



# Diagnostic and therapeutic approach to children with Nijmegen breakage syndrome in relation to development of lymphoid malignancies

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## Abstract

**Introduction and objective.** Nijmegen breakage syndrome (NBS) is a rare chromosomal instability disorder. The majority of patients carry founder mutation in the *NBN* gene (c.657\_661del5). Characteristic features of the NBS include progressive microcephaly, dysmorphic facial features, immunodeficiency, and high predisposition to malignancy with cumulative cancer incidence by the age of 20 years, and amounted to over 70%.

The aim of study is to present the latest methods of diagnosis, potential cancer risk factors and treatment of lymphoid malignancies in children with NBS.

**Review methods.** To review the evidence using PubMed and Google Scholar search which included articles published between 2009–2021, focusing on articles published between 2013–2021.

**Abbreviated description of the state of knowledge.** The average delay in diagnosis of NBS ranges from 4–5 years. Neonatal screening of T-cell excision circles (TREC) and kappa-deleting recombination excision circles (KREC) seems favourable in NBS. There are no specific protocols for the treatment of lymphoid malignancies in children with NBS, and full-dose chemotherapy is the most frequently applied method. Reducing the doses of chemotherapy does not significantly reduce the toxicity. Main cause of death is cancer progression and treatment-related mortality mostly associated with infectious complications. Patients with diagnosed cancer who received haematopoietic stem cell transplantation (HSCT) had significantly higher 20-year OS than those who did not (42.7% vs. 30.3%).

**Summary.** Further meta-analysis is essential to establish the best monitoring and treatment regimen in patients with NBS and lymphoid malignancies.

## Key words

diagnosis, lymphoma, antineoplastic agents, Nijmegen Breakage Syndrome, leukemia, Haematopoietic Stem Cell Transplantation

## INTRODUCTION

Nijmegen breakage syndrome (NBS) is a rare chromosomal instability disorder. The gene responsible for the disease, *NBN*, is located on chromosome 8q21. The majority of patients are diagnosed with five base pair deletion in *NBN* gene (c.657\_661del5) which is a founder mutation in Slavic populations [1]. NBS seems to occur worldwide, but most patients originated from Poland and Russia [2, 3].

The *NBN* gene codes for nibrin which is involved in DNA double-strand break repair [4]. Due to mutation, V(D)J rearrangement of T cells is affected and circulating T cells have signs of senescence in the young patients with NBS [5, 6].

The characteristic clinical features of the NBS include progressive microcephaly, dysmorphic facial features

(including sloping forehead, prominent nose, long mandible, long philtrum), mild growth delay and premature ovarian insufficiency. Profound immunodeficiency both cellular and humoral response develops among most patients [4]. Patients with NBS have cumulative cancer incidence by the age of 20 years and amounted to over 70% [2, 3]. Non-Hodgkin lymphomas (NHLs) are most common with a similar number of lymphomas developed from abnormal T-cells and B-cells. T-cell acute lymphoblastic leukemia (T-ALL) is most frequently diagnosed among leukemias [2, 3, 7, 8, 9].

## OBJECTIVE

The aim of the study is to present the current state of knowledge on NBS and lymphoid malignancies with a focus on the peculiarities of diagnosis, cancer risk factors and treatment.

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## REVIEW METHODS

To review the evidence, use was made of a PubMed and Google Scholar search on 30 June 2021 which included articles published between 2009–2021, focusing on articles published between 2013–2021. Combinations of the following key words were used: ‘Nijmegen Breakage syndrome’, ‘diagnosis’, ‘TREC’s’, ‘KREC’s’, ‘primary immunodeficiency’, ‘cancer risk factors’, ‘lymphoid malignancies’, ‘chemotherapy’, ‘ECIL-8’, ‘HSCT’, ‘immunotherapy’, ‘second malignancies’ (key words: Nijmegen Breakage syndrome, leukemia, lymphoma, diagnosis, Haematopoietic Stem Cell Transplantation, antineoplastic agents) to review the evidence. The articles were in English. After compiling a list of potentially relevant articles, the full text of each paper was appraised, with particular emphasis on articles presenting patients with Nijmegen Breakage syndrome and lymphoid malignancies. Due to the rarity of the disease, case reports have been also included in the review.

## STATE OF KNOWLEDGE

**Epidemiology of NBS.** The disease likely occurs worldwide, but the highest occurrence of NBS has been described in Central and Eastern Europe due to the homozygous c.657\_661del5 mutation in *NBN* gene. Most patients with NBS originate from Poland (137 patients), followed by Russia (81 patients), Ukraine (56 patients), the Czech Republic (33 patients), Belarus (22 patients) and Slovakia (14 patients) [1, 2, 3]. An unexpectedly high incidence of NBS patients has also been reported in regions of southeast Germany, which is linked to Slavic origin and migration in the Middle Ages [10]. Single cases of such a genetic disorder have also been reported in Turkey, the United Kingdom, the United States of America, Austria, and Macedonia [2, 11]. Overall, analysis shows the highest prevalence of NBS in Poland and the Czech Republic [1, 2]. The manner of the reported prevalence can be explained that founder mutation c.657del5 most likely occurred less than 300 generations ago, which confirms that the original mutation predated the historic split and subsequent spread of the ‘Slavic people’. Moreover, the massive population movement after World War II influenced the geographical distribution of NBS [1]. Three patients with NBS have been described in Pakistan and are homozygous carriers of an other mutation c.1089C>A [2].

**Diagnosis of NBS.** The diagnosis of NBS is based on clinical manifestations and is confirmed by genetic analysis. The hallmark symptom of NBS is progressive microcephaly, which influences facial phenotype. The sloping forehead, the small nose, the receding chin, and the relatively large ears are the most common. Clinical features are also mild growth retardation, general lack of neurological symptoms, recurrent respiratory tract infections and malignancies [4]. Median age at diagnosis of NBS ranged from 4–5 years. This is about average for all the countries [2, 3]. Delay in diagnosis can result from the absence of unambiguous symptoms of NBS. There is no data on the most common symptoms at which diagnosis was started; notably, children with the mildest form of NBS remain underdiagnosed.

Microcephaly is a frequent clinical sign in various disease entities. The putative etiology for microcephaly is ascertained in 59% of all patients [12]. Deripapa et al. observed that

microcephaly was diagnosed at birth or shortly thereafter in 80% patients with NBS; however, the average delay of NBS diagnosis was 6.5 years [8]. Wolska-Kuśnierz et al. reported that prevalence of sporadic respiratory tract infections differs slightly from the incidence of severe respiratory problems. The most frequent infections are pneumonia and acute bronchitis. Surprisingly, the incidence of opportunistic infections is low, affecting only about 5% [7]. The distribution of infections does not differ significantly from the general paediatric population and makes the decision about starting diagnosis more difficult. Patients born in large cities and the capitals of Eastern Slavic countries are diagnosed earlier in comparison to those from rural regions, which may be partly explained by accessibility of specialist medical care [2, 3].

For Slavic populations, sequencing of exon 6 of the *NBN* gene to detect founder mutation c.657\_661del5 on both alleles is the method of choice [2, 3, 4]. Although genotyping common founder mutations in NBS is the basic diagnostic method, it results in delays in diagnosis. T-cell excision circles (TREC’s) and kappa-deleting recombination excision circles (KREC’s) can be potentially useful for the detection of NBS patients during neonatal screening [13]. Measurements of TREC’s, KREC’s and beta-actin as quality markers of DNA based on real-time polymerase chain reaction are used for screening severe primary immunodeficiency diseases (PID), mainly severe combined immunodeficiency. Routine screening is currently performed in the USA, Israel, New Zealand, Norway, Taiwan, several provinces in Canada, Switzerland, Iceland, Sweden, Italy (Tuscany), Spain (Catalonia), and in some regions in Australia [14]. In European countries, pilot projects have been announced in France, Spain, Norway, and the Netherlands [15]. In the USA, 2 cases of NBS were identified with solely low KREC’s copy numbers [13], and 2 cases of NBS were reported with low TREC’s [16, 17]. In the first population screening study for T and/or B immunodeficiency in Central and Eastern Europe, which included 44,287 newborns, one case of NBS with decreased values of TREC’s and KREC’s was identified [15].

Prophylaxis against viral, fungal, antibacterial and *Pneumocystis jiroveci* infections is not routinely recommended or implemented [7].

If there is lack of early diagnosis, children with NBS are exposed to ionizing radiation. Homozygous c.657–661del mutation affects cell radiosensitivity. Moreover, it is associated with disturbance in the activation of cell cycle checkpoints and with defects in DNA repair [18]. However, some patients were irradiated prior to NBS diagnosis with no unexpected toxicity or second tumour diagnosis [8].

Patients with undiagnosed NBS are exposed to adverse events related to immunization involving administration of viable vaccines and demand modifications in vaccination schedules. The association of immunodeficiency-related vaccine-derived rubella virus with cutaneous granulomatous disease has been reported. Granulomas are resistant to various types of therapy and have been resolved after haematopoietic stem cell transplantation (HSCT) in some patients with NBS [8, 19]. However, the use of attenuated BCG vaccine is favourable if the annual tuberculosis incidence is between 0.1–1%, and none of NBS patients experienced any adverse effects [7, 20].

**Factors of increased risk of lymphoid malignancies.** Attempts are being made to find the causes of increased carcinogenesis. The *MRE11*, *RAD50*, and *NBN* genes encode

proteins of the MRN complex which is involved in various DNA damage repair pathways. Some specific alternations of the *RAD 50* may be associated with childhood ALL [21]. Biallelic 657del5 mutation of *NBN* is associated with significant risk of cancers [22]. Interestingly, heterozygous carriers of mutation of the *NBN* gene have increased risk of development cancers, especially breast, prostate, and colorectal cancers, leukemias, and NHL. Among mutations in *NBN* gene, the 657 del5, the I171V and the R215W mutations are the most common [23]. However, the contribution of heterozygous p.I171V and p.R215W *NBN* mutations to cancer initiation is uncertain. The germline p.I171V may be considered as a risk factor in childhood ALL [21]. Significant differences in the number of chromatid breaks are observed only for mutant cells c. 657-661del, compared to control cells [24].

Chromosomal instability is associated with the development of complex genetic markers in pre-cancer cells. Despite the karyotype of NBS patients is generally normal, several abnormalities in the form of aneuploidies, structural rearrangements and marker chromosomes may be observed in 10–60% of cells [4].

Because of *NBN* mutation, T-cell development is affected at the stage of thymic output and V(D)J rearrangement. Circulating T cells of patients with NBS have signs of senescence which is already present in the youngest age group. Patients have a decreased level of CD4+ and CD8+ T cells. Although recent thymic emigrants and naïve T lymphocyte cell populations are significantly lower, the generation of antigen-primed (effector) T-cells is similar or even greater in NBS patients than in healthy control. There is high frequency of the more differentiated T cells expressing the senescent cell marker CD57, KLRG1, PD1 and do not express co-stimulatory molecule CD28 [5, 6]. Such properties might be related to increased susceptibility to lymphoid malignancies.

The extremely high cancer incidence in NBS can be explained by impaired telomeric repair which synergize with telomere attrition. Alternatively, cultivated lymphoblastoid cells derived from NBS patients with long survival times (>12 years) display shorter telomeres and lower caspase 7 activities, compared to cells derived from patients with short survival times (<3 years), suggesting low apoptosis rates and increased senescence rates with effective tumour suppression. In view of the bifunctional role of telomere attrition, an unambiguous role in carcinogenesis is difficult to identify [25].

Viral infections and monoclonal gammopathy are analyzed for early detection of changes preceding the development of malignancy. The predominance of EBV and monoclonal gammopathy, mainly of the IgM type, accompanied by an elevated level of IgM, are most often reported in patients who developed B-NHL. Clonal Ig (IgH, IgK or IgL) and/or TCR (BG and/or D) gene rearrangements are observed in 73.9% of patients and precede the diagnosis of lymphoma, on average, by 3 years. However, the strength of these associations needs to be further evaluated [26].

Another cause that directly affects carcinogenesis is chronic oxidative stress which is caused by endogenous reactive oxygen species overproduction and impairment of mitochondrial homeostasis. Oxidative stress leads to structural changes in DNA, proteins, and lipids [27]. Maciejczak et al. reported only significantly increased activity of catalase, decreased total antioxidant capacity and greater oxidative damage to proteins (advanced glycation end products, advanced

oxidation protein products), lipids (4-hydroxynonenal protein adducts, 8-isoprostanes) and DNA (8-hydroxy-2'-deoxyguanosine) [28]. Patients with NBS display statistically lower total antioxidant status. CoQ10, which deficiency of which may contribute to abnormal function of mitochondria during cancer, is significantly lower. There is a need for diagnosing oxidative stress as a potential disease biomarker. Supplementation with tocopherol and CoQ10 may enhance treatment efficacy in NBS diseases [29].

In literature, the search for cancer prevention is made in NBS patients. Antisense oligonucleotides as a potential cancer prophylaxis have been demonstrated. Antisense oligonucleotides have been used to enforce alternative splicing in humanized NBS murine mouse model to remove exon 6 and 7, and efficiently generate p-80 protein. This p80-nibrin contributes to a milder phenotype and improves survival [30].

**Characteristic of tumours in NBS patients.** The cumulative cancer incidence by the age of 10 years amounted to 41.6% [3] and 40.21% [2], and dramatically increased further to 71.6% [3] and 77.78% [2] by the age of 20 years. The first episode of malignancy occurred at a median age of 6.0–10.7 years. Male predominance is observed in patients with cancers and NBS [2, 3, 7, 9, 31], with the exception of one study with female predominance [8]. With respect to tumour type, NHLs represent most malignancies with a similar number of lymphoid tumours of B-cell origin and lymphomas developed from abnormal T-cells. Diffuse large B-cell lymphoma (DLBCL) was the most common among B-cell lymphomas and T-cell lymphoblastic lymphoma (T-LBL) among T-cell lymphomas. Interestingly, peripheral T-cell lymphomas (PTCL) are diagnosed in patients with NBS, which are tumours rarely diagnosed in the non-syndromic paediatric population [32]. Most patients are classified as stage III or IV. T-ALL is most often diagnosed in acute leukemia. Most patients are classified as high-risk group. Among other types of solid malignancies, there were diagnosed medulloblastoma, neuroblastoma, thyroid carcinoma, gastric carcinoma, low-grade glioma, rhabdomyosarcoma, unspecified liver tumour, dysgerminoma and ganglioglioma [2, 3, 7, 33]. Data on neoplasms in NBS patients are presented in Table 1.

**Treatment strategies of lymphoid malignancies in NBS patients.** There are no specific protocols for the treatment of lymphoid malignancies in children with NBS. The decision to reduce cytostatic agents depends on the oncologist and is based on the detailed medical history of every patient. Full-dose chemotherapy is the most frequently applied method [3, 8, 21]. In previous studies, it was suggested to administer 80% of dose chemotherapy, particularly alkylating agents as cyclophosphamide, DNA topoisomerase II, inhibitors, and antimetabolites such as methotrexate, and increase doses when well toleration is observed [11, 34]. Similarly, Aleinikova et al. reported that unrecognized NBS is the main reason for non-reduced dose regimens during induction therapy, and can be linked with increased toxicity of chemotherapy [9, 35]. Interestingly, Schutte et al. reduced over 25% of doses of chemotherapy agents, methotrexate, daunorubicin, doxorubicin, cyclophosphamide; however, severe adverse effects were observed [33].

Patients with a decreased immunoglobulins level should receive intravenous immunoglobulins (IVIG). According to current guidelines for immunoglobulin supplementation in

**Table 1.** Types of neoplasms in patients with NBS

References	Wolska-Kuśnierz (2015) [7]	Aleinikova (2015) [9]	Attarbaschi (2016) [31]	Deripapa (2017) [8]	Wolska-Kuśnierz (2020) [2]	Sharapova (2021) [3]
No. of patients with NBS	149	19	-	35	241	136
No. of patients with malignancies	63	9	26	20	151	62
Median age (years) at diagnosis	10.25	10.7	9.95	6.0	9.31	-
<b>B-NHL</b>	<b>30</b>	-	<b>12</b>	<b>11</b>	<b>49</b>	<b>20</b>
DLBCL	-	-	8	10	25	16
other	-	-	4	1	24	4
<b>T-NHL</b>	<b>21</b>	<b>7</b>	<b>11</b>	<b>4</b>	<b>46</b>	<b>18</b>
T-LBL	-	-	5	2	43	13
PTCL	-	-	6	2	3	5
ALCL	-	-	1	-	2	1
Other NHL	-	1	2	-	-	3
HL	7	-	-	-	9	2
<b>AL</b>	<b>17</b>	-	-	<b>7</b>	<b>32</b>	<b>12</b>
T-ALL	10	-	-	-	23	9
BCP-ALL	-	-	-	-	3	1
AML	-	-	-	-	1	1
other	7	1	-	1	5	1
Other solid malignancies	5	-	-	1	11	4

- no data; **NBS** – Nijmegen breakage syndrome; **B-NHL** – B-cell non-Hodgkin lymphoma; **DLBCL** – diffuse large B-cell lymphoma; **T-NHL** – T-cell non-Hodgkin lymphoma; **T-LBL** – T-cell lymphoblastic lymphoma; **PTCL** – peripheral t-cell lymphoma; **ALCL** – anaplastic large cell lymphoma; **HL** – Hodgkin lymphoma; **AL** – acute leukemia; **T-ALL** – T-cell acute lymphoblastic leukaemia; **BCP-ALL** – B-cell progenitor acute lymphoblastic leukemia; **AML** – acute myeloid leukemia.

primary immunodeficiencies, NBS patients should receive IVIG to maintain a serum IgG level over 5.0 g/l [36]. However, Deripapa et al. postulate that most (if not all) patients require IVIG substitution, and this decision should be made based on antibody-specific response. The ECIL-8 (European Conference on Infections in Leukaemia) group does not recommend routine antibacterial prophylaxis for patients at high risk, which contrasts with a previous international paediatric specific guideline. However, careful risk-benefit evaluation might favour antibacterial prophylaxis in individual patients, depending on their circumstances [37]. Due to musculoskeletal problems and central nervous system adverse events (CNS-AEs), the European Medicines Agency issued warning against the use of fluoroquinolones [37]. The previously recommend short time of oral prophylaxis in most immunodeficient patients, should not be implemented [36]. Reduction of invasive fungal infections is crucial for reduction of mortality during chemotherapy in NBS patients. Therefore, consistently with ECIL-8, fluconazole and posaconazole in patients over the age of 13 years older should be administered [38].

There is limited data on the frequency of toxicities in NBS patients with malignancies. In the Belarusian experience, all patients on chemotherapy experienced grade 3–4 complications according to the National cancer Institute Common Toxicity Criteria. The most common were neutropenic infections and haemostatic disorders [9, 35, 39].

Zawitkowska et al. reported that complications occur mostly in the induction phase of therapy, and in the reinduction phase. Among the analyzed toxicities in children with ALL, the most common included infection, gastrointestinal toxicity, and hepatotoxicity [40]. Dexamethasone during the induction phase of EURO-LB 02 may prevent central nervous system relapse more efficiently than prednisone, but produces significantly more grade 3 and 4 toxicities [41].

The main cause of death is cancer progression and treatment-related mortality, mostly associated with infectious complications [2, 3, 7, 8, 9, 31].

The overall survival rates for patients with DNA damage repair defect and without any cancer susceptibility syndromes are 9.7% and 68.7 %, respectively [42]. Mortality rate in patients with NBS and malignancies ranges from 22.6%-67.6% (Tab. 2). This divergence might be explained by delivery of full-dose chemotherapy along with IVIG treatment [7, 8]. The next contributing factor might be supplementation of immunoglobulins, irrespective of IgG levels. Low IgG levels were observed as a laboratory predictor of severe infections which is related to poor prognosis [7]. KRECs numbers are also reliable predictors of severe infections, oncological, and possible autoimmune complications [8]. TRECs has been reported as indicator of thymic output of naïve T-cells [5]; however, Deripapa et al. observed no correlation between TRECs and severity of infections. Moreover, complex aberrations related to chromosomal instability which are

**Table 2.** Outcome of patients with Nijmegen breakage syndrome and lymphoid malignancies

	Wolska-Kuśnierz (2015) [7]	Attarbaschi (2016) [31]	Deripapa (2017) [8]	Wolska-Kuśnierz (2020) [2]	Sharapova (2021)[3]
No. of patients with malignancies	63	26	20	151	62
CR	-	10 (38%)	-	-	35(71.4%)
Relapse/progression	-	10 (38%)	5 (25%)	-	12 (24.5%)
Death	28 (44%)	16 (62%)	5 (25%)	101 (67.6 %)	14 (22.6%)

- no data; **CR** – complete remission

revealed in cytogenic karyotype and microarray tests, trigger poor outcome [43]. The next factor for poor prognosis is not performing a transplantation [8].

About a quarter of patients with NBS have relapse [3, 8]. Tomasiak et al. observed that heterozygous carriers of the c.657\_661del 5 mutation in the *NBN* gene are at higher risk of BCP-ALL relapse within the CNS, compared to NBS patients [44].

**Possibilities of molecular targeted therapy for children with NBS.** Considering the poor outcome of patients with NBS and malignancies, new strategies of therapy are still needed to reduce clonality of lymphocytes. There are limited data on the use of molecular targeted therapy in the treatment of children with PIDs and lymphoid malignancies. Therapeutic approach should be considered in relation to the nature of the NBS as a DNA repair disorder, and to the remaining immune function available in the patient's body, which is required to enhance immunotherapy. Antitumoural targeted therapies, such as monoclonal antibodies (mAb), chimeric antigen receptor (CAR) T-cells and small-molecule inhibitors, potentially offer significant opportunities for optimizing treatment of lymphoid malignancies [45, 46].

Monoclonal antibodies have shown great advances since their first member rituximab in 1997, respectively. The next-generation of antibodies aim to boost antibody-dependent cellular cytotoxicity and component-dependent cytotoxicity, for example, obinutuzumab. MAbs may also be combined with antineoplastic drugs intensifying its concentration in tumour cells, such as gemtuzumab ozogamicin. Bi-specific T-cell engaging (BiTE) Abs recruit polyclonal T lymphocyte population, avoiding the need for antigen specificity and putative tumour immune evasion through HLA downregulation. MAbs can also interfere with immune checkpoint pathways and allow potentiate anti-tumour responses, for instance, pembrolizumab -anti-programmed cell-death 1 – PD1. The CAR T-cells, which are autologous T-cells from patients and genetically engineered to express tumor-reactive CARs, are the most recent method of treatment [46, 47]. The next group are inhibitors of PI3K/

AKT/mTOR pathway, such as sirolimus. In EBV- driven lymphoid malignancies anti-EBV cytotoxic T-lymphocytes are considered as a salvage therapy in refractory disease [45]. There are reported cases that therapy was well tolerated in patients with ataxia telangiectasia [48].

Therapies that target cell surface antigens are currently undergoing clinical trials with most common indication for refractory/relapse lymphoid malignancy [46]. Agents that may potentially be used in the treatment of patients with NBS and lymphoid malignancies are listed in Table 3 [45, 46, 47]. There are no supportive data on the use of inhibitors of the PI3K/ AKT/mTOR pathway in patients with DNA repair defects [45].

Referring to the early age of cancer development in NBS, there are data on blinatumomab in the treatment of children aged one year or older [46]. Patients with NBS and leukemias are a potential group among other PIDs who can benefit from treatment with blinatumomab due to the persistent residual function of T cells [45]. However, the use of molecular targeted therapies in patients with NBS has some limitations due to the nature of immunodeficiency. Reinfusing genetically engineered autologous T lymphocytes prove challenging, but it might be overcome by the development of universal allogeneic CAR T-cells derived from healthy donors [49]. Compared blinatumomab with CAR T-cells, blinatumomab has a few advantages: it is an off-the shelf product; it is less expensive, and the short half-life enables the precise control of serum levels [50].

Despite promising outcomes of targeted therapies, patients should be carefully observed due to potential toxicities. Invasive fungal disease is common in patients treated with blinatumomab, gemtuzumab ozogamicin and tisagenlecleucel, and uncommon in patients treated with brentuximab vedotin and pembrolizumab [46]. These data are particularly important for patients with immunodeficiencies and highlight the need to intensify antifungal prophylaxis. Cytokine release syndrome (CRS) is another toxicity, and many patients require administration of tocilizumab, an anti-interleukin 6 receptor antagonist [51]. Neurotoxicity after blinatumomab cannot be ignored in patients with NBS and prone to neurovegetative disorders [45, 52].

**Table 3.** Possibilities of molecular targeted therapy for children with NBS [45–47]

Target	Immunotherapy	Lymphoid malignancy
	denituzumab mafodotin	r/r B-ALL and r/r B-NHL*
CD19	BiTEs blinatumomab	Ph-negative CD19-positive B-ALL r/r or after HSCT; Relapsed B-ALL*
	CAR-T cell therapy tisagenlecleucel	B-ALL and DLBCL; r/r B-ALL*
	CAR-T cell therapy axicabtagene ciloleucel	r/r pre-B-ALL and r/r B-cell NHL*
CD20	ofatumumab	Recurrent lymphoblastic lymphoma and ALL in CR*
	obinutuzumab	Lymphoma*
CD22	epratuzumab	relapsed ALL*
CD30	brentuximab vedotin	r/r HL, systemic ALCL, r/r HL*
CD33	gemtuzumab ozogamicin	AML and r/r AML*
CTLA-4	ipilimumab	r/r lymphoma*
	nivolumab	
PD-1	pembrolizumab	r/r lymphoma*
	atezolizumab	

\* ongoing clinical trial. **NBS** – Nijmegen breakage syndrome; **CD** – cluster of differentiation; **ALL** – Acute Lymphoblastic Leukemia; **NHL** – non-Hodgkin lymphoma; **HSCT** – Hematopoietic Stem Cell Transplantation; **Ph** – Philadelphia chromosome; **DLBCL** – diffuse large B-cell lymphoma; **C** – complete remission; **HL** – Hodgkin lymphoma; **ALCL** – anaplastic large cell lymphoma; **AML** – Acute Myeloblastic Leukemia; **r/r** – relapsed or refractory; **CTLA-4** – cytotoxic T-lymphocyte-associated protein 4; **PD-1** – programmed cell death protein 1; **PD-L1** – programmed death-ligand 1 protein

**Table 4.** Characteristics of patients with NBS who underwent HSCT

	Wolska- Kuśnierz (2015) [7]	Deripapa (2017) [8]	Wolska-Kuśnierz (2020) [2]	Sharapova (2021) [3]
No. of patients with NBS	149	35	241	136
No. of patients with HSCT	14 (9.4%)	15 (42.3 %)	49 (20.3 %)	13 (9.6 %)
Median age of HSCT	8.8	-	9.0	12.3
HSCT due to malignancy	11 (78.6%)	8 (53.3%)	35 (71.4 %)	7 (53.8 %)
HSCT due to immunodeficiency	3 (21.4%)	5 (33.3 %)	14 (28.6%)	5 (38.5%)
Death from relapse or infections	5 (35.7%)	2 (13.3 %)	13 (26.5 %)	5 (38.5%)

- no data NBS - Nijmegen breakage syndrome  
 HSCT - haematopoietic stem cell transplantation

**HSCT in NBS patients.** The general reasons of HSCT are defined as infection (severe or recurrent infections), malignancy, bone marrow failure (defined as leukopenia, anaemia, or thrombocytopenia, without the presence of infection or malignancy), autoimmunity or pre-emptive [53].

Transplantation in NBS patients is recommended in first complete remission (CR) of lymphoma or leukemia. Some patients were transplanted after secondary tumours. Another indication was clinically relevant immunodeficiency with severe or chronic infections, or immune dysregulation [2]. Patients with NBS who underwent HSCT are presented in Table 4. The reasons why not all patients have transplantations are the low percentage of patients who achieve complete remissions and the relatively high mortality due to infections during chemotherapy. On the other hand, there is disproportion between the severity of the infection's symptoms and underlying immunodeficiency. Research is needed to validate the transplantation in asymptomatic patients with NBS [2, 3].

Despite the immunodeficiency, rejection and poor stem cell engraftment are likely without preparative conditioning. There are myeloablative conditioning with alkylating agents or total body irradiation and reduced intensity conditioning (RIC) without alkylating agents and with fludarabine  $\leq 150$  mg/m<sup>2</sup> and cyclophosphamide  $\leq 40$  mg/kg [53]. The use of an RIC in comparison to a myeloablative conditioning regimen demonstrates beneficial HSCT outcomes in patients with NBS. Sharapova et al. reported a patient who received myeloablative conditioning but died soon after HSCT [3]. Most patients received modified Fanconi RIC based on fludarabine, and cyclophosphamide with or without anti-thymocyte globulin therapy [2, 3, 53, 54]. Laberko et al. reported that a conditioning regimen with treosulfan 30 g/m<sup>2</sup> demonstrated a low level of early transplanted-associated toxicity, and enhanced graft function with stable donor chimerism in comparison to regimen with busulfan [55].

In the literature the incidence of acute and chronic graft versus host disease (GvHD) is inconclusive, showing a higher rate of acute GvHD. Sharapova et al. observed acute GvHD in 38.5% patients [3]. However, there are reports that none of patients developed GvHD [8, 48]. This does not seem to be related with use of matched, unrelated donors and the presence of malignancies. The most common non-GvHD complication is viraemia due to adenovirus, cytomegalovirus, Epstein-Barr virus, and BK virus. Visceral toxicity is rarely observed in NBS patients [2]. There is a report on a patient who developed cholelithiasis and recto-sigmoid ulcers [56]. Some patients fully rejected the graft [8]. Fatal infections after HSCT are the most common cause of death. The next causes of death after HSCT are cancer progression and

relapses, particularly of primary lymphomas [2, 3, 7, 8]. Patients who undergo pre-emptive transplantation had a more favorable outcome. Wolska-Kuśnierz et al. reported no transplant-related deaths in this group [2]. Sharapova et al. observed that 67% of patients survived [3].

In a cohort study, a 20-year OS in a transplanted group did not significantly differ from that observed in the non-transplanted group. However, NBS patients with diagnosed cancer who received HSCT had a significantly higher 20-year OS than those who did not. (42.7% vs. 30.3%) [2]. Similarly, Sharapova et al. reported that from the time of NBS diagnosis, the probability of 15-year OS in patients who underwent HSCT is significantly higher, in contrast to those who did not undergo HSCT. (61.5% vs 36.9%) [3]. Despite the favourable outcome of patients who undergo HSCT due to malignancies, there is a need for the further monitoring of HSCT as a pre-emptive therapeutic approach. Cancer-free survival outcome should be evaluated in these patients.

**Second malignancies in NBS patients.** Second malignant neoplasm (SMN) is defined as a malignancy which occurs more than 5 years from diagnosis of the first malignancy among patients who achieved complete remission. Frequency of patients with NBS who developed subsequent malignancy ranges from 13.2%-19%. However, it includes neoplasms which occur less than 5 years from the first diagnosis of malignancy, which makes this statistic less comparable with the other [2, 3, 31]. The median age of diagnosis is 18 years [2] and the most frequent second malignancies are lymphomas [2, 3, 7]. Wolska-Kuśnierz et al. confirmed T-cell leukemia / lymphoma as half of secondary tumours [2] The most common at primary diagnosis were B-cell lymphoma / leukemia with DLBCL as most frequent [2, 3, 31]. The general risk factors for the development of second neoplasms are acute-lymphoblastic-leukemia-type therapy, cumulative anthracycline dose, CNS-involvement, female gender, and cancer-predisposing conditions [57].

On the other hand, reduced chemotherapy does not prevent the development of SMN [2]. Patients with NBS have a higher risk of SMN, compared to other pre-existing disorders, but much less than patients with constitutional mismatch repair disease [31, 58]. In the general population, Attarbaschi et al. reported that lymphoid malignancies are the most frequent SMN; however, Moser et al. observed predominantly carcinomas [57, 58].

The survival rate 10 years after diagnosis of SMN is 47%. Patients with a known cancer predisposing syndrome have a much lower survival rate of only 11.1% [57]. The majority of patients with NBS and SMN die due to tumour progression and infectious complications during chemotherapy. In the

literature, there are reports of patients who were successfully transplanted and remained in the remission of cancer for more than 5 years. Analysis of the impact of HSCT on the cumulative risk of death in the presence of secondary malignancy as a competing event, showed that the risk of death was reduced twofold in transplanted patients compared with non-transplanted individuals [2].

## CONCLUSIONS

Further meta-analysis is essential to establish the best monitoring and treatment regimen in patients with NBS and lymphoid malignancies. TRECs and KRECs can be potentially useful for the detection of NBS patients during neonatal screening. Oxidative stress should be diagnosed as potential disease marker. Supplementation with tocopherol and CoQ10 may enhance treatment efficacy in NBS diseases. Patients with NBS and lymphoid malignancies are particularly vulnerable to a high incidence progression of disease and therapy-associated toxicity. However, full-dose chemotherapy is the most frequently applied method. HSCT has a significant positive effect on survival.

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