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1 Clothing-mediated exposures to chemicals and particles

Revised for Environmental Science & Technology

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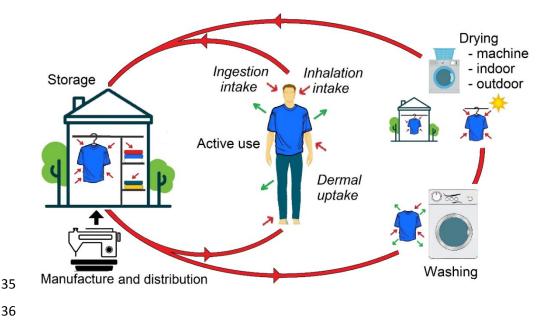
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17 Abstract

A growing body of evidence identifies clothing as an important mediator of human exposure to 18 chemicals and particles, which may have public health significance. This paper reviews and 19 20 critically assesses the state of knowledge regarding how clothing, during wear, influences exposure to molecular chemicals, abiotic particles, and biotic particles, including microbes and 21 allergens. The underlying processes that govern the acquisition, retention and transmission of 22 23 clothing-associated contaminants and the consequences of these for subsequent exposures are explored. Chemicals of concern have been identified in clothing, including byproducts of their 24 manufacture and chemicals that adhere to clothing during use and care. Analogously, clothing 25 acts as a reservoir for biotic and abiotic particles acquired from occupational and environmental 26 27 sources. Evidence suggests that while clothing can be protective by acting as a physical or chemical barrier, clothing-mediated exposures can be substantial in certain circumstances and 28 may have adverse health consequences. This complex process is influenced by the type and 29 30 history of the clothing, the nature of the contaminant and by wear, care and storage practices. 31 Future research efforts are warranted to better quantify, predict and control clothing-related 32 exposures.

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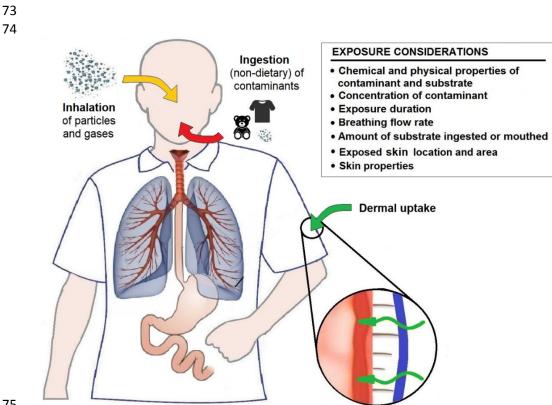


37 **1.** Introduction

38 Diverse chemicals, particles and microbes are found on clothing. Some are present at the time clothing is purchased, and some are acquired during the care, storage, and use of garments. 39 People spend most of their lives in intimate contact with clothing. They are exposed to the 40 species found on and in their clothing via inhalation, ingestion and dermal absorption (Figure 1). 41 More specifically, humans inhale species that desorb or are released from their clothing, ingest 42 clothing-associated chemicals and particles when clothing materials enter their mouths, and 43 acquire species on their skin from the clothing they wear. Once in the lungs, in the 44 gastrointestinal system or on the skin, chemicals from clothing may be absorbed into the body.¹ 45 46 As we show in this review, the resulting exposures are influenced by factors inherent to clothing, such as fiber type, weave, morphology, dveing process, color and chemical treatment (including 47 incorporation of flame retardants, stain repellants, and anti-wrinkle agents). Exposures are also 48 influenced by external factors such as washing, drying, storage, and usage patterns. Clothing-49

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50	mediated exposures can contribute to irritation, allergic reactions, and infections as well as risks
51	for adverse health effects as diverse as cancer, birth defects and heavy-metal poisoning. ²⁻⁴
52	Studies related to clothing-mediated exposures have been conducted by a diverse set of
53	researchers in the textile industry, government laboratories and academia. ^{2,3,5,6} While many
54	results have been summarized in reviews, government reports and books, the findings have yet to
55	be summarized within a framework that focuses on the ways in which clothing mediates
56	exposures to chemicals and particles. This review aims to provide a critical summary from such a
57	perspective. We present the review in two main sections, considering clothing-mediated
58	exposures first to chemicals and second to biotic and abiotic particles. Within these sections, we
59	summarize evidence for the influence of clothing on exposure to chemicals (§2.1) and particles
60	(§3.1). We review the occurrence and persistence of chemicals in clothing (§2.2), outline a
61	framework for quantifying clothing-mediated particle exposures (§3.2), discuss mechanisms of
62	accumulation and transfer of chemicals (§2.3), and review factors that influence clothing-
63	associated exposures to particles (§3.3). We discuss situations where the underlying factors
64	influencing chemical and particle exposures are similar, while also recording fundamental ways
65	that they differ. Whereas the potential influence on health risks is the key ultimate reason to
66	better understand clothing-mediated exposures, a detailed examination to quantify clothing-
67	associated health risks is beyond the scope of this review. We conclude (§4) with an examination
68	of knowledge gaps that currently limit the ability to predict or mitigate clothing-related
69	exposures to chemicals and particles. We suggest some research directions that could reduce
70	these limitations. Overall, we find that the influence of clothing on environmental exposures is
71	often substantial and so additional research efforts are warranted to better understand how
72	clothing influences human exposures, and ultimately human health and well-being.



75 76

Figure 1. Non-dietary routes of human exposure for contaminants of concern: Inhalation,ingestion and dermal absorption.

79 **2.** Chemical exposures

80 **2.1 Evidence of clothing-associated exposure to chemicals**

81 2.1.1 Clothing-associated chemicals in skin, blood and urine

82 Human exposure and uptake of organic compounds by means of transfer from treated fabrics has

- been investigated for several decades. For example, in the late 1970s, Blum et al.⁷ reported
- 84 finding metabolites of the flame retardant tris(2,3-dibromopropyl)phosphate (tris) in the urine of
- children who had worn clothing treated with this chemical. Radiolabeled tris in treated and dried
- cloth was shown to penetrate clipped skin of rabbits. Moistening the cloth with simulated sweat
- did not increase absorption.⁸ Earlier, Armstrong et al.⁹ and Brown¹⁰ reported instances of infant
- poisoning attributable to use of phenolic disinfectants in improperly laundered hospital fabrics.

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Recently. forestry workers wearing permethrin-treated, tick-proof pants were shown to have 89 significantly elevated levels of a permethrin metabolite in their urine.¹¹ Moreover, absorption of 90 ethylene oxide (a fumigant), glyphosate (an herbicide), malathion (an insecticide) and 91 benzothiazole (used as dye, biocide, herbicide, and fungicide) from fabric into skin or a skin-92 mimicking membrane has been demonstrated in studies using an *in vitro* diffusion cell.^{12–14} 93 Measurements of polychlorinated dibenzo-p-dioxins and dibenzofurans (PCDD/F) in the stratum 94 corneum, epidermis and subcutis of eight volunteers as well as in a variety of new fabric 95 swatches showed that some textiles are contaminated and can be an important source of exposure 96 to these chemicals.¹⁵ The PCDD/F species were shown to diffuse through the stratum corneum 97 into the deeper layers of the skin. Stratum corneum concentrations were substantially higher after 98 99 wearing contaminated shirts rather than uncontaminated shirts. Skin contamination was heterogeneous, both among individuals and among sites on the same individual. However, when 100 identical, homogeneously contaminated T-shirts were used in a companion study, relatively little 101 spatial and interpersonal variability was observed.¹⁶ Uptake from polyester was found to be an 102 103 order-of-magnitude lower than from cotton. Wearing fabrics that were previously worn enhanced 104 transfer. Residual sweat and lipid compounds may have served as transfer vehicles, or possibly 105 weakened the binding interaction between the fabric and PCDD/F. Heavy perspiration during 106 intense physical activity also increased the migration rate of a textile dye, Dianix[®], onto the skin of volunteer subjects, while contact time was found to be less important.¹⁷ 107 Clothing can act as a means of transporting pollutants from one environment to another. This 108 phenomenon has been studied in the context of health concerns related to "para-occupational" 109 110 exposures. Certain hazardous chemicals, such as lead, beryllium, polychlorinated biphenyls

111 (PCB) and pesticides, can be transferred from a work site to the worker's home via clothing and

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thereby contribute to elevated levels in the blood and urine, or even to direct adverse health 112 effects.^{18–20} For example, women who laundered agricultural work clothes had up to 42% higher 113 serum levels of dichlorodiphenyltrichloroethane (DDT) and hexachlorobenzene compared to 114 women who did not.^{21,22} Similarly, women living in homes in which agricultural workers wore 115 their work clothes had higher levels of most of the organochlorine pesticides that were being 116 used.²² Multivariate analyses by Park et al.²³ indicated an association between serum levels of 117 polybrominated diphenyl ethers in California firefighters and the storage and cleaning practices 118 used for protective gear. The authors suggested that these flame retardants can be transported to 119 120 fire stations via fireborne dust on soiled turnout gear and that good housekeeping practices can reduce subsequent exposure. (See also \$3.1.4.) 121

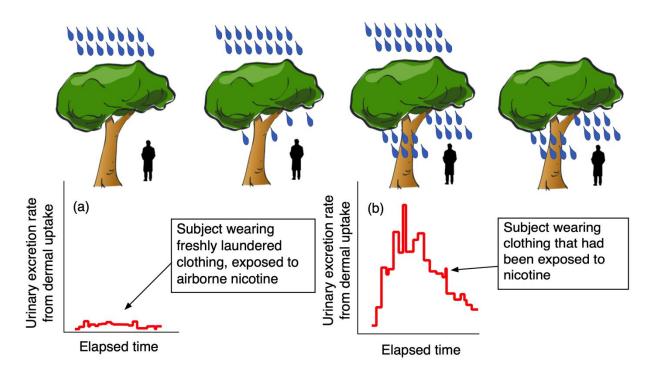
122 2.1.2 Influence of clothing on dermal uptake of airborne chemicals

Until recently, the influence of clothing on dermal uptake of airborne organic compounds 123 received relatively little attention. Initial studies examined a few chemicals, primarily volatile 124 organic compounds in occupational settings. Piotrowski²⁴ found that clothing reduced dermal 125 uptake of airborne nitrobenzene by about 20-30%, but that clothing had no observable effect on 126 phenol absorption.²⁵ Recent efforts have addressed dermal exposures to semivolatile organic 127 compounds common in everyday indoor settings. For example, Morrison et al.²⁶ measured the 128 uptake of two airborne phthalates, diethyl phthalate (DEP) and di-n-butyl phthalate (DnBP), by 129 an individual wearing either clean clothes or clothes previously air-exposed in a chamber with 130 elevated phthalate concentrations. When compared with dermal uptake for bare-skinned 131 individuals under otherwise identical experimental conditions,²⁷ clean clothes decreased 132 transdermal uptake by factors of 3–6, whereas previously exposed clothes increased dermal 133 uptake by factors of 3 and 6 for DEP and DnBP, respectively. Analogous results were obtained 134

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135	for nicotine. ^{28,29} This role of clothing as either "protector" or "amplifier" of dermal uptake is
136	illustrated in Figure 2. In another study, three subjects exhibited elevated urinary excretion rates
137	of the UV filter benzophenone-3 (BP-3) and its metabolite benzophenone-1 shortly after donning
138	T-shirts previously exposed to air with elevated BP-3 levels. ³⁰ The authors suggested that dermal
139	uptake of BP-3 from clothing could meaningfully contribute to overall body burdens.
140	The protective effect of uncontaminated clothing has also been indicated by reduced phthalate
141	and halogenated flame retardant concentrations in skin wipe samples taken from body parts
142	covered with clothing compared to uncovered skin. ^{31,32} However, clothing did not provide total
143	protection in these studies. In vitro experiments demonstrated reduced absorption of
144	organophosphates through a cotton shirt as compared to unclothed skin. ³³ However, common
145	clothing is reported to have little effect on dermal exposure to certain gases in hazardous material
146	incidents, such as methyl bromide, sulfuryl fluoride, chloropicrin and ethylene oxide.34,35

147



148

Figure 2. Dynamic urinary excretion rates due to dermal uptake for nicotine and two urinary 149 metabolites (summed) after exposing participants, who were wearing breathing hoods, to 150 airborne nicotine.²⁹ (a) Freshly laundered clothing, unexposed to nicotine, is protective; this is 151 analogous to a tree at the beginning of a rainstorm that protects a person from getting wet. (b) 152 Clothing that has been previously exposed to airborne nicotine dramatically increases urinary 153 excretion rate for days after wearing the clothing, just as a standing under tree after a long 154 rainstorm is most certain to get the person wet. This exposure occurs while wearing the clothes 155 in the environment containing the contaminant (third tree from left) and can continue after 156 157 leaving this environment.

- 158
- 159 2.1.3 Health effects as evidence of exposure

160 Studies of health effects related to hazardous substances in textiles further suggest clothing-

- 161 associated exposures. These works have mainly focused on dermatitis caused by textile dyes and
- 162 finishing resins.^{36–41} A limited literature also exists on carcinogenic, mutagenic and reprotoxic
- substances in textile articles. These effects have been suggested for certain dyes, especially azo
- 164 dyes^{42–45} and for some antibacterial agents, such as triclosan.^{46,47} Brominated flame retardants,
- 165 phthalates and degradation products of highly fluorinated polymeric water repellents and stain

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repellents, which can be present in textile articles, have been associated with reproductive and
developmental toxicity.⁴⁸⁻⁵² Evidence of direct health effects of such clothing-related exposures
is lacking. Comprehensive reviews of textile-related health studies can be found in the Swedish
Chemicals Agency's report³ and in the opinion statement of the German Federal Institute for
Risk Assessment.²

171 **2.2** Occurrence, persistence and accumulation of chemicals in clothing

172 The chemicals present in clothing are a mix of those present at the time of purchase (possibly

attenuating with time) and those acquired post-purchase. This mix changes with cleaning

174 practices, storage and wear.

175 2.2.1 Chemicals present at time of purchase

176 Most of the chemicals that have been measured in clothing at the time of purchase are a consequence of manufacturing processes (e.g., dyeing, bleaching, finishing) or have been 177 deliberately added and are intended to be retained during the life of the garment. The latter 178 group, referred to as "auxiliaries," includes antiwrinkling resins, flame retardants, antimicrobial 179 agents, pesticides, surfactants and other coating chemicals. Dyeing involves the largest range of 180 chemicals, with an estimated 800 dyes currently in use.² A move towards more environmentally 181 benign textile dyeing is altering the mix of chemicals used in dyeing.⁵³ Some chemicals in 182 clothing fabrics are present as a consequence of packaging, transport, storage and other processes 183 184 that occur between manufacture and purchase.

185 Chemicals that have been identified on newly purchased clothing include trace elements such as

heavy metals;^{54–62} residual aromatic amines associated with certain azo dyes;^{63,64} quinoline and

187 substituted quinolines;^{65–69} alkylphenol ethoxylates, alkylphenols, bisphenols and

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188	benzophenones; ^{70,71} benzothiazoles and benzotriazoles, ^{68,72-74} dioxins and furans; ¹⁵ PCBs; ^{19,75}
189	organo-phosphorous flame retardants and pesticides; ⁷⁶ halogenated flame retardants; ^{7,8,77}
190	fluorinated surfactants; ⁷⁸⁻⁸¹ phthalate ester plasticizers; ^{82,83} glycol solvents; ⁸⁴ formaldehyde from
191	antiwrinkle resins;40,41,85,86 and common petrochemical fuel constituents such as linear and
192	branched C ₁₀ –C ₁₆ alkanes, C ₃ alkylbenzenes, and straight-chained C ₇ –C ₁₀ aldehydes. ⁸⁷
193	Relatively new chemical analysis techniques are being applied to assess chemicals in clothing.
194	Antal et al.65 described the use of direct analysis in real-time (DART) mass spectrometry to
195	measure more than 40 chemicals in clothing items including alkylphenol ethoxylates, phthalate
196	esters, alkyl amines, aniline, pyridine, quinoline and substituted quinoline. In a recent review,
197	Rovira and Domingo ⁴ reported on chemicals that have a high probability of being detected on
198	clothing, with a focus of the health risks posed by these species. Of special note are extensive
199	government reports from Denmark, ⁸⁴ the Netherlands, ⁵ Germany ² and Sweden ³ that review and
200	critically discuss chemicals found in clothing, especially the chemicals that may be present at the
201	time of purchase.

202 2.2.2 Chemicals acquired post-purchase

Chemicals present in air, especially indoor air, can also be present on clothing exposed to that 203 air.^{30,88–95} A commonly encountered example of chemical uptake from air occurs when clothing 204 205 is exposed to environmental tobacco smoke (ETS). Up to a milligram of nicotine can be sorbed by a square meter of cotton fabric during just a few hours of exposure.⁹⁶ Odors derived from ETS 206 constituents can linger on clothing for hours to days. More generally, how much or how little of 207 a chemical is transferred from air to clothing depends on several factors. One key factor is the 208 partition coefficient between clothing and air (K_{ca}) for the fabric in question. As a rule of thumb, 209 the more an airborne chemical resembles the chemical nature of the fabric that constitutes the 210

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211	clothing, the larger is the value of K_{ca} and consequently the greater is the sorptive partitioning of
212	that chemical to the clothing. The octanol/air partition coefficient (K_{oa}) is a good predictor of K_{ca}
213	for cotton, since cotton is cellulosic, for which octanol is a reasonable surrogate. ^{89–91,97} Values of
214	K_{ca} in relation to vapor pressure for several different fiber types have been reported. ⁹¹ Still
215	needed are systematic investigations of K_{ca} for an array of environmental chemicals to a range of
216	clothing fibers, including wool, polyester, nylon, rayon, and other synthetics, as well as to
217	blends, to better estimate the sorption of airborne chemicals to these fiber types.
218	Clothing can acquire chemicals while in closets, storage containers, and chests. A well-known
219	example is sorption of the chemical agents used as moth repellants: naphthalene, camphor, and
220	<i>p</i> -dichlorobenzene. ^{98,99} Similarly, one would anticipate that phthalate esters or alternative
221	plasticizers would be sorbed to clothing stored in polyvinylchloride (PVC) storage boxes or bags.
222	Contact with surfaces can transfer chemicals to clothing. Such chemicals can migrate through
223	clothing becoming available for dermal uptake. Personal care products and fragrances applied to
224	the skin or hair can also be transferred to clothing via contact. ^{100,101} Clothing can retain certain
225	chemicals transferred from personal care products, exposing the wearer and, in principle, those
226	sharing indoor spaces to such chemicals during storage and during repeat wearings until the item
227	is effectively cleaned.
228	Laundering and dry-cleaning removes certain chemicals from clothing but can add others. The

fraction of a chemical that is removed from clothing during cleaning varies with the nature of the chemical as well as with the cleaning practices, including the detergent or dry-cleaning solvent that is employed. Gong et al.³¹ found that the efficiency with which machine washing removed phthalates from cotton jeans increased with the octanol/water partition coefficient (K_{ow}) of the

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phthalate. During dry-cleaning, clothing can retain chemicals from cleaning solvents that 233 subsequently contribute to personal exposures.^{102,103} During laundering, clothing may acquire 234 scents (e.g., synthetic musks²) and other detergent constituents (e.g., alkylphenol ethoxylates¹⁰⁴). 235 Following the wash cycle, clothing is either air-dried or mechanically dried. When air-dried, the 236 clothing can sorb chemicals from the air in which it is dried. When mechanically dried, some 237 238 chemicals can be thermally desorbed while other chemicals (e.g., fabric softeners introduced using 'dryer sheets') may be sorbed by the clothing. Laundering also results in chemicals being 239 transferred among the differing items that are washed or dried together. Cross-contamination of 240 fabrics during laundering and storage has been reported for permethrin-treated garments.¹⁰⁵ 241 Chemicals on clothing can be chemically transformed to other species. Of longstanding concern 242 243 are the abiotic and microbial reduction of azo dyes to carcinogenic aromatic amines such as aniline, benzidine, and 2-naphthylamine.^{106–111} For example, analysis of 86 textile products 244 purchased in Japan detected aromatic amines at low concentrations in socks, undershorts, pants 245 and other garments.⁶⁴ Oxidants can also degrade azo dyes, as shown by reactions initiated by the 246 hydroxy radical, generating benzene and substituted benzenes.¹¹² Photolytic debromination has 247 been shown to produce low levels of polybrominated dibenzofurans¹¹³ when clothing containing 248

the flame retardant hexabromocyclodecane (HBCD) is dried in the sun.

During wear, clothing acquires skin oils, whose constituents can be altered via microbial activity. Different fiber types promote the growth of different microbes, influencing malodor generation from microbial metabolism of apocrine and sebaceous secretions.^{114,115} Squalene, a major constituent of skin oil,¹¹⁶ has been shown to react with ozone on T-shirts generating products with a range of volatilities.^{117–119} The less volatile products remain on the apparel item, exposing the wearer to species such as C_{27} -pentaenal, C_{22} -tetraenal, C_{17} -trienal and their carboxylic acid

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counterparts.¹²⁰ Squalene also reacts with HOCl, the active ingredient in chlorine bleach, to 256 generate chlorinated squalene products. Three to four chlorine atoms become covalently 257 incorporated into the squalene molecule during a one-hour exposure to 1 ppb HOCl.¹²¹ Such 258 species may not be fully removed from clothing during washing. More generally, bleach oxidizes 259 chemicals on clothing, increasing the water solubility of the contaminants but perhaps leaving 260 behind oxidized and chlorinated residues. Numerous low volatility oxidation products, starting 261 with primary carbonyls and carboxylic acids and evolving to products with high O to C ratios, 262 result when ozone reacts with terpenes or sesquiterpenes transferred to clothing from personal 263 264 care products. Other examples of chemicals generated via reactions that occur on clothing include nonylphenol, a known endocrine disruptor, from the degradation of nonylphenol 265 ethoxylate detergent residues⁶⁵ and formaldehyde from urea-formaldehyde and 266 melamine/formaldehyde resins used as antiwrinkling agents.^{40,41,86} The potential for chemical 267 transformations to occur on clothing is commonly overlooked during assessments of exposures 268 to environmental chemicals. 269

270 **2.3** Mechanisms, quantification and prediction of exposure and transfer of chemicals

Clothing influences chemical exposure by a variety of mechanisms, including some that are 271 complex and poorly characterized. Organizations such as the US Environmental Protection 272 Agency, the World Health Organization and the European Chemicals Agency provide guidance 273 on estimating exposure from consumer articles;^{122–124} however, such recommendations are based 274 on a far-from-complete understanding and are therefore of limited utility in accurately 275 characterizing complex chemical exposures mediated by clothing. Notwithstanding their 276 limitations, these recommendations and models can be combined with stochastic representations 277 of exposure factors and behaviors to estimate population distributions of exposure.¹²⁵ 278

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279 2.3.1 Dermal transfer and absorption

280 Most exposure models of skin contact transfer of chemicals from surfaces are conservative by design, i.e. they account, realistically, for the maximum potential exposure for risk assessment 281 282 and risk management purposes. Exposure is derived from factors including the skin area in 283 contact, the concentration of the chemical in the material, the number, frequency or duration of contact events, the type of contact and a transfer efficiency.^{122,124} The transfer efficiency is the 284 285 fraction of the chemical in the material that transfers during contact events. Experimental measurements of the transfer of pesticides¹²⁶ and fluorescent tracers¹²⁷ from carpet and of 286 permethrin from military uniforms¹²⁸ have been used to quantify transfer efficiency of residues 287 from textiles. Some experimental results used to derive residue transfer efficiency are based on 288 289 low-volatility chemicals directly applied to the side of the textile in contact with the skin. Therefore, the residue transfer model may inaccurately characterize exposure from clothing that 290 has volatile or semivolatile chemicals distributed throughout the fabric. Recognizing that 291 diffusive migration can occur within consumer materials, it has been proposed that a transfer 292 efficiency can be derived from the amount that can diffuse from a thin "contact layer" of the 293 material. The thickness of this layer can be specified for consumer products or can be estimated 294 if diffusion coefficients are known for specific chemical-material combinations.¹²⁹ These models 295 generally do not account for the uptake resistance of skin itself.¹³⁰ 296

Models of sweat-mediated transfer of chemicals from clothing also use a transfer efficiency approach. The leachable fraction is derived from experiments using artificial sweat to extract substances such as dyes^{17,43} and trace elements.⁵⁷ (See also §3.1.3.) Often, the extracted fraction is assumed to be entirely transferred to the skin. Such an approach is likely to overestimate exposures, since only a fraction of the sweat will return to the skin from clothing. For example,

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Meinke et al.¹⁷ extracted fluorescent dyes from a polyester/cotton blend shirt using a sweat simulant and compared the predicted exposure (based on 100% transfer) to that observed in volunteers wearing the shirt during 30 minutes of exercise or for 12 hours of normal activity. In these experiments, less than 1% of the estimated amount of a dye was transferred to volunteers during normal wear or sweating.

Indirect (non-contact) exposure to environmental contaminants can also be influenced by 307 clothing. Clothing has been observed to reduce the transfer of airborne insecticides,¹³¹ phthalate 308 esters²⁶ and organophosphate flame retardants³³ to skin. Some models of indirect dermal 309 exposure to airborne contaminants have assumed that clothing is fully permeable.¹³² Other 310 models assume that clothing is fully impermeable. For example, in estimating dermal uptake of 311 polycyclic aromatic hydrocarbons from barbeque fumes to bare skin. Lao et al.⁸⁸ assumed that 312 areas covered by clothing were fully protected. Between these extremes, a mechanistic modeling 313 approach has been introduced that accounts for the history of clothing, contaminant-transfer 314 315 between clothing and the environment, sorptive partitioning of chemicals to clothing, diffusive 316 and advective transfer through clothing and to skin lipids, as well as resistance to uptake through skin.^{133,134} The clothing component of these models is similar to that used to assess clothing for 317 318 chemical protection¹³⁵ and can account for uptake through clothing from air as well as exposure 319 to contaminants present in clothing when donned. Predictions using such mechanistic models agree reasonably well with urinary excretion rate measurements for the limited number of 320 321 human-subject studies in which adequate information is available to populate the model parameters.^{30,134,136} These models indicate that clothing can either reduce or increase dermal 322 uptake relative to bare-skin uptake (Figure 2). The extent of exposure is predicted to be sensitive 323 to a chemical's partition coefficient between clothing and air (K_{ca}) , ^{89–93,137,138} the efficiency of 324

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chemical removal during laundering,^{31,68,92,139} the air-gap between fabric and skin, laundering
frequency, and the history of the clothing items prior to wear.¹³³ A key advantage of dynamic
mechanistic models is that they can predict how clothing accumulates chemicals under nonequilibrium conditions. Such models can be used to derive simpler exposure heuristics for
classes of chemicals, types of clothing and exposure scenarios for risk assessment purposes.¹⁴⁰

330 2.3.2 Inhalation

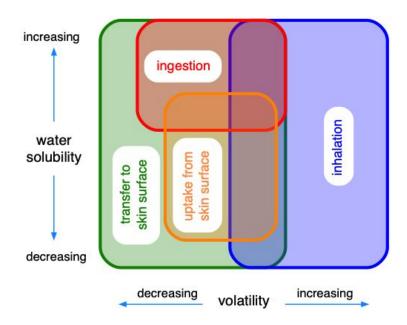
Inhalation exposures for clothing-associated chemicals can be modeled using methods similar to 331 those used to estimate inhalation of chemicals emitted by consumer products. For example, the 332 emission rate of dry-cleaning solvents from clothing hung in a closet can be combined with 333 building air-exchange rates to predict indoor air concentrations,¹⁰³ which are then used to assess 334 inhalation exposures. Inhalation exposure from the emissions that are generated while wearing 335 an article of clothing may be enhanced owing to the "personal cloud" effect, as described for 336 337 particles in §3.1.5. For gaseous pollutants, personal-cloud type alterations have been illustrated in climate chamber experiments investigating transport and pollutant distribution in the breathing 338 zone,¹⁴¹ as well as using computational fluid dynamics to predict breathing zone concentrations 339 of volatile products that result from ozone reactions with the surface of the body and 340 clothing.^{142,143} For a seated person under typical indoor conditions, inhalation exposure to 341 volatile ozone reaction products with skin oils was predicted to be up to 2.5 times higher than the 342 corresponding value for room-average concentrations. Predicted exposure to ozone itself was 343 estimated to be 0.6 to 0.9 times the corresponding condition for room-average concentrations.¹⁴² 344 Simulations are currently limited to simple scenarios, such as stationary seated or standing 345 individuals. Experimental validation of personal cloud effects for clothing-associated chemical 346 exposures are lacking. 347

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348 2.3.3 Ingestion

349 Ingestion by mouthing of fabrics can be a significant exposure pathway, especially for young children. Exposure estimation requires information on the extractability of compounds in saliva, 350 the frequency of mouthing clothing and the area of the fabric mouthed. Extractability can be 351 quantified using a broader set of *in vitro* bioavailability methods, ¹⁴⁴ which have been applied to 352 determine extractability in saliva simulants of azo dyes¹⁴⁵ and for silver from nanoparticles.¹⁴⁶ 353 354 For highly water-soluble species, upper bounds on exposure can be established by assuming that the chemical is completely extractable. In an evaluation of indirect exposure to environmental 355 356 airborne methamphetamine in former residential methamphetamine labs, mouthing of cotton fabric by toddlers was predicted to generate intakes approximately 10 times greater than all other 357 358 exposure pathways combined.⁹⁰

A diagramatic summary of these exposure pathways as influenced by physical-chemical 359 properties is shown in Figure 3. Excepting particles and particle-associated chemicals, inhalation 360 requires a chemical to be volatile enough to become airborne. Ingestion is important for more 361 water-soluble chemicals. Most species can be transferred to skin by contact or transfer through 362 the clothing-skin air gap. For both ingestion and transfer to skin surface, the chemical must be of 363 lower volatility to be present in clothing at meaningful concentrations. Transdermal uptake from 364 the skin surface tends to be highest for chemicals with intermediate volatilities and relatively low 365 water-solubilities. 366



367

Figure 3. The relative importance of clothing-associated exposure pathways based on a chemical's volatility and water solubility.

370

371 **3.** Exposures to particles

372 **3.1 Clothing-associated exposures to biotic and abiotic particles**

373 Ample evidence from environmental and occupational exposure studies indicate that clothing

374 can act as an important source of particle-borne agents that contribute to human exposures.

375 Clothing-associated exposures have been observed for biotic and abiotic particles, with varied

acquisition, retention and release mechanisms, exposure routes and potential health outcomes.

377 This section consolidates evidence from relevant empirical and field studies in the context of a

- 378 providing an overview of exposure to biotic and abiotic particles associated with clothing.
- 379 3.1.1 Allergens
- 380 Exposure to allergenic biological particles from clothing has been well-studied, including cat
- allergen (Fel d 1), dog allergen (Can d 1), dust-mite allergen, and pollens. Tovey et al.¹⁴⁷ were
- among the first to identify clothing as a significant source of inhalable allergens. They showed

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that allergenic dust particles can become resuspended directly from clothing by body movement 383 and can travel to the wearer's breathing zone by means of the thermal plume, thus causing 384 increased allergenic exposures. Other studies found that exposures to mite and cat allergens were 385 closely related to the quantity of particle-bound allergen found on wearer's clothing, suggesting 386 that personal clothing could be an important factor influencing both mite and cat allergen 387 exposure.^{148,149} Evidence of allergen exposure also has been reported for people that are not in 388 direct contact or proximity to any allergenic source.^{148,150} These studies identified clothing as an 389 important indirect exposure vector, transporting particle-borne allergens from one space to 390 391 another.

Much prior evidence concerning clothing-mediated exposures to pet allergens has focused on 392 school environments.^{151,152} Studies have found that allergens can be transported on children's 393 clothing from homes to schools, including both the cat allergen, Fel d 1,^{148,153} and the dog 394 allergen, Can f 1.154 Children without pets can also acquire allergens while in school and 395 subsequently bring them back to their homes.¹⁵⁵ A study focusing on exposure interventions 396 397 found that the level of airborne cat allergens in schools could be effectively mitigated either by pet ownership prohibition or by using school uniforms.¹⁵⁶ Additional evidence has shown that 398 clothing can be a transport vector for the mite allergens, Der f 1 and Der p 1,^{154,157} and for 399 allergenic pollen.^{158–160} Taken together, this body of research persuasively documents that 400 clothing can be an important secondary source of allergenic exposures in buildings, including 401 environments that are free of direct allergenic sources. 402

403 3.1.2 Pathogenic microbes

A second category of clothing-related biological particles is pathogenic microorganisms that
 pose threats for the transmission of infectious diseases. Most research about clothing-associated

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406	pathogens has focused on health-care settings, owing to concern about hospital-acquired
407	infections. ^{6,161,162} Studies have identified pathogenic bacteria on physicians' white coats, ^{163–165}
408	on neckties, ^{166,167} on gloves, ¹⁶⁸ on nurses' uniforms, ^{169,170} and on the coats of medical
409	students. ^{171,172} A commonly detected pathogen on health-care apparel is methicillin-resistant
410	Staphylococcus aureus (MRSA). ^{164,165,168,170,173–176} Other pathogenic bacteria found on
411	healthcare workers' uniforms have included Clostridium difficile ¹⁷⁰ and vancomycin-resistant
412	Enterococcus (VRE). ^{165,168,170,176} In addition to bacterial pathogens, analysis of clothing samples
413	worn by caregivers and visitors has revealed the presence of respiratory syncytial virus, a major
414	cause of respiratory infections among premature infants. ¹⁷⁷
415	Other studies have provided evidence that links bacterial occurrence in clothing with subsequent
415 416	Other studies have provided evidence that links bacterial occurrence in clothing with subsequent exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air
416	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air
416 417	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air has been identified in operating theatres, ^{178,179} in isolation wards ¹⁸⁰ and in hospital storage
416 417 418	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air has been identified in operating theatres, ^{178,179} in isolation wards ¹⁸⁰ and in hospital storage rooms. ¹⁸¹ Early research indicates that pathogen liberation from clothing into air can occur by
416 417 418 419	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air has been identified in operating theatres, ^{178,179} in isolation wards ¹⁸⁰ and in hospital storage rooms. ¹⁸¹ Early research indicates that pathogen liberation from clothing into air can occur by human movement and by frictional interactions between clothing fibers and skin. ^{178,182–184} A
416 417 418 419 420	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air has been identified in operating theatres, ^{178,179} in isolation wards ¹⁸⁰ and in hospital storage rooms. ¹⁸¹ Early research indicates that pathogen liberation from clothing into air can occur by human movement and by frictional interactions between clothing fibers and skin. ^{178,182–184} A seminal study by Duguid and Wallace ¹⁸² found that clothing can liberate pathogenic microbes by
416 417 418 419 420 421	exposure. The direct dispersal of <i>Staphylococcus aureus</i> and other bacteria from clothing into air has been identified in operating theatres, ^{178,179} in isolation wards ¹⁸⁰ and in hospital storage rooms. ¹⁸¹ Early research indicates that pathogen liberation from clothing into air can occur by human movement and by frictional interactions between clothing fibers and skin. ^{178,182–184} A seminal study by Duguid and Wallace ¹⁸² found that clothing can liberate pathogenic microbes by promoting skin shedding. That same study also showed that sterile, dust-proof fabrics can act as

During the past few decades, embedded nanomaterials have emerged as a class of technological innovations for improving certain features of clothing fabrics, such as reducing microbial growth and survival, protecting against ultraviolet radiation, and improving water repellency. To achieve specific targeted functions, prevalent nanostructured clothing additives have included titanium dioxide (TiO₂), silver, zinc oxide (ZnO), gold, copper, carbon nanotubes, and nanoclays.¹⁸⁵ An

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429	emerging consensus indicates that excessive exposure to nanomaterials can contribute to
430	detrimental health outcomes, including pulmonary inflammation, carcinogenicity, genotoxicity
431	and circulatory effects. ¹⁸⁶ The effects of nanomaterial additives in clothing on human exposure
432	and consequent health effects remain a subject of debate. Such materials have the potential to be
433	released from clothing fabrics and contribute to exposures of their wearers and others. The
434	mechanisms of release from clothing are different for nanomaterials as compared with biological
435	particles. For example, in addition to mechanical abrasion, nanoparticles can potentially be
436	released from clothing by migrating into human sweat and saliva. ¹⁴⁶
437	To date, most exposure-related studies have focused on the migration of silver nanoparticles
438	from clothing into human sweat, 146, 187, 188 their release during laundering, 189, 190 and their
439	antimicrobial properties. ¹⁹¹ Dermal exposure to clothing-embedded nanoparticles has not been
440	rigorously investigated. One group of studies reported that TiO_2 and ZnO nanoparticles do not
441	penetrate deeply into the skin. ¹⁹² To the contrary, there is evidence of the increase of the ⁶⁸ Zn
442	isotope in the blood of a healthy adult after exposure to ⁶⁸ ZnO nanoparticles in a sunscreen
443	formulation. ¹⁹³ One study reported that healthy skin is a more effective barrier for silver
444	nanoparticles than damaged skin. ¹⁹⁴

445 Overall, there is a need for more research to characterize the influence of antimicrobial agents, 446 including nanoparticles, on microbial diversity in clothing and on the development of microbial 447 resistance over time. Whether the presence of nanomaterials on fabrics in contact with the skin 448 could alter the local skin microbiota remains a key open question.

449 3.1.4 Para-occupational exposures

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450	Studies have reported instances of para-occupational (or "take-home") exposures to hazardous
451	particles encountered in workplaces. Most such studies have focused on asbestos. As reviewed
452	by Donovan et al. ¹⁹⁵ and Goswami et al., ¹⁹⁶ there is abundant evidence for increased risks of
453	mesothelioma and lung cancer owing to para-occupational exposure to asbestos fibers and
454	asbestos-containing dust on workers' clothing. However, relatively little research provides
455	quantitative evidence that mechanistically links workplace encounters with subsequent
456	household exposures. Sahmel et al. ¹⁹⁷ found that handling clothes contaminated with chrysotile
457	asbestos resuspends 0.2–1.4% of the material. Sanon and Watkins ¹⁷⁶ demonstrated that
458	healthcare uniforms can act as a vector for pathogen transmission outside of hospitals. Overall,
459	the take-home effect for particles and microbial exposure via clothing seems to be a plausible
460	route of transmission worthy of increased attention.

461 3.1.5 Personal cloud

An enhancement of inhalation exposure to particles beyond the room-average levels may occur
for clothing-associated particle releases. This feature, termed the "personal cloud," was
introduced for clothing-mediated chemical exposures in §2.3.2.

There are multiple dimensions to the clothing-associated personal cloud effect. Key determinants 465 involve size-dependent emission rates of particles from clothing, proximity of clothing to the 466 breathing zone, and local air movement in relation to personal activities. Several studies suggest 467 that direct shedding from clothing surfaces may be a noteworthy source of coarse-mode particles 468 and bioaerosols indoors, 198-201 but none of them quantified contributions to the personal-cloud 469 effect. A recent study by Licina et al.²⁰² reported that clothing movement can release coarse 470 particles into the perihuman space of a seated person, which can then be transported upwards by 471 means of the metabolically induced thermal plume. In that study, the contribution of such 472

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releases to the personal cloud was substantial — from 2 to 13 μ g/m³ in the particle diameter 473 range 1-10 µm. The contribution of clothing-associated particle release to a personal cloud effect 474 was observed only for seated occupants and specifically not observed during walking. The study 475 suggests that the personal cloud is contingent on physical activities and that manipulating the 476 metabolic thermal plume could alter exposure to clothing-released particles. Additionally, during 477 478 more intensive clothing manipulations, such as putting on and taking off a shirt, or folding and unfolding a shirt, sharp peaks in the breathing zone PM_{10} mass concentration were detected, at 479 times exceeding 40 μ g/m³. Overall, the emerging evidence regarding the personal cloud 480 481 combined with evidence that clothing can harbor allergens, potentially pathogenic microorganisms, and other harmful substances suggest that clothing surfaces may be an 482 underappreciated factor influencing particle exposure, possibly with public-health relevance. 483

484 **3.2 Toward quantifying clothing-mediated particle exposures**

The previous section summarized evidence that clothing-mediated exposures to particles are potentially meaningful in diverse circumstances. It is important to characterize exposures quantitatively and – in as far as it is possible – mechanistically, so that one is able to extract generalizable findings from limited experimental evidence. In this section, we outline a framework that could guide and support systematic knowledge acquisition for better understanding how clothing influences inhalation exposures to biotic and abiotic particles.

The central element in this framework is the determination of size-resolved and composition specific emission rates of particles associated with clothing. Such emission rates can be expressed in terms of particle mass per time or particle number per time. Composition is key in relation to health outcomes of concern: allergenic particles, infectious microbes, and abiotic particles each contribute to increased yet distinct adverse health risks.

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496	For known clothing-associated particle emission rates, contributions to exposures can be
497	assessed. For example, particle emission rates from clothing can be incorporated into material
498	balance models to estimate the component of exposure associated with increased indoor air
499	concentrations. ²⁰³ Alternatively, the intake fraction approach can be applied to estimate mass or
500	particle number inhalation increments directly from emission rate information. ²⁰⁴ Additional
501	contributions to exposure from the personal-cloud effect can be assessed based on
502	experimental ²⁰² or numerical ¹⁴³ evidence.
502	experimental ²⁰² or numerical ¹⁴³ evidence.
502 503	experimental ²⁰² or numerical ¹⁴³ evidence. It is reasonable to expect that particle exposures associated with clothing occur mainly indoors.
	-
503	It is reasonable to expect that particle exposures associated with clothing occur mainly indoors.

507 activities.^{199,202} It is also plausible to infer emission rates from field observations; however, doing

so for clothing-associated particle emissions poses the challenge of separately accounting for

resuspension from flooring, commonly an important source of coarse particles indoors.²⁰⁵

In assessing clothing-associated emissions, it is worthwhile to differentiate broadly among three 510 particle source categories (see Figure 4). One category is skin flakes, known as squames, 511 generated through frictional interaction between clothing and skin. These squames consist of 512 skin fragments with associated microbes, especially bacteria. A second category would be 513 particles endogenous to the clothing fabric, such as fabric fragments and nanoparticle additives. 514 A third category, the broadest, is exogenous particles that become associated with clothing 515 articles by means of environmental transfer. The first category has been studied most carefully in 516 517 connection with concerns about hospital-acquired infections. Concern about the second category is increasing, in part due to the emerging use of nanoparticle fabric treatments. The third 518

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- category would be relevant for concerns as diverse as allergen exposure, para-occupational
- 520 exposure, and general enhancements of airborne particles via the personal cloud.

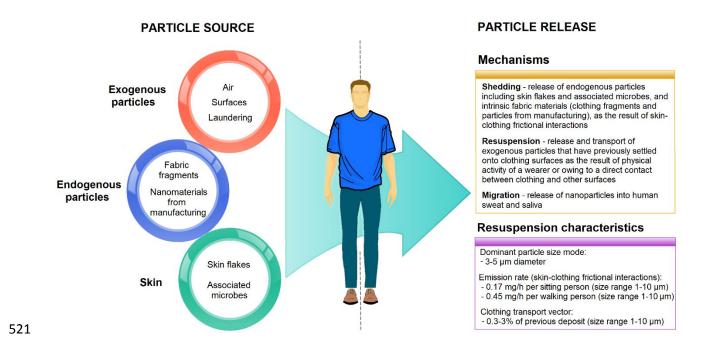


Figure 4. Particle source categories associated with clothing (left); and mechanisms of size
 dependent particle release and resuspension characteristics (right). Corresponding references:
 dominant particle size mode reported by Bhangar et al.¹⁹⁹; size-resolved emission rates from
 sitting and walking person reported by Licina et al.²⁰²; release of previously deposited particles
 from clothing (transport vector effect) reported by Licina and Nazaroff.²⁰⁶

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519

528	For squame emissions associated with clothing, key factors would include the state of the skin
529	surface (dryness, for example), the nature and intensity of frictional interaction between fabric
530	and skin, and the tightness of the weave. Variability in the emissions of skin-associated
531	Staphyloccus aureus among individuals has been demonstrated to be large, and systematically
532	higher for men than for women. ¹⁸⁴ Notwithstanding a long history of studies, the issue of what
533	should be worn by medical staff in the operating theatre to minimize surgical site infections
534	remains a subject of debate. ²⁰⁷ For endogenous particle emissions (e.g., nanoparticle additives),

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one expects that important factors affecting emissions would include initial particle loading of
the fabric, the nature of bonding with fibers, the nature and intensity of movement generating
frictional forces, and the overall wear of the fabric.

For emissions of exogenous particles, one might envision clothing articles as environmental 538 reservoirs and aim to account for the net movement of particles between these reservoirs and the 539 surroundings. Consider an article of clothing, such as a T-shirt, passing through a cycle starting 540 with laundering. The washing cycle might effectively remove previously deposited particles, but 541 could conceivably add particles from dissolved salts in the wash water and from detergent 542 residue. A tumble-dry cycle could effectively add some airborne exogenous particles filtered by 543 the clothing items from the drying air that passes through the drum. The clothing article might 544 545 then lose some of these particles, and contribute an increment of exposure, during the postlaundry handling of folding and placing in storage. When worn, the T-shirt can acquire 546 exogenous particles by deposition from the air and by direct contact with particle-laden surfaces. 547 548 Exogenous particles may also be acquired during storage intervals, especially if exposed in a 549 manner that would be influenced by settling dust. The accumulation of particles during these 550 processes could be quantified through deposition assessments, for example through the 551 multiplicative combination of exposure concentrations of particles, a suitable deposition velocity, 552 and duration of exposure. Knowing the size-resolved and composition-specific quantities of 553 exogenous particles on a clothing article, one could assess the emission rate through the use of 554 loss-rate coefficients. An analogous approach has been used to systematically investigate particle resuspension from walking.²⁰⁵ 555

556 **3.3 Factors affecting clothing-mediated particle exposures**

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The previous section outlined an approach that could be used to systematically assess clothingassociated exposures to particles. Specific information about relevant factors and processes is sparse. This section describes what is known from prior experimental investigations about the major factors that influence the size-dependent emissions of particles associated with clothing, emphasizing the relationship to inhalation exposures.

Early studies revealed important findings about clothing-skin surface interaction as a means of 562 liberating bacteria-laden skin flakes.^{178,179,182–184,208–211} Recent advances in DNA-based 563 measurements have enabled rapid progress in characterizing the human microbiome, including 564 detailed descriptions of diverse communities of bacteria^{212,213} and fungi²¹⁴ present on human 565 skin. Analyses of clothing surface samples or air exposed to clothing have revealed populations 566 of pathogenic bacteria,^{164,176} respiratory syncytial virus,¹⁷⁷ fungi,^{215,216} dust-mite and cat 567 allergens, endotoxins^{157,217} and allergenic pollen.^{158–160} Some quantitative evidence exists 568 documenting microbial transfer to clothing from skin²¹⁰ and by hands,^{218,219} although more 569 570 studies are needed to quantify this phenomenon and to better characterize the process 571 mechanistically.

Available evidence suggests that the rate of particle release from clothing fibers is influenced by 572 a combination of three main factors: properties of clothing, environmental conditions, and human 573 factors. A dominant factor influencing release is the intensity of movement. Up to an order of 574 575 magnitude higher emission rates have been observed during vigorous bodily movement compared to slight activity, presumably owing to increased frictional interactions between 576 clothing fibers and skin.^{182,199,202,220} Men have been found to release significantly more particles 577 compared to women.^{184,211,221–224} Application of skin lotion has been linked to reduced dispersal 578 rate of biotic particles.^{201,223} Some studies,^{221,225,226} but not all,^{223,227} have found that the emission 579

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rate of biotic particles from clothing-skin interactions increases within an hour after showering.

581 Transport of particles through clothing surfaces and subsequent dispersal can be reduced by

582 wearing tightly-woven and non-woven fabrics.^{179,224,228}

A few recent studies have applied a material-balance approach to infer size-resolved biotic 583 particle emission rates associated with human occupancy. Qian et al.²²⁹ used quantitative PCR to 584 infer that a single university classroom occupant contributes effective emissions of 37 million 585 bacterial genomes per hour, with a modal aerodynamic diameter of 3–5 µm. However, that study 586 could not differentiate between emissions associated with clothing and those from other sources 587 such as resuspension from a carpeted floor. Bhangar et al.²³⁰ applied a laser-induced fluorescence 588 technique to quantify the per person emission rate of fluorescent biological aerosol particles 589 590 (FBAP) in the size range $1-15 \,\mu\text{m}$ diameter in an uncarpeted university classroom. Their work, which again did not isolate the contribution of clothing, yielded an average emission rate of 2 591 million FBAP per hour with modal diameters of $3-4 \mu m$. In a subsequent chamber study, 592 Bhangar et al.¹⁹⁹ found that at least 60-70% of occupancy-associated FBAP emissions originated 593 594 from the floor. However, they also found that "clothing, or its frictional interaction with human 595 skin, was ... a source of coarse particles, and especially of the highly fluorescent fraction." That 596 study also revealed a dominant size mode for FBAP of 3-5 µm diameter.

When considering the specific issue of infectious disease transmission in relation to clothing, the persistence and survival of infectious agents on fabrics needs to be considered. Variation in building environmental conditions and properties of clothing fabrics produce various effects on microbial persistence and survival.^{231,232} Longitudinal assessment of bacteria survival across different studies showed a remarkably high persistence — from several days up to more than 90 days for isolates of VRE and MRSA.^{176,233,234} Survival and persistence of viruses and fungi on

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clothing fabrics has similar days- to months-long time scales.^{215,235} Among different factors
influencing survival and persistence, relative humidity and fabric material have been explored.
Increased relative humidity (from 35 to 78%) has been linked to reduced stability of both
bacterial and viral strains in clothing.^{235,236} Survival and persistence of bacteria, virus and fungi
is higher on commonly used polyester and wool fabrics compared to cotton materials.^{215,235–237}

Another common theme in the literature concerning clothing-mediated exposure to pathogenic 608 microbes considers the effectiveness of laundering practices such as washing, drying and ironing. 609 Mechanical removal includes fabric agitation assisted by surfactant properties of detergents, 610 while inactivation processes can occur as a consequence of elevated water temperature combined 611 with laundry additives such as sodium hypochlorite. Among relevant studies, Callewaert et al.²³⁸ 612 documented microbial exchanges among clothing articles during washing. Nordstrom et al.²³⁹ 613 found that home-washed hospital scrubs had increased prevalence of bacterial species compared 614 to those laundered in hospitals, presumably due to low temperature washing. A 7-log reduction 615 in bacterial load can be achieved by 10-minutes of washing with 60 °C water.²⁴⁰ Adding sodium 616 617 hypochlorite to a detergent is an effective way to eliminate bacteria and inactivate enteric and respiratory viruses;^{241,242} however, it might also lead to increases in the abundance of chlorinated 618 619 organic compounds on clothing.¹²¹ Detergents free of bleach can reduce the prevalence of *Staphyloccus aureus*,²⁴³ while adding bleach-enriched detergents completely eliminates the 620 621 same. Recent adjustments in laundering procedures include addition of enzymes, reduced water use, lower water temperature and bleach-free detergents.^{244–246} 622

Both biotic and abiotic material can be deposited onto clothing surfaces from various

- 624 environmental sources including outdoor air,^{158,159,247} grassland,¹⁶⁰ residential air,²⁴⁸ public
- transport microenvironments,²⁴⁹ and from physical contact with items such as furniture, storage

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surfaces and car seats.^{250–252} The rate of deposition from air to clothing can be described using
the deposition velocity concept.²⁵³ Studies have found that particle size and local air movement
are dominant influencing factors.^{249,254}

Research has clearly documented that previously deposited material can be released into air from clothing.^{197,206,255,256} For example, using a controlled chamber study approach, Licina and Nazaroff²⁰⁶ found that 0.3-3% of deposited particles (size range 1-10 μ m) deposited through settling could be released via fabric motion. In that work, the release fraction monotonically increased with particle size.

The degree of particle binding to clothing fibers and the rate of resuspension may arise from a combined influence of different forces acting upon the fibers. The forces governing the release of clothing-embedded particles are abrasive actions between clothing surfaces – a consequence of physical activity of a wearer.^{182,184} Forces influencing release are strongly linked to particle size. Because detachment forces increase more strongly with particle diameter than do adhesion forces, clothing-associated emissions are more discernible among coarse-mode than fine-mode particles.^{199–202}

Common clothing fibers are wool, cotton and polyester. Wool has been reported to have particle release rates up to 10× times higher than the other two materials;^{149,257} cotton exhibits higher emissions than polyester.²²⁰ The higher particle emissions from wool garments could be linked to different surface roughness and weave pattern,²⁵⁸ but also to less frequent laundering as compared to cotton and polyester fabrics.²⁵⁷ Other clothing conditions found to increase particle release rate include increased clothing age²⁵⁹ and reduced cleanliness.^{1496,257} While it is generally understood that adhesion forces acting on particles increase with relative humidity, we know of

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only two studies that have examined its effect on clothing-associated emissions. Yoon and
 Brimblecombe²⁵⁷ found an association between low relative humidity and increased particle
 emission rate, whereas Zhou et al.²⁰¹ reported an insignificant influence.

651 **4. Future outlook**

There is ample evidence that clothing influences human exposure to chemicals and particles.
Yet, only a few studies have quantified clothing-mediated exposure by means of direct
measurements.^{7,11,15,24–26,28–30,149,202}

We know surprisingly little about the occurrence of contaminants acquired by everyday clothing after purchase. For a relatively low cost, we could learn a large amount from simply assessing the occurrence, concentrations and extractability (e.g., by sweat and saliva) of chemicals and particles in everyday clothing. Cross-sectional exposure studies would greatly benefit from the addition of clothing analyses, potentially identifying direct connections between clothingassociated exposure and health.

The diversity of clothing, environmental and human factors make predicting exposures 661 challenging. Therefore, it will be important to reduce the many variables to those that are most 662 influential. Progress can be achieved through models, laboratory and field investigations of 663 human exposure and uptake. In addition to chemical properties, important factors affecting 664 exposures may include textile materials, weave, thickness and permeability; wear, care and 665 666 storage practices; environmental conditions; intensity and types of activities; skin-oil transfer to clothing and its aging; human physiology (skin integrity, lipid generation, sweating) and personal 667 hygiene habits. Simulated exposures with human subjects also should consider pollutant transfer 668

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669	from textiles other than clothing (e.g. pillows, quilts, bed linen). The sleeping environment is
670	potentially of great importance in this matter given the large proportion of time spent in bed.
671	Predicting and controlling exposure rely on adequate understanding of underlying mechanisms.
672	A robust literature describes transport mechanisms for chemicals among environmental
673	reservoirs. Reasonable approaches have been proposed for assessing risk and exposure to
674	chemicals in clothing. However, we have limited in vivo evaluations of such assessments.
675	Compared with chemical transport, mechanisms of particle uptake and subsequent release from
676	clothing are even less well understood. Further quantitative investigations of factors that drive
677	acquisition, retention and transmission of biotic and abiotic particles in clothing are needed to
678	better link such processes to clothing associated exposures. We also need to better understand the
679	extent to which clothing plays a role in the spread of infectious disease. Considerable research
680	has focused on textile innovations and personal protective clothing designed to limit the spread
681	of infectious agents in hospital environments. Researchers could usefully build upon lessons
682	learned and consider the potential utility of incorporating such innovations in everyday clothing.
683	One should anticipate that future changes in clothing will influence exposure. The useful lifetime
684	of some clothing has become shorter. High turnover (short ownership time) might yield greater
685	exposure to chemicals that are present in newly purchased clothing, with proportionately less
686	exposure to environmental chemicals that require a long period to equilibrate (e.g., high
687	molecular weight phthalates). Similarly, increased use of antimicrobial agents as coatings on
688	clothing articles may increase uptake of nanoparticles by the human body and lead to altered
689	toxicological effects. Worth noting is that people in Western countries commonly have closets
690	full of clothes that are rarely worn. These articles may have sufficient time to equilibrate with the
691	chemicals present in their storage environment. Worldwide, demand for synthetic fabrics is

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- 692 increasing.²⁶⁰ Synthetics have chemical partitioning behaviors and moisture holding capacities
- 693 that differ from those of natural fibers, altering the capacity to be reservoirs of contaminants.
- Advances in materials and adjustments in laundering procedures may also influence how
- clothing is cared for and how chemicals and particles are acquired and retained in clothing.
- 696 Increased recycling and re-use of clothing can influence tertiary exposures.
- 697 People spend nearly their entire lives in intimate contact with clothing and other textiles. The
- 698 evidence reviewed in this article supports a view that this environmental compartment plays
- 699 important roles in exposure and health risk. Consequently, clothing as a mediator of chemical
- and particle exposure deserves substantial attention from the environmental science research and
- 701 regulatory communities.

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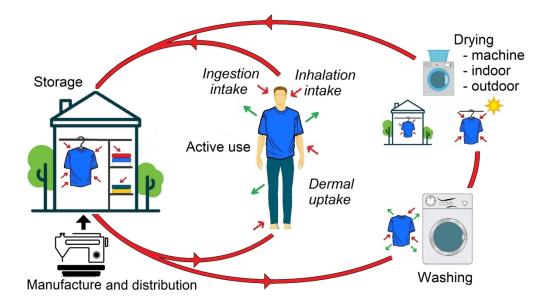
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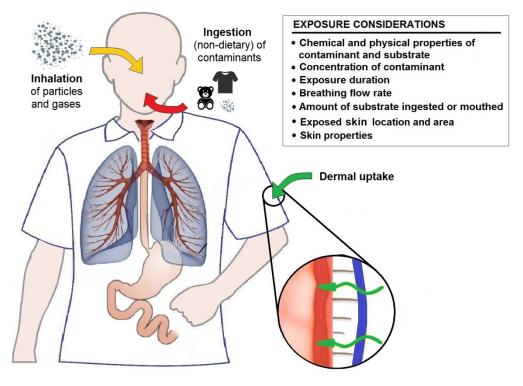


Figure 1. Non-dietary routes of human exposure for contaminants of concern: Inhalation, ingestion and dermal absorption.

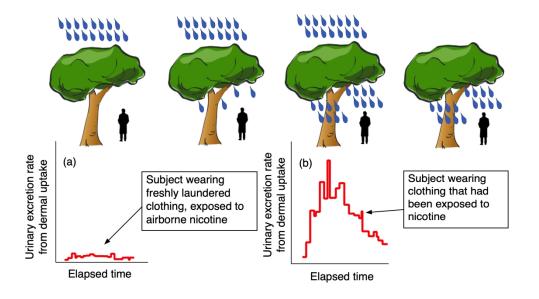


Figure 2. Dynamic urinary excretion rates due to dermal uptake for nicotine and two urinary metabolites (summed) after exposing participants, who were wearing breathing hoods, to airborne nicotine.²⁹ (a) Freshly laundered clothing, unexposed to nicotine, is protective; this is analogous to a tree at the beginning of a rainstorm that protects a person from getting wet. (b) Clothing that has been previously exposed to airborne nicotine dramatically increases urinary excretion rate for days after wearing the clothing, just as a standing under tree after a long rainstorm is most certain to get the person wet. This exposure occurs while wearing the clothes in the environment containing the contaminant (third tree from left) and can continue after leaving this environment

224x129mm (300 x 300 DPI)

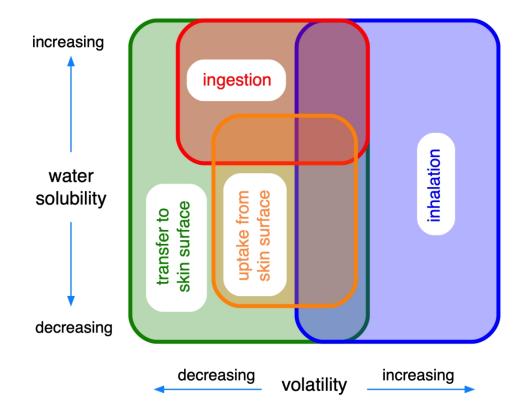


Figure 3. The relative importance of clothing-associated exposure pathways based on a chemical's volatility and water solubility.

133x106mm (300 x 300 DPI)

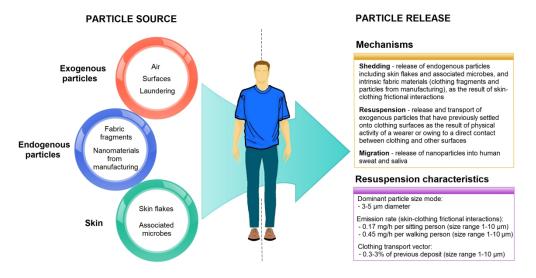


Figure 4. Particle source categories associated with clothing (left); and mechanisms of size dependent particle release and resuspension characteristics (right). Corresponding references: dominant particle size mode reported by Bhangar et al.¹⁹⁹; size-resolved emission rates from sitting and walking person reported by Licina et al.²⁰²; release of previously deposited particles from clothing (transport vector effect) reported by Licina and Nazaroff.²⁰⁶