

abstract

Recovery Kinetics of Serum Albumin in Pediatric Cancer Patients: It's Determinants and Correlation with Outcomes

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Recovery Kinetics of Serum Albumin in Pediatric Cancer Patients: It's Determinants and Correlation with Outcomes

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Introduction: Hypoalbuminemia and cancer-cachexia predict poor survival and chemotherapy-toxicity in adult-cancers, with limited literature in pediatrics. Also, the recovery-kinetics of serum-albumin and its effect on cancer-outcomes is unknown. This study investigated trends and determinants of serum-albumin at various time-points and their influence on 3-year-survival of pediatric-cancers.

Methodology: The study included cancer patients ≤ 18 yrs and excluded those with liver/kidney diseases and relapsed-malignancies. Baseline systemic symptoms, stage/risk-group, nutritional status, serum-albumin at baseline, 3-months, 6-months and 12-months of therapy were recorded along with outcomes like relapse/death and toxicity. Analysis was done to identify the determinants and impact of serial albumin values.

Results: A total of 402 patients were enrolled. Median age was 4yrs(2.5mo-14yrs), majority were hemato-lymphoid cancers(57.71%), with 50%, 19% and 31% belonging to high-risk, intermediate-risk and low/standard-risk respectively. While 59.20% had systemic-symptoms, 46.97% had malnutrition. The optimum cut-off albumin that predicted survival was 4.0g/dl. Prevalence of hypoalbuminemia at baseline, 3m and 6m were 46.52%, 26.45% and 22.0% respectively. The 3-year overall-survival(OS) was lower for patients with hypoalbuminemia at baseline(44.9% vs 69.8%; HR=2.1, P=0.00), 3m(HR=2.28; P=0.00) or 6m (HR=2.02; P=0.001) in univariate-analysis.

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On multivariate-analysis, only risk- group ($HR=1.74; P=0.01$), 6m-albumin ($HR=3.33; P=0.00$) and first 6m albumin-trend ($HR=1.84; P=0.02$) were independent predictors of OS, but not baseline-albumin($HR=1.23; P=0.57$) or 3m albumin($HR=1.18; P=0.63$).

Only risk-group ($HR=1.59; P=0.02$) and 6m-albumin ($HR=2.91; P=0.00$) were significant predictors for event-free-survival(EFS). Liver/spleen size($r=-0.39$) and serum-globulin($r=-0.37$) had moderate negative correlation with baseline- albumin. On multivariate-analysis, systemic-symptoms($OR=8.01, P=0.01$) and baseline-malnutrition ($OR=2.08; P=0.00$) were independently associated with baseline-hypoalbuminemia. In contrast, 6-months albumin was dependent only on baseline-systemic-symptoms($OR=5.28; P=0.01$), but independent of baseline-malnutrition($OR=1.72; P=0.09$). Also, hypoalbuminemia at baseline, 3m or 6m was associated with increased risk of febrile neutropenia($OR=1.29; P=0.01$) and hospitalization($OR=2.86; P=0.00$).

Conclusion: Baseline risk-group and 6m-albumin are independent predictors of OS/EFS, rather than baseline-albumin. While baseline-albumin is influenced by malnutrition, systemic-symptoms, inflammatory state(hypergammaglobulinemia) and disease-bulk(organomegaly), the 6-months-albumin is independent of baseline-malnutrition and likely a better indicator of the cancer-inflammation-cachexia-cascade to predict disease-outcomes in pediatric-cancers.