Clinicopathology Figures and Survival of Non-Hodgkin’s Lymphoma in Iran

Safa Najafi,1 Mehrdad Payandeh,2 and Masoud Sadeghi1,2*
1Breast Diseases Department, Breast Cancer Research Centre, ACECR, Iran
2Cancer Research Center, Kermanshah University of Medical Sciences, Kermanshah, Iran
*Corresponding author: Masoud Sadeghi, Cancer Research Center, Kermanshah University of Medical Sciences, Kermanshah, Iran. Tel: +98-9185960644, Fax: +98-8334276471, E-mail: sadeghi_mbrc@yahoo.com

Received 2016 January 01; Revised 2016 February 22; Accepted 2017 January 03.

Abstract

Background: Non-Hodgkin’s lymphoma (NHL) is the fifth most frequently diagnosed cancer whose incidence has risen by at least 100 percent over the past five decades especially in the West.

Objectives: The aim of this study is to investigate Clinicopathology figures, overall survival (OS) and Progression-free survival (PFS) in patients with NHL in Iran.

Methods: In a descriptive study, 143 patients referred to Clinic of Hematology in two centers between 2005 and 2014. We checked age, sex, types and subtypes of NHL, recurrence, Ki-67, organomegaly, lymphadenopathy, radiotherapy, OS and PFS in the patients. The mean follow-up was 49 months. All patients received R-CHOP regimen for 6 to 8 cycles.

Results: The mean age at diagnosis for the patients was 46 years (range, 16 - 82) and 76.9% had age < 60 years and 62.2% were male. 58% patients had primary nodal NHL and out of 139 patients, 88.5% were aggressive NHL. Organomegaly and lymphadenopathy were in 12 patients (8.4%) and 80 (55.9%), respectively. Of all patients, 106 patients (74.1%) were treated with radiotherapy. 2-year OS rate was 91.5% and 5-year OS rate was 85%. 2-and 5-year PFS rates were 79.6% and 63.3%, respectively.

Conclusions: Percentage of NHL in males is more than females and the median age is around 45 years. Also, the prevalence of nodal is more compared with extranodal in NHL patients.

Keywords: Non-Hodgkin’s Lymphoma, Overall Survival, Progression-Free Survival, Lymphadenopathy

1. Background

Non-Hodgkin’s lymphomas (NHLs) are a heterogeneous group of malignancies of the lymphoid system (1, 2) and account for almost 3.4% of cancer deaths in the US (3). This disease is the fifth most frequently diagnosed cancer and its incidence has risen by at least 100% over the past five decades especially in Western countries (4). Based on WHO’s classification of hematological and lymphoid tumors, these diseases have been divided as B-and T-cell neoplasms. B-cell lymphomas account for approximately 90% of all lymphomas, and the 2 most common histological disease entities are follicular lymphoma and diffuse large B-cell lymphoma (DLBCL) (5). Compared with Western countries, high-grade NHLs are more prevalent in Iran (6). Expression of Ki-67, a nuclear antigen protein present in all cycling cells, is used to distinguish the growth fraction of tumors (7). General prognosis of NHL depends on various tumor and host-related factors including age of patients and performance status, histological subtype, grade and stage of lymphoma (8). Other risk factors are sex, race, primary or hereditary immune deficiency syndrome, autoimmune diseases such as Sjogren’s, Celiac disease, Rheumatoid arthritis, systemic Lupus Erythematosus, certain infections (such as EBV, HCV), genetic factors and family history of lymphoma in first-degree relatives (4). Rituximab is the monoclonal antibody responsible for all clinical improvement noted to date. The addition of this drug to cyclophosphamide, doxorubicin, vincristine, prednisone (R-CHOP) improves the response rate, progression-free survival (PFS) and overall survival (OS) in NHL, especially DLBCL (9).

2. Objectives

The aim of this study is to investigate Clinicopathology figures, OS and PFS in patients with NHL in Iran.

3. Methods

3.1. Patients

Between 2005 and 2014 in a descriptive study, 143 patients referred to Clinic of hematology in two centers. We checked age, sex, types and subtypes of NHL, recurrence, Ki-67, organomegaly, lymphadenopathy, radiotherapy, OS and PFS in the patients. The mean follow-up was 49 months and during this time, there was 32 deaths and 4 patients...
lost follow-up and were censored from our study. OS was defined from the date of diagnosis until death from any cause or the date of the last follow-up. PFS was defined as the time elapsed between treatment initiation and (3) tumor progression (growth of the tumor, new tumors, metastasis etc) (4) death from any cause or the date of the last follow-up, whichever came first, with censoring of patients who are lost to follow-up. All patients received R-CHOP regimen for 6 to 8 cycles.

3.2. Statistical Analysis

The curves of OS and PFS were plotted by GraphPad Prism 5 software. Date analysis was done by SPSS version 19 software.

4. Results

The median age at diagnosis for the patients was 46 ± 16 years (range, 16 - 82) that 110 patients (76.9%) had age < 60 years and 89 patients (62.2%) were male (Table 1). Eighty-three patients (58%) had primary nodal NHL and out of 139 patients, 123 patients (88.5%) were aggressive NHL. In follow-up time, 45 patients (31.5%) had recurrence. Out of 87 patients, Ki-67 < 65% was in 51 patients (58.6%). Organomegaly and lymphadenopathy were in 12 patients (8.4%) and 80 (55.9%), respectively. Of all patients, 106 patients (74.1%) were treated with radiotherapy.

Table 2 shows the correlation between the type of NHL and a number of variables in patients. There was just a significant correlation between lymphadenopathy with the type of NHL (P = 0.000). Therefore, lymphadenopathy was more in nodal NHL compared to extranodal NHL.

5. Discussion

The prevalence of all NHLs coded as being of extranodal origin is between 25% and 35% in most countries that stomach, skin and small intestine were the most common extranodal sites (10). The risk factor(s) of lymphoma has not been determined or clear yet (11). The most common types of lymphoma have originated from B cells (12). In a study (13), out of 77 patients with newly diagnosed primary
**Table 2.** The Relationship Between Type of Non-Hodgkin Lymphoma and a Number of Variables in Patients (n = 143)

| Variables                        | Nodal, n = 83 | Extranodal, n = 60 | P Value |
|----------------------------------|---------------|--------------------|---------|
| **Age, years**                   |               |                    |         |
| Mean                             | 44.2          | 48.8               | 0.084   |
| <60                              | 66 (79.5%)    | 44 (73.3%)         | 0.252   |
| **Sex**                          |               |                    |         |
| Male                             | 48 (57.8%)    | 41 (68.3%)         | 0.135   |
| **Subtype of primary NHL, n = 139** |             |                    |         |
| Aggressive                       | 73 (89%)      | 50 (87.7%)         | 0.508   |
| **Recurrence**                   |               |                    |         |
| Yes                              | 22 (26.5%)    | 23 (38.3%)         | 0.094   |
| <65%                             | 30 (58.8%)    | 21 (58.3%)         | 0.569   |
| **Organomegaly**                 |               |                    |         |
| Yes                              | 11 (12%)      | 2 (3.3%)           | 0.057   |
| **Lymphadenopathy**              |               |                    |         |
| Yes                              | 65 (78.3%)    | 15 (25%)           | 0.000   |

NHL of the bone, 56 patients (72.7%) were male; the median age was 41.8 years, with a range of 16 - 84 years. In another study (14), among 387 NHL patients, the median age was 55 years with a male to female (M: F) ratio of 1.9:1. Naz et al. (15) reported that in NHL patients, 67% were male and M: F ratio was 2.6:1 with the mean age of 43.9 years (range, 6 to 80 years). In our study, the median age was 46 years, 62.2% male that M: F ratio was 1.7:1. Therefore, the percentage of NHL in males is more than females and the median age is around between 45 years.

In a research on NHL patients, the extranodal involvement was seen in 40.3% cases, while 59.7% cases showed nodal involvement (15). Padhi et al. (16) showed that primary extranodal constituted 22.0% of all NHLs. Also, Otter et al. (17) reported that primary extranodal lymphoma was 41% in their study. In this study, 58% patients had nodal NHL. Therefore, the prevalence of nodal is more than extranodal. A study (14) showed that in their patients with NHL, the aggressive histological subtypes predominate. Our study confirmed this result.

In a study in our area (18), 67.9% patients with NHL had Ki-67 < 60% that in our study, Ki-67 < 65% is 58.6%. The results of two studies are almost similar.

The 5-year OS rate and 5-year PFS rate were significantly higher in B-cell group than in T-cell group (69.5% vs. 35.5%, 53.3% vs. 28.9%) (9). In patients with DLBCL, The 5-year event-free survival was 61% for patients receiving chemotherapy alone and 64% for patients receiving CHOP plus radiotherapy; the 5-year OS were 72% and 68%, respectively (20). A study in primary gastric lymphoma patients showed that 5-year disease-free survival and OS rates were 60% and 70%, respectively (8). Hauptrick et al. (21) in 1666 treated patients with rituximab monotherapy, reported that median PFS was 23.5 months. Coiffier et al. (22) compared 8 cycles R-CHOP with CHOP in elderly patients with DLBCL resulted in a significant survival benefit for patients in the R-CHOP arm (OS 37 months). In a research on 2745 NHL patients (23), 2-year PFS for subtypes of DLBCL was 64%, BL (Burkitt’s lymphoma) 56%, lymphoma intermediate between DLBCL and BL 70%. Also, 2-year OS rates for DLBCL was 76.9%, BL 56%, lymphoma intermediate between DLBCL and BL 80%. Among patients of DLBCL subgroup, 2-year PFS and OS according to different chemotherapy regimens were MCP-842 with 51% and 69.2%, CHOP: 65% and 80%, R-CHOP: 83.3%, and 88.7% (23). Another study (14) showed that 5-year OS rate was 65% for patients. Two studies in our area showed the 3-year, 5-year and 10-year OS rates for nodal NHL patients were 65%, 54.2% and 51%, respectively (24) and for extranodal were 70%, 62.2% and 60.8%, respectively (25). In our study that more patients treated with R-CHOP plus radiotherapy, 2 and 5-year OS rates for all patients were 91.5% and 85% and also 2 and 5-year PFS were 79.6% and 63.3%, respectively. Also, 2 and 5-year OS rates for aggressive NHL patients were 91% and 79.1%, respectively; whereas 2 and 5-year PFS were 79% and 63%, respectively. Therefore, the results show that treated patients with R-
2-year OS rate was 91.5% and mean OS was 20 months. Also, 5-year OS rate was 85% and mean OS was 36.1 months; whereas, for aggressive NHL patients, 2-year OS rate was 91% and mean OS was 20 months and also 5-year OS rate was 79.1% and mean OS was 35.5 months.

5.1. Conclusions

Percentage of NHL in males is more than females and the median age of NHL patients is around 45 years. Also, the prevalence of nodal is more compared with extranodal in NHL patients.

Acknowledgments

There is no acknowledgement.

Footnotes

**Authors’ Contribution:** Safa Najafi and Mehrdad Payande were supervisor and determined patients; Masoud Sadeghi was the corresponding author and revised the article.

**Conflict of Interest:** There is no conflict of interest.

**Financial Disclosure:** There is no financial disclosure.
Figure 3. Shows 2-and 5-Year PFS in NHL Patients That 2-and 5-Year PFS Rates Were 79.6% and 63.3%, Respectively.

Also, mean 2-and 5-year PFS were 18.7 and 32.3 months, respectively; whereas for aggressive NHL patients, 2-year PFS rate was 79% and mean PFS was 18.7 months and also 5-year PFS rate was 63% and mean PFS was 31.5 months.

References
1. Memar B, Aledavood A, Shahidsales S, Ahadi M, Farzadnia M, Raziee HR, et al. The Prognostic Role of Tumor Marker CA-125 in B-Cell non-Hodgkin’s Lymphoma. Iran J Cancer Prev. 2015;8(1):42-6. [PubMed: 25825970].
2. Aminian O, Abedi A, Chavoshi F, Ghasemi M, Rahmatt-Najarkolaei F. Evaluation of Occupational Risk Factors in Non-Hodgkin Lymphoma and Hodgkin’s Disease in Iranian Men. Iran J Cancer Prev. 2012;5(4):889-93. [PubMed: 2352969].
3. Alexander DD, Mink PJ, Adami HO, Chang ET, Cole P, Mandel JS, et al. The non-Hodgkin lymphomas: a review of the epidemiologic literature. Int J Cancer. 2007;120 Suppl 12:39. doi: 10.1002/ijc.22719. [PubMed: 17405021].
4. Muller AM, Iborst G, Mertelsmann R, Engelhardt M. Epidemiology of non-Hodgkin’s lymphoma (NHL): trends, geographic distribution, and etiology. Ann Hematol. 2005;84(3):12. doi: 10.1007/s00277-004-0939-7. [PubMed: 15480663].
5. Ansell SM, Armitage J. Non-Hodgkin lymphoma: diagnosis and treatment. Mayo Clin Proc. 2005;80(8):1087-97. doi: 10.4065/80.8.1087. [PubMed: 16092591].
6. Hashemi-Bahremani M, Parwaresch MR, Tabrizchi H, Gupta RK, Raffi MR. Lymphomas in Iran. Arch Iran Med. 2007;10(3):343-8. [PubMed: 17604472].
7. Broyde A, Boycov O, Strenov Y, Okon E, Shpilberg O, Bairey O. Role and prognostic significance of the Ki-67 index in non-Hodgkin’s lymphoma. Am J Hematol. 2009;84(6):338–43. doi: 10.1002/ajh.21406. [PubMed: 19384938].
8. Hosseini S, Dehghan P. Primary non-hodgkin lymphoma of the stomach: clinicopathological characteristics and prognostic factors in Iranian patients. Iran J Cancer Prev. 2014;7(4):219-24. [PubMed: 25628842].
9. Kahl B. Chemotherapy combinations with monoclonal antibodies in non-Hodgkin’s lymphoma. Semin Hematol. 2008;45(2):90-4. doi: 10.1053/j.seminhemat.2008.02.003. [PubMed: 18381031].
10. Newton R, Ferlay J, Beral V, Devesa SS. The epidemiology of non-Hodgkin’s lymphoma: comparison of nodal and extra-nodal sites. Int
11. Irshaid F, Tarawneh K, Alshdefat A, Dilmi F, Jaran A, Al-Hadithi R, et al. Loss of P16 Protein Expression and Its Association with Epstein-Barr Virus LMP1 Expression in Hodgkin’s Lymphoma. Iran J Cancer Prev. 2013;6(2):78–84. [PubMed: 25250915].

12. Aledavood SA, Ghavam-Nasiri MR, Ghaffarzadegan K, Raziee HR, Saboori G, Anvari K, et al. Hepatitis-C Infection Incidence Among the non-Hodgkin’s B-cell Lymphoma Patients in the Northeast of Iran. Iran J Cancer Prev. 2014;7(3):147–51. [PubMed: 25250166].

13. Barbieri E, Cammelli S, Mauro F, Perini F, Cazzola A, Neri S, et al. Primary non-Hodgkin’s lymphoma of the bone: treatment and analysis of prognostic factors for Stage I and Stage II. Int J Radiat Oncol Biol Phys. 2004;59(3):760–4. doi: 10.1016/j.ijrobp.2003.11.020. [PubMed: 15183479].

14. Economopoulos T, Asprou N, Stathakis N, Fountzilas G, Pavlidis N, Papaspyrou S, et al. Primary extranodal non-Hodgkin’s lymphoma of the head and neck. Oncology. 1992;49(6):484–8. [PubMed: 1465289].

15. Naz E, Mirza T, Aziz S, Danish F, Siddiqui ST, Ali A. Frequency and clinicopathologic correlation of different types of non Hodgkin’s lymphoma according to WHO classification. J Pak Med Assoc. 2011;61(3):260–3. [PubMed: 21465941].

16. Padhi S, Paul TR, Challa S, Prayaga AK, Rajappa S, Raghunadharao D, et al. Primary extra nodal non Hodgkin lymphoma: a 5 year retrospective analysis. Asian Pac J Cancer Prev. 2012;13(10):4889–95. [PubMed: 23244076].

17. Otter R, Gerrits WB, vd Sandt MM, Hermans J, Willemze R. Primary extranodal and nodal non-Hodgkin’s lymphoma. A survey of a population-based registry. Eur J Cancer Clin Oncol. 1989;25(8):1203–10. [PubMed: 2767099].

18. Payandeh M, Sadeghi M, Sadeghi E. The ki-67 index in non-hodgkin’s lymphoma: Role and prognostic significance. American J Cancer Prevention. 2015;3(5):100–2.

19. Zou GR, Zhang YJ, Xie FY, Zheng W, Li HX, Xia YF, et al. [Prognosis and treatment strategies of primary B-cell and NK/T-cell nasopharyngeal non-Hodgkin’s lymphoma at early stage]. Ai Zheng. 2006;25(12):1543–9. [PubMed: 1796636].

20. Friedberg JW, Fisher RI. Diffuse large B-cell lymphoma. Hematol Oncol Clin North Am. 2008;22(5):941–52. doi: 10.1016/j.hoc.2008.07.002. [PubMed: 18954744].

21. Hauptrock B, Hess G. Rituximab in the treatment of non-Hodgkin’s lymphoma. Biologics. 2008;2(4):619–33. [PubMed: 19707443].

22. Coiffier B, Lepage E, Briere J, Herbrecht R, Tilly H, Bouabdallah R, et al. CHOP chemotherapy plus rituximab compared with CHOP alone in elderly patients with diffuse large-B-cell lymphoma. N Engl J Med. 2002;346(4):235–42. doi: 10.1056/NEJMoa011795. [PubMed: 11807147].

23. Sengar M, Akhade A, Nair R, Menon H, Shet T, Gujral S, et al. A retrospective audit of clinicopathological attributes and treatment outcomes of adolescent and young adult non-Hodgkin lymphomas from a tertiary care center. Indian J Med Paediatr Oncol. 2011;32(4):197–203. doi: 10.4103/0971-5851.95140. [PubMed: 22563152].

24. Payandeh M, Sadeghi M, Shahriari-Ahmadi A, Sadeghi E. The survival of nodal non-hodgkin’s lymphoma patients in the west of Iran. American J Cancer Prevention. 2015;3(5):95–8.