



**SAN DIEGO COUNTY
DEPARTMENT OF ENVIRONMENTAL HEALTH AND QUALITY
HAZARDOUS MATERIALS DIVISION**

**California Accidental Release Prevention Program (CalARP)
Risk Management Plan Development & Submittal Guidelines**

**Revised
March 2023**

It is recommended that the user read the entire contents of this document prior to proceeding in the development of their Risk Management Plan (RMP).

These guidelines are intended to assist stationary sources with preparing a Risk Management Program. They are not intended to replace the regulations found in Chapter 4.5, Division 2, Title 19 of the California Code of Regulations.

TABLE OF CONTENTS

RMP Coordination, Consultation, Review & Cost Recovery _____	5
RMP Public Document _____	6
1. Registration / Data Elements _____	6
2. Executive Summary _____	6
3. Offsite Consequences Analysis _____	7
4. Accident History _____	8
5. Prevention Program 2 (if applicable) _____	8
6. Prevention Program 3 (if applicable) _____	8
7. External Events & Seismic Assessment _____	9
8. Emergency Response Program _____	9
9. Certification _____	10
10. Updates _____	10
11. Corrections _____	10
 Guidelines for Technical Studies/Supporting Records _____	 11
I. Offsite Consequence Analysis _____	12
II. Program 2 - Prevention Program _____	19
III. Program 3 - Prevention Program _____	24
IV. External Events including Seismic Assessment _____	34
V. Emergency Response Plans _____	35
 Appendices	
Risk Management Plan Registration / Work Plan _____	A-1
Useful Web Links _____	A-2
Sample Incident Investigation _____	A-3
Sample Audit Checklist for Safety Information _____	A-4
Public Review Information _____	A-5
Record of Revision for RMP (samples) _____	A-6
Seismic Issue Summary _____	A-7
What-If/Checklist (sample) _____	A-8
Trade Secret/Confidential Information _____	A-9
Quick Reference Worst Case Scenario Requirements _____	A-10
Alternative Release Scenario Analysis _____	A-11
Fees _____	A-12
OCA Data Table (sample) _____	A-13

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

Coordination, Consultation, Review Cost Recovery

California Code of Regulations, Title 19, Division 2, Chapter 4.5, Article 3 requires the Owners and Operators of a Stationary Source to coordinate with the Certified Unified Program Agency (CUPA)/Administering Agency (AA) in the development and implementation of Risk Management Plans/Programs. The Hazardous Materials Division (HMD) is a division within the Department of Environmental Health and Quality (DEHQ). DEHQ is the CUPA in San Diego County. In San Diego County, the terms AA, HMD and CUPA are used interchangeably. Coordination and consultation early in the RMP development process is essential for proper RMP implementation.

- 19 CCR Section 2785.1 (a) The owner or operator of a stationary source shall closely coordinate with the Unified Program Agency (UPA) to ensure that appropriate technical standards are applied to the implementation of this chapter.
- 19 CCR Section 2735.5 General Requirements. Coordination. The owner or operator of a stationary source shall closely coordinate with the UPA to implement the requirements of this chapter and to determine the appropriate level of documentation required for an RMP to comply with Sections 2745.3 through 2745.9 of this chapter. This requirement shall not preclude public access to RMP information. Classified information need not be included in the RMP but shall be made available to the UPA to the extent allowable by law. Trade secrets are protected pursuant to Section 25538 of HSC.

To facilitate coordination, the DEHQ requires submission of a Risk Management Plan (RMP) Registration/Work Plan Form HM-9240. (See **Appendix A-1.**) The RMP Registration/Work Plan submission should be made prior to development of the RMP. The RMP Registration/Work Plan will be reviewed by DEHQ Specialists with the intent of establishing an initial high-level understanding of the stationary source and the covered process. The DEHQ will review the RMP Registration/Work Plan for agreement of applicability of the CalARP regulations, and the appropriate Program (i.e., 1, 2, or 3*). Review of the RMP Registration/Work Plan may result in questions about the covered process, that require responses from the Owner Operator and/or Consultant.

The CUPA uses a direct cost recovery process for specific activities related to coordination, consultation, and review related to CalARP covered processes. (See **Appendix A-12 Fees.**)

*CalARP regulations include a Program 4 Prevention Program for Refineries. Currently there are no refineries in San Diego County, nor are there plans to add any.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

The Risk Management Plan (RMP) Public Document

The Risk Management Plan Public Document should reflect a facility's overall effort in the management and prevention of risks associated with the storage, use and/or processing of regulated substances. (See **Appendix A-5** for more information on Public Review of RMP.)

The RMP Public Document is supported by the following technical studies:

- ◆ Off-Site Consequence Analysis
- ◆ Program 2 or Program 3 Prevention Program (if applicable)
- ◆ Hazard review or Process Hazard Analysis (depending on Prevention Program)
- ◆ External Events Analysis & Seismic Assessment

The RMP Public Document shall be in the form of a **single volume for all Regulated Substances** handled unless otherwise instructed by the Hazardous Materials Division (HMD).

Submitting to HMD – (regulated substances greater than state or federal thresholds)

Per section 2745.1(a) The Owner Operator shall submit, to the CUPA, a single RMP public document. HMD requests the single submittal be indexed and tabbed. HMD requests the submittal be in a single PDF, with individual tabbed sections as follows:

1. **RISK MANAGEMENT REGISTRATION/ DATA ELEMENTS (CCR 2740.1) -**
Complete and submit the CalARP Registration/Work Plan Form HM-9240. (See **Appendix A-1**.) If you are also submitting an RMP to USEPA, then you may use the DATA Elements form you've prepared for USEPA as the data elements section of the RMP Public Document for the HMD.
2. **RMP EXECUTIVE SUMMARY (CCR 2745.3) -** The RMP Executive Summary should be brief and concise, no more than four pages in length. A more detailed description is to be provided in the body of the report. Your executive summary shall include:
 - (a) The accidental release prevention and emergency policies in place at your facility.
 - (b) A description of your regulated processes and regulated substances handled. This information may be presented in a paragraph or as bullets. The information should include the following:
 - Primary activities (e.g., water treatment, power generation, metal work).
 - Use of regulated substances (e.g., chlorine used to treat drinking water, ammonia for SCR, HF for metal preparation).
 - Total quantities handled or stored in pounds (and in gallons if a liquid at ambient temperature).
 - Quantity in the largest vessel (in pounds). This is used for the offsite consequence analysis.
 - (c) The general accidental release prevention program and chemical-specific prevention steps. For example, you may state that you are in compliance with the OSHA PSM rule and the CalARP requirements. Consider highlighting general or specific steps that you believe are key to your prevention program. These steps may be either technological (e.g., backup systems) or procedural/managerial (e.g., improved maintenance or training).
 - (d) The five-year accident history. This should be a summary (e.g., we have had five accidental releases of chlorine in the past five years; the largest release was 1500 pounds. No one offsite was injured, but several houses were evacuated as a precautionary measure during the releases). Do not present this information in a table format.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

- (e) A summary of the emergency response program. State here whether you are a “responding facility” (respond to control or mitigate releases of the regulated substance) or a “non-responding facility” (do not respond in PPE to mitigate releases of the regulated substance).
- (f) Planned changes to improve safety.
- (g) Safety (Local Requirement) – List and describe all safety features (equipment (sensors, alarms, PRD, administrative, passive, and active, etc.)) that are in place to make this a safe process or are to be implemented. Include the dates of implementation. A Safety Data Sheet is required.

3. OFFSITE CONSEQUENCE ANALYSIS COMPONENT (CCR 2745.4)

The Offsite Consequence Analysis (OCA) describes the potential exposure levels for surrounding populations and environmental receptors from accidental releases of **regulated substances**. Provide the following information for each Program 1, 2, or 3 covered process:

- (a) **For the worst-case scenarios**, describe the largest vessel or pipeline in the process and the regulated substance used for the worst-case, assumptions and parameters used. Assumptions should include any passive mitigation that used to limit the quantity that could be released. State the distance to end point. Documentation should include the anticipated effect of the controls and mitigation on the release quantity and rate. Use a summary table to present the parameters. (A sample OCA Data Table is provided in **Appendix A-13**.)
- (b) **For the alternative release scenarios** (not needed in Program 1), describe the scenarios identified, assumptions and parameters used, and the rationale for the selection of specific scenarios. Assumptions shall include use of any active and passive mitigation that was assumed to limit the quantity that could be released. State the predicted distance to end point. Documentation shall include the effect of the controls and mitigation on the release quantity and rate. Use a summary table to present the parameters. (**Appendix A-13** – Sample OCA Data Table.)
- (c) For each scenario state the chemical name, quantity in pounds, and physical state (toxic only).
- (d) Describe the methodology. Give the computer air model used (if applicable) or state which USEPA Guideline (include version date) was used. If a USEPA Guideline was used, identify the specific tables referenced. If a computer air model was used, describe the known limitations of the air model.
- (e) Document the estimated quantity released, release rate, and duration of the release.
- (f) Document the wind speed and atmospheric stability class (toxic only).
- (g) Document the topography i.e., rural or urban (toxic only).
- (h) Document passive and/or active mitigation (alternative scenario only) considered.
- (i) Provide the data used to estimate the potentially affected population and environmental receptors.
- (j) Provide a list of the following within the zone of vulnerability: the estimated residential population; presence of any schools, hospitals, long term health care facilities, child day care facilities; parks and recreational areas; major commercial, office or industrial areas; and prisons. Populations estimated need only include residential populations and may be rounded to two significant digits (e.g., 5,500; 11,000). Identify and list any environmental receptors, National or state parks, forests, or

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

monuments; officially designated wildlife sanctuaries, preserves, or refuges; and federal wilderness areas.

- (k) Provide a map showing the location of the facility, the vulnerable zone within the radius equal to the distance to the toxic or flammable endpoint for the worst case and alternative scenarios. Identify within the vulnerable zone offsite residences, institutions (e.g., schools, hospitals), industrial, commercial, and office buildings, parks, or recreational areas inhabited or occupied by the public and any identified environmental receptor.

4. ACCIDENT HISTORY/INVESTIGATION (CCR 2745.5) - Provide the following information:

- (a) Identify who is responsible (by title) for investigating accidents.
- (b) Describe management's involvement.
- (c) State if the accident investigation program addresses "near misses". A "near miss", is any incident, which would have resulted in an unintentional release of a Regulated Substance, if action outside the scope of normal operating procedures had not occurred.
- (d) Provide the date, time, and approximate duration of the release.
- (e) The estimated quantity in pounds and the name of the regulated substance released.
- (f) The NAICS code that most closely corresponds to the process.
- (g) The type of release event and its source.
- (h) Weather conditions, if known.
- (i) On-site impacts, offsite impacts & whether offsite impacts were notified.
- (j) Initiating event and contributing factors, if known.
- (k) Operational or process changes that resulted from the investigation.

5. PREVENTION PROGRAM 2 (CCR 2735.5, 2745.6 & Article 5) – For each Program 2 process, provide the following information:

- (a) CalARP Program Management System (CCR 2735.6). Summarize the following items regarding management accountability: Titles of individuals responsible for implementing the RMP and associated programs; Chain-of-command and responsibilities; and the designated RMP coordinator.
- (b) Provide a summary of the Prevention Program 2 elements: safety information, hazard review, operating procedures, training program, maintenance, compliance audits, and incident investigations. The summary should include enough detail to clearly explain how each program element will be managed.
- (c) Provide a specific list of the Federal or state regulations or industry-specific codes and standards used to demonstrate compliance with the safety information requirements. Include the date of the most recent review or revision of the safety information.
- (d) Provide a table of detection and monitoring devices and methods. Include their sensitivities.
- (e) Provide a list of the standard operating procedures in place for the process.
- (f) Provide a list of maintenance procedures in place. Provide a list of the major equipment components of the process that are inspected or tested.
- (g) Provide a list of any changes to the process (SOPs, maintenance, training, etc.) which resulted out of the triennial compliance audit. A person knowledgeable in the process must conduct a compliance audit at least once every three years.

6. PREVENTION PROGRAM 3 (CCR 2735.5, 2745.7 & Article 6) – For each Program 3 process, provide the following information:

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

- (a) CalARP Program Management System (CCR 2735.6). Summarize the following items regarding management accountability: Titles of individuals responsible for implementing the RMP and associated programs; Chain-of-command and responsibilities; and the designated RMP coordinator.
- (b) Provide a summary of your Prevention Program 3 elements: process safety information, operating procedures, training program, mechanical integrity, management of change, pre-start up review, compliance audit, incident investigations, employee participation, hot work permit, and contractors. The summary should include enough detail to clearly explain how each program element will be managed.
- (c) Provide a specific list of the Federal or state regulations or industry-specific codes and standards used to demonstrate compliance with the process safety information requirements. Include the date of the most recent review or revision of the process safety information.
- (d) Provide a table of detection and monitoring devices and methods. Include their sensitivities.
- (e) Provide a list of the standard operating procedures in place for the process.
- (f) Provide a list of maintenance procedures in place. Provide a list of the major equipment components of the process that are inspected or tested.
- (g) Provide a list of any changes to the process (SOPs, maintenance, training, etc.) which resulted out of the triennial compliance audit. A person knowledgeable in the process must conduct a compliance audit at least once every three years.
- (h) Briefly describe the plan you have in place to ensure your employees and their representatives have access to process hazard analysis(s) and other Risk Management Plan information.
- (i) Describe your management of change procedures you have in place to manage changes (other than “replacement in kind”) to process chemicals, technology, equipment, and procedures.
- (j) Summarize your and the contractor’s responsibilities where contractors perform maintenance or repair, turnaround, major renovation, or specialty work on or adjacent to a covered process.

7. EXTERNAL EVENTS & SEISMIC ASSESSMENT - Provide the following information regarding external events that were reviewed as part of or separate from your hazard review or process hazard analysis:

- (a) A list of the types of natural or human caused external events considered.
- (b) A description of the parameters used in considering a seismic analysis.
- (c) Provide the edition of the Uniform Building Code that was used when the process was designed.
- (d) Provide for each external event, with a potential to create a release of a regulated substance that will reach an endpoint offsite: the expected date of completion of any changes to mitigate the potential release; a description of the major hazards identified; process controls and mitigation in place; monitoring and detection systems in use.
- (e) Seismic Assessment (See **Appendix A-2 Useful Weblinks**)

8. EMERGENCY RESPONSE PLAN (Article 7) – Provide the following information [a copy of your Hazardous Materials Business Plan *may* suffice for (a) and (b)].

- (a) Describe what emergency response procedures you have in place in the event of a release of a regulated substance. This must describe the actions to be taken by employees and other individuals on-site over the entire course of the release event including at least the following:
 - Activation of alarm system and interpretation of signals.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

- Safe evacuation, assembly, and return.
 - Selection of response strategies and incident command structure.
 - Use of response equipment and other release mitigation activities; and
 - Post-release equipment and personnel cleanup and decontamination.
- (b) Describe the offsite response assistance you require for potential release scenarios, including firefighting, security, and notification of the public.
- (c) Describe who will oversee the response operation and how will authority be delegated down the internal and offsite chain of command.
- (d) Describe any planned drills with emergency responders. Provide dates.
- (e) List all other federal or state emergency plan requirements to which your facility is subject.

9. **CERTIFICATION** – The RMP shall be certified as follows:

- (a) **Program 1**, certify in the RMP the following: “Based on the criteria in Section 2735.4 of Title 19 CCR, the distance to the specified endpoint for the worst-case accidental release scenario for the following process(es) is less than the distance to the nearest public receptor: [list process(es)]. Within the past five years, the process(es) has (have) had no accidental release that caused offsite impacts provided in the risk management program Section 2735.4(c)(1). No additional measures are necessary to prevent offsite impacts from accidental releases. In the event of fire, explosion, or a release of a regulated substance from the process(es), entry within the distance to the specified endpoints may pose a danger to public emergency responders. Therefore, public emergency responders should not enter this area except as arranged with the emergency contact indicated in the RMP. The undersigned certifies that, to the best of my knowledge, information, and belief, formed after reasonable inquiry, the information submitted is true, accurate, and complete”. This certification must be signed by the owner or operator of the stationary source, and shall include his or her title, and the date signed.
- (b) **Program 2 and 3**, the owner or operator shall submit in the RMP a single certification that, to the best of the signer’s knowledge, information, and belief formed after reasonable inquiry, the information submitted is true, accurate, and complete. This certification must be signed by the owner or operator of the stationary source, and shall include his or her title, and the date signed.

10. **Updates (CCR 2745.10)** - The RMP must be updated when the following exist:

- At least every five years from the initial submittal date
- Within six months of a process change that requires a new process hazard analysis
- Within six months of a change that causes the offsite consequence to change by a factor of two
- Within six months of a change that results in a Prevention Program level change

11. **Corrections (CCR 2745.10.5)** - The following corrections must be submitted to include in the RMP:

- Information on any accidental releases that require an incident investigation.
- Emergency contact information must be updated within 30 days of a change in personnel. (Use **Appendix A-1** - CalARP Registration Form HM-9240.)
- RMP submittals, corrections and updates go directly to the CUPA, not through CERS currently.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

GUIDELINES FOR TECHNICAL SUPPORTING RECORDS AND STUDIES

CALIFORNIA ACCIDENTAL RELEASE PREVENTION PROGRAM (CalARP)

Separate records for supporting technical information must be prepared and maintained in the custody of the facility. The County of San Diego, Department of Environmental Health & Quality, Hazardous Materials Division (HMD) may choose to review the facility's supporting technical information by either requesting submittal of such information or requesting to review such information during an audit of the facility.

Facilities are to categorize the records of their supporting technical information as follows:

- I. Offsite Consequence Analysis
- II. Prevention Program
 - a. Program 2 (if applicable)
 - b. Program 3 (if applicable)
- III. Hazard Review or Process Hazard Analysis (depending on Program)
- IV. External Events (including Seismic Analysis)

In developing your Risk Management Program, the HMD recommends the following USEPA Guidance Documents: (Download them from the EPA web site: <http://www2.epa.gov/rmp/guidance-facilities-risk-management-programs-rmp>)

Accidental Release Information Program- all facilities

Risk Management Program and Plan for Ammonia Refrigeration - Ammonia Refrigeration facilities

Risk Management Program and Plan for Water Treatment - Water Treatment facilities

Risk Management Program and Plan for Propane Users and Small Retailers - Propane facilities

Risk Management Program and Plan for Propane Storage Facilities – Propane facilities

Risk Management Program and Plan for Wastewater Treatment Plants – Wastewater Treatment facilities

Risk Management Program and Plan for Warehouses – Warehouse facilities

Risk Management Program and Plan for Chemical Distributors – Chemical Distribution facilities

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

I. OFFSITE CONSEQUENCE ANALYSIS

The Offsite Consequence Analysis (OCA) describes the potential exposure levels for surrounding populations and environmental receptors from accidental releases of regulated substances. Specific release scenarios and dispersion models are used to generate a circular area of theoretical exposure around the point of accidental release. This generated exposure area is referred to as a “zone of vulnerability.”

Computer air dispersion models or tables such as those provided in USEPA’s “*RMP Offsite Consequence Analysis Guidance*” documents are used to calculate the air concentration of a regulated substance as a function of distance from the point of release. A “toxic endpoint” concentration value for the regulated substance is input into a computer model, or looked up in table, for the selected release scenario to calculate the “zone of vulnerability.”

A. General Requirements

Section 2745.4 in Title 19 CCR, requires that the following offsite consequence scenarios be performed:

Program 1

Worst-Case

One worst-case release scenario for each Program 1 process must be evaluated.

Program 2 or 3

Worst-Case

A minimum of one worst-case release scenario to represent all toxic regulated substances held above the threshold quantity and one worst-case release scenario to represent all flammable regulated substances held above the threshold quantity. Additional worst-case release scenarios may be needed depending on the regulated substances in process and their location within the facility.

Alternative

A minimum of one alternative release scenario for each toxic regulated substance held above the threshold quantity and one alternative release scenario to represent all flammable regulated substances held above the threshold quantity.

B. Air Dispersion Models

The USEPA has developed a “RMP Offsite Consequence Analysis Guidance” document that includes tables for calculating air dispersion of toxic regulated substances and flammable regulated substances. One model accepted is RMP Comp. This is available to download at no cost from:

<http://www2.epa.gov/rmp/general-rmp-guidance-chapter-4-offsite-consequence-analysis>

Proprietary computer air models must be approved by the HMD. Prior to approval for use, the HMD may request a copy of the proprietary computer air model from the facility or their consultant for review.

C. Worst-Case Release Scenario Analysis

1. Definition of Worst-Case Scenario

USEPA has defined worst-case release as the release of the largest quantity of a regulated substance from a vessel or process line failure that results in the greatest distance to a specified endpoint. For the worst-case analysis, you do not need to

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

consider the possible causes of the worst-case release or the probability that such a release might occur; the release is simply assumed to take place.

2. Determining the Quantity for the Worst-Case Scenario

For analysis of the worst-case scenario, you must consider the largest quantity of a regulated substance handled on site in a single vessel or process line failure at any one time. A summary of worst-case release scenario requirements is provided in the appendices.

It is the HMD's opinion that administrative controls to limit vessel inventories are often disregarded by facilities under special circumstances when additional inventory is needed. Therefore, the HMD does not allow administrative controls to be considered when determining worst-case analysis.

3. Selecting a Worst-Case Scenario

The hazard assessment requires a single offsite consequence analysis of the worst-case scenario for substances in each hazard category (i.e., one for regulated toxic substances and one for regulated flammable substances). Only the hazard for which the substance is listed needs to be considered (i.e., substances on the list of regulated toxic substances that are also flammable should only be analyzed for their toxic hazard; substances on the list of regulated flammable substances should be considered only for flammability).

The regulated substance chosen for the consequence analysis for each hazard should be the regulated substance that has the potential to cause the greatest offsite consequence. Choosing the toxic regulated substance that might lead to the greatest offsite consequence may require a screening analysis of all the regulated toxic substances on site, because the potential consequences are dependent on a number of factors, including quantity, toxicity, and volatility. Location (distance to fence line) and conditions of processing or storage (e.g., a high temperature process) also should be considered.

For regulated flammable substances, the consequences of a vapor cloud explosion must be considered in the analysis. The severity of the consequences of a vapor cloud explosion depends on the quantity of the released regulated substance in the vapor cloud and its heat of combustion. In most cases, the analysis probably should be based on the regulated flammable substance present in the greatest quantity; however, a substance with a high heat of combustion may have a greater potential offsite impact than a larger quantity of a substance with a lower heat of combustion. In some cases, a regulated flammable substance that is close to the fence line might have a greater potential offsite impact than a larger quantity farther from the fence line.

For worst-case scenarios, you are allowed to consider passive mitigation system, such as dikes, firewalls, blast walls, enclosures, etc. Active mitigation systems are not allowed to be considered for worst-case scenarios.

4. Release Rates for Worst-Case Scenarios

Toxic Substances

Toxic Gases (Normally Gases at Ambient Temperature)

Toxic gases include all regulated toxic substances that are gases at ambient temperature (temperature 25°C, 77°F). For consequence analysis, a gaseous release of

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

the total quantity is assumed to occur in 10 minutes. The release rate shall be assumed to be the total quantity divided by 10.

Toxic Gases (Refrigerated Liquids at Ambient Pressure)

For regulated toxic substances handled as refrigerated liquids at ambient pressure, if the released regulated toxic substance is not contained by passive mitigation systems or if the contained pool would have a depth of 1 centimeter or less, you must assume that the regulated toxic substance is released as a gas in 10 minutes. If the released regulated toxic substance is contained by passive mitigation systems in a pool with a depth greater than 1 centimeter (0.39-inch), you may assume that the quantity of regulated toxic substance is spilled instantaneously to form a liquid pool. The volatilization rate (release rate) shall be calculated at the boiling point of the regulated toxic substance.

Toxic Liquids

For regulated toxic substances handled as liquids, the total quantity in a vessel is assumed to be spilled onto a flat, non-absorbing surface. (NOTE: if the release would occur onto a surface that is not paved or smooth, you may take into account the actual surface characteristics). For toxic liquids carried in pipelines, the quantity that may be released from the pipeline is assumed to form a pool.

The surface area of the pool shall be determined by assuming that the liquid spreads to 1 centimeter (0.39-inch) unless passive mitigation systems are in place that serves to contain the spill and limit the surface area. Where passive mitigation is in place, the surface area of the contained liquid shall be used to calculate the volatilization rate.

The volatilization rate shall account for the highest daily maximum temperature occurring in the past three years, the temperature of the substance in the vessel, and the concentration of the substance if the liquid spilled is a mixture or solution.

The rate of release to air shall be determined from the volatilization rate of the liquid pool. You may use the methodology in USEPA's "*RMP Offsite Consequence Analysis Guidance*" or any other publicly available techniques that account for the modeling conditions and are recognized by industry as applicable as part of current practices.

Flammables

For regulated flammable substances, including both flammable gases and volatile flammable liquids, the worst-case release is assumed to result in a vapor cloud containing the total quantity of the regulated substance that could be released from a vessel or pipeline. The entire quantity of the cloud is assumed to be between the upper and lower flammable limits of the regulated substance. A yield rate of 10 percent of the available energy released in the explosion shall be used to determine the distance to the explosion endpoint if the model used is based on TNT-equivalent methods.

Required Parameters for Modeling a Worst-Case Scenario

Endpoints

Toxic - The toxic endpoints to be used in a worst-case analysis shall be taken from the most current Tables listed in Appendix A Title 19 CCR. These endpoints are based on the Emergency Response Planning Guidelines (ERPG) developed by the American Industrial Hygiene Association. The ERPPG 2 is the maximum airborne concentration below which it is believed nearly all individuals could be exposed for up to one hour without experiencing or developing irreversible or other serious health effects or symptoms which could impair an individual's ability to take protective action.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

Flammables - The flammable endpoint used in a worst-case analysis is an overpressure of 1 pound per square inch (psi), a radiant heat/exposure time of 5kw/square meter, or the lower flammability limit. USEPA chose this endpoint as the threshold for potential serious injuries to people as a result of property damage caused by an explosion such as injuries from flying glass from shattered windows or falling debris from damaged houses.

Wind Speed/Atmospheric Stability

For worst-case analysis use a wind speed of 1.5 meters per second and an F atmospheric stability class unless you can demonstrate that local meteorological data applicable to the site show a higher minimum wind speed or less stable atmosphere at all times during the previous three years.

Ambient Temperature/Humidity

For worst-case analysis of regulated toxic substances use the highest daily maximum temperature in the previous three years and average humidity for the site, based on temperature/humidity data gathered at the stationary source or at a local meteorological station. If you use 25°C and 50% humidity as values for these variables if you are using the USEPA's RMP Offsite Consequence Analysis Guidance or RMP Comp as your air model.

Height of Release

For worst-case analysis of regulated toxic substances assume a ground level (0 feet) release.

Surface Roughness (Topography)

For worst-case analysis use either urban or rural topography, as appropriate. Urban means that there are many obstacles in the immediate area and the terrain is generally flat and unobstructed. Rural means there are no buildings in the immediate area and the terrain is generally flat and unobstructed.

Dense or Neutrally Buoyant Gases

For worst-case analysis tables or models used for dispersion of regulated toxic substances must appropriately account for gas density.

Temperature of the Released Substance

For worst-case analysis consider liquids (other than gases liquefied by refrigeration) to be released at the highest daily maximum temperature, based on data for the previous three years, or at process temperature, whichever is higher. Assume gases liquefied by refrigeration at atmospheric pressure are released at their boiling points.

D. Alternative Release Scenario Analysis

1. Selecting Alternative Release Scenarios

You are required to analyze at least one alternative release scenario for each listed regulated toxic substance you have in a **Program 2** or **Program 3** process above its threshold quantity. You are also required to analyze one alternative release scenario to represent all regulated flammable substances covered in your Program 2 or Program 3 processes. You do not need to analyze an alternative release scenario for each regulated flammable substance. For example, if you have five regulated substances – ammonia, chlorine, hydrogen chloride, acetylene, and propane – above the threshold quantity in either Program 2 or Program 3 processes, you will need to analyze one alternative scenario each for ammonia, chlorine, and hydrogen chloride; and a single alternative scenario to cover both acetylene and propane.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

For alternative release scenarios, you are allowed to consider active mitigation systems, such as scrubber systems, shutdown systems, pressure relieving devices, emergency isolation systems, fire sprinklers, and deluge systems, as well as passive mitigation systems.

Alternative Release Scenarios for Regulated Toxic Substances

Alternative release scenarios for regulated toxic substances should be those that will reach an endpoint offsite, unless no such scenario exists. Those releases that have the potential to reach the public are of the greatest concern.

You should consider the following when selecting an alternative release scenario:

- You may use your worst-case release scenario and apply your active mitigation system to limit the quantity released and the duration of the release.
- You may use information from your *process hazard analysis.
- You may use an actual event based on your five-year accident history review.
- You may use an actual event based on industry accident history as it relates to your process.

*If you use information for your process hazard analysis to select your alternative scenario, you should at a minimum consider the following: (a) transfer hose releases due to splits or sudden hose uncoupling; (b) process piping releases from failures at flanges, joints, welds, valves, and valve seals, and drains or bleeds; (c) process vessel or pump releases due to cracks, seal failure, or drain, bleed, or plug failure; (d) vessel overfilling and spill, or over pressurization an venting through relief valves or rupture disks; and (e) shipping containers mishandling and breakage or puncturing leading to a spill. In addition, if you use your process hazard analysis to select an alternative scenario, you must justify your choice either qualitatively or quantitatively.

Alternative Scenarios for Regulated Flammable Substances

Alternative release scenarios for regulated flammable substances are somewhat more complicated than those release scenarios for regulated toxic substances because the consequences of a release and the endpoint of concern may vary. For the worst-case, the consequence of concern is a vapor cloud explosion, with an overpressure endpoint. For alternative scenarios (e.g., fires), other endpoints (e.g., heat radiation) may need to be considered. Possible scenarios to consider that would involve regulated flammable substances include:

- Vapor cloud fires (flash fires) resulting from dispersion of a cloud of flammable vapor and ignition of the cloud following dispersion. Such a fire could flash back and could represent a severe heat radiation hazard to anyone in the area of the cloud.
- A pool fire, with potential radiant heat effects, resulting from a spill of a flammable liquid.
- A boiling liquid, expanding vapor explosion (BLEVE), leading to a fireball that may produce intense heat and may occur if a vessel containing a flammable material ruptures explosively as a result of exposure to fire. Heat radiation from the fireball is the primary hazard; vessel fragments and overpressure from the explosion also can result. BLEVEs are generally considered unlikely events.

2. Release Rates for Alternative Scenarios

Refer to USEPA's "RMP Offsite Consequence Analysis Guidance" to determine appropriate release rates.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

3. Required Parameters for Modeling an Alternative Scenario Endpoints

Toxic

The toxic endpoints used in a worst-case analysis shall be taken from the most current Tables listed in Section 2770.5 of Title 19 CCR.

Flammables

The flammable endpoint to be used in an alternative analysis varies according to the scenarios studied and may be one of the following: (a) an overpressure of 1 pound per square inch (psi); (b) a radiant heat level of 5 kilowatts per square meter (kW/m²) for 40 seconds for heat from fires (or equivalent dose); or (c) lower flammability limit (LFL) as specified in NFPA documents or other generally recognized sources.

Wind Speed/Atmospheric Stability

For site-specific modeling, use typical meteorological conditions for your site. If you use USEPA's *"RMP Offsite Consequence Analysis Guidance"* you may assume wind speed of 3 meters per second and a D atmospheric stability class.

Ambient Temperature/Humidity

For site-specific modeling, use average temperature/humidity data gathered at the site or at a local meteorological station. Assume 25°C and 50% humidity as values if you are using the USEPA's *"RMP Offsite Consequence Analysis Guidance"* as your air model.

Height of Release

Release height may be determined by the release scenario or by assuming ground level.

Surface Roughness (Topography)

Use either urban or rural topography, as appropriate. Urban means that there are many obstacles in the immediate area and the terrain is generally flat and unobstructed. Rural means there are no buildings in the immediate area and the terrain is generally flat and unobstructed.

Dense or Neutrally Buoyant Gases

Tables or models used for dispersion of regulated toxic substances must appropriately account for gas density. Most computer models automatically take this into account.

Temperature of the Released Substance

Substances may be considered to be released at a process or ambient temperature that is appropriate to the scenario. If you are using the USEPA's *"RMP Offsite Consequence Analysis Guidance"* as your air model 25°C or the boiling point of the released substance may be used.

E. Defining Offsite Impacts to the Population/Environment

For each release scenario, estimate the population within a circle (zone of vulnerability) with its center at the point of the release and a radius determined by the distance to the endpoint. Population shall include residential population. To estimate the population potentially affected, use the most recent Census data or other more accurate information if it is available. Population data shall be estimated to two significant digits.

Include within the zone of vulnerability the presence of any schools, hospitals, long-term health care facilities, child day care facilities, and prisons. Identify and list any environmental receptors, parks and recreational areas, major commercial, office, and industrial buildings. Environmental receptors can be determined from local United States Geological Survey (USGS) maps.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

F. Offsite Consequence Analysis Technical Document

The following records shall be maintained on the offsite consequence analyses in a Technical Document. This Technical Document will be subject to submittal upon HMD's request and/or by onsite auditing by HMD:

1. Include a table of contents.
2. Place divider tabs or pages between sections of the OCA.
3. For the worst-case scenarios, describe the vessel or pipeline and substance selected as worst-case, assumptions and parameters used, and the rationale for selection. Assumptions should include any passive mitigation that was assumed to limit the quantity that could be released.
4. For the alternative release scenarios, describe the scenarios identified, assumptions and parameters used, and the rationale for the selection of specific scenarios. Assumptions shall include use of any active and passive mitigation that was assumed to limit the quantity that could be released
5. Include the same information required in the RMP Public Document in this section.
6. If using a computer air model, include the computer-generated runs of the scenario(s). Be prepared to provide a copy of the air modeling software if not using the available USEPA RMP Comp or ALOHA.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

II. PREVENTION PROGRAM 2

Most Program 2 processes are likely to be relatively simple and may be located at small businesses. EPA developed the Program 2 prevention program by identifying the basic elements that are the foundation of sound prevention practices - safety information, hazard review, operating procedures, training, maintenance, compliance audits, and accident investigation. As important as each of the elements is, you will not gain the full benefit from them unless you integrate them into a risk management system that you implement on an on-going basis. For example, the hazard review must be built on the safety information; the results of the hazard review should be used to revise and update operating and maintenance procedures. Workers must be trained in these procedures and must use them every day.

There are seven elements in the Program 2 prevention program found in Article 5 of Title 19 CCR.

SUMMARY OF PROGRAM 2 PREVENTION PROGRAM

Title 19 Div 2 Ch 45	Section Title
Section 2755.1	Safety Information
Section 2755.2	Hazard Review
Section 2755.3	Operating Procedures
Section 2755.4	Training
Section 2755.5	Maintenance
Section 2755.6	Compliance Audits
Section 2755.7	Incident Investigation

You must integrate these seven elements into a risk management program that you and your employees implement daily. Understanding and managing risks must become part of the way you operate. Doing so will provide benefits beyond accident prevention as well. Preventive maintenance and routine inspections will lessen the number of equipment failures and down time; well trained workers, aware of optimum operating parameters, will allow you to gain the most efficient use of your substances.

A. Safety Information

You must compile and maintain safety information related to the regulated substances and process equipment for each Program 2 process. You probably have much of this information already because you would have developed it to comply with OSHA or other rules. EPA has limited the information to what is likely to apply to the processes covered under the Program 2 release prevention program.

SAFETY INFORMATION REQUIREMENTS

Information you must compile and maintain:	You must ensure:	You must update the safety information if:
<ul style="list-style-type: none"> ✓ Safety Data Sheets ✓ Maximum intended inventory ✓ Safe upper and lower parameters ✓ Equipment specifications ✓ Codes & standards used to design, build, and operate the process. 	<ul style="list-style-type: none"> ✓ That the process is designed in compliance with recognized codes and standards 	<ul style="list-style-type: none"> ✓ There is a major change at your business that makes the safety information inaccurate

After you have documented your safety information, you should double-check it to be sure that the files you have reflect the equipment you are currently using. It is important to keep this

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

information up to date. (**Appendix A-4** is a Sample Audit Checklist for Safety Information.) Whenever you replace equipment, be sure that you put the new equipment specifications in the file and consider whether any of your other prevention elements need to be reviewed to reflect the new equipment.

B. Hazard Review

You do not have to perform a full Process Hazard Analysis for a Program 2 process, but you must conduct a hazard review. The hazard review will help you determine whether you're meeting applicable codes and standards, identify and evaluate the types of potential failures, and focus your emergency response planning efforts. The hazard review is key to understanding your operation and continuing to operate safely. You must identify and review specific hazards and safeguards of your Program 2 processes. The HMD requires at a minimum a "What-If" hazard evaluation as the hazard review methodology for Program 2 processes.

HAZARD REVIEW REQUIREMENTS

Conduct a review & identify...	Use a guide for conducting the review	Document results & resolve problems	Update your hazard review
<ul style="list-style-type: none"> ✓ The hazards associated with the Program 2 process & regulated substances. ✓ Opportunities for equipment malfunction or human error that could cause a release. ✓ Safeguards that will control the hazards or prevent the malfunction or error. ✓ Steps to detect or monitor releases. 	<ul style="list-style-type: none"> ✓ You may use any what-if/checklist (such as you might in a model risk management program) to conduct the review. ✓ For a process designed to industry standards like NFPA-58 or Federal/state design rules, check the equipment to make sure that it's fabricated, installed and operated properly. 	<ul style="list-style-type: none"> ✓ Your hazard review must be documented, and you must show that you have addressed problems. 	<ul style="list-style-type: none"> ✓ You must update your review at least once every five years or whenever there is a major change in the process. ✓ You must resolve problems identified in the new review before you start up the changed process.

Record the Results

The team scribe should record the results during the "What-If" Hazard Evaluation process. An example of the recording format is provided in **Appendix A-10**.

"What-If" Hazard Evaluation Document

The Hazard Evaluation Document **shall include the following:**

- ◆ Include a table of contents.
- ◆ Place dividers and tabs between the sections of the "What-If" Hazard Evaluation document
- ◆ Describe the regulated substance process(es) studied including a review of the chemistry and chemical reactions that take place in the system.
- ◆ Provide a copy of the process flow diagram and color code regulated substance lines.
- ◆ List the individual pieces of equipment (e.g., pumps, reactors, heat exchangers) and piping that were studied.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

- ◆ Provide a copy of the "What-If" questions used in the evaluation.
- ◆ Provide copies of the "What-If" session worksheets.
- ◆ Include a table of the individuals involved in the "What-If" Hazard Evaluation and the role of each individual.
- ◆ Include a table of all the procedures reviewed during the "What-If" Hazard Evaluation.
- ◆ Include a table of all the documents reviewed during the "What-If" Hazard Evaluation.
- ◆ Define the extent of on-site interviewing of operators used to estimate human/operator error probabilities.
- ◆ Describe the database or sources used to estimate equipment failure.
- ◆ Provide a table of all the recommendations from the "What-If" Hazard Evaluation by individual equipment according to process flow and in order of priority. Separate the recommendations that will be addressed from those that will not be addressed. For all recommendations that will not be addressed explain why they will not be addressed.
- ◆ Provide a table of references used in the hazard analysis.

Remember that you have a general duty to prevent accidents and ensure safety at your source, which may require you to take steps beyond those explicitly, specified in the risk management program rule. (CA HSC 25531.2.(b))

C. Operating Procedures

You must prepare written operating procedures that give workers clear instruction for safely conducting activities involving a covered process. You may use standardized procedures developed by industry groups or provided in model risk management programs as the basis for your operating procedures but be sure to check that these standard procedures are appropriate for your activities. If necessary, you must update your Program 2 operating procedures whenever there is a major change and before you start up the changed process. The following table briefly summarizes what your operating procedures must address.

OPERATING PROCEDURES REQUIREMENTS

Steps for each operating phase	Operating limits
<ul style="list-style-type: none"> ✓ Initial startup ✓ Normal operations ✓ Temporary operations ✓ Emergency shutdown ✓ Emergency operations ✓ Normal shutdown ✓ Startup following a normal or emergency shutdown or a major change 	<ul style="list-style-type: none"> ✓ Consequences of deviating ✓ Steps to avoid, correct deviations
<p>Local requirements:</p> <ul style="list-style-type: none"> ✓ Delivery Procedures ✓ Cylinder change out procedures ✓ Release investigation procedures 	<ul style="list-style-type: none"> ✓ Equipment components must be properly identified to avoid operating errors

You must update your procedures whenever you change your process in a way that alters the

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

steps needed to operate safely. If you add new equipment, you will need to expand your procedures or develop a separate set to cover the new items. Whenever you change your safety information you should review your procedures to be sure that they are still appropriate. Anytime you conduct a hazard review, check your operating procedures as you implement changes to address hazards.

You must maintain your current set of operating procedures. You are not required to keep old versions; in fact, you should avoid doing so because keeping copies of outdated procedures may cause confusion. You should date all procedures so you will know when they were last updated.

D. Training

Training programs often provide immediate benefits because trained workers have fewer accidents, damage less equipment, and improve operational efficiency. Training gives workers the information they need to understand how to operate safely and why safe operations are necessary. A training program, including refresher training, is the key to ensuring that the rest of your prevention program is effective. You already have some type of training program because you must conduct training to comply with OSHA's Hazard Communication standard (29 CFR 1910.1200).

The following lists things that you may find useful in developing your training program.

- ◆ Clearly identify the employees who need to be trained and the subjects to be covered.
- ◆ Specify learning objectives, and write them in clear, measurable terms before training begins. Remember that training must address the process operating procedures and the maintenance procedures.
- ◆ Tailor the specific training modules or segments to the training objectives. Enhance learning by including hands-on training like using simulators whenever appropriate. Make the training environment as much like the working environment as you can, consistent with safety. Allow your employees to practice their skills and demonstrate what they know.
- ◆ Evaluate your training program periodically to see if your employees have the skills and know the routines required under your operating procedures. Make sure that language or presentations are not barriers to learning. Decide how you will measure your employee's competence.
- ◆ Make sure all workers - including maintenance and contract employees - receive initial and refresher training. If you make changes to process chemicals, equipment, and technology, make sure that involved workers understand the changes and the effects on their job.

Keep documentation of your training program. An attendance log for any formal training courses and refresher training is required to ensure that everyone who needs to be trained is trained. Such logs will help you when you do a compliance audit.

E. Maintenance/Mechanical Integrity

You must prepare and implement written procedures for maintaining the mechanical integrity of process equipment and train your workers in the maintenance procedures. You may use procedures or instructions from equipment vendors, in Federal or State regulations, or in industry codes as the basis of your maintenance program. You must develop a schedule for inspecting and testing your equipment based on manufacturers' recommendations, industry standards or your own experience. The following is a summary of the elements of a maintenance program:

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

Written procedures	Training	Inspection & testing
<ul style="list-style-type: none"> ✓ You may use someone else's procedures as the basis for your program. If you choose to develop your own, you must write them down. 	<ul style="list-style-type: none"> ✓ Train process maintenance employees in process hazards and how to avoid or correct an unsafe condition. ✓ Make sure this training covers the procedures applicable to safe job performance. 	<ul style="list-style-type: none"> ✓ Inspect & test process equipment. ✓ Use recognized and generally accepted good engineering practices. ✓ Follow a schedule that matches the manufacturer's recommendations or that prior operating experience indicates is necessary.

Keep your written procedures and schedules as well as any agreements you have with contractors. Keep training logs or maintenance logs. Without some record, you will have to rely on workers' memories about when something was last checked. As workers leave or change jobs at your company, it can be difficult to keep track of when inspections and tests were done. Maintaining a record of when something was last done or is scheduled to be done next can help keep your program working smoothly.

F. Compliance Audits

At least every three years, you must certify that you have evaluated your compliance with the requirements for the prevention program for each covered process. At least one person on your audit team must be knowledgeable about the covered process. You must develop a report of your findings, determine, and document an appropriate response to each finding, and document that you have corrected any deficiency. Coordinate with the CUPA on timelines for addressing compliance audit recommendations. The CUPA believes that recommendations from compliance audits should be addressed within one (1) year from the audit date. If you need more time, coordinate with the CUPA on a timeline.

You must keep a written record of the findings and actions. Maintain copies of the last two compliance audits.

G. Incident Investigation

You must investigate each incident which resulted in or could have resulted in a "catastrophic release" of a regulated substance." A catastrophic release is one that presents an imminent and substantial endangerment to public health and the environment. The following table briefly summarizes the steps you must take for investigating incidents. You should also consider investigating minor accidents or near misses because they may help you identify problems that could lead to more serious accidents.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

INCIDENT INVESTIGATION REQUIREMENTS

✓ Initiate an investigation promptly.	Begin investigating no later than 48 hours following the incident.
✓ Summarize the investigation in a report.	Among other things, this report will include the factors contributing to the incident. Remember that identifying the root cause may be more important than identifying the initiating event. Remember, also, that the purpose of the report is to help management take corrective action.
✓ Address the team's findings and recommendations	Establish a system to address the incident report findings and recommendations and document resolutions and corrective actions.
✓ Review the report with your staff and contractors	You must share the report - its findings and recommendations - with affected workers whose job tasks are relevant to the incident.
✓ Retain the report.	Keep incident investigation summaries for five years.

You must maintain the summary of the accident, recommendations, and actions. A sample is in **Appendix A-3**. Note that the sample includes accident data that you will need for the five-year accident history. These data are not necessarily part of the incident investigation report but including them will create a record you can use later to create the accident history.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

III. PREVENTION PROGRAM 3

If you already have the OSHA Process Safety Management (PSM) program in place you will need to do little that is new to comply with the Program 3 prevention program. Whether you're building on to the PSM standard or creating a new program, keep these things in mind.

- ◆ CalARP and OSHA have different authority. If you are already complying with the PSM standard, your Process Hazard Analysis (PHA) team may have to assess new hazards that could affect the public or the environment offsite. Protection measures that are suitable for workers may be the very kind of thing that imperils the public.
- ◆ Integrate the elements of your prevention program. You must ensure that a change in any single element of your program leads to a review of other elements to identify any effect caused by the change.
- ◆ Most important, make accident prevention an institution at your site. Like the entire risk management program, a prevention program is more than a collection of written documents. It is a way to make safe operations and accident prevention the way you do business every day.

There are twelve elements in the Program 3 prevention program. Two OSHA elements are not included. Emergency Response is addressed separately in CalARP; the OSHA Trade Secrets requirement (provision of trade secret information to employees) is beyond the CalARP statutory authority.

SUMMARY OF PROGRAM 3 PREVENTION PROGRAM

19 CCR Section	Title	OSHA PSM Reference
2760.1	Process Safety Information	PSM Standard Section 1910.119(d)
2760.2	Process Hazard Analysis (PHA)	PSM Standard Section 1910.119(e)
2760.3	Operating Procedures	PSM Standard Section 1910.119(f)
2760.4	Training	PSM Standard Section 1910.119(g)
2760.5	Mechanical Integrity	PSM Standard Section 1910.119(j)
2760.6	Management of Change	PSM Standard Section 1910.119(l)
2760.7	Pre-Startup Safety Review	PSM Standard Section 1910.119(i)
2760.8	Compliance Audits	PSM Standard Section 1910.119(o)
2760.9	Incident Investigation	PSM Standard Section 1910.119(m)
2760.10	Employee Participation	PSM Standard Section 1910.119(c)
2760.11	Hot Work Permits	PSM Standard Section 1910.119(k)
2760.12	Contractors	PSM Standard Section 1910.119(h)

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

A. Process Safety Information

The following table summarizes the safety information requirements.

PROCESS SAFETY INFORMATION

For chemicals, you must complete information on:	For process technology, you must provide:	For equipment in the process, you must include:
<ul style="list-style-type: none">✓ Toxicity✓ Permissible exposure limits✓ Physical data✓ Reactivity✓ Corrosivity✓ Thermal & chemical stability✓ Hazardous affects you can foresee if you mixed materials together accidentally	<ul style="list-style-type: none">✓ A block diagram or simplified process flow diagram✓ Information on process chemistry✓ Maximum intended inventory of the CalARP-regulated chemical✓ Safe upper & lower limits for such items as temperature, pressure, flows, or composition✓ An evaluation of the consequences of deviation	<ul style="list-style-type: none">✓ Materials of construction✓ Block flow diagram✓ Piping & instrumentation diagrams (P&IDs)✓ Electrical classification✓ Relief system design & design basis✓ Ventilation system design✓ Design codes & standards employed✓ Safety systems

B. Process Hazard Analysis

A process hazard analysis (PHA) is one of the most important elements of the process safety management program. A PHA is an organized and systematic effort to identify and analyze the significance of potential hazards associated with the processing or handling of highly hazardous chemicals. A PHA provides information that will assist employers and employees in making decisions for improving safety and reducing the consequences of unwanted or unplanned releases of hazardous chemicals.

A PHA is directed toward analyzing potential causes and consequences of fires, explosions, releases of toxic or flammable chemicals and major spills of hazardous chemicals. The PHA focuses on equipment, instrumentation, utilities, human actions (routine and non-routine), and external factors that might impact the process. These considerations assist in determining the hazards and potential failure points or failure modes in a process.

Selection of a PHA methodology

In San Diego County most facilities that conducted a PHA under the previous Risk Management and Prevention Program used either a What-If/Checklist method or Hazard and Operability Analysis (HAZOP). For more detailed information regarding these techniques refer to *"Guidelines for Hazard Evaluation Procedures, 2nd Ed."*, published by Center for Chemical Process Safety of the American Institute of Chemical Engineers.

Offsite impacts

You must consider offsite impacts when you conduct a PHA under CalARP. A well-done PHA should identify all failure scenarios that could lead to significant exposure of workers, the public, or the environment. The only issue that is likely to require consideration above what you have done already for the PSM standard is whether any protection measures that were adequate for worker safety are inadequate for public and environmental safety.

Consider two circumstances - one where PSM and the risk management program rule should lead to the same result, and another where protecting workers could mean endangering the public and the environment. For flammables, any scenario that could affect

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

the public almost certainly would have the potential to affect workers; measures taken to protect your employees likely will protect the public and the environment. On the other hand, for toxics under PSM, you may plan to address a loss of containment by venting toxic vapors to the outside air. In each circumstance, a PHA should define the failure sequence. However, for toxics, the PHA team must reassess venting as an appropriate mitigation measure.

Rejecting team recommendations

You may not always agree with your PHA team's recommendations and may wish to reject a recommendation. You may decline a team recommendation if you can document one of the following:

- (1) the analyses upon which the recommendations are based contain factual errors.
- (2) the recommendation is not necessary to protect the health of employees, the public or contractors.
- (3) an alternative measure would provide a sufficient level of protection; or
- (4) the recommendation is infeasible.

Updating and revalidating your PHA

For CalARP, you must complete the initial PHA for each Program 3 process prior to bringing a threshold quantity of a regulated substance in a process on site and update it at least once every five years. You may use an OSHA PHA as your initial PHA, and update and revalidate it every five years on the OSHA schedule.

Revising your PHA

You should revise your PHA whenever there is a new hazard or risk created by changes to your process. Such changes might include introducing a new process, process equipment, or regulated substance; altering process chemistry that results in any change to safe operating limits; or other alteration that introduces a new hazard. However, EPA recommends that you consider revising your PHA whenever adjoining processes create a hazard.

Remember that you have a general duty to prevent accidents and ensure safety at your source, which may require you to take steps beyond those explicitly, specified in the risk management program rule. (CA HSC 25531.2.(b))

C. Operating Procedures

You must prepare written operating procedures that give workers clear instruction for safely conducting activities involving a covered process. You may use standardized procedures developed by industry groups or provided in model risk management programs as the basis for your operating procedures but be sure to check that these standard procedures are appropriate for your activities. If necessary, you must update your Program 3 operating procedures whenever there is a major change and before you start up the changed process. The following table briefly summarizes what your operating procedures must address:

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

OPERATING PROCEDURES REQUIREMENTS

Steps for each operating phase	Operating limits	Safety & health considerations	Safety systems & their functions
<ul style="list-style-type: none"> ✓ Initial start-up ✓ Normal operations ✓ Temporary operations ✓ Emergency shutdown ✓ Normal shutdown ✓ Start-up following a turnaround or emergency shutdown 	<ul style="list-style-type: none"> ✓ Consequences of deviating ✓ Steps to avoid, correct deviations 	<ul style="list-style-type: none"> ✓ Chemical properties & hazards ✓ Precautions for preventing chemical exposure ✓ Control measures for exposure ✓ QC for raw materials and chemical inventory ✓ Special or unique hazards 	<ul style="list-style-type: none"> ✓ Address whatever is applicable
Local requirements (as applicable): <ul style="list-style-type: none"> ✓ Delivery Procedures ✓ Cylinder change out procedures ✓ Release investigation procedures 		<ul style="list-style-type: none"> ✓ Equipment components must be properly identified to avoid operating errors 	

Update the procedures whenever you change your process in a way that alters the steps needed to operate safely. If you add new equipment, you must either expand your procedures or develop a separate set to cover the new items. Whenever you change your safety information you must review your procedures to be sure that they are still appropriate. Anytime you conduct a process hazard analysis, check your operating procedures as you implement changes to address hazards.

Maintain the current set of operating procedures. You are not required to keep old versions; in fact, you should avoid doing so because keeping copies of outdated procedures may cause confusion. You should date all procedures so you will know when they were last updated. Assure that operating procedures are certified annually by an appropriate person.

D. Training

Training programs often provide immediate benefits because trained workers have fewer accidents, damage less equipment, and improve operational efficiency. Training gives workers the information they need to understand how to operate safely and why safe operations are necessary. A training program, including refresher training, is the key to ensuring that the rest of your prevention program is effective.

The following lists things that you may find useful in developing your training program:

- ◆ Clearly identify the employees who need to be trained and the subjects to be covered.
- ◆ Specify learning objectives, and write them in clear, measurable terms before training begins. Remember that training must address the process operating procedures and maintenance procedures.
- ◆ Tailor the specific training modules or segments to the training objectives. Enhance learning by including hands-on training like using simulators whenever appropriate. Make the training environment as much like the working environment as you can, consistent with safety. Allow your employees to practice their skills and demonstrate what they know.
- ◆ Evaluate your training program periodically to see if your employees have the skills and know the routines required under your operating procedures. Make sure that language or presentations are not barriers to learning. Decide how you will measure your employee's competence.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

- ◆ Make sure all workers - including maintenance and contract employees - receive initial and refresher training. If you make changes to process chemicals, equipment, and technology, make sure that involved workers understand the changes and the effects on their job. Careful consideration must be given to assure that employees including maintenance and contract employees receive current and updated training.

Keep documentation of your training program. An attendance log for any formal training courses and refresher training is required to ensure that everyone who needs to be trained is trained. Such logs will help you when you do a compliance audit.

E. Mechanical Integrity

You must have a mechanical integrity program for pressure vessels and storage tanks, piping systems, relief and vent systems and devices, emergency shutdown systems, controls, and pumps. The following table summarizes other requirements of a mechanical integrity program.

MECHANICAL INTEGRITY

Written procedures	Training	Inspection & testing	Equipment deficiencies	Quality assurance
<ul style="list-style-type: none"> ✓ Establish & implement written procedures to maintain the integrity of process equipment 	<ul style="list-style-type: none"> ✓ Train process maintenance employees in an overview of the process and its hazards. ✓ Make sure this training covers the procedures applicable to safe job performance 	<ul style="list-style-type: none"> ✓ Inspect & test process equipment ✓ Use recognized and generally accepted good engineering practices ✓ Follow a schedule that matches the manufacturer's recommendations; industry standards or that prior operating experience indicates is necessary ✓ Document each inspection & test 	<ul style="list-style-type: none"> ✓ Correct equipment deficiencies before further uses of process equipment or whenever necessary to ensure safety 	<ul style="list-style-type: none"> ✓ Establish a QA program for new construction & equipment, newly installed equipment, maintenance materials, and spare parts & equipment

Keep your written procedures and schedules as well as any agreements you have with contractors. You should also keep training logs, inspection & testing logs, and maintenance logs. Without some record, you will have to rely on workers' memories about when something was last checked. As workers leave or change jobs at your company, it can be difficult to keep track of when inspections and tests were done. Maintaining a record of when something was last done or is scheduled to be done next can help keep your program working smoothly.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

F. Management of Change

The following table summarizes the Management of Change requirements:

MANAGEMENT OF CHANGE (MOC) REQUIREMENTS

MOC procedures must address:	Employees affected by the change must:	Update process safety information if:	Update operating procedures if:
<ul style="list-style-type: none"> ✓ Technical basis for the change ✓ Impact on safety and health ✓ Modifications to operating procedures ✓ Necessary time period for the change 	<ul style="list-style-type: none"> ✓ Be informed of the change before startup ✓ Trained in the change before startup 	<ul style="list-style-type: none"> ✓ A change covered by MOC procedures result in a change in any PSI required under EPA's rule 	<ul style="list-style-type: none"> ✓ A change covered by MOC procedures results in a change in any operating procedure required under EPA's rule

To properly manage changes to process chemicals, technology, equipment, and facilities, one must define what is meant by change. In this process safety management standard, change includes all modifications to equipment, procedures, raw materials, and processing conditions other than "replacement in kind." Copies of process changes need to be kept in an accessible location to ensure that design changes are available to operating personnel as well as to PHA team members when a PHA is being done or one is being updated.

G. Pre-Startup Safety Review

You must conduct a pre-startup review before you introduce a regulated substance into a process. The following table lists items you must address.

PRE-STARTUP SAFETY REVIEW REQUIREMENTS

Design Specifications	Adequate Procedures	Training
<ul style="list-style-type: none"> ✓ Confirm that new or modified construction and equipment meet design specifications 	<ul style="list-style-type: none"> ✓ Ensure that procedures for safety, operating, maintenance, and emergencies are adequate and in place 	<ul style="list-style-type: none"> ✓ Confirm that each employee involved in the process has been trained completely

New Processes

The initial startup procedures and normal operating procedures need to be fully evaluated as part of the pre-startup review to assure a safe transfer into the normal operating mode for meeting the process parameters.

Existing Processes

For existing processes that have been shut down for turnaround, or modification, etc., the employer must assure that any changes other than "replacement in kind" made to the process during shutdown go through the management of change procedures. Update P&IDS, operating procedures, and instructions as necessary. If the changes made to the process during shutdown are significant and impact the training program, then operating personnel as well as employees engaged in routine and non-routine work in the process area may need some refresher or additional training in light of the changes. Any incident investigation recommendations, compliance audits or PHA recommendations need to be reviewed as well to see what impacts they may have on the process before beginning the startup.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

F. Compliance Audits

At least every three years, you must certify that you have evaluated compliance with the requirements for the prevention program for each covered process. At least one person on your audit team must be knowledgeable about the covered process. You must develop a report of your findings, determine, and document an appropriate response to each finding, and document that you have corrected any deficiency. Coordinate with the CUPA on timelines for addressing compliance audit recommendations. The CUPA believes that recommendations from compliance audits should be addressed within one (1) year from the audit date. If you need more time, coordinate with the CUPA on a timeline.

You must keep a written record of the findings and actions. Maintain copies of the last two compliance audits.

G. Incident Investigation

You must investigate each incident that resulted in or could have resulted in a "catastrophic release of a regulated substance." A catastrophic release is one that presents an imminent and substantial endangerment to public health and the environment. Although the CalARP requires you to investigate only those incidents that resulted in, or could reasonably have resulted in a catastrophic release, you are encouraged to investigate all accidental releases. Investigating minor accidents or near misses can help you identify problems that could result in major releases if not addressed.

The following is a summary of the steps you must take for investigating an incident:

- ◆ Initiate the investigation promptly. Begin investigating no later than 48 hours following the incident.
- ◆ Establish a knowledgeable investigation team. Establish an investigation team to gather the facts, analyze the event, and develop the "how" and "why" of what went wrong. At least one team member must have knowledge of the process. Consider adding other workers familiar with the process to the incident team. Their additional knowledge will assist in the fullest insight into the incident.
- ◆ Summarize the investigation in report. Among other things, this report will include the factors contributing to the incident. Remember that identifying the root cause may be more important than identifying the initiating event. Also, remember that the purpose of the report is to help management take corrective action.
- ◆ Address the team's findings and recommendations. Establish a system to address the incident report findings and recommendations and document resolutions and corrective actions.
- ◆ Review the report with your staff and contractors. You must share the report –its findings and recommendations- with affected workers whose job tasks are relevant to the incident.
- ◆ Retain the report. Keep incident investigation reports for five years.

H. Employee Participation

Section 2760.10 in Title 19 of the California Code of Regulations states that employers are to consult with their employees and their representatives regarding the employer's efforts in the development and implementation of the process safety management program elements and hazard assessments. Many employers, under their safety and health programs, have already established means and methods to keep employees and their representatives informed about relevant safety and health issues and employers may be able to adapt these practices and procedures to meet their obligations under this section. Employers who have not implemented an occupational safety and health program may wish to form a safety and health committee of

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

employees and management representatives to help the employer meet the obligations specified by this standard. This committee can become a significant ally in helping the employer to implement and maintain an effective process safety management program for all employees.

The following is a summary of what you must do:

- **Write a plan.** Develop a written plan of action regarding how you will implement employee participation.
- **Consult with employees.** Consult your employees and their representatives regarding conducting and developing PHAs and other elements of process safety management and the risk management program.
- **Provide access to information.** Ensure that your employees and their representatives have access to PHAs and all other information required under the CalARP.

I. Hot Work Permits

Non-routine work that is conducted in process areas needs to be controlled by the employer in a consistent manner. The hazards identified involving the work that is to be accomplished must be communicated to those doing the work, but also to those operating personnel whose work could affect the safety of the process. A work authorization notice, or permit must have a procedure that describes the steps the maintenance supervisor, contractor representative or other person needs to follow to obtain the necessary clearance to get the job started. The work authorization procedures need to reference and coordinate, as applicable, lockout/tag out procedures, line breaking procedures, confined space entry procedures and hot work authorizations. This procedure also needs to provide clear steps to follow once the job is completed. These steps must provide closure for those that need to know the job is now completed and equipment can be returned to normal.

The following summarizes how to meet the hot work permit requirements:

Issue a hot work permit. You must issue this permit for hot work conducted on or near a covered process

- ◆ **Implement fire prevention and protection.** You must ensure that the fire prevention and protection requirements in 8 CCR 5189 are implemented before the hot work begins. The permit must document this.
- ◆ **Indicate the appropriate dates.** The permit must indicate the dates authorized for hot work.
- ◆ **Identify the work.** The permit must identify the object on which hot work is to be performed.
- ◆ **Maintain the permit on file.** You must keep the permit on file until workers have completed the hot work operations.

J. Contractors

Employers, who use contractors to perform work in and around processes that involve highly hazardous chemicals, will need to establish a screening process so that they hire and use contractors who accomplish the desired job tasks without compromising the safety and health of employees at a facility. For contractors, whose safety performance on the job is not known to the hiring employer, the employer will need to obtain information on injury and illness rates and experience and should obtain contractor references. Additionally, the employer must assure that the contractor has the appropriate job skills, knowledge, and certifications (such as for pressure vessel welders). Contractor work methods and experiences should be evaluated.

Contract employees must perform their work safely. Considering that contractors often perform

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

very specialized and potentially hazardous tasks such as confined space entry activities and non-routine repair activities it is quite important that their activities be controlled while they are working on or near a covered process. A permit system or work authorization system for these activities would also be helpful to all affected employers. The use of a work authorization system keeps an employer informed of contract employee activities, and as a benefit the employer will have better coordination and more management control over the work being performed in the process area. A well run and well-maintained process where employee safety is fully recognized will benefit all of those who work in the facility whether they be contract employees or employees of the owner.

While HMD has no direct authority to require that you maintain an occupational injury and illness log for contract employees. However, OSHA does have this authority, and the PSM standard sets this requirement. Therefore, maintaining an occupational injury and illness log for contract employees is a recognized standard industry practice.

The following table summarizes both your and the contractors' responsibilities:

You must:	Your contractor must:
<ul style="list-style-type: none"> ✓ Check safety performance. You must evaluate the safety performance of the contractor ✓ Provide safety and hazard information. You must inform the contractor of potential fire, explosion or toxic release hazards; and of your emergency response activities as they relate to the contractor's work and the process. ✓ Ensure safe practices. You must assure that you have safe work practices such as controlling the entrance, presence, and exit of contract employees in covered process areas. ✓ Verify that the contractor acts responsibly. You must verify that the contractor is fulfilling its responsibility to provide appropriate health, safety, and craft training. 	<ul style="list-style-type: none"> ✓ Ensure training for its employees. The contractor must train and supervise contract employees to ensure that they perform their jobs safely and in accordance with your source's safety procedures. ✓ Ensure its employees know process hazards and applicable emergency actions. The contractor must assure that contract employees are aware of hazards and emergency procedures relating to the employees' work ✓ Document training. The contractor must prepare a record documenting and verifying adequate employee training. ✓ Inform you of hazards. The contractor must tell you of any unique hazards presented by its work or of any hazards it finds during performance.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

IV. EXTERNAL EVENTS INCLUDING SEISMIC ASSESSMENT

For Program 2 and Program 3 processes you must evaluate, as part of your Process Hazardous Analysis (PHA) or Hazard Review, any potential natural or human caused external events. At a minimum a seismic assessment must be performed. The following are guidelines to use when conducting a seismic assessment.

A. Seismic Assessment General Requirements

The primary purpose of a seismic assessment is to determine types of regulated substance releases that might occur due to an earthquake. These guidelines have been developed for businesses that have a relatively simple process. In general, the approach toward seismic analysis should be qualitative unless findings indicate the need for further evaluation by an experienced structural engineer.

The evaluation should be geared towards finding, evaluating, and, if necessary, strengthening governing elements in the process system. Emphasis should be placed on pipes and hoses. Visually inspect the piping system of concern to evaluate flexibility, support, and guide adequate reinforcement. Many of the failures can be traced to one of the following causes:

- ◆ Lack of flexibility between piping anchor points. In earthquakes, vessel and piping anchor points can grossly displace relative to each other.
- ◆ Branch lines do not have adequate flexibility to accommodate seismic movement of the main line.
- ◆ Lack of adequate piping guides or lateral restraints allows a pipe to slide off its supports.
- ◆ Lack of reinforcement between branch lines and the main header.

B. Seismic Assessment Resource

The HMD recommends that facilities conducting a seismic assessment refer to the *Guidance For California Accidental Release Prevention (CalARP) Program Seismic Assessments*. This document was prepared for the UPA REGION I LOCAL EMERGENCY PLANNING COMMITTEE (LEPC). This document was last updated January 2019 and is located on the CUPA website.

This manual can be obtained at the following web page:

<https://www.sandiegocounty.gov/content/dam/sdc/deh/hmd/pdf/CalARPSismicAssessments.pdf>

This manual can be used by your business to evaluate the potential of an earthquake to cause accidental releases of regulated substances and to assist your business in determining appropriate mitigation measures to prevent and/or minimize those potential accidental releases.

C. External Event Documentation

The following external event records shall be maintained in a Technical Document. This Technical Document will be subject to submittal upon HMD's request and/or onsite auditing by HMD:

1. Include a table of contents.
2. Place dividers and tabs between the sections of the document.
3. Describe the types of external events (including seismic) that were evaluated and any potential release of regulated substance that likely could occur; emergency mitigation

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

systems and measures in place; and recommended mitigation or measures that will be implemented to reduce the likelihood of a release.

4. For the seismic analysis list:

- Any buildings and structures that were evaluated.
- All the nonstructural components (i.e., piping, tanks, etc.) that were evaluated.
- Any potential regulated substance releases.
- The emergency mitigation systems and measures in place to prevent a release of a regulated substance should an earthquake occur.
- Any recommended mitigation systems or measures that will be implemented as a result of the seismic analysis.
- List the edition of the Uniform Building Code that was used when the process was designed.

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

V. EMERGENCY RESPONSE PLANS

Responding agencies must prepare a written emergency response plan. This plan should explain how accidental releases are handled and actions to take during an external event that can affect a covered process or any other operations at a stationary source. The plan should be detailed enough to provide employees with the proper actions to take during an emergency but allow flexibility to respond in the most appropriate way. An emergency response plan must contain the following:

- Procedures for interfacing with the public and local emergency response agencies about accidental releases, emergency planning, and emergency response
- Documentation of proper first aid and emergency medical treatment necessary to treat accidental human exposures. This can be found in the Safety Data Sheet.
- Procedures for the use of emergency response equipment and for its inspection, testing, and maintenance
- Training for all employees in relevant aspects of the Incident Command System (ICS).
- Procedures to review and update, as appropriate, the emergency response plan to reflect changes at the stationary source and ensure that employees are informed of changes.

If you need any clarification on this guidance document or the CalARP Program, call the County of San Diego and ask for the CalARP Program Manager at 858-505-6880.

APPENDICIES

RMP (PUBLIC DOCUMENT) DEVELOPMENT AND SUBMITTAL GUIDELINES

Appendix A -1
CalARP Registration / Work Plan
HMD Form HM-9240



SAN DIEGO COUNTY CUPA
DEPARTMENT OF ENVIRONMENTAL HEALTH AND QUALITY
HAZARDOUS MATERIALS DIVISION
P.O. BOX 129261, SAN DIEGO, CA 92112-9261
(858) 505-6880 FAX (858) 505-6848
<http://www.sdcdelh.org>

CalARP RMP REGISTRATION 2740.1(d) / WORK PLAN

I. STATIONARY SOURCE INFORMATION

NAME OF STATIONARY SOURCE		UNIFIED PROGRAM FACILITY PERMIT NUMBER	
SITE ADDRESS		CITY	CA ZIP CODE
LATITUDE AND LONGITUDE & METHOD FOR OBTAINING THIS			NAICS CODE
STATIONARY SOURCE USEPA IDENTIFIER	STATIONARY SOURCE DUN & BRADSTREET #	# FULLTIME EMPLOYEES	
CORPORATE/PARENT COMPANY NAME		CORPORATE/PARENT DUN & BRADSTREET #	
WEBSITE			

II. STATIONARY SOURCE OWNER/OPERATOR & RMP CONTACT INFORMATION

NAME OF OWNER/OPERATOR		OWNER/OPERATOR PHONE#	
NAME OF RMP CONTACT	TITLE OF RMP CONTACT	PHONE # RMP CONTACT	
MAILING ADDRESS FOR RMP CONTACT	E-MAIL OF RMP CONTACT	CELL # RMP CONTACT	
CITY RMP CONTACT	STATE	ZIP CODE RMP CONTACT	
24-HR EMERGENCY CONTACT PRIMARY	TITLE	24 HR PHONE #	E-MAIL
24-HR EMERGENCY CONTACT ALTERNATE	TITLE	24 HR PHONE #	E-MAIL

III. RMP CONSULTANT CONTACT INFORMATION (IF APPLICABLE)

COMPANY NAME	CONSULTANT'S NAME	CONSULTANT'S PHONE #
CONSULTANT'S ADDRESS	CONSULTANT'S E-MAIL	CONSULTANT'S CELL #
CONSULTANT'S CITY	STATE	CONSULTANT'S ZIP CODE

IV. PROCESS INFORMATION

NAME OF REGULATED SUBSTANCE (one sheet per item)	CAS NUMBER	MAX QUANTITY (in Lbs.)	RMP PROGRAM LEVEL <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3
IS PROCESS SUBJECT TO PSM TITLE 8 SEC. 5189? <input type="checkbox"/> YES <input type="checkbox"/> NO	IS SS SUBJECT TO PART 355 OF TITLE 40 CFR? <input type="checkbox"/> YES <input type="checkbox"/> NO	IS PROCESS SUBJECT TO TITLE V PERMIT? <input type="checkbox"/> YES <input type="checkbox"/> NO	
PROCESS INSTAL/MODIFICATION DATE: _____ / _____ / _____	LAST SAFETY INSPECTION BY: <input type="checkbox"/> FEDERAL <input type="checkbox"/> STATE <input type="checkbox"/> LOCAL DATE OF INSPECTION: _____ / _____ / _____ Specify:		

V. RMP TECHNICAL STUDIES

METHODS FOR AIR DISPERSION MODELING: Manual Calculation (Tables) <input type="checkbox"/> YES <input type="checkbox"/> NO or Computerized Air Model <input type="checkbox"/> YES <input type="checkbox"/> NO	TYPE OF PHA CONDUCTED:	DATE OF SEISMIC WALKTHROUGH: _____ / _____ / _____
NAME OF COMPUTERIZED AIR MODEL (IF APPLICABLE):		
PASSIVE MITIGATION CONSIDERED FOR WORST-CASE SCENARIO (SPECIFY):		
REASON FOR RMP CHANGE: <input type="checkbox"/> 5-year update <input type="checkbox"/> corrections (see back) <input type="checkbox"/> new stationary source <input type="checkbox"/> modified stationary source		
CERTIFIED BY:	PRINTED NAME:	DATE _____ / _____ / _____

CALIFORNIA ACCIDENTAL RELEASE PREVENTION (CalARP) PROGRAM RISK MANAGEMENT PLAN (RMP) REGISTRATION / WORK PLAN - INSTRUCTIONS

I. STATIONARY SOURCE INFORMATION

NAME OF STATIONARY SOURCE- Enter the full legal name of the business. This is the same as the term "Facility Name" or "DBA" – Doing Business As.
STATIONARY SOURCE UPF PERMIT NUMBER- Enter the Unified Program Facility Permit (UPFP) number.
STATIONARY SOURCE SITE ADDRESS- This is the site (physical) address of the stationary source.
LATITUDE AND LONGITUDE- Enter the latitude and longitude of the stationary source & the method used for obtaining the latitude and longitude.
NAICS CODE- Enter the 6-digit North American Industry Classification System (NAICS) code.
STATIONARY SOURCE USEPA IDENTIFIER- Enter the stationary source USEPA ID number, if applicable
STATIONARY SOURCE DUN AND BRADSTREET (D&B) #- Enter the stationary source Dun and Bradstreet number
CORPORATE/PARENT COMPANY NAME- Enter the name of parent or corporate owner.
OF FULL-TIME EMPLOYEES- Enter the number of full-time employees at the stationary source.
CORPORATE/PARENT COMPANY D&B#- Enter the Dun and Bradstreet number of the parent or corporate owner.
WEBSITE- Web address (URL) for the corporate/parent organization.

II. STATIONARY SOURCE RMP CONTACT, OWNER/OPERATOR INFORMATION

NAME & PHONE# OF OWNER/OPERATOR- Enter the name and phone number of the owner or operator of the stationary source
NAME OF RMP CONTACT- Enter the name of the person with the overall responsibility for RMP elements, integration, and implementation. This is the primary RMP contact for the stationary source.
TITLE OF RMP CONTACT- Enter the title of the person with the overall responsibility for RMP elements and implementation.
MAILING ADDRESS FOR RMP CONTACT- Enter the primary RMP contact's mailing address.
PHONE# OF RMP CONTACT- Enter the phone number for the primary RMP contact.
CELL PHONE# OF RMP CONTACT- Enter a cell phone number for the primary RMP contact.
E-MAIL OF RMP CONTACT- Enter the e-mail address for the primary RMP contact.
24-HOUR EMERGENCY CONTACT PRIMARY- Enter the name of a primary person available for RMP-related emergencies 24 hours a day.
TITLE- Title of the 24-hour primary RMP emergency contact person.
PHONE#- Enter the 24-hour phone number for the primary 24-hour RMP emergency contact person.
E-MAIL- Enter the e-mail address for the primary RMP emergency contact person.
24-HOUR EMERGENCY CONTACT ALTERANTE- Enter the name of an alternate person available for RMP-related emergencies 24 hours a day.
TITLE- Title of the 24-hour alternate RMP emergency contact person.
PHONE#- Enter the 24-hour phone number for the alternate 24-hour RMP emergency contact person.
E-MAIL- Enter the e-mail address for the alternate 24-hour RMP emergency contact person.

III. RMP CONSULTANT CONTACT INFORMATION, IF APPLICABLE

COMPANY NAME- Enter the company name of the RMP consultant or project coordinator, if applicable.
CONSULTANT'S NAME- Enter the name of the primary RMP consultant or project coordinator.
CONSULTANT'S ADDRESS INCLUDING CITY, STATE and ZIP CODE- Enter the complete address of the RMP consultant or project coordinator.
CONSULTANT'S PHONE#- Enter the phone number of the RMP consultant or project coordinator.
CONSULTANT'S CELL PHONE#- Enter the cell phone number of the RMP consultant or project coordinator.
CONSULTANT'S E-MAIL- Enter the e-mail address of the primary RMP consultant or project coordinator.

IV. PROCESS INFORMATION

NAME OF REGULATED SUBSTANCE- Enter the name, concentration & quantity (in pounds) of the regulated substance in the process. One registration per process.
CAS NUMBER- Enter the chemical abstract service number for the regulated substance above.
RMP PROGRAM LEVEL- Mark the program level for the RMP.
PROCESS SUBJECT TO PSM- Mark yes if the process is subject to OSHA Process Safety Management (PSM) Title 8 Section 5189, or no if not subject to PSM.
IS SS SUBJECT TO PART 355 OF TITLE 40 OF CFR- Mark yes if the stationary source is subject to chemical disclosure under title 40 CFR Part 355 or mark no if quantity onsite is below federal threshold planning quantities.
PROCESS SUBJECT TO TITLE V PERMIT- Mark yes if the process is subject to the Title V CAA permit requirements, or no if not subject to Title V.
DATE OF SEISMIC WALKTHROUGH- Enter the date of the most recent seismic walkthrough.
PROCESS INSTALLATION DATE- Enter the process installation date. If adding a new process or modifying an existing process, enter the planned startup date for the new or modified process.
DATE OF THE LAST SAFETY INSPECTION- Enter the date and the name of the agency that last performed a safety inspection of the stationary source. Mark if the safety inspection was performed by a federal, state, or local agency

V. RMP TECHNICAL STUDIES

METHODS OF AIR DISPERSION MODELING: MANUAL CALCULATIONS (TABLES)- If you used [will use] manual calculations or EPA look-up tables for your offsite consequence analysis check yes or no, as applicable.
TYPE OF PROCESS HAZARD ANALYSIS [TO BE] CONDUCTED- Enter the name of the type of Process Hazard Analysis (PHA) or Hazard Review [to be] conducted, i.e., what-if/checklist, HAZOP, etc. For Program 1 mark "none."
COMPUTERIZED AIR MODEL- If you used a computerized air model for your offsite consequence analyses check yes, if not, check no. If you used a computer air model Enter the name and version.
PASSIVE MITIGATION FOR WORST-CASE (SPECIFY)- Specify the type of passive mitigation you used [plan to use] for the worst-case offsite consequence. if you did not use [or do not plan to use] passive mitigation state "none".
MARK THE REASON FOR RMP CHANGE- Mark the reason for the RMP change i.e., 5-year update, correction, new or modified stationary source. Updates and re-submissions required under Section 2745.10(a) or (b); or corrections under Section 2745.10.5 or for purposes of correcting minor clerical errors, updating administrative information, providing missing data elements, or reflecting facility ownership changes, and which do not require an update and re-submission.
CERTIFIED BY- This line must be signed by the person certifying that the information provided is true and accurate.
PRINTED NAME- Print the name of the person certifying that the information provided is true and accurate.
DATE- Enter the date the registration or work plan was certified and signed.

A-2 Useful Web Links

Federal Industry RMP Guidance: A list of RMP guidance documents for specific businesses.

<http://www2.epa.gov/rmp/guidance-facilities-risk-management-programs-rmp>

RMP Submit link

<http://www2.epa.gov/rmp/rmpsubmit>

California RMP Guidance: California State guidance on the CalARP program.

[http://www.caloes.ca.gov/HazardousMaterials/Pages/California-Accidental-Release-Prevention-\(CalARP\).aspx](http://www.caloes.ca.gov/HazardousMaterials/Pages/California-Accidental-Release-Prevention-(CalARP).aspx)

Seismic Assessment Guidance

[Guidance for CalARP Program Seismic Assessments Updated January 2019](#)

Marplot Mapping. Free mapping software from the U.S. EPA.

<http://www2.epa.gov/cameo/cameo-downloading-marplot>

EPA RMP Comp Free software for calculating affected areas during release scenarios.

<http://www2.epa.gov/rmp/rmpcomp>

U.S. Census Bureau. Data for estimating populations.

<http://www.census.gov/>

A-3 Sample Incident Investigation

Anhydrous Ammonia Tank Release

Date: XX/XX/XXXX at 08:35

Substance: Anhydrous ammonia

Quantity: 350 lbs

Duration: 1 hour

Weather: Overcast, cool, approximately 55 degrees, light wind from the west at about 5 mph.

Description: During a delivery of anhydrous ammonia, the product delivery hose split and released ammonia. The delivery agent and all facility employees were away from the tank for at least 35 minutes when the release occurred. The ammonia pooled around the tank then evaporated. The ammonia spread throughout the outside of the facility and drifted to the adjacent lot, where outside workers at XXX Company were exposed to ammonia gas. The XYZ Company called 911 and sent an employee to investigate where the ammonia was coming from and notified our business of the outside ammonia release. We immediately implemented our Emergency Response Plan. All ignition sources were shut off and we called 911 to notify emergency responders of the release. The Emergency Response Team was called and they began to prepare for a response. After determining that they could approach the truck from up wind and not be exposed to *over* 250 ppm as measured on a Dräger colorimetric tube, our response team, wearing protective clothing and proper respiratory protection, shut off the delivery valve on the ammonia truck within 15 minutes. The County Hazardous Incident Response Team (HIRT) arrived and ordered our facility and XYZ Company to evacuate upwind of the ammonia spill while the remaining ammonia evaporated into the atmosphere.

1. **Findings:** After mitigating the release, we investigated the causes. A 2-inch delivery hose had burst during delivery causing the release. Upon closer inspection the hose was old and deteriorated. The hose was not inspected prior to use. Also, no operators or employees from our company stayed with the truck during delivery. This allowed the release to continue for a longer period of time than if an attendant were present.
2. **Recommendations:** The delivery hose needs to be inspected and replaced at regular intervals to prevent future releases. Also, higher-pressure hoses are available and should be used. Operational procedures need to be changed so that the ammonia tank and delivery truck are never left unattended during ammonia deliveries. The Emergency Response Team should always be put on alert and secure their personal equipment when ammonia deliveries occur. Investigate installing an ammonia sensor near the ammonia tank.
3. **Actions:** Replace the damaged delivery hose with a higher-pressure hose. Assure that the hose is inspected before each weekly delivery. Have one member of the Emergency Response Team put the other members on alert before the delivery begins. Assign one member of the Team to be present during the entire delivery process. Conduct emergency response refresher training with all employees. Install an ammonia sensor to warn of any ammonia leaks near the ammonia tank by XX/XX/XXXX.

A-4
Sample Audit Checklist for Safety Information

Element	Yes/No/NA	Action/Completion Date
SDSs up to date?		
Maximum Inventory determined?		
Determine		
• Safe upper and lower temperature for materials		
• Safe upper and lower pressures		
• Safe process flow rates		
• Compatible equipment with the materials used?		
Storage Tank specifications met?		
Pressure relief valves functioning?		
Emergency shut off valves present and working?		
Gauges working?		
Pumps working and serviced per manufacture's recommendations?		
Compressors serviced and functioning?		
Hoses inspected and in good repair?		
All equipment install to manufacturer's specifications and industry standards?		
Have inspections all been documented?		
Have inspections been conducted after each major change?		

A-5 Public Review of Risk Management Plans

Federal Requirements

Stationary sources that are required to complete a Risk Management Plan (RMP) must submit their RMP to both the USEPA and the San Diego County Department of Environmental Health & Quality, Hazardous Materials Division (HMD).

NOTE: Stationary sources that only meet the State threshold planning quantities are not required to submit their RMP to the USEPA.

California Requirements

California Health and Safety Code Section 25535.2 requires the HMD to make your RMP available to the public for review within 15 days after determining an initial RMP submittal to be complete. The public review comment period is for 45 days. Once an RMP is determined to be complete by the HMD, the CUPA e-mails the local Fire Marshal and posts on its "[public notice website](#)" for 45 days stating that the RMP Public Document is available for public review.

All comments received by the HMD during the public comment period will be reviewed and considered prior to the HMD's final review of the RMP. The HMD will notify the stationary source of any deficiencies after the public review period. The stationary source will then have 30 days to correct any deficiencies noted by the HMD.

Availability of Information to the Public (19 CCR 2775.5)

- (a) The RMP required under Article 3 of this chapter shall be available to the public pursuant to Section 25534.05(a)(4) of HSC, except for offsite consequence analysis data, pursuant to (b).
- (b) The UPA shall insure that any member of the public has access, by appointment, to a copy of the offsite consequence analysis data, pursuant to Section 2745.4. The member of the public may read, but not remove, reproduce, print, scan or image the documents. The UPA may require personal photo identification issued by a Federal, State or local government agency to the person, and may require the person's signature on a sign-in sheet. The UPA may limit a person's access to offsite consequence analysis data to 10 stationary sources in any calendar month.

RMP Updates

When a 5-year RMP update is submitted to the HMD, the RMP is once again determined to be complete. The CUPA e-mails the local Fire Marshal and posts on its "[public notice website](#)" for 45 days stating that the RMP Public Document is available for public review.

ISSUE SUMMARY: Seismic Assessment Above Ground Storage Tanks

Problem:

- Cylindrical tanks are vulnerable to several types of failure in earthquakes.
- Leaks can occur due to the failure of internal baffles.
- The top of the outside tank wall can be damaged by battering from the floating roof.
- "Elephant's foot" buckling failures at the base of tanks are caused by horizontal forces and can result in the complete loss of contents.
- Sliding is common for unanchored tanks. If there is enough flexibility in the connecting pipelines, such movement can occur without any loss of contents. However, attached pipelines often break. Even anchored tanks can move, but such movement is usually minor and rarely results in loss of contents.
- Corrosion at the base of tanks can also be a problem. In particular, "pitting corrosion" at the tank base, combined with earthquake forces, can cause the tanks to fail, losing their contents.
- Tank walls can be damaged due to inadequate detailing at connections with external pipes, valves and ladders, and due to improper welding.
- Finally, elevated tanks can topple if inadequately supported. Elevated tanks typically have more performance problems in earthquakes than ground-mounted structures.

Mitigation Options:

Increasing the thickness of tank walls at their base can be used to help prevent "elephant's foot" buckling. However, thick base walls can simply create buckling in the next higher tier of metal if that section is inadequate. Also, the upper sections of the walls can be thickened or otherwise strengthened to accommodate the forces caused by sloshing or the impact of the floating roof.

The factor of safety use in the design of tank walls containing hazardous materials is greater than for those containing water. In some instances, the safety factor for water systems may need to be increased, such as where water may be critical in an emergency or where excessive amounts of water spraying in an area could impede access to critical areas.

Tank/foundation connections should be carefully designed, particularly if associated piping is subject to failure if the tank moves. Often, the use of a larger number of smaller anchor bolts are preferable to fewer larger bolts. Additional coating can be used to help bond fiber-reinforced plastic tanks to their supports to prevent the rolling and shifting of the tank.

Particular attention should be paid to the detailing of connections with external pipes, valves and ladders, for these are frequently weak points. Welds should be inspected to make sure that weld quality and penetration are sufficient. A specific instance of concern is when piping attached to unanchored tank lacks flexibility, such as when it exits the tank and goes directly into the ground or through a wall.

Elevated tank supports can be inadequate because of the tendency to stop the leg at the tank base. Often, it is preferable to extend the support up the tank wall a foot or two and put a ring of additional material for added strength at the tank base. Adequate diagonal cross bracing should also be used on

supports for these elevated tanks. If possible, such tanks should be redesigned to function without being elevated.

In addition, elevated tanks may be supported on reinforced concrete frame structures with no redundancy. These structures may be substandard, especially in the ductile detailing of the connections, and pose a significant risk of catastrophic collapse in a -large earthquake. A qualified structural engineer should evaluate this type of support structure. Retrofits may include adding steel bracing.

The possibility of tall or elevated tanks falling on adjacent buildings and equipment should be a part of the decision on the design and location of those tanks.

Finally, tanks should be adequately maintained to detect corrosion and other tank deterioration before an earthquake exacerbates any problem. Particular care should be used in inspecting the bases of tanks.

A-8

WHAT-IF/CHECKLIST FORM

Analysis Team: _____ Date of Analysis: _____

Area of Investigation: _____ Page: _____

Scenario No.	Checklist No. OR Drawing No.	What-if?	Consequences	Likelihood	Existing Controls	Recommendation

Trade Secrets/Confidential Information

Trade Secret

If a business believes that information required or requested involves the release of a trade secret, the business shall provide the Hazardous Materials Division (HMD) with a notification in writing that the information is considered a trade secret. Upon receipt of a claim of a trade secret related to the Risk Management Plan (RMP), the HMD will review the claim and will segregate properly substantiated trade secret information from other information that is otherwise disclosable to the public upon request in accordance with the California Public Records Act (Chapter 3.5 commencing with Section 6250, Division 7, Title 1 of the Government Code).

Trade secret is defined in subdivision (d) of Section 6254.7 of the Government Code and Section 1060 of the Evidence Code.

The HMD may disclose trade secrets to authorized officers or employees of other governmental agencies only in connection with the official duties of those officers or employees pursuant to any law for the protection of health and safety.

Confidential Information

Information that identifies where hazardous materials are used, handled, or stored at a facility, such as site maps, is considered confidential. It is the business's responsibility to ensure that such information is identified within the RMP document and is marked confidential.

A-10

Quick Reference Worst Case. Release Scenario Requirements

Type of Chemical	Assume Time for Total Release	Release Rate (Pounds/minute)
Toxic gases at ambient temperature (handled as a gas or as a liquid under pressure)	Quantity in the vessel or pipe is released as a gas over 10 minutes.	If no passive mitigation systems are in place, total quantity released divided by 10.
		If passive mitigation systems are in place, total quantity released divided by 10, then multiplied by 0.55 (mitigation factor). ¹
Toxic gases at ambient pressure (handled as refrigerated liquids)	If no passive mitigation or if the contained pool would <i>have</i> a depth of 1 cm or less: released as a gas in 10 minutes.	Total quantity released divided by 10.
	If contained by passive mitigation in a pool with a depth greater than 1 cm: assume the quantity in the vessel or pipe is spilled instantaneously to form a liquid pool.	The volatilization rate (release rate) shall be calculated at the boiling point of the substance and at the conditions specified in "Toxic liquids" below.
Toxic liquids at ambient temperature	Assume that the quantity is spilled instantaneously to form a liquid pool. <ul style="list-style-type: none"> • Undiked: Pool will spread until it is 1 cm deep. • Diked (passive mitigation): Pool will have surface area defined by the area within the dike. 	Calculated by a model that includes volatilization rate, surface area, maximum temperature and concentration of the chemical in the pool, and the surface characteristics of the substrate underneath the spill.
Flammables (liquids or gases)	Make appropriate assumptions based on facility conditions. Consider both <i>active</i> and passive mitigation systems. ^{2,3}	
Solids	Assume one-hour release.	Use USEPA, OES or California Air Resources Board approved model. (Currently OES has not identified an air dispersion model for solids. The AA may want to confer with the local air quality management district or air pollution control district on appropriate air dispersion modeling.)

From the California Office of Emergency Services CalARP Program AA Guidance, January 31, 2005

¹ USEPA's Off-site Consequence Analysis Guidance Document, April 1999, Section 3.1.2

² Ibid, Section 1.5.3

³ USEPA's General Guidance for Risk Management Programs, Chapter 4, Section 4-9

A-11

Alternative Release Scenario Analysis (Section 2750.4)

The facility must identify at least one alternative release scenario for each toxic chemical and one alternative release scenario for all flammable chemicals.

Each selected alternative release scenario must:

- Be more likely to occur than the worst-case release scenario above, and
- Potentially reach an endpoint offsite, unless no such scenario exists.

Potential alternative release scenarios might include:

- Transfer hose releases;
- Process piping releases;
- Process vessel or pump releases;
- Vessel overfilling and spill, or vessel over-pressurization and venting through relief valves or rupture disks; or,
- Shipping container mishandling; breakage or puncture leading to a spill.

Active and passive mitigation systems may be considered if they can withstand the event that triggered the release and remain functional.

The facility must consider the following in selecting alternative release scenarios:

- The five-year accident history required by Section 2750.9; and
- Failure scenarios identified under Section 2755.2 or 2760.2.

From the California State Office of Emergency Services CalARP Program AA Guidance, January 31, 2005.

A-12 Fees

The County of San Diego Department of Environmental Health & Quality, Hazardous Materials Division (HMD):

California Code of Regulations, Title 19, Division 2, Chapter 4.5, Article 3 requires the Owners and Operators of a Stationary Source to coordinate with the Certified Unified Program Agency (CUPA)/Administering Agency (AA) in the development and implementation of Risk Management Plans/Programs.

Pursuant to the San Diego County Code of Regulatory Ordinances Title 6, Division 5, Section 65.107 (k)(13) & (14), a business is charged an hourly rate for all time involving consultation and Risk Management Plan (RMP) review.

The HMD estimated the RMP review process to involve between 25 and 100 hours. The actual time involved will depend on the scope and complexity of the project. Submitting accurate, well written RMP documents following the guidance document will expedite the review process and consequently save the facility money.

Businesses are billed approximately quarterly or at a case closure, whichever comes first. Additionally, hourly charges are used to cover the costs incurred by the HMD in carrying out other elements of the California Accidental Release Prevention (CalARP) Program such as participation in Process Hazard Analysis and Hazard Review Studies.

All Program 1, 2 or 3 stationary sources are subject to an annual permit fee along with the other HMD fees.

California Environmental Protection Agency (CalEPA):

The CUPA will add the CalEPA CalARP service fee to each stationary source subject to the CalARP Program. Service fees are required by the State and collected by the Certified Unified Program Agencies (CUPAs) and submitted to the CalEPA. The DEHQ is the CUPA for San Diego County.

Questions

If you have questions concerning the Risk Management Program fee policy, contact the County of San Diego, ask for the CalARP Program Manager at 858-505-6880.

A-13

SAMPLE OCA Data Table		
	Worst Case Scenario	Alternative Case Scenario
Location: 33.XXXXXXX °N, -117.XXXXXXX °W		
Input		
Total Quantity Held Onsite (include physical state and concentration where applicable):		
Total Release Quantity		
Release Rate - total		
Release Rate to Atmosphere		
Release Time		
Release Direction		
Release Temperature (Daily Max. Temp. - WC)		
Release Pressure		
Ambient Temperature - Daily Max. Temp.		
Ambient Pressure		
Relative Humidity		
Stability Class		
Wind Speed		
Surface Roughness (Urban or Rural)		
Averaging Time		
Level of Concern		
Output		
Estimated Population Affected (CAMEO-MARPLOT Version X.X.X Mapping Software)		
Population Reported to CalARP/EPA Reporting (2 Significant Figures)		
Release Rate - EPA Guidance Document		
Distance at Endpoint: General Guidance on Risk Management Programs for Chemical Accident Prevention		