

EXHIBIT S7 TO DECLARATION OF
STEPHEN G. SCHWARZ IN SUPPORT OF
PLAINTIFFS' MOTION FOR CLASS
CERTIFICATION



Certified Mail

September 27, 2007

NO CBI

Document Processing Center
EPA East - Room 6428 Attn: Section 8(e)
Office of Pollution Prevention and Toxics
US EPA
1200 Pennsylvania Avenue NW
Washington DC 20460-0001

Re: TSCA 8(e) Substantial Risk Notice: Supplemental to Docket No. EHQ-0598-373;
Sulfonate-based and Carboxylic-based Fluorochemicals

To whom it may concern:

3M is submitting this notice to supplement its previous submissions on sulfonate-based and carboxylic-based fluorochemicals. Enclosed is a final report detailing the results of a cohort mortality study of workers at the former 3M ammonium perfluorooctanoate (APFO) production facility in Cottage Grove, MN. While 3M does not believe that these data are reportable under the TSCA 8(e) reporting criteria, a decision was made to provide this report to the EPA, recognizing the ongoing work to assess fluorochemical exposure pathways and potential risks.

The purpose of cohort mortality study was to examine possible associations between working in jobs with varying exposure to APFO and specific causes of death. Conclusions of the authors were as follows:

"APFO exposed workers did not have an elevated risk of death when compared to the population of the state of Minnesota, however, within the cohort risk of death from prostate cancer and cerebrovascular disease was elevated for workers with higher estimated exposure. Interpreting the somewhat contradictory results requires caution and consideration of several assumptions. *A priori* causes of death selected based on animal toxicology studies, liver, pancreatic and testicular cancer and cirrhosis of the liver, were not observed to be associated with APFO exposure."

If you have any questions, please contact Deanna Luebker at (651) 737-1374 or djluebker@mm.com.

Sincerely,

Katherine E. Reed
Staff Vice President, Environmental, Health and Safety Operations

Enclosure

TABULAR LIST

- 429.4 Rupture of papillary muscle
- ① 429.7 Certain sequelae of myocardial infarction, not elsewhere classified
 Use additional code to identify the associated myocardial infarction:
 with onset of 3 weeks or less (429.80-429.82)
 with onset of more than 3 weeks (429.83)
- Excludes:** congenital defects of heart (742, 743)
 coronary aneurysm (424.22)
 disorders of papillary muscle (429.6, 429.8V)
 postmyocardial infarction syndrome (421.0)
 rupture of chordae tendinae (429.2)
- 429.70 Acquired cardiac septal defect
- Excludes:** acute septal infarction (429.80-429.82)
- 429.78 Other
 Mixed (ventricular septal) (ventricular), acquired, following myocardial infarction
- ① 429.8 Other ill-defined heart diseases
- 429.80 Other disorders of papillary muscle
- | | |
|-------------------|-------------------|
| Papillary muscle: | Papillary muscle: |
| atrophy | incompetence |
| degeneration | incoordination |
| dysfunction | scarring |
- 429.82 Hypertensive heart disease
- 429.88 Other
 Carditis
- Excludes:** that due to hypertension (402.0-402.9)
- 429.9 Heart disease, unspecified
 Heart disease (organic) NOS
 Morbus cordis NOS
- Excludes:** that due to hypertension (402.0-402.9)

CEREBROVASCULAR DISEASE (430-438)

- Includes:** with mention of hypertension (conditions classifiable to 401-402)
- Use additional code, if desired, to identify presence of hypertension
- Excludes:** any condition classifiable to 430-434, 436, 437 occurring during pregnancy, childbirth, or the puerperium, or specified as postpartum (674.0)
- 430 Subarachnoid hemorrhage
- Meningeal hemorrhage
- Eloquent
- berry aneurysm
 (congenital) cerebral aneurysm NOS
- Excludes:** ruptured ruptured cerebral aneurysm (544.87)
- 431 Intracerebral hemorrhage
- | | |
|------------------|-----------------------------------|
| Hemorrhage (of): | Hemorrhage (of): |
| basilar | supratentorial |
| bulbar | posterior |
| cerebellar | subarachnoid |
| cerebral | ventricular |
| corticostriatal | Eloquent of blood vessel in brain |
| cortical | |
| internal capsule | |
- ① 432 Other and unspecified intracranial hemorrhage
- 432.0 Nontraumatic extradural hemorrhage
 Nontraumatic epidural hemorrhage
- 432.1 Subdural hemorrhage
 Subdural hematoma, nontraumatic
- 432.8 Unspecified intracranial hemorrhage
 Intracranial hemorrhage NOS

- ① 433 Occlusion and stenosis of 2
 The following fifth-digit code:
 0 without mention of
 1 with cerebral infarct
- Includes:**
 embolism
 narrowing
 obstruction
 thrombosis
- Excludes:** insufficiency 2
- ① 433.8 Basilar artery
- ① 433.1 Carotid artery
- ① 433.2 Vertebral artery
- ① 433.3 Multiple and bilateral
- ① 433.4 Other specified artery
- ① 433.9 Unspecified proximal
 Proximal artery 2
- ① 434 Occlusion of cerebral artery
 The following fifth-digit code:
 0 without mention of
 1 with cerebral infarct
- ① 434.0 Cerebral thrombosis
 Thrombosis of cereb
- ① 434.1 Cerebral embolism
- ① 434.9 Cerebral artery with
- ① 435 Transient cerebral ischemia
Includes: arteriovascular
 episodes
 insufficiency 0
 spasm of cereb
- Excludes:** acute cerebrom
 that due to any
- 435.0 Basilar artery syndrome
- 435.1 Vertebral artery syndrome
- 435.2 Subclavian steal syndrome
- 435.3 Vertebrobasilar artery
- 435.4 Other specified trans
- 435.9 Unspecified transient
 Impending cerebrov
 Ischemic cerebral
 Transient ischemic 2
- ① 436 Aneurysm, but ill-defined, cerebral
 Aneurysm, unspecified
 NOS
 atretic
 cerebral
 aneurysm
- Excludes:** any condition 1
- ① 437 Other and ill-defined cerebral
- 437.0 Cerebral arteriovenous
 Arteriovenous of cerebral
 Cerebral arteriovenous
- ① 437.1 Other specified for
 Acute arteriovenous
 Cerebral arteriovenous
- 437.2 Hypertensive aneurysm

CIRCULATORY SYSTEM

433 Occlusion and stenosis of procrural arteries

The following fifth-digit subclassification is for use with category 433:

0 without mention of cerebral infarction

1 with cerebral infarction

Includes:

<ul style="list-style-type: none"> stenosis narrowing obstruction thrombosis 	}	of basilar, carotid, and vertebral arteries
--	---	---

Excludes: insufficiency NOS of procrural arteries (423.0-423.9)

- 433.0 Basilar artery
- 433.1 Carotid artery
- 433.2 Vertebral artery
- 433.3 Multiple and bilateral
- 433.8 Other specified procrural artery
- 433.9 Unspecified procrural artery
Procrural artery NOS

434 Occlusion of cerebral arteries

The following fifth-digit subclassification is for use with category 434:

0 without mention of cerebral infarction

1 with cerebral infarction

- 434.0 Cerebral thrombosis
Thrombosis of cerebral arteries
- 434.1 Cerebral embolism
- 434.9 Cerebral artery occlusion, unspecified

435 Transient cerebral ischemia

Includes: cerebrovascular insufficiency (acute) with transient focal neurological signs and symptoms
insufficiency of basilar, carotid, and vertebral arteries
spasm of cerebral arteries**Excludes:** acute cerebrovascular insufficiency NOS (437.2)
that due to any condition classifiable to 433 (433.0-433.9)

- 435.0 Basilar artery syndrome
- 435.1 Vertebral artery syndrome
- 435.2 Subclavian steal syndrome
- 435.3 Vertebrobasilar artery syndrome
- 435.8 Other specified transient cerebral ischemia
- 435.9 Unspecified transient cerebral ischemia
Impending cerebrovascular accident
Intermittent cerebral ischemia
Transient ischemic attack (TIA)

436 Acute, but ill-defined, cerebrovascular disease

<ul style="list-style-type: none"> Apparent, cryptic: NOS attack cerebral attack 	<ul style="list-style-type: none"> Cerebral attack Cerebrovascular accident (CVA) NOS Stroke
---	---

Excludes: any condition classifiable to categories 430-433

437 Other and ill-defined cerebrovascular disease

- 437.0 Cerebral arteriovenous
Anomalous of cerebral arteries
Cerebral arteriovenous
- 437.1 Other generalized ischemic cerebrovascular disease
Acute cerebrovascular insufficiency NOS
Cerebral ischemia (chronic)
- 437.2 Hypertensive encephalopathy

Add 4th or 5th digit	Nonspecific code	Unspecified code	Morbidity code	291
----------------------	------------------	------------------	----------------	-----

TABULAR LIST

- 437.3 Cerebral aneurysm, unruptured
 Internal carotid artery, intracranial portion
 Internal carotid artery NOS
Excludes congenital cerebral aneurysm, unruptured (747.81)
 internal carotid artery, extracranial portion (942.81)

437.4 Cerebral arteritis

437.5 Moyamoya disease

437.6 Nonpyogenic thrombosis of intracranial venous sinus

Excludes pyogenic (122)

437.7 Transient global amnesia

437.8 Other

437.9 Unspecified

Cerebrovascular disease or lesion NOS

438 Late effects of cerebrovascular disease

Note: This category is to be used to indicate conditions in 430-437 as the cause of late effects. The "late effects" include conditions specified as such, or sequelae, which may occur at any time after the onset of the causal condition.

438.0 Cognitive deficits

⊖ 438.1 Speech and language deficits

438.10 Speech and language deficit, unspecified

438.11 Aphasia

438.12 Dysphasia

438.19 Other speech and language deficits

⊖ 438.2 Hemiplegia/hemiparesis

438.20 Hemiplegia affecting unspecified side

438.21 Hemiplegia affecting dominant side

438.22 Hemiplegia affecting nondominant side

⊖ 438.3 Monoplegia of upper limb

438.30 Monoplegia of upper limb affecting unspecified side

438.31 Monoplegia of upper limb affecting dominant side

438.32 Monoplegia of upper limb affecting nondominant side

⊖ 438.4 Monoplegia of lower limb

438.40 Monoplegia of lower limb affecting unspecified side

438.41 Monoplegia of lower limb affecting dominant side

438.42 Monoplegia of lower limb affecting nondominant side

⊖ 438.5 Other paralytic syndromes

Use additional code to identify type of paralytic syndrome, such as:
 locked-in state (344.11)
 quadriplegia (344.00-344.09)

Excludes late effects of cerebrovascular accident with:
 hemiplegia/hemiparesis (438.20-438.22)
 monoplegia of lower limb (438.40-438.42)
 monoplegia of upper limb (438.30-438.32)

438.50 Other paralytic syndromes affecting unspecified side

438.51 Other paralytic syndromes affecting dominant side

438.52 Other paralytic syndromes affecting nondominant side

438.59 Other paralytic syndromes, bilateral

⊖ 438.6 Other late effects of cerebrovascular disease

438.61 Apraxia

438.62 Dysphagia

438.69 Other late effects of cerebrovascular disease

Use additional code to identify the late effect

438.9 Unspecified late effects of cerebrovascular disease

DISEASES OF ARTERIES, ART

440 Atherosclerosis

Includes: arteriosclerosis
 arteriosclerosis /
 arteriosclerosis +
 atherosclerosis
 degenerative:
 arterial
 arterio-vascular
 vascular
 arteriosclerosis
 arterio-arteriosclerosis
 arterio-arteriosclerosis

Excludes arteriosclerosis

440.0 Of aorta

440.1 Of renal artery

Excludes arteriosclerosis /

⊖ 440.2 Of native arteries of l

Excludes arteriosclerosis /

440.20 Atherosclerosis

440.21 Atherosclerosis

440.22 Atherosclerosis

Includes: an

440.23 Atherosclerosis

Includes: an

Use additional

440.24 Atherosclerosis

Includes: an

with ischem

Excludes post ganglion /

440.29 Other

⊖ 440.3 Of bypass graft of th

Excludes arteriosclerosis /

anastomosis
 anastomosis
 of graft (996.74)

440.30 Of unspecified

440.31 Of aortopathy

440.32 Of vasculitis

440.8 Of other specified art

Excludes basilar (433.0)

coronary (432.1)

coronary (437.0)

coronary (434.1)

mesenteric (522)

pericardial (432)

pulmonary (431)

vertebral (433.1)

440.9 Generalized and unsp

Arteriosclerosis NOS

Excludes arteriosclerosis

441 Aortic aneurysm and dissec

Excludes cystic aortic

aneurysm

⊖ 441.0 Dissection of aorta

Dissecting aortic

441.00 Unspecified

441.01 Thoracic

441.02 Abdominal

TABLEULAR LIST

CONGENIT

747.41	Total anomalous pulmonary venous connection Total anomalous pulmonary venous return (TAPVR) subdiaphragmatic supradiaphragmatic	
747.42	Partial anomalous pulmonary venous connection Partial anomalous pulmonary venous return	
747.49	Other anomalies of great veins Absence Congenital aneurysm Fistula left posterior cardinal vein left superior vena cava Scimitar syndrome Transposition of pulmonary veins (NOI)	of vein (see [diabetic] response)
747.5	Absence or hypoplasia of artificial artery Single artificial artery	
3 747.6	Other anomalies of peripheral vascular system Absence Anomaly Atresia Arteriovenous aneurysm (peripheral) Arteriovenous malformation of the peripheral vascular system Congenital aneurysm (peripheral) phlebotomy aneurysm, artery vein Multiple renal arteries	of artery or vein, not elsewhere classified
	Excludes anomaly of: cerebral vessels (747.81) pulmonary artery (747.3) congenital renal aneurysm (743.29) hemangioma (238.00-238.09) lymphangioma (228.7)	
747.60	Anomaly of the peripheral vascular system, unspecified site	
747.61	Gastrointestinal vessel anomaly	
747.62	Renal vessel anomaly	
747.63	Upper limb vessel anomaly	
747.64	Lower limb vessel anomaly	
747.69	Anomalies of other specified sites of peripheral vascular system	
3 747.8	Other specified anomalies of circulatory system 747.81 Anomalies of cerebrovascular system Arteriovenous malformation of brain Cerebral arteriovenous aneurysm, congenital Congenital anomalies of cerebral vessels Includes ruptured cerebral (arteriovenous) aneurysm (430) 747.82 Spinal vessel anomaly Arteriovenous malformation of spinal vessel 747.89 Other Aneurysm, congenital, specified site not elsewhere classified Excludes congenital aneurysm: coronary (745.27) peripheral (747.6) pulmonary (747.3) renal (743.30)	
747.9	Unspecified anomaly of circulatory system	
3 748	Congenital anomalies of respiratory system Excludes congenital defect of diaphragm (739.4)	
748.0	Choanal atresia Absence Congenital stenosis	of nares (nasal) (posterior)

748.1	Other anomalies of nose Absence nose Accessory nose Cleft nose Deficiency of wall of nasal cavity Excludes congenital deviation of nasal septum	C
748.2	Web of larynx Web of larynx: NOI glottis subglottic	
748.3	Other anomalies of larynx, trachea Absence or agenesia of: trachea larynx trachea Anomaly (soft): cricoid cartilage epiglottis thyroid cartilage laryngeal cartilage Absence (NOI): epiglottis glottis larynx trachea Cleft thyroid cartilage, congenital	C
748.4	Congenital cystic lung Divertic lung Cystic, congenital pulmonary, congenital Excludes acquired or unspecified cyst	I
748.5	Agenesis, hypoplasia, and dysplasia Absence of lung (Ductus) (Duct) Aplasia of lung Hypoplasia of lung (Duct) Sequestration of lung	
3 748.6	Other anomalies of lung 748.60 Anomaly of lung, unspecified 748.61 Congenital bronchomalacia 748.69 Other Accessory lung (Duct) Accessory lobe (Ductus), 4	I
748.8	Other specified anomalies of respiratory system Abnormal communication between Anomaly, pleural folds Atria of nasopharynx Congenital cyst of mediastinum	
748.9	Unspecified anomaly of respiratory system Anomaly of respiratory system, 7	
3 749	Cleft palate and cleft lip 3 749.0 Cleft palate 749.00 Cleft palate, unspecified 749.01 Unilateral, complete 749.02 Unilateral, incomplete Cleft uvula 749.03 Bilateral, complete 749.04 Bilateral, incomplete	
3 749.1	Cleft lip Chelidactyls Congenital fissure of lip	

430 ● Code new to this edition ▲ Revision of existing code ④ ⑤ Fourth or fifth digit required

3 749.1 Add 4th or 5th digit **3 749.1** Nonspecific code

TABULAR LIST

- 441.0 **Thoracoabdominal**
- 441.1 Thoracic aneurysm, ruptured
- 441.2 Thoracic aneurysm without mention of rupture
- 441.3 Abdominal aneurysm, ruptured
- 441.4 Abdominal aneurysm without mention of rupture
- 441.5 Aortic aneurysm of unspecified site, ruptured
Rupture of aorta NOS
- 441.6 Thoracoabdominal aneurysm, ruptured
- 441.7 Thoracoabdominal aneurysm, without mention of rupture
- 441.9 Aortic aneurysm of unspecified site without mention of rupture
Aneurysm
Dissection
Hyaline aneurysm | of aorta
- 442 Other aneurysm**
Includes: aneurysm (ruptured) (closed) (false) (vascular)
aneurysmal cysts
Excludes: arteriovenous aneurysm or fistula
acquired (447.0)
congenital (747.80-747.89)
traumatic (900.0-904.8)
- 442.0 Of artery of upper extremity
- 442.1 Of radial artery
- 442.2 Of iliac artery
- 442.3 Of artery of lower extremity
Aneurysm
Internal
popliteal | artery
- 442.4 Of other specified artery
- 442.51 Artery of neck
Aneurysm of carotid artery (common) (internal) (external, external portion)
Excludes: internal carotid artery, nonvascular portion (437.2)
- 442.52 Subclavian artery
- 442.53 Splenic artery
- 442.54 Other visceral artery
Aneurysm
celiac
gastrohepatic
gastrosplenic
hepatic
pancreatoduodenal
superior mesenteric | artery
- 442.59 Other
Aneurysm
mediastinal
splenic | artery
- Excludes: cerebral (unruptured) (437.2)
congenital (747.81)
ruptured (430)
secondary (414.17)
heart (414.00)
pulmonary (417.1)
- 442.9 Of unspecified site
- 443 Other peripheral vascular disease**
- 443.0 Raynaud's syndrome
Raynaud's
disease
phosphenes (secondary)
Use additional code, if desired, to identify ganges (785.4)
- 443.1 Thromboangiitis obliterans (Buerger's disease)
Frodo's ganges

CIRCULAT

- 443.2 Other specified peripheral vasculitis
- 443.3 Peripheral angiopathy in diabetes mellitus (250.7)
- 443.39 Other
Acromegaly
Acroparesthesia
Mongle (Schaban's type)
vasomotor (Nikolsky's)
Erythrocyanosis
Erythromelalgia
Excludes: cellulitis (861.1)
psoriasis (710.2-710.3)
scurvy (261.4)
- 443.9 Peripheral vascular disease, unspecified
Inconsistent classification NOS
Peripheral
angiopathy NOS
vascular disease NOS
Spasm of artery
Excludes: arterioarteriosclerosis of the anterior
portion of cerebral artery (433)
- 444 Arterial embolism and thrombosis**
Includes: embolism
embolic
thrombotic
occlusion
Excludes: the complicating
abortion (634.42) with A, B,
or C or molar pregnancy /
pregnancy, childbirth, or the
- 444.0 Of abdominal aorta
Aortic fibrillation syndrome Left
Aorticiliac obstruction Right
- 444.1 Of thoracic aorta
Ectasia or dilatation of aorta (86)
- 444.2 Of arteries of the extremities
- 444.21 Upper extremity
- 444.22 Lower extremity
Arterial embolism or thrombosis
Internal
popliteal NOS
popliteal
Excludes: bifurcated (444.81)
- 444.3 Of other specified artery
- 444.31 Iliac artery
- 444.39 Other
Excludes: basilar (433.0)
cervical (433.1)
cervical (434.0-434.9)
coronary (410.00-410.92)
mesenteric (537.0)
ophthalmic (362.30-362.34)
peroneal (413.0-413.9)
pulmonary (415.19)
renal (582.81)
renal (582.80-582.84)
vertebral (433.2)
- 444.9 Of unspecified artery
- 445 Polyarteritis nodosa and related conditions**

Final Report

Mortality Study of Workers Employed at the 3M Cottage Grove Facility

Submitted by:

Bruce H. Alexander, PhD

Division of Environmental and Occupational Health

School of Public Health

University of Minnesota

Mayo Mail Code 807

Mayo Building

420 Delaware Street S.E.

Minneapolis, MN 55455

April 26, 2001

Summary

Objective:

To determine whether occupational exposure to perfluorooctanoic acid (PFOA) and other fluorochemicals is related to the mortality experience of employees of the 3M facility in Cottage Grove, Minnesota.

Methods:

All employees who accrued at least one year of employment at Cottage Grove were eligible for inclusion in the study. Cohort members were assigned to one of three exposure groups based on their work history: non-exposed, probable exposure to PFOA, and definite exposure to PFOA. The cohort was followed through December 31, 1997. Death certificates were obtained for all known deaths and coded for analysis. Standardized mortality ratios were estimated for all cause and cause specific mortality using mortality rates from the general population of Minnesota as a reference. SMR estimates were made for the sub-cohorts ever exposed to PFOA, by exposure category, and for a minimum of one year of exposure.

Results:

There were 3,992 eligible cohort members who accrued 108,198 person-years of follow-up and 607 deaths. Forty-six of the deaths occurred in the sub-cohort with definite PFOA exposure. The all cause and all cancer mortality rates for the entire study population, and for the exposure sub-cohorts were less than expected in the general population. There was no association between exposure to PFOA or other fluorochemicals and cancer of the liver, kidney, or prostate or cirrhosis of the liver. A modest increased risk of death from cerebrovascular disease (CVD) was observed in the definite PFOA exposure subcohort (5 observed, 1.94 expected, SMR=2.58, 95% CI=0.84-6.03). A dose response relationship between PFOA or other fluorochemical exposure and CVD was not apparent.

Conclusion:

Employees of the Cottage Grove facility were not observed to have an excess risk of mortality from cancer in relation to PFOA and other fluorochemical exposure. The association observed between PFOA exposure and CVD was modest, but unexpected. The association may be due to some occupational exposure although there is no biologically plausible mechanism identified at this time.

Introduction

The Cottage Grove manufacturing facility of the Minnesota Mining and Manufacturing Corporation (3M) has produced perfluorinated compounds since 1947. A primary product from this plant is ammonium perfluorooctanoate ($\text{CF}_3(\text{CF}_2)_6\text{CO}_2\text{NH}_4^+$, APFO), a potent synthetic surfactant used in industrial applications. APFO rapidly dissociates in biologic media to perfluorooctanoate ($\text{CF}_3(\text{CF}_2)_6\text{CO}_2^-$, PFOA) which is the anion of perfluorooctanoic acid ($\text{CF}_3(\text{CF}_2)_6\text{COOH}$). In laboratory animals, PFOA and its salts are: 1) absorbed by ingestion, inhalation or dermal contact;¹⁻³ 2) distributed primarily in the liver and blood;⁴ 3) not biotransformed, conjugated or incorporated into lipids;^{3,8} and 4) eliminated in the female rat at a greater rate of excretion than the male rat.⁷ In rats, administration of APFO resulted in peroxisome proliferation, uncoupling of mitochondrial oxidative phosphorylation and altered lipid metabolism.^{4,9} In lifetime feeding bioassays of rats, APFO in the diet at 300 ppm (daily dose of 15 mg/kg/day) increased the incidence of liver Leydig cell and pancreas acinar cell adenomas.¹⁰ The liver tumors most likely occurred via the nongenotoxic mechanisms of oxidative stress and apoptosis. Increased hepatic aromatase activity may have resulted in a hormone-mediated mechanism (increased estradiol) for the formation of the Leydig cell tumors.^{11,12} The pancreas acinar cell adenomas have been hypothesized to be a result of a mild but sustained increase in cholecystokin (CCK) levels secondary to hepatic cholestasis.¹³ In a 90-day gavage study of rhesus monkeys, mortality was pronounced prior to end of study in the 100 mg/kg/day and 30 mg/kg/day dose groups.¹⁴ Histopathologic examination revealed marked diffuse lipid depletion in the adrenals, slight to moderate hypocellularity of bone marrow and moderate atrophy of lymphoid follicles. No histopathologic changes were reported in the 3 and 10 mg/kg/day dose groups. A recently completed 6-month gavage study of cynomolgus primates demonstrated a steep dose response curve.¹⁵ Both the low (3 mg/kg/day) and mid (10 mg/kg/day)

dose groups resulted solely in increased liver weights. The highest dose group (30/20 kg/mg/day) resulted in severe toxicity, which required the removal of treatment for some of the high dose group animals. The exact mechanism of toxicity in the primate remains to be elucidated.

Hepatic toxicity, hypolipidemia and abnormal hormone levels (e.g., estradiol) have not been associated with the PFOA levels measured in male APFO production workers.¹⁸⁻²⁰ However, it should be noted that the serum concentration (50 ppm) associated with liver enlargement in the 3.0 mg/kg/day dose group of the cynomolgus primate study is within the range experienced by workers with higher occupational exposure.^{15,18,20} A retrospective cohort mortality study of workers engaged in APFO production at the Cottage Grove facility found no significantly increased cause-specific standardized mortality ratio although a two-fold nonsignificant increase in prostate cancer mortality, based on 4 observed deaths was reported.²¹ This report summarizes the results of an update of that cohort mortality study with a specific emphasis on exposure to PFOA.

Methods

Cohort Enumeration

The cohort for this study was enumerated using employment records from the Cottage Grove facility. Workers accruing at least 1 year of cumulative employment at the Cottage Grove facility as of December 31, 1997 were eligible for inclusion in the cohort. A review of employee work history records by 3M personnel identified workers eligible for the cohort. The records of any Cottage Grove employee with at least one year of cumulative employment were abstracted to record the worker's name, Social Security Number, 3M identification number, date of birth, and the dates of any entry on the work history record, including layoffs and leaves of absence.

Information about each job was abstracted wherever available, including the department codes, and job classifications. The names, Social Security Numbers, 3M identification numbers, and dates of birth were recorded for workers with less than one year of cumulative employment for comparison with the original cohort. The abstracted data were entered into a computer database and provided to University of Minnesota investigators.

The newly enumerated cohort was linked to records from the original cohort to update the employment information and verify names, Social Security Numbers dates of birth and dates of death. Discrepancies identified in the records were resolved using TRW/Experian, a credit reporting agency, and the Social Security Administration service for epidemiologic research studies. The latter reports the most recent account activity of an individual and whether they are recorded as deceased in the Social Security Death Index (SSDI). Duplicate records due to name changes or incorrect data were eliminated.

Investigators at the University of Minnesota reviewed the eligibility for inclusion in the cohort. To be eligible for the cohort a worker had to accrue at least 365 days of cumulative employment at the Cottage Grove site. The eligibility of each cohort member was determined by summing his or her dates of employment, exclusive of periods of absence due to illness, military leave, maternity leave, or layoff. Currently employed workers were assigned December 31, 1997 as their last date of employment.

Follow-up and Determination of Vital Status

Eligible cohort members were followed from the day they accrued 365 days of cumulative employment till December 31, 1997 or their date of death. Vital records searches were performed for all cohort members not employed by 3M on December 31, 1997, or for whom a death certificate was not obtained in the original study. The National Death Index (NDI) was searched for all workers in the original study and new workers included in the cohort. The Social Security Administration data and/or the SSDI were searched to verify the vital status of workers who terminated employment before 1979.

The records of cohort members identified as deceased through the NDI or SSDI were reviewed by hand to ensure a valid match and a copy of the death certificate was requested from the state of record. A licensed nosologist coded the death certificates to the International Classification of Disease Version 8. A second licensed nosologist coded the death certificate using the rules for the ICD version in effect at the time of death. This second coding was used for verification and enabled the use of actual (unadjusted) mortality reference data.

Exposure Assessment

The goal of this study was to describe mortality experience in relation to fluorochemical exposure. Of particular interest was exposure to PFOA. The areas in the Cottage Grove facility where PFOA and other fluorochemicals were produced changed over the years. Because the department codes used to classify the work areas also changed over years it was not possible to assign the workers to exposure categories on work history information alone. To ascertain exposure status the department codes were reviewed to determine the building and division

assigned to each code. These lists were then reviewed independently by a panel of veteran workers and plant industrial hygienists to determine where fluorochemical production or the development of fluorochemical products took place throughout the history of the Cottage Grove Facility. The responses of the individual reviews were summarized and the panel was convened as a group to discuss the exposure assignments. The available information limited the panel's ability to classify each department with certainty, thus general classifications of exposure were adopted. The job history information was classified into the three following groups;

- Definite PFOA exposure (potentially high). These jobs included workers employed in the areas where cell generation, drying, shipping, and packaging of PFOA occurred throughout the history of the plant.
- Probable PFOA exposure. These jobs include other chemical division jobs where exposure to PFOA was possible, but with lower or transient exposures.
- Not exposed to fluorochemicals. Primarily non-chemical division jobs.

Hereafter these exposure subgroups will be referred to as definite exposure, probable exposure and nonexposed, respectively.

Potential for exposure to other fluorochemicals was possible in the chemical division. In particular, there has been production of some salts of perfluorooctane sulfonic acid, which may disassociate to perfluorooctanesulfonate (PFOS, $C_8F_{17}SO_2$). However, this did not usually occur where definite PFOA exposure in the workplace was likely, as the production areas were in different buildings. It is feasible that employees who had worked in jobs with definite PFOA exposure may have transferred to this other building in the course of their career.

Analysis

The mortality experience of the Cottage Grove cohort was compared to that of the general population of the state of Minnesota. Mortality reference rates from seven regional counties (Hennepin, Ramsey, Anoka, Carver, Dakota, Scott, and Washington) were also used to rule out large variations based on regional mortality reporting differences. Reference data were obtained from the Mortality Population Data System (MPDS) center at the University of Pittsburgh. These data are derived from National Center for Health Statistics data and provide all cause mortality and malignant neoplasm rates back to 1940, and non-malignant cause specific death rates from 1962 forward. These reference data are age (5 year), gender, race, and calendar period (5 year) specific and are coded using the rules for the ICD version in effect for the calendar period.

Standardized mortality ratios (SMR) were computed for all cause and cause specific deaths using the Minnesota reference data. The expected number of deaths for all cause and malignant neoplasm deaths were estimated for all years. The expected number of deaths from non-malignant causes was computed for the years 1962-1997. Observed deaths and person-years in the denominator occurring before the reference data were available were excluded from the analysis. The SMRs and appropriate 95% confidence intervals were computed using the PC Life Table Analysis System (PCLTAS) software developed by the National Institutes of Occupational Safety and Health (NIOSH).²² This program computes age, gender, and race specific SMR using standard life table methods. The expected number of deaths are estimated by multiplying the age, gender, race, and calendar period tabulated person-years of follow-up to the corresponding cause specific mortality reference rates. No data on race were available for the cohort; therefore, the reference data were limited to the mortality rates for white Minnesotans.

The all cause and cause specific SMRs were initially computed for the entire cohort and the subcohorts with definite exposure, probable exposure, and not exposed to fluorochemicals. A more exposure specific analysis stratified workers based on a one-year minimum employment in jobs with definite PFOA exposure and definite or probable exposure. The latter included workers who accrued one year of employment with a combination of definite and probable exposed jobs.

Four causes of death potentially related to PFOA exposure based on laboratory animal data and the earlier cohort study, prostate cancer, liver cancer, kidney cancer, and cirrhosis of the liver, were analyzed by duration of exposure in each fluorochemical exposure subgroup. Other causes of death that appeared to be in excess in one or more of the fluorochemical exposed groups were also evaluated by duration of exposure.

Results

Of the 6678 individual workers identified at the Cottage Grove plant, 3992 employees met the one year inclusion criteria. Of these, 12 percent (492) worked at least one day in areas where definite exposure to PFOA occurred. Forty-two percent (1685) had probable exposure, but not definite PFOA exposure, and the remaining 45% (1815) were not exposed to fluorochemicals (Table 1). The latter are non-chemical division jobs at Cottage Grove. Male workers made up 80% of the cohort, but were 92% of the PFOA exposed cohort. The average age at follow up was slightly less in the PFOA exposed cohort, but the average duration of employment at Cottage Grove was slightly longer. There were 607 deaths identified in the cohort, 46 deaths in the definite PFOA exposure group and 267 in the probable PFOA exposure group. Death certificates were obtained for 590 of the decedents (97%). Six of the missing death certificates were in the probable PFOA exposure group and the rest were in the non-exposed group. More extensive exposure to PFOA, based on a one-year minimum employment in definitely exposed jobs, occurred to 182 workers (17 deaths), and 1673 workers had definite or probable exposure for at least one year (219 deaths) (Table 2).

The all cause and cause specific mortality rates for the entire cohort were lower than expected compared to the general population: 607 observed and 715 expected (SMR=0.85, 95% CI=0.78-0.92) (Table 3). A similar pattern was observed for all deaths from cancer; 172 observed, 204 expected (SMR=0.84, 95% CI = 0.72-0.98). Deaths from all causes and all cancers were fewer than expected for the exposure subcohorts (Table 4-6), and for the strata limited to workers with a minimum of one year of definite exposure (Table 7), or a combination of definite or probable exposure (Table 8).

There was no association observed between fluorochemical exposure and cancer of the prostate, liver, kidney, or from cirrhosis of the liver (Table 4-8). In the definite PFOA exposure subcohort only one death from prostate cancer was observed (0.77 expected). Five deaths from prostate cancer were observed in the probable PFOA exposure group (5.8 expected), and another 2 observed in the non-exposed sub-cohort (6.8 expected). Only one cancer of the liver was observed and that was in the probable PFOA exposure group. Again, limiting the cohort to a minimum of one year of exposure did not alter the results.

Overall, nonmalignant causes of death did not exceed that expected in Minnesota. Deaths from cerebrovascular disease (CVD) did exceed the number of expected in the definite PFOA exposed cohort; 5 observed and 1.94 expected (SMR=2.58, 95% CI 0.84-6.03). Deaths from CVD were not elevated in the rest of the cohort. Three CVD deaths occurred in the subcohort with definite exposure for at least one year, where 0.89 deaths were expected (SMR=3.36, 95% CI=0.69, 9.82). It is plausible that the coding of CVD deaths varies by region, where a CVD death in an older person may be reported as "Natural Causes" on the death certificate, which would receive a different ICD code. To verify these results the CVD deaths were compared to the local county mortality rates. The results were essentially the same.

To further evaluate the distribution of CVD deaths, the SMRs in the definite PFOA exposure sub-cohort were stratified by duration of employment in PFOA exposed jobs (Table 9), and exposure weighted time of employment (Table 10). In the exposure-weighted time of employment weights of 0, 1 and 3 were assigned respectively to the non-exposed, probable

PFOA exposed, and definite PFOA exposed jobs in the work histories. The weighted time of exposure was derived by multiplying the duration of employment in the exposed jobs, in days, by the weighting factor. The results for the years of employment in PFOA exposed jobs did not reveal a dose-response relationship between PFOA exposure and the risk of CVD; however, the SMR for high PFOA exposure for five or more years was 6.9 (95% CI = 1.39-20.24). This, however, was based on only three cases, and no deaths from CVD occurred among workers with ten or more years in high PFOA jobs (Table 9). The weighted exposure analysis, which includes information from the workers with probable exposure to PFOA indicated that the fluorochemical exposed workers with less than 10,000 exposure days experienced fewer than expected deaths from CVD. The SMR for those with 10,000 or more exposure days was 3.32 (95% CI=0.89-8.49). An exposure days value of 10,000 equates to 27 years of exposure in probable PFOA exposed jobs or 9 years in definite PFOA exposed jobs.

The number of deaths from traumatic injuries was less than expected for the entire cohort, however, the frequency of deaths from violence was modestly elevated in the definite exposure cohort. Five of the six violent deaths were suicides (2.1 expected, SMR 2.33, 95% CI=0.76-5.45) (Table 4).

Discussion

This updated mortality study evaluated the mortality experience of workers with at least one year of employment at the 3M Cottage Grove facility, with specific attention to exposure to PFOA.

No excess mortality was observed for malignant neoplasms or for all causes of death. There were modest elevations in the risk of death from cerebrovascular disease and deaths due to violence in the higher PFOA exposed workers, but these results are based on very few cases.

Some limitations must be considered when interpreting the results of this mortality analysis.

Although several methods of follow-up were employed to ascertain deaths in this cohort, the possibility remains that some deaths were not accounted for in the analysis. A death certificate was not obtained for seventeen known decedents; thus they were not included in the cause specific death analysis. The extent to which these limitations would affect the results is unknown, however most of the missing death certificates were in the nonexposed sub-cohort.

Another limitation of this study is the lack of employee specific exposure data for PFOA and other fluorochemicals. The determination of potential exposure to these compounds was made using all available information from work histories and expert input from veteran workers and plant industrial hygienists. Nevertheless, some misclassification of exposure was likely.

Maintenance and other mobile workers not specifically identified as definitely PFOA exposed workers may have routinely entered the PFOA exposed sites, and a few workers assigned to the PFOA exposure areas may not have spent much time in those areas. The extent to which this misclassification occurred and the attendant effects on the study results remain unknown.

This study differs from the analysis published by Gilliland and Mandel²³ by the study inclusion criteria and the exposure definition. The earlier study required six months of cumulative

employment for inclusion, while the current study required one year. The change was made primarily to exclude the relatively large number of short-term workers. Workers who left after only six months on the job were likely to have different underlying risk factors than the long-term workers. The Gilliland and Mandel analysis limited the exposure assignment to Chemical Division/Non-Chemical Division and assumed duration of employment in the chemical division equated with exposure to PFOA. The current analysis was driven by more recent toxicological evaluations of the compound, and specifically categorizes PFOA exposure as definite and probable within the Chemical Division as only certain areas and tasks within the Chemical Division would have led to high exposure to PFOA. . . Another difference of note between the two analyses is the inclusion of 169 additional cohort members in the current study that, according to available employment data, were eligible for both studies.

The two analyses of this cohort differ by the results for prostate cancer and cerebrovascular disease. The previous analysis identified 6 cases of prostate cancer, four of which were in the Chemical Division. The current analysis identified eight cases of prostate cancer, one in the definite PFOA exposure group and five in the probable PFOA exposure group. In neither of the exposure groups did the number of prostate cancer deaths exceed the expected number. One case of prostate cancer identified in the previous study was not included in the current analysis because the worker did not meet the one-year minimum employment criteria.

Although Gilliland and Mandel considered an association between PFOA exposure and prostate cancer as biologically plausible based on the animal and human data,^{17,20} subsequent research,¹⁸

as well as the present study findings would argue that, at this time, there is not an association observed in this workforce.

The result for cerebrovascular disease is difficult to interpret. There were 13 CVD deaths in the previous analysis and 36 in the current study. There was no excess of CVD deaths in the non-exposed group or the probable PFOA exposure group. In fact the SMRs were well below 1.0. There was not an apparent dose-response relationship, however such an analysis was hampered by the relatively few cases available to analyze. The lack of an association in the probable-PFOA exposure group and the absence of a change by using the local counties as a reference suggest that this is not an artifact of death certificate coding. CVD may be related to life style factors including smoking. It is noteworthy that the SMRs for all heart disease (1.08) and lung cancer (1.17) were at or above unity in the PFOA exposed sub-cohort. These diseases may be markers for smoking related illness. Heart disease and CVD are almost always below unity in epidemiologic studies of chemical workers. Therefore, these SMRs reported in the definite PFOA exposure group are unexpected. The observed association may be due to some occupational exposure at the Cottage Grove facility, although there is no biologically plausible mechanism identified. At this time a causal association cannot be drawn between exposure to PFOA and death from cerebrovascular disease.

The absence of measurable adverse health effects from PFOA exposure was also reported in earlier studies on this population. PFOA exposure did not alter circulating levels of reproductive hormones¹⁸ and a study of the effects of PFOA on markers of liver function did not detect frank hepatotoxic effect.^{18,20}

Recommendations

There does not appear to be a clear association between employment at the Cottage Grove plant and risk of mortality from cancer or other causes. However, due to the previous observation of an association with prostate cancer, the apparent excess occurrence of death from cerebrovascular disease, and the evolving understanding of the toxicology of PFOA, continued mortality follow-up of this cohort is warranted.

References

1. Griffith FD, Long JE. Animal toxicity studies with ammonium perfluorooctanoate. *Am Ind Hyg Assoc J* 1980;41:576-583.
2. Kennedy G. Dermal toxicity of ammonium perfluorooctanoate. *Toxicol Appl Pharm* 1985;81:348-355.
3. Kennedy G, Hall G, Britelli J, Chen H. Inhalation toxicity of ammonium perfluorooctanoate. *Fd Chem Toxicol* 1986;24:1325-1329.
4. Vanden Heuvel J, Kuslikis B, Van Refelghem M, Peterson R. Tissue distribution, metabolism and elimination of perfluorooctanoic acid. *J Biochem Toxicol* 1991;6:83-92.
5. Orphaug R, Singer L. Metabolic handling of perfluorooctanoic acid in rats. *Proc Soc Exp Biol Med* 1980;163:19-23.
6. Pastoor T, Lee K, Perri M, Gillies P. Biochemical and morphological studies of ammonium perfluorooctanoate-induced hepatomegaly and peroxisome proliferation. *Exp Mol Path* 1987;47:98-109.
7. Hanhijarvi H, Phaug R, Singer L. The sex-related difference in perfluorooctanoate excretion in the rat. *Proc Soc Exp Biol Med* 1982;171:51-55.
8. Keller B, Marxman D, Popp J, Thurman R. Several nongenotoxic carcinogens uncouple mitochondrial phosphorylation. *Biochim Biophys Acta* 1991;1102:237-244.
9. Haugbom B, Spydevold O. The mechanism underlying the hypolipemic effect of perfluorooctanoic acid, perfluorooctane sulphonic acid (PFOSA) and clofibrate acid. *Biochimica et Biophysica Acta* 1992;1128(1):65-72.
10. Sibinski L. Two-year oral (diet) toxicity/carcinogenicity study of fluorochemical FC-143 in rats. St. Paul: Riker Laboratories, 1987.
11. Cook J, Murray S, Frame S, Hartt M. Induction of Leydig cell adenomas by ammonium perfluorooctanoate: a possible endocrine-related mechanism. *Toxicol Appl Pharm* 1992;113:209-217.
12. Biegel L, Liu R, Hartt M, Cook J. Effects of ammonium perfluorooctanoate on Leydig cell function: in vitro, in vivo, and ex vivo studies. *Toxicol Appl Pharm* 1995;134:18-25.
13. Obourn J, Frame S, Bell R, Longnecker D, Elliott G, Cook J. Mechanisms for the pancreatic oncogenic effects of the peroxisome proliferator Wyeth-14,643. *Toxicol Appl Pharm* 1997;145:425-436.
14. Goldenthal E, Jessup D, Geil R, Mehring J. Ninety-day subacute rhesus monkey toxicity study. Mattawan, MI: International Research Development Corp, 1987.

15. Butenhoff J, Costa G, Elcombe C, Farrar D, Hansen K, Iwai H, Jung R, Kennedy G, Lieder P, Olsen G, Thornford P. Toxicity of ammonium perfluorooctanoate (APFO) in cynomolgus monkeys after 26 weeks of oral dosing. *Toxicol Sci* 2001;in preparation.
16. Ubel F, Sorenson S, Roach D. Health status of plant workers exposed to fluorochemicals: a preliminary report. *Am Ind Hyg Assoc* 1980;41:584-589.
17. Gilliland F. Fluorocarbons and Human Health: Studies in Occupational Cohort [Doctoral Dissertation]. University of Minnesota, 1992.
18. Gilliland F, Mandel J. Serum perfluorooctanoic acid and hepatic enzymes, lipoproteins and cholesterol: a study of occupationally exposed men. *Am J Ind Med* 1996;29:560-568.
19. Olsen GW, Gilliland FD, Burlew MM, Burris JM, Mandel JS, Mandel JH. An epidemiologic investigation of reproductive hormones in men with occupational exposure to perfluorooctanoic acid. *Journal of Occupational & Environmental Medicine* 1998;40(7):614-22.
20. Olsen GW, Burris JM, Burlew MM, Mandel JH. Plasma cholecystokinin and hepatic enzymes, cholesterol and lipoproteins in ammonium perfluorooctanoate production workers. *Drug and Chemical Toxicology* 2000;23:603-620.
21. Gilliland FD, Mandel JS. Mortality among employees of a perfluorooctanoic acid production plant. *Journal of Occupational Medicine* 1993;35(9):950-4.
22. National Institutes for Occupational Safety and Health. PC LTAS: Life table analysis system for use on the PC. Cincinnati: U.S. Department of Health and Human Services, 1998.

Table 1. Characteristics of 3M employees with one or more years of employment at Cottage Grove.

	Definite PFOA exposure ^a	Probable PFOA Exposure ^b	Non-exposed ^c	Total
Total	492	1685	1815	3992
Gender				
M	452 (92%)	1387 (83%)	1344 (74%)	3183 (80%)
F	40 (8%)	298 (17%)	471 (26%)	809 (20%)
Mean age at follow-up	52.0	57.4	57.0	56.6
Median age at follow-up	50.6	57.8	57.6	57.0
Mean year at birth	1944	1938	1938	1938
Median year at birth	1946	1938	1938	1939
Mean years at CG	16.6	14.5	8.6	12.1
Median years at CG	14.2	10.7	4.5	7.2
Person years of follow-up	10703	44295	49188	108198
Deaths	46	267	294	607

a: Ever employed in job with definite (high) PFOA exposure

b: Ever employed in a job with probable to other fluorochemicals including low PFOA exposure, but never in a job with definite exposure.

c: Primarily non-Chemical Division.

Table 2. Characteristics of Cottage Grove workers with definite PFOA exposure and definite or probable PFOA exposure for a minimum of one year.

	Definite PFOA exposure ^a	Definite or Probable PFOA exposure ^b
Total	182	1673
Deaths	17	219
Person years	3897	41487
Gender		
M	168 (92%)	1442 (86%)
F	14 (8%)	231 (14%)
Mean age at follow-up	53	56
Mean year of birth	1943	1940
Mean years at Cottage Grove	17.8	15.7
Mean years of exposure	6.2	9.4

a: Definite (high) PFOA exposure for at least one year.

b: Definite or Probable PFOA or other fluorochemical exposure for at least one year. Includes workers who accrued one year of exposure with definite and probable jobs combined.

Table 3. Cause specific deaths and standardized mortality ratios for selected causes of death for all Cottage Grove employees.

Cause	Observed	Expected	SMR	95% CI
All Deaths	607	715.13	0.85	0.78-0.92
Cancers				
All Malignant Neoplasms	172	203.96	0.84	0.72-0.98
Buccal Cavity and Pharynx	2	4.16	0.48	0.06-1.74
Digestive Organs and Peritoneum	42	50.38	0.83	0.60-1.13
Esophagus	3	5.34	0.56	0.12-1.64
Stomach	4	6.31	0.63	0.17-1.62
Large Intestine	19	18.18	1.04	0.63-1.63
Rectum	2	3.93	0.51	0.06-1.84
Biliary Passages and Liver Primary	1	4.43	0.23	0.01-1.25
Pancreas	12	10.78	1.11	0.57-1.94
All Other Digestive	1	1.40	0.71	0.02-3.97
Respiratory System	56	61.44	0.91	0.69-1.18
Larynx	2	1.80	1.11	0.13-4.02
Bronchus, Trachea, Lung	53	58.97	0.90	0.67-1.18
Breast	6	8.60	0.70	0.25-1.52
Female Reproductive	3	5.23	0.57	0.12-1.68
Male Reproductive	9	14.27	0.63	0.29-1.20
Prostate	8	13.41	0.60	0.26-1.18
Testis and Other Male Genital Organs	1	0.86	1.16	0.03-6.47
Urinary Organs	8	9.89	0.81	0.35-1.59
Kidney	3	6.06	0.49	0.10-1.45
Bladder and Other Urinary Organs	5	3.83	1.31	0.42-3.05
Malignant Melanoma of Skin	4	3.24	1.24	0.34-3.16
All Lymphatic and Hematopoietic Tissue	18	23.47	0.77	0.45-1.21
Non-malignant causes				
Cerebrovascular Disease	26	35.91	0.72	0.47-1.06
All Heart Disease	195	234.49	0.83	0.72-0.96
Nonmalignant Respiratory Disease	29	45.92	0.63	0.42-0.91
Cirrhosis of Liver	11	14.33	0.77	0.38-1.37
Nephritis and Nephrosis	2	4.12	0.48	0.06-1.75
Accidents	32	46.32	0.69	0.47-0.98
Motor Vehicle Accidents	19	22.08	0.86	0.52-1.34
All Other Accidents	13	24.24	0.54	0.29-0.92
Violence	17	22.45	0.76	0.44-1.21
Suicides	13	18.23	0.71	0.38-1.22
Homicides	4	4.22	0.95	0.26-2.42

Table 4. Cause specific deaths and standardized mortality ratios for selected causes of death for Cottage Grove employees ever employed in jobs with definite PFOA exposure.

Cause	Observed	Expected	SMR	95% CI
All Deaths	46	50.14	0.92	0.67-1.22
<u>Cancers</u>				
All Malignant Neoplasms	11	13.79	0.80	0.40-1.43
Buccal Cavity and Pharynx	0	0.31	0.00	0.00-11.81
Digestive Organs and Peritoneum	3	3.44	0.87	0.18-2.55
Esophagus	0	0.42	0.00	0.00-8.86
Stomach	0	0.42	0.00	0.00-8.85
Large Intestine	2	1.20	1.67	0.20-6.02
Rectum	0	0.26	0.00	0.00-13.97
Biliary Passages and Liver Primary	0	0.30	0.00	0.00-12.12
Pancreas	1	0.75	1.34	0.03-7.42
All Other Digestive	0	0.09	0.00	0.00-40.54
Respiratory System	5	4.45	1.12	0.36-2.63
Larynx	0	0.13	0.00	0.00-27.97
Bronchus, Trachea, Lung	5	4.26	1.17	0.38-2.74
Breast	0	0.18	0.00	0.00-20.31
Female Reproductive	0	0.09	0.00	0.00-40.65
Male Reproductive	1	0.85	1.17	0.03-6.51
Prostate	1	0.77	1.30	0.03-7.20
Testis and Other Male Genital Organs	0	0.08	0.00	0.00-45.03
Urinary Organs	0	0.71	0.00	0.00-5.22
Kidney	0	0.47	0.00	0.00-7.82
Bladder and Other Urinary Organs	0	0.23	0.00	0.00-15.72
Malignant Melanoma of Skin	0	0.30	0.00	0.00-12.27
All Lymphatic and Hematopoietic Tissue	0	1.70	0.00	0.00-2.17
<u>Non-malignant causes</u>				
Cerebrovascular Disease	5	1.94	2.58	0.84-6.03
All Heart Disease	17	15.69	1.08	0.63-1.73
Nonmalignant Respiratory Disease	1	2.57	0.39	0.01-2.16
Cirrhosis of Liver	0	1.18	0.00	0.00-3.14
Nephritis and Nephrosis	0	0.23	0.00	0.00-16.01
Accidents	5	4.79	1.04	0.34-2.44
Motor Vehicle Accidents	2	2.43	0.82	0.10-2.97
All Other Accidents	3	2.35	1.28	0.26-3.73
Violence	6	2.64	2.27	0.83-4.95
Suicides	5	2.14	2.33	0.76-5.45
Homicides	1	0.49	2.02	0.05-11.23

Table 5. Cause specific deaths and standardized mortality ratios for selected causes of death for Cottage Grove employees ever employed in jobs with probable PFOA exposure, but did not hold jobs with definite PFOA exposure.

Cause	Observed	Expected	SMR	95% CI
All Deaths	267	314.73	0.85	0.75-0.96
<u>Cancers</u>				
All Malignant Neoplasms	80	90.13	0.89	0.70-1.10
Buccal Cavity and Pharynx	1	1.85	0.54	0.01-3.00
Digestive Organs and Peritoneum	19	22.46	0.85	0.51-1.32
Esophagus	1	2.40	0.42	0.01-2.32
Stomach	1	2.78	0.36	0.01-2.00
Large Intestine	8	8.11	0.99	0.43-1.94
Rectum	2	1.74	1.15	0.14-4.15
Biliary Passages and Liver Primary	1	1.99	0.50	0.01-2.80
Pancreas	6	4.84	1.24	0.45-2.70
All Other Digestive	0	0.61	0.00	0.00-6.00
Respiratory System	26	27.45	0.95	0.62-1.39
Larynx	1	0.80	1.25	0.03-6.93
Bronchus, Trachea, Lung	25	26.36	0.95	0.61-1.40
Breast	2	3.58	0.56	0.07-2.02
Female Reproductive	2	2.22	0.90	0.11-3.26
Male Reproductive	6	6.15	0.98	0.36-2.12
Prostate	5	5.78	0.86	0.28-2.02
Testis and Other Male Genital Organs	1	0.36	2.75	0.07-15.30
Urinary Organs	3	4.38	0.68	0.14-2.00
Kidney	2	2.70	0.74	0.09-2.67
Bladder and Other Urinary Organs	1	1.68	0.59	0.02-3.30
Malignant Melanoma of Skin	2	1.41	1.42	0.17-5.11
All Lymphatic and Hematopoietic Tissue	8	10.34	0.77	0.33-1.52
<u>Non-malignant causes</u>				
Cerebrovascular Disease	10	15.70	0.64	0.30-1.17
All Heart Disease	81	104.04	0.78	0.62-0.97
Nonmalignant Respiratory Disease	12	20.17	0.60	0.31-1.04
Cirrhosis of Liver	6	6.35	0.95	0.35-2.06
Nephritis and Nephrosis	1	1.79	0.56	0.01-3.10
Accidents	16	19.87	0.81	0.46-1.31
Motor Vehicle Accidents	12	9.39	1.28	0.66-2.23
All Other Accidents	4	10.48	0.38	0.10-0.98
Violence	6	9.55	0.63	0.23-1.37
Suicides	6	7.77	0.77	0.28-1.68
Homicides	0	1.78	0.00	0.00-2.07

Table 6. Cause specific deaths and standardized mortality ratios for selected causes of death for Cottage Grove employees never exposed to PFOA or other fluorochemicals (non-chemical division).

Cause	Observed	Expected	SMR	95% CI
All Deaths	294	342.46	0.86	0.76-0.96
<u>Cancers</u>				
All Malignant Neoplasms	81	98.17	0.83	0.66-1.03
Buccal Cavity and Pharynx	1	1.96	0.51	0.01-2.83
Digestive Organs and Peritoneum	20	24.05	0.83	0.51-1.28
Esophagus	2	2.49	0.80	0.10-2.90
Stomach	3	3.04	0.99	0.20-2.88
Large Intestine	9	8.74	1.03	0.47-1.96
Rectum	0	1.88	0.00	0.00-1.96
Biliary Passages and Liver Primary	0	2.11	0.00	0.00-1.75
Pancreas	5	5.12	0.98	0.32-2.28
All Other Digestive	1	0.67	1.49	0.04-8.26
Respiratory System	25	29.18	0.86	0.55-1.27
Larynx	1	0.85	1.18	0.03-6.53
Bronchus, Trachea, Lung	23	28.01	0.82	0.52-1.23
Breast	4	4.70	0.85	0.23-2.18
Female Reproductive	1	2.81	0.36	0.01-1.98
Male Reproductive	2	7.19	0.28	0.03-1.00
Prostate	2	6.82	0.29	0.04-1.06
Testis and Other Male Genital Organs	0	0.36	0.00	0.00-10.12
Urinary Organs	5	4.73	1.06	0.34-2.47
Kidney	1	2.84	0.35	0.01-1.96
Bladder and Other Urinary Organs	4	1.89	2.11	0.58-5.40
Malignant Melanoma of Skin	2	1.48	1.35	0.16-4.89
All Lymphatic and Hematopoietic Tissue	10	11.10	0.90	0.43-1.66
<u>Non-malignant causes</u>				
Cerebrovascular Disease	11	18.21	0.60	0.30-1.08
All Heart Disease	103	114.39	0.90	0.73-1.09
Nonmalignant Respiratory Disease	17	23.11	0.74	0.43-1.18
Cirrhosis of Liver	6	6.74	0.89	0.32-1.94
Nephritis and Nephrosis	1	2.10	0.48	0.01-2.64
Accidents	17	20.71	0.82	0.48-1.31
Motor Vehicle Accidents	10	9.64	1.04	0.50-1.91
All Other Accidents	7	11.07	0.63	0.25-1.30
Violence	6	9.87	0.61	0.22-1.32
Suicides	2	8.04	0.25	0.03-0.90
Homicides	4	1.83	2.19	0.60-5.60

Table 7. Cause specific standardized mortality ratios for Cottage Grove employees with a minimum of one year of employment in a job with definite PFOA exposure.

Cause	Observed	Expected	SMR	95% CI
All Deaths	17	22.25	0.76	0.44-1.22
Cancers				
All Malignant Neoplasms	4	6.33	0.63	0.17-1.62
Digestive Organs and Peritoneum	1	1.59	0.63	0.02-3.48
Esophagus	0	0.20	0.00	0.00-18.87
Stomach	0	0.19	0.00	0.00-19.33
Large Intestine	1	0.56	1.79	0.05-9.94
Rectum	0	0.12	0.00	0.00-30.22
Biliary Passages and Liver Primary	0	0.14	0.00	0.00-26.38
Pancreas	0	0.35	0.00	0.00-10.67
All Other Digestive	0	0.04	0.00	0.00-89.75
Respiratory System	1	2.09	0.48	0.01-2.66
Bronchus, Trachea, Lung	1	2.01	0.50	0.01-2.77
All Other Respiratory	0	0.02	0.00	0.00-160.98
Breast	0	0.07	0.00	0.00-51.86
Prostate	1	0.38	2.63	0.07-14.62
Urinary Organs	0	0.32	0.00	0.00-11.39
Kidney	0	0.21	0.00	0.00-17.37
Bladder and Other Urinary Organs	0	0.11	0.00	0.00-33.12
Malignant Melanoma of Skin	0	0.12	0.00	0.00-29.99
Thyroid and Other Endocrine Glands	0	0.02	0.00	0.00-155.50
All Lymphatic and Hematopoietic Tissue	0	0.75	0.00	0.00-4.90
Non-malignant causes				
Cerebrovascular Disease	3	0.89	3.36	0.69-9.82
All Heart Disease	7	7.28	0.96	0.39-1.98
Other Nonmalignant Respiratory	0	0.70	0.00	0.00-5.30
Cirrhosis of Liver	0	0.52	0.00	0.00-7.11
Accidents	1	1.74	0.58	0.01-3.20
Motor Vehicle Accidents	0	0.84	0.00	0.00-4.41
All Other Accidents	1	0.90	1.11	0.03-6.19
Violence	2	0.93	2.15	0.26-7.75
Suicides	2	0.76	2.62	0.32-9.45
Homicides	0	0.17	0.00	0.00-22.07

Table 8. Cause specific standardized mortality ratios for Cottage Grove employees with a minimum of one year of employment in a job with definite or probable PFOA^a.

Cause	Observed	Expected	SMR	95% CI
All Deaths	219	274.36	0.80	0.70-0.91
Cancers				
All Malignant Neoplasms	68	77.33	0.88	0.68-1.11
Digestive Organs and Peritoneum	21	19.40	1.08	0.67-1.65
Esophagus	1	2.16	0.46	0.01-2.57
Stomach	1	2.42	0.41	0.01-2.29
Large Intestine	10	6.91	1.45	0.69-2.66
Rectum	2	1.51	1.32	0.16-4.78
Biliary Passages and Liver Primary	1	1.70	0.59	0.01-3.27
Pancreas	6	4.17	1.44	0.53-3.13
All Other Digestive	0	0.52	0.00	0.00-7.06
Respiratory System	23	24.20	0.95	0.60-1.43
Bronchus, Trachea, Lung	22	23.22	0.95	0.59-1.43
Breast	0	2.09	0.00	0.00-1.77
Prostate	6	5.19	1.16	0.42-2.52
Urinary Organs	2	3.88	0.52	0.06-1.86
Kidney	1	2.41	0.42	0.01-2.31
Bladder and Other Urinary Organs	1	1.47	0.68	0.02-3.79
Malignant Melanoma of Skin	2	1.30	1.54	0.19-5.55
Thyroid and Other Endocrine Glands	0	0.28	0.00	0.00-13.34
All Lymphatic and Hematopoietic Tissue	4	9.03	0.44	0.12-1.13
Nonmalignant causes				
Cerebrovascular Disease	11	13.03	0.84	0.42-1.51
All Heart Disease	68	90.90	0.75	0.58-0.95
Nonmalignant Respiratory Disease	6	17.09	0.35	0.13-0.76
Cirrhosis of Liver	4	5.69	0.70	0.19-1.80
Accidents	13	18.75	0.69	0.37-1.19
Motor Vehicle Accidents	10	8.95	1.12	0.53-2.06
All Other Accidents	3	9.80	0.31	0.06-0.90
Violence	7	9.35	0.75	0.30-1.54
Suicides	7	7.62	0.92	0.37-1.89
Homicides	0	1.74	0.00	0.00-2.13

a: Includes workers who accrued one year of exposure with definite and probable jobs combined.

Table 9. Observed and expected deaths from cerebrovascular disease with SMRs and 95% CI by years of employment in jobs with definite PFOA exposure.

Years of PFOA Exposure	OBS	EXP	SMR	95% CI
<1	2	1.05	1.91	0.22-6.91
1-5	0	0.46	0.00	0.0-8.02
5-10	3	0.19	15.03	3.02-43.91
> 10	0	0.23	0.0	0.0-15.17
Total	5	1.94	2.58	0.83-6.03

Table 10. Observed and expected deaths from cerebrovascular disease with SMRs and 95% CI by cumulative exposure.

<u>Weighted Exposure^a</u>	<u>OBS</u>	<u>EXP</u>	<u>SMR</u>	<u>95% CI</u>
>0-2499	8	9.67	0.83	0.36-1.63
2500-4999	2	2.80	0.71	0.08-2.58
5000-7499	1	2.32	0.43	0.01-2.40
7500-9999	0	1.76	0.00	0-2.08
10000-& Over	4	1.21	3.31	0.89-8.46
<u>Total</u>	<u>15</u>	<u>17.75</u>	<u>0.85</u>	<u>0.47-1.39</u>

a: Duration of employment (days)*exposure weighting factor.