UV Light for Processing Foods

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ABSTRACT
Ultraviolet light (UV) light holds considerable promise in food processing as an alternative to traditional thermal processing. Its applications include pasteurization of juices, post lethality treatment for meats, treatment of food contact surfaces and to extend the shelf-life of fresh produce. This paper will review published studies and commercial applications that utilize UV treatment for solid and liquid foods. Designs of UV reactors that were tested in the author's lab for juice and apple cider treatment are discussed. Future research needed to extend the range of UV light applications in food processing industry is presented.

Key Words: UV light, food, juices, pasteurization, absorption, food pathogens, vitamin C

INTRODUCTION
The use of ultraviolet light (UV) light is well established for water treatment, air disinfection and surface decontamination. With the growing negative public reaction over chemicals added to foods, UV light holds considerable promise in food processing. As a physical preservation method, UV irradiation has a positive consumer image. The US Food and Drug Administration (FDA) and US Department of Agriculture (USDA) have concluded that the use of UV irradiation is safe. In 2000, the FDA approved UV-light as alternative treatment to thermal pasteurization of fresh juice products (US FDA 2000). The performance criterion defined by FDA for fruit and vegetable juice processing is a $5\text{-log}_{10}$ reduction in the number of the target pathogen of concern (US FDA 2000). In addition, the definition of “pasteurization” for foods was recently revised and now includes any process, treatment, or combination thereof, which is applied to food to reduce the most microorganism(s) of public health significance (Food Chemical News 2004). The processes and technologies examined in the above mentioned report include UV irradiation as an alternative to heat that can be used for pasteurization purposes.

This paper will provide a general review of the applications and efficacy of UV light treatment of foods. Consideration will be given to research that describes UV treatment as an alternative preservation method for solid and liquid foods. In addition, the emphasis is given to the future research required to extend the range of UV light applications in the food processing industry.

UV surface treatment
UV light is used in the food industry for disinfecting surfaces. Applications include decontamination of surfaces of equipment in bakeries, cheese and meat plants, as an adjunct to usual cleaning and sanitizing practices, and for decontamination of conveyor surfaces and packaging containers such as boxes, caps, bottles, cartons, tubes, films and foils. Despite the efficacy of UV light to disinfect smooth surfaces, there are relatively few applications of this technology in the food processing industry. The restricted range of commercially available equipment for disinfecting solids may contribute to its limited use. In addition, most kinetic data of microbial inactivation were obtained in suspension in aqueous media or air. These data are of limited use in predicting the surface disinfection rate. Since complex interactions may occur between microorganisms and surface materials, such as shielding effects from incident UV, efficacy of UV light depends on surface structure or topography.

The recent outbreaks of Listeria in Ready-To-Eat (RTE) meats prompted the USDA to implement a regulation to control Listeria in facilities producing RTE products (FSIS 2003). Alternatives 1 and 2 of this regulation include the use of a post lethality treatment and/or an antimicrobial agent together or separately to reduce or eliminate the bacterium on the product. The ability of UV light to disinfect surfaces of meat products has been reported in the literature. A few studies have evaluated the use of UV irradiation to reduce levels of Escherichia coli and Salmonella on pork skin and muscle (Wong et al.1998); Listeria monocytogenes on chicken meat (Kim et al. 2002), and Salmonella Typhimurium on poultry carcasses (Wallner-Pendelton et al. 1994). Despite the known limited ability of UV light to penetrate rough food surfaces such as meats, these studies demonstrated that UV light has the potential to reduce bacterial contamination on food surfaces and therefore has the potential to be used as post lethality treatment to control L. monocytogenes and other pathogens of concern.
An example of a commercially available system to decontaminate surfaces of foods is the UV tumbling process that was developed by C&S Equipment Company (www.cs-equipment.com) (Chapman 2003). The company incorporated either a rotating drum or screw conveyor that lifts and tumbles the product to ensure exposure to the UV source. The unit can be used to treat fresh products (vegetables, fruits, meats, etc.), frozen products (vegetables, fruits, meats, seafood, bakery products, etc.), and cooked, refrigerated products (pasta, cheese, etc.). The C&S Equipment Company designs solutions for these problems using the patented technology of Steril-Aire™ UV Emitters. These emitters are sleeved in plastic to meet the food safety requirements of food processing facilities. The patented design allows emitters to work efficiently in the cold environment of refrigerated or chiller coils, where competitive units lose their effectiveness. Examples of UV units currently used in commercial processing facilities include: a) UV tumbling drum in operation for chicken and beef fajita strips (cooked and IQF frozen) with a capacity 6,000 – 7,000 lb / h; b) cooked and IQF frozen hamburger patty treatment (hooded conveyor with turn over), with a capacity 3,000 lb/h; and c) a deli meat system (custom conveyor with UVC hood) for formed deli ham logs, with a capacity of 10,000 lb/h.

UV treatment has also been applied to prolong shelf life of wrapped partially baked baguettes to minimize post baking contamination (Doula et al. 2000). UV light has been documented be effective in reducing various bacterial populations on egg shell surfaces including total aerobic plate count (Chavez et al. 2001), S. Typhimurium and E. coli (Coufal et al. 2002), and Yersinia enterocolitica (Favier et al. 2001). Despite the urgent need to improve egg safety, UV treatment of eggs has not been yet commercially implemented.

Liquid foods and beverages

UV light has considerable promise to reduce the levels of microbial contamination for a wide range of liquid foods and beverages. Due to the presence of color compounds, organic solutes and suspended matter, liquid foods such as fresh juice products and beverages transmit relatively little UV light, and this low transmission lowers the performance efficiency of the UV pasteurization processes.

Comparison of absorption coefficients (Figure 1) indicates that the absorbance of fresh juices is significantly higher that that for water. In addition, the absorbance and turbidity of clear fresh juices and juices with pulp varies considerably. Clear apple juice has a low absorbance, with absorption coefficients about 11 cm⁻¹, whereas orange juice can have absorbances close to 50 cm⁻¹ (Koutchma, Keller et. al. 2004). The turbidity of juices arises from the presence of suspended solids and can be in a range from 1000 NTU for apple and other clear juices to > 4000 NTU for opaque varieties such as carrot, orange and pineapple juices. The juices also have different Brix (soluble solids content) and pH levels, as well as varying viscosities (Figure 2). An increase in viscosity significantly increases the power requirements to maintain the unique and desirable fluid flow characteristics of the individual reactor designs.

![Figure 1: Comparison of the absorption coefficients of water and fresh juices.](image1)

![Figure 2: Comparison of the viscosity of water and fresh juices.](image2)

Absorption coefficients, soluble solids content (deg Brix), pH, color (L, a and b – values) and vitamin C content of three brands of clear apple juices are summarized in Table 1. Three brands of packaged apple juice (pasteurized, no preservatives) were purchased locally and stored at 4°C for the trial.

1. Ocean Spray, plastic bottle (Ocean Spray, Lakeville-Middleboro, MA, abbreviated as OS)
2. Sahara Burst, aseptic box package (Sysco, Houston, TX, abbreviated as SB)

3. Gordon Food Service, aseptic box package (Gordon Food Service, Grand Rapids, MI, abbreviated as GFS)

Sahara Burst and Gordon Food Service brands were enriched with Vitamin C.

A small difference of pH and Brix was observed among juices. SB juice had the highest absorption coefficient and OS juice was the least absorptive in terms of UV light. The correlation between vitamin C content and absorption coefficient can be seen from the data in Table 1. The juices enriched with vitamin C such as SB and OS had the higher absorption coefficients. It was also observed that OS juice was not enriched with vitamin C and had the lowest magnitude of absorption coefficient and L (lightness), a (yellowness), and b (greenness) values. The increase of absorbance of buffer solution with addition of vitamin C is shown in Figure 3. Vitamin C is a light-sensitive vitamin in apple, carrot, orange and vegetable blended juices and can be degraded by UV treatment. The destruction of the vitamin C at varied initial concentration in buffer is shown in Figure 4 demonstrating that destruction kinetics of vitamin C and the effect of vitamin C on absorption coefficient had to be included in the numerical simulation of UV fluence. Juices enriched with vitamin C require higher UV fluence.

<table>
<thead>
<tr>
<th>Apple Juice</th>
<th>pH</th>
<th>Brix</th>
<th>Absorption coefficient mm⁻¹</th>
<th>Vitamin C mg/ml</th>
<th>L-Values</th>
<th>a</th>
<th>b</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sahara Burst</td>
<td>3.49</td>
<td>11.9</td>
<td>3.91</td>
<td>0.30</td>
<td>0.12</td>
<td>5.74</td>
<td>-0.667 ± 0.022</td>
</tr>
<tr>
<td>(SB)</td>
<td></td>
<td></td>
<td>±0.09</td>
<td>±0.25</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ocean Spray</td>
<td>3.44</td>
<td>11.65</td>
<td>0.71</td>
<td>0</td>
<td>0</td>
<td>4.03</td>
<td>-0.35 ± 0.017</td>
</tr>
<tr>
<td>(OS)</td>
<td></td>
<td></td>
<td>±0.12</td>
<td>±0.04</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>GFS</td>
<td>3.51</td>
<td>11.75</td>
<td>3.71</td>
<td>0.45</td>
<td>0.22</td>
<td>±4.67 ± 0.42</td>
<td>-0.39 ± 0.07</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>±0.12</td>
<td>±0.12</td>
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</tbody>
</table>

Table 1: Physical properties of clear apple juices

![Figure 3: Effect of concentration of vitamin C on absorbance of buffer solution.](image)

![Figure 4: Degradation of vitamin in the single lamp annular reactor.](image)
This means that the combination of physical and nutritional properties, such as liquid density and viscosity, vitamin C content must be considered to meet the required pasteurization standard of a $5\text{-log}_{10}$ reduction in the number of the target pathogen of concern for fresh juices. Exposure of microorganisms to UV-light or residence time in the UV reactor should be sufficient to achieve the required level of inactivation.

**Reactor designs for UV treatment of juices**

The correct UV reactor design can reduce the interference of high UV absorbance and viscosity associated with some food products and therefore improves the inactivation efficiency. The flow pattern inside the UV reactor strongly influences the total applied UV dose, since the position and the residence time of the microorganisms in certain regions of the irradiance field can vary significantly. Currently, different continuous flow UV reactor designs are being evaluated for use in fresh juice pasteurization. The first design approach uses an extremely thin film UV reactor to decrease the path length and thus avoid problems associated with lack of penetration. Thin film reactors are characterized by laminar flow with a parabolic velocity profile. The maximum velocity of the liquid is observed in the center, which is twice as fast as the average velocity of the liquid; this results in non-uniform processing conditions (Koutchma and Parisi 2004). The two laminar flow designs shown in Figure 5 are a thin film CiderSure reactor (FPE Inc., Macedon, NY) and the Taylor-Couette flow UV reactor (Forney and Pierson 2004). In the CiderSure unit (Figure 5a), low-pressure mercury arc lamps are mounted within a quartz sleeve running centrally through the reactor. Juice is pumped from a reservoir through a 0.08 cm annular gap between the inner surface of the chamber and the outer surface of the quartz sleeve. Forney and Pierson (2004) developed a UV reactor that pumps fluid through the annular gap between two concentric cylinders, as shown in Figure 5b. To provide sufficient exposure and to reduce the fluid boundary layer thickness next to the UV radiation source contained within the outer stationary cylinder, the smaller inner cylinder (called Taylor-Couette flow) consists of laminar vortices that both fill the annular gap of several millimeters and circumscribe the inner cylinder.

A second design approach increases the turbulence within a UV reactor to bring all material into close proximity of the UV light during the treatment. The higher flow rates achieved under turbulent conditions provide improved homogeneity of the flow when the fastest flowing particle travels 1.1-1.2 times faster than the volume averaged particle, and each volume of the product will be exposed to UV light due to better mixing. Unfortunately, as turbulence increases, the pressure drops across the reactor, and the high flow rate to ensure turbulent flow is coupled to a reduced fluid residence time which can lead to complications scale-up. In the Aquionics UV reactor (Hanovia Ltd, Slough, England), treatment is achieved by passing liquid through a stainless steel chamber containing UV emitting low-pressure arc-tubes (Figure 6a). Each single arc-tube is mounted in a quartz sleeve and fitted within the chamber allowing the liquid to pass the sleeve on all sides (Koutchma et al. 2004). The UV module (Salcor Inc, CA) shown in Figure 4b contains a coiled Teflon tube with 24 ultraviolet lamps and reflectors. The coiled tube promotes additional turbulence and causes a secondary eddy flow effect, also known as a Dean effect, and results in a more uniform velocity and residence time distribution. The lamps and reflectors are placed both inside and outside the coiled tube, increasing not only UV irradiance of the flowing liquid, but its uniformity as well.

![Figure 5: Schematics of (a) a laminar thin film reactor (Cider Sure) and (b) a laminar Taylor - Couette UV reactor](image-url)
for the same time period and all microorganisms would receive an equivalent UV dose, if the UV irradiance were equal at all points. However, it is important to recognize that treatment of some high viscosity fluids or fluids with pulp will be incompatible with some of the reactor designs.

UV light sources

Several alternative UV sources types, such as continuous UV low-pressure and medium-pressure mercury lamps, pulsed-UV, and excimer lamp technologies have been developed and can be applied to foods. However, the efficacy and specific characteristics of common UV light sources that are used today for water treatment have not been evaluated for food applications. Traditional low pressure mercury UV lamps at 254 nm were used for applications for disinfection of food surfaces and food liquid treatments discussed previously.

There are no reports available, except for the study made by Warriner et al. (2002), of applications of excimer lamp technology for foods. UV–excimer lamps can produce monochromatic output that can be tuned to the wavelength of interest by the combination with gases. Excimer lamps also have an advantage of extremely low output and are able to operate at much lower surface temperatures. Thus they can provide an advantage by avoiding fouling behavior by liquid foods. Warriner et al. (2002) demonstrated that UV-excimer light was effectively used for sterilization of the packaging carton surfaces.

A few studies recently reported an application of UV pulsed light for foods. A pulsed xenon UV-light treatment was applied to inactivate spores of Aspergillus niger in corn meal. However, low penetration power and excessive heat buildup inside the chamber was reported (Jun et al. 2003). Pulsed UV light was found to be effective in inactivating Saccharomyces cerevisiae (Takeshita et al. 2003). In addition, the pulsed UV light was used to control microbial levels on fresh processed lettuce. Allende and Artes (2003) reported that pulsed UV light was effective for reducing the levels of psychrotrophic and coliform bacteria as well as yeast without adversely affecting the sensory quality of lettuce. Sharma and Demirci (2003) demonstrated that pulsed UV light holds promise for eliminating pathogens such as E. coli O157:H7 from alfalfa seeds.

CONCLUSIONS

The recent advances in the science and engineering of UV light irradiation have made it a viable option for commercial application in food processing. As a nonthermal alternative to traditional thermal processing, UV light has a potential to be used for pasteurization of juices and beverages, as a post lethality treatment in controlling microbial contamination on meats and shell eggs surfaces, and as a means for the shelf life extension of fresh produce. UV light processing can improve safety of selected solid and liquid foods without appreciable loss in quality or nutrient content.

However, to improve the efficacy of UV light for food application, the following areas of research need to be conducted. To predict UV disinfection rates on food surfaces, more kinetic inactivation data need to be obtained for pathogen and spoilage microorganisms, taking into account interactions between microorganisms and surface materials, such as shielding effects from incident UV and their dependency on surface structure or topography. In addition, novel methods and models need to be developed to measure the inactivation rates or dose–response behavior of food pathogens in highly absorptive and viscous food liquids, such as juices and beverages. The correct choice and/or design of the UV reactor, its flow characteristics and UV source can reduce the interference of high UV absorptivity and viscosity associated with some liquid food products and improves inactivation efficiency. The development of validation methods for food processing facilities requires identification of surrogate microorganisms or suitable actinometers for pathogens. Research in the indicated areas can ensure the effectiveness of UV light for microbial inactivation, stimulate the growing interest in the nonthermal technologies, and assist in the successful commercialization of UV light for food processing applications.
REFERENCES


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