Research Article

Clinical presentation and outcomes in children with retinoblastoma managed at the Uganda Cancer Institute

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Background. The majority of patients with retinoblastoma, the most common intraocular cancer of childhood, are found in low- and middle-income countries (LMICs), with leukocoria being the most common initial presenting sign and indication for referral. Findings from the current study serve to augment earlier findings on the clinical presentation and outcomes of children with retinoblastoma in Uganda. Methods. This was a retrospective study in which we reviewed records of children admitted with a diagnosis of retinoblastoma at the Uganda Cancer Institute from January 2009 to February 2020. From the electronic database, using admission numbers, files were retrieved. Patient information was recorded in a data extraction tool. Results. A total of 90 retinoblastoma patients were studied, with a mean age at the first Uganda Cancer Institute (UCI) presentation of 36.7 months. There were more males (57.8%) than females, with a male to female ratio of 1.37 : 1. The majority (54.4%) had retinoblastoma treatment prior to UCI admission. The most common presenting symptoms were leukocoria (85.6%), eye reddening (64.4%), and eye swelling (63.3%). At 3 years of follow-up after index admission at UCI, 36.7% of the patients had died, 41.1% were alive, and 22.2% had been lost to follow-up. The median 3-year survival for children with retinoblastoma in our study was 2.18 years. Significant predictors of survival in the multivariate analysis were follow-up duration (P < 0.001), features of metastatic spread (P = 0.001), history of eye swelling (P = 0.012), and bilateral enucleation (P = 0.011). Conclusions. The majority of children who presented to the Uganda Cancer Institute were referred with advanced retinoblastoma, and there was a high mortality rate. Retinoblastoma management requires a multidisciplinary team that should include paediatric ophthalmologists, paediatric oncologists, ocular oncologists, radiation oncologists, and nurses.

1. Introduction

Retinoblastoma is a primary intraocular childhood malignancy arising from the embryonal cells of the retina and the most common primary intraocular malignancy of childhood, with over 90% of cases diagnosed before the age of five [1–3]. The retinoblastoma incidence in the USA is 1 in 15,000–20,000 live births, with 60% of cases presenting with unilateral disease [4]. It is estimated that 250 to 300 new cases in the United States of America and approximately 8,000 to 9,000 cases worldwide of retinoblastoma are diagnosed every year [3, 5]. However, the annual incidence of retinoblastoma is the same across geographical regions, and the annual prevalence of retinoblastoma is higher in Africa than in higher income countries [3, 6].

Studies show that over 95% of children with retinoblastoma in the United States and other developed nations survive their malignancy, whereas approximately 50% survive
of survival among children with retinoblastoma. The pattern of clinical presentation, outcomes and predictors in oncology facilities in East Africa. With the increasing number of children with retinoblastoma who needed chemotherapy as the treatment of choice for each participant. Treatment consisted of one or a combination of the following treatment modalities: chemotherapy, surgery (unilateral or bilateral enucleation), and external beam radiotherapy.

2.2. Data Management and Analysis. The extracted data were entered into an electronic database using Epidata 4.2 by the double entry method. The database was checked and cleaned for consistency. The cleaned data were exported into STATA 17 for analysis. Continuous variables such as age and lag time were summarized as the mean ± standard deviation. Categorical variables such as sex, clinical signs and symptoms, laterality, and vital signs were summarized in proportions or frequencies and presented as tables or graphs. The time-specific survival rate (3-year survival) with a 95% confidence interval (CI) was estimated by the Kaplan–Meier method and assessed by a log-rank test. Three-year survival was defined as the period of follow-up of 3 years from the date of index admission at UCI. Cox regression analysis was used to assess the predictors of outcome and presented as hazard ratios with their 95% confidence intervals at both bivariate and multivariate levels. A P value <0.2 was considered at bivariate analysis, and all variables...
that had \( P < 0.2 \) were entered at multivariate Cox regression analysis. A \( P \) value <0.05 was considered significant.

3. Results

A total of 97 charts with retinoblastoma diagnoses were retrieved from the records department of the Uganda Cancer Institute, and 90 charts that had confirmed retinoblastoma diagnoses were studied. There was a mean age of 33.3 months (SD 25.3) for bilateral retinoblastoma and 38 months (SD 25.6) for unilateral retinoblastoma, with an overall mean age at first UCI presentation of 36.7 months (SD 25.5). There were more males than females, with a male to female ratio of 1.37:1. The majority of study subjects (36.7%, \( n = 33 \)) came from the central region of the country, and 13.3% (\( n = 12 \)) of children were nationals of neighboring countries (Table 1).

Leukocoria was the most common retinoblastoma sign (77, 85.6%) reported by parents prior to the first UCI visit, followed by eye reddening (58, 64.4%) and eye swelling (57, 63.3%). There was a family history of retinoblastoma among first-degree relatives in 4.4% of children. The majority of patients had prior retinoblastoma treatment at eye departments of tertiary hospitals before UCI admission, with 46.7% having had enucleation of at least one eye, as shown in Table 1. A significant number (37, 41%) of children had a lag time to UCI admission of at least 12 months (Table 1).

Using the international intraocular retinoblastoma classification scheme (IIRC), most eyes in our study presented with advanced disease of IIRC groups D (20%) and E (53%). Twenty eyes (17%) had no record of their IIRC classification (Table 2).

Sixty-five patients (72.3%) presented with unilateral retinoblastoma, and proptosis was the most common (45%, \( n = 41 \)) presenting sign, including children with recurrent orbital retinoblastoma, followed by leukocoria 36 (40%).

The majority (55, 97.8%) of children with unilateral retinoblastoma whose IIRC staging were performed had Group E (81.8%) classification, followed by D (18.1%) classification (Table 2).

At the 3-year follow-up after admission at UCI with a diagnosis of retinoblastoma, 41.1% of the patients were still alive, and 36.7% of the patients had died. Furthermore, 54.4% and 3.3% had unilateral and bilateral eye loss, respectively (Table 3). The median survival time in years for children with retinoblastoma in our study was 2.18 years. (Figure 1).

The factors that were statistically significant predictors of mortality in the multivariate analysis were lag time between first sign and UCI visit, history of eye swelling (Figure 2), features of metastatic spread (Figure 3), and bilateral enucleation. Children with metastatic spread (AHR 8.93, CI 3.23–24.69, \( P = 0.001 \)) were 8.9 times more likely to die than those without metastatic spread. Children who presented with a history of eye swelling prior to any retinoblastoma treatment (AHR 4.39, CI 1.38–14.01, \( P = 0.012 \)) were 4.4 times more likely to die than those who had no history of eye swelling. In addition, children who underwent bilateral enucleation (AHR 3.53, CI 1.33–9.36, \( P = 0.011 \)) were more likely to die than those who did not undergo enucleation (Table 4).

| Variable                                      | Frequency | Percentage (%) |
|-----------------------------------------------|-----------|----------------|
| Gender                                        |           |                |
| Female                                        | 38        | 42.2           |
| Male                                          | 52        | 57.8           |
| Age group at first sign in months             |           |                |
| 0 to 12                                       | 40        | 40             |
| 13 to 36                                      | 29        | 32.2           |
| Over 36                                       | 21        | 23.3           |
| Age group at first UCI visit in months         |           |                |
| 0 to 12                                       | 5         | 5.6            |
| 13 to 36                                      | 55        | 61.1           |
| Over 36                                       | 30        | 33.3           |
| Symptoms reported by parents                  |           |                |
| Leukocoria                                    | 77        | 85.6           |
| Eye reddening                                 | 58        | 64.4           |
| Eye swelling                                  | 57        | 63.3           |
| Jerky eye movements                           | 8         | 8.9            |
| Others                                        | 7         | 7.8            |
| Family history of retinoblastoma              | 4         | 4.4            |
| Had prior retinoblastoma treatment.           | 49        | 54.4           |
| Enucleation                                   | 42        | 46.7           |
| Radiotherapy                                  | 5         | 5.6            |
| Chemotherapy                                  | 10        | 11.1           |
| Lag time between first sign and UCI visit in months |   |                |
| 0 to 6                                        | 38        | 42.2           |
| 7 to 12                                       | 15        | 16.7           |
| Above 12                                      | 37        | 41.1           |
| Laterality                                    |           |                |
| Unilateral                                    | 65        | 72.3           |
| Bilateral                                     | 25        | 27.8           |
| Vision in the affected eye(s) at first UCI visit (\( n = 73 \) eyes) |   |                |
| No                                            | 66        | 90.4           |
| Yes                                           | 7         | 9.6            |
| Clinical signs at first UCI visit             |           |                |
| Leukocoria                                    | 36        | 40             |
| Proptosis                                     | 41        | 45.6           |
| Enucleated with no visible tumour in socket   | 13        | 14.4           |
| Disease extent                                |           |                |
| Intraocular involvement only                  | 27        | 30             |
| Features of extraocular spread                | 63        | 70             |
| Mean age at first UCI visit (months): overall  |           |                |
| Unilateral                                    | 38.0      | 15.6           |
| Bilateral                                     | 33.3      | 25.3           |

Table 1: Social demographic and clinical characteristics.
There was no record of IIRC classification for 12 (13.3%) children. This finding complicates the management and follow-up of children with retinoblastoma. This was found mostly in children who underwent enucleation prior to UCI admission. The majority of children had the worst affected eyes with Group E (n = 61, 67.8%) IIRC classification, followed by Group D (n = 16, 17.8%). The majority (n = 55, 97.8%) of children with unilateral Rb whose IIRC staging was performed had Group E (81.8%), followed by Group D (18.1%). This finding is comparable to other studies conducted in less developed countries [9, 10, 14–16].

Sixty-five (72.3%) children presented with unilateral retinoblastoma, whereas 25 (27.8%) presented with bilateral disease. These findings were similar to those of other studies, which showed higher unilateral retinoblastoma cases than bilateral retinoblastoma. For instance, studies conducted in India by Chawla et al. and in Kinshasa by Kazadi et al. found that 67.6% and 77.5% of children presented with unilateral retinoblastoma [8, 9].

In our study, the most common retinoblastoma sign that prompted parents to seek eye health attention was leukocoria, reported in 77 (85.6%) patients, followed by eye reddening in 58 (64.4%) and eye swelling in 57 (63.3%), all of which are features of advanced disease. Our findings were similar to those from other studies that revealed leukocoria as the most common sign noted by parents of children with retinoblastoma in India (83%), Singapore (70.6%), and the Democratic Republic of Congo (DRC) (67.5%) [8, 9, 13]. Eye reddening (58, 64.4%) and eye swelling (57, 63.3%) were the second most common signs noted by parents in our study. Proptosis was shown to be the second most common sign noticed by parents in studies in India (17%) and in the DRC (15%) [8, 9, 13]. The percentage of children with proptosis as one of the most common signs was higher in our study than in other studies. This shows that most children in our study sought their first medical attention with advanced disease.

At first presentation at the Uganda Cancer Institute, proptosis was the most common presenting sign reported in 41 (45.6%) children, followed by leukocoria reported in 36 (40%) children. This finding is similar to studies by Kazadi Lukusa et al. in the DRC, with 55% and 25% of children with retinoblastoma presenting with proptosis and leukocoria, respectively [9]. In addition, proptosis was the most common presenting sign (56%) at the National Cancer Institute in Gezira, followed by leukocoria (32%) [12].

A family history of retinoblastoma in first-degree relatives (parents, offspring, and siblings) was reported in 4.4% (n = 4) of children who were admitted at UCI. Our findings are consistent with findings from similar studies in Iran and Taiwan, where a family history of retinoblastoma was reported in 4.8% and 3.1% of patients, respectively [18, 19]. In contrast, a family history of retinoblastoma was reported in 11.0% of children treated between 1981 and 2004 at a medical facility in Turkey [17]. The low percentage of children with a family history of retinoblastoma in our study is possibly due to a lack of knowledge about eye cancers in our community. In addition, patients do not survive to child bearing age.

### Table 2: IIRC classification of children with retinoblastoma at UCI based on worst affected eye(s).

| IIRC group | Total (N = 90) | Bilateral Rb Frequency | Unilateral Rb Frequency |
|------------|---------------|------------------------|-------------------------|
| C          | 1 (1.1%)      | 0 0                    | 1 1.5                   |
| D          | 16 (17.8%)    | 6 24                   | 10 15.4                 |
| E          | 61 (67.8%)    | 16 64                  | 45 69.2                 |
| Staging not done | 12 (13.3%) | 3 12               | 9 13.8                 |

### Table 3: Outcomes of children with retinoblastoma at the last follow-up contact.

| Variable                      | Frequency (N = 90) | Percentage (%) |
|-------------------------------|--------------------|----------------|
| Metastatic spread             |                    |                |
| No                            | 46                 | 51.1           |
| Yes                           | 44                 | 48.9           |
| Visual outcome                |                    |                |
| Vision loss in one eye        | 64                 | 71.1           |
| Vision loss in both eyes      | 12                 | 13.3           |
| Sign of vision present        | 14                 | 15.6           |
| Eye ball loss                 |                    |                |
| No eye loss                   | 38                 | 42.2           |
| Unilateral eye loss           | 49                 | 54.4           |
| Bilateral eye loss            | 3                  | 3.3            |
| Vital status                  |                    |                |
| Alive                         | 37                 | 41.1           |
| Died                          | 33                 | 36.7           |
| Lost to follow-up             | 20                 | 22.2           |

### 4. Discussion

Charts for 90 children with retinoblastoma diagnosis were reviewed. There was a female to male ratio of 1:1.4. With the exception of a study in Iran, which showed a higher female to male ratio, the findings in our study were similar to those of other studies performed in India, Singapore, Eastern Nepal, Kinshasa (DRC), and Gezira-Sudan, which revealed a higher male to female ratio [8, 9, 12, 13]. The majority of the patients (95%) had their first presentation at UCI beyond 12 months of age, with 33% (30 children) at least 3 years old. This is because the largest percentage of children underwent retinoblastoma management prior to admission.

In our study, the overall mean age at the first UCI visit, 36.7 months (SD ± 25.5), was slightly higher than that from a study performed in Eastern Nepal [13]. Our study also revealed a lower difference in age at first UCI presentation between unilateral (38.0(SD ± 25.6) and bilateral (33.3(SD ± 25.3) retinoblastoma cases than that reported in other studies [14, 15]. The disparity in age at first presentation is because a significant number of children (n = 49, 54.4%) admitted at UCI had prior retinoblastoma management.
The mean lag time between the first sign noted by parents and the first UCI visit was 11.2 months (SD 10.8), with a mean lag time of 14.5 months and 9.9 months for bilateral and unilateral retinoblastoma, respectively. A significant number of children (41.1%) had a longer lag time (beyond 12 months). This is because most children were managed at local eye units prior to referral to UCI. Consequently, lag time was a significant predictor of 3-year survival, as patients with retinoblastoma who reported to UCI more than 7 months after developing the first sign of retinoblastoma were more likely to die within 3-years compared to patients who presented within at least 6 months of developing the first sign of retinoblastoma (AHR 0.10, CI 0.04–0.25, \( P = 0.001 \)).

In our study, the majority of children had the worst affected eye, with IIRC Group E 61 (67.8%), followed by Group D 16 (17.8%). Additionally, most children with unilateral Rb (\( n = 55, 97.8\% \)) whose IIRC staging was performed had Group E staging (81.8%) followed by D staging (18.1%). The majority of children (63, 70%) had features of extraocular spread at diagnosis at the first UCI visit. These findings were similar to a study by Lim et al., which showed a higher percentage of children (88.6%) with Group E and D staging [18]. In a study by Waddell et al., nearly all diagnoses (\( n = 282 \)) in the first affected eye were Group E or already had extraocular disease, with only 2 children having IIRC Group C eyes [10]. The lower percentage of findings in our study compared to that of Waddell et al. may be attributed to the awareness campaign for primary health care workers together with patient facilitation, which was recommended.

In our study, management included all treatment received by the children in the study prior to and after UCI admission. Forty-nine (54.4%) had received treatment prior to UCI admission, with 42 (46.7%) having undergone

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**Figure 1:** Graph showing the 3-year survival of children with retinoblastoma from the time of admission to the Uganda Cancer Institute.

**Figure 2:** Graph showing the 3-year survival of children with retinoblastoma by history of eye swelling.

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In our study, management included all treatment received by the children in the study prior to and after UCI admission. Forty-nine (54.4%) had received treatment prior to UCI admission, with 42 (46.7%) having undergone
enucleation. At the last follow-up, an additional 10 (11.1%) children underwent enucleation after UCI admission.

Vision and globe salvage were attempted in 13 (11.3%) and 15 (13%) eyes, respectively. However, the only treatment option utilized was chemotherapy. The majority (68, 75.6%) of children received chemotherapy, 43 (47.8%) received neo-adjuvant chemotherapy, and 18 (20%) underwent radiotherapy. Seventy-five (83.3%) out of the 90 patients admitted

| Variable                                  | CHR (95% CI) | AHR (95% CI) | P value|
|-------------------------------------------|--------------|--------------|--------|
| Age group at first sign in months         |              |              |        |
| 0 to 12                                   | 1            | 1            |        |
| 13 to 36                                  | 1.08 (0.48 – 2.40) | 0.89 (0.33 – 2.26) | 0.772 |
| Over 36                                   | 1.08 (0.45 – 2.58) | 0.40 (0.14 – 1.13) | 0.084 |
| Sex: Male                                 | 1.04 (0.52 – 2.04) | 1.12 (0.54 – 2.31) | 0.769 |
| Eye loss                                  |              |              |        |
| No                                        | 1            | 1            |        |
| 1 eye enucleated                          | 1.40 (0.67 – 2.94) | 1.12 (0.51 – 2.45) | 0.779 |
| Both eyes enucleated                      | 3.94 (1.64 – 9.46) | 3.53 (1.33 – 9.36) | 0.011 |
| Metastatic spread                         |              |              |        |
| No                                        | 1            | 1            |        |
| Yes                                       | 9.24 (3.88 – 22.02) | 8.93 (3.23 – 24.69) | 0.001 |
| Lag time between first sign and UCI visit in months |              |              |        |
| 0 to 6                                    | 1            | 1            |        |
| 7 and above                               | 0.18 (0.07 – 0.42) | 0.10 (0.04 – 0.25) | 0.001 |
| Proptosis                                 |              |              |        |
| No                                        | 1            | 1            |        |
| Yes                                       | 1.65 (0.82 – 3.31) | 1.35 (0.52 – 3.48) | 0.535 |
| Eye swelling                              |              |              |        |
| No                                        | 1            | 1            |        |
| Yes                                       | 2.62 (1.10 – 6.27) | 4.39 (1.38 – 14.01) | 0.012 |
| Eye reddening                             |              |              |        |
| No                                        | 1            | 1            |        |
| Yes                                       | 1.65 (0.73 – 3.70) | 0.36 (0.14 – 0.94) | 0.036 |

Figure 3: Graph showing the 3-year survival of children with retinoblastoma by metastasis.
into care at UCI were initiated on prescribed treatment. The follow-up duration for the majority (59, 65.6%) of children was less than 6 months from the time of the first UCI visit. The utilization of chemotherapy for Rb treatment, alongside other treatment modalities, has been embraced at UCI for the majority of children, which corresponds with current treatment recommendations [19].

At the 3-year follow-up from the first UCI visit, 37 (41.1%) of all children in our study were alive, and 33 (36.7%) of children with retinoblastoma had died. The number of those lost to follow-up was 20 (22.2%). The 3-year survival observed in our study (41.1%) was significantly lower than that reported in Taiwan (64.41%), Turkey (89.6%), and Iran (94.8%) [17, 20, 21]. The significantly low 3-year survival in our study could be attributed to advanced disease at first UCI admission. Although enucleation was one of the most utilized treatment modalities in our study (57.8%, n = 52), it was lower than that in studies in India by Chawla et al. (95.7%), in a country-wide study in Uganda by Waddell et al. (96%), and in Iran by Nabie et al. (95.7%) [8, 10, 16]. This finding can be explained by the increased utilization of neoadjuvant chemotherapy at UCI. Forty-nine (54.4%) and three (3.3%) of the children in our study had unilateral and bilateral enucleation, respectively. The lower percentage of enucleation in our study may be attributed to the noncompletion of prescribed treatment options, which was recorded in 61 (67.8%) children.

At 3 years of follow-up from first admission at UCI, 12 (13.3%) and 64 (71.1%) children in our study had bilateral and unilateral vision loss, respectively, including those who had enucleation in the affected eye(s). The significant number of children with vision loss in our study is attributed to the unavailability of vision salvage treatment modalities at the Uganda Cancer Institute during the study period.

The number of children lost to follow-up in our study (20, 22%) was slightly lower than that reported in a previous retrospective study (32%) of childhood cancer patients at the Uganda Cancer Institute by Mutyaba et al. [22]. This may be attributed to the recent introduction of a functional oculocutaneous oncology unit and personnel at UCI.

In bivariate analysis, the predictors of survival in our study were a history of eye swelling, a lag time of 7 months or more, extra ocular retinoblastoma in the affected eye(s), advanced disease stage (IIRC Group E) in the worst affected eye, completion of prescribed treatment, metastatic retinoblastoma, and bilateral enucleation. These predictors were similar to findings in a study in China by Gao et al., which showed that extraocular disease, treatment abandonment, bilateral disease, and advanced tumour stage were predictors of survival in children with retinoblastoma [23]. The above predictors indicate that most children with retinoblastoma seek medical attention at the advanced stages of disease and are referred to tertiary oncology centres with metastatic disease.

5. Conclusion

Children with retinoblastoma were referred to the Uganda Cancer Institute with advanced disease. In addition, the mortality of retinoblastoma patients at UCI was high, as evidenced by the very low 3-year survival. Thus, there is an urgent need for the early involvement of a multidisciplinary team of health workers in the management of retinoblastoma, including paediatric ophthalmologists, paediatric oncologists, oculist oncologists, radiation oncologists, and nurses, for better treatment outcomes.

Data Availability

Data is available on request through the author (email: kabikarakuth@gmail.com).

Ethical Approval

This study was approved by Makerere University School of Medicine Research Ethics Committee (#REC REF 2020-031).

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this study.

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