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Integrating the results of the CULPRIT-SHOCK trial in the 2017 ESC STEMI Guidelines: Viewpoint of the Task Force

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Spirit of the viewpoint

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3 The recent 2017 European Society of Cardiology (ESC) Guidelines for the management
4 of acute myocardial infarction in patients presenting with ST-segment elevation (STEMI
5 GL).¹ included 159 recommendations based on 477 references. Although the field of
6 acute myocardial infarction is highly evidence-based many treatment options have never
7 been tested in prospective randomized clinical trials (RCTs). Under these circumstances
8 the guideline TF is expected to develop clinically useful recommendations using
9 consensus based on available evidence from small trials or observational studies or plain
10 clinical experience (level of evidence (LOE) C). In the recent STEMI guidelines 49% of
11 the recommendations were labeled as LOE C. Many of these LOE C recommendations
12 were acknowledged as relevant areas for future research in the 2017 STEMI GL
13 document.¹

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22 Regarding the management of patients with cardiogenic shock complicating STEMI and
23 with severe stenosis apart from the infarct-related artery (IRA), the recent 2017 STEMI
24 GL¹ favoured complete revascularization during the index primary percutaneous
25 coronary intervention (PCI), allocating a class of recommendation IIa with LOE C. After
26 the publication of the STEMI GL,¹ the “Culprit Lesion Only PCI versus Multivessel PCI
27 in Cardiogenic Shock” (CULPRIT-SHOCK) trial demonstrated that routine complete
28 revascularization during index PCI procedure in this population is harmful.² In light of
29 this outcome, the 2017 STEMI GL task force (TF) considers it important to provide the
30 cardiology community with a viewpoint that can help readers to place these apparent
31 contradictory messages in context and help physicians taking the best therapeutic decision
32 for their patients. A selected group of members of the 2017 STEMI GL TF leading the
33 chapters related to this topic decided to write this document. To get the broadest view of
34 this relevant topic, additional authors were invited to participate in this document,
35 including the principal investigator of CULPRIT-SHOCK (H.T.), the chair of the ESC
36 Committee for Practice Guidelines (CPG) (S.W.), and the chair of the upcoming 2018
37 ESC Myocardial Revascularization GL (F.-J.N.).

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**Complete revascularization in STEMI patients with multi-vessel disease but no
cardiogenic shock: evidence leading to the 2017 recommendation**

1 The recommendation for non-IRA PCI in STEMI was significantly modified from the
2 2012³ to the 2017 STEMI GL.¹ In the 2012 GL, it was recommended that primary PCI
3 should be limited to the culprit vessel with the exception of cardiogenic shock and
4 persistent ischaemia after PCI of the supposed culprit lesion (class IIa LOE B).³ Thus,
5 routine PCI of non-IRA was not recommended. The 2017 document proposed that routine
6 PCI of non-IRA severe stenoses should be considered before hospital discharge (Class
7 IIa, LOE A).¹ Justification for this change was based on the results of four medium-sized
8 randomized controlled trials (RCTs) and several meta-analyses comparing IRA-only PCI
9 vs. complete revascularization in stable STEMI patients⁴⁻⁷ none of which included
10 patients with cardiogenic shock or resuscitated from cardiac arrest. The primary outcome
11 measure was mainly driven by a reduction in repeat revascularisation rates. While this
12 might be seen as a self-fulfilled prophecy (i.e. revascularisations already done upfront in
13 the active treatment), meta-analyses also revealed ischemic outcomes (death or
14 myocardial infarction) were numerically lower in the non-IRA PCI group in most of the
15 trials.⁸ The overall low event rate in conjunction with the relatively small size of all trials
16 precluded the demonstration of a clinical benefit beyond repeat revascularisations. For
17 this reason, the class of recommendation for non-IRA PCI before hospital discharge was
18 IIa and not in I.

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35 **Evidence to recommend non-IRA PCI during index procedure in STEMI patients**
36 **with multivessel disease and cardiogenic shock**

37 In patients with STEMI and cardiogenic shock, early revascularization of the IRA
38 improves outcomes.^{9, 10} Up to 80% of patients with STEMI and shock have multivessel
39 disease, and the mortality in these patients is higher than that of those with single-vessel
40 disease.¹¹ Prior to the publication of CULPRIT-SHOCK trial, late in 2017,² the evidence
41 available on the clinical benefit of complete vs. IRA-only PCI in this population was
42 based on indirect evidence and observational studies. In the SHOCK trial, 40% of patients
43 underwent coronary artery bypass grafting (CABG) likely extending beyond the IRA
44 revascularisation.¹² In the Manitoba Cardiogenic Shock Registry, a retrospective
45 multicentre cohort of patients with cardiogenic shock undergoing coronary angiography,
46 complete revascularization was identified as an independent predictor for hospital
47 survival in the subgroup of STEMI.¹³ In the absence of prospective RCTs, these data was
48 considered by the 2012 STEMI GL to recommend non-IRA PCI in STEMI patients with
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1 persistent shock after IRA PCI.³ Similarly, the most recent U.S. appropriate-use criteria
2 defined as appropriate to perform immediate PCI of a non-IRA if cardiogenic shock
3 persisted after IRA PCI.¹⁴ Owing to the mentioned benefits of non-IRA PCI in STEMI
4 patients without cardiogenic shock of and the absence of evidence of harm, the 2017
5 STEMI GL document maintained the 2012 recommendation for non-IRA in STEMI
6 patients with multi-vessel disease in cardiogenic shock although the wording was less
7 stringent by eliminating the consideration that this should be done in patients with
8 persistent shock after IRA PCI.
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18 **The CULPRIT-SHOCK trial**

19 The CULPRIT-SHOCK trial is the largest RCT in cardiogenic shock complicating
20 myocardial infarction (62% STEMI) to date comparing IRA-only PCI with immediate
21 PCI of all severe lesions.² The CULPRIT-SHOCK trial addressed a contemporary, very
22 high-risk patient population. Roughly half of enrolled patients had been resuscitated prior
23 to randomization, and almost one third received some form of hemodynamic support. The
24 primary endpoint (composite of death or severe renal failure leading to renal-replacement
25 therapy at one month) was higher in the immediate multivessel PCI than in the IRA-only
26 PCI group (55.4% vs. 45.9%; P=0.01).² The results were mainly driven by an absolute
27 8.2% difference in 30-day all-cause mortality (51.5% vs. 43.3%, P=0.03). Results were
28 consistent across pre-specified subgroups including all age groups, sex, presence/absence
29 of diabetes, presence/absence of hypertension, STEMI or non-STEMI, anterior/non-
30 anterior STEMI, previous/no previous infarction²/3-vessel disease, or presence/absence
31 of chronic total occlusion (CTO). In the multivessel PCI group, complete
32 revascularization was achieved in 81.0% of the patients. Staged revascularization was
33 performed in 17.7% of the patients in the IRA -only PCI group, and the cross-over rate
34 was limited (12.5% in the IRA -only PCI group and 9.4% in the multivessel PCI group).
35 The consistent risk estimates for the primary end point in the intention-to-treat, per-
36 protocol, and as-treated analyses support the robustness of the findings.
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56 It is well known that presence of a CTO is frequent in cardiogenic shock and associated
57 with high mortality.¹⁵ At least one CTO was present in 22.4% in the IRA-only PCI arm
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1 and in 24.0% in the immediate multivessel PCI arm. In CULPRIT-SHOCK, immediate
2 CTO recanalization was attempted in roughly 50% of patients in the immediate
3 multivessel PCI group and was successful in approximately one third of attempts. The
4 results for the primary study endpoint were consistent for CTO presence or absence. The
5 mechanisms leading to the higher 30-day mortality in the immediate multivessel PCI
6 group in CULPRIT-SHOCK might be related to the significantly higher amount of
7 contrast medium given (250 cc versus 190 cc; $p<0.001$) with subsequent impairment of
8 renal function. There was a lower estimated glomerular filtration rate in the immediate
9 multivessel PCI group at days 3 and 4, although differences in the incidence of severe
10 renal failure leading to renal replacement therapy failed to reach conventional levels of
11 statistical significance (11.6% versus 16.4%; $p=0.07$). The higher dose of contrast
12 medium in the immediate multivessel PCI group may also have led to acute left
13 ventricular volume overload with a negative effect on myocardial function and recovery.
14 In addition, the prolonged duration of the multivessel PCI procedure may be hazardous
15 at a time when the patient is hemodynamically compromised. Additional myocardial
16 damage may also have been induced by PCI in non-IRA stable lesions.

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29 Interestingly, the 30-day mortality rate in the IRA-only PCI group was nearly identical
30 with that of the SHOCK (Should We Emergently Revascularize Occluded Coronaries for
31 Cardiogenic Shock) trial performed 2 decades ago.^{9, 12} This similar mortality rate may be
32 partly explained by the higher risk profile in the CULPRIT-SHOCK trial because patients
33 with single vessel coronary artery disease were excluded.

41 **Position of the TF after the publication of CULPRIT SHOCK**

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43 Based on the new robust evidence from the adequately powered CULPRIT-SHOCK trial,
44 it is now the opinion of the 2017 STEMI TF that in patients with cardiogenic shock
45 complicating STEMI, primary PCI should be restricted to the IRA. Immediate multivessel
46 PCI may be justified in the rare cases where the IRA is difficult to identify or incorrectly
47 defined initially or when multiple culprit lesions are identified. Selected cases in which
48 there is a very severe flow-limiting non-IRA stenosis irrigating a large myocardial area
49 may also justify immediate non-IRA PCI. Staged non-IRA PCI might be an option,
50 carefully balancing the benefits and risks of a new procedure with additional contrast
51 loading and risk of complications. The new edition of the ESC/EACTS guidelines on
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1 myocardial revascularization will be released this year, incorporating the results of all
2 published data so far. In the meantime, decision making in STEMI patients with
3 cardiogenic shock and multi-vessel disease should be based on available data from
4 CULPRIT SHOCK trial taking into consideration the individual patient using medical
5 judgement based on the available evidence.
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11 **Conclusion**

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16 Evidence regarding the best approach for cardiogenic shock complicating STEMI and
17 multivessel disease is constantly being generated, and several metaanalyses from non-
18 RCT are being published with mix, even contradictory results to each other.¹⁶⁻¹⁸ We have
19 learnt that data from non-randomized, retrospective and observational studies is
20 potentially affected by important bias and might not represent a real effect. Data from
21 RCT represent the best evidence to guide therapies. The CULPRIT-SHOCK trial is the
22 only RCT performed addressing this issue, and demonstrated that routine multivessel PCI
23 during the index procedure in STEMI patients and cardiogenic shock is not safe.
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31 **Summary Key Points**

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36 In patients with cardiogenic shock complicating STEMI and NSTEMI:

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38 Primary PCI should routinely be restricted to the IRA

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40 Immediate multivessel PCI may be justified if the IRA is difficult to identify or
41 incorrectly defined initially or when multiple culprit lesions are identified.

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43 Immediate multivessel PCI may be justified in selected cases in which there is a
44 flow-limiting non-IRA very severe stenosis irrigating a large myocardial area.

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46 Staged non-IRA PCI might be an option, carefully balancing the benefits and
47 risks.
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53 **Disclosures**

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56 Holger Thiele was the PI of the CULPRIT-SHOCK trial, which was funded the
57 European Union 7th Framework Program (FP7/2007-2013) and by the German Heart
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1 Research Foundation and the German Cardiac Society. All other authors have nothing
2 to declare in relation to this work.
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