Pre-bacute lymphoblastic leukemia: case report and literature review

* Agustín Tiol-Carrillo, Patricia Enzaldo-de la Cruz*

ABSTRACT

Leukemia is one of the most common oncological disorders found in the first decade of life. It is characterized by the excessive production of non-functional immature lymphocytic cells, called blasts, which invade the bloodstream causing fatal consequences. Unless timely treated, it is a deadly, severe condition. Its more common manifestations are the following: generalized paleness, spontaneous hemorrhaging, vascular lesions such as bruises and petechiae, general malaise, weight loss, and specific stomatological manifestations such as oral mucosa paleness, leukocyte infiltrate gingivorrhage or apparition of petechiae in some locations of the mouth. The present article documents the case of a five year old girl, diagnosed with B-cell precursor acute lymphoblastic leukemia. The child was subjected to oral rehabilitation under general anesthesia due to the fact that she exhibited multiple infectious foci in her teeth which contraindicated initiation of chemotherapy treatment.

Key words: Hospital dentistry, leukemia, chemotherapy, oncologic cancer patients in dentistry.

INTRODUCTION

Leukemia is the most common oncologic disorder found in childhood. It is characterized by a bone marrow disorder which generates excessive production of immature cells called blasts. In order to be classified, leukemia exhibits great diversity of characteristics based on type of producing cells or maturation degree of these cells.

According to producing cell lineage leukemia can be lymphoblastic or myeloblastic. Lymphoblastic leukemia is that whose cellularity derives from the lymphoid lineage, that is to say, Lymphocytes B and T respectively, whereas myeloblastic leukemia derives from myeloid lineage cells such as red blood cells, neutrophils, basophils, eosinophils and platelets (Figure 1). Thus, according to cell functionality, they can be differentiated into acute and chronic.

Acute leukemia is characterized by possessing a non-functional cell population, since cells are totally immature, differing from chronic leukemia where cells exhibit greater maturation degrees. From this premise we can infer that acute leukemia is generally more aggressive than chronic leukemia. The origin of this condition is strongly associated to genetic syndromes such as Down’s syndrome.

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Lifraumeni’s syndrome, Klinefelter syndrome, Wiskitt Aldrich disease, Fanconi anemia, early age exposition to ionizing radiations and chemical agents as well as specific cytogenetic predisposition such as presence of Philadelphia chromosome.\textsuperscript{1,2} Lymphoblastic acute leukemia precursor of B cells is the most commonly found in Mexican pediatric population.

**CLINICAL CASE**

A five year old female, coming from León, Guanajuato, with previous diagnosis of B cell precursor acute lymphoblastic leukemia was referred to receive dental treatment. The oncologist deemed dental treatment urgent so as to achieve dental rehabilitation previous to initiation of chemotherapy treatment (Figure 2). Medical history revealed that the patient had begun with generalized paleness, weakness, bone-ache, generalized petechiae in the back, thorax and limbs. After initial pediatric assessment she was immediately referred to the oncologist on suspicion of leukemia. Due to the nil cooperation of the patient, to the complexity of her condition, as well as the urgency present, it was decided to rehabilitate the patient under general anesthesia (Figure 3).

Before dental treatment under general anesthesia, relevant studies were conducted to prepare the patient pre-operatively. Laboratory tests of blood count and clotting times were requested.

Blood count tests revealed anemia with 9 g/dL hemoglobin, neutropenia with neutrophil count of 500/mm\(^3\) and thrombocytopenia with platelet count of 50,000/mm\(^3\).
Previous to procedure initiation, the patient received a platelet apheresis to obtain maximum regulation of platelet levels, achieving elevation to up to 90,000/\( \text{mm}^3 \). Once in the operating theatre, dosed antibiotic prophylaxis was administered (20 mg/kg/dose) with intravenous clindamycin. The patient was intubated by naso-tracheal route.

Sterile fields were conventionally placed and arches were fully isolated with a rubber dam (Figure 4). The following dental treatments were performed: pulpotomies of upper first molars, stainless steel crowns in upper and lower first molars, resin restoration in lower second molars, and pit and fissure sealers in upper second molars.

In views of the fact that the patient was to be subjected to a chemotherapy treatment which would significantly immunosuppress her, it was decided to preventively extract upper lateral and central incisors, so as to avoid latent risk of odontogenic infection which could aggravate her health circumstances. At that time, it was decided against placement of esthetic devices to replace extracted teeth, since appliances could act as bacteria reservoir, which at a given time could complicate the patient’s clinical evolution (Figure 5).

Because of preventive care exerted before treatment, the patient evolved satisfactorily, and was remitted to the recovery room in order to monitor vital signs.

Once the procedure was completed, the patient was ready to initiate chemotherapy as soon as possible, after healing of wounds caused by extractions.

**DISCUSSION**

All leukemia types belong to the group of proliferative malignant diseases. They represent the most common cancer disorders found in childhood, in addition to retinoblastoma, osteosarcoma and neuroblastoma.\(^3\) As mentioned before, leukemia can be classified according to producing cell lineage and to functionality and cell differentiation degree.\(^1,3\)

According to statistical studies conducted by the Health Ministry (Secretaría de Salud) in 2001, approximately 13,500 patients were admitted in hospitals with leukemia diagnosis. These cases were fatal in 232 patients aged one to four years, and 558 fatalities were recorded in the group of children aged five to fourteen years.\(^4,5\) Acute lymphoblastic leukemia is characterized by having its onset in the first decade of life, nevertheless, it is possible that the frequency might increase in senior citizens. Immature cells, which under normal circumstances would differentiate from any cell of lymphoid or myeloid lineage are called blasts.\(^6\) These cells, being non-functional, should not
be found in the bloodstream, nevertheless, when the patient first displays leukemia, an excessive blast population appears which invades the bone marrow of long bones, preventing thus normal reproduction of other well-differentiated cells and their installation in the space within the bone marrow, generating thus in the patient a decrease in cell counts (pancytopenia) as well as pain in arms and legs. Since platelets and hemoglobin experience a sharp decrease, the patient experiences anemia and thrombocytopenia, which clinically translates into generalized paleness, asthenia, adynamia, and vascular lesions in the skin and oral mucosa. All patients first experiencing leukemia suffer certain associated syndromes, which can be: febrile, infiltrative, hemorrhagic, anemic, tumor and consumptive syndromes (Table I).

Since this is a disease derived from white blood cells, it is common to find leukocyte infiltrate in the oral mucosa; this represents the invasion of leukocyte neoplastic cells which penetrate into the oral mucosa to then destroy it. Clinically, they are observed as whitish lesions that do not detach when scraping. Leukocyte infiltrate can be a subclinical manifestation that a patient is suffering leukemia (Figure 6).

Medical diagnosis of leukemia is based on characteristic symptomatology as well as laboratory studies prescribed for the patient (Table II). Bone marrow aspiration is generally indicated to corroborate diagnosis, in order to assess whether blast invasion is absent or present; the aforementioned diagnosis might be emitted in cases where there is medullar blastosis exceeding 30% of total cell population. Beside cell counts revealed by laboratory tests, characterized by exhibiting high leukocyte numbers (exceeding 50,000 mm$^3$) as well as reported presence of blasts, there is important decrease in the count of platelets, neutrophils, hemoglobin and erythrocytes.

Cytogenetics is very important to determine leukemia prognosis. When cell with larger numbers of chromosomes (called hyperdiploid) are observed in a cytogenetic study, patients enjoy greater possibilities of healing, differing from those patients with lesser amounts of chromosomes (hypodiploid) who generally suffer worse prognosis. Likewise, the presence of the so-called Philadelphia chromosome is a factor which determines a poor prognosis for this disease.$^4,^5,^7$

Nevertheless, besides cytogenetics there are some other factors which are related with leukemia prognosis (Table III).

Once the patient has been diagnosed, chemotherapy treatment must be undertaken as soon as possible, nevertheless this treatment cannot be initiated in cases when the patient presents infectious foci in the mouth, therefore, the dentist plays an essential role in the treatment of these patients.$^8$

Dentists must be familiar with chemotherapy phases to which the patient with this disease will be subjected to, since, at each given phase there are specific drugs an sequels which might contraindicate of warrant a modification in the dental treatment course chosen for the patient. When the child has not been subjected to chemotherapy treatment, the dentist must evaluate blood

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**Table I. Early signs and symptoms of patients with leukemia.**

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Signs and symptoms</th>
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<tbody>
<tr>
<td>Febrile</td>
<td>Constant fever up to 40 °C which does not abate with antipyretic drug administration. Associated to infection susceptibility and oral ulcers</td>
</tr>
<tr>
<td>Infiltrative</td>
<td>Invasion of leukemia cells to different tissues and organs such as bones, kidney or liver</td>
</tr>
<tr>
<td>Hemorrhagic</td>
<td>Due to the severe thrombocytopenia exhibited by the patient it is common to suffer epistaxis, ecchymosis, petequeia in oral mucosa and in conjunctive eye’s as well as gingivorrhagia</td>
</tr>
<tr>
<td>Tumoral</td>
<td>Hepatosplenomegaly caused by blast invasion in the liver, spleen, lymphs and gonads. Presence of adenomegalia in more than two locations</td>
</tr>
<tr>
<td>Consumptive or due to wear</td>
<td>Excessive loss of corporal weight, weakness and high fever</td>
</tr>
</tbody>
</table>

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**Figure 6.** Leukocyte infiltrate in the gingival mucosa of a 15 year old patient who started with acute B-lymphocytic leukemia. All orthodontic appliances were removed from this patient before initiation of chemotherapy treatment.
count and coagulation (clotting) times to assess whether the patient requires platelet apheresis, or prophylactic antibiotic administration before invasive dental treatments due to the important decrease of defense cells. At this stage it is also necessary to completely remove any orthopedic or orthodontic device worn by the patient, since they could become a plaque reservoir which might affect the patient’s evolution once immunosuppression is triggered by chemotherapy treatment.

The first immunotherapy phase is called remission induction.2,3,8 During this phase, the patient receives high doses of corticosteroids, such as methylprednisolone, with the sole aim of immunosuppressing him and initially arrest the disease’s evolution; if the patient evolves satisfactorily, the next stage will be undertaken. This stage is called consolidation or intensification stage,2,8 and in it, in addition to administering corticosteroids, chemotherapy drugs are also administered, among these drugs we can count vincristine, citarabine, daunorubicin, L-asparaginase, etoposides, cyclophosphamide, and methotrexate. After having ingested these drugs, the patient experiences specific sequels such as anagen effluvium, weight loss, nausea, vomit, and generalized paleness (Figure 7). After a week of ingesting these drugs, the patient will experience an ailment called seventh day neutropenia, or nadir phase, which is characterized by under 200 neutrophil counts, sometimes even reaching 0 neutrophils per cubic millimeter.

During this phase the patient might exhibit the most painful chemotherapy sequel in the mouth, called mucositis. Mucositis tends to appear as a consequence of methotrexate administration and is characterized by ulceration and desquamation of all the digestive tract mucosa, from the mouth to the anus. The patient experiences decrease to oral route tolerance in a hundred per cent, the dentist should prescribe use of palliative treatment to help the patient tolerate oral intake. The treatment can be mucosa coaters or topical antihistamines which will decrease local inflammation of the ulcerated oral mucosa (Figure 8).9-12

It is at this precise moment due to the fragile circumstances of the child, when the dentist must suppress all treatment. Any ailment must be treated palliatively (pain killers, antibiotics or antifungal in necessary cases) waiting for a normalization of cell count in the following days.10,13

Once this phase is overcome, the patient is subjected to a new bone marrow aspiration and another cell count, where important reduction of

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### Table II. Diagnostic studies performed in a leukemia-affected patient.

<table>
<thead>
<tr>
<th>Diagnostic study</th>
<th>Study justification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blood cell count</td>
<td>Assessment of different types of cells and their microscopic appearance</td>
</tr>
<tr>
<td>Bone marrow aspiration and biopsy</td>
<td>To be obtained from the iliac bone to observe or discard the presence of blasts in the bone marrow</td>
</tr>
<tr>
<td>Other blood tests</td>
<td>Hepatic and renal function tests in order to discard leukemic cell invasion in these organs</td>
</tr>
<tr>
<td>Lumbar puncture</td>
<td>To discard presence of leukemic cells in the cerebrospinal fluid</td>
</tr>
<tr>
<td>Flow cytometry</td>
<td>Determine with high specificity the type of leukemia afflicting the patient through the use of antibodies exclusively adhering to certain types of leukemic cells</td>
</tr>
<tr>
<td>Cytogenetics</td>
<td>Detection of chromosomal anomalies such as translocation, assistance to establish disease’s prognosis, observing whether cytogenetically cells are sensitive or resistant to chemotherapy</td>
</tr>
</tbody>
</table>

### Table III. Factors affecting leukemia patients’ prognoses.

<table>
<thead>
<tr>
<th>Prognosis factor</th>
<th>Justification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>Children under 1 year and over 10 years of age suffer worse prognosis than children aged 1-9 years</td>
</tr>
<tr>
<td>Gender</td>
<td>Males suffer worse prognosis than females due to the possibility of relapse in the testicles</td>
</tr>
<tr>
<td>Ethnicity</td>
<td>Children of African or Hispanic ethnicity suffer worse prognosis than Caucasian children</td>
</tr>
<tr>
<td>White cell count</td>
<td>A count of 50,000 mm³ or over is indicative of a poor prognosis</td>
</tr>
<tr>
<td>Extra-medullar disease</td>
<td>In cases when there is invasion of leukemic cells into central nerve system or testicles</td>
</tr>
<tr>
<td>Response to treatment</td>
<td>Favorable prognosis can be assumed when the child suitably responds to the early phases of treatment.</td>
</tr>
<tr>
<td></td>
<td>Conversely, poor prognosis can be established when the child does not respond to initial treatment</td>
</tr>
</tbody>
</table>
blast cells is expected, which would indicate suitable response to the treatment.

After this, the patient passes to the following and next to last phase called maintenance phase. In it corticosteroid and chemotherapeutic agents are periodically administered in order to control the disease. A patient can last several years in this phase, and obviously under sporadic laboratory control studies (blood count, bone marrow aspiration and lumbar punctures). This patient can be treated in the dental office, always taking into account possible sequels to administered drugs, antibiotic prophylactic coverage might be required in some cases.

When the patient has overcome the disease, and having spent a minimum of two years with low doses of drugs and total absence of blasts (as revealed in periodical tests), the last phase of the oncological treatment is undertaken, this is the surveillance (monitoring) phase. All subjects in their monitoring phase must undergo yearly periodical revision for the rest of their life. Nevertheless, it must be pointed out that a patient in monitoring phase, can be treated as a healthy patient from the dental point of view.

Leukemia, once overcome, exhibits an important relapse risk; this is due to specific situations which must be foreseen before initiating chemotherapy treatment. Leukemic cells tend to penetrate into the so-called «sanctuary organs», which are the central nervous system and the testicles in males. These are zones where chemotherapeutic drugs are ineffective, and thus become the anatomical areas suffering higher risks of relapse.\textsuperscript{2,3,14} Therefore intrathecal therapy is mandatory in the treatment of a leukemia patient, as well as performance of several lumbar punctures during treatment, conducted with the aim of discarding blast invasion in the central nervous system.\textsuperscript{14,15}

CONCLUSIONS

Dentists play a key role in the diagnosis and treatment of a leukemia patient. Many leukemia-indicating early lesions can appear in the mouth, for this reason, dental personnel is obliged to be familiar with them. Dental intervention is of the utmost importance due to the need to eradicate infectious foci before initiating cancer therapy, as well as treating oral and dental problems which might arise as a consequence of chemotherapy. Knowledge of the chemotherapy phases to which a patient will be subjected, to a great extent determines the dental and stomatological conduct that must be followed.

REFERENCES


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