



A: I would first advise you to remove the URL if you don't need it. I was able to build the example dataset into a Google Fusion Table to meet the criteria you mention: I used the following settings: Google Fusion Table Server URL: Google Fusion Table ID: SimpleNotebook Location: Somewhere in the EU (here in Switzerland) I have no recommendation on how to tune the LAS opts. Your VLOOKUP gets the dataset from the Time variable, which is the time of the day (the midpoint of the cell number). The Time needs to start from noon as the time is important in this column. The * Needs to be on the Time column, while the (1,)* needs to be on the 2nd column. VLOOKUP(G2,C2:R16,1,0) So, to summarize: Remove the URL. Use VLOOKUP(G2,C2:R16,1,0) and it should work with a reference range of "Reference:" on your sheet. Note: Since the VLOOKUP was not automated I couldn't get the Reference to be part of the formula. I also included a step to add some rows to the dataset to assure the range will fit the reference range. The result: The proposed research will focus on the mechanisms of developmental apoptosis and the associated aging in the context of the two highly conserved, but non-equivalent processes, apoptosis and innate immunity. Whereas aging is a homogeneous process, apoptosis appears to result from a wide variety of intrinsic and extrinsic signals, each of which is likely to produce a response that promotes the survival or death of the organism. The specific aims of the proposal are to test the hypothesis that apoptosis in the developing nervous system is one mechanism by which a variety of extrinsic signals and intrinsic danger signals trigger a response that prevents tissues from being overwhelmed by a developmentally inappropriate excess of cells. This hypothesis will be addressed through three specific aims. First, the involvement of p53 and its tumor suppressor gene will be examined in axon growth, synaptogenesis and synaptic pruning as a result of neurotrophic factor withdrawal. Second, pro-apoptotic molecules will be investigated as mediators of developmental or age-related apopt

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