

Research

Open Access

A clinical evaluation committee assessment of recombinant human tissue factor pathway inhibitor (tifacogin) in patients with severe community-acquired pneumonia

Pierre-François Laterre¹, Steven M Opal², Edward Abraham³, Steven P LaRosa², Abla A Creasey⁴, Fang Xie⁵, Lona Poole⁵ and Richard G Wunderink⁶

¹St Luc University Hospital, Université Catholique de Louvain, Avenue Hippocrate 10, 1200 Brussels, Belgium

²Division of Infectious Diseases, Rhode Island Hospital, POB Suite #330, 593 Eddy Street, Providence, RI 02903, USA

³Department of Medicine, University of Alabama at Birmingham School of Medicine, 420 Boshell Building, 1808 7th Avenue South, Birmingham, AL 35294, USA

⁴Alza Corporation, Johnson & Johnson, 1900 Charleston Road, Mountain View, CA 94042, USA

⁵Novartis, 4560 Horton Street, Emeryville, CA 94608, USA

⁶Division of Pulmonary and Critical Care Medicine, Northwestern University Feinberg School of Medicine, 240 E. Huron Street, McGaw M300, Chicago, IL 60611, USA

Corresponding author: Pierre-François Laterre, laterre@rean.ucl.ac.be

Received: 8 Feb 2008 Revisions requested: 14 Mar 2008 Revisions received: 22 Oct 2008 Accepted: 15 Mar 2009 Published: 15 Mar 2009

Critical Care 2009, **13**:R36 (doi:10.1186/cc7747)

This article is online at: <http://ccforum.com/content/13/2/R36>

© 2009 Laterre *et al.*; licensee BioMed Central Ltd.

This is an open access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Introduction The purpose of this analysis was to determine the potential efficacy of recombinant human tissue factor pathway inhibitor (tifacogin) in a subpopulation of patients with community-acquired pneumonia (CAP) from a phase III study of severe sepsis.

Methods A retrospective review of patients with suspected pneumonia was conducted by an independent clinical evaluation committee (CEC) blinded to treatment assignment. The CEC reanalyzed data from patients enrolled in an international multicenter clinical trial of sepsis who had a diagnosis of pneumonia as the probable source of sepsis. The primary efficacy measure was all-cause 28-day mortality.

Results Of 847 patients identified on case report forms with a clinical diagnosis of pneumonia, 780 (92%) were confirmed by the CEC to have pneumonia. Of confirmed pneumonia cases, 496 (63.6%) met the definition for CAP. In the CEC CAP population, the mortality rates of the tifacogin and placebo

groups were 70/251 (27.9%) and 80/245 (32.7%), respectively. The strongest signals were seen in patients with CAP not receiving concomitant heparin, having microbiologically confirmed infection, or having the combination of documented infection and no heparin. The reduction in mortality in this narrowly defined subgroup when treated with tifacogin compared with placebo was statistically significant (17/58 [29.3%] with tifacogin and 28/54 [51.9%] with placebo; unadjusted *P* value of less than 0.02).

Conclusions Tifacogin administration did not significantly reduce mortality in any severe CAP patient. Exploratory analyses showed an improved survival in patients who did not receive concomitant heparin with microbiologically confirmed infections. These data support the rationale of an ongoing phase III study exploring the potential benefit of tifacogin in severe CAP.

Trial Registration ClinicalTrials.gov identifier NCT00084071.

Introduction

Sepsis is a systemic response to infection associated with significant mortality and substantial direct patient care costs [1]. Community-acquired pneumonia (CAP) is the most common

cause of sepsis [2-5]. CAP mortality rates are significant and have not changed significantly over several decades despite the availability of improved broad-spectrum antibiotics [6]. While successful outcome from severe CAP requires

APACHE II: Acute Physiology and Chronic Health Evaluation II; aPC: activated protein C; CAP: community-acquired pneumonia; CEC: clinical evaluation committee; CRF: case report form; HAP: hospital-acquired pneumonia; LPS: lipopolysaccharide; OPTIMIST: Optimized Phase III Tifacogin in Multicenter International Sepsis Trial; PCT: procalcitonin; TF: tissue factor; TFPI: tissue factor pathway inhibitor.

adequate treatment of the infection, antimicrobial agents alone have only limited capacity to reduce the mortality rate associated with severe CAP and adjunctive measures are required to treat organ dysfunction such as respiratory failure [6].

Likely contributors to organ dysfunction and death are intravascular and intrapulmonary generation of thrombin and deposition of fibrin due to break down in hemostatic regulation. Increased cell surface expression of tissue factor (TF) in severe CAP induces thrombin generation and fibrin formation [7,8]. TF expression in the lungs of pneumonia patients leads to a proinflammatory and procoagulant environment as well as to decreased fibrinolysis [9].

TF pathway inhibitor (TFPI) regulates coagulation initiated by TF. Expression of TF and TFPI is imbalanced in acute lung injury [10]. Administration of recombinant TFPI or factor VIIa antagonists reduces lung injury and systemic cytokine responses in infection models [11-14]. Therefore, TF inhibition may have beneficial effects in disease states such as acute lung injury or pneumonia in which coagulation and inflammation play prominent roles [9].

Safety and efficacy of tifacogin, a recombinant form of human TFPI, were assessed in a phase III study (TFP007 OPTIMIST [Optimized Phase III Tifacogin in Multicenter International Sepsis Trial]) in patients with severe sepsis [15]. Although efficacy of the primary endpoint of 28-day all-cause mortality was not shown, treatment benefit in a subset of patients with pneumonia with microbiological documentation and not receiving heparin within 24 hours prior to and/or during study drug infusion was observed in *post hoc* analysis. However, these analyses were based on case report forms (CRFs) in which investigators were allowed to list multiple sites of infection and any positive cultures. Not all positive cultures grew pathogens, and the organisms grown were not necessarily consistent with the suspected infection site.

Concern regarding the accuracy of subgroup classification in TFP007 prompted the creation of a clinical evaluation committee (CEC) to validate the CRF-based analyses. CECs have previously been engaged to evaluate negative trials of adjuvant agents in critical illness in order to determine a target population for further study [16,17]. The CEC was specifically charged with determining (a) the validity of the pneumonia diagnosis, (b) whether the pneumonia was CAP, hospital-acquired pneumonia (HAP), or other diagnoses, and (c) the level of evidence of a microbiological etiology of CAP.

Materials and methods

A detailed description of the study was previously published [15]. The OPTIMIST study was approved by the ethics committee of each individual participating center, and written consent was obtained from each patient or next of kin. The CEC retrospective study was approved by the ethics committee of

St Luc University Hospital (Brussels, Belgium). Initial analyses of the TFP007 patient subgroup with pneumonia used a programmatic definition of CAP that allowed a maximum of 2 days of hospitalization prior to the start of study drug for the pneumonia to be classified as CAP. Patients hospitalized longer than 2 days were classified as having HAP.

The CEC consisted of critical care, pulmonary disease, and infectious disease specialists who remained blinded to treatment throughout the evaluation. A charter incorporating a predetermined set of clinical and microbiological classification rules was used to ensure uniformity of this retrospective assessment [18]. Criteria to be classified as CAP included all five of the following: (a) the clinical and radiographic evidence was consistent with pneumonia, (b) microbiology (when provided) was consistent with a CAP pathogen, (c) the primary reason for hospital admission was pneumonia, (d) there was no evidence of aspiration or major immunocompromised state, and (e) the patient was not a known nursing home resident or transfer from another institution. Chest x-ray protocol provided by a radiologist at each investigator site was used to define evidence of CAP. In the CEC analysis, the CAP time window was expanded to 4 days between hospital admission and start of study drug infusion for cases with signs and symptoms of CAP on admission. This interval was chosen based on the time windows used for patient enrollment [19] and to include CAP patients who deteriorated after admission [20,21].

TFP007 investigators classified 847 patients as having CAP on the CRFs. Cases in which pneumonia was not listed by the investigator as a potential site of infection were not reviewed. The CEC reviewed all available information on CRFs from the locked TFP007 database for those subjects. Each case was independently reviewed by one member, and then the CEC met to reach a consensus on all problematic cases. No adjudication of any outcome data was performed.

The CEC assessment forms were tabulated, and the tifacogin arm was compared with placebo for all CEC-confirmed CAP cases. Additional analyses were carried out for microbiological evidence (based on culture only), heparin use, serious bleeding events and contributing causes, and the Acute Physiology and Chronic Health Evaluation II (APACHE II) score quartiles [4,22]. The results of the CEC evaluation were compared with those of the original programmatic classification.

Elevated procalcitonin (PCT) (>0.5 ng/mL) levels are associated with a bacterial etiology in patients presenting with suspected pulmonary infection [23]. PCT levels were measured in plasma specimens prospectively collected (Brahms AG, Hennigsdorf, Germany). The CEC classification was performed without knowledge of PCT values. PCT levels were evaluated in the CEC-designated CAP population. Chi-square tests were used to compare treatment groups for dichotomous variables. All *P* values are unadjusted for multiple testing. Logistic

Table 1**Baseline demographics for clinical evaluation committee community-acquired pneumonia patients**

	Tifacogin n = 251	Placebo n = 245	Selected P value
Age in years, mean \pm standard deviation	60.5 \pm 15.8	60.2 \pm 15.1	
Gender, number (percentage)			
Female	91 (36)	100 (41)	
Male	160 (64)	145 (59)	
Ethnicity, number (percentage)			
Caucasian	207 (82)	199 (81)	
Black	26 (10)	20 (8)	
Hispanic	9 (4)	12 (5)	
Asian	5 (2)	7 (3)	
Other	4 (2)	7 (3)	
Baseline APACHE II score			
Mean \pm standard deviation	25.6 \pm 7.0	25.2 \pm 6.7	
Baseline interleukin-6	434.6	478.9	
Geometric mean (95% CI)	(329.2, 573.6)	(355.1, 645.9)	
Baseline procalcitonin			
Geometric mean (95% CI)	8.89 (7.15, 11.05)	8.09 (6.35, 10.31)	
<2 ng/mL, number (percentage)	50 (20)	60 (25)	
Shock, number (percentage)	163 (65)	176 (72)	0.10
Ventilatory support, number (percentage)	193 (77)	200 (82)	0.19
Number of organ dysfunctions, number (percentage)			
Two or less	87 (35)	72 (29)	0.21
Three or more	164 (65)	173 (71)	
Heparin use, number (percentage)	172 (69)	167 (68)	
Organism identified, number (percentage)	170 (68)	154 (63)	

APACHE II, Acute Physiology and Chronic Health Evaluation II; CI, confidence interval.

regression models were also used to adjust for baseline APACHE II score, PCT level, shock, and use of ventilator support.

Results

Confirmation of pneumonia and community-acquired pneumonia diagnosis

In its review of 847 patients identified on CRFs with a diagnosis of pneumonia, the CEC concurred that pneumonia was the cause of sepsis in 780 cases (92%). One patient could not be evaluated. Of the 780 confirmed pneumonia cases, 496 were classified by the CEC as CAP (251 in the tifacogin group and 245 in the placebo group) and 259 were classified as HAP (132 in the tifacogin group and 127 in the placebo group). Because of a major immunocompromised state, aspiration, or radiation pneumonitis, 25 patients did not meet the standard definition of CAP and therefore were excluded. The CRF-

defined pneumonia subgroup identified pneumonia patients misclassified as having CAP when they actually had HAP and vice versa.

Demographics of the 496 CEC CAP patients are presented in Table 1. Sixty-eight percent received heparin and 65% had a microbiologically confirmed infection. Baseline characteristics in the placebo and tifacogin groups were similar. For both documented CAP and no-heparin-use subgroups, baseline APACHE II scores and presence of shock did not differ between TFPI and placebo.

The spectrum of etiologic microorganisms is presented in Table 2. *Streptococcus pneumoniae* was the most common pathogenic organism. Twelve cases had only sputum Gram stain evidence of pneumonia. Baseline PCT levels above 2 ng/mL were present in the majority (78%) of CEC CAP patients.

Table 2

Clinical evaluation committee classification of causative microorganisms in community-acquired pneumonia patients

Organism identified	Number (percentage)	Tifacogin n = 251		Placebo N = 245	
		Number (percentage)	Mortality, number (percentage)	Number (percentage)	Mortality, number (percentage)
<i>Streptococcus pneumoniae</i>	69 (27.5)	14 (20.3)	70 (28.6)	19 (27.1)	
<i>Staphylococcus aureus</i>	33 (13.1)	14 (42.4)	19 (7.8)	12 (63.2)	
<i>Haemophilus influenzae</i>	26 (10.4)	5 (19.2)	10 (4.1)	5 (50)	
Other respiratory organisms	24 (9.6)	9 (37.5)	23 (9.4)	6 (26.1)	
Enteric Gram-negative	20 (8.0)	6 (30)	16 (6.5)	9 (56.3)	
Gram stain only	9 (3.6)	3 (33.3)	9 (3.7)	4 (44.4)	
Legionella species	6 (2.4)	1 (16.7)	10 (4.1)	2 (20)	
<i>Pseudomonas aeruginosa</i>	6 (2.4)	1 (16.7)	12 (4.9)	5 (41.7)	
Respiratory viruses	2 (0.8)	1 (50)	0	0	
<i>Chlamydophila pneumoniae</i>	0	0	2 (0.8)	0	
No organism identified	81 (32.3)		91 (37.1)		

Only 60/245 (25%) of placebo and 50/251 (20%) tifacogin-treated patients had PCT levels of less than 2 ng/mL.

Effect of tifacogin in clinical evaluation committee community-acquired pneumonia cohort

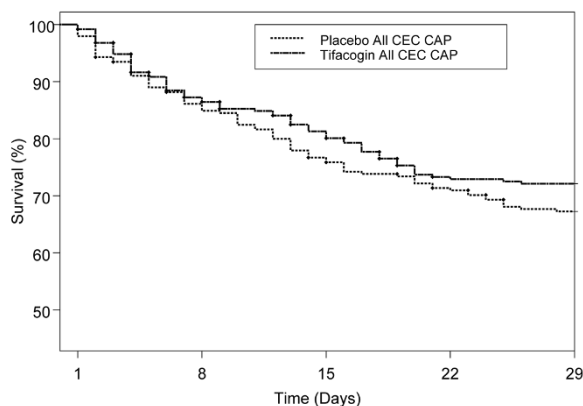
Kaplan-Meier plots of 28-day cumulative survival data for all CEC CAP patients (Figure 1) and for CEC CAP subjects who did not receive heparin and who also had microbiological evidence of an infectious etiology for their pneumonia (Figure 2) are shown. The CEC CAP patients treated with tifacogin had lower 28-day all-cause mortality compared with the placebo group (27.9% versus 32.7%; $P = 0.25$, Pearson chi-square test; $P = 0.22$, logistic regression model). The largest difference in 28-day mortality (Table 3 and Figure 2) occurred in the

subgroup of patients with microbiological evidence of infection in the no-heparin CAP cohort (29.3%, tifacogin; 51.9%, placebo; $P = 0.02$, Pearson chi-square test; $P = 0.02$, logistic regression model).

Effect of pathogen class and causative microorganism on mortality

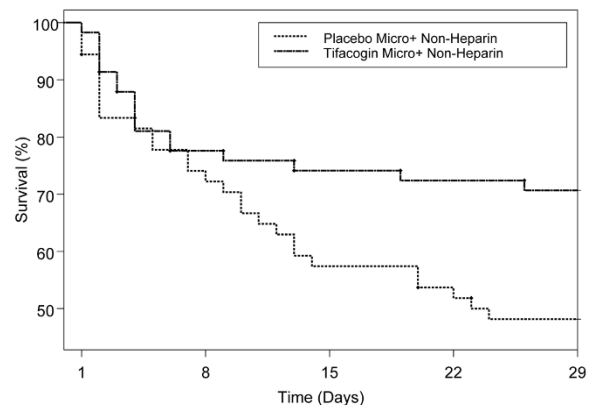
The observed mortality of the tifacogin-treated group was lower than that of the placebo group for all pathogen classes (Gram-positive, Gram-negative, mixed, and other) (Figure 3) and when analyzed by individual pathogen, except unusual respiratory pathogens and respiratory viruses (Table 2). For *S. pneumoniae*, the observed 28-day mortality in tifacogin-treated subjects was 20.3% versus 27.1% in the placebo arm.

Figure 1



Kaplan-Meier survival analysis for all clinical evaluation committee (CEC) community-acquired pneumonia (CAP) patients. P value = 0.25.

Figure 2



Kaplan-Meier survival analysis for clinical evaluation committee community-acquired pneumonia patients in the non-heparin cohort with microorganism identified. P value = 0.02.

Table 3**Mortality (28-day) in tifacogin and placebo groups for all patients and by microbiology status and heparin use**

	Tifacogin		Placebo		P value
	All Number	Mortality Number Percentage	All Number	Mortality Number Percentage	
All community-acquired pneumonia patients	251	70 27.9	245	80 32.7	0.25
Microbiology status					
Organism identified	170	46 27.1	154	55 35.7	0.09
Organism not identified	81	24 29.6	91	25 27.5	0.75
Procalcitonin level					
<2	50	15 30.0	60	18 30.0	1.00
≥ 2	200	55 27.5	183	62 33.9	0.18
Heparin use					
No Heparin	79	23 29.1	78	33 42.3	0.08
Heparin	172	47 27.3	167	47 28.1	0.87
Microbiology status and heparin use					
No heparin/Organism identified	58	17 29.3	54	28 51.9	0.02
No heparin/Organism not identified	21	6 28.6	24	5 20.8	0.54
Shock					
Yes	163	50 30.7	176	64 36.4	0.27
No	88	20 22.7	69	16 23.2	0.95
Ventilatory support					
Yes	193	60 31.1	200	75 37.5	0.18
No	58	10 17.2	45	5 11.1	0.38
Number of organ dysfunctions					
Two or less	87	17 19.5	72	13 18.1	0.81
Three or more	164	53 32.3	173	67 38.7	0.22

When analyzed by PCT levels of less than 2 ng/mL and of greater than or equal to 2 ng/mL, the observed mortality in the tifacogin-treated cohort was improved in subjects with higher PCT levels (Table 3).

Tifacogin treatment effect and APACHE II score

Based on PROWESS (Recombinant Human Activated Protein C Worldwide Evaluation in Severe Sepsis) [4,22], CEC CAP patients were segregated into quartiles of APACHE II scores of less than 20 (n = 91), 20 to 24 (n = 149), 25 to 29 (n = 127), and greater than 29 (n = 128) (Figure 4). In both tifacogin and placebo groups, mortality increased with higher APACHE II scores. Mortality for patients receiving tifacogin was lower than mortality in the placebo group in all four APACHE II score quartiles.

Safety

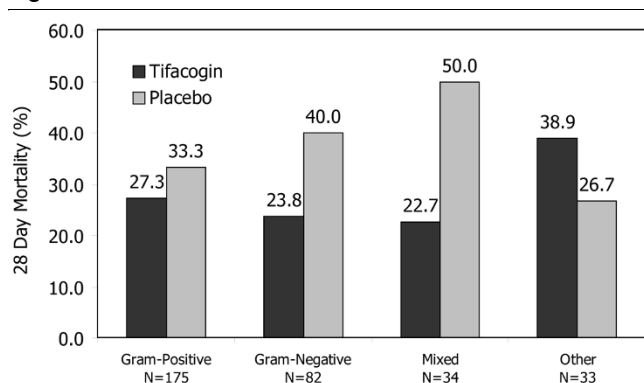
The overall incidence of all adverse events was similar in the tifacogin group (93%) and the placebo group (91%). Serious adverse event rates, likewise, were similar (41% versus 52%,

respectively). Since tifacogin is an anticoagulant, the incidence of events involving bleeding was scrutinized. Tifacogin-treated CAP patients had higher rates of bleeding events (23% versus 18%) and serious bleeding events (6% versus 2%) compared with placebo CAP patients. The most common sites of bleeding were the gastrointestinal and respiratory tracts in the tifacogin group and the gastrointestinal tract and skin (ecchymoses) in the placebo group. Higher rates of bleeding events occurred in subgroups receiving concomitant heparin (Tables 4 and 5) than in those not treated with heparin.

Discussion

Retrospective subgroup analyses may identify potential target populations for future trials. The OPTIMIST trial [15] showed no improvement in mortality with tifacogin compared with placebo. Overall, the 28-day survival in patients with CAP treated with tifacogin was higher compared with placebo but the difference did not reach statistical significance. However, subgroup analysis of this study suggested that patients not receiving heparin and/or with microbiological evidence of

Figure 3



Mortality by bacterial Gram stain morphology in clinical evaluation committee community-acquired pneumonia patients.

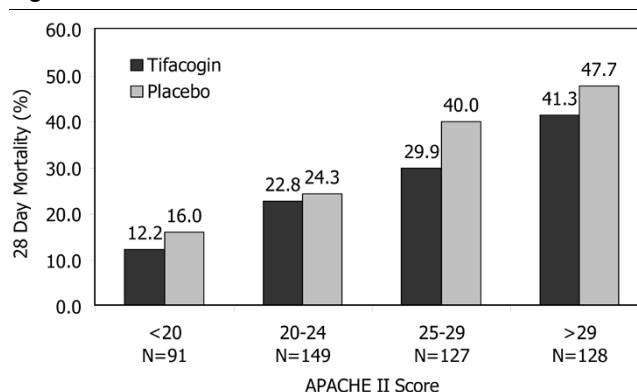
pneumonia appeared to benefit from tifacogin. Using blinded and stringent evaluations, the CEC strengthened the database used for reanalysis, demonstrating an important role for CECs in retrospective review. The CEC analysis corroborated the initial analysis by showing a reduction in mortality in the tifacogin-treated CAP subgroup, not receiving heparin, with microbiologically confirmed infection, or when these two conditions were present.

Benefit of an agent affecting the coagulation pathway in a population with pneumonia versus other sources of infection has biological plausibility. In animal models of acute bronchopneumonia, activation of coagulation can be readily demonstrated [24,25]. Bronchoalveolar lavage specimens from patients with acute lung injury also indicate activation of coagulation [26]. Recombinant human activated protein C (aPC), an anticoagulant approved for the treatment of severe sepsis, had its greatest benefit in the population with severe CAP in a similar CEC analysis [19].

PCT has been shown to be consistently elevated in bacterial infections [23,27]. The beneficial effect of tifacogin in patients with levels above 2 ng/mL reinforces the need for the phase III confirmatory study to emphasize documented bacterial CAP. Both microbiological data and the PCT levels suggest that tifacogin may have a disproportionate beneficial effect in microbiologically confirmed cases of CAP.

The finding of a beneficial effect of tifacogin in patients with microbiologically confirmed infection has several potential explanations. Opal and colleagues [28] demonstrated that patients with severe sepsis with a microbiologically confirmed infection had greater perturbations of their coagulation and inflammatory parameters compared with patients with culture-negative severe sepsis. The ability to recover an organism may indicate a patient with greater activation of the coagulation system, a more pronounced proinflammatory stimulus, or both. Recombinant human TFPI binds to lipopolysaccharides (LPSs) and blocks LPS interaction with LPS-binding protein

Figure 4



Mortality by Acute Physiology and Chronic Health Evaluation II (APACHE II) score quartiles in clinical evaluation committee community-acquired pneumonia patients treated with tifacogin or placebo. The quartiles were determined by PROWESS (Recombinant Human Activated Protein C Worldwide Evaluation in Severe Sepsis) trial results.

[29]. Endotoxemia may occur in both Gram-positive and Gram-negative cases of pneumonia [30]. This finding raises the possibility that tifacogin exerts a beneficial effect via immune signaling activities. Finally, tifacogin could potentially play a role in aiding bacterial clearance, which would explain this differential benefit in culture-positive cases [31]. Therefore, three potential mechanisms of action whereby tifacogin may benefit patients with severe CAP are (a) coagulation regulation, (b) immune modulation, and (c) bacterial clearance. The clinical relevance of this hypothesis remains unknown.

In contrast to results in the no-heparin cohort, no benefit of tifacogin was found in CAP patients who received heparin. This result can possibly be explained by potential interactions of tifacogin and heparin. TFPI is most active when expressed on the surface of the cell [32]. Heparin initiates intracellular signaling that results in the transfer of endothelial cell surface-bound TFPI to intracellular storage vesicles, decreasing activity. Heparin could also result in TFPI release into the bloodstream, where it is eventually degraded and is no longer active. In addition, the heparin-binding site on TFPI overlaps the LPS-binding site in the third Kunitz region and carboxyl terminus and competes with TFPI LPS binding [30]. Such an effect could interfere with tifacogin biological activity, suggesting the possibility of a true drug-drug interaction to explain the neutralization of beneficial effect of tifacogin by heparin.

An apparent mortality benefit of heparin use in the placebo group has been noted in several sepsis trials using anticoagulant therapies. However, patients were not randomly assigned to heparin or no-heparin treatment; they were randomly assigned to the study drug only. Investigators used heparin at their discretion and it is reasonable to assume that heparin use would be selected for patients who were less critically ill and less likely to have major coagulopathies. Patients who died early in the course of their illness after random assignment did

Table 4**Incidence of serious bleeding adverse events in patients with and without concomitant heparin use**

MedDRA system organ class	Number (percentage) of subjects ^a			
	Heparin cohort		Non-heparin cohort	
	Tifacogin (n = 172)	Placebo (n = 167)	0.025 TFPI (n = 79)	Placebo (n = 78)
Any serious adverse event	11 (6%)	4 (2%)	3 (4%)	2 (3%)
Gastrointestinal disorders	1 (1%)	2 (1%)	2 (3%)	2 (3%)
General disorders and administration site condition	1 (1%)	1 (1%)	0	0
Injury and poisoning	0	1 (1%)	0	0
Nervous system disorders	3 (2%)	0	0	0
Respiratory, thoracic, and mediastinal disorders	1 (1%)	0	0	0
Surgical and medical procedures	2 (1%)	0	0	0
Vascular disorders	3 (2%)	0	1 (1%)	0

^aNumber and percentage of subjects with one or more events that map to each MedDRA system organ class. Hence, MedDRA system organ class counts may not equate with overall counts. MedDRA, Medical Dictionary for Regulatory Activities; TFPI, tissue factor pathway inhibitor.

not have the opportunity to receive heparin. While the benefit of heparin is likely due to the unequal allocation and selection bias [17], a beneficial effect of heparin alone cannot be excluded. A randomized controlled trial of unfractionated heparin for sepsis is currently under way (ClinicalTrials.gov identifier NCT00100308). However, because of both potential confounding and the possible drug-drug interaction, the phase III confirmation study will require exclusion of heparin therapy during the time of active treatment. TFPI has not been demonstrated to be efficacious for the prevention of deep venous thrombosis in critically ill patients. Mechanical compression devices, an acceptable alternative for critically ill patients at increased risk of bleeding (American College of Chest Physicians guidelines), would therefore be required for both treated and placebo groups.

An additional finding in the CEC CAP subgroup is the apparent absence of a disease severity interaction. Though not

reaching statistical significance, the mortality rates in the tifacogin-treated arm were consistently lower than those in the placebo arm in all four APACHE II score quartiles. This finding is unlike results of other clinical trials involving anti-inflammatory compounds and aPC [33].

Incidence rates of adverse events and events associated with bleeding in CAP patients receiving tifacogin were similar to those in the original TFP007 patient population [15]. Bleeding risk increased in CAP patients receiving both heparin and tifacogin, further emphasizing that tifacogin should not be coadministered with heparin. Most patients who experienced serious bleeding events had pre-existing conditions that put them at increased risk for hemorrhagic complications.

CEC analyses of large phase III databases have recognized limitations. These evaluations are retrospective in nature and

Table 5**Incidence of central nervous system bleeding events in placebo- and tifacogin-treated patients with and without concomitant heparin use**

MedDRA system organ class	Number (percentage) of subjects ^a			
	Heparin cohort		Non-heparin cohort	
	Tifacogin (n = 172)	Placebo (n = 167)	0.025 TFPI (n = 79)	Placebo (n = 78)
Any serious adverse event	5 (3%)	0	1 (1%)	0
Nervous system disorders	3 (2%)	0	0	0
Vascular disorders	2 (1%)	0	1 (1%)	0

^aNumber and percentage of subjects with one or more events that map to each MedDRA system organ class. Hence, MedDRA system organ class counts may not equate with overall counts. MedDRA, Medical Dictionary for Regulatory Activities; TFPI, tissue factor pathway inhibitor.

are based on progressively smaller subgroup sizes, leading to an increasing potential for error. Retrospective analyses of CAP patients' data include an additional hazard: they lack assessment of adequacy of antimicrobial therapy. As is typical of retrospective subgroup analyses, this analysis of a small subgroup of severe CAP patients is solely hypothesis-generating. A subsequent study to test the hypothesis developed by subgroup analysis is more likely to succeed if underlying biological principles support the use of the molecule in that defined population. While the statistical tests are not corrected for the number of subgroups examined, these data and supportive evidence from the literature strengthen the hypothesis that the best target for tifacogin is a population with severe CAP in the absence of concomitant heparin use.

Conclusions

From this retrospective review of patients with severe CAP evaluating the role of tifacogin administration, exploratory analyses showed an improved survival in patients with documented infections who did not receive concomitant heparin. These data support the rationale of the phase III double-blind randomized controlled study exploring the potential benefit of tifacogin in patients with severe CAP admitted to the intensive care unit.

Key messages

- From this retrospective analysis, tissue factor pathway inhibitor seems to improve outcome in severe documented community-acquired pneumonia.
- Concomitant heparin use seems to suppress this observed benefit.
- A prospective randomized controlled study is warranted to confirm this hypothesis.

Competing interests

P-FL has been a consultant for, has participated in advisory boards to, and has received lecture fees from Eli Lilly and Company (Indianapolis, IN, USA), Novartis (formerly Chiron, Emeryville, CA, USA), and GlaxoSmithKline (Uxbridge, Middlesex, UK). SMO is funded by Wyeth Research (Madison, NJ, USA) for preclinical research. He serves as an investigator for the Ocean State Clinical Coordinating Center (Providence, RI, USA), which is funded by Novartis (East Hanover, NJ, USA) and Eisai Medical Research (Woodcliff Lake, NJ, USA) for the conduct of clinical trials for the treatment of sepsis. EA was one of the principal investigators for the TFP007 study, and his institution received a contract from Chiron for patient enrollment. Since 2004, he has not received any consulting income or any other funds from Chiron/Novartis or any entity with interest in the subject of this manuscript. SPL has received consulting fees from Eisai Medical Research and Chiron/Novartis for serving on the CEC and has received investigator grants from these companies for serving on the Ocean State Clinical

Coordinating Center. AAC, FX, and LP are current or former Chiron/Novartis employees. RGW has been paid on an hourly basis for work on the CEC and has also received an investigator-initiated grant from Chiron/Novartis.

Authors' contributions

P-FL, SMO, SPL, and RGW participated in the study design, in the acquisition and interpretation of the data, and in the drafting of the manuscript. EA, AAC, FX, and LP participated in the interpretation of the data and in the drafting of the manuscript. All authors read and approved the final manuscript.

Authors' information

This work was performed at Novartis (formerly Chiron, Emeryville, CA, USA).

Acknowledgements

The authors would like to thank the following Novartis employees: Connie D Louie for her contribution in organizing the materials for the CEC review, Alan Nakamoto for programming the analyses, and Christian Zwingelstein, Jo Ellen Schweinle, and Steve Hardy for their critical review of the manuscript. Editorial assistance of the manuscript was provided by Patrice Ferriola, whose work was financially supported by Novartis.

References

1. Angus DC, Linde-Zwirble WT, Lidicker J, Clermont G, Carcillo J, Pinsky MR: **Epidemiology of severe sepsis in the United States: analysis of incidence, outcome, and associated costs of care.** *Crit Care Med* 2001, **29**:1303-1310.
2. Opal S, Laterre PF, Abraham E, Francois B, Wittebole X, Lowry S, Dhainaut JF, Warren B, Dugernier T, Lopez A, Sanchez M, Demeyer I, Jauregui L, Lorente JA, McGee W, Reinhart K, Kljucar S, Souza S, Pribble J, Controlled Mortality Trial of Platelet-Activating Factor Acetylhydrolase in Severe Sepsis Investigators: **Recombinant human platelet-activating factor acetylhydrolase for the treatment of severe sepsis: results of a phase III, multicenter, randomized, double-blind, placebo-controlled, clinical trial.** *Crit Care Med* 2004, **32**:332-341.
3. Warren BL, Eid A, Singer P, Pillay SS, Carl P, Novak I, Chalupa P, Atherstone A, Pénzens I, Kübler A, Knaub S, Keinecke HO, Heinrichs H, Schindel F, Juers M, Bone RC, Opal SM, KyberSept Trial Study Group: **High-dose antithrombin III in severe sepsis (the KyberSept trial).** *JAMA* 2001, **286**:1869-1887.
4. Bernard G, Vincent JL, Laterre PF, LaRosa SP, Dhainaut JF, Lopez-Rodriguez A, Steingrub JS, Garber GE, Helterbrand JD, Ely EW, Fisher CJ Jr: **Efficacy and safety of recombinant human activated protein C for severe sepsis.** *N Engl J Med* 2001, **344**:699-709.
5. Annane D, Sebille V, Charpentier C, Bollaert PE, François B, Korach JM, Capellier G, Cohen Y, Azoulay E, Troché G, Chaumet-Riffaut P, Bellissant E: **Effect of treatment with low doses of hydrocortisone and fludrocortisone on mortality in patients with septic shock.** *JAMA* 2002, **288**:862-871.
6. Bartlett JG, Dowell SF, Mandell LA, File TM Jr, Musher DM, Fine MJ: **Practice guidelines for the management of community-acquired pneumonia in adults.** *Infectious Diseases Society of America.* *Clin Infect Dis* 2000, **31**:347-382.
7. Camerer E, Kolsto AB, Prydz H: **Cell biology of tissue factor, the principal initiator of blood coagulation.** *Thromb Res* 1996, **81**:1-41.
8. Levi M, Poll T van der, ten Cate H: **Tissue factor in infection and severe inflammation.** *Semin Thromb Hemost* 2006, **32**:33-39.
9. Choi G, Schultz MJ, van Till JW, Bresser P, Zee JS van der, Boermeester MA, Levi M, Poll T van der: **Disturbed alveolar fibrin turnover during pneumonia is restricted to the site of infection.** *Eur Respir J* 2004, **24**:786-789.

10. Gando S, Kameue T, Matsuda N, Hayakawa M, Morimoto Y, Ishitani T, Kemmotsu O: **Imbalances between the levels of tissue factor and tissue factor pathway inhibitor in ARDS patients.** *Thromb Res* 2003, **109**:119-124.
11. Welty-Wolf KE, Carraway MS, Miller DL, Ortel TL, Ezban M, Ghio AJ, Idell S, Piantadosi CA: **Coagulation blockade prevents sepsis-induced respiratory and renal failure in baboons.** *Am J Respir Crit Care Med* 2001, **164**:1988-1996.
12. Miller DL, Welty-Wolf K, Carraway MS, Ezban M, Ghio A, Suliman H, Piantadosi CA: **Extrinsic coagulation blockade attenuates lung injury and proinflammatory cytokine release after intratracheal lipopolysaccharide.** *Am J Respir Cell Mol Biol* 2002, **26**:650-658.
13. Enkhbaatar P, Okajima K, Murakami K, Uchiba M, Okabe H, Okabe K, Yamaguchi Y: **Recombinant tissue factor pathway inhibitor reduces lipopolysaccharide-induced pulmonary vascular injury by inhibiting leukocyte activation.** *Am J Respir Crit Care Med* 2000, **162**:1752-1759.
14. Bajaj MS, Birktoft JJ, Steer SA, Bajaj SP: **Structure and biology of tissue factor pathway inhibitor.** *Thromb Haemost* 2001, **86**:959-972.
15. Abraham E, Reinhart K, Opal S, Demeyer I, Doig C, Rodriguez AL, Beale R, Svoboda P, Laterre PF, Simon S, Light B, Spapen H, Stone J, Seibert A, Peckelsen C, De Deyne C, Postier R, Pettita V, Artigas A, Percell SR, Shu V, Zwingelstein C, Tobias J, Poole L, Stolzenbach JC, Creasey AA, OPTIMIST Trial Study Group: **Efficacy and safety of tifacogin (recombinant tissue factor pathway inhibitor) in severe sepsis: a randomized controlled trial.** *JAMA* 2003, **290**:238-247.
16. Sprung CL, Finch RG, Thijs LG, Glauser MP: **International Sepsis Trial (INTERSEPT): role and impact of a clinical evaluation Committee.** *Crit Care Med* 1996, **24**:1441-1447.
17. Opal SM, Fisher C, Dhainaut JF, Vincent JL, Brase R, Lowry SF, Sadoff JC, Slotman GJ, Levy H, Balk RA, Shelly MP, Pribble JP, LaBrecque JF, Lookabaugh J, Donovan H, Dubin H, Baughman R, Norman J, DeMaria E, Matzel K, Abraham E, Seneff M: **Confirmatory interleukin-1 receptor antagonist trial in severe sepsis: a phase III, randomized, double-blind, placebo-controlled, multicenter trial. The Interleukin-1 Receptor Antagonist Investigator Group.** *Crit Care Med* 1997, **25**:1115-1124.
18. Dhainaut JF, Laterre PF, LaRosa SP, Levy H, Garber GE, Heiselman D, Kinasevitz GT, Light RB, Morris P, Schein R, Sollet JP, Bates BM, Utterback BG, Maki D: **The clinical evaluation committee in a large multicenter phase 3 trial of drotrecogin alfa (activated) in patients with severe sepsis (PROWESS): role, methodology, and results.** *Crit Care Med* 2003, **31**:2291-2301.
19. Laterre PF, Garber G, Levy H, Wunderink R, Kinasevitz GT, Sollet JP, Maki DG, Bates B, Yan SC, Dhainaut JF, PROWESS Clinical Evaluation Committee: **Severe community-acquired pneumonia as a cause of severe sepsis: data from the PROWESS study.** *Crit Care Med* 2005, **33**:952-961.
20. Rosón B, Carratalà J, Fernández-Sabé N, Tubau F, Manresa F, Gudiol F: **Causes and factors associated with early failure in hospitalized patients with community-acquired pneumonia.** *Arch Intern Med* 2004, **164**:502-508.
21. Menéndez R, Torres A, Zalacain R, Aspa J, Martín Villasclaras JJ, Borderías L, Benítez Moya JM, Ruiz-Manzano J, Rodríguez de Castro F, Blanquer J, Pérez D, Puzo C, Sánchez Gascón F, Gallardo J, Alvarez C, Molinos L, Neumofail Group: **Risk factors of treatment failure in community acquired pneumonia: implications for disease outcome.** *Thorax* 2004, **59**:960-965.
22. Siegel JP: **Assessing the use of activated protein C in the treatment of severe sepsis.** *N Engl J Med* 2002, **347**:1030-1034.
23. Christ-Crain M, Jaccard-Stolz D, Bingisser R, Gencay MM, Huber PR, Tamm M, Müller B: **Effect of procalcitonin-guided treatment on antibiotic use and outcome in lower respiratory tract infections: cluster-randomised, single-blinded intervention trial.** *Lancet* 2004, **363**:600-607.
24. Gross TJ, Simon RH, Sitrin RG: **Tissue factor procoagulant expression by rat alveolar epithelial cells.** *Am J Respir Cell Mol Biol* 1992, **6**:397-403.
25. McGee MP, Wallin R, Wheeler FB, Rothberger H: **Initiation of the extrinsic pathway of coagulation by human and rabbit alveolar macrophages: a kinetic study.** *Blood* 1989, **74**:1583-1590.
26. Levi M, Schultz MJ, Rijneveld AW, Poll T van der: **Bronchoalveolar coagulation and fibrinolysis in endotoxemia and pneumonia.** *Crit Care Med* 2003, **31**:S238-S242.
27. Luyt CE, Guerin V, Combes A, Trouillet JL, Ayed SB, Bernard M, Gibert C, Chastre J: **Procalcitonin kinetics as a prognostic marker of ventilator-associated pneumonia.** *Am J Respir Crit Care Med* 2005, **171**:48-53.
28. Opal SM, Garber GE, LaRosa SP, Maki DG, Freebairn RC, Kinasevitz GT, Dhainaut JF, Yan SB, Williams MD, Graham DE, Nelson DR, Levy H, Bernard GR: **Systemic host responses in severe sepsis analyzed by causative microorganism and treatment effects of drotrecogin alfa (activated).** *Clin Infect Dis* 2003, **37**:50-58.
29. Park CT, Creasey AA, Wright SD: **Tissue factor pathway inhibitor blocks cellular effects of endotoxin by binding to endotoxin and interfering with transfer to CD14.** *Blood* 1997, **89**:4268-4274.
30. Opal SM, Scannon PJ, Vincent JL, White M, Carroll SF, Palardy JE, Parejo NA, Pribble JP, Lemke JH: **Relationship between plasma levels of lipopolysaccharide (LPS) and LPS-binding protein in patients with severe sepsis and septic shock.** *J Infect Dis* 1999, **180**:1584-1589.
31. Hardy S, Schirm S, Liu X, Dai Y: **Tifacogin increases bacterial clearance from blood.** *Crit Care* 2006, **10**:P155.
32. Lupu C, Poulsen E, Roquefeuil S, Westmuckett AD, Kakkar VV, Lupu F: **Cellular effects of heparin on the production and release of tissue factor pathway inhibitor in human endothelial cells in culture.** *Arterioscler Thromb Vasc Biol* 1999, **19**:2251-2262.
33. Eichacker PQ, Parent C, Kalil A, Esposito C, Cui X, Banks SM, Gertenberger EP, Fitz Y, Danner RL, Natanson C: **Risk and efficacy of anti-inflammatory agents: retrospective and confirmatory studies of sepsis.** *Am J Respir Crit Care Med* 2002, **166**:1197-1205.