An update of clinical management of acute intermittent porphyria

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Abstract: Acute intermittent porphyria (AIP) is due to a deficiency of the third enzyme, the hydroxymethylbilane synthase, in heme biosynthesis. It manifests with occasional neuropsychiatric crises associated with overproduction of porphyrin precursors, aminolevulinic acid and porphobilinogen. The clinical criteria of an acute attack include the paroxysmal nature and various combinations of symptoms, such as abdominal pain, autonomic dysfunction, hyponatremia, muscle weakness, or mental symptoms, in the absence of other obvious causes. Intensive abdominal pain without peritoneal signs, acute peripheral neuropathy, and encephalopathy usually with seizures or psychosis are the key symptoms indicating possible acute porphyria. More than fivefold elevation of urinary porphobilinogen excretion together with typical symptoms of an acute attack is sufficient to start a treatment. Currently, the prognosis of the patients with AIP is good, but physicians should be aware of a potentially fatal outcome of the disease. Mutation screening and identification of type of acute porphyria can be done at the quiescent phase of the disease. The management of patients with AIP include following strategies: A, during an acute attack: 1) treatment with heme preparations, if an acute attack is severe or moderate; 2) symptomatic treatment of autonomic dysfunctions, polyneuropathy and encephalopathy; 3) exclusion of precipitating factors; and 4) adequate nutrition and fluid therapy. B, during remission: 1) exclusion of precipitating factors (education of patients and family doctors), 2) information about on-line drug lists, and 3) mutation screening for family members and education about precipitating factors in mutation-positive family members. C, management of patients with recurrent attacks: 1) evaluation of the lifestyle, 2) evaluation of hormonal therapy in women, 3) prophylactic heme therapy, and 4) liver transplantation in patients with severe recurrent attacks. D, follow-up of the AIP patients for long-term complications: chronic hypertension, chronic kidney insufficiency, chronic pain syndrome, and hepatocellular carcinoma.

Keywords: porphyria, acute intermittent porphyria, neuropathy, treatment, heme

Introduction

Acute intermittent porphyria (AIP) is a rare inherited metabolic disease due to a deficiency of the hydroxymethylbilane synthase (HMBS) in heme biosynthesis.1 AIP manifests after the puberty with occasional neuropsychiatric crises associated with accumulation of porphyrin precursors such as δ-aminolevulinic acid (ALA) and porphobilinogen (PBG) which are released from the liver into the circulation1,2 (Figure 1). The diagnosis is often delayed, because clinical manifestations are unspecific2 and commonly mimic acute encephalopathy or abdominal crisis of other origin.3

AIP is the most common type of acute porphyrias in most of the countries worldwide.1 The major clinical manifestation of AIP is an acute attack, which is clinically indistinguishable from those caused by other acute porphyrias: variegate...
porphyria (VP) and hereditary coproporphyria (HCP). The management of an acute attack in each disease is similar, and thus, more specific classification of an acute porphyria can be done in remission. Photosensitivity and skin fragility found in patients with cutaneous porphyrrias, VP and HCP, occur independently of acute attacks and do not occur in AIP.

When acute porphyria is suspected, biochemical plasma or urinalyses of porphyrin precursors are mandatory to confirm the diagnosis of an acute porphyrionic attack. More than five-fold elevation of urinary PBG excretion is sufficient to start a treatment, but in each case other causes of abdominal crisis must be excluded before a specific treatment of an acute attack is administered. Plasma porphyrin spectrum is a valuable tool to confirm the diagnosis in the early phase and helps to identify different subtypes of acute porphyrrias during an acute attack. In AIP emission peak is only transient, and if taken late, may be negative leading to misdiagnosis. AIP and HCP cannot be distinguished by plasma analysis. Mutation screening can be done at the quiescent phase of the disease (Table 1).

Currently, the prognosis of patients with AIP is good even in severe attacks, but physicians should be aware of a potentially fatal outcome of the disease. During remission the majority of the patients experience no clinical symptoms. Hypertension, chronic kidney insufficiency, chronic pain syndromes, and hepatocellular carcinoma (HCC) may be long-term complications of AIP.

The management of patients with AIP include following strategies:

A. During an acute attack: 1) treatment with heme preparations, if an acute attack is severe or moderate; 2) symptomatic treatment of autonomic dysfunctions, sensorimotor neuropathy and encephalopathy; 3) exclusion of precipitating factors; and 4) adequate nutrition and fluid therapy.

B. During remission: 1) exclusion of precipitating factors (education of patients and family doctors), 2) information about on-line drug lists, and 3) mutation screening for family members and education about precipitating factors in mutation-positive family members which can diminish mortality and prevent subsequent attacks among them.
During an acute attack no need to test in acute porphyria. Always elevated but less than U-PBG in AiP: αδ-aminolevulinic acid in urine; U-PBG, porphobilinogen in urine; U-ALA, δ-aminolevulinic acid in urine; B-, whole-blood metabolites; P-, metabolites in plasma; U-, metabolites in urine; F-, metabolites in feces; HCP, hereditary coproporphyria; VP, variegate porphyria; HMBS, hydroxymethylbilane synthase; Erc-HMBS, HMBS activity in erythrocytes; PPOX, protoporphyrinogen oxidase; Ly-PPOX, PPOX activity in lymphocytes; CoproOX, coproporphyrinogen oxidase; Erc-CoproOX, CoproOX activity in erythrocytes.

Table 1 The laboratory investigations to confirm AIP and other acute porphyrias

| I. Screening tests during an acute attack | U-PBG (always elevated, at least > fivefold, usually >50-fold compared to reference values) P-porphyrin emission spectrum (peak at 615–620 nm in AIP and HCP, 624–627 nm in VP) |
| II. Additional tests (elevated in an acute attack) | During an acute attack | Always elevated but less than U-PBG |
| U-ALA | in AIP: | U-uroerythroblastic porphyrin (I and III) > U-coproporphyrin (I and III) |
| U-porphyrins | in VP: | U-coproporphyrin (I and III) > U-uroerythroblastic porphyrin (I and III) |
| F-porphyrins | in AiP: | F-protoporphyrin (increased in 20% of the patients with AIP) |
| | in VP: | F-protoporphyrin > F-coproporphyrin |
| B-porphyrin | in HCP: | F-coproporphyrin (III:I>2) |
| B-porphyrin | no need to test in acute porphyria |
| III. Specific tests in remission | Enzymes: | Erc-HMBS (<50% in ~84%–95% of AIP patients, can be normal) |
| | in AIP: | Ly-PPOX |
| | in VP: | Erc-CoproOX |
| DNA analysis for specific mutations: | in AIP: | HMBS gene |
| | in VP: | PPOX gene |
| | in HCP: | CoproOX gene |

Abbreviations: AIP, acute intermittent porphyria; VP, variegate porphyria; U-PBG, porphobilinogen in urine; U-ALA, δ-aminolevulinic acid in urine; B-, whole-blood metabolites; P-, metabolites in plasma; U-, metabolites in urine; F-, metabolites in feces; HCP, hereditary coproporphyria; VP, variegate porphyria; HMBS, hydroxymethylbilane synthase; Erc-HMBS, HMBS activity in erythrocytes; PPOX, protoporphyrinogen oxidase; Ly-PPOX, PPOX activity in lymphocytes; CoproOX, coproporphyrinogen oxidase; Erc-CoproOX, CoproOX activity in erythrocytes.

C. Management of patients with recurrent attacks: 1) evaluation of the lifestyle, 2) evaluation of hormonal therapy in women, 3) prophylactic heme therapy, and 4) liver transplantation in AIP patients with severe recurrent attacks.

D. Follow-up of the AIP patients for long-term complications: chronic hypertension, chronic kidney insufficiency, chronic pain syndrome, and HCC.

Clinical manifestations and pathogenesis of an acute attack

The majority of acute attacks manifest as a combination of abdominal pain, mild mental symptoms, and autonomic dysfunction. Both acute peripheral neuropathy and severe encephalopathy may develop if an acute attack proceeds. It is usually iatrogenic, mainly due to administration of porphyrinogenic drugs when the diagnosis of acute porphyria is delayed. Both endogenous and exogenous factors, such as certain medications, alcohol, infections, low caloric intake, or changes in sex hormone balance during the menstrual cycle or pregnancy, can provoke clinical manifestations in AIP. All these factors induce heme synthesis either directly or indirectly via activation of ALA synthase in the liver resulting in accumulation of porphyrins and their precursors in the tissues and circulation.

Excess of ALA is the most potential candidate to cause neuronal damage and could be responsible for autonomic and peripheral neuropathy and encephalopathy via multiple mechanisms. Results from both the experimental and clinical data support the direct neurotoxicity of ALA, but also modification of γ-aminobutyric acid (GABA)-ergic system due to the structural similarity of ALA and GABA/glutamate, as well as formation of free radicals and reactive oxygen species from ALA may play a role in the pathogenesis of an acute attack.

Heme preparations, in the current treatment, lead to a rapid decrease of synthesis of porphyrin precursors via negative feedback (Figure 1). Reduced transcription of ALA synthase in the liver achieved by heme results in cessation of an acute attack within few days. Dose-dependent administration of glucose has also been shown to downregulate ALA synthase in experimental conditions via peroxisome-proliferator-activated receptor γ coactivator 1α (PGC-1α), a protein which directly induces transcription of ALA synthase.
Subsequently, glucose infusions have been used to prevent fasting and may be sufficient in mild attacks. Despite the fact that recombinant human-HMBS-enzyme (rh-HMBS) therapy decreased the plasma level of PBG rapidly, it had no effect on the ALA level \(^{23}\) or the patients’ acute symptoms. In contrast, liver transplantation immediately corrects porphyrin metabolism to normal demonstrating the dominant role of liver as a source of ALA. \(^{24}\)

The exact mechanism of cyclic attacks in women is unknown. Despite the level of sex hormones is at the highest during the second and third trimester, acute attacks are rare during pregnancy. \(^{3,8,17,25}\) Although, especially, progesterone is known to be a potent inducer of ALA synthase, \(^{26}\) the direct role of sex hormones as sole precipitating factors is unlikely. \(^{27}\) Moreover, cyclical attacks occur mainly in premenstruum when the levels of estrogen and progesterone fluctuate the most, and usually are resolved during early menstruation. \(^{8,17}\)

Individual variation in the progesterone metabolism may play a role in clinical manifestations of AIP. \(^{28}\) Cytochrome-P450 activities in the liver also vary individually and can result in an abnormal level or ratio of sex hormones affecting the feedback mechanism to hypothalamus. Several neurotransmitters control the menstrual cycle through the regulation of pulsatile release of gonadotropin-releasing hormone (GnRH) and other clock mechanisms in the hypothalamus. \(^{29}\) This interaction may activate abnormal liver metabolism, \(^{30}\) and consequently precipitate acute attacks by the central mechanism making AIP a central nervous system disorder in addition to a liver disease. \(^{27}\)

### Diagnosis of AIP

The clinical criteria of an acute attack include the paroxysmal nature of the symptoms with abdominal or back pain associated with one or more signs of autonomic dysfunction, hyponatremia, muscle weakness, or mental symptoms \(^{6}\) (Table 2).

The biochemical criteria of an acute attack include more than a fivefold increase of urinary PBG excretion (Table 1).

**Figure 2** Staging of an acute attack in connection with precipitating factors and recommendations of heme therapy.

**Note:** Treatment with heme preparations: recommended (thin solid arrow), usually unnecessary (dotted arrow), influence of precipitating factors and connection with staging of acute attack (thick gray arrow).

**Abbreviations:** CNS, central nervous system; d, days.

**Table 2** The key symptoms indicating acute porphyria

| Intensive acute or recurrent abdominal pain without peritoneal signs or other obvious causes, especially if associated with |
| Dysautonomia (tachycardia, hypertension, nausea, vomiting, constipation) |
| Dark urine (due to porphyrins) |
| Any neurological or psychiatric manifestation |
| Hyponatremia |
| Hypertensive crisis |
| Associated with abdominal pain |
| Acute or recurrent motor/sensorimotor polyneuropathies, especially if associated with |
| Abdominal pain or dysautonomia, commonly preceding neuropathy |
| Signs of central nervous system involvement |
| Hepatopathy or hyponatremia |
| Mainly axonopathy on nerve conduction studies |
| Normal cerebrospinal fluid |
| Acute encephalopathy with no obvious etiology, especially if associated with |
| Dysautonomia or pain |
| Posterior reversible encephalopathy syndrome (PRES) on brain magnetic resonance imaging (MRI) or normal MRI |
| Normal cerebrospinal fluid |
| Epileptic seizures or acute psychosis as a sign of acute encephalopathy with no obvious etiology, especially if associated with |
| Hyponatremia |
| Drug resistant |
| Premenstrual |
| Rhabdomyolysis with no obvious etiology, especially if associated with |
| Abdominal pain |
| Hyponatremia |
which can be detected by a simple Watson-Schwartz or Hoesch qualitative test. The results should be confirmed by a quantitative measurement of urinary PBG, since false positive results in these screening tests are possible, especially, if perchloric acid instead of amyl alcohol is used as an extract. If the urine samples are not sheltered from the light, urinalysis may become false negative.

Urinary excretion of PBG is elevated in 88% of the patients with AIP in remission. During an acute attack PBG excretion increases commonly at least two- to fourfold from the values found in remission. Urinary ALA is always increased during an acute attack but remains elevated only in 61% of AIP cases in remission. In AIP, urinary excretion of uroporphyrins is increased, including both I and III isomers, and exceeds that of coproporphyrins I and III.

Of note, abnormal metabolism of porphyrin and their precursors may also be detected in patients with hepatopathy or heavy metal intoxications. The clinical manifestations may even resemble AIP, but biochemically only a mild to moderate coproporphyrinuria is present, and porphyrin precursors are commonly only transiently elevated. If urinary ALA level exceeds that of PBG significantly, lead intoxication should be excluded.

In AIP, plasma porphyrin emission spectrum test with excitation wave length of 405 nm shows a peak at 615–620 nm during an acute attack similar to HCP but it can be less frequently found in remission. Emission peak is due to porphyrins’ ability to absorb light at wave length around 400 nm and their emission as red fluorescence, at around 600 nm. Plasma emission spectrum test is used mainly to exclude symptomatic VP, since it has a unique 624–627 nm spectrum due to protein-associated plasma porphyrins. Of note, emission peak is commonly negative at the asymptomatic phase of VP.

Around 20% of the patients with AIP have moderately increased excretion of fecal protoporphyrin, which is less prominent than that of VP patients. Since fecal coproporphyrin level is usually normal in AIP, more than tenfold excretion of coproporphyrin (isoform III:1 >2) in feces together with protoporphyrin suggests HCP.

In 84%–95% of patients with AIP, erythrocyte HMBS (Erc-HMBS) activity has been lower than normal. In the variant form of AIP (5%–16% of all patients), Erc-HMBS activity was normal due to an alternative splicing of the HMBS gene in erythroid cells. HMBS activity should be assayed in remission, since erythropoiesis may be enhanced during an attack as well as in hypochromic or hemolytic anemias and hepatopathy. In contrast, it can be decreased in non-porphyric individuals with sideropenia.

DNA analysis is the most reliable method to confirm AIP in the patients and their symptom-free relatives. The direct sequencing of the HMBS gene is used to identify a mutation in the proband and the asymptomatic gene carriers among the family members. The sensitivity of the mutation analysis is 90%–100%. To date, 391 mutations have been reported in the HMBS gene, and therefore DNA testing in an index case of a family is perhaps more laborious and time consuming, but afterward mutation analysis may easily reveal several family members at risk.

Treatment of an acute attack
Current treatment options include heme preparations during an acute attack, which may be life-saving, especially if encephalopathy or polyneuropathy develop. The treatment should be started immediately during a severe or moderate acute attack after the demonstration of typical symptoms of acute porphyria and more than fivefold elevation of urine PBG shown by qualitative tests. Other causes of abdominal crises and neuropsychiatric symptoms often demanding other specific and rapid interventions should always be excluded.

Only around 30%–50% of the patients with a mutation in the HMBS gene have mild or moderate clinical symptoms of AIP during their life span. Less commonly, around 3%–5% of the patients with AIP have recurrent severe attacks, and no time for neuronal recovery. These patients are at a high risk for chronic pain syndrome. The onset and clinical outcome of an attack is commonly influenced by several exogenous factors simultaneously, and endogenous factors, such as the residual activity of a mutated protein, individual differences in other metabolic pathways in the liver and in neuronal protection capacity, may modify the clinical outcome.

Heme
Heme preparations have been used for acute attacks for more than three decades without tolerance. Hemin is isolated and purified from human red cell concentrates. In Europe, Asia, and South Africa, hemin is commercially available as heme arginate (Normosang®, Orphan Europe SARL, Puteaux, France) and in Northern America as lypo-philitized hematin (Panhematin®, Ovation Pharmaceuticals Inc., Deerfield, IL, USA).

In an open series of 22 patients, the patients treated with heme arginate recovered more rapidly in comparison with
those treated with glucose infusions in the earlier series. Safety and efficiency of lyophilized hematin has also been demonstrated in six open-labeled studies involving over 200 AIP patients. The only study using a placebo-controlled series found insignificant benefit of heme arginate for the analgesic requirement, pain score, and duration of the hospital stay. The validity of the trial has been questioned due to small number of patients (eleven patients treated with heme and ten with a placebo), a high proportion of the patients with peripheral neuropathy in this series (43%), delayed administration of preparation (>2 days after admission), and difficulties in arranging a placebo, which resembles heme arginate.

Heme arginate and lyophilized hematin are usually infused daily (3–4 mg/kg) into a large peripheral vein or venous access port for 3–4 consequent days, but a repetitive course may be required if porphyric symptoms are still progressing. (Table 3). Concentrated solution of heme arginate is mixed with 100 mL physiological saline, and lyophilized hematin is reconstituted with sterile water before infusion. Lyophilized hematin should be used immediately after reconstitution. Heme arginate should also be used soon after dilution, since it becomes unstable and may aggregate. In both cases, addition of human serum albumin may be beneficial in order to diminish the risk of phlebitis at the site of infusion. After infusion, the vein should be washed with saline for 10–15 minutes. For the long-term use of heme, a central access catheter (tunneled catheter or portacat) may be useful. Low molecular weight heparin can be used subcutaneously to prevent or treat thrombophlebitis.

The treatment with heme preparations should be started without delay, but even in the late stage of progressing motor neuropathy, it is efficient. If a patient’s neurological condition has stabilized (plateau phase, Figure 2), an additional treatment with heme is rarely necessary but other causes such as infections may deteriorate the patient’s clinical condition and should be treated properly.

The reported side effects include mild coagulopathy, thrombophlebitis, and anaphylactic shock in one case. 

### High carbohydrate loading and supportive treatment

The treatment of acute attacks with high carbohydrate diet or infusions (300–500 g/day) has been in use in order to downregulate the activity of ALA synthase and prevent fasting. High dose of glucose should be infused continuously 24 hours per day with an automatic syringe, and blood sugar level should be monitored regularly to avoid hyper- or hypoglycemia causing additional neurological complications.

If a patient is able to eat, carbohydrate rich meals may similarly have a beneficial effect.

Currently, the use of glucose is limited only to mild attacks (i.e., mild pain, no paresis, seizures, or hyponatremia) according to the guidelines for the treatment of an acute attack in the USA and South Africa, or if heme arginate or hematin are not available locally. (Table 3). In mild cases, the dose of glucose can be lower than discussed earlier. If glucose infusions do not result in clinical remission within a day or two,
or if an acute attack is severe at the onset, heme preparations should be used. Mild or clearly resolving attacks with minor pain or anxiety may be treated symptomatically. 

Low to moderate dose of glucose and saline infusions should be used as supportive treatment to prevent dehydration and during fasting if the patient is unable to drink. The amount of daily fluids may vary from restricted fluid intake in a case of inappropriate antidiuretic hormone secretion to rapid restoration of intravascular volume and correction of electrolyte disturbances in rhabdomyolysis-induced renal failure. Thus, fluid restoration must be tailored individually and careful monitoring of water and electrolyte balance, including sodium, potassium, and magnesium, and renal function should be done (Table 4). Mild to severe hyponatremia is a rather common phenomenon (25%–60%) during an acute attack, and should be corrected slowly (<12 mmol/L/24 h) because of potential pontine myelinolysis, a condition also described in a patient with AIP.

Rhabdomyolysis during an acute attack is often neglected, but can be easily diagnosed by measurement of plasma myoglobin or creatinine kinase level and should be treated according to the guidelines. Acute kidney insufficiency is a rare but a severe complication of an acute attack, and may proceed to hemodialysis.

**Elimination of precipitating factors**

All potentially precipitating factors such as drugs, smoking, and alcohol should be eliminated during an acute attack. Infections should be treated promptly and caloric intake of a patient should be sufficient to avoid fasting. Administration of porphyrinogenic drugs commonly results in proceeding of an acute attack to neuropathy or encephalopathy (Figure 2). Several lists of potentially safe and unsafe drugs are available on the Internet.

Some drugs, such as barbiturates and sulfonamides, are strictly forbidden since their use has been associated with several severe attacks. Most of the drugs, however, are classified as potentially porphyrinogenic, since clinical data about their safety in acute porphyria is lacking. The majority of the patients, especially asymptomatic, tolerate many drugs well; thus, total avoidance of drugs for the safety reasons leads to inappropriate treatment of patients’ other diseases. Antibiotics excluding sulfonamides, drugs for cardiovascular diseases, and pain killers are usually well tolerated in addition to preparations used in oncology. The follow-up of urinary excretion of porphyrin precursors may elucidate the effect of a drug on the heme biosynthesis but increased excretion of porphyrin metabolites without clinical symptoms should not solely determine its use. In each case the potential risk and advantage of a drug should be evaluated by a clinician.

**Symptomatic therapy**

Symptomatic therapy for pain, hypertension, tachycardia, nausea, and vomiting is commonly required (Table 3). Abdominal pain is usually intensive (visual analog scale, VAS > 7 cm, scale from 0 to 10 cm) and opiates are needed. Morphine, pethidine, oxycodone, tramadol, and fentanyl have been used without complication. Paracetamol and anti-inflammatory drugs can be used in mild cases.

Beta blockers are commonly used for tachycardia and to prevent arrhythmia. If a hypertensive crisis develops, it can be treated with beta blockers or clonidine or other drugs recommended by the current guidelines. Nausea and vomiting can be controlled by olanzapine, lorazepam, or prochlorperazine. Domperidone has been used during an acute attack, but interactions with opiates and other drugs increasing the risk for arrhythmias may prevent its use. Metoclopramide has been associated with neuropathy and encephalopathy during a few acute attacks, but some patients have used it without complications. Urinary retention is quite common and can be treated with catheterization. Blood pressure, heart rate, pain score by visual analog scale, and severity of the muscle weakness should be evaluated daily at the bed side. Bulbar paresis, proceeding muscle weakness, and arrhythmia are signs of progression of an acute attack and the patient should be transferred to the intensive care unit. Respiratory insufficiency as a sign of motor neuropathy increases risk of
pneumonia and may require early mechanical ventilation. Patients with paresis due to neuropathy or encephalopathy require rehabilitation therapy even in the early phase.

Epileptic seizures, which are usually generalized, can be treated with intravenous diazepam, gabapentin, levetiracetam, or propofol if status epilepticus develops.2,49,59 Usually there is a single or few transient seizures, which associate with acute encephalopathy visualized as posterior reversible syndrome in brain magnetic resonance imaging (MRI) and does not require anticonvulsive treatment in the follow-up.16 Correction of hyponatremia may be beneficial in patients with seizures (Tables 3 and 4). Insomnia and anxiety are usually mild and do not require additional medications but can be treated with benzodiazepines, such as lorazepam.49 Hallucinations are also signs of acute encephalopathy and should be treated with phenothiazine or olanzapine.

**Prognosis of an acute attack**

Duration of the diagnostic delay of an acute attack correlates with a fatal outcome10 mainly due to the administration of drugs known to precipitate AIP for a misdiagnosed attack. The causes of death are related to complications of prolonged ventilation and cardiac arrest.10,11,50,51 The majority of patients display full functional recovery even after a severe attack.10 The mortality has decreased dramatically during the last decades among diagnosed AIP patients by 5%–20% during an acute attack,8,17 but it is still a potentially fatal disease.

**Prevention of acute attacks**

**Avoidance of precipitating factors**

In remission, patients may tolerate medications classified as potentially unsafe and alcohol, but if porphyric symptoms appear their use should be restricted.8,17,49 The education of patients and their family doctors about precipitating factors, including the information of safe and unsafe drugs, necessity of prompt treatment of infections, healthy lifestyle with regular normocaloric diet, avoidance of alcohol, if porphyric symptoms occur, and smoking, reduction of excessive stress are essential for patients’ prognosis.8,17,49 In the treatment of obesity, the patients with AIP should include carbohydrates into their diet, and the weight reduction should be done slowly.

DNA diagnostics among family members is recommended before the adulthood, since it decreases the likelihood of an acute attack to 5% in patients diagnosed at the presymptomatic phase.12 The prenatal screening in families with AIP is not recommended because of a low risk of severe attacks among mutation carriers12 and good therapy options even if acute attacks would develop.49

**Prevention of cyclical recurrent attacks**

In women with AIP, the menstrual cycle is the most common precipitating factor (10%–39%) manifesting usually with 1–3 months interval in premenstruum.8,17 In the majority cases, these attacks are mild and do not require hospitalization. Irregular cycles due to hormonal imbalance may predispose to cyclical attacks. However, in 3%–5% of the women cyclical attacks are severe and frequent (up to 2–4 weeks interval),17,47 which disable their lives. If the patient has recurrent attacks preventive therapy is needed.

**Evaluation of the lifestyle**

Usually these women have several precipitating factors simultaneously. It is important to first correct lifestyle, indicating cessation of smoking and use of alcohol or any medication or homeopathic drug which may be porphyrinogenic. Since body mass index (BMI) is commonly at the low or low normal level among women with recurrent attacks, gaining weight, at least a few kilos, may be beneficial to balance energy metabolism. Patients should avoid fasting and potential hypoglycemia by regular eating habits.

**Evaluation of hormonal therapy in women**

The exogenous hormonal therapy can be used, if lifestyle changes are insufficient to prevent attacks. Contraceptive pills have been used for prophylaxis of recurrent acute attacks for many years,8,71 but not all women respond to this therapy even after 3 months of use. We have used ethinyl estradiol-levonorgestrel preparation successfully, and the responders have tolerated them well, even for years. About 46%–58% of the Finnish and Swedish women with AIP and VP have used hormonal pills for contraception,8,17 and the majority of them have had no complications in remission despite use of various progesterone compounds in combination with estrogen preparations. Intrauterine devices including levonorgestrel have not been reported to cause porphyrinic symptoms.17 Based on our results, we have let our patients use hormonal contraception under supervision.

Of note, contraceptive pills can also provoke acute attacks in 5%–14% of the women with both latent and manifest AIP (24% of selected women with previously manifest AIP),8,17 and as such many guidelines recommend not to use...
Tolerance to progesterone preparations may vary individually but estrogen preparations, especially used for menopausal symptoms, do not usually precipitate porphyric symptoms. Thus, it is important that the risk for potential complications and benefits of hormonal therapy are evaluated individually by a clinician. Excretion of porphyrin precursors should be controlled and hormonal preparations should be stopped, if symptoms suggestive for acute porphyria occur during their use.

In contrast, GnRH analogs inducing ovarian suppression are not porphyrinogenic and have been reported to diminish the severity and frequency of attacks in 60%–80% of the women. The responders have more frequently had regular menstrual cycles and pronounce decrease in the estradiol level after the treatment with GnRH analogs when compared to the women who have been nonresponders or have responded only to a high dose of the preparation. Drug-induced menopause decreased the excretion of porphyrin precursors to 60% of the previous values in both groups.

The treatment should first be continued up to a few months, and thereafter its efficiency should be evaluated. If a longer period for treatment is needed, GnRH agonist can be combined with estrogen preparations to avoid osteopenia and other menopausal symptoms. Estrogen patches are preferred to avoid the first passage metabolism in the liver. Progesterone should be administered regularly to avoid increased risk of endometrial cancer during the postmenopausal estrogen therapy, but they may also induce acute attacks during the combination treatment. Intrauterine device with levonorgestrel has been used without complications in this compound hormonal therapy, and another option is to lower the dose of GnRH analogs to sustain natural sex-hormone level, which could prevent the long-term side effects of the GnRH agonist. The patients should be followed carefully and efficiently, and side effects and need for the combination therapy should be evaluated at regular intervals. Women with cyclical attacks may also become asymptomatic after their natural menopause.

Screening of urinary excretions of PBG and ALA may elucidate the biochemical activity of the disease, and maybe useful to follow.

Prophylactic heme therapy

Regular heme infusions commonly alleviate the severity and frequency of recurrent attacks. The aim of this treatment is to decrease substantially the level of porphyrin precursors in plasma. The majority of patients respond well but the long-term treatment may induce dependence on the exogenic heme. As a result, a patient may have increased need for heme from monthly to twice a week infusions and withdrawal of the treatment is difficult due to severe porphyric symptoms. Frequent administrations of heme preparations may lead to thrombotic complications of superficial veins, and assessment of a permanent central venous catheter may be necessary. Moreover, long-term treatment of heme may lead to iron overload and organ damage due to hemosiderosis. Progressing hepatopathy, heart failure, and endocrinopathies may develop. Follow-up of plasma ferritin and transferrin saturation levels is necessary, and computed tomography (CT) or MRI reveal iron load in organs. Venesections are usually poorly tolerated but iron chelates may be used.

Liver transplantation in AIP patients with severe recurrent attacks

If the standard treatment is unsuccessful or quality of life is unbearable, liver transplantation is an option for severely affected patients. Currently, more than ten AIP patients have undergone liver (or liver-kidney) transplantation since 2004. It has been successful in the majority of cases resulting in immediate correction of abnormal porphyrin metabolism and cessation of attacks and chronic pain syndrome. Immunosuppressive drugs have been tolerated well by the patients, and their quality of life has improved dramatically. A few patients have died soon after liver transplantation mainly due to infections. Timing is also important in the liver transplantation. Patients should not wait too long in a too poor condition since the recovery from the operation and immunosuppression may associate with early complications such as infections or thromboembolic complications. If chronic motor neuropathy and encephalopathy have been present for years, the full recovery of neuronal damage may not be reached even after total normalization of porphyrin metabolism. Of note, partial liver transplantation has not corrected biochemical abnormalities or clinical manifestations permanently and should not be done. Moreover, transplantation of the HMBS deficient liver induced an abnormal porphyrin metabolism in a recipient.

Management of pregnancies

Previously, pregnancies were commonly complicated by acute attacks mainly during the first trimester and postpartum, and women with AIP were advised to avoid pregnancy. Currently, overall prognosis for pregnancy in AIP is good, especially if the diagnosis is known in advance and no porphyrinogenic drugs are used.
Cyclical attacks do not predict attacks during pregnancy, and pregnancy nowadays seldom precipitates acute attacks. Heme preparations have been used during pregnancy without fetal or maternal complications. In some women, who have experienced cyclical attacks for many years, recurrent attacks have stopped after the first trimester of pregnancy, and these women have been asymptomatic thereafter. Thus, pregnancy may act as a hormonal therapy and could be even considered as an option for the women with cyclical attacks.

Long-term complications of AIP

Hypertension and renal insufficiency

The prevalence of hypertension is significantly higher in patients with manifest AIP than in general population. Thus, early monitoring of blood pressure and efficient treatment with antihypertensive drugs is mandatory to avoid organ complications. This is especially crucial if renal failure is already present.

Monitoring of creatinine level in the follow-up of patients with manifest AIP shows mild to moderate elevation in 10%–50% of the cases. This is significantly more common than in general population. Tubulointerstitial kidney disease has been the main presentation in a biopsy of the investigated cases. It has been commonly associated with recurrent acute attacks, hypertension, and use of anti-inflammatory drugs. The main recommendation for renal protection include adequate drinking regimen, blood pressure control, and avoidance of anti-inflammatory or other nephrotoxic drugs according to the general guidelines. Patients with AIP and nephropathy, even mild, have an increased risk of deterioration of renal functions during an attack, and should be treated carefully with fluid therapy and hemodialysis if necessary.

Chronic hemodialysis may induce cutaneous symptoms in patients with AIP mimicking porphyria cutanea tarda (PCT). Since mainly ALA and PBG but not porphyrins have been filtered during dialysis, recurrent attacks were transformed to cutaneous manifestations in one patient. Use of high-flux hemodialysis and erythropoietin substitution might be of help since they remove plasma porphyrins and have alleviated clinical manifestation in PCT patients with end-stage renal failure. Kidney transplantation or combined liver and kidney transplantation have been done successfully and should be done whenever needed. The patients who have undergone kidney or liver transplantation should be followed up regularly because of increased risks of renal failure, hypertension, and cardiovascular complications. Metabolic changes and osteoporosis may also develop. Skin cancers and lymphomas are more common than in the general population and thus, extensive exposure to the sunlight and smoking should be avoided. Long-term complications are often multifactorial but commonly related to immunosuppressive drugs in use.

Hepatocellular carcinoma

Several population-based studies of AIP patients have shown an evidence of a significantly increased incidence of HCC in AIP compared to general population. The highest incidence of HCC (23%–27%, more than 60-fold increased risk compared to general population) was shown in the Finnish and Swedish patients with AIP. In contrast to renal insufficiency, HCC has developed in patients with both symptomatic and asymptomatic AIP, although the patients with manifested AIP are likely at higher risk. Only in one-third of the cases, HCC has coexisted with liver cirrhosis or well-known risk factors for HCC, such as viral hepatitis or heavy alcohol consumption. Hepatoma should always be considered if a patient develops acute attacks or an abdominal pain after long-term remission or abnormally high porphyrin excretion pattern even at the old age. Hepatomas can be solitary or multiple in AIP patients, and should be treated according to current guidelines (ie, hepatic resection, local ablation therapies, multikinase inhibitors) whenever possible.

Since 10% of the patients with AIP die of hepatoma, the patients with AIP most likely benefit from regular screening for HCC after 50 years of age. Early diagnosis of HCC is crucial for the prognosis and can be achieved only by accurate radiological imaging.

Chronic pain

Among AIP patients with recurrent acute attacks, chronic neuropathic, myalgic, or abdominal pain usually accompanied by fatigue is common (18%–22%). This could be due to repetitive autonomic and peripheral nerve damage during acute attacks, and the long-term hypocaloric nutrition inducing catabolic stage and muscle atrophy. The mouse model of AIP, which is in contrast to human autosomal-dominant AIP caused by compound heterozygous mutations in HMBS gene (HMBS) and severe deficiency of HMBS enzyme, has developed chronic axonal motor neuropathy even in the absence of acute attacks and elevated ALA levels. In humans, a homozygous AIP patient has developed a chronic sensorimotor neuropathy associated with severe neurodegenerative encephalopathy and mental retardation at his early childhood but no chronic pain syndrome has been present.
The treatment options include prevention of acute attacks as described earlier and prolonged treatment with gabapentin or antidepressants, such as fluoxetine. Prophylactic heme infusions commonly alleviate the chronic pain syndrome. Regular use of opiates induces tolerance to these drugs and high doses may increase the risk of side effects such as addiction, somnolence, and apnea.

**Future therapies**

rh-HMBS preparation has been shown to lower plasma PBG levels in symptom-free patients with increased excretion of urinary PBG. The similar biochemical effect of rh-HMBS therapy was demonstrated in the HMBS (−/−) mouse model. The rh-HMBS-enzyme replacement therapy, however, had no effect on the patients’ symptoms during acute attacks mainly due to constantly high plasma ALA levels. This may be due to the fact that the rh-HMBS included only the erythroid specific form of the enzyme and was not targeted to the liver, the main site of ALA overproduction. The potential development of rh-HMBS-enzyme replacement therapy should be focused on the delivery of an active enzyme into the liver restoring the normal HMBS activity and heme biosynthesis.

Both adeno-viral-mediated and nonviral HMBS gene transfers corrected the metabolic defect in the cell lines of AIP patients and HMBS (−/−) mice. Only virus-mediated DNA transfer into the liver was successful and corrected metabolic defect in HMBS (−/−) mice at least for a month. Adenoviral cytotoxicity, especially, for hepatocytes, the immunological response to viral antigens, and only transient transduction of the therapeutic gene limit potential applications of this method.

Adeno-associated viral (AAV) vector encoding HMBS and driven by liver-specific regulatory elements, such as α1-microglobulin enhancer and α1-antitrypsin promoter, is a promising alternative to the first-generation adenoviral vectors, since it demonstrated long-term transgene expression of HMBS in the liver with reduced risk of side effects. AAV vector encoding HMBS (recombinant AAV serotype 5-codon-optimized human HMBS, rAAV5-cohHMBS) has been used in macaques. The safety and efficiency of AAV vector containing the HMBS gene is currently investigated in the clinical trial (Phase I).

Small interfering RNA targeting liver ALA synthase results in reduction of ALA synthase mRNA and prevents acute attacks in AIP mouse model, and it is under development for human studies. This demonstrates that downregulation of liver heme biosynthesis via direct ALA synthase inhibitor could result in a promising future therapy.

**Conclusion**

Before the glucose and hematin treatment of an acute attack has been introduced, the mortality rate during an acute attack of AIP was up to 66%. Currently, the improved diagnostics, the early and effective treatment with heme preparations, and the prevention of acute attacks have decreased the mortality substantially. However, the morbidity and mortality (5%–20%) of an acute attack is still significant if the diagnosis of AIP is delayed. Thus, it is important that AIP patients are treated in the experienced centers with facilities that enable monitoring of their clinical and biochemical profile.

Current options to prevent acute attack include patients and family doctor education and family screening for HMBS mutation carriers. In patients with recurrent attacks, ovarian suppression using hormonal interventions, prophylactic heme therapy, or liver transplantation should be considered to improve the prognosis and quality of their lives. Future therapies, such as enzyme or gene delivery, should demonstrate clinical efficiency and safety before they can be applied to the clinical use. In the long-term follow-up, impaired renal function and hypertension should be treated carefully, and screening for hepatomas should be considered after 50 years of age.

**Disclosure**

The authors have participated in clinical trials organized by Zymenex, Alnylam, Clinuvel, and Boehringer Ingelheim, and as speakers in seminars organized by Orphan Europe. The authors report no other conflicts of interest in this work.

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