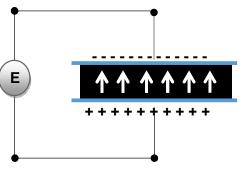


Piezotech Processing's guides

- How to pole -

In order to activate a copolymer material, a poling step is necessary. Using an increasing low-frequency voltage, an electric field above the coercive field value (*i.e.* 50V/ μ m) has to be applied. Depending on the film thickness, its surface, and the response precision needed, different methods can be used.



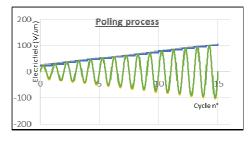
Orientation of Crystalline Phases

The application of the electric field and the hysteresis curve characterization can be done using one of the following:

- A direct application of the electric field through the electrodes with a voltage generator (in this case there is no measurement of the induced electrical polarization (charge displacement))
- A Sawyer-Tower circuit (electrical charges measurement, hysteresis curve)
- A ferroelectric tester (*i.e., Precision MultiFerroelectric Radiant Technology*) with an external amplifier if a high electric field is needed (depending on the film thickness)

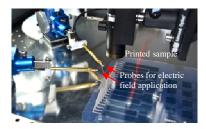
When a voltage generator is coupled with a signal generator, the electric field can be applied according to the following parameters:

- Number of cycles to reach E_{max}: 15
- Frequency: 0,5Hz (higher is possible)
- Signal: sinusoidal
- E_{max} > 2E_c
- Typically E_{max}=100V/μm

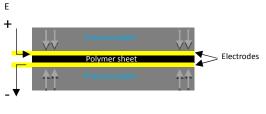


Typical poling process used for copolymer polarization

For printed devices (mainly thin layers), the poling electric field can be applied directly through the film electrodes by using a direct probes contact system. For non-printed films (thick layers), poling can be done through contacting and pressing the film between two electrodes. For large surfaces, a Corona poling can be used.



Direct probes poling process



Contact poling process



Safety and Storage

Please refer to the safety datasheet

Contact Information

Piezotech Arkema-CRRA Rue Henri Moissan 69496 Pierre-Benite Cedex Info.piezotech@arkema.com Tel : +33 4 72 39 97 03 www.piezotech.eu

Disclaimer

The statements, technical information and recommendations contained herein are believed to be accurate as of the date hereof. Since the conditions and methods of use of the product and of the information referred to herein are beyond our control, ARKEMA expressly disclaims any and all liability as to any results obtained or arising from any use of the product on reliance on such information; NO WARRANTY OF FITNESS FOR ANY PARTICULAR PURPOSE, WARRANTY OF MERCHANTABILITY OR ANY OTHER WARRANTY, EXPRESSED OR IMPLIED, IS MADE CONCERNING THE GOODS DESCRIBED OR THE INFORMATION PROVIDED HEREIN. The information provided herein relates only to the specific product designated and may not be applicable when such product is used in combination with other materials or in any process. The user should thoroughly test any application before to take appropriate steps to be sure that any proposed use of the product will not result in patent infringement. See SDS for Health & Safety Considerations. Arkema has implemented a Medical Policy regarding the use of Arkema products in Medical Devices applications that are in contact with the body or circulating bodily fluids (http://www.arkema.com/en/social-responsibility/responsible-product-management/medical-device-policy/index.html) Arkema has designated Medical paces to be suce of rough Medical

(http://www.arkema.com/en/social-responsibility/responsible-product-management/medical-device-policy/index.html) Arkema has designated Medical grades to be used for such Medical Device applications. Products that have not been designated as Medical grades are not authorized by Arkema for use in Medical Device applications that are in contact with body or circulating bodily fluids. In addition, Arkema strictly prohibits the use of any Arkema products in Medical Device applications that are in contact with bodily fluids or tissues for greater than 30 days. The Arkema the Arkema and the Arkema anales and the Arkema anales and the Arkema anales or temporary implantable devices, and customers shall not represent to anyone else, that Arkema allows, endorses or permits the use of Arkema products in such medical devices.

on temporary implantable devices, and distolers shall not represent to anyone else, that Arkenta anows, endotses of permits the use of Arkenta products in solution medical devices. It is the sole responsibility of the manufacturer of the medical device to determine the suitability (including biocompatibility) of all raw materials, products and components, including any medical grade Arkenta products, in order to ensure that the final end-use product is safe for its end use; performs or functions as intended; and complies with all applicable legal and regulatory requirements (FDA or other national drug agencies). It is the sole responsibility of the manufacturer of the medical device to conduct all necessary tests and inspections and to evaluate the medical device under actual end-use requirements and to adequately advise and warn purchasers, users, and/or learned intermediaries (such as physicians) of pertinent risks and fulfill any postmarket surveillance obligations. Any decision regarding the appropriateness of a particular Arkema material in a particular medical device should be based on the judgment of the manufacturer, seller, the competent authority, and the treating physician. ¹

Piezotech SAS. Registered in the R.C.S of Lyon (France) under the number 391 321 122. Arkema-CRRA Rue Henri Moissan 69493 Pierre-Benite Cedex / France