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The SBLA SUBMISSION WITH FDA follows in the back of recently announced positive TMB-302 ResultsMontreal, Dec. 06, 2021 (Globe Newswire) - Theratechnologies Inc. (Theratechnologies or Company) (TSX: TH) (NASDAQ: THTX), a biopharmaceutical Company focused on the development and marketing of innovative therapies is pleased to announce the application of a supplementary biology license (SBLA) application to the U.S. Food and Drug Administration (FDA) for the company's intravenous administration (iv) form. TROGARZOÂ®. The IV Push TrogarzoÂ® continues the company's commitment to improving the lives and outcomes of treatment for patients living with HIV. The FDA's submission © the next step in the advanced push program TROGARZOÂ® IV TROGARZOÂ®. As recently© announced, the TMB-302 study demonstrated that there was no difference in pharmacokinetics (PK) between the IV Push and IV infusion and was conducted by the company's partner, Timed Biologics (Taimed). TROGARZOÂ® IV push, a more convenient form of administration, can be infused within 30 seconds without dilution in commendable with the infusion time of 15 minutes of the original IV infusion. Theratecology believes that this mode of administration will represent a marked improvement for patients.Aditionalmente, TMB-302 also demonstrated that there were no serious adverse events observed and drug©-related adverse events were considered mild to moderate. The secondary endpoints were also allocated confirming no differences in HIV©-1 viral load due to the change of the IV infusion to push IV. Also there were no anti©trogarzointibodies© or immunogenicity concerns of TROGARZOÂ® detected. About TROGARZOÂ® (IBalizumab-Uiyk) InjectiontrogarzoÂ® A© an HIV-1 inhibitor directed by CD4. TROGARZOÂ® is © for commercialization in the United States and the European Union. In the United States, trogarzoÂ®, in combination with other antiretrovirals, is © for the Get Type 1 virus infection (HIV-1) in adults highly tested with multi-drug disease (MDR) infection by HIV-1 failing their current anti retroviral regimen. In Europe, Trogarzoe © is approved for the treatment of adults infected with HIV-1 MDR for which no © It is possible to build a suppressive antiviral regime. Important safety information Before receiving the Thunderstorm ©, inform your health care provider if you are pregnant or plan to become pregnant, as it is not known whether Trogarzoe © may harm your foetus or if you are breast-feeding or plan to breast-feed, as it is not known whether Trogarzoe © passes into breast milk. Your immune system can get stronger and start fighting infections that have been hidden in your body for a long time. Tell your health care provider immediately if you start having new symptoms after starting your HIV-1 medicine. The most common side effects of Trogarzos © include: diarrhoea, dizziness, consumption and cutaneous eruptions. These are not all possible side effects of Trojan ©. For more information, ask your health care provider or pharmacist. Full information on prescription available at www.trogarzo. With About Theratechnologies Theratechnologies (TSX: TH) (NASDAQ: THTX) © a biopharmaceutical company focused on the development and marketing of innovative therapies that respond to the human needs © Not satisfied tips. Further information on Theratechnologies is available on the Enterprise website at www.theratech. with, in SEDAR at www.sedar. with and in EDGAR at www.sec. govForward-Looking InformationThis press release contains © with forward-looking statements and information, or forward-looking information, Prospective Declarations, in the sense of the applicable title laws, which are based on our beliefs and management assumptions and information currently available for our management. You can identify Prospective Declarations in terms of "can", "will", "will", "will", "should", "could", and wouldldds. "perspective", "believe," "plan," "predict," "anticipate," "wait" and "esteem," or the negatives of these terms, or variations thereof. The Prospective Declarations contained in this press release include, but are not limited to, statements regarding the expected improvement of the way in which management IV push es patients. Although the Prospective Declarations contained in this press release are based on what the Company considers to be reasonable assumptions in the light of the information currently available, investors are warned not to unduly trust these statements, since the real results may vary from the Prospective Declarations. Some assumptions made in the preparation of the Prospective Declarations include that: the FDA will approve the BLA, the patients and the hands © dices will accept form IV Trojan admin push ©, the current pandemic will not negatively affect patient access to their mother © dices and their treatments and Trogarzoan © The assumptions of the Prospective Declarations are subject to one alone. © range of risks and uncertainties, many of which are beyond © the control of Theratechnologies which could make the real results materially different from those which are disclosed or implied in the said Prospective Declarations. These risks and uncertainties include, among others, those related to or resulting from the adverse impact of the ongoing COVID-19 pandemic on patient access to mother © dices and clans, not approved by the FDA of BLA, the withdrawal of Trojan © and the acceptance by patients and mother © dices of this new way of administration section of our Annual Information Form dated February 24, 2021 available at SEDAR in www.sedar.com and EDGAR in www.sec.gov as an exhibition to our 40-F report dated February 25, 2021 under Theratechnologies' public records for additional risks related to the Company. The reader is © warned to consider these and other risks and uncertainties carefully and not put undue trust in forward-looking declarations. Forward-looking statements reflect current expectations in terms of future events and speak only from the date of this press release and represent our expectations from that date. We do not undertake any obligation to update or revise the information contained in this press release, whether as a result of new information, future events or circumstances or otherwise, except as required by applicable law. For investors© and media: Philippe DubucSenior Vice President and Head of Communications Financeiras@theratech.com514-336-7800 Telecommunications es@theratech.com514-336-7800

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