Honeywell

Recommended Medical Treatment for Hydrofluoric Acid Exposure



Hydrofluoric Acid Treatment Quick Reference Chart Fold OUT HERE

This Booklet describes the special First Aid and Medical Treatment measures necessary following exposure to or injury from HYDROFLUORIC ACID.

However, it must be emphasized that

PREVENTION

of exposure or injury must be the primary goal.

Preventive measures include:

- 1. Making everyone who handles or uses HF aware of its properties and dangers.
- 2. Training everyone who uses HF in proper handling and safety precautions.
- 3. Utilizing all appropriate engineering controls, and making sure that the controls are maintained and functioning properly.
- 4. Requiring everyone who handles or uses HF to have available the proper safety and personal protective equipment, to be trained to use the equipment, and to always use the equipment when necessary.
- 5. Arranging <u>ahead of time</u> to provide first aid or medical treatment measures if necessary.

If you have questions, comments or suggestions, please write to:

Technical Service Manager - Hydrofluoric Acid Honeywell P. O. Box 1053 101 Columbia Road Morristown, New Jersey 07962-1053

TREATMENT OF HYDROFL

NOTE: In addition to the usual medical history, the physician will find it helpful to obtain the following inform exposed/affected, first aid measures instituted (what, when, how long). Injuries due to dilute HF solutions or low

SKIN E	BURNS	EYE EXPOSURE
		FIRS
CONCENTRATED HF	DILUTE HF	ALL HF
Water Wash THEN Iced Benzalkonium Chloride* 0.13% Soaks OR Calcium Gluconate 2.5% Gel	Water Wash THEN Iced Benzalkonium Chloride* 0.13% Soaks OR Calcium Gluconate 2.5% Gel	Water Wash OR Saline Wash
		MEDICAL
CONCENTRATED HF	DILUTE HF	MEDICAL

¹ This is a brief summary of First Aid and Medical Treatment measures. The text of the brochure "RECOMMENDED MEDICAL TREATMENT FOR HYDROFLUORIC ACID EXPOSURE" must be consulted for more complete information.

² 5% calcium gluconate injections must be used if the soaks or gel do not significantly relieve pain in 30-40 minutes. Injections may also be used as the primary treatment, especially for larger and/or deeper burns.

³ Systemic effects include hypocalcemia, hypomagnesemia, hyperkalemia, cardiac arrhythmias, and altered pulmonary hemodynamics. TREATMENT includes cardiac monitoring, monitoring serum calcium, magnesium, and electrolytes; administration of IV calcium gluconate, correcting magnesium and electrolyte imbalance, and, in extreme cases, hemodialysis.

⁴ Calcium gluconate is normally supplied in ampules containing 10% calcium gluconate. Concentrations less than 10% are obtained by diluting with normal saline.
 * Benzalkonium chloride is a high molecular weight quatemary ammonium compound available as Zephiran* a Registered Trademark of Sanofi Pharmaceuticals, New York, NY 10010 + Registered trademark, Johnson - Merck, Fort Washington, PA 19034

JORIC ACID (HF) EXPOSURE EFERENCE

nation: concentration of HE, date and time of exposure, duration of exposure, how exposure occurred, body parts concentrations of vapors may result in delays in clinical presentation up to 24 hours following exposure.

INHAI	LATION	INGESTION		
T AID				
CONCENTRATED HF	DILUTE HF	ALL HF		
Oxygen AND 2.5% Calcium Gluconate⁴ by Nebulizer	Oxygen THEN Consider 2.5% Calcium Gluconate⁴ by Nebulizer	Do Not Induce Vomiting Milk or Water THEN Milk of Magnesia OR Mylanta®+		
TREATMENT	REATMENT			
CONCENTRATED HF	DILUTE HF	ALL HF		
Observe AND Prophylactic Inhalational Steroids THEN Treat (if necessary) Bronchoconstriction, Pulmonary Edema, Systemic Effects ³	Observe Serious Effects Unlikely Inhalation of HF Fumes from Diluted Acid is Uncommon	Lavage with Calcium Chloride or Calcium Gluconate AND Treat Systemic Effects ³		

For additional reference charts or information on properties, storage and handling, or medical treatment for hydrofluoric acid, contact:

> Honeywell Industrial Fluorines P.O. Box 1053 Morristown, NJ 07962-1053

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This foldout chart is also available as a laminated 15" x 23" wall poster:

In the event of an emergency with this product, call the 24-hour Honeywell emergency telephone number: **800-707-4555 or 602-365-4980**



TABLE OF CONTENTS

PREVENTION
TREATMENT OF HF EXPOSURE: QUICK REFERENCEInside Cover
TABLE OF CONTENTS
INTRODUCTION
ACUTE TOXICITY2Skin Contact2Systemic Toxicity3Eye Contact4Inhalation4Ingestion4
CHRONIC TOXICITY
FIRST AID TREATMENT FOR HYDROFLUORIC ACID BURNS.5Skin Contact.5Eye Contact.6Inhalation.6Ingestion.7
MEDICAL TREATMENT FOR HYDROFLUORIC ACID BURNS7Burns of the Skin – General7Quaternary Ammonium Compounds7Calcium Gluconate Gel8Calcium Gluconate Injections8Calcium Gluconate Solution9Burns of the Fingers and Nails9Intra-arterial and Intravenous Calcium Infusion9Additional Measures10Other and Unproven Therapies10Systemic Absorption and Metabolic Effects10Hemodialysis11Inhalation Injuries12Ingestion Injuries12
REFERENCES
APPENDIX - FIRST AID AND MEDICAL SUPPLIES
INDEX



INTRODUCTION

Because the medical treatment of hydrofluoric acid exposure is so specialized and differs from the treatment of other inorganic acid exposures, not all physicians may be aware of appropriate treatment measures. It is recommended that HF users <u>make arrangements ahead of time</u> with local medical resources to be sure that users are familiar with first aid measures and that professional personnel are familiar with the toxicity of HF and the treatment of HF exposure. This would include, at a minimum, thoroughly reviewing this booklet and making sure that treatment facilities and supplies are available.

Hydrofluoric acid (CAS # 7664-39-3) is very aggressive physiologically because of the fluoride ion. Both anhydrous hydrofluoric acid (hydrogen fluoride) and its solutions are clear, colorless liquids. When exposed to air, concentrated solutions and anhydrous hydrofluoric acid produce **pungent fumes which are especially dangerous**. Unless heated, dilute concentrations of hydrofluoric acid in water (e.g., less than 40% HF) do not produce significant vapor concentrations.

NOTE: Persons unfamiliar with <u>hydrofluoric acid</u> often mistake it for, or confuse it with, hydro<u>ch</u>loric acid. Although hydrofluoric acid (HF) and hydrochloric acid (HCl) have similar sounding names, the toxicity of these two acids is very different. To decrease or avoid confusion, we recommend that *HYDROFLUORIC ACID* and *HYDROGEN FLUORIDE* be referred to as "HF".

HF is primarily an industrial raw material. It is used in stainless steel manufacturing, iron and steel foundries, metal finishing, aluminum manufacturing, inorganic and organic chemical manufacturing, petroleum refining, mineral processing, glassmaking and electronic components manufacturing. It is also used in certain industrial and consumer cleaning compounds. However, its use in consumer products is discouraged because of the hazards described herein.

Most non-industrial burns are caused by dilute concentrations of HF (e.g, less than 15% HF). Most of the HF used in the electronics industry is less than 50%. However, many industrial uses of HF involve concentrated (50-100%) HF.

WARNING: BURNS WITH CONCENTRATED HF ARE USUALLY VERY SERIOUS, WITH THE POTENTIAL

FOR SIGNIFICANT COMPLICATIONS DUE TO FLU-ORIDE TOXICITY. CONCENTRATED HF, LIQUID OR VAPOR, MAY CAUSE SEVERE BURNS, METABOLIC IMBALANCES, PULMONARY EDEMA AND LIFE THREATENING CARDIAC ARRYTHMIAS. EVEN MODERATE EXPOSURES TO <u>CONCENTRATED</u> HF MAY RAPIDLY PROGRESS TO FATALITY IF LEFT UNTREATED.

Honeywell is the world's leading supplier of hydrofluoric acid. The recommended medical procedures described in this brochure are based on a review of the available literature, shared experiences with others who have dealt with the health effects of HF, the personal knowledge and experiences of Honeywell physicians, nurses and other professionals in dealing with the unique hazards of this product, and experimental laboratory work sponsored by Honeywell.

Every effort must be made to prevent exposure to hydrofluoric acid. If exposure does occur, the specialized procedures which follow are recommended to avoid the very serious consequences that might otherwise occur.

ACUTE TOXICITY Skin Contact

Hydrofluoric acid (HF) can cause serious, painful burns of the skin. Specialized first aid and medical treatment is required. **Burns larger than 25** square inches (160 square cm) may result in serious systemic toxicity.

Hydrofluoric acid is a highly corrosive acid which can severely burn skin, eyes, and mucous membranes. The vapors from anhydrous hydrofluoric acid or its concentrated solutions can also burn these tissues.

Hydrofluoric acid is similar to other acids in that the initial extent of a burn depends on the concentration, the temperature and the duration of contact with the acid. Hydrofluoric acid differs, however, from other acids because the fluoride ion readily penetrates the skin, causing destruction of deep tissue layers. Unlike other acids which are rapidly neutralized, this process may continue for days if left untreated.

Strong acid concentrations (over 50%), and particularly anhydrous HF (AHF or 100% HF), cause immediate, severe, burning pain and a whitish discoloration of the skin which usually proceeds to blister formation.



Exposure to HF vapors can also result in similar burns.

In contrast to the immediate effects of concentrated HF, the effects of contact with more dilute hydrofluoric acid or its vapors may be delayed, and this is one of the problems with the recognition of some HF burns. Skin contact with acid concentrations in the 20% to 50% range may not produce clinical signs or symptoms for one to eight hours. With concentrations less than 20%, the latent period may be up to twenty-four hours. HF concentrations as low as 2% may cause symptoms if the skin contact time is long enough. (1)

HF skin burns are usually accompanied by severe, throbbing pain which is thought to be due to irritation of nerve endings by increased levels of potassium ions entering the extracellular space to compensate for the reduced levels of calcium ions, which have been bound to the fluoride. <u>Relief of pain is an important guide to the success of treatment.</u>

The usual initial signs of an HF burn are redness, edema, and blistering. With more concentrated acids, a blanched white area appears. The fluoride ion penetrates the upper layers of the skin. A thick granular exudate may form under blisters due to liquefaction necrosis. In rare (and untreated) cases, there may be penetration to underlying bone with decalcification. **HF burns require immediate and specialized first aid and medical treatment** (2, 3, 4, 5, 6, 7) differing from the treatment of other chemical burns. If untreated or if improperly treated, permanent damage, disability or death may result. (8) If, however, the burns are promptly and properly recognized and managed, the results of treatment are generally favorable.

Treatment is directed toward binding the fluoride ions to prevent tissue destruction. High molecular weight quaternary ammonium compounds, e.g. benzalkonium chloride (Zephiran®), are used as soaking agents. (9, 10, 11) Calcium gluconate as a gel or ointment can be applied locally, and calcium gluconate solution may be injected (subcutaneously, intravenously, or intra-arterially), inhaled, or used as an irrigant. (3, 12, 13, 14, 15)

Speed is of the essence. Delays in first aid care or medical treatment or improper medical treatment will likely result in greater damage or may, in some cases, result in a fatal outcome.

Systemic Toxicity

To produce HF, calcium fluoride is reacted with sulfuric acid:

$$CaF_2 + H_2SO_4 \rightarrow 2HF + CaSO_4$$

This production process requires a great deal of energy to accomplish. On the other hand, in the body:

$$2HF + Ca^{++} \rightarrow CaF_2$$

This process releases energy, and therefore occurs very readily. The toxic effect of HF on body calcium is certainly more complicated than this. There is some evidence that fluoride may combine with calcium and phosphate, so that five calcium ions are tied up for each fluoride ion (e.g. $Ca_5F(PO_4)_3$), rather than two. There is also some evidence that there may be high intracellular levels of calcium in some tissues, rather than low levels, as would intuitively be expected. (16) However, the reaction of fluoride with body calcium is one of the major toxic effects and forms the basis for many treatment recommendations.

One of the most serious consequences of severe exposure to HF by any route is the marked lowering of serum calcium (hypocalcemia) and other metabolic changes, which may result in a fatal outcome if not recognized and treated. Hypocalcemia should be considered a potential risk in all instances of inhalation or ingestion, and whenever skin burns exceed 25 square inches, (160 square centimeters). Serum magnesium may also be lowered, and elevations in serum potassium have been reported to further complicate the metabolic imbalances which will need to be monitored and corrected. (16, 17, 18) High levels of fluorides have been noted both in the blood and body organs. Hemodialysis has been reported to be effective therapy for cases of severe systemic intoxication. (19, 20, 21) Treatment for shock may also be required as for other severe injuries.

Other effects reported from fluoride exposure include coagulation defects and inhibition of a number of enzymes, including preglycolytic enzymes, phosphatases and cholinesterase. The results of this enzyme inhibition include inhibition of cellular glucose phosphorylation and subsequent glycolysis, inhibition of respiration, and increased sensitivity of cholinergic mechanisms to acetyl cholinesterase. (22) While hypocalcemia has been traditionally considered the major systemic effect of severe poisoning with HF, it is apparent that hypomagnesemia, hyperkalemia, the cardiodepressing and vasodilating effects of fluoride and effects on pulmonary hemodynamics and systemic capacitance vessels, including an increase in pulmonary vascular resistance, all play a role in systemic toxicity. Although some of these effects have been described, the implications for therapeutic measures have not been well defined. (23, 24)

Eye Contact

Hydrofluoric acid can cause severe eye burns with destruction or opacification of the cornea. Blindness may result from severe or untreated exposures. Immediate first aid and specialized medical care is required. (3,13)

Inhalation

Hydrofluoric acid fumes may cause laryngospasm, laryngeal edema, bronchospasm and/or acute pulmonary edema. Acute symptoms may include coughing, choking, chest tightness, chills, fever and cyanosis. Many reported fatalities from HF exposures have been due to severe pulmonary edema (coupled with systemic toxicity) that did not respond to medical treatment.

Burns from vapors or liquid contact to the oropharyngeal mucosa or upper airway may cause severe swelling to the point of requiring a tracheostomy. It is recommended that all patients with such exposures be hospitalized for observation and/or treatment.

Because of the strong irritant nature of hydrofluoric acid, an individual inhaling HF vapors or fumes will usually experience upper respiratory injury, with mucous membrane irritation and inflammation as well as cough. All individuals suspected of having inhaled HF should be observed for pulmonary effects. This would include those individuals with significant upper respiratory irritation, bronchoconstriction by pulmonary auscultation or spirometry, and any individual with HF exposure to the head, chest or neck areas. It has been reported that pulmonary edema may be delayed for several hours and even up to two days. If there is no initial upper respiratory irritation, significant inhalation exposure can generally be ruled out.

The Permissible Exposure Limit (PEL) set by the U.S. Occupational Safety and Health Administration

(OSHA) is a time weighted average exposure for 8 hours of 3 ppm. (25) The American Conference of Governmental Industrial Hygienists (ACGIH) recommends a ceiling level of 3 ppm or 2.3 mg/m³. (26, 27) The National Institute for Occupational Safety and Health (NIOSH) has established the level that is immediately dangerous to life and health (IDLH) at 30 ppm. (28, 29) The American Industrial Hygiene Association has published an Emergency Response Planning Guideline setting 50 ppm as the maximum level below which nearly all individuals could be exposed for one hour without experiencing or developing life-threatening health effects (ERPG-3), 20 ppm as the maximum level below which nearly all individuals could be exposed for one hour without developing irreversible health effects or symptoms which would impair taking protective action (ERPG-2), and 2 ppm as the maximum level below which nearly all individuals could be exposed up to one hour without experiencing other than mild, transient adverse health effects (ERPG-1). (30)

Ingestion

If hydrofluoric acid is ingested, severe burns to the mouth, esophagus and stomach may occur. Severe systemic effects usually also occur. Ingestion of even small amounts of dilute HF have resulted in death. (31)

CHRONIC TOXICITY

Because it is a strong irritant, HF has not been found to cause chronic toxicity nor has it been the subject of long term toxicity studies or testing. Once HF enters the body, it is expected that the fluoride ion would be the major concern from a chronic toxicity standpoint. Chronic toxicity from long term, high exposure to fluoride salts has been reported to result in tooth mottling in children, bone fluorosis and sometimes osteosclerosis in adults and children.

Skeletal fluorosis is known to be associated with excessive exposure to fluoride compounds. Cases of skeletal fluorosis have been reported in populations exposed to naturally occurring drinking water containing greater than 10 ppm of fluoride ion and in individuals exposed to high levels of fluoride containing dusts. However, skeletal fluorosis has not been reported as a consequence of hydrofluoric acid exposure.

Because of the use of fluoride to prevent dental caries, there is ongoing evaluation of fluorides for

Honeywell Ver. 1.0 the potential to cause cancer. There is no evidence that fluoride is genotoxic except in some in vitro assays at cytotoxic concentrations. Epidemiological studies have not demonstrated an association between fluoride in drinking water and an increase in cancer. The International Agency for Research on Cancer (IARC) has not classified hydrogen fluoride as to its human carcinogenicity, and neither fluorides nor HF are listed by IARC, NTP, OSHA, ACGIH, NIOSH, the State of California or other governmental agencies as causing cancer. (32, 33, 34) In animal studies, fluoride salts have caused effects in progeny only at high, maternally toxic levels. Some animal studies have shown effects on male fertility, e.g. decreased sperm counts. (34) Fluoride exposures should be kept below recommended levels to assure no adverse effects to the developing fetal skeletal system or teeth.

Monitoring of urine for fluorides is an accepted method of determining exposure. (35) Urine fluoride levels above 3 mg/liter at the beginning of a workshift, or above 10 mg/liter at the end of a workshift, may indicate excessive absorption of fluoride. It should be noted that fluorides are often present in significant amounts in persons not occupationally exposed (because of dietary sources of fluoride such as tea), and that the urine fluoride determination is not specific for HF. (36, 37)

FIRST AID TREATMENT FOR HYDROFLUORIC ACID BURNS

In Case of Contact or Suspected Contact with Hydrofluoric Acid:

A. Skin Contact

- 1. Move victim immediately under safety shower or other water source and flush affected area thoroughly with large amounts of running water. Speed and thoroughness in washing off the acid is of primary importance.
- 2. Begin flushing even before removing clothing. Remove all contaminated clothing while continuing to flush with water.
- 3. Rinse with large amounts of running water. If 0.13% benzalkonium chloride (Zephiran[®]) solution or 2.5% calcium gluconate gel are available, the rinsing may be limited to 5 minutes, with the soaks or gel applied as soon as the rinsing is

stopped. If benzalkonium chloride (Zephiran[®]) or calcium gluconate gel is not available, rinsing must continue until medical treatment is rendered.

- 4. While the victim is being rinsed with water, someone should alert first aid or medical personnel and arrange for subsequent treatment.
- 5. Immediately after thorough washing, use <u>one</u> of the measures below:
 - a. Begin soaking the affected areas in iced <u>0.13%</u> <u>benzalkonium chloride (Zephiran[®]) solution</u>.

Use ice cubes, <u>not</u> shaved ice, in order to prevent frostbite.

If immersion is not practical, towels should be soaked with iced 0.13% benzalkonium chloride (Zephiran[®]) solution and used as compresses for the burned area. Compresses should be changed every two to four minutes.

Do not use benzalkonium chloride (Zephiran[®]) solution for burns of the eyes. Exercise caution when using benzalkonium chloride (Zephiran[®]) solution near the eyes as it is an eye irritant.

Benzalkonium chloride (Zephiran[®]) soaks or compresses should be continued until pain is relieved or until more definitive medical treatment is provided.

b. Start massaging <u>2.5% calcium gluconate gel</u> into the burn site.

Apply gel frequently and massage continuously until pain and/or redness disappear or until more definitive medical care is given.

It is advisable for the individual applying the calcium gluconate gel to wear surgical gloves to prevent a possible secondary HF burn.

NOTE: Clinical experience has shown that both benzalkonium chloride (Zephiran[®]) and calcium gluconate gel are effective when used correctly in appropriate situations. In an animal model, benzalkonium chloride (Zephiran[®]) soaks are superior to calcium gluconate gel under the experimental conditions used. (38, 39)

- 6. After treatment of burned areas is begun, the victim should be examined to ensure there are no other burn sites which have been overlooked.
- 7. Arrange to have the victim seen by a physician. (If burns are small and/or caused by weak acid,



and treatment has been provided by an experienced individual, evaluation by a physician may not be necessary.) During transportation to a medical facility or while waiting for a physician to see the victim, continue the benzalkonium chloride (Zephiran®) soaks or compresses or continue massaging calcium gluconate gel. In many situations, particularly for minor burns covering a small skin area or for burns caused by dilute HF, continued treatment with soaks or gel may be effective as the sole type of medical care. All persons with extensive burns or burns with significant blister formation or with the appearance of whitish or dead skin need to be seen by a physician. All persons with HF burns which do not respond to either calcium gluconate gel or benzalkonium chloride (Zephiran®) soaks or compresses within 30 minutes should be evaluated by a physician.

- 8. The physician may advise continuation of benzalkonium chloride (Zephiran[®]) soaks or *calcium gluconate gel.*
 - a. If the physician advises continued treatment with benzalkonium chloride (Zephiran®) soaks or compresses, the soaks or compresses are usually required for 2 to 4 hours. Significant relief of pain should be noted within the first 30 minutes. If this does not occur, the victim must be seen by a physician and more definitive care instituted. If the pain is substantially relieved within the first 30 minutes. continue the treatment for a total of two hours. After that time, discontinue treatment and observe for the recurrence of pain. If pain recurs, continue soaks or compresses until relief of pain occurs. Soaking for six hours is sometimes needed. (Note: Because prolonged immersion in the ice bath may result in discomfort, relief may be obtained by removing the part from the bath every ten minutes for a minute or so and then reimmersing it. After the initial 30-60 minutes of treatment. less ice can be used so the bath is cool rather than cold.)
 - b. *Calcium gluconate gel* may be used for several hours or even repeated over a period of a few days. However, if significant relief of pain does not occur within 30 to 40 minutes, more <u>definitive</u> treatment will be required. For small burns, or burns of the face, ears, and near

mucous membranes, calcium gluconate gel may be very useful. The gel is applied frequently and massaged into the burned area. This is continued until relief is obtained or further medical care is available.

9. For serious burns, medical attention must be provided as quickly as possible.

For minor burns, if first aid treatment does not alleviate symptoms or if symptoms persist or recur, medical attention must be sought.

B. Eye Contact

- 1. Immediately flush the eyes for at least 15 minutes with large amounts of gently flowing water. Hold the eyelids open and away from the eye during irrigation to allow thorough flushing of the eyes. Do not use the benzalkonium chloride (Zephiran*) solutions described for skin treatment. If the person is wearing contact lenses, the lenses should be removed, if possible. However, flushing with water should not be interrupted, and the lenses should be removed by a person who is qualified to do so. If sterile 1% calcium gluconate solution is available, washing may be limited to 5 minutes, after which the 1% calcium gluconate solution should be used repeatedly to irrigate the eye using a syringe.
- 2. Take the victim to a doctor, preferably an eye specialist, as soon as possible. Ice water compresses may be applied to the eyes while transporting the victim to the doctor.
- 3. If a physician is not immediately available, apply one or two drops of 0.5% tetracaine hydrochloride, 0.5% proparacaine, or other aqueous, topical ophthalmic anesthetic and continue irrigation. Use no other medications unless instructed to do so by a physician. Rubbing of the eyes is to be avoided.
- 4. An Eye Irrigator[™] may be useful in delivering flushing or treating solutions, including calcium gluconate solution, to the eye. (See page 15)

C. For Inhalation of Vapors:

- 1. Immediately move victim to fresh air and get medical attention.
- 2. Keep victim warm, quiet and comfortable.
- 3. If breathing has stopped, start artificial respiration at once. Make sure mouth and throat are free of foreign material.



- 4. Oxygen should be administered as soon as possible by a trained individual. Continue oxygen while awaiting medical attention unless instructed otherwise by a physician.
- 5. A nebulized solution of 2.5% calcium gluconate may be administered with oxygen by inhalation.
- 6. Do not give stimulants unless instructed to do so by a physician.
- 7. The victim should be examined by a physician and held under observation for at least a 24 hour period.
- 8. Vapor exposures can cause skin and mucous membrane burns as well as damage to pulmonary tissue. Vapor burns to the skin are treated the same as liquid HF burns.

D. If Acid is Ingested:

- 1. Have the victim drink large amounts of water as quickly as possible to dilute the acid. Do not induce vomiting. Do not give emetics or baking soda. Never give anything by mouth to an unconscious person.
- Give several glasses of milk or several ounces of milk of magnesia, Mylanta[®], Maalox[®], etc or grind up and administer up to 30 Tums[™], Caltrate[™] or other antacid tablets with water. The calcium or magnesium in these compounds may act as an antidote.
- 3. Get immediate medical attention. Ingestion of HF is a life-threatening emergency.

MEDICAL TREATMENT FOR HYDROFLUORIC ACID BURNS

Burns of the Skin – General

Burns from dilute acid are difficult to distinguish from other chemical burns and usually appear as areas of erythema. However, they may progress, if not treated, to areas of blistering, necrosis or ulceration. Burns from more concentrated acid have a rather characteristic appearance and present as severely reddened, swollen areas with blanched, whitish regions which rapidly progress to blistering and necrosis. A thick granular exudate usually appears under these blisters which requires debridement and removal.

Hydrofluoric acid burns cause extreme pain. The pain is thought to result from nerve ending irritation due to increased levels of potassium ions in extracellular spaces to compensate for the reduced levels of calcium ions which have been bound by the fluoride. **Relief** of pain is an excellent indication of the success of treatment and, therefore, local anesthetics should be avoided.

Many different types of therapies have been suggested for HF burns. The aim of all treatment is to chemically sequester the fluoride ion and to prevent extensive, deep-tissue destruction. (38,39)

After treatment of recognized burned areas is begun, the victim should be carefully examined to insure there are no other burn sites which may have been overlooked.

Quaternary Ammonium Compounds

Most HF burns can be satisfactorily treated by immersion of the burned part in an iced, aqueous solution of a quaternary ammonium compound. Two solutions have been clinically successful, 0.13% benzalkonium chloride (e.g. Zephiran®) or 0.2% benzethonium chloride (e.g., Hyamine® 1622). Because of its availability as a non-prescription drug, benzalkonium chloride (Zephiran®) is recommended in the United States.

The solutions should be cooled with ice cubes. (Shaved or crushed ice may cause excessive cooling, with the danger of frostbite.)

If immersion in the solution is not practical, soaked compresses of the same iced solution should be applied to the burned area. The immersion or compresses should be used **for at least two hours**. Compresses should be changed or soaked with additional solution approximately every two to four minutes.

If blisters are present, they should be opened and drained and necrotic tissue should be debrided by a physician or qualified health care practitioner as soon as possible. However, immersion in benzalkonium chloride (Zephiran[®]) or use of compresses should not be delayed if debridement cannot be accomplished immediately.

Prolonged immersion in the iced benzalkonium chloride (Zephiran[®]) bath may result in discomfort due to excess chilling; relief may be obtained by removing the part from the bath every ten to fifteen minutes for a few minutes and then reimmersing it. After the initial 30-60 minutes of treatment, less ice can be used so the bath is cool rather than cold. The success of this treatment is indicated by relief of the severe pain in the burned area. If there is no significant relief of pain within 30 to 40 minutes, the use of 5% calcium gluconate injections may be necessary. If pain recurs when the treatment is stopped at the end of the first two hours, immersion or compresses should be resumed until pain is relieved. A total of four to six hours immersion or use of compresses of benzalkonium chloride (Zephiran[®]) is usually required for the treatment of most burns. No further treatment will be required in many instances. The use of iced quaternary ammonium compound solutions offers several advantages:

- reduction of local pain
- · possible slowing of the rate of tissue destruction
- possible slowing of the passage of the fluoride ion into tissues and into the bloodstream

Large burns, serious burns due to concentrated HF, or burns with delayed treatment will probably require the use of calcium gluconate injections in addition to or instead of the benzalkonium chloride (Zephiran[®]) soaks.

Quaternary ammonium compounds should not be used for burns on the face, ears or other sensitive areas due to their irritating nature. It is preferable to use calcium gluconate gel or calcium gluconate injection in these areas.

Calcium Gluconate Gel

Calcium gluconate gel, consisting of 2.5% USP calcium gluconate in a surgical water soluble lubricant, is widely used for first aid and/or primary treatment of HF burns of the skin. The gel is convenient to carry and can be used to initially treat small burns that might occur away from medical care. (The gel is not recommended for burns with concentrated HF except as a first aid measure.) The gel is used by massaging it promptly and repeatedly into the burned area, until pain is relieved. If possible, surgical gloves should be worn during initial application of the gel, so the person providing treatment will not receive a secondary burn. This treatment can be started without waiting for medical direction.

If used as the only method of treatment, liberal quantities of calcium gluconate gel must be massaged into the burned area continuously for up to several hours. Relief of pain can be used to assess the efficacy of this treatment. If good relief of pain is not obtained after 30-40 minutes, alternate methods of treatment such as calcium gluconate injections or benzalkonium chloride (Zephiran[®]) soaks should be considered.

The gel is especially useful for burns on the face, particularly near the mouth and eyes or on the ears. It may be convenient to use the gel for very small burns where the victim can easily apply and massage the gel into the burned area. Use of the gel may be more convenient for dilute acid burns such as occur with commercial products like rust removers, aluminum cleaners or etching solutions.

Calcium Gluconate Injections

After first aid measures have been taken, injection of a 5% calcium gluconate solution is indicated as the primary medical treatment for large burns (over 25 square inches or 160 square centimeters). For smaller burns, if benzalkonium chloride (Zephiran[®]) soaks or calcium gluconate gel do not result significant relief of pain within 30 to 40 minutes, injection of calcium gluconate solution is indicated. Injection of calcium gluconate solution may also be indicated for burns in which treatment has been delayed.

The physician should inject sterile 5% aqueous calcium gluconate beneath, around and into the burned area. Calcium gluconate is packaged as a 10% solution, and must be diluted 50-50 (equal parts) with normal saline. (Note: DO NOT USE calcium chloride, which is corrosive and may result in additional damage.)

If subcutaneous calcium gluconate injections are used, the amount injected initially is small and should not exceed 0.5 cc per square centimeter of affected skin surface. The injections should not distort the appearance of the skin. A small-gauge needle (#27 - #30) should be used, and the burned area should be injected through multiple sites. With successful treatment, pain relief following injection of 5% calcium gluconate solution is very rapid. The patient can usually advise when the pain stops, and this is an indicator of adequate treatment. Multiple injections in skin that has compromised integrity may increase the risk of infection, and the use of antibiotic creams such as Silvadene® (silver sulfadiazine) or Garamycin[®] (gentamicin sulfate cream) should be considered following such treatment. Local anesthetics should not be used since they mask pain relief which is an important indication of adequacy of treatment.

Some physicians prefer using calcium gluconate injections initially as the primary treatment, instead of using quaternary ammonium compound soaks or compresses or using calcium gluconate gel. Injections often are not necessary when there has been early and adequate treatment with soaks or gel.

Calcium Gluconate Solution

In some instances, a 5% or 10% calcium gluconate solution may be used in compresses or for irrigation. For example, irrigating with a calcium gluconate solution may be the best treatment should HF enter the external ear canal. In this instance, referral to an otolaryngologist may also be needed.

Burns of the Fingers and Nails

Burns of the fingers often create special problems in treatment. Finger and toe nails permit penetration of fluoride ions but prevent soaks or gels from being effective. It may be necessary to drill, split or even remove nails to allow the topical methods of treatment to be effective. One author has cautioned that removal of the nail should rarely be necessary in the case of dilute HF acid (less than 10%) burns. (40) The treating physician must consider the morbidity associated with removal of the nail versus the need to treat the HF exposure.

If immersion in benzalkonium chloride (Zephiran[®]) solution is started immediately, it may be possible to avoid removing the nail. Sometimes better penetration under the nail can be successfully accomplished by splitting the nail or by drilling several burr holes in the nail using a large gauge needle or a nail drill. If calcium gluconate injection is used as treatment, the nail may still need to be split or removed. If nail removal is necessary, using a short acting regional or ring-block anesthetic may facilitate this procedure and not interfere with using pain relief as an indicator of effective treatment. When using calcium gluconate injections in the digits, care must be taken to inject the solution cautiously so as to avoid compromising the circulation in these areas.

As an alternative to using benzalkonium chloride (Zephiran[®]) soaks, experience has shown that some finger or hand burns can be treated by using a glove filled with calcium gluconate gel. Initially, calcium gluconate gel should be massaged into the burned area. Following this, an oversize surgical glove should be partially filled with calcium gluconate gel,

and the hand inserted into the glove. The gloved hand may be immersed in ice water, if available, which may aid pain relief. This treatment works best for burns where there is no blistering, or after the burns have been debrided. As in other cases where calcium gluconate gel is used, alternate methods of treatment should be considered if good relief of pain is not achieved within 30 - 45 minutes. If pain is relieved, the glove should remain in place for three to for hours.

Intra-arterial and Intravenous Calcium Infusion

Reports in the literature have described the use of intra-arterial injection or infusion of dilute calcium gluconate solutions to treat HF burns of the hand and digits, usually from prolonged contact with dilute HF, which do not respond to other methods, either due to inadequate or improper treatment, or in cases where treatment has been greatly delayed. The method is described as follows:

"A long catheter was inserted percutaneously into the radial artery using standard aseptic technique. Intra-arterial catheter placement was confirmed by pressure transducer and oscilloscope. If the burn involved only the thumb, index, or long fingers, the catheter was advanced only a few centimeters proximally in preparation for digital subtraction arteriography. If the burn involved the ring or small fingers, the catheter was advanced proximally into the brachial artery because access to the ulnar circulation was necessary.

Following satisfactory placement of the arterial catheter, we performed digital subtraction arteriography on all patients in our series to identify the origin of vascular supply to digits involved.

Once the tip of the arterial catheter was in the desired location, a dilute preparation of calcium [gluconate] (10 ml of a 10% solution mixed in 40 to 50 ml 5% dextrose) was infused with a pump apparatus into the catheter over four hours. We generally have used calcium gluconate... Each patient was observed closely during the infusion period for progression of symptoms and potential complications of the procedure, such as alterations of distal vascular supply.

Following the four-hour infusion, the arterial catheter was maintained in place in the usual

manner while the patient underwent an observation period. If typical HF pain returned within four hours, a second calcium infusion was repeated until the patient was pain free four hours following completion of the calcium infusion". (14)

This method, although rather involved, should be considered in selected cases, especially where inadequate or delayed treatment has occurred.

There are now several reports of the successful use of intravenous calcium gluconate to treat HF burns of the upper extremity. (41, 42, 43) Graudins, et al. describe their method:

An intravenous catheter was placed on the dorsum of the affected hand. The superficial veins were exsanguinated by elevation. A double-cuffed pneumatic tourniquet was applied above the elbow, inflated to 100 mm Hg above systolic blood pressure, and 10 ml of 10% calcium gluconate diluted with 30 to 40 ml of 0.9% saline solution was then infused. Ischemia was maintained for 25 minutes; the cuff was sequentially released over 3 to 5 minutes.

This method was most successful for burns due to dilute acid. If the use of intravenous calcium gluconate was not successful in relieving pain (which occurred with burns due to 49% HF, the highest concentration seen in the series of patients), Graudins et al. turned to intra-arterial calcium gluconate.

Additional Measures

Where blistering and/or necrosis occur, early debridement may facilitate healing.

In instances of extensive burns, skin grafting has occasionally been required, but the need for this treatment should be markedly reduced by immediate and aggressive primary treatment.

Follow-up care requires monitoring to prevent secondary infections. The use of antibiotic creams such as Silvadene[®] or Garamycin[®] has proven effective. HF burns may heal slowly, but if properly treated most heal with little or no scarring in 14 to 28 days.

Other and Unproven Therapies

The use of intravenous calcium gluconate is discussed above. Both Williams, <u>et al.</u> (44) and Cox, <u>et al.</u> (45) have discussed the use of intravenous magnesium sulfate to treat localized moderate to serious skin burns. Using either a rat or a rabbit model, the authors administered intravenous magnesium sulfate. Cox used a 0.2 mEq bolus over two minutes, followed by a slower infusion of 0.2 mEq per hour for four hours, with a total of 1.0 mEq/kg magnesium sulfate administered. Williams administered 8 mg/kg over five minutes or 160 mg/kg over 10 minutes. These authors compare this dose to the amount of magnesium sulfate, infused more slowly, used in the treatment of eclampsia.

Dunn, <u>et al</u>. (38) have shown effectiveness of locally applied calcium acetate solution, 10% in water at room temperature, in an animal model.

Seyb et al. (47), performed an experiment in rats using a topically applied solution of 50% aqueous dimethyl sulfoxide (DMSO) containing calcium gluconate (20% wt/vol). This treatment gave results comparable to injecting 10% calcium gluconate or 10% magnesium sulfate, and was superior to calcium gluconate gel in treating experimental HF burns.

It should be noted that many of these therapies, while promising, have been tested to a limited degree, if at all, in humans.

A product developed in France, "Hexafluorine" (46), has been marketed in Europe and the United States for the emergency rinsing of HF skin and eye exposure. Documentation of effectiveness and experience with this product are lacking.

Systemic Absorption and Metabolic Effects

Significant amounts of fluoride ion may be absorbed by skin contact, inhalation, or by ingestion. If systemic absorption of fluoride occurs, hypocalcemia, hypomagnesemia and hyperkalemia may also occur. All of these parameters need to be monitored and appropriate therapeutic measures instituted. The patient should be observed for clinical signs of hypocalcemia following ingestion or inhalation or following extensive burns greater than 25 square inches. Serum calcium determinations must be performed immediately and periodically to monitor and treat hypocalcemia. Severe lowering of serum calcium levels can occur within one to two hours even with HF burns covering less than 2.5% of body surface area. (8)Continuous EKG monitoring to observe prolongation of the Q-T interval may be useful to detect early changes in serum calcium, although profound hypocalcemia following HF exposure has been reported in the



absence of EKG changes or in the absence of other signs of tetany.

The fall in serum calcium may occur precipitously following HF exposure. In two reported cases of exposure to anhydrous HF, the serum calcium fell to levels around 3 milliequivalents per liter (mEq/L) [normal = 8.8 - 10.3 mEq/L] within one to three hours of exposure. (8)

If necessary, aqueous calcium gluconate may be given intravenously. Calcium gluconate as a 10% solution must be given <u>slowly</u> since excess calcium can produce vagal bradycardia, ventricular arrhythmias and ventricular fibrillation. The IV calcium gluconate should be repeated until serum calcium levels return to, and remain at, normal levels. In one fatal case, 280 mEq of calcium over four hours was not sufficient to correct the profound hypocalcemia. (8) Without additional measures such as hemodialysis, it may not be possible to correct extreme hypocalcemia.

Serum magnesium levels should also be monitored and magnesium loss should be replaced intravenously if indicated. Yamaura, et al. have reported a case of HF exposure in which prolonged QT interval occurred, in which ionized calcium levels were relatively high but the magnesium level was low. (48) Serum potassium must also be carefully monitored. Significant elevations of serum potassium have been noted in cases of fluoride toxicity and also in laboratory studies. Hyperkalemia has also been implicated as a causative factor in cardiovascular collapse. The use of quinidine may be helpful in preventing this serious complication. (20)

Hemodialysis with fluoride free water (and normal to low potassium and slightly higher calcium concentrations), in conjunction with other treatments mentioned, should be considered in all cases of serious burns and may need to be repeated if indicated. (19, 20, 21) Serum fluoride levels should be monitored. Normal plasma fluoride levels may differ because of various methodologies and analytical techniques. The decision to use dialysis should be based on the HF exposure (concentration, body surface area) and the clinical condition of the patient, including the serum levels of fluoride, calcium and potassium.

Primary excision has been recommended by some practitioners as a method of reducing systemic absorption of fluoride. (49) While this could in some instances be life saving, it is a rather drastic measure. It is likely that renal dialysis could be

used to effectively treat systemic toxicity and would not result in the disfigurement, disability, or morbidity which could be associated with primary excision.

Eye Injuries

HF can cause severe eye burns, which, if not properly treated, may result in scarring and blindness. The prognosis is not good if first aid treatment is delayed or inadequate. After first aid treatment (see FIRST AID section) the following medical treatment may be provided:

If the individual wears contact lenses, it is usually best to remove the lenses before additional eye irrigation.

For minor exposures with very dilute HF, the following treatment has been successful:

Mix 10 ml of 10% calcium gluconate with 100 ml of normal saline to give approximately a 1% calcium gluconate solution. With a syringe, irrigate the eye intermittently for a period of 15 to 30 minutes or until relief of pain occurs.

With more serious HF eye burns, good results have been reported with the following procedure:

Mix 50 ml of 10% calcium gluconate with 500 ml of normal saline to give approximately a 1% calcium gluconate solution. After administering local anesthetic eye drops, use an eye clamp and IV infusion set or a two pronged nasal oxygen cannula to instill the solution over a period of one to two hours. More prolonged use of the solution could possibly damage the cornea. Consultation with an ophthalmologist to consider the use of steroids, antibiotics, cycloplegics or additional treatment is recommended.

Notes:

Various approaches to the treatment of ocular exposures to HF have been recommended. Some authorities recommend the use of Lactated Ringer's solution rather than normal saline for eye irrigation. (50)

Previous recommendations have included the use of a Morgan Lens. A Morgan Lens may limit the delivery of solution to the cornea and sclera, while the methods discussed above allow the solution to contact the lids and surrounding tissues as well. A newer device, the Eye Irrigator[™], may be useful in delivering the irrigating solution to an injured eye. All therapies must be based on the individual case and on the experience and skills of the physician.

Inhalation Injuries

Patients with inhalation exposures should also be observed for signs of systemic absorption and fluoride toxicity.

Exposure to hydrofluoric acid fumes can cause acute respiratory irritation, bronchospasm, and/or **pulmonary edema**. Medical personnel should also be alert to the possibility of development of pulmonary edema when extensive burns of the face, neck or chest have occurred. Intubation should be avoided, if possible.

The victim should be removed from exposure and administered 100% oxygen immediately. The use of 2.5% aqueous calcium gluconate given by nebulizer with 100% oxygen, or with intermittent positive pressure, has been recommended. Theoretically, this should reduce toxicity and damage from the fluoride ion and should be seriously considered in cases of inhalation exposure.

Burns of the oral mucosa or upper airway may cause severe swelling and necessitate a tracheostomy. It is, therefore, recommended that all such patients be admitted to a hospital for observation.

Because inhalation of HF may be associated with significant bronchospasm, inhaled, oral or parenteral bronchodilators should be administered as necessary. Even in the absence of symptoms, the prophylactic administration of inhalational steroids (e.g. beclomethasone dipropionate) may be indicated. (21) Pulmonary function testing may be helpful in assessing the degree and progress of pulmonary injury. Specific measures may be needed to treat pulmonary edema. High doses of parenteral steroids may be needed along with the administration of appropriate diuretics. Caution should be taken not to administer excessive fluid. Hemoconcentration may require treatment by phlebotomy. The management of pulmonary edema may result in renal failure due to reduced fluid volume, and this may be another indication for hemodialysis.

If it is necessary to relieve anxiety, use general measures and do not use sedatives which could cause central nervous system depression or hypoventilation. Although right heart failure is uncommon in chemically-induced pulmonary edema, monitoring of pulmonary pressure, arterial pressure, and central venous pressure may be indicated.

Secondary infections must be treated. It is preferable to start antibiotics at the first signs of infection, such as fever or tachycardia. Periodic blood cultures may be advisable. Prophylactic use of antibiotics is not advised.

Ingestion Injuries

After first aid is completed (drinking several glasses of water followed by two glasses of milk or two ounces of milk of magnesia, Mylanta[®], or other calcium or magnesium containing antacids), the stomach may be lavaged with a solution of a calcium containing antacid. The Levin tube must be passed with care to prevent perforation. Treatment for the corrosive effects is the same as for ingestion of other strong acids. <u>Systemic toxicity is very likely to occur and may require aggressive treatment</u>.

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APPENDIX FIRST AID AND MEDICAL SUPPLIES

The following supplies should be maintained in a dispensary or first aid station near hydrofluoric acid handling and storage areas:

- 1. Benzalkonium chloride (Zephiran[®]) solution*
 - a. For soaks and compresses, 3 to 4 gallons of 0.13% water solution (1:750) of benzalkonium chloride (Zephiran®). It is recommended that the 1:750 (0.13%) solution be purchased. It is available as a non-prescription drug in gallon containers. The solution should be obtained in advance. It should replaced before the expiration date on the label. It is recommended that it be stored in properly labeled light-resistant containers.

Benzalkonium chloride (Zephiran[®]) is also available as a 17% solution. If this concentrate is used to make a 0.13% (1:750) solution, the dilution should be performed by a qualified individual, such as a registered pharmacist. The shelf life of the diluted solution is uncertain, and it should be replaced annually.

Benzalkonium chloride (Zephiran[®]) should be available as a non-prescription drug through most local pharmacies. The local pharmacies obtain it from pharmaceutical wholesale distributors such as McKesson Pharmaceuticals, Cardinal Health Inc., or the local pharmaceutical wholesaler. It can also be obtained from:

> Towne Pharmacy 103 Ridgedale Ave. Cedar Knolls, NJ 07927 Tel: 973-538-6787 Fax: 973-538-6791

In addition to benzalkonium chloride (Zephiran[®]), benzethonium chloride (Hyamine 1622[®]) has also been used successfully to treat HF burns. Because of its availability as a non-prescription drug, benzalkonium chloride (Zephiran[®]) is recommended.

- b. Ice cubes (not crushed or shaved ice).
- c. Assorted basins (for immersing burned areas in benzalkonium chloride (Zephiran®) solution).

- d. Towels (for use as wet compresses).
- 2. Calcium gluconate gel, 2.5%

Calcium gluconate gel is available commercially from:

Pharmascience Inc. 6111 Royalmount Ave. Montreal, Quebec H4P 2T4 Canada Telephone: 800.207.4477 514.340.9735 Fax 514.340.9290 www:pharmascience.com

It may also be made by mixing one ampule of 10% calcium gluconate solution for each ounce of lubricating jelly (e.g., K-Y[®] Brand Lubricating Jelly) using 40 cc per 4 ounce tube. Although this makes a somewhat "soupy" mixture, it has the advantage that the ingredients may be readily available. In addition, the ingredients may be stored separately until needed, and shelf life is less of a concern.

Calcium gluconate gel (2.5% calcium gluconate in a water soluble base) may also be formulated by a pharmacist by dissolving 3.2 grams of calcium gluconate USP in 5 cc of sterile water, and then mixing with 120 cc (4 oz. tube) of K-Y[®] Jelly or other water soluble lubricant (2.5 grams per 100 cc lubricant). The shelf life is uncertain and replacement every six months is recommended.

- 3. <u>Aqueous calcium gluconate, 10% USP, 10 cc</u> <u>ampules</u> (4.5 mEq calcium or 93 mg elemental calcium per 10 cc)
 - a. To make calcium gluconate gel, or
 - b. To mix with sterile saline for eye irrigation (5 ampules 10% calcium gluconate per 500 cc sterile normal saline for a 1% solution), or
 - c. To mix with sterile saline for administration with oxygen by nebulization (10 cc 10% calcium gluconate in 30 cc sterile saline for a 2.5% solution), or
 - d. To be administered by a physician. When injected subcutaneously, 10% calcium gluconate must be diluted half and half with normal saline to produce a 5% solution.



4. Sterile 0.9% saline

- a. Vials, (e.g. 10 cc, 30 cc, or 50 cc) to dilute 10% calcium gluconate to 5% for injection, or to 2.5% for nebulization.
- b. 500 cc IV to dilute 10% calcium gluconate to 1% for eye irrigation.
- 5. <u>0.5% tetracaine hydrochloride</u> solution to counteract blepharospasm and facilitate eye irrigation.
- 6. Medical oxygen.
- 7. <u>Nebulizer</u>, to administer 2.5% calcium gluconate with oxygen.
- 8. <u>Beta adrenergic bronchodilators and steroids</u> for inhalation.
- 9. <u>Surgical gloves</u>.

10.Syringes and needles (27-30 gauge).

11. Eye Irrigator[™] Ocular Irrigating System

American Health and Safety 6250 Nesbitt Road P. O. Box 46340 Madison, WI 53744-6340 Tel: 800.522.7554 Fax: 800.326.3245

The FIRST AID AND MEDICAL TREATMENTS AND SUPPLIES recommended in this brochure are based on information reported in the medical literature and the personal experience of physicians with Honeywell. It should be noted that there are no medications in the U.S. for which the specific indication is the treatment of hydrofluoric acid burns. The physician has the dilemma of using prescription drugs in a non-approved manner, or of using substances which are not approved drugs but which have been proven effective for medical treatment. Given the choice between recommending effective treatment, or recommending the use of only drugs which are approved, we have chosen to recommend the effective treatment.

Benzalkonium chloride (Zephiran[®]) is available in the U.S. as a non-prescription drug. It is a surface active agent sold for use as a disinfectant. It is available in a 1:750 aqueous solution, a 17% concentrate, and a tinted tincture. The concentrated 17% solution must be diluted. The tinted tincture is not recommended to treat HF exposures. Benzethonium chloride (Hyamine[®] 1622) has been used in veterinary medicine as an antiseptic for wounds and infections, but it is not available as a drug. Care should be taken that Hyamine[®] 1622 is used, not hyamine with other numeric or alphanumeric modifiers.

CALCIUM GLUCONATE INJECTION, USP (one gram in 10 ml, 10% solution) is labeled for intravenous use only. Experience has shown that when diluted to 5% with normal saline, and used as described in this brochure, it is a safe and effective treatment for HF skin exposure. When diluted to 2.5% and used as described, it is safe for nebulization and inhalation, and when diluted to 1.0% and used as described, it is safe for eye irrigation.

Notes

Caltrate[®] is a Registered Trademark of Lederle Consumer Health, Madison, NJ 07940

Garamycin[®] is a Registered Trademark of Schering Corporation, Kenilworth, NJ 07033

Hyamine $^{\rm *}$ 1622 is a Registered Trademark of Lonza, Inc., Fairlawn, NJ 07410

K-Y[®] Brand Lubricating Jelly is a Registered Trademark of Johnson & Johnson Products, Inc., Skillman, NJ 08558

Maalox[®] is a Registered Trademark of Novartis Consumer Health, Woodbridge, NJ 07095

Mylanta[®] is a Registered Trademark of Johnson & Johnson - Merck, Fort Washington, PA 19034

Silvadene[®] is a Registered Trademark of Hoechst Marion Roussel, Kansas City, MO 64134

Tums[®] is a Registered Trademark of SmithKline Beecham Consumer Healthcare, L.P., Pittsburgh, PA 15230

Zephiran[®] is a Registered Trademark of Sanofi Pharmaceuticals, New York, NY 10016



INDEX

ACGIH	4
Acute toxicity	3
Benzalkonium chloride	
see Zephiran	
Benzethonium chloride	
see Hyamine	
Blisters	7
Brochodilators	12
Brochospasm	12
Calcium acetate	10
Calcium gluconate gel	
first aid	5,6
medical treatment	8
mixing	14
ordering	14
Calcium gluconate injections	8, 14, F
Calcium gluconate solution	
1%, for eyes	6, 11, 14
2.5%, by nebulizer	6, 14
5%, subcutaneous	14
10% ampules	14
compresses, irrigation	9
intra-arterial	9
intravenous	9, 10, 11
Calcium, serum	10, 11
Cancer	5
CAS number	2
Chronic toxicity	4
Concentrated HF	2
Consumer products	2
Contact lenses	6
Debridement	7
Dilute HF	2, 3
Dimethyl sulfoxide (DMSO)	10
DMSO	10
EKG monitoring	10
Emergency Contact	F
ERPG	4
Excision, primary	11
Eye, first aid	6
Eye Irrigator	6, 15
Eye, medical treatment	11
Eye toxicity	4
Face	8
Fingernails	9
anesthesia	9
burr holes	9
Ca gluconate gel	9
Ca gluconate injections	9
removal	9
Fingers	9
First aid	
calcium gluconate gel	5,6
eyes	6

ingestion	7
inhalation	6
skin contact	5
Zephiran Soaks	5,6
Fluorides in urine	5
Fluorosis, skeletal	4
Frostbite	7
Fumes of HF	3
Hemodialysis	3, 11
Hexafluorine	10
Hyamine	14
Hyperkalemia	11
Hypocalcemia	3, 11
Hypomagnesemia	11
IDLH	4
Immersion	7
Ingestion	4
first aid	7
medical treatment	12
Inhalation	
first aid	6
medical treatment	11
toxicity	4
Intra-arterial infusion Ca solution	-
Intravenous infusion Ca solution	
Lactated Ringer's eye	110,11
Large burns	7
Magnesium sulfate	11
Magnesium, serum	11
Medical treatment	7
blisters	7
	7 7, F
calcium gluconate gel calcium gluconate	7, 1
	0 E
5% injections concentrated HF	8, F 9 F
	8, F 7
debridement	7 7 F
dilute HF	7, F
eye	11, F
Ingestion	12, F
Inhalation	12, F
pain relief	7
Quick Reference	F
skin burns Zambinan Saaba	7, F 7 F
Zephiran Soaks	7, F
Nails	9
Nebulizer	7, 12, 14
NIOSH	4
OSHA	4
Oxygen	12
Pain relief	3, 7, 8
Permissible Exposure Limit	4 *
Preventative Measures	
Primary excision	11
Proparacaine	6
Pulmonary edema	4, 12

Quaternary ammonium	-
compounds	7
Quick reference to treatment	F
References	12
Saline solution	15
Sensitive areas,,no Zephiran	7
Serum calcium determinations	11
Serum magnesium	11
Skin contact	2
first aid	5
medical treatment	7
Skin grafting	10
Systemic toxicity	2, 3
absorption and	
metabolic effects	11
hemodialysis	11
treatment	11
Tetany	11
Tetracaine HCl	6, 15
m · ·	
Toxicity	
acute	2
5	2 4
acute chronic	
acute	4
acute chronic eye	4 4
acute chronic eye inhalation	4 4 4 2
acute chronic eye inhalation skin contact	4 4 4
acute chronic eye inhalation skin contact systemic	4 4 2 2, 3
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF	4 4 2 2, 3 5
acute chronic eye inhalation skin contact systemic Urine fluorides	4 4 2 2, 3 5 2
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran	4 4 2 2, 3 5 2 15
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran 1:750 solution	4 4 2 2, 3 5 2 15 14
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran 1:750 solution 17% solution	4 4 2 2, 3 5 2 15 14 14
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran 1:750 solution 17% solution about	4 4 2 2, 3 5 2 15 14 14 14 14 14
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran 1:750 solution 17% solution about dilution	4 4 2 2, 3 5 2 15 14 14 14
acute chronic eye inhalation skin contact systemic Urine fluorides Uses of HF Zephiran 1:750 solution 17% solution about dilution first aid	4 4 2, 3 5 2 15 14 14 14 14 14 5, 6

F Foldout



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We will be one of the world's premier companies, distinctive and successful in everything we do.
We will become a Total Quality Company by continuously improving all our work processes to satisfy our internal and external customers.
Customers Our first priority is to satisfy customers
Integrity We are committed to the highest level of ethical conduct wherever we operate. We obey all laws, produce safe products, protect the environment, practice equal employment, and are socially responsible.
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Honeywell Fluorine Products Product Stewardship Policy

Honeywell, in its continuing quest to be one of the world's premier companies, subscribes to the Chemical Manufacturer's Association's "Responsible Care" Program. Product Stewardship is an integral part of that program and enables us to make health, safety and environmental protection a key part of all product related activities.

At Honeywell Fluorine Products, Product Stewardship encompasses all aspects of the product life cycle including design, manufacture, marketing, distribution, use recycling and disposal of our products. It involves working closely with our worldwide customers, suppliers, employees, distributors, wholesalers, tollers and contractors to meet these goals. Product Stewardship is not a one time effort designed only to comply with regulations, but a continuous, long-term process that is applied throughout all of our business operations.

Fluorine Products is committed to:

- Giving high priority to health, safety and environmental considerations in our business planning encompassing proper selection of raw materials through Materials Management, Customer Linked Commercialization for all new products, processes and waste materials and our Customer Linked Manufacturing endeavors to improve existing operation processes.
- Guiding customers on the safe use, transportation and disposal of our products.
- Reporting promptly to customers, employees, government officials and the public of any new information on product related issues.
- Providing technical assistance on various uses and applications for our products.
- Sharing pertinent information and experiences with others who produce, handle, use, transport or dispose of our products.

Product Stewardship is a part of everyone's job. By each employee supporting and implementing the Product Stewardship practices outlined above, we will move ahead of our competition, be viewed by our customers as a premier partner and create growth for Fluorine Products.

Fluorine Products



HF Products

FOR ADDITIONAL INFORMATION

For additional assistance, including technical information covering all aspects of hydrofluoric acid, safe handling, use and disposal write:

Honeywell Industrial Fluorines P.O. Box 1053 101 Columbia Road Morristown, NJ 07962-1053

FAX: 973-455-6141

In the event of an emergency with this product, call the 24-hour Honeywell emergency telephone number: **800 707-4555 or 602-365-4980**

To place an order, obtain prices or product availablity information, call toll free:

- From within the continental United States 800-522-8001 973-455-6300
- From any location in Canada 800-553-9749

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