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| Chairperson | Prof Surjit Singh, Head of Pediatrics |
| Representative of Financial Advisor | Sr. AO (P.T.) |
| Legal Representative | Law Officer |
| General Public Representative | S. Karanvir Singh Sibia/Ms Jagdeep Kaur |
| Members of Committee | Prof Vanita Suri-Head, Obstetrics and Gynecology,  Prof Amit Rawat- Pediatric Medicine,  Prof Deepak Bansal- Pediatric Medicine  Dr Naveen Sankhyan-Pediatric Medicine |
| Member Convener | Dr Ranjitpal Singh Bhogal  Assistant Professor, Hospital Administration and Deputy Medical superintendent |
| Co- Opted Members |  |
| Hematological disorders | Prof Amit Trehan/Dr Prateek Bhatia, Prof Pankaj Malhotra  OR  Representative of Head, PediatricHematology/Hematology |
| Neuromuscular disorders | Prof Manish Modi; Dr RenuSuthar  Or  (Representative of Head, Pediatric Neurology/Neurology |
| Lysosomal Storage disorders | Prof InushaPanigrah/Dr Anupriya Kaur  Or  (Representative of Head, Genetics) |
| Nephrology | Dr KaralanghinTiewsoh/Dr LesaDawman  Or  (Representative of Head, Pediatric Nephrology/Nephrology) |
| Primary Immune Deficiency | Prof Deepti Suri/Dr VigneshPandhiarajan  Or  (Representative of Head, Pediatric Immunology) |
| Neurometabolic disorders | Prof Savita Verma/Dr.Arushi G Saini  Or  (Representative of Head, Pediatric Neurology) |
| Other Neurological disorders | Dr Jitender K Sahu  Or  (Representative of Head, Pediatric Neurology) |
| Endocrinal Disorders | Prof Devi Dayal/Prof Rakesh Kumar; Prof Sanjay Bhadada  Or  (Representative of Head, PediatricEndocrinolgy/Endocrinology) |
| Prenatal Diagnosis | Prof RashmiBagga, Dr Priyanka Srivastva  Or  (Representative of Head, Obstetrics &Gynecology/ Head, Genetics) |
| Other Disorders | Consultant to be co-opted on case-to-case basis |
| Representative of Parent Support Groups | Sh. Prashant Kumar Choubey/Sh Suresh Kumar (Other representative that may be suggested by committee members) |

**Committee on Rare Diseases:** A committee of following members has been constituted for implementation of National Health Policy for Rare Diseases,2021: