

# MATERIAL SAFETY DATA SHEET



Consumer Care Division  
 Bayer Corporation, 1884 Miles Avenue, Elkhart, IN 46514  
 219 264 -8400

For accidental ingestion, or medical  
 emergency, call 1-800-800-4793

HMIS Code		
Health	1	Fire 0
Reactivity	0	Pers. Prot. 0

## SECTION 1 - IDENTITY

Common Name: (used on label)

(Trade Name & Synonyms) Original ALKA-SELTZER Antacid & Pain Reliever

Product Code: 4028, 4002  
 4024, 4022, 4019  
 4017, 4012, 4011

Chemical Name:  
 Active Ingredients: Aspirin 325 mg, Sodium Bicarbonate  
 1916 mg, Citric Acid 1000 mg (See Section 8).

Chemical Family  
 None

Formula  
 Effervescent mixture.

CAS No.  
 None

## SECTION 2 - HAZARDOUS INGREDIENTS

Hazardous Components - (chemical & common names)

Aspirin

Hazard  
 Ingestion (oral LD50 in rats = 5g or greater/kg).

TLV Units

For additional information see Section 4.

## SECTION 3 - PHYSICAL & CHEMICAL CHARACTERISTICS (Fire & Explosion Data)

Boiling Point

Not applicable.

Specific Gravity (H<sub>2</sub>O = 1)

Not applicable.

Vapor Pressure (mm hg)

Not applicable.

Percent Volatile by Volume (%)

Not applicable.

Vapor Density (Air = 1)

Not applicable.

Evaporation Rate (= 1)

Not applicable.

Solubility in Water

Not applicable.

Reactivity in Water/Air

Effervesces in contact with water, producing CO<sub>2</sub>.

Appearance and Odor

Effervescent white tablet.

Flash Point

Not applicable.

Flammable Limits in Air % by Volume

Lower Upper

Not applicable.

Auto-ignition Temperature

Not applicable.

Extinguisher Media

Water

Special Fire Fighting Procedures

None

Unusual Fire and Explosion Hazards

None

## SECTION 4 - HEALTH HAZARDS

OSHA Permissible Exposure Limit

Not established.

ACGIH Threshold Limit Value

Not established.

Other Exposure Limit Used

FDA - Aspirin 4000 mg/24 hrs for 10 days in a 70 kg person; acute toxicity may follow ingestions of 150 mg/kg.

Signs and Symptoms - Acute Overexposure

Nausea, vomiting, ringing in the ears, fever, coma, respiratory alkalosis, metabolic acidosis, convulsions (see Section 8).

Signs and Symptoms - Chronic Overexposure

Ringing in the ears, diminished hearing, confusion, agitation, lethargy, pulmonary edema, cardiovascular collapse (see Section 8).

Medical Conditions Generally Aggravated by Exposure

Ulcers (aspirin), asthma (aspirin), predisposition to bleeding & decreased platelet function (aspirin), congestive heart failure (sodium), hypertension (sodium).

Primary Route(s) of Exposure

Oral

Emergency and First Aid Procedures

Contact your regional poison control center or physician immediately. Additional information may be obtained by contacting Bayer Corporation at 1-800-800-4793

Hygienic Practices

Normal clinical.

Chemical Listed as Carcinogen or Potential Carcinogen

None

Common Name: Original ALKA-SELTZER Antacid & Pain Reliever

### SECTION 5 - PHYSICAL HAZARDS

Stability: Unstable  Conditions to avoid  
Stable

Incompatibility  
(Materials to Avoid) None

Hazardous  
Decomposition Products None

Hazardous Polymerization May Occur  Conditions to Avoid  
Will Not Occur

### SECTION 6 - SPECIAL PROTECTION INFORMATION

Respiratory Protection  
None for normal use.

Ventilation None for normal use. Work Practices  
Not applicable.

Protective Gloves None for normal use. Eye Protection  
None for normal use.

Other Protective  
Clothing or Equipment None for normal use.

### SECTION 7 - SPECIAL PRECAUTIONS AND SPILL/LEAK PROCEDURES

Precautions to be Taken  
in Handling and Storage  
None

Steps to be Taken in Case  
Material is Released or Spilled  
Contain for disposal.

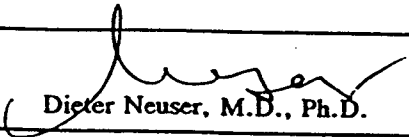
Waste Disposal  
Methods  
Dispose as normal solid waste.

### SECTION 8 - ADDITIONAL INFORMATION

Alka-Seltzer in water contains principally the antacid sodium citrate and the analgesic sodium acetylsalicylate.

Classic signs and symptoms of acute and chronic salicylate overdose are listed in Section 4. However, studies indicate that plasma salicylate will not reach toxic levels following ingestion of highly buffered aspirin in solution, regardless of the dose taken. This is due to alkalinization of the urine by the citrate buffer which increases the excretion of salicylate. Therefore, the classic signs and symptoms of salicylate toxicity listed in Section 4 may not be seen with overdose of this product.

Signature of Person  
Responsible for Preparation

  
Dieter Neuser, M.D., Ph.D.

Date  
Prepared March 1991

Telephone No.  
219/264-8142

Date  
Revised November 1991

The opinions expressed herein are those of qualified experts within Bayer Corporation. We believe that the information contained herein is current as of the date of this Material Safety Data Sheet. Since the use of this information and these opinions and the conditions of the product are not within the control of Bayer Corporation, it is the user's obligation to determine the conditions of safe use of the product.



MATERIAL SAFETY DATA SHEET

BAYER CORPORATION
CONSUMER CARE DIVISION
36 Columbia Road
Morristown, NJ 07962-1910

TRANSPORTATION EMERGENCY
CALL CHEMTREC: 800-424-9300
INTERNATIONAL: 703-527-3887

NON-TRANSPORTATION
BAYER EMERGENCY PHONE...: (800) 743-5423
BAYER INFORMATION PHONE.: (800) 743-5423

1. CHEMICAL PRODUCT IDENTIFICATION:

PRODUCT NAME.....: Original Bayer (R) Aspirin
PRODUCT CODE.....: 114,110,111,117,112,120,113,181
CHEMICAL FAMILY.....: Analgesic

2. COMPOSITION/INFORMATION ON INGREDIENTS:

INGREDIENT NAME /CAS NUMBER EXPOSURE LIMITS CONCENTRATION (%)

\*\*\*\*\* HAZARDOUS INGREDIENTS \*\*\*\*\*

This pharmaceutical product, available without a prescription, is for human use. These materials are not subject to the Federal OSHA Hazard Communication Standard 29 CFR 1910.1200.

\*\*\*\*\* OTHER INGREDIENTS \*\*\*\*\*

Aspirin
50-78-2 OSHA : 5.00 mg/m3 TWA Greater than 25 %
ACGIH: 5.00 mg/m3 TWA
\* See Section 3 for potential health effects.

3. HAZARDS IDENTIFICATION:

```

*****
*
*                               EMERGENCY OVERVIEW                               *
*
* Color: White; Form: Solid; Tablets/Caplets; Odor: Slight *
* acidic; Product poses little or no hazard if spilled and no *
* unusual hazard if involved in a fire; See Potential Health *
* Effects if the recommended dosage is exceeded. *
*****

```

POTENTIAL HEALTH EFFECTS:

ROUTE(S) OF ENTRY.....: Appropriate route of entry: oral

HUMAN EFFECTS AND SYMPTOMS OF OVEREXPOSURE:

NOTE: This is a pharmaceutical material available without a prescription - use only as directed. See product packaging for further information concerning adverse effects and drug interaction precautions.

ACUTE EFFECTS OF EXPOSURE.....: Acute overexposure to this product may cause rapid or deep breathing, confusion, agitation, nausea, vomiting, diminished hearing, ringing in the ears, hemorrhage, acid/base imbalances, coma, seizures, hypotension or cardiac arrhythmias. Allergic reactions are possible with symptoms of reddening, itching, rash, swelling of the face, throat or tongue, and breathing problems. If swelling of the face, throat, tongue or breathing problems occur, seek medical attention immediately.

CHRONIC EFFECTS OF EXPOSURE....: Chronic overexposure to this product may cause effects described under acute exposure. In addition, liver and/or kidney dysfunction may result.

CARCINOGENICITY.....: The components of this product are not listed by NTP, IARC or regulated as a carcinogen by OSHA.

MEDICAL CONDITIONS

AGGRAVATED BY EXPOSURE.....: Persons with asthma, preexisting hypersensitivity to the components of this product or other pain releivers, young children, the elderly, and pregnant women may be more susceptible to the effects of this product. In addition, children and teenagers with chickenpox or flu symptoms, persons who are at risk for hemorrhage, those with a history of gastrointestinal ulcers or bleeding, who have impaired renal function, who are taking other prescription medications, or who consume more than 3 alcohol containing drinks per day may also be more susceptible to the effects of this product.

EXPOSURE LIMITS.....: FDA -- Aspirin 4000 mg/24 hrs. for 10 days in a 70 kg person; acute toxicity may follow ingestions of 150 mg/kg.

-----  
4. FIRST AID MEASURES:  
-----

FIRST AID FOR EYES.....: In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Call a physician.  
FIRST AID FOR SKIN.....: Flush skin with plenty of soap and water. Contact a physician if irritation develops.  
FIRST AID FOR INHALATION: Not applicable.  
FIRST AID FOR INGESTION.: In case of overdose, contact your regional poison control center or physician immediately. For additional information, contact Bayer Corporation at 1-800-800-4793.

-----  
5. FIRE FIGHTING MEASURES:  
-----

FLASH POINT.....: Not Applicable  
AUTO-IGNITION TEMPERATURE.....: Not Applicable  
EXTINGUISHING MEDIA.....: Water  
SPECIAL FIRE FIGHTING PROCEDURES: Firefighters should be equipped with self-contained breathing apparatus to protect against potentially toxic and irritating fumes.

-----  
6. ACCIDENTAL RELEASE MEASURES:  
-----

SPILL OR LEAK PROCEDURES.....: Spills should be swept up and placed in appropriate containers for disposal. Avoid creating dusty conditions.

-----  
7. HANDLING AND STORAGE:  
-----

STORAGE TEMPERATURE (MIN/MAX): Room temperature.  
SHELF LIFE.....: Do not use after expiration date.  
SPECIAL SENSITIVITY.....: None known.  
HANDLING/STORAGE PRECAUTIONS: Keep this and all drugs out of the reach of children. Avoid contact with eyes and skin. Wash thoroughly after handling. Store in a dry place away from excessive heat. Reseal containers immediately after use. Use normal precautions for storage of a drug.

-----  
8. PERSONAL PROTECTION;  
-----

EYE PROTECTION REQUIREMENTS.....: None for normal use.  
SKIN PROTECTION REQUIREMENTS.....: None for normal use.  
VENTILATION REQUIREMENTS.....: Under normal conditions of use, special  
ventilation is not required.  
RESPIRATOR REQUIREMENTS.....: Under normal conditions of use,  
respiratory protection is not required.  
WORK PRACTICES.....: Normal clinical practice. Use good  
personal hygiene - wash hands and exposed skin thoroughly with soap and  
water after each use.  
ADDITIONAL PROTECTIVE MEASURES.....: Employers shall provide handwashing  
facilities which are readily accessible to employees. Educate and train  
employees in the safe use and handling of this product.

-----  
9. PHYSICAL AND CHEMICAL PROPERTIES;  
-----

PHYSICAL FORM.....: Solid  
APPEARANCE.....: Tablets/Caplets  
COLOR.....: White  
ODOR.....: Slight acidic  
PH.....: Not Established  
BOILING POINT.....: Not Applicable  
MELTING/FREEZING POINT.....: Not Applicable  
SOLUBILITY IN WATER.....: Soluble  
SPECIFIC GRAVITY.....: Not Established  
BULK DENSITY.....: Not Established  
VAPOR PRESSURE.....: Not Applicable

-----  
10. STABILITY AND REACTIVITY;  
-----

STABILITY.....: This is a stable material.  
HAZARDOUS POLYMERIZATION...: Will not occur.  
INCOMPATIBILITIES.....: See product packaging for drug interaction.  
INSTABILITY CONDITIONS.....: None known.  
DECOMPOSITION PRODUCTS.....: Not Applicable.

-----  
11. TOXICOLOGICAL INFORMATION:  
-----

TOXICITY DATA FOR: Aspirin

ACUTE TOXICITY

ORAL LD50.....: Greater than 1,500 mg/kg (rat)

-----  
12. ECOLOGICAL INFORMATION:  
-----

NO ECOLOGICAL INFORMATION AVAILABLE

-----  
13. DISPOSAL CONSIDERATIONS  
-----

WASTE DISPOSAL METHOD.....: Waste disposal should be in accordance with existing federal, state and local environmental control laws.

-----  
14. TRANSPORTATION INFORMATION:  
-----

TECHNICAL SHIPPING NAME.....: Analgesic

PRODUCT LABEL.....: Original Bayer (R) Aspirin

DOT (DOMESTIC SURFACE)  
-----

HAZARD CLASS OR DIVISION .....: Non-Regulated

IMO / IMDG CODE (OCEAN)  
-----

HAZARD CLASS DIVISION NUMBER...: Non-Regulated

ICAO / IATA (AIR)  
-----

HAZARD CLASS DIVISION NUMBER...: Non-Regulated

-----  
15. REGULATORY INFORMATION:  
-----

OSHA STATUS.....: This material is not subject to the OSHA Hazard Communication Standard as noted in 29 CFR 1910.1200(b) (6) (vii).

Product Code: 114,110,111,117,112,120,113,181  
Approval date: 04/10/2002

MSDS Page 5  
Continued on next page

15. REGULATORY INFORMATION (Continued)

TSCA STATUS.....: This product is exempt from TSCA Regulation under Section 3 (2)(B)(vi) when used for pharmaceutical application.

CERCLA REPORTABLE QUANTITY...: None

SARA TITLE III:

SECTION 302 EXTREMELY HAZARDOUS SUBSTANCES...: None

SECTION 311/312 HAZARD CATEGORIES.....: Exempt from SARA Section 311/312

SECTION 313 TOXIC CHEMICALS.....: None

RCRA STATUS.....: If discarded in its purchased form, this product would not be a hazardous waste either by listing or by characteristic. However, under RCRA, it is the responsibility of the product user to determine at the time of disposal, whether a material containing the product or derived from the product should be classified as a hazardous waste. (40 CFR 261.20-24)

The following chemicals are specifically listed by individual states; other product specific health and safety data in other sections of the MSDS may also be applicable for state requirements. For details on your regulatory requirements you should contact the appropriate agency in your state.

COMPONENT NAME /CAS NUMBER	CONCENTRATION	STATE CODE
Aspirin 50-78-2	Greater than 25 %	PA1, CA , MA, NJ1
Corn starch 9005-25-8	10-25 %	PA1, MA, NJ4

CA = California Proposition 65  
 MA = Massachusetts Hazardous Substance List  
 NJ1 = New Jersey Hazardous Substance List  
 NJ4 = New Jersey Other - included in 5 predominant ingredients > 1%  
 PA1 = Pennsylvania Hazardous Substance List

ADDITIONAL INFORMATION: ACTIVE INGREDIENT (per tablet) Aspirin, 325 mg  
 INACTIVE INGREDIENTS: Hydroxypropyl Methylcellulose, Starch, Triacetin.

16. OTHER INFORMATION:

HMIS RATINGS: Health 1 Flammability 0 Reactivity 0

Product Code: 114,110,111,117,112,120,113,181  
 Approval date: 04/10/2002

MSDS Page 6  
 Continued on next page



16. OTHER INFORMATION (Continued)

-----  
0=Minimal 1=Slight 2=Moderate 3=Serious 4=Severe

Bayer's method of hazard communication is comprised of Product Labels and Material Safety Data Sheets. HMIS ratings are provided by Bayer as a customer service.

REASON FOR ISSUE.....: Name Change;was Genuine Bayer (R) Aspirin  
PREPARED BY.....: S. Van Volkenburg  
APPROVED BY.....: Llew C. Williams  
APPROVAL DATE.....: 04/10/2002  
SUPERSEDES DATE.....: 06/12/1998  
MSDS NUMBER.....: 34102

-----  
This information is furnished without warranty, expressed or implied, except that it is accurate to the best knowledge of Bayer Corporation. The data on this sheet relates only to the specific material designated herein. Bayer Corporation assumes no legal responsibility for use or reliance upon these data.  
-----

Product Code: 114,110,111,117,112,120,113,181  
Approval date: 04/10/2002

MSDS Page 7  
Last page



MATERIAL SAFETY DATA SHEET

BAYER CORPORATION
CONSUMER CARE DIVISION
36 Columbia Road
Morristown, NJ 07962-1910

TRANSPORTATION EMERGENCY CALL CHEMTREC: 800-424-9300
INTERNATIONAL: 703-527-3887
NON-TRANSPORTATION BAYER EMERGENCY PHONE...: (800) 743-5423
BAYER INFORMATION PHONE.: (800) 743-5423

1. CHEMICAL PRODUCT IDENTIFICATION:

PRODUCT NAME.....: Aleve
PRODUCT CODE.....: 501; 503; 505; 004
CHEMICAL FAMILY.....: Analgesic
NDC NUMBER.....: 0280-6010-24; 0280-6010-50; 0280-6010-01; 0280-6010-15

2. COMPOSITION/INFORMATION ON INGREDIENTS:

INGREDIENT NAME /CAS NUMBER EXPOSURE LIMITS CONCENTRATION (%)

\*\*\*\*\* HAZARDOUS INGREDIENTS \*\*\*\*\*

This pharmaceutical product, available without a prescription, is for human use. These materials are not subject to the Federal OSHA Hazard Communication Standard 29 CFR 1910.1200. This Material Safety Data Sheet is not intended for industrial exposures.

3. HAZARDS IDENTIFICATION:

\*\*\*\*\*
\* EMERGENCY OVERVIEW \*
\* Color: Light blue; Form: Solid; Caplet; Odor: Odorless; \*
\* Product poses little or no hazard if spilled and no unusual \*
\* hazard if involved in a fire; See Potential Health Effects \*
\* if the recommended dosage is exceeded. \*
\*\*\*\*\*

POTENTIAL HEALTH EFFECTS:

Product Code: 501; 503; 505; 004
Approval date: 06/30/1997

MSDS Page 1
Continued on next page

## 3. HAZARDS IDENTIFICATION (Continued)

ROUTE(S) OF ENTRY.....: Appropriate route of entry: oral

## HUMAN EFFECTS AND SYMPTOMS OF OVEREXPOSURE:

NOTE: This is a pharmaceutical material available without a prescription - use only as directed. See product packaging for further information concerning adverse effects and drug interaction precautions.

ACUTE EFFECTS OF EXPOSURE.....: Overdose may cause nausea, vomiting, confusion, ringing in the ears, irregular heartbeat, headache, drowsiness, and blood pressure effects.

CHRONIC EFFECTS OF EXPOSURE...: Chronic overexposure to this product may cause effects as described under acute exposure. An allergic reaction is possible with symptoms of reddening, rash, and itching.

CARCINOGENICITY.....: The components of this product are not listed by NTP, IARC or regulated as a carcinogen by OSHA.

## MEDICAL CONDITIONS

AGGRAVATED BY EXPOSURE.....: Persons with preexisting hypersensitivity to the components of this product and pregnant women may be more susceptible to the effects of this product. Persons consuming alcohol may also be more susceptible to the effects of this product.

## 4. FIRST AID MEASURES:

FIRST AID FOR EYES.....: In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Call a physician.

FIRST AID FOR SKIN.....: Flush skin with plenty of soap and water. Contact a physician if irritation develops.

FIRST AID FOR INHALATION: Not applicable.

FIRST AID FOR INGESTION.: In case of overdose, contact your regional poison control center or physician immediately. For additional information, contact Bayer Corporation at 1-800-800-4793.

## 5. FIRE FIGHTING MEASURES:

FLASH POINT.....: Not Applicable

EXTINGUISHING MEDIA.....: All extinguishing media are suitable.

SPECIAL FIRE FIGHTING PROCEDURES: Firefighters should be equipped with self-contained breathing apparatus to protect against potentially toxic and irritating fumes.

-----  
6. ACCIDENTAL RELEASE MEASURES:  
-----

SPILL OR LEAK PROCEDURES.....: Spills should be swept up and placed in appropriate containers for disposal. Avoid creating dusty conditions.

-----  
7. HANDLING AND STORAGE:  
-----

STORAGE TEMPERATURE (MIN/MAX): Room temperature.  
SHELF LIFE.....: Do not use after expiration date.  
SPECIAL SENSITIVITY.....: Avoid direct sunlight.  
HANDLING/STORAGE PRECAUTIONS: Keep this and all drugs out of the reach of children. Avoid contact with eyes and skin. Wash thoroughly after handling. Store in a dry place away from excessive heat. Reseal containers immediately after use. Use normal precautions for storage of a drug.

-----  
8. PERSONAL PROTECTION:  
-----

EYE PROTECTION REQUIREMENTS.....: None for normal use.  
SKIN PROTECTION REQUIREMENTS.....: None for normal use.  
VENTILATION REQUIREMENTS.....: Under normal conditions of use, special ventilation is not required.  
RESPIRATOR REQUIREMENTS.....: Under normal conditions of use, respiratory protection is not required.  
WORK PRACTICES.....: Normal clinical practice. Use good personal hygiene - wash hands and exposed skin thoroughly with soap and water after each use.  
ADDITIONAL PROTECTIVE MEASURES.....: Employers shall provide handwashing facilities which are readily accessible to employees. Educate and train employees in the safe use and handling of this product.

-----  
9. PHYSICAL AND CHEMICAL PROPERTIES:  
-----

PHYSICAL FORM.....: Solid  
APPEARANCE.....: Caplet  
COLOR.....: Light blue  
ODOR.....: Odorless  
pH.....: Not Established  
BOILING POINT.....: Not Applicable  
MELTING/FREEZING POINT.....: Not Applicable  
SOLUBILITY IN WATER.....: Soluble

## 9. PHYSICAL AND CHEMICAL PROPERTIES (Continued)

-----  
SPECIFIC GRAVITY .....: Not Established  
BULK DENSITY.....: Not Established  
VAPOR PRESSURE .....: Not Applicable  
-----

## 10. STABILITY AND REACTIVITY:

-----  
STABILITY.....: This is a stable material.  
HAZARDOUS POLYMERIZATION...: Will not occur.  
INCOMPATIBILITIES.....: See product packaging and the Physicians' Desk  
Reference (PDR) for drug interaction.  
INSTABILITY CONDITIONS.....: None known.  
DECOMPOSITION PRODUCTS.....: Not Applicable.  
-----

## 11. TOXICOLOGICAL INFORMATION:

-----  
NO ANIMAL TOXICITY INFORMATION AVAILABLE  
-----

## 12. ECOLOGICAL INFORMATION:

-----  
NO ECOLOGICAL INFORMATION AVAILABLE  
-----

## 13. DISPOSAL CONSIDERATIONS

-----  
WASTE DISPOSAL METHOD.....: Waste disposal should be in accordance with  
existing federal, state and local environmental control laws.  
-----

## 14. TRANSPORTATION INFORMATION:

-----  
TECHNICAL SHIPPING NAME.....: Analgesic  
PRODUCT LABEL.....: Aleve  
-----

DOT (DOMESTIC SURFACE)  
-----

HAZARD CLASS OR DIVISION .....: Non-Regulated  
-----

## 14. TRANSPORTATION INFORMATION (Continued)

-----  
 DOT (continued)  
 -----

IMO / IMDG CODE (OCEAN)  
 -----

HAZARD CLASS DIVISION NUMBER...: Non-Regulated

ICAO / IATA (AIR)  
 -----

HAZARD CLASS DIVISION NUMBER...: Non-Regulated

-----  
 15. REGULATORY INFORMATION:  
 -----

OSHA STATUS.....: This material is not subject to the OSHA Hazard  
 Communication Standard as noted in 29 CFR  
 1910.1200(b)(6)(vii).

TSCA STATUS.....: This product is exempt from TSCA Regulation under  
 Section 3.(2)(B)(vi) when used for pharmaceutical  
 application.

CERCLA REPORTABLE QUANTITY...: None

SARA TITLE III:

SECTION 302 EXTREMELY

HAZARDOUS SUBSTANCES...: None

SECTION 311/312

HAZARD CATEGORIES.....: Exempt from SARA Section 311/312

SECTION 313

TOXIC CHEMICALS.....: None

RCRA STATUS.....: If discarded in its purchased form, this product  
 would not be a hazardous waste either by listing  
 or by characteristic. However, under RCRA, it is  
 the responsibility of the product user to  
 determine at the time of disposal, whether a  
 material containing the product or derived from  
 the product should be classified as a hazardous  
 waste. (40 CFR 261.20-24)

ADDITIONAL INFORMATION: ACTIVE INGREDIENT (per caplet): Naproxen Sodium, 220  
 mg. INACTIVE INGREDIENTS: Magnesium Stearate, Microcrystalline cellulose,  
 Opadry, Povidone, Talc.

-----  
 16. OTHER INFORMATION:  
 -----

HMIS RATINGS:

Health	Flammability	Reactivity
1	0	0
0=Minimal	1=Slight	2=Moderate
		3=Serious
		4=Severe

Product Code: 501; 503; 505; 004  
 Approval date: 06/30/1997

MSDS Page 5  
 Continued on next page

## 16. OTHER INFORMATION (Continued)

-----  
Bayer's method of hazard communication is comprised of Product Labels and Material Safety Data Sheets. HMIS ratings are provided by Bayer as a customer service.

REASON FOR ISSUE.....: Update format  
PREPARED BY.....: R. Ruppel-Kerr  
APPROVED BY.....: Llew C. Williams  
APPROVAL DATE.....: 06/30/1997  
SUPERSEDES DATE.....: 05/04/1995  
MSDS NUMBER.....: 29404

-----  
This information is furnished without warranty, expressed or implied, except that it is accurate to the best knowledge of Bayer Corporation. The data on this sheet relates only to the specific material designated herein. Bayer Corporation assumes no legal responsibility for use or reliance upon these data.  
-----

Product Code: 501; 503; 505; 004  
Approval date: 06/30/1997

MSDS Page 6  
Last page