

EMERGENCY D-Virus

Plan of Care

CELLULAR THERAPY

DIETARY SUPPLEMENT PRODUCTS

AnterFerron-1, AnterFerron-2, ImunStem, and Aktiffvate

COMPANY

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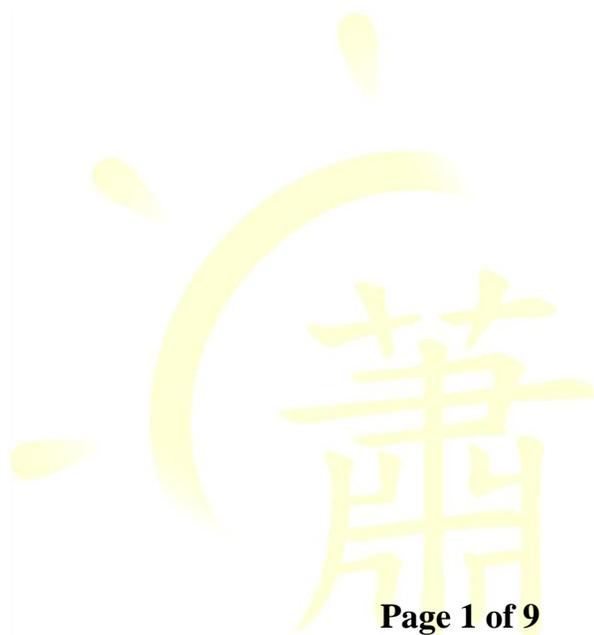
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“TREATMENT”

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1.0 INTRODUCTION

Stephen R. MEIS, M.D., Board Certified (Dr. Meis), I strongly recommend Golden Sunrise Nutraceutical Incorporation herbal products *ImunStem* and *Aktiffvate*, along with their *AnterFeerons* product, as uniquely qualified to treat and modify the course of the **virus** epidemic in CHINA and other countries. Patients with late stages of *Hepatitis C*, and *AIDS / HIV* have responded with greatly improved quality-of-life and extending their lives when treated with *ImunStem* and *Aktiffvate*. For viral colds, *Aktiffvate*, when given in frequent dosing, as frequent as every one half (½) hour to one (1) hour, will not only alleviate the symptoms quickly, but stop the cold virus itself, usually in less than 2 – 3 days. Now *AnterFeerons* has been added to the *ImunStem* and *Aktiffvate* and shown added improvement for a variety of infections associated with cancer patients and the chronically ill, whether it be viral or bacterial.

ImunStem, *Aktiffvate*, and *AnterFeerons* herbs are the basis of the whole cellular therapy developed by Golden Sunrise Nutraceutical. They have proven themselves to the United States Food & Drug Administration (FDA). *ImunStem*, an herbal product, was the first dietary supplement in the United States to be approved as a prescription medicine and also for the indication to treat ***Serious or Life-threatening*** conditions. It qualified for both of these under the Regenerative Medicine Advance Therapy (RMAT) designation in the 2016 Cures Act, enacted by the 114th United States Congress. This designation acknowledges not only the effectiveness of these herbs, usually only associated with pharmaceutical drugs, but also causing no side effects, a quality of dietary supplements.

Golden Sunrise Nutraceutical metabolic therapies will treat ***Serious or Life-threatening*** conditions. These conditions result from an accumulation of toxins in the body from food additives, preservatives, pesticides, prescription drugs, and industrial pollution that disrupt the immune system and cell metabolism. Regenerating the cellular metabolic abnormalities with plant based botanicals found in the ***EMERGENCY D-Virus Plan of Care*** is the basis for the remarkable improvement for human health.

The technology developed by Golden Sunrise Nutraceutical is the key for the effectiveness of the herbs on the immune system and cellular metabolism. They have immune stimulating properties. In-vivo studies on treated patients demonstrate increasing phagocytic activity and synthesis of helper cell function. The cells possess a bipolarity and lipophilicity that facilitates molecular diffusion through permeable and selective membranes, including crossing the blood / brain barrier. Golden Sunrise Nutraceutical herbal supplements are able to penetrate the cells at the cellular level without any disruption or damage to the cells because they are recognized as food. This food provides the cells with the necessary building blocks for the cells to repair and rejuvenate themselves and flush out the accumulated toxins in the cells.

Golden Sunrise Nutraceutical products and treatments improve genetic ***Telomeres*** for cellular regeneration which increases the overall-health of the body and can increase human longevity.

2.0 INSTRUCTIONS FOR THE TREATMENT

General dosing recommendations: *ImunStem* and *Aktiffvate* should be taken throughout the course of the illness. Recommended starting dose for those without any viral symptoms would be twice a day administration. But for with symptoms or with chronic health issues three (3) or four (4) times a day is advised. For someone who is exhibiting symptoms, after three (3) days of the *ImunStem* and

Aktiffvate, the *Aktiffvate* alone should be increased to a quarter dropperful every thirty (30) to sixty (60) minutes in addition to be *ImunStem / Aktiffvate* combination already being taken by the patient. With this increased dosing of *Aktiffvate*, disappearance of viral symptoms is expected within two (2) to four (4) days. If symptoms however do not resolve with this regimen or the person is steadily worsening, the *AnterFeerons* should be used as directed. For the infirmed or elderly, the *AnterFeerons* must be administered under direct observation of health professionals, because it will bring on nausea and or vomiting as part of utilizing the body’s natural shedding mechanism of the illness / virus. It will cause some dehydration which must be compensated either by an electrolyte solution (*Pedialyte, Gatorade, etc.*) or *IV fluids*.

2.1 Administration and Dosage of *IMUNSTEM* and *AKTIFFVATE*

Upon the first visit it is suggested that once the medical evaluation of the patient is completed and the medical staff deems it appropriate, then patients will receive ½ – ¾ of a dropperful of *ImunStem* and *Aktiffvate* 1–4 times a day. The medical staff should monitor the patient for at least ten (10) minutes to help with any effects that might need other medical attention. For example, *ImunStem* can open and improve blood flow throughout the body and the patient might experience a feeling of warmth and begin having nasal mucus discharge. After ten (10) minutes if the patient is stable, the ½–¾ dropperful of *Aktiffvate* should be administered with similar monitoring.

ImunStem and *Aktiffvate* are liquid form:

Product	Dose	Dose Size of a Dropper	Dose Per day
<i>ImunStem</i>	1 ml	½ - ¾	1–4
<i>Aktiffvate</i>	1 ml	½ - ¾	1–4

2.2 Administration and Dosage of *ANTERFEERONS*

Take one (1 fl.oz.) fluid ounce of *AnterFeeron-1*. Then in forty-five (45) minutes to one (1) hour following ingestion of *AnterFeeron-1*, take one (1 fl.oz.) fluid ounce of *AnterFeeron-2*, administered in the same dose. Each bottle must be emptied into a small glass and swallowed quickly all at once (not sipped or sniffed), followed by water to wash it down.

AnterFeeron-1 and *AnterFeeron-2* are liquid form:

Product	Dose
<i>AnterFeeron-1</i>	1 fl.oz.
<i>AnterFeeron-2</i>	1 fl.oz.

2.3 Ongoing Treatment

ImunStem, Aktiffvate, and *AnterFeerons* are available now and once they are started, they will help alleviate the people immediately with the acute illness of the virus. But this treatment, which is tailored for this acute emergency, is the beginning of Golden Sunrise Nutraceutical’s *METABOLIC Plan of Care*. The *METABOLIC Plan of Care* is a carefully planned course of therapy with herbal treatments to treat at a cellular level chronic illnesses such as hypertension, diabetes, peripheral neuropathies, parkinsonism, multiple sclerosis, gastrointestinal conditions, schizophrenia, Lyme’s disease, to name a few. It is a preventative for cancer, which primarily is a metabolic problem like our other diseases. For those who start with the *EMERGENCY D-Virus Plan of Care*, it would be ideal to continue with Golden Sunrise’s *METABOLIC Plan of Care*.

3.0 WARNING AND PRECAUTIONS

ADMINISTRATION OF *IMUNSTEM*, *AKTIFFVATE*, *ANTERFEERON-1*, and *ANTERFEERON-2*, SHOULD ALWAYS BE UNDER THE SUPERVISION OF A PHYSICIAN. RECOMMENDATION FOR GOLDEN SUNRISE NUTRACEUTICAL **EMERGENCY D-VIRUS PLAN OF CARE** IS BASED ON MEDICAL EVALUATION OF THE PATIENT. THESE PRODUCTS CAN LOWER BLOOD SUGAR IN DIABETICS AND LOWER BLOOD PRESSURE IN HYPER-TENSIVE PATIENTS. BLOOD SUGAR AND BLOOD PRESSURE SHOULD BE MONITORED FOR THOSE WITH A HISTORY OF HIGH SENSIVITY TO MEDICATIONS. A LOWER STARTING DOSE MIGHT BE APPROPRIATE.

4.0 DRUG INTERACTIONS

ImunStem, *Aktiffvate*, and *AnterFeerons* have not been reported, so far, to have had any interactions with single or multiple combination of prescription drugs. You should always read product labels. If you have a medical condition, or are taking other prescription drugs, herbs, or dietary supplements, you should speak with a qualified healthcare provider before starting a new therapy.

5.0 THERAPEUTIC RESPONSE

5.1 Adverse Sensitivity Response of *ImunStem* and *Aktiffvate*

- *ImunStem* and *Aktiffvate* can cause severe allergic skin rashes.
- Vomiting.
- In rare circumstances an adverse sensitivity response in the mouth, such as mild blisters, have occurred.
- A burning sensation in the throat in the beginning of oral treatment may occur, but it subsides. If the burning sensation persists, gelatin capsules used for administration, may be substituted as an alternative.

6.0 CLINICAL PHARMACOLOGY

6.1 Mechanism of Action

EMERGENCY D-Virus Plan of Care are herbal (botanical) products which help to regenerate the cell and promote the cellular metabolism to return to homeostasis in the human body.

It is postulated that in arresting abnormal cellular mutation, the precursors for normal and enhanced cellular regeneration and cellular division can be accelerated in relation to the selective existing disease pathology. It is therefore theorized that the combination of the body's thermal energy, particularly during variance mutation and inflammatory processes, potentiates the self-propelling of molecules by possible passive membrane transportation or passive diffusion of the **EMERGENCY D-Virus Plan of Care** intact across all cell membranes, including the blood-brain barrier, which allows its efficacy for the central nervous system.

EMERGENCY D-Virus Plan of Care possess a bipolarity and a lipophilicity that facilitates molecular diffusion through various permeable and selective membranes. In-vivo studies on test subjects have indicated that the cell membrane integrity remains intact and is not disrupted or destroyed in the process of assimilating into the cell, thus ensuring long-term effectiveness.

EMERGENCY D-Virus Plan of Care are the key for the effectiveness of the products on the immune system and cellular metabolism. They have immune-stimulating properties. In-vivo studies on treated patients demonstrate increasing phagocytic activity and synthesis of helper cell function. **EMERGENCY D-Virus Plan of Care** have shown to transform Deoxyribonucleic Acid / Ribonucleic Acid (DNA / RNA) repair, before, during and after chemotherapy drugs, prescription drugs, toxic exposure and chemical induced damage. These herbals have a variety of effects including antioxidant activity and anti-inflammatory properties.

EMERGENCY D-Virus Plan of Care can play an integral role in hypothalamic activity, easing both parasympathetic and sympathetic nervous systems towards a state of homeostasis. For example, blood pressure level is directly linked to the sympathetic nervous system and the normalizing of blood pressure levels exhibited following administration of **EMERGENCY D-Virus Plan of Care** are compatible with the stabilization of the sympathetic nervous system activity. **EMERGENCY D-Virus Plan of Care** reduce inflammation by affecting immune responsiveness through neuroendocrine factors. Improvement of balance problems, a side-effect of some chemotherapies, by Golden Sunrise Nutraceutical products is thought to be from improvement of vestibular function.

6.2 Pharmacokinetics

EMERGENCY D-Virus Plan of Care are readily absorbed into soft tissue when taken as an oral dose. This is accomplished because **EMERGENCY D-Virus Plan of Care** readily passes through membrane tissue by passive diffusion.

The **EMERGENCY D-Virus Plan of Care** dietary supplements are metabolized by the body to more hydrophilic products that are excreted in the urine and feces.

7.0 RESULTS OF PATIENTS AFTER TREATMENT

The group of patient with the virus found that **EMERGENCY D-Virus Plan of Care** improved the immune system and alertness immediately. Physicians have observed that using **EMERGENCY D-Virus Plan of Care** provokes a significant response, i.e., a reduction in symptoms in patients with the virus.

On April 2020, six (6) patients administered **EMERGENCY D-Virus Plan of Care** over the course of time from 1 – 2 weeks, controlled under supervision of a physician “**Clinical Assessment and Progress Note**” included observations by Golden Sunrise Nutraceutical medical staff.

Safety was monitored and throughout these six (6) patients no adverse side-effects were noticed. A total of six (6) patients were diagnosed with the virus that required close monitoring by both the attending physician, specialist of that field and by Golden Sunrise Nutraceutical medical staff.

Six (6) PATIENTS’ Histories and EMERGENCY D-Virus Plan of Care Treatment of the virus.

Four (4) patients who were confirmed with the virus illness, were treated with the **EMERGENCY D-VIRUS Plan of Care**. They were all treated in their homes. They showed improvement by day number

two (#2) or day number three (#3) of the treatment. All were asymptomatic by day number seven (#7) to day number nine (#9) of treatment.

- a) M.T. is a 49-year-old female Caucasian nurse with heart arrhythmia. Her symptoms first began on Thursday, 04/02/2020 with ninety-nine point six (99.6°F) degrees Fahrenheit fever, some body aches and bad headaches. She quickly progressed to fevers, tightness in her chest, shortness of breath, severe fatigue, generalized muscle aches and pains, no appetite, headaches, sore throat, and severe dry cough.

She started **EMERGENCY D-Virus Plan of Care** treatment on Saturday, 04/04/2020 (her virus test resulted positive on Sunday, 04/05/2020). The next day she 'sensed' improvement, but uncertain of any improvement. By day number five (#5), on Wednesday, 04/08/2020, she exclaimed "I can take a deep breath" and she "slept like a baby". By Sunday, 04/12/2020, day number nine (#9), she was completely asymptomatic of all of her symptoms, and **repeat virus** test was done on **Friday, 04/17/2020. It was positive.** The next day, day number fifteen (#15) on Saturday, 04/18/2020, she had a fever of ninety-nine point nine (99.9°F) degrees Fahrenheit and increasing cough, no other symptoms, energy level was still good. She proceeded to the **METABOLIC Plan of Care** that evening. On day number sixteen (#16) and day number seventeen (#17), on Sunday, 04/19/2020 and Monday, 04/20/2020, no fevers and the cough continues improvement on the **METABOLIC Plan of Care.**

- b) R.H. is a 50-year-old male Hispanic insulin-using diabetic and asthmatic. He was confirmed the **virus positive** on Thursday, 04/02/2020. He **failed** a five (5) days course of **Hydroxychloroquine** and **Azithromycin**. He was still experiencing fevers up to one hundred and three point five (103.5°F) degrees Fahrenheit, chills, extreme fatigue, severe generalized muscle achiness and pain, severe headaches, tightness in chest, frequent dry coughing, throat irritation, poor appetite with loss of about five (5 lbs) pounds, shortness of breath, bad dry cough, chest tightness, severe generalized muscle pains, extreme fatigue, and insomnia, and some diarrhea (from the **Hydroxychloroquine** most likely per patient).

Patient underlying condition of Diabetes Mellitus Type II was diagnosed fifteen (15) years earlier and treated with **Insulin**. As a result of this underlying condition, patient suffers from peripheral neuropathy, chronic tinnitus, and balance problems. Patient also suffers from Asthma, for which he treats with a **Symbicort** inhaler, twice a day. There were no other underlying conditions.

Following testing, his reported to his physician, who started the patient on a five (5) day course of treatment with **Hydroxychloroquine** and **Azithromycin**. Upon completion of the five (5) day course of treatment, patient's symptoms had not improved. His physician proceeded to refer the patient to Dr. Meis on Wednesday, 04/08/2020.

On Wednesday, 04/08/2020, Dr. Meis commenced treating the patient with **EMERGENCY D-Virus Plan of Care**. By the next day his fever was coming down. By day number six (#6) the patient no longer suffered with fever. On day number eight (#8) **AnterFeerons** was added to the patient's treatment regimen, after which (approximately two (2) days later) the patient was essentially asymptomatic, with the exception of a persistent cough and chest

tightness. Five (5) days following the introduction of *AnterFeerons*, the patient continued to be asymptomatic and still **positive** with the **virus**. However, the patient was able to **jog two (2) miles** by Saturday, 04/18/2020.

On Saturday, 04/18/2020 the patient started treatment with *METABOLIC Plan of Care*, including *LunCov* and *Gordon-PXL* products. On Wednesday, 04/22/2020 *DetoxHerb-1* and *DetoxHerb-2* were introduced. By Friday, 04/24/2020 the patient had **virus (not detected) negative**, was asymptomatic, feeling very good, with no fever. There was only a slight tightness of chest consistent with his underlying Asthma.

Patient has continued treatment with *ImunStem* and *Aktiffvate*, and is symptom free, as well as **negative (not detected) for virus**.

- c) D.M. is a 64-year-old healthy male Caucasian who had persisting symptoms for three (3) weeks,. He returned with his wife, from NETHERLANDS Holland on Tuesday, 03/17/2020. He was in bed for two (2) days with fever, chills, a cough – dry and also productive at times, chest tightness, loss of his sense of taste and smell, no appetite, slight headaches, burning in his nasal / sinus areas, and mild muscle aches. He was not able to completely recover. He continued to suffer from low energy, poor sense of taste and smell, mild dry cough, some tightness in his chest and inability to take a deep breath, and recurrent mild headaches. He could become winded and light-headed doing outdoor chores. He was **virus** confirmed on Thursday, 04/09/2020.

On Thursday, 04/09/2020 he started the *EMERGENCY D-Virus Plan of Care*. Steadily his symptoms improved until he was asymptomatic by day number seven (#7) of treatment, on Wednesday, 04/15/2020. By Thursday, 04/30/2020 the patient had **virus (not detected) negative**, was asymptomatic, feeling very good, with no fever, and no cough.

Patient has continued treatment with *ImunStem* and *Aktiffvate*, and is symptom free, as well as **negative (not detected) for virus**.

- d) N.M. is a 62-year-old healthy female Caucasian who returned from her trip with her husband from NETHERLANDS Holland on Tuesday, 03/17/2020. She similarly like her husband battled with persisting symptoms for about three (3) weeks. About on Thursday, 03/19/2020 she was in bed for two (2) days. She had a low-grade fever, mild headaches, generalized muscle achiness, bad dry cough, tightness in the chest, shortness of breath, fatigue, poor appetite, and poor sense of taste and smell. She slowly improved, but she had persisting mild / moderate dry and productive cough, chest tightness, fatigue, and some improved appetite. She was having periodic fever as well, none of them ever higher than one hundred (100°F) degrees Fahrenheit. She failed a course of *Ciprofloxacin* and then had tested positive for the **virus** on Saturday, 04/04/2020. On Wednesday, 04/08/2020, she took only one (1) dose each of *Hydroxychloroquine* and *Azithromycin*, then chose the *EMERGENCY D-Virus Plan of Care*.

On Wednesday, 04/08/2020 she started *EMERGENCY D-Virus Plan of Care* (day number one (#1) of treatment). By the next day, on Thursday, 04/09/2020, day number two (#2) of treatment, she already had improved energy and ability to concentrate. Her symptoms steadily improved until she was symptom-free by Wednesday, 04/15/2020, day number eight (#8) of treatment.

Patient has continued treatment with *ImunStem* and *Aktiffvate*, and is symptom free, *COVID-19* test, on Monday, 04/27/2020, was **inconclusive**.

- e) A **fifth virus** positive patient on Saturday, 04/11/2020, started receiving the **EMERGENCY D-Virus Plan of Care** on Wednesday, 04/15/2020.

N.R. is a 28-year-old, healthy Hispanic male, who started symptoms on Friday, 04/10/2020 with fevers up to one hundred and two (102°F) degrees Fahrenheit, shortness of breath, sinus / nasal congestion and pressure, “non-stop” phlegm production, no appetite, extreme fatigue, generalized muscle pain, insomnia, and poor taste and smell.

On Wednesday, 04/15/2020 he started **EMERGENCY D-Virus Plan of Care**. By the next day, day number two (#2) of treatment, on Thursday, 04/16/2020, the patient was already noticing improvement in his taste and smell, and most of his other symptoms, i.e. muscle pain etc. He has been afebrile since day number four (#4), on Saturday, 04/18/2020. Follow up in coming.

- f) A **sixth virus** positive on Tuesday, 04/14/2020, started receiving **EMERGENCY D-Virus Plan of Care** on Tuesday, 04/28/2020.

S.R. is an 86-year-old. Hispanic male who was admitted to the hospital from a nursing home with the **virus** on Tuesday, 04/14/2020. He was in the Intensive Care Unit (ICU) and required the **ventilator** for eight (8) days until Wednesday, 04/22/2020.

He was started on the **EMERGENCY D-Virus Plan of Care in the hospital** on Tuesday, 04/28/2020 and **discharged to home twenty-four (24) hours later**. He is gaining strength and he regained his appetite. The family members are amazed with his quick improvement once he started the **EMERGENCY D-Virus Plan of Care**. He is currently on day number three (#3) of treatment.

Summary:

All patients have become completely asymptomatic by day number seven (#7) to day number nine (#9) of treatment with the **EMERGENCY D-Virus Plan of Care**. Once people are afebrile for three (3) days and with improved cough, current policy allows discontinuation of self-quarantine measures. Up until now, because there has been no effective treatment, the effort of controlling the viral pandemic has necessitated a preventative approach, utilizing social isolation measures and testing. Success with these measures come at great cost both socially and economically. Now with the **EMERGENCY D-Virus Plan of Care** showing effective treatment for the **virus**, the focus can change, at it should, from prevention to treatment. Social isolation and **viral** testing can be significantly adjusted with treatment taking the primary approach of controlling the **virus**. Prompt administration of this treatment will significantly diminish the occurrence of serious cases and need for hospitalization. Confidence can be restored and people can return much more quickly, more likely in a matter of seven (7) to nine (9) days instead of weeks, to a more normal life style.

8.0 STORAGE, HANDLING, AND PRODUCTS**8.1 Storage and Stability**

STORAGE: Store materials at controlled room temperature 20°C (68°F).

STABILITY: *EMERGENCY D-Virus Plan of Care* is chemically stable for two (2) years at room temperature. Do not freeze.

8.2 Product Classification

Dietary Supplement.

9.0 ATTACHMENT LABELS

ImunStem, Aktiffvate, AnterFerron-1, and AnterFerron-2

10.0 HOW SUPPLIED**10.1 Packaging**

PRODUCT	CONTAINER CONTENT	NET CONTENT
<i>AnterFerron-1</i>	1 bottle	1 fl.oz. Liquid
<i>AnterFerron-2</i>	1 bottle	1 fl.oz. Liquid
<i>ImunStem</i>	2 bottle	1 fl.oz. Liquid
<i>Aktiffvate</i>	2 bottle	1 fl.oz. Liquid

AnterFeeron-1

Dietary Supplement

WARNING

Keep out of reach of children
do not use if safety seal is damaged or missing

SUPPLEMENT FACTS		
Serving Size: (1 fl.oz.) (491.50mg)		
Serving Per Container: One (1) serving		
Amount Per Serving		%DV
Bilberry leaf	40mg	**
Graviola	120mg	**
Goldenseal	80mg	**
** Daily Value (DV) not established.		

Other Ingredients: solvents, organic compounds, Chuchuhuasi, Cayenne, Maca, and Turmeric.

STRUCTURE FUNCTION

“Support Immunity” and “Boost Stamina”

“Helps Maintain Joint Health and Flexibility”

“Helps Maintain Cardiovascular Function and a Healthy Circulatory System”

The *AnterFeeron-1* has no side-effects. It will promote the body’s natural cleansing process which may include purging responses such as nausea, diarrhea, vomiting and mucus discharges. Other possible symptoms a person can experience may depend on the person’s previous health issues, which may include headaches, migraines, weakness, muscle aches, joint pain, heart palpitations, inflammation of the throat, excessive bloating, gas, and shortness of breath. These symptoms are only temporary at the time that the patient is being treated with *AnterFeeron*. ONLY USE UNDER THE SUPERVISION OF A PHYSICIAN’S CARE.

Administration: Empty entire contents of *AnterFeeron-1* into a glass cup and swallow entire contents.

Dosage: Take one fluid ounce (1 fl.oz.)

AnterFeeron-1 dietary supplement may support immunity, improve overall health for the human body and maintain good well-being.

WARNING: Not recommended for use by pregnant or nursing women. Should you have any questions regarding the use of *AnterFeeron-1*, please consult your doctor or call the product hot line in U.S.A. at 1.559.781.0658 or 1.559.361.0097. Keep out of reach of children. To be kept in a dry and cool place.

* These statements have not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure or prevent any disease.

AnterFeeron-2

Dietary Supplement

WARNING

Keep out of reach of children
do not use if safety seal is damaged or missing

SUPPLEMENT FACTS		
Serving Size: (1 fl.oz.) (491.50mg)		
Serving Per Container: One (1) serving		
Amount Per Serving		%DV
Astragalus	20mg	**
Reishi	95mg	**
Mistletoe	45mg	**
** Daily Value (DV) not established.		

Other Ingredients: Cat's claw, organic compounds, Echinacea, and Cordyceps.

STRUCTURE FUNCTION

"Promote Bowel Movements"

The *AnterFeeron-2* has no side-effects. It will promote the body's natural cleansing process which may include purging responses such as nausea, diarrhea, vomiting and mucus discharges. Other possible symptoms a person can experience may depend on the person's previous health issues, which may include headaches, migraines, weakness, muscle aches, joint pain, heart palpitations, inflammation of the throat, excessive bloating, gas, and shortness of breath. These symptoms are only temporary at the time that the patient is being treated with *AnterFeeron*. **ONLY USE UNDER THE SUPERVISION OF A PHYSICIAN'S CARE.**

Administration: Empty entire contents of *AnterFeeron-2* into a glass cup and swallow entire contents.

Dosage: Take one fluid ounce (1 fl.oz.)

AnterFeeron-2 dietary supplement may support immunity, improve overall health for the human body and maintain good well-being.

WARNING: Not recommended for use by pregnant or nursing women. Should you have any questions regarding the use of *AnterFeeron-2*, please consult your doctor or call the product hot line in U.S.A. at 1.559.781.0658 or 1.559.361.0097. Keep out of reach of children. To be kept in a dry and cool place.

* These statements have not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure or prevent any disease.

ImunStem[®]

Dietary Supplement

WARNING

Keep out of reach of children
do not use if safety seal is damaged or missing

SUPPLEMENT FACTS		
Serving Size: (0.50ml) (491.50mg)		
Serving Per Container: 25 servings		
Amount Per Serving		%DV
Olive Leaf extract	260mg	**
Yarrow extract	52mg	**
Rosemary extract	63mg	**
** Daily Value (DV) not established.		

Other Ingredients: Organic compounds and solvents, monoterpene, Cassia oil, and Yucca.

STRUCTURE FUNCTION

“Support Immunity” and “Boost Stamina”

“For the Relief of Occasional Sleeplessness”

“Maintains Healthy Lung Function”

“Helps Restore Mental Alertness or Wakefulness when Experiencing Fatigue or Drowsiness”

“Helps You Relax”

ADVERSE ACTIONS

- * In rare circumstances an adverse reaction in the mouth such as “mild blisters” have occurred.
- * A burning sensation in the throat in the beginning of oral treatment may occur, but subsides. If the burning sensation persists, filling gelatin capsules and swallowing may be substituted as an alternative.
- * Vomiting.
- * Yarrow flowers can cause severe allergic skin rashes.

Shake bottle well before using and use dropper to place ½ – ¾ dropperful of *ImunStem* under the tongue. Leave under the tongue for approximately forty (40) seconds and then swallow with a drink of water.

Dosage: Take ½ – ¾ dropperful, 1–4 times a day, as frequently as every 1–3 hours.

ImunStem dietary supplement may support immunity, improve overall health for the human body and maintain good well-being.

WARNING: Not recommended for use by pregnant or nursing women. Should you have any questions regarding the use of *ImunStem*, please consult your doctor or call the product hot line in U.S. at 1.559.781.0658 or 1.559.361.0097. Keep out of reach of children. To be kept in a dry and cool place.

* These statements have not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure or prevent any disease.

Aktiffvate®

Dietary Supplement

WARNING

Keep out of reach of children
do not use if safety seal is damaged or missing

SUPPLEMENT FACTS

Serving Size: (0.50ml) (491.50mg)

Serving Per Container: 25 servings

Amount Per Serving		%DV
Turmeric extract	175mg	**
Cayenne extract	40mg	**
Eucalyptus extract	20mg	**

** Daily Value (DV) not established.

Other Ingredients: Wintergreen, solvents, organic compounds, Yucca, and Olive leaf.

STRUCTURE FUNCTION

“Support Immunity” and “Boost Stamina”

“For the Relief of Occasional Sleeplessness”

“Maintains Healthy Lung Function”

“Helps Restore Mental Alertness or Wakefulness when Experiencing Fatigue or Drowsiness”

“Helps You Relax”

“Helps Maintain Cardiovascular Function and a Healthy Circulatory System”

“Reduces Stress and Frustration”

ADVERSE ACTIONS

* In rare circumstances an adverse reaction in the mouth such as “mild blisters” have occurred.

* A burning sensation in the throat in the beginning of oral treatment may occur, but subsides. If the burning sensation persists, filling gelatin capsules and swallowing may be substituted as an alternative.

* Vomiting.

Shake bottle well before using and use dropper to place ½ – ¾ dropperful of *Aktiffvate* under the tongue. Leave under the tongue for approximately forty (40) seconds and then swallow with a drink of water.

Dosage: Take ½ – ¾ of a dropperful, 1–4 times a day, as frequently as every 1–3 hours.

Aktiffvate dietary supplement may support immunity, improve overall health for the human body and maintain good well-being.

WARNING: Not recommended for use by pregnant or nursing women. Should you have any questions regarding the use of *Aktiffvate*, please consult your doctor or call the product hot line in U.S. at 1.559.781.0658 or 1.559.361.0097. Keep out of reach of children. To be kept in a dry and cool place.

* These statements have not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure or prevent any disease.