Form Approved: OIMB No. 0920-0109 Exp. Date: Nov. 30, 2007

## 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:

[C.3]	Manufacturer Data: [C.2] To	⁄pe of A	pplication	New	,				
Does	your organization currently hold any NI	OSH ap	provals?	0	Yes	0	No	]	
Is thi	s a CBRN application?			0	Yes	0	No	]CBR1	√ Туре
Is thi	s a SEI joint application? (CBRN/NFPA)	)		0	Yes	0	No	]	
Is thi	s a SEI nethofit nesonator?			0	Yes	0	No	]	
[C.5]	Manufacturer: Status of Facility: Application Representative: Address:								
[C.3]	Telephone:	[C.3] Internet Address: [C.15] Shipping Number:							
	Manufacturing Site Name, if different from above:	[CLL)	o rappare r	· unibe	<b>.</b>				
	Has your organization submitted a reque at this manufacturing site at any time in				espirat <b>Yes</b>	or pro			

Public reporting burden of this collection of information is estimated to average 64 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-24, Atlanta, Georgia 30333; ATTN: PRA (0920-0109). Do not send the completed form to this address.

## Standard Application Form for the Approval of Respirators Version 7

[C.1] Applicant-Assigned Reference Number:

	ate of Application:  (B.2.4) Previous Task#(if resubmittal):								
[C.7] T <sub>3</sub>	ype of Product:								
[C.8]	Is this an amended application? O Yes O No								
[C.12/ C.8]	Is this device intended for mine use? O Yes O No								
[C.8]	Is the approval of this application dependent upon the approval of an application O Yes O No that is in process?								
	If yes, enter the reference number of the application in process?								
[C.9] [C.10]	Reason for Application: Approval History:								
[C.R]	Is testing required? O Yes O No Do you want test samples returned? O Yes O No								
	If testing is not required, state why:  If no, NIOSH will dispose of samples.								
[C.11]	Respirator Description:								
[C.12]	Intended Protection and Safe Design:								
[C.13]	Pre-submission tests that have been performed								
[C.14]	Model Numbers:								
[C.15]	Test Samples:								
[C.16]	Quality Assurance Documentation:								
[C.17]	Fee Data:								
 [C.24]	Summary of Related Documents:								

Standard App	lication F	orm for th	e Approval	of Respirators	Version	7
[C.1]	Applicant-Ass	igned Referenc	e Number:			

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