Abstract
Although hyperferritinemia is not specific, it may be diagnostically very contributing if well interpreted by the physician. Physiologically, ferritin is a protein synthesized essentially by the liver and intended to store iron in the liver, macrophage and erythrocyte. Despite this, the increase in ferritinemia does not necessarily mean iron overload or liver disease. A global approach integrating the degree of ferritinemia, anamnesis and clinical context makes possible in the vast majority of cases to find the cause of hyperferritinemia without going through the saturation factor of transferrin and thorough liver investigations. The use of transferrin saturation coefficient, liver imaging, genetic tests or even liver biopsy proves to be necessary only in rare cases where the most obvious causes could not be identified.

Introduction
Unlike hypoferritinemia, which almost constantly reflects iron deficiency, hyperferritinemia is often difficult to interpret in the absence of an obvious clinical context. In medical practice, major hyperferritinemia greater than 1000 μg/l is a classic situation. Moore et al. had published in 2013 the results of a 2-year study in a department of internal medicine that found a major hyperferritinemia in 627 patients with the most common causes of cancer [1]. Vardi et al. found similar results with a pejorative prognostic value of severe hyperferritinemia, regardless underlying cause [2]. Another much older study by Lee et al. found similar frequencies of hyperferritinemia in the previous two studies with high frequency of etiological associations in the same patients [3]. The high frequency of hyperferritinemia, their wide etiological spectrum and the frequent concomitant presence of several etiologies reflect the need to address hyperferritinemia based on a global vision prioritizing the most obvious and serious causes.

Iron metabolism and some definitions according gender
Hyperferritinemia is defined by serum values >400 μg/l in men and >300 μg/l in women. According to NHANES III, values generally found in the general population rarely exceed 200 μg/l in men and 150 μg/l in women. Furthermore, in a study by Murtagh et al. on patients with confirmed hereditary haemochromatosis, ferritin levels were between 300 and 3000 μg/l for men and between 250 and 3000 μg/l for women. The interpretation of the limit values for ferritinemia must therefore be extremely cautious, especially in the absence of a plausible condition explaining these values. A ferritinemia between 300 and 400 μg/l in men and between 200 and 300 μg/l in women can be synonymous with a real overload martial imposing a close biological monitoring or a thorough assessment from the outset. On the other hand, a ferritinemia less than 100 μg/l in an inflammatory context or any other hyperferritinemic situation should be considered a potential iron deficiency and managed as such [4,5].

A small amount of exogenous iron not exceeding 2 mg/day is absorbed by the duodenum and partially stored in the enterocytes as ferritin. Most of the body’s iron comes from a recycling phenomenon through macrophages that provide physiological hemolysis and degradation of senescent hemoglobin. This iron is subsequently transported to the bone marrow at a rate of 20 mg per day but also and especially to the hepatocytes which are the main storage site for ferritin. Excretion is through enterocytes and menstruation. Ferritin is a protein synthesized by the liver in the form of an egg about 12 nm in diameter able to store up to 5000 iron atoms. It is present mainly in hepatocytes, macrophages and cells of the red line but remains extremely ubiquitous topographically and functionally, which explains its elevation in several etiologies including hepatopathies, inflammation and cytolysis in addition to overload martial [6].

Main causes of hyperferritinemia
Hereditary haemochromatosis
The absorption of dietary iron is done by the enterocyte through the carriers "DMT1" and "Ferroportin". Subsequently, passage through the circulation, transferrin binding and interaction with the transferrin receptor (TFR) of the cryptic cell and the hepatic HFE protein occur. Ferroportin is also present on the surface of the macrophage. In the healthy person, there is a sufficient production of hepcidin which has as a function the negative regulation of the activity of DMT1 and ferroportin and therefore the limitation of the passage of iron in the blood from the enterocyte and macrophage. In the person with HFE mutation, there are 3 essential abnormalities: 1) Alteration of the HFE-TFR interaction resulting in less iron entry into the cryptic cell and the emission of a false deficiency signal resulting in an absorptive response of iron; 2) A release of enterocyte iron to the plasma by hyperactivity DMT1 and ferroportin; 3) A release of macrophagic iron by hyperactivity of macrophagic ferroportin. The consequence of these

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abnormalities is an increase initially in the transferrin saturation coefficient (TSC) and then in ferritin with an accumulation of tissue iron and the occurrence of organic damage, particularly liver, joint, glandular, cardiac and cutaneous explaining the characteristic triad “cirrhosis-diabetes-melanoderma”. The first manifestations of the disease occur in adulthood with later onset in women because of the protective role played by menstruation. In practice, the diagnosis is based on the elevation of the TSC, the elimination of other obvious causes of this elevation such as cytosis and hepatopathy and the demonstration of a homozygous mutation HFE C282Y or composite C282Y/H63D. The search for other mutations on the genes of the HFE or TFR receptors will be considered in rare cases where the table is typical and very evocative with negativity of the usual mutations. Hepatic biopsy has been less and less necessary since the advent of genetic tests, as well as liver MRI, which is very efficient in assessing hepatic parenchymal iron overload [7].

Metabolic syndrome

Abdominal obesity, arterial hypertension, glycemic and lipid disorders and microalbuminuria are the criteria taken into account by WHO to define the metabolic syndrome whose main pathophysiological axis is insulin resistance. The very high prevalence of hyperferritinemia in patients with metabolic syndrome is currently established. The COALUS study published in 2008 focused on 3063 women and 2752 men and had clearly demonstrated this finding (2.2% vs. 11.4% for women and 25% vs. 40.6% for men). This close epidemiological association is supported by the study of Brudevold et al., which showed a proportional increase in serum ferritin levels as a function of the level of peptide C, apart from any other cause of iron overload [8].

One of the main conditions associated with the metabolic syndrome is steatosis and non-alcoholic steatohepatitis where it is currently demonstrated the presence of several hepatocyte disorders related to insulin resistance with a cellular uptake of iron clearly superior to the externalization. The occurrence of hepatic inflammation in the context of steatohepatitis gives rise to cytosis and may in this way aggravate hyperferritinemia. The predominance of tissue and cell abnormalities explains why the hyperferritinemia of the metabolic syndrome is only rarely accompanied by an increase in the CST, whereas hepatic MRI often reveals an obvious overload [9].

Cytolysis

The hyperferritinemia of cytosis is explained by the ubiquitous presence of ferritin in hepatocytes, monocytes, erythrocytes and muscle cells, as well as pancreatic, testicular, bronchial, placental and renal cells. Hyperferrinemias of cytosis are often multifactorial and should not be “easy” on cell lysis before ruling out other potential associated causes. In a study of Kotok et al. performed on 100 patients with acute hepatitis, the ferritin level correlated closely with that of ALT but with a particularly severe elevation in the group of viral hepatitis thus implying a participation of anti-macrophage activation. Another study had shown elevated ferritin levels in patients with dermatomyositis but with a greater increase in the subgroup with more than one interstitial lung disease (790 vs 186 ng/ml, P<0.0001). Finally, hyperferritinemia of hemolytic anemia is more marked in thalassemia than in other causes of hemolysis because of the particularities of iron hyperabsorption and transfusion requirements in thalassemic individuals. TSC during cytosis is classically elevated due to massive plasma release of intracellular iron, particularly hepatic and erythrocyte iron [10-12].

Inflammation

Increase of ferritinemia is related to both direct stimulation by prostaglandins and lipopolysaccharides and indirect through the increase of NO and oxidative stress induced by pro-inflammatory cytokines. In a recent study of Vanarsa et al. in lupus patients, it was clearly demonstrated in the active forms of lupus that ferritin was increased not only in the serum (p<0.0001) but also in urinary sites, particularly in urine, in patients with nephropathy (p<0.0001). Inflammatory ferritinemia does not typically exceed 1000 ng/ml and is not accompanied by an elevation of the TSC. Two situations are important to differentiate: on the one hand hyperferritinemias higher than 1000 ng/ml which would orient more towards the Still’s disease, the serious septic states and the hemophagocytosis syndromes and on the other hand the inflammations with “normal” ferritin which must lead to take out masked marital deficiency [13-16].

Alcohol consumption

The link between alcohol and hyperferritinemia has been known for a long time and is well understood physiopathologically. Alcohol leads to direct stimulation of ferritin synthesis regardless of its pathogenic effect on the liver. Significant alcohol consumption should be sought in any patient with hyperferritinemia. Alcohol withdrawal leads to a rapid decline in ASAT, GGT and ferritinemia within 15 days of cessation of alcohol consumption. This test is both diagnostic and therapeutic. The biological phenotype of alcoholic hyperferritinemia is a classic absence of increased TSC with inconsistent hepatic iron overload on MRI, whether or not there is already established alcoholic liver disease [17].

Still’s disease, Anti-phospholipids and Haemophagocytic syndrome

A major hyperferritinemia greater than 1000 is usually directed to particular etiologies like still disease, macrophage activation syndrome and catastrophic syndrome of anti-phospholipid antibodies (CAPS). In Still’s disease, functional prognosis is related to joint damage and life-threatening hepatic involvement, DIC, SAM, myocardial involvement and amyloidosis. The absence of major hyperferritinemia in Still’s disease simply requires rethinking the diagnosis. The TSC has no value because the clinic is immediately very evocative. On the other hand, glycosylated ferritin less than 20% is a relevant and highly characteristic element that is one of the major diagnostic criteria for the disease (Table 1) [18].

In the macrophage activation syndrome, there is a real cytokine storm linked to an initial overactivation of T lymphocytes of genetic, infectious, tumoral or autoimmune origin, an activation in the second time of macrophages under the effect of interferon, TNF alpha, interleukin-6 and M-CSF and finally increased phagocytosis by activated macrophages of mature cells including figured elements of the blood. This macrophage activation in turn results in a positive feedback on T lymphocytes and a loop co-stimulation process that maintains the inflammatory and phagocytic phenomena. In the same way as in still disease, the clinical picture here is very obvious and diagnosis can be made without TSC. In practice, the clinical picture is that of a febrile tumor syndrome with pancytopenia, hypertryglyceridemia, elevation of LDH and signs of haemophagocytosis in the medullary study or lymph node biopsy (Table 1). With a mortality approaching 60%, the simple suspicion of a SAM should lead to an immediate and the widest possible treatment targeting the most common causes namely bacterial, fungal, herpetic and tuberculous infections as well as malignancy including lymphomas [19].
Table 1. Diagnostic criteria of still disease, CAPS and hemophagocytosis syndrome

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<td>Major criteria</td>
<td>3 organs damage</td>
<td>Fever≥38.5, ou≥7 days</td>
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<tr>
<td>Fever</td>
<td>Simultaneous occurrence in less than a week</td>
<td>Cytopения≥2</td>
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<td>Joint pain</td>
<td>Histological microvascular occlusion</td>
<td>Ferritin≥1000</td>
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<tr>
<td>Intermittent erythema</td>
<td>Presence of antiphospholipid antibodies</td>
<td>LDH≥1000</td>
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<tr>
<td>Pharyngitis</td>
<td>Diagnosis if presence of the 4 criteria</td>
<td>Bone marrow, splenic, hepatic or lymphoid</td>
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<td>Neutrophilia≥80%</td>
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<td>haemophagocytosis,</td>
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<tr>
<td>Glycated ferritin≥20%</td>
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<td>Main causes: CMV, VHI, HSV, EBV, Mycobacteria,</td>
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<tr>
<td>Minor criteria</td>
<td></td>
<td>Lymphoma, Solid malignancy</td>
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<td>Maculopapular rash</td>
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<tr>
<td>Leucocytosis≥10000/mm³</td>
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<td>4 major criteria or 3 major+2 minor criteria</td>
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Finally, the CAPS carry out a thrombotic microangiopathy table which clinically results in a multi-organ failure related to a generalized activation of the micro and macrovascular thrombotic cascade [20]. Mortality is also very important even with a rapid and complete therapeutic intervention.

Table 2 summarize in a simplified and didactic way the main pathophysiological mechanisms of hyperferritinemias according to the etiologies as well as their implications on the paraclinical profile of patients.

Conclusion

It is necessary to separate the etiologies of hyperferritinemia into 3 main essential categories:

- Causes that are quickly detectable by interrogation, clinic and simple paraclinical examinations and whose management is often easy and allows a fairly rapid reversibility by simple measures of eviction and etiological treatment.
- Causes whose clinical presentation is very telling and quickly leads to a rather narrow and specific etiological spectrum. These are most often serious causes that must be sought first.
- Chronic causes often pauciymtomatic at the beginning and can evolve long low noise making it then indispensable the use of other parameters such as CST, hepatic MRI and the search for mutations of hereditary hemochromatosis or even other examinations Molecular and cytogenetic more advanced.

References


