



# Papillary thyroid cancer after hematopoietic stem cell transplantation in young age



<sup>1</sup>M. Snajderova, <sup>2</sup>P. Keslova, <sup>3</sup>P. Sykorova, <sup>2</sup>P. Sedlacek, <sup>2</sup>R. Formankova and <sup>2</sup>J. Stary  
<sup>1</sup>Department of Paediatrics, <sup>2</sup>Department of Paediatric Haematology and Oncology, <sup>3</sup>Department of Nuclear Medicine and Endocrinology, University Hospital Motol, Prague, Czech Republic

Nothing to disclose

## Introduction

- Increasing number of survivors following hematopoietic stem cell transplantation (HSCT) leads to necessity to focus also on careful monitoring for late effects. High dose chemotherapy and total body irradiation (TBI) is used for conditioning regimen in many patients.
- Thyreopathies** belong to the most frequent among late endocrinopathies.

## Objective

Aim of the study was to evaluate **prevalence of secondary thyroid malignancies after HSCT in young age, especially after TBI.**

## Subjects

**HSCT performed 1989 – 2012**

### Inclusion criteria

- HSCT in childhood or in adolescence
- Patients surviving more than 1 year after HSCT

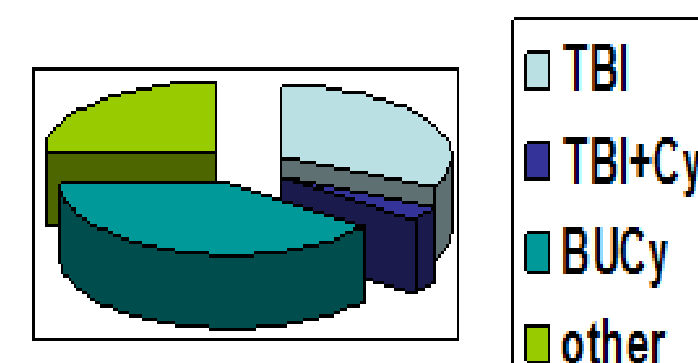
### Characteristics of patients

Patients	n	288
Female/Male	n	110/178
Malignant diagnosis	n	177 (61.5%)
TBI 10-14 Gy	n	91 (32%)
Age at HSCT Median (range)	years	8.2 (0.2 – 20.5)
Age at evaluation Median (range)	years	17.9 (1.8 – 40.5)
Time after HSCT Median (range)	years	8.5 (1.3 – 24.3)

## Transplant characteristics of study group

Preparative regimen

- TBI : 92
- TBI+Busulphan : 10
- Bu/Cy : 114
- Other : 72



Bu: busulphan Cy: cyclophosphamide

## Methods

- All patients were screened yearly for:** Serum fT4, TSH, thyroid antibodies, thyroid function
- thyroid ultrasound (US) was performed in 62/288 (21.5%) patients with a Toshiba Nemio 17 ultrasound machine, transducer 7.5 MHz

## Results

### Malignant tumours

**4 patients (1.4%)** - 2 male, 2 female  
**all with papillary thyroid carcinoma (PTC) micronodular, T1 or T2 stage**  
**all of these patients survive after PTC, disease free**

PTC diagnosed after HSCT	8.7 years (5.3-15.2)
HSCT for malignant diagnosis	n = 3
TBI-based regimen	all 4 patients
Therapy of PTC	thyroidectomy and replacement thyroid hormone in all 1 with additional <sup>131</sup> I treatment
Previous therapy for thyreopathy	n=3 AITD n=1 hypothyroidism

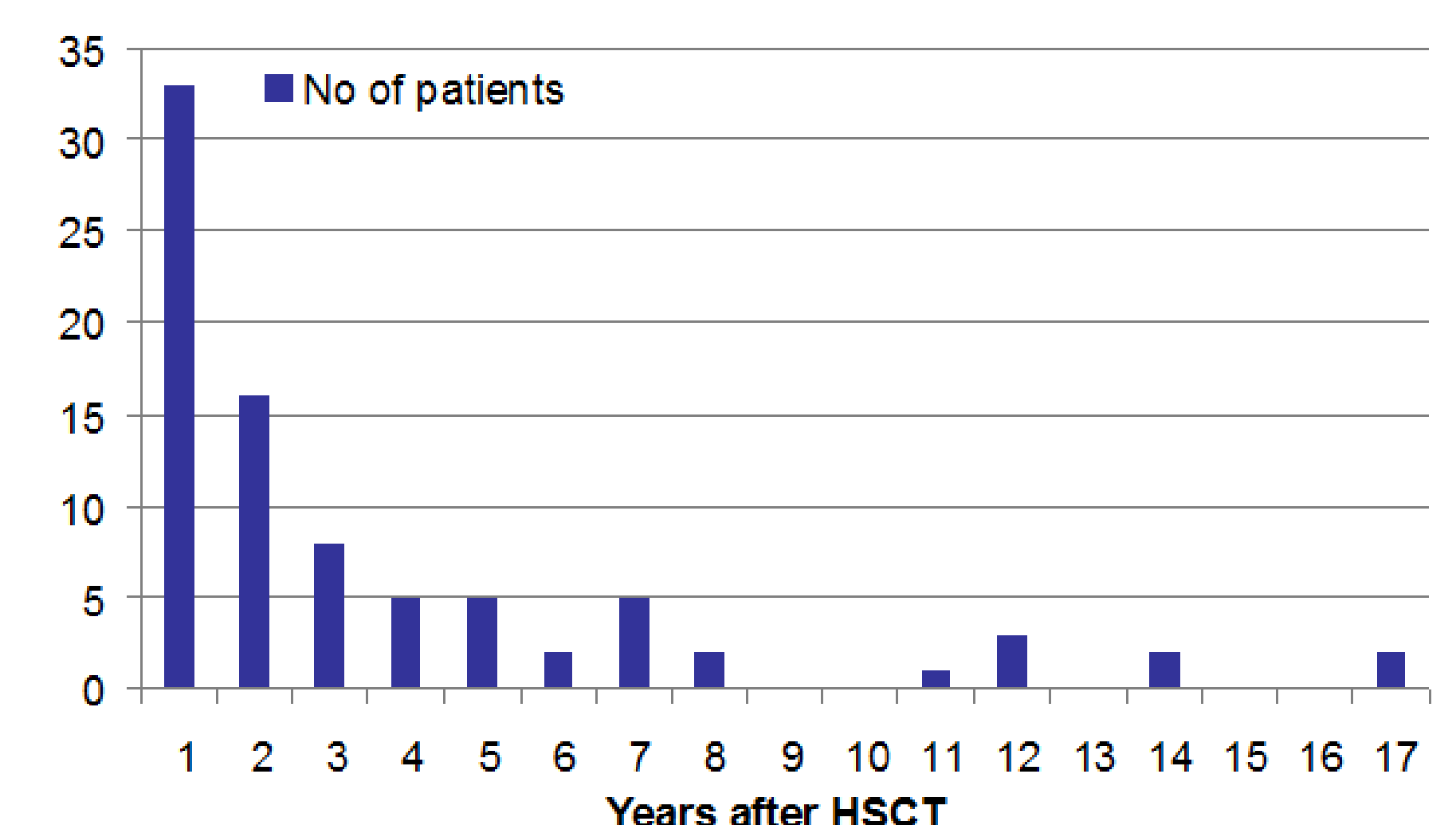
## Benign thyroid nodules (FNAB confirmed)

**9 patients (3.1%)**  
 7 after TBI-based regimen  
 4 treated for AITD  
 all are regularly monitored

## Thyroid dysfunction (TDF) after HSCT

Patients with TDF	n (%)	83/288 (29.0%)
Onset of TDF after HCST	years	3.4 (0.5-16.3)
Hypothyroidism subclinical	n (%)	77 (28.5%)
overt primary		5
Hyperthyroidism	n (%)	2 (0.7%)
AITD (autoimmune thyroiditis)	n (%)	18 (6.2%)

## Onset of thyroid dysfunction after HSCT



## Conclusions

- Risk of secondary malignancies after HSCT is increasing within the time.**
- Long-life late effects monitoring as an important part of post-transplant care is necessary.**
- Regular sonographic evaluation of thyroid gland and neck is very important especially more than 5 years after HSCT and namely in all patients after TBI.**
- Regular monitoring of thyroid function, laboratory parameters and ultrasound is highly recommended.**