

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

UGUHAKANA IGIKORWA CO GUSUBIRA GUPIMISHA UMWANA AKIVUKA

Jewe/Twebwe, _____, Umuvyeyi/Umurezi wa
 Amazina y'umuvyeyi/umurezi _____, yavutse kuwa _____
 Amazina y'uruyoya Itariki y'amavuko _____

Aho yavukiye
 Ko ata kibazo cokika amagara yiwe kikandurukamwo urupfu, ubumuga canke ubundi burwayi. Turatahura ko ibipima vyakozwe ubwambere bitari bihagije canke vyasabwe ko hakorwa ibindi bipimo vy'amaraso. Ivyihwejwe mu bipimonvy'amaraso harimwo ibi mirongo itatu na bitatu bikurikira: Nda(Tura)tahura ko Ubushikiranganji bujejwe amagara y'abantu bw'I Vermont buhimiriza kw'inzoya zose zikurikiranwa mu minsi ya mbere hisunzwe ibi bikurikira:

<i>3-Methylcrotonyl-CoA carboxylase deficiency (3MCC)</i>	<i>Maple syrup urine disease (MSUD)</i>
<i>3-OH 3-CH₃ glutaric aciduria (HMG)</i>	<i>Medium-chain acyl-CoA dehydrogenase deficiency (MCAD)</i>
<i>Argininosuccinic acidemia (ASA)</i>	<i>Methylmalonic acidemia (Cbl A, B)</i>
<i>Beta-ketothiolase deficiency (BKT)</i>	<i>Methylmalonic acidemia (MUT)</i>
<i>Biotinidase deficiency (BIOT)</i>	<i>Mucopolysaccharidosis type I (MPS I)</i>
<i>Carnitine uptake defect (CUD)</i>	<i>Multiple carboxylase deficiency (MCD)</i>
<i>Citrullinemia (CIT)</i>	<i>Phenylketonuria (PKU)</i>
<i>Congenital adrenal hyperplasia (CAH)</i>	<i>Pompe disease</i>
<i>Congenital hypothyroidism (HYPOTH)</i>	<i>Propionic acidemia (PROP)</i>
<i>Cystic fibrosis (CF)</i>	<i>Severe Combined Immunodeficiency (SCID)</i>
<i>Galactosemia (GALT)</i>	<i>Sickle cell anemia (SCA)</i>
<i>Glutaric acidemia type I (GA I)</i>	<i>Spinal muscular atrophy (SMA)</i>
<i>Hb S/Beta-thalassemia (Hb S/Th)</i>	<i>Trifunctional protein deficiency (TFP)</i>
<i>Hb S/C disease (Hb S/C)</i>	<i>Tyrosinemia type I (TYR I)</i>
<i>Homocystinuria (HCY)</i>	<i>Very long-chain acyl-CoA dehydrogenase deficiency (VLCAD)</i>
<i>Isovaleric acidemia (IVA)</i>	<i>X-linked adrenoleukodystrophy (X-ALD)</i>
<i>Long-chain L-3-OH acyl-CoA dehydrogenase deficiency (LCHAD)</i>	

Ibindi bipimo vy'amaraso bishobora gukorerwa muhira canke kwa muganga harimwo n'igipimo c'amatwi hamwe n'igipimo c'ukwo umutima utera mukuraba kw'atakindi kibazo c'umutima yoba afise.

~Na/ (Twa)rasomye agatabu ndangabikorwa k'Ubushikiranganji bw'Amagara meza bw'I Vermont mu gisata c'ugupima inzoya mu minsi ya mbere.

~Ndatahura/Turatahura ko gisata c'ugupima inzoya c'I Vermont gihimiriza ko ibipimo bikurikiranwa kubera hihwejwe ibipimo vyakozwe kuwa _____ vyerekanye ko _____.

~Ndumva/Turumva ko muri twebwe twabwiwe vyose vyerekeye uguSubiramwo ibipimo vy'umwana/abana bacu, dufata n'ingingo y'ukudasubira.

~Jewe/Twebwe, ntitwipfuza gusubira kuyaga n'abatanga ubufasha mu vyerekeye ibipimo vy'amaraso ku ruyoya canke n'umuganga ajejwe uruyoya rwacu canke abandi bose bafasha kuvy'amagara meza bafise umwanya wo kwishura ibibazo biberekeye twobaza.

~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.

Igikumu c'umuvyeyi/Umurezi

Itariki

Igikumu c'icabona

Itariki