Press and Press communication

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Electron beams for rapid, safe vaccine production
Pathogens in the firing line

As part of a multi-disciplinary project, researchers from four Fraunhofer Institutes have developed a technology to quickly and safely destroy pathogens used in vaccine manufacture.

These days, it is not just goods and people which travel around the world within a few hours. In the age of globalization, pathogens can also spread extremely rapidly. New vaccines must be manufactured quickly to prevent epidemics.

A new procedure, developed by an interdisciplinary team from the Fraunhofer Society, is helping to quickly and completely destroy the pathogens used in the manufacture of vaccines, without damaging the external form of the viruses, bacteria or parasites. The inactivated vaccine that is created is not a threat to the body. Instead, it causes the immune system to produce antibodies which protect the body against an outbreak of the disease.

As IPA team leader Martin Thoma explains, until now, pathogens in vaccines have mainly been inactivated using chemicals. However, this method has disadvantages: “Chemical treatment is time-consuming and cannot be integrated into a continuous production process. Moreover, chemical residues remain in the vaccine, which then get into the human body”.

For five years, the IPA team, along with groups from three further Fraunhofer Institutes, have been experimenting with new “procedures for the inactivation of pathogens using low-energy electron beams”. The research received USD 1.8 million in funding from the Bill & Melinda Gates Foundation.

“The interdisciplinary cooperation was key to its success”, underlines Thoma. The experts from the Fraunhofer Institute for Organic Electronics, Electron Beam and Plasma Technology FEP developed the system which precisely measures out the low energy electrons so that they destroy the pathogens’ genetic material in very small quantities of liquid. Researchers at the Fraunhofer Institute for Cell Therapy and Immunology IZI cultivated various pathogens, including bird flu and equine influenza pathogens, and verified whether they were completely inactivated following radiation and whether it
makes for an effective vaccination. The team at the Fraunhofer Institute for Interfacial Engineering and Biotechnology IGB investigated the dose-dependent inactivation. Finally, Thoma’s team at IPA developed the modules which ensure that the liquid containing the pathogens is filtered through to the electron beam at precisely the right layer height and speed.

“The automation of this process was a particular challenge because a sufficiently homogeneous dose distribution in liquids during radiation with low energy electron beams (ca. 200keV) can only be guaranteed up to a penetration depth of ca. 200 micrometers. Because of this, you have to guide the probes through the rays at a very controlled layer height and speed”, reports Thoma. A module was specially developed for small probe sizes which carried out the pathogen suspension in pouches below the electron source. Another module, the “Liquid roll”, was developed to inactivate pathogens in large liquid volumes: Here, a roll is dampened with the pathogen suspension and then wiped after radiation. This method can be integrated into industrial vaccine production. An initial research and test facility has been recently put into operation at Fraunhofer IZI in Leipzig, which now enables the teams to continue to optimize the technology and, together with their industry partners, test the production of various vaccines.

Profile

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<tr>
<th>Project name</th>
<th>MAVO-Project ELVIRA</th>
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<tr>
<td>Title</td>
<td>The development of a method for the electron beam-based inactivation of viruses and bacteria for vaccine manufacture</td>
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<td>01 March 2014 – 31 March 2018</td>
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<td>Project partners</td>
<td>Fraunhofer Institute for Interfacial Engineering and Biotechnology IGB, Fraunhofer Institute for Organic Electronic, Electron Beam and Plasma Technology FEP, Fraunhofer Institute for Manufacturing Engineering and Automation IPA, Fraunhofer Institute for Cell Therapy and Immunology IZI</td>
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<td>Coordination</td>
<td>Dr. Sebastian Ulbert, Fraunhofer Institute for Cell Therapy and Immunology IZI</td>
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Module for continuous inactivation of pathogen suspension.

The module shown above is mounted below an electron beam source and ensures a continuous and extremely defined fluid flow of infectious pathogen suspension through the electron beam. With this prototype, a volume of approx. 3.5L/h can already be inactivated today, with 15L corresponding to approx. 1 million vaccine doses. The process is easily scalable to significantly higher throughputs.
(Source: Fraunhofer IPA)