

Fungi, Bacteria, Nano-particulates, Mycotoxins and Human Health in Water-Damaged Indoor Environments

Jack Dwayne Thrasher*

Advisory Committee, Chemical Impact Project, Tides Foundation, MillValley, CA, United States

*Corresponding author: Jack Dwayne Thrasher, Advisory Committee, Chemical Impact Project, Tides Foundation, MillValley, CA, United States, Tel: 575-937-1150; Fax: 916-827-2520; E-mail: toxicologist1@msn.com

Received date: Feb 29, 2016; Accepted date: Mar 03, 2016; Published date: Mar 10, 2016

Copyright: © 2016 Thrasher JD. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Abstract

Nine types of biocontaminants in damp indoor environments from microbial growth are discussed: (1) indicator molds; (2) Gram negative and positive bacteria; (3) microbial particulates; (4) mycotoxins; (5) volatile organic compounds, both microbial (MVOCs) and non-microbial (VOCs); (6) proteins; (7) galactomannans; (8) 1-3-b-D-glucans (glucans) and (9) lipopolysaccharides LPS (endotoxins). When mold species exceed those outdoors contamination is deduced. However, there are no current recommendations by the EPA, OSHA, NIOSH, WHO and the Medical and Toxicology professions as to what constitutes a safe level of indoor molds and bacteria and their toxins in a water-damaged indoor environment. The thrust of his review is to discuss the role of fungi and their toxins on the health of occupants of damp indoor spaces.

Keywords: Bacteria; Fungi; Nano-particulates; Mycotoxin

Introduction

Damp or wet building materials occur from a variety construction defects, roof leaks, HVAC condensation, water intrusion from floods, hurricanes, leaking appliances and plumbing, poorly designed foundations, e.g. basement walls that allow water seepage from wet soils, slope of the building lot leading to water accumulation under concrete slabs. We have been involved in homes with cracked cement slabs, bent aluminium window framing, highly contaminated wall cavities, poorly installed roofing, improperly sealed fireplaces, to mention a few. All of these situations lead to both hidden as well visual signs of fungal and bacterial growth. For simplicity, "water intrusion" will be used as all encompassing term [1-18].

Signs of Microbial Growth

Signs of water intrusion include, but are not necessarily limited to: (1) water stains on ceilings, walls and around windows; (2) increased moisture content using a moisture meter with penetrating electrodes on dry wall (e.g. wall cavities), crawl space, attics and carpeting; (3) visible fungal growth on surface of dry wall, insulation, e.g. crawl space attic, clothing, shoes and other wearing apparel, bedding, under side of carpeting; (4) musty odor from microbial volatile organic compounds; (5) The E.P.A. cautions that approximately 50% of the fungal growth can be hidden, therefore hidden from view. The identification of airborne mold spores only reveals what is present at the time of testing, not 24/7. Airborne mold testing does not necessarily reveal hidden mold, e.g. wall cavities, attic, under carpeting, ventilation ducts.

Water activity

Water concentration of substrate upon which fungi and bacteria grow dictates the species of fungi that are present. Thus, the water

concentration is defined as water activity (a_w) as follows: Water activity is the **partial** vapour pressure of water in a substance divided by the standard state partial vapour pressure of water. The standard state is most often defined as the partial vapour pressure of pure water at the same temperature.

Using this particular definition, pure distilled water has a water activity of exactly one. Higher a_w substances tend to support more microorganisms. Bacteria usually require at least 95+, and fungi at $0.7 \geq 0.95$. The fungi that are present at different water concentrations are listed in Table 1.

Xerophilic ($a_w < 0.8$)

Species of *Aspergillus* and *Penicillium*, and *Eurotium amstelodami* and other fungi grow at this water activity. Several of these species produce mycotoxins that include *Ochratoxin A*, *Aflatoxins*, *Sterigmatocystin*, *Aflatoxins* and *Gliotoxin*. Some species of *Aspergillus* (*fumigatus*, *niger*, *terreus* and *flavus*) are potential human pathogens, particularly *A. fumigatus*.

Mesophilic ($a_w 0.8-0.9$)

Species of *Alternaria*, *Cladosporium*, *Phoma* and *Ulocladium* as well as *Epicoccum nigrum*. This group of fungi also produces mycotoxins, but they are not as toxic as those produced by the xerophilic and hydrophilic fungi.

Hydrophilic ($a_w > 0.9$)

The fungi at this water concentration include *Chaetomium globosum*, species of *Fusarium*, *Trichoderma*, *Stachybotrys chartarum*, *Memnoniella echinata* and *Rhizopus stolonifer*. Trichothecenes are produced by *Fusarium*, *Trichoderma*, and *Stachybotrys* and *Memnoniella*, while *Chaetomium* produces chaetoglobosins A and C. All of these mycotoxins are very toxic to humans and animals.

Fungi Growing on Building Materials and Their a_w			
Colonizer Group	a_w Range	Classification	Fungal Example
Primary Colonizers (storage fungi)	< 0.80	Xerophilic/ Xerotolerant	<i>Penicillium chrysogenum</i> and <i>Aspergillus versicolor</i> : the most common ones; <i>A. fumigatus</i> , <i>niger</i> , <i>sydowii</i> , <i>ustus</i> , <i>Eurotium spp.</i> , <i>P. brevicompactum</i> , <i>commune</i> , <i>corylophilum</i> , <i>palitans</i> , <i>variotta</i> , <i>Paecilomyces Wallemia sebi</i>
Secondary Colonizers	0.80-0.9	Mesophilic	<i>Alternaria spp.</i> , <i>Cladosporium spp.</i> , <i>Epicoccum nigrum</i> , <i>Phoma spp.</i> , and <i>Ulocladium spp.</i>
Tertiary Colonizers	> 0.9	Hydrophilic	<i>Chaetomium globosum</i> , <i>Fusarium</i> , <i>Memnoniella echinata</i> , <i>Rhizopus stolonifer</i> , <i>Stachybotrys chartarum</i> , <i>Trichoderma spp.</i> (<i>T. atroviride</i> , <i>T. citrinoviride</i> , <i>T. harzianum</i> , and <i>T. longibrachiatum</i>)

Table 1: This table lists the species of fungi and their water activity requirements for growth.

Bacteria

A variety of bacteria have been identified in water damaged indoor environments. They require a_w of ≥ 0.95 . They include several species Gram positive *Bacilli* and *Cocci* (*Streptococcus*, *Micrococcus*, and *Staphylococcus*) as well as Gram negative bacteria. Several of these bacteria are pathogens, while the Gram negative bacteria release endotoxins into the indoor environment. Other bacteria that have been identified in water-damaged indoor environment are the Actinobacteria: species of *Streptomyces*, *Mycobacterium* and *Nocardia*. The Actinobacteria produce exotoxins, e.g. Valinomycin, a mitochondrial poison. The nontuberculin Mycobacteria can cause Mycobacterium Avium Complex (MAC) in humans as reviewed by Griffin et al and published on the website of the American Thoracic Society

Particulates

Particulates shed from molds include spores, fragments of mycelia and nano-particulates. Of the mold particulates the greatest concern is the nano-particulates at or less than 0.3 microns shed from mold colonies. Field studies of water-damaged homes have shown concentrations of nano-particulates in indoor dust that are at least 1000 times or greater than the indoor air mold spore counts [12-18]. These particulates contain 1, 3-beta glucans, a variety of fungal proteins that include substrate enzymes as well as mycotoxins.

The translocation of nano-particulates with their attached toxins occurs by two mechanisms: the nervous system and the surfactants of the alveoli. The nano-particulates enter the surfactants of the lungs and are then transported across the alveolar cell membranes and enter the systemic circulation. In the alveoli they are taken up by alveolar macrophages and alveolar Type I cells. They cause the generation of reactive oxygen species and nitrogen species, release of proinflammatory cytokines and injury to nuclear DNA. Morphological changes include emphysema and granulomatous and fibrotic lesions [19-21]. The other mode of transportation is via the olfactory neurons. Nano-particulates attach to the nasal mucosa and are transported up the olfactory nerve through the cribiform plate and enter the hypothalamus/pituitary axis and spreading throughout the brain [22-25].

Mycotoxins

Mycotoxins produced by various fungi have been identified in damaged building materials, wall cavities, and dust samples from

HVAC ducts (return and supply air) and refrigerator compressor (Table 2).

The fungi present in the dust samples were identified by ERMI-36 and are listed in Table 3. *Stachybotrys chartarum* was identified in all dust samples along with other mycotoxin producing fungal species (work in progress).

Home I.D.	HVAC & Refrigerator	Aflatoxin s	Ochratoxin A	Trichothecenes
1	Supply Air (Mstr bath)	0	0	2.659
	Supply Air (Bdrm)	0	0	0.109
	Supply Air (Off/Den)	0	0	4.082
	Refrigerator Coil Dust	0	0	1.36
2	Return Air	0	0.04	1.15
	Supply air (Mstr Bdrm)	0	0.63	2.634
3	Return Air	0	0	0.491
	Supply Air (Mstr Bdrm)	0	0	0.305
	Refrigerator Compressor	0	0	2.652
4	Return Air	0	0	0.661
5	Return Air	0	0	7.417
6	Return Air	0	0	0.257
7	Return Air	0	0.5	7.753
8	HVAC Supply Air	0	0	26.511
9	Refrigerator Coil Dust	0	0.7	0.86

Table 2: HVAC ducts, Refrigerator compressor and HVAC Air Filter. This table summarizes the concentrations of mycotoxins in ppb detected in HVAC ducts, dust from the refrigerator compressor and one home with a dirty filter in the intake of the HVAC duct.

Mycotoxins and molds have been identified in urine samples from patients with ME/CFS and in autopsy materials from deceased individuals exposed to fungi in their water-damaged homes [25-34]. Three of the case studies will be briefly reviewed below.

Home I.D	Mold Species identified by PCR-DNA IERMI-36 Tests
1	<i>Aspergillus fumigatus</i> , versicolor, penicillioides, niger, restrictus; <i>Eurotium amstelodami</i> ; <i>Stachybotrys chartarum</i> ; <i>Chaetomium globosum</i> ; <i>Aureobasidium pullulans</i> , <i>Paecilomyces variotta</i> ; <i>Penicillium variable</i> ; <i>Trichoderma viride</i> ; <i>Wallemia sebi</i> ; <i>Scopulariopsis brevicaulis</i>
2	<i>Aspergillus fumigatus</i> , restrictus, niger; <i>Cladosporium sphaerospermum</i> ; <i>Aureobasidium pullulans</i> ; <i>Scopulariopsis chartarum</i>
3	<i>Aspergillus fumigatus</i> , restrictus, niger, <i>Cladosporium sphaerospermum</i> ; <i>Eurotium amstelodami</i> ; <i>Penicillium purpurogenum</i> ; <i>Stachybotrys chartarum</i> ; <i>Aureobasidium pullulans</i> ; <i>Wallemia sebi</i>
4	<i>Aspergillus fumigatus</i> , penicillioides; <i>Stachybotrys chartarum</i> ; <i>Cladosporium sphaerospermum</i> ; <i>Penicillium brevicompactum</i> ; <i>Wallemia sebi</i>
5	<i>Aspergillus fumigatus</i> , penicillioides; <i>Cladosporium sphaerospermum</i> ; <i>Stachybotrys chartarum</i> ; <i>Scopulariopsis chartarum</i> ; <i>Paecilomyces variotta</i> ; <i>Scopulariopsis chartarum</i> ; <i>Penicillium brevicompactum</i>
6	<i>Aspergillus flavus</i> , niger, sydowii, unguis, <i>Chaetomium globosum</i> ; <i>Cladosporium sphaerospermum</i> ; <i>Paecilomyces variotta</i> ; <i>Scopulariopsis chartarum</i> ; <i>Stachybotrys chartarum</i>
7	<i>Aspergillus fumigatus</i> , restrictus, penicillioides, flavus, niger; <i>Stachybotrys chartarum</i> ; <i>Aureobasidium pullulans</i> ; <i>Paecilomyces variotta</i> ; <i>Wallemia sebi</i>
8	<i>Aspergillus flavus</i> , fumigatus, niger, <i>Ochraceus penicillioides</i> , ustus, versicolor; <i>Eurotium amstelodami</i> ; <i>Penicillium chrysogenum</i> ; <i>Scopulariopsis chartarum</i> ; <i>Stachybotrys chartarum</i> ; <i>Wallemia sebi</i>
9	<i>Aspergillus fumigatus</i> , ochraceus, versicolor, niger, <i>Stachybotrys chartarum</i> ; <i>Eurotium amstelodami</i> ; <i>Aureobasidium pullulans</i> ; <i>Penicillium crustosum</i> , <i>Scopulariopsis chartarum</i> ; <i>Trichoderma viride</i> ; <i>Wallemia sebi</i>

Table 3: This table summarizes the indicator species of molds identified by ERMI-36 in the dust taken from various sources in Table 1. Only the species of molds in the highest concentration are listed. Note that although *Stachybotrys chartarum* was not detected in home I.D. 2, it is noted that it does not readily shed its spores unless its colonies are disturbed or desiccated.

Tables 4 and 5 are from reference 30. The family of five (parents and 3 children) was exposed to molds and mycotoxins. The father and a daughter developed fungal sinusitis requiring surgical removal. The mother was pregnant during the first and second trimesters of the third child. The family moved out of the home during her third trimester. The infant was born with a total body flare, developed Neurofibromatosis Type I, resulting in at least over 90 skin pigment

blotches over her body. There was no family history of NFT1. The pet dog developed over 90 skin lesions predominantly lipomas. Family symptoms included: persistent coughing, throat irritation, headaches, severe fatigue, decreased concentration and memory, nose bleeds, loss of libido (parents), shortness of breath with wheezing. Trichothecenes ochratoxin were detected in samples of dust and other samples from various areas of the home (Table 4).

Sample	Trichothecenes (ppb)	Aflatoxins (ppb)	Ochratoxin A (ppb)
Towel Master Bath	11.7	NP	4.9
Sandal Master Bdrm	0.47	NP	3.4
Wood Trust-Crawl Space	1.68	3.5	5.8
Gravel-Crawl Space	7.7	NP	7.7
Dirt-Crawl Space	2.1	NP	2.1
Plastic Sheet-Crawl Space	NP	NP	2.8

Table 4: This table summarizes the detection of trichothecenes, aflatoxins and ochratoxin A present in bulk samples taken from the master bath, master bedroom (sandal) and crawl space. The reported date are in ppb per mycotoxin, NP: Not Present, Limit of Detection: Trichothecenes (0.2 ppb); Aflatoxins (1.0 ppb); Ochratoxin A (2.0 ppb).

Mycotoxins were present in nasal secretions, urine, mothers breast milk, placenta and umbilical cord. The dog had trichothecenes in its urine and skin lesions. The urine of the newborn was negative for mycotoxins. However, the presence of mycotoxins in the placenta, breast milk and umbilical cord most likely are indicative of fetal exposure (Table 5).

Figure 1 and Table 6 are from reference 33. This case involved a 55 year old woman who was exposed to molds in a water-damaged office

building. She developed a mass in her sphenoid sinus that lead to severe headaches and fatigue.

Diagnostic imaging revealed a mass that six different pathologists diagnosed as a malignancy. Sphenoid surgery, radiation, chemotherapy and radiation did not remedy the situation. Slides of the biopsy were sent to Dr. Dumanov (one of the authors). He performed differential staining that revealed fungal growth.

PCR-DNA test identified *Aspergillus terreus*. She was then seen by Dr. Gray, Benson, Arizona, He treated intranasally with voriconazole and cyclosporine. MRI imaging demonstrated that the mass had resolved. During the antifungal treatment her urine was tested for mycotoxins.

Specimen	Trichothecenes	Aflatoxins	Ochratoxin A
Urine - Father	NP	NP	18.2
Nasal Secretion-Father ₁	NP	0.5	13
Urine - Mother	NP	NP	18.2
Nasal Secretion - Mother	1.02	1.2	1.6
Urine - Daughter	0.23	NP	28.0
Nasal Secretion- Daughter ₂	4.68	NP	3.8
Urine - Son	0.2	NP	18.9
Nasal Secretion - Son	ND	ND	ND
Breast Milk	0.18	0.9	2.7
Placenta	NP	NP	4.2
Umbilical Cord	NP	NP	NP
Urine - New Born	NP	NP	NP
Urine - Dog	1.49	NP	25.9
Ear Mass - Dog	23.07	0	2.2
Liooma - Dog	20.9	0	1.4

Table 5: Mycotoxins present in body fluids and tissues of the family and pet dog. The newborn's urine sample was negative with respect to mycotoxins. It is noted that the amniotic fluids were lost with the birth of the baby, Limits of Detection: Trichothecenes (0.2 ppb); Aflatoxins (1.0 ppb) Ochratoxin (2.0 ppb), ND – Not Done, NP – Not Present, 1: *Pseudomonas aureoginosa* and *Penicillium* were cultured from the nasal secretions. These data represent two different tests, 2: *Acinetobacter* sp and *Aspergillus fumigatus* were cultured left sphenoid sinus surgical specimens.

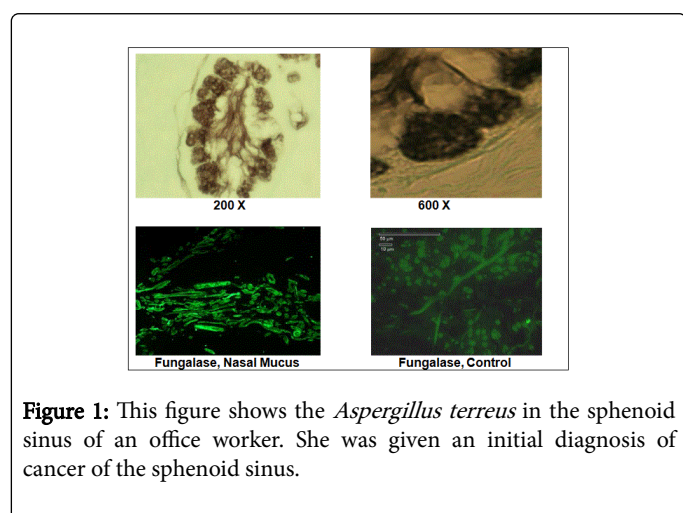


Figure 1: This figure shows the *Aspergillus terreus* in the sphenoid sinus of an office worker. She was given an initial diagnosis of cancer of the sphenoid sinus.

Tables 7-9 are from reference 34. This case study is a family of five (parents and three daughters. Prior to renting a home the health

history consisted of common colds and annual flu. Upon occupying a water-damaged home all became ill with a variety of symptoms.

Date	Aflatoxins (ppb) ¹	Trichothecenes (ppb)
3/27/2007	21	1.51
3/11/2007	5	0.53
6/3/2007	4	1.38
7/10/2007	12	3.44
7/14/2007	9	0.73
8/21/2007	5	21.3
12/12/2007	20	2.9

Table 6: This table summarizes the mycotoxins detected in the urine on several dates after removal of the fungal aspergilloma. The data show that detoxification probably results from the liberation of stored

mycotoxins, Limit of Detection Aflatoxins (1.0 ppb) Trichothecenes (0.2 ppb).

The most concerning to the family was the presence of persistent ME/CFS. ERMI-36 testing of dust samples from the refrigerator compressor/insulation dust identified species of *Aspergillus*, *Penicillium*, *Trichoderma viride*, *Chaetomium globosum* and *Stachybotrys chartarum* (Table 7).

Ochratoxin and trichothecenes were identified in the dust sample from the refrigerator and in urine specimens from all family members (Table 8). The diagnosis of ME / CFS was confirmed by sleep monitoring, demonstrating periods of waking, restless and un-refreshing sleep. In addition, a variety of Gram negative and positive bacteria were identified in a dust sample taken from the master bedspread at 8,400,000 CFU/g (data not shown). Thus, bacteria must also be considered as contributing factors in the ME / CFS.

Fungal Species	Number of Spores/mg of dust
<i>Aspergillus flavus</i>	8
<i>Aspergillus fumigatus</i>	33
<i>Aspergillus versicolor</i>	78
<i>Aspergillus ochraceus</i>	151
<i>Aspergillus niger</i>	33
<i>Aspergillus sydowii</i>	10
<i>Eurotium amstelodami</i>	785
<i>Penicillium purpurogenum</i>	7
<i>Aureobasidium pullulans</i>	114
<i>Penicillium corylophilum</i>	27
<i>Penicillium crustosum</i>	1,946
<i>Scopulariopsis chartarum</i>	44
<i>Trichoderma viride</i>	12
<i>Wallemia sebi</i>	25
<i>Chaetomium globosum</i>	3
<i>Stachybotrys chartarum</i>	1

Table 7: This table summarizes the results of the ERMI-36 test performed on the dust sample taken from the area of the refrigerator compressor and insulation. The species of each mold is given as the number of spores per milligram of dust.

Sample I.D.	Aflatoxin	Ochratoxin	Trichothecene
Refrigerator Dust	0	0.7	3.86
Urine (Father) 32 y	0	1.4	0.81
Urine (Mother) 31 y	0	2.1	1.26
Daughter 15 y	0	2.8	0.91
Daughter 12 y	0	1.4	0.97
Daughter 5 y	0	1.4	0.22

Table 8: This table summarizes the mycotoxins identified in the dust sample from the refrigerator compressor area and in the urine of the five occupants of the house. The concentrations are in ppb, Urine Ochratoxin References: < 1.8 ppb (negative), 1.8-2.0 (equivocal, 2.0 ppb (positive)). Trichothecene References: ≥ 0.2 ppb (positive).

Event	Father, 36 y	Mother, 34 y	Daughter, 17 y	Daughter, 15 y	Daughter, 7 y
Sleep Minutes	477 ± 62.5	508.2 ± 68.1	407 ± 67.7	431.6 ± 45.9	551.4 ± 62.2
Times Awake	3 ± 1.6	3.4 ± 1.3	2.2 ± 1.3	0.8 ± 1.3	1.5 ± 1.5
Restless	19.3 ± 3.2	19.6 ± 9.5	13 ± 6.5	9.8 ± 5.3	14.8 ± 6.2
Un-refreshing Sleep	42.8 ± 6.7	38 ± 20.8	25.2 ± 18.4	32.6 ± 12.2	26.6 ± 12.4

Table 9: This table summarizes the results of sleep monitoring events in each family member using FitBit Surge monitor.

Discussion

A short review of the literature

Three case studies have been briefly reviewed with respect to chronic illness resulting from water-damage and fungal biocontaminants. However, the peer-reviewed literature is replete with published papers regarding adverse health effects on occupants in, water-damaged indoor environments with demonstrable fungal and bacterial growth. These include, but are not necessarily limited to the following:

Upper and lower respiratory infections, bronchitis and lung disease

The health effects include infections, asthma, and hypersensitivity pneumonitis [7-10, 35-40].

Fungal sinusitis

Fungi as well as bacteria in damp indoor environments do cause sinusitis as well as adversely affecting the endocrine functions of the hypothalamus/pituitary axis. In addition, intracranial invasion occurs in immunocompetent patients [41-47].

Sarcoidosis

This is systemic inflammatory illness that can affect one or all organs. It is associated with nano-particulates shed by mold that contain 1, 3-beta glucans, mycotoxins and a variety of antigenic proteins [14,21,25,48-55].

Nervous system

Neurological damage includes the following: decrease in short and long memory in adults and children, autistic spectrum disorder in young children, peripheral neuropathy, loss of balance, facial pain, glossopharyngeal neuralgia, head and neck myalgias, movement disorders, and decreased visual acuity [56-64].

Kidney disease

Kidney disease has been associated with exposure to ochratoxin A produced by several species of *Aspergillus*. In European Asian counties it is referred to as endemic Balkan nephropathy associated with oral consumption of affected foods. In addition, a recent review of ochratoxin A has associated inhalational exposure to kidney disease, including focal segmental glomerulosclerosis [65,66].

Conclusion

Exposure to water-damaged indoor environments and subsequent fungal and bacterial growth leads to a variety of symptoms that are often overlooked by the medical profession. Most likely this results from the fact that a medical doctor with a busy practice has not kept up with the peer reviewed literature on this subject. Often, the office and/or field nurses are the first individuals to interview and interact with the patient. With respect to the nurse in the field, I highly recommend that awareness of water intrusion and microbial growth that are often present in homes and retirement facilities. Therefore, you are encouraged to look for signs of water-damaged and fungal growth. For example, I just had a conversation with a mother, age 32, who has a ten month old infant with chronic rasping cough, Also, her two older children have repeated upper and lower respiratory infections. The mother has ME / CFS as well as chronic sinus-nasal congestion. The rented home has had water intrusion via a faulty roof and plumbing leaks.

References

1. Campbell AW, Thrasher JD, Gray MR, Vojdani A (2004) Mold and mycotoxins: effects on the neurological and immune systems in humans. *Adv Appl Microbiol* 55: 375-406.
2. Indoor air Pollution (2009) Dampness and Mould. World Health Organization.
3. NIOSH Mold Alert (2013) Preventing Occupational Respiratory Disease from Exposures Caused by Dampness in Office Buildings, Schools, and other Nonindustrial Buildings.
4. Thrasher JD, Crawley S (2009) The biocontaminants and complexity of damp indoor spaces: More than meets the eyes. *Toxicology Industrial Health* 25: 583-615.
5. Damp Indoor Spaces and Health (2004) Institute of Medicine of the National Academies. The National Academic Press.
6. Pestka JJ, Yike I, Dearborn DG, Ward MD (2008) Harkema *Stachybotrys chartarum*, trichothecene mycotoxins, and damp building-related illness: new insights into a public health enigma. *Toxicol Sci* 104: 4-26.
7. Fisk WJ, Eliseeva EA, Mendell MJ (2010) Association of residential dampness and mold with respiratory tract infections and bronchitis: a meta-analysis. *Environ Health* 9: 72.
8. Fisk WJ, Lei-Gomez Q, Mendell MJ. (2007) Meta-Analyses of the associations of respiratory health effects with dampness and mold in homes. *Indoor Air* 17: 284-296.
9. Park JH, Cox-Ganser JM (2011) Mold exposure and respiratory health in damp indoor environments. *Front Biosci (Elite Ed)* 3: 757-771.
10. Park JH, Cox-Ganser JM (2011) Mold exposure and respiratory health in damp indoor environments. *Frontiers Biosci E3*: 757-7571.
11. Gosh BH, Lal JH, Srivastava A (2015) Review of bioaerosols in indoor environment with special reference to sampling, analysis and control mechanisms. *Environmental International* 85: 254-272.

12. Górný RL, Reponen T, Willeke K, Schmechel D, Robine E, et al. (2002) Fungal fragments as indoor air biocontaminants. *Appl Environ Microbiol* 68: 3522-3531.
13. Cho S-H, Seo S-C, Schmechel D, Grinshpun SA, Reponen T (2005) Aerodynamic characteristics and respiratory deposition of fungal fragments. *Atmos Environ* 39: 5454-5465.
14. Reponen T, Seo S-C, Grimsley F, Lee T, Crawford C, et al. (2007) Fungal fragments in moldy houses: A field study in homes in New Orleans and Southern Ohio. *Atmos Environ* 41: 8140-8149.
15. Górný RL, Reponen T, Willeke K, Schmechel D, Robine E, et al. (2002) Fungal fragments as indoor air biocontaminants. *Appl Environ Microbiol* 68: 3522-3531.
16. Adhikari A, Jung J, Reponen T, Lewis JS, DeGrasse EC, et al. (2009) Aerosolization of fungi, (1->3)-beta-D glucan, and endotoxin from flood-affected materials collected in New Orleans homes. *Environ Res* 109: 215-224.
17. Thrasher JD, Crawley S (2009) The biocontaminants and complexity of damp indoor spaces: more than what meets the eyes. *Toxicol Ind Health* 25: 583-615.
18. Straus DC (2009) Molds, mycotoxins, and sick building syndrome. *Toxicol Ind Health* 25: 617-635.
19. Mühlfeld C, Rothen-Rutishauser B, Blank F, Vanhecke D, Ochs M, et al. (2008) Interactions of nanoparticles with pulmonary structures and cellular responses. *Am J Physiol Lung Cell Mol Physiol* 294: L817-829.
20. Peters A, Veronesi B, Calderon-Garciduenas L, Gerhr P, Chen LC, Geiser M, et al. (2006) Translocation and potential neurological effects of fine and ultrafine particles a critical update. *Particle Fibre Toxicol* 3: 13.
21. Rylander R (1997) Investigations of the relationship between disease and airborne (1->3)-beta-D-glucan in buildings. *Mediators Inflamm* 6: 275-277.
22. Genc S, Zadeoglulari Z, Fuss SH, Genc K (2012) The adverse effects of air pollution on the nervous system. *J Toxicol* 2012: 782462.
23. Block ML, Calderón-Garcidueñas L (2009) Air pollution: mechanisms of neuroinflammation and CNS disease. *Trends Neurosci* 32: 506-516.
24. Calderon-Garciduenas L, Franco-Lira M, Torres-Jardon R, Henriquez-Roland c, Barragán-Mejía G, et al (2007) Pediatric respiratory and systemic effects of chronic air pollution exposure: Nose, lung heart, and brain pathology. *Toxicol Pathol* 35: 154-162.
25. Calderon-Garciduenas L, Solt AC, Henriquez-Roldan C, Torres-Jordan R, Nuse V, et al. (2008) Long-term air pollution is associated with neuroinflammation, an altered innate immune response, disruption of the blood-brain barrier, ultrafine particulate deposition, an accumulation of β -42 and α -synuclein in children and young adults. *Toxicol Pathol* 36: 289-310.
26. Gray MR, Thrasher JD, Crago R, Madison RA, Campbell AW, et al. (2003) Mixed mold mycotoxicosis: Immunological changes in humans following exposure in water-damaged buildings. *Arch Environ Health* 59: 410-420.
27. Hooper DG, Bolton VE, Guilford FT, Straus DC (2009) Mycotoxin detection in human samples from patients exposed to environmental molds. *Int J Mol Sci* 10: 1465-1475.
28. Brewer JH, Thrasher JD, Straus DC, Madison RA, Hooper D (2013) Detection of mycotoxins in patients with chronic fatigue syndrome. *Toxins (Basel)* 5: 605-617.
29. Gray MR, Hooper D, Thrasher JD (2014) Molds and mycotoxins in autopsy specimens in a death related to fungal pneumonia and pancytopenia, marijuana usage and a water-damaged home: A case report. *Internat J Clin Toxicol* 2: 11-20
30. Thrasher JD, Gray MR, Kilburn KH, Dennis DP, Yu A (2012) A water-damaged home and health of occupants: a case study. *J Environ Public Health* 2012: 312836.
31. Thrasher JD, Hooper DH, Taber J (2014) Family of six, their health and the death of a 16 month old male from pulmonary hemorrhage: Identification of mycotoxins and mold in the home, lungs, liver and brain of the deceased infant. *Inter J Clin Toxicol* 2: 1-10.
32. Gray MR, Thrasher JD, Hooper D, Crago R (2014) A case of Reye's-like syndrome in a 68-day old infant: Water damage home, mold, bacteria and aflatoxins. *Inter J Clin Toxicol* 2: 42-54
33. Gray MR, Thrasher JD, Dennis D, Dumanov M, Cravens H, et al. (2015) Sphenoid Aspergilloma: Diagnosed as a malignancy: A case report. *Otolaryngology* 2015: 3
34. Thrasher JD, Prokop C, Roberts C, Hooper D (2015) Family with ME/CFS following exposure to molds, mycotoxins and bacteria in a water-damaged home: A case report. *Inter J Clin Toxicol* 3: 18-28.
35. Kurup VP, Zacharisen MC, Fink JN (2006) Hypersensitivity pneumonitis. *Indian J Chest Dis Allied Sci* 48: 115-128.
36. Vesper SJ, Wymer L, Kennedy S, Grimsley LF (2013) Decreased pulmonary function measured in children exposed to high environmental relative moldiness index homes. *The Open Respir Med J* 7: 83-86.
37. Cox-Ganser JM, White SK, Jones R, Hilsbos K, Storey E, et al. (2005) Respiratory morbidity in office workers in a water-damaged building. *Environ Health Perspect* 113: 485-490.
38. Blanc PD, Quinlan PJ, Katz PP, Balmes JR, Trupin L (2013) Higher environmental relative moldiness index values measured in homes of adults with asthma, rhinitis, or both conditions. *Environ Health* 122: 98-101.
39. Tercelj M, Salobir B, Narancsik Z, Kriznar K, Brzetic-Romcevic T, et al. (2012) Nocturnal asthma and domestic exposure to fungi. *Indoor Built Environ* 22: 876-880.
40. White SK, Cox-Ganser JM, Benaise LG, Kreiss K (2013) Work-related peak flow and asthma symptoms in a damp building. *Occup Med (Lond)* 63: 287-290.
41. Dennis DP (2003) Chronic sinusitis: defective T-cells responding to superantigens, treated by reduction of fungi in the nose and air. *Arch Environ Health* 58: 433-441.
42. Dennis D, Robertson D, Curtis L, Black J (2009) Fungal exposure endocrinopathy in sinusitis with growth hormone deficiency: Dennis-Robertson syndrome. *Toxicol Ind Health* 25: 669-680.
43. Gorovoy IR, Kazanjian M, Kersten RC, Kim HJ, Vagefi MR (2012) Fungal rhinosinusitis and imaging modalities. *Saudi J Ophthalmol* 26: 419-426.
44. Mossa-Basha M, Ilica AT, Maluf F, Karakoç Ö, Izbudak I, et al. (2013) The many faces of fungal disease of the paranasal sinuses: CT and MRI findings. *Diagn Interv Radiol* 19: 195-200.
45. Siddiqui AA, Shah AA, Bashir SH (2004) Craniocerebral aspergillosis of sinonasal origin in immunocompetent patients: clinical spectrum and outcome in 25 cases. *Neurosurgery* 55: 602-611.
46. Srinivasan US (2008) Intracranial aspergilloma in immunocompetent patients successfully treated with, radical surgical intervention and antifungal therapy – Case series. *Ann Acad Med Singapore* 37: 783-787.
47. Gupta AK, Gupta AK (2009) Postgraduate institute management protocol for invasive *Aspergillus flavus*: *Internat J Infectious Dis* 13: 134-139.
48. TerÄelj M, Salobir B, Harlander M, Rylander R (2011) Fungal exposure in homes of patients with sarcoidosis - an environmental exposure study. *Environ Health* 10: 8.
49. TerÄelj M, Salobir B, Zupancic M, Wraber B, Rylander R (2014) Inflammatory markers and pulmonary granuloma infiltration in sarcoidosis. *Respirology* 19: 225-230.
50. TerÄelj M, StopinÄjek S, Ihan A, Salobir B, SimÄiÄ S, et al. (2011) In vitro and in vivo reactivity to fungal cell wall agents in sarcoidosis. *Clin Exp Immunol* 166: 87-93.
51. Tercelj M, Salobir B, Zupancic M, Rylander R (2011) Antifungal medication is efficient in the treatment of sarcoidosis. *Ther Adv Respir Dis* 5: 157-162.
52. Gerke AL (2014) Morbidity and mortality in sarcoidosis. *Curr Opin Pulm Med* 20: 472-478.
53. Rao DA, Dellaripa PF (2013) Extrapulmonary manifestations of sarcoidosis. *Rheum Dis Clin North Am* 39: 277-297.

54. Engelhard SB, Patel V1, Reddy AK1 (2015) Intermediate uveitis, posterior uveitis, and panuveitis in the Mid-Atlantic USA. *Clin Ophthalmol* 9: 1549-1555.
55. Wilson NJ, King CM (1998) Cutaneous sarcoidosis. *Postgrad Med J* 74: 649-652.
56. Anyanwu EC, Campbell AW, Vojdani A (2003) Neurophysiological effects of chronic indoor environmental toxic mold exposure on children. *ScientificWorldJournal* 3: 281-290.
57. Kilburn KH, Thrasher JD, Immers NB (2009) Do terbutaline- and mold-associated impairments of the brain and lung relate to autism? *Toxicol Ind Health* 25: 703-710.
58. Kilburn KH (2009) Neurobehavioral and pulmonary impairment in 105 adults with indoor exposure to molds compared to 100 exposed to chemicals. *Toxicol Ind Health* 25: 681-692.
59. Empting LD (2009) Neurologic and neuropsychiatric syndrome features of mold and mycotoxin exposure. *Toxicol Ind Health* 25: 577-581.
60. Campbell AW, Thrasher JD, Gray MR, Vojdani A (2004) Mold and mycotoxins: effects on the neurological and immune systems in humans. *Adv Appl Microbiol* 55: 375-406.
61. Carey SA, Plopper DG, Hyde SM, Islam Z, Pestka JJ, et al. (2012) Satratoxin-G from the black mold *Stachybotrys chartarum* induces rhinitis and apoptosis of olfactory neurons in the nasal airways of Rhesus monkeys. *Toxicol Pathol* 40: 887-898.
62. Jedrychowski W, Maugeri U, Stigter L, Jankowski J, Butscher M, et al. (2011) Cognitive function of 6-year old children exposed to mold-contaminated homes in early postnatal period, prospective birth control study in Poland. *Physiol Behav* 104: 989-995.
63. Campbell AW, Thrasher JD, Madison RA, Vojdani A, Gray MR, et al. (2003) Neural autoantibodies and neurophysiologic abnormalities in patients exposed to molds in water-damaged buildings. *Arch Environ Health* 58: 464-474.
64. Doi K, Uetsuka K (2011) Mechanisms of mycotoxin-induced neurotoxicity through oxidative stress-associated pathways. *Int J Mol Sci* 12: 5213-5237.
65. Grollman AF, Jelkovic B (2007) Role of environmental toxins in endemic (Balkan) nephropathy. *J Am Soc Nephrol* 18: 2817-2883.
66. Hope J, Hope BE (2012) A review of the diagnosis and treatment of ochratoxin A inhalational exposure associated with human illness and kidney disease including focal segmental glomerulosclerosis. *J Environ Pub Health*.