										Approval ex	xpires XX/XX/XXXX
Protection Agency Spe				Confidentia	ntial Statement of Product pecifications - DRAFT				Health	1	
				Speci					Canada	P	age of
Please rea	ad the instruction	าร	Please type	e or print clearly in blac	ck ink	Leave sh	aded areas blanl	K	This form	can be saved using A	lobe Acrobat
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Preliminary Questions	2. 3. PMRA For Canadian Registration			rmulation # & Version # 4. EPA Basic F			ormulation O		Date Rece		sion/File No.
	US Registration			OR Alternate Formulation N			ormulation No	. L			
	5. Is the product a Repack? 6.				product a M	icrobial?		7. Does t	he product have	e food uses?	
	8. What is the product type? Technical Grade Active Ingredient Integrated Systems Product End Use Product Manufacturing Concentrate										
					General	Information					
	9. Product Na	ame		10. Product Regn No. 11. Name of Applicant/Registrant							
Product											
	12. Formulation Type Code			13. Specific Gravity/Density Units 14. @ Temp Units			15. Weight/	/Formulated Piece	Units		
Properties				to							
Troperties	16. Flash Point Units 17. Flame Extension				n	Units 18. Viscosity (mPa(s))			19. pH Range		
Certification of Approving Official Company Code											
			"I cer	tify that all the inforn	mation conta	ined within this	form is true an	d complete"			
20. Is this CS	SPS being provid	ded on behalf of an	other Applicant/F	Registrant (3 rd Party (Confidential)	? Yes No					
21. Position	Title		22. N	ame				23. Signature			
24. Address								25. City			
26. Province	/State							27. Country		28. Postal	Code/ZIP
	,										
29. Date	30	. Phone No.	31. Fa	ax	32. ema	nil					
YYYY-MM-DD XXX-XXX-XXXX XXX-XXXX											
					Agency/	Office Review					
		Reviewer			7.50	- I - I - I - I - I - I - I - I - I - I		Povioveor			
	Date	Keviewei				Date					
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by

OMB Control No. 20XX-XXXX

OMB Control No. 20XX-XXXX Approval expires XX/XX/XXXX

Product Nam	е	Product Registration No. Date Received Submission/File No. 1.						
	YYYY-MM-DD Page							
Components								
Names		ne (ISO Proposed or Accepted) 38. Chemical Name (IUPAC or CAS name if applicable) 46						
Informatio n and Limits	39. Reg. or Sub/File No. 40.%Purity 41. CAS# 42. Purpose in Formulation	43. % LCL 44. % Nominal 45. % UCL %v						
Label Guarantee	47. Label Guarantee 48. Value 49. Units Certified Limits of Label Guarantee (for pure active ingredient) 50. LCL 51. UCL							
Microbial and Other	52. Culture Collection Deposit 53. Potency 54. Viability	55. Other information						
Admin.	ACT IN ACT OUT NACT LIST ACTIONS	PC Codes						
Names	34. Active Formulant/Inert Impurity 35. Is this a member of a set of multiple active guarantees, or alternate formulants/inerts? Yes Set 36. Trade Name 37. Common Name 37. C	ne (ISO Proposed or Accepted) 38. Chemical Name (IUPAC or CAS name if applicable) 46	6.					
Informatio n and Limits	39. Reg. or Sub/File No. 40.%Purity 41. CAS# 42. Purpose in Formulation	43. % LCL 44. % Nominal 45. % UCL %v						
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Admin.	ACT IN ACT OUT NACT LIST ACTIONS	PC Codes						
		56. Total weight (%)						

OMB Control No. 20XX-XXXX Approval expires XX/XX/XXXX

egistration No. Date Received	Submission/File No.	1
YYYY-MM-DD		Page of

			Sites & Suppliers		
57. Manufacturing Site Formulating Site Formulant Supplier	58. Row Number(s) of the Component(s)	59. Name 62. Province/State	60. Address 63. Country	61. City 64. Postal Code/ZIP	Company Code
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