MEDISCA®
YOUR TRUSTED PARTNER IN COMPOUNDING

Date: 01/15/2021

ALLERGEN DECLARATION

PRODUCT NAME: PLURONIC GEL 30% (F127)

1. The product is FREE FROM the following:

- ☒ Cereals containing gluten and products thereof
- ☒ Corn and products thereof
- ☒ Soybeans and products thereof
- ☒ Celery and products thereof
- ☒ Mustard and products thereof
- ☒ Yeast
- ☒ Egg and products thereof
- ☒ Dairy Products (Milk, Lactose, Caseinates, Whey)
- ☒ Peanut and products thereof
- ☒ Tree nuts and products thereof
- ☒ Sesame Seeds and products thereof
- ☒ Sunflower Oil
- ☒ Sulfites >10ppm
- ☒ Artificial Colours
- ☒ Artificial Flavours
- ☒ Dyes
- ☒ Genetically Modified organisms (GMO)
- ☒ Beef/Chicken/Pork derivatives
- ☒ Fish/Shellfish derivatives
- ☐ Preservatives
- ☒ Parabens
- ☒ Latex

2. The starting materials were FREE FROM the following:

- ☒ Cereals containing gluten and products thereof
- ☒ Corn and products thereof
- ☒ Soybeans and products thereof
- ☒ Celery and products thereof
- ☒ Mustard and products thereof
- ☒ Yeast
- ☒ Egg and products thereof
- ☒ Dairy Products (Milk, Lactose, Caseinates, Whey)
- ☒ Peanut and products thereof
- ☒ Tree nuts and products thereof
- ☒ Sesame Seeds and products thereof
- ☒ Sunflower Oil
- ☒ Sulfites >10ppm
- ☒ Artificial Colours
- ☒ Artificial Flavours
- ☒ Dyes
- ☒ Genetically Modified organisms (GMO)
- ☒ Beef/Chicken/Pork derivatives
- ☒ Fish/Shellfish derivatives
- ☐ Preservatives
- ☒ Parabens
- ☒ Latex

Please be advised that Medisca has not tested for the presence of the above stated allergens. Therefore, this statement is provided for informational purposes only, as instructed to Medisca by the manufacturer, and is not meant to be a guarantee of absence of the above stated allergens.

Please note that there may be several steps before the raw material is produced and access to data on manufacturing raw materials is not easily obtained for the starting materials and intermediates used in the production process.

No raw material sourced or derived from animals are used in the manufacturing. Hence, the materials used in the production of have demonstrated compliance with the note for the guidance on minimizing the risk of transmitting animal spongiform encephalopathy agents via human and veterinary medicinal products.

It is the responsibility of the pharmacist to verify state and federal regulations prior to repackaging and/or dispensing the product.

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Signature: