Methods: Consecutive patients undergoing stem cell transplant from November 2009 to May 2010 were assessed using QLQ-C30 (self assessment questionnaire) on admission to the transplant ward just before undergoing stem cell transplant. Written consent was obtained. Demographic data as well as disease details and status of patients are collected at assessment. Another assessment of the QLQ-C30 was done after the transplant (from D+30 until D+180). The QLQ-C30 was scored according to the global health, functional level and symptomatology.

Results: A total of 29 patients answered the survey.

Global health status and functional abilities (physical, role, cognitive, emotional, social) of our patients seemed to be lower than the general population. Our patients seemed to have similar global health, physical and emotional status but lower role function and also cognitive function compared to the "all cancer" reference population. Global Health Status was similar in pre and post allogenic transplant patients with only autologous transplant patients appeared to have better scores post transplant. In the early post transplant period, most of the allogenic transplant patients had worse scores while most of the auto pts have better scores.

Conclusion: Haematology patients who are planned to undergo transplant have similar QOL as other cancer patients. Allogenic stem cell transplant patients have a worse QOL in the early post transplant period. Autologous transplant patients seemed to have better QOL in the early post transplant period.

	reyed (Pre transplant) = 29
Auto	13, Allo 15, Cord 1
Gend	ler = 20 female, 9 male
Mari	tal status = 13 married, 16 single
Child	ren = 7 have children, 22 no children
Race	= 12 Chinese, 16 Malay, 1 Indian
ECOG per	formance status
0 = 4	patients
1 = 2	3 patients
2 = 1	patient
3 = 1	patient
Median A	ge = 28.3 years
ducation	Level = 16 Secondary school, 13 tertiery education

Diagnosis	***************************************
Diagnosis Aplas	
Diagnosis Aplas Acute	stic Anaemia = 2
Diagnosis Aplas Acute Acute	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6
Diagnosis Aplas Acute Acute Acute	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8
Diagnosis Aplas Acute Acute Acute Chro	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1
Acute Acute Acute Acute Chro	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2
Acute Acute Acute Acute Chro Diffu	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2 se Large B Cell Lymphoma = 1
Acute Acute Acute Chro Diffu Follic	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2 se Large B Cell Lymphoma = 1 ular Lymphoma = 1
Diagnosiss Aplas Acute Acute Acute Chro Diffu Follic Hodg	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2 se Large B Cell Lymphoma = 1 ular Lymphoma = 1 kin Lymphoma = 5
Diagnosis Aplas Acute Acute Chro Diffu Follic Hodg Myel	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2 se Large B Cell Lymphoma = 1 ular Lymphoma = 1 kin Lymphoma = 5 odysplastic Syndrome = 1
Diagnosis Aplas Acute Acute Chro Diffu Follic Hodg Myel Mult	stic Anaemia = 2 e Lymphoblastic Leukaemia = 6 e Myeloid Leukaemia = 8 e Promyelocytic Leukaemia = 1 nic Myeloid Leukaemia = 2 se Large B Cell Lymphoma = 1 ular Lymphoma = 1 kin Lymphoma = 5 odysplastic Syndrome = 1 iple Myeloma = 2

(Median scores)	Global health Status	Functional Scale – Physical Function	Functional Scale – Role Function	Functional Scale – Emotional Function	Functional Scale – Cognitive Function	Functional Scale – Social Function
Normal population	75	100	100	83.3	100	100
All cancer	66.7	80	80	75	83.3	83.3
All pre transplant(29)	66.7	80	66.7	66.7	66.7	66.7
All post transplant(13)	66.7	80	83.3	83.3	66.7	66.7

P715

Stem cell transplantation: a 10-year single.centre experience

B. Georgievski, Z. Stojanoski, S. Genadieva-Stavrik, A. Pivkova, M. Ivanovski, O. Karanfilski, L. Cevreska, S. Krstevska-Balkanov, S. Trajkova

Bone Marrow Transplantation Unit (Skopje, MK)

Stem cell transplantation is the treatment of choice for patients with hematological malignant diseases. Aim of this study is to evaluate the results in 10 years experience with this procedure in our center. During a this period we have treated 195 patients with different malignant hematological diseases.107 male, 88 female with median age of 34 years. Allogeneic sibling transplantation were performed in 56 patients, and autologous transplantation in 139 patients. According to diagnose: AML:91 ALL:10 CML:7 CLL:1 MP:1 NHL:20 HD:27 MM:34 AA:2 Ewing sarcoma: 1. Bone marrow was used as a source of stem cells in 28, PBSC in 168 patients. Conditioning regimen consisted chemotherapy with: Bu-Cy2,Bu-Cy-Mel, BEAM, high dose Melphalan, high dose ICE, Flu/Mel. Engraftment was reached on day +12 (10-24). Median value of MNC in BMT was 3,8x10/Kg (2,5-4,5), while in PBSCT MNC was 4,1x10 (2,8-12,0). TRM in allogeneic recipients was 16%, with non-relapse mortality 10%, and in autologous recipients TRM was 3,8% with non-relapse mortality 2,0%. Primary disease was cause of death after transplantation in 40% in allogeneic, and 66% in autologous transplantation. 35% of allogeneic and 28% of autologous transplantation were transplanted in active disease.

P716

First Russian experience of calcium phosphate mouth rinse usage for treatment of children with oral mucositis undergoing haematopoietic stem cell transplantation

E.V. Skorobogatova (1), Z.M. Dyshlevaia (1), A.I. Karachunsky (2), E.D. Pashanov (2)

(1)Central Children Hospital (Moscow, RU); (2)Federal Centre for Pediatric Hematology, Oncology, and Immunology (Moscow, RU)

Objective: High-dose chemotherapy administered as conditioning regimen prior to hematopoietic stem cell transplantation (HSCT) leads to injury or disruption of oral and gastro-intestinal mucosa. Mucositis is associated with increased risk of infections, mortality of patients (pts), duration and doses of morphine, duration of hospitalization, higher costs of treatment. As previously described, Caphosol, a neutral, super-saturated, calcium and phosphate solution for mouth rinse, is effective and safe for prevention of oral mucositis (OM) in adults undergoing HSCT and in head and neck cancer pts receiving chemo- and radiotherapy. However, the effect of Caphosol on OM in children is not yet established. We decided to first evaluate efficacy, safety and tolerability of Caphosol added to standard OM management in children with OM after HSCT.

Methods: Five children (4-16 years; median 13) with AML (n=2; allo-HSCT), neuroblastoma (n=2; auto-HSCT), Ewing sarcoma (n=1; auto-HSCT) were included. All pts received high-dose chemotherapy before HSCT and had OM grade 1 (n=4) or 4 (n=1) before Caphosol administration. Pts used Caphosol rinse four times daily, 30 ml each time. The OM was assessed according to Oral Mucositis Assessement Scale (OMAP) published by Sonis et al. in 1999.

Results: We detected the regression of OM in all pts (8-15 days; median 12). The duration of morphine administration was 0 (refusal by the patient), 8,10,12, and 15 days, with progressive decrease of the dose. Four of 5 pts had pain decrease during first hours after first Caphosol administration. The fever with no positive microbiology tests has been developed in 3/5 pts with duration of 3, 13, and 15 days. No adverse events nor bad taste or other unpleasant sensations on behalf of pts have been observed.