### Objective
To maintain the quality control of NIOSH approved respiratory protective devices in the field by investigating and resolving reports of product nonconformance issues in a timely manner.

### Stakeholders
- Respirator users
- Enforcement agencies
- Emergency Responders
- Respirator manufacturers worldwide

### Key Partners
- NFPA
- IAFF
- SEI
- MSHA
- OSHA
- FDA

### Project Scope
- Respond to external and internal reports.
- Establish and enforce accountability for nonconformances.
- Review and evaluate proposed resolutions from to ensure conformance 42 CFR, Part 84.

### FY17 Milestones
- Timely initiation of CPIPs in response to nonconformance reports, ongoing.
- Develop means for obtaining more permanent support for investigations and outreach.
- Improve database capabilities for metrics.

### Outputs CY2016
- 12 CPIPs opened
- 17 CPIPs completed

### Outputs CY2017 (through 2/28/17)
- 8 CPIP opened
- 4 CPIP completed

### Outcomes
- Ensure that NIOSH approved respiratory devices continue to meet or exceed the criteria on which the approval was based.
- Improved respirator components.

Updated: 28 Feb 2017