

Latex Content in Nalge Products

Allergic reactions to latex, or natural rubbers, is a health risk. NNI takes such a risk seriously, and has taken measures to make sure that our products do not contain these compounds. The FDA requires labeling of medical devices that contain latex or natural rubber, 21CFR section 801.437.

In compliance with the United States Food & Drug Administration (FDA) regulations published in the Code of Federal Regulations (CFR), Title 21, § 801.437 "User labeling for devices that contain natural rubber", this is to document that NALGENE® brand products and packaging do not contain any:

1. Natural rubber latex, or
2. Dry natural rubber, or
3. Synthetic latex or synthetic rubber that contains natural rubber in its formulation.

The only exception is the Filter Stopper, catalog number DS0396-0080 (Filter stopper, No. 8, rubber). This product is also a component supplied with the Filter Funnel with Clamp, catalog number 315-0047, and the Microbiological Monitors, catalog numbers 136-4547, 136-4556, 137-4547 and 137-4556. It should be noted that these products are not listed as medical devices.