Germicidal ultraviolet irradiation in air conditioning systems: effect on office worker health and wellbeing: a pilot study

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Abstract

Objectives—The indoor environment of modern office buildings represents a new ecosystem that has been created totally by humans. Bacteria and fungi may contaminate this indoor environment, including the ventilation systems themselves, which in turn may result in adverse health effects. The objectives of this study were to test whether installation and operation of germicidal ultraviolet (GUV) lights in central ventilation systems would be feasible, without adverse effects, undetected by building occupants, and effective in eliminating microbial contamination.

Methods—GUV lights were installed in the ventilation systems serving three floors of an office building, and were turned on and off during a total of four alternating 3 week blocks. Workers reported their environmental satisfaction, symptoms, as well as sickness absence, without knowledge of whether GUV lights were on or off. The indoor environment was measured in detail including airborne and surface bacteria and fungi.

Results-Airborne bacteria and fungi were not significantly different whether GUV lights were on or off, but were virtually eliminated from the surfaces of the ventilation system after 3 weeks of operation of GUV light. Of the other environmental variables measured, only total airborne particulates were significantly different under the two experimental conditions-higher with GUV lights on than off. Of 113 eligible workers, 104 (87%) participated; their environmental satisfaction ratings were not different whether GUV lights were on or off. Headache, difficulty concentrating, and eye irritation occurred less often with GUV lights on whereas skin rash or irritation was more common. Overall, the average number of work related symptoms reported was 1.1 with GUV lights off compared with 0.9 with GUV lights on.

Conclusion—Installation and operation of GUV lights in central heating, ventilation and air conditioning systems of office buildings is feasible, cannot be detected by workers, and does not seem to result in any adverse effects.

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Keywords: indoor environment; indoor air pollution; indoor air quality; environmental health; building related illness; germicidal ultraviolet light Most modern office buildings and virtually all high rise office towers have sealed exterior shells with mechanical ventilation systems. These ventilation systems are the main means by which the indoor environment is heated or cooled, humidified or dehumidified, and kept pollution free by exchange of indoor with outdoor air. Over the past 20 years, there have been numerous reports of health problems arising among workers in such buildings.12 These problems-mainly non-specific symptoms including headache, fatigue, difficulty concentrating, as well as mucosal irritative symptoms-are estimated to occur in 20%-30% of office workers.3 4 Although in some cases a specific causative agent or problem is identified, in most instances no cause can be found.12

There is evidence to implicate bacteria and fungi in the pathogenesis of non-specific building related illnesses. These microorganisms have been detected in high concentrations on cooling coils,⁵ filters,⁶ drip pans,⁵ humidification systems,⁷ and in the ductwork of the supply air.8 9 Microbial contamination of ventilation systems has been responsible for several specific building related illnesses through toxic, allergic, hypersensitive, or infectious mechanisms.10 In five large scale cross sectional surveys involving >20 000 workers in 103 buildings, air conditioning was consistently associated with excess prevalence of non-specific building related illnesses11-15 This has been attributed to microbial contamination.11

The germicidal efficacy of ultraviolet (UV) light has been known for many decades,^{16 17} and is used in hospitals, food processing plants, pharmaceutical manufacturing, and other areas where microbial decontamination is important. In the past, UV lights were not used in ventilation systems of office buildings because of technical limitations. However, newly developed high intensity lamps have overcome these limitations, making this approach feasible and safe because there would be no direct exposure to occupants of the building.

To study the efficacy of GUV lights installed in central heating, ventilation, and air conditioning (HVAC) systems, we first conducted a pilot study to assess safety, feasibility, duration of intervention periods, and possibility of detection of operation of GUV lights by building occupants.

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Methods

DESIGN

A cross over design was used. Ultraviolet lights were installed within the ventilation systems serving three floors of one downtown office building, and were turned on, or off, for a total of four alternating blocks of 3 weeks each. In the 3rd week of each block, workers completed self administered questionnaires and the indoor environment was measured in detail. Two complete sets of self administered questionnaires and environmental measurements were taken with the GUV lights off, and two sets with them on, allowing an analysis of effect within subjects.

INTERVENTION

Ultraviolet lights from three different manufacturers were installed in the ventilation systems serving separate floors of the study building (one manufacturer per floor). For manufacturer A, eight 37 W J-shaped bulbs which were 0.7 m in total length were installed. These bulbs produced UV light with an intensity of $131 \,\mu$ W/cm², at a distance of 1 m.

For manufacturer B, two bulbs 0.6 m long and two bulbs 0.46 m long were installed in each system for a total of 2.1 m of UV light generation. These bulbs produced 158 μ W/cm² at a distance of one metre.

For manufacturer C, 18 U shaped bulbs, each with total length of 0.4 m were installed in each system. Each bulb produced UV light with an intensity of 25 μ W/cm² at a distance of one metre.

All UV bulbs produced light at 254 nm wavelength and were installed in the central air supply ducting just downstream from, and shining directly on the cooling coils, walls, and floor of the air supply ducting, and drip pans below the cooling coils.

STUDY POPULATION

For this pilot study, we selected a building on the basis of the characteristics of the building and ventilation system, not because of known microbial contamination, or previous problems of increased symptoms or complaints about indoor air quality. The building was an office tower constructed 15 years earlier which had a sealed exterior shell. Separate ventilation systems serving each floor had air conditioning, bag filter systems rated at 80%–90% efficiency, and steam humidification.

Workers were potentially eligible if their worksite was on one of the three floors served by the ventilation systems in which the UV lights were installed. We excluded workers who were not present throughout the entire study period, did not have a fixed worksite—for example, messengers or couriers—or who worked within the study building for <2 days a week on average—for example, sales representatives. Eligible workers who agreed to participate signed an informed consent form. This study was approved by an ethics committee of the Montreal Chest Institute Research Centre.

QUESTIONNAIRES

Before beginning the study, a baseline questionnaire was completed for demographic characteristics, personal, medical, and smoking history, as well as work and worksite characteristics. On four occasions, a weekly questionnaire was completed. This questionnaire asked respondents to indicate their environmental ratings for several variables on indoor thermal and physical comfort, and air quality. Workers reported the occurrence of symptoms that day, whether they had started since arriving at work (defined as work related), and the impact of these symptoms on their ability to work. Finally, workers were asked to indicate whether they had missed hours or days due to sickness of any type and whether they attributed any such illnesses to problems of indoor air quality.

ENVIRONMENTAL MEASURES

In the same week that the questionnaires were completed, environmental measures were taken with the following instruments and methods:

Temperature, humidity, and air velocity were measured with a direct reading VELOCI-CALC hot wire anometer.

Carbon dioxide was measured at the same sites and at the same times with an ADC infrared direct reading monitor.

Carbon monoxide was measured at the same times and sites as CO_2 with a direct reading NEOTOX electrochemical sensor.

Total volatile organic compounds (TVOCs) were collected onto a charcoal tube with an SKC volumetric pump operating at 200 ml/min over 8 hours, desorbed with carbon disulphide, and measured by a flame ionisation detection (FID) method—NIOSH method 1510 modified.¹⁸

Nitrogen dioxide (NO₂) was collected with SKC volumetric air samplers operating at 75 ml/min over 8 hours onto a solid sorbent sampling tube containing triethanolamine impregnated molecular surface and analysed with the NIOSH method P and CAM 231.¹⁸

Ozone (O₃) was collected by bubbling air at 1 l/min over 8 hours through an ozone absorbency solution, and analysed with NIOSH method P and CAM 154.¹⁸

Formaldehyde was collected with SKC passive samplers over an 8 hour period, analysed with the ASTM method D 5014-89.¹⁹

Total airborne dust was collected with SKC volumetric samplers operating at 1.5 l/min for 8 hours. Total dust weight was estimated by comparing dry filter weights before and after collection.

Airborne fungal colony forming units were collected with Burkhard volumetric samplers operating for 15 minutes and impacting onto Petri dishes containing Sabhourad or maltose extract agar (MEA) media. Airborne bacteria were collected with the same sampling method impacting onto blood agar (for all bacteria) and Columbia CNA (for gram positive bacteria) media. Colony forming units were counted after 48–72 hours of growth at 37°C.^{20 21}

Surface swabs were taken from the interior walls of the main ventilation system where the

Table 1Data of study population (n (%))

Participation (n):	
Total eligible identified	119
Refused or incomplete	15
Participants (% of eligible)	104 (87)
Average participation each week (% of eligible)	97 (82)
Characteristics of participants (104):	
Age (mean (SD))	45 (9)
Sex $(n(\%))$:	
Male	43 (41)
Female	61 (59)
Smoking (currently) (n(%)):	
Yes	18 (17)
No	86 (83)
History of atopy (n(%)):	37 (38)
Job type $(n(\%))$:	
Clerical	24 (23)
Management or professional	80 (77)

UV lights were installed, cooling coils, and drip pans. The swab was dipped in a solution of 0.1% peptone with 0.01% Tween 80, and then pulled straight along the surface for 15 cm at an angle of 45°. The swab was then rubbed in a back and forth motion covering evenly one quadrant of a Petri dish containing Sabhourad or MEA media for fungi, or blood agar plates or Columbia CNA for bacteria. The swab was discarded and a sterile loop used to spread from the edge of this first quadrant into the second quadrant, then from the edge of the second quadrant into the third quadrant, and finally, from the edge of the third into the fourth quadrant. All plates were cultured for 48 to 72 hours at 37°C, after which colony forming units were estimated with a semiquantitative scale from 0 for no growth at all to 4 for confluent growth. Colony counts were made from the last quadrant where there was growth.

All variables were measured in outdoor air on the same days as indoor measurements were taken. Temperature, humidity, air velocity, carbon dioxide, and carbon monoxide were measured in return air and supply air of each HVAC system, as well as at the level of the ceiling diffusers and breathing zone of four randomly selected workers on each floor in the morning and afternoon of the day of completing the questionnaire (total about 300 measures of each variable). The TVOCs, NO₂, O₃, and formaldehyde were measured in supply air and at the level of ceiling diffusers and breathing zones of one worker on each floor in each week (total 40 measures of each variable). Airborne bacteria and fungi were sampled in return air, supply air, and at the level of ceiling

Table 2 Results of environmental measures under two experimental conditions

	GUV on		GUV off	
	Outdoor air	Sites+	Outdoor air	Sites†
Temperature (°C)	8.3	22.5	7.4	22.6
Relative humidity (%)	44.2	36.3	46.3	34.9
Air velocity (m.s ⁻¹)	-	0.02	-	0.02
Carbon dioxide (ppm)	395	624	353	684
Carbon monoxide (ppm)	< 2	< 2	< 2	< 2
TVOC (µg/m ³)	31	298	360	203
NO ₂ (ppb)	0.46	0.30	0.25	0.18
Ozone (ppb)	0.01	0.01	0	0
Formaldehyde (ppm)	0	0.03	0.03	0.03
Total airborne particulates (µg/m ³)	0	91**	12	31

** p <0.01 For total airborne particulates.

Difference in environmental variables.

-No significant difference for all variables except particulates.

GUV=germicidal ultraviolet lights.

diffusers and worksites of two workers per floor (total 100 samples for each variable) and six surface swabs were taken for bacteria and fungi on each floor (total of 72 samples for variable) over the course of the study.

Ultraviolet light intensity was measured with a hand held photometer with a sensor calibrated to measure ultraviolet C (UVC) at 254 nm wavelength (International Light model No IL1400A with model SEL240 sensor).

Results

As shown in table 1, 104 of 119 (87%) eligible workers participated—defined as completion of at least one weekly questionnaire with UV on, and at least one with UV off. Most participants were women who worked in management positions and in closed offices.

Results of environmental measures are summarised in table 2. Of particular interest, the operation of UV light did not result in increased concentrations of ozone, or TVOCs. Variation in outdoor temperature and humidity resulted in changes in indoor humidity and CO₂ concentrations-because of changes in outdoor air supply. All variables were within recommended limits with the exception of total airborne particulates which were above the United States Environmental Protection Agency limits of 75 μ g/m³ for 24 hour outdoor exposure on 36% of all occasions and 58% of occasions when measurements were taken with UV lights on. Airborne particulates were the only environmental contaminants that significantly increased in concentration in weeks when the GUV lights were on.

The results of fungal and bacterial cultures are shown in table 3. There was little apparent impact of operation of GUV lights on airborne microbial concentrations, although the concentrations in supply air were very low even before the UV lights were turned on, because of highly efficient bag filters used in these ventilation systems. An interesting and unexpected finding was that airborne concentrations of bacteria and fungi in the air supply was much higher when measured at the ceiling diffusers than in samples taken 10-15 feet from the GUV lights in the central air supply ducting. This phenomenon was found in all weeks and on all floors and presumably shows recontamination from microbial growth within the duct work itself. At the level of the worksite, the bacterial concentrations were somewhat higher than at diffuser level, indicating possible human source contamination. Operation of the UV lights resulted in virtual elimination of all bacterial and fungal growth on surfaces within the HVAC system. The airborne and surface fungal and bacterial concentrations were similar on all three floors-that is, there was no detectable difference in germicidal efficacy of GUV lights produced by the three manufacturers. The intensity of GUV light, measured at the cooling coils, exceeded 550 μ W/cm² for all three manufacturers.

Surface contamination was no longer detectable in the second experimental block even 3 weeks after the UV lights were turned off, suggesting that three weeks was not sufficient time

 Table 3
 Total viable (culturable) fungi and bacteria under two experimental conditions

	Bacteria	Bacteria		Fungi	
	GUV on	GUV off	GUV on	GUV off	
Airborne (cfu/m ³):					
Outdoor air	135	58	50	6	
Return air	57	34	6	3	
Supply air	13	14	1	0	
Diffuser	94	98	24	18	
Worksite	100	132	34	23	
HVAC surfaces (cfu/swabs):					
Drip pans	1.0	5	0	1.5	
Cooling coils	0.8	62	0.8	6.5	
Walls	0.3	5	0	1	

GUV=germicidal ultraviolet lights.

for recontamination to occur. This may have been because the study was conducted in October to December when outdoor microbial concentrations and humidity are low, resulting in less optimum conditions for microbial growth. After 2 months without GUV lights surface swabs showed that some recontamination had occurred by February 1998.

As shown in table 4, the weekly environmental satisfaction ratings were not different whether the UV lights were on or off. With GUV lights on, 82% of respondents thought that the ventilation system was functioning adequately to meet their needs, compared with 77% with GUV lights off. Based on all measures of workers' subjective appraisal of the indoor environment, it seemed that no worker was able to detect whether the UV lights were on or off.

Overall, 60% of workers reported some work related symptoms with the GUV lights off compared with 58% with GUV lights on, although the average number of work related symptoms reported was about 20% lower with GUV lights on than off (table 5). There was substantial fluctuation in sickness absence from week to week; in total, with GUV lights on, 5% of workers missed some work because of illness in the preceding week, compared with 5% with GUV lights off. When GUV lights were on, no work was missed due to illnesses that the workers attributed to indoor air quality problems, whereas when the GUV lights were off, a total of 15 hours were missed in the preceding week because of such illnesses.

Table 4 Weekly environmental satisfaction ratings *

	Overall	Overall	
	GUV on	GUV off	
Questionnaires (n): Thermal comfort:	196	203	
Temperature	3.1	3.2	
Humidity	3.0	3.0	
Air movement	3.2	3.1	
Physical environment:			
Lighting	3.4	3.3	
Surrounding noises	3.6	3.5	
Working space	3.6	3.5	
Indoor air quality:			
Dust	2.2	2.1	
Odours	2.8	2.8	
Overall air quality	2.5	2.4	

*All ratings scored 0 (far from ideal) to 4 (ideal).

GUV=germicidal ultraviolet lights.

Table 5Workers (%) reporting work related symptomsunder two experimental conditions

	Overall	
	GUV on	GUV off
Questionnaires completed (n)	196	203
Any symptoms	58	60
Symptoms of the CNS:	27	33
Headache	15	20
Difficulty concentrating	14	14
Trouble staying awake	10	13
Mucosal symptoms:	42	44
Irritated eyes	17	24
Dry or irritated throat	21	20
Dry or irritated nose	18	17
Stuffy, congested, or runny nose	14	17
Cough or difficulty breathing	7	7
General symptoms	30	27
Musculoskeletal	13	14
Skin rash or irritation	22	15
Average number of symptoms:		
Not work related	1.5	1.6
Work related	0.9	1.1
Work related and reducing work		
capacity (moderately or greatly)	0.2	0.2

GUV=germicidal ultraviolet lights.

Discussion

Installation and operation of GUV lights in the central HVAC system proved to be feasible. On the basis of symptoms and measurements of O_3 and TVOCs, GUV lights seemed to be safe, and on the basis of environmental satisfaction the workers seemed to remain blinded. Operation of the UV lights eliminated surface bacteria and fungi, although the initial levels were low, but had no effect on airborne concentrations which were influenced more strongly by the presence of highly efficient bag filters and by apparent recontamination during passage through the supply air duct work.

Inferences from this study are limited by the few workers involved, low initial concentrations of microbial contaminants, limited microbial characterisation, and use of GUV lights produced by three different manufacturers. The small size of the study was intentional. We had calculated that a sample size of 98 participants would be required to detect with 80% power (β =0.8), and 95% sensitivity (α =0.05) that the number of workers who thought that the environment was worse or unacceptable increased from 15% to 30%.22 In fact, more workers felt that the ventilation was acceptable with GUV lights on, than when they were off, and satisfaction ratings of indoor air quality and thermal comfort were not different. These findings suggest that there was no adverse effect on perceived indoor environmental quality and the workers remained blinded to the study intervention.

The study building was selected on the basis of HVAC characteristics of air conditioning and humidification which are risk factors for microbial contamination,^{5 7 23} as well as for nonspecific building related illnesses,^{11 12 14 15 24} and not because it was a sick building—a very poorly defined term.²⁵ Although it may be argued that the building should have been selected on the basis of HVAC microbial contamination, there have not been any systematic surveys of this phenomenon and therefore the concentrations of HVAC microbial contamination that can be

considered normal or safe as opposed to hazardous, have not been defined.

There is enormous diversity and complexity of fungal and bacterial organisms which have been found in this indoor environment. This complexity of microbial flora has contributed to the difficulty in population based studies (non-outbreak situations), of detecting consistent relations between specific bacteria or fungi and occupants' symptoms of non-specific building related illnesses. The financial and logistic constraints of the present pilot study meant that characterisation of microbial contamination was crude-being limited to counts of total viable bacteria and fungi. However, use of GUV lights as the study intervention has the advantage that GUV lights is non-specific in its germicidal action-killing a wide range of microbes, making identification less important.

We used lights produced by three different manufacturers for this pilot study. There were no significant differences in concentrations of TVOCs or ozone, nor in eradication of surface microbial contamination in the central HVAC systems between the lights produced by the three manufacturers. Given the few workers, we were unable to compare reporting of symptoms or environmental ratings between groups of workers exposed to air irradiated by the lights from the three different manufacturers.

The trend in reduction of symptoms and sickness absence in weeks when the GUV lights were operating is encouraging, but raises the question of biological plausibility. The rationale for proposing GUV lights in central HVAC systems is based on indirect evidence. (1) There is abundant evidence that exposure in the home¹⁰ or outdoor²⁶⁻²⁸ environments to the same microbes results in health effects. (2) Microbial contamination of HVAC systems has been documented, wherever there is water condensation.⁵ (3) Cross sectional surveys have consistently identified air conditioning as a risk factor for non-specific building related symptoms among office workers^{11 12 14 24}—this has been attributed to microbial contamination.¹¹ (4) The HVAC microbial contamination has been associated with symptoms in allergic people.^{29 30} (5) The HVAC microbial contamination has been implicated in many outbreaks of specific building related illness.^{1 23 31 32} (6) In two well documented outbreaks, there was a wide range of manifestations. A few people had abnormalities of gas exchange, lung function, and chest radiography, but most had nonspecific symptoms, and manv were asymptomatic.^{23 32} All of these studies provide evidence that microbial contamination could be responsible for non-specific symptoms. These would probably be labelled as sick building syndrome if there were no sentinel case with objective health effects which indicated that a specific building related illnesses existed. On the other hand, several cross sectional surveys in non-problem buildings or non-outbreak situations have failed to detect any relation between fungal concentrations and occupants' symptoms.¹

We hypothesise that microbial contamination of central HVAC systems may play an important part in the pathogenesis of nonspecific building related illnesses. Source control remains the most important method of prevention of such illnesses. However, if source control measures have failed or as an adjunct measure, germicidal ultraviolet lights are a relatively low cost intervention that could eradicate this contamination. Our study has shown that it is feasible to install and operate these GUV lights in existing ventilation systems without detection or adverse effects among workers. These findings plus a trend to reduction in symptoms support the need for a larger scale study of GUV lights involving more workers in more buildings.

For this type of study, buildings must be selected on the basis of risk factors for microbial contamination, which include air conditioning,^{5 7 23} humidification,⁵ low effifilters,³⁰ ciency poor cleaning and maintenance,³³ or evaporative type of humidification.^{14 24} As well, the study design must incorporate longer cross over periodsthat is, more time between periods of operation of GUV lights, because in this pilot study 2 months was required for recontamination to occur. Such longer intervals would have the potential disadvantage of greater losses to the study population, but may result in less reduction in symptoms over time-a problem noted in this and other short term cross over studies.3 34 If such a study showed a considerable beneficial effect of GUV light, then this technology, which is safe and relatively low cost, could be installed in existing buildings throughout North America. Also, such a result would provide further evidence for the causal role of microorganisms in the pathogenesis of non-specific illnesses related to buildings.

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