```
/*****************************
                      Semana 3: Problem Set 2
                      Universidad de San Andrés
                          Economía Aplicada
                                                      2022
    ********************
* Source: https://www.aeaweb.org/articles?id=10.1257/app.20200204
/**********************************
Este archivo sigue la siguiente estructura:
0) Set up environment and globals
1) Regressions
* 0) Set up environment
                          // Set Version number for backward compatibility
      version 16
      set more off
                          // Disable partitioned output
                           // Start with a clean slate
// Line size limit to make output more readable
      clear all
      set linesize 80
gl main "C:\Users\Anzony\Documents\GitHub\Applied_Econometrics\PS2"
gl input "$main/input"
gl output "$main/output"
* Open data set
use "$input/measures.dta", clear
* Global with control variables
global covs_eva "male i.eva_fu"
global covs_ent "male i.ent_fu"
* 1) Regressions
*-----*
* PANEL A (Child's cognitive skills at follow up)
       local bayley "b_tot_cog b_tot_lr b_tot_le b_tot_mf"
      local i = 1
       foreach y of local bayley{
       local append append
       if "'y'"=="b tot cog" local append replace
             cap drop V*
reg `y'1_st treat `y'0_st $covs_eva , cluster(cod_dane)
est sto panel_A_`i'
local i = `i' + 1
              estadd scalar N1 = e(N)
       }
```

estadd scalar N1 = $e(\overline{N})$ est sto panel D_`i' local i = `i' + 1

}

```
// aux reg for extra column
        reg home name1 st
        estadd scalar \overline{N}1 = .
        est sto panel D 6
******************
* Replicated Table
************
                * Making table for mortality
                #delimit ;
                       global note "Notes: All scores have been internally standardiz
> ed nonparametrically
                                        for age and are expressed in standard deviatio
> n
                                        units (see online Appendix B for details about
                                        the measures and the standardization procedure
> ).
                                       Measures followed by (-) have been reversed so
> that a higher
                                        score refers to better behavior. The effects r
> elating
                                        to the latent factors are in log points. Coeff
> icients and
                                        standard errors clustered at the municipality
> level (in
                                        parentheses) are from a regression of the depe
> ndent variable
                                        measured at follow-up on an indicator for whet
> her
                                        the child received any psychosocial stimulatio
> n and controlling
                                        for the child's sex, tester effects, and basel
> ine
                                        level of the outcome.";
                #delimit cr
                # delimit ;
                        esttab panel A 1 panel A 2 panel A 3
                        panel_A_4 panel_A_5 panel_A_6 using "${output}/table2 replicat
> ion.tex", replace
                        cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$8.3f)
> 6.3f)))
                        starlevels(* 0.10 ** 0.05 *** 0.01)
                        s(N1 , label( "N" ) fmt(%9.0gc) )
                        collabels (none) nostar noobs nonote
                        nonumbers eqlabels ( none )
                        nonote
                       keep( treat )
                        varlabels( treat "Treatment" )
                        mtitle(
                                "\shortstack{ \\ Bayley: \\ Cognitive}"
"\shortstack{ \\ Bayley: \\ Receptive language}"
                                "\shortstack{ \\ Bayley: \\ Expressive language}"
"\shortstack{ \\ Bayley: \\ Fine motor}"
                                "\shortstack{ \\ MacArthur: \\Words the child can say
> }"
                                "\shortstack{ \\ MacArthur: \\ Complex phrases \\ the
> child can say}")
                        mgroups( "\underline{ Panel A.} \textbf{ Child's cognitive ski
> lls at follow-up }"
, pattern( 1 0 0 0 0 0
> ) prefix(\multicolumn{@span}{c}{) suffix(}) span end(\hline) )
                        prehead("\begin{table} \small \centering
                                \protect \captionsetup{justification=centering}
```

```
\caption{\label{tab:table1} Treatment Impacts on Raw M
> easures and Latent Factors }"
                                   "\noindent\resizebox{\textwidth}{!}{ \begin{threepartt}
> able}"
                                   "\begin{tabular}{lcccccc}" \toprule)
                          posthead(\hline) prefoot(\midrule) postfoot(\midrule);
                          esttab panel_B_1 panel_B_2 panel_B_3
                          panel B 4 panel B 5 panel B 6 using "${output}/table2 replica
> tion.tex", append
                          cells(b(label(coef.) star fmt(%8.3f) ) se(label((z)) par fmt(%
> 6.3f)))
                          starlevels(* 0.10 ** 0.05 *** 0.01)
                          s(N1 , label( "N" ) fmt(%9.0gc) )
                          collabels ( none )
                         nostar noobs nonote
                          nonumbers eqlabels ( none )
                          keep( treat )
                         mgroups( "
                                   \underline{ Panel B.}
                                   \textbf{ Child's socio-emotional skills at follow-up }
> "
                                   , pattern( 1 0 0 0 0 0 )
                                   prefix(\multicolumn{@span}{c}{) suffix(})
                                   span end(\hline))
                          mtitle(
                                  "\shortstack{ \\ ICQ: \\Difficult (-)}"
"\shortstack{ \\ ICQ: \\ Unsociable (-)}"
"\shortstack{ \\ ICQ: \\ Unstoppable (-)}"
                                   "\shortstack{ \\ ECBQ: \\ Inhibitory control}"
"\shortstack{ \\ ECBQ: \\ Attentional focusing}")
                         nonote
                         varlabels( treat "Treatment" )
prehead( "" )
                           posthead( \hline )
                           prefoot(\midrule)
                           postfoot("")
                           delim("&") nonumbers
                          esttab panel_C_1 panel_C_2 panel_C_3 panel_C_4 panel_C_5 panel_C_6 using "${output}/table2_replica
> tion.tex", append
                          cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$
> 6.3f)))
                          starlevels(* 0.10 ** 0.05 *** 0.01)
                          s(N1 , label("N" ) fmt(%9.0gc) )
                          collabels(none) nostar noobs
                         nonumbers eqlabels( none )
mgroups( "
                                   \underline{ Panel C.}
                                   \textbf{ Material investments at follow-up }"
                                   , pattern( 1 0 0 0 0 0 )
                                   prefix(\multicolumn{@span}{c}{) suffix(})
                                   span end(\hline) )
                          mtitle(
                                   > erials}"
                                   "\shortstack{ \\ FCI: \\ Number of coloring \\ and dra
> wing books}"
                                   "\shortstack{ \\ FCI: \\ Number of toys \\ to learn mo
> vement}"
                                   "\shortstack{ \\ FCI: \\ Number of toys \\ to learn sh
> apes}"
                                   "\shortstack{ \\ FCI: \\ Number of \\ shop-bought toys
> }"
                                  "")
                          nonote keep( treat )
                          varlabels( treat "Treatment" )
```

```
prehead( \hline )
                       posthead( \hline )
                       prefoot(\midrule)
                       postfoot("")
                       delim("&") nonumbers
                      esttab panel D 1 panel D 2 panel D 3
                      panel_D_4 panel_D_5 panel_D_6 using "${output}/table2_replica
> tion.tex", append
                      cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$
> 6.3f)))
                      starlevels(* 0.10 ** 0.05 *** 0.01)
                      s(N1 , label("N") fmt(%9.0gc))
                      collabels (none) nostar noobs nonote
                      nonumbers eqlabels( none ) keep( treat )
                      mgroups("
                              \underline{ Panel D.}
                             \textbf{ Time investments at follow-up }"
                             , pattern(10000)
                             prefix(\multicolumn{@span}{c}{) suffix(})
                             span end(\hline) )
                      mtitle(
                             > ivities \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times told \\ a sto
> ry to child \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times read \\ to ch
> ild \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times \\ played wit
> h toys \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times \\ named thin
> gs to child \\ in last 3 days}"
                             "")
                      varlabels( treat "Treatment" )
                       prehead( \hline )
                       posthead( \hline )
                       prefoot(\midrule)
                       delim("&") nonumbers
                       postfoot( \hline \end{tabular}
                              \begin{tablenotes}
                             \begin{footnotesize}
                             ${note}
                              \end{footnotesize}
                             "\end{tablenotes} \end{threeparttable} } \end{table}")
> ;
               #delimit cr
* 2) Modification
*-----*
* 2) P-Values Correction
* PANEL A (Child's cognitive skills at follow up)
       * Define number of hypothesis
       scalar hyp = 6
       * Define level of significance
       scalar signif = 0.05
```

> ==0

```
local bayley "b tot cog b tot lr b tot le b tot mf"
local i = 1
local group_regressions = ""
mat p values = J(hyp,1,.)
foreach y of local bayley{
local append append
if "`y'"=="b_tot_cog" local append replace cap drop V*
         reg `y'1_st treat `y'0_st $covs_eva , cluster(cod_dane) eststo panel_A_`i': test treat = 0 mat p_values[`i',1]=r(p)
          scalar p_value = r(p)
          scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
          estadd scalar N1 = e(N)
          local est_name panel_A_`i'
local group_regressions `group_regressions' `est_name'
          local i = `i' + 1
local macarthur "mac words mac phrases"
foreach y of local macarthur{
          cap drop V*
          reg `y'l st treat mac words0 st $covs ent , cluster(cod dane)
          eststo panel A `i': test treat = 0
mat p_values[`i',1]=r(p)
          scalar p value = r(p)
          scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
          estadd scalar N1 = e(N)
          local est_name panel_A_`i'
local group_regressions `group_regressions' `est_name'
          local i = `i' + 1
}
          // Modification of p-vals
          preserve
                    // * Define number of hypothesis
                    // scalar hyp = hyp
                    // * Define level of significance
                    // scalar signif = signif
                    * Holm Correction
                              // Bring the p_{values} matrix as column dta
                              symat p values
                              // Identify the original variable outcome
                              gen outcome_order_var = _n
                              // Sort values
                              sort p_{values1} // Indicate the rank of the variable
                              gen rank = _n
// Identify if its significan or note
                              gen alpha corr = signif/(hyp+1-rank)
                             gen significant_Holm = (p_values1<alpha_corr)
replace significant_Holm = 0 if significant_Holm[_n-1]</pre>
                              // Sort again based on outcome order
                              sort outcome_order_var
                              // Export the result as a matrix
                              mkmat significant Holm, matrix(holm cor)
                              keep p_values1 outcome_order_var
```

```
* Benjamini et al.
                                  rename (p values1 outcome order var) (pval outcome)
                                  quietly sum pval
                                  local total pvals = r(N)
                                  * Sort the p-values in ascending order and generate a
> variable that codes each p-value's rank
                                  quietly gen int original_sorting_order = _n
                                  quietly sort pval
                                  quietly gen int rank = _n if pval~=.
                                  * Set the initial counter to 1
                                  local qval = 1
                                  * Generate the variable that will contain the BKY (200
> 6) sharpened q-values
                                  gen bky06 gval = 1 if pval~=.
                                  * Set up a loop that begins by checking which hypothes
> es are rejected at q = 1.000, then checks which hypotheses are rejected at q = 0.999
> , then checks which hypotheses are rejected at q = 0.998, etc. The loop ends by che
> cking which hypotheses are rejected at q = 0.001.
                                  local totalpvals = ${totalpvals}
                                 local qval = 1
                                         .
qval' > 0 {
                                  while
                                          * First Stage
                                          * Generate the adjusted first stage q level we
> are testing: q' = q/1+q
                                          local qval_adj = `qval'/(1+`qval')
                                          * Generate value q'*r/M
                                          gen fdr_temp1 = `qval_adj'*rank/`totalpvals'
                                          * Generate binary variable checking condition
> p(r) <= q'*r/M
                                          gen reject_temp1 = (fdr_temp1>=pval) if pval~=
                                          * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                          gen reject_rank1 = reject_temp1*rank
                                          * Record t\overline{h}e rank of the \overline{l}argest p-value that
> meets above condition
                                          egen total rejected1 = max(reject rank1)
                                          * Second Stage
                                          ^{\star} Generate the second stage q level that accou
> nts for hypotheses rejected in first stage: q_2st = q'*(M/m0) local qval_2st = `qval_adj'*(`totalpvals'/(`to
> talpvals'-total rejected1[1]))
                                          * Generate value q_2st*r/M
gen fdr_temp2 = `qval_2st'*rank/`totalpvals'
                                           Generate binary variable checking condition
> p(r) \le q_2st*r/M
                                          gen reject temp2 = (fdr temp2>=pval) if pval~=
                                          * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                          gen reject_rank2 = reject_temp2*rank
                                          * Record the rank of the largest p-value that
> meets above condition
                                          egen total rejected2 = max(reject rank2)
```

```
* A p-value has been rejected at level q if it
> s rank is less than or
                                             * equal to the rank of the max p-value that me
> ets the above condition
                                             replace bky06 qval = `qval' if rank <= total r</pre>
> ejected2 & rank~=.
                                              * Reduce q by 0.001 and repeat loop
                                             drop fdr_temp* reject_temp* reject_rank* total
> rejected*
                                             local qval = `qval' - .001
                                    quietly sort original sorting order
                                    pause off
                                    mkmat bky06 qval, matrix(pval bky06)
                           scalar i = 1
                           foreach reg store of local group regressions{
                                    scalar holm cor val = holm cor[i, 1]
                                    scalar pval_bky06[i, 1]
                                    if holm cor val == 1 {
                                             est restore `reg_store'
estadd local pholm "Significant"
                                             estadd scalar bky_06 = pval_bky06_val
                                    if holm_cor_val == 0 {
                                             est restore `reg store'
                                             estadd local pholm "No Significant"
                                             estadd scalar bky_06 = pval_bky06_val
                                    }
                  restore
* PANEL B (Child's socio-emotional skills at follow up)
         * Define number of hypothesis
         scalar hyp = 5
         * Define level of significance
         scalar signif = 0.05
         local bates "bates_difficult bates_unsociable bates_unstoppable"
         local i = 1
         local group_regressions = ""
        mat p values = J(hyp,1,.)
         foreach y of local bates{
                  cap drop V*
                 reg `y'1_st treat `y'0_st $covs_ent, cl(cod_dane)
eststo panel_B_`i': test treat = 0
mat p_values[`i',1]=r(p)
scalar p_value = r(p)
                  scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
                  estadd scalar N1 = e(N)
                  local est_name panel_B_`i'
local group_regressions `group_regressions' `est_name'
                  local i = i' + 1
         }
```

> ==0

> 6) sharpened q-values

```
local roth "roth inhibit roth attention"
         foreach y of local roth{
                   cap drop V*
                  reg `y'1_st treat bates_difficult0_st $covs_ent , cluster(cod_dane) eststo panel_B_`i': test treat = 0 mat p_values[`i',1]=r(p)
                  scalar p_value = r(p)
scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
                  estadd scalar N1 = e(N)
                  local est name panel B `i'
                  local group_regressions `group_regressions' `est_name'
                  local i = `i' + 1
         reg home_name1_st
         estadd s\overline{c}alar \overline{N}1 =
         est sto panel B 6
         // Modification of p-vals
                  preserve
                            // * Define number of hypothesis
                            // scalar hyp = hyp
                            // * Define level of significance
                            // scalar signif = signif
                            * Holm Correction
                                      clear
                                      // Bring the p_values matrix as column dta
                                      svmat p_values
                                      // Identify the original variable outcome
                                      gen outcome_order_var = _n
                                      // Sort values
                                      sort p_values1
                                      // Indicate the rank of the variable
                                      gen rank = n
                                      // Identify if its significan or note
                                      gen alpha_corr = signif/(hyp+1-rank)
gen significant_Holm = (p_values1<alpha_corr)
replace significant_Holm = 0 if significant_Holm[_n-1]</pre>
                                      // Sort again based on outcome order
                                      sort outcome_order_var
                                      // Export the result as a matrix
                                      mkmat significant_Holm, matrix(holm_cor)
                                      keep p values1 outcome order var
                            * Benjamini et al.
                                      rename (p_values1 outcome_order_var) (pval outcome)
                                      quietly sum pval
                                      local total pvals = r(N)
                                      * Sort the p-values in ascending order and generate a
> variable that codes each p-value's rank
                                      quietly gen int original sorting order = n
                                      quietly sort pval
                                      quietly gen int rank = _n if pval~=.
                                      * Set the initial counter to 1
```

local qval = 1

gen bky06 qval = 1 if $pval\sim=$.

* Generate the variable that will contain the BKY (200

```
* Set up a loop that begins by checking which hypothes
> es are rejected at q = 1.000, then checks which hypotheses are rejected at q = 0.999
> , then checks which hypotheses are rejected at q = 0.998, etc. The loop ends by che
> cking which hypotheses are rejected at q = 0.001.
                                   local totalpvals = ${totalpvals}
                                   local qval = 1
while `qval' >
                                          `qval' > 0 {
                                            * First Stage
                                            * Generate the adjusted first stage q level we
> are testing: q' = q/1+q
                                            local qval_adj = `qval'/(1+`qval')
                                            * Generate value q'*r/M
gen fdr_temp1 = `qval_adj'*rank/`totalpvals'
* Generate binary variable checking condition
> p(r) \le q'*r/M
                                            gen reject_temp1 = (fdr_temp1>=pval) if pval~=
                                            * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                            gen reject_rank1 = reject_temp1*rank
* Record the rank of the largest p-value that
> meets above condition
                                            egen total_rejected1 = max(reject_rank1)
                                            * Second Stage
                                            * Generate the second stage q level that accou
> nts for hypotheses rejected in first stage: q_2st = q'*(M/m0) local qval_2st = `qval_adj'*(`totalpvals'/(`to
> talpvals'-total rejected1[1]))
                                            * Generate value q_2st*r/M gen fdr_temp2 = `qval_2st'*rank/`totalpvals'
                                            * Generate binary variable checking condition
> p(r) \le q 2st*r/M
                                            gen reject_temp2 = (fdr_temp2>=pval) if pval~=
                                            * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                            gen reject_rank2 = reject_temp2*rank
                                            * Record the rank of the Targest p-value that
> meets above condition
                                            egen total rejected2 = max(reject rank2)
                                            * A p-value has been rejected at level q if it
> s rank is less than or
                                            * equal to the rank of the max p-value that me
> ets the above condition
                                            replace bky06 qval = `qval' if rank <= total r</pre>
> ejected2 & rank~=.
                                            * Reduce q by 0.001 and repeat loop
                                            drop fdr_temp* reject_temp* reject_rank* total
> rejected*
                                            local qval = `qval' - .001
                                   }
                                   quietly sort original_sorting_order
                                   pause off
                                   mkmat bky06 qval, matrix(pval bky06)
                          scalar i = 1
                          foreach reg store of local group regressions{
                                   scalar holm cor val = holm cor[i, 1]
```

scalar pval bky $\overline{0}6$ val = pval bky06[i, 1]

```
if holm cor val == 1 {
                                           est restore `reg_store'
                                           estadd local pholm "Significant"
                                           estadd scalar bky 06 = pval bky06 val
                                  if holm_cor_val == 0 {
        est restore `reg_store'
                                           estadd local pholm "No Significant"
                                           estadd scalar bky_06 = pval_bky06_val
                                  }
                 restore
*************************
* PANEL C (Material investments)
****************
        * Define number of hypothesis
        scalar hyp = 5
        * Define level of significance
        scalar signif = 0.05
        local fcimat "fci play mat type Npaintbooks Nthingsmove Ntoysshape Ntoysbought
        local i = 1
        local group_regressions = ""
        mat p values = J(hyp, 1, .)
        foreach y of local fcimat{
                 cap drop V*
                 reg `y'1_st treat fci_play_mat_type0_st $covs_ent , cluster(cod_dane)
                eststo panel_C_`i': test treat = 0
mat p_values[`i',1]=r(p)
                 scalar p_value = r(p)
                 scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
                 estadd scalar N1 = e(N)
                 local est_name panel_C_`i'
local group_regressions `group_regressions' `est_name'
                 local i = `i' + 1
        reg home name1 st
        estadd scalar \overline{N}1 = .
        est sto panel C 6
        // Modification of p-vals
                 preserve
                          // * Define number of hypothesis
                         // scalar hyp = hyp
                          // * Define level of significance
                          // scalar signif = signif
                          * Holm Correction
                                  clear
                                  // Bring the p_values matrix as column dta
                                  symat p values
                                  // Identify the original variable outcome
                                  gen outcome_order_var = _n
                                  // Sort values
                                  sort p_values1
                                  // Indicate the rank of the variable
                                  gen rank = n
                                  // Identify if its significan or note
                                  gen alpha_corr = signif/(hyp+1-rank)
gen significant_Holm = (p_values1<alpha_corr)
replace significant_Holm = 0 if significant_Holm[_n-1]</pre>
```

```
// Sort again based on outcome order
                                   sort outcome order var
                                   // Export the result as a matrix
                                   mkmat significant Holm, matrix (holm cor)
                                   keep p_values1 outcome_order_var
                           * Benjamini et al.
                                   rename (p values1 outcome order var) (pval outcome)
                                   quietly sum pval
                                   local total pvals = r(N)
                                   * Sort the p-values in ascending order and generate a
> variable that codes each p-value's rank
                                   quietly gen int original sorting order = n
                                   quietly sort pval
                                   quietly gen int rank = n if pval~=.
                                   * Set the initial counter to 1
                                   local qval = 1
                                   * Generate the variable that will contain the BKY (200
> 6) sharpened q-values
                                   gen bky06 qval = 1 if pval~=.
                                   * Set up a loop that begins by checking which hypothes
> es are rejected at q = 1.000, then checks which hypotheses are rejected at q = 0.999 >, then checks which hypotheses are rejected at q = 0.998, etc. The loop ends by che
> cking which hypotheses are rejected at q = 0.001.
                                   local totalpvals = ${totalpvals}
                                   local qval = 1
                                           .
qval' > 0 {
                                   while
                                            * First Stage
                                            * Generate the adjusted first stage q level we
> are testing: q' = q/1+q
                                            local qval_adj = `qval'/(1+`qval')
                                            * Generate value q'*r/M gen fdr_temp1 = `qval_adj'*rank/`totalpvals'
                                            * Generate binary variable checking condition
> p(r) \le q'*r/M
                                            gen reject temp1 = (fdr temp1>=pval) if pval~=
                                            * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                            gen reject_rank1 = reject temp1*rank
                                            \star Record the rank of the largest p-value that
> meets above condition
                                            egen total_rejected1 = max(reject_rank1)
                                            * Second Stage
                                            ^{\star} Generate the second stage q level that accou
> nts for hypotheses rejected in first stage: q_2st = q'*(M/m0) local qval_2st = `qval_adj'*(`totalpvals'/(`totalpvals'))
> talpvals'-total rejected1[1]))
                                            * Generate value q_2st*r/M
gen fdr_temp2 = `qval_2st'*rank/`totalpvals'
                                             * Generate binary variable checking condition
> p(r) \le q 2st*r/M
                                            gen reject temp2 = (fdr temp2>=pval) if pval~=
                                            * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                            gen reject_rank2 = reject_temp2*rank
                                             * Record the rank of the largest p-value that
> meets above condition
                                            egen total rejected2 = max(reject rank2)
```

```
* A p-value has been rejected at level q if it
> s rank is less than or
                                         * equal to the rank of the max p-value that me
> ets the above condition
                                         replace bky06 qval = `qval' if rank <= total r</pre>
> ejected2 & rank~=.
                                         * Reduce q by 0.001 and repeat loop
                                         drop fdr_temp* reject_temp* reject_rank* total
> rejected*
                                         local qval = `qval' - .001
                                quietly sort original sorting order
                                pause off
                                mkmat bky06 qval, matrix(pval bky06)
                        scalar i = 1
                        foreach reg store of local group regressions{
                                scalar holm cor val = holm cor[i, 1]
                                scalar pval_bky06[i, 1]
                                if holm cor val == 1 {
                                        est restore `reg_store' estadd local pholm "Significant"
                                         estadd scalar bky_06 = pval_bky06_val
                                if holm_cor_val == 0 {
                                         est restore `reg store'
                                         estadd local pholm "No Significant"
                                         estadd scalar bky_06 = pval_bky06_val
                                }
                restore
*************************
* PANEL D (Time investments)
******************
        * Define number of hypothesis
        scalar hyp = 5
        * Define level of significance
        scalar signif = 0.05
        local fcitime "fci play act home stories home read home toys home name"
        local i = 1
        local group_regressions = ""
       mat p values = J(hyp,1,.)
        foreach y of local fcitime{
                cap drop V*
                reg `y'1_st treat fci_play_act0_st $covs_ent , cluster(cod_dane)
                eststo panel_D_`i': test treat = 0
mat p_values[`i',1]=r(p)
scalar p_value = r(p)
                scalar corr_p_value = min(1,r(p)*hyp)
estadd scalar bonferroni = corr_p_value
                estadd scalar N1 = e(N)
                local est_name panel_D_`i'
local group_regressions `group_regressions' `est_name'
```

```
local i = `i' + 1
         // aux reg for extra column
        reg home_name1_st
        estadd s\overline{c}alar \overline{N}1 =
        est sto panel_D_6
// Modification of p-vals
                 preserve
                           // * Define number of hypothesis
                           // scalar hyp = hyp
                           // * Define level of significance
                           // scalar signif = signif
                           * Holm Correction
                                    clear
                                    // Bring the p_values matrix as column dta
                                    svmat p_values
                                    // Identify the original variable outcome
                                    gen outcome order var = n
                                    // Sort values
                                    sort p values1
                                    // Indicate the rank of the variable
                                    gen rank = _n
// Identify if its significan or note
                                   gen alpha_corr = signif/(hyp+1-rank)
gen significant_Holm = (p_values1<alpha_corr)
replace significant_Holm = 0 if significant_Holm[_n-1]</pre>
> ==0
                                    // Sort again based on outcome order
                                    sort outcome order var
                                    // Export the result as a matrix
                                   mkmat significant_Holm, matrix(holm_cor)
                                    keep p values1 outcome order var
                           * Benjamini et al.
                                    rename (p_values1 outcome_order_var) (pval outcome)
                                    quietly sum pval local totalpvals = r(N)
                                    * Sort the p-values in ascending order and generate a
> variable that codes each p-value's rank
                                    quietly gen int original_sorting_order = _n
                                    quietly sort pval
                                    quietly gen int rank = n if pval~=.
                                    * Set the initial counter to 1
                                    local qval = 1
                                    * Generate the variable that will contain the BKY (200
> 6) sharpened q-values
                                    gen bky06 qval = 1 if pval\sim=.
                                    * Set up a loop that begins by checking which hypothes
> es are rejected at q = 1.000, then checks which hypotheses are rejected at q = 0.999
> , then checks which hypotheses are rejected at q = 0.998, etc. The loop ends by che
> cking which hypotheses are rejected at q = 0.001.
```

```
local totalpvals = ${totalpvals}
                                  local qval = 1
                                  while `qval' > 0 {
                                           * First Stage
                                           * Generate the adjusted first stage q level we
> are testing: q' = q/1+q
                                          local qval_adj = `qval'/(1+`qval')
* Generate value q'*r/M
                                          gen fdr_temp1 = `qval adj'*rank/`totalpvals'
                                           * Generate binary variable checking condition
> p(r) \le q'*r/M
                                          gen reject temp1 = (fdr temp1>=pval) if pval~=
                                           * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                          gen reject rank1 = reject temp1*rank
                                           * Record the rank of the largest p-value that
> meets above condition
                                          egen total rejected1 = max(reject rank1)
                                           * Second Stage
                                           * Generate the second stage q level that accou
> nts for hypotheses rejected in first stage: q_2st = q'*(M/m0)
                                          local qval_2st = `qval_adj'*(`totalpvals'/(`to
> talpvals'-total rejected1[1]))
                                          * Generate value q_2st*r/M
gen fdr_temp2 = `qval_2st'*rank/`totalpvals'
                                           * Generate binary variable checking condition
> p(r) \le q 2st*r/M
                                          gen reject temp2 = (fdr temp2>=pval) if pval~=
                                           * Generate variable containing p-value ranks f
> or all p-values that meet above condition
                                          gen reject_rank2 = reject_temp2*rank
* Record the rank of the largest p-value that
> meets above condition
                                          egen total rejected2 = max(reject rank2)
                                          * A p-value has been rejected at level q if it
> s rank is less than or
                                           * equal to the rank of the max p-value that me
> ets the above condition
                                          replace bky06 qval = `qval' if rank <= total r</pre>
> ejected2 & rank~=.
                                          * Reduce q by 0.001 and repeat loop
                                          drop fdr_temp* reject_temp* reject_rank* total
> rejected*
                                          local qval = `qval' - .001
                                  }
                                  quietly sort original sorting order
                                  pause off
                                  mkmat bky06 qval, matrix(pval bky06)
                         scalar i = 1
                         foreach reg store of local group regressions{
                                  scalar holm cor val = holm cor[i, 1]
                                  scalar pval_bky06_val = pval_bky06[i, 1]
```

```
if holm cor val == 1 {
                                         est restore `reg_store'
                                         estadd local pholm "Significant"
                                         estadd scalar bky 06 = pval bky06 val
                                 if holm_cor_val == 0 {
        est restore `reg_store'
                                         estadd local pholm "No Significant"
                                         estadd scalar bky_06 = pval_bky06_val
                                 }
                restore
* Replicated Table
******************
                 * Making table for mortality
                #delimit ;
                         global note "Notes: All scores have been internally standardiz
> ed nonparametrically
                                         for age and are expressed in standard deviatio
> n
                                         units (see online Appendix B for details about
                                         the measures and the standardization procedure
> ).
                                         Measures followed by (-) have been reversed so
> that a higher
                                         score refers to better behavior. The effects r
> elating
                                         to the latent factors are in log points. Coeff
> icients and
                                         standard errors clustered at the municipality
> level (in
                                         parentheses) are from a regression of the depe
> ndent variable
                                         measured at follow-up on an indicator for whet
> her
                                         the child received any psychosocial stimulatio
> n and controlling
                                         for the child's sex, tester effects, and basel
> ine
                                         level of the outcome.";
                #delimit cr
                # delimit ;
                         esttab panel A 1 panel A 2 panel A 3
                         panel_A_4 panel_A_5 panel_A_6 using "${output}/table2_replicat
> ion newpval.tex", replace
                         cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$
> 6.3f)))
                         starlevels(* 0.10 ** 0.05 *** 0.01)
                         s(N1 bonferroni pholm bky_06, label( "N" "Bonferroni P-val" "
> Holm P-value" "BKY P-val") fmt(%9.0gc) )
                        collabels (none) nostar noobs nonote
                         nonumbers eqlabels( none )
                         nonote
                         keep( treat )
                         varlabels( treat "Treatment" )
                        mtitle(
                                 "\shortstack{ \\ Bayley: \\ Cognitive}"
"\shortstack{ \\ Bayley: \\ Receptive language}"
"\shortstack{ \\ Bayley: \\ Expressive language}"
```

```
"\shortstack{ \\ Bayley: \\ Fine motor}"
                                   "\shortstack{ \\ MacArthur: \\Words the child can say
> }"
                                   "\shortstack{ \\ MacArthur: \\ Complex phrases \\ the
> child can say}")
                          mgroups( "\underline{ Panel A.} \textbf{ Child's cognitive ski
> lls at follow-up }"
, pattern( 1 0 0 0 0 0
> ) prefix(\multicolumn{@span}{c}{) suffix(}) span end(\hline) )
                          prehead("\begin{table} \small \centering
                                   \protect \captionsetup{justification=centering}
                                   \caption{\label{tab:table1} Treatment Impacts on Raw M
> easures and Latent Factors - New Inference Value }"
                                   "\noindent\resizebox{\textwidth}{!}{ \begin{threepartt}
> able \} "
                                   "\begin{tabular}{lcccccc}" \toprule)
                          posthead(\hline) prefoot(\midrule) postfoot(\midrule);
                          esttab panel_B_1 panel_B_2 panel_B_3
                          panel_B_4 panel_B_5 panel_B_6 using "${output}/table2 replica
> tion newpval.tex", append
                          cells(b(label(coef.) star fmt(%8.3f) ) se(label((z)) par fmt(%
> 6.3f)))
                          starlevels(* 0.10 ** 0.05 *** 0.01)
                          s ( N1 bonferroni pholm bky 06,
                                   label ( "N" "Bonferroni P-val" "Holm P-value" "BKY P-va
> 1")
                                   fmt(0 3 3 ) )
                          collabels ( none )
                          nostar noobs nonote
                           nonumbers eqlabels ( none )
                          keep( treat )
                          mgroups ( "
                                   \underline{ Panel B.}
                                   \textbf{ Child's socio-emotional skills at follow-up }
> "
                                   , pattern( 1 0 0 0 0 0 )
                                   prefix(\multicolumn{@span}{c}{) suffix(})
                                   span end(\hline) )
                          mtitle(
                                   "\shortstack{ \\ ICQ: \\Difficult (-)}"
                                   "\shortstack{ \\ ICQ: \\ Unsociable (-)}"
"\shortstack{ \\ ICQ: \\ Unstoppable (-)}"
"\shortstack{ \\ ECBQ: \\ Inhibitory control}"
"\shortstack{ \\ ECBQ: \\ Attentional focusing}" )
                          nonote
                          varlabels( treat "Treatment" )
prehead( "" )
                           posthead( \hline )
                           prefoot(\midrule)
                           postfoot("")
                           delim("&") nonumbers
                          esttab panel C 1 panel C 2 panel C 3
                          panel C 4 panel C 5 panel C 6 using "${output}/table2 replica
> tion newpval.tex", append
                          cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$
> 6.3f)))
                          starlevels(* 0.10 ** 0.05 *** 0.01)
                          s ( N1 bonferroni pholm bky 06,
                                   label ("N" "Bonferroni P-val" "Holm P-value" "BKY P-va
> 1")
                                   fmt(0 3 3 ))
                          collabels (none) nostar noobs
                          nonumbers eqlabels ( none )
                          mgroups( "
                                   \underline{ Panel C.}
```

```
\textbf{ Material investments at follow-up }"
                             , pattern( 1 0 0 0 0 0 )
                             prefix(\multicolumn{@span}{c}{) suffix(})
                             span end(\hline) )
                     mtitle(
                             > erials}"
                             "\shortstack{ \\ FCI: \\ Number of coloring \\ and dra
> wing books}"
                             > vement}"
                             "\shortstack{ \\ FCI: \\ Number of toys \\ to learn sh
> apes}"
                             "\shortstack{ \\ FCI: \\ Number of \\ shop-bought toys
> }"
                             "")
                     nonote keep( treat )
                     varlabels( treat "Treatment" )
                      prehead( \hline )
                      posthead( \hline )
                      prefoot(\midrule)
                      postfoot("")
                      delim("&") nonumbers
                     esttab panel_D_1 panel_D_2 panel_D_3
                     panel D 4 panel D 5 panel D 6 using "${output}/table2 replica
> tion newpval.tex", append
                     cells(b(label(coef.) star fmt(\$8.3f)) se(label((z)) par fmt(\$
> 6.3f)))
                     starlevels(* 0.10 ** 0.05 *** 0.01)
                     s ( N1 bonferroni pholm bky 06,
                             label ("N" "Bonferroni P-val" "Holm P-value" "BKY P-va
> 1")
                             fmt(0 3 3 ))
                     collabels(none) nostar noobs nonote
                      nonumbers eqlabels( none ) keep( treat )
                     mgroups( "
                             \underline{ Panel D.}
                             \textbf{ Time investments at follow-up }"
                             , pattern( 1 0 0 0 0 )
                             prefix(\multicolumn{@span}{c}{) suffix(})
                             span end(\hline) )
                     mtitle(
                             > ivities \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times told \\ a sto
> ry to child \\ in last 3 days}"
                             > ild \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times \\ played wit
> h toys \\ in last 3 days}"
                             "\shortstack{ \\ FCI: \\ Number of times \\ named thin
> gs to child \\ in last 3 days}"
                             "")
                     varlabels( treat "Treatment" )
                      prehead( \hline )
                      posthead( \hline )
                      prefoot(\midrule)
                      delim("&") nonumbers
postfoot(\hline \end{tabular}
                             \begin{tablenotes}
                             \begin{footnotesize}
                             ${note}
                             \end{footnotesize}
                             "\end{tablenotes} \end{threeparttable} } \end{table}")
> ;
```