1016/j.medin.2017.08.009](https://doi.org/10.1016/j.medin.2017.08.009).

## References

1. Fox KAA, Steg PG, Eagle KA, Goodman SG, Anderson FA Jr, Granger CB, et al. Decline in rates of death and heart failure in acute coronary syndromes, 1999–2006. *JAMA*. 2007;297:1892–900.
2. Thiele H, Zeymer U, Neumann FJ, Ferenc M, Olbrich HG, Hausleiter J, et al. IABP-SHOCK II Trial Investigators: intraor-

- tic balloon support for myocardial infarction with cardiogenic shock. *N Engl J Med.* 2012;367:1287–96.
3. Musial R, Moncznik P, Smialek P, Stolinski J, Sadowski J, Drwila R. Experience in application of therapies VA ECMO as short-term mechanical support of circulatory system of adult patients in cardiogenic shock. *Kardiol Pol.* 2016;74:1477–84.
  4. Toller WG, Stranz C. Levosimendan a new inotropic and vasodilator agent. *Anesthesiology.* 2006;104:556–69.
  5. Sorsa T, Pollesello P, Solaro RJ. The contractile apparatus as a target for drugs against heart failure: interaction of levosimendan, a calcium sensitiser, with cardiac troponin c. *Mol Cell Biochem.* 2004;266:87–107.
  6. Papp Z, Csapo K, Pollesello P, Haikala H, Edes I. Pharmacological mechanisms contributing to the clinical efficacy of levosimendan. *Cardiovasc Drug Rev.* 2005;23:71–98. Spring.
  7. Greco T, Calabrò MG, Covello RD, Greco M, Pasin L, Morelli A, et al. A Bayesian network meta-analysis on the effect of inodilatory agents on mortality. *Br J Anaesth.* 2015;114:746–56.
  8. Higgins JP, Green S. *Cochrane handbook for systematic reviews of interventions.* Version 5. The cochrane collaboration, 2011; 2011.
  9. Stang A. Critical evaluation of the Newcastle–Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol.* 2010;25:603–5.
  10. Fuhrmann JT, Schmeisser A, Schulze MR, Wunderlich C, Schoen SP, Rauwolf T, et al. Levosimendan is superior to enoximone in refractory cardiogenic shock complicating acute myocardial infarction. *Crit Care Med.* 2008;36:2257–66.
  11. Husebye T, Eritsland J, Muller C, Sandvik L, Arnesen H, Seljeflot I, et al. Levosimendan in acute heart failure following primary percutaneous coronary intervention treated acute ST-elevation myocardial infarction. Results from the LEAF trial: a randomized, placebo-controlled study. *Eur J Heart Fail.* 2013;15:565–72.
  12. Samimi-Fard S, Garcia-Gonzalez MJ, Dominguez-Rodriguez A, Abreu-Gonzalez P. Effects of levosimendan versus dobutamine on long-term survival of patients with cardiogenic shock after primary coronary angioplasty. *Int J Cardiol.* 2008;127:284–7.
  13. Peng L, Rui L, Yongguang Z, Wei H, Jianhui X, Jianye X, et al. Levosimendan for treating acute myocardial infarction complicated by cardiogenic shock in 54 cases. *China Pharm.* 2015;9:46–7.
  14. 罗毅, 刘路培, 杨振宇等. 左西孟旦对急性心肌梗死并发性休克患者的疗效. *实用医学杂志* 2013; 第29卷19期: 3229–30.
  15. Affronti A, di Bella I, Carino D, Ragni T. Levosimendan may improve weaning outcomes in venoarterial ECMO patients. *ASAIO J.* 2013;59:554–7.
  16. Omerovic E, Ramunddal T, Albertsson P, Holmberg M, Hallgren P, Boren J, et al. Levosimendan neither improves nor worsens mortality in patients with cardiogenic shock due to ST-elevation myocardial infarction. *Vasc Health Risk Manage.* 2010;7:657–63.
  17. Christoph A, Prondzinsky R, Russ M, Janusch M, Schlitt A, Lemm H, et al. Early and sustained haemodynamic improvement with levosimendan compared to intraaortic balloon counterpulsation (IABP) in cardiogenic shock complicating acute myocardial infarction. *Acute Cardiac Care.* 2008;10:49–57.
  18. Soos P, Becker D, Barczy G, Szabo G, Zima E, Fulop G, et al. Levosimendan therapy does not improve survival of post-resuscitation cardiogenic shock patients. *Crit Care.* 2009;13 Suppl. 1:172.
  19. Poli M, Trambaiolo P, Mustili M, De Luca M, Lukic V, Basso V, et al. Mixed venous oxygen saturation. Its role in the assessment of patients developing after ST-elevation myocardial infarction (STEMI). *Giornale Italiano di Cardiologia.* 2011;5 Suppl. 1:S39–40.
  20. Caetano F, Almeida I, Silva J, Botelho A, Mota P, Leitao Marques A. Cardiogenic shock in acute myocardial infarction: still looking for the best inotrope. *Eur Heart J: Acute Cardiovasc Care.* 2012;1 Suppl. 1:S28–9.
  21. Katsytadze I, Amosova E, Prudkyi I, Kyrylova A. Long term effects of levosimendan therapy in patients with cardiogenic shock. *Resuscitation.* 2013;84 Suppl.:S11.
  22. Mancone M, Pennacchi M, Lucisano L, Calcagno S, D'Ambrosi A, Marceca A, et al. Hemodynamic improvement following levosimendan treatment in acute myocardial infarction complicated by cardiogenic shock patients treated with primary PCI and IABP. *JACC.* 2013;10:E770.
  23. Koster F, Wetterslev J, Gluud C, Zijstra JG, Scheeren TW, van der Horst IC, et al. Effects of levosimendan for low cardiac output syndrome in critically ill patients: systematic review with meta-analysis and trial sequential analysis. *Intensive Care Med.* 2015;41:203–21.
  24. Lim JY, Deo SV, Rababa'h A, Altarabsheh SE, Cho YH, Hand D, et al. Levosimendan reduces mortality in adults with left ventricular dysfunction undergoing cardiac surgery: a systematic review and meta-analysis. *J Card Surg.* 2015;30:547–54.
  25. Kass DA, Solaro RJ. Mechanisms and use of calcium-sensitizing agents in the failing heart. *Circulation.* 2006;113:305–15.
  26. Zangrillo A, Alvaro G, Pisano A, Guarracino F, Lobreglio R, Bradic N, et al. A randomized controlled trial of levosimendan to reduce mortality in high-risk cardiac surgery patients (CHEETAH): rationale and design. *Am Heart J.* 2016;177:66–73.
  27. Hasenfuss G, Pieske B, Castell M, Kretschmann B, Maier LS, Just H. Influence of the novel inotropic agent levosimendan on isometric tension and calcium cycling in failing human myocardium. *Circulation.* 1998;98:2141–7.
  28. Ukkonen H, Saraste M, Akkila J, Knuuti MJ, Lehtikainen P, Nagren K, et al. Myocardial efficiency during calcium sensitization with levosimendan: a noninvasive study with positron emission tomography and echocardiography in healthy volunteers. *Clin Pharmacol Ther.* 1997;61:596–607.
  29. Kaheinen P, Pollesello P, Levijoki J, Haikala H. Effects of levosimendan and milrinone on oxygen consumption in isolated guinea pig heart. *J Cardiovasc Pharmacol.* 2004;43:555–61.
  30. Ukkonen H, Saraste M, Akkila J, Knuuti MJ, Lehtikainen P, Nagren K, et al. Myocardial efficiency during calcium sensitization with levosimendan: a noninvasive study with positron emission tomography and echocardiography in healthy volunteers. *Clin Pharmacol Ther.* 1997;61:596–607.
  31. Ukkonen H, Saraste M, Akkila J, Knuuti J, Karanko M, Iida H, et al. Myocardial efficiency during levosimendan infusion in congestive heart failure. *Clin Pharmacol Ther.* 2000;68:522–31.
  32. Franco F, Monteiro P, Correia J, Roque C, Goncalves F, Castro G, et al. Levosimendan is safe and effective in patients with severe low cardiac output heart failure and critical hypotension. *J Am Coll Cardiol.* 2004;43 Suppl. A:191A.
  33. McLean A, Huang S, Stewart D, Nalos M, Tang B. Efficacy of levosimendan in shock. (Abstract). *Crit Care.* 2004;8 Suppl. 1:83.