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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
I. DR	UGS & PHARMAC	EUTICALS		
1.	Raw Materials & Bulk Drugs (General Tests)	Description	IP 2014, Ph. Eur.8.0, 2014 USP 37, BP 2009	Qualitative
	(General Tests)	Solubility	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	Qualitative
		Identification by UV	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	Qualitative
		Melting Range	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	30°C to 300°C
		pН	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	1 to 12
		Specific Optical Rotation	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	-200° to 200°
		Refractive Index	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	0.25 to 2.00
		Sulphated Ash/ Residue on ignition	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	0.05% to 30%
		Viscosity at 25°C	IP 2014, Ph. Eur.8.0, 2014,	1.0 cps to 100000 cps

USP 37, BP 2014

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Assay by Potentiometry	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	80% to 102%
		Heavy Metals	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	Qualitative
		Limit tests for chlorides	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	Qualitative
		Water (By KF)	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	0.5% to 40%
		Loss on Drying	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	0.1% to 20%
		Particle Size Analysis	IP 2014	0.02 to 2000 microns
2.	Sodium Methyl Paraben	Description	IP 2014	Qualitative
		Identification A (By IR)	IP 2014	Qualitative
		Appearance of Solution	IP 2014	Qualitative
		Related Substances by HPLC	IP 2014	Qualitative
3.	Sodium propyl Paraben	Description	IP 2014	Qualitative
		Identification A (By IR)	IP 2014	Qualitative
		Appearance of Solution	IP 2014	Qualitative
		Related Substances(By HPLC)	IP 2014	Qualitative

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4.	Disodium EDTA	Description	IP 2014	Qualitative
		Identification A (By IR) Identification B & C (By Chemical)	IP 2014	Qualitative
		Impurity A by HPLC	IP 2014	Qualitative
		Assay by Titrimetry	IP 2014	98.5% to 101.0%
5.	Sorbitol	Description	IP 2014	Qualitative
		Identification A (By HPLC) Identification B & C (By Chemical)	IP 2014	Qualitative
		Appearance of Solution	IP 2014	Qualitative
		Acidity or Alkalinity	IP 2014	Qualitative
		Related substances by HPLC	IP 2014	Qualitative
		Reducing sugars by Titrimetry	IP 2014	0.1% to 0.4%
		Assay by HPLC	IP 2014	98% to 101%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
6.	Mannitol	Description	IP 2014	Qualitative
**		Identification A(By IR), Identification B & C (By Chemical)	IP 2014	Qualitative
		Appearance of Solution	IP 2014	Qualitative
		Acidity or Alkalinity	IP 2014	Qualitative
		Reducing sugars by Titrimetry	IP 2014	0.1% to 0.4%
		Sorbitol by TLC	IP 2014	Qualitative
		Assay by HPLC	IP 2014	98.0% to 102.0%
7.	Paracetamol	Description	IP 2014	Qualitative
		Identification A by IR, Identification C by Chemical Identification A by IR	IP 2014	Qualitative
		Related substances by HPLC (Total impurities)	IP 2014	Qualitative
		Assay by Titrimetry	IP 2014	99.0% to 101.0%

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Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
Ibuprofen	Description	IP 2014	Qualitative
	Identification A by IR, Identification C by TLC	IP 2014	Qualitative
	Appearance of Solution	IP 2014	Qualitative
	Related substances by HPLC	IP 2014	Qualitative
	Assay by titrimetry	IP 2014	98.5% to 101.0%
Niacinamide	Identification A by IR	USP-37	Qualitative
	Readily carbonizable substances	USP-37	Qualitative
	Assay by HPLC	USP-37	98.5% to 101.5%
Pyridoxine HCl	Description	IP 2014	Qualitative
	Identification A by IR Identification C by TLC Identification D by chemical	IP 2014	Qualitative
	Appearance of solution	IP 2014	Qualitative
	Assay by Titrimetry	IP 2014	99.0% to 101.0%
	Material of Test Ibuprofen Niacinamide	Ibuprofen Description Identification A by IR, Identification C by TLC Appearance of Solution Related substances by HPLC Assay by titrimetry Niacinamide Identification A by IR Readily carbonizable substances Assay by HPLC Pyridoxine HCI Description Identification A by IR Identification A by IR Identification C by TLC Identification D by chemical Appearance of solution	Material of Test Description IP 2014 Identification A by IR, IP 2014 Appearance of Solution IP 2014 Related substances by HPLC IP 2014 Assay by titrimetry IP 2014 Niacinamide Identification A by IR USP-37 Readily carbonizable substances USP-37 Assay by HPLC USP-37 Pyridoxine HCl Description IP 2014 Identification A by IR IP 2014 Identification A by IR IP 2014 Identification C by TLC Identification D by chemical Appearance of solution IP 2014 IP

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
11.	Cyanocobalamin	Description	IP 2014	Qualitative
		Identification B by TLC, Identification C by chemical	IP 2014	Qualitative
		Related substances by HPLC	IP 2014	Qualitative
		Assay by UV	IP 2014	96.0% to 102.0%
12.	Nebivolol Hcl	Palladium(Pd) by ICP-OES	ICP-MVP-004/2011	0.1 ppm to 100 ppm
13.	Montelukast Sodium	Palladium(Pd) by ICP-OES	ICP-MVP-005/2011	0.1 ppm to 100 ppm
14.	Fosaprepitant Dimeglumine	Palladium(Pd) by ICP-OES	ICP-MVP-008/2011	0.3 ppm to 100 ppm
15.	Voriconazole	Lithium(Li) by ICP-OES Nickel(Ni) by ICP-OES Palladium(Pd) by ICP-OES	ICP-MVP-029/2012	0.1 ppm to 200 ppm 0.15 ppm to 100 ppm 0.12 ppm to 100 ppm
16.	Capsules General Tests	Description	IP 2014, Ph. Eur.8.0, 2014 USP 37, BP 2014	Qualitative
		Tablet Disintegration	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	1 min to 60 min

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6.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
17.	Omeprazole Capsules	Identification by HPLC	IP 2014	Qualitative
	Capsures	Dissolution	IP 2014	Qualitative
		Uniformity of content by HPLC	IP 2014	Qualitative
		Assay by HPLC	IP 2014	80% to 120.0%
18.	Tablets General Tests	Description	IP 2014, Ph. Eur.8.0, 2014 USP 37, BP 2014	Qualitative
		Tablet Disintegration	IP 2014, Ph. Eur.8.0, 2014, USP 37, BP 2014	1 min to 60 min
19.	Folic acid tablets	Identification A by TLC, Identification B by Chemical	IP 2014	Qualitative
		Hydrolysis products by HPLC	IP 2014	Qualitative
		Uniformity of content by HPLC	IP 2014	40% to 140%
		Assay by HPLC	IP 2014	90.0% to 115.0%
20.	Liquid Orals General Tests	Description	IP 2014, Ph. Eur.8.0, 2014 USP 37, BP 2014	Qualitative
		Viscosity	IP 2014	1 cps to 100000 cps

Laboratory	DKR Labs, Plot No. 121, IDA, Prashanthi Nagar, Kukatpally, Hyderabad,
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
21.	Paracetamol Syrup	Identification A by TLC, Identification B by HPLC	IP 2014	Qualitative
		4-Aminophenol by HPLC	IP 2014	Qualitative
		Assay by HPLC	IP 2014	90.0% to 110.0%