

Chapter NR 465

NATIONAL EMISSION STANDARDS FOR HAZARDOUS AIR POLLUTANTS FOR WOOD FURNITURE MANUFACTURING OPERATIONS

NR 465.01	Applicability; purpose.	NR 465.06	Compliance methods and procedures.
NR 465.02	Definitions.	NR 465.07	Initial compliance demonstration.
NR 465.03	Symbols.	NR 465.08	Continuous compliance demonstration.
NR 465.04	Emission limits.	NR 465.09	Performance test methods.
NR 465.05	Work practice standards.	NR 465.10	Recordkeeping requirements.
NR 465.055	Compliance dates.	NR 465.11	Reporting requirements.

NR 465.01 Applicability; purpose. (1) APPLICABILITY.

(a) This chapter applies to each facility that is engaged, either in part or in whole, in the manufacture of wood furniture or wood furniture components and that is located at a plant site that is a major source of hazardous air pollutants.

(b) An incidental wood furniture manufacturer shall maintain purchase or usage records demonstrating the source meets the criteria specified in s. NR 465.02(33), but the source is not subject to any other provisions of this chapter.

(c) A source that qualifies as an area source under this paragraph is not subject to any provisions of this chapter other than those in this paragraph. For subs. 1. and 2., finishing materials, adhesives, cleaning solvents and washoff solvents used for wood furniture or wood furniture component manufacturing operations shall account for at least 90% of annual HAP emissions at the plant site, and if the plant site has HAP emissions that do not originate from the listed materials, the owner or operator shall keep any records necessary to demonstrate that the 90% criterion is being met. A source that initially relies on the limits and criteria specified in subd. 1., 2. or 3. to become an area source, but subsequently exceeds the relevant limit, without first obtaining and complying with other limits that keep its potential to emit hazardous air pollutants below major source levels, becomes a major source and shall comply thereafter with all applicable provisions of this chapter starting on the applicable compliance date in s. NR 465.055. Nothing in this paragraph is intended to preclude a source from limiting its potential to emit through other appropriate mechanisms. A source qualifies as an area source for the purposes of this chapter if the criteria in one of the following subdivisions are met:

1. The owner or operator of the source uses no more than a total of 250 gallons per month, for every month, of coating, adhesive, cleaning material and washoff materials at the source, including materials used for source categories other than wood furniture, but excluding materials used in routine janitorial or facility grounds maintenance, personal uses by employees or other persons, the use of products for the purpose of maintaining motor vehicles operated by the facility, the use of toxic chemicals contained in intake water used for processing or noncontact cooling and intake air used either as compressed air or for combustion. The owner or operator shall maintain records for 5 years of the total gallons of coating, adhesive, cleaning material and washoff material used each month, and upon request submit the records to the department.

2. The owner or operator of the source uses no more than 3,000 gallons per rolling 12-month period, for every 12-month period, of coating, adhesive, cleaning material and washoff material at the source, including materials used for source categories other than wood furniture, but excluding materials used in routine janitorial or facility grounds maintenance, personal uses by employees or other persons, the use of products for the purpose of maintaining motor vehicles operated by the facility, the use of toxic chemicals contained in intake water used for processing or noncontact cooling and intake air used either as compressed air or

for combustion. The owner or operator of the source shall maintain records of the total gallons of coating, adhesive, cleaning material and washoff material used each month and the total gallons used each previous month, and upon request submit the records to the department. The owner or operator shall keep monthly records beginning no less than one year before the compliance date specified in s. NR 465.055 and maintain the records for 5 years.

3. The source emits no more than 4.5 Mg (5 tons) of any one HAP per rolling 12-month period and no more than 11.4 Mg (12.5 tons) of any combination of HAP per rolling 12-month period, and at least 90% of the plantwide HAP emissions per rolling 12-month period are associated with the manufacture of wood furniture or wood furniture components. The owner or operator shall maintain records for 5 years that demonstrate that annual emissions do not exceed these levels, including monthly usage records and certified product data sheets for all finishing material, adhesive, cleaning material and washoff material, and any other records necessary to document emissions from source categories other than wood furniture. The owner or operator shall submit the records to the department upon request.

(d) This chapter does not apply to research or laboratory equipment for which the primary purpose is to conduct research and development into new processes and products, where the equipment is operated under the close supervision of technically trained personnel and is not engaged in the manufacture of products for commercial sale in commerce, except in a de minimis manner.

(e) The owner or operator of an affected source shall comply with the requirements of ch. NR 460, according to the applicability of ch. NR 460 to the sources, as identified in ch. NR 460 Appendix JJ.

(f) Reconstructed affected sources are subject to the requirements for new affected sources. The costs associated with the purchase and installation of air pollution control equipment are not considered in determining whether the facility has been reconstructed, unless the control equipment is part of the process. Additionally, the costs of retrofitting and replacement of equipment that is installed specifically to comply with this chapter are not considered reconstruction costs.

Note: An example of when control equipment is part of a process is where the equipment is used to recover product or raw material. An example of costs for equipment replacement which would not be considered reconstruction costs is the replacement of storage tanks, mix equipment and transfer lines to accommodate conversion to waterborne coatings where the purpose of the conversion is to comply with this chapter.

(2) PURPOSE. This chapter is adopted under ss. 285.27(2) and 285.65, Stats., to establish emission standards for hazardous air pollutants for wood furniture and wood furniture component manufacturing operations.

Note: This chapter is based on the federal regulations contained in 40 CFR part 63 Subpart JJ, created Dec. 7, 1995, as last revised on Dec. 28, 1998.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.02 Definitions. For terms not defined in this section, the definitions contained in chs. NR 400 and 460 apply to the

terms used in this chapter, with definitions in ch. NR 460 taking priority over definitions in ch. NR 400. If this section defines a term which is also defined in ch. NR 400 or 460, the definition in this section applies in this chapter. In this chapter:

(1) "Adhesive" means any chemical substance that is applied for the purpose of bonding 2 surfaces together other than by mechanical means. Products used on humans and animals, adhesive tape, contact paper or any other product with an adhesive incorporated onto or in an inert substrate are not considered adhesives under this chapter.

(2) "Aerosol adhesive" means an adhesive that is dispensed from a pressurized container as a suspension of fine solid or liquid particles in gas.

(3) "Affected source" means a wood furniture manufacturing facility that is engaged, either in part or in whole, in the manufacture of wood furniture or wood furniture components and that is located at a plant site that is a major source of hazardous air pollutants.

(4) "Alternative method" means any method of sampling and analyzing for an air pollutant that is not a reference or equivalent method but has been demonstrated to the administrator's satisfaction to, in specific cases, produce results adequate for a determination of compliance.

(5) "As applied" means the HAP and solids content of the coating or contact adhesive that is actually used for coating or gluing the substrate. It includes the contribution of materials used for in-house dilution of the coating or contact adhesive.

(6) "Basecoat" means a coat of colored material, usually opaque, that is applied before graining inks, glazing coats or other opaque finishing materials, and is usually topcoated for protection.

(7) "Capture device" means a hood, enclosed room, floor sweep or other means of collecting solvent emissions or other pollutants into a duct so that the pollutant can be directed to a pollution control device such as an incinerator or carbon adsorber.

(8) "Capture efficiency" means the fraction of all organic vapors generated by a process that are directed to a control device.

(9) "Certified product data sheet" or "CPDS" means documentation furnished by coating or adhesive suppliers or an outside laboratory and measured in accord with s. NR 465.09(1) that provides all of the following:

(a) The VHAP content of a finishing material, contact adhesive or solvent, by percent by weight, for each VHAP present in concentrations greater than or equal to 1.0% by weight, or 0.1% for VHAP that are carcinogens as defined by the occupational safety and health administration hazard communication standard in 29 CFR part 1910 Subpart Z, incorporated by reference in s. NR 484.03(1).

(b) The solids content of a finishing material or contact adhesive by percent by weight.

(c) The density of the finishing material, adhesive or solvent.

Note: Because the optimum analytical conditions under Method 311, used pursuant to s. NR 465.09(1), vary by coating, the coating or adhesive supplier may also choose to include on the CPDS the optimum analytical conditions for analysis of the coating, adhesive or solvent using EPA Method 311. Information may include, but not be limited to, separation column, oven temperature, carrier gas, injection port temperature, extraction solvent and internal standard.

(10) "Cleaning operations" means operations in which organic HAP solvent is used to remove coating materials or adhesives from equipment used in wood furniture manufacturing operations.

(11) "Coating" means a protective, decorative or functional film applied in a thin layer to a surface. Coating materials include paints, topcoats, varnishes, sealers, stains, washcoats, basecoats, enamels, inks and temporary protective coatings. Adhesives and aerosol spray used for touch-up and repair are not considered coatings under this chapter.

(12) "Coating solids" or "solids" means the part of the coating which remains after the coating is dried or cured.

(13) "Contact adhesive" means an adhesive that is applied to 2 substrates, dried and then mated under only enough pressure to provide sufficient contact so that the resulting bond is immediate and sufficiently strong to hold pieces together without further clamping, pressure or airing.

(14) "Continuous coater" means a finishing system that continuously applies finishing materials onto furniture parts moving along a conveyor. Examples of application methods that can be used with a continuous coater include spraying, curtain coating, roll coating, dip coating and flow coating.

(15) "Continuous compliance" means that the affected source is meeting the emission limitations and other requirements of the rule at all times and is fulfilling all monitoring and recordkeeping provisions of the rule in order to demonstrate compliance.

(16) "Control device" means any equipment that reduces the quantity of a pollutant that is emitted to the air.

Note: The device may destroy or secure the pollutant for subsequent recovery.

(17) "Control device efficiency" means the ratio of the amount of the pollutant reduced by a control device and the amount of the pollutant introduced to the control device.

(18) "Control system" means the combination of capture and control devices used to reduce emissions to the atmosphere.

(19) "Conventional air spray" means a spray coating method in which the coating is atomized by mixing it with compressed air and applied at an air pressure greater than 10 psig at the point of atomization.

Note: Airless and air assisted airless spray technologies are not conventional air spray because the coating is not atomized by mixing it with compressed air. Electrostatic spray technology is also not considered conventional air spray because an electrostatic charge is employed to attract the coating to the workpiece.

(20) "Day" means a period of 24 consecutive hours beginning at midnight local time, or beginning at a time consistent with a facility's operating schedule.

(21) "Enamel" means a coat of colored material, usually opaque, that is applied as a protective topcoat over a basecoat, primer or previously applied enamel coats. In some cases, another finishing material may be applied as a topcoat over the enamel.

(22) "Existing", when used to modify affected source, area source or source, means construction or reconstruction which is commenced before December 6, 1994.

(23) "Finishing material" means a coating used in the wood furniture industry. "Finishing material" includes stains, basecoats, washcoats, enamels, sealers and topcoats.

(24) "Finishing operation" means those operations in which a finishing material is applied to a substrate and is subsequently air-dried, cured in an oven or cured by radiation.

(25) "Foam adhesive" means a contact adhesive used for gluing foam to fabric, foam to foam and fabric to wood.

(26) "Gluing operations" means those operations in which adhesives are used to join components, for example, to apply a laminate to a wood substrate or foam to fabric.

(27) "Incidental wood furniture manufacturer" means a major source that is primarily engaged in the manufacture of products other than wood furniture or wood furniture components and that uses no more than 100 gallons per month of finishing material and no more than 100 gallons per month of adhesives in the manufacture of wood furniture or wood furniture components.

(28) "Material safety data sheet" or "MSDS" means the documentation required for hazardous chemicals by the occupational safety and health administration hazard communication standard in 29 CFR part 1910 Subpart Z, incorporated by reference in s. NR 484.03(1), for a solvent, cleaning material, contact adhesive, coating or other material that identifies select reportable hazardous ingredients of the material, safety and health considerations, and handling procedures.

(29) “New”, when used to modify affected source, area source or source, means construction or reconstruction which is commenced on or after December 6, 1994.

(30) “Nonporous substrate” means a surface that is impermeable to liquids. Examples include metal, rigid plastic, flexible vinyl and rubber.

(31) “Operating parameter value” means a minimum or maximum value established for a control device or process parameter that, if achieved by itself or in combination with one or more other operating parameter values, determines that an owner or operator has complied with an applicable emission limit.

(32) “Organic HAP solvent” means a HAP that is a volatile organic liquid used for dissolving or dispersing constituents in a coating or contact adhesive, adjusting the viscosity of a coating or contact adhesive, or cleaning equipment. When used in a coating or contact adhesive, the organic HAP solvent evaporates during drying and does not become a part of the dried film.

(33) “Overall control efficiency” means the efficiency of a control system, calculated as the product of the capture and control device efficiencies, expressed as a percentage.

(34) “Sealer” means a finishing material used to seal the pores of a wood substrate before additional coats of finishing material are applied. “Sealer” does not include special purpose finishing materials that are used in some finishing systems to optimize aesthetics.

(35) “Stain” means any color coat having a solids content by weight of no more than 8.0% that is applied in single or multiple coats directly to the substrate. “Stain” includes nongrain raising stains, equalizer stains, prestains, sap stains, body stains, no-wipe stains, penetrating stains and toners.

(36) “Storage containers” means vessels or tanks, including mix equipment, used to hold finishing, gluing, cleaning or wash-off materials.

(37) “Strippable spray booth coating” means a coating that meets all of the following:

(a) Is applied to a spray booth wall to provide a protective film to receive overspray during finishing operations.

(b) Is subsequently peeled off and disposed.

(c) Reduces or eliminates the need to use organic HAP solvents to clean spray booth walls.

(38) “Thinner” means a volatile liquid that is used to dilute coatings or contact adhesives to reduce viscosity, color strength and solids, or to modify drying conditions.

(39) “Topcoat” means the last film-building finishing material that is applied in a finishing system.

(40) “Touchup and repair” means the application of finishing materials to cover minor finishing imperfections.

(41) “VHAP” means any volatile hazardous air pollutant listed in Table 1.

(42) “VHAP of potential concern” means any VHAP from the list in Table 5.

(43) “Washcoat” means a transparent special purpose finishing material having a solids content by weight of 12.0% by weight or less.

Note: Washcoats are applied over initial stains to protect, to control color and to stiffen the wood fibers in order to aid sanding.

(44) “Washoff operations” means those operations in which organic HAP solvent is used to remove coating from wood furniture or a wood furniture component.

(45) “Wood furniture” means any product made of wood, a wood product such as rattan or wicker, or an engineered wood product such as particleboard that is manufactured under any of the following standard industrial classification codes, as described in the standard industrial classification manual, 1987, incorporated by reference in s. NR 484.05(1): 2434, 2511, 2512, 2517, 2519, 2521, 2531, 2541, 2599 or 5712.

(46) “Wood furniture component” means any part that is used in the manufacture of wood furniture. Examples include drawer sides, cabinet doors, seat cushions and laminated tops.

(47) “Wood furniture manufacturing operations” means the finishing, gluing, cleaning and washoff operations associated with the production of wood furniture or wood furniture components.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.03 Symbols. The symbols used in this chapter have the following meanings:

(1) “ac” means after the control system is installed and operated.

(2) “A_k” is the area of each natural draft opening k in a total enclosure, in square meters.

(3) “bc” means before control.

(4) “C_{aj}” is the concentration of VHAP in gas stream j exiting the control device, in parts per million by volume.

(5) “C_{bi}” is the concentration of VHAP in gas stream i entering the control device, in parts per million by volume.

(6) “C_c” is the VHAP content of a finishing material c in kilograms of VHAP per kilogram of coating solids (kg VHAP/kg solids), as supplied. Also given in pounds of VHAP per pound of coating solids (lb VHAP/lb solids).

(7) “C_{di}” is the concentration of VHAP in gas stream i entering the control device from the affected source, in parts per million by volume.

(8) “C_{fk}” is the concentration of VHAP in uncontrolled gas stream k emitted directly to the atmosphere from the affected source, in parts per million by volume.

(9) “E” is the emission limit achieved by an emission point or a set of emission points, in kg VHAP/kg solids (lb VHAP/lb solids).

(10) “F” is the control device efficiency, expressed as a fraction.

(11) “FV” is the average inward face velocity across all natural draft openings in a total enclosure, in meters per hour.

(12) “G” is the VHAP content of a contact adhesive, in kg VHAP/kg solids (lb VHAP/lb solids), as applied.

(13) “M_c” is the mass of solids in finishing material c used monthly, kg solids/month (lb solids/month).

(14) “N” is the capture efficiency, expressed as a fraction.

(15) “Q_{aj}” is the volumetric flow rate of gas stream j exiting the control device, in dry standard cubic meters per hour.

(16) “Q_{bi}” is the volumetric flow rate of gas stream i entering the control device, in dry standard cubic meters per hour.

(17) “Q_{di}” is the volumetric flow rate of gas stream i entering the control device from the emission point, in dry standard cubic meters per hour.

(18) “Q_{fk}” is the volumetric flow rate of uncontrolled gas stream k emitted directly to the atmosphere from the emission point, in dry standard cubic meters per hour.

(19) “Q_{ini}” is the volumetric flow rate of gas stream i entering the total enclosure through a forced makeup air duct, in standard cubic meters per hour, wet basis.

(20) “Q_{outj}” is the volumetric flow rate of gas stream j exiting the total enclosure through an exhaust duct or hood, in standard cubic meters per hour, wet basis.

(21) “R” is the overall efficiency of the control system, expressed as a percentage.

(22) “S” is the VHAP content of a solvent, expressed as a weight fraction, added to finishing materials.

(23) “W” is the amount of solvent, in kilograms (pounds), added to finishing materials during the monthly averaging period.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.04 Emission limits. (1) The owner or operator of an existing affected source shall do all of the following:

(a) Limit VHAP emissions from finishing operations to no more than the emission limitations for existing sources presented in Table 2, using any of the compliance methods in s. NR 465.06(1)(a). To determine VHAP emissions from a finishing material containing formaldehyde or styrene, the owner or operator of the affected source shall use the methods presented in s. NR 465.05(12)(a)2. for determining styrene and formaldehyde usage.

(b) Limit VHAP emissions from contact adhesives to no more than the following as appropriate:

1. 1.8 kg VHAP/kg solids (1.8 lb VHAP/lb solids), as applied, for foam adhesives used in products that meet the upholstered seating flammability requirements of California technical bulletin 116, 117 or 133, incorporated by reference in s. NR 484.06(5), the business and institutional furniture manufacturers association's BIFMA X5.7-1991, incorporated by reference in s. NR 484.11(7), upholstered furniture action council flammability test methods, incorporated by reference in s. NR 484.11(10), or any similar requirements from local, state or federal fire regulatory agencies.

2. 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids), as applied, for all other contact adhesives, including foam adhesives used in products that do not meet the standards in subd. 1., but excluding aerosol adhesives and excluding contact adhesives applied to non-porous substrates.

(c) Limit HAP emissions from strippable spray booth coatings by using coatings that contain no more than 0.8 kg VOC/kg solids (0.8 lb VOC/lb solids), as applied.

(2) The owner or operator of a new affected source shall do all of the following:

(a) Limit VHAP emissions from finishing operations to no more than the emission limitations for new sources presented in Table 2 using any of the compliance methods in s. NR 465.06(2)(a). To determine VHAP emissions from a finishing material containing formaldehyde or styrene, the owner or operator of the affected source shall use the methods presented in s. NR 465.05(12)(a)2. for determining styrene and formaldehyde usage.

(b) Limit VHAP emissions from contact adhesives, excluding aerosol adhesives and excluding contact adhesives applied to non-porous substrates, to no more than 0.2 kg VHAP/kg solids (0.2 lb VHAP/lb solids), as applied, using either of the compliance methods in s. NR 465.06(2)(b).

(c) Limit HAP emissions from strippable spray booth coatings by using coatings that contain no more than 0.8 kg VOC/kg solids (0.8 lb VOC/lb solids), as applied.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.05 Work practice standards. (1) WORK PRACTICE IMPLEMENTATION PLAN. The owner or operator of an affected source shall prepare and maintain a written work practice implementation plan that defines environmentally desirable work practices for each wood furniture manufacturing operation and addresses each of the work practice standards in subs. (2) to (12). The plan shall be developed no more than 60 days after the applicable compliance date in s. NR 465.055. The written work practice implementation plan shall be available for inspection by the department upon request. If the department determines that the work practice implementation plan does not adequately address each of the topics specified in subs. (2) to (12), or that the plan does not include sufficient mechanisms for ensuring that the work practice standards are being implemented, the department may require that the owner or operator of the affected source modify the plan. Revisions or modifications to the plan do not require a revision of a permit issued under ch. NR 407.

(2) OPERATOR TRAINING COURSE. The owner or operator of an affected source shall train all personnel, including contract personnel, who are involved in finishing, gluing, cleaning or washoff

operations, use of manufacturing equipment or implementation of the requirements of this chapter. Personnel hired on or after the compliance date of the standard shall be trained upon hiring. Personnel hired before the compliance date shall be trained within 6 months of the compliance date of the standard. All personnel shall be given refresher training annually. The owner or operator of an affected source shall maintain a copy of the training program with the work practice implementation plan. The training program shall include, at a minimum, all of the following:

(a) A list of all current personnel by name and job description that are required to be trained.

(b) An outline of the subjects to be covered in the initial and refresher training for each position or group of personnel.

(c) Lesson plans for courses to be given at the initial and the annual refresher training that include, at a minimum, appropriate application techniques, appropriate cleaning and washoff procedures, appropriate equipment setup and adjustment to minimize finishing material usage and overspray and appropriate management of cleanup wastes.

(d) A description of the methods to be used to demonstrate and document that personnel have successfully completed the initial and refresher training.

(3) INSPECTION AND MAINTENANCE PLAN. The owner or operator of an affected source shall prepare and maintain, with the work practice implementation plan, a written leak inspection and maintenance plan for all pumps, valves, flanges and other equipment used to transfer or apply coatings, adhesives or organic HAP solvents that specifies all of the following:

(a) An inspection schedule which specifies a minimum visual inspection frequency of once per month.

(b) Methods for documenting the date and results of each inspection and any repairs that were made.

(c) The timeframe between identifying the leak and making the repair, which adheres, at a minimum, to the following schedule:

1. A first attempt at repair shall be made no later than 5 calendar days after the leak is detected.

2. Final repairs shall be made within 15 calendar days after the leak is detected, unless the leaking equipment is to be replaced by a new purchase, in which case repairs shall be completed within 3 months.

(4) CLEANING AND WASHOFF SOLVENT ACCOUNTING SYSTEM. The owner or operator of an affected source shall develop an organic HAP solvent accounting form to record all of the following:

(a) The quantity and type of organic HAP solvent used each month for washoff and cleaning operations.

(b) The number of pieces washed off, and the reason for the washoff.

(c) The quantity of spent organic HAP solvent generated from each washoff and cleaning operation each month, and whether it is reused onsite in a process other than cleaning or washoff or disposed of outside of the facility boundaries.

(5) CHEMICAL COMPOSITION OF CLEANING AND WASHOFF SOLVENTS. The owner or operator of an affected source may not use solvents for cleaning or washoff operations that contain any of the pollutants listed in Table 3 in concentrations that require inclusion on an MSDS in accordance with the occupational safety and health administration hazard communication standard in 29 CFR part 1910 Subpart Z, incorporated by reference in s. NR 484.03(1).

(6) SPRAY BOOTH CLEANING. The owner or operator of an affected source may not use compounds containing more than 8.0% by weight of VOC for cleaning spray booth components other than conveyors, continuous coaters and their enclosures, or metal or plastic filters. If the spray booth coating or other protective material used to cover the booth is being replaced, the owner or operator may not use more than 1.0 gallon of organic HAP sol-

vent per booth to prepare the surface of the booth prior to applying the booth coating.

(7) STORAGE REQUIREMENTS. The owner or operator of an affected source shall store finishing, gluing, cleaning and washoff materials in closed containers.

(8) APPLICATION EQUIPMENT REQUIREMENTS. The owner or operator of an affected source may use conventional air spray guns to apply finishing materials only under any of the following circumstances:

(a) To apply finishing materials that have a VOC content no greater than 1.0 kg VOC/kg solids (1.0 lb VOC/lb solids), as applied.

(b) For touchup and repair under any of the following conditions:

1. The touchup and repair occurs after completion of the finishing operation.

2. The touchup and repair occurs after the application of stain and before the application of any other type of finishing material, and the materials used for touchup and repair are applied from a container that has a volume of no more than 2.0 gallons.

(c) When the spray gun is aimed and triggered automatically.

(d) When emissions from the finishing application station are directed to a control device.

(e) When the cumulative total usage of finishing materials applied with conventional air spray guns is no more than 5.0% of the total gallons of finishing material used during that semiannual period.

(f) When the conventional air gun is used to apply stain on a part for which it is technically or economically infeasible to use any other spray application technology. The owner or operator of an affected source shall demonstrate technical or economic infeasibility by submitting to the department a videotape, a technical report or other documentation that supports the affected source's claim of technical or economic infeasibility. The owner or operator shall use one or both of the following criteria to support a claim that no other spray application technology is technically or economically feasible:

1. The production speed is too high or the part shape is too complex for one operator to coat the part and the application station is not large enough to accommodate an additional operator.

2. The excessively large vertical spray area of the part makes it difficult to avoid sagging or runs in the stain.

(9) LINE CLEANING. The owner or operator of an affected source shall pump or drain all organic HAP solvent used for line cleaning into a container which shall be kept covered except when actively adding solvent to or removing solvent from the container.

(10) GUN CLEANING. The owner or operator of an affected source shall collect all organic HAP solvent used to clean spray guns into a container which shall be kept covered except when actively adding solvent to or removing solvent from the container.

(11) WASHOFF OPERATIONS. The owner or operator of an affected source shall control emissions from washoff operations by doing both of the following:

(a) Equipping any tank used for washoff operations with a cover and keeping the cover closed whenever the tank is not being used.

(b) Minimizing dripping by tilting or rotating the part to drain as much solvent as possible.

(12) FORMULATION ASSESSMENT PLAN FOR FINISHING OPERATIONS. (a) The owner or operator of an affected source shall prepare and maintain with the work practice implementation plan a formulation assessment plan that does all of the following:

1. Identifies VHAP from the list presented in Table 4 that are being used in finishing operations by the affected source.

2. Establishes a baseline level of usage by the affected source for each VHAP identified in subd. 1. The baseline usage level

shall be the highest annual usage from 1994, 1995 or 1996, for each VHAP identified in subd. 1., or from another year approved by the department if annual usage data for those years is not available or if the affected source was not in operation during those years. For formaldehyde, the baseline level of usage shall be based on the amount of free formaldehyde present in the finishing material when it is applied. For styrene, the baseline level of usage shall be an estimate of unreacted styrene, which shall be calculated by multiplying the amount of styrene monomer in the finishing material, when it is applied, by a factor of 0.16. For sources using a control device to reduce emissions, an adjusted usage may be calculated based on the overall control efficiency of the control system.

3. Tracks the annual usage of each VHAP identified in subd. 1. that is present in amounts that require inclusion on an MSDS in accordance with the occupational safety and health administration hazard communication standard in 29 CFR part 1910 Subpart Z, incorporated by reference in s. NR 484.03(1).

(b) If, after November 1998, the annual usage of the VHAP identified under par. (a)1. exceeds the baseline level established under par. (a)2., the owner or operator of the affected source shall provide a written notification to the department that describes the amount of the increase and explains the reasons for exceedance of the baseline level. Any of the following explanations relieve the owner or operator from further action, unless the affected source is not in compliance with any state regulations or requirements for that VHAP:

1. The exceedance is no more than 15.0% above the baseline level.

2. Usage of the VHAP is below the de minimis level presented in Table 4 for that VHAP. For sources using a control device to reduce emissions, an adjusted usage based on the overall control efficiency of the control system may be calculated and used to demonstrate that the source does not exceed the de minimis level in Table 4.

3. The affected source is in compliance with ch. NR 400 to 499 for the VHAP.

4. The source of the pollutant is a finishing material with a VOC content of no more than 1.0 kg VOC/kg solids (1.0 lb VOC/lb solids), as applied.

(c) If none of the explanations in par. (b) are the reason for the increase, the owner or operator shall confer with the department to discuss the reason for the increase and whether there are practical and reasonable technology-based solutions for reducing the usage. The evaluation of whether a technology is reasonable and practical shall be based on cost, quality and marketability of the product, whether the technology is being used successfully by other wood furniture manufacturing operations, or other criteria mutually agreed upon by the department and owner or operator. If there are no practical and reasonable solutions, the owner or operator need take no further action. If there are solutions, the owner or operator shall develop a plan to reduce usage of the pollutant to the extent feasible. The plan shall address the approach to be used to reduce emissions, a timetable for implementing the plan and a schedule for submitting notification of progress.

(d) If, after November 1998, the owner or operator of an affected source uses a VHAP of potential concern listed in Table 5 for which a baseline level has not been previously established, the baseline level shall be established as the de minimis level provided in Table 5 for that chemical. The owner or operator shall track the annual usage of each VHAP of potential concern identified in this paragraph that is present in amounts that require inclusion on an MSDS in accordance with the occupational safety and health administration hazard communication standard in 29 CFR part 1910 Subpart Z, incorporated by reference in s. NR 484.03(1). If usage of the VHAP of potential concern exceeds the de minimis level listed in Table 5 for that chemical, the owner or operator shall provide an explanation to the department that docu-

ments the reason for exceedance of the de minimis level. If the explanation is not one of those listed in par. (b), the owner or operator shall follow the procedures in par. (c).

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.055 Compliance dates. (1) EXISTING SOURCES. The compliance date for existing affected sources that emit less than 50 tons per year of HAP in 1996 is December 7, 1998. The compliance date for existing affected sources that emit 50 tons or more of hazardous air pollutants in 1996 is November 21, 1997. The owner or operator of an existing area source that increases its emissions of, or its potential to emit, HAP such that the source becomes a major source that is subject to this chapter shall comply with this chapter one year after becoming a major source.

(2) NEW SOURCES. The compliance date for new affected sources is immediately upon startup or by December 7, 1995, whichever is later. The compliance date for new area sources that become major sources is immediately upon becoming a major source.

Note: Compliance dates are federally enforceable under 40 CFR 63.800 prior to the effective date of this section.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.06 Compliance methods and procedures.

(1) EXISTING SOURCES. (a) Finishing materials. The owner or operator of an existing affected source shall comply with s. NR 465.04 (1)(a) by using any of the following methods:

1. Calculate the average VHAP content for all finishing materials used at the facility using Equation 1, and maintain a value of E no greater than 1.0.

$$E = (M_{c1}C_{c1} + M_{c2}C_{c2} + \dots + M_{cn}C_{cn} + S_1W_1 + S_2W_2 + \dots + S_nW_n) / (M_{c1} + M_{c2} + \dots + M_{cn}) \quad \text{(Equation 1)}$$

2. Use compliant finishing materials according to the following criteria:

a. Demonstrate that each stain, sealer and topcoat has a VHAP content of no more than 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids), as applied, and each thinner contains no more than 10.0% VHAP by weight by maintaining certified product data sheets for each coating and thinner.

b. Demonstrate that each washcoat, basecoat and enamel that is not formulated at the affected source by thinning another finishing material has a VHAP content of no more than 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids), as applied, and each thinner contains no more than 10.0% VHAP by weight by maintaining certified product data sheets for each coating and thinner.

c. Demonstrate that each washcoat, basecoat and enamel that is formulated at the affected source is formulated using a finishing material containing no more than 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids) and a thinner containing no more than 3.0% VHAP by weight.

3. Use a control system with an overall control efficiency (R) such that the value of E_{ac} in Equation 2 is no greater than 1.0. The value of E_{bc} in Equation 2 shall be calculated using Equation 1 in subd. 1.

$$R = [(E_{bc} - E_{ac})/E_{bc}](100) \quad \text{(Equation 2)}$$

4. Use any combination of the methods in subds. 1. to 3.

(b) *Foam adhesives.* The owner or operator of an existing affected source shall comply with s. NR 465.04(1)(b)1. by using foam adhesives with a VHAP content no greater than 1.8 kg VHAP/kg solids (1.8 lb VHAP/lb solids), as applied.

(c) *Other contact adhesives.* The owner or operator of an existing affected source shall comply with s. NR 465.04(1)(b)2. by using either of the following methods:

1. Use contact adhesives with a VHAP content no greater than 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids), as applied.

2. Use a control system with an overall control efficiency (R) such that the value of G_{ac} in Equation 3 is no greater than 1.0.

$$R = [(G_{bc} - G_{ac})/G_{bc}](100) \quad \text{(Equation 3)}$$

(2) NEW SOURCES. (a) Finishing materials. The owner or operator of a new affected source shall comply with s. NR 465.04(2)(a) by using any of the following methods:

1. Calculate the average VHAP content across all finishing materials used at the facility using Equation 1 in sub. (1)(a)1., and maintain a value of E no greater than 0.8.

2. Use compliant finishing materials according to the following criteria:

a. Demonstrate that each sealer and topcoat has a VHAP content of no more than 0.8 kg VHAP/kg solids (0.8 lb VHAP/lb sol-

ids), as applied, each stain has a VHAP content of no more than 1.0 kg VHAP/kg solids (1.0 lb VHAP/lb solids), as applied, and each thinner contains no more than 10.0% VHAP by weight by maintaining certified product data sheets for each coating and thinner.

b. Demonstrate that each washcoat, basecoat and enamel that is not formulated at the affected source by thinning another finishing material has a VHAP content of no more than 0.8 kg VHAP/kg solids (0.8 lb VHAP/lb solids), as applied, and each thinner contains no more than 10.0% VHAP by weight by maintaining certified product data sheets for each coating and thinner.

c. Demonstrate that each washcoat, basecoat and enamel that is formulated at the affected source is formulated using a finishing material containing no more than 0.8 kg VHAP/kg solids (0.8 lb VHAP/lb solids) and a thinner containing no more than 3.0% VHAP by weight.

3. Use a control system with an overall control efficiency (R) such that the value of E_{ac} in Equation 4 is no greater than 0.8. The value of E_{bc} in Equation 4 shall be calculated using Equation 1 in sub. (1)(a)1.

$$R = [(E_{bc} - E_{ac})/E_{bc}](100) \quad \text{(Equation 4)}$$

4. Use any combination of the methods in subds. 1. to 3.

(b) *Contact adhesives.* The owner or operator of a new affected source shall comply with s. NR 465.04(2)(b) by using either of the following methods:

1. Use contact adhesives with a VHAP content no greater than 0.2 kg VHAP/kg solids (0.2 lb VHAP/lb solids), as applied.

2. Use a control system with an overall control efficiency (R) such that the value of G_{ac} in Equation 3 in sub. (1)(c)2. is no greater than 0.2.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.07 Initial compliance demonstration.

(1) FINISHING MATERIALS. The owner or operator of an affected source subject to the provisions of s. NR 465.04(1)(a) or (2)(a) shall demonstrate initial compliance according to the following procedures as appropriate:

(a) If complying by using the methods in s. NR 465.06(1)(a)1. or (2)(a)1., submit the results of the averaging calculation using Equation 1 in s. NR 465.06(1)(a)1. for the first month with the initial compliance status report required by s. NR 465.11(2). The first month's calculation shall include data for the entire month in which the compliance date falls.

(b) If complying by using the methods s. NR 465.06 (1) (a) 2. or (2) (a) 2., state in the initial compliance status report under s. NR 465.11 (2) that compliant stains, washcoats, sealers, topcoats, basecoats, enamels and thinners, as applicable, are being used by the affected source.

(c) If complying by using the methods in s. NR 465.06(1)(a)2. or (2)(a)2. and applying coatings using continuous coaters, do one of the following:

1. State in the initial compliance status report under s. NR 465.11(2) that compliant coatings, as determined by the VHAP content of the coating in the reservoir and the VHAP content as calculated from records, and compliant thinners are being used.

2. State in the initial compliance status report under s. NR 465.11(2) that compliant coatings, as determined by the VHAP content of the coating in the reservoir, are being used; the viscosity of the coating in the reservoir is being monitored; and compliant thinners are being used. The owner or operator shall also submit data that demonstrate that viscosity is an appropriate parameter for demonstrating compliance.

(d) If complying by using the methods in s. NR 465.06(1)(a)3. or (2)(a)3., do all of the following:

1. Submit a monitoring plan that identifies each operating parameter to be monitored for the capture device and discusses why each parameter is appropriate for demonstrating continuous compliance.

2. Conduct an initial performance test as required under s. NR 460.06 using the procedures and test methods listed in ss. NR 460.06 and 465.09(3) and (4).

3. Calculate the overall control efficiency (R) following the procedures in s. NR 465.09(4).

4. Determine those operating conditions critical to determining compliance and establish one or more operating parameters that will ensure compliance with the standard. Operating parameters shall include the following:

a. For a thermal incinerator, minimum combustion temperature shall be the operating parameter.

b. For a catalytic incinerator equipped with a fixed catalyst bed, the minimum gas temperature both upstream and downstream of the catalyst bed shall be the operating parameters.

c. For a catalytic incinerator equipped with a fluidized catalyst bed, the minimum gas temperature upstream of the catalyst bed and the pressure drop across the catalyst bed shall be the operating parameters.

d. For a carbon adsorber, the total regeneration mass stream flow for each regeneration cycle and the carbon bed temperature after each regeneration, or the concentration level of organic compounds exiting the adsorber, shall be the operating parameters, unless the owner or operator requests and receives approval from the administrator to establish other operating parameters.

e. For a control device not listed in this subdivision, one or more operating parameter values shall be established as follows: The owner or operator shall submit, for the administrator's approval, a description of the device, test data verifying the performance of the device, and appropriate site-specific operating parameters that will be monitored to demonstrate continuous compliance with the standard.

(e) The value for each site-specific operating parameter in par. (d)4. shall be calculated as the arithmetic average of the maximum or minimum operating parameter values, as appropriate, that demonstrate compliance with the standards, during the 3 test runs required by s. NR 465.09(3)(a).

(2) CONTACT ADHESIVES. The owner or operator of an affected source subject to the provisions of s. NR 465.04(1)(b) or (2)(b) shall demonstrate initial compliance according to the following procedures as appropriate:

(a) If complying by using the methods in s. NR 465.06(1)(b), (c)1. or (2)(b)1., state in the initial compliance status report under s. NR 465.11(2) that compliant contact adhesives are being used by the affected source.

(b) If complying by using the methods in s. NR 465.06(1)(c)2. or (2)(b)2., do all of the following:

1. Submit a monitoring plan that identifies each operating parameter to be monitored for the capture device and discusses why each parameter is appropriate for demonstrating continuous compliance.

2. Conduct an initial performance test as required under s. NR 460.06 using the procedures and test methods listed in ss. NR 460.06 and 465.09(3) and (4).

3. Calculate the overall control efficiency (R) following the procedures in s. NR 465.09(4).

4. Determine those operating conditions critical to determining compliance and establish one or more operating parameters that will ensure compliance with the standard. Operating parameters shall include the following:

a. For a thermal incinerator, minimum combustion temperature shall be the operating parameter.

b. For a catalytic incinerator equipped with a fixed catalyst bed, the minimum gas temperature both upstream and downstream of the catalyst shall be the operating parameters.

c. For a catalytic incinerator equipped with a fluidized catalyst bed, the minimum gas temperature upstream of the catalyst bed and the pressure drop across the catalyst bed shall be the operating parameters.

(c) The value for each site-specific operating parameter in par. (b)4. shall be calculated as the arithmetic average of the maximum or minimum operating values, as appropriate, that demonstrate compliance with the standards, during the 3 test runs required by s. NR 465.09(3)(a).

(3) STRIPPABLE SPRAY BOOTH COATINGS. The owner or operator of an affected source subject to the provisions s. NR 465.04(1)(c) or (2)(c) shall demonstrate initial compliance by stating in the initial compliance status report under s. NR 465.11(2) that compliant strippable spray booth coatings are being used.

(4) WORK PRACTICE STANDARDS. The owner or operator of an affected source subject to the work practice standards in s. NR 465.05 shall demonstrate initial compliance by stating in the initial compliance status report under s. NR 465.11(2) that the work practice implementation plan has been developed and procedures have been established for implementing the provisions of the plan.

History: CR 00–160: cr. Register August 2001 No. 548, eff. 9–1–01.

NR 465.08 Continuous compliance demonstration.

(1) FINISHING MATERIALS. The owner or operator of an affected source subject to the provisions of s. NR 465.04(1)(a) or (2)(a) shall demonstrate continuous compliance according to the following procedures as appropriate:

(a) If complying by using the methods in s. NR 465.06(1)(a)1. or (2)(a)1., submit the results of the averaging calculation using Equation 1 in s. NR 465.06(1)(a)1. for each month within that semiannual period, and submit a compliance certification in accord with s. NR 465.11(3) which states that the value of E, as calculated using Equation 1 in s. NR 465.06(1)(a)1., is no greater than 1.0 for existing sources or 0.8 for new sources. An affected source is in violation of the standard if the value of E is greater than 1.0 for existing sources or 0.8 for new sources for any month. A violation of the monthly average is a separate violation of the standard for each day of operation during the month, unless the affected source can demonstrate through records that the violation of the monthly average can be attributed to a particular day or days during the period.

(b) If complying by using the methods in s. NR 465.06(1)(a)2. or (2)(a)2., except as provided for in par. (c), submit a compliance certification in accord with s. NR 465.11(3) which states that compliant stains, washcoats, sealers, topcoats, basecoats, enamels and thinners, as applicable, have been used each day in the semiannual reporting period or shall otherwise identify the periods of non-compliance and the reasons for noncompliance. An affected source is in violation of the standard whenever a noncompliant

coating, as demonstrated by records or by a sample of the coating, is used.

(c) If complying by using the methods in s. NR 465.06(1)(a)2. or (2)(a)2. and applying coatings using continuous coaters, do one of the following:

1. Use compliant coatings, as determined by the VHAP content of the coating in the reservoir and the VHAP content as calculated from records, use compliant thinners and submit a compliance certification in accord with s. NR 465.11(3) which states that compliant coatings have been used each day in the semi-annual reporting period, or shall otherwise identify the days of noncompliance and the reasons for noncompliance. An affected source is in violation of the standard whenever a noncompliant coating, as determined by records or by a sample of the coating, is used. Use of a noncompliant coating is a separate violation for each day the noncompliant coating is used.

2. Use compliant coatings, as determined by the VHAP content of the coating in the reservoir, use compliant thinners, maintain a viscosity of the coating in the reservoir that is no less than the viscosity of the initial coating by monitoring the viscosity with a viscosity meter or by testing the viscosity of the initial coating and retesting the coating in the reservoir each time solvent is added, maintain records of solvent additions and submit a compliance certification with the semiannual report required by s. NR 465.11(3). The compliance certification shall be signed by a responsible official of the company that owns or operates the affected source and shall state that compliant coatings, as determined by the VHAP content of the coating in the reservoir, have been used each day in the semiannual reporting period. Additionally, the certification shall state that the viscosity of the coating in the reservoir has not been less than the viscosity of the initial coating, that is, the coating that is initially mixed and placed in the reservoir, for any day in the semiannual reporting period. An affected source is in violation of the standard when a sample of the as-applied coating exceeds the applicable limit established in s. NR 465.06(1)(a)2. or (2)(a)2., as determined using Method 311, in 40 CFR part 63, Appendix A, incorporated by reference in s. NR 438.04, or the viscosity of the coating in the reservoir is less than the viscosity of the initial coating.

(d) If complying by using the methods in s. NR 465.06(1)(a)3. or (2)(a)3., install, calibrate, maintain and operate equipment according to manufacturer's specifications to monitor each site-specific operating parameter established in accordance with s. NR 465.07(2)(b)1. The owner or operator shall also submit the excess emissions and continuous monitoring system performance report and summary report required by ss. NR 465.11(4) and 460.09(5). The appropriate monitoring equipment and related requirements include the following:

1. For an incinerator, a temperature monitoring device equipped with a continuous recorder shall be used as follows:

a. Where a thermal incinerator is used, the temperature monitoring device shall be installed in the firebox or in the ductwork immediately downstream of the firebox in a position before any substantial heat exchange occurs.

b. Where a catalytic incinerator equipped with a fixed catalyst bed is used, temperature monitoring devices shall be installed in the gas stream immediately before and after the catalyst bed.

c. Where a catalytic incinerator equipped with a fluidized catalyst bed is used, the temperature monitoring device shall be installed in the gas stream immediately before the bed. In addition, a pressure monitoring device shall be installed to determine the pressure drop across the catalyst bed. The pressure drop shall be measured monthly at a constant flow rate.

2. For a carbon adsorber, one of the following shall be used:

a. An integrating stream flow monitoring device, having an accuracy of $\pm 10\%$, capable of recording the total regeneration stream mass flow for each regeneration cycle; and a carbon bed temperature monitoring device, having an accuracy of $\pm 1\%$ of the

temperature being monitored or ± 0.5 °C, whichever is greater, and capable of recording the carbon bed temperature after each regeneration and within 15 minutes of completing any cooling cycle.

b. An organic compound monitoring device, equipped with a continuous recorder, to indicate the concentration level of organic compounds exiting the carbon adsorber.

c. Any other monitoring device that has been approved by the administrator in accordance with s. NR 465.07(1)(d)4.d.

3. The owner or operator of an affected source may not operate a capture or control device at a daily average value greater than or less than, as appropriate, the operating parameter values. The daily average value shall be calculated as the average of all values for a monitored parameter recorded during the operating day.

4. The owner or operator of an affected source that is complying through the use of a catalytic incinerator equipped with a fluidized catalyst bed shall maintain a constant pressure drop, measured monthly, across the catalyst bed.

5. An owner or operator using a control device not listed in s. NR 465.07(1)(d) shall submit, for the administrator's approval, a description of the device, test data verifying the performance of the device, and appropriate site-specific operating parameters that will be monitored to demonstrate continuous compliance with the standard.

(2) CONTACT ADHESIVES. The owner or operator of an affected source subject to the provisions of s. NR 465.04(1)(b) or (2)(b) shall demonstrate continuous compliance according to the following procedures as appropriate:

(a) If complying by using the methods in s. NR 465.06(1)(b), (c)1. or (2)(b)1., submit a compliance certification in accord with s. NR 465.11(3) which states that compliant contact adhesives, including foam adhesives, have been used each day in the semi-annual reporting period, or should otherwise identify each day noncompliant contact or foam adhesives were used. Each day a noncompliant contact or foam adhesive is used is a single violation of the standard.

(b) If complying by using the methods in s. NR 465.06(1)(c)2. or (2)(b)2., install, calibrate, maintain and operate equipment according to the manufacturer's specifications to monitor each site-specific operating parameter established in accordance with s. NR 465.07(2)(b)1. The owner or operator shall also submit the excess emissions and continuous monitoring system performance report and summary report required by ss. NR 465.11(4) and 460.09(5). The appropriate monitoring equipment and related requirements include the following:

1. For an incinerator, a temperature monitoring device equipped with a continuous recorder shall be used as follows:

a. Where a thermal incinerator is used, the temperature monitoring device shall be installed in the firebox or in the ductwork immediately downstream of the firebox in a position before any substantial heat exchange occurs.

b. Where a catalytic incinerator equipped with a fixed catalyst bed is used, temperature monitoring devices shall be installed in the gas stream immediately before and after the catalyst bed.

c. Where a catalytic incinerator equipped with a fluidized catalyst bed is used, the temperature monitoring device shall be installed in the gas stream immediately before the bed. In addition, a pressure monitoring device shall be installed to measure the pressure drop across the catalyst bed. The pressure drop shall be measured monthly at a constant flow rate.

2. For a carbon adsorber, one of the following shall be used:

a. An integrating stream flow monitoring device having an accuracy of $\pm 10\%$, capable of recording the total regeneration stream mass flow for each regeneration cycle; and a carbon bed temperature monitoring device, having an accuracy of $\pm 1\%$ of the temperature being monitored or ± 0.5 °C, whichever is greater, and capable of recording the carbon bed temperature after each regeneration and within 15 minutes of completing any cooling cycle.

b. An organic compound monitoring device, equipped with a continuous recorder, to indicate the concentration level of organic compounds exiting the carbon adsorber.

c. Any other monitoring device that has been approved by the administrator in accordance with s. NR 465.07(1)(d)4.d.

3. The owner or operator of an affected source may not operate a capture or control device at a daily average value greater than or less than, as appropriate, the operating parameter values. The daily average value shall be calculated as the average of all values for a monitored parameter recorded during the operating day.

4. The owner or operator of an affected source that is complying through the use of a catalytic incinerator equipped with a fluidized catalyst bed shall maintain a constant pressure drop, measured monthly, across the catalyst bed.

5. An owner or operator using a control device not listed in s. NR 465.07(1)(d) shall submit for the administrator's approval a description of the device, test data verifying the performance of the device, and appropriate site specific operating parameters that will be monitored to demonstrate continuous compliance with the standard.

(3) STRIPPABLE SPRAY BOOTH COATINGS. The owner or operator of an affected source subject to the provisions s. NR 465.04(1)(c) or (2)(c) shall demonstrate continuous compliance by submitting a compliance certification in accord with s. NR 465.11(3) which states that compliant strippable spray booth coatings have been used each day in the semiannual reporting period, or otherwise identifies each day noncompliant materials were used. Each day a noncompliant strippable booth coating is used is a single violation of the standard.

(4) WORK PRACTICE STANDARDS. The owner or operator of an affected source shall demonstrate continuous compliance with the work practice standards in s. NR 465.05 by submitting a compliance certification in accord with s. NR 465.11(3) which states that the work practice implementation plan is being followed, or otherwise identifies the provisions of the plan that have not been implemented and each day the provisions were not implemented. During any period of time that an owner or operator is required to implement the provisions of the plan, each failure to implement an obligation under the plan during any particular day is a violation.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.09 Performance test methods. (1) (a) Except as provided in par.(c), the owner or operator of an affected source shall use Method 311 in 40 CFR part 63, Appendix A, incorporated by reference in s. NR 484.04(25), in conjunction with formulation data, to determine the VHAP content of liquid coatings. Formulation data shall be used to identify VHAP present in the coating. Method 311 shall then be used to quantify those VHAP identified through formulation data. Method 311 may not be used to quantify HAP such as styrene and formaldehyde that are emitted during the cure.

(b) Except as provided in par.(c), the owner or operator of an affected source shall use Method 24 in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), to determine the solids content by weight and the density of coatings.

(c) If it is demonstrated to the satisfaction of the administrator that a coating does not release VOC or HAP byproducts during the cure, batch formulation information shall be accepted for determining the VHAP or VOC content of the coating. The owner or operator of an affected source may request approval from the administrator to use an alternative method for determining the VHAP content of the coating. In the event of any inconsistency between Method 24 or Method 311 test data for a material and formulation data for the same material, the applicable test method shall govern unless, after consultation, the owner or operator can demonstrate to the satisfaction of the department that the formulation data are correct.

(2) An owner or operator demonstrating compliance in accordance with s. NR 465.07(1)(d) or (2)(b) or 465.08(1)(d) or (2)(b), or complying with any of the other emission limits of s. NR 465.04 by operating a capture and control device, shall determine the overall control efficiency of the control system (R) as the product of the capture and control device efficiency, using the test methods in sub. (3) and the procedures in sub. (4).

(3) When an initial compliance demonstration is required by s. NR 465.07(1)(d) or (2)(b), the following test methods shall be used:

(a) Method 18 in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be used to determine the HAP concentration of gaseous air streams. The test shall consist of 3 separate runs, each lasting a minimum of 30 minutes.

(b) Method 1 or 1A in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be used for sample and velocity traverses.

(c) Method 2, 2A, 2C or 2D in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be used to measure velocity and volumetric flow rates.

(d) Method 3 in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be used to analyze the exhaust gases.

(e) Method 4 in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be used to measure the moisture in the stack gas.

(f) Methods 2, 2A, 2C, 2D, 3 and 4 in 40 CFR part 60, Appendix A, incorporated by reference in s. NR 484.04(13), shall be performed, as applicable, at least twice during each test period.

(4) An owner or operator of an affected source demonstrating compliance in accordance with s. NR 465.07(1)(d) or (2)(b) shall perform a gaseous emission test using the following procedures:

(a) Construct the overall HAP emission reduction system so that all volumetric flow rates and total HAP emissions can be accurately determined by the applicable test methods specified in sub.(3).

(b) Determine capture efficiency from the affected emission points by capturing, venting and measuring all HAP emissions from the affected emission points. The owner or operator shall isolate affected emission points located in an area with other non-affected gaseous emission sources from all other gaseous emission points and shall determine capture efficiency according to the following procedures as appropriate:

1. For temporary total enclosures, capture efficiency shall be determined according to Method 204, and as applicable, Methods 204A through 204F, in 40 CFR part 51, Appendix M, incorporated by reference in s. NR 484.04(9).

2. For permanent total enclosures built around the affected emissions points, and where the building that houses the process is used as the enclosure, capture efficiency shall be assumed to be 100% when the requirements for a permanent total enclosure in Method 204, in 40 CFR part 51, Appendix M, incorporated by reference in s. NR 484.04(9), are satisfied. Notwithstanding par. (e), when a permanent total enclosure is confirmed through Method 204, a value of one shall be used for N when determining R under sub. (5), (6), (7) or (8).

3. Use any alternative protocol and test method provided they meet either the requirements of the data quality objective (DQO) approach or the lower confidence level (LCL) approach in 40 CFR part 63, Subpart KK, Appendix A, incorporated by reference in s. NR 484.04(24).

4. Shut down all nonaffected HAP emission points and continue to exhaust fugitive emissions from the affected emission points through any building ventilation system and other room exhausts such as drying ovens. All exhaust air shall be vented through stacks suitable for testing.

5. Use another methodology approved by the administrator provided it complies with the criteria for acceptance under Method 301 in 40 CFR part 63, Appendix A, incorporated by reference in s. NR 484.04(25).

(c) Operate the control device with all affected emission points that will subsequently be delivered to the control device connected and operating at maximum production rate.

(d) Determine the efficiency (F) of the control device using the following equation:

$$F = \frac{\sum_{i=1}^n Q_{bi} C_{bi} - \sum_{j=1}^p Q_{aj} C_{aj}}{\sum_{i=1}^n Q_{bi} C_{bi}} \quad (\text{Equation 5})$$

(e) Determine the efficiency (N) of the capture system using the following equation:

$$N = \frac{\sum_{i=1}^n Q_{di} C_{di}}{\sum_{i=1}^n Q_{di} C_{di} + \sum_{k=1}^p Q_{fk} C_{fk}} \quad (\text{Equation 6})$$

(5) For an owner or operator complying with s. NR 465.04(1)(a) in accordance with s. NR 465.06(1)(a)3., compliance is demonstrated if the product of (F x N)(100) yields a value (R) such that the value of E_{ac} in Equation 2 in s. NR 465.06(1)(a)3. is no greater than 1.0.

(6) For an owner or operator complying s. NR 465.04(2)(a) in accordance with s. NR 465.06(2)(a)3., compliance is demonstrated if the product of (F x N)(100) yields a value (R) such that the value of E_{ac} in Equation 4 in s. NR 465.06(2)(a)3. is no greater than 0.8.

(7) For an owner or operator complying with s. NR 465.04(1)(b)2. in accordance with s. NR 465.06(1)(c)2., compliance is demonstrated if the product of (F x N)(100) yields a value (R) such that the value of G_{ac} in Equation 3 in s. NR 465.06(1)(c)2. is no greater than 1.0.

(8) For an owner or operator complying with s. NR 465.04(2)(b) in accordance with s. NR 465.06(2)(b)2., compliance is demonstrated if the product of (F x N)(100) yields a value (R) such that the value of G_{ac} in Equation 3 s. NR 465.06(1)(c)2. is no greater than 0.2.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01; **reprinted to correct error in (4) (d) and (e), Register October 2001 No. 550.**

NR 465.10 Recordkeeping requirements. (1) The owner or operator of an affected source shall fulfill all recordkeeping requirements of s. NR 460.09 according to the applicability criteria in s. NR 465.01(1)(e).

(2) The owner or operator of an affected source subject to the emission limits in s. NR 465.04 shall maintain records of the following as appropriate:

(a) A certified product data sheet for each finishing material, thinner, contact adhesive and strippable spray booth coating subject to the emission limits in s. NR 465.04.

(b) The VHAP content, in kg VHAP/kg solids (lb VHAP/lb solids), as applied, of each finishing material and contact adhesive subject to the emission limits in s. NR 465.04.

(c) The VOC content, in kg VOC/kg solids (lb VOC/lb solids), as applied, of each strippable spray booth coating subject to the emission limits in s. NR 465.04(1)(c) or (2)(c).

(3) The owner or operator of an affected source following the compliance method in s. NR 465.06(1)(a)1. or (2)(a)1. shall maintain copies of the averaging calculation for each month following the compliance date, as well as the data on the quantity of coatings and thinners used that is necessary to support the calculation of E in Equation 1 in s. NR 465.06(1)(a)1.

(4) The owner or operator of an affected source following the compliance procedures of s. NR 465.07(1)(c)2. or 465.08(1)(c)2. shall maintain the records required by s. NR 465.10(2) as well as records of all of the following:

(a) Solvent and coating additions to the continuous coater reservoir.

(b) Viscosity measurements.

(c) Data demonstrating that viscosity is an appropriate parameter for demonstrating compliance.

(5) The owner or operator of an affected source subject to the work practice standards in s. NR 465.05 shall maintain onsite a work practice implementation plan and all records associated with fulfilling the requirements of that plan, including, but not limited to, all of the following:

(a) Records demonstrating that the operator training program required by s. NR 465.05(2) is in place.

(b) Records collected in accordance with the inspection and maintenance plan required by s. NR 465.05(3).

(c) Records associated with the cleaning solvent accounting system required by s. NR 465.05(4).

(d) Records associated with the limitation on the use of conventional air spray guns showing total finishing material usage and the percentage of finishing materials applied with conventional air spray guns for each semiannual period as required by s. NR 465.05(8)(e).

(e) Records associated with the formulation assessment plan required by s. NR 465.05(12).

(f) Copies of documentation such as logs developed to demonstrate that the other provisions of the work practice implementation plan are followed.

(6) The owner or operator of an affected source following the compliance method of s. NR 465.07(1)(d) or 465.08(1)(d) shall maintain copies of the calculations demonstrating that the overall control efficiency (R) of the control system results in the value of E_{ac} required by Equation 2 in s. NR 465.06(1)(a)3. or Equation 4 in s. NR 465.06(2)(a)3., records of the operating parameter values, and copies of the semiannual compliance reports required by s. NR 465.11(4).

(7) The owner or operator of an affected source following the compliance method of s. NR 465.07(2)(b) or 465.08(2)(b) shall maintain copies of the calculations demonstrating that the overall control efficiency (R) of the control system results in the applicable value of G_{ac} calculated using Equation 3 in s. NR 465.06(1)(c)2., records of the operating parameter values, and copies of the semiannual compliance reports required by s. NR 465.11(4).

(8) The owner or operator of an affected source subject to the emission limits in s. NR 465.04 and following the compliance provisions of s. NR 465.07(1)(a) to (c), (2)(a), (3) or (4) or 465.08(1)(a) to (c), (2)(a), (3) or (4) shall maintain records of the compliance certifications submitted in accordance with s. NR 465.11(3) for each semiannual period following the compliance date.

(9) The owner or operator of an affected source shall maintain records of all other information submitted with the compliance

status report required by ss. NR 460.08(8) and 465.11(2) and the semiannual reports required by s. NR 465.11(3).

(10) The owner or operator of an affected source shall maintain all records in accordance with the requirements of s. NR 460.09(2)(a).

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

NR 465.11 Reporting requirements. (1) The owner or operator of an affected source subject to this chapter shall fulfill all reporting requirements of ss. NR 460.06 to 460.09 according to the applicability criteria in s. NR 465.01(1)(e).

(2) The owner or operator of an affected source demonstrating compliance in accordance with s. NR 465.07(1)(a) to (c), (2)(a), (3) or (4) shall submit the compliance status report required by s. NR 460.08(8) no later than 60 days after the compliance date. The report shall include the information required by s. NR 465.07 (1) (a) to (c), (2)(a), (3) or (4).

(3) The owner or operator of an affected source demonstrating compliance in accordance with s. NR 465.08(1)(a) to (c), (2)(a), (3) or (4) shall submit a report covering the previous 6 months of wood furniture manufacturing operations as follows:

(a) The first report shall be submitted no later than 30 calendar days after the end of the first 6-month period following the compliance date.

(b) Subsequent reports shall be submitted no later than 30 calendar days after the end of each 6-month period following the first report.

(c) The semiannual reports shall include the information required by s. NR 465.08(1)(a) to (c), (2)(a), (3) and (4), a compliance certification stating whether the affected source was in compliance or noncompliance, and, if the affected source was in noncompliance, the measures taken to bring the affected source into compliance. The compliance certification shall be signed by a responsible official of the company that owns or operates the affected source.

(4) The owner or operator of an affected source demonstrating compliance in accordance with s. NR 465.08(1)(d) or (2)(b) shall submit the excess emissions and continuous monitoring system performance report and summary report required by s. NR 460.09(5). The report shall include the monitored operating parameter values required by s. NR 465.08(1)(d) or (2)(b). If the source experiences excess emissions, the report shall be submitted quarterly for at least one year after the excess emissions occur and until a request to reduce reporting frequency is approved, as indicated in s. NR 460.09(5)(c)1.c. If no excess emissions occur, the report shall be submitted semiannually.

(5) The owner or operator of an affected source required to provide a written notification under s. NR 465.05(12)(b) shall include in the notification one or more statements that explains the reasons for the usage increase. The notification shall be submitted no later than 30 calendar days after the end of the annual period in which the usage increase occurred.

History: CR 00-160: cr. Register August 2001 No. 548, eff. 9-1-01.

Table 1
List of Volatile Hazardous Air Pollutants

Chemical Name	CAS Number		
		DDE (1,1-Dichloro-2,2-bis(p-chlorophenyl) ethylene)	72559
		Diazomethane	334883
		Dibenzofuran	132649
		1,2-Dibromo-3-chloropropane	96128
		Dibutylphthalate	84742
		1,4-Dichlorobenzene	106467
		3,3'-Dichlorobenzidine	91941
		Dichloroethyl ether (Bis(2-chloroethyl)ether)	111444
		1,3-Dichloropropene	542756
		Diethanolamine	111422
		N,N-Dimethylaniline	121697
		Diethyl sulfate	64675
		3,3'-Dimethoxybenzidine	119904
		4-Dimethylaminoazobenzene	60117
		3,3'-Dimethylbenzidine	119937
		Dimethylcarbamoyl chloride	79447
		N,N-Dimethylformamide	68122
		1,1-Dimethylhydrazine	57147
		Dimethyl phthalate	131113
		Dimethyl sulfate	77781
		4,6-Dinitro-o-cresol, and salts	534521
		2,4-Dinitrophenol	51285
		2,4-Dinitrotoluene	121142
		1,4-Dioxane (1,4-Diethyleneoxide)	123911
		1,2-Diphenylhydrazine	122667
		Epichlorohydrin (1-Chloro-2,3-epoxypropane)	106898
		1,2-Epoxybutane	106887
		Ethyl acrylate	140885
		Ethylbenzene	100414
		Ethyl carbamate (Urethane)	51796
		Ethyl chloride (Chloroethane)	75003
		Ethylene dibromide (Dibromoethane)	106934
		Ethylene dichloride (1,2-Dichloroethane)	107062
		Ethylene glycol	107211
		Ethylene oxide	75218
		Ethylenethiourea	96457
		Ethylidene dichloride (1,1-Dichloroethane)	75343
		Formaldehyde	50000
		Glycol ethers ^a	
		Hexachlorobenzene	118741
		Hexachloro-1,3-butadiene	87683
Acetaldehyde	75070		
Acetamide	60355		
Acetonitrile	75058		
Acetophenone	98862		
2-Acetylaminofluorine	53963		
Acrolein	107028		
Acrylamide	79061		
Acrylic acid	79107		
Acrylonitrile	107131		
Allyl chloride	107051		
4-Aminobiphenyl	92671		
Aniline	62533		
o-Anisidine	90040		
Benzene	71432		
Benzidine	92875		
Benzotrichloride	98077		
Benzyl chloride	100447		
Biphenyl	92524		
Bis(2-ethylhexyl) phthalate (DEHP)	117817		
Bis(chloromethyl) ether	542881		
Bromoform	75252		
1,3-Butadiene	106990		
Carbon disulfide	75150		
Carbon tetrachloride	56235		
Carbonyl sulfide	463581		
Catechol	120809		
Chloroacetic acid	79118		
2-Chloroacetophenone	532274		
Chlorobenzene	108907		
Chloroform	67663		
Chloromethyl methyl ether	107302		
Chloroprene	126998		
Cresols (isomers and mixture)	1319773		
o-Cresol	95487		
m-Cresol	108394		
p-Cresol	106445		
Cumene	98828		
2,4-D (2,4-Dichlorophenoxyacetic acid, including salts and esters)	94757		

Table 1 (Continued)
List of Volatile Hazardous Air Pollutants

Chemical Name	CAS Number
Hexachloroethane	67721
Hexamethylene-1,6-diisocyanate	822060
Hexamethylphosphoramide	680319
Hexane	110543
Hydrazine	302012
Hydroquinone	123319
Isophorone	78591
Maleic anhydride	108316
Methanol	67561
Methyl bromide (Bromomethane)	74839
Methyl chloride (Chloromethane)	74873
Methyl chloroform (1,1,1-Trichloroethane)	71556
Methyl ethyl ketone (2-Butanone)	78933
Methylhydrazine	60344
Methyl iodide (Iodomethane)	74884
Methyl isobutyl ketone (Hexone)	108101
Methyl isocyanate	624839
Methyl methacrylate	80626
Methyl tert-butyl ether	1634044
4,4'-Methylene bis(2-chloroaniline)	101144
Methylene chloride (Dichloromethane)	75092
4,4'-Methylenediphenyl diisocyanate (MDI)	101688
4,4'-Methylenedianiline	101779
Naphthalene	91203
Nitrobenzene	98953
4-Nitrobiphenyl	92933
4-Nitrophenol	100027
2-Nitropropane	79469
N-Nitroso-N-methylurea	684935
N-Nitrosodimethylamine	62759
N-Nitrosomorpholine	59892
Phenol	108952
p-Phenylenediamine	106503
Phosgene	75445
Phthalic anhydride	85449
Polychlorinated biphenyls (Aroclors)	1336363

Polycyclic Organic Matter^b	
1,3-Propane sultone	1120714
β-Propiolactone	57578
Propionaldehyde	123386
Propoxur (Baygon)	114261
Propylene dichloride (1,2-Dichloropropane)	78875
Propylene oxide	75569
1,2-Propylenimine (2-Methyl aziridine)	75558
Quinone	106514
Styrene	100425
Styrene oxide	96093
2,3,7,8-Tetrachlorodibenzo-p-dioxin	1746016
1,1,2,2-Tetrachloroethane	79345
Tetrachloroethylene (Perchloroethylene)	127184
Toluene	108883
2,4-Toluenediamine	95807
Toluene-2,4-diisocyanate	584849
o-Toluidine	95534
1,2,4-Trichlorobenzene	120821
1,1,2-Trichloroethane	79005
Trichloroethylene	79016
2,4,5-Trichlorophenol	95954
2,4,6-Trichlorophenol	88062
Triethylamine	121448
Trifluralin	1582098
2,2,4-Trimethylpentane	540841
Vinyl acetate	108054
Vinyl bromide	593602
Vinyl chloride	75014
Vinylidene chloride (1,1-Dichloroethylene)	75354
Xylenes (isomers and mixture)	1330207
o-Xylene	95476
m-Xylene	108383
p-Xylene	106423

^aIncludes mono- and di-ethers of ethylene glycol, diethylene glycols and triethyl-ene glycol; R-(OCH₂CH₂)_n-OR' where:

n = 1, 2 or 3

R = alkyl or aryl groups

R' = R, H, or groups which, when removed, yield glycol ethers with the structure: R-(OCH₂CH₂)_n-OH. Polymers are excluded from the glycol category.

^bIncludes organic compounds with more than one benzene ring, and which have a boiling point greater than or equal to 100 °C.

Table 2
Summary of Emission Limits and Compliance Demonstration Methods

Emission Point	Existing Source	New Source
(1) Finishing Operations:		
(a) Achieve a monthly weighted average VHAP content across all finishing materials (maximum kg VHAP/kg solids [lb VHAP/lb solids], as applied).	1.0	0.8
(b) Use compliant finishing materials (maximum kg VHAP/kg solids [lb VHAP/lb solids], as applied):		
1. Stains	^a 1.0	^a 1.0
2. Sealers	^a 1.0	^a 0.8
3. Topcoats	^a 1.0	^a 0.8
4. Washcoats	^b 1.0	^b 0.8
5. Basecoats	^b 1.0	^b 0.8
6. Enamels	^b 1.0	^b 0.8
(c) Use a control device.	^c 1.0	^c 0.8
(d) Use any combination of (a), (b) and (c).	1.0	0.8
(2) Cleaning Operation. Use compliant strippable spray booth coatings (maximum kg VOC/kg solids [lb VOC/lb solids], as applied)	0.8	0.8
(3) Contact Adhesives:		
(a) Use compliant contact adhesives (maximum kg VHAP/kg solids [lb VHAP/lb solids], as applied):		
1. For aerosol adhesive, and for contact adhesives applied to nonporous substrates.	^d NA	^d NA
2. For foam adhesives used in products that meet flammability requirements.	1.8	0.2
3. For all other contact adhesives.	1.0	0.2
(b) Use a control device	^e 1.0	^e 0.2

^a Any thinner used onsite shall contain no more than 10.0% VHAP by weight.

^b Any thinner used onsite shall contain no more than 10.0% VHAP by weight if the finishing material is purchased premade, that is, if it is not formulated onsite by thinning other finishing materials, i.e., thinning a sealer to use as a washcoat. If formulated onsite, these shall be formulated using compliant finishing materials, i.e., those that meet the limits specified in this table, and thinners containing no more than 3.0% VHAP by weight.

^c The control device shall be operated at an efficiency that is equivalent to no greater than 1.0 pound (or 0.8 pounds) of VHAP being emitted from the affected emission source per pound of solids used.

^d There is no limit on the VHAP content of these adhesives.

^e The control device shall be operated at an efficiency that is equivalent to no greater than 1.0 pounds (or 0.2 pounds) of VHAP being emitted from the affected emission source per pound of solids used.

Table 3
Pollutants Excluded From Use in Cleaning
and Washoff Solvents

Chemical Name	CAS Number		
Acetaldehyde	75070	Dimethyl carbamoyl chloride	79447
Acetamide	60355	Dimethyl formamide	68122
2-Acetylaminoflourine	53963	1,1-Dimethyl hydrazine	57147
Acrylamide	79061	2,4-Dinitrotoluene	121142
Acrylonitrile	107131	1,4-Dioxane (1,4-Diethyleneoxide)	123911
4-Aminobiphenyl	92671	1,2-Diphenylhydrazine	122667
Aniline	62533	Epichlorohydrin	106898
o-Anisidine	90040	Ethyl acrylate	140885
Antimony trioxide	1309644	Ethyl carbamate (Urethane)	51796
Arsenic and inorganic arsenic compounds	7440382	Ethylene dibromide (1,2-Dibromoethane)	106934
Benz(c)acridine	225514	Ethylene dichloride (1,2-Dichloroethane)	107062
Benzene	71432	Ethylene oxide	75218
Benzidine	92875	Ethylene thiourea	96457
Benzo(a)anthracene	56553	Formaldehyde	50000
Benzo(b)fluoranthene	205992	Heptachlor	76448
Benzo(a)pyrene	50328	Hexachlorobenzene	118741
Beryllium compounds	7440417	Hexamethylphosphoramide	680319
Beryllium salts		Hydrazine	302012
Bis(chloromethyl)ether	542881	Indeno(1,2,3-cd)pyrene	193395
Bis(2-ethylhexyl)phthalate (DEHP)	117817	Lindane (Hexachlorocyclohexane, gamma)	58899
Bromoform	75252	Methyl hydrazine	60344
1,3-Butadiene	106990	Methylene chloride (Dichloromethane)	75092
Cadmium compounds		4,4'-Methylenedianiline	101779
Captan	133062	Nickel refinery dust	
Carbon tetrachloride	56235	Nickel subsulfide	12035722
Chlordane	57749	2-Nitropropane	79469
Chlorobenzilate	510156	N-Nitrosodimethylamine	62759
Chloroform	67663	N-Nitroso-N-methylurea	684935
Chromium compounds (hexavalent)		N-Nitrosomorpholine	59892
Chrysene	218019	Pentachlorophenol	87865
Coke oven emissions		Polychlorinated biphenyls (Aroclors)	1336363
DDE (1,1-Dichloro-2,2-bis(p-chlorophenyl) ethylene) 2559	72559	1,3-Propane sultone	1120714
Dibenz(ah)anthracene	53703	Propoxur	114261
1,2:7,8-Dibenzopyrene	189559	Propylene dichloride (1,2-Dichloropropane)	78875
1,2-Dibromo-3-chloropropane	96128	Propylene oxide	75569
1,4-Dichlorobenzene(p)	106467	1,2-Propylenimine (2-Methyl aziridine)	75558
3,3'-Dichlorobenzidine	53963	Selenium sulfide (mono- and di-)	7488564
Dichloroethyl ether (Bis(2-chloroethyl) ether)	111444	Styrene oxide	96093
1,3-Dichloropropene	542756	2,3,7,8-Tetrachlorodibenzo-p-dioxin	1746016
Dichlorvos	62737	2,4-Toluene diamine	95807
Diethyl sulfate	64675	o-Toluidine	95534
3,3'-Dimethoxybenzidine	119904	Toxaphene (Chlorinated camphene)	8001352
Dimethyl aminoazobenzene	60117	2,4,6-Trichlorophenol	88062
7,12-Dimethylbenz(a)anthracene	57976	Vinyl bromide (Bromoethene)	593602
3,3'-Dimethyl benzidine	119937	Vinyl chloride	75014

Table 4
List of VHAP of Potential Concern Identified by Industry

Chemical Name	CAS Number	de minimis (tons/yr)
Diethanolamine	111422	5.0
Dimethyl formamide	68122	1.0
2-Ethoxyethyl acetate	111159	10.0
Formaldehyde	50000	0.2
Isophorone	78591	0.7
2-Methoxyethanol	109864	10.0
Methylene chloride	75092	4.0
2-Nitropropane	79469	1.0
Phenol	108952	0.1
Styrene monomer	100425	1.0

Table 5
VHAP of Potential Concern

Chemical Name	CAS Number	de minimis (tons/yr)*
Acetaldehyde	75070	0.9
Acetamine	60355	1.0
Acetophenone	98862	1.0
2-Acetylaminoflourine	53963	0.0005
Acrolein	107028	0.04
Acrylamide	79061	0.002
Acrylonitrile	107131	0.03
Allyl chloride	107051	1.0
4-Aminobipheny	192671	1.0
Aniline	62533	0.1
o-Anisidine	90040	1.0
Benz(c)acridine	225514	0.01
Benzene	71432	0.2
Benzidine	92875	0.00003
Benzo(a)anthracene	56553	0.01
Benzo(b)fluoranthene	205992	0.01
Benzo(a)pyrene	50328	0.001
Benzotrichloride	98077	0.0006
Benzyl chloride	100447	0.04
Bis(chloromethyl) ether	542881	0.00003
Bis(2-ethylhexyl) phthalate (DEHP)	117817	0.5
Bromoform	75252	2.0
1,3-Butadiene	106990	0.007
Captan	133062	2.0
Carbon disulfide	75150	1.0
Carbon tetrachloride	56235	0.1
Carbonyl sulfide	463581	5.0
Catechol	120809	5.0
Chloramben	133904	1.0
Chlordane	57749	0.005
Chloroacetic acid	79118	0.1
2-Chloroacetophenone	532274	0.06
Chlorobenzilate	510156	0.04
Chloroform	67663	0.09
Chloromethyl methyl ether	107302	0.1
Chloroprene	126998	1.0
Chrysene	218019	0.01
Cobalt carbonyl	10210681	0.1
m-Cresol	108394	1.0
o-Cresol	95487	1.0
p-Cresol	106445	1.0
Cresols/Cresylic acid (isomers and mixture)	1319773	1.0
DDE (1,1-Dichloro-2,2-bis(p-chlorophenyl) ethylene)	72559	0.01
Diazomethane	334883	1.0
Dibenz(ah)anthracene	53703	0.01
Dibenzofurans	132649	5.0
1,2:7,8-Dibenzopyrene	189559	0.01
1,2-Dibromo-3-chloropropane	96128	0.001
1,4-Dichlorobenzene(p)	106467	0.3
3,3'-Dichlorobenzidine	91941	0.02
Dichloroethyl ether (Bis(2-chloroethyl)ether)	111444	0.006
1,3-Dichloropropene	542756	1.0
Dichlorvos	62737	0.02
Diethanolamine	111422	5.0
Diethyl sulfate	64675	1.0
3,3'-Dimethoxybenzidine	119904	0.01
Dimethyl aminoazobenzene	60117	1.0
N,N-Dimethylaniline	121697	1.0
7,12-Dimethylbenz(a)anthracene	57976	0.01
3,3'-Dimethyl benzidine	119937	0.001
Dimethyl carbamoyl chloride	79447	0.002
Dimethyl formamide	68122	1.0
1,1-Dimethyl hydrazine	57147	0.0008
Dimethyl sulfate	77781	0.1
4,6-Dinitro-o-cresol, and salts	534521	0.1
2,4-Dinitrophenol	51285	1.0
2,4-Dinitrotoluene	121142	0.002
1,4-Dioxane (1,4-Diethylenoxide)	123911	0.6
1,2-Diphenylhydrazine	122667	0.009
Epichlorohydrin	106898	2.0
1,2-Epoxybutane	106887	1.0
Ethyl acrylate	140885	0.1
Ethyl carbamate (Urethane)	51796	0.08
Ethylene dibromide (1,2-Dibromoethane)	106934	0.01

Table 5 (Continued)
VHAP of Potential Concern

Chemical Name	CAS Number	de minimis (tons/yr)			
Ethylene dichloride (1,2-Dichloroethane)	107062	0.08	2-Nitropropane	79469	1.0
Ethylene imine	151564	0.0003	N-Nitrosodimethylamine	62759	0.0001
Ethylene oxide	75218	0.09	N-Nitroso-N-methylurea	684935	0.00002
Ethylene thiourea	96457	0.06	N-Nitrosomorpholine	59892	1.0
Ethylidene dichloride (1,1-Dichloroethane)	75343	1.0	Parathion	56382	0.1
Fluomine	62207765	0.1	Pentachloronitrobenzene (Quintobenzene)	82688	0.03
Formaldehyde	50000	0.2	Pentachlorophenol	87865	0.07
Glycol ethers ^a		5.0	Phenol	108952	0.1
Heptachlor	76448	0.002	Phthalic anhydride	85449	5.0
Hexachlorobenzene	118741	0.004	Polychlorinated biphenyls (Aroclors)	1336363	0.0009
Hexachlorobutadiene	87683	0.09	Polycyclic organic matter ^b		0.01
Hexachlorocyclopentadiene	77474	0.1	1,3-Propane sultone	1120714	0.003
Hexachloroethane	67721	0.5	β-Propiolactone	57578	0.1
Hexamethylene-1,6-diisocyanate	822060	5.0	Propionaldehyde	123386	5.0
Hexamethylphosphoramide	680319	0.01	Propoxur	114261	2.0
Hydroquinone	123319	1.0	Propylene dichloride (1,2-Dichloropropane)	78875	0.1
Indeno(1,2,3-cd)pyrene	193395	0.01	Propylene oxide	75569	0.5
Isophorone	78591	0.07	1,2-Propylenimine (2-Methylaziridine)	75558	10.0003
Lindane (Hexachlorocyclohexane, gamma)	58899	0.005	Quinoline	91225	0.0006
Maleic anhydride	108316	1.0	Quinone	106514	5.0
Mercury, (acetato-o)phenyl- (Phenylmercuric Acetate)	62384	0.01	Styrene	100425	1.0
2-Methoxy ethanol	109864	10.0	Styrene oxide	96093	1.0
Methyl bromide (Bromomethane)	74839	10.0	2,3,7,8-Tetrachlorodibenzo-p-dioxin	1746016	0.00000006
Methyl chloride (Chloromethane)	74873	1.0	1,1,2,2-Tetrachloroethane	79345	0.03
Methylcyclopentadienyl manganese	12108133	0.1	Tetrachloroethylene (Perchloroethylene)	127184	4.0
4,4'-Methylene bis(2-chloroaniline)	101144	0.02	Tetraethyl lead	78002	0.01
Methylene chloride (Dichloromethane)	75092	4.0	Tetramethyl lead	75741	0.01
4,4'-Methylenedianiline	101779	1.0	2,4-Toluene diamine	95807	0.002
Methylene diphenyl diisocyanate	101688	0.1	2,4-Toluene diisocyanate	584849	0.1
Methyl hydrazine	60344	0.006	o-Toluidine	95534	0.4
Methyl iodide (Iodomethane)	74884	1.0	Toxaphene (Chlorinated camphene)	8001352	0.006
Methyl isocyanate	624839	0.1	1,1,2-Trichloroethane	79005	0.1
Nickel carbonyl	13463393	0.1	Trichloroethylene	79016	1.0
Nitrobenzene	98953	1.0	2,4,5-Trichlorophenol	95954	1.0
4-Nitrobiphenyl	92933	1.0	2,4,6-Trichlorophenol	88062	0.6
4-Nitrophenol	100027	5.0	Trifluralin	1582098	0.9
			2,2,4-Trimethylpentane	540841	5.0
			Vinyl acetate	108054	1.0
			Vinyl bromide (Bromoethene)	593602	0.06
			Vinyl chloride	75014	0.02
			Vinylidene chloride (1,1-Dichloroethylene)	75354	0.04

* These values are based on the de minimis levels provided in the proposed rule-making, as published in the Federal Register on March 14, 1995 (60 FR 13664), pursuant to section 112(g) of the act using a 70-year lifetime exposure duration for all VHAP. Default assumptions and the de minimis values based on inhalation reference doses (RfC) are not changed by this adjustment.

^a Except for ethylene glycol butyl ether, ethylene glycol ethyl ether (2-ethoxy ethanol), ethylene glycol hexyl ether, ethylene glycol methyl ether (2-ethoxyethanol), ethylene glycol phenyl ether, ethylene glycol propyl ether, ethylene glycol mono-2-ethylhexyl ether, diethylene glycol butyl ether, diethylene glycol ethyl ether, diethylene glycol methyl ether, diethylene glycol hexyl ether, diethylene glycol phenyl ether, diethylene glycol propyl ether, triethylene glycol butyl ether, triethylene glycol ethyl ether, triethylene glycol methyl ether, triethylene glycol propyl ether, ethylene glycol butyl ether acetate, ethylene glycol ethyl ether acetate, and diethylene glycol ethyl ether acetate.

^b Except for benzo(b)fluoranthene, benzo(a)anthracene, benzo(a)pyrene, 7,12-dimethylbenz(a)anthracene, benz(c)acridine, chrysene, dibenz(ah)anthracene, 1,2:7,8-dibenzopyrene, indeno(1,2,3-cd)pyrene, but including dioxins and furans.