Webinar - Medical Device Clinical Investigations in Türkiye

As it is known, Medical Device Regulation (MDR), which came into force on 02 June 2021 (26 May 2021), brings more stringent regulations along with some new requirements in the context of clinical evaluation and clinical investigation.

While clinical investigations have been made mandatory for some device classes and types in order to obtain sufficient clinical data when placing a device on the market, post-marketing studies have come to the fore for all devices within the scope of post-market surveillance obligation.

In addition, the main objective for the "Pharmaceuticals and Medical Devices" sector, which is among the priority sectors in our Eleventh Development Plan, has been determined as increasing our competitiveness in the global market and carrying our country to a higher position in the value chain. One of our main policies is "To ensure that our country becomes the leading country in the region in clinical research." In addition to increasing the share we receive from global clinical research, both in number and financially, it is also aimed to increase clinical research to be conducted with innovative products to be developed in our country.

In this respect; in line with the transition period to MDR and our strategic goals, an international webinar will be held in cooperation with ARTED on 28 September 2022 Wednesday at 16:00 (UCT+3, TR time) in order to introduce the clinical investigation ecosystem in Türkiye and to provide information about the current legislation and infrastructure opportunities.

Necessary information about registration and program content is given below.
Announced to all interested parties.

Agenda

	28 September 2022 Wednesday 16:00 (UCT+3, TR time)
16:00 – 16:10	Opening Speeches
16:10 – 16:30	**Clinical investigation regulations in Türkiye** *Mol. Bio. Gökhan ÖZKAN* *TİTCK*
16:30 – 16:50	**Clinical investigation infrastructure of the City Hospitals in Türkiye** *Prof. Dr. Fuat Emre CANPOLAT* *Ministry of Health Ankara City Hospital*
16:50 – 17:10	**The clinical investigation environment in Türkiye from a Manufacturer/Sponsor perspective** *Elif ÖZMAN PUSAT* *ARTED Secretary General*
17:10 – 17:45	Question&Answer

Registration

Interested participants who want to attend the meeting must register on the http://www.canlikonsey.tv/klinikarastirma website. The participation links related to the meeting together with the confirmation information will be sent to the e-mail address or phone number that is provided during registration.