

Lipid-Free, Stabilized Emulsions for Delivering Anesthesia and Other Hydrophobic Drugs

View U.S. Patent No. 10,758,483 in PDF format.

WARF: P140337US02

Inventors: Sandro Mecozzi, Robert Pearce, William Tucker

The Wisconsin Alumni Research Foundation (WARF) is seeking commercial partners interested in developing a method for formulating non-lipid nanoemulsions that are especially stable and less susceptible to bacterial/fungal contamination.

Overview

Emulsions containing soybean oil have been used intravenously for more than 40 years. Recently they have been employed in the delivery of certain hydrophobic drugs such as anti-inflammatory agents and anesthetics like propofol (i.e., Diprivan).

Propofol is used extensively to induce and maintain general anesthesia as well as for sedation purposes. However, the oil used in the emulsion is highly susceptible to bacterial and fungal growth. To combat contamination, tubing and open vials of propofol must be replaced every 12 hours. The current formulation also may cause pain, lipid intolerance and other serious complications in some patients.

The Invention

UW-Madison researchers have developed non-lipid nanoemulsions for delivering propofol and other hydrophobic compounds. The formulations contain miniscule droplets of semifluorinated block copolymers and perhalogenated fluorous compounds, such as perfluorooctyl bromide or perfluorodecalin.

These ingredients are capable of forming a stable nanoemulsion without the need for conventional lipid components (e.g., soybean oil) that support bacterial and/or fungal growth. The emulsions have enhanced stability with respect to droplet size due to decreased particle coarsening, coagulation and/or phase separation.

Applications

Nanoemulsions for formulating, delivering and releasing hydrophobic drugs including propofol

Key Benefits

- · Lipid-free and very stable
- · Solves the problem of bacterial/fungal contamination
- Highly versatile
- · Supports controlled drug release
- Nanoemulsions are intrinsically more stable and absorbable in vivo.

We use cookies on this site to enhance your experience and improve our marketing efforts. By continuing to browse without changing your browser settings to block or delete cookies, you agree to the storing of cookies and related technologies on your device. See our privacy policy

The new emulsions have been tested in animals.



Additional Information

For More Information About the Inventors

- Sandro Mecozzi
- Robert Pearce

Tech Fields

Drug Delivery : Other drug delivery technologies

For current licensing status, please contact Rafael Diaz at rdiaz@warf.org or 608-960-9847

We use cookies on this site to enhance your experience and improve our marketing efforts. By continuing to browse without changing your browser settings to block or delete cookies, you agree to the storing of cookies and related technologies on your device. See our privacy policy

