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## Hydrocortisone Therapy for Patients with Septic Shock

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### ABSTRACT

#### BACKGROUND

Hydrocortisone is widely used in patients with septic shock even though a survival benefit has been reported only in patients who remained hypotensive after fluid and vasopressor resuscitation and whose plasma cortisol levels did not rise appropriately after the administration of corticotropin.

#### METHODS

In this multicenter, randomized, double-blind, placebo-controlled trial, we assigned 251 patients to receive 50 mg of intravenous hydrocortisone and 248 patients to receive placebo every 6 hours for 5 days; the dose was then tapered during a 6-day period. At 28 days, the primary outcome was death among patients who did not have a response to a corticotropin test.

#### RESULTS

Of the 499 patients in the study, 233 (46.7%) did not have a response to corticotropin (125 in the hydrocortisone group and 108 in the placebo group). At 28 days, there was no significant difference in mortality between patients in the two study groups who did not have a response to corticotropin (39.2% in the hydrocortisone group and 36.1% in the placebo group,  $P=0.69$ ) or between those who had a response to corticotropin (28.8% in the hydrocortisone group and 28.7% in the placebo group,  $P=1.00$ ). At 28 days, 86 of 251 patients in the hydrocortisone group (34.3%) and 78 of 248 patients in the placebo group (31.5%) had died ( $P=0.51$ ). In the hydrocortisone group, shock was reversed more quickly than in the placebo group. However, there were more episodes of superinfection, including new sepsis and septic shock.

#### CONCLUSIONS

Hydrocortisone did not improve survival or reversal of shock in patients with septic shock, either overall or in patients who did not have a response to corticotropin, although hydrocortisone hastened reversal of shock in patients in whom shock was reversed. (ClinicalTrials.gov number, NCT00147004.)

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**S**EVERE SEPSIS IS A MAJOR CAUSE OF MORTALITY and morbidity worldwide.<sup>1,2</sup> Septic shock, the most severe manifestation, occurs in 2 to 20% of inpatients.<sup>3</sup> The incidence of the condition has been rising,<sup>4</sup> and a death rate of 33 to 61% has been reported in the placebo groups of multicenter trials.<sup>5-8</sup>

The use of corticosteroids as an adjunctive therapy has been controversial for decades.<sup>9</sup> After the study by Schumer,<sup>10</sup> a short course of high-dose corticosteroids became accepted therapy. Subsequent studies, however, did not confirm a survival benefit with this regimen and suggested an increase in superinfection-related mortality.<sup>11-13</sup> Studies that have used lower doses of hydrocortisone (200 to 300 mg per day) for longer durations have reported earlier reversal of shock<sup>14-18</sup> and improved survival.<sup>14,16</sup> The prognostic importance of the response to corticotropin had been recognized in critical illness previously,<sup>19,20</sup> and the results in the hydrocortisone studies were particularly apparent in patients who did not have a response to a corticotropin test. Meta-analyses,<sup>21,22</sup> reviews,<sup>20</sup> and guidelines<sup>23</sup> have advocated the use of low-dose hydrocortisone in patients with septic shock. These recommendations were based primarily on a study of patients with septic shock who remained hypotensive after at least 1 hour of resuscitation with fluids and vasopressors.<sup>16</sup> In this study, a survival benefit was seen in patients with no response to corticotropin who received hydrocortisone and fludrocortisone. Our trial, called the Corticosteroid Therapy of Septic Shock (CORTICUS) study, evaluated the efficacy and safety of low-dose hydrocortisone therapy in a broad population of patients with septic shock — in particular, patients who had had a response to a corticotropin test, in whom a benefit was unproven.<sup>9</sup>

## METHODS

### STUDY DESIGN

In this multicenter, randomized, double-blind, placebo-controlled study, the protocol was approved by the ethics committee at each of the 52 participating intensive care units (ICUs). Patients were enrolled from March 2002 to November 2005, after providing written informed consent. In cases in which a patient lacked mental competency, consent was obtained either from a surrogate, the next of kin, or a legal representative

(with retrospective consent obtained from patients who regained competency), according to national regulations. An independent data and safety monitoring board met after each of three interim analyses. At the end of the study, a clinical evaluation committee whose members were unaware of study-group assignments assessed the appropriateness of antiinfective treatments.

The authors designed the study, gathered and analyzed the data, and vouch for the completeness and accuracy of the data and the analysis. The sponsors had no role in the design and conduct of the study, in the collection, management, analysis, or interpretation of the data, or in the preparation, review, or approval of the manuscript.

### PATIENTS

Patients who were 18 years of age or older and had been hospitalized in participating ICUs were prospectively enrolled in the study if they met all eligibility criteria. (For details, see Table 1 of the Supplementary Appendix, available with the full text of this article at [www.nejm.org](http://www.nejm.org).) Inclusion criteria were clinical evidence of infection, evidence of a systemic response to infection, and the onset of shock within the previous 72 hours (as defined by a systolic blood pressure of <90 mm Hg despite adequate fluid replacement or a need for vasopressors for at least 1 hour) and hypoperfusion or organ dysfunction attributable to sepsis. Notable exclusion criteria included underlying disease with a poor prognosis, a life expectancy of less than 24 hours, immunosuppression, and treatment with long-term corticosteroids within the past 6 months or short-term corticosteroids within the past 4 weeks.

### RANDOMIZATION

Randomization (in a 1:1 ratio) was stratified according to study center in blocks of four with the use of a computerized random-number generator list provided by a statistician who was not involved in the determination of eligibility, administration of a study drug, or an assessment of outcomes. In each center, the study drug (hydrocortisone or placebo) was sealed in sequentially numbered, identical boxes that contained the entire treatment for each patient to be administered sequentially. The sequence was concealed from the investigators. All patients, medical and nursing staff members, pharmacists, investigators,

and members of the monitoring board remained unaware of study-group assignments throughout the study period.

#### STUDY DRUGS

Hydrocortisone (Rotexmedica) was prepared in vials containing 100 mg of hydrocortisone hemisuccinate powder with ampules containing 2 ml of sterile water diluent; the vials were then coded and masked centrally (Klocke Verpackungs Service). Vials containing placebo were identical to those containing hydrocortisone. The study drugs were administered as a 50-mg intravenous bolus every 6 hours for 5 days, then tapered to 50 mg intravenously every 12 hours for days 6 to 8, 50 mg every 24 hours for days 9 to 11, and then stopped. A total of 29 doses were given. Evidence-based guidelines for the treatment of patients were encouraged.<sup>24</sup>

#### DEFINITIONS

Organ-system failure was defined for each of the six major organ systems as a Sequential Organ Failure Assessment (SOFA) score of 3 or 4 points (on a scale ranging from 0 to 4 for each organ system, for an aggregate score of 0 to 24, with higher scores indicating more severe organ dysfunction).<sup>25</sup> Reversal of shock was defined as the maintenance of a systolic blood pressure of at least 90 mm Hg without vasopressor support for at least 24 hours. Superinfection was defined as a new infection occurring 48 hours or more after the initiation of a study drug.<sup>26</sup> New sepsis was defined as a new septic episode with or without microbiologic confirmation. New septic shock was defined as a new episode of septic shock after reversal of the initial episode. The absence of a response to a corticotropin test was defined as an increase in the cortisol level of no more than 9  $\mu$ g per deciliter (248 nmol per liter).<sup>16</sup>

#### DATA COLLECTION

##### *Clinical Evaluation*

The following data were recorded: general characteristics of the patients, including demographic data, diagnoses, and recent surgery; the severity of illness, as assessed by vital signs, the Simplified Acute Physiology Score (SAPS) II (on a scale from 0 to 163, with higher scores indicating more severe organ dysfunction),<sup>27</sup> and the SOFA score<sup>25</sup>; and interventions, including the type and doses of vasopressors, antibiotics, and adjunctive treatments such as corticosteroids and etomidate.

#### *Laboratory Measures*

Hematologic and chemical data, blood gas analyses, and cultures of blood and other specimens that were obtained from potential sites of infection were recorded. A short corticotropin test was performed with the use of blood samples taken immediately before and 60 minutes after an intravenous bolus of 0.25 mg of cosyntropin (Novartis or Alliance). After centrifugation, serum samples were stored at a temperature no higher than  $-20^{\circ}\text{C}$  until assayed. To reduce heterogeneity in the determination of cortisol levels, all samples were measured blindly and serially before interim and final analyses in a central laboratory with the use of the Elecsys cortisol assay (Roche Diagnostics).

#### FOLLOW-UP

During the 28-day period after randomization, data were collected regarding vital signs, results from laboratory tests and cultures of specimens drawn from any new site of infection, and any major interventions that were performed. Rates of death at 28 days, in the ICU, in the hospital, and at 1 year after randomization were recorded.

#### END POINTS

The primary end point was the rate of death at 28 days in patients who did not have a response to corticotropin. Secondary end points were the rates of death at 28 days in patients who had a response to corticotropin and in all patients, the rates of death in the ICU and in the hospital, the rates of death at 1 year after randomization, a reversal of organ system failure (including shock), and the duration of the stay in the ICU and the hospital.

Safety was assessed by recording adverse events, particularly superinfection, gastrointestinal bleeding, hyperglycemia, hypernatremia, clinical muscular weakness, stroke, acute myocardial infarction, and peripheral ischemia. Methods to enhance the quality of measurements included holding biannual meetings of investigators, sending newsletters, and conducting random quality-assurance evaluations.

#### STATISTICAL ANALYSIS

A sample size of 800 patients (400 per group) was needed to achieve a statistical power of 80%

to detect an absolute decrease in mortality of 10% from an existing death rate of 50% in patients who did not have a response to corticotropin (40% of the total group). All analyses were performed according to a prespecified plan. The population was analyzed according to an intention-to-treat principle. The rate of death from all causes at 28 days was analyzed with the use of Fisher's exact test for differences between study groups. A maximum overall two-sided probability of a type I error of 5% was accepted. The test result was corrected for two interim analyses for efficacy. Splitting the alpha error function was performed according to the O'Brien-Fleming method ( $P=0.0006$ ,  $P=0.005$ , and  $P=0.047$  for the first, second, and final analysis, respectively).

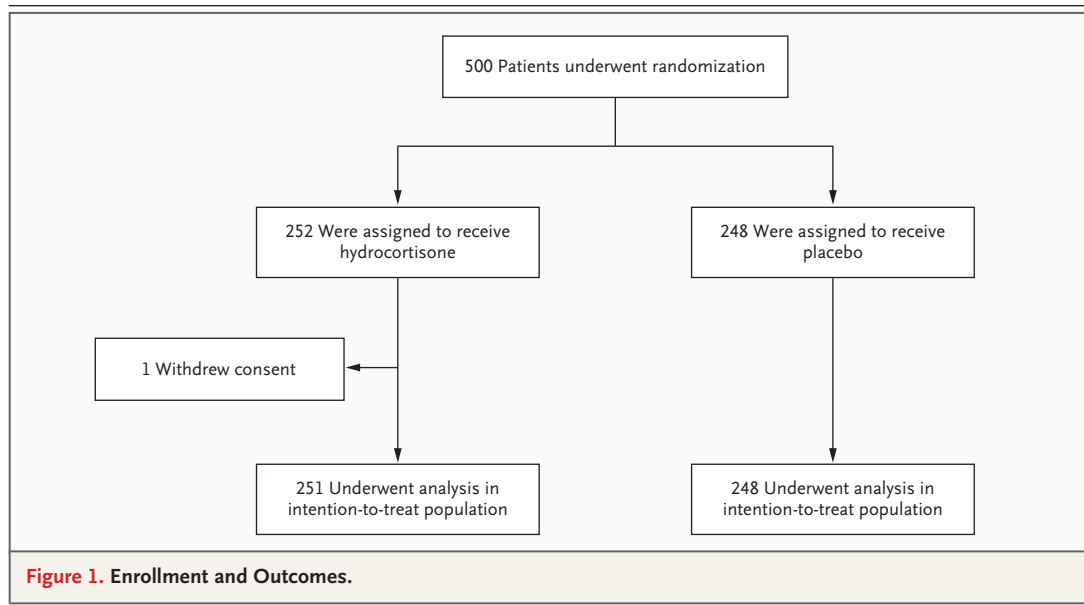
The difference in the rate of death at 28 days between the two study groups was considered to have statistical significance if the stopping criteria of the interim analysis were met or the two-sided  $P$  value of the final analysis was less than 0.047. The differences between all other secondary efficacy variables were assumed to have statistical significance for  $P$  values of less than 0.05. Kaplan-Meier curves for cumulative survival during the 28-day observation period were constructed and compared with the use of the log-rank test. The median time until the reversal of septic shock was calculated with the use of Kaplan-Meier analysis. Adverse events were reported for the per-protocol population.

## RESULTS

## PATIENTS

Five hundred patients were enrolled in the study (Fig. 1). One patient in the hydrocortisone group was excluded because consent was withdrawn. Of the remaining 499 patients, all met the entry criteria, although 15 also fulfilled the exclusion criteria (8 patients in the hydrocortisone group and 7 in the placebo group) since 14 had received previous corticosteroid therapy and 1 had undergone previous cardiopulmonary resuscitation. Eighty-seven percent of patients in both the hydrocortisone group and the placebo group received at least 90% of the doses of a study drug.

Of 499 patients in the study, 233 (46.7%) did not have a response to corticotropin (125 in the hydrocortisone group and 108 in the placebo group); 254 patients (50.9%) did have a response to corticotropin (118 in the hydrocortisone group and 136 in the placebo group). Results were unknown for eight patients in the hydrocortisone group and four in the placebo group (2.4%). Etomidate was used in 51 of 251 patients in the hydrocortisone group (20.3%) and in 45 of 248 patients in the placebo group (18.1%) before study entry and in 22 patients (8.8%) and 20 patients (8.1%), respectively, after study enrollment. Among the 96 patients who had received etomidate, 58 did not have a response to corticotropin (60.4%), as compared with 175 of 403 who did



not receive etomidate (43.4%,  $P=0.004$ ). The median time between the last dose of etomidate and enrollment was 14 hours (range, 1 to 67).

At baseline, the two study groups were well balanced with regard to demographic characteristics (Table 1), clinical characteristics (Table 2), and the type and site of infection and infecting organisms (Table 2 of the Supplementary Appendix).

**PRIMARY END POINTS**

*Mortality*

There was no significant difference between the two study groups in the primary outcome, the rate of death at 28 days among patients who did not have a response to corticotropin. There were 49 deaths in 125 patients in the hydrocortisone group (39.2%; 95% confidence interval [CI], 30.5 to 47.9) and 39 deaths in 108 patients in the placebo group (36.1%; 95% CI, 26.9 to 45.3;  $P=0.69$ ).

Likewise, there was no significant difference in the rate of death at 28 days in patients who had a response to corticotropin. There were 34 deaths in 118 patients in the hydrocortisone group (28.8%; 95% CI, 20.6 to 37.0) and 39 deaths among 136 patients in the placebo group (28.7%; 95% CI, 21.1 to 36.3;  $P=1.00$ ). Overall, there were 86 deaths in the hydrocortisone group (34.3%; 95% CI, 28.3 to 40.2) and 78 deaths in the placebo group (31.5%; 95% CI, 25.6 to 37.3;  $P=0.51$ ).

No differences in mortality were seen between the study groups or between the corticotropin subgroups at any other time point (Fig. 2 and Table 3). Twenty-one post hoc analyses were performed to elucidate the reasons for the rates of death at 28 days in subgroups of patients. These analyses showed that among patients with a systolic blood pressure persisting at less than 90 mm Hg within 30 hours after study entry, 31 of 69 patients in the hydrocortisone group (44.9%)

**Table 1. Demographic Characteristics of the Patients, According to Subgroup.\***

Characteristic	No Response to Corticotropin		Response to Corticotropin		All Patients	
	Hydrocortisone (N=125)	Placebo (N=108)	Hydrocortisone (N=118)	Placebo (N=136)	Hydrocortisone (N=251)	Placebo (N=248)
Age — yr	63±13	63±15	62±14	64±16	63±14	63±15
Sex — no. (%)						
Male	85 (68)	69 (64)	76 (64)	95 (70)	166 (66)	166 (67)
Female	40 (32)	39 (36)	42 (36)	41 (30)	85 (34)	82 (33)
White race — %†	119 (95)	101 (94)	110 (93)	125 (92)	236 (94)	228 (92)
Previous disease — no./total no. (%)						
Hypertension	48 (38)	37 (34)	39 (33)	60/133 (45)	89 (35)	98/245 (40)
Coronary artery disease	20 (16)	26 (24)	17 (14)	21/133 (16)	37 (15)	47/245 (19)
Congestive heart failure	5 (4)	8 (7)	4 (3)	12/133 (9)	10 (4)	20/245 (8)
Neurologic	19 (15)	10 (9)	14 (12)	14/133 (11)	33 (13)	25/245 (10)
Chronic obstructive pulmonary disease	14 (11)	12 (11)	11 (9)	17/133 (13)	27 (11)	29/245 (12)
Other pulmonary disorder	6 (5)	12 (11)	17 (14)	12/133 (9)	23 (9)	24/245 (10)
Cancer	27 (22)	21 (19)	18 (15)	16/133 (12)	47 (19)	37/245 (15)
Diabetes	22 (18)	19 (18)	28 (24)	37/133 (28)	51 (20)	56/245 (23)
Liver	14 (11)	10 (9)	9 (8)	7/133 (5)	23 (9)	17/245 (7)
Chronic renal failure	12 (10)	11 (10)	10 (9)	10/133 (8)	22 (9)	21/245 (9)
Admission category — no./total no. (%)						
Medical	39 (31)	35/107 (33)	37/116 (32)	57/135 (42)	80/249 (32)	93/246 (38)
Emergency surgery	69 (55)	63/107 (59)	66/116 (57)	67/135 (50)	138/249 (55)	132/246 (54)
Elective surgery	17 (14)	9/107 (8)	13/116 (11)	11/135 (8)	31/249 (12)	21/246 (9)

\* Plus-minus values are means ±SD.

† Race was reported by the investigators.

**Table 2. Clinical Characteristics of the Patients at Baseline, According to Subgroup.\***

Variable	No Response to Corticotropin			Response to Corticotropin			All Patients			
	No. of Patients	Hydrocortisone (N=125)	Placebo (N=108)	No. of Patients	Hydrocortisone (N=118)	Placebo (N=136)	No. of Patients	Hydrocortisone (N=251)	Placebo (N=248)	
Temperature — °C	124	37.7±1.6	108	116	38.0±1.4	135	248	37.9±1.5	247	38.0±1.4
Heart rate — bpm	124	121±24	108	115	116±29	136	247	119±26	248	118±25
Systolic blood pressure — mm Hg	124	92±22	108	116	94±24	136	248	94±23	248	95±27
SAPS II score†	125	50.7±17.8	108	117	47.9±18.0	136	250	49.5±17.8	248	48.6±16.7
SOPA score‡	125	11.0±3.4	108	118	10.3±3.4	136	251	10.6±3.4	248	10.6±3.2
Leukocytes — thousands/mm <sup>3</sup>	121	15.8±11.2	106	114	14.3±8.1	133	243	14.9±9.8	243	14.7±9.8
Platelets — thousands/mm <sup>3</sup>	121	205±131	106	114	218±140	132	243	212±135	242	202±133
Glucose — mg/dl	117	140±65	102	111	139±59	133	235	140±65	239	137±50
Arterial lactate — mmol/liter	102	4.6±4.0	94	94	3.1±3.0	114	202	3.9±3.6	212	4.1±4.1
Ratio of partial pressure of arterial oxygen to fraction of inspired oxygen	107	159±89	94	99	162±82	119	214	162±89	216	154±73
Cortisol — µg/dl	125		108	118		136	243		244	
Before corticotropin		30±20			27±19			28±20		29±20
60 min after corticotropin		33±19			46±22			39±22		39±22
Response to corticotropin test		3±4			18±11			11±11		10±8
Receiving vasopressor or inotrope at baseline — no. (%)		125 (100)			117 (99)			249 (99)		243 (98)
Type of vasopressor§										
Norepinephrine — no. (%)		116 (93)			103 (87)			224 (89)		231 (93)
Maximum dose — µg/kg/min		0.5±0.5			0.4±0.7			0.5±0.6		0.4±0.5
Epinephrine — no. (%)		19 (15)			14 (12)			35 (14)		22 (9)
Maximum dose — µg/kg/min		0.8±1.6			0.3±0.4			0.6±1.2		0.9±2.6
Dopamine — no. (%)		10 (8)			16 (14)			27 (11)		29 (12)
Maximum dose — µg/kg/min		12.9±9.6			9.8±6.1			10.4±7.5		7.9±6.6

Ventilatory support at baseline																				
Mechanical ventilation — no. (%)	125	113 (90)	108	99 (92)	118	108 (92)	134	110 (82)	251	228 (91)	246	212 (86)								
Tidal volume — ml/kg	99	7.6±2.1	80	7.5±2.1	95	7.6±2.2	97	7.7±2.2	201	7.7±2.1	180	7.6±2.1								
Fraction of inspired oxygen — %	117	67±25	104	64±25	106	60±24	125	63±24	231	64±25	232	63±24								
Positive end-expiratory pressure — cm of water	108	8±4	92	8±3	99	9±4	114	9±4	215	9±4	209	9±4								
Activated protein C — no. (%)¶	125	11 (9)	108	13 (12)	118	6 (5)	136	7 (5)	251	17 (7)	248	20 (8)								
Antithrombin — no. (%)¶	125	24 (19)	108	17 (16)	118	15 (13)	136	19 (14)	251	40 (16)	248	36 (15)								

\* Plus-minus values are means ±SD. To convert the values for cortisol to nanomoles per liter, multiply by 27.59. To convert the values for glucose to millimoles per liter, multiply by 0.05551.  
 † The Simplified Acute Physiology Score (SAPS) II ranges from 0 to 163, with higher scores indicating more severe organ dysfunction.  
 ‡ Scores on the Sequential Organ Failure Assessment (SOFA) range from 0 to 24, with higher scores indicating more severe organ dysfunction.  
 § Five patients received vasopressin, four in the hydrocortisone group and one in the placebo group.  
 ¶ Patients may have received these drugs after baseline.

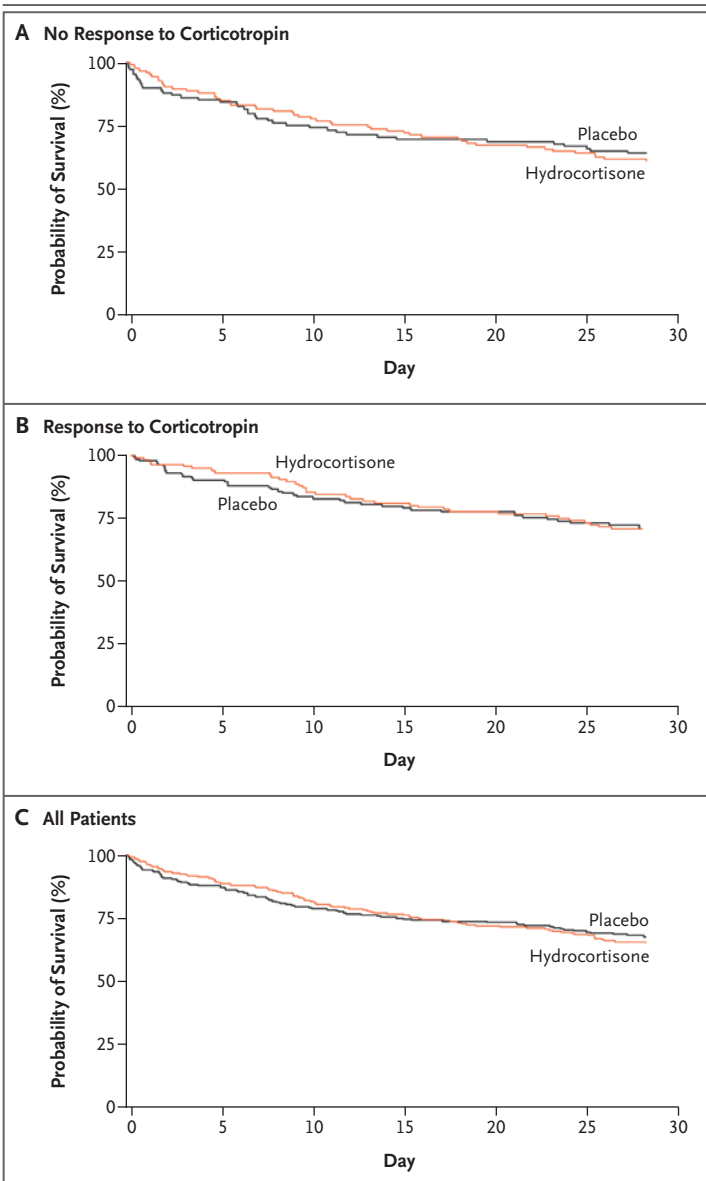
and 32 of 57 patients in the placebo group (56.1%) had died, with an absolute difference in the hydrocortisone group of -11.2% (95% CI, -18.6 to 6.2; P=0.28); among those whose systolic blood pressure was measured at 90 mm Hg or more within 30 hours after study entry, 55 of 181 in the hydrocortisone group (30.4%) and 46 of 189 in the placebo group (24.3%) died, with an absolute difference in the hydrocortisone group of 6.1% (95% CI, -3.0 to 15.1; P=0.20). In post hoc analyses, the rates of death among the 384 patients who received a study drug within 12 hours after baseline were similar in the two groups: 73 of 198 patients in the hydrocortisone group (36.9%) and 58 of 186 patients in the placebo group (31.2%, P=0.28) (Table 3 of the Supplementary Appendix).

Post hoc analyses also revealed an increased rate of death at 28 days among patients who received etomidate before randomization in both groups (23 of 51 in the hydrocortisone group [45.1%] and 18 of 45 in the placebo group [40.0%]), as compared with patients who did not receive etomidate (63 of 200 in the hydrocortisone group [31.5%] and 60 of 203 in the placebo group [29.6%, P=0.03]).

**REVERSAL OF SHOCK**

The proportions of patients who underwent a reversal of shock were similar among patients who did not have a response to corticotropin: 95 of 125 in the hydrocortisone group (76.0%; 95% CI, 68.5 to 83.5) and 76 of 108 in the placebo group (70.4%; 95% CI, 61.8 to 79.0; P=0.41); among patients who had a response to corticotropin: 100 of 118 patients in the hydrocortisone group (84.7%; 95% CI, 78.3 to 91.2) and 104 of 136 in the placebo group (76.5%; 95% CI, 69.3 to 83.6; P=0.13); and among all patients: 200 of 251 in the hydrocortisone group (79.7%; 95% CI, 74.7 to 84.7) and 184 of 248 in the placebo group (74.2%; 95% CI, 68.7 to 79.6; P=0.18).

The duration of time until the reversal of shock was significantly shorter among patients receiving hydrocortisone for all patients (P<0.001), for those who had a response to corticotropin (P<0.001), and for those who did not have a response to corticotropin (P=0.06) (Fig. 3). The median time until reversal of shock was also shorter in the hydrocortisone group than in the placebo group: for all patients, 3.3 days (95% CI, 2.9 to 3.9) versus 5.8 days (95% CI, 5.2 to 6.9);



**Figure 2.** Kaplan–Meier Curves for Survival at 28 Days.

For the comparison between patients with septic shock who received hydrocortisone and those who received placebo, there was no significant difference among those who did not have a response to a corticotropin test (Panel A), those who had a response to corticotropin (Panel B), and all patients who underwent evaluation (Panel C).

for those who had a response to corticotropin, 2.8 days (95% CI, 2.1 to 3.3) versus 5.8 days (95% CI, 5.2 to 6.9); and for those who did not have a response, 3.9 days (95% CI, 3.0 to 5.2) versus 6.0 days (95% CI, 4.9 to 9.0).

The number of extubated patients on day 28 was similar in the two study groups: 119 of 228

patients who underwent ventilation at baseline in the hydrocortisone group (52%) and 113 of 212 patients in the placebo group (53%). For the 357 patients with cultured pathogens for their primary infection, the clinical evaluation committee determined that appropriate antimicrobial therapy was given to 126 of 173 in the hydrocortisone group (72.8%) and 145 of 184 in the placebo group (78.8%). There was no significant difference in outcome between study groups among patients receiving appropriate antibiotic therapy and those receiving inappropriate therapy.

#### USE OF CORTICOSTEROIDS AND OTHER DRUGS

Eleven patients in the hydrocortisone group (4.4%) and 10 patients in the placebo group (4.0%) received corticosteroids after study enrollment for allergic reactions, laryngeal edema, bronchospasm, brain edema, replacement of long-term corticosteroid therapy whose history was unknown at enrollment, acute respiratory distress syndrome, and septic shock. Five patients received corticosteroids for septic shock after completion of the course of a study drug. The numbers of patients in both groups who received activated protein C, antithrombin, or both were similar (Table 2).

#### ADVERSE EVENTS

In the hydrocortisone group, there was an increased incidence of superinfections, including new episodes of sepsis or septic shock, with a combined odds ratio of 1.37 (95% CI, 1.05 to 1.79); there was also an increased incidence of hyperglycemia and hypernatremia (Table 4). Neuromuscular weakness was rarely reported.

#### DISCUSSION

In our study, the use of low-dose hydrocortisone had no significant effect on the rate of death in patients with septic shock at 28 days, regardless of the patients' adrenal responsiveness to corticotropin. The proportion of patients in whom reversal of shock was achieved was similar in the two groups, though this goal was achieved earlier in patients who received hydrocortisone.

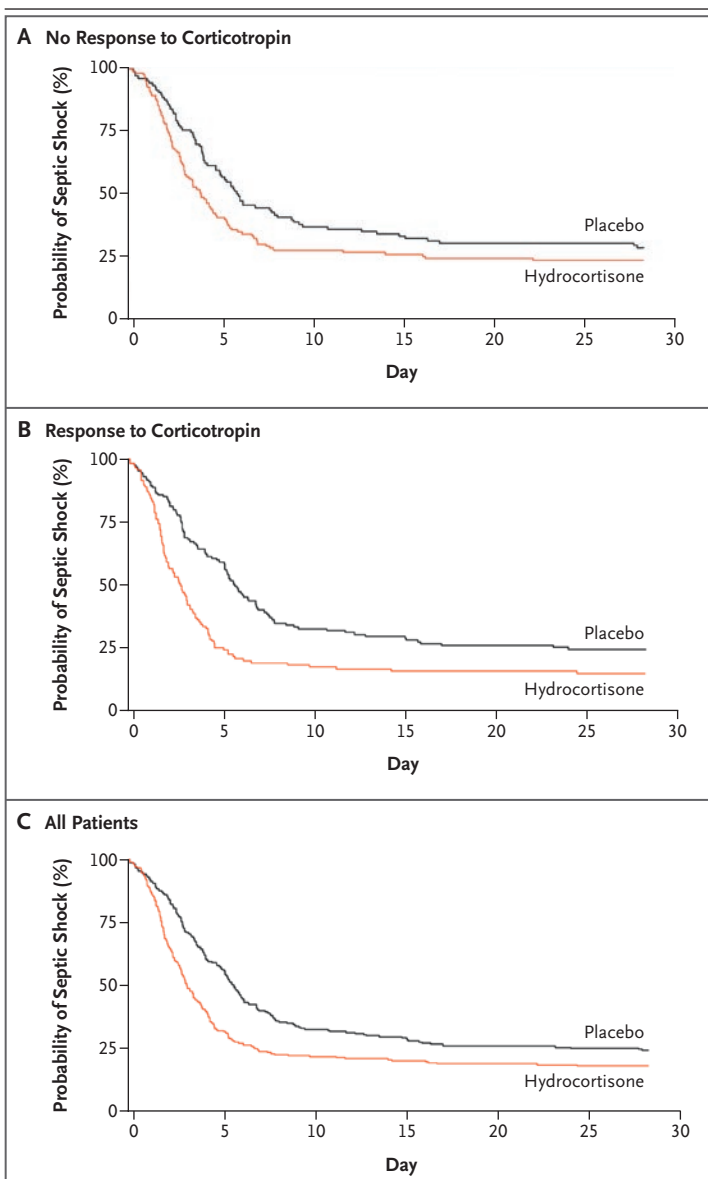
These results are in marked contrast to those of the study by Annane et al.,<sup>16</sup> in which both improved survival and reversal of shock were reported in patients with no response to corticotropin who received hydrocortisone plus fludro-



**Table 3. Outcomes According to Subgroup.\***

Variable	No Response to Corticotropin Hydrocortisone (N=125)	Placebo (N=108)	P Value	Response to Corticotropin Hydrocortisone (N=118)	Placebo (N=136)	P Value	All Patients Hydrocortisone (N=251)	Placebo (N=248)	P Value
Death within 28 days — no. (%)	49 (39.2)	39 (36.1)	0.69	34 (28.8)	39 (28.7)	1.00	86 (34.3)	78 (31.5)	0.51
Relative risk (95% CI)	1.09 (0.77 to 1.52)			1.00 (0.68 to 1.49)			1.09 (0.84 to 1.41)		
Absolute difference — % (95% CI)	3.1 (-9.5 to 15.7)			0.1 (-11.2 to 11.4)			2.8 (-5.5 to 11.2)		
Death in ICU — no./total no. (%)	58/125 (46.4)	44/108 (40.7)	0.43	41/118 (34.7)	45/135 (33.3)	0.89	102/251 (40.6)	89/247 (36.0)	0.31
Relative risk (95% CI)	1.14 (0.85 to 1.53)			1.04 (0.74 to 1.47)			1.13 (0.90 to 1.41)		
Absolute difference — % (95% CI)	5.7 (-7.1 to 18.4)			1.4 (-10.3 to 13.1)			4.6 (-3.9 to 13.1)		
Death during hospitalization — no./total no. (%)	60/125 (48.0)	50/108 (46.3)	0.90	48/118 (40.7)	50/133 (37.6)	0.70	111/251 (44.2)	100/245 (40.8)	0.47
Relative risk (95% CI)	1.04 (0.79 to 1.36)			1.08 (0.79 to 1.47)			1.08 (0.88 to 1.33)		
Absolute difference — % (95% CI)	1.7 (-11.1 to 14.6)			3.1 (-9.0 to 15.2)			3.4 (-5.3 to 12.1)		
Death at 1 yr — no./total no. (%)	73/124 (58.9)	60/105 (57.1)	0.89	61/111 (55.0)	67/126 (53.2)	0.80	137/242 (56.6)	127/235 (54.0)	0.58
Relative risk (95% CI)	1.03 (0.83 to 1.29)			1.03 (0.82 to 1.31)			1.05 (0.89 to 1.23)		
Length of stay — days									
In ICU	17±19	17±17	0.47	18±22	19±16†	0.26	19±31	18±17†	0.51
In hospital	29±26	31±27	0.82	36±40	35±43‡	0.68	34±41	34±37‡	0.47

\* Relative risks and percent differences are for the comparison between the hydrocortisone group and the placebo group. P values for categorical variables were calculated with the use of Fisher's exact test. P values for continuous variables were calculated with the use of the Wilcoxon rank-sum test. ICU denotes intensive care unit.  
 † Data were missing for one patient.  
 ‡ Data were missing for three patients.



**Figure 3.** Kaplan–Meier Curves for the Time to Reversal of Shock.

For the comparison between patients with septic shock who received hydrocortisone and those who received placebo,  $P=0.06$  for patients who did not have a response to a corticotropin test (Panel A) and  $P<0.001$  both for patients who had a response to corticotropin (Panel B) and for all patients (Panel C).

cortisone. Differences between the two studies may stem from several factors. First, the studies involved dissimilar populations of patients. In the Annane study, the patients had higher SAPS II scores at baseline, and the entry requirement for systolic blood pressure was less than 90 mm Hg for more than 1 hour despite fluid and vasopres-

sor therapy; there also was a much higher rate of death at 28 days in the placebo group (61%, as compared with 32% in our study). Second, enrollment in the Annane study was allowed only within 8 hours after fulfilling entry criteria, as compared with a 72-hour window in our study. Third, fludrocortisone was not given to patients in our study, since 200 mg of hydrocortisone should provide adequate mineralocorticoid activity.<sup>28</sup> Furthermore, absorption of oral fludrocortisone is variable in the shock state. Although an analysis of patients in our study who had a systolic blood pressure that persisted below 90 mm Hg at 1 day after fluid and vasopressor resuscitation showed a rate of death of 56.1% in the placebo group and an absolute reduction in mortality of 11.2% in the hydrocortisone group (results that are similar to those reported by Annane et al.), the subgroups of patients who received a study drug within 12 hours after baseline did not show any significant differences in outcome.

As reported previously,<sup>14,15,18</sup> our study showed a decrease in the time to reversal of shock in the hydrocortisone group. However, the total number of patients who underwent reversal of shock was unaffected. It remains unclear why vascular tone improves in some patients but not in others. In an unexpected finding, the earlier rate of reversal of shock was greater in patients who had had a response to corticotropin but was not associated either with survival benefit or a reduction in duration of stay in either the ICU or the hospital. These findings may be unrelated to adrenal insufficiency but could instead result from a direct interaction with mechanisms producing vascular hyporeactivity.<sup>29,30</sup> Alternatively, the effect may be due to a more widespread antiinflammatory action of corticosteroids, which inhibit the expression of proinflammatory cytokines, mediators, and receptors.<sup>31</sup>

The duration of the administration of corticosteroids may be pertinent, with the possibility that any gain that was achieved by an earlier reversal of shock was counterbalanced by later complications.<sup>9</sup> In the Annane study, corticosteroid treatment was stopped abruptly after 7 days, whereas in our study, therapy was tapered from day 5 to day 11. Tapering was used because of the increase in proinflammatory mediators and hemodynamic deterioration after abrupt cessation

**Table 4. Adverse Events (Per-Protocol Population).\***

Event	Hydrocortisone (N = 234)	Placebo (N = 232)	Relative Risk (95% CI)
	<i>no. of patients (%)</i>		
Superinfection	78 (33)	61 (26)	1.27 (0.96–1.68)
Catheter-related	3 (1)	3 (1)	0.99 (0.20–4.86)
Lung	34 (15)	30 (13)	1.12 (0.71–1.77)
Gastrointestinal	22 (9)	19 (8)	1.15 (0.64–2.06)
Urinary tract	11 (5)	10 (4)	1.09 (0.47–2.52)
Wound	9 (4)	7 (3)	1.27 (0.48–3.37)
Other	16 (7)	8 (3)	1.98 (0.87–4.54)
New sepsis	6 (3)	2 (1)	2.97 (0.61–14.59)
New septic shock	14 (6)	5 (2)	2.78 (1.02–7.58)
Other adverse event			
Anastomotic leak	4 (2)	4 (2)	0.99 (0.25–3.92)
Wound dehiscence	2 (1)	2 (1)	0.99 (0.14–6.98)
Repeat shock	72 (31)	57 (25)	1.25 (0.93–1.68)
Bleeding			
Any	21 (9)	16 (7)	1.30 (0.70–2.43)
Gastrointestinal	15 (6)	13 (6)	1.14 (0.56–2.35)
Polyneuropathy	2 (1)	4 (2)	0.50 (0.09–2.68)
Multiple organ system failure	34 (15)	33 (14)	1.02 (0.66–1.59)
Refractory shock	20 (9)	25 (11)	0.79 (0.45–1.39)
Pulmonary	8 (3)	13 (6)	0.61 (0.26–1.44)
Renal	7 (3)	6 (3)	1.16 (0.39–3.39)
Neurologic	1 (<1)	1 (<1)	0.99 (0.06–15.76)
Hyperglycemia (glucose $\geq$ 150 mg/dl on any day from day 1 to day 7) <sup>†</sup>	186 (85)	161 (72)	1.18 (1.07–1.31)
Hypernatremia (sodium $\geq$ 150 mmol/liter on any day from day 1 to day 7) <sup>‡</sup>	67 (29)	42 (18)	1.58 (1.13–2.22)
Possibly related to shock			
Stroke	3 (1)	1 (<1)	2.97 (0.31–28.39)
Acute myocardial infarction	14 (6)	13 (6)	1.24 (0.34–4.56)
Peripheral limb ischemia	0	1 (<1)	

\* Some patients had more than one adverse event. Relative risks are for the comparison between the hydrocortisone group and the placebo group. To convert values for glucose to millimoles per liter, multiply by 0.05551.

<sup>†</sup> For the diagnosis of hyperglycemia, 220 patients were evaluated in the hydrocortisone group and 225 patients in the placebo group.

<sup>‡</sup> For the diagnosis of hypernatremia, 231 patients were evaluated in the hydrocortisone group and 229 patients in the placebo group.

of corticosteroids.<sup>17</sup> Our study showed an increased incidence of superinfection, including new episodes of sepsis or septic shock, in the hydrocortisone group. Previous studies with high-dose corticosteroids have shown similar find-

ings,<sup>11</sup> whereas the study conducted by the Acute Respiratory Distress Syndrome (ARDS) Network<sup>32</sup> using higher doses of corticosteroids and meta-analyses of studies that used low doses<sup>21,22</sup> did not report higher rates of infectious complications.

Studies involving critically ill patients have reported an association between corticosteroid therapy and the incidence of neuromuscular weakness.<sup>32,33</sup> We did not see this in our study, although electrophysiological testing was not performed. However, the duration of mechanical ventilation was similar in the two study groups. Finally, the increased glucose levels in the hydrocortisone group may have contributed to increased mortality.<sup>34</sup>

The use of etomidate for induction of anesthesia in our study (in 26% of patients) was similar to that in the Annane study (24%). Etomidate has a low profile of cardiovascular complications,<sup>35</sup> but a single dose can inhibit the metabolism of corticosteroids for at least 24 hours in patients who are critically ill.<sup>36</sup> An association between etomidate and the likelihood of adrenal hypo-responsiveness was also found in our study.

The prognostic importance of adrenal insufficiency in septic shock is well described.<sup>19</sup> Routine testing of adrenal function has been advocated to guide corticosteroid therapy.<sup>16,18-21</sup> In our study, a modest increase in the rate of death at 28 days was seen in patients who did not have a response to corticotropin (38%), as compared with those who had a response (29%). However, there was no difference in outcome in either subgroup of the hydrocortisone group. The short corticotropin test does not appear to be useful for determining the advisability of corticosteroid treatment in patients with septic shock, and our results call into question the definition of relative adrenal insufficiency. Indeed, significant variability in cortisol levels has been described, depending on the measurement methods used.<sup>37</sup> Studies have described the poor relationship between total and free cortisol levels<sup>38</sup> and other issues concerning the dose, timing, and type of corticotropin.<sup>39</sup>

The strengths of our study include the fact that it was initiated by the investigators and involved 52 ICUs in nine countries. Limitations include the lack of adequate power, since only 500 patients were enrolled rather than the projected 800. This was due to a combination of slow recruitment (which was probably related to a loss of equipoise in view of the various guidelines recommending corticosteroid use<sup>23</sup>), termination of funding, and time expiry of the trial drug. On

the basis of the current data, however, the likelihood of seeing any difference in outcomes between the two study groups was unlikely. Finally, 21 patients received open-label corticosteroids (4.2%), although this finding was unlikely to have had a material effect on the outcome.

In summary, the use of hydrocortisone did not decrease mortality in a general population of patients with septic shock, even though the drug hastened reversal of shock. This lack of improvement may be related to an increased incidence of superinfection and new septic episodes. No benefit was seen in a subgroup of patients who had had no response to corticotropin, as was shown previously for patients with severe septic shock. This finding may be related to methodologic issues surrounding the accurate diagnosis of adrenal insufficiency in critically ill patients or to a decreased prognostic importance of this phenomenon in less severe shock. On the basis of these findings, hydrocortisone cannot be recommended as general adjuvant therapy for septic shock (vasopressor responsive), nor can corticotropin testing be recommended to determine which patients should receive hydrocortisone therapy. Hydrocortisone may have a role among patients who are treated early after the onset of septic shock who remain hypotensive despite the administration of high-dose vasopressors (vasopressor unresponsive).<sup>16</sup>

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## APPENDIX

In addition to the authors, the following investigators and institutions participated in this study: **Austria:** Landeskrankenhaus Feldkirch, Feldkirch: P. Fae; Krankenhaus Barmherzige Schwestern, Linz: J. Reisinger; Universitätsklinik fuer Innere Medizin II, Vienna: G. Heinz; **Belgium:** Hôpital St. Joseph, Arlon: M. Simon; St. Luc University Hospital, Brussels: X. Wittebole, M.N. France; University Hospital Erasme, Université de Bruxelles, Brussels: J.-L. Vincent, D. DeBacker; Centre Hospitalier Universitaire Charleroi, Charleroi: P. Biston; **France:** Hôpital de Caen, Caen: C. Daubin; Hôpital Raymond Poincaré, Garches: D. Lipiner, V. Maxime; Hôpital Huriez, Lille: P.A. Rodie Talbere, B. Vallet; Hôpital Caremeau, Nîmes: J.Y. Lefrant; Hôpital Saint-Antoine, Paris: G. Offenstadt; Hôpital Lariboisière, Paris: A.C. Lukaszewicz; **Germany:** Zentralklinikum Augsburg, Augsburg: G. Neeser, Y. Barth; Charité Universitätsmedizin Berlin, Campus Virchow-Klinikum, Berlin: J. Langrehr, M. Oppert; Campus Charité Mitte, Berlin: C. Spies, S. Rosseau; Campus Benjamin Franklin, Berlin: J. Weimann; Evangelisches Waldkrankenhaus Spandau, Berlin: M. Reyle-Hahn; St. Joseph Krankenhaus, Berlin: M. Schmutzler; Vivantes Klinikum Spandau, Berlin: K.J. Slama; Vivantes Klinikum Neukoelln, Berlin: H. Gerlach; Vivantes Klinikum im Friedrichshain, Berlin: S. Veit; Institute for Anesthesia and Operative Intensive Care Medicine, Darmstadt: M. Welte, L. Von Beck; University Hospital Carl Gustav Carus, Dresden: C. Marx; Krankenhaus Hennigsdorf, Hennigsdorf: A. Lange; Friedrich Schiller Universität, Jena: F. Bloos, F. Brunkhorst; Klinikum Kempten-Oberallgaeu, Kempten: M. Haller; Klinikum Landshut, Landshut: U. Helms; Klinikum Mannheim, Mannheim: F. Fiedler; Universitätsklinikum Marburg, Marburg: M. Max; Klinikum der Universität, Ludwig Maximilians Universität, Munich: W. Hartl; Staedisches Krankenhaus Muenchen Harlaching, Munich: M. Klimmer, T. Helmer; Universität Erlangen-Namberg, Nuerenberg: M. Baumgaertel; Klinikum Ernst von Bergman, Potsdam: D. Pappert; **Israel:** Haemek Hospital, Afula: A. Lev; Hadassah Medical Organization, Jerusalem: O. Shatz; Belinson Medical Centre, Petach Tikva: P. Singer; Ichilov Hospital, Tel Aviv: A. Hallam; **Italy:** Policlinico di Tor Vergata, Rome: S. Natoli; Centro di Rianimazione, Ospedale S. Eugenio, Rome: F. Turani; **the Netherlands:** Erasmus University Medical Center, Rotterdam: B. Van der Hoven; **Portugal:** Hospital de St. Antonio do Capuchos, Lisbon: R. Matos; **United Kingdom:** Aberdeen Royal Infirmary, Aberdeen: S. Roughton; Ipswich Hospital National Health Service (NHS), Ipswich: M. Garfield; General Infirmary at Leeds, Leeds: A. Mallick; University College London Hospitals NHS Foundation Trust, London: M. McKendry; Southampton General Hospital, Southampton: T. Woodcock.

Committee members for the study were as follows: **Steering Committee:** C. Sprung (chair), D. Annane, J. Briegel, D. Keh, R. Moreno, D. Pittet, M. Singer, Y. Weiss; **Safety and Efficacy Monitoring Committee:** J. Cohen (chair), C. Dore, T. Evans, N. Soni, F. Sorenson (Analytica International); **Study Coordinating Center:** C. Sprung (physician coordinator), J. Benbenishty (nurse coordinator), A. Avidan, E. Ludmir, J. Kabiri, K. Furmanov, B. Hain, O. Kalugin, I. Zack; **Clinical Evaluation Committee:** Y. Weiss (chair), D. Annane, J. Briegel, S. Goodman, D. Keh, R. Moreno, M. Singer, C. Sprung; **Berlin Coordinating Center:** D. Keh (chair), A. Goessinger; **French Coordinating Center:** D. Annane (chair), N. Zinsou, D. Friedman; **Munich Central Laboratory Harmonization:** J. Briegel (chair), M. Vogeser; **Statistical Analysis:** Analytica International, F. Sorenson, K. Freivogel.

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## CORRESPONDENCE



## Corticosteroids for Septic Shock

**TO THE EDITOR:** On the basis of the Corticosteroid Therapy of Septic Shock (CORTICUS) study, Sprung et al. (Jan. 10 issue)<sup>1</sup> report that hydrocortisone does not improve survival among patients with septic shock. However, it is disturbing that of the 357 study patients with culture-positive sepsis, 86 (24%) were considered not to have received appropriate antimicrobial therapy. This was determined retrospectively by a clinical evaluation committee. The actual mortality rates among patients receiving appropriate antimicrobial therapy as compared with those receiving inappropriate antimicrobial therapy are not provided. Antimicrobial agents remain the cornerstone of therapy for sepsis.<sup>2</sup> However, this study shows that basic antimicrobial therapy continues to be clinically problematic in sepsis. A trial of septic shock in which one quarter of the culture-positive patients receive inappropriate antimicrobial therapy seems to defeat the hypothesized purpose of adjunctive corticosteroid use. It is unfortunate that the rec-

ommendations of the clinical evaluation committee regarding antimicrobial therapy were not available and were applied prospectively instead of retrospectively in this trial.

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1. Sprung CL, Annane D, Keh D, et al. Hydrocortisone therapy for patients with septic shock. *N Engl J Med* 2008;358:111-24.
2. Bochud PY, Glauser MP, Calandra T. Antibiotics in sepsis. *Intensive Care Med* 2001;27:Suppl 1:S33-S48.

**TO THE EDITOR:** The failure of the CORTICUS study group to reach the enrollment target of 800 patients may have resulted from physician bias against enrolling more severely ill patients, given previously published data on the benefits of corticosteroids in septic shock.<sup>1-3</sup> The extent of this bias cannot be assessed, because the number of patients screened before randomization is not provided. The largest previous trial of low-dose corticosteroids for the treatment of septic shock enrolled patients in whom hypotension had persisted for 1 hour despite fluid resuscitation and vasopressor treatment.<sup>1</sup> In contrast, the CORTICUS study group enrolled patients who were not as severely ill, requiring only a systolic blood pressure of less than 90 mm Hg for 1 hour despite adequate fluid resuscitation or need for vasopressors.

The benefit of antiinflammatory agents in sepsis is directly proportional to the severity of illness.<sup>4</sup> The control-group mortality was 36% in the CORTICUS study, as compared with 63% in the study by Annane et al.<sup>1</sup> The lack of a treatment benefit in the CORTICUS study may be due to the enrollment of a population of patients with a

## THIS WEEK'S LETTERS

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lower mortality, for whom the risk of side effects of corticosteroid therapy outweighs its potential benefit.

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1. Annane D, Sébille V, Charpentier C, et al. Effect of treatment with low doses of hydrocortisone and fludrocortisone on mortality in patients with septic shock. *JAMA* 2002;288:862-71.
2. Minneci PC, Deans KJ, Banks SM, et al. Meta-analysis: the effect of steroids on survival and shock during sepsis depends on the dose. *Ann Intern Med* 2004;141:47-56.
3. Annane D, Bellissant E, Bollaert PE, Briegel J, Keh D, Kupfer Y. Corticosteroids for severe sepsis and septic shock: a systematic review and meta-analysis. *BMJ* 2004;329:480-8.
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**TO THE EDITOR:** Sprung et al. report the use of hydrocortisone at a physiologic dose to treat patients with septic shock. The study population was divided into two groups on the basis of their response to a 250- $\mu$ g corticotropin stimulation test. The absence of a response to corticotropin stimulation was defined as an increase in the serum cortisol level of less than 9  $\mu$ g per deciliter (248 nmol per liter). We have several points of concern. First, 24.8% of the nonresponders received etomidate, an anesthetic agent that also blocks the adrenal enzyme 11 $\beta$ -hydroxylase and cortisol synthesis.<sup>1</sup> We think that absolute cortisol levels and the cortisol area under the curve may be more precise ways to define adrenal insufficiency in septic shock.<sup>2,3</sup> In fact, basal cortisol levels were higher in the group without a response than in the group with a response. It is evident that the mean basal cortisol level was high in the nonresponders and that the standard deviation was very large. This suggests that the group was heterogeneous, clearly including not only patients with real adrenal insufficiency but also a sizable number of patients with normal cortisol secretion.

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1. Jackson WL Jr. Should we use etomidate as an induction agent for endotracheal intubation in patients with septic shock? A critical appraisal. *Chest* 2005;127:1031-8.
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**TO THE EDITOR:** Sprung et al. found no improvement in survival or shock reversal and an increase in superinfections. These conclusions contrast with those of a previous multicenter study<sup>1</sup> and of two meta-analyses<sup>2,3</sup> and should not lead to the abandonment of corticosteroids in this setting for several reasons. First, as emphasized in the accompanying editorial, the study was underpowered to detect an important effect of treatment on mortality. In addition, in a subgroup of patients whose illness was more severe than that in the entire study population, an absolute reduction in mortality of 11.2% was observed. Second, the rate of shock reversal at 28 days is less relevant than the rate observed earlier, since it underestimates what happens during the first days of treatment. Corticosteroids increase the pressor response to catecholamines,<sup>4</sup> which explains, at least in part, why a noticeable effect on shock reversal was observed on day 7 in all studies (including the current study reported by Sprung et al.). Third, since new episodes of sepsis and septic shock appeared to account for the increase in superinfections, we wonder how many episodes were confirmed by microbiologic assessment and how they were distinguished from possible shock rebound after corticosteroid treatment.

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**TO THE EDITOR:** We believe that selection bias confounds interpretation of the CORTICUS study findings for three reasons. First, the demographic characteristics of the enrolled patients differ from those of patients enrolled in the major sepsis trials of the past decade. Second, it appears that only 1 of 25 eligible patients was enrolled in the CORTICUS study.<sup>1</sup> Third, the study was terminated prematurely because of slow recruitment, which is odd, given that septic shock is the most com-



mon admission diagnosis in intensive care units (ICUs) worldwide.<sup>1</sup> The high prevalence of septic shock<sup>1</sup> and the overwhelming use of corticosteroids in its management<sup>2</sup> are critically disconnected from the stated inability to enroll patients in this study; this suggests that many of those screened — but not eligible — were ineligible because they were already receiving corticosteroids. We therefore believe that the fairest conclusion might be that corticosteroids were not likely to benefit patients with septic shock when their physicians had already decided that they were not sufficiently ill to warrant such therapy.

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**TO THE EDITOR:** Sprung et al. report that patients with septic shock who received etomidate had increased mortality. Etomidate decreases serum cortisol levels by inhibiting 11  $\beta$ -hydroxylase, and continuous etomidate infusions have been found to increase mortality among critically injured patients.<sup>1</sup> Retrospective studies of mortality and the use of single-dose etomidate in sepsis have yielded conflicting results,<sup>2,3</sup> and etomidate is less likely to cause hypotension and cardiac depression than other induction agents.<sup>3</sup> Although the assessment of hydrocortisone by Sprung et al. was prospective, their assessment of etomidate was not. Clinicians who are aware of etomidate's benign hemodynamic profile and potentially adverse metabolic effects may reserve it for patients who are most hemodynamically unstable at intubation. These patients may be more likely to die, regardless of treatment. In addition, it is notable that patients who received etomidate and hydrocortisone in the CORTICUS trial were at least as likely to die as pa-

tients who received etomidate but did not receive corticosteroid replacement. These considerations argue for a well-designed prospective trial of single-dose etomidate for intubation of patients with septic shock.

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**THE AUTHORS REPLY:** In response to Daley: 24% of patients were classified as not receiving appropriate antimicrobial therapy because of the strict criteria used by the clinical evaluation committee and frequent inability to confirm appropriateness. This included cases in which laboratory sensitivities were not reported for pathogens isolated or for the class of antibiotic used and cases in which prescribed antibiotics did not cover all of the multiple pathogens isolated. It is difficult to establish prospective antimicrobial guidelines since bacterial sensitivity patterns vary among ICUs and hospitals. A clinical evaluation committee is in the position of making retrospective analyses, when pathogens and sensitivity patterns are known. We believe that most of our patients did receive appropriate antibiotics, although some may have been classified as having received inappropriate antibiotics because of the lack of data. A higher mortality was seen among patients classified as receiving appropriate antimicrobial agents as compared with those not receiving appropriate antibiotics (35% vs. 23%).

We agree with Seam that the patients in the CORTICUS study were not as severely ill as those in the study by Annane et al.,<sup>1</sup> with lower mortality in the placebo group, lower scores on the Simplified Acute Physiology Score II, and fewer patients who did not have a response to corticotropin. Although corticosteroids may be effective if given early in a population of severely ill patients who do not have a response to vasopressor therapy,<sup>1</sup> the CORTICUS population represents patients who are more typical of those usually

seen. We appreciate Luboshitzky and Qupti's comments, but the definition and optimal diagnostic testing of adrenal insufficiency in septic shock remain controversial and elusive.

In response to Bollaert: although our study was underpowered, it is still, to our knowledge, the largest performed to date. Given that the relative risk of death was increased with corticosteroid use, the likelihood of a positive finding for hydrocortisone would have been low even if the targeted number of patients had been enrolled. It is important to assess subgroups that have been defined prospectively, since post hoc subgroup analyses can be misleading. The majority of patients with superinfections did not have new sepsis or septic shock. Among patients with new episodes of sepsis, microbiologic confirmation occurred in more than 50%. Although we acknowledge the difficulty in discriminating between rebound inflammation and new sepsis, repeat shock episodes in the hydrocortisone and placebo groups were similar.

In response to Marik et al., the patients in the CORTICUS study actually had 28-day and ICU mortality rates that were similar not only to those in the trial they cite but also to those in other contemporary trials involving patients with septic shock.<sup>2</sup> Consequently, our patients were sufficiently ill to warrant corticosteroid therapy. Sepsis is not septic shock. In the Sepsis Occurrence in Acutely Ill Patients (SOAP) study,<sup>3</sup> only 15% of patients had septic shock at some time during their ICU admission (7.7% on admission) and many would not have met our inclusion criteria. We acknowledge that we lack precise information for eligible patients who were not enrolled. The slow-

ness of recruitment was due in part to the many patients who were ineligible because they had already been given corticosteroids on the basis of recommendations for their use, but the study investigators had clinical equipoise for the use of corticosteroids in enrolled patients. As we noted, hydrocortisone use may benefit the small subgroup of patients with early, unresponsive septic shock. However, on the basis of currently available evidence, corticosteroids cannot be recommended for the majority of patients with septic shock who do have a response to vasopressors,<sup>4</sup> particularly in view of the increased risk of harmful complications.

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## Insulin and Pentastarch for Severe Sepsis

**TO THE EDITOR:** Brunkhorst and colleagues (Jan. 10 issue)<sup>1</sup> report that the Efficacy of Volume Substitution and Insulin Therapy in Severe Sepsis (VISEP) study established “that intensive insulin therapy has no measurable, consistent benefit in critically ill patients” in a medical intensive care unit (ICU). This interpretation appears too conclusive, since the Supplementary Appendix of the article (available with the full text of the article at [www.nejm.org](http://www.nejm.org)) indicates that the goal of normoglycemia was reached in less than 50% of the patients in the intensive insulin-therapy group.

In addition to the intrinsic properties of an intensive insulin-therapy protocol, heterogeneity in the extrinsic factors (e.g., the level of training of nurses in the protocol and their workload) among centers participating in multicenter trials further influences the risk of failure to achieve normoglycemia.<sup>1,2</sup> The intensive-therapy protocol used by Brunkhorst et al. was previously used in two single-center trials, with a 70% rate of successful glycemic control.<sup>3-5</sup> To assess, in large multicenter trials, whether normoglycemia influences outcomes, the newly implemented protocol