

## PRODUCT DATASHEET IGNITOL WMO 8

## Meets the requirements of the following specifications:

- US FDA 21 CFR 172.878 & 21 CFR 178.3620(a), White Mineral Oil.
- USP 40/ NF35 (US Pharmacopoeia/ National Formulary), Light Mineral Oil.
- Light Liquid Paraffin British Pharmacopoeia and European Pharmacopoeia.

S.	Characteristic	Unit	Test Method	Typical D Min.	
No.	\/;1		\		Max.
1.	Visual	-	Visual	A colourless, transparent, oily	
				liquid free from fluorescence in day light. Practically insoluble	
				in water, slightly soluble in	
				ethanol (96%), miscible with	
				hydrocarbons.	
2.	Colour, Saybolt	-	ASTM D 156	+ 30	
3.	Odour	-	Olfactory	Almost odourless	
4.	Kinematic Viscosity at 40 <sup>O</sup> C	Cst	ASTM D 445/D	7.0	9.0
	5 1 11 5 11 100 ()00		7042	0.040	2.22
5.	Relative Density at 20 <sup>O</sup> C	-	BP/EP	0.810	0.865
6.	Specific Gravity at 25/25 <sup>O</sup> C	-	NF/ USP	0.815	0.870
7.	Flash Point	ОС	ASTM D 92	150	200
8.	Pour Point	OC	ASTM D 97		-28
9.	Acidity	-	USP	Not more than 1 ml of 0.01N NaOH	
10.	Limit of Polycyclic Aromatic Hydrocarbons	-	NF/USP/BP/EuP/IP	Pass	
11.	Readily Carbonizable Subsances	-	NF/USP/BP/EuP/IP	Pass	
12.	Solid Paraffin	-	NF/USP/BP/EuP/IP	Pass	
13.	Sulphur Compounds	-	NF/USP/IP	Pass	

**IGNITOL WMO 8** – Light Mineral Oil – NF/USP/BP/EuPh is a highly pure grade of White Mineral Oil specially formulated from chosen severely hydro treated and highly refined Paraffinic Oils, thus qualifying for the severe requirements stipulated under NF/ United States Pharmacopoeia