

## **Quercetin Summary for Intravenous use**

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### **INTRAVENOUS QUERCETIN:**

Intravenous Quercetin has studied potential for increased bioavailability [1,2,4] as well as potent potential anti-tumor activity [4-5]. Intravenous data in human subjects shows it to be tolerated and safe [1-3]. Data available suggest multiple mechanisms of action in Tyrosine Kinase inhibition [4] as well as tumor growth suppression [5]. Two years of clinical use has revealed no adverse events when used under standard dose and administration guidelines [3].

### **INTRAVENOUS USE GUIDELINES:**

#### **Dose: [1-4]**

- Test dose at 1 mg/kg IV on the first day
- Subsequent doses could increase to 140 mg/kg if tolerated two times weekly
- Generally tolerated doses are between 500 and 1000 mg. 2500 mg have been given IV.

#### **Administration:**

- Intravenous dosing via either a central or peripheral line.
  - Use a filtered line or add on filter set
- Carrier solutions:
  - Per compounding pharmacy instructions, usually NS or D5W
  - Dilute not more concentrated than 100 mg per 100 mL solution
- Rate of administration: Start at a rate of 30 minutes per 100 mg and ramp up to tolerance. Tolerance varies greatly.
  - Monitor for signs of nausea which can be the first sign of a non-tolerated dose [3]
  - For allergic / anaphylactic reaction treat per standard protocol.
- Other IV compatibility:
  - Generally incompatible with other IV solutions in the same IV container

#### **Screening:**

- Intolerance to oral Quercetin is a caution and may exclude use in the IV setting

- Lab studies:
  - CBC, Chemistry panel (Metabolic panel including electrolytes, bilirubin, AST/ALT/GGT, eGFR/BUN/CRE).

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**References:**

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2. Graefe EU, Derendorf H, Veit M. Pharmacokinetics and bioavailability of the flavonol quercetin in humans. *Int J Clin Pharmacol Ther.* 1999 May;37(5):219-33. PMID: 10363620
3. Anderson P, Cochran B. Personal experiences with the clinical use of intravenous therapies. AMSA, BIORC and Private clinic data. Seattle Washington, 2014
4. Ferry D R, Smith A, Malkhandi J, et al. Phase I clinical trial of the flavonoid quercetin: pharmacokinetics and evidence for in vivo tyrosine kinase inhibition. *Clin Cancer Res* 1996;2:659-668.
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