Liver dysfunction associated with artificial nutrition in critically ill patients

Teodoro Grau¹, Alfonso Bonet², Mercedes Rubio³, Dolores Mateo⁴, Mercé Farré⁵, José Antonio Acosta⁶, Antonio Blesa⁷, Juan Carlos Montejo⁸, Abelardo García de Lorenzo⁹, Alfonso Mesejo¹⁰ and the Working Group on Nutrition and Metabolism of the Spanish Society of Critical Care

¹Intensive Care Unit, Hospital Severo Ochoa. Av. Orellana s/n, 28911 Leganés, Madrid, Spain
²Intensive Care Unit, Hospital Josep Trueta. Av. de Francia s/n, 17007 Girona, Spain
³Cardiovascular Intensive Care Unit, Hospital Universitario 12 de Octubre. Av. de Córdoba s/n, 28041 Madrid, Spain
⁴Intensive Care Unit, Newham University Hospital NHS Trust. Glen Road, Plaistow London E13 8SL, UK
⁵Intensive Care Unit, Hospital Universitari Vall d'Hebró. Paseo Vall d'Hebró 119-129, 08035 Barcelona, Spain
⁶Intensive Care Unit, General de Alicante. Maestro Alonso 109, 03010 Alicante, Spain
⁷Intensive Care Unit, Hospital Clínico San Carlos. Profesor Martin Lagos s/n, 28040 Madrid, Spain
⁸Intensive Care Unit, Hospital Universitario Doce de Octubre.Av. de Córdoba s/n, 28041 Madrid, Spain
⁹Intensive Care Unit, Hospital Universitario La Paz. Paseo de la Castellana 261, 28046 Madrid, Spain
¹⁰Intensive Care Unit, Hospital Universitario La Fe. Av. Campanar 21, 46009 Valencia, Spain

Corresponding author: Teodoro Grau, tgrau.hdoc@salud.madrid.org

Received: 20 Jul 2006 Revisions requested: 1 Sep 2006 Revisions received: 30 Nov 2006 Accepted: 25 Jan 2007 Published: 25 Jan 2007

Critical Care 2007, 11:R10 (doi:10.1186/cc5670)

This article is online at: http://ccforum.com/content/11/1/R10

© 2007 Grau 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 Liver dysfunction associated with artificial nutrition in critically ill patients is a complication that seems to be frequent, but it has not been assessed previously in a large cohort of critically ill patients.

Methods We conducted a prospective cohort study of incidence in 40 intensive care units. Different liver dysfunction patterns were defined: (a) cholestasis: alkaline phosphatase of more than 280 IU/l, gamma-glutamyl-transferase of more than 50 IU/l, or bilirubin of more than 1.2 mg/dl; (b) liver necrosis: aspartate aminotransferase of more than 40 IU/l or alanine aminotransferase of more than 40 IU/l or alanine aminotransferase of more than 42 IU/l, plus bilirubin of more than 1.2 mg/dl or international normalized ratio of more than 1.4; and (c) mixed pattern: alkaline phosphatase of more than 280 IU/l or gamma-glutamyl-transferase of more than 50 IU/l, plus aspartate aminotransferase of more than 40 IU/l or alanine aminotransferase of more than 42 IU/l.

Results Seven hundred and twenty-five of 3,409 patients received artificial nutrition: 303 received total parenteral nutrition (TPN) and 422 received enteral nutrition (EN). Twenty-three percent of patients developed liver dysfunction: 30% in the TPN group and 18% in the EN group. The univariate analysis showed an association between liver dysfunction and TPN (p < 0.001), Multiple Organ Dysfunction Score on admission (p < 0.001), sepsis (p < 0.001), early use of artificial nutrition (p < 0.03), and malnutrition (p < 0.01). In the multivariate analysis, liver dysfunction was associated with TPN (p < 0.001), sepsis (p < 0.02), early use of artificial nutrition (p < 0.03), and calculated energy requirements of more than 25 kcal/kg per day (p < 0.05).

Conclusion TPN, sepsis, and excessive calculated energy requirements appear as risk factors for developing liver dysfunction. Septic critically ill patients should not be fed with excessive caloric amounts, particularly when TPN is employed. Administering artificial nutrition in the first 24 hours after admission seems to have a protective effect.

APACHE II = Acute Physiology and Chronic Health Evaluation II; CI = confidence interval; EN = enteral nutrition; ICU = intensive care unit; INR = international normalized ratio; IQ = interquartile; LCT = long-chain triglyceride; LD = liver dysfunction; MCT = medium-chain triglyceride; MODS = Multiple Organ Dysfunction Score; OR = odds ratio; TPN = total parenteral nutrition.
Introduction
Artificial nutrition support is part of the standard of care in critically ill patients [1]. Some of these patients have sepsis or systemic inflammatory response syndrome, which produce hypermetabolism, accelerated lipolysis, insulin resistance, and protein catabolism. These phenomena, associated with the lack of oral intake, can lead to malnutrition. Artificial nutrition usually does not reverse these metabolic derangements but can decrease the depletion of the lean body mass [2]. Hepatobiliary complications related to artificial nutrition have been widely reported, particularly in patients receiving total parenteral nutrition (TPN), and less frequently in patients receiving enteral nutrition (EN) [3]. There are many potential causes of liver dysfunction (LD) related to artificial nutrition, but the etiology is unclear and there are few data on the prevalence in critically ill patients. Moreover, these patients can present hepatic dysfunction as part of the multiple organ failure syndrome [4]. The aim of this study was to assess the prevalence of hepatobiliary complications related to artificial nutrition, the risk factors associated with these complications, and their influence on the prognosis in critically ill patients.

Materials and methods
Design
This study was designed as a multicenter prospective cohort study of incidence of LD in patients admitted to any of the 40 participating intensive care units (ICUs) from tertiary hospitals in Spain between 1 March and 15 April 2000. Patients were enrolled consecutively when the treating physician expected them to need artificial nutrition for five days or more. The protocol and definitions of LD were established previously in a meeting with the participants. The institutional review board of each participating hospital approved the study. Informed consent was waived according to these boards and Spanish law. Our funding sources had no role in the acquisition, analysis, or interpretation of data or in the submission of this report.

Patients
Patients entered in the study were followed prospectively until hospital discharge or 28 days after ICU admission to check mortality at that time. Age, gender, weight, primary diagnosis, group (medical, surgical, or trauma), APACHE II (Acute Physiology and Chronic Health Evaluation II) score [5], Multiple Organ Dysfunction Score (MODS) [4], the need for mechanical ventilation, and the presence and origin of sepsis and/or septic shock were recorded on admission. The diagnosis of sepsis or septic shock on admission was made according to previously published criteria [6]. Sepsis was defined when a patient had a confirmed infection with two or more of the following criteria: (a) temperature greater than 38°C or less than 36°C, (b) heart rate greater than 90 beats per minute, (c) respiratory rate greater than 20 respirations per minute or PaCO₂ (partial pressure of carbon dioxide) less than 32 mm Hg, and (d) leukocytes greater than 12,000 per cubic millimeter or greater than 10% band neutrophils. Septic shock was defined as arterial hypotension induced by sepsis, which persists in spite of the adequate replacement of fluids and associated with hypoperfusion and organ dysfunction. Exclusion criteria were age of less than 18 years, expected survival of less than 24 hours, or previous cardiopulmonary resuscitation. Patients with previously recognized liver disease were excluded by the following criteria: (a) portal hypertension with gastrointestinal bleeding at the time of admission and/or transfer, (b) clinically apparent ascites on a hepatocellular basis, (c) total bilirubin of more than 3 mg/dl or aspartate aminotransferase of more than 40 IU/l or on a hepatocellular basis, (d) serum albumin of less than 0.03 g/l with portal hypertension, (e) encephalopathy of grade II or greater, and (f) clinical diagnosis of alcoholic hepatitis [7].

Choice of the type of nutrition
The clinician responsible for the patient chose the type of nutrition, the administration route, and the type of diet following the published recommendations [8]. The protocol was discussed in previous meetings with the researchers. The use of early artificial nutrition was encouraged to the participants. EN was recommended as the preferred route for feedings if the patient’s gastrointestinal system was preserved. Clinicians could switch to TPN if the patient did not tolerate EN due to gastrointestinal complications or if 75% of the caloric requirements were not achieved after three days of enteral feedings. Also, clinicians were allowed to administer EN for as long as the gastrointestinal function was recovered. In both cases, the amount of calories was limited to the planned caloric intake. TPN was administered through a central venous catheter, with the use of ‘all in one’ ternary mixtures, by means of a continuous pump infusion. The TPN bag was replaced every 24 hours. EN was administered through a nasogastric or nasojejunal tube at the doctor’s discretion and continuously through an infusion pump in accordance with a previously established protocol [9]. The systems used for EN administration were replaced at least once a day, and the feeding tube was flushed on a shift basis three times a day with 20 ml of distilled water. Malnutrition was assessed by means of the Subjective Global Assessment [10]. The calculated nutritional requirements were 25 kcal/kg per day (using the actual weight) with an intake of 1 to 1.5 g of protein/kg per day and a ratio of carbohydrates/fat of 60:40, in agreement with the recommendations published by the SEMICYUC (Spanish Society of Intensive Care) [11]. Fats used in the TPN group were long-chain triglyceride (LCT) or a physical admixture of medium-chain triglyceride (MCT)/LCT, according to the practice of each center. Enteral diets used in the EN group were always polymeric. Once the nutrition had been started, the following parameters were recorded: blood sugar and glucosuria every six hours; urea, creatinine, sodium, potassium, and chloride every 24 hours; and a weekly analysis that included cholesterol, triglycerides, phosphorus, calcium, magnesium, and osmolality. Liver function tests (total and direct bilirubin, aspartate aminotransferase, alanine aminotransferase, gamma-
glutamyl-transferase, and alkaline phosphatase), prothrombin time, and international normalized ratio (INR) were recorded on admission and twice a week (on Tuesday and Friday). The withdrawal of artificial nutrition was defined as the definitive suppression of artificial nutrition, and suspension was defined as a temporary cancellation not longer than 24 hours.

Definitions
The criteria used in this study to define the patterns of LD were the following: (a) cholestasis: alkaline phosphatase of more than 280 IU/l, gamma-glutamyl-transferase of more than 50 IU/l, or bilirubin of more than 1.2 mg/dl; (b) liver necrosis: aspartate aminotransferase of more than 40 IU/l, alanine aminotransferase of more than 42 IU/l, or INR of more than 1.4; and (c) mixed pattern: alkaline phosphatase of more than 280 IU/l, gamma-glutamyl-transferase of more than 50 IU/l, or bilirubin of more than 1.2 mg/dl, plus aspartate aminotransferase of more than 40 IU/l, alanine aminotransferase of more than 42 IU/l, or INR of more than 1.4. These boundaries represent a 10% increase of the normal values in the reference laboratories used. LD was diagnosed when any of the previously defined enzymatic alterations were present. The diagnosis of acalculous cholecystitis was based on clinical criteria and ultrasound. Liver biopsies were not carried out in this study.

Statistical analysis
An intention-to-treat analysis was done for both types of nutrition, TPN and EN. The newly created database was centralized and managed by the main researchers. Any doubts about application of the protocol were discussed with the participants, and the main researchers made the final decision. Once the time of the study was over, the database was closed down. The analysis was blind to the type of nutrition used. The statistical analysis was performed using the SPSS v12 program (SPSS Inc, Chicago, Illinois, USA). The quantitative values were analyzed for normality. The values with normal distribution were compared using the Student’s t test, and the others using non-parametric tests (Kruskall-Wallis test). The qualitative values were analyzed for normality. The values with normal distribution were compared using the Student’s t test, and the others using non-parametric tests (Kruskall-Wallis test). The qualitative values were compared using Fisher’s uncorrected chi-square test, and we calculated the relative risk with the confidence interval (CI) set at 95%. Statistical significance was set at p < 0.05. The quantitative data were expressed as a median and interquartile (IQ) range, and the qualitative data were expressed in absolute values and percentages. The multivariate analysis for LD was carried out by means of a 'stepwise forward' logistical regression model with the most important demographic variables and those that reached statistical significance in the univariate analysis. Time free of LD was analyzed using the Kaplan-Meyer test.

Results

Description of the population
Three thousand four hundred and nine patients were admitted during the study. Seven hundred and fifty-six patients received nutrition in some form, whether TPN or EN, but 31 were excluded and 725 were studied (Table 1). Four hundred and eighty-eight were men and 237 were women. Three hundred and three patients (41.8%) received TPN and 422 (58.2%) received EN as the initial treatment. The patients who received TPN were older than those treated with EN (66 years, IQ range 48 to 73 years, versus 61 years, IQ range 45 to 71 years; p < 0.01) and mainly were women (38% versus 29%; p < 0.05). TPN was mostly used in surgical patients (175/264 versus 89/264; p < 0.001). Two hundred and eight patients had sepsis on admission; of these patients, 105 had septic shock. In both cases, TPN was used more frequently than EN. APACHE II score was higher in the group of patients who received EN (19, IQ range 13 to 23, versus 17, IQ range 12 to 22), without reaching statistical significance. More patients in the EN group required mechanical ventilation (91% versus 79%; p < 0.001). Also, ICU length of stay was longer in patients who received EN (12 days, IQ range 7 to 21 days, versus 8 days, IQ range 5 to 17 days; p < 0.001). Mortality, assessed 28 days after admission, showed no significant differences in either group (Table 2).

The nutritional parameters were different in the two groups of patients. There was a significant statistical association between TPN and severe malnutrition (36% versus 15%; p < 0.001). The calculated energy requirements were similar in both groups as well as the days of artificial nutrition. Nutrition was started early after admission in both groups (median: 1 day, IQ range: 0 to 2 days), without differences between them. The duration of artificial nutrition was also similar in both groups (median: 9 days, IQ range: 5 to 8 days). One hundred and twenty-two patients assigned to the TPN group received EN when the gastrointestinal function recovered, and EN was stopped in 67 because they were unable to achieve the caloric requirements at day 3 or because they had EN-related complications. MCT/LCT admixtures were used in both groups when receiving TPN, without differences between them. Patients with EN received significantly fewer calories per kilogram on day 1 (20.8, IQ range 15.7 to 25, versus 22.9, IQ range 217.57 to 27.67; p < 0.01) and day 3 of the study (22.5, IQ range 17.65 to 26.87, versus 24.1, IQ range 20 to 29.45; p < 0.005) (Table 3).

LD and artificial nutrition
One hundred and sixty-six patients (23%) had LD. There was a significant statistical association between the appearance of LD and age (p < 0.01), the MODS score (p < 0.001), in surgical (35%) and trauma patients (41%) (p < 0.03), if they had sepsis (p < 0.001) or septic shock on admission (p < 0.02), and in patients who were mechanically ventilated (p < 0.02). The stay in the ICU (16 days, IQ range 8 to 28 days, versus 9 days, IQ range 5 to 17 days; p < 0.001) and in the hospital (28 days, IQ range 17 to 29 days, versus 23 days, IQ range 14 to 28 days; p < 0.01) was longer in the group with LD. No difference in mortality was shown between the two groups (Table 4). The patients with LD were less nourished (33% versus...
21%; \( p < 0.01 \) and were treated mostly with TPN (30% versus 18%; \( p < 0.001 \)) for more days (13 days, IQ range 8 to 25, versus 8 days, IQ range 4 to 16 days; \( p < 0.001 \)). Patients fed early had significantly less LD. The use of MCT/LCT admixtures was similar in patients with or without LD, but the calculated energy requirements were higher (25.54 kcal/kg per day, IQ range 24.49 to 30 kcal/kg per day, versus 25 kcal/kg per day, IQ range 23.33 to 29.41 kcal/kg per day; \( p < 0.05 \)) (Table 5).

LD, TPN, and type of patients
In the univariate analysis, 91 patients treated with TPN developed some form of LD but only 75 in the EN group did (odds ratio [OR] 1.7, 95% CI 1.3 to 2.2) (Table 6). Surgical patients (31% versus 16%; OR 1.8, 95% CI 1.02 to 3.1) and trauma patients (52% versus 23%; OR 2.1, 95% CI 1.1 to 4) treated with TPN had more LD. This association was maintained for all types of LD: cholestasis (OR 1.7, 95% CI 1.04 to 2.9), liver necrosis (OR 1.95, 95% CI 1.1 to 3.42), and mixed pattern (OR 1.8, 95% CI 1.3 to 2.6). The patients with sepsis and TPN showed a higher incidence of LD than the group treated with EN (39% versus 24%; OR 1.6, 95% CI 1.02 to 2.4), although no type of LD was greater in either group. When looking at the time free of LD, EN increased the time free of disease in surgical patients only in the Kaplan-Meyer survival test (Figure 1). Only three patients were diagnosed with acalculous cholecystitis.

Multivariate analysis
The risk factors associated with LD in the multivariate analysis were TPN (OR 1.96, 95% CI 1.3 to 2.97, \( p < 0.001 \)), the early use of artificial nutrition (TPN or EN) the first day after admission (OR 0.6, 95% CI 0.4 to 0.9, \( p < 0.01 \)), MODS (OR 1.1, 95% CI 1.04 to 1.2, \( p < 0.001 \)), and the diagnosis of sepsis on admission (OR 1.76, 95% CI 1.08 to 2.9, \( p < 0.02 \)). The rest of the variables analyzed, such as age, gender, APACHE II score, septic shock on admission, medical patients, surgical patients, mechanical ventilation, the use of MCT/LCT admixtures, or severe malnutrition, did not reach statistical significance in the logistical regression model (Table 7).

Discussion
Our study shows that the incidence of LD associated with artificial nutrition in seriously ill patients is low (23%) and is more frequent in patients who received TPN, with sepsis on admission, and when the planned calculated caloric intake was higher than 25 kcal/kg per day. LD is a widely recognized complication associated with the use of artificial nutrition, particularly TPN, with an incidence of between 25% and 100% [12,13]. Acalculous cholecystitis was diagnosed in only three patients who received TPN, with an incidence of close to the 4% published elsewhere [12].

Multiple factors are related to LD associated with TPN, linked to the type of formulation or the appearance of nutritional deficiencies with the use of TPN [13-16]. Some of these factors are shortage of essential fatty acids [17,18], excessive caloric intake [19], imbalance in the composition of the amino acids [20] or of the non-protein substrates [21], fat deposit in the liver [22], a cholestatic effect of the amino acids [24], the absence of choline [25], production of endotoxins and lithocholic acid due to intestinal bacterial overgrowth [26], shortage of carnitine [27], or the absence of enteral nutritional intake [28,29].

However, few studies examine the risk factors attributable to the clinical state of the patient. The aims of this study were to identify the relationship between the appearance of LD and the use of artificial nutrition and to identify the contributing factors specific to the critically ill patient (severity scores, associated co-morbidity such as sepsis, and mechanical ventilation) which can act as confusion factors. Many studies have demonstrated the superiority of EN over TPN, both in surgical patients [30-33] and in patients admitted to the ICU [34,35]. Our results show that patients who received EN had a lower incidence of LD. Most patients who received EN were medi-
### Table 2

#### Demographic data

|                          | TPN   | EN   | Total | p   |
|--------------------------|-------|------|-------|-----|
| Number of patients       | 303   | 422  | 725   |     |
| Women                    | 114 (38%) | 123 (29%) | 237 (33%) | 0.02 |
| Age in years             | 66 (48–73) | 61 (45–71) | 63 (47–72) | 0.01 |
| APACHE II score          | 17 (12–22) | 19 (13–23) | 18 (12–22) | 0.08 |
| MODS                     | 5 (3–8) | 5 (3–7) | 5 (3–7) | 0.95 |
| Primary diagnosis        |       |       | 0.001 |     |
| Gastrointestinal surgery | 145   | 33   | 178   |     |
| Respiratory failure      | 21    | 112  | 133   |     |
| Stroke                   | 22    | 103  | 125   |     |
| Cardiovascular           | 36    | 50   | 86    |     |
| Trauma                   | 19    | 64   | 83    |     |
| Infections in non-immunosuppressed patients | 18 | 22 | 40 |     |
| Infections in immunosuppressed patients | 4 | 7 | 11 |     |
| Metabolic diseases       | 5     | 5    | 10    |     |
| Urology                  | 4     | 6    | 10    |     |
| Hematology               | 7     | 2    | 9     |     |
| Poisoning                | 4     | 4    | 8     |     |
| Obstetrics/Gynecology    | 6     | 1    | 7     |     |
| AIDS                     | 1     | 1    | 2     |     |
| Other                    | 11    | 12   | 23    |     |
| Type of patients         |       |       | 0.001 |     |
| Medical                  | 105   | 257  | 362   |     |
| Surgical                 | 175   | 89   | 264   |     |
| Trauma                   | 23    | 76   | 99    |     |
| Sepsis on admission      | 122 (40%) | 86 (20%) | 208 (29%) | 0.001 |
| Septic shock on admission | 70 (23%) | 35 (8%) | 105 (15%) | 0.001 |
| Patients on mechanical ventilation | 239 (79%) | 382 (91%) | 621 (86%) | 0.001 |
| Days of mechanical ventilation | 7 (2–16) | 9 (4–17) | 8 (3–16) | 0.001 |
| Intensive care unit length of stay in days | 8 (5–17) | 12 (7–21) | 10 (6–20) | 0.001 |
| Hospital length of stay in days | 25 (15–29) | 25 (15–28) | 25 (15–29) | 0.6 |
| Mortality at 28 days     | 85 (28.1%) | 119 (28.2%) | 204 (28%) | 0.9 |

Parenthetical values indicate range or percentage. APACHE II, Acute Physiology and Chronic Health Evaluation II; EN, enteral nutrition; MODS, Multiple Organ Dysfunction Score; TPN, total parenteral nutrition.
were more in need of mechanical ventilation, and had a longer stay in the ICU but showed less LD (18% in the EN group versus 30% in the TPN group). This result is strong enough because we have performed an ‘intention to treat analysis,’ and the 16% of the patients on EN also received TPN. We have found that other factors, such as previous gastrointestinal surgery or sepsis on admission, can explain the greater incidence of LD shown in the results of our study and in other studies [36,37].

Our study shows that cholestasis and the mixed pattern are the two most frequent types of LD. The elevations of serum transaminases, alkaline phosphatase, and bilirubin are the changes most often associated with the use of TPN [38,39]. Although the increase of serum transaminases usually takes place in the first two or three weeks of TPN, it is unusual to observe a significant increase of bilirubin in this period, at least in adult patients [40-42]. In many cases, these enzymatic alterations are mild and transient, even without the interruption of TPN, and only occasionally lead to liver steatosis. Fat infiltration and intrahepatic cholestasis are the typical findings in these patients [28,43,44]. Liver biopsies showed that the predominant finding in patients with enzymatic alterations is liver steatosis [3,11]. When biopsies are carried out in different periods of time, steatosis is an early and sometimes transient phenomenon, whereas cholestasis is a later finding and generally persists during the TPN. Nevertheless, there are contradictory data between an abnormal level of the hepatic enzymes and steatosis or cholestasis [43,44]. Interestingly, our data show that the early use of artificial nutrition, TPN or EN, can delay the

### Table 3

**Nutritional parameters**

|                        | TPN     | EN      | Total    | $p$  |
|------------------------|---------|---------|----------|------|
| Weight                 | 70 (63–80) | 73 (65–80) | 72 (65–80) | 0.2  |
| Nutritional status     |         |         |          | 0.001|
| Moderate malnutrition  | 76 (25%)| 49 (12%)| 125 (17%)|      |
| Severe malnutrition    | 33 (11%)| 14 (3%) | 47 (7%)  |      |
| Energy requirements per kilogram | 25 (23.29–29.37) | 25 (23.76–30) | 25 (23.64–29.74) | 0.7  |
| Patients receiving TPN | -       | 67      |          |      |
| Patients receiving EN  | 122     | -       |          |      |
| Patients receiving MCT/LCT on TPN | 186 (61%) | 47 (71%) | 233 (63%) | 0.2  |
| Days of artificial nutrition | 8 (4–18) | 10 (5–19) | 9 (5–8) | 0.2  |
| Days on EN             | 1 (0–1) | 9 (5–18) | 6 (1–13) | 0.001|
| Days on TPN            | 7 (3–11)| 0 (0–1) | 1 (0–7)  | 0.001|
| Starting time after ICU admission in days | 1 (0–2) | 1 (0–2) | 1 (0–2) | 0.6  |
| Prescribed caloric intake per kilogram on day 1 | 24.65 (18.77–28.57) | 23.53 (20.00–26.67) | 24 (19.3–27.64) | 0.09  |
| Administered caloric intake per kilogram on day 1 | 22.92 (17.57–27.67) | 20.8 (15.72–25) | 21.43 (16.36–26.28) | 0.01  |
| Prescribed caloric intake per kilogram on day 3 | 25 (21.25–30) | 25 (21.25–28.57) | 25 (21.25–29.36) | 0.3  |
| Administered caloric intake per kilogram on day 3 | 24.17 (20–29.45) | 22.5 (17.65–28.87) | 23.14 (18.69–27.99) | 0.003|
| Prescribed caloric intake per kilogram on day 7 | 25.84 (22.22–29.94) | 25.35 (21.43–30) | 25.66 (21.43–30) | 0.6  |
| Administered caloric intake per kilogram on day 7 | 24.72 (20–29.46) | 24.06 (19.63–28.57) | 24.31 (19.76–28.61) | 0.2  |

Parenthetical values indicate range or percentage. EN, enteral nutrition; ICU, intensive care unit; MCT/LCT, medium-chain triglyceride/long-chain triglyceride; TPN, total parenteral nutrition.
Table 4

Demographic data in patients with and without liver dysfunction

|                                | With liver dysfunction | Without liver dysfunction | Total |  \( p \)  |
|--------------------------------|------------------------|---------------------------|-------|----------|
| **Number of patients**         | 166 (23%)              | 559 (77%)                 | 725   |          |
| **Women**                      | 54 (33%)               | 183 (33%)                 | 237 (33%) | 0.9  |
| **Age in years**               | 63 (47–72)             | 63 (44–73)                | 63 (47–72) | 0.8  |
| **APACHE II score**            | 18 (14–23)             | 18 (12–22)                | 18 (12–22) | 0.2  |
| **MODS**                       | 6 (4–8)                | 5 (3–7)                   | 5 (3–7) | 0.001 |
| **Primary diagnosis**          |                        |                           |       | 0.4     |
| Gastrointestinal surgery       | 52                     | 126                       | 178   |          |
| Respiratory failure            | 26                     | 107                       | 133   |          |
| Stroke                         | 27                     | 98                        | 125   |          |
| Cardiovascular                 | 14                     | 72                        | 86    |          |
| Trauma                         | 16                     | 67                        | 83    |          |
| Infections in non-immunosuppressed patients | 12 | 28 | 40 | |
| Infections in immunosuppressed patients | 3 | 8 | 11 | |
| Metabolic diseases             | 1                      | 9                         | 10    |          |
| Urology                        | 3                      | 7                         | 10    |          |
| Hematology                     | 1                      | 8                         | 9     |          |
| Poisoning                      | 2                      | 6                         | 8     |          |
| Obstetrics/Gynecology          | 3                      | 4                         | 7     |          |
| AIDS                           | 1                      | 1                         | 2     |          |
| Other                          | 5                      | 18                        | 23    |          |
| **Type of patients**           |                        |                           |       | 0.03    |
| Medical                        | 68                     | 294                       | 362   |          |
| Surgical                       | 69                     | 195                       | 264   |          |
| Trauma                         | 29                     | 70                        | 99    |          |
| Sepsis on admission            | 68 (41%)               | 140 (25%)                 | 208 (29%) | 0.001 |
| Septic shock on admission      | 33 (20%)               | 72 (13%)                  | 105 (15%) | 0.02 |
| Patients on mechanical ventilation | 152 (92%)             | 469 (84%)                | 621 (86%) | 0.01 |
| Days of mechanical ventilation | 13 (6–24)              | 7 (3–14)                  | 8 (3–16) | 0.001 |
| Intensive care unit length of stay in days | 16 (8–28)          | 9 (5–17)                  | 10 (6–20) | 0.001 |
| Hospital length of stay in days | 28 (17–29)          | 23 (14–28)                | 25 (15–29) | 0.01 |
| Mortality at 28 days           | 47 (28.3%)             | 157 (28.1%)               | 204 (28%) | 0.9   |

Parenthetical values indicate range or percentage. APACHE II, Acute Physiology and Chronic Health Evaluation II; MODS, Multiple Organ Dysfunction Score.
appearance of any type of LD and can avoid permanent liver damage in these patients.

Another factor that could contribute to the low incidence of LD found in our group is related to the composition of the TPN. There are studies that emphasize the effect of overfeeding on the hepatic metabolism [45-47] or suggest that a lipid mixture containing MCTs (MCT/LCT) could decrease the risk of steatosis or liver cholestasis [48]. Our results do not confirm this protective effect of the MCT/LCT lipid admixture. The energy requirements of our patients were calculated at 25 kcal/kg per day. We have noted a significant difference in the administered calories in the TPN group compared with the EN group on the first and third days of follow-up, as well as a larger energy intake administered the first day of nutrition in the group of patients who developed LD. The carbohydrate/fat ratio (60:40) that we used in this study seems to be safe and can prevent the abnormalities in liver tests [49].

**Conclusion**

Our results show that the patients who developed LD have a characteristic profile in the multivariate analysis. They had a higher MODS on admission, they were septic, and they were treated with TPN. The assessment of multiple organ dysfunction includes among its parameters an LD based on high levels of bilirubin, so this association should be expected. The liver is the key organ in the starting and development of multiple organ dysfunction in the septic patient and plays an essential role in the metabolism of bilirubin.
role by clearing endotoxins, bacteria, and derived vasoactive substances. Sepsis and inflammation can increase the production of cytokines, which are potent inhibitors of bile secretion, and the consequent development of cholestasis that can be enhanced by TPN. Although the negative effects that both TPN and sepsis exert on hepatic metabolism have previously been studied independently, this study shows that there is a greater effect when both conditions, TPN and sepsis, are present. Also, early artificial nutrition seems to exert a beneficial effect. Notwithstanding prevention and treatment measures, the presence of sepsis and multiple organ failure should compel to clinicians to strictly control the caloric intake of seriously ill patients, start artificial nutrition early, and frequently monitor their liver function.

### Competing interests
B. Braun Medical S.A., Cta de Tarrasa 121, 08191 Barcelona, Spain has financially supported the data acquisition, but without access to the database or results, and will support the article-processing charge. TG is a member (vice-coordinator) of the Spanish Working Group on Metabolism and Nutrition (section of the Spanish Society of Critical Care). ABo is the

### Table 6
Incidencia de la función hepática

|                  | TPN | EN | Total | \( p \) | OR (95% CI) |
|------------------|-----|----|-------|--------|-------------|
| **Overall patients** | 303 | 422 | 725   |        |             |
| Liver dysfunction | 91  | 75 | 166   | 0.001  | 1.7 (1.3–2.2) |
| Cholestasis       | 31  | 25 | 56    | 0.03   | 1.7 (1.04–2.9) |
| Hepatic necrosis  | 28  | 20 | 48    | 0.02   | 1.95 (1.1–3.4) |
| Mixed pattern     | 56  | 43 | 99    | 0.001  | 1.8 (1.3–2.6) |

|                  | Medical | Surgical | Trauma |
|------------------|---------|----------|--------|
| Liver dysfunction| 24 (23%)| 55 (31%) | 12 (52%)|
| Cholestasis       | 6 (7%)  | 14 (16%) | 17 (23%)|
| Hepatic necrosis  | 3 (4%)  | 10 (8%)  | 28 (9%) |
| Mixed pattern     | 14 (16%)| 25 (10%) | 31 (10%)|

|                  | Medical | Surgical | Trauma |
|------------------|---------|----------|--------|
| Liver dysfunction| 24 (23%)| 55 (31%) | 12 (52%)|
| Cholestasis       | 6 (7%)  | 14 (16%) | 17 (23%)|
| Hepatic necrosis  | 3 (4%)  | 10 (8%)  | 28 (9%) |
| Mixed pattern     | 14 (16%)| 25 (10%) | 31 (10%)|

CI, confidence interval; EN, enteral nutrition; OR, odds ratio; TPN, total parenteral nutrition.

### Figure 1

![Time free of liver dysfunction](image)

Time free of liver dysfunction in surgical patients treated with Enteral Nutrition or Total Parenteral Nutrition. EN, enteral nutrition; TPN, total parenteral nutrition; AN days, days on artificial nutrition.
coordinator of the Spanish Working Group on Metabolism and Nutrition (section of the Spanish Society of Critical Care). The other authors declare that they have no competing interests.

Authors’ contributions
TG and ABo conceived the study, participated in its design and coordination, and helped to draft the manuscript. TG performed the statistical analysis. MR and DM were involved in drafting the manuscript or revising it critically for important intellectual content. ABI, MF, JAA, and JCM participated in the design of the study and they coordinated the meetings with the participants. AG and AM have given final approval of the version to be published. All authors read and approved the final manuscript.

Acknowledgements
The following members of the Working Group on Nutrition and Metabolism of the Spanish Society of Critical Care participated in the study: Zabarte M (Hospital Na Sra de Aranzazu, San Sebastián), Bonet Sarís A., Sirvent Calvera JM (Hospitral Joseph Trueta, Gerona) Farré Vladrich M, Salvador Salvat J (Hospital Universitari de La Vall D’Hebron, Barcelona), Acosta Escribano JA (Hospital Universitario de Alicante, Alicante), Blesa Malpica A (Hospital Clinico San Carlos, Madrid), Montejo González JC (Med-Surg ICU, Hospital 12 De Octubre, Madrid), Jiménez Jiménez J, Ortiz Leyba C (Hospital Virgen Del Rocío, Sevilla), Cuñat J, Arguedas J (Hospital Universitari de la Fe, Valencia), Abella A, Blanco J (Hospital Universitario de Getafe, Madrid), Sanchez-Izquierdo R JA (Trauma ICU, Hospital 12 de Octubre, Madrid), Iturralde Yánez J (Hospital de Navarra), Ruiz Santana S, Peña Morant V (Hospital Universitario Dr Negrín, Las Palmas de Gran Canaria), Morán García V (Hospital de León, León), Albert Bonamusa I (Hospital Universitario de Alicante, Alicante), Garcia de Lorenzo y Mateos A (Hospital Virgen De La Candelaria, Tenerife), Martínez García P (Hospital Universitario De Puerto Real, Cadiz), Palacios Rubio V (Hospital Miguel Server, Zaragoza), Martin Garcia P (Hospital Universitario De San Juan, Alicante), López Martínez J (Hospital Severo Ochoa, Madrid), Rodríguez A, Serviá L (Hospital Universitari Arnau De Vilanova, Lleida), Tejada Artigas A (Trauma ICU, Hospital Miguel Server, Zaragoza), Jara Clemente F (Hospital Mutua de Terrassa), De La Fuente O’Connor E (Hospital Princeps de Asturias, Madrid), Masdeu Eixarch G (Hospital Verge De La Cinta, Tortosa), Fernandez Ortega JF (Hospital Universitario Carlos Haya, Málaga), Casanovas Taltavull M (Hospital Universitari Arnau De Vilanova, Lleida), Tejada Artigas A (Trauma ICU, Hospital Miguel Server, Zaragoza), Jara Clemente F (Hospital Mutua de Terrassa), De La Fuente O’Connor E (Hospital Princeps de Asturias, Madrid), Masdeu Eixarch G (Hospital Verge De La Cinta, Tortosa), Fernandez Ortega JF (Hospital Universitario Carlos Haya, Málaga), Casanovas Taltavull M (Hospital General de Segovia, Segovia), Ortellis Huerta X (Hospital Marina Alta, Alicante), Herrera Morillas F
References

1. ASPEN Board of Directors and the Clinical Guidelines Task Force: Guidelines for the use of parenteral and enteral nutrition in adult and pediatric patients. JPEN J Parenter Enteral Nutr 2002, 26(1 Suppl):1S-138S5A.

2. Cerra FB: Hypermetabolism, organ failure and metabolic support. Surgery 1987, 101:1-14.

3. Shattuck KE, Klein GL: Hepatobiliary complications of parenteral nutrition. In Enteral and Tube Feeding 3rd edition. Edited by: Rombeau JL, Rolandiell RH. Philadelphia: WB Saunders; 1997:141-156.

4. Marshall JC, Cook DJ, Christou NV, Bernard GR, Sprung CL, Sibbald WJ: Multitele sepsis dysfunction score: a reliable descriptor of a complex clinical outcome. Crit Care Med 1995, 23:1638-1652.

5. Knaus WA, Draper EA, Zimmerman JE: APACHE II: a severity of disease classification system. Crit Care Med 1985, 13:818-829.

6. American College of Chest Physicians/Society of Critical Care Medicine Consensus Conference: Definitions for sepsis and multiple organ failure and guidelines for the use of innovative therapies in sepsis. Crit Care Med 1992, 20:864-874.

7. Buzby GP, Kjos LT, Spoutz DA, Olesen N, Haakenson CM, McNeal GE, Page CP, Peterson OL, Reinhardt GF, Willford WO: Study protocol: a randomized clinical trial of total parenteral nutrition in malnourished surgical patients. Am J Clin Nutr 1984, 40(2):364-381.

8. American Society for Parenteral and Enteral Nutrition: Clinical Pathways and Algorithms for Delivery of Parenteral and Enteral Nutrition Support in Adults Silver Spring, MD: American Society for Parenteral and Enteral Nutrition; 1998.

9. Montijo JC: Enteral nutrition-related gastrointestinal complications in critically ill patients: a multicenter study. The Nutritional and Metabolic Working Group of the Spanish Society of Intensive Care Medicine and Coronary Units. Crit Care Med 1999, 27:1447-1453.

10. Deterre JP, Deterys AS, Wesson DE, Wolman SL, Stewart S, Whitney J, Langer B, Jessebeyoh KN: Nutritional assessment: a comparison of clinical judgement and objective measurements. N Engl J Med 1982, 306:969-972.

11. Bonet A, Sanchez Alvarez C, Nuñez Ruiz R: Protocolo de nutricion parenteral. In Guías de Práctica Clínica en Medicina Intensiva. [Total parenteral nutrition protocol. Clinical Practice Guidelines in Critical Care] Edited by: Latorre FJ, Ibáñez J. Madrid: Medetex; 1996:1-7.

12. Quiqley EM, Marsh MN, Shaffer JL, Markin RS: Hepatobiliary complications of total parenteral nutrition. Gastroenterology 1993, 104:286-301.

13. Briones ER, Iber FL: Liver and biliary tract changes and injury associated with total parenteral nutrition: pathogenesis and prevention. J Am Coll Nutr 1995, 14:219-228.

14. Lowey SJ, Lowry SF: Parenteral nutrition and liver dysfunction – new insight? JPEN J Parenter Enteral Nutr 1991, 15:54-59.

15. Braxton C, Lowry SF: Parenteral nutrition and liver dysfunction – new insight? JPEN J Parenter Enteral Nutr 1995, 19:3-4.

16. de Pablo MA, Angeles Puertollano M, Álvarez de Cienfuegos G: Immune cell functions, lips and host natural resistance. FEMS Immunol Med Microbiol 2000, 29:323-328.

17. Buzby GP, Kjos LT, Spoutz DA, Olesen N, Haakenson CM, McNeal GE, Page CP, Peterson OL, Reinhardt GF, Willford WO: Study protocol: a randomized clinical trial of total parenteral nutrition in malnourished surgical patients. Am J Clin Nutr 1984, 40(2):364-381.

18. Keim NL: Nutritional effectors of hepatic steatosis induced by parenteral nutrition in the rat. JPEN J Parenter Enteral Nutr 1997, 21:18-22.

19. Sheldon GF, Petersen SR, Sanders R: Hepatic dysfunction during hyperalimentation. Arch Surg 1978, 113:504-508.

20. Buzby G, Mullen JL, Stein TP, Rosato EF: Manipulation of TPN caloric substrate and fatty infiltration of the liver. J Surg Res 1981, 31:46-54.

21. Burke JF, Wolfe RR, Mullany CJ, Mathews BE, Bier DM: Glucose requirements following burn injury. Parameters of optimal glucose infusion and possible hepatic and respiratory abnormalities following excessive glucose intake. Ann Surg 1979, 180:274-285.

22. Thompson SW: Hepatic toxicity of intravenous fat emulsions. In Fat Emulsions in Parenteral Nutrition Edited by: Meng HC, Willmore DW. Chicago: American Medical Association; 1976:90-95.

23. Preiss R, Rentrop B: Biliary transport and cholestatic effects of amino acids. Gastroenterology 1977, 73:1240-1246.

24. Burt ME, Hanin I, Brennan MF: Choline deficiency associated with total parenteral nutrition. Lancet 1980, 2:638-639.

25. Fourni-Fontenet H, Le Quernec L, Erlinger S, Lerebours E, Colan R: Hepatic alterations during total parenteral nutrition in patients with inflammatory bowel disease: a possible consequence of lathiocholate toxicity. Gastroenterology 1982, 82:932-937.

26. Penn D, Schmidt-Sommerfeld E, Pascu F: Decreased tissue carnitine concentrations in newborn infants receiving total enteral nutrition. J Pediatr 1981, 98:976-978.

27. Zamir O, Nussbaum MS, Bhadra S, Subbiah MT, Rafferty JF, Fischer JE: Effect of enteral feeding on hepatic steatosis induced by total parenteral nutrition. JPEN J Parenter Enteral Nutr 1994, 18:20-25.

28. Pallarès R, Sitgues-Serra A, Fuentes J: Cholestasis associated with total parenteral nutrition. Lancet 1983, 1:758-762.

29. Pacelli F, Bossola M, Papa V, Malerba M, Modesti C, Sgardari A, Bellantone R, Doglietto GB, Modesti C, EN-TPN Study Group: Enteral vs parenteral nutrition in elderly patients undergoing major surgical procedures: is it even match. Arch Surg 2001, 136:933-938.

30. Woodcock NP, Zeiger D, Palmer DM, Buckley P, Mitchell CJ, MacFie J: Enteral versus parenteral nutrition: a pragmatic study. Nutrition 2001, 17:1-12.

31. Borum ML, Lynn J, Zhong Z, Roth K, Connors AF Jr, Desbois NA, Phillips RS, Dawson NV: The effect of nutritional supplementation on survival in seriously ill hospitalized adults: an evaluation of the SUPPORT data. Study to Understand Prognoses and Preferences for Outcomes and Risks of Treatments. J Am Coll Nutr 2003.

32. Braga M, Gianotti L, Gentilini O, Parisi V, Salis C, Di Carlo V: Early post-operative enteral nutrition improves gut oxygenation and reduces costs compared with total parenteral nutrition. Crit Care Med 2001, 29:242-248.

33. MacRae PE, Pinsky M: Death by parenteral nutrition. Intensive Care Med 2003, 29(6):867-869.

34. Heyland DK, MacDonald S, Keefe L, Drover JW: Total parenteral nutrition in the critically ill patient: a meta-analysis. JAMA 1998, 280:2013-2019.

35. Hidaka K, Ikeda S, Homma T, Mitaka T, Furuhata T, Katsuramaki T, Hata F, Makiya M: Sepsis and cholestasis: basic findings in the sinuous and bile canaliculi. J Hepatobiliary Pancreat Surg 2001, 8:20-26.

36. Pallarès R, Sitgues-Serra A, Fuentes J, Laurieta E, Guardia J, Fernández-Nogués F, Sitges-Serra A: Factori etiopatogénicos posiblemente implicados en la disfunción hepática asociada a la nutrición parenteral: estudio prospectivo de 104 pacientes adultos. [Etiopathogenic factors possibly implicated in hepatic dysfunction associated with parenteral nutrition: prospective study of 104 adult patients]. Med Clin (Barc) 1984, 83:832-836.

37. Buchman A: Total parenteral nutrition-associated liver disease. JPEN J Parenter Enteral Nutr 2002, 26:543-548.

38. Bookhu IS, Jarvis C, Everson GT: Total parenteral nutrition and cholestasis. Clin Liver Dis 1999, 3:489-508.

39. Cavicchi M, Amato D, Crenn P, Degott C, Messing B: Prevalence of liver disease and contributing factors in patients receiving home parenteral nutrition for permanent intestinal failure. Ann Intern Med 2000, 132:526-532.

40. Spilsbod JD, Kalfaretos P: Total parenteral nutrition-associated liver dysfunction. Nutrition 1994, 10:255-260.

41. Angelico M, Della Guardia P: Review article: hepatobiliary complications associated with total parenteral nutrition. Aliment Pharmacol Ther 2000, 14(suppl 1):S4-S7.

42. Morán Penco JM, Sánchez-Martinez J, Maciá Botejara J: ¿Qué sucede en el hígado durante la alimentación artificial? [What happens to the liver during artificial feeding?]
happens with the liver during artificial feeding]. Nutr Hosp 2001, 16:145-151.

44. Sax HC, Talamini MA, Brackett K, Fisher JE: Hepatic steatosis in total parenteral nutrition: failure of fatty infiltration to correlate with abnormal serum hepatic enzyme levels. Surgery 1986, 100:697-704.

45. Pallarés R, Sancho S, Sitges-Serra A, Jaurrieta E, Cañadas E, Guardia J, Fernández-Nogués F, Sitges-Creus A: Estudio clínico-morfológico de la disfunción hepática asociada a la nutrición parenteral en adultos: a propósito de 15 casos. [Clinico-morphologic study of hepatic dysfunction associated with parenteral nutrition in adults: apropos of 15 cases]. Med Clin (Barc) 1984, 83:837-841.

46. Grant JP, Cox CE, Kleinman LM, Maher MM, Pittman MA, Tangrea JA, Brown JH, Gross E, Beazley RM, Jones RS: Serum hepatic enzyme and bilirubin elevations during parenteral nutrition. Surg Gynecol Obstet 1977, 145:573-580.

47. Buchmiller CE, Kleiman-Wexler RL, Ephgrave KS, Booth B, Hensley CE 2nd: Liver dysfunction and energy source: results of a randomized clinical trial. JPNEN J Parenter Enteral Nutr 1993, 17:301-306.

48. Chandra S, Mehendale HM: Nutritional modulation of the final outcome of hepatotoxic injury by energy substrates: a hypothesis for the mechanism. Med Hypotheses 1996, 46:261-268.

49. Carpentier YA, Dubois DY, Siderova VS, Richelle M: Exogenous lipids and hepatic function. In Organ Metabolism and Nutrition: Ideas for Future Critical Care Edited by: Kinney JM, Tucker HN. New York: Raven Press, Ltd; 1994:349-367.