

classified as severe (27 patients) or moderate (12 patients) with FVIII:C dosage <3%. After collection, samples were processed for genomic DNA extraction. First, patients were genotyped for intron 22 inversion by inverse PCR. Patients who were negative for intron 22 inversion were genotyped for intron 1 inversion. Samples negative for both inversions will be full-length NGS sequencing. Among 39 patients, 21 (54%) were diagnosed with intron 22 inversion and one (2%) with intron 1 inversion. In addition, among 12 patients previously diagnosed with moderate HA, seven (58%) patients were reclassified as severe HA because they presented intron 22 inversion. The molecular diagnosis proved to be important for the correct classification of the severity of HA patients. Due to the diagnosis normally based only on residual activity of plasmatic FVIII, the disease severity classification may be incorrect. It is believed that the high variability of coagulometric techniques and the treatment of patients prior to laboratory diagnosis may affect the correct classification of HA patient severity. This is an ongoing study and the number of participants needs to be increased in order to draw conclusions. We thank all participants, Fapemig, Hemominas, and the CGSH of the Ministry of Health.

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QUALITY OF LIFE OF HEMOPHILIA A FEMALE CARRIERS OF INTRON 22 INVERSION

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Assessing the quality of life of family members of patients with hemophilia A (HA) is necessary given the emotional overload and dedicated care of their family members. The 36-item Short Form Health Survey (SF-36) measure quality of life related to health using eight health concepts: physical functioning (PF), limitations due to physical health (PH), pain (P), general health (GH), energy/fatigue (EF), social functioning (SF), limitations due to emotional problems (EP), and emotional well-being (EW). The aim of this study was to analyze the quality of life profile of family members of patients with severe hemophilia A with intron 22 inversion mutation treated at Fundação Hemominas. To date, sixteen participants have been enrolled in the study after signing an informed consent form. Participants answered a 36-item short health questionnaire and collected peripheral blood samples in citrate tubes. Samples were processed for DNA extraction and genotyping. Participants were divided into two groups: Carriers of the intron 22 inversion and participants who were not carriers of the intron 22 inversion mutation. Among carriers the median (MED) scores were: 88% (IQR 77.5%-96%) for PF; 100% (IQR 56.3%-100%) for PH; 61% (IQR 61%-72%) for P; 87% (IQR 77%-92%) for GH; 60% (IQR 38.8% - 66.3%) for EF; 69% (IQR 47% - 75%) for SF; 100% (IQR 24.8% - 100%) for EP; and 70% (IQR 59% - 76%) for EW. Among participants who were not carriers, 93% (IQR 81.3% - 95%) for PF; 88% (IQR 18.8% - 100%) for PH; 72% (IQR 58.3% - 88%) for P; 67% (IQR

47%-80.3%) for GH; 58% (IQR 41.3%-70%) for EF; 69% (IQR 52.8%-91%) for SF; 84% (IQR 0%-100%) for EP; and 68% (IQR 54%-78%) for EW. Interestingly, carriers presented a significant higher score in general health domain ($p=0.03$). In conclusion, this study is important to assess the quality of life in relatives (carriers and non-carriers) of HA patients. This is an ongoing project and the number of participants needs to be increased in order to draw conclusions. We thank all participants, Fapemig, Fundação Hemominas, and the CGSH of the Ministry of Health.

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INVESTIGATION OF INHERITANCE BY MOLECULAR DIAGNOSIS IN A FAMILY WITH NO FAMILY HISTORY OF HEMOPHILIA A: CASE REPORT

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Hemophilia A (HA) is classified according to residual circulating FVIII (FVIII:C). Patients with FVIII:C 5%-40% are classified as mild HA; patients with FVIII:C 1%-5% are considered moderate; and patients with FVIII:C <1% are classified as severe HA. Patients with severe HA have two common mutations: Inversions of introns 1 and 22 of the F8 gene. The aim of this study is to report the case of a patient diagnosed with HA without any family history of the disease and treated in Fundação Hemominas in 2022. Male patient, NB, presented an extensive hematoma on the back after 24 hours of delivery. A blood count was performed, which reveals a hemoglobin value of 5 g/dL (reference value approximately 16 g/dL). Patient is admitted to the ICU where a blood transfusion and coagulometric tests are performed, indicating HA. The patient is referred to the pediatric hematology unit of Fundação Hemominas. Coagulation tests confirmed the diagnosis of severe HA. The patient was referred for molecular diagnosis of the disease. In addition, three members of the patient's family agreed to participate in the molecular diagnosis. The family members answered two questionnaires: 1) bleeding assessment score - ISTH (BAT-ISTH); and 2) arthralgia assessment. Peripheral blood was drawn from all participants in a citrate tube, and samples were processed for genomic DNA extraction. Genotyping of the intron 22 inversion mutation was performed by inverse PCR. The patient was positive for intron-22 inversion, confirming the diagnosis of severe hemophilia A. The patient's mother (34 years old), with a bleeding score of 3, without joint pain, was diagnosed with the same inversion and received genetic counseling when the report was handed over. The patient's aunt (30 years old), with a bleeding score of 0, without joint pain, had a negative result for intron inversion 22. The patient's grandmother, 59 years old, with a bleeding score of 1, who presented right knee pain, also had a negative result. Evaluation of the patient's three-generation pedigree suggests that the patient inherited the de