

Cancer Association of South Africa (CANSA)



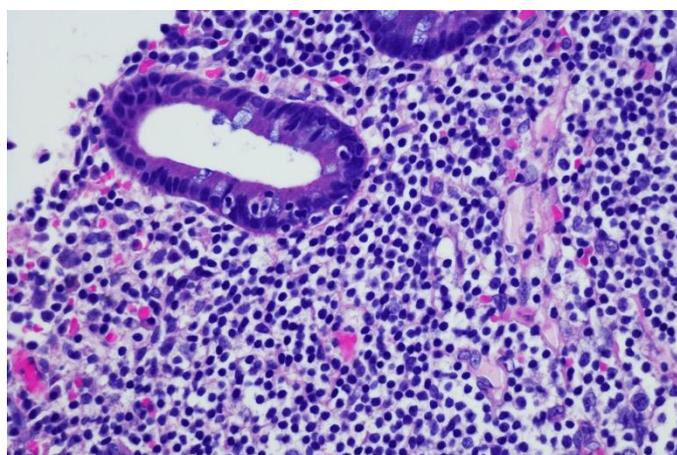
Fact Sheet on Nodal Marginal Zone Lymphoma

Introduction

Lymphoma is a type of cancer. It can happen when growth of a lymphocyte (white blood cell) population goes out of control. Marginal zone lymphomas develop from B lymphocytes (B cells) that are normally found in the 'marginal zone'. The marginal zone is at the edge of the area of lymphoid tissue and is where B cells are normally found.

[Picture Credit: NMLZ]

Lymphoid tissue is part of the immune system, for example the lymph nodes or spleen.



Other types of marginal zone lymphomas are MALT Lymphoma (extranodal marginal zone lymphoma) and splenic marginal zone lymphoma.

Nodal Marginal Zone Lymphoma

Nodal marginal zone lymphoma (nodal MZL) is a rare type of low-grade (slow-growing) non-Hodgkin's Lymphoma (NHL). It is a rare type of lymphoma – fewer than 2 in 100 cases of NHL are Nodal Marginal Zonal Lymphoma.

Two clinicopathological forms of Nodal Marginal Zone Lymphoma are recognised: adult type and paediatric-type. Nodal Marginal Zone Lymphomas show overlapping features with other types of Marginal Zone Lymphoma, but distinctive features as well. It remains an enigmatic entity with accompanying difficulties in diagnosis and a lack of knowledge of prognosis and treatment.

Nodal marginal zone lymphoma is more common in older adults. People with this disease are usually diagnosed when they are 60 years old or older. It is more common in women than in men. More than 70% of people are stage 3 or 4 at the time of diagnosis.

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019

Page 1

Bron, D., Meuleman, N. & Eurobloodnet for rare diseases and EHA SWG 'Aging and Hematology'. 2019.

PURPOSE OF REVIEW: Choosing an optimal treatment in older patients with indolent lymphomas is a challenge for hematologists. They must concomitantly treat some potentially curable entities, manage other symptomatic incurable diseases and protect their patients from life-threatening toxicities. Specific recommendations for older patients with different subtypes of marginal zone lymphomas are thus required in terms of treatment and supportive care.

RECENT FINDINGS: All the data in the literature agree that the therapeutic approach of older patients with malignant hemopathies should include the appraisal of their life expectancy and of the prognostic factors of their tumor, the evaluation of their physiological and cognitive functions and their socioeconomic environment, and their expectancy in terms of quality of life. Major progresses have, therefore, been achieved in the management of lymphoma patients of 80 years and older.

SUMMARY: With an optimal 'geriatric assessment', most of the recommended treatments are also appropriate in older marginal zone lymphoma patients. Extranodal MALT lymphoma: eradication of the pathogen is a major part of the first-line therapy. Prognosis is excellent in early stages. In advanced stages, observation and anti-CD20 antibodies with or without cytostatic drugs are recommended. Nodal MZL: Usually confined to lymph nodes, bone marrow and peripheral blood, they should be managed as follicular lymphomas. Splenic MZL: in this unique entity involving the spleen, the bone marrow and the peripheral blood. Hepatitis infection should be eradicated before considering treatment. Only symptomatic patients require to be treated by splenectomy and/or anti-CD20 antibodies.

Magro, C.M., Momtahan, S., Coleman, M. & Grossman, M.E. 2019.

"Epidermotropic B cell lymphoma represents a rare form of marginal zone lymphoma presenting as a disseminated skin rash resembling pityriasis rosea. To date there are 8 reported cases. In addition to the widespread nature of the skin rash, there is a proclivity for spleen and bone marrow involvement raising consideration regarding its categorization as a systemic lymphoma. We present an 89-year-old man with epidermotropic B cell lymphoma, who presented with a pityriasis rosea-like skin rash. An initial diagnosis of diffuse large cell B cell lymphoma was made based on the extent of dermal-based large cell infiltration. However, after recognizing the epidermotropic component and the distinctive clinical presentation, a diagnosis of epidermotropic B cell lymphoma was rendered. There was minimal bone marrow involvement based only on flow cytometric analysis, but there was no apparent bone marrow or splenic involvement on routine light microscopic assessment. Remission was achieved with single agent rituximab chemotherapy and the patient remained symptom free. The neoplastic CD20 positive epidermotropic B lymphocytes expressed CXCR3. Similar to the prior reported cases by the authors, the neoplastic cells expressed CXCR3, a chemokine whose organ and tissue specific ligands could contribute to its relatively indolent clinical course."

Luminari, S., Merli, M., Rattotti, S., Tarantino, V., Marcheselli, L., Cavallo, F., Varettoni, M., Bianchi, B., Merli, F., Tedeschi, A., Cabras, G., Re, F., Visco, C., Torresan Delamain, M., Cencini, E., Spina, M., Ferrero, S., Ferrari, A., Deodato, M., Mannina, D., Annibali, O., Rago, A., Orsucci, L., Defrancesco, I., Frigeni, M., Cesaretti, M. & Arcaini, L. 2019.

"Marginal zone lymphomas (MZLs) are indolent nonfollicular B-cell lymphomas (INFLs) and have heterogeneous clinical behavior. Recently, time to progression of disease at 24 months (POD24) was identified to stratify overall survival (OS) in follicular non-Hodgkin lymphoma and in INFL. Here, we examined the ability of POD24 to predict subsequent OS in a large, international cohort of MZL as part of the NF10 prospective international registry headed by Fondazione Italiana Linfomi (FIL). POD24 was only calculated for MZL patients requiring immediate therapy and was defined as experiencing lymphoma progression within 24 months from diagnosis. Among the 1325 patients enrolled in the NF10 study, we identified 321 patients with MZL for whom

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019

Page 2

immediate therapy was planned right after lymphoma diagnosis. Overall, POD24 was confirmed in 59 patients (18%). Three-year OS for patients with POD24 was 53% with a hazard ratio of 19.5 (95% confidence interval, 8.4-45) compared with patients without POD24 (3-year OS, 95%). Association of POD24 with OS was confirmed for the subgroup of splenic and extranodal MZLs. Assessment of POD24 stratifies subsequent outcome in MZL and identifies a high-risk population. This trial was registered at www.clinicaltrials.gov as [#NCT02904577](#)."

Incidence of Nodal Marginal Zone Lymphoma

The National Cancer Registry (2014) does not provide any information regarding the incidence of Nodal Marginal Zone Lymphoma.

Signs and Symptoms of Nodal Marginal Zone Lymphoma

The most common symptom is a painless swelling in the neck, armpit or groin. This is caused by the lymphoma cells building up in the lymph nodes, making them bigger.

Other symptoms may include:

- tiredness
- unexplained weight loss
- drenching night sweats
- high temperatures (fevers).

Diagnosis of Nodal Marginal Zone Lymphoma

Nodal Marginal Zone Lymphoma (NMZL) is frequently a diagnosis of exclusion, including reactive hyperplasia and indolent small B cell lymphoma. Morphologically, the tumour cells surround reactive follicles and expand into the interfollicular areas. Follicular colonisation may be present. In cases with a diffuse pattern, follicle remnants may be detected with stains for follicular dendritic cells and germinal centre markers.

In paediatric NMZL, the tumour is similar to that seen in adults except that there are often progressively transformed germinal centres in which the outer border of the follicles is disrupted and infiltrated by tumour cells.

Treatment of Nodal Marginal Zone Lymphoma

Experts do not really know how best to treat nodal marginal zone lymphoma. One may be offered one or more of the following treatments:

Watchful waiting (also called active surveillance) - may be an option because nodal marginal zone lymphoma develops slowly and may not need to be treated right away. The healthcare team will monitor the person with nodal marginal zone lymphoma carefully and start treatment when symptoms appear or there are signs that the disease is progressing more quickly.

Chemotherapy - Nodal marginal zone lymphoma usually responds well to chemotherapy. It may be given as a single drug or a combination of drugs and is often given with a targeted therapy drug.

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019

Targeted therapy - uses drugs to target specific molecules (such as proteins) on the surface of cancer cells. These molecules help send signals that tell cells to grow or divide. By targeting these molecules, the drugs stop the growth and spread of cancer cells while limiting harm to normal cells.

Immunotherapy - helps to strengthen or restore the immune system's ability to fight cancer.

Radiation therapy - External beam radiation therapy may be given to the lymph nodes to treat early stage (stage 1 or stage 2) nodal marginal zone lymphoma. It is used when this type of lymphoma affects only 1 or 2 areas of lymph nodes in the body.

De Luca, M.L., Lomnardi, L., Tartaglia, G., Fazio, F., Di Prima, A., Serrao, A., Canichella, M. & Pulsoni, A. 2019. "Many epidemiological, biological and therapeutic studies have extensively investigated the etiological link between HCV infection and B-cell Non-Hodgkin Lymphoma (NHL). Large experiences in the literature demonstrated HCV-related indolent NHL regression after antiviral therapy. While the response to interferon-ribavirin-based antiviral therapy is well documented, evidence of the efficacy of interferon-free Direct-Acting Antivirals (DAAs) in this subset of lymphoma is also currently increasing. Splenic and Nodal Marginal zone Lymphoma (MZL) are frequently associated with HCV chronic infection. In this article we report two cases of HCV-related MZL with an unusual presentation, subcutaneous "lipoma-like" nodules, treated with DAAs. Both patients, a 59-years-old woman and a 72-years-old man, were affected by HCV chronic infection since several years and were referred to our Institute for a diagnosis of MZL with subcutaneous presentation. Given the possible etiological link with HCV infection, both patients were treated with DAAs. A Sustained virological response (SVR) was reached after few weeks of therapy and at the end of treatment a clinically and radiologically documented reduction of MZL localizations, persisting to date, were obtained in both patients. The two clinical cases presented in this article confirm the efficacy of DAA's as first-line treatment in HCV related NHL, also in this rare entity of MZL with subcutaneous presentation."

Ayyappan, S. & William, B.M. 2018.

PURPOSE OF REVIEW: The purpose of the study is to summarize the current conundrums in the management of marginal zone lymphomas (MZL).

RECENT FINDINGS: In 2017, the US Food and Drug Administration (FDA) approved ibrutinib, a first in class Bruton Tyrosine Kinase inhibitor, for the treatment of relapsed/refractory MZL based on pivotal open-label phase II trial demonstrating an overall response rates of 48%. Clinical trials design utilizing chemotherapy-free regimens for relapsed/refractory disease are gaining popularity. Recent studies have identified multiple genetic biomarkers that helped characterize and prognosticate different subtypes of MZL. MZLs are heterogeneous, mostly indolent, malignancies derived from B lymphocytes. Three disease subtypes are recognized, extranodal, nodal, and splenic. The disease characteristics, clinical picture, and treatment algorithms vary considerably based on subtype and site of involvement. Recent discoveries have enhanced our knowledge of the pathogenesis of MZLs leading to development of more accurate prognostic models as well as novel targeted systemic therapies.

About Clinical Trials

Clinical trials are research studies that involve people. They are conducted under controlled conditions. Only about 10% of all drugs started in human clinical trials become an approved drug.

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019

Clinical trials include:

- Trials to test effectiveness of new treatments
- Trials to test new ways of using current treatments
- Tests new interventions that may lower the risk of developing certain types of cancers
- Tests to find new ways of screening for cancer

The [South African National Clinical Trials Register](#) provides the public with updated information on clinical trials on human participants being conducted in South Africa. The Register provides information on the purpose of the clinical trial; who can participate, where the trial is located, and contact details.

For additional information, please visit: www.sanctr.gov.za/

Medical Disclaimer

This Fact Sheet is intended to provide general information only and, as such, should not be considered as a substitute for advice, medically or otherwise, covering any specific situation. Users should seek appropriate advice before taking or refraining from taking any action in reliance on any information contained in this Fact Sheet. So far as permissible by law, the Cancer Association of South Africa (CANSA) does not accept any liability to any person (or his/her dependants/estate/heirs) relating to the use of any information contained in this Fact Sheet.

Whilst the Cancer Association of South Africa (CANSA) has taken every precaution in compiling this Fact Sheet, neither it, nor any contributor(s) to this Fact Sheet can be held responsible for any action (or the lack thereof) taken by any person or organisation wherever they shall be based, as a result, direct or otherwise, of information contained in, or accessed through, this Fact Sheet.



Sources and References Consulted and/or Utilised

Ayyappan, S. & William, B.M. 2018. Marginal zone lymphoma: clinicopathologic variations and approaches to therapy. *Curr Oncol Rep.* 2018 Mar 23;20(4):33. doi: 10.1007/s11912-018-0687-9.

Bron, D., Meuleman, N. & Eurobloodnet for rare diseases and EHA SWG 'Aging and Hematology'. 2019. Marginal zone lymphomas: second most common lymphomas in older patients. *Curr Opin Oncol.* 2019 Sep;31(5):386-393. doi: 10.1097/CCO.0000000000000554.

De Luca, M.L., Lomnardi, L., Tartaglia, G., Fazio, F., Di Prima, A., Serrao, A., Canichella, M. & Pulsoni, A. 2019. Response to Interferon-free Direct Antivirals (DAAS) treatment in HCV-related subcutaneous marginal zone B-cell lymphoma with lipoma-like presentation: report of two cases. *Mediterr J Hematol Infect Dis.* 2019 Sep 1;11(1):e2019053. doi: 10.4084/MJHID.2019.053. eCollection 2019.

Luminari, S., Merli, M., Rattotti, S., Tarantino, V., Marcheselli, L., Cavallo, F., Varettoni, M., Bianchi, B., Merli, F., Tedeschi, A., Cabras, G., Re, F., Visco, C., Torresan Delamain, M., Cencini, E., Spina, M., Ferrero, S., Ferrari, A., Deodato, M., Mannina, D., Annibali, O., Rago, A., Orsucci, L., Defrancesco, I., Frigeni, M., Cesaretti, M. & Arcaini, L. 2019. Early progression as a predictor of survival in marginal zone lymphomas: an analysis from the FIL-NF10 study. *Blood.* 2019 Sep 5;134(10):798-801. doi: 10.1182/blood.2019001088. Epub 2019 Jul 10.

Magro, C.M., Momtahn, S., Coleman, M. & Grossman, M.E. 2019. Epidermotropic CXCR3 positive marginal zone lymphoma: a distinctive clinical histopathological entity potentially originating in the skin: it does not always indicate splenic marginal zone lymphoma. *Dermatol Online J.* 2019 Jul 15;25(7). pii: 13030/qt4207n83g.

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019

Page 5

NMLZ

<https://www.oncologynurseadvisor.com/home/cancer-types/lymphoma/risk-factors-for-higher-grade-transformation-in-marginal-zone-lymphomas-identified/>

Nodal Marginal Zone Lymphoma

<https://lymphoma-action.org.uk/types-lymphoma-non-hodgkin-lymphoma/nodal-marginal-zone-lymphoma>

<https://www.uptodate.com/contents/nodal-marginal-zone-lymphoma>

<http://atlasgeneticsoncology.org/Anomalies/NMZLID2145.html>

<https://www.macmillan.org.uk/information-and-support/lymphoma/lymphoma-non-hodgkin/understanding-cancer/types-of-non-hodgkin-lymphoma/nodal-marginal-zone-b-cell-lymphoma.html>

<https://www.cancer.ca/en/cancer-information/cancer-type/non-hodgkin-lymphoma/non-hodgkin-lymphoma/more-types-of-nhl/nodal-marginal-zone-lymphoma/?region=on>

Researched and Authored by Prof Michael C Herbst

[D Litt et Phil (Health Studies); D N Ed; M Art et Scien; B A Cur; Dip Occupational Health; Dip Genetic Counselling; Dip Audiometry and Noise Measurement; Diagnostic Radiographer; Medical Ethicist]

Approved by Ms Elize Joubert, Chief Executive Officer [BA Social Work (cum laude); MA Social Work]

October 2019