Epidemiology of Hematologic Malignancies in Real-World Settings: Findings From the Hemato-Oncology Latin America Observational Registry Study

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PURPOSE Limited information is available on multiple myeloma (MM), chronic lymphocytic leukemia (CLL), and non-Hodgkin lymphoma (NHL) management in Latin America. The primary objective of the Hemato-Oncology Latin America (HOLA) study was to describe patient characteristics and treatment patterns of Latin American patients with MM, CLL, and NHL.

METHODS This study was a multicenter, retrospective, medical chart review of patients with MM, CLL, and NHL in Latin America identified between January 1, 2006, and December 31, 2015. Included were adults with at least 1 year of follow-up (except in cases of death within 1 year of diagnosis) treated at 30 oncology hospitals (Argentina, 5; Brazil, 9; Chile, 1; Colombia, 5; Mexico, 6; Panama/Guatemala, 4).

RESULTS Of 5,140 patients, 2,967 (57.7%) had NHL, 1,518 (29.5%) MM, and 655 (12.7%) CLL. Median follow-up was 2.2 years for MM, 3.0 years for CLL, and 2.2 years for NHL, and approximately 26% died during the study observation period. Most patients had at least one comorbidity at diagnosis. The most frequent induction regimen was thalidomide-based chemotherapy for MM and chlorambucil with or without prednisone for CLL. Most patients with NHL had diffuse large B-cell lymphoma (DLBCL; 49.1%) or follicular lymphoma (FL; 19.5%). The majority of patients with DLBCL or FL received rituximab plus cyclophosphamide, doxorubicin, vincristine, and prednisone.

CONCLUSION The HOLA study generated an unprecedented level of high-quality, real-world evidence on characteristics and treatment patterns of patients with hematologic malignancies. Regional disparities in patient characteristics may reflect differences in ethnoracial identity and level of access to care. These data provide needed real-world evidence to understand the disease landscape in Latin America and may be used to inform clinical and health policy decision making.

INTRODUCTION Hematologic malignancies (HMs) originate from uncontrolled growth of hematopoietic and lymphoid tissues. These biologically and clinically heterogeneous disorders account for 6.5% of all cancers around the world, including approximately 9.0% in the United States and Europe. Less is known about HM epidemiology in Latin America (Central and South America). The region has witnessed a considerable increase in the amount of hematology and oncology research; however, most of the contributions were observed in just a few countries (Brazil, Argentina, Mexico, Peru, Chile, and Uruguay). In the multinational CONCORD program, which estimates survival from cancer in 1.9 million adults from 101 population-based cancer registries in 31 countries on 5 continents, only 2 of the participating countries were located in Latin America. Moreover, according to the WHO, only 8% of Latin American populations are covered by cancer registries. Real-world data on HMs in Latin America are needed to inform decisions about patient care and health
CONTEXT

Key Objective
Despite improved assessment and management of multiple myeloma, chronic lymphocytic leukemia (CLL), and non-Hodgkin lymphoma (NHL) around the world, there is a paucity of high-quality, real-world data on these diseases in Latin America. The aim of the HOLA study is to investigate patient and treatment characteristics to understand the unmet needs in cancer care in Latin America.

Knowledge Generated
More than 5,000 patients were included from 7 Latin American countries, with the median age at diagnosis being slightly different from other regions’ reports (range, 57 years in patients with NHL to 67 years in patients with CLL). Most patients had comorbid conditions, and there was considerable regional heterogeneity in characteristics related to both patients and their management.

Relevance
Findings from this study can fill an important gap for physicians, patients, and health authorities in Latin America for the management of hematologic malignancies and can be used to understand the disease landscape in this region for further improvement of patient care.

METHODS

In this multicenter, observational study, the medical records of Latin American patients with CLL, MM, or NHL were retrospectively reviewed (ClinicalTrials.gov identifier: NCT02559583). Adults with HMs were studied at 30 oncology specialty hospitals in 7 countries: Argentina (4 private, 1 public setting), Brazil (1 private, 8 public settings), Chile (1 public setting), Colombia (5 private settings), Mexico (1 private, 5 public settings), Panama (1 private, 2 public settings), and Guatemala (1 public setting). The hospitals were selected based on their experience in providing clinical treatment to patients with CLL, MM, or NHL; geographical representation; the type of practice; interest in participating in the study; and fulfillment of the study requirements. All study sites were eligible to include all 3 types of cancer.

Patient population inclusion criteria were incident or prevalent CLL, MM, or NHL diagnosed between January 1, 2006, and December 31, 2015; age ≥ 18 years at the time of first observed diagnosis of these HMs (whether incident or prevalent); ≥ 1 year of patient data after first observed diagnosis (except in the event of patient death within 1 year of being diagnosed); and ability and willingness to provide informed consent (except if a waiver of informed consent was obtained). There were no exclusion criteria.

The study was conducted under the Guidelines for Good Pharmacoepidemiology Practices issued by the International Society for Pharmacoepidemiology, the Declaration of Helsinki and its amendments, and any applicable national guidelines. Each study was approved by its local ethics committee and followed local country requirements (Brazil, approval by central ethics committee; Colombia, notification to Instituto Nacional de Vigilancia de Medicamentos y Alimentos; Argentina, approval by Dirección Nacional de Protección de Datos Personales; Mexico, approval by Secretaría de Salud/COFEPRIS, Comisión Nacional de Investigación Científicas; and Guatemala, notification to Ministerio de Salud Publica).

Each study site selected and prepared medical records for chart abstractors to review, and the principal investigator provided clarification in cases of doubt. The study employed chart abstractors with previous experience in performing medical record reviews, and the chart abstractors were centrally trained by Janssen Cilag personnel on the electronic data capture system, study protocol, and how to complete the electronic case report forms (CRFs). The data fields, such as laboratory results, diagnosis specification, and other clinical criteria in the CRFs, had predefined ranges in clickable fields as opposed to free-text fields. This was designed to ensure that the abstractor looked only for the required information in the medical chart and selected the field that contained the specific range in the CRF where the data fit. Each trained chart abstractor signed a training form to document training module completed and date of training. All training materials and records were available for review, and future chart abstractors were required to undergo the same training.under Source document verification visits
were performed at selected sites, which represented 60% of all study sites. Demographic data, including sex, country of residence, age at diagnosis, and age category (<30, 30 to <40, 40 to <50, 50 to <60, 60 to <70, 70 to <80, ≥80 years) were collected. Clinical data included comorbidity status, disease stage, Eastern Cooperative Oncology Group (ECOG) performance status (PS), and first-contact physician type at diagnosis. Disease stage at diagnosis was based on International Staging System (ISS) for MM, Ann Arbor staging for NHL, and Binet/Rai stage for CLL. The subtype of NHL was also recorded. Treatment data included the time of treatment initiation and the treatment regimen of the induction therapy. Information about a patient's transplantation history was also collected from patients diagnosed with MM.

Descriptive statistical analyses were stratified by cancer type and country. Patient characteristics and treatment patterns were summarized using descriptive statistics, including measures of central tendency (mean, median) and spread (variance, range, minimum, maximum) for continuous variables (eg, age at diagnosis) and frequency distributions (number, %) for categorical variables (eg, sex). Because of low overall enrollment, data from Panama and Guatemala were pooled for analysis. Treatment characteristics of patients with MM were further stratified by their history of transplantation (yes/no). Statistical analyses were conducted using SAS 9.2 software (SAS Institute, Cary, NC).

RESULTS
Of a total of 5,140 patients, 2,967 (57.7%) had NHL, 1,518 (29.5%) had MM, and 655 (12.7%) had CLL. Among the original 5,140 patients, 2,715 (52.8%) were still being followed at the study end date (Fig 1).

MM
Among 1,518 initially eligible individuals with MM, 713 (47.0%) were still being followed at the time of study end.

**FIG 1.** Disposition of (A) all patients and those with (B) multiple myeloma (MM), (C) chronic lymphocytic leukemia (CLL), and (D) non-Hodgkin lymphoma (NHL).
with a median follow-up of 2.2 years (range, < 0.1-12.1 years). One third of the patients with MM (506; 33.3%) died during the observation period (range, 18.4% in Panama/Guatemala to 54.4% in Brazil; Table 1). The median age at diagnosis was 61 years (range, 23-91 years). The most frequent age categories were 60 to < 70 years (31.1%) and 50 to < 60 years (28.3%). There was a slight predominance of males (50.7%) over females (49.3%), and the sex balance varied across regions (Table 2).

MM was highly comorbid with a range of chronic and other conditions, with approximately 53% of patients with one or two comorbidities and approximately 15% with more than two comorbidities. Hypertension was reported most frequently in 36.5% of patients (range, 28.2% in Argentina to 49.2% in Brazil). Other frequent comorbidities were diabetes and heart disease in 12.6% and 11.5% of patients, respectively. Potential manifestations of target organ damage included a history of bone disorders in 7.2% of patients and renal disease in 11.1% (Table 2).

At diagnosis, patients had an ECOG PS of 0 (27.2%), 1 (31.7%), or 2 (20.4%). Most patients were ISS stage II (29.4%) or III (48.8%). Stage III MM was the most frequently observed disease stage in Mexico (62.5%), Chile (60.0%), Brazil (49.3%), and Colombia (45.8%). Most Panamanian/Guatemalan patients (57.1%) had stage II, and most Argentinean patients had stage I (38.2%). Of patients with MM, 35% had renal impairment, which was fairly constant across countries. Of all patients with MM,

| Characteristic | Panama/Guatemala (n = 178) | Chile (n = 571) | Argentina (n = 1,009) | Colombia (n = 918) | Mexico (n = 1,345) | Brazil (n = 1,119) | Total (N = 5,140) |
|---------------|----------------------------|----------------|-----------------------|--------------------|-------------------|-------------------|------------------|
| MM Patients   | 49 (27.5)                  | 99 (17.3)      | 284 (28.1)            | 338 (36.8)         | 364 (27.1)        | 384 (34.3)        | 1,518 (29.5)     |
| Duration of follow-up, * years | 44                     | 94             | 260                   | 327                | 350               | 367               | 1,442            |
| Median        | 2.16                       | 1.79           | 2.98                  | 1.92               | 2.38              | 2.06              | 2.21             |
| Minimum-maximum | 0.1-9.7                  | 0.1-11.7       | < 0.1-12.1            | < 0.1-9.4          | < 0.1-10.2        | < 0.1-9.7         | < 0.1-12.1       |
| Missing       | 5                          | 5              | 24                    | 11                 | 14                | 17                | 76               |
| Patients who died | 9 (18.4)               | 41 (41.4)      | 70 (24.6)             | 103 (30.5)         | 74 (20.3)         | 209 (54.4)        | 506 (33.3)       |
| Patients still being followed at end of study | 28 (57.1)          | 23 (23.2)      | 173 (60.9)            | 128 (37.9)         | 204 (56.0)        | 157 (40.9)        | 713 (47.0)       |
| CLL Patients  | 16 (9.0)                   | 19 (3.3)       | 121 (12.0)            | 87 (9.5)           | 138 (10.3)        | 274 (24.5)        | 655 (12.7)       |
| Duration of follow-up, * years | 16                     | 19             | 113                   | 76                 | 135               | 261               | 620              |
| Median        | 1.64                       | 3.43           | 2.99                  | 2.45               | 3.83              | 3.01              | 3.03             |
| Minimum-maximum | 0.9-8.6                  | 0.6-7.4        | 0.3-9.2               | < 0.1-8.1          | < 0.1-9.6         | 0.1-10.1          | < 0.1-10.1       |
| Missing       | 0                          | 0              | 8                     | 11                 | 3                 | 13                | 35               |
| Patients who died | 2 (12.5)                | 4 (21.1)       | 12 (9.9)              | 15 (17.2)          | 12 (8.7)          | 85 (31.0)         | 130 (19.8)       |
| Patients still being followed at end of study | 12 (75.0)          | 14 (73.7)      | 104 (86.0)            | 43 (49.4)          | 90 (65.2)         | 168 (61.3)        | 431 (65.8)       |
| NHL Patients  | 113 (63.5)                 | 453 (79.3)     | 604 (59.9)            | 493 (53.7)         | 843 (62.7)        | 461 (41.2)        | 2,967 (57.7)     |
| Duration of follow-up, * years | 104                   | 442            | 577                   | 471                | 796               | 431               | 2,821            |
| Median        | 2.76                       | 2.04           | 3.07                  | 1.77               | 1.97              | 2.40              | 2.20             |
| Minimum-maximum | < 0.1-9.9                | < 0.1-8.8      | < 0.1-11.8            | < 0.1-8.4          | < 0.1-10.1        | < 0.1-9.7         | < 0.1-11.8       |
| Missing       | 9                          | 11             | 27                    | 22                 | 47                | 30                | 146              |
| Patients who died | 6 (5.3)                   | 60 (13.2)      | 120 (19.9)            | 130 (26.4)         | 176 (20.9)        | 199 (43.2)        | 691 (23.3)       |
| Patients still being followed at end of study | 90 (79.6)          | 173 (38.2)     | 440 (72.8)            | 191 (38.7)         | 451 (53.5)        | 226 (49.0)        | 1,571 (52.9)     |

Abbreviations: CLL, chronic lymphocytic leukemia; MM, multiple myeloma; NHL, non-Hodgkin lymphoma.

*Follow-up defined as patients who were lost to follow-up or with ongoing follow-up from diagnosis to last consult date; for patients who died, from diagnosis to date of death.
### TABLE 2. Demographics, Baseline Clinical Characteristics, and Treatment Patterns of Patients With Multiple Myeloma

| Characteristic | Panama/Guatemala (n = 49) | Chile (n = 99) | Argentina (n = 284) | Colombia (n = 338) | Mexico (n = 364) | Brazil (n = 384) | Total (n = 1,518) |
|----------------|---------------------------|---------------|---------------------|-------------------|-----------------|-----------------|------------------|
| **Age at diagnosis, years** | | | | | | | |
| Mean (SD) | 57.4 (13.0) | 63.8 (11.0) | 58.9 (11.1) | 61.7 (11.9) | 59.7 (11.2) | 61.8 (10.8) | 60.7 (11.4) |
| Median (range) | 57 (29-85) | 65 (27-85) | 60 (25-91) | 61 (23-91) | 60 (28-88) | 62 (27-90) | 61 (23-91) |
| **Age category at diagnosis, years** | | | | | | | |
| < 50 | 15 (30.6) | 11 (11.1) | 60 (21.1) | 50 (14.8) | 66 (18.2) | 55 (14.3) | 257 (16.9) |
| 50 to < 60 | 14 (28.6) | 24 (24.2) | 81 (28.5) | 101 (29.9) | 112 (30.9) | 97 (25.3) | 429 (28.3) |
| 60 to < 70 | 10 (20.4) | 31 (31.3) | 101 (35.6) | 88 (26.0) | 107 (29.5) | 135 (35.2) | 472 (31.1) |
| 70 to < 80 | 7 (14.3) | 30 (30.3) | 29 (10.2) | 78 (23.1) | 68 (18.7) | 78 (20.3) | 290 (19.1) |
| ≥ 80 | 3 (6.1) | 3 (3.0) | 13 (4.6) | 21 (6.2) | 10 (2.8) | 19 (4.9) | 69 (4.5) |
| **Sex** | | | | | | | |
| Male | 31 (63.3) | 48 (48.5) | 135 (47.5) | 158 (46.7) | 195 (53.6) | 202 (52.6) | 769 (50.7) |
| Female | 18 (36.7) | 51 (51.5) | 149 (52.5) | 180 (53.3) | 169 (46.4) | 182 (47.4) | 749 (49.3) |
| **No. of comorbidities at diagnosis** | | | | | | | |
| 0 | 17 (34.7) | 14 (14.1) | 102 (35.9) | 82 (24.3) | 177 (48.6) | 4 (24.5) | 486 (32.0) |
| 1-2 | 28 (57.1) | 61 (61.6) | 146 (51.4) | 194 (57.4) | 155 (42.6) | 225 (58.6) | 809 (53.3) |
| > 2 | 4 (8.2) | 24 (24.2) | 36 (12.7) | 62 (18.3) | 32 (8.8) | 65 (16.9) | 223 (14.7) |
| **Most frequent comorbidities at diagnosis** | | | | | | | |
| Hypertension | 17 (34.7) | 41 (41.4) | 80 (28.2) | 113 (33.4) | 114 (31.3) | 189 (49.2) | 554 (36.5) |
| Diabetes | 7 (14.3) | 20 (20.2) | 15 (5.3) | 31 (9.2) | 58 (15.9) | 61 (15.9) | 192 (12.6) |
| Heart disease | 5 (10.2) | 14 (14.1) | 45 (15.8) | 40 (11.8) | 28 (7.7) | 43 (11.2) | 175 (11.5) |
| Renal | 5 (10.2) | 22 (22.2) | 18 (6.3) | 39 (11.5) | 13 (3.6) | 71 (18.5) | 168 (11.1) |
| Bone | 4 (8.2) | 3 (3.0) | 24 (8.5) | 52 (15.4) | 6 (1.6) | 20 (5.2) | 109 (7.2) |
| **ECOG PS at diagnosis** | | | | | | | |
| 0 | 5 (41.7) | 20 (28.2) | 7 (53.8) | 15 (27.3) | 9 (12.5) | 16 (38.1) | 72 (27.2) |
| 1 | 1 (6.3) | 28 (39.4) | 3 (23.1) | 24 (43.6) | 16 (22.2) | 12 (28.6) | 84 (31.7) |
| 2 | 3 (25.0) | 10 (14.1) | 0 (0.0) | 11 (20.0) | 24 (33.3) | 6 (14.3) | 54 (20.4) |
| 3 | 3 (25.0) | 10 (14.1) | 2 (15.4) | 2 (3.6) | 16 (22.2) | 7 (16.7) | 40 (15.1) |
| ≥ 4 | 0 (0.0) | 3 (4.2) | 1 (7.7) | 3 (5.5) | 7 (9.7) | 1 (2.4) | 15 (5.7) |
| Not available | 37 | 28 | 271 | 283 | 292 | 342 | 1,253 |
| **ISS at diagnosis** | | | | | | | |
| I | 1 (7.1) | 18 (25.7) | 58 (38.2) | 31 (18.7) | 28 (14.0) | 39 (19.4) | 175 (21.8) |
| II | 8 (57.1) | 10 (14.3) | 49 (32.2) | 59 (35.5) | 47 (23.5) | 63 (31.3) | 236 (29.4) |
| III | 5 (35.7) | 42 (60.0) | 45 (29.6) | 76 (45.8) | 125 (62.5) | 99 (49.3) | 392 (48.8) |
| NE | 2 | 0 | 2 | 1 | 0 | 1 | 6 |
| Not available | 33 | 29 | 130 | 171 | 164 | 182 | 709 |
| **Most frequent first-contact physician** | | | | | | | |
| Hematologist | 9 (18.8) | 25 (25.3) | 101 (35.6) | 121 (35.8) | 45 (12.4) | 76 (19.8) | 377 (24.9) |
| Internist | 6 (12.5) | 28 (28.3) | 27 (9.5) | 32 (9.5) | 99 (27.2) | 42 (10.9) | 234 (15.4) |
| General physician | 7 (14.6) | 9 (9.1) | 60 (21.1) | 45 (13.3) | 33 (9.1) | 79 (20.6) | 233 (15.4) |

| Patients who received a transplantation | | | | | | | |
| No. of patients who received chemotherapy | 15 | 3 | 196 | 132 | 48 | 103 | 497 |
| Time from initial diagnosis to treatment, days | 12 | 2 | 144 | 102 | 24 | 88 | 372 |
| Mean (SD) | 245.7 (806.70) | 28.0 (26.87) | 117.3 (310.72) | 42.0 (91.17) | 67.0 (243.56) | 68.0 (142.40) | 85.4 (262.80) |
| Median (range) | 6.0 (0-2806) | 28.0 (9-47) | 28.5 (0-284) | 7.5 (0-467) | 6.0 (0-1201) | 21.5 (0-849) | 18.0 (0-2806) |
| Missing | 3 | 1 | 52 | 30 | 24 | 15 | 125 |

(Continued on following page)
The first physician contact was with hematologists in 24.9% followed by 15.4% each with internists and general physicians.

In the entire cohort, 497 patients with MM (32.7%) underwent autologous hematopoietic stem-cell transplantation (ASCT); however, the proportion of patients who underwent transplantation varied among countries. The highest rate was 69.0% in Argentina, and the lowest was 3.0% in Chile. Median time from diagnosis to initial treatment was 18.0 days (range, 6.0 days in Panama/Guatemala and Mexico to 28.5 days in Argentina). The 497 transplantation recipients who received induction chemotherapy were treated predominantly with thalidomide-based (151; 30.4%) and bortezomib-based (125; 25.2%) regimens. In these patients, median time from diagnosis to initial treatment was 15.0 days (range, 4.0 days in Colombia to 58.0 days in Chile). Most patients with MM who did not undergo transplantation received thalidomide-based chemotherapy (308 of 975; 31.6%) or melphalan, thalidomide, and steroid treatment (164 of 975; 16.8%) at induction. Thalidomide-based chemotherapy was the preferred treatment in Argentina (28.2%), Mexico (49.5%), and Brazil (28.3%).

**CLL**

Among 655 initially eligible individuals with CLL, 431 (65.8%) were still being followed at the time of study end, with a median follow-up of 3 years (< 0.1-10.1 years; Table 1). The proportion of patients with CLL who died during the observation period was 19.8% (range, 8.7% in Mexico to 31.0% in Brazil). Median age at diagnosis was 67 years (range, 23-96 years), which was relatively consistent across countries (Table 2). The most frequent age categories were 60 to < 70 years (34.6%) and 70 to < 80 years (29.1%). There was a slight predominance of male (54.2%) over female (45.8%) patients with CLL.

A total of 468 (71.5%) of the 655 patients with CLL had comorbidities at diagnosis (Table 3). The most frequent comorbidity was hypertension observed in 46.1% (range, 37.5% in Panama/Guatemala to 73.7% in Chile); other common comorbidities were heart disease (15.7%) and diabetes (15.1%).

Approximately two thirds of patients with CLL presented with Binet stage A (66.5%) and/or Rai stage 0 or 1 (63.0%). Most patients (92.3%) had an ECOG PS of 0 or 1 (range, 81.3% in Chile to 93.6% in Brazil) at diagnosis. Compared with other specialties, the first physician contacted most frequently was with a hematologist (33.8%; range, 18.1% in Mexico to 84.2% in Chile). Other physicians were general physicians (23.6%) and internists (13.8%).

Of the 655 patients with CLL, 415 (63.4%) received chemotherapy. The most common initial treatment of CLL was chlorambucil with or without prednisone (191; 46.0%) followed by fludarabine, cyclophosphamide, and rituximab (FCR; 87; 21.0%) and fludarabine and cyclophosphamide (34; 8.2%). The majority of patients in Brazil (108 of 181; 59.7%) and Mexico (57 of 95; 60.0%) received chlorambucil with or without prednisone. FCR was the most frequent initial chemotherapy regimen in Argentina (53.6%), Panama/Guatemala (50.0%), Colombia (40%), and Chile (38.5%).

**Table 2.** Demographics, Baseline Clinical Characteristics, and Treatment Patterns of Patients With Multiple Myeloma (Continued)

| Characteristic                          | Panama/Guatemala (n = 49) | Chile (n = 99) | Argentina (n = 284) | Colombia (n = 338) | Mexico (n = 364) | Brazil (n = 384) | Total (n = 1,518) |
|----------------------------------------|---------------------------|---------------|---------------------|-------------------|-----------------|----------------|-----------------|
| Three most-used chemotherapy treatments |                           |               |                     |                   |                 |                |                 |
| Thalidomide based                       | 1 (6.7)                   | 1 (33.3)      | 91 (46.4)           | 12 (9.1)          | 15 (31.3)       | 31 (30.1)       | 151 (30.4)      |
| Bortezomib based                        | 11 (73.3)                 | 0 (0.0)       | 40 (20.4)           | 18 (13.6)         | 6 (12.5)        | 5 (4.9)         | 125 (25.2)      |
| CTD                                    | 0 (0.0)                   | 1 (33.3)      | 12 (6.1)            | 3 (2.3)           | 6 (12.5)        | 31 (30.1)       | 53 (10.7)       |
| Patients who did not receive a transplantation |                   |               |                     |                   |                 |                |                 |
| No. of patients who received chemotherapy | 33                        | 88            | 78                   | 200               | 307             | 269            | 975             |
| Time from initial diagnosis to treatment, days | 19                        | 66            | 65                   | 154               | 243             | 233            | 780             |
| Mean (SD)                              | 72.1 (155.70)             | 109.2 (160.96)| 56.6 (121.67)       | 28.7 (75.89)      | 34.6 (112.73)   | 62.5 (113.10)  | 50.8 (115.53)   |
| Median (range)                         | 16.0 (1.0-680.0)          | 58.0 (0-869.0)| 23.0 (0-838.0)      | 4.0 (0-536.0)     | 9.0 (0-1461.0)  | 20.0 (0-784.0) | 15.0 (0-1461.0) |
| Missing                                 | 14                        | 22            | 13                   | 46                | 64              | 36             | 195             |
| Three most-used chemotherapy treatments |                           |               |                     |                   |                 |                |                 |
| Thalidomide based                       | 3 (9.1)                   | 26 (9.5)      | 22 (28.2)           | 29 (14.5)         | 152 (49.5)      | 76 (28.3)       | 308 (31.6)      |
| Melphalan and thalidomide plus steroid  | 1 (3.0)                   | 14 (15.9)     | 21 (26.9)           | 27 (13.5)         | 52 (16.9)       | 49 (18.2)       | 164 (16.8)      |
| Melphalan plus steroid                  | 1 (3.0)                   | 18 (20.5)     | 6 (7.7)             | 10 (5.0)          | 17 (5.5)        | 36 (13.4)       | 88 (9.0)        |

Abbreviations: CTD, cyclophosphamide, thalidomide, and dexamethasone; ECOG PS, Eastern Cooperative Oncology Group performance status; ISS, International Staging System; NE, not estimated; SD, standard deviation.
| Characteristic | Panama/Guatemala (n = 16) | Chile (n = 19) | Argentina (n = 121) | Colombia (n = 87) | Mexico (n = 138) | Brazil (n = 274) | Total (n = 655) |
|---------------|---------------------------|----------------|---------------------|------------------|----------------|----------------|----------------|
| Age at diagnosis, year | 16 | 19 | 121 | 87 | 137 | 273 | 653 |
| Mean (SD) | 67.3 (10.4) | 66.3 (9.2) | 64.7 (10.4) | 66.1 (12.7) | 67.3 (10.0) | 65.9 (11.0) | 66.1 (10.9) |
| Median (range) | 67 (47-82) | 68 (46-88) | 65 (39-95) | 68 (33-91) | 67 (38-93) | 66 (23-96) | 67 (23-96) |
| Unknown | 0 | 0 | 0 | 0 | 1 | 1 | 2 |
| Age category at diagnosis, years | | | | | | | |
| < 50 | 1 (6.3) | 1 (5.3) | 6 (5.0) | 12 (13.8) | 7 (5.1) | 21 (7.7) | 48 (7.4) |
| 50 to < 60 | 2 (12.5) | 3 (15.8) | 31 (25.6) | 14 (16.1) | 25 (18.2) | 49 (17.9) | 124 (19.0) |
| 60 to < 70 | 7 (43.8) | 8 (42.1) | 46 (38.0) | 22 (25.3) | 45 (32.8) | 98 (35.9) | 226 (34.6) |
| 70 to < 80 | 4 (25.0) | 6 (31.6) | 28 (23.1) | 25 (28.7) | 47 (34.3) | 80 (29.3) | 190 (29.1) |
| ≥ 80 | 2 (12.5) | 1 (5.3) | 10 (8.3) | 14 (16.1) | 13 (9.5) | 25 (9.2) | 65 (10.0) |
| Unknown | 0 | 0 | 0 | 0 | 1 | 1 | 2 |
| Sex | | | | | | | |
| Male | 11 (68.8) | 5 (26.3) | 80 (66.1) | 48 (55.2) | 64 (46.4) | 147 (53.6) | 355 (54.2) |
| Female | 5 (31.3) | 14 (73.7) | 41 (33.9) | 39 (44.8) | 74 (53.6) | 127 (46.4) | 300 (45.8) |
| No. of comorbidities at diagnosis | | | | | | | |
| 0 | 5 (31.3) | 1 (5.3) | 36 (29.8) | 23 (26.4) | 46 (33.3) | 76 (27.7) | 187 (28.5) |
| 1-2 | 8 (50.0) | 11 (57.9) | 56 (46.3) | 52 (59.8) | 77 (55.8) | 161 (58.8) | 365 (55.7) |
| > 2 | 3 (18.8) | 7 (36.8) | 29 (24.0) | 12 (13.8) | 15 (10.9) | 37 (13.5) | 103 (15.7) |
| Most frequent comorbidities at diagnosis | | | | | | | |
| Hypertension | 6 (37.5) | 14 (73.7) | 52 (43.0) | 38 (43.7) | 56 (40.6) | 136 (49.6) | 302 (46.1) |
| Heart disease | 1 (6.3) | 1 (5.3) | 49 (40.5) | 9 (10.3) | 15 (10.9) | 28 (10.2) | 103 (15.7) |
| Diabetes | 3 (18.8) | 7 (36.8) | 12 (9.9) | 9 (10.3) | 31 (22.5) | 37 (13.5) | 99 (15.1) |
| ECOG PS at diagnosis | | | | | | | |
| 0 | 0 (0.0) | 11 (68.8) | 10 (71.4) | 3 (42.9) | 15 (75.0) | 34 (72.3) | 73 (70.2) |
| 1 | 0 (0.0) | 2 (12.5) | 3 (21.4) | 3 (42.9) | 5 (25.0) | 10 (21.3) | 23 (22.1) |
| 2 | 0 (0.0) | 2 (12.5) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 2 (4.3) | 4 (3.8) |
| 3 | 0 (0.0) | 1 (6.3) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 1 (2.1) | 2 (1.9) |
| ≥ 4 | 0 (0.0) | 0 (0.0) | 1 (7.1) | 1 (14.3) | 0 (0.0) | 0 (0.0) | 2 (1.9) |
| Not available | 16 | 3 | 107 | 80 | 118 | 227 | 551 |
| Binet staging at diagnosis | | | | | | | |
| A | 3 (60.0) | 6 (35.3) | 39 (73.6) | 49 (77.8) | 56 (70.0) | 115 (62.2) | 268 (66.5) |
| B | 0 (0.0) | 6 (35.3) | 8 (15.1) | 3 (4.8) | 9 (11.3) | 32 (17.3) | 58 (14.4) |
| C | 2 (40.0) | 5 (29.4) | 6 (11.3) | 11 (17.5) | 15 (18.8) | 38 (20.5) | 77 (19.1) |
| NE | 0 | 0 | 0 | 1 | 4 | 0 | 5 |
| Not available | 11 | 2 | 68 | 23 | 54 | 89 | 247 |
| Rai staging at diagnosis | | | | | | | |
| 0 | 4 (33.3) | 5 (27.8) | 23 (42.6) | 12 (37.5) | 40 (41.2) | 54 (33.1) | 138 (36.7) |
| 1 | 5 (41.7) | 3 (16.7) | 15 (27.8) | 8 (25.0) | 24 (24.7) | 44 (27.0) | 99 (26.3) |

(Continued on following page)
Among 2,967 eligible individuals with NHL, 1,571 (52.9%) were still being followed at the time of study end, with a median follow-up of 2.2 years (range, 0.1-11.8 years). The proportion of patients with NHL who died during the observation period was 23.3% (range, 5.3% in Panama/Guatemala to 43.2% in Brazil; Table 1). More than two thirds of patients with NHL (68.6%) had either diffuse large B-cell lymphoma (DLBCL; 1,457; 49.1%) or follicular lymphoma (FL; 578; 19.5%). This section focuses on these subpopulations. Additional descriptive data on individuals with mantle cell lymphoma and mucosa-associated lymphoid tissue (MALT) lymphoma are listed in Appendix Tables A1 and A2.

Median age at diagnosis was 58 years (range, 18-95 years) for DLBCL and 57 years (range, 18-92 years) for FL. These values showed considerable variation across countries. Sex was approximately evenly distributed in patients with DLBCL and slightly imbalanced toward females with FL (Tables 4 and 5).

Comorbidities were present in 59.1% of patients with DLBCL and 54.0% with FL. The most frequent comorbidity for patients with DLBCL was hypertension (29.0%) followed by diabetes (12.8%) and heart disease (8.0%). For patients with FL, the most frequent comorbidity was hypertension (27.7%) followed by diabetes (10.2%) and heart disease (6.6%).

At diagnosis, patients with DLBCL had an ECOG PS of 0 (46.9%) or 1 (32.0%). Approximately one third of patients with DLBCL (33.4%) who had available Ann Arbor disease staging data were classified as having stage IV; 24.4% had stage II disease, 24.3% had stage III, and 17.8% had stage I. A total of 29.4% of patients with FL had an ECOG PS, of whom 68.8% had a PS of 0 and 22.4% had a PS of 1. A total of 520 (90.0%) of 578 patients with FL had available staging data, with 33.1% having stage IV, 34.0% stage III, 19.6% stage II, and 13.3% stage I disease.

Among patients with DLBCL and FL, hematologists were most frequently the first physician contacted (28.9%).
TABLE 4. Demographics, Baseline Clinical Characteristics, and Treatment Patterns of Patients With Diffuse Large B-Cell Lymphoma

| Characteristic                        | Panama/ Guatemala (n = 52) | Chile (n = 244) | Argentina (n = 252) | Colombia (n = 266) | Mexico (n = 442) | Brazil (n = 201) | Total (n = 1,457) |
|---------------------------------------|---------------------------|-----------------|--------------------|-------------------|-----------------|-----------------|------------------|
| Age at diagnosis, years               | 51                        | 244             | 252                | 266               | 442             | 200             | 1,455           |
| Mean (SD)                             | 53.3 (16.3)               | 58.7 (15.7)     | 57.4 (15.0)        | 56.0 (16.3)       | 56.9 (15.1)     | 55.7 (16.5)     | 56.8 (15.7)     |
| Median (range)                        | 57 (18-81)                | 61 (18-95)      | 60 (22-87)         | 58 (18-88)        | 56 (21-92)      | 57 (20-90)      | 58 (18-95)      |
| Unknown                               | 1                         | 0               | 0                  | 0                 | 1               | 1               | 2               |
| Age category at diagnosis, years      |                           |                 |                    |                   |                 |                 |                  |
| < 50                                  | 19 (37.3)                 | 63 (25.8)       | 71 (28.2)          | 89 (33.4)         | 146 (33.0)      | 69 (30.0)       | 448 (30.8)      |
| 50 to < 60                            | 9 (17.6)                  | 47 (19.3)       | 53 (21.0)          | 52 (19.5)         | 102 (23.1)      | 51 (25.5)       | 314 (21.6)      |
| 60 to < 70                            | 16 (31.4)                 | 73 (29.9)       | 73 (29.0)          | 67 (25.2)         | 93 (21.0)       | 51 (25.5)       | 373 (25.6)      |
| 70 to < 80                            | 6 (11.8)                  | 47 (19.3)       | 44 (17.5)          | 45 (16.9)         | 72 (16.3)       | 22 (11.0)       | 236 (16.2)      |
| ≥ 80                                  | 1 (2.0)                   | 14 (5.7)        | 11 (4.4)           | 13 (4.9)          | 29 (6.6)        | 16 (8.0)        | 84 (5.8)        |
| Unknown                               | 1                         | 0               | 0                  | 0                 | 0               | 1               | 2               |
| Sex                                   |                           |                 |                    |                   |                 |                 |                  |
| Male                                  | 27 (51.9)                 | 126 (51.6)      | 134 (53.2)         | 137 (51.5)        | 210 (47.5)      | 86 (42.8)       | 720 (49.4)      |
| Female                                | 25 (48.1)                 | 118 (48.4)      | 118 (46.8)         | 129 (48.5)        | 232 (52.5)      | 115 (57.2)      | 737 (50.6)      |
| No. of comorbidities at diagnosis     |                           |                 |                    |                   |                 |                 |                  |
| 0                                     | 25 (48.1)                 | 77 (31.6)       | 91 (36.1)          | 73 (27.4)         | 242 (54.8)      | 88 (43.8)       | 596 (40.9)      |
| 1-2                                   | 24 (46.2)                 | 142 (58.2)      | 116 (46.0)         | 152 (57.1)        | 176 (39.8)      | 98 (48.8)       | 708 (48.6)      |
| > 2                                   | 3 (5.8)                   | 25 (10.2)       | 45 (17.9)          | 41 (15.4)         | 24 (5.4)        | 15 (7.5)        | 153 (10.5)      |
| Most frequent comorbidities at diagnosis |                       |                 |                    |                   |                 |                 |                  |
| Hypertension                          | 19 (36.5)                 | 91 (37.3)       | 73 (29.0)          | 85 (32.0)         | 93 (21.0)       | 62 (30.8)       | 423 (29.0)      |
| Diabetes                              | 7 (13.5)                  | 41 (16.8)       | 20 (7.9)           | 25 (9.4)          | 73 (16.5)       | 20 (10.0)       | 186 (12.8)      |
| Heart disease                         | 2 (3.8)                   | 13 (5.3)        | 49 (19.4)          | 23 (8.6)          | 15 (3.4)        | 15 (7.5)        | 117 (8.0)       |
| Other neoplasia                       | 0 (0.0)                   | 11 (4.5)        | 30 (11.9)          | 24 (9.0)          | 10 (2.3)        | 0 (0.0)         | 75 (5.1)        |
| ECOG PS at diagnosis                  |                           |                 |                    |                   |                 |                 |                  |
| 0                                     | 7 (100)                   | 114 (55.9)      | 33 (45.2)          | 25 (34.2)         | 38 (33.0)       | 25 (56.8)       | 242 (46.9)      |
| 1                                     | 0 (0.0)                   | 67 (32.8)       | 19 (26.0)          | 28 (38.4)         | 40 (34.8)       | 11 (25.0)       | 165 (32.0)      |
| 2                                     | 0 (0.0)                   | 14 (6.9)        | 12 (16.4)          | 13 (17.8)         | 19 (16.5)       | 5 (11.4)        | 63 (12.2)       |
| 3                                     | 0 (0.0)                   | 6 (2.9)         | 3 (4.1)            | 6 (8.2)           | 12 (10.4)       | 3 (6.8)         | 30 (5.8)        |
| ≥ 4                                   | 0 (0.0)                   | 3 (1.5)         | 6 (8.2)            | 1 (1.4)           | 6 (5.2)         | 0 (0.0)         | 16 (3.1)        |
| Not available                         | 45                        | 40              | 179                | 193               | 327             | 157             | 941             |
| Ann Arbor stage                       |                           |                 |                    |                   |                 |                 |                  |
| I                                     | 6 (17.6)                  | 47 (19.8)       | 36 (17.5)          | 33 (15.3)         | 77 (19.9)       | 17 (12.8)       | 216 (17.8)      |
| II                                    | 10 (29.4)                 | 68 (28.7)       | 47 (22.8)          | 56 (25.9)         | 86 (22.3)       | 29 (21.8)       | 296 (24.4)      |
| III                                   | 8 (23.5)                  | 56 (23.6)       | 61 (29.6)          | 49 (22.7)         | 91 (23.6)       | 30 (22.6)       | 295 (24.3)      |
| IV                                    | 10 (29.4)                 | 66 (27.8)       | 62 (30.1)          | 78 (36.1)         | 132 (34.2)      | 57 (42.9)       | 405 (33.4)      |
| NE                                    | 7                         | 0               | 40                 | 24                | 26              | 0               | 97              |
| Unknown                               | 11                        | 7               | 6                  | 26                | 30              | 68              | 148             |

(Continued on following page)
followed by internists/general physicians (21.0%) and surgeons (13.7%). Only 8%-12% of patients in Brazil and Panama/Guatemala first consulted with a hematologist.

For patients with DLBCL, the 2 most frequent regimens were rituximab plus cyclophosphamide, doxorubicin, vincristine, and prednisone (R-CHOP; 70.3%) and cyclophosphamide, doxorubicin, vincristine, and prednisone (CHOP; 14.6%). All other regimens were used by 5% of patients. Similar trends were observed across countries.

The majority of patients with FL received chemotherapy (546 of 578; 94.5%); the 3 most frequent regimens were R-CHOP (280 of 546; 51.3%), rituximab plus cyclophosphamide, vincristine, and prednisone (96 of 546; 17.6%), and CHOP (70 of 546; 12.8%). CHOP was the most frequently reported therapy in Chile (26 of 68; 38.2%) and Brazil (17 of 55; 30.9%).

**DISCUSSION**

To our knowledge, the HOLA registry—unprecedented in size and scope—provides high-quality, real-world data from > 5,000 patients with HMs observed for 8 years in Latin America. Despite considerable heterogeneity across the 7 participating countries, certain common themes are of potential concern to health care providers and policy-makers. First, large proportions of patients with HMs had comorbidities, which can limit therapeutic options. Second, considerable numbers of patients also had high-grade and other aggressive forms of disease (eg, DLBCL), and approximately 26% of patients died during the observation period.

Our study identified potential regional access-to-care issues. Between 28% and 50% of patients with MM received thalidomide-based chemotherapy in Argentina, Brazil, Mexico, and Chile, and 36% of those in Colombia received bortezomib. Furthermore, there was wide variation between countries in the proportion of patients with MM who had undergone transplantation. In Argentina, 20-fold more patients underwent ASCT than in Chile. The treatment regimens in Latin American were different compared with those of other countries in the world. For example, the majority of transplantation recipients with MM were administered thalidomide-based (30.4%) or bortezomib-based (25.2%) treatment in the studied regions, while lenalidomide, bortezomib, and dexamethasone chemotherapy is the most commonly used frontline treatment of transplantation-eligible patients with MM in the United States. Differences in treatment regimens were probably due to the restricted access and lack of approved novel agents in these studied regions.

With regard to comorbidities, these included hypertension (range across HMs, 29.0%-46.1%), diabetes (12.6%-15.1%), and heart disease (8.0%-15.7%). In one study, patients with MM who received treatment with thalidomide experienced a 23% increase in grade 3-4 adverse events, including, most frequently, cardiovascular disease followed by other hematologic conditions, thromboembolic events, infection, and neuropathy. 

A Latin American observational study found the median age of patients with MM to be 67.4 years, and 70.7%...
| Characteristic                              | Panama/Guatemala (n = 21) | Chile (n = 75) | Argentina (n = 185) | Colombia (n = 62) | Mexico (n = 175) | Brazil (n = 60) | Total (n = 578) |
|--------------------------------------------|---------------------------|---------------|---------------------|-------------------|-----------------|----------------|-----------------|
| **Age at diagnosis, years**                | 20                        | 75            | 185                 | 62                | 174             | 60             | 576             |
| **Mean (SD)**                              | 58.1 (12.0)               | 61.1 (13.7)   | 58.8 (11.8)         | 52.7 (12.8)       | 54.6 (12.4)     | 59.4 (13.6)    | 57.2 (12.8)     |
| **Median (range)**                         | 61 (34-81)                | 61 (18-84)    | 59 (26-90)          | 53 (29-82)        | 55 (21-92)      | 57 (37-92)     | 57 (18-92)      |
| **Unknown**                                | 1                         | 0             | 0                   | 0                 | 1               | 0              | 2               |
| **Sex**                                    | Male 10 (47.6)            | 35 (46.7)     | 82 (44.3)           | 31 (50.0)         | 81 (46.3)       | 21 (35.0)      | 260 (45.0)      |
| **Female**                                 | 11 (52.4)                 | 40 (53.3)     | 103 (55.7)          | 31 (50.0)         | 94 (53.7)       | 39 (65.0)      | 318 (55.0)      |
| **No. of comorbidities at diagnosis**      | 0                         | 7 (33.3)      | 26 (34.7)           | 69 (37.3)         | 31 (50.0)       | 106 (60.6)     | 27 (45.0)       |
| **1-2**                                    | 12 (57.1)                 | 40 (53.3)     | 95 (51.4)           | 26 (41.9)         | 65 (37.1)       | 26 (43.3)      | 264 (45.7)      |
| **> 2**                                    | 2 (9.5)                   | 9 (12.0)      | 21 (11.4)           | 5 (8.1)           | 4 (2.3)         | 7 (11.7)       | 48 (8.3)        |
| **Most frequent comorbidities at diagnosis**| Hypertension 13 (61.9)   | 30 (40.0)     | 55 (29.7)           | 10 (16.1)         | 30 (17.1)       | 22 (36.7)      | 160 (27.7)      |
| **Diabetes**                               | 1 (4.8)                   | 9 (12.0)      | 13 (7.0)            | 4 (6.5)           | 26 (14.9)       | 6 (10.0)       | 59 (10.2)       |
| **Heart disease**                          | 1 (4.8)                   | 3 (4.0)       | 28 (15.1)           | 2 (3.2)           | 3 (1.7)         | 1 (1.7)        | 38 (6.6)        |
| **ECOG PS at diagnosis**                   | 0                         | 4 (100)       | 50 (75.8)           | 32 (80.0)         | 8 (47.1)        | 17 (54.8)      | 6 (50.0)        |
| **1**                                      | 0 (0.0)                   | 12 (18.2)     | 7 (17.5)            | 6 (35.3)          | 9 (29.0)        | 4 (33.3)       | 38 (22.4)       |
| **2**                                      | 0 (0.0)                   | 3 (4.5)       | 1 (2.5)             | 2 (11.8)          | 5 (16.1)        | 2 (16.7)       | 13 (7.6)        |
| **3**                                      | 0 (0.0)                   | 0 (0.0)       | 0 (0.0)             | 1 (5.9)           | 0 (0.0)         | 0 (0.0)        | 1 (0.6)         |
| **≥ 4**                                    | 0 (0.0)                   | 1 (1.5)       | 0 (0.0)             | 0 (0.0)           | 0 (0.0)         | 0 (0.0)        | 1 (0.6)         |
| **Not available**                          | 17                        | 9             | 145                 | 45                | 144             | 48             | 408             |
| **Ann Arbor stage at diagnosis**           | I 6 (37.5)                | 9 (12.2)      | 29 (17.5)           | 5 (8.6)           | 16 (10.0)       | 4 (8.7)        | 69 (13.3)       |
|                                            | II 6 (37.5)               | 15 (20.3)     | 32 (19.3)           | 11 (19.0)         | 28 (17.5)       | 10 (21.7)      | 102 (19.6)      |
|                                            | III 3 (18.8)              | 24 (32.4)     | 51 (30.7)           | 26 (44.8)         | 59 (36.9)       | 14 (30.4)      | 177 (34.0)      |
|                                            | IV 1 (6.3)                | 26 (35.1)     | 54 (32.5)           | 16 (27.6)         | 57 (35.6)       | 18 (39.1)      | 172 (33.1)      |
|                                            | NE 0                      | 0             | 17                  | 1                 | 3               | 0              | 21              |

(Continued on following page)
TABLE 5. Demographic, Baseline Clinical Characteristics, and Treatment Patterns of Patients With Follicular Lymphoma (Continued)

| Characteristic                        | Panama/Guatemala (n = 21) | Chile (n = 75) | Argentina (n = 185) | Colombia (n = 62) | Mexico (n = 175) | Brazil (n = 60) | Total (n = 578) |
|---------------------------------------|---------------------------|---------------|---------------------|------------------|-----------------|---------------|-----------------|
| Unknown                               | 5                         | 1             | 2                   | 3                | 12              | 14            | 37              |
| Missing, No.                          | 0                         | 0             | 0                   | 0                | 0               | 0             | 0               |
| Most frequent first-contact physician |                           |               |                     |                  |                 |               |                 |
| Hematologist                          | 2 (9.5)                   | 57 (76.0)     | 50 (27.0)           | 32 (51.6)        | 22 (12.6)       | 5 (8.3)       | 168 (29.1)      |
| General physician                     | 7 (33.3)                  | 3 (4.0)       | 48 (25.9)           | 13 (21.0)        | 34 (19.4)       | 19 (31.7)     | 124 (21.5)      |
| Surgeon                               | 3 (14.3)                  | 4 (5.3)       | 30 (16.2)           | 8 (12.9)         | 37 (21.1)       | 5 (8.3)       | 87 (15.1)       |
| Time from initial diagnosis to treatment, months |               |               |                     |                  |                 |               |                 |
| Mean (SD)                             | 2.5 (3.44)                | 2.8 (3.99)    | 5.5 (12.25)         | 2.2 (4.30)       | 1.9 (5.74)      | 2.7 (4.04)    | 3.3 (8.25)      |
| Median (range)                        | 1.2 (0.1-13.4)            | 2.2 (< 0.1-30.6) | 1.2 (< 0.1-74.7) | 0.8 (< 0.1-20.8) | 0.7 (0-60.3) | 1.5 (< 0.1-21.8) | 1.1 (0-74.7) |
| Missing                               | 3                         | 6             | 16                  | 18               | 24              | 12            | 79              |
| Three most-used chemotherapy treatments |                           |               |                     |                  |                 |               |                 |
| R-CHOP                                | 12 (57.1)                 | 19 (27.9)     | 101 (59.1)          | 35 (57.4)        | 97 (57.1)       | 16 (29.1)     | 280 (51.3)      |
| RCVP                                  | 5 (23.8)                  | 10 (14.7)     | 38 (22.2)           | 18 (29.5)        | 20 (11.8)       | 5 (9.1)       | 96 (17.6)       |
| CHOP                                  | 2 (9.5)                   | 26 (38.2)     | 3 (1.8)             | 0 (0.0)          | 22 (12.9)       | 17 (30.9)     | 70 (12.8)       |

Abbreviations: CHOP, cyclophosphamide, doxorubicin, vincristine, and prednisone; ECOG PS, Eastern Cooperative Oncology Group performance status; NE, not estimated; R-CHOP, rituximab plus cyclophosphamide, doxorubicin, vincristine, and prednisone; RCVP, rituximab, cyclophosphamide, vincristine, prednisone; SD, standard deviation.

studied were in ISS stage II or III. Approximately one third of patients received thalidomide as frontline therapy, and 26.9% received ASCT.\textsuperscript{15} An analysis of medical records for 9,120 patients at a hematologic clinic in Puebla, Mexico, over 20 years identified 855 individuals with HMs, including 66 (7.7%) with MM; median age was 66 years, and 51% survived through 540 days. The authors contended that a range of immunoproliferative disorders are less frequent in the indigenous Mexican population (mestizos) compared with whites, including CLL (5 times less frequent) and monoclonal gammopathy of undetermined significance (4 times less frequent).\textsuperscript{16}

In a series of 1,028 patients with NHL in Central America (Guatemala) and South America (Argentina, Brazil, Chile, and Peru), Laurini et al\textsuperscript{17} had findings consistent with those of the HOLA study. Forty percent of patients with NHL had DLBCL. In the HOLA study, 49.1% of patients with NHL had DLBCL (63.5% in Panama/Guatemala). In all, 20.4% of Central and South American patients with NHL had FL; 6.9% had marginal zone B-cell lymphoma, MALT type; and 6.8% had peripheral T-cell lymphomas. Median age of Latin American patients with NHL was 59 years compared with 58 (in patients with DLBCL) in the HOLA study. Other studies in Guatemala,\textsuperscript{17} Mexico,\textsuperscript{18} and Chile\textsuperscript{19} echo the findings from the HOLA study, showing that DLBCL was the most frequent subtype of NHL and revealing that most patients had a B-cell rather than a T-cell lineage. The Guatemalan study of 226 consecutive biopsy samples showed that 44% of patients with NHL had DLBCL (median age, 58.5 years).\textsuperscript{17}

The combination of the HOLA cohort size (\(N = 5,140\)) and broad scope of follow-up data (8 years) are study strengths. Given the liberal eligibility criteria and observational nature of the study, the findings should be generalizable to most Latin American practitioners’ treatment populations and settings. Chart abstractors were centrally trained and had considerable expertise in using clearly prespecified disease definitions. On the other hand, the cohort represented a convenience population, which potentially reduces the generalizability of the findings. Because of potential medical surveillance bias, the prevailing tertiary care nature of the treatment milieus may have resulted in over-estimations of the percentage of patients with HMs, comorbidities, and treatment patterns compared with less-specialized clinics. Given that Latin American cancer care delivery systems are largely skewed toward urban settings,
individuals who reside in rural locales may have been underrepresented. Rural workers may experience greater vocational exposure to insecticides and other lymphomagens. The International Classification of Diseases for Oncology codes used to identify patients with MM, CLL, and NHL were developed for reimbursement, not for case ascertainment purposes. Numbers of centers, and hence overall denominators in calculations, were somewhat small, especially in Chile and Panama/Guatemala.

In conclusion, the HOLA study generated an unprecedented level of high-quality, real-world evidence on the disease and treatment characteristics of patients with HMs. Considerable regional variations in HM management were observed, and the findings can likely be ascribed to heterogeneity in both patient characteristics and treatment patterns. Results from this study can be used to understand the disease landscape in Latin America. Future research is needed to explicitly associate observed trends with treatment outcomes.

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15 Hospital Italiano La Plata, Buenos Aires, Argentina
16 Hospital das Clínicas da Universidade Federal de Goiás, Goiânia, Brazil
17 Hospital das Clínicas de Porto Alegre, Porto Alegre, Brazil
18 Star Hospitals, Buenos Aires, Argentina
19 Hospital Italiano La Plata, Buenos Aires, Argentina
20 Hospital das Clínicas da Universidade Federal de Goiás, Goiânia, Brazil
21 Hospital das Clínicas de Porto Alegre, Porto Alegre, Brazil
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**EQUAL CONTRIBUTION**

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**DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST**

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**AUTHORS’ DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST**

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REFERENCES

1. Dighiero G, Hamblin TJ: Chronic lymphocytic leukaemia. Lancet 371:1017-1029, 2008
2. Hossain MS, Iqbal MS, Khan MA, et al: Diagnosed hematological malignancies in Bangladesh - a retrospective analysis of over 5000 cases from 10 specialized hospitals. BMC Cancer 14:438, 2014
3. Acevedo AM, Gómez A, Becerra HA, et al: Distribution and trends of hematology and oncology research in Latin America: A decade of uncertainty. Cancer 120:1237-1245, 2014
4. Curado MP, de Souza DL: Cancer burden in Latin America and the Caribbean. Ann Glob Heal 80:370-377, 2014
5. International Agency for Research on Cancer: CIA: Cancer Incidence in Five Continents. Lyon, France, IARC, 2014
6. Burger JA, Tedeschi A, Barr PM, et al: Ibrutinib as initial therapy for patients with chronic lymphocytic leukemia. N Engl J Med 373:2425-2437, 2015
7. Hungria VTM, Maiolino A, Martinez G, et al: Confirmation of the utility of the International Staging System and identification of a unique pattern of disease in Brazilian patients with multiple myeloma. Haematologica 93:791-792, 2008
8. Eichhorst B, Robak T, Ghan P, et al: Chronic lymphocytic leukaemia: ESMO clinical practice guidelines for diagnosis, treatment and follow-up. Ann Oncol 26:v78-v84, 2015
9. Ludwig H, Sonneveld P, Davies F, et al: European perspective on multiple myeloma treatment strategies in 2014. Oncologist 19:829-844, 2014
10. Rajkumar SV, Dimopoulos MA, Palumbo A, et al: International Myeloma Working Group updated criteria for the diagnosis of multiple myeloma. Lancet Oncol 15:e538-e548, 2014
11. Palumbo A, Rajkumar SV, San Miguel JF, et al: International Myeloma Working Group consensus statement for the management, treatment, and supportive care of patients with myeloma not eligible for standard autologous stem-cell transplantation. J Clin Oncol 32:587-600, 2014
12. Hallek M, Cheson BD, Catovsky D, et al: Guidelines for the diagnosis and treatment of chronic lymphocytic leukemia: A report from the International Workshop on Chronic Lymphocytic Leukemia updating the National Cancer Institute-Working Group 1996 guidelines. Blood 111:5446-5456, 2008
13. Brenner H, Gondos A, Pulte D: Recent major improvement in long-term survival of younger patients with multiple myeloma. Blood 111:2521-2526, 2008
14. Hungria VTD, Crusoe EQ, Quero AA, et al: Guidelines on the diagnosis and management of multiple myeloma treatment: Associação Brasileira de Hematologia e Hemoterapia e Terapia Celular Project guidelines: Associação Médica Brasileira - 2012. Rev Bras Hematol Hemoter 35:201-217, 2013
15. Hungria VTM, Maiolino A, Martinez G, et al: Observational study of multiple myeloma in Latin America. Ann Hematol 96:65-72, 2017
16. Ruiz-Argüelles GJ, Gómez-Rangel JD, Ruiz-Delgado GJ, et al: Multiple myeloma in Mexico: A 20-year experience at a single institution. Arch Med Res 35:163-167, 2004
17. Laurini JA, Perry AM, Boilesen E, et al: Classification of non-Hodgkin lymphoma in Central and South America: A review of 1028 cases. Blood 120:4795-4801, 2012
18. Perry AM, Molina-Kirsch H, Nathwani BN, et al: Classification of non-Hodgkin lymphomas in Guatemala according to the World Health Organization system. Leuk Lymphoma 52:1681-1688, 2011
19. Alavanja MC, Hofmann JN, Lynch CF, et al: Non-Hodgkin lymphoma risk and insecticide, fungicide and fumigant use in the Agricultural Health Study. PLoS One 9:e109332, 2014
20. Vlaanderen J, Lan Q, Kromhout H, et al: Occupational benzene exposure and the risk of lymphoma subtypes: A meta-analysis of cohort studies incorporating three study quality dimensions. Environ Health Perspect 119:159-167, 2011
### APPENDIX

#### TABLE A1. Baseline Characteristics of Patients With Mantle Cell Lymphoma

| Characteristic | Panama/Guatemala (n = 9) | Chile (n = 20) | Argentina (n = 37) | Colombia (n = 38) | Mexico (n = 38) | Brazil (n = 41) | Total (n = 183) |
|----------------|--------------------------|----------------|-------------------|-------------------|----------------|----------------|----------------|
| **Age at diagnosis, years** | | | | | | | |
| Mean (SD) | 92 (16.3) | 60.4 (11.5) | 59.4 (12.4) | 62.9 (10.7) | 61.6 (15.2) | 63.2 (11.4) | 61.8 (12.5) |
| Median (range) | 61 (38-93) | 58 (39-85) | 60 (39-82) | 63 (36-88) | 63 (27-84) | 63 (39-87) | 61 (27-93) |
| **Age category at diagnosis, years** | | | | | | | |
| < 50 | 2 (22.2) | 2 (10.0) | 11 (29.7) | 4 (10.5) | 7 (18.5) | 6 (14.6) | 32 (17.5) |
| 50 to < 60 | 2 (22.2) | 9 (45.0) | 7 (18.9) | 9 (23.7) | 7 (18.4) | 11 (26.8) | 45 (24.6) |
| 60 to < 70 | 3 (33.3) | 5 (25.0) | 11 (29.7) | 15 (39.5) | 11 (28.9) | 12 (29.3) | 57 (31.1) |
| 70 to < 80 | 1 (11.1) | 4 (15.0) | 4 (10.8) | 9 (23.7) | 9 (23.7) | 9 (22.0) | 35 (19.1) |
| ≥ 80 | 1 (11.1) | 1 (5.0) | 4 (10.8) | 1 (2.6) | 4 (10.5) | 3 (7.3) | 14 (7.7) |
| **Sex** | | | | | | | |
| Male | 7 (77.8) | 12 (60.0) | 30 (81.1) | 27 (71.1) | 26 (68.4) | 26 (63.4) | 128 (69.9) |
| Female | 2 (22.2) | 8 (40.0) | 7 (18.9) | 11 (28.9) | 12 (31.6) | 15 (36.6) | 55 (30.1) |
| **No. of comorbidities at diagnosis** | | | | | | | |
| 0 | 1 (11.1) | 6 (30.0) | 15 (40.5) | 17 (44.7) | 21 (55.3) | 14 (34.1) | 74 (40.4) |
| 1-2 | 7 (77.8) | 10 (50.0) | 16 (43.2) | 18 (47.4) | 12 (31.6) | 24 (58.5) | 87 (47.5) |
| > 2 | 1 (11.1) | 4 (20.0) | 6 (16.2) | 3 (7.9) | 5 (13.2) | 3 (7.3) | 22 (12.0) |
| **Most frequent comorbidities at diagnosis** | | | | | | | |
| Hypertension | 5 (55.6) | 8 (40.0) | 12 (32.4) | 8 (21.1) | 9 (23.7) | 15 (36.6) | 57 (31.1) |
| Diabetes | 3 (33.3) | 6 (30.0) | 4 (10.8) | 4 (10.5) | 5 (13.2) | 3 (7.3) | 25 (13.7) |
| Heart disease | 2 (22.2) | 1 (5.0) | 9 (24.3) | 4 (10.5) | 4 (10.5) | 2 (4.9) | 22 (12.0) |
| **ECOG PS at diagnosis** | | | | | | | |
| 0 | 0 (0.0) | 12 (66.7) | 9 (81.8) | 4 (66.7) | 5 (41.7) | 3 (75.0) | 33 (63.5) |
| 1 | 0 (0.0) | 5 (27.8) | 0 (0.0) | 0 (0.0) | 6 (50.0) | 0 (0.0) | 11 (21.2) |
| 2 | 1 (100) | 1 (5.6) | 2 (18.2) | 2 (33.3) | 1 (8.3) | 1 (25.0) | 8 (15.4) |
| 3 | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) |
| ≥ 4 | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) | 0 (0.0) |
| **Not available** | | | | | | | |
| 0 | 8 | 2 | 26 | 32 | 26 | 37 | 131 |
| **Ann Arbor stage at diagnosis** | | | | | | | |
| I | 1 (50.0) | 4 (21.1) | 1 (3.6) | 2 (6.1) | 2 (6.1) | 2 (5.4) | 12 (7.9) |
| II | 0 (0.0) | 3 (15.8) | 1 (3.6) | 2 (6.1) | 2 (6.1) | 2 (5.4) | 10 (6.6) |
| III | 0 (0.0) | 5 (26.3) | 4 (14.3) | 5 (15.2) | 7 (21.2) | 2 (5.4) | 23 (5.1) |

(Continued on following page)
| Characteristic                        | Panama/Guatemala (n = 9) | Chile (n = 20) | Argentina (n = 37) | Colombia (n = 38) | Mexico (n = 38) | Brazil (n = 41) | Total (n = 183) |
|--------------------------------------|--------------------------|----------------|--------------------|-------------------|----------------|----------------|----------------|
| IV                                   | 1 (50.0)                 | 7 (36.8)       | 22 (78.6)          | 24 (72.7)         | 22 (66.7)      | 31 (83.8)      | 107 (70.4)     |
| NE                                   | 3                        | 0              | 9                  | 3                 | 1              | 0              | 16             |
| Unknown                              | 4                        | 1              | 0                  | 2                 | 4              | 4              | 15             |

Most frequent first-contact physician

- **Hematologist**: 3 (33.3) 15 (75.0) 13 (35.1) 16 (42.1) 6 (15.8) 4 (9.8) 57 (31.1)
- **General physician**: 1 (11.1) 3 (15.0) 12 (32.4) 11 (28.9) 4 (10.5) 6 (14.6) 37 (20.2)
- **Surgeon**: 2 (22.2) 2 (10.0) 7 (18.9) 3 (7.9) 5 (13.2) 1 (2.4) 20 (10.9)

Time from initial diagnosis to treatment, months

- **Mean (SD)**: 0.4 (0.4) 1.5 (0.7) 4.6 (12.5) 0.7 (1.3) 1.3 (1.4) 2.3 (1.8) 2.1 (5.9)
- **Median (range)**: 0.3 (<0.1-0.2) 1.6 (<0.1-2.9) 1.1 (<0.1-63.3) 0.2 (0-6.0) 0.7 (<0.1-4.2) 1.6 (<0.1-6.3) 0.9 (0-63.3)
- **Missing**: 3 5 4 10 13 7 42

Three most-used chemotherapy treatments

- **R-CHOP**: 3 (33.3) 5 (26.3) 13 (41.9) 20 (54.1) 15 (41.7) 15 (37.5) 71 (41.3)
- **CHOP**: 2 (22.2) 11 (57.9) 0 (0.0) 3 (8.1) 0 (0.0) 12 (30.0) 28 (16.3)
- **Hyper CVAD**: 1 (11.1) 2 (10.5) 1 (3.2) 1 (2.7) 8 (22.2) 5 (12.5) 18 (10.5)

Abbreviations: CHOP, cyclophosphamide, doxorubicin, vincristine, and prednisone; CVAD, cyclophosphamide, vincristine, doxorubicin, and dexamethasone; ECOG PS, Eastern Cooperative Oncology Group performance status; NE, not estimated; R-CHOP, rituximab plus cyclophosphamide, doxorubicin, vincristine, and prednisone; SD, standard deviation.
| Characteristic                          | Panama/Guatemala (n = 4) | Chile (n = 14) | Argentina (n = 17) | Colombia (n = 12) | Mexico (n = 23) | Brazil (n = 14) | Total (n = 84) |
|----------------------------------------|--------------------------|----------------|--------------------|-------------------|-----------------|---------------|---------------|
| Age at diagnosis, years                |                          |                |                    |                   |                 |               |               |
| Mean (SD)                              | 67.8 (8.7)               | 64.2 (17.2)    | 56.4 (17.2)        | 57.8 (13.1)       | 49.4 (17.9)     | 61.6 (14.5)    | 57.4 (16.8)   |
| Median (range)                         | 65 (61-80)               | 67 (21-86)     | 55 (29-88)         | 63 (31-74)        | 47 (18-82)      | 63 (35-77)     | 62 (18-88)    |
| Age category at diagnosis, years       |                          |                |                    |                   |                 |               |               |
| < 50                                   | 0 (0.0)                  | 2 (14.3)       | 7 (41.2)           | 3 (25.0)          | 13 (56.5)       | 3 (21.4)       | 28 (33.3)     |
| 50 to < 60                             | 0 (0.0)                  | 2 (14.3)       | 3 (17.6)           | 2 (16.7)          | 3 (13.0)        | 2 (14.3)       | 12 (14.3)     |
| 60 to < 70                             | 3 (75.0)                 | 3 (21.4)       | 1 (5.9)            | 5 (41.7)          | 4 (17.4)        | 3 (21.4)       | 19 (22.6)     |
| 70 to < 80                             | 0 (0.0)                  | 5 (35.7)       | 5 (29.4)           | 2 (16.7)          | 2 (8.7)         | 6 (42.9)       | 20 (23.8)     |
| ≥ 80                                   | 1 (25.0)                 | 2 (14.3)       | 1 (5.9)            | 0                 | 1 (4.3)         | 0             | 5 (6.0)       |
| Sex                                    |                          |                |                    |                   |                 |               |               |
| Male                                   | 1 (25.0)                 | 8 (57.1)       | 10 (58.8)          | 4 (33.3)          | 6 (26.1)        | 7 (50.0)       | 36 (42.9)     |
| Female                                 | 3 (75.0)                 | 6 (42.9)       | 7 (41.2)           | 8 (66.7)          | 17 (73.9)       | 7 (50.0)       | 48 (57.1)     |
| No. of comorbidities at diagnosis      |                          |                |                    |                   |                 |               |               |
| 0                                      | 0 (0.0)                  | 4 (28.6)       | 5 (29.4)           | 1 (8.3)           | 18 (78.3)       | 1 (7.1)        | 29 (34.5)     |
| 1-2                                    | 4 (100)                  | 9 (64.3)       | 10 (58.8)          | 11 (91.7)         | 5 (21.7)        | 12 (85.7)      | 51 (60.7)     |
| > 2                                    | 0 (0.0)                  | 1 (7.1)        | 2 (11.8)           | 0 (0.0)           | 0 (0.0)         | 1 (7.1)        | 4 (4.8)       |
| Most frequent comorbidities at diagnosis|                        |                |                    |                   |                 |               |               |
| Hypertension                           | 4 (100)                  | 7 (50.0)       | 4 (23.5)           | 2 (16.7)          | 4 (17.4)        | 7 (50.0)       | 28 (33.3)     |
| Diabetes                               | 1 (25.0)                 | 4 (28.6)       | 1 (5.9)            | 1 (8.3)           | 1 (4.3)         | 5 (35.7)       | 13 (15.5)     |
| Heart disease                          | 1 (25.0)                 | 0 (0.0)        | 3 (17.6)           | 1 (8.3)           | 0 (0.0)         | 1 (7.1)        | 6 (7.1)       |
| Infections                             | 1 (25.0)                 | 0 (0.0)        | 2 (11.8)           | 2 (16.7)          | 1 (4.3)         | 0 (0.0)        | 6 (7.1)       |
| ECOG PS at diagnosis                   |                          |                |                    |                   |                 |               |               |
| 0                                      | 0 (0.0)                  | 6 (60.0)       | 5 (71.4)           | 0 (0.0)           | 4 (66.7)        | 1 (100)        | 16 (64.0)     |
| 1                                      | 0 (0.0)                  | 4 (40.0)       | 0 (0.0)            | 0 (0.0)           | 1 (16.7)        | 0 (0.0)        | 5 (20.0)      |
| 2                                      | 0 (0.0)                  | 0 (0.0)        | 2 (28.6)           | 1 (100)           | 0 (0.0)         | 0 (0.0)        | 3 (12.0)      |
| 3                                      | 0 (0.0)                  | 0 (0.0)        | 0 (0.0)            | 0 (0.0)           | 1 (16.7)        | 0 (0.0)        | 1 (4.0)       |
| ≥ 4                                    | 0 (0.0)                  | 0 (0.0)        | 0 (0.0)            | 0 (0.0)           | 0 (0.0)         | 0 (0.0)        | 0 (0.0)       |
| Not available                          | 4 (100)                  | 4 (28.6)       | 3 (33.3)           | 1 (25.0)          | 4 (17.4)        | 1 (16.7)       | 13 (26.0)     |
| Ann Arbor stage at diagnosis           |                          |                |                    |                   |                 |               |               |
| I                                      | 0 (0.0)                  | 6 (42.9)       | 2 (22.2)           | 1 (25.0)          | 9 (52.9)        | 0 (0.0)        | 18 (36.0)     |
| II                                     | 0 (0.0)                  | 4 (33.3)       | 3 (33.3)           | 1 (25.0)          | 4 (23.5)        | 1 (16.7)       | 13 (26.0)     |
| III                                    | 0 (0.0)                  | 3 (21.4)       | 3 (33.3)           | 0 (0.0)           | 1 (5.9)         | 0 (0.0)        | 7 (14.0)      |
| IV                                     | 0 (0.0)                  | 1 (7.1)        | 1 (11.1)           | 2 (50.0)          | 3 (17.6)        | 5 (83.3)       | 12 (24.0)     |
| NE                                     | 3 (75.0)                 | 0 (0.0)        | 8 (57.1)           | 3 (25.0)          | 1 (8.3)         | 0 (0.0)        | 15 (36.0)     |
| Unknown                                | 1 (25.0)                 | 0 (0.0)        | 5 (33.3)           | 5 (41.7)          | 4 (17.4)        | 3 (21.4)       | 19 (22.6)     |

(Continued on following page)
### TABLE A2. Baseline Characteristics of Patients With Mucosa-Associated Lymphoid Tissue Lymphoma (Continued)

| Characteristic | Panama/Guatemala (n = 4) | Chile (n = 14) | Argentina (n = 17) | Colombia (n = 12) | Mexico (n = 23) | Brazil (n = 14) | Total (n = 84) |
|---------------|--------------------------|---------------|-------------------|------------------|----------------|---------------|---------------|
| **Most frequent first-contact physician** |                         |               |                   |                  |                |               |               |
| Hematologist  | 0 (0.0)                  | 10 (71.4)     | 7 (41.2)          | 6 (50.0)         | 2 (8.7)        | 0 (0.0)       | 25 (29.8)     |
| General physician | 3 (75.0)                 | 0 (0.0)       | 1 (5.9)           | 2 (16.7)         | 5 (21.7)       | 3 (21.4)       | 14 (16.7)     |
| Surgeon       | 0 (0.0)                  | 2 (14.3)      | 8 (47.1)          | 1 (8.3)          | 1 (4.3)        | 0 (0.0)       | 12 (14.3)     |
| **Time from initial diagnosis to treatment, months** |                         |               |                   |                  |                |               |               |
| Mean (SD)     | 2.1 (1.20)               | 3.8 (2.11)    | 9.8 (13.79)       | 1.4 (1.44)       | 2.1 (1.89)     | 9.3 (12.92)   | 4.8 (8.29)    |
| Median (range)| 2.2 (0.9-3.3)            | 3.4 (1.1-7.8) | 2.2 (<0.1-41.9)   | 0.9 (<0.1-4.8)   | 1.1 (<0.1-6.2) | 3.9 (0.5-42.1)| 2.3 (<0.1-42.1)|
| Missing       | 1                        | 2             | 2                 | 0                | 4              | 1             | 10            |
| **Three most-used chemotherapy treatments** |                         |               |                   |                  |                |               |               |
| R-CHOP        | 1 (25.0)                 | 1 (8.3)       | 6 (50.0)          | 3 (30.0)         | 11 (57.9)      | 2 (20.0)      | 24 (35.8)     |
| CHOP          | 0 (0.0)                  | 8 (66.7)      | 0 (0.0)           | 0 (0.0)          | 3 (15.8)       | 1 (10.0)      | 12 (17.9)     |
| RCVP          | 3 (75.0)                 | 0 (0.0)       | 1 (8.3)           | 3 (30.0)         | 1 (5.3)        | 1 (10.0)      | 9 (13.4)      |

Abbreviations: CHOP, cyclophosphamide, doxorubicin, vincristine, and prednisone; ECOG PS, Eastern Cooperative Oncology Group performance status; NE, not estimated; R-CHOP, rituximab plus cyclophosphamide, doxorubicin, vincristine, and prednisone; RCVP, rituximab, cyclophosphamide, vincristine, and prednisone; SD, standard deviation.