SYSTEMIC PHLEBOTROPIC DRUGS IN PHARMACOTHERAPY OF CHRONIC VENOUS INSUFFICIENCY OF THE LOWER EXTREMITIES

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The modern aspects of pharmacotherapy of chronic venous insufficiency (CVI) of the lower extremities are presented in the article. Phlebotropic drugs are medications for basic pharmacotherapy of CVI. A common characteristic of pharmacodynamics of these drugs is the ability to increase the venous tone, and reduce severity of specific symptoms and syndromes of CVI. The efficiency of the most widespread venotonic drugs has been analyzed. The largest number of preclinical and clinical research among all groups of vено-active drugs has been devoted to the study of Micronized Purified Flavonoid Fraction (MPFF). It has been determined that the therapeutic effect of MPFF is higher compared to the therapeutic effect of non-micronised Diosmin. An important feature of the action of MPFF is a rapid onset of the venotonic effect, the effect is shown already from the first hour after administration and lasts during the day. This action is successfully complemented by improved rheological properties of the blood, reduction in blood viscosity, stimulation of the drainage function of lymphatic vessels. Treatment should be given by courses, which duration depends on disease severity, and is for 2-3 months, but not less than twice a year. The continuous therapeutic scheme can be used in the case of refractory forms of CVI. MPFF is characterized by a high clinical efficacy, high bioavailability and safety. It does not cause serious adverse reactions, well tolerated and can be used as a drug of the first choice.

Chronic venous insufficiency (CVI) of the lower extremities is a syndrome characterized by disorders of the venous outflow on the macrohemodynamic level, it leads to disorganization of the regional microcirculation system [6, 16]. CVI is considered to be the most common peripheral vascular disease. The high level of morbidity of the working-age population reaching 40-50% by expert estimations allows to conclude that CVI is an important medical and social problem. Furthermore, if previously the disease was predominantly a problem of persons of the senior age group (over 50 years), at the present time the first signs of venous reflux are found in 10-15% of 12-13 years school children [5, 24].

The main predisposing factors of CVI are:

• hereditary or acquired valvular insufficiency of the veins;
• decrease of the tone of the varicose vein wall;
• high pressure due to the blood reflux from the deep venous system to the surface venous system [7, 21, 22].

CVI of the lower extremities is primarily characterized by stagnation or redistribution of the venous blood flow in the venous system. The mechanism of its development is associated with hypertension in the inferior vena cava forming as a result of the varicose transformation of the venous wall or its thrombus burden [6, 22]. Initial macrohemodynamic disorders lead to the serious changes at the tissue level. The result of this process is development of microcirculatory disorders in the form of trophic disorders.

Until recently, the sufficient attention has not been paid to pharmacotherapy of CVI in Ukraine. The cause for this situation is the absence of effective, available and safe drugs. The situation has radically changed in the last two decades when the improvement of pharmaceutical technologies has led to the appearance a new generation of phlebotropic drugs at the pharmaceutical market [4, 5, 7].

Medications of the basic pharmacotherapy CVI are phlebotropic drugs (PHD) (synonyms are vено-active drugs, phleboprotector drugs, venotonic drugs). They are a heterogeneous group of biologically active substances obtained by processing of plant raw material or by chemical synthesis. The ability to increase the venous tone and reduce the venospecific symptoms and syndromes of CVI is a common feature of the pharmacodynamics of these drugs [1, 18].

It should be noted that more than 20 PHD have been registered and are used in Ukraine. The main phlebotropic drugs and their daily doses are shown in Table 1.

Flavonoids are the most studied among the basic venotonic drugs. Drugs based on Diosmin, which are especially widely used in the international phlebological practice, are prescribed more often than others [1, 2, 13]. Most experts call Micronized Purified Flavonoid Fraction (MPFF), which has the brand name “Detrelex”, the reference phleboprotector drug because of its multitargeted mechanism of action, ease of use, high bioavailability and safety [3, 12, 20].

An important feature of micronized Diosmin is a rapid onset of the venotonic effect, the effect is shown already from the first hour after administration, and lasts during the day [1, 15]. This action is successfully
complemented by improved rheological properties of the blood, reduction in blood viscosity [2, 3], stimulation of the drainage function of lymphatic vessels [11, 23].

Most PHD increase the tone of peripheral veins and lymphatic vessels due to the effect on the norepinephrine-dependent mechanism, and also due to some specific effects of individual drugs [8]. Requirements for PHD are not limited only by increase of the venous wall tone. Modern veno-active drugs should have the ability to stimulate the lymphatic drainage and improve microcirculation [9, 15]. Basic clinical effects of different PHD are presented in Table 2.

The main indication for the use of PHD is the symptoms associated with chronic venous insufficiency such as heavy legs, discomfort, itching, pain along the varicose veins, plethora, night cramps, and other veno-specific complaints, chronic venous edema, and trophic skin disorders, including venous ulcers. The results of numerous clinical studies suggest that at early stages of the disease all PHD have a good therapeutic effect on subjective symptoms of CVI, but not the objective symptoms of CVI such as telangiectasia, varicose reticular and subcutaneous veins [10, 13]. At the same time in choosing the drug therapy at early stages of CVI the preference should be given to PHD with the efficacy and safety proven in randomized controlled clinical studies. Chronic venous edema is an absolute indication for PHD [14]. According to the meta-analysis data MPFF shows the most evident anti-edematous effect, which is the drug of first choice in chronic venous edema [15].

Hydroxyethyl rutosides (Venoruton) and the extract of Ruscus aculeatus (Cyclo 3 fort) also reduce the chronic venous edema and can be the second line drugs. Efficiency of non-micronized (native) Diosmin in regard to the chronic venous edema is not significantly different from placebo. To assess the efficiency of other phlebotropic drugs in the chronic venous edema it is necessary to conduct numerous randomized controlled clinical studies according to standardized protocols [11, 29].

Phlebotropic drugs should be administered in adequate doses, which are recommended by drug manufacturers, as well as the standard therapeutic schemes and duration of treatment should be observed. The excess of the standard daily doses does not necessarily improve the clinical effect, but increases the frequency of unwanted side effects that can break the started course of pharmacotherapy. However, results of certain clinical research indicate that in some clinical situations

Table 1

| Basic phlebotropic drugs |
|---------------------------|
| **Trade name of the drug** | **Chemical group** | **Active component** | **Daily dose, mg** |
| Herbion aesculus | α-Benzopyrones | Coumarin | 100-400 |
| Anavenol, Venoruton | Flavonoids | Hydroxyethyl rutoside | 1000-3500 |
| Venorutinol, Troxevasin | | Troxerutin | 1000-3500 |
| Detralex, Phlebodia | | Diosmin | 600-1000 |
| Endotolon | Pycnogenols | Procyanidolic oligomers (grape seed extract) | 300 |
| Aescusan, L-Lysine Aescinat | | Escin (horse chestnut extract) | 60-120 |
| Gotu Cola | Saponins | Centella asiatica (“clever herb”) | 400–800 |
| Ginkor Fort, Tanakan | Saponins and flavonoids | Ginkgo biloba | 60-120 |
| Cyclo 3 Fort | | Ruscus aculeatus | 300-450 |
| Tonoress | Ergot alkaloids | Dihydroergotamine | 2.5-7.5 |
| Redergin | | Dihydroergocristine | 2.5-7.5 |
| Glyvenol | Synthetic compounds | Tribenoside | 800 |
| Doxi-Chem | | Calcium dobesilate | 1000-1500 |

Table 2

| Therapeutic effects and mechanisms of action | PHD |
|-------------------------------------------|-----|
| Antiedematous and capillary protective | All phlebotropic drugs |
| Analgesic (venous pain) | MPFF, grape seed extract |
| Venotonic | All phlebotropic drugs |
| Lymphotropic | MPFF, Ruscus aculeatus extract |
| Rheological | MPFF, Troxerutin, Rutin |
| Profibrinolytic | MPFF, Troxerutin, Diosmin |
| Anti-inflammatory | MPFF, Diosmin, Ginkgo biloba extract |
| Inhibition of leukocyte-endothelial adhesion | MPFF |
| Protection of the venous valves | MPFF |
| Protection of the venous wall:  
- inhibition of lysosomal enzymes  
- stabilization of collagen  
- suppression of free radicals  
- normalization of PGE2 synthesis  
- improvement of circulation in the vasa vasorum | MPFF, oligomers  
MPFF, oligomers  
Ginkgo biloba extract  
MPFF  
Hydroxyethyl rutoside |
The increase in the dose of the phlebotropic drug can increase its therapeutic efficacy. In this connection, in situations when the benefit of the drug outweighs the potential risk, the dose of the phlebotropic drug can be increased after documentary evidence and obtaining of the informed consent of the patient [17].

Phlebotropic drugs can be prescribed as mono-therapy or fixed combinations (multi-drug) since the simultaneous use of two or more drugs belonging to related chemical classes does not intensify the therapeutic effect, but increases the probability of undesirable side reactions.

PHD are relatively well tolerated by most patients even if taken them for long time. The undesirable dyspeptic disorders (stomach pain, diarrhea, vomiting, etc.) and vegetative disorders (insomnia, dizziness, etc.) are observed in no more than 5% of patients [23]. The typical undesirable side reactions of various PHD are presented in Table 3.

Treatment should be given by courses, which duration depends on disease severity, and lasts 2-3 months, but not less than twice a year. The interval between courses should be used for physical therapy or SPA treatment. The continuous therapeutic scheme can be used in the case of refractory forms of CVI [7, 25]. MPFF has the highest safety profile in the long-term use (6-12 months) [1, 3, 19, 20].

According to standard protocols adapted to each stage and form of CVI the randomized controlled clinical studies should be conducted for all new phlebotropic and phleboprotector drugs in order to assess their efficacy and safety [16, 25]. Formal bioequivalence of generic and original drugs can not guarantee the similar clinical effect and safety [1, 7, 20].

CONCLUSIONS

In conclusion, it should be noted that the rational pharmacological therapy allows to perform effective pathogenetic and symptomatic treatment of various forms and complications of CVI. The promising development of this method is implemented in several directions. Among them the following ones should be distinguished:

- the search of new chemical substances which selectively block the synthesis of adhesion molecules, leukocyte activation factors and tissue metalloproteases;
- the study of drugs that stimulate and control the synthesis of the connective tissue;
- optimization of delivery methods of pharmacological agents in the affected area with the help of nanotechnology.

The successful implementation of these programmes in the future will allow to speak about the possibility of the effective control of CVI.

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| Active component          | Skin reactions | Dyspeptic reactions | Other adverse drug reactions |
|---------------------------|----------------|---------------------|-----------------------------|
| Oxerutin and rutosides   | Often          | Often               |                             |
| Escin (horse chestnut extract) | +              | +                   | Dizziness, headache, rash  |
| Ruscus aculeatus extract | +              | +                   | Dizziness, headache, rash, tachycardia |
| Grape seed extract       | +              | Often               | Dizziness, headache, rash, tachycardia |
| Ginkgo biloba extract    | +              | +                   |                             |
| Troxerutin               | Often          | +                   |                             |
| Diosmin                  | +              | +                   | Dizziness, tachycardia, weakness |
| MPFF                     | +              | +                   | Rare vegetative reactions such as dizziness and weakness |
| Calcium dobesilate       | +              | +                   | Agranulocytosis             |
СИСТЕМНИ ФЛЕБОТРОПНІ ПРЕПАРАТИ У ФАРМАКОТЕРАПІЇ ХРОНИЧНОЇ ВЕНОЗНОЇ НЕДОСТАТОЧНОСТІ НИЖНІХ КІНЦІВКОВ
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Ключові слова: хронічна венозна недостатність; флеботропні препарати; мікронізована очищена фракція флавоноїдів
Наведені сучасні аспекти фармакотерапії хронічної венозної недостатності (ХВН) нижніх кінцівок. Засобами базисної фармакотерапії ХВН служать флеботропні лікарські препарати. Загальна особливість їх фармакодинамики – здатність підвищувати венозний тонус, а також зменшувати вираженість веноспецифічних симптомів і синдромів. Проаналізовано ефективність найбільш поширених венотонічних препаратів. Серед усіх груп веноактивних препаратів найбільшу кількість досліджень стосується вивчення мікронізованої очищеної фракції флавоноїдів (МОФФ). Встановлено, що терапевтичний ефект МОФФ вище в порівнянні з немікронізованим діосміном. Важлива особливість дії МОФФ – швидкий, вже з першої години після прийому початок венотонічного ефекту, який зберігається протягом доби. Ця дія вдається доповнюється поліпшенням реологічних властивостей крові, зниженням її в'язкості, стимуляцією дренажної функції лімфатичних судин. Лікування слід проводити курсами, тривалість яких залежить від тяжкості захворювання і складає 2-3 міс., не рідше двох разів на рік. У разі рефрактерних форм ХВН може бути використана безперервна схема фармакотерапії. МОФФ характеризується високою клінічною ефективністю, має високу біодоступність та безпеку, не викликає серйозних побічних реакцій, добре переноситься хворими і може застосовуватися як препарат першого вибору.

СИСТЕМНІ ФЛЕБОТРОПНІ ПРЕПАРАТИ В ФАРМАКОТЕРАПІЇ ХРОНИЧНОЇ ВЕНОЗНОЇ НЕДОСТАТОЧНОСТІ НИЖНІХ КОНЕЧНОСТЕЙ
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Приведені сучасні аспекти фармакотерапії хронічної венозної недостатності (ХВН) нижніх конечностей. Средствами базисной фармакотерапії ХВН служать флеботропні лекарські препарати. Общая особенность их фармакодинамики – способность повысить венозный тонус, а также уменьшать выраженность веноспецифичных симптомов и синдромов. Проанализирована эффективность наиболее распространенных венотонических препаратов. Среди всех груп веноактивных препаратов наибольшее количество исследований касается изучения мікронізованої очищеної фракції флавоноїдів (МОФФ). Установлено, что терапевтический эффект МОФФ выше по сравнению с немікронізованим діосміном. Важная особенность действия МОФФ – быстрое, уже с первого часа после приема начало венотонизирующего эффекта, сохраняющегося в течение суток. Это действие удачно дополняется улучшением реологических свойств крови, снижением её вязкости, стимуляцией дренажной функции лимфатических сосудов. Лечение следует проводить курсами, продолжительность которых зависит от тяжести заболевания и составляет 2-3 мес., не реже двух раз в год. В случае рефрактерных форм ХВН может быть использована непрерывная схема фармакотерапии. МОФФ характеризуется высокой клинической эффективностью, имеет высокую биодоступность и безопасность, не вызывает серьезных побочных реакций, хорошо переносится больными и может применяться как препарат первого выбора.