Form Approved: OMB No. 0920-0109 Expiration Date: Xxx XX, 20XX

National Institute for Occupational Safety and Health National Personal Protective Technology Laboratory Respirator Branch



Standard Application Form for the Approval of Respirators Version 7

[C.1] Applicant-Assigned Reference Number: AAA_Sample

[C.3] Manufacturer Data: [C.2] Type of Applic	eation: New							
Does your organization currently hold any NIOSH approv	als? O Yes	• N	0					
Is this a CBRN application?	O Yes	• N	CBRN Type N/A					
Is this a SEI joint application? (CBRN/NFPA)	O Yes	• N	0					
Is this a SEI retrofit respirator?	O Yes	No	0					
[C.3] Manufacturer: Sample Status of Facility: Approval Holder [C.5] Application Representative: Sample								
[C.3] Address: Sample Sample Sample								
[C.3] Telephone: 111111 [C.3] Inte	rnet Address:							
[C.3] FAX: [C.15] Ship	[C.15] Shipping Number:							
[C.4] Manufacturing Site Name, if different from above:								
Has your organization submitted a request for appro at this manufacturing site at any time in the last 3 ye	• •	tor produ						

Public reporting burden of this collection of information is estimated to average 229 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333: ATTN: PRA(0920-0 109). Do not send the completed form to this address.

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[C.1] Applicant-Assigned Reference Number: AAA_Sample

[C.6] D	ate of Applic	ation:	07/14/2014											
[C.2] T	ype of Applic	ation:	New				[B.	2.4]	Previo	us Ta	ısk# (if	resubmittal):	
C.7] T	ype of Produc	et:	Air-Purifyii	ng										
[C.8]	Is this an a	mended	application?)	0	Yes	•	No						
[C.12/ C.8]	Is this dev	ice inter	nded for mine	e use?	0	Yes	•	No						
[C.8]	Is the approval of this application dependent upon the approval of an application that is in process? \bigcirc Yes													
	If yes, er	iter the	reference nur	nber of the	e app	olicatio	on in p	roces	ss?					
C.9]	Reason for A	Applicat	ion: Sample	application	on									
[C.10]	Approval Hi	story:												
[C.15]	Is testing r	equired	? • Yes	O No	<u> </u>	Do	you w	ant to	est sam	ples	returne	d? O Yes	• No	-
	If testing is	not req	uired, state w	hy:		If n	o, NI	OSH	will dis	spose	of sam	ples.		
	Type of AP I	Respirat	oe: Regular or: Particula oe: Filtering	nte Filteri	ng	nit (R	PU)							
			d? Non-pow	_	•									
			it: Tight fit											
	I	s this re	spirator fit-cl	neckable?	(• Ye	es	\circ N	0					
	rec	uires us	rator is fit-che e of ancillary for approval te	equipment										
			pirator conta ents been app							_		O No		
	I	Does thi	s respirator h	ave an exh	nalat	ion va	lve?	•	Yes	0	No			
	I	Does thi	s respirator h	ave an inh	alati	on val	ve?	0	Yes	•	No			

Standard Application Form for the Approval of Respirators Version 7 [C.1] Applicant-Assigned Reference Number: AAA_Sample Number of Filters: 1 Location of Filter: Facepiece-mounted Is the filter replaceable? O Yes No Comments: [C.12] Intended Protection and Safe Design: **Series and Level of Protection** N95 [C.13] Pre-submission tests that have been performed [C.14] Model Numbers: [C.15] Test Samples: [C.16] Quality Assurance Documentation: Title of QA Manual: Sample Revision: 1 Date of QA Manual: Has the QA Manual been previously accepted? O Yes ○ In Process No If in process, under which reference number was the QA Manual previously submitted? [C.17] Fee Data: Check Number: Check Date: Check Amount:

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[C.1] Applicant-Assigned Reference Number: AAA_Sample

[C.24] **Summary of Related Documents:**

I certify the information contained in this application is correct and that if approved, no further changes will be made to the product(s) without prior written approval of the National Institute for Occupational Safety and Health, Respirator Branch.

Signature of Authorized Representative