

## Catalog # 10-5047 Fludarabine

CAS# 21679-14-1

9-β-D-Arabinofuranosyl-2-fluoro-9H-purin-6-amine; F-ara-A; NSC118218 Lot # X109662

Fludarabine is a synthetic adenosine analog that inhibits DNA biosynthesis and is a clinically useful antineoplastic agent. In cells fludarabine accumulates as its 5'-triphosphate (F-ara-ATP) for which the rate-limiting step in formation is the conversion of fludarabine to its monophosphate. F-ara-ATP has multiple mechanisms of action including inhibition of ribonucleotide reductase, DNA polymerase, ligase and primase. A frequently used agent in myeloablative conditioning regimens for allogeneic hematopoietic cell transplantation. Immunosuppressive effects are mediated via inhibition of TNF $\alpha$ -stimulated production of IL-2 and IFN- $\gamma$  through inactivation of NF $\kappa$ B. Antagonist at adenosine A1 receptors.

- 1) Ross et al. (1993), Fludarabine. A review of its pharmacological properties and therapeutic potential in malignancy; Drugs, 45 737
- 2) Gandhi and Plunkett (2002), Cellular and clinical pharmacology of fludarabine; Pharmacokinet., 41 93
- 3) Langenhorst et al. (2019), Fludarabine exposure in the conditioning prior to allogeneic hematopoietic cell transplantation predicts outcomes; Blood Adv., 3 2179
- 4) Nishioka et al. (2008), Fludarabine induces growth arrest and apoptosis of cytokine- or alloantigen-stimulated peripheral blood mononuclear cells and decreases production of Th1 cytokines via inhibition of nuclear factor kappaB; Bone Marrow Transplant., 41 303
- Jensen et al. (2012), Cytotoxic purine nucleoside analogues bind to A1, A2A and A3 adenosine receptors; Naunyn Schmiedebergs Arch. Pharmacol., 385 519

## **PHYSICAL DATA**

Molecular Weight: 285.23

NMR: (Conforms)

Solubility: DMSO (up to 30 mg/ml)

Physical Description: White solid

Storage and Stability: Store as supplied desiccated at -20°C for up to 1 year from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 3 months.

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