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```
IPipelineMode { internal
    readonly
    IAsyncPipelineImpl
    pipeline; internal AsyncPi
    pipelineMode(IPipelineImpl
    pipeline, string name) {
    this.pipeline = pipeline;
this.name = name; } /// ///
Flush the current pipeline.
    /// public void Flush() {
        var result =
            pipeline.Flush();
OnFlush(); } /// /// Execute
the pipeline. /// ///
```

Command to be executed. /// Cancellation token. 50b96ab0b6

EAXA Carenado-195P base model Limulus amoebocyte lysate test to evaluate the appropriateness of dose escalation in radiotherapy of benign breast disease. The value of the limulus amoebocyte lysate test (LAL test) has been studied in a series of breast cancer patients treated with moderate hypofractionation (45 Gy in 30 fractions) in order to avoid the increased risk of skin reactions after large fraction sizes. One group of patients received the LAL test before and after radiotherapy (RT), another group before RT only. The value of the LAL test as an indicator of acute radiation damage was examined by measuring the increase in clotting activity of the patient's plasma and the analysis of its components. The LAL test can be used to evaluate the degree of acute damage to tissues in vivo due to an underlying disease or environmental factor (in this case hypofractionated RT), since it indicates the amount of circulating fibrinogen with its fibrin degradation products (FDPs) in a patient's plasma. Our results do not show any increase in FDPs after hypofractionated RT in either group. At the same time, an increase in the clotting activity of the patient's plasma is detected after RT. Also, the LAL test is more sensitive than the RTOG score in detecting early adverse effects of RT due to a higher sensitivity of LAL compared to the RTOG score. This indicates that the LAL test is more suitable than the RTOG score for the detection of early effects of RT, and probably also for the development of normal tissue tolerance. This is important for the development of radiotherapy planning aimed at sparing critical organs at risk at the expense of increased morbidity in the breast. Approximately 1.5 million individuals are affected by schistosomiasis in the U.S. A more effective vaccine would be an important tool for disease control. We have shown that an experimental vaccine can protect mice from liver and intestinal damage induced by schistosomes, and that this protection correlates with the induction of highly protective anti-schistosome Th1-type cellular immune responses. In this proposal we plan to develop a new mouse-based schistosome vaccine using recombinant *S. mansoni* antigens expressed in a vaccine delivery system that provides the necessary adjuvant activity to induce protective immune responses. We will also evaluate humoral responses by developing assays that correlate with protection in mice. The

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