# DEVELOPMENT OF COATINGS FOR NONWOVEN GARMENTS TO PROVIDE BOTH REASONABLE COMFORT AND PROTECTION AGAINST BLOODBORNE PATHOGENS Gemma T. Kennedy **Biomedical Development Corporation** San Antonio, TX **Bakul M. Bhatt** University of Texas Health Science Center and **Biomedical Development Corporation** San Antonio, TX **Charles J. Gauntt and H. Ralph Rawls** University of Texas Health Science Center San Antonio, TX **Tyrone L. Vigo USDA-ARS** Southern Regional Research Center New Orleans, LA

#### Abstract

A microporous polyvinylidene fluoride film sandwiched between non-woven fabric demonstrated adequate comfort and blocked penetration by bacteria and a 150-200 nm blood-borne virus, but allowed a 33 nm virus to pass. Improvements to prevent pinholes and other film defects are expected to lead to protective garments that meet OSHA standards for health care workers while providing adequate comfort.

# **Introduction**

The 1992 OSHA standard mandates that protective clothing designed to protect against bloodborne pathogens must provide an efficient barrier to blood and body fluid penetration ("strike-through") of such viruses as HIV (~100 nm) and Hepatitis B (~50 nm) (USDL 1992). Unfortunately, thermal comfort must be sacrificed to prevent penetration by the smallest viruses (Vigo 1994; Schoenberger; White and Blood 1993). A polyvinylidene fluoride (PVDF)-based fabric coating can form a flexible, hydrophobic, microporous membrane (Kaminska et al., 200). We hypothesized that such a film could be applied to either *span the spaces between fibers* or be placed as a membrane between fabric layers, and that such a membrane (having a tortuous capillary bed porosity) would allow moisture vapor to pass while blocking passage of blood and other body fluids. The idea is to exclude bacteria & viruses while allowing air and water vapor to pass, thus keeping personnel comfortable but protected from "blood-borne" pathogens.

## **Materials and Methods**

#### **Independent Variables**

*Fabric construction:* non-woven (PGI 94Q5: 55/45 cellulose/polyester, 3.7 oz/yd<sup>2</sup>). *Fiber surface energy:* hydrophilic *vs.* hydrophobic (use of Zony® fluoro-surfactants). *Formulation:* PVDF copolymers, polymer blends, solvent (acetone & n-butyl acetate), concentration. *Coating technique:* Cast directly on fabric *vs.* form film and apply as membrane sandwiched between fabric layers.

#### **Dependent Variables**

Water repellency (contact angle), comfort (drape, vapor transmission), low-pressure water penetration (dye "strike through"), and microorganism penetration — resistance to penetration by large and small viruses under high hydraulic pressure (ASTM F1670-95; 2psi, 60 min).

#### **Results**

## **Comfort**

Conditions were found for which little or no reduction occurred for either drape/suppleness or moisture vapor transmission rate (MVTR) / breathability, for fabrics prepared by both direct and indirect coating techniques.

# **Biological Safety / Direct application**

Conditions were found with adequate drape and MVTR, which passed the high-pressure strike through test and blocked transmission of *S. epidermidis* (~ 1  $\mu$ m), but failed to block penetration of *Herpes Simplex* (100 nm) and polio (30 nm) viruses. Indirect application: Samples prepared with membranes cast from 15% polymer had adequate drape and MVTR, passed the high-pressure strike through test, blocked *Herpes Simplex*, but failed to stop penetration of the smaller poliovirus.

## **References**

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<u>Note</u>

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# Moisture Vapor Transmission Rate

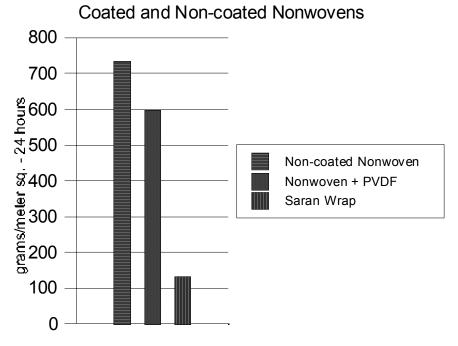
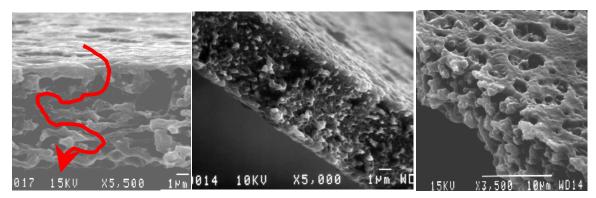


Figure 1. Moisture Vapor Transmission Rate.



Pores range up to approximately 1 μm. Figure 2. Scanning Electron Micrograph of freeze-fractured PVDF-based coating.