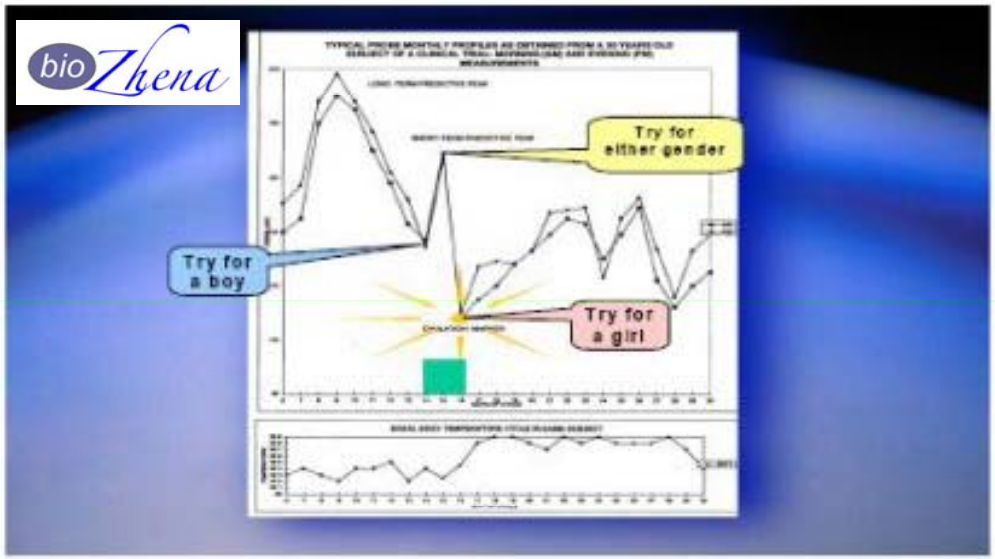
Summary of the first application



The personal home-use device determines the 3 days during which conception can occur , and it generates a new type of profile for use by the medical profession

Citing from FDA 510K Letter Number K973860: ... *The Monitor serves as an independent information* ***aid to the woman*** *by helping her to define the fertile window... whereby she may choose the* ***proper timing*** *for vaginal intercourse. ... The Monitor may serve to provide the user* ***and her physician*** *with data to better time artificial insemination or other interventional techniques.*