# Using Cvt2Mae to Convert Incyte Array Data for MAExplorer

http://www.lecb.ncifcrf.gov/Cvt2Mae

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Cvt2Mae version 0.60

# Accessing Arrays with MAExplorer

- MAExplorer works with any arrays using the schema (see Appendix C of MAExplorer Reference Manual for details)
- All data files are tab-delimited text files
- Databases could be constructed with tools like Excel for editing user data into the schema format
- The Cvt2Mae array data converter "Wizard" tool converts non-standard <User-defined> academic or commercial data to MAExplorer format
- Affymetrix, **Incyte**, GenePix, Scanalyze, and other array data formats may be converted using predefined "Array Layouts"

# S.1 MAExplorer Data Schema

- MAExplorer works with any array data using our data schema

- The schema is described in detail in MAExplorer Reference Manual Appendix C.

- Data Schema: tab-delimited experiment data files:
  1. GIPO (Gene In Plate Order or "print" file)
  2. List of hybridized samples in database
  3. Configuration data describing the array and conventions
  4. Separate spot quantification data files

- The Cvt2Mae "wizard" tool converts user array data to this schema

# S.1.1 MAExplorer GIPO or Print File

- GIPO file maps a spot on the array to a particular gene

- Contains:
  1. location or grid-geometry
  2. one or more genomic identifiers (e.g., Clone ID, GenBank ID, LocusID, etc.)
  3. gene description as Gene Name (or other description)
  4. Optional: global spot quality (QualCheck)
  5. optional: plate coordinates for clones

# S.1.2 MAExplorer Samples Database File

- List of hybridized samples file SamplesDB.txt file contains:

1. full sample description

2. base file name of quantification file (without .quant file extension)

3. optional sample ID number

4. other data you wish to carry with the samples (used in array reports)

# S.1.3 MAExplorer Configuration Database File

- Configuration data file MaeConfig.txt describes particular type of array and hybridization labeling you are using. This includes:

- grid-geometry - # of replicate fields, grids, rows/grid, columns/grid

- spot hybridization labeling - intensity or ratio data, dye names

- various presentation options - use pseudo-array or actual (x,y) coordinates, etc.

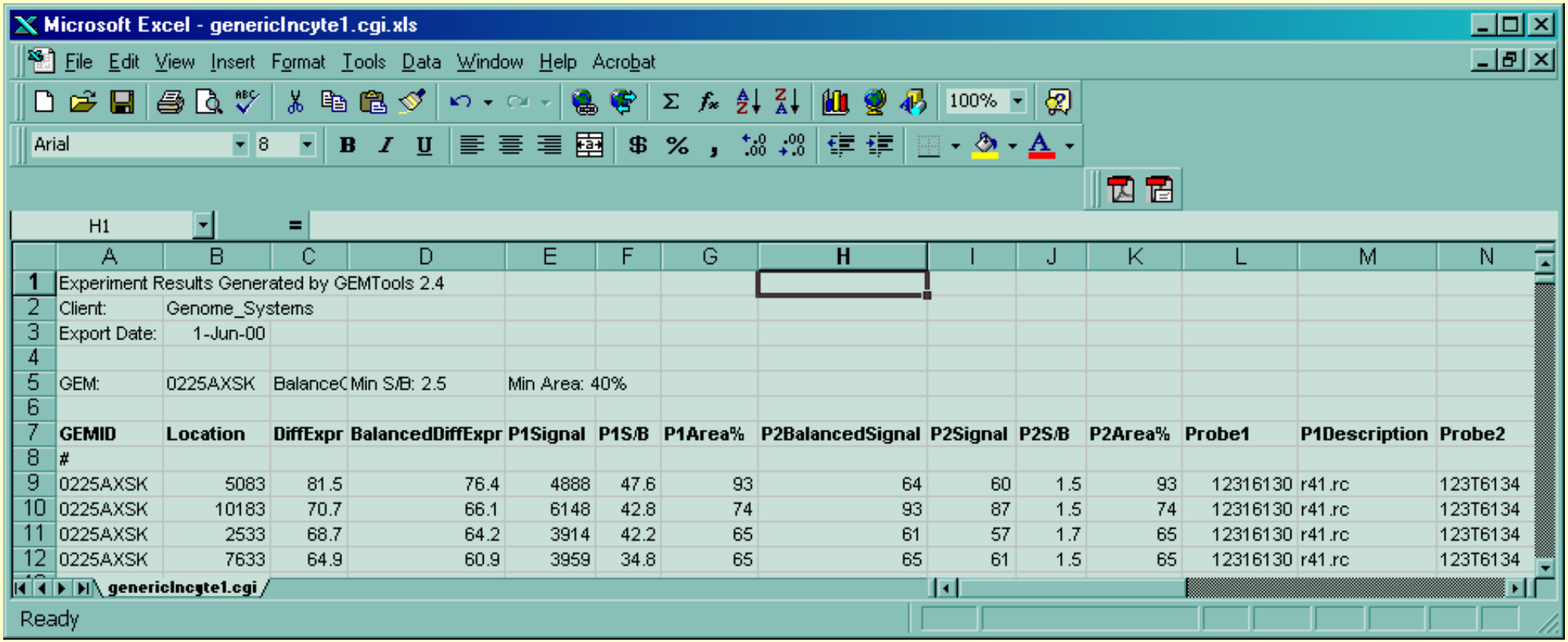
# S.1.4 MAExplorer Spot Quantification Files

- Separate spot quantification data files (with .quant file extension) are used for each hybridized sample
- $^{33}P$ or biotin labeled samples are specified as one hybridization intensity information per file
- Fluorescent Cy3/Cy5-dye labeled samples are specified as two channels of hybridization intensity information per file
- Intensity background data is optional
- Spot quality (QualCheck) data is optional
- Grid-coordinates are specified the same as for GIPO file

# S.2 Assumptions About User Data - Array Layout

- User data is tab-delimited ASCII text files (could generate with Excel)
- If the array geometry (#fields, grids, rows/grid, columns/grid) is known, that geometry may be used in MAExplorer
- Otherwise, a pseudo-array geometry is generated for visual use in MAExplorer from the total # of spots in the user data
- An Array Layout describes the user data. It may be edited and saved for subsequent use in converting other array data files of the same type
- The <User-defined> array layout gives users complete flexibility in describing the array

# S.3 Example of tab-delimited Incyte Data



Microsoft Excel - genericIncyte1.cgi.xls

File Edit View Insert Format Tools Data Window Help Acrobat

Arial 8 100%

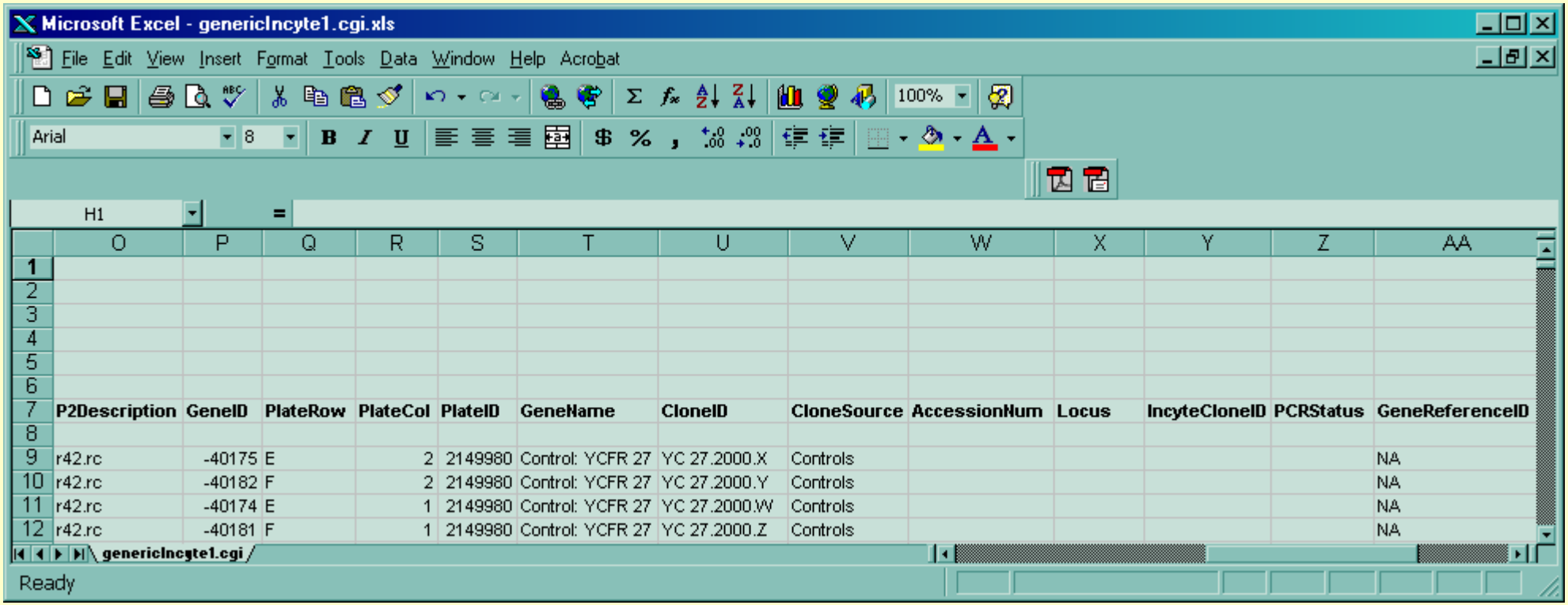
H1 =

| | A | B | C | D | E | F | G | H | I | J | K | L | M | N |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 1 | Experiment Results Generated by GEMTools 2.4 | | | | | | | | | | | | | |
| 2 | Client: | Genome_Systems | | | | | | | | | | | | |
| 3 | Export Date: | 1-Jun-00 | | | | | | | | | | | | |
| 4 | | | | | | | | | | | | | | |
| 5 | GEM: | 0225AXSK | BalanceC | Min S/B: 2.5 | Min Area: 40% | | | | | | | | | |
| 6 | | | | | | | | | | | | | | |
| 7 | **GEMID** | **Location** | **DiffExpr** | **BalancedDiffExpr** | **P1Signal** | **P1S/B** | **P1Area%** | **P2BalancedSignal** | **P2Signal** | **P2S/B** | **P2Area%** | **Probe1** | **P1Description** | **Probe2** |
| 8 | # | | | | | | | | | | | | | |
| 9 | 0225AXSK | 5083 | 81.5 | 76.4 | 4888 | 47.6 | 93 | 64 | 60 | 1.5 | 93 | 12316130 | r41.rc | 123T6134 |
| 10 | 0225AXSK | 10183 | 70.7 | 66.1 | 6148 | 42.8 | 74 | 93 | 87 | 1.5 | 74 | 12316130 | r41.rc | 123T6134 |
| 11 | 0225AXSK | 2533 | 68.7 | 64.2 | 3914 | 42.2 | 65 | 61 | 57 | 1.7 | 65 | 12316130 | r41.rc | 123T6134 |
| 12 | 0225AXSK | 7633 | 64.9 | 60.9 | 3959 | 34.8 | 65 | 65 | 61 | 1.5 | 65 | 12316130 | r41.rc | 123T6134 |

genericIncyte1.cgi

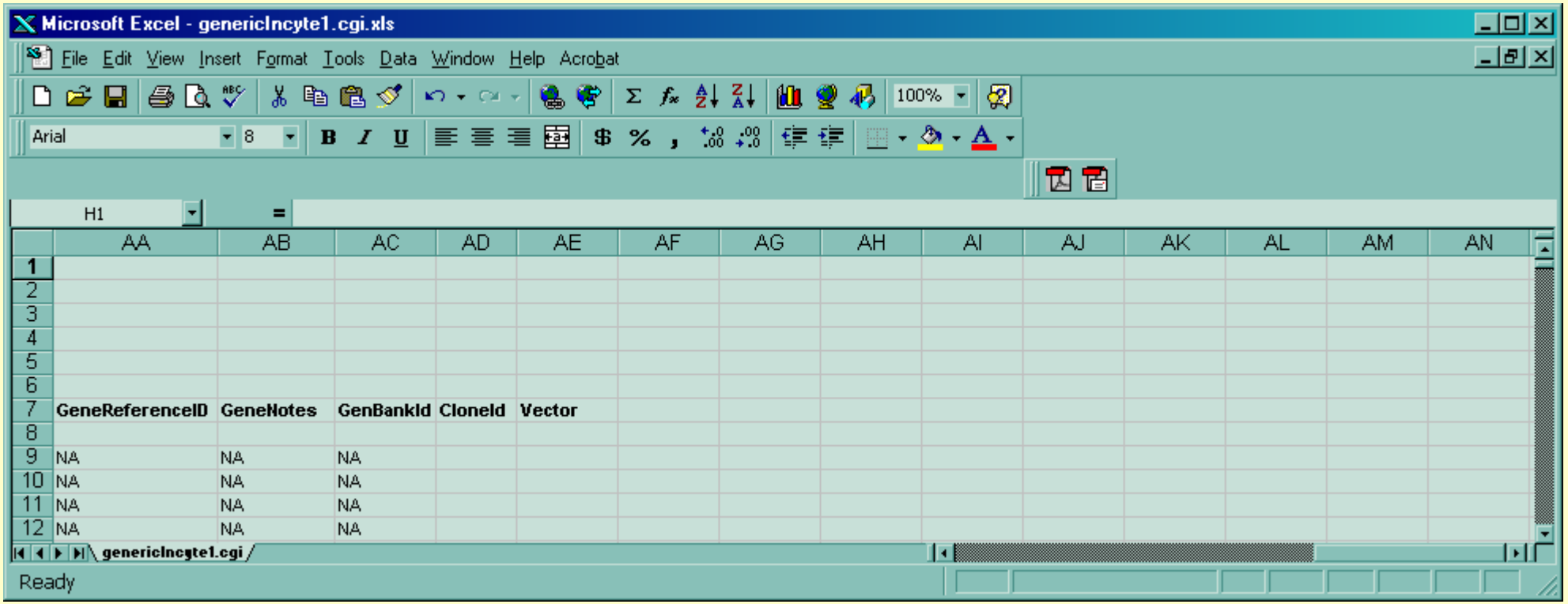
Ready

# S.3.1 Example of tab-delimited Incyte Data




| | O | P | Q | R | S | T | U | V | W | X | Y | Z | AA |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 1 | | | | | | | | | | | | | |
| 2 | | | | | | | | | | | | | |
| 3 | | | | | | | | | | | | | |
| 4 | | | | | | | | | | | | | |
| 5 | | | | | | | | | | | | | |
| 6 | | | | | | | | | | | | | |
| 7 | **P2Description** | **GeneID** | **PlateRow** | **PlateCol** | **PlateID** | **GeneName** | **CloneID** | **CloneSource** | **AccessionNum** | **Locus** | **IncyteCloneID** | **PCRStatus** | **GeneReferenceID** |
| 8 | | | | | | | | | | | | | |
| 9 | r42.rc | -40175 | E | 2 | 2149980 | Control: YCFR 27 | YC 27.2000.X | Controls | | | | | NA |
| 10 | r42.rc | -40182 | F | 2 | 2149980 | Control: YCFR 27 | YC 27.2000.Y | Controls | | | | | NA |
| 11 | r42.rc | -40174 | E | 1 | 2149980 | Control: YCFR 27 | YC 27.2000.W | Controls | | | | | NA |
| 12 | r42.rc | -40181 | F | 1 | 2149980 | Control: YCFR 27 | YC 27.2000.Z | Controls | | | | | NA |

# S.3.2 Example of tab-delimited Incyte Data



# I. Procedure: Convert Data for Array Layouts

1. Select the Chip Set array layout (**Incyte - generic**) if in list, otherwise pick <User-defined>)

2. Select 1 or more input files using the "Browse filename" .

3. You may edit or change various array layout parameters at this time
   3.1 you may edit the array layout with "Edit Layout"
   3.2 you may "Assign GIPO fields" in user data file
   3.3 you may "Assign Quantification fields" in user data file
   3.4 if you changed any array layout parameters, you may save it with "Save Layout"

4. Select the project output directory (i.e., folder) to save generated files

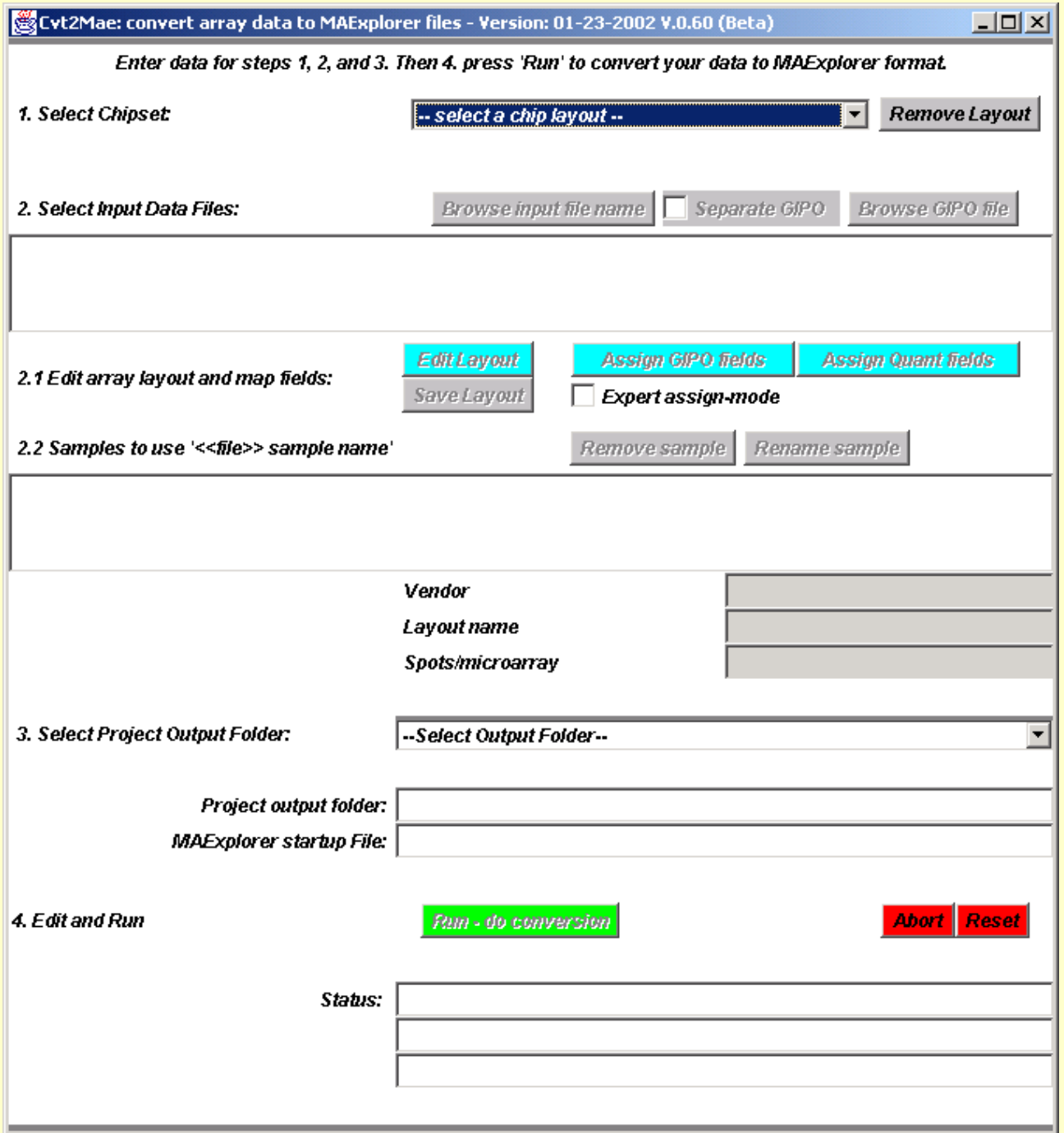
# I. Procedure: continued...

5. Press “Run” to convert the data

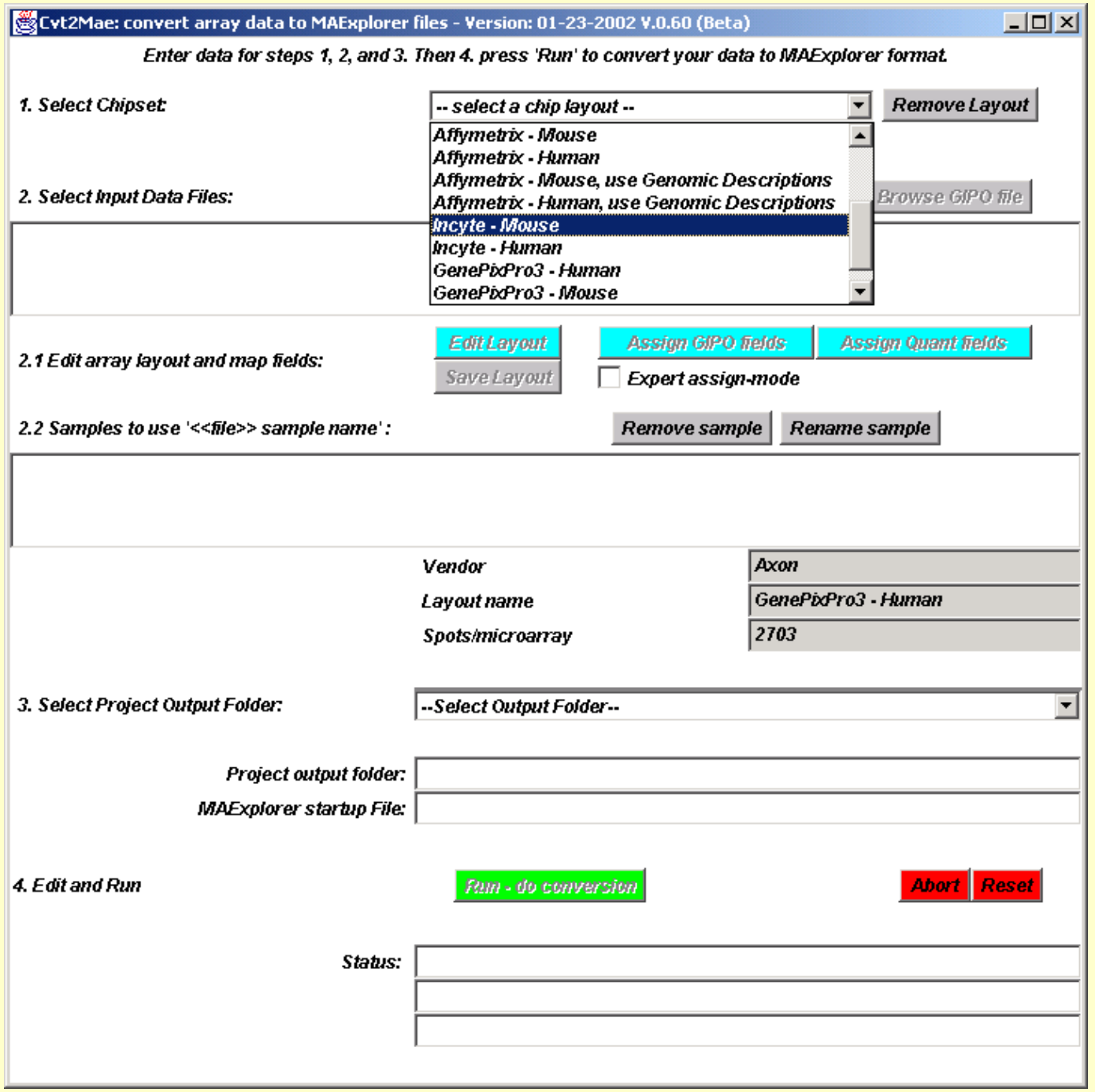
6. Press “Done” when it is finished.

7. Go to the project directory and then to the MAE sub-directory, click on the Start.mae file to start MAExplorer on the new data

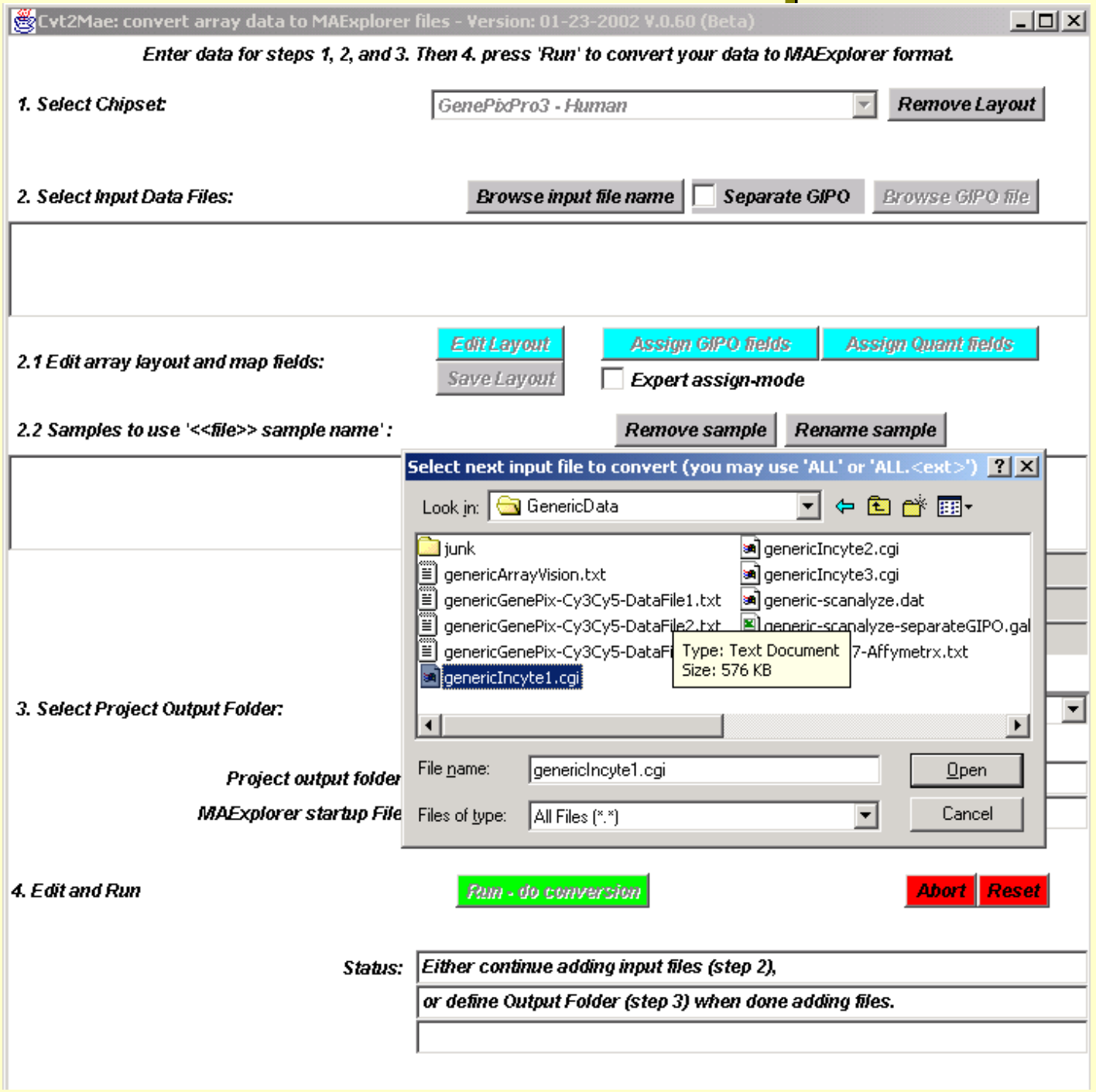
# 1. Initial State of Cvt2Mae Program



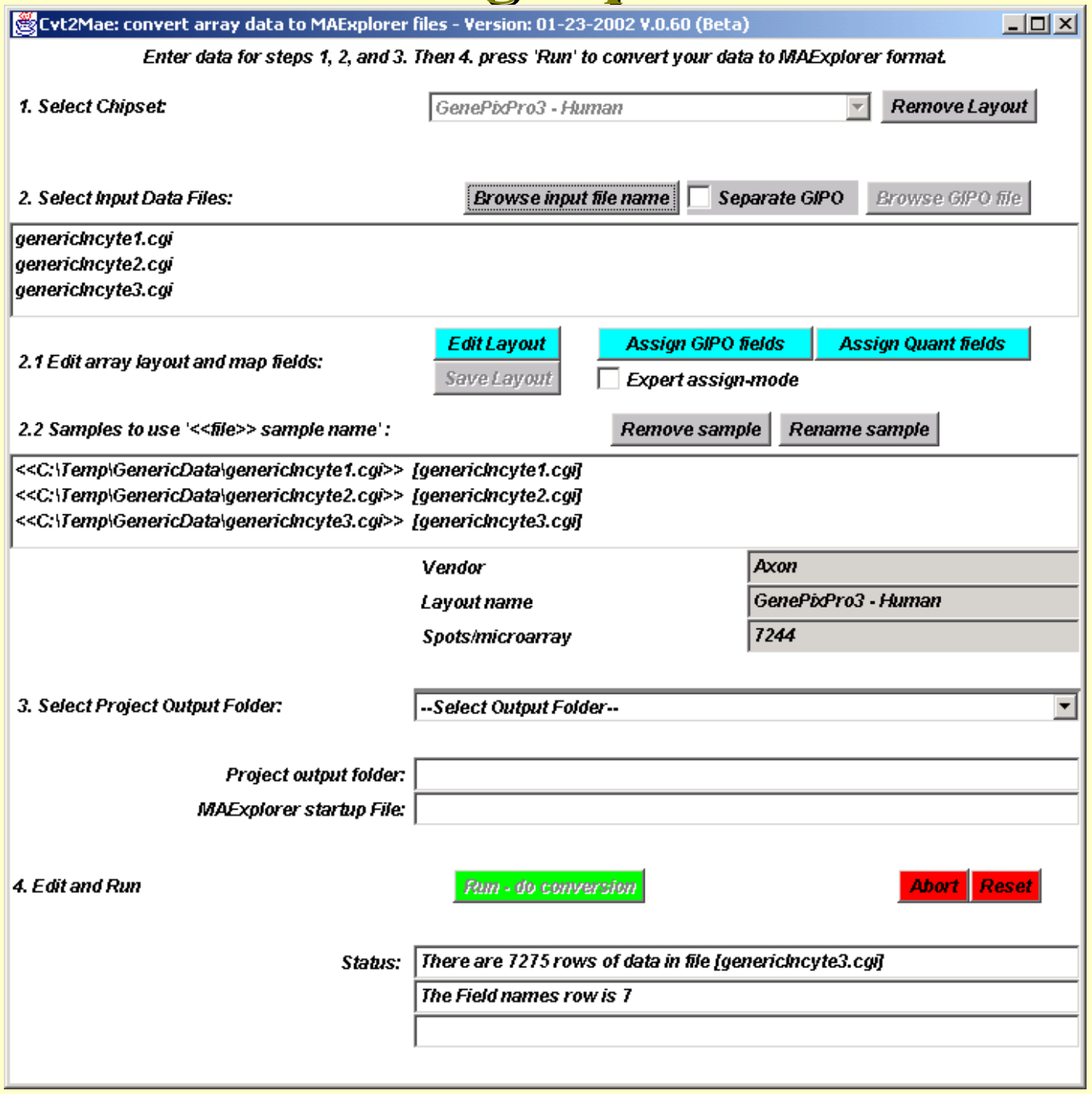
# 2. Selecting Incyte Chipset Array-Layout



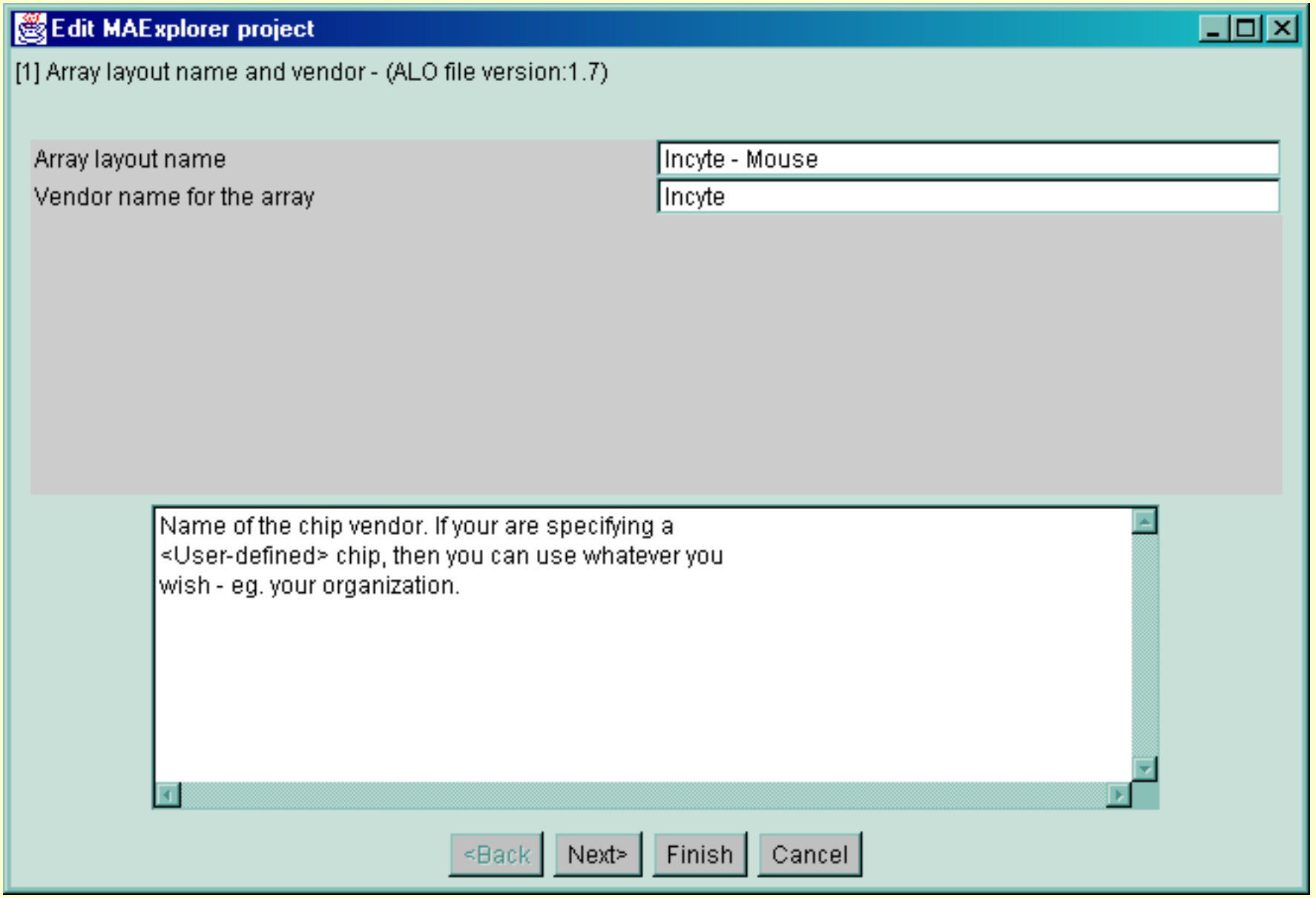
# 3. Select Files with "Browse input file" Name



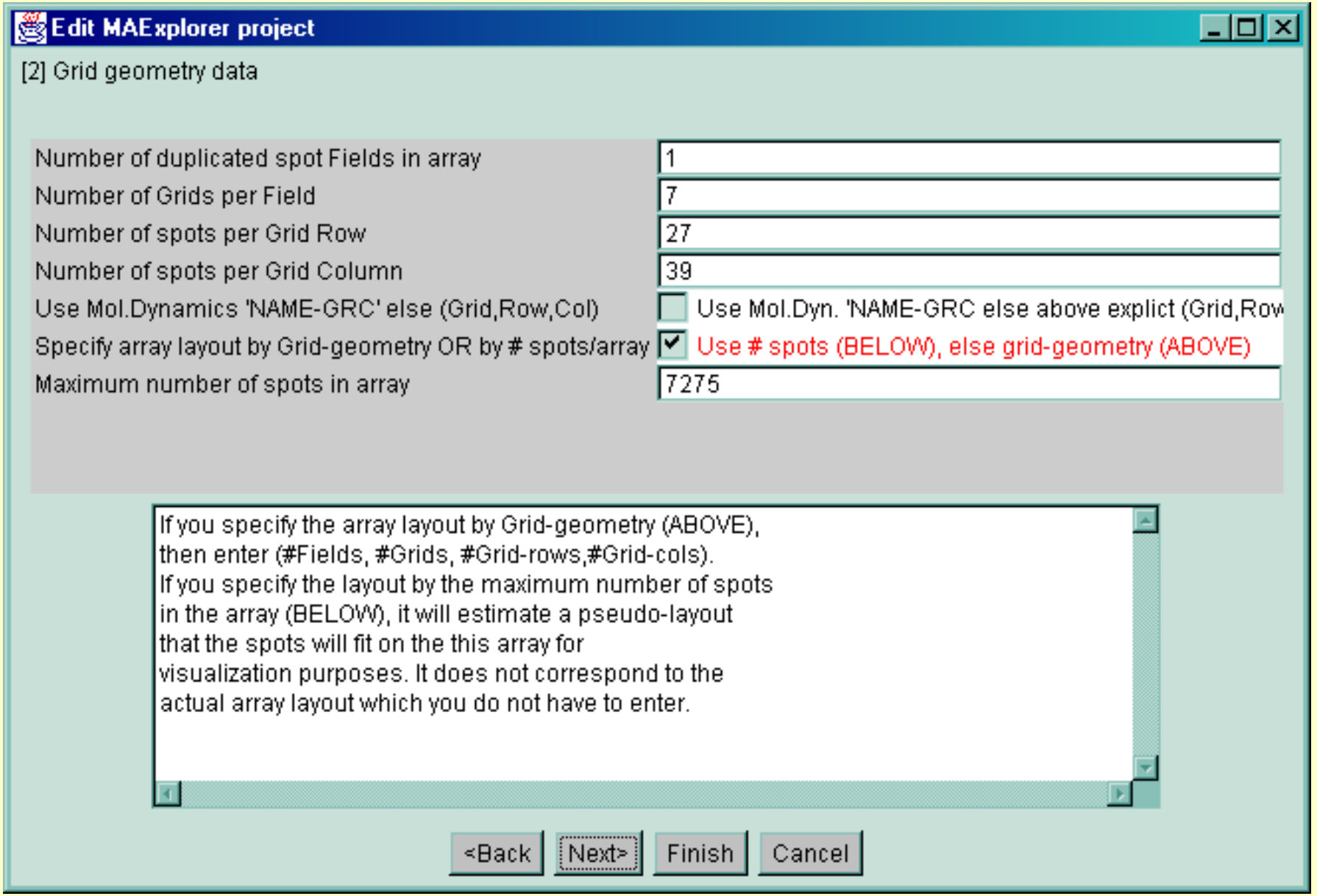
# 4. Continue Adding Input Files If Needed



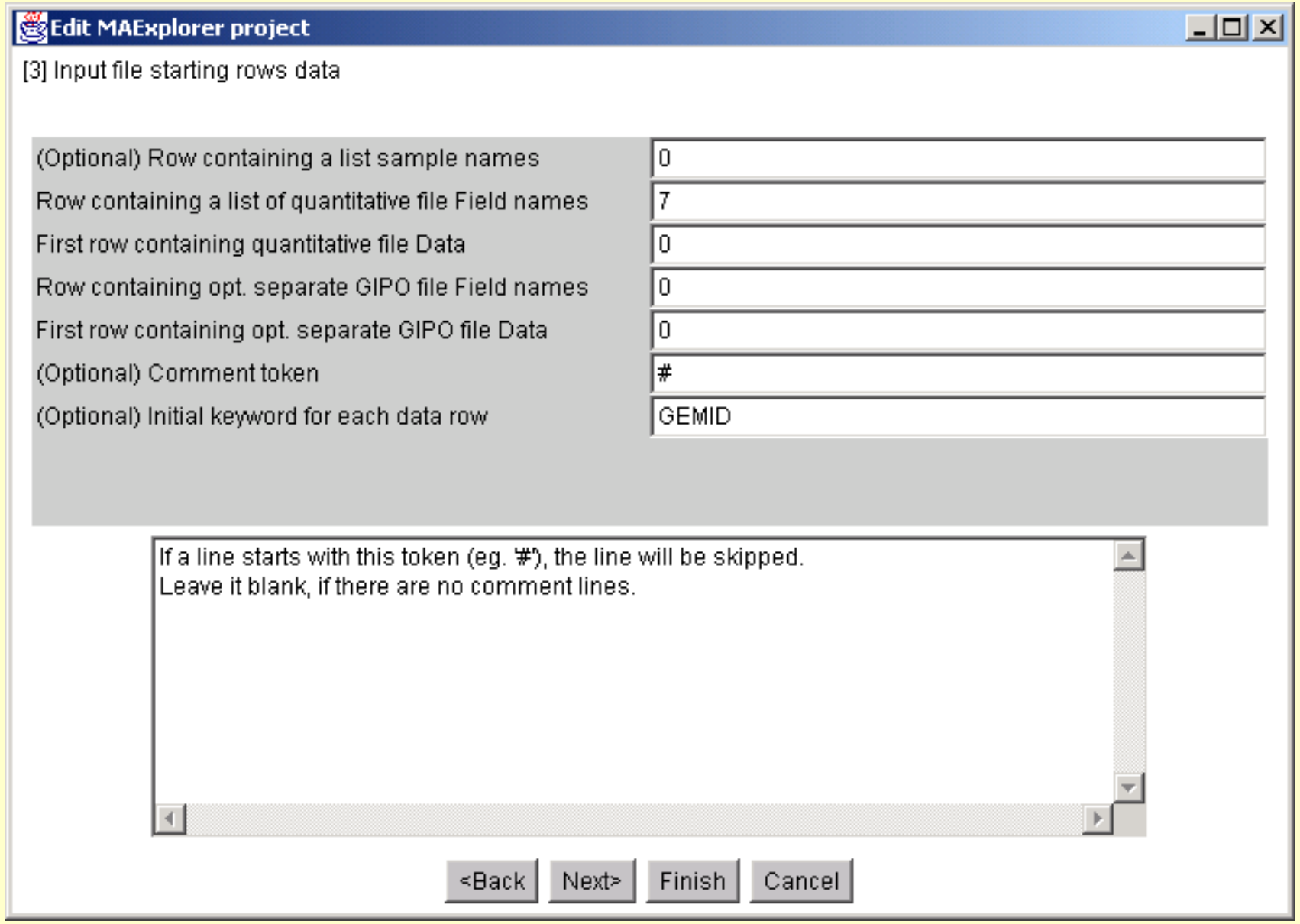
# 5.1 Edit Layout 'Wizard' Values for This Array



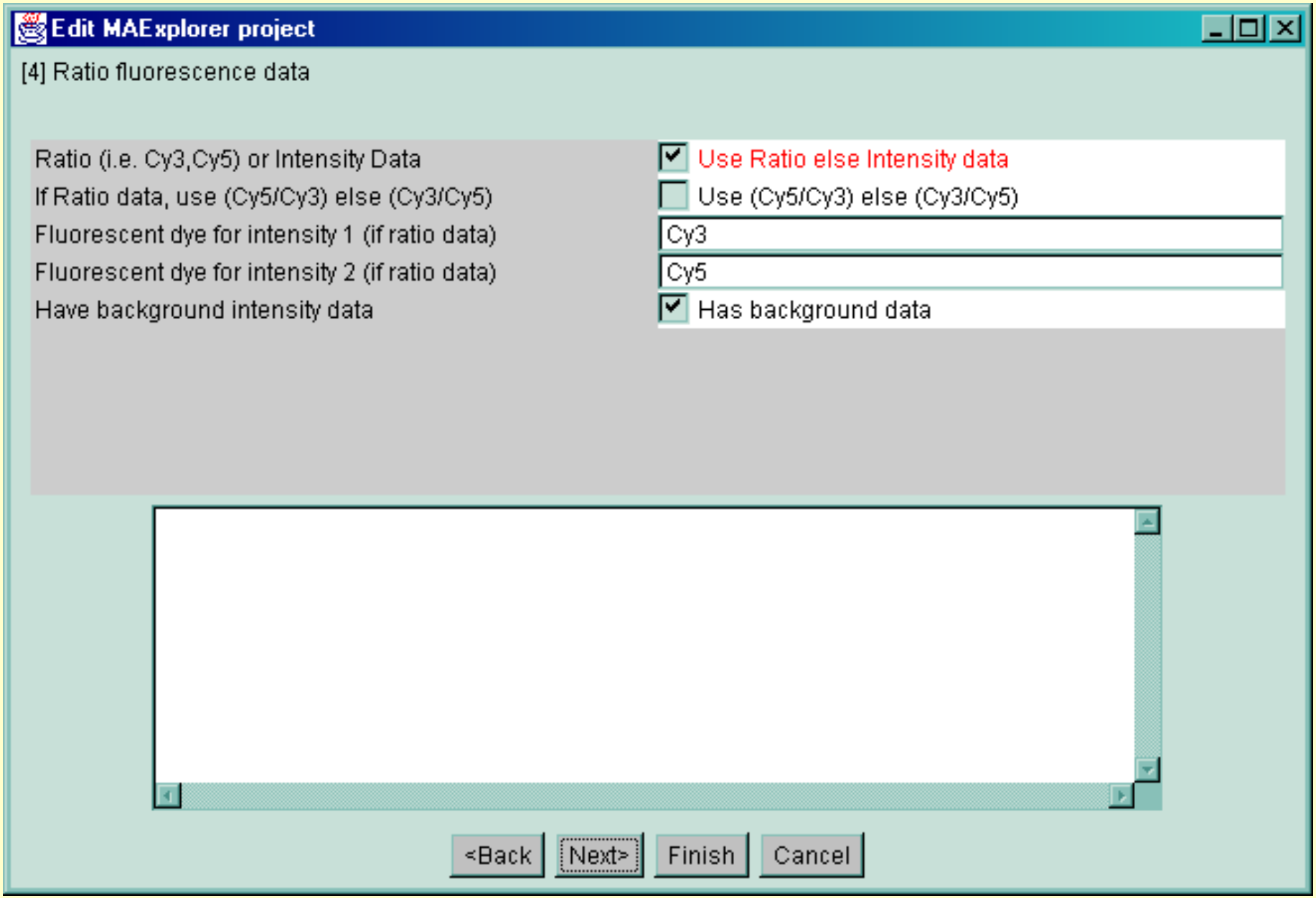
# 5.2 Edit Layout 'Wizard' - Grid Geometry. Enter (Grid, Rows/Grid, Columns/Grid) Values



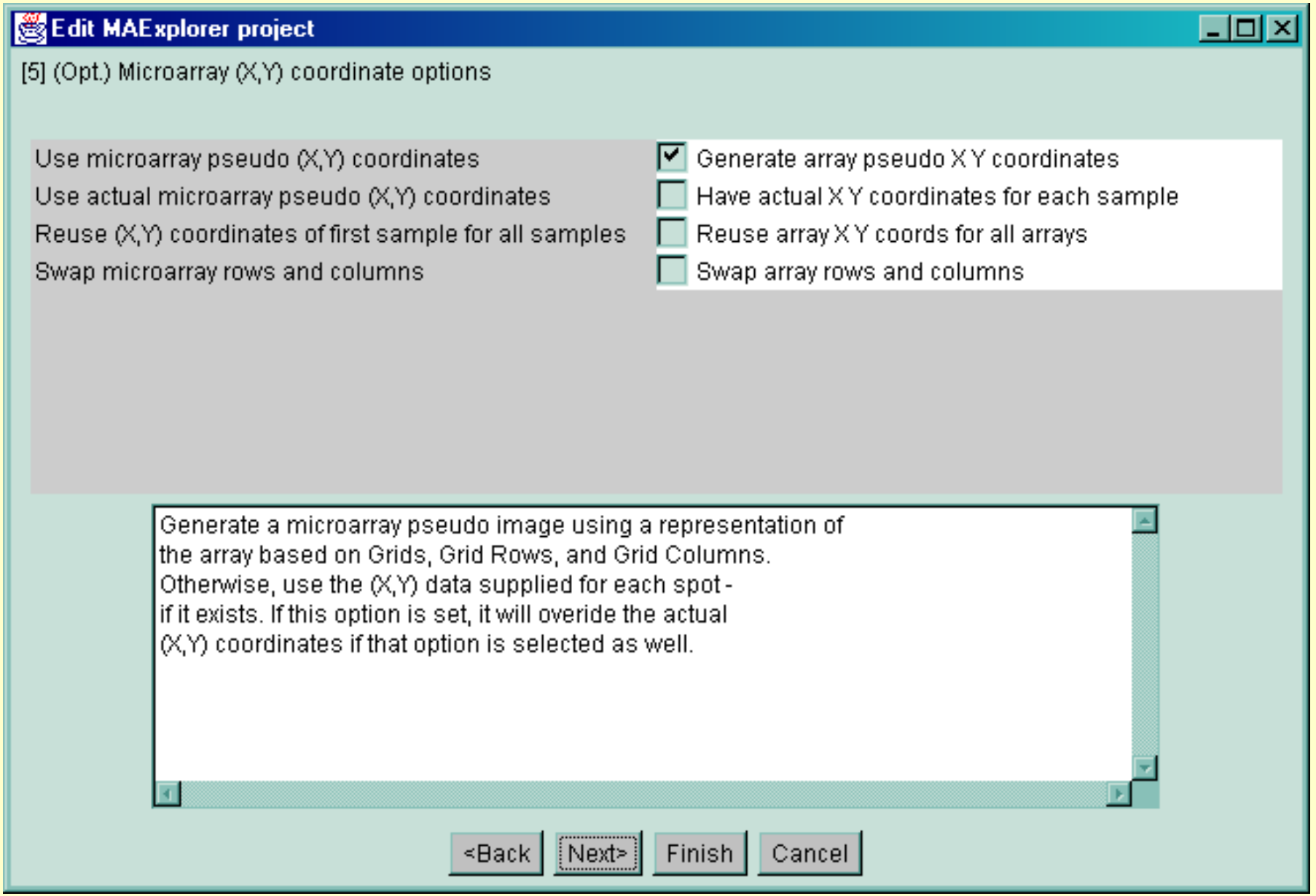
# 5.3 Edit Layout 'Wizard' Input File Row Values. Verify Row Where Field Names Are Defined



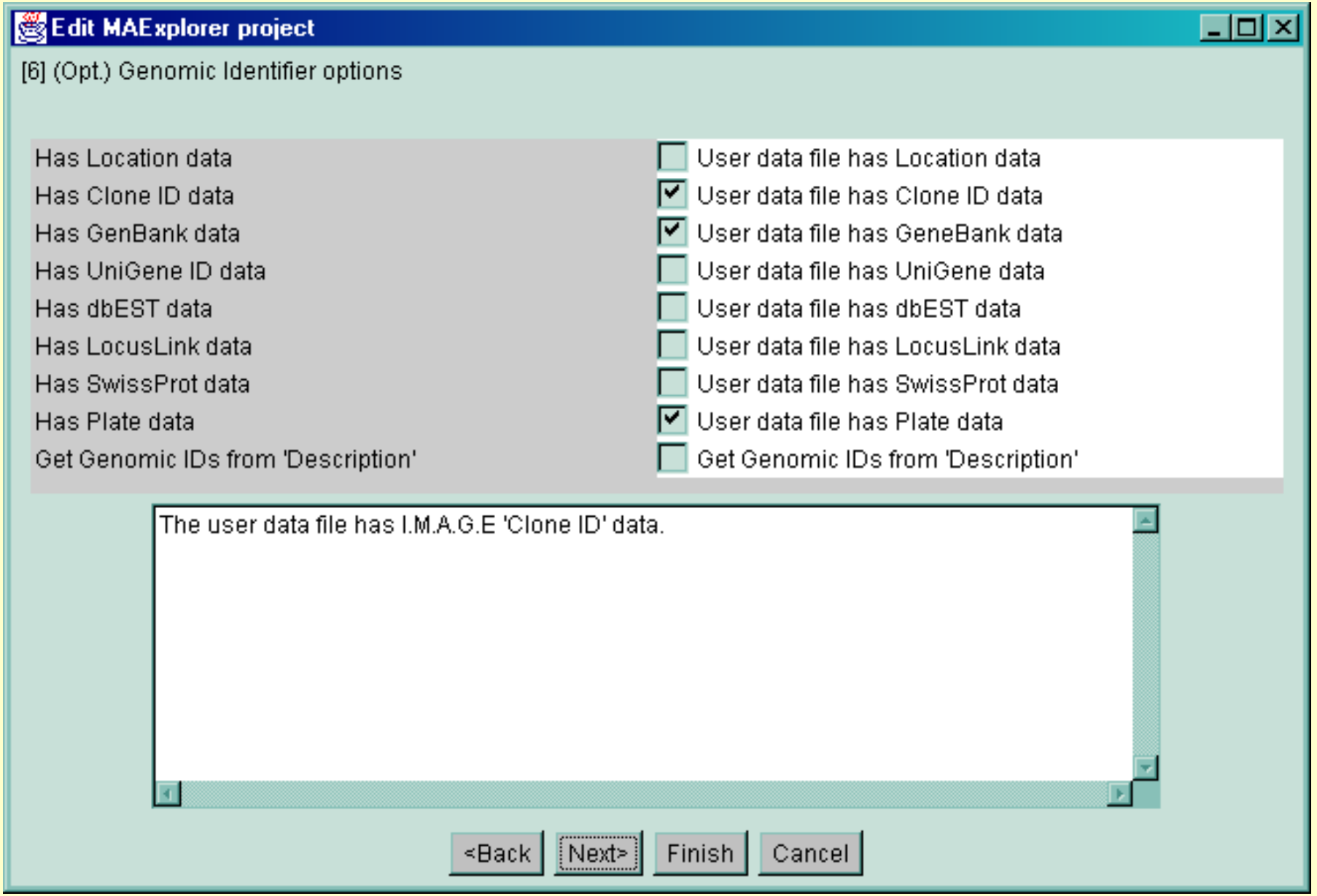
# 5.4 Edit Layout 'Wizard' Ratio or Intensity Values



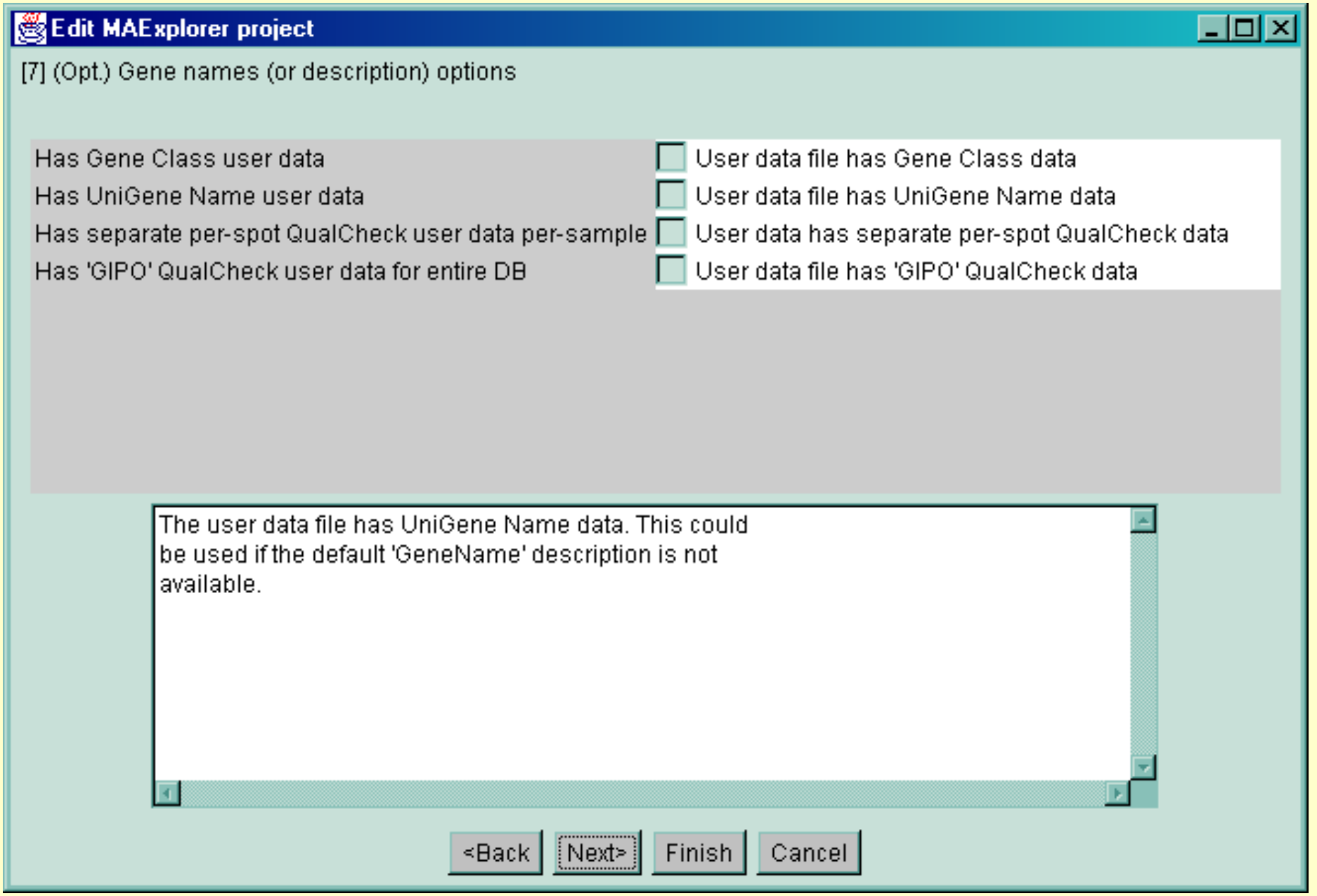
# 5.5 Edit Layout 'Wizard' optional (X,Y) Coordinate Values



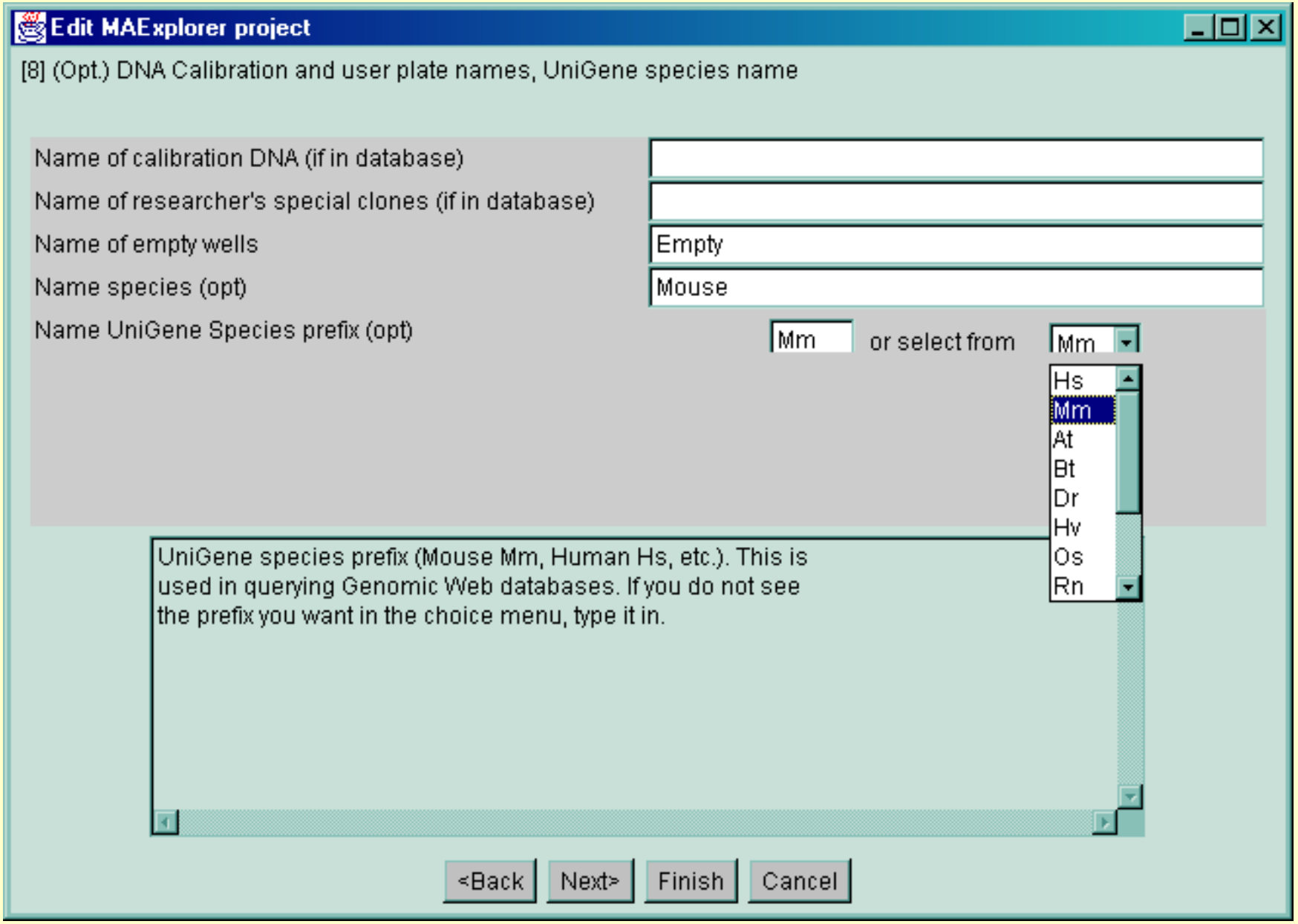
# 5.6 Edit Layout ‘Wizard’ Genomic ID Values



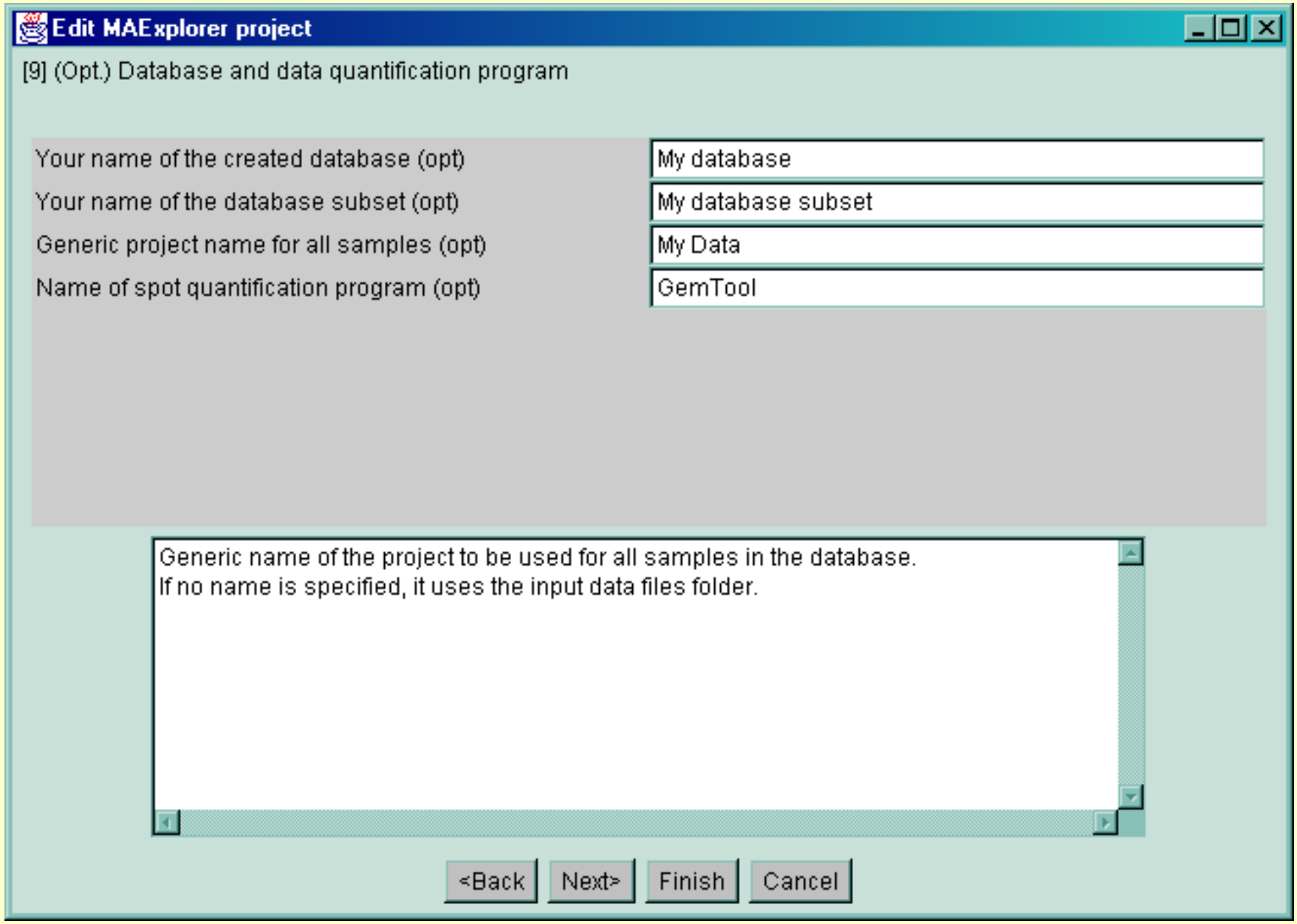
# 5.7 Edit Layout ‘Wizard’ Gene Names Description



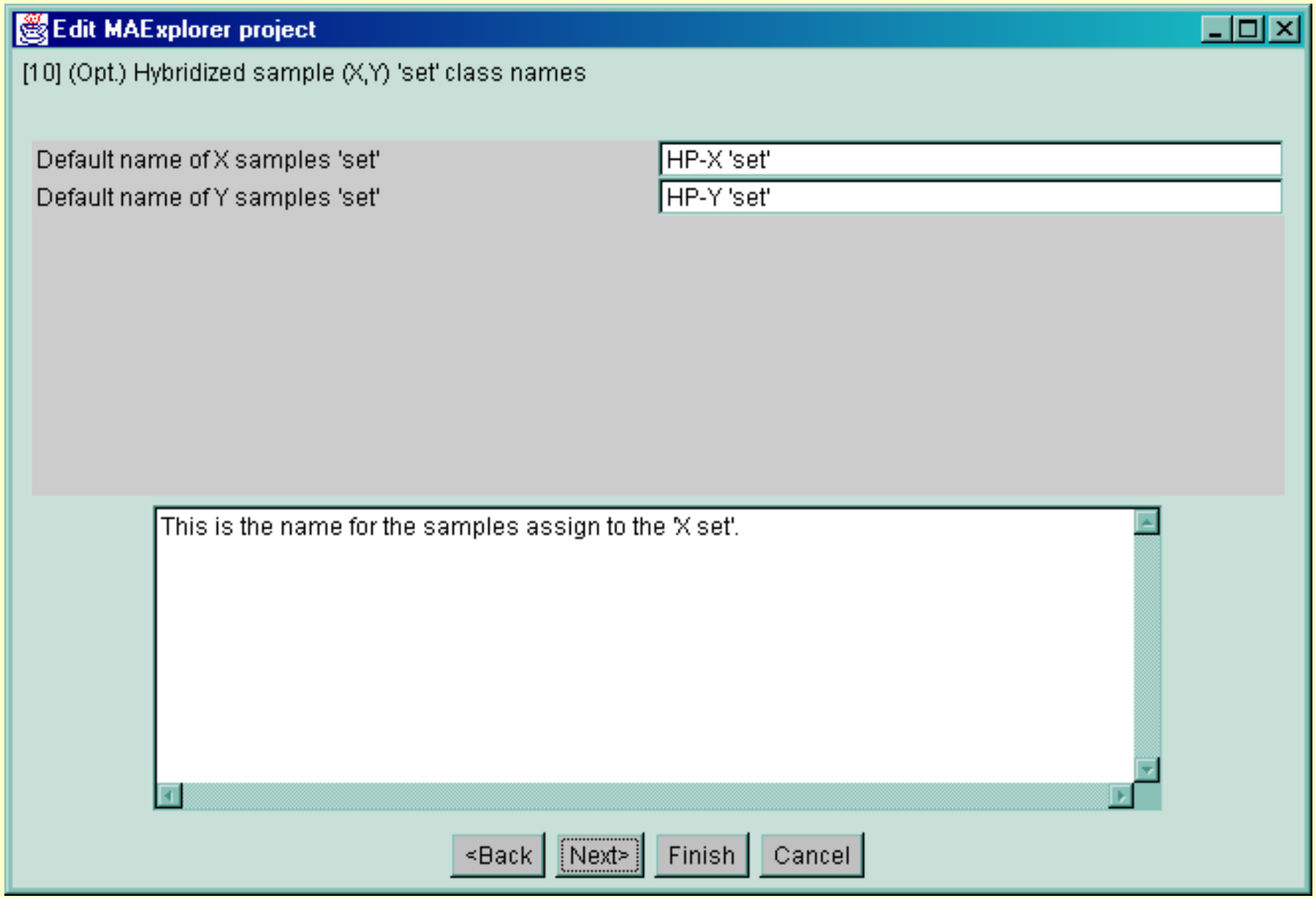
# 5.8 Edit Layout 'Wizard' Calibration Values. Define UniGene Species prefix



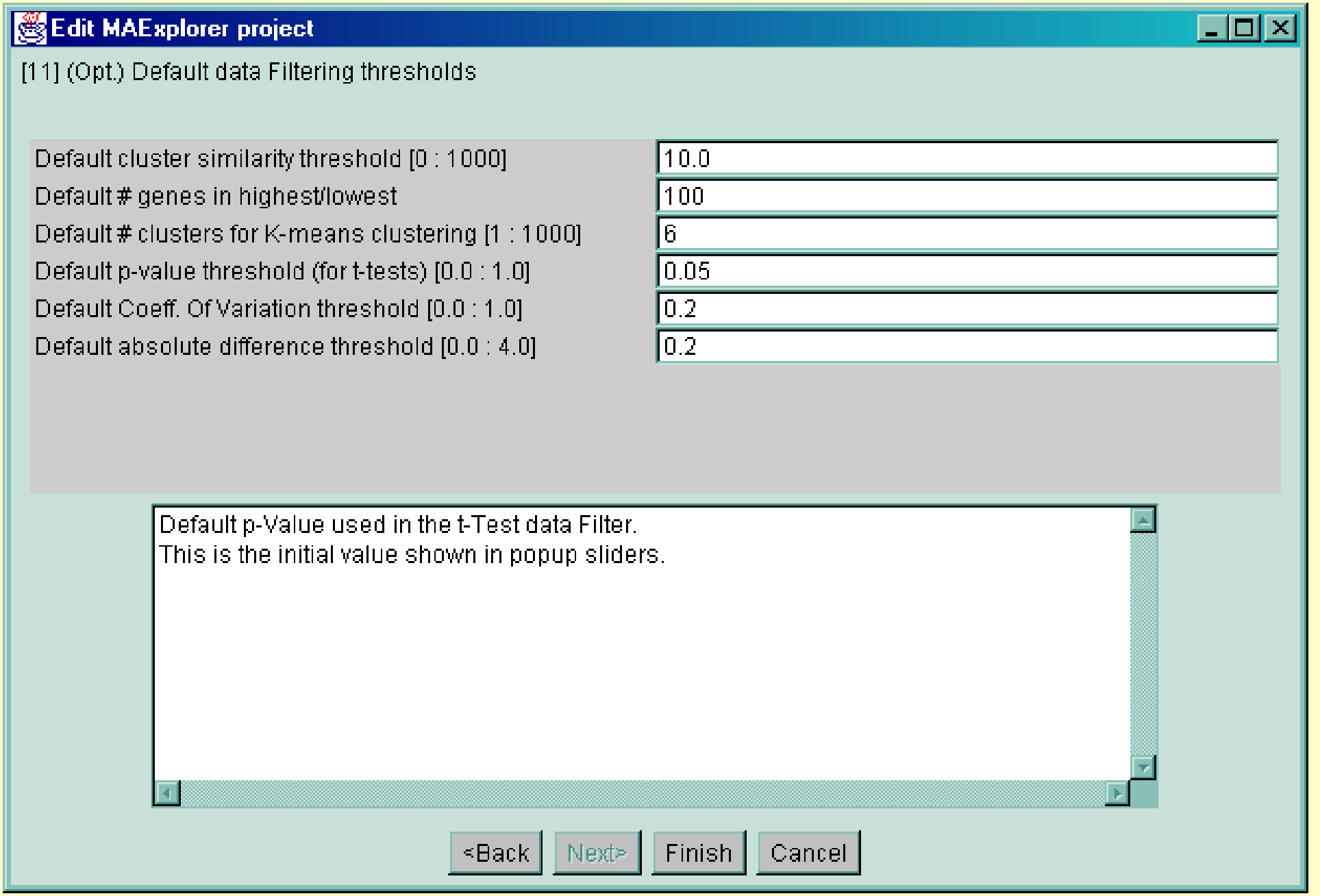
# 5.9 Edit Layout 'Wizard' Database Name Values. Define Optional Names for Database



# 5.10 Edit Layout ‘Wizard’ HP-X,-Y Class Names



# 5.11 Edit Layout 'Wizard' Default Thresholds

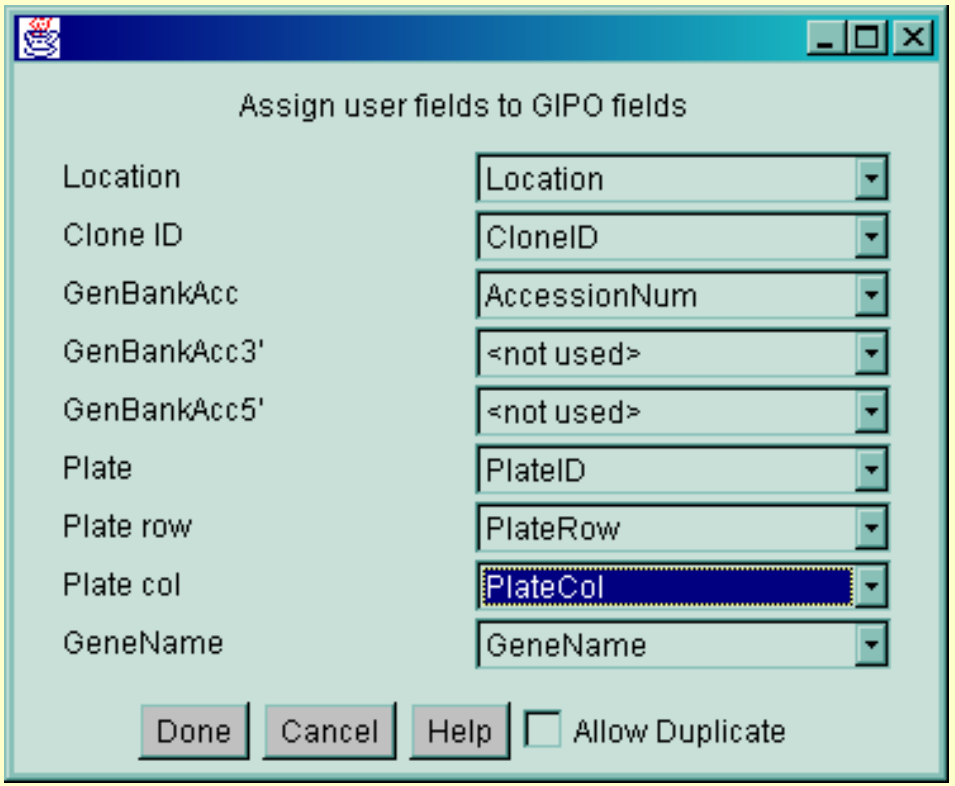


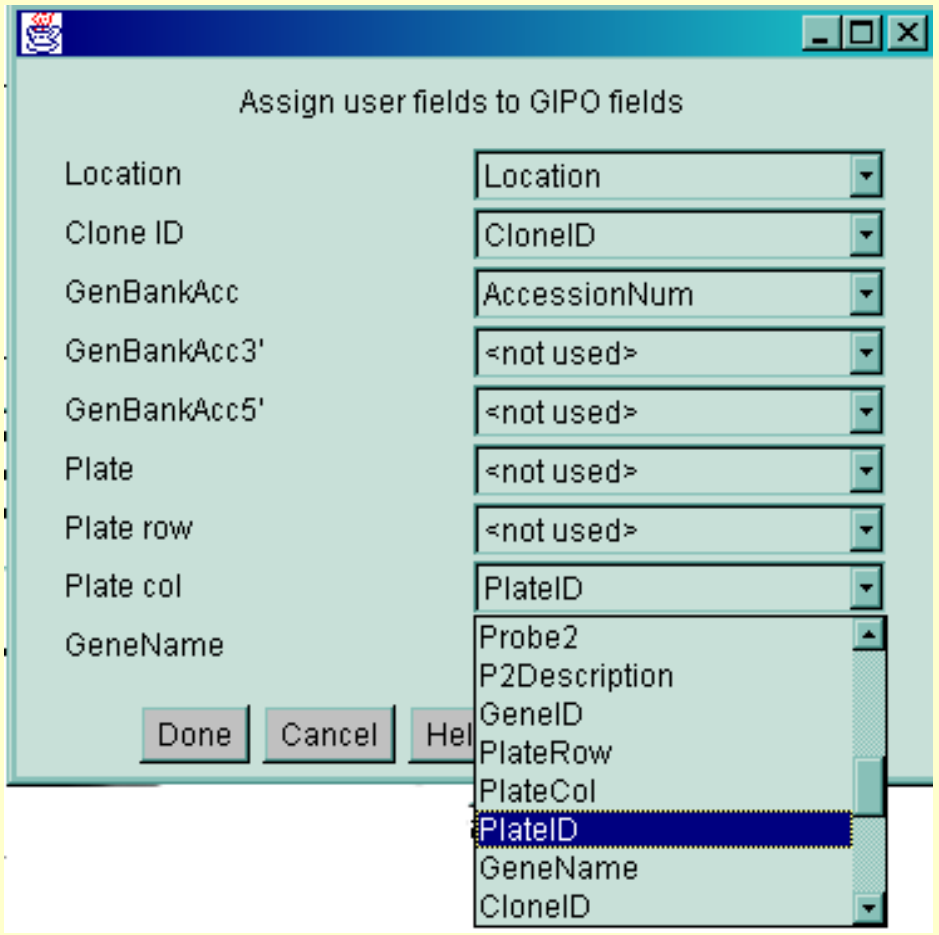
# 6. Other Options - Assigning User Data Fields to MAExplorer Fields

- GIPO (Gene In Plate Order or "array print" table) - assigns genes to positions on the array as well as GeneBank ID, Clone ID, LocusID (if available), Gene Name, etc.

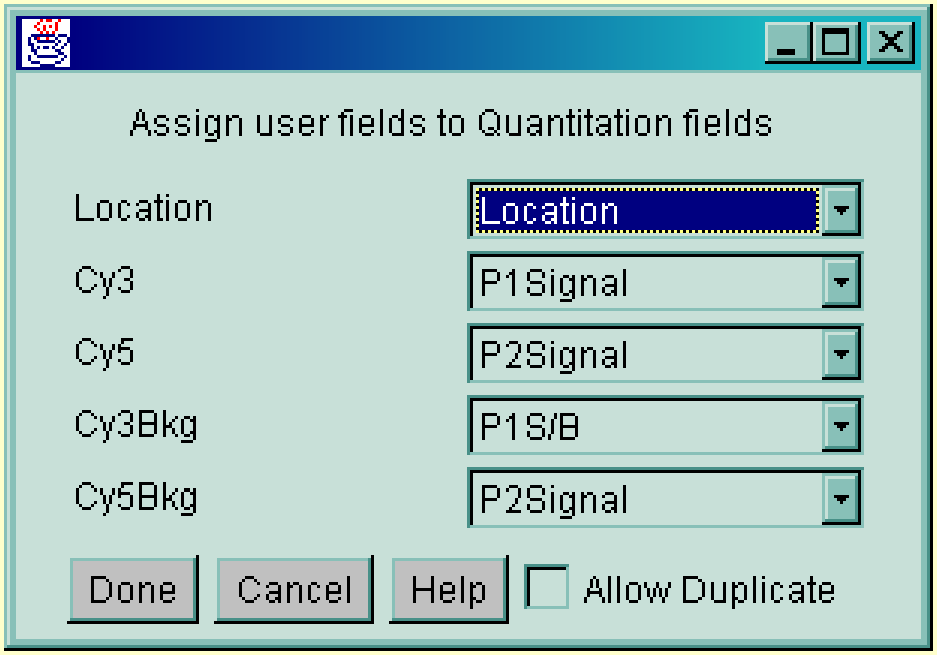
- Quant data - assigns names of quantified data in the user file to MAExplorer data (e.g. Cy3 intensity to RawIntensity1, Cy5 to RawIntensity2, etc).

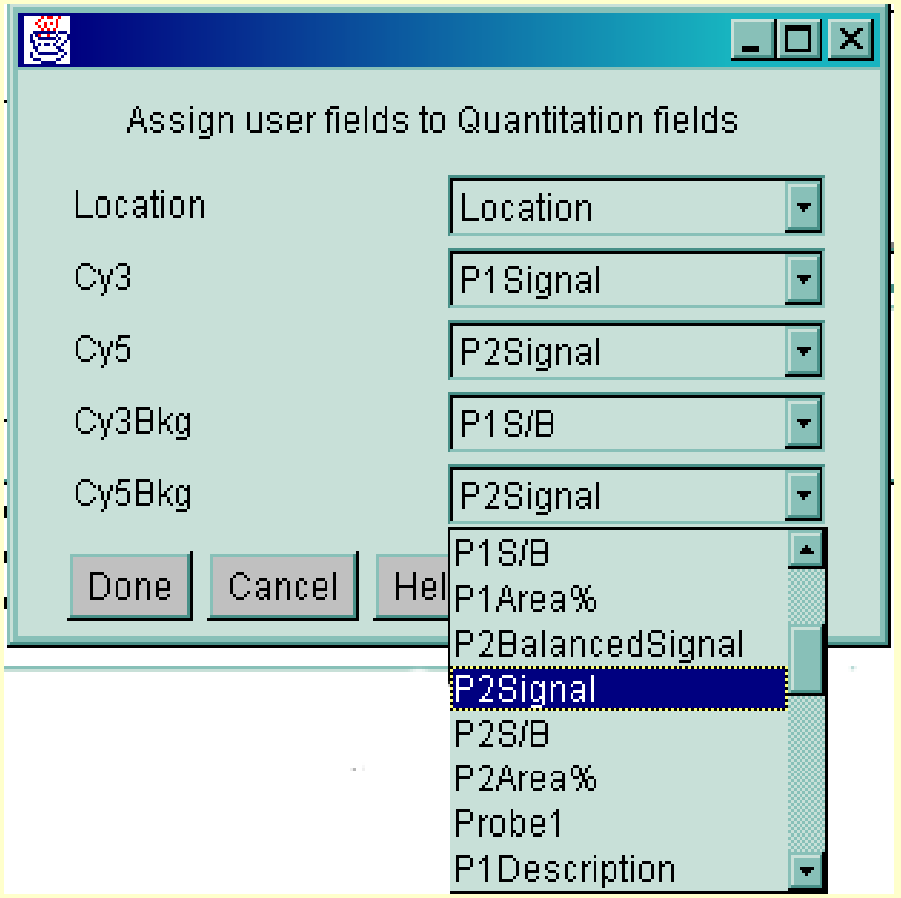
# 6.1 “Assign user fields to GIPO fields”



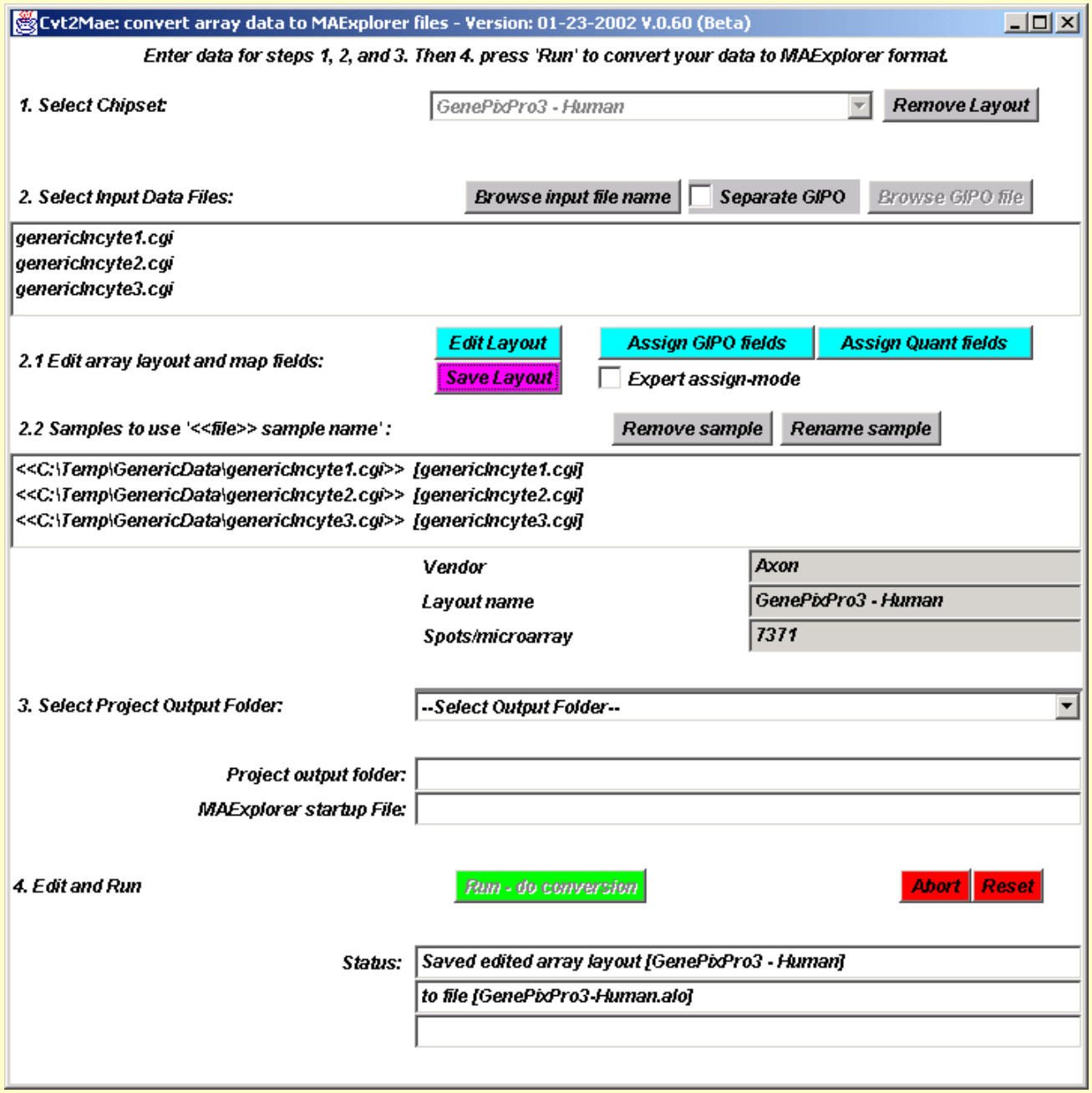



# 6.2 “Assign user fields to GIPO fields”

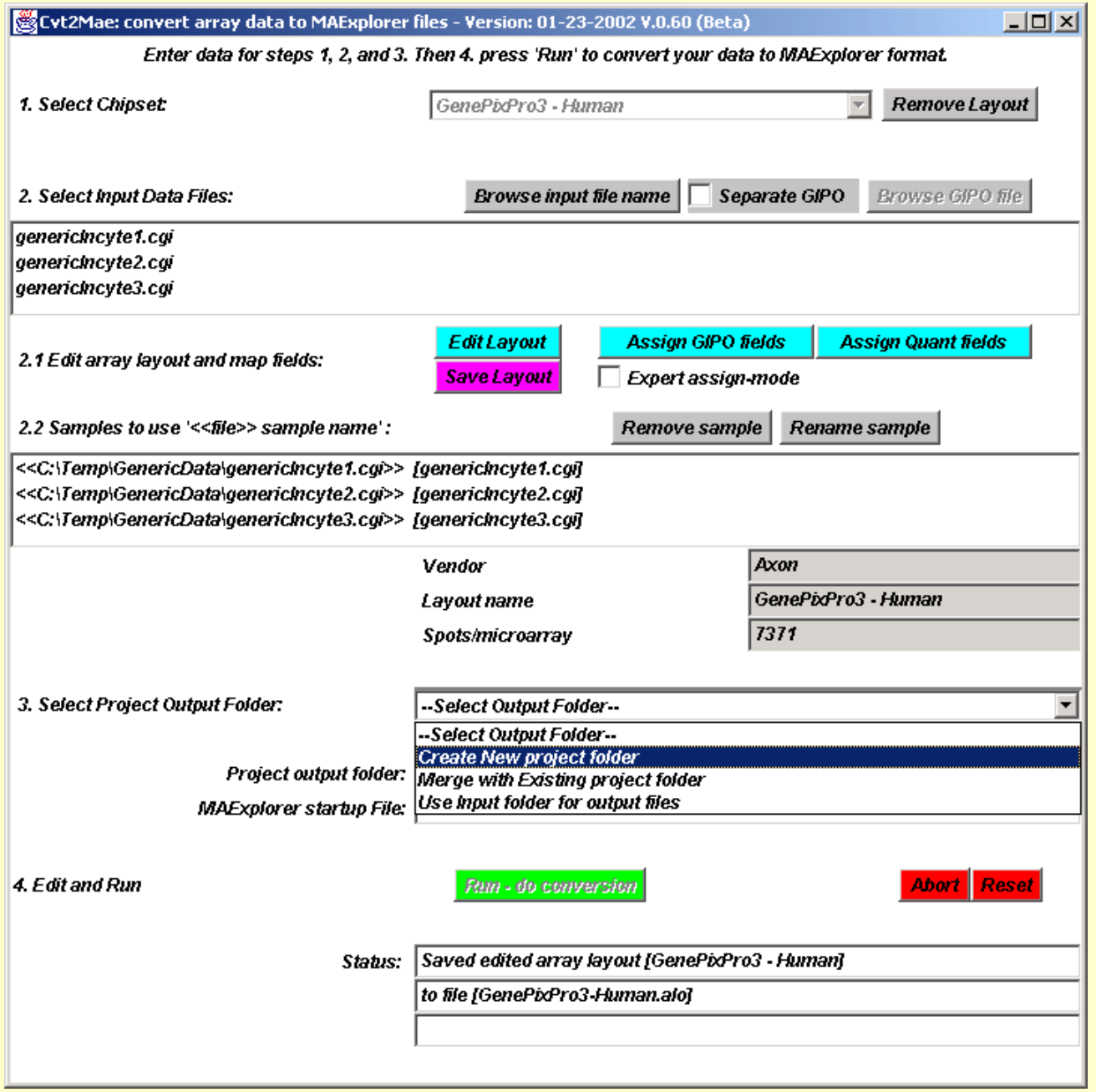


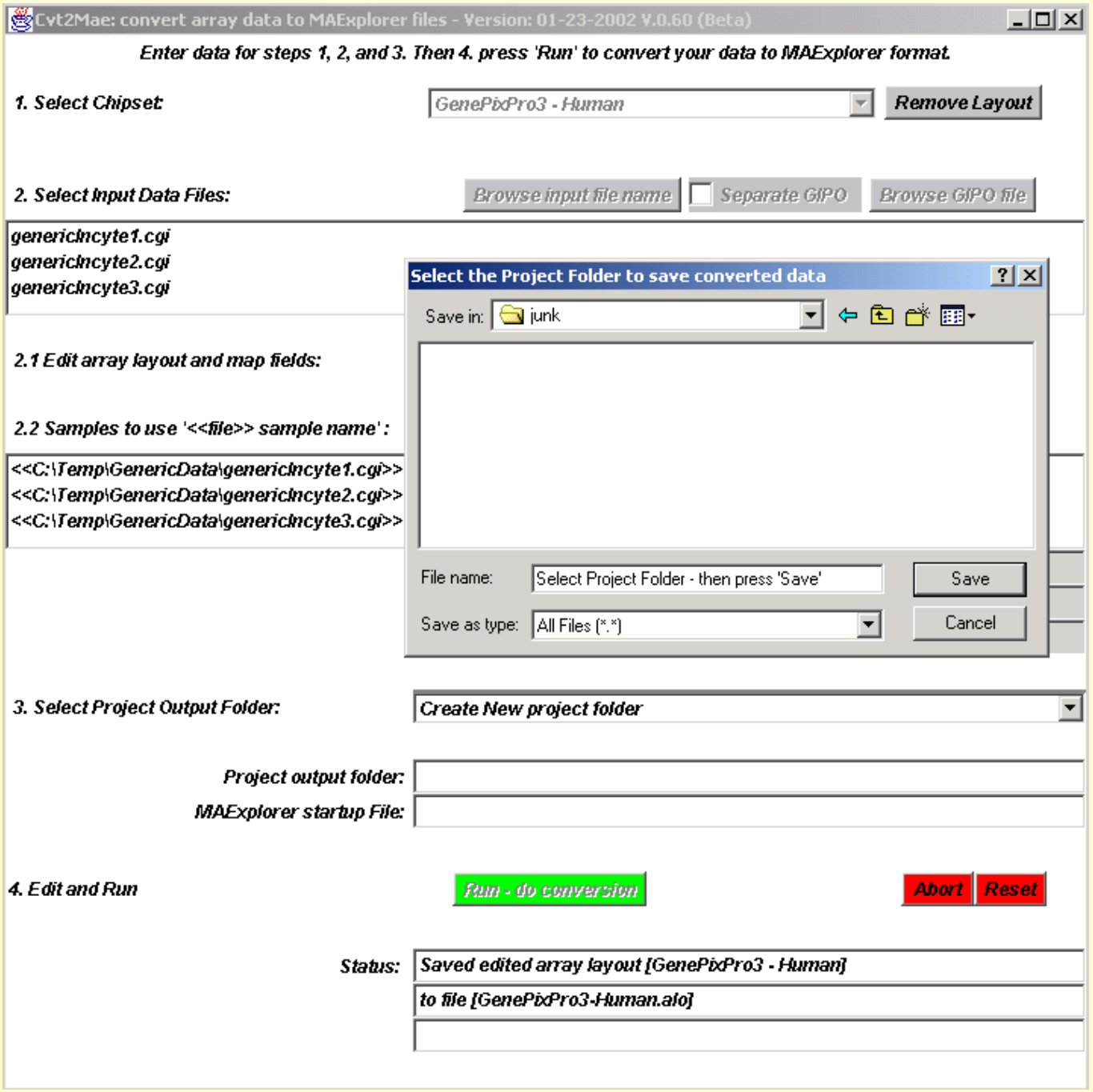
# 7. Optional "Save Layout" to Array Layout Database After Edit Layout and Assign fields



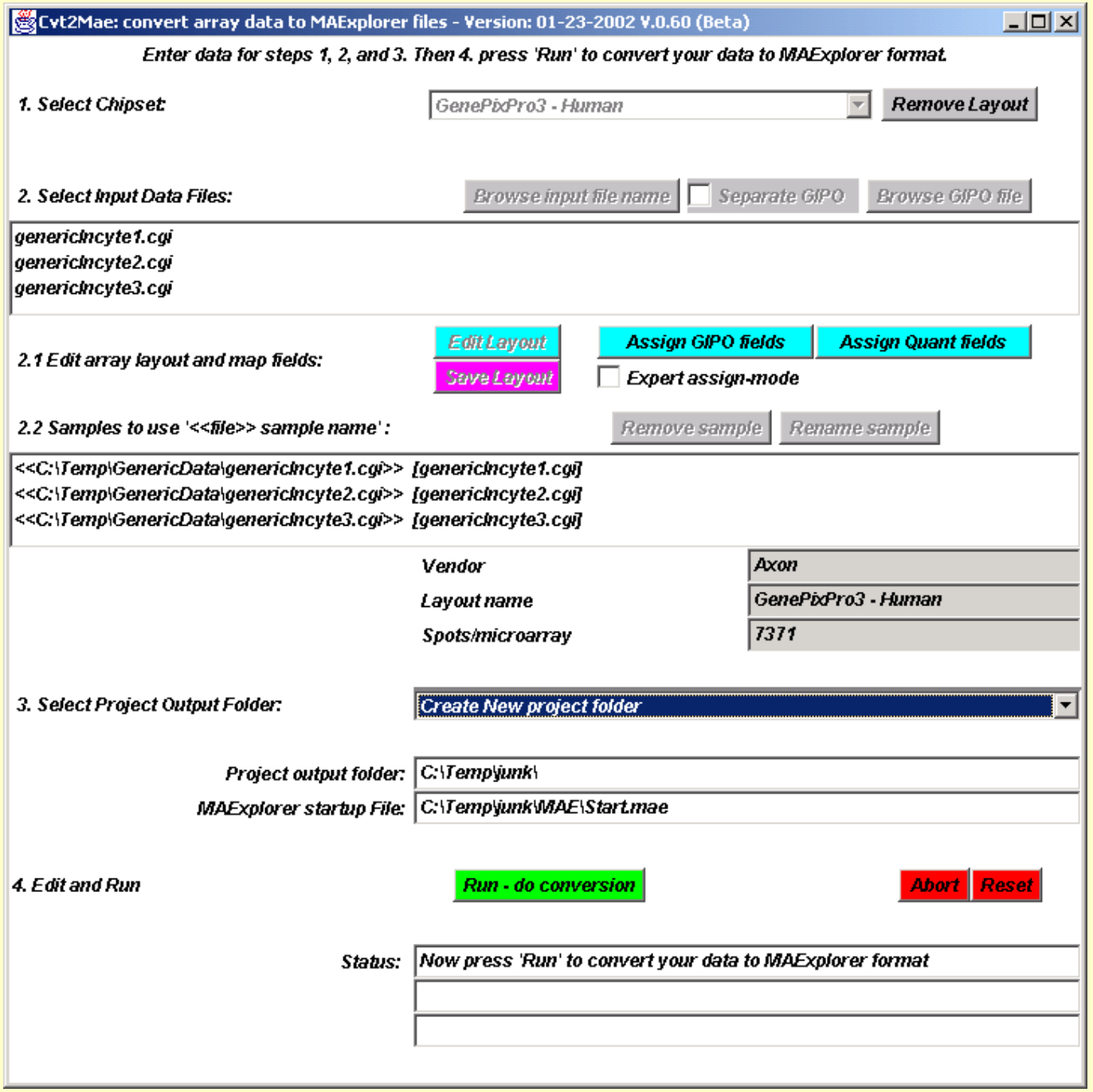
# 8. Specifying “Create new project folder” Option Where Generated Database Will Be Saved



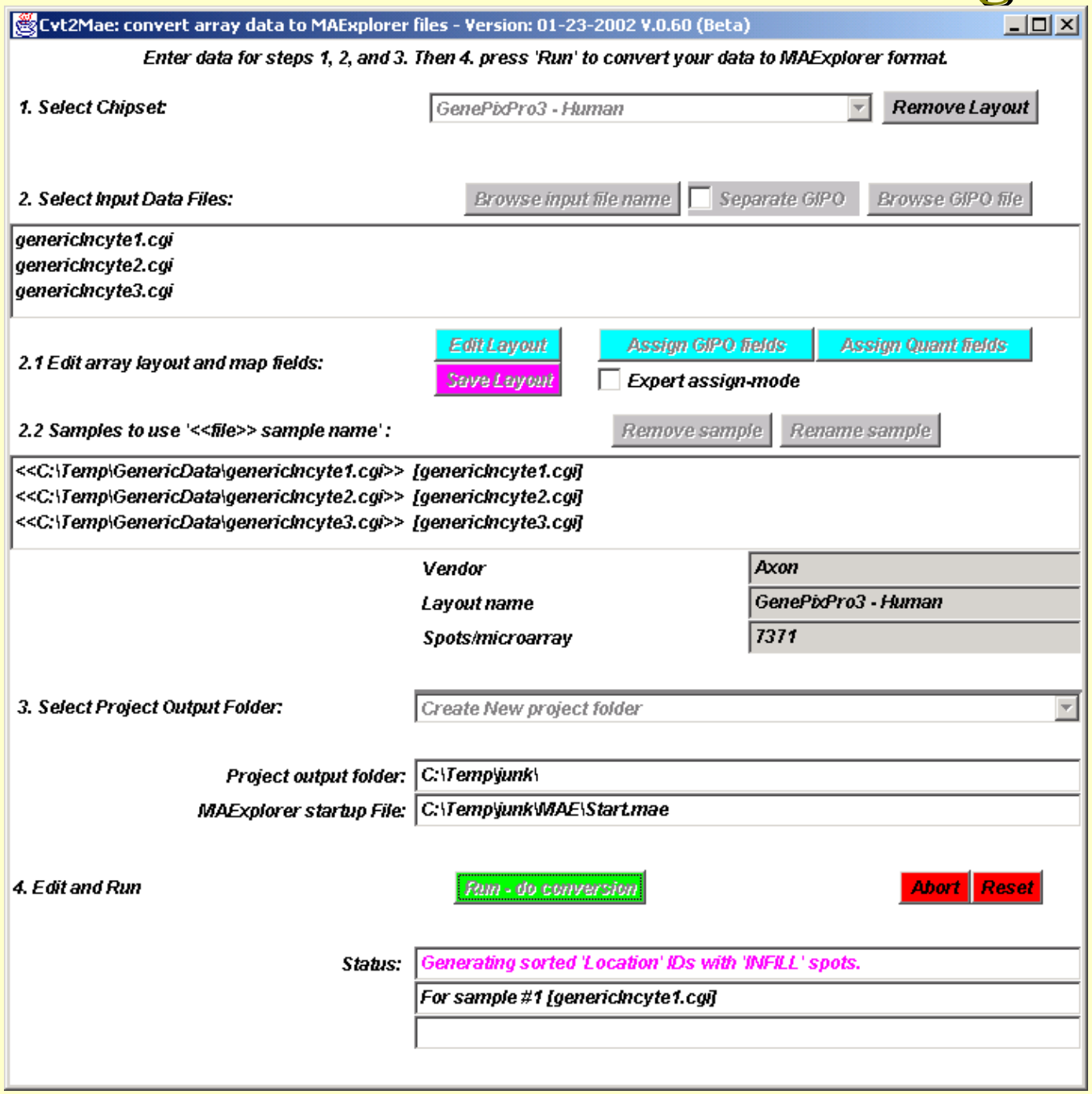
# 8.1 Specifying New "Project Output Folder"



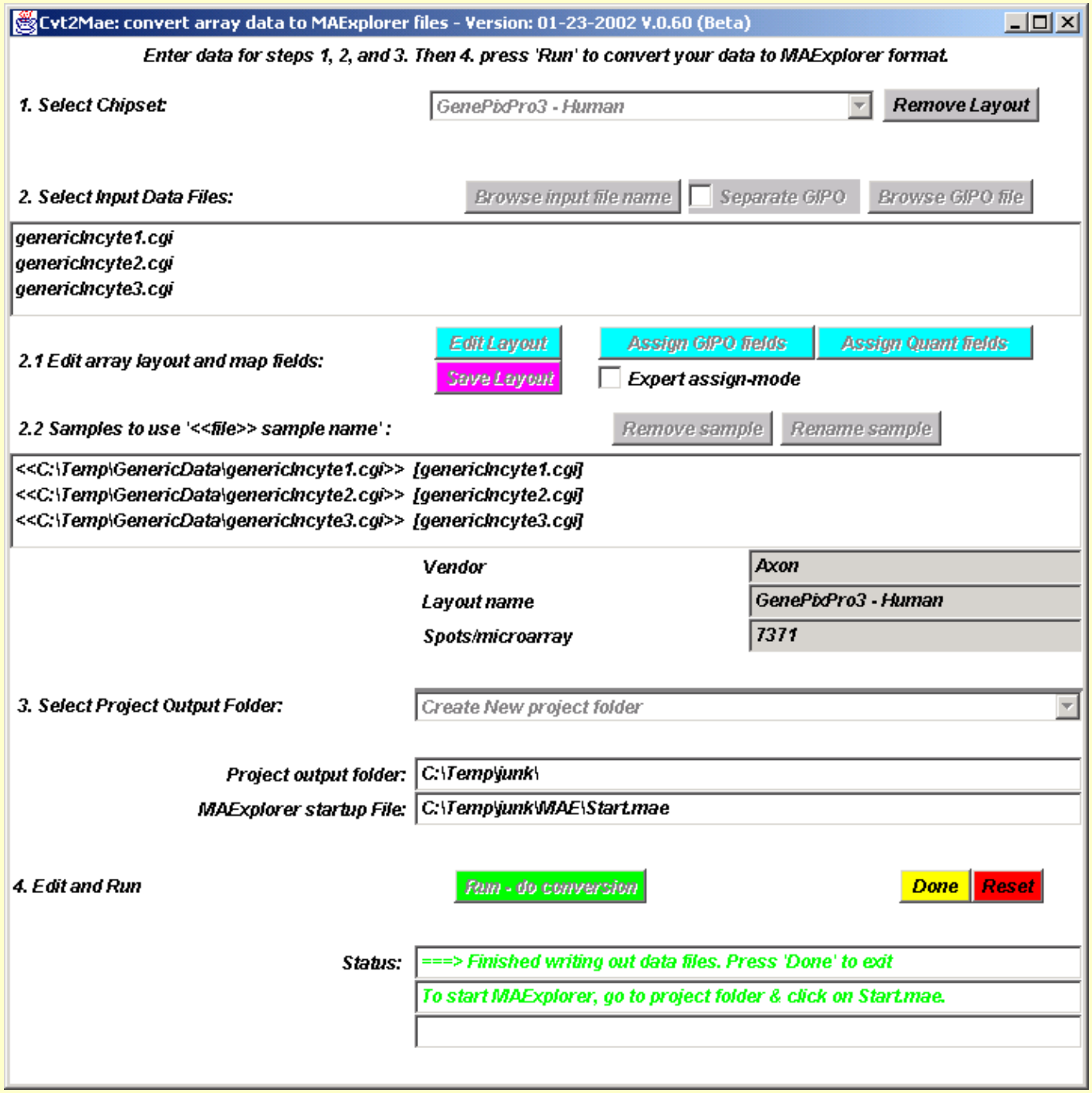
# 8.2 “Project Output Folder” & MAE startup file



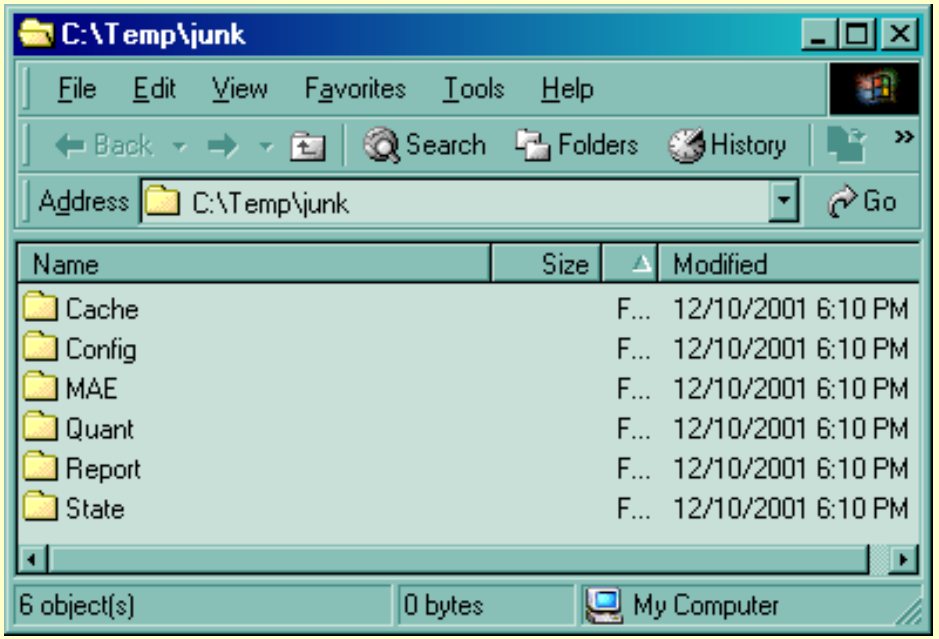
# 9. Conversion in Process After Pressing "RUN"

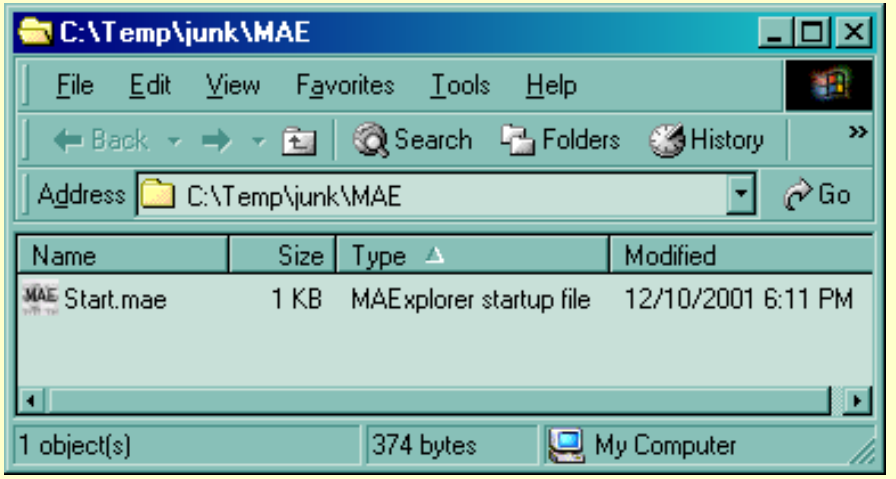


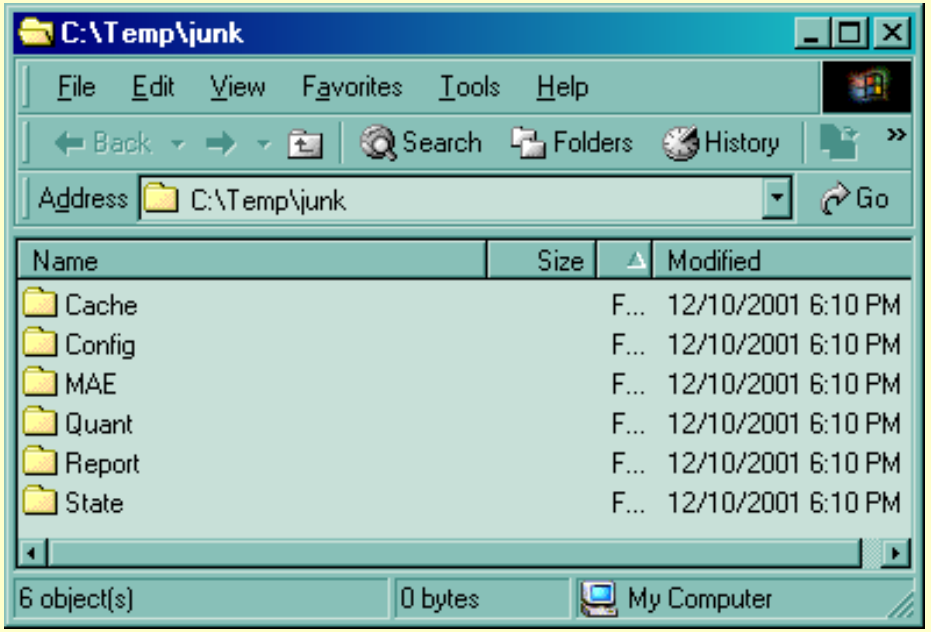
# 10. Notification that Conversion is Finished

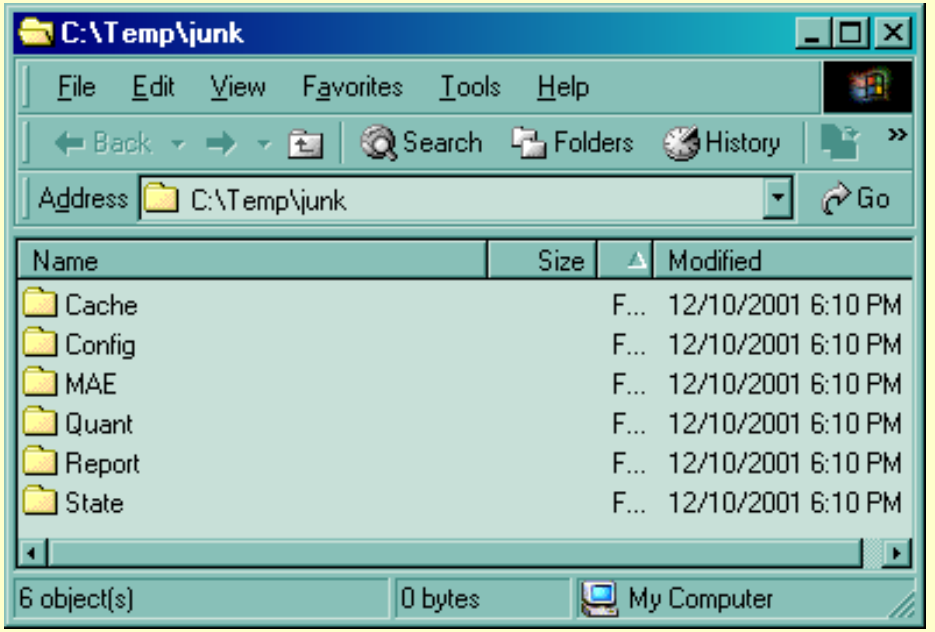


# 11. MAExplorer Data Created By Cvt2Mae



# 12. Running MAExplorer on the Converted Data