

Scenario 1: Use as intermediate

Section 1	Exposure Scenario Title
Title	2-Pyrrolidone use as chemical intermediate; CAS RN616-45-5
Use Descriptor	Sector of Use: Industrial (SU3)
	Process Categories: PROC2, PROC3, PROC8b, PROC9
	Environmental Release Categories: ERC6A
Processes, tasks, activities covered	Use as intermediate
Section 2	Operational conditions and risk management
	measures
<i>Field for additional statements to explain scenario if required.</i>	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product up to 100 % (unless stated differently) [G13].
Amounts used	<i>Not applicable</i>
Frequency and duration of use	Covers daily exposures up to 8 hours (unless stated differently) [G2]
Human factors not influenced by risk management	Not applicable
Other Operational Conditions affecting worker exposure	Assumes use at not $> 20^{\circ}$ C above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1].
Contributing Scenarios	Risk Management Measures Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1. Technical measures to prevent release, 2. Technical measures to prevent dispersion, 3. Organisational measures, 4. Personal protection. Phrases between brackets are good practice advice only, beyond REACH Chemical Safety Assessment and may be communicated in Section 5 of the ES or within the main sections of the SDS.

General exposures (closed systems) [CS15].	No specific measures identified [EI18]. {Handle
With sample collection [CS56]. With occasional	substance within a predominantly closed system
controlled exposure [CS137]	provided with extract ventilation [E49]}. {Provide a
	good standard of general or controlled ventilation
	(10 to 15 air changes per hour) [E40]}.;
	{Ensure material transfers are under containment or
	extract ventilation [E66]}. {Wear suitable gloves
	tested to EN374 [PPE15]}.
General exposures (closed systems) [CS15]. Use in	No specific measures identified [EI18]. {Provide a
contained batch processes [CS37].	good standard of general or controlled ventilation
	(10 to 15 air changes per hour) [E40]}. {Wear
	suitable gloves tested to EN374 [PPE15]}.
Process sampling [CS2].	Provide a good standard of general or controlled
	ventilation (10 to 15 air changes per hour)
	[E40].{Ensure operatives are trained to minimise
	exposures [EI119]}. {Wear suitable gloves tested to
	EN374 [PPE15]}.
Bulk transfers [CS14].	Provide a good standard of general or controlled
(open systems) [CS108]With potential for aerosol	ventilation (10 to 15 air changes per hour)
generation [CS138].	[E40].{Ensure operatives are trained to minimise
	exposures [EI119]}. {Wear suitable gloves tested to
	EN374 [PPE15]}.
Bulk transfers [CS14].	Provide a good standard of general or controlled
(closed systems) [CS107];	ventilation (10 to 15 air changes per hour)
	[E40].{Ensure operatives are trained to minimise
	exposures [EI119]}. {Wear suitable gloves tested to
	EN374 [PPE15]}.
Process sampling [CS2].	Provide extract ventilation to points where
	emissions occur [E54]. {Clear up spills
	immediately and dispose of waste safely [EI9]}.
	;{Avoid manual contact with wet work pieces
	[EI17]}. {Wear suitable gloves tested to EN374
	[PPE15]}.
Bulk transfers [CS14].	Provide extract ventilation to points where
	emissions occur [E54]. {Clear up spills
	immediately and dispose of waste safely [EI9]}.
	;{Avoid manual contact with wet work pieces
	[EI17]}. {Wear suitable gloves tested to EN374
	[PPE15]}.
Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried out in
	accordance to article 14.3, the registrant concludes
	that the substance does not meet the criteria for
	classification as dangerous for the environment;
	therefore risk characterisations for environmental
Section 2	endpoints were not developed.
Section 3	Exposure Estimation

3.1. Health	When the recommended risk management measures	
	(RMMs) and operational conditions (OCs) are	
	observed, exposures are not expected to exceed the	
	predicted DNELs and the resulting risk	
	characterisation ratios are expected to be less than	
	1.	
3.2. Environment	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes	
	that the substance does not meet the criteria for	
	classification as dangerous for the environment;	
	therefore risk characterisations for environmental	
	endpoints were not developed.	
Section 4	Guidance to check compliance with the	
	Exposure Scenario	
4.1. Health	Confirm that RMMs and OCs are as described.	
4.2. Environment	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes	
	that the substance does not meet the criteria for	
	classification as dangerous for the environment;	
	therefore risk characterisations for environmental	
	endpoints were not developed.	
Section 5	Additional good practice advice beyond the	
	REACH Chemical Safety Assessment	
Note: The measures reported in this section have not been taken into ac above. They are not subject to obligation laid down in Article 37 (4) of 1		
Control of Worker Exposure		
Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are {indicated} and	
	incorporated within the ES Section 2 or	
	consolidated into the main sections of the SDS.	
Control of environmental exposure		
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes	
	that the substance does not meet the criteria for	
	classification as dangerous for the environment;	
	therefore risk characterisations for environmental	
	endpoints were not developed.	

Scenario 2: Distribution of 2-Pyrrolidone

Table 2	Descri	ntion of	ES 2
I abic #	DUSCIT	phon of	

Section 1	Exposure Scenario Title
Title	Distribution of 2-Pyrrolidone;
	CAS RN616-45-5
Use Descriptor	Sector of Use: Industrial (SU8, SU9)
	Process Categories: PROC1, PROC2, PROC3, PROC4,
	PROC8a, PROC8b, PROC9, PROC15
	Environmental Release Categories: ERC1 (loading) ERC2
	(repacking)
Processes, tasks, activities covered	Loading (including marine vessel/barge, rail/road car and IBC
	loading) and repacking (including drums and small packs) of

	substance, including its distribution and associated laboratory activities
Section 2	Operational conditions and risk management measures
Field for additional statements to explain scenario if required.	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product up to 100 % (unless stated differently) [G13].
Amounts used	Not applicable
Frequency and duration of use	Covers daily exposures up to 8 hours (unless stated differently) [G2]
Human factors not influenced by risk management	Not applicable
Other Operational Conditions affecting worker exposure	Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1].
Contributing Scenarios	Risk Management Measures Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1. Technical measures to prevent release, 2. Technical measures to prevent dispersion, 3. Organisational measures, 4. Personal protection. Phrases between brackets are good practice advice only, beyond REACH Chemical Safety Assessment and may be communicated in Section 5 of the ES or within the main sections of the SDS.
General exposures (closed systems) [CS15]. e.g. In-line additive dosing equipment, in-line filter cleaning	No specific measures identified [EI18]. {Wear suitable gloves tested to EN374 [PPE15]}.
General exposures (closed systems) [CS15]. ; With sample collection [CS56]. With occasional controlled exposure [CS137]	No specific measures identified [EI18]. {Handle substance within a predominantly closed system provided with extract ventilation [E49]}. {Provide a good standard of general or controlled ventilation (10 to 15 air changes per hour) [E40]}. ; {Ensure material transfers are under containment or extract ventilation [E66]}. {Wear suitable gloves tested to EN374 [PPE15]}.
General exposures (closed systems) [CS15]. Use in contained batch processes [CS37]. General exposures (open systems) [CS16]. Batch process [CS55]. ; With sample collection [CS56].	No specific measures identified [EI18]. {Provide a good standard of general or controlled ventilation (10 to 15 air changes per hour) [E40]}. {Wear suitable gloves tested to EN374 [PPE15]}. Provide a good standard of general ventilation (not less than 3 to 5 air changes per hour). [E11].{Wear suitable gloves tested to EN374 [PPE15]}.
Process sampling [CS2].	No specific measures identified [EI18]. {Avoid dip sampling [E42]}. {Provide a good standard of general or controlled ventilation (10 to 15 air changes per hour) [E40]}. {Wear suitable gloves tested to EN374 [PPE15]}.

Laboratory activities [CS36].	No specific measures identified [EI18]. {Handle in a fume cupboard or under extract ventilation [E83]}.{Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14].;	Provide a good standard of general or controlled ventilation (10
(closed systems) [CS107]	to 15 air changes per hour) [E40].{Clear transfer lines prior to de- coupling [E39]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14].;	Provide a good standard of general or controlled ventilation (10
(open systems) [CS108]	to 15 air changes per hour) [E40].Ensure operatives are trained to minimise exposures [EI119]. {Clear transfer lines prior to de- coupling [E39]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Drum and small package filling [CS6].	Fill containers/cans at dedicated fill points supplied with local extract ventilation [E51]Ensure material transfers are under containment or extract ventilation [E66]. {Clear transfer lines prior to de-coupling [E39]}. {Provide a good standard of general or controlled ventilation (10 to 15 air changes per hour) [E40]}. {Put lids on containers immediately after use [E9]}.; {Clear spills immediately [C&H13]}.
	{Wear suitable gloves tested to EN374 [PPE15]}.
Equipment cleaning and maintenance [CS39].	Drain down and flush system prior to equipment break-in or maintenance [E55].{Transfer via enclosed lines [E52]}. {Apply vessel entry procedures including use of forced supplied air [AP15]}. {Wear suitable gloves tested to EN374 [PPE15]}. {Retain drain downs in sealed storage pending disposal or for subsequent recycle [ENVT4]}.
Storage [CS67]With occasional controlled exposure [CS137]	Store substance within a closed system [E84].{Locate bulk storage outdoors [E88]}.{Wear suitable gloves tested to EN374 [PPE15]}.
Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.
Section 3	Exposure Estimation
3.1. Health	When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1.
3.2. Environment	As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.
Section 4	Guidance to check compliance with the Exposure Scenario
4.1. Health	Confirm that RMMs and OCs are as described.

4.2. Environment	As a result of the hazard assessment carried out in accordance to		
4.2. Environment			
	article 14.3, the registrant concludes that the substance does not		
	meet the criteria for classification as dangerous for the		
	environment; therefore risk characterisations for environmental		
	endpoints were not developed.		
Section 5	Additional good practice advice beyond the REACH		
	Chemical Safety Assessment		
Note: The measures reported in this section have not b	een taken into account in the exposure estimates related to the exposure scenario		
above. They are not subject to obligation laid down in	above. They are not subject to obligation laid down in Article 37 (4) of REACH.		
Control of Worker Exposure			
Selection of relevant Contributing	Good practice RMM phrases are {indicated} and incorporated		
Scenario phrases	within the ES Section 2 or consolidated into the main sections of		
*	the SDS.		
Control of environmental exposure			
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried out in accordance to		
	article 14.3, the registrant concludes that the substance does not		
	meet the criteria for classification as dangerous for the		
	environment; therefore risk characterisations for environmental		
	endpoints were not developed.		

Scenario 3: Use in coatings (industrial)

Section 1	Exposure Scenario Title
Title	Use of 2-Pyrrolidone in Coatings
	(Industrial); CAS RN616-45-5
Use Descriptor	Sector of Use: Industrial (SU3)
	Process Categories: PROC1, PROC2,
	PROC3, PROC4, PROC5, PROC7,
	PROC8a, PROC8b, PROC10, PROC13,
	PROC15
	Environmental Release Categories: ERC 4
Processes, tasks, activities covered	Covers the use in coatings (paints, inks,
	adhesives, etc) including exposures during
	use (including materials receipt, storage,
	preparation and transfer from bulk and
	semi-bulk, application by spray, roller,
	spreader, dip, flow, fluidised bed on
	production lines and film formation) and
	equipment cleaning, maintenance and
	associated laboratory activities.
Section 2	Operational conditions and risk
	management measures
Field for additional statements to explain scenario if required.	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product
	up to 100 % (unless stated differently)
	[G13].

Amounts used	Not applicable
Frequency and duration of use	Covers daily exposures up to 8 hours (unless stated differently) [G2]
Human factors not influenced by risk management	Not applicable
Other Operational Conditions affecting worker exposure	Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1].
Contributing Scenarios	Risk Management Measures
Contributing Scenarios	Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1. Technical measures to prevent release, 2. Technical measures to prevent dispersion, 3. Organisational measures, 4. Personal protection. Phrases between brackets are good practice advice only, beyond REACH Chemical Safety Assessment and may be communicated in Section 5 of the ES or within the main sections of the SDS.
General exposures (closed systems) [CS15].	No specific measures identified [EI18]. {Wear suitable gloves tested to EN374 [PPE15]}.
General exposures (closed systems) [CS15]. With sample collection [CS56]. ; Use in contained systems [CS38].	No specific measures identified [EI18]. {Handle substance within a predominantly closed system provided with extract ventilation [E49]}. {Ensure material transfers are under containment or extract ventilation [E66]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Film formation - force drying (50 - 100°C). Stoving (>100°C). UV/EB radiation curing [CS94]	No specific measures identified [EI18]. {Ensure material transfers are under containment or extract ventilation [E66]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Mixing operations (closed systems) [CS29]. General exposures (closed systems) [CS15].	No specific measures identified [EI18]. {Ensure material transfers are under containment or extract ventilation [E66]}. {Wear suitable gloves tested to EN374 [PPE15]}.;
Film formation - air drying [CS95]	{Provide extract ventilation to points where emissions occur [E54]}. {Avoid manual contact with wet work pieces [EI17]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Preparation of material for application [CS96]Mixing operations (open systems) [CS30].	Provide extract ventilation to points where emissions occur [E54]. {Avoid manual
(open systems) [Coso].	ennissions occur [E34]. (Avoiu manual

	contect with wet work pieces [E117])
	contact with wet work pieces [EI17]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Spraying (automatic/robotic) [CS97]	Provide a good standard of general or
	controlled ventilation (10 to 15 air changes
	per hour) [E40]. {Carry out in a vented
	booth provided with laminar airflow
	[E59]}. {Wear chemically resistant gloves
	(tested to EN374) in combination with
	'basic' employee training [PPE16]}.;
	{Wear a respirator conforming to EN140
	with Type A filter or better [PPE22]}.
Manual [CS34]. Spraying [CS10].	Provide a good standard of general or
	controlled ventilation (10 to 15 air changes
	per hour) [E40]. {Wear chemically resistant
	gloves (tested to EN374) in combination
	with 'basic' employee training [PPE16]}.;
	{Wear a respirator conforming to EN140
	with Type A filter or better [PPE22]}.
Material transfers [CS3]. Non-dedicated facility [CS82]	Clear transfer lines prior to de-coupling
	[E39].{Provide extract ventilation to points
	where emissions occur [E54]}. {Wear
	suitable gloves tested to EN374 [PPE15]}.
Material transfers [CS3]. Dedicated facility [CS81]	Clear transfer lines prior to de-coupling
	[E39].{Provide extract ventilation to points
	where emissions occur [E54]}. {Wear
	suitable gloves tested to EN374 [PPE15]}.
Roller, spreader, flow application [CS98]	Minimise exposure by partial enclosure of
Koner, spreader, now appreadon [esse]	the operation or equipment and provide
	extract ventilation at openings [E60]. {Use
	long handled tools where possible [E50]}.
	{Carefully pour from containers [E62]}.
	{Wear suitable gloves tested to EN374
	[PPE15]].
Dipping, immersion and pouring [CS4].	Provide extract ventilation to points where
Dipping, minicision and pouring [C54].	emissions occur [E54]. {Clear up spills
	immediately and dispose of waste safely
	[EI9]}.;
	{Avoid manual contact with wet work pieces [EI17]}. {Wear suitable gloves
Laboratory activities [CS26]	tested to EN374 [PPE15]}.
Laboratory activities [CS36].	No specific measures identified [EI18].
	{Provide a good standard of general or
	controlled ventilation (10 to 15 air changes
	per hour) [E40]}.;
	{Handle in a fume cupboard or under
	extract ventilation [E83]}.{Wear suitable
	gloves tested to EN374 [PPE15]}.

Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 3	Exposure Estimation
3.1. Health	When the recommended risk management
5.1. Heath	measures (RMMs) and operational
	conditions (OCs) are observed, exposures
	are not expected to exceed the predicted
	DNELs and the resulting risk
	characterisation ratios are expected to be less than 1.
2.2 Engineering	As a result of the hazard assessment carried
3.2. Environment	
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 4	Guidance to check compliance with the
	Exposure Scenario
4.1. Health	Confirm that RMMs and OCs are as
	described.
4.2. Environment	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 5	Additional good practice advice beyond
	the REACH Chemical Safety Assessment
Note: The measures reported in this section have not been taken into account in above. They are not subject to obligation laid down in Article 37 (4) of REACH.	the exposure estimates related to the exposure scenario
Control of Worker Exposure	
Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are
	{indicated} and incorporated within the ES
	Section 2 or consolidated into the main
	sections of the SDS.
Control of environmental exposure	
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	TEVISITATIL CONCINCES THAT THE SHONIANCE OPEN I
	registrant concludes that the substance does
	not meet the criteria for classification as
R0718346 Version 1	-

risk characterisations for environmental
endpoints were not developed.

Scenario 4: Use in coatings (professional)

Table 4 Description of ES 4

Section 1	Exposure Scenario Title
Title	Use of 2-Pyrrolidone in Coatings (Prof.);
	CAS RN616-45-5
Use Descriptor	Sector of Use: Professional (SU22)
*	Process Categories: PROC1, PROC2,
	PROC3, PROC4, PROC5, PROC8a,
	PROC8b, PROC10, PROC11, PROC13,
	PROC15, PROC19
	Environmental Release Categories: ERC
	8A, ERC 8C, ERC 8D, ERC 8F
Processes, tasks, activities covered	Covers the use in coatings (paints, inks,
	adhesives, etc) including exposures during
	use (including materials receipt, storage,
	preparation and transfer from bulk and
	semi-bulk, application by spray, roller,
	spreader, dip, flow, fluidised bed on
	production lines and film formation) and
	equipment cleaning, maintenance and
	associated laboratory activities.
Section 2	Operational conditions and risk
	management measures
Field for additional statements to explain scenario if required.	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product
	up to 100 % (unless stated differently)
	[G13].
Amounts used	Not applicable
Frequency and duration of use	
Trequency and duration of use	Covers daily exposures up to 8 hours
requercy and duration of use	Covers daily exposures up to 8 hours (unless stated differently) [G2]
Human factors not influenced by risk management	
	(unless stated differently) [G2]
Human factors not influenced by risk management	(unless stated differently) [G2] Not applicable
Human factors not influenced by risk management	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient
Human factors not influenced by risk management	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];Assumes a good basic standard of
Human factors not influenced by risk management	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];Assumes a good basic standard ofoccupational hygiene is implemented [G1].
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];Assumes a good basic standard ofoccupational hygiene is implemented [G1].Risk Management Measures
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2] Not applicable Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1]. Risk Management Measures Note: list RMM standard phrases
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];Assumes a good basic standard ofoccupational hygiene is implemented [G1].Risk Management MeasuresNote: list RMM standard phrasesaccording to the control hierarchy
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2] Not applicable Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1]. Risk Management Measures Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1.
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2] Not applicable Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1]. Risk Management Measures Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1. Technical measures to prevent release, 2.
Human factors not influenced by risk management Other Operational Conditions affecting worker exposure	(unless stated differently) [G2]Not applicableAssumes use at not > 20oC above ambient[G15];Assumes a good basic standard ofoccupational hygiene is implemented [G1].Risk Management MeasuresNote: list RMM standard phrasesaccording to the control hierarchyindicated in the ECHA template: 1.Technical measures to prevent release, 2.Technical measures to prevent dispersion,

	good practice advice only, beyond REACH
	Chemical Safety Assessment and may be
	communicated in Section 5 of the ES or
	within the main sections of the SDS.
General exposures (closed systems) [CS15].	No specific measures identified [EI18].
	{Wear suitable gloves tested to EN374
	[PPE15]}.
General exposures (closed systems) [CS15]. With sample	No specific measures identified [EI18].
collection [CS56].;	{Ensure material transfers are under
Use in contained systems [CS38].	containment or extract ventilation [E66]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Film formation - force drying (50 - 100°C). Stoving (>100°C).	No specific measures identified [EI18].
UV/EB radiation curing [CS94]	{Ensure material transfers are under
	containment or extract ventilation [E66]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Mixing operations (closed systems) [CS29]. General exposures	No specific measures identified [EI18].
(closed systems) [CS15].	{Ensure material transfers are under
	containment or extract ventilation [E66]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.;
Film formation - air drying [CS95]	{Provide extract ventilation to points where
	emissions occur [E54]}. {Avoid manual
	contact with wet work pieces [EI17]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Preparation of material for application [CS96]Mixing operations	Provide extract ventilation to points where
(open systems) [CS30].	emissions occur [E54]. {Avoid manual
	contact with wet work pieces [EI17]}.
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Spraying (automatic/robotic) [CS97]	Clear transfer lines prior to de-coupling
	[E39].{Provide extract ventilation to points
	where emissions occur [E54]}. {Wear
	suitable gloves tested to EN374 [PPE15]}.
Manual [CS34]. Spraying [CS10].	Clear transfer lines prior to de-coupling
Ivianuar [C554]. Spraying [C510].	[E39].{Provide extract ventilation to points
	where emissions occur [E54]}. {Wear
	suitable gloves tested to EN374 [PPE15]}.
Material transfers [CS3]. Non-dedicated facility [CS82]	Clear transfer lines prior to de-coupling
	1 1 0
	[E39].{Use long handled tools where
	possible [E50]}. {Carefully pour from
	containers [E62]}. {Wear suitable gloves
Material transform [CS2] Dedicated facility [CS21]	tested to EN374 [PPE15]}.
Material transfers [CS3]. Dedicated facility [CS81]	Provide a good standard of general or
	controlled ventilation (10 to 15 air changes
	per hour) [E40]. {Provide extract ventilation

	to points where emissions occur [E54]}. {Wear suitable gloves tested to EN374
	[PPE15]}.
Roller, spreader, flow application [CS98]	Minimise exposure by partial enclosure of
	the operation or equipment and provide
	extract ventilation at openings [E60].
	{Wear suitable gloves tested to EN374
	[PPE15]}.
Dipping, immersion and pouring [CS4].	Provide extract ventilation to points where
	emissions occur [E54]. {Clear up spills
	immediately and dispose of waste safely
	[EI9]}.;
	{Avoid manual contact with wet work
	pieces [EI17]}. {Wear suitable gloves
	tested to EN374 [PPE15]}.
Laboratory activities [CS36].	Avoid manual contact with wet work
	pieces [EI17]. {Provide a good standard of
	general or controlled ventilation (10 to 15
	air changes per hour) [E40]}.;
	{Handle in a fume cupboard or under
	extract ventilation [E83]}.{Wear suitable
	gloves tested to EN374 [PPE15]}.
Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 3	Exposure Estimation
3.1. Health	When the recommended risk management
	measures (RMMs) and operational
	conditions (OCs) are observed, exposures
	are not expected to exceed the predicted
	DNELs and the resulting risk
	characterisation ratios are expected to be
	less than 1.
3.2. Environment	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 4	Guidance to check compliance with the
	Exposure Scenario

4.1. Health	Confirm that RMMs and OCs are as
	described.
4.2. Environment	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Section 5	Additional good practice advice beyond
	the REACH Chemical Safety Assessment
Note: The measures reported in this section have not been taken into accou above. They are not subject to obligation laid down in Article 37 (4) of REA	
Control of Worker Exposure	
Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are
	{indicated} and incorporated within the ES
	Section 2 or consolidated into the main
	sections of the SDS.
Control of environmental exposure	
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore
	risk characterisations for environmental
	endpoints were not developed.
Soonaria 5. Usa in asstings (consumpt)	enupoints were not developed.

Scenario 5: Use in coatings (consumer)

Table 5 Description of ES 5

Section 1	Exposure Scenario Title
Title	GES USES:
	Consumer Use of 2-Pyrrolidone in Inks and Toners (PC 18)
Sector of Use (SU code)	21 (Consumer Use)
Use Descriptor (PC codes)	PC 18
Processes, tasks, activities covered	DESCRIPTION:
	Use of 2-Pyrrolidone in Inks and Toners
Environmental Release Category	Refer to section 9.0 for description of relevant ERCs
Specific Environmental Release	No SPERCs used in this assessment
Category	
Section 2	Operational conditions and risk management measures
Field for additional statements to	As described below.
explain scenario if required -	
pending better understanding from	
ECHA	
Section 2.1	Control of consumer exposure
Product characteristics	
Physical form of product	liquid
Vapour pressure (Pa)	2

Concentration of substance in		Unless otherwise stated, cover concentrations up to 10%
product		[ConsOC1]
Amounts used		Unless otherwise stated, covers use amounts up to40g
		[ConsOC2];covers skin contact area up to 71,4cm2 [ConsOC5]
Frequency and duration of		Unless otherwise stated, covers use frequency up to 0 days per
use/exposure		year [ConsOC3];Unless otherwise stated, covers use frequency
		up to 1 times per day [ConsOC4];
Other Operational Conditions		Unless otherwise stated assumes use at ambient temperatures
affecting exposure		[ConsOC15]; assumes use in a 20 m ³ room [ConsOC11];
		assumes use with typical ventilation [ConsOC8].
Section 2.1.1 Product categories		
PC18_n: Ink and tonersInks and	OC	Exposure time: 2.2 h/day, once per day. One cartridge (40 g) per
toners.		day (printing of several hundred pages daily)
	RMM	Exposure modifier: Air exchange rate 0.6/h;

Scenario 6: Use in laboratories (industrial)

Table 6 Description of ES 6

Section 1	Exposure Scenario Title
Title	Use of Small Quantities 2-Pyrrolidone in
	Laboratory Settings (Industrial);
	CAS RN616-45-5
Use Descriptor	Sector of Use: Industrial (SU3)
	Process Categories: PROC10, PROC15
	Environmental Release Categories: ERC 4
Processes, tasks, activities covered	Use of the substance within laboratory
	settings, including material transfers and
	equipment cleaning
Section 2	Operational conditions and risk
	management measures
Field for additional statements to explain scenario if required.	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product
	up to 100 % (unless stated differently) [G13].
Amounts used	Not applicable
Frequency and duration of use	Covers daily exposures up to 8 hours (unless
	stated differently) [G2]
Human factors not influenced by risk management	Not applicable
Other Operational Conditions affecting worker exposure	Assumes use at not > 20 ^o C above ambient
	[G15];
	Assumes a good basic standard of
	occupational hygiene is implemented [G1].
Contributing Scenarios	Risk Management Measures
	Note: list RMM standard phrases according
	to the control hierarchy indicated in the
	ECHA template: 1. Technical measures to
	prevent release, 2. Technical measures to
	prevent dispersion, 3. Organisational
R0718346 Version 1	Revision date: 2017-04-25 Page 14 of 2

	magging 4 Dangen al most action Dhaga as
	measures, 4. Personal protection. Phrases
	between brackets are good practice advice
	only, beyond REACH Chemical Safety
	Assessment and may be communicated in
	Section 5 of the ES or within the main
	sections of the SDS.
Laboratory activities [CS36]. Small scale [CS61]. Handling	Handle in a fume cupboard or under extract
small quantities (<1000ml) for more than 4 hours/day - inside	ventilation [E83]. {Wear suitable gloves
fume cupboard.	tested to EN374 [PPE15]}.
Cleaning [CS47]. Rolling, Brushing [CS51].;	Provide a good standard of general or
Vessel and container cleaning [CS103]Cleaning equiment,	controlled ventilation (5 to 15 air changes per
glassware etc under general ventilation for 15 min - 1 hour/day	hour) [E40]. {Use long handled tools where
	possible [E50]}. {Carefully pour from
	containers [E62]}. {Wear chemically
	resistant gloves (tested to EN374) in
	combination with 'basic' employee training
	[PPE16]}.
Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried
	out in accordance to article 14.3, the
	registrant concludes that the substance does
	not meet the criteria for classification as
	dangerous for the environment; therefore risk
	characterisations for environmental
Section 3	endpoints were not developed.
Section 3 3.1. Health	endpoints were not developed. Exposure Estimation
Section 3 3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management
	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions
	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not
	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and
	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1.
	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints
3.1. Health 3.2. Environment	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.
3.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the
3.1. Health 3.2. Environment	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario
3.1. Health 3.2. Environment	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the
3.1. Health 3.2. Environment Section 4	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario
3.1. Health 3.2. Environment Section 4	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario Confirm that RMMs and OCs are as
3.1. Health 3.2. Environment Section 4 4.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario Confirm that RMMs and OCs are as described. As a result of the hazard assessment carried
3.1. Health 3.2. Environment Section 4 4.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario Confirm that RMMs and OCs are as described. As a result of the hazard assessment carried out in accordance to article 14.3, the
3.1. Health 3.2. Environment Section 4 4.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1.As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.Guidance to check compliance with the Exposure ScenarioConfirm that RMMs and OCs are as described.As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does
3.1. Health 3.2. Environment Section 4 4.1. Health	endpoints were not developed. Exposure Estimation When the recommended risk management measures (RMMs) and operational conditions (OCs) are observed, exposures are not expected to exceed the predicted DNELs and the resulting risk characterisation ratios are expected to be less than 1. As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed. Guidance to check compliance with the Exposure Scenario Confirm that RMMs and OCs are as described. As a result of the hazard assessment carried out in accordance to article 14.3, the

	characterisations for environmental endpoints were not developed.
Section 5	Additional good practice advice beyond the REACH Chemical Safety Assessment
Note: The measures reported in this section have not been taken into accor above. They are not subject to obligation laid down in Article 37 (4) of RE	ount in the exposure estimates related to the exposure scenario
Control of Worker Exposure	
Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are {indicated} and incorporated within the ES Section 2 or consolidated into the main sections of the SDS.
Control of environmental exposure	
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.

Scenario 7: Use in laboratories (professional)

Section 1	Exposure Scenario Title
Title	Use of Small Quantities of 2-Pyrrolidone in
	Laboratory Settings (Professional);
	CAS RN 616-45-5
Use Descriptor	Sector of Use: Professional (SU22)
	Process Categories: PROC10, PROC15
	Environmental Release Categories: ERC 8A
Processes, tasks, activities covered	Use of small quantities within laboratory
	settings, including material transfers and
	equipment cleaning.
Section 2	Operational conditions and risk
	management measures
Field for additional statements to explain scenario if required.	As described below.
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product up
	to 100 % (unless stated differently) [G13].
Amounts used	Not applicable
Frequency and duration of use	Covers daily exposures up to 8 hours (unless
	stated differently) [G2]
Human factors not influenced by risk management	Not applicable
Other Operational Conditions affecting worker exposure	Assumes use at not > 20 above ambient
	[G15];
	Assumes a good basic standard of
	occupational hygiene is implemented [G1].

Contributing Scenarios	Risk Management Measures
	Note: list RMM standard phrases according to
	the control hierarchy indicated in the ECHA
	template: 1. Technical measures to prevent
	release, 2. Technical measures to prevent
	dispersion, 3. Organisational measures, 4.
	Personal protection. Phrases between
	brackets are good practice advice only,
	beyond REACH Chemical Safety Assessment
	and may be communicated in Section 5 of the
	ES or within the main sections of the SDS.
Laboratory activities [CS36]. Small scale [CS61].;	Handle in a fume cupboard or under extract
Fume-cupboard Activity [CS139].	ventilation [E83]. {Wear suitable gloves tested
	to EN374 [PPE15]}.
Cleaning [CS47]. Rolling, Brushing [CS51].;	Provide a good standard of general or
Vessel and container cleaning [CS103]	controlled ventilation (5 to 15 air changes per
	hour) [E40]. {Use long handled tools where
	possible [E50]}. {Carefully pour from
	containers [E62]}. {Wear suitable gloves
	tested to EN374 [PPE15]}.
Section 2.2	Control of environmental exposure
	As a result of the hazard assessment carried
	out in accordance to article 14.3, the registrant
	concludes that the substance does not meet the
	criteria for classification as dangerous for the
	environment; therefore risk characterisations
	for environmental endpoints were not
	developed.
Section 3	Exposure Estimation
3.1. Health	When the recommended risk management
	measures (RMMs) and operational conditions
	(OCs) are observed, exposures are not
	expected to exceed the predicted DNELs and
	the resulting risk characterisation ratios are
	expected to be less than 1.
3.2. Environment	As a result of the hazard assessment carried
	out in accordance to article 14.3, the registrant
	concludes that the substance does not meet the
	criteria for classification as dangerous for the
	environment; therefore risk characterisations
	for any ironmontal and points wars not
	for environmental endpoints were not
Section 4	developed.
Section 4	developed.Guidance to check compliance with the
	developed.Guidance to check compliance with the Exposure Scenario
Section 4 4.1. Health	developed.Guidance to check compliance with the Exposure ScenarioConfirm that RMMs and OCs are as
4.1. Health	developed.Guidance to check compliance with the Exposure ScenarioConfirm that RMMs and OCs are as described.
	developed.Guidance to check compliance with the Exposure ScenarioConfirm that RMMs and OCs are as

	concludes that the substance does not meet the	
	criteria for classification as dangerous for the	
	environment; therefore risk characterisations	
	for environmental endpoints were not	
	developed.	
Section 5	Additional good practice advice beyond the	
	REACH Chemical Safety Assessment	
Note: The measures reported in this section have not been taken into account in the exposure estimates related to the exposure scenario above. They are not subject to obligation laid down in Article 37 (4) of REACH.		
Control of Worker Exposure		
Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are {indicated}	
	and incorporated within the ES Section 2 or	
	consolidated into the main sections of the	
	SDS.	
Control of environmental exposure		
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried	
	out in accordance to article 14.3, the registrant	
	concludes that the substance does not meet the	
	criteria for classification as dangerous for the	
	environment; therefore risk characterisations	
	for environmental endpoints were not	
	developed.	

Scenario 8: Use in polymer processing

Table 8 Description of ES 8	
Section 1	Exposure Scenario Title
Title	Use of 2_Pyrrolidone in Polymer Processing; CAS
	RN 616-45-5
Use Descriptor	Sector of Use: Industrial (SU10)
	Process Categories: PROC1, PROC2, PROC3, PROC4,
	PROC5, PROC6, PROC8a, PROC8b, PROC9,
	PROC13, PROC14
	Environmental Release Categories: ERC6D
Processes, tasks, activities covered	Polymer processing
Section 2	Operational conditions and risk management
	measures
Field for additional statements to explain scenario if	As described below.
required.	
Section 2.1	Control of worker exposure
Product characteristics	
Physical form of product	Liquid, vapour pressure < 0.5 kPa [OC3].
Concentration of substance in product	Covers percentage substance in the product up to 100 %
-	(unless stated differently) [G13].
Amounts used	Not applicable
Frequency and duration of use	Covers daily exposures up to 8 hours (unless stated
	differently) [G2]
Human factors not influenced by risk management	Not applicable

Other Operational Conditions affecting worker exposure	Assumes use at not > 20oC above ambient [G15]; Assumes a good basic standard of occupational hygiene is implemented [G1].
Contributing Scenarios	Risk Management Measures Note: list RMM standard phrases according to the control hierarchy indicated in the ECHA template: 1. Technical measures to prevent release, 2. Technical measures to prevent dispersion, 3. Organisational measures, 4. Personal protection. Phrases between brackets are good practice advice only, beyond REACH Chemical Safety Assessment and may be communicated in Section 5 of the ES or within the main sections of the SDS.
Bulk transfers [CS14]. (closed systems) [CS107].Bulk transfers of polymer prill/pellet etc to/from storage	No specific measures identified [EI18]. {Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14]. (closed systems) [CS107].Bulk transfers of polymer prill/pellet etc to/from storage	Ensure material transfers are under containment or extract ventilation [E66]. {Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14]. Semi-bulk transfers to/from storage e.g. IBCs, big bags	Provide a good standard of general or controlled ventilation (5 to 15 air changes per hour) [E40].{Clear transfer lines prior to de-coupling [E39]}. {Wear suitable gloves tested to EN374 [PPE15]}.
General exposures (closed systems) [CS15]. With sample collection [CS56]. In-line weighing of (bulk) polymer additives	No specific measures identified [EI18]. {Wear suitable gloves tested to EN374 [PPE15]}.
With sample collection [CS56]. small scale weighing of polymer additives	Provide extract ventilation to points where emissions occur [E54]. {Clear up spills immediately and dispose of waste safely [EI9]}. ;{Avoid manual contact with wet work pieces [EI17]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Additive premixing [CS92].Pre-mixing of polymer additives e.g. polyols	Provide extract ventilation to points where emissions occur [E54]. {Wear suitable gloves tested to EN374 [PPE15]}.
Additive premixing [CS92].batch pre-mixing of additives	Provide extract ventilation to points where emissions occur [E54]. {Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14]. transfer of polymer additives to calendars	Provide a good standard of general or controlled ventilation (5 to 15 air changes per hour) [E40].{Clear transfer lines prior to de-coupling [E39]}. {Wear suitable gloves tested to EN374 [PPE15]}.
Bulk transfers [CS14]. transfer of polymer additives to calendars	Provide extract ventilation to points where emissions occur [E54]. {Wear suitable gloves tested to EN374 [PPE15]}.
Calendering (including Banburys) [CS64].Calendaring activities	Provide extract ventilation to points where emissions occur [E54]. {Wear suitable gloves tested to EN374 [PPE15]}.

Production of articles by dipping and pouring	Provide a good standard of general or controlled	
[CS113].Formation of articles via polyol processes	ventilation (5 to 15 air changes per hour) [E40]. {Wear	
	suitable gloves tested to EN374 [PPE15]}.	
Extrusion and masterbatching [CS88].Extrusion	Ensure material transfers are under containment or	
and masterbatching of finished/ formulated polymer	extract ventilation [E66]. {Wear suitable gloves tested	
	to EN374 [PPE15]}.	
Injection moulding of articles [CS89].Article	No specific measures identified [EI18]. {Wear suitable	
formation (injection moulding)	gloves tested to EN374 [PPE15]}.	
Equipment maintenance [CS5]. build up and	Drain down system prior to equipment break-in or	
finishing of articles (dermal exposures)	maintenance [E65]. {Wear suitable gloves tested to	
	EN374 [PPE15]}.	
Storage [CS67]. With occasional controlled exposure	No specific measures identified [EI18]. {Avoid dip	
[CS137]	sampling [E42]}. {Provide extract ventilation to	
	material transfer points and other openings	
	[E82]}.{Wear suitable gloves tested to EN374	
	[PPE15]}.	
Section 2.2	Control of environmental exposure	
	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes that	
	the substance does not meet the criteria for classification	
	as dangerous for the environment; therefore risk	
	characterisations for environmental endpoints were not	
	developed.	
Section 3	Exposure Estimation	
3.1. Health	When the recommended risk management measures	
	(RMMs) and operational conditions (OCs) are observed,	
	exposures are not expected to exceed the predicted	
	DNELs and the resulting risk characterisation ratios are	
	expected to be less than 1.	
3.2. Environment	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes that	
	the substance does not meet the criteria for classification	
	as dangerous for the environment; therefore risk	
	characterisations for environmental endpoints were not	
	developed.	
Section 4	Guidance to check compliance with the Exposure	
4 1 H14h	Scenario	
4.1. Health	Confirm that RMMs and OCs are as described.	
4.2. Environment	As a result of the hazard assessment carried out in	
	accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification	
	as dangerous for the environment; therefore risk	
	characterisations for environmental endpoints were not developed.	
Section 5	A	
	Additional good practice advice beyond the REACH Chemical Safety Assessment -	
Note: The measures reported in this section have not been taken in	to account in the exposure estimates related to the exposure scenario	
above. They are not subject to obligation laid down in Article 37 (4) of REACH.		
Control of Worker Exposure		
	Version 1 Revision date: 2017-04-25 Page 20 of 21	

Selection of relevant Contributing Scenario phrases	Good practice RMM phrases are {indicated} and incorporated within the ES Section 2 or consolidated into the main sections of the SDS.
Control of environmental exposure	
Selection of relevant RMM Core Phrases	As a result of the hazard assessment carried out in accordance to article 14.3, the registrant concludes that the substance does not meet the criteria for classification as dangerous for the environment; therefore risk characterisations for environmental endpoints were not developed.