



Igisata kijejwe kwitaho abantu
 Igisata citaho amagara meza y'umwana n'umuvyeyi
 Abana bakenera ubundi bufasha bwiharije
 Igisata kijejwe gupima abana bakivuka(inzoya)

UGUHAKANA IGIKORWA CO GUPIMISHA ABANA BAKIVUKA

Jewe/twebwe, _____, Umuvyeyi/Umurezi wa
 Amazina y'umuvyeyi/y'umurezi

_____, yavutse kuwa _____
 Amazina y'uruyoya itariki y'amavuko

_____, mpakanye(duhakanye) k'umwana wacu afatwa amaraso ku mvo z'ukumenya
 Aho yavukiye

Ko ata kibazo cokika amagara yiwe kikandurukamwo urupfu,ubumuga canke ubundi burwayi. Mu vyashoboye gukurikiranwa n'ugupimwa harimwo ibi mirongo itatu na bitatu vyanditse aha hepfo. Vyarasiguwe ko Ubushikirangaji bujewe amagara y'abantu I Vermont buhimiriza ko inzoya zose zopimwa mu minsi ya mbere hisunzwe ivyakurikiranywe.

3-Methylcrotonyl-CoA carboxylase deficiency (3MCC)

3-OH 3-CH3 glutaric aciduria (HMG)

Argininosuccinic acidemia (ASA)

Beta-ketothiolase deficiency (BKT)

Biotinidase deficiency (BIOT)

Carnitine uptake defect (CUD)

Citrullinemia (CIT)

Congenital adrenal hyperplasia (CAH)

Congenital hypothyroidism (HYPOTH)

Cystic fibrosis (CF)

Galactosemia (GALT)

Glutaric acidemia type I (GA I)

Hb S/Beta-thalassemia (Hb S/Th)

Hb S/C disease (Hb S/C)

Homocystinuria (HCY)

Isovaleric acidemia (IVA)

Long-chain L-3-OH acyl-CoA dehydrogenase deficiency (LCHAD)

Maple syrup urine disease (MSUD)

Medium-chain acyl-CoA dehydrogenase deficiency (MCAD)

Methylmalonic acidemia (Cbl A, B)

Methylmalonic acidemia (MUT)

Mucopolysaccharidosis type I (MPS I)

Multiple carboxylase deficiency (MCD)

Phenylketonuria (PKU)

Pompe disease

Propionic acidemia (PROP)

Severe Combined Immunodeficiency (SCID)

Sickle cell anemia (SCA)

Spinal muscular atrophy (SMA)

Trifunctional protein deficiency (TFP)

Tyrosinemia type I (TYR I)

Very long-chain acyl-CoA dehydrogenase deficiency (VLCAD)

X-linked adrenoleukodystrophy (X-ALD)

Ubundi bwoko bw'ibipimo ni: Igipimo c'amatwi hamwe n'igipimo c'ukwo umutima utera mukuraba kw'atakindi kibazo c'umutima yoba afise.

~Na/(Twa)ramenyeshewe yukwo mu bikorwa harimwo n'ugutera urushinge muni y'ikirenge mu gufata amaraso.

~Na/(Twa)raronse akaryo ko kuvugana ivy'ibipimo na muganga w'umwana wacu,abandi baganga eka n'abandi bafasha kwa muganga, twaranyuzwe n'inyishu zose batwishuye.

~Nda/(Tura)tahura ko mu gihe umwana avukanye ikibazo na kimwe murivyo twavuze, mu gihe adapimwe ngo avurwe murico minsi akivuka amahirwe ni menshi y'ukugira amagara mabi,ubwenge buke canke urupfu.

~Ndemeza/Turemeza ko runo rupapuro nujuje ruzoshirwa hamwe n'ibindi vy'umwana wacu,ivyafotowe bikajanywa ku bamuvura no ku bushikiranganji bw'Amagara y'Abantu bw'I Vermont.

Igikumu c'umuvyeyi/Umurezi

Itariki

Igikumu c'icabona

Itariki

Ibifasha mu kwuzuzwa:

1. Runo rupapuro rutegerezwa kwuzuzwa kubwa kira ruyoya mugihe umuvyeyi canke umurezi banse ibipimo vy'uruyoya. Urwashizweko umukono rugashirwa muri dosiye yo kwa muganga y'umwana canke mu gihe avukiye muhira, kugira bibonwe n'uwumwibarutsa.
2. Ivyafotowe birangikwa kubatanga ubuvuzi butangura gukenerwa hamwe no mu Gisata c'I Vermont kijejwe gupima abana b'inzoya. Agasandugu ka Posita 70, 108 Cherry St., Burlington, VT 05402. Mufise ico mubaza, hamagara kuri (802) 951-5180