

# Bromocriptine for the treatment of peripartum cardiomyopathy: a multicentre randomized study

Denise Hilfiker-Kleiner<sup>1\*†</sup>, Arash Haghikia<sup>1,2†</sup>, Dominik Berliner<sup>1</sup>, Jens Vogel-Claussen<sup>3</sup>, Johannes Schwab<sup>4</sup>, Annegret Franke<sup>5</sup>, Marziel Schwarzkopf<sup>5</sup>, Philipp Ehlermann<sup>6</sup>, Roman Pfister<sup>7</sup>, Guido Michels<sup>7</sup>, Ralf Westenfeld<sup>8</sup>, Verena Stangl<sup>9</sup>, Ingrid Kindermann<sup>10</sup>, Uwe Kühl<sup>2</sup>, Christiane E. Angermann<sup>11</sup>, Axel Schlitt<sup>12</sup>, Dieter Fischer<sup>1</sup>, Edith Podewski<sup>1</sup>, Michael Böhm<sup>10</sup>, Karen Sliwa<sup>13</sup>, and Johann Bauersachs<sup>1\*</sup>

<sup>1</sup>Department of Cardiology and Angiology, Hannover Medical School, Carl-Neuberg-Str. 1, D-30625 Hannover, Germany; <sup>2</sup>Department of Cardiology, Charité-Universitätsmedizin Berlin, Campus Benjamin Franklin, Hindenburgdamm 30, 12203 Berlin, Germany; <sup>3</sup>Institute for Diagnostic and Interventional Radiology, Hannover Medical School, Carl-Neuberg-Str. 1, 30625 Hannover, Germany; <sup>4</sup>Universitätsklinik für Innere Medizin 8 – Schwerpunkt Kardiologie und Institut für Radiologie und Neuroradiologie, Klinikum Nürnberg Süd, Paracelsus, Medizinische Privatuniversität Nürnberg, Breslauer Str. 201, 90471 Nürnberg, Germany; <sup>5</sup>Faculty of Medicine, University Leipzig, Clinical Trial Centre (KKS), ZKS Leipzig, Haertelstr. 16-18, D-04103 Leipzig, Germany; <sup>6</sup>Department of Cardiology, Angiology, and Pneumology, University of Heidelberg, Im Neuenheimer Feld 410, 69120 Heidelberg, Germany; <sup>7</sup>Department of Cardiology, Pulmonology, and Vascular Medicine, University of Cologne, Cologne, Kerpenerstr. 62, 50937 Köln, Germany; <sup>8</sup>Department of Cardiology, Pulmonology, and Vascular Medicine, Medical Faculty, University Duesseldorf, Duesseldorf, Germany; <sup>9</sup>Department for Cardiology and Angiology, Angiology, Center for Cardiovascular Research (CCR), Charité-Universitätsmedizin Berlin, Charitéplatz 1, 10117 Berlin, Germany; <sup>10</sup>Department of Internal Medicine III, University Hospital of the Saarland, 66421 Homburg/Saar, Germany; <sup>11</sup>Department of Internal Medicine I, University Hospital Wuerzburg, Am Schwarzenberg 15, Haus A1597078 Würzburg, Germany; <sup>12</sup>Department of Medicine III, Medizinische Fakultät der Martin Luther-Universität Halle-Wittenberg, Magdeburger Straße 8, 06108 Halle, Germany; and <sup>13</sup>Faculty of Health Sciences, Hatter Institute of Cardiology Research in Africa, 2 Anzio Road, Chris Barnard Building, 4th Floor, OBSERVATORY 7925, South Africa

Received 10 April 2017; revised 16 May 2017; editorial decision 2 June 2017; accepted 23 June 2017; online publish-ahead-of-print 27 July 2017

See page 2680 for the editorial comment on this article (doi: 10.1093/eurheartj/ehx428)

## Aims

An anti-angiogenic cleaved prolactin fragment is considered causal for peripartum cardiomyopathy (PPCM). Experimental and first clinical observations suggested beneficial effects of the prolactin release inhibitor bromocriptine in PPCM.

## Methods and results

In this multicentre trial, 63 PPCM patients with left ventricular ejection fraction (LVEF)  $\leq 35\%$  were randomly assigned to short-term (1W: bromocriptine, 2.5 mg, 7 days) or long-term bromocriptine treatment (8W: 5 mg for 2 weeks followed by 2.5 mg for 6 weeks) in addition to standard heart failure therapy. Primary end point was LVEF change (delta) from baseline to 6 months assessed by magnetic resonance imaging. Bromocriptine was well tolerated. Left ventricular ejection fraction increased from  $28 \pm 10\%$  to  $49 \pm 12\%$  with a delta-LVEF of  $+21 \pm 11\%$  in the 1W-group, and from  $27 \pm 10\%$  to  $51 \pm 10\%$  with a delta-LVEF of  $+24 \pm 11\%$  in the 8W-group (delta-LVEF:  $P = 0.381$ ). Full-recovery (LVEF  $\geq 50\%$ ) was present in 52% of the 1W- and in 68% of the 8W-group with no differences in secondary end points between both groups (hospitalizations for heart failure: 1W: 9.7% vs. 8W: 6.5%,  $P = 0.651$ ). The risk within the 8W-group to fail full-recovery after 6 months tended to be lower. No patient in the study needed heart transplantation, LV assist device or died.

## Conclusion

Bromocriptine treatment was associated with high rate of full LV-recovery and low morbidity and mortality in PPCM patients compared with other PPCM cohorts not treated with bromocriptine. No significant differences were observed between 1W and 8W treatment suggesting that 1-week addition of bromocriptine to standard heart failure treatment is already beneficial with a trend for better full-recovery in the 8W group.

## Clinical trial registration:

ClinicalTrials.gov, study number: NCT00998556.

\* Corresponding authors. Tel: +49 511532 2531, Fax: +49 511532 3263, Email: hilfiker.denise@mh-hannover.de; Tel: +49 511 532 3841, Fax: +49 511 5325412, Email: bauersachs.johann@mh-hannover.de

† The first two authors contributed equally as first authors.

© The Author 2017. Published by Oxford University Press on behalf of the European Society of Cardiology.

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0/>), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited. For commercial re-use, please contact journals.permissions@oup.com

## Keywords

Peripartum cardiomyopathy • Bromocriptine • Prolactin • heart failure

## Introduction

Peripartum cardiomyopathy (PPCM) is increasingly recognized as a major cause of pregnancy related heart failure with high morbidity and mortality.<sup>1–4</sup> The reported incidence of PPCM is regionally different ranging between 1 per 100 in Nigeria, 1 in 299 in Haiti to 1 in 1149–3189 live births in the USA.<sup>1,5–7</sup> To date no evidence-based disease specific therapy is available for PPCM. Recommended treatment for PPCM is similar to that of heart failure from other aetiologies based on registry data and expert opinion.<sup>2,4,8–10</sup> Although the aetiology of PPCM is still under investigation, in recent years a number of contributory mechanisms have been recognized to initiate and propagate the disease.<sup>1–4</sup> Hereby, high levels of the nursing hormone prolactin and the production of a cleaved 16kDa N-terminal fragment of prolactin have emerged as potential key factors in the pathophysiology of PPCM. Experimental studies suggest that 16kDa prolactin induces profound endothelial damage and subsequent cardiomyocyte dysfunction and full-length prolactin promotes inflammation in PPCM while inhibition of prolactin release by the dopamine-D2-receptor agonist bromocriptine prevents onset of PPCM.<sup>11–14</sup> Bromocriptine has been used for many years to stop lactation in postpartum women. In addition, prolactin-independent cytoprotective effects of bromocriptine were also shown in various organs including the heart.<sup>14,15</sup> The strong improvement of cardiac function in previous reports including a clinical pilot study and register-based data demonstrating that prolonged treatment of PPCM patients with bromocriptine is feasible and may improve left ventricular (LV) recovery and clinical outcome<sup>12,16,17</sup> lend support to the hypothesis that prolonged bromocriptine treatment beyond the cumulative dose to stop lactation might be needed to achieve maximum clinical benefits. We therefore designed this randomized multicentre trial to compare the effects of prolonged bromocriptine treatment vs. short-term treatment sufficient to stop lactation in addition to guideline-based heart failure therapy<sup>9,10</sup> on LV function and clinical outcomes in patients with PPCM.<sup>18</sup>

## Methods

The study design has been reported previously.<sup>18</sup>

Detailed information on the study protocol and methods are provided in the Supplementary material online.

## Results

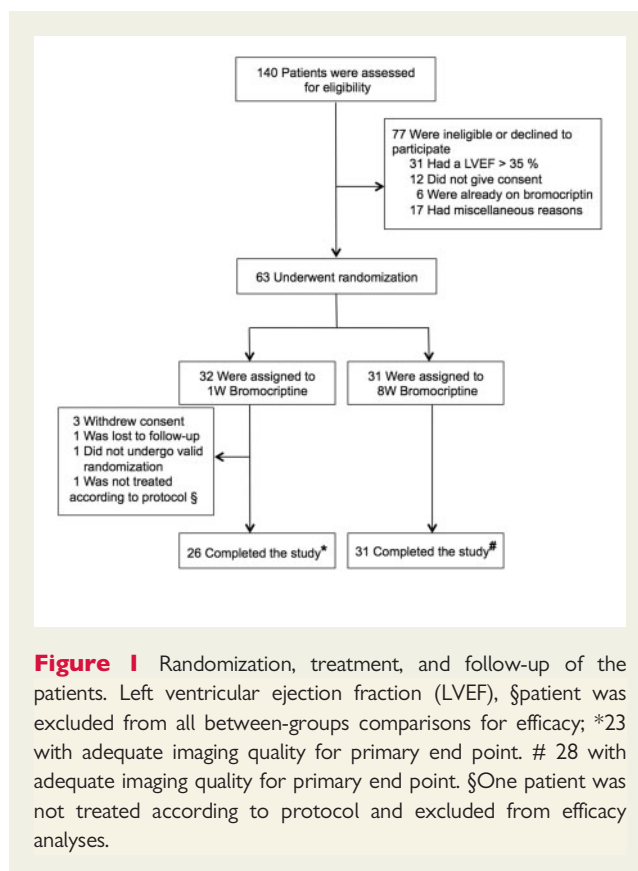
### Screening, inclusion criteria, and randomization of study patients

From June 2010 until September 2015, 140 patients at 12 centres were screened for eligibility. Of these patients, 63 were included in the study while 77 patients did not fulfil the criteria for randomization. Reasons for exclusion were LVEF > 35%, other reasons for heart failure, time since delivery more than 6 months, social reasons,

not agreed to participate in the study, drug abuse, noncompliance and other reasons. All patients included in the study were postpartum. Accordingly, 32 patients were randomly assigned to receive 2.5 mg bromocriptine for 1 week (1W group) and 31 to receive 5 mg bromocriptine for 2 weeks followed by 2.5 mg for 6 weeks (8W group). In the 1W group, three patients withdrew consent after randomization, one was lost to follow-up, one did not undergo valid randomization. An additional patient of the 1W group was not treated according to the protocol but received higher dose of bromocriptine (up to 10 mg) and relevantly longer so that the clinical event committee (CEC) decided to exclude her from all between-groups comparisons of efficacy end points. Therefore, 31 patients per group comprised the efficacy analysis set of patients. In total, 26 patients from the 1W group and 31 from the 8W group completed the study with follow-up data of 6 months after diagnosis (Figure 1). Of these patients, 23 patients from the 1W and 28 patients from the 8W group met the imaging quality standards evaluated by the CEC in order to be taken into consideration for the primary end point (trial profile is shown in Figure 1).

### Baseline characteristics of study patients

The characteristics of all randomized patients at baseline (Visit 1) are shown in Table 1. More details about the baseline characteristics are



**Table 1** Baseline characteristics of all randomized patients

Characteristic	1W bromocriptine (N = 32)	8W bromocriptine (N = 31)
Age—year	33.8 ± 5.8	34.0 ± 4.5
Median gravida (range)	1.5 (1–7)	2 (1–5)
Median parity (range)	1.5 (1–7)	2 (1–6)
Race or ethnic group—number (%) <sup>a</sup>		
Caucasian	32 (100)	30 (97)
Black	0 (0)	1 (3)
Systolic blood pressure—mmHg	117 ± 18	111 ± 17
Heart rate—beats per minute	90 ± 16	84 ± 12
Body mass index <sup>b</sup>	28.3 ± 6.8	29.0 ± 7.9
Serum creatinine—mg/dL	0.8 ± 0.2	0.9 ± 0.2
Clinical features of heart failure		
Left ventricular ejection fraction—(%)	28 ± 10	27 ± 9
Right ventricular ejection fraction—(%) <sup>c</sup>	48 ± 12	42 ± 13
Median NT-proBNP—pg/mL	2164 (1290–3066)	2437 (1423–4158)
NYHA functional class—number (%) <sup>d</sup>		
I	0 (0)	1 (3.2)
II	4 (12.5)	4 (12.9)
III	10 (31.3)	9 (29.0)
IV	18 (56.3)	17 (54.8)
Medical history—number (%)		
Hypertension	7 (21.9)	11 (35.5)
Diabetes	2 (6.3)	1 (3.2)
Smoker or former smoker	16 (50)	15 (48)
Pregnancy related conditions—number (%)		
Preeclampsia	5 (15.6)	8 (25.8)
Gestational diabetes	3 (9.4)	2 (6.5)
Treatment at randomization—number (%)		
ACE inhibitor	30 (93.8)	27 (87.1)
Angiotensin receptor blocker	3 (9.4)	3 (9.7)
Mineralcorticoid antagonist	27 (84.6)	25 (80.6)
Beta-blocker	29 (90.6)	31 (100)
Diuretic	28 (88.5)	28 (90.3)

Plus-minus values are means ± SD. There were no significant differences between the two groups. Data were missing for the following characteristics: N-terminal pro-B-type natriuretic peptide for 1 patient in the 1W group and 1 in the 8W group, respectively, body-mass index, for 1 patient in the 1W group. Percentages may not total 100 because of rounding. To convert the values for creatinine to micromoles per litre, multiply by 88.4.

IQR, interquartile range; NYHA, New York Heart Association.

<sup>a</sup>Race or ethnic group was reported by the investigators.

<sup>b</sup>The body mass index is the weight in kilograms divided by the square of the height in meters.

<sup>c</sup>The data for right ventricular ejection fraction were available for  $n = 44$  patients (1W:  $n = 20$ ; 8W:  $n = 24$ ).

<sup>d</sup>The data for NYHA class reflect the status of patients at the time of randomization.

provided in the Supplementary material online, *Appendix Tables S1–S8*. The groups were balanced with respect to baseline characteristics (age, gravida, parity, race, body mass index, haemodynamics, cardiac function New York Heart Association class), cardiovascular and pregnancy-related risk factors. Data for right ventricular ejection fraction (RVEF) as assessed by CMR were available for 44 patients. Baseline RVEF did not differ significantly between 1W ( $48 \pm 12\%$ ,  $n = 20$ ) and 8W ( $42 \pm 13\%$ ,  $n = 24$ ; 1W vs. 8W:  $P = 0.20$ ). The prevalence of impaired RV function (RVEF < 40%) did not differ significantly between both groups (20% in 1W vs. 42% in 8W,  $P = 0.19$ ) although patients in the 8W group tended to have more often impaired RV function at baseline. No patient was diagnosed with PPCM in a

previous pregnancy or was diagnosed during pregnancy. Diagnosis was made  $1.6 \pm 1.6$  months after delivery and average time between first symptoms and diagnosis was  $0.8 \pm 1.2$  months with no difference between groups. All patients were randomized postpartum. Randomization was done no more than 7 days after first diagnosis of PPCM. One patient was of African race, all other were of Caucasian race. Notably, use of guideline-based heart failure therapy including ACE inhibition or angiotensin receptor blocker (ARB), beta-blockade and mineralocorticoid receptor antagonist (MRA)<sup>9,10</sup> during the study was high and did not differ between both groups (*Table 1*). No patient in the 1W group resumed lactation after bromocriptine treatment.

## Change in left ventricular function as primary end point

The LVEF increased from a mean of  $28 \pm 10\%$  at baseline to  $49 \pm 12\%$  at 6 months in the 1W group ( $n = 23$ ), and from  $27 \pm 10\%$  to  $51 \pm 10\%$  in the 8W group ( $n = 28$ ) with a between-groups difference at 6-months follow-up of 2.0 [95% confidence limit: -4.2; 8.2]% in favour of the 8W groups. Individual courses are presented in Figure 2A. Delta LVEF was slightly higher in the 8W group ( $+24 \pm 11\%$ ) compared with the 1W group ( $+21 \pm 11\%$ ) but this was not statistically significant ( $P = 0.381$ ) with 2.5 [95% confidence limit: -3.2; 8.3]%. In order to analyse effects of the two treatment concepts in more critically ill patients, a subgroup analysis was performed including only patients in whom baseline LVEF was  $<30\%$ . As shown in Figure 2B, the LVEF increased from a mean of  $21 \pm 6\%$  at baseline to  $45 \pm 14\%$  at 6 months in the 1W group ( $n = 14$ ), and from  $21 \pm 6\%$  to  $50 \pm 11\%$  in the 8W group ( $n = 18$ ) again showing a slightly higher delta LVEF in the 8W subgroup ( $+29 \pm 10\%$ ) as compared with delta LVEF of the 1W subgroup ( $+24 \pm 11\%$ ) which was not statistically significant (8W vs. 1W:  $P = 0.222$ ). Between-groups differences at 6-months follow-up of LVEF change of 4.3 [95% confidence limit: -4.6; 13.2]% and for LVEF change of 4.7 [95% confidence limit: -2.9; 12.4]% in favour of the 8W groups were observed.

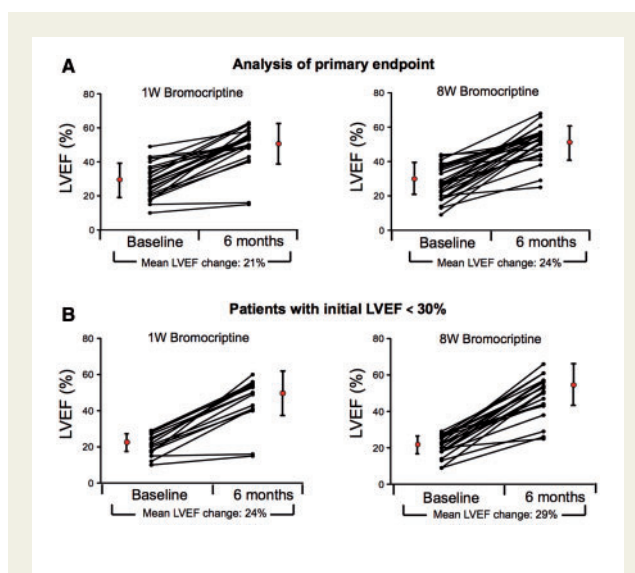
## Recovery rate of left ventricular function in peripartum cardiomyopathy patients after 6-months follow-up

After 6 months follow-up 52% of the 1W-patients ( $n = 32$ ) showed full functional LV recovery (LVEF  $\geq 50\%$ ), 21% partial recovery (LVEF between 35% and  $<50\%$ ) and 28% no recovery (LVEF  $<35\%$ ), prematurely terminated the trial or had missing LVEF data. In the 8W arm ( $n = 31$ ), the respective rates were 68% for full functional recovery, 25% for partial recovery, and 7% for no recovery. Full recovery rates showed a descriptive benefit for the 8W group (68% compared with 52% in 1W group).

Fisher exact test (two-sided) resulted in  $P = 0.283$  and revealed an OR = 0.508 [confidence limit: 0.173, 1.490] showing that—although not significant—the risk within the 8W group to fail ‘full recovery’ after 6 months is reduced.

## Change in left ventricular function over time as assessed by echocardiography

We also analysed LVEF determined by echocardiography, which was performed immediately after patients’ inclusion (Visit 1) with sequential performance at 2 weeks (Visit 2), 1 month (Visit 3), 3 months (Visit 4) and 6 months (Visit 5) after randomization (Figure 3B). Echocardiographic recordings meeting the quality standards of the core lab were available from 45 patients displaying a mean baseline LVEF of  $23 \pm 7\%$ . There was no difference in baseline LVEF measured by echocardiography between the 1W ( $23 \pm 7\%$ ,  $n = 21$ ) and the 8W ( $23 \pm 8\%$ ,  $n = 24$ ) group. After the 6 months follow-up, the absolute improvement (delta LVEF) in the 1W group was  $+25 \pm 12\%$  and  $+27 \pm 11\%$  in the 8W group, which was not statistically significant ( $P = 0.581$ ).



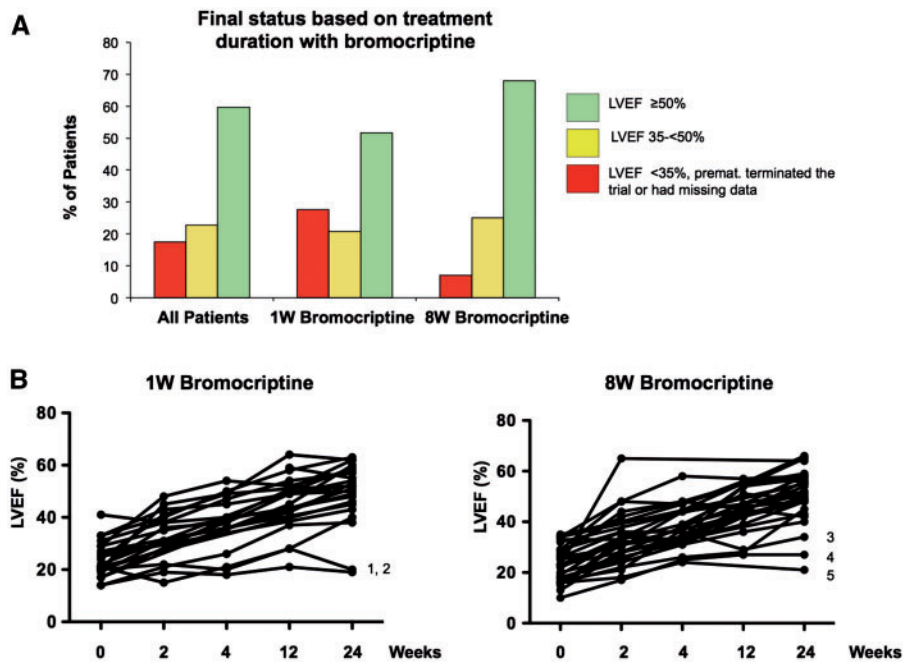
**Figure 2** Analyses of global left ventricular ejection fraction (LVEF) change from baseline to 6 months follow-up determined by CMR. (A) Individual courses of LVEF change from baseline to 6-months follow-up in the 1W group ( $n = 23$ ) and 8W group ( $n = 28$ ) with a between-groups difference at 6-months follow-up of 2.0% in favour of the 8W groups ( $P = 0.38$ ). (B) Individual courses of LVEF change from baseline to 6-months follow-up for the subgroup of patients with LVEF  $<30\%$  at study entry in the 1W group ( $n = 14$ ) and the 8W group ( $n = 18$ ) with between-groups differences at 6-months follow-up of 4.3% and for LVEF change of 4.7% in favour of the 8W groups ( $P = 0.22$ ).

## Secondary end points of study outcome

Secondary end points were hospitalization for heart failure, cardiac transplantation, and death of patient during trial, or combinations of these conditions (Table 2). Among the patients of the 1W group 3 of 31 (=9.7%) and among the patients of the 8W group 2 of 31 (=6.5%) were hospitalized for heart failure until the end of study (Table 2). Only one patient was listed for transplantation during the trial but was removed from the waiting list at follow-up due to improvement of the clinical condition and cardiac function. None of the patients received a left ventricular assist device (LVAD) or a heart transplantation, and no patient died. Altogether, the secondary end points of clinical events did not significantly differ between treatment groups ( $P = 0.651$ ).

## Safety of study treatment

A total of six serious adverse events (SAEs) in four patients were reported, and all SAEs occurred in patients of the 1W group (Table 3). Therefore, no dose relationship can be concluded from the results. For three of the SAEs, causality was considered possible with respect to treatment with bromocriptine. In all three cases, SAE occurred after termination of bromocriptine therapy. These SAEs included two venous embolisms and one peripheral artery occlusion. A complete list of all adverse events (AEs) and adverse reactions is provided in the Supplementary material online, Appendix Tables S12–S14.



**Figure 3** Outcome of patients at 6-months follow-up. (A) Left ventricular ejection fraction (LVEF) at 6-months follow-up according to predefined categories in all patients of the present study (treated 1W,  $n = 32$  or 8W,  $n = 31$  with bromocriptine, baseline LVEF <35%). Red columns illustrate the percentage of patients with no recovery (event or final LVEF <35%, prematurely terminated the trial or had missing LVEF data), yellow columns illustrate the percentage of patients with partial recovery (final LVEF 35% to <50%) and green columns depict percentage of women with complete recovery (final LVEF  $\geq 50\%$ ). (B) Step-wise change in LVEF measured by echocardiography during follow-up period in the 1W ( $n = 21$ ) and the 8W group ( $n = 24$ ). The number 1–5 marks time course of the five patients who did not recover LVEF >35% after 6-months. However, after  $\geq 12$  months Number 1 displayed a LVEF = 62%, 2 a LVEF = 47%, 3 and 4 a LVEF = 50%, and 5 a LVEF = 15%.

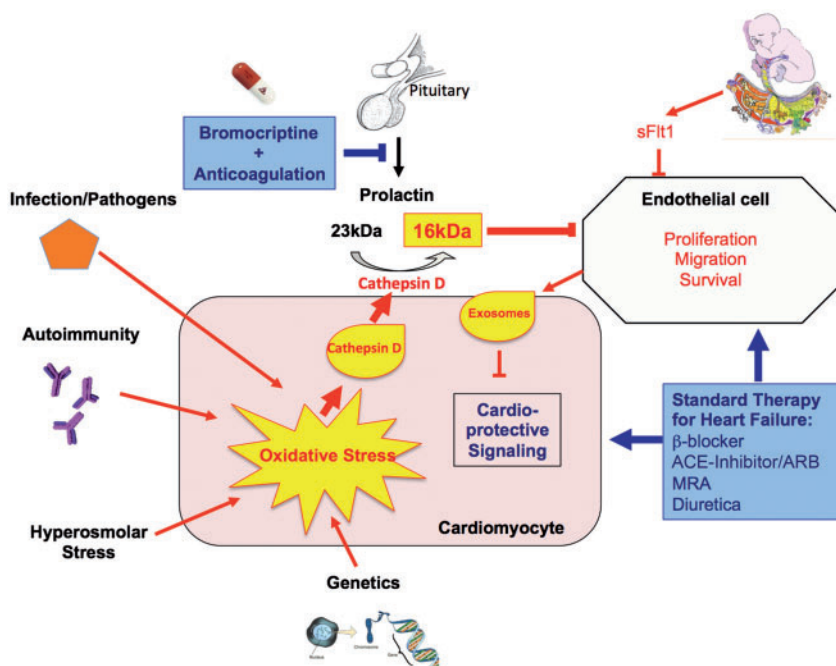
### Comparison of outcome with IPAC study

As a limitation of our study, we do not have a placebo control group because it was considered unethical given the results of previous registry data and pilot studies<sup>12,16,17</sup> and the risk for mastitis for stopping lactation without medical support. We therefore compared a subgroup of our collective characterized by a baseline LVEF <30% (echocardiography) with a subgroup of PPCM patients with a baseline LVEF <30% (echocardiography) from the Investigation in Pregnancy Associate Cardiomyopathy (IPAC) collective, which has been analysed over the same time period and obtaining comparable standard therapy for heart failure but no bromocriptine.<sup>19</sup> In the IPAC subgroup ( $n = 27$  patients), 37% remained in severe heart failure after a follow-up period of 6–12 months. In addition, the rate of transplantation and/or use of LVAD was 19%, and the mortality rate was 15% (Table 4). In the present study, 37 patients ( $n = 18$  in the 1W and  $n = 19$  in the 8W group) displayed a baseline LVEF <30% with at least 6-months follow-up. No patient in the 1W or the 8W group died, needed heart transplantation or a LVAD (Table 4). In the 1W group two and in the 8W group three patients did not recover LVEF above 35% after 6 months. Follow-up analyses after  $\geq 12$  months were available for these five patients showing that three patients displayed complete recovery (LVEF  $\geq 50\%$ ), and one patient showed substantial recovery with an LVEF of 47% and only one patient in the

8W group remained in severe heart failure (Table 4). However, this patient showed poor adherence to standard heart failure medication during follow-up.

### Discussion

Our study is the first prospective, randomized, and multicentre trial at adequate size to test short- vs. long-term inhibition of prolactin by bromocriptine as a causal therapy for PPCM. It demonstrates that inhibition of prolactin release with long-term or short-term bromocriptine in addition to standard therapy for heart failure in patients with severe forms of PPCM is associated with a high recovery rate and very low rate of adverse outcome. In fact no patient died or needed a heart transplantation or a LVAD and full recovery rate was higher than in any ever published PPCM collective.<sup>3,6,16,20–26</sup> In addition, our study population tolerated bromocriptine well suggesting that both, short- and long-term application appear to be safe since attention was paid to ensure sufficient anticoagulant therapy during treatment. The study did not detect a significant benefit for prolonged inhibition of prolactin release with bromocriptine in addition to standard therapy for heart failure in increasing LVEF or reducing hospitalization for heart failure compared with short-term



**Figure 4** Disease specific therapy with bromocriptine. Scheme depicting inducers of increased oxidative stress in the peripartum maternal heart that mediate the generation of antiangiogenic 16kDa prolactin from the nursing hormone prolactin which drives PPCM as well as the hypothetical mechanisms how the prolactin release blocker bromocriptine together with standard heart failure medication interferes with it.

**Table 2** Effect of treatment on secondary end points

End point	1W bromocriptine (N = 31) <sup>a</sup>	8W bromocriptine (N = 31)	P-value
Hospitalization for heart failure	3 (9.7)	2 (6.5)	0.651
Cardiac transplantation	0 (0)	0 (0)	n.a.
Death of patient during trial period	0 (0)	0 (0)	n.a.
Combination of hospitalization for heart failure, cardiac transplantation or death	3 (9.7)	2 (6.5)	0.651

Data are numbers of events (%).

<sup>a</sup>One patient of the 1W group who was not treated according to the protocol but received higher dose of bromocriptine (up to 10 mg) for a relevantly longer time was excluded from the endpoint analysis.

bromocriptine application sufficient to stop lactation in patients with PPCM. However, we observed a trend for more patients reaching full recovery after 6 months in the 8W group suggesting a small benefit of prolonged bromocriptine treatment.

Despite advances in understanding the pathomechanisms of PPCM, clinical trials testing disease-specific therapeutics beyond standard therapy for heart failure in this condition are scarce.<sup>2,4,8–10</sup> The nursing hormone prolactin and specifically its cleaved 16kDa form is considered to play a key role in the pathophysiology of PPCM (Figure 4).<sup>12,13,16,17</sup> However, unlike the placebo controlled study by Sliwa et al.<sup>17</sup> the present study was not designed to compare bromocriptine therapy with placebo but to compare two different dose

regimes of bromocriptine in severely diseased PPCM patients: A short-term regime that is sufficient to suppress lactation and a long-term regime that may exert additional cardioprotective effects.

The primary end point of our study, the global improvement in LVEF at 6 months, was not significantly different among the short-term and long-term bromocriptine groups, although the long-term treatment descriptively showed a higher rate of predefined full LV recovery. It is important to note, that the overall outcome among all patients enrolled in this study was better than the outcome of any prospective study on PPCM reported so far: Only 7% of all patients remaining in the study with complete data sets were still in severe LV dysfunction (EF <35%) at 6-months follow-up and only 3% remained

**Table 3** Incidence of serious adverse events during the study by system organ class

System organ class	1W bromocriptine (N = 32) <sup>a</sup>	Relation to bromocriptine	8W bromocriptine (N = 31)	P-value
Cardiac disorders				
Coronary artery occlusion <sup>b</sup>	1 (3.2)	Not related	0(0)	1.000
Musculoskeletal disorders				
Chest pain	1 (3.2)	Unlikely related	0(0)	1.000
Respiratory disorders				
Dyspnoea	1 (3.2)	Unlikely related	0(0)	1.000
Vascular disorders				
Venous Embolism	2 (6.4)	Possibly related	0(0)	0.491
Peripheral artery occlusion	1 (3.2)	Possibly related	0(0)	1.000

Data are numbers of events (%).

<sup>a</sup>The one patient of the 1W group who was not treated according to the protocol but received higher dose of bromocriptine (up to 10 mg) for a relevantly longer time was included in the safety analyses.

<sup>b</sup>Occlusion was due to air embolism as a complication during coronary angiography.

**Table 4** Effect of treatment on outcome in peripartum cardiomyopathy patients with left ventricular ejection fraction <30% in the bromocriptine study (treated either with 1W or 8W bromocriptine) compared with the IPAC study without bromocriptine treatment<sup>19</sup>

Follow-up characteristics	1W bromocriptine baseline LVEF <30% (n = 18)	8W bromocriptine baseline LVEF <30% (n = 19)	1W and 8W bromocriptine baseline LVEF <30% (n = 37)	IPAC study placebo baseline LVEF <30% (n = 27)
LVEF <35%	0% (0/18)	5% (1/19)	3% (1/37)	37%
LVEF 35–49%	22% (6/18)	37% (7/19)	35% (13/37)	26%
Full recovery, LVEF ≥50	67% (12/18)	58% (11/19)	62% (23/37)	37%
LVAD and HTX	0% (0/18)	0% (0/19)	0% (0/37)	19% (5/27)
Death	0% (0/18)	0% (0/19)	0% (0/37)	15% (4/27)

LVEF was analysed by echocardiography in the core labs of both studies. Follow-up in the IPAC study was 12 months, follow-up in our study was at 6–36 months. LVEF, left ventricular ejection fraction; LVAD, left ventricular assist device; HTX, heart transplantation.

in heart failure thereafter. Moreover, none of the patients experienced a major adverse event including death, cardiac transplantation or LVAD implantation. It has to be noted that all patients were randomized no longer than 7 days after first diagnosis in a severe disease state (LVEF ≤35%), thereby no selection for more stable patients was ensured. In a comparable group of patients in Germany, we previously reported a 15% treatment failure in a German registry with prospective data of 96 PPCM patients despite standard heart failure drug therapy.<sup>1,16</sup> Importantly, in the German registry the percentage of patients with persistent severe LV dysfunction was significantly higher among those patients who did not receive bromocriptine. The rate of maternal major adverse events among all patients in the German registry was 10% with a mortality rate of 2%. Interestingly, in the recent prospective North American IPAC study including PPCM patients with LVEF <45% a comparable event rate of 7% and a mortality rate of about 4% were reported.<sup>19</sup> In the IPAC study a subgroup of severely diseased patients with an LVEF <30% was analysed of whom 37% reached an LVEF ≥50% (full recovery) and an equal number, 37% experienced a major event (transplantation/LVAD: 19% or death: 15%) or a final LVEF <35%.<sup>19</sup> We

identified a similar subgroup in our study that presented with a baseline LVEF <30%, that however displayed a substantially better outcome (62% full recovery and 0% LVAD, HTX or death, only 3% with persistently LVEF <35%). Although this indirect comparison has to be interpreted with caution given some differences in the population characteristics, in particular the proportion of patients with African origin (2% in our study vs. 27% in the IPAC study) for whom prognosis might be worse,<sup>21</sup> it seems to further support the view that there is a benefit of bromocriptine treatment. In fact, also in non-African countries, i.e. for example in Turkey, mortality rates for PPCM between 25 and 30% were reported.<sup>20</sup> In view of these studies, a treatment concept that combines short-term low dose bromocriptine with standard heart failure therapy, as evaluated in the present study, appears to be associated with a better outcome, although a number of factors have to be taken into account which may have an impact on patient outcome, such as different ethnicities with potentially different genetic dispositions and different medical care opportunities.

An important reason for the beneficial effect of bromocriptine treatment may also be the associated ab lactation which enables early

optimal heart failure treatment with beta-blockade, ACE inhibition, MRA, and ivabradine.<sup>2,8,27</sup> No controlled studies have ever analysed the potential adverse effects of heart failure medication transferred to the infant in the breast milk, whereas normal growth percentiles and no adverse outcome were observed for the infants of PPCM patients in South Africa who terminated breastfeeding.<sup>17</sup> Thus, terminating breastfeeding with bromocriptine appears to be safe for the child and may enable implementation of early intensive heart failure therapy at high dosages associated with faster recovery of the mother enabling her to better care for her newborn.

Some concerns have been raised about a potential risk for cerebral and cardiovascular complications in patients treated with high doses of bromocriptine.<sup>28,29</sup> However, our observation that no adverse event associated with bromocriptine occurred during bromocriptine treatment in both groups suggests that bromocriptine (together with anticoagulant therapy) is safe and potentially effective in PPCM patients although the optimal dosage and duration of therapy to achieve maximal cardioprotective effects still remains a matter of investigation. Future studies should also explore whether extended prophylactic anticoagulation beyond the period of bromocriptine treatment is required in these patients with a considerable risk of thromboembolism.

Recent nation-wide observations and population-based studies estimated a rise in incidence of PPCM worldwide,<sup>5–7</sup> emphasizing that future research should focus on investigating preventive strategies. Moreover, modern echocardiography and cardiac magnetic resonance imaging have broadened the phenotypic profile of PPCM showing for example, that in about one-third of PPCM patients the right ventricle is affected by the disease with lower likelihood to recover despite optimal therapy.<sup>30</sup> The phenotypic heterogeneity of this disease with variable outcome may reflect additional contributing factors such as genetic or (auto-)immune mechanisms<sup>31,32</sup> that may determine the course of disease and the clinical outcome. Thus, the challenge of future research will be to elaborate specific therapeutic concepts for patients who are likely to be refractory to current treatment options. Since improvement of LV function was similar between the two bromocriptine regimens, an outcome study testing the concept appears worthwhile. Currently, our own experience suggests that critically ill patients (baseline LVEF <25%, cardiogenic shock) may profit from a prolonged treatment with a higher initial dosage of bromocriptine (see Supplementary material online, Figure S1), a feature that is also supported by the trend for faster full-recovery in the 8W group. This hypothesis needs to be scrutinized in a prospective randomized outcome trial.

In conclusion, the findings of our study further support a potential benefit of bromocriptine in addition to standard heart failure therapy and best supportive care in PPCM patients (Figure 4). It appears that a short low dose bromocriptine therapy aiming to stop lactation is sufficient in most forms of PPCM. However, the limited numbers of patients available in single countries warrant large international prospective registries, i.e. PPCM EORP registry [ESC EURObservational Research Programme (<http://www.eorp.org>)], to get further insights into the efficiency of treatment strategies.

## Manufacturer of the study drug

AbZ-Pharma GmbH

## Supplementary material

Supplementary material is available at *European Heart Journal* online.

## Acknowledgements

We thank S. Gutzke, B. Ritter, S. Tammen, M. Kasten, and M. Jungesblut for technical assistance.

## Funding

This trial was funded by the German Federal Ministry of Education and Research (BMBF), the cluster of excellence Rebirth II (German Research Foundation), and the Foundation Leducq and the 'Junge Akademie' Programme of the Hannover Medical School to A.H. ClinicalTrials.gov number: NCT00998556.

**Conflict of interest:** None of the members of the study team and study committees has a conflict of interest or financial interest, arrangement or affiliation with the manufacturer of the study drug. *Steering Committee:* D.H.-K. (authorized representative of the sponsor), J.B. (coordinating investigator), A.H. (representative of the coordinating investigator), and M.B. (University of Saarland). *Safety Monitoring Board Members:* J. McMurray (University of Glasgow), A. Koch (Hannover Medical School), G. P. Meyer (Hannover) and N. Marx (University of Aachen). *Clinical Event Committee:* Stefan Anker (Göttingen), Kenneth Dickstein (University of Bergen) and Dirk J. van Veldhuisen (University of Groningen), *Study management Clinical Trial Centre* at the University of Leipzig (ZKS Leipzig).

## References

1. Sliwa K, Fett J, Elkayam U. Peripartum cardiomyopathy. *Lancet* 2006;**368**:687–693.
2. Hilfiker-Kleiner D, Haghikia A, Nonhoff J, Bauersachs J. Peripartum cardiomyopathy: current management and future perspectives. *Eur Heart J* 2015;**36**:1090–1097.
3. Hilfiker-Kleiner D, Sliwa K. Pathophysiology and epidemiology of peripartum cardiomyopathy. *Nat Rev Cardiol* 2014;**11**:364–370.
4. Sliwa K, Hilfiker-Kleiner D, Petrie MC, Mebazaa A, Pieske B, Buchmann E, Regitz-Zagrosek V, Schaufelberger M, Tavazzi L, van Veldhuisen DJ, Watkins H, Shah AJ, Seferovic PM, Elkayam U, Pankuweit S, Papp Z, Mouquet F, McMurray JJ. Current state of knowledge on aetiology, diagnosis, management, and therapy of peripartum cardiomyopathy: a position statement from the Heart Failure Association of the European Society of Cardiology Working Group on peripartum cardiomyopathy. *Eur J Heart Fail* 2010;**12**:767–778.
5. Brar SS, Khan SS, Sandhu GK, Jorgensen MB, Parikh N, Hsu JW, Shen AY. Incidence, mortality, and racial differences in peripartum cardiomyopathy. *Am J Cardiol* 2007;**100**:302–304.
6. Kolte D, Khera S, Aronow WS, Palaniswamy C, Mujib M, Ahn C, Jain D, Gass A, Ahmed A, Panza JA, Fonarow GC. Temporal trends in incidence and outcomes of peripartum cardiomyopathy in the United States: a nationwide population-based study. *J Am Heart Assoc* 2014;**3**:e001056.
7. Sliwa K, Boehm M. Incidence and prevalence of pregnancy associated heart disease. *Cardiovasc Res* 2014;**101**:554–560.
8. Bauersachs J, Arrigo M, Hilfiker-Kleiner D, Veltmann C, Coats AJ, Crespo-Leiro MG, De Boer RA, van der Meer P, Maack C, Mouquet F, Petrie MC, Piepoli MF, Regitz-Zagrosek V, Schaufelberger M, Seferovic P, Tavazzi L, Ruschitzka F, Mebazaa A, Sliwa K. Current management of patients with severe acute peripartum cardiomyopathy: practical guidance from the Heart Failure Association of the European Society of Cardiology Study Group on peripartum cardiomyopathy. *Eur J Heart Fail* 2016;**18**:1096–1105.
9. Ponikowski P, Voors AA, Anker SD, Bueno H, Cleland JG, Coats AJ, Falk V, Gonzalez-Juanatey JR, Harjola VP, Jankowska EA, Jessup M, Linde C, Nihoyannopoulos P, Parissis JT, Pieske B, Riley JP, Rosano GM, Ruilope LM, Ruschitzka F, Rutten FH, van der Meer P; Authors/Task Force M, Document R. 2016 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure: the Task Force for the diagnosis and treatment of acute and chronic heart failure of the European Society of Cardiology (ESC). Developed with the special contribution of the Heart Failure Association (HFA) of the ESC. *Eur J Heart Fail* 2016;**18**:891–975.



10. Regitz-Zagrosek V, Blomstrom Lundqvist C, Borghi C, Cifkova R, Ferreira R, Foidart JM, Gibbs JS, Gohlke-Baerwolf C, Gorenek B, Iung B, Kirby M, Maas AH, Morais J, Nihoyannopoulos P, Pieper PG, Presbitero P, Roos-Hesselink JW, Schaufelberger M, Seeland U, Torracca L, Bax J, Auricchio A, Baumgartner H, Ceconi C, Dean V, Deaton C, Fagard R, Funck-Brentano C, Hasdai D, Hoes A, Knuuti J, Kolh P, McDonagh T, Moulin C, Poldermans D, Popescu BA, Reiner Z, Sechtem U, Sirnes PA, Torbicki A, Vahanian A, Windecker S, Aguiar C, Al-Attar N, Garcia AA, Antoniou A, Coman I, Elkayam U, Gomez-Sanchez MA, Gotcheva N, Hilfiker-Kleiner D, Kiss RG, Kitsiou A, Konings KT, Lip GY, Manolis A, Mebazaa A, Mintale I, Morice MC, Mulder BJ, Pasquet A, Price S, Puri SG, Salvador MJ, Shotan A, Silversides CK, Skouby SO, Stein JJ, Tornos P, Vejlstrup N, Walker F, Warnes C. ESC Guidelines on the management of cardiovascular diseases during pregnancy: the Task Force on the Management of Cardiovascular Diseases during Pregnancy of the European Society of Cardiology (ESC). *Eur Heart J* 2011;**32**:3147–3197.
11. Halkein J, Tabruyn SP, Ricke-Hoch M, Haghikia A, Nguyen NQ, Scherr M, Castermans K, Malvaux L, Lambert V, Thiry M, Sliwa K, Noel A, Martial JA, Hilfiker-Kleiner D, Struman I. MicroRNA-146a is a therapeutic target and biomarker for peripartum cardiomyopathy. *J Clin Invest* 2013;**123**:2143–2154.
12. Hilfiker-Kleiner D, Kaminski K, Podewski E, Bonda T, Schaefer A, Sliwa K, Forster O, Quint A, Landmesser U, Doerries C, Luchtefeld M, Poli V, Schneider MD, Balligand JL, Desjardins F, Ansari A, Struman I, Nguyen NQ, Zschemisch NH, Klein G, Heusch G, Schulz R, Hilfiker A, Drexler H. A cathepsin D-cleaved 16 kDa form of prolactin mediates postpartum cardiomyopathy. *Cell* 2007;**128**:589–600.
13. Patten IS, Rana S, Shahul S, Rowe GC, Jang C, Liu L, Hacker MR, Rhee JS, Mitchell J, Mahmood F, Hess P, Farrell C, Koullis N, Khankin EV, Burke SD, Tudorache I, Bauersachs J, del Monte F, Hilfiker-Kleiner D, Karumanchi SA, Arany Z. Cardiac angiogenic imbalance leads to peripartum cardiomyopathy. *Nature* 2012;**485**:333–338.
14. Ricke-Hoch M, Bultmann I, Stapel B, Condorelli G, Rinas U, Sliwa K, Scherr M, Hilfiker-Kleiner D. Opposing roles of Akt and STAT3 in the protection of the maternal heart from peripartum stress. *Cardiovasc Res* 2014;**101**:587–596.
15. Mejia-Rodriguez O, Herrera-Abarca JE, Ceballos-Reyes G, Avila-Diaz M, Prado-Urbe C, Belio-Caro F, Salinas-Gonzalez A, Vega-Gomez H, Alvarez-Aguilar C, Lindholm B, Garcia-Lopez E, Paniagua R. Cardiovascular and renal effects of bromocriptine in diabetic patients with stage 4 chronic kidney disease. *Biomed Res Int* 2013;**2013**:104059.
16. Haghikia A, Podewski E, Libhaber E, Labidi S, Fischer D, Roentgen P, Tsikas D, Jordan J, Lichtinghagen R, von Kaisenberg CS, Struman I, Bovy N, Sliwa K, Bauersachs J, Hilfiker-Kleiner D. Phenotyping and outcome on contemporary management in a German cohort of patients with peripartum cardiomyopathy. *Basic Res Cardiol* 2013;**108**:366.
17. Sliwa K, Blauwet L, Tibazarwa K, Libhaber E, Smedema JP, Becker A, McMurray J, Yamac H, Labidi S, Struman I, Hilfiker-Kleiner D. Evaluation of bromocriptine in the treatment of acute severe peripartum cardiomyopathy: a proof-of-concept pilot study. *Circulation* 2010;**121**:1465–1473.
18. Haghikia A, Podewski E, Berliner D, Sonnenschein K, Fischer D, Angermann CE, Bohm M, Rontgen P, Bauersachs J, Hilfiker-Kleiner D. Rationale and design of a randomized, controlled multicentre clinical trial to evaluate the effect of bromocriptine on left ventricular function in women with peripartum cardiomyopathy. *Clin Res Cardiol* 2015;**104**:911–917.
19. McNamara DM, Elkayam U, Alharethi R, Damp J, Hsieh E, Ewald G, Modi K, Alexis JD, Ramani GV, Semigran MJ, Haythe J, Markham DW, Marek J, Gorcsan J 3rd, Wu WC, Lin Y, Halder I, Pisarcik J, Cooper LT, Fett JD; Investigators I. Clinical outcomes for peripartum cardiomyopathy in North America: results of the IPAC Study (Investigations of Pregnancy-Associated Cardiomyopathy). *J Am Coll Cardiol* 2015;**66**:905–914.
20. Biteker M. Peripartum cardiomyopathy in Turkey. *Int J Cardiol* 2012;**158**:e60–e61.
21. Blauwet LA, Libhaber E, Forster O, Tibazarwa K, Mebazaa A, Hilfiker-Kleiner D, Sliwa K. Predictors of outcome in 176 South African patients with peripartum cardiomyopathy. *Heart* 2013;**99**:308–313.
22. Fett JD. Peripartum cardiomyopathy. Insights from Haiti regarding a disease of unknown etiology. *Minn Med* 2002;**85**:46–48.
23. Fett JD, Christie LG, Carraway RD, Murphy JG. Five-year prospective study of the incidence and prognosis of peripartum cardiomyopathy at a single institution. *Mayo Clin Proc* 2005;**80**:1602–1606.
24. Kamiya CA, Kitakaze M, Ishibashi-Ueda H, Nakatani S, Murohara T, Tomoike H, Ikeda T. Different characteristics of peripartum cardiomyopathy between patients complicated with and without hypertensive disorders. Results from the Japanese Nationwide survey of peripartum cardiomyopathy. *Circ J* 2011;**75**:1975–1981.
25. Pillarisetti J, Kondur A, Alani A, Reddy M, Reddy M, Vacek J, Weiner CP, Ellerbeck E, Schreiber T, Lakkireddy D. Peripartum cardiomyopathy: predictors of recovery and current state of implantable cardioverter-defibrillator use. *J Am Coll Cardiol* 2014;**63**(25 Pt A):2831–2839.
26. Sliwa K, Forster O, Tibazarwa K, Libhaber E, Becker A, Yip A, Hilfiker-Kleiner D. Long-term outcome of peripartum cardiomyopathy in a population with high seropositivity for human immunodeficiency virus. *Int J Cardiol* 2011;**147**:202–208.
27. Haghikia A, Tongers J, Berliner D, König T, Schafer A, Brehm M, Bohm M, Hilfiker-Kleiner D, Bauersachs J. Early ivabradine treatment in patients with acute peripartum cardiomyopathy: subanalysis of the German PPCM registry. *Int J Cardiol* 2016;**216**:165–167.
28. Hopp L, Haider B, Iffy L. Myocardial infarction postpartum in patients taking bromocriptine for the prevention of breast engorgement. *Int J Cardiol* 1996;**57**:227–232.
29. Iffy L, O'donnell J, Correia J, Hopp L. Severe cardiac dysrhythmia in patients using bromocriptine postpartum. *Am J Ther* 1998;**5**:111–115.
30. Haghikia A, Rontgen P, Vogel-Claussen J, Schwab J, Westenfeld R, Ehlermann P, Berliner D, Podewski E, Hilfiker-Kleiner D, Bauersachs J. Prognostic implication of right ventricular involvement in peripartum cardiomyopathy: a cardiovascular magnetic resonance study. *ESC Heart Fail* 2015;**2**:139–149.
31. Haghikia A, Kaya Z, Schwab J, Westenfeld R, Ehlermann P, Bachelier K, Oettl R, von Kaisenberg CS, Katus HA, Bauersachs J, Hilfiker-Kleiner D. Evidence of auto-antibodies against cardiac troponin I and sarcomeric myosin in peripartum cardiomyopathy. *Basic Res Cardiol* 2015;**110**:60.
32. Ware JS, Li J, Mazaika E, Yasso CM, DeSouza T, Cappola TP, Tsai EJ, Hilfiker-Kleiner D, Kamiya CA, Mazzarotto F, Cook SA, Halder I, Prasad SK, Pisarcik J, Hanley-Yanez K, Alharethi R, Damp J, Hsieh E, Elkayam U, Sheppard R, Kealey A, Alexis J, Ramani G, Safirstein J, Boehmer J, Pauly DF, Wittstein IS, Thohan V, Zucker MJ, Liu P, Gorcsan J 3rd, McNamara DM, Seidman CE, Seidman JG, Arany Z; Imac, Investigators I. Shared genetic predisposition in peripartum and dilated cardiomyopathies. *N Engl J Med* 2016;**374**:233–241.