Burkitt’s lymphoma: The prevalence of HIV/AIDS and the outcome of treatment

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The prevalence of HIV in Burkitt’s lymphoma (BL) patients and the outcome of treatment in Cameroon were unknown. Records of all patients diagnosed with BL at three Cameroon Baptist Convention hospitals were reviewed to ascertain the recorded HIV status and outcome of treatment. Of 979 patients diagnosed with BL, 717 were tested for HIV and 11 (1.5%) were HIV-positive. Three of eight patients treated with both cyclophosphamide (CPM)-based chemotherapy and antiretrovirals were alive at 62, 96 and 111 months, respectively. The HIV rate was comparable to that of 1% for the general population of children aged <15 years. Low-cost high-frequency CPM was the only available treatment option for BL and was associated with 37.5% long-term survival in a resource-limited setting.

The population of Cameroon in 2014 was 22 773 014, of whom ~40% (9 109 205) were aged <15 years. It was estimated that 94 000 of these children (1%) were infected with HIV, and only 11% of these received antiretroviral (ARV) therapy. In 2013, the prevalence of HIV infection in pregnant women in the north-west and south-west regions was 4.6% and 6.8%, respectively, and the mother-to-child transmission rate in 18-month-old breastfed infants was 25%. Children diagnosed with Burkitt’s lymphoma (BL) between 2003 and 2015 at Banso and Mbingo Baptist hospitals in the north-west and Mutengene Baptist Hospital in the south-west were all treated with cyclophosphamide-based chemotherapy regimens, with an overall long-term survival rate of 55%.

Objective
To record the prevalence of HIV and the outcome of treatment in a large cohort of patients with BL.

Methods
Our database is a Pediatric Oncology Network Database (POND) cancer registry for the period 2003 - 2013. The number of patients in whom enzyme-linked immunosorbent assay was positive for HIV, the CD4+ count when available, and whether ARV treatment was given or not were extracted from the database. The diagnosis of BL was based on a fine-needle tumour aspirate, a bone marrow aspirate, a cytological cerebrospinal fluid examination and an abdominal ultrasound scan. The long-term outcome was established by personal follow-up of every HIV-positive patient or their families. Chemotherapy induction treatment consisted of cyclophosphamide (CPM) 40 mg/kg on days 1, 8 and 15, followed by consolidation chemotherapy with 1 - 3 more doses of CPM with or without intravenous (IV) vincristine 1.5 mg/m² and IV methotrexate 1.0 g/m² for advanced or non-responsive disease. The duration of chemotherapy was 2 months. Rescue treatment of early relapses consisted of 3 weekly pulses of CPM and vincristine, and relapses beyond 1 year were treated in full again. ARV treatment was not freely available in the early years of the study. More recent treatment always consisted of three drugs, based on World Health Organization guidelines and the national protocol. Examples of first-line treatment are as follows: age <3 years – abacavir + lamivudine + lopinavir/ritonavir; age 3 - 10 years and adolescents weighing <35 kg – abacavir + lamivudine + efavirenz; and age 11 - 19 years or weight >35 kg – tenofovir + lamivudine (or emtricitabine) + efavirenz.

Ethical approval for the BL treatment protocols was obtained from the Institutional Review Board of the Cameroon Baptist Convention Health Board.

Results
Of 979 patients diagnosed with BL, 717 (73.2%) were tested for HIV and 11 (1.5%) tested positive. Their ages ranged from 2 to 13 years (median 7 years). The age at diagnosis of BL, St Jude stage of the BL, CD4+ count if done, ARV treatment given, current status and duration of long-term survival are listed in Table 1. On 30 October 2016, three of eight patients (37.5%) who had received the recommended ARV treatment and were still on ARV maintenance treatment were long-term survivors at 111, 96 and 62 months, respectively. Three patients who were not treated with ARVs died within 3 - 15 months, and five patients treated with ARVs died within 3 - 39 months after the onset of treatment from progressive disease. There was no chemotherapy-related death. The CD4+ count at diagnosis was recorded in only four patients.

Discussion
Children with HIV have an increased risk of developing malignancies, and BL accounted for 21% of malignancies in HIV-positive children in South Africa, with a projected cure rate of 54.6% at 60 months using modern chemotherapy. Modern chemotherapy for BL has a potential cure rate of >90% in HIV-negative patients, but at considerable cost and the risk of significant morbidity. Our BL
patients were treated with chemotherapy of much lower intensity because of limited access to drugs and supportive care. Long-term survival of 5 - 9 years in three of eight patients (37.5%) who received both chemotherapy and ARVs is encouraging. The documented HIV prevalence rate of 1.5% in this large cohort of BL patients is comparable to the 1% HIV prevalence rate for the general population of children in Cameroon.

**Conclusion**

HIV-positive children with BL are best treated with a context-appropriate chemotherapy regimen.

**Acknowledgements.** Poster presentation at the 48th congress of the International Society of Paediatric Oncology (SIOP), Dublin, October 2016.

**Author contributions.** PBH: Principal investigator, FK: physician in charge of the children’s cancer service, EK: physician in charge of the HIV service of the Cameroon Baptist Convention Health Board, GM: research assistant responsible for the cancer registry and data co-ordination, PW: involved in planning and follow-up.

**Funding.** Funding was received from the Beryl Thyer Memorial Africa Trust (UK), World Child Cancer (UK) and the Swiss Cancer League.

**Conflicts of interest.** None.

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**Accepted 5 September 2017.**

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Table 1. Patient details and outcome

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**ARV = antiretroviral.**