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SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are	Range of Testing / Limits of Detection
			performed	

BIOLOGICAL TESTING

Ι.	DURGS AND PHAR	RMACETICALS		
1.	Raw Material for P	harmaceutical Preparation	n	
a.	Active Pharmaceuticals Ingredients	Total Viable Aerobic count	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	1 cfu per g or mL to 10⁵cfu per g or mL
		Total Yeast and Mold count	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	1 cfu per g or mL to 10⁵cfu per g or mL
		Staphylococcus aureus	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	Present or Absent per mL or g
		E coli	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	Present or Absent per mL or g
		Pseudomonas aeruginosas	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	Present or Absent per mL or g
		Salmonella	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	Present or Absent per 10 mL or g
2.	Finished Pharmace	eutical Preparation		
а.	Tablet, Capsule, Syrup	Total Viable Aerobic count	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	1 cfu per g or mL to 10⁵cfu per g or mL
		Total Yeast and Mold count	IP 2014 (Vol. I), 2.2.9, USP 39 <61>, <62>, BP 2017, Appendix XVIB	1 cfu per g or mL to 10⁵cfu per g or mL

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Specific Test SI. **Test Method Specification** Range of Testing / **Product / Material** Performed against which tests are Limits of Detection of Test performed IP 2014 (Vol. I), 2.2.9, Present or Absent per mL Staphylococcus aureus USP 39 <61>, <62>, or g BP 2017, Appendix XVIB E coli IP 2014 (Vol. I), 2.2.9, Present or Absent per mL USP 39 <61>, <62>, or g BP 2017, Appendix XVIB IP 2014 (Vol. I), 2.2.9, Pseudomonas Present or Absent per mL USP 39 <61>, <62>, aeruginosas or g BP 2017, Appendix XVIB Salmonella IP 2014 (Vol. I), 2.2.9, Present or Absent per mL USP 39 <61>, <62>, or g BP 2017, Appendix XVIB 3. **Antibiotics (Finished Product)** Amikacin IP 2014, 10 mg/mL to 550 mg/mL a. Assav **Sulphate Injection** (Cup Plate Method) (Vol. II) Page No.1028 IP b. Erythromycin IP 2014. 10 mg/ to 550 mg/Tablet Assav (Vol. II) Page No.1684 Stearate tablet IP (Cup Plate Method) 10 mg/mL to 40 mg/mL Gentamycin IP 2014, C. Assav Sulphate Injection (Cup Plate Method) (Vol. II) Page No.1859 IP 4. **Sterility Tests** Injection, Sterility Test IP 2014, 2.2.11, Qualitative a. **Ophthalmic** USP 39, <71>, (Complies/ Does not Preparations, BP 2017, Appendix XVI A comply) Water for Injection, Disposable Syringe 5. **Bacterial Endotoxins Test** IP 2014, 2.2.3, Injection, Water Bacterial endotoxin Qualitative a. for Injection (By Gel clot method) BP 2017, Appendix XIVC, (Complies/ Does not USP 39, <85> comply)

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	of Test	Performed	against which tests are	Limits of Detection
			performed	

CHEMICAL TESTING

I.	DRUGS AND PHA			
1.	Raw Material (General Test)	Description	IP 2014 BP 2017 USP 39	Qualitative
		Solubility	IP 2014, Page No. 174 BP 2017, Page No. V-11 USP 39, Page No. 1536	Qualitative
		Identification by IR	IP 2014, Page No. 337 BP 2017, Page No. V-A 159, USP 39, Page No. 144	Qualitative
		Identification by UV	IP 2014, Page No. 139 BP 2017, Page No. V-A 165, USP 39, Page No. 144	Qualitative
		Identification by Chemical	IP 2014 Page No. 87 BP 2017 Page No. V-A 225 USP 39 Page No. 140	Qualitative
		Loss on Drying	IP 2014 Page No. 162 BP 2017 Page No. V-A 296 USP 39 Page No. 369	1 % to 40 %
		Viscosity (HPMC)	IP 2014 Page No. 203 BP 2017 Page No. V-A 243 USP 39 Page No. 494	0.5 CPS to 200000 CPS
		Arsenic	IP 2014, Page No. 95 BP 2017, Page No. V-A 259, USP 39, Page No. 155	Qualitative
		Sulphate	IP 2014, Page No. 98 BP 2017, Page No. V-A 258 USP 39, Page No. 157	Qualitative

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SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Sulphated Ash (Lactose)	IP 2014, Page No. 98 BP 2017, Page No. V-A 293 USP 39, Page No. 176	0.01 % to 50 %
		Melting Range/Point (Urea)	IP 2014, Page No. 164-166 BP 2017, Page No. V-A 235 USP 39, Page No. 378	40 °C to 300 °C
		pH (Amoxycillin Trihydrate)	IP 2014, Page No. 169 – 170 BP 2017, Page No. V-A 248 USP 39, Page No. 402	0 to 14
		Refractive Index (Dicloro Methane)	IP 2014, Page No. 203 BP 2017, Page No. V-A 237 USP 39, Page No. 475	1.30 to 1.70
		Specific Optical Rotation (Amoxycillin Trihydrate)	IP 2014, Page No. 167-168 BP 2017, Page No. V-A 241 USP 39, Page No. 391	(-) 360° to (+) 360°
		Chloride	IP 2014, Page No. 96 BP 2017, Page No. V-A 256 USP 39, Page No. 157	Qualitative
2.	Drug Substance (API)			
a.	Parácetamol	Assay Dissolution	IP 2014, Page No. 2430 BP 2017, Page No. II-499 USP 39, Page No. 1564	50 % to 120 %
b.	Caffeine	Assay	IP 2014, Page No. 2058 BP 2017, Page No. I-344 USP 39, Page No. 3820	50 % to 120 %

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SI. **Test Method Specification** Range of Testing / Product / Material **Specific Test** Performed against which tests are Limits of Detection of Test performed Amoxycillin Water IP 2014, Page No. 1292 0.1 % to 40 % c. BP 2017, Page No. I-427 Trihydrate USP 39, Page No. 2181 d. Chloramphenicol IP 2014, Page No. 1349 Assay 50 % to 120 % BP 2017, Page No. I-496 USP 39, Page No. 2277 Nicotinamide IP 2014, Page No. 2332 Quantitative Assay е. BP 2017, Page No. II-366 f. IP 2014, Page No. 1016 Alprazolam **Related Substance** Qualitative BP 2017, Page No. I-104 USP 39, Page No. 1654 Diclofenac IP 2014, Page No. 1551 50 % to 120 % Assay g. Sodium **Related Substance** IP 2014, Page No. 1551 0 to 2.0 % Ibuprofen IP 2014, Page No. 1943 h. 50 % to 120 % Assay BP 2017, Page No. I-1165 **Tetracycline HCL** IP 2014, Page No. 2849 50 % to 120 % i. Assay **Related Substance** IP 2014, Page No. 2848-Qualitative 2849 j. Glycerin Ethylene Glycol, IP 2014, Page No. 1870 Qualitative Diethylene Glycol & BP 2017, Page No. I-1063 Related Substance USP 39, Page No. 3167 Water IP 2014, Page No. 1870 0.1 % to 40 % BP 2017, Page No. I-1063 USP 39, Page No. 3167 k. Sodium Carboxy 2.0 % to 15.0 % Assay IP 2014, Page No. 1281 Methyl Cellulose 3. Pharmaceutical Identification by IR IP 2014, Page No. 134 -Qualitative Dosage Forms/ 139 BP 2017, Page No. V-Preparations A 159 USP 39, Page No. (General Test) 144 IP 2014, Page No. 169 pН 2 to 12 BP 2017, Page No. V-A 248 USP 39, Page No. 343

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		Disintegration Time	IP 2014, Page No. 251 -253 BP 2017, Page No. V-A 333 USP 39, Page No. 342	0 to 180 min
		Uniformity of Weight	IP 2014, Page No. 256 BP 2017, Page No. V-A 357 USP 39, Page No. 491	As per specification
		Dissolution	IP 2014, Page No. 253-256 BP 2017, Page No. V-A 344, USP 39, Page No. 344	70 % to 110 %
4.	Pharmaceutical Dos	sage Forms		
1.	Chloroamphenicol Capsules	Assay	IP 2014, Page No. 1349 USP 39, Page No. 2327	50 % to 120 %
2.	Paracetamol Tablet	Assay	IP 2014, Page No. 2434 BP 2017, Page No. III-191- 920 USP 39, Page No. 1568	50 % to 120 %
3.	Ciprofloxacin HCL Tablets	Assay	IP 2014, Page No. 1404 BP 2017, Page No. III-310 USP 39, Page No. 2355	50 % to 120 %
4.	Alprazolam Tablets	Uniformity of Content	IP 2014, Page No. 1017 USP 39, Page No. 1656	50 % to 120 %
		Assay	IP 2014, Page No. 1018 USP 39, Page No. 1656	50 % to 120 %
II.	FOOD AND AGRICU	JLTURAL PRODUCTS		
1.	Milk & Milk Products	Test for Adulterants & Preservatives	FSSAI Lab Manual 1	
		Cane Sugar	FSSAI Lab Manual 1 :2016 Method 1.2.1	Present/Absent

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SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Starch	FSSAI Lab Manual 1 :2016 Method 1.2.2	Present/Absent
		Cellulose	FSSAI Lab Manual 1 :2016 Method 1.2.3	Present/Absent
		Added Urea	FSSAI Lab Manual 1 :2016 Method 1.2.4	Present
		Ammonium Sulphate	FSSAI Lab Manual 1 :2016 Method 1.2.5	Present/Absent
		Glucose	FSSAI Lab Manual 1 :2016 Method 1.2.7	Present/Absent
		Sodium Chloride	FSSAI Lab Manual 1 :2016 Method 1.2.8	Present/Absent
		Saccharin	FSSAI Lab Manual 1 :2012 Method 1.2.9	Present/Absent
		Salicylic Acid	FSSAI Lab Manual 1 :2016 Method 1.2.20	Present/Absent
		Dulcin	FSSAI Lab Manual 1 :2012 Method 1.2.10	Present/Absent
		Neutralizers	FSSAI Lab Manual 1 :2016 Method 1.2.11	Present/Absent
		Formalin	FSSAI Lab Manual 1 :2016 Method 1.2.17	Present/Absent
		Hydrogen Peroxide	FSSAI Lab Manual 1 :2016 Method 1.2.18	Present/Absent
		Boric Acid & Borates	FSSAI Lab Manual 1 :2016 Method 1.2.19	Present/Absent
2.	Food Grains, Pulses and Their	Foreign matter	Clause 6.2 of IS 4333 (Part 1): 2002 (RA 2012)	0 to 25 g/100g
	By Products	Weeviled grains	Clause 6.3 of IS 4333 (Part 1): 2002(RA 2012)	0 to 25 g/100g
		Damaged grains	Clause 6.3 of IS 4333 (Part 1): 2002 (RA 2012)	0 to 25 g/100g

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SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Insect damaged grains	Clause 6.3 of IS 4333 (Part 1): 2002 (RA 2012)	0 to 25 g/100g
		Broken Matter	Clause 6.3 of IS 4333 (Part 1): 2002 (RA 2012)	0 to 25 g/100g
		Other edible grains	Clause 6.3 of IS 4333 (Part 1): 1996 (RA 2012)	0 to 25 g/100g
		Moisture	Appendix A of IS 1155:1968 (RA 2010)	1 g/100g to 30 g/100g
		Total ash	Appendix B of IS 1155:1968 (RA 2010)	0.05 g/100g to 10 g/100g
		Gluten	Appendix D of IS 1155:1968 (RA 2010)	2 g/100g to 20 g/100g
		Protein	IS 7219:1973 (RA 2009)	0.5 g/100g to 40 g/100g
		Fat	Appendix F of IS 4684:1975 (RA 2009)	0.5 g/100g to 10 g/100g
		Carbohydrates	AOAC (18 th Ed) Official Method 986.25 (By Computation)	50 g/100g to 80 g/100g
		Energy (Calorific value)	IS 14433 (Part 1): 2000 (RA 2005) (By Computation)	325 Kcal/100g to 450 Kcal/100g
3.	Animal Feed & Pet Food	Moisture	Clause 4 of IS 7874 (Part I): 1975 (RA 2009)	0.5 g/100g to 25 g/100g
		Ash	Clause 9 of IS 7874 (Part I): 1975 (RA 2009)	0.5 g/100g to 10 g/100g
		Crude Fat	Clause 7 of IS 7874 (Part I): 1975 (RA 2009)	0.5 g/100g to 25 g/100g
		Crude Fibre	Clause 8 of IS 7874 (Part I): 1975 (RA 2009)	0.5 g/100g to 5 g/100g
		Protein	IS 7219:1973 (RA 2009)	0.5 g/100g to 60 g/100g
		Carbohydrate	AOAC (18 th Ed) Official Method 986.25 (By Computation)	30 g/100g to 70 g/100g

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4.	Bakery Products	Moisture	Annexure B of IS 1011:2002 (RA 2009)	0.5 % to 50 %
		Total Ash	Annexure C of IS 1011:2002 (RA 2009)	0.5 % to 10 %
		Acid Insoluble ash	Annexure C of IS 1011:2002 (RA 2009)	0.05 % to 2.0 %
		Fat	Clause 10 of IS 12711: 1989 (RA 2009)	0.5 % to 20 %
		Protein	IS 7219: 1973 (RA 2009)	1 % to 40 %
		Sugar	Appendix E of IS 2650:1975 (RA 2005)	1 % to 50 %
		Acidity of extracted fat	Annexure D of IS 1011:2002 (RA 2009)	0.1 % to 0.5 %
		Carbohydrate	AOAC (18 th Ed) Official Method 986.25 (By Computation)	50 g/100g to 80 g/100g
		Energy (Calorific value)	IS 14433 (Part 1): 1997 (RA 2005) (By Computation)	400 Kcal/100g to 500 Kcal/100g
5.	Snacks & Extruded Snacks	Moisture	Annexure B of IS 15271:2003 (RA 2009)	0.5 % to 10 %
	(Namkeens)	Fat	Annexure D of IS 15271:2003 (RA 2009)	0.5 % to 60 %
		Ash	Annexure C of IS 15271:2003 (RA 2009)	0.5 % to 10 %
		Acid Insoluble Ash	Annex C of IS 15271:2003 (RA 2009)	0.05 % to 2.0 %
		Protein	IS 7219: 1973 (RA 2009)	1 % to 20 %
		Acid value of extracted fat	Annex E of IS 15271:2003 (RA 2009)	0.1 % to 2.0 %
		Carbohydrate	AOAC (18 th Ed) Official Method 986.25 (By Computation)	30 g/100g to 70 g/100g

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		Energy (Calorific value)	IS 14433 (Part 1): 1997 (RA 2005) (By Computation)	450 Kcal/100g to 600 Kcal/100g
		Energy (calorific value)	IS 14433 (Part 1): 1997 (RA2005) (By Computation)	350 Kcal/100g to 500 Kcal/100g
6.	Honey	Moisture	Annexure-B-2 of IS 4941- 1994 (RA 2002) Amd. 2011	13 g/100g to 25g/100g
		Optical Density at 660 nm	Annexure-H of IS 4941- 1994 (RA 2002) Amd.2011	0.01 to 0.799
		Total reducing sugar	Annexure-C-1 of IS 4941- 1994 (RA 2002) Amd.2011	50 g/100g to 85 g/100g
		Sucrose	Annexure-C-2 of IS 4941- 1994 (RA 2002) Amd.2011	0.40 g/100g to 10 g/100g
		Ash	Annexure-D of IS 4941- 1994 (RA 2002) Amd.2011	0.002 g/100g to 5 g/100g
		Acidity (as Formic acid)	Annexure-E of IS 4941- 1994 (RA 2002) Amd.2011	0.01 g/100g to 0.5 g/100g
7.	Spices & Condiments	Extraneous matter	Clause 4 of IS 1797:1985 (RA 2009)	0.1 g/100g to 30 g/100gm
		Moisture /Loss on Drying	Clause 9 of IS 1797:1985 (RA 2009)	1 g/100g to 30 g/100g
		Total Ash	Clause 6 of IS 1797:1985 (RA 2009)	1 g/100g to 30 g/100g
		Acid insoluble ash	Clause 8 of IS 1797:1985 (RA 2009)	0.05 g/100g to 20 g/100g
		Water insoluble ash	Clause 7 of IS 1797:1985 (RA 2009)	0.5 m/100gm to 5 m/100gm
		Cold Water soluble extract	Clause 11 of IS 1797:1985 (RA 2009)	0.04 g/100g to 50 g/100g
		Alcohol soluble extract	Clause 10 of IS 1797:1985 (RA 2009)	0.04 g/100g to 50 g/100g
		Crude fibre	Clause 13 of IS 1797:1985 (RA 2009)	0.2 g/100g to 35 g/100g

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51.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Starch	Clause 9 of IS 4706 (Part 2): 1978 (RA 2009)	1 g/100g to 90 g/100g
		Acidity	Annexure A of IS 13242: 1991 (RA 2012)	0.5 % to 30 %
		Curcumin Content in Turmeric Powder/ Whole	Appendix B of IS 10925: 1984 (RA 2012)	1 % to 6 %
III.	WATER			
1.	Ground Water, Packaged	рН	IS 3025 (Part 11): 1983 (RA 2006)	2 to 12
	Drinking Water, Packaged Natural Mineral Water	Total Residue (Total solids-Dissolved & Suspended)	IS 3025 (Part 15): 1984 (RA 2009)	20 mg/l to 1500 mg/l
		Total dissolved solid	IS 3025 (Part 16): 1984 (RA 2006)	20 mg/l to 1500 mg/l
		Total alkalinity (as CaCO ₃)	IS 3025 (Part 23): 1986 (RA 2009)	10 mg/l to 500mg/l
		Total Hardness (as CaCO ₃)	IS 3025 (Part 21): 1983 (RA 2006)	5 mg/l to 1000 mg/l
		Calcium (as Ca)	IS 3025 (Part 40): 1991 (RA 2009)	2 mg/l to 500 mg/l
		Magnesium (as Mg)	IS 3025 (Part 46): 1994 (RA 2009) By Calculation	2 mg/l to 500 mg/l
		Chloride (as Cl)	IS 3025 (Part 32): 1988 (RA 2009)	5 mg/l to 400 mg/l