Sir,

Increasing antibiotic resistance in bacteria is a cause for concern in the treatment of infections, particularly so in hematopoietic stem cell transplant (HSCT) patients who have a greater propensity toward acquiring infections because of the underlying immunosuppression. The patients’ endogenous flora from the bowel, mouth and skin may be responsible for most of the serious bacterial infections that may occur and, although the debatable practice of routine prophylaxis is followed in many settings, the emerging issue of increasing antibiotic resistance can influence this. Surveillance cultures from common organism-harboring sites can help identify resident multidrug-resistant endogenous bacteria.

We conducted a study to determine the prevalence of multidrug-resistance bacteria in surveillance cultures in our HSCT patients. Ninety-nine patients who were to undergo HSCT at our oncology center for the treatment of various malignant conditions during the period 2009–2010 were included. Surveillance cultures were performed from these patients when they were admitted. Swabs were collected from the anterior nares and axilla to look for the presence of methicillin-resistant *Staphylococcus aureus* (MRSA). Feces cultures were performed to screen for the presence of organisms like multidrug-resistant Gram negative bacilli including extended-spectrum beta lactamase-producing organisms (ESBL) and other carbapenem-resistant organisms. They were also screened for the presence of vancomycin-resistant Enterococci (VRE). Swabs were also collected from insertion and exit sites of Hickman’s catheters from these patients. All samples were processed according to the standard microbiology laboratory protocol and antibiotic selection and susceptibility interpretation was as per the Clinical Laboratory Standards Research Institute (CLSI) guidelines.

A total of 14 *S. aureus* isolates were recovered from the nasal and axillary swabs, of which seven (50%) were found to be methicillin resistant, with an overall MRSA incidence of 7%. Insertion and exit site swabs from Hickman catheters grew coagulase-negative Staphylococci in 4% and MRSA in 1% of the cases. In addition, *Pseudomonas aeruginosa*, *Pseudomonas spp* and *Acinetobacter spp* were recovered in 6% of the swabs. Of the 98 fecal cultures, a total of 68 grew *Escherichia coli*, of which 76.5% were resistant to the third-generation cephalosporins ceftazidime and cefotaxime; 36.8% were resistant to cefoperazone–sulfactam and 8.2% were resistant to imipenem and meropenem. Twenty fecal cultures grew *Klebsiella pneumoniae*, of which 60% were resistant to cefotaxime and ceftazidime; 35% were resistant to cefoperazone–sulfactam and 5% were resistant to the carbapenems tested. Forty-five percent of the *Klebsiella pneumoniae* and 61.8% of the *E. coli* strains were ESBL producers. Fifteen strains of *Pseudomonas aeruginosa/spp* were isolated, which showed a higher resistance to piperacillin–tazobactam (86.6%), ceftazidime (80%) and cefoperazone–sulfactam (80%), and also to the carbapenems (73.3%). Twenty percent of the Enterococci were vancomycin resistant.

Thus, a high level of antibiotic resistance in the endogenous organisms of our HSCT patients was demonstrated by surveillance cultures, which is evident from isolations of MRSA, VREs and ESBL-producing and carbapenem-resistant Gram negative bacteria. Patients who are colonized with multidrug-resistant bacteria may serve as a source of later infection in the same patient and also as infection risk for neighboring patients if not isolated.
Surveillance cultures may be helpful in identifying these multidrug-resistant bacteria, which may have a bearing on further patient management.

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Sir,

Omega-3 polyunsaturated fatty acid (n-3 PUFA) and branch chain amino acid (BCAA) supplementation is a presently widely used nutritional supplementation to help promote immunity. There is no doubt that this immunonutrition is helpful for the patients. However, its effect on biochemistry is not often mentioned. Here, the authors present a case of breast cancer with liver, lung and lymph node metastasis. The patient is a 55-year-old female patient, and is consulted for immunonutritional management. At the starting point, the laboratory parameters for this case are as the following: Insulin-like growth factor (IGF)-1=166; Alpha-fetoprotein (AFP)=3.80; Carcinoembryonic antigen (CEA)=8.63; and CA15-3=25.5. After 1 month of supplementation of n-3 PUFA and BCAA once a week by intravenous infusion, the laboratory parameters for this case are as the follows: IGF-1=113; AFP=2.87; CEA=6.99; and CA15-3=21.43. A reduction of the tumor markers can be seen. Indeed, the fluctuation of the tumor markers in cancerous patient is an important concern in cancer therapy. Some conditions such as heart failure are confirmed in induction of fluctuation.[1] However, the effect of the immunonutrition is limited. Theoretically, omega supplementation can induce apoptosis of cancerous cells, and this can affect the level of tumor markers.[2] In this letter, an example case of reduced tumor markers in cancerous patients receiving omega supplementation is presented. This might reflect the usefulness of the supplementation.[3] However, this does not mean that supplementation can destroy the cancer cell. The decreased level might be a fluctuation that is only a laboratory observation. Further studies to clarify this topic are recommended.[3]

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