

**LISTADO DE FORMATOS Y REFERENCIAS:**

**PRENATAL & POSTNATAL ANALYSIS**

<b>PRODUCTO</b>	<b>REFERENCIA</b>	<b>FORMATO</b>
MLPA Aneuploidies	P095	100 rxn
MLPA Mental Retardation - Telomers	P036B	100 rxn
MLPA Mental Retardation - Telomers	P069	100 rxn
MLPA Mental Retardation – Telomers	P070	100 rxn
MLPA Mental Retardation syndroms – MR1	P064	100 rxn
MLPA Mental Retardation syndroms – MR2	P096	100 rxn
MLPA Monosomy 1p36	P147	100 rxn

**VARIOUS SYNDROMES AND DISEASES :**

<b>PRODUCTO</b>	<b>REFERENCIA</b>	<b>FORMATO</b>
MLPA Alagille syndrome	P184	100 rxn
MLPA Alzheimer early-onset	P170	100 rxn
MLPA Aniridia/Optic atrophy 1	P097	100 rxn
MLPA Aortic aneurysm syndrome	P148	100 rxn
MLPA a.d Dopa-responsive dystonia	P099	100 rxn
MLPa Arrhythmogenic right ventricular cardiomyopathy (ARVC)	P168	100 rxn
MLPA Ataxia telangiectasia syndrome	P123	100 rxn
MLPA Branchio-otorenal dysplasia syndrome (BOR)	P153	100 rxn

MLPA Bugada/long QT	P108	100 rxn
MLPA Canavan	P025	100 rxn
MLPA Cerebral Carvenous Malformations	P130/P131	100 rxn
MLPA Charcot Marie Tooth (CMT)	P033B	100 rxn
MLPA Classic galactosemia	P156	100 rxn
MLPA CMT2A/1B	P143	100 rxn
MLPA Congenital Adrenal Hyperplasia	P050	100 rxn
MLPA Congenital long QT syndrome	P114	100 rxn
MLPA Cornelia de Lange syndrome	P141/P142	100 rxn
MLPA Craniofacial disorders	P080	100 rxn
MLPA Cystic Fibrosys (CTFR)	P091	100 rxn
MLPA DiGeorge/Velocardiofacial/Cat eye	P023	100 rxn
MLPA Duchene/Becker	P034/P035	100 rxn
MLPA Duchene/Becker on Agarose	P071-P078	100 rxn
MLPA Fabry disease	P159	100 rxn
MLPA Fanconi Anemia	P031/P032	100 rxn
MLPA Fanconi Anemia Group B	P113	100 rxn
MLPA Gitelman syndrome	P136	100 rxn
MLPA GLUT1 deficiency syndrome	P138	100 rxn
MLPA Haemophilia A	P178	100 rxn
MLPA Hereditary spastic paraplegia	P165	100 rxn
MLPA Hirschprung	P169	100 rxn
MLPA Hunter Syndrome	P164	100 rxn

MLPA Hypercholesterolemia (LDLR)	P062	100 rxn
MLPA Hypertrophic cardiomyopathy	P100	100 rxn
MLPA Ichthiosis x-linked	P160	100 rxn
MLPA Intersex	P185	100 rxn
MLPA Lissencephaly	P061	100 rxn
MLPA Limb malformations-1	P179	100 rxn
MLPA Limb Malformations-2/Heart	P180	100 rxn
MLPA Limb girdle muscular dystrophies (LGMD)	P176	100 rxn
MLPA LMNA (several syndromes)	P048	100 rxn
MLPA LPAC syndrome (ABCB4 gene)	P109	100 rxn
MLPA Marfan syndrome	P065/P066	100 rxn
MLPA MEFV	P094	100 rxn
MLPA Menkes disease	P104	100 rxn
MLPA Mental Retardation syndromes - 1	P064	100 rxn
MLPA Mental Retardation syndromes- 2	P096	100 rxn
MLPA Mental retardation X-linked	P106	100 rxn
MLPA HMGS deficiency	P068	100 rxn
MLPA Mitochondrien	P125	100 rxn
MLPA Monosomy 1p36	P147	100 rxn
MLPA Myoclonic epilepsy	P137	100 rxn
MLPA Ophtalmogenetic Anomalies	P054	100 rxn
MLPA Ornithine catbamoyltransferase (OTC)	P079	100 rxn

MLPA Osler-Weber-Rendu syndrome (HHT)	P093	100 rxn
MLPA Parkinson (Familial)	P051/P052	100 rxn
MLPA Pelizaeus Merzbacher Disease	P022	100 rxn
MLPA Phenilketonuria (PAH)	P055	100 rxn
MLPA Pyruvate kinase (PK) deficiency	P203	100 rxn
MLPA Renitis Pitmentosa	P115	100 rxn
MLPA RETT syndrome	P015	100 rxn
MLPA Prader Willi/Angelman syndrome	ME028	100 rxn
MLPA PROS1 deficienty	P112	100 rxn
MLPA Pseudoxanthoma elasticum	P092	100 rxn
MLPA Simpson Golabi- Behmel syndrome	P154	100 rxn
MLPA SOTOS syndrome	P026B	100 rxn
MLPA Spynal Muscular Atrophy (SMA)	P021	100 rxn
MLPA Stargardt macular dystrophy (ABCA4)	P051/P052	100 rxn
MLPA Thalasemias	P140	100 rxn
MLPA Tuberous sclerosis TSC2	P046	100 rxn
MLPA Tuberous sclerosis TSC1	P124	100 rxn
MLPA Turner/Klinefelter/Triple X	P018	100 rxn
MLPA Waardenburg syndrome (WS)	P186	100 rxn
MLPA Williams Beuren Syndrome	P029	100 rxn
MLPA Willson disease	P098	100 rxn
MLPA X-linked hypohidrotic ectodermal dysplasia	P183	100 rxn

**CANCER HEREDITARIO:**

<b>PRODUCTO</b>	<b>REFERENCIA</b>	<b>FORMATO</b>
MLPA Ataxia teleangiectasia	P123	100 rxn
MLPA Breast Cancer	P002B	100 rxn
MLPA Breast Cancer	P087	100 rxn
MLPA Breast Cancer	P045	100 rxn
MLPA Colon Cancers (HNPCC)	P003	100 rxn
MLPA Colon Cancers (HNPCC)	P008	100 rxn
MLPA Colon Cancer (polyposis)	P043	100 rxn
MLPA Fanconi Anemia	P031/P032	100 rxn
MLPA Gorlin syndrome	P067	100 rxn
MLPA Li-Fraumeni syndrome	P056	100 rxn
MLPA Metastasis CDH1	P083	100 rxn
MLPA Multiple endocrine neoplasia	P017	100 rxn
MLPA Neurofibromatosis type 1	P122	100 rxn
MLPA Neurofibromatosis type 1	P081/P082	100 rxn
MLPA Neurofibromatosis type 2	P044	100 rxn
MLPA Peutz-Jeghers syndrome	P101	100 rxn
MLPA Retinoblastoma	P047	100 rxn
MLPA Tuberous sclerosis	P046	100 rxn
MLPA Von Hippel-Lindau syndrome	P016B	100 rxn
MLPA Various types of tumours	P024	100 rxn
MLPA Wilms Tumour	P118	100 rxn

**SUPRESION DE TUMORES:**

<b>PRODUCTO</b>	<b>REFERENCIA</b>	<b>FORMATO</b>
MLPA Breast, ovarian, cervical (ERBB2)	P004	100 rxn
MLPA Breast, ovarian, cervical in Agarose	P012	100 rxn
MLPA Chronic Lymphocytic Leukemia	P037/P038	100 rxn
MLPA Chromosome 8 kit	P014	100 rxn
MLPA CDKN2A-CDKN2B (P16)	P024	100 rxn
MLPA Gain1, Gain2 & Gain3	P171-P173	100 rxn
MLPA HNSCC1	P084	100 rxn
MLPA Human Chromosomal aberration1	P005	100 rxn
MLPA Human Chromosomal aberration2	P006	100 rxn
MLPA Human chromosomal aberration3	P007	100 rxn
MLPA Lung Cancer	P126/P127	100 rxn
MLPA Neuroblastoma	P161/P162	100 rxn
MLPA Oligodendroglioma1	P088	100 rxn
MLPA Oligodendroglioma2	P105	100 rxn
MLPA Tumor suppressor	ME001	100 rxn