






	Patient Population	Pre-clinical	Phase 1/2	Phase 3	Regulatory Approval
<b>Setmelanotide</b> <i>daily formulation</i>	POMC, PCSK1 or LEPR deficiency				US, Canada*
	Bardet-Biedl syndrome				US, Canada*
<b>Setmelanotide</b> <i>daily formulation</i>	Hypothalamic obesity			Enrollment complete	
	 Obesity and Hunger Clinical Trial				
	Pediatrics (age 2 to <6 years, biallelic POMC, PCSK1, or LEPR deficiency or Bardet-Biedl syndrome)				
	 Obesity and Hunger Clinical Trial				
<b>Setmelanotide</b> <i>weekly formulation</i>	Switch study with patients previously on setmelanotide				
<b>LB54640</b>	Hypothalamic obesity				
<b>RM-718</b>	Rare MC4R pathway diseases				Ph1 trial 1H2024
<b>Pre-clinical</b>	Congenital Hyperinsulinism (CHI)				Lead identification underway

 Complete
  Denotes trial underway
  Denotes planned trial

\*Other countries include EU, Great Britain, Israel

LEPR, leptin receptor; MC4R, melanocortin-4 receptor; PCSK1, proprotein convertase subtilisin/kexin type 1; POMC, proopiomelanocortin.