Virally-Induced, Intracellular Biofilms; Novel Findings in Molluscum Contagiosum

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Received date: October 10, 2017; Accepted date: October 24, 2017; Published date: October 26, 2017

Abstract
Both the presence and impact of biofilms have proved to be groundbreaking regarding early diagnosis and treatment in acute and chronic cutaneous, neurological and other internal diseases. Further, biofilms and the activation of the innate immune system have added clarity to the pathogenesis of those diseases. Our current observations are the first to demonstrate viral biofilms in skin disease; these observations are also the first to demonstrate intracellular biofilms in skin disease. We have observed these in lesions of Molluscum contagiosum (MC). The only previous observation of viral-induced biofilms has been with the HTLV-1 virus. The essential elements of a biofilm are extracellular polysaccharides (EPS) that form the bulk of the biomass, and amyloid fibers that form the proteinaceous infrastructure of the biofilm. Histopathology of skin lesions revealed positive periodic acid Schiff (PAS) and positive Congo red (CR) and crystal violet (CV) stains within MC lesions. The PAS stains the EPS while the Congo red and crystal violet stain the amyloid. Similar staining was absent both in controls and in surrounding skin; this is strong support for the hypothesis that the virus "hi-jacks" the cell's DNA and makes "intra"cellular biofilms.

Keywords: Viral biofilms; Intracellular; Molluscum contagiosum; Dermatopathology

Introduction
We present, for the very first time, observations that a virus, Molluscum contagiosum (MC), makes biofilms in a skin disease. Only once in the medical literature has a virus been shown to make biofilms, namely the HTLV-1 virus [1]. Further, we present the novel finding that the MC virus causes biofilms to be made inside epidermal cells which is the first instance of "intra"cellular biofilms in dermatologic disease. We have shown this with histopathology of MC and have identified the principal elements of a biofilm (polysaccharides and amyloid) in molluscum lesions by utilizing periodic acid Schiff (PAS) and Congo red (CR) stains [2]. The amyloid was further confirmed with crystal violet (CV) staining. The MC lesions from ten patients were studied.

Methods
This study was approved by the Drexel University College of Medicine Institutional Review Board.

Ten histologic specimens of MC were examined microscopically by four dermatopathologists. Hematoxylin and eosin (H+E), PAS, CR and CV staining was performed. Ten control specimens from cutaneous lesions of atopic dermatitis were examined with the same staining procedures. Each of the stained specimens was examined by four dermatopathologists.

Results
The diagnosis of MC on routine H+E staining was readily apparent with each specimen demonstrating the hallmark features of a molluscum lesion: central punctum, lesional epidermal hyperplasia, and typical Henderson-Patterson bodies. The PAS stains revealed focal intracytoplasmic staining in the Malpighian cells just above the basal zone. (Figure 1) The CR stains were positive in a similar location, as were the CV stains. (Figure 2) The atopic dermatitis specimens showed no similar intracytoplasmic staining in the Malpighian cells; they did show staining in the distal sweat ducts (which is the pathologic signature in atopic dermatitis) [2]. The positive staining was present in 10/10 specimens (P=0.001, $\chi^2$). Each specimen also served as its own control with the lesions showing changes and the adjacent (normal) epidermis demonstrating no similar findings.

Figure1: PAS positive deposits in the Malpighian cells above the basal layer (arrows). This represents staining of the polysaccharides in a biofilm. Similar deposits are not found in the adjacent epidermis. Molluscum bodies are clearly present (PAS stain 10X).
those lesions.

organisms.

biofilm

biofilms

exporter cells, and water channels [3].

polysaccharides, amyloid, DNA, fatty acids, protein, dead cells, intracytoplasmic; the concept of "intra" cellular polysaccharides in the

ISSN:2327-5073

Clin Microbiol, an open access journal

3 nm.

a "quorum" and begin forming a

organisms where ten microbes (in any direction) are necessary to make

contains 250 organisms [7]. Whether the same parameters exist in

understood.

That

biofilm

would be 0.003 µ given the size of the MC organisms as 1 × 2 ×

calculation would lead to a total volume for that

biofilm, on the other hand, is clearly intracytoplasmic; the concept of "intra" cellular biofilms is well established and has been seen in such diverse conditions as urinary tract disease, pneumonitis and Alzheimer's disease [8]. The intracellular microbes in a biofilm are doubly protected from the immune system and other disease modifying agents. Viruses have been shown to assist bacteria (filamentous phages and Pseudomonas) in biofilm production and to enhance biofilm growth (Pseudomonas and respiratory syncytial virus) via dysregulation of immunity from alteration of iron metabolism [9,10]. This may be a potential role for the Herpes simplex virus that has been found in the brains of Alzheimer's patients [11] namely assisting the spirochetes that are present there to create biofilms and activate the innate immune system [3].

Treatment that might arise from this observation could possibly involve a biofilm dispersing agent [12] but whether that agent would penetrate through the punctum to the depths of the lesion and then through the cell wall, is questionable. This work does offer a protocol to investigate other viral diseases to discover whether those create biofilms. This is compelling in that most organisms prefer to live in communities rather than in the planktonic state [13].

References