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GSK provides update on regulatory filings for Zunrisa/Rezonic

GlaxoSmithKline (GSK) has made the difficult decision to discontinue the regulatory filings for Zunrisa/Rezonic (casopitant), which was being developed for chemotherapy-induced and post-operative nausea and vomiting. Regulatory applications were filed in a number of countries worldwide and GSK is currently communicating with those authorities and all study investigators.

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GSK has made this decision, after regulatory consultation, based on the company's assessment that significant further safety data would be required to support the registration of casopitant on a worldwide basis, which would take a considerable time to produce. Consequently, all on-going regulatory files for casopitant are being withdrawn.

Azurity Pharmaceuticals, Inc. EXHIBIT - 1036

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