

NIOSH Respiratory Protective Device Information

**NIOSH CA 2025-1099
September 2025**

Effective Immediately: Voluntary Rescission of 16 approvals for Interspiro USA, Inc.



**U.S. Centers for Disease
Control and Prevention**
National Institute for
Occupational Safety and Health

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The National Institute for Occupational Safety and Health (NIOSH) has honored a request by **Interspiro USA, Inc.** to rescind 16 approvals.

As of September 15, 2025, **any respirator marked with a NIOSH approval label and approval number listed in the table below is no longer NIOSH approved.** The NIOSH [Certified Equipment List](#) no longer includes these approval numbers:

Approval Number	Product Line or Model
13F - 0132	Spiromatic/Spirolite 4515
13F - 0132CBRN	Spiromatic SCBA Models S3, S4, S5, S6, 9030, 6630, 4530 and 4515
13F - 0133	Spiromatic
13F - 0133CBRN	Spiromatic, CBRN and NFPA 1981, 2007 Edition
13F - 0197	Spiromatic
13F - 0197CBRN	Spiromatic, CBRN and NFPA 1981, 2007 Edition
13F - 0199	Spiromatic/Spirolite 9030
13F - 0213	Spiromatic
13F - 0213CBRN	Spiromatic, CBRN and NFPA 1981, 2007 Edition
13F - 0375	SPIROTEK T4
13F - 0375CBRN	SPIROTEK T4, CBRN and NFPA 1981, 2007 Edition
13F - 0381	UNITOR 4515 (industrial) and SPIROTEK 4515 (NFPA)
13F - 0420	SPIROTEK
13F - 0420CBRN	SPIROTEK, CBRN and NFPA 1981, 2007 Edition
13F - 0421	SPIROTEK
13F - 0421CBRN	SPIROTEK, CBRN and NFPA 1981, 2007 Edition

Due to the voluntary rescission of these NIOSH approvals, **respirators bearing these NIOSH approval numbers may no longer be used, manufactured, assembled, sold, or distributed.**

Please reach out to Interspiro USA, Inc.. The [Certified Equipment List](#) can be used to locate other NIOSH Approved® respirators.

NIOSH Approved® is a certification mark of the U.S. Department of Health and Human Services (HHS) registered in the United States and several international jurisdictions