

Partitioning: sample covariance matrix has transpose of each other within the partitioned matrix. **Number of Elements:** $\frac{n(n+1)}{2}$ unique elements in a cov or corr matrix of n variables. **Generalized Sample Variance:** $|S|$ is product of eigenvalues of S , $|S| = 0$ if any eigenvalue is 0 (multi co-linearity). **Mahalanobis distance:** multivariate version of a z-score; $\delta^2 = (y_i - \mu)^T \Sigma^{-1} (y_i - \mu)$ is in quadratic form and return s a scalar; **variables that have a high degree of variation will contribute less to the overall Mahalanobis distance;** how far a measurement is from the mean vector relative to what a typical deviation from the mean is. **Properties of Multivariate Normal:** y is $N_p(y_i, \Sigma)$; $z \sim N(a^T \mu, a^T \Sigma a)$ and individual y 's must be normal too; $\text{rank}(A) = q \leq p$ then $z \sim N_q(A\mu, A^T \Sigma A)$; if $\mu = 0$ and $\Sigma = I$ then $Ay \sim N(0, I)$. **Independent Random Variables:** y_j and y_k are independent if and only if covariance $s_{jk} = 0$ or stated in terms of correlation of row jk ; **Implication goes both ways since property of MV normal distribution.** *Handout 5: Disprove p vars are jointly MV normally distributed:* null hypothesis of these tests is that variables do follow a MV normal distribution. **Multivariate CLT:** $\bar{y} \sim N_p(\mu, \frac{\Sigma}{n})$ for a large enough sample size n . **Sample Variance and Cov Matrix Distribution:** univariate follows a chi-squared distribution; sample var/cov matrix follows a Wishart distribution: $(n-1)S \sim \text{Wishart}(n-1, \Sigma)$ if y follows a MV normal distribution. **Univariate 1-Sample T Test:** $t = \frac{\bar{y} - \mu_0}{s/\sqrt{n}}$. **Hotelling's 1-Sample T^2 Test:** $T^2 = n(\bar{y} - \mu_0)^T S^{-1} (\bar{y} - \mu_0)$; measures how far observed \bar{y} is from its expected value, if the null were true, while taking into account the variance/cov of the sample mean vector. **Hotelling's T^2 Density:** no upper bound; reject when T^2 is large. **P-value accuracy:** data values must have been sampled from MV normal dist, S must be non-singular, and $n > p$ (observations > variables). **Steps to Take After Rejecting Null:** check multivariate normality of data, conduct univariate tests on each variable. **Benefits of MV Tests:** using p univariate tests inflates the type I error rate (rejecting null when it is true); $p = 4$ and $\alpha = 0.05$ then $1 - (1 - 0.05)^4 = 0.19$ probability and quickly increases with p ; does not ignore correlation structure between p vars; more power (prob of rejecting null when null is true) and high power is good; small deviations may be statistically significant when combined. **Limitations of MV Tests:** interpretations difficult without univariate tests when statistically significant MV test; **non-directional so null hypothesis for MV is 2-tailed.**— *Handout 6: Univariate 2-Sample t-Test:* $H_0: \mu_1 = \mu_2$; $H_a: \mu_1 \neq \mu_2, \mu_1 > \mu_2, \mu_1 < \mu_2$; $t = \frac{\bar{y}_1 - \bar{y}_2}{s_{pl} \sqrt{\frac{1}{n_1} + \frac{1}{n_2}}}$;

$s_{pl}^2 = \frac{(n_1-1)s_1^2 + (n_2-1)s_2^2}{n_1 + n_2 - 2}$; b/c assuming equal variances; validity of p-val: sampled from normal dist. but diff means. **MV 2-Sample Hotelling's T Test:** $H_0: \mu_1 = \mu_2$; $H_a: \mu_1 \neq \mu_2$; $T^2 = (\bar{y}_1 - \bar{y}_2)^T [S_{pl} l(\frac{1}{n_1} + \frac{1}{n_2})]^{-1} (\bar{y}_1 - \bar{y}_2)$; is the Maha. distance between mean vectors; $S_{pl} = \frac{(n_1-1)S_1 + (n_2-1)S_2}{n_1 + n_2 - 2}$; validity of p-val is MV normal dist. with equal cov. matrices but diff. means; y_{ij} is the j th obs. in group i and vectors with p entries; follow up with univariate t-tests to see where differences lie. **Paired MV Data:** Difference = Treatment - Control; MV and Paired (same subject); Apply 1-Sample Hotelling T Test; $H_0: \mu_d = 0$; $H_a: \mu_d \neq 0$; $T^2 = \bar{d}^T [S_d(\frac{1}{n})]^{-1} \bar{d}$; Maha. dist between \bar{d} and 0; validity of p-val is MV normal dist. with $N_z(\mu_d, \Sigma_d)$; follow up with univariate t-tests to see which dist. from 0 is significant. *Handout 7: Uni 1-Way ANOVA:* $y_{i.}$ = sum across all measurements in sample i , $\bar{y}_{i.}$ = sample mean of all measurements in sample i , $\bar{y}_{..}$ = (grand) sample mean of all measurements across all samples; statistical model: $y_{ij} = \mu_i + \alpha_i + \epsilon_{ij} = \mu_i + \epsilon_{ij}$, where (1) is overall pop mean response, (2) effect on mean resp. due to pop i , (3) population mean response for pop i , (4) random error assoc. with j th response in pop i ; assume: (1) $\epsilon_{ij} \sim N(0, \sigma^2)$, (2) $E(y_{ij}) = \mu_i$, (3) $\text{Var}(y_{ij}) = \sigma^2$; var equal; $H_0: \mu_1 = \mu_2 = \dots = \mu_k$; H_a : at least 1 $\mu_i \neq \mu_j$; estimate of σ^2 : **within** $MSE = \frac{\sum_{i=1}^k \sum_{j=1}^{n_i} (y_{ij} - \bar{y}_{i.})^2}{nk - k} = \frac{SSE}{nk - k}$, thought of as pooled estimate of σ^2 ; **between** $MSH = \frac{n \sum_{i=1}^k (\bar{y}_{i.} - \bar{y}_{..})^2}{k-1} = \frac{SSH}{k-1}$, thought of as estimate of σ^2 ; $F = \frac{MSH}{MSE}$, which is explained variation / unexp. variation (within is noise and between is signal); reject when F is large, look at graphs, sd of within group (noise). **1-Way MANOVA:** y_{ij} is a vector of y_{ij1}, \dots, y_{ijp} ; same stat model but with vectors and $y_{i.}$ estimate of μ ; $H_0: \mu_1 = \mu_2 = \dots = \mu_k$; H_a : all pops. do not have the same mean vectors; **between** $H = n \sum_{i=1}^k (\bar{y}_{i.} - \bar{y}_{..})(\bar{y}_{i.} - \bar{y}_{..})^T$; **within** $E = \sum_{i=1}^k \sum_{j=1}^{n_i} (y_{ij} - \bar{y}_{i.})(y_{ij} - \bar{y}_{i.})^T$; $E + H$ is total sample covariance; $\text{Det}()$ of each give generalized within/total variation: $\Lambda = \frac{|E|}{|E+H|}$; Wilk's Λ stat which is noise / total; reject when Λ is small; Transform Λ to F and when large reject; **manova()** then **summary()** with "Wilks", $\$SS$ gives H and $\$Residuals$ gives E ; follow up with univariate ANOVAs to see which var has diff means across groups; **TukeyHSD()** with Bonferroni correction: which groups are different; p-adj bumped up to protect from Type I error; compare p-values to α/p to see which are significant, compare sample means this way. *Handout 8: MANOVA Test Statistics:* All functions of the eigenvalues of $E^{-1}H$, $s = \min(k-1, p)$ is the rank of it; Wilk's $\Lambda = \prod_{i=1}^s \frac{\lambda_i}{1+\lambda_i}$, Roy's largest root $\frac{1}{1+\lambda_1}$, Pillai's $\sum_{i=1}^s \frac{1}{1+\lambda_i}$ (for hetero in cov matrix), Lawley-Hotelling $\sum_{i=1}^s \lambda_i$. **MANOVA Type 1 Error Rate:** As Corr between vars incr. type 1 error rate remains below 0.05. **Profile Analysis on MANOVA:** H_{01} : k profiles are parallel (slope from var to var is same across groups: $\mu_{12} - \mu_{11} = \mu_{22} - \mu_{21} = \dots$), H_{02} : k profiles are at the same level (average of the mean elements are eq across groups (sums equal): $\sum \mu_{1i} = \sum \mu_{2i} = \dots$) **offset in graph, sum of endpoints with 2 lines**, H_{03} : k profiles are flat (p vars avgs across groups are the same: $\mu_{11} + \mu_{21} + \dots = \mu_{12} + \mu_{22} + \dots$) **stacked points avgs are the same with 2 lines**; start with a MANOVA; Parallel Interp: moderate evidence that mean weights change at different rate depending on treatment (reject), Levels Interp: average of the mean weights may be the same across treatments (fail to reject), Flat Interp: mean weights do change regardless of group (reject); Repeated Measures Data: same experimental unit yields multiple observations (obsiv unit not same as exp unit), use profile analysis and MANOVA for this data. *Handout 9: Single Pop Cov Test:* $H_0: \Sigma = \Sigma_0$; $H_a: \Sigma \neq \Sigma_0$; **Test Stat:** $u = (n-1)[\ln|\Sigma_0| - \ln|S| - \text{tr}(S\Sigma_0^{-1}) - p]$; S close to Σ_0 then logs are similar, $S\Sigma_0^{-1} \sim I$, so trace is p , so $u \sim 0$; reject when stat is large; **DF for X^2 dist is the expected value;** Stat and P-val appropriate when n is relatively large and follows MV dist. (Mardia test). **Several Pop. Cov Test:** need for Hotelling's (2-Sample) and MANOVA; $H_0: \Sigma_1 = \Sigma_2 = \dots = \Sigma_k$; H_a : all pop. do not have same covariance matrix; Box M Test Conditions are that k independent samples of data and each k comes from MV dist $N(\mu_i, \Sigma_i)$; Handout 6 pooled sample cov eq but now for several pops is the weighted avg of sample cov. matrices, weighting by sample size; Λ is ration of det of S_i to det of S_{pl} , when 1 then lambda is around 1; Transform to M-test and reject when M-stat is large; **DF is the expected value. Test for Indep. of P Vars:** Σ is diagonal and 0 elsewhere means p vars indep. of each other (can be Σ_0); P_ρ is pop. corr. matrix where diagonal are 1 and 0 elsewhere means p vars indep.; $H_0: P_\rho = I$; $H_a: P_\rho \neq I$; $|R|$ is det of sample corr. matrix and $|R| = 1$ if indep. and 0 if dep., $0-1$ measures degree of evidence for/against null; $X^2 = -[(n-1) - \frac{2p+5}{6}]\ln|R|$ with df being $\frac{p(p-1)}{2}$ which is the expected value. *Handout 10: Discriminant Analysis:* follow up to MANOVA; MANOVA and Hotelling's can distinguish linear combos of vars across groups; maximally separate z-bars of groups; $z = a'y = [S_{pl}^{-1}(\bar{y}_1 - \bar{y}_2)]^T y$, where a' is a scalar; t test on z scores reveals significant differences. **Disc. Analysis on Several Pops.:** $E^{-1}H$ eigenvector or largest eigenvalue is optimal disc. func. to maximally distinguish; handout 8 first line gives number of

funcs; s disc. funcs. define s indep vars that are funcs. of original p vars.; **Relative Importance Formula:** $\frac{\lambda_i}{\sum_{i=1}^s \lambda_i}$, how much variability you retain between groups; \bar{z} of each group is pushed as far as possible from each other. *Handout 11: Stand. Coeffs. of Disc. Funcs.:* `scale()` to get z scores to compare relative importance of vars. **Stat. Signif. of Disc. Funcs.:** $\Lambda_m = \prod_{i=m}^s \frac{1}{1+\lambda_i}$, $V_m = -[N - \frac{p+k+1}{2}] \ln(\Lambda_m)$, V_m transforms to F-stat with $df = (p - m + 1)(k - m)$ as expected value; rejecting MANOVA tests whether at least first eigenvector provides significant dimension of separation; conditions: MV normal and equal cov. matrices across groups. **Stat. Sign. of Vars:** test if vars contribute sign. in group separation; $\Lambda = \frac{\Lambda_p}{\Lambda_{p-1}}$ where top is all vars and bottom is without var, useful if result is small (same if not useful); $F = \frac{1-\Lambda}{\Lambda} \frac{N-p-k+1}{p+k-1}$, large when var is useful, reject null that var does not contribute to group separation; partial F test: compare p-vals with Bonferroni correction of α/p . **Classification Trees:** want high node homo or low hetero; measure node hetero using misclassification rates; `rpart()` with output as `root`, `n`, `loss (misclassified)`, `(Group1, ...)`; large stretch in tree means helpful in split. *Handout 12: Fisher's Procedure:* after `lda()` and `predict()` compute \bar{z} of all groups, `predict()` on new y to get z score, find group using $D^2 = (z_1 - \bar{z}_{i1})^2 + (z_2 - \bar{z}_{i2})^2$. **Conf. Matrix:** accuracy: prop of obs. that classified correctly, misclass. rate is $1 - A$; sensitivity: prop of a given class that is classified as that class; specificity: prop of records not of a given class that are not classified as that class (ex 98/100). **Linear Class Func:** assume groups have same cov. matrix; minimize Maha. distance foild to maximize $L_i(y) = \bar{y}_i^T S_{pl}^{-1} y - (1/2) \bar{y}_i^T S_{pl}^{-1} y_i$. **Quad Class Func:** same minimize Maha dist. but with indiv cov matrix so less power (less data) b/c no assumption. **Steps on Analysis:** (1) Graphic and stats (2) mvn test (3) BoxM test for equal cov matrices across groups (4) standardize for disc analysis (5) `lda()` confusion matrix (6) `qda()` conf matrix. **LOO CV:** Fisher/QDA not good on new data; Training and Testing Set; CV by omit 1st and do lda on rest, but use model on all data and CV as honest estimator of how well model does; Interp: using CV estimates of accuracy/misc rates, sens, spec, one method does a bit better than the other; after CV use model for all predictions (not CV).

Handout 13: Univariate Linear Regression: setup is 1 quant resp var and q quant explanatory vars; random error allows random variation around trend. **Assumptions:** residuals are normal with mean 0, σ^2 constant; variance of resid is σ^2 and cov is 0; so cov matrix is $\sigma^2 I$; expected value of resid is 0, expected value of y is $X\beta$. **Estimation:** $\hat{\beta} = (X^T X)^{-1} X^T y$; $\hat{y} = X\hat{\beta}$; minimize sum of squared residuals; sample variance of residuals is $MSE = s^2 = \frac{SSE}{n-q-1}$; df of SSE is $n - (q + 1)$, where q+1 is the number of β 's. **Coeff of Determ:** statistical significance of R^2 is tested with F-stat **MV Linear Regression:** modeled as $y = X\beta + \epsilon$. **Assumptions:** $E(\epsilon) = 0$, which means $E(y) = X\beta$; $cov(y_i) = \Sigma$, meaning cov between y's for same observation; $cov(y_i, y_j) = 0$ for $i \neq j$, meaning y's for diff observations is 0; both these imply that measurements of y on teh same observation can be correlated, but with the same cov structure for all obsv, measurements taken on diff obsv are uncorrelated; $\Sigma_i \sim N_p(0, \Sigma)$; **Estimation:** Dimension of β is $(q + 1) \times p$. **Estimate Common Cov Matrix:** $S_e = \frac{1}{n-q-1} E$, where E is like SSE and is the MV version of unexplained variability in data (error SS); total = $E + H = (Y^T Y - \hat{\beta}^T X^T Y) + (\hat{\beta}^T X^T Y - n\bar{y}^T \bar{y})$. **Test of Overall Regression:** $\lambda = \frac{|E|}{|E+H|}$, which is unexplained over total; H_0 : no linear association between any of the x's (vector) and any of the y's (vector); H_0 : $\beta_1 = 0$, where β_1 is the part of the matrix with the predictors (without intercept); reject when λ is small. **Conclusion:** lambda small: there is little unexplained variation. regression model explains a significant amount. At least one of the x's is linearly associated with at least one of y's; then look at univariate tests to see which x's are significant. *Handout 14: Canonical Correlation:* also the MV correlation coeff; partition data into μ and Σ , which has cov matrix of y and x; sample mean vector is \bar{y} and \bar{x} ; sample cov matrix is S_{yy} , $2 S_{yx}$, and S_{xx} . **Intuition:** find 2 linear combs of y and x that will maximize Pearson's sample corr between observed transformed values; Find a_1 and b_1 such that $\mu_1 = a_1^T y$ and $\nu_1 = b_1^T x$ have max corr. **Solution:** a_1 is first eigenvector of $S_{yy}^{-1} S_{yx} S_{xx}^{-1} S_{yx}$ and b_1 is first eigenvector of $S_{xx}^{-1} S_{xy} S_{yy}^{-1} S_{xy}$; max corr coeff is first eigenvalue of any of the above eqs; μ and ν are the canonical variates; 2 matrices should have same eigenvalues but different vectors. **Bivariate Graph:** first canonical corr should be at least max corr between any pair of y and x in graph. **Correlation:** squaring first eigenvalue gives correlation coeff; in R `corr(all.u, all.v)`; largest λ_1 is the largest possible squared corr of any 2 linear comb of x and y vars; $\lambda_1 = (corr(\mu_1, \nu_1))^2$. **Definition:** First canonical corr r_1 is $\sqrt{\lambda_1} = |corr(\mu_1, \nu_1)|$, magnitude of largest possible corr coeff; s = min(p, q) eigenvalues so s canonical corr coeff. **Test of Significance of Canonical Corr:** (Test of Overall Regression); Wilk's $\Lambda = \prod_{i=1}^s (1 - r_i^2)$, **can test sign of next canonical corr coef by starting product at $i + 1$;** in R `linearHypothesis()`; reversing the order of x and y will give same test of sign but diff H and E. *Handout 15: Intuition Comp Analysis:* goal is dimension reduction; SE are large and lack of significance b/c of multicoll; reduce vars to increase precision (lower SEs) estimates and predictions; can be not significant b/c of covering 0 in CI interval; benefit from having exp vars with larger variance. **Principal Comp Analysis:** vars will be uncorrelated and max variance; **Derivation:** $z = Ay$ and $\Sigma = A\Sigma_y A^T$, where cov matrix of z is diagonal: 0s elsewhere, $cov(z_i, z_j) = 0$ and $var(z_i) = \sigma_{z_i}^2$. **Spectral Decomp:** $\Sigma_y = CDC^T$ where C is the normalized eigenvectors and D contains eigenvalues of Σ_y , so $D = C^T \Sigma_y C$, which is a diagonalized sigma matrix; $z = C^T y$ so $\Sigma_z = C^T \Sigma_y C = D$; $tr(\Sigma_y)$ is the total population variance; total variance of 1st k princ comps / total var of all y's = $\frac{\sum_{i=1}^k \lambda_i}{\sum_{i=1}^p \lambda_i}$, bottom is $tr(\Sigma_y)$; z's have all of the var/cov of the y's in terms of generalized and total variance. **Implementation:** use sample corr matrix R_y when (1) some of vars have much larger variance than others (2) y-vars on much diff scales; use as many components to retain at least 80% of og vars total variance; Hotelling's test to see if they differ by class. **Handout 16: Objective of Cluster Analysis:** only 1 way to put all obsv into 1 cluster, only 1 way to put data into 16 clusters; 10 billion ways to cluster these 16 observations example. **Hierarchical Agglomerative:** start with m any small then join most similar, g suitable degree of homo, stop until 1 cluster will all obsv; use euclidean distance; single linkage or nearest neighbor: uses distance between 2 nearest obsv or vectors as distance between clusters; complete linkage or farthest: distance between 2 clusters is max possible dist between obsv (poor); average: average distance between obsv in clusters; centroid: distance between sample mean vector of obsv in clusters; Ward's: measure within cluster variability by sums of squares, combine 2 clusters with smallest increase in within-cl variability, minimize $SSE_{AB} - (SSE_A + SSE_B)$; dendrogram for linkage. **Hierarchical Divisive:** few large and successively divide into more and more similar to make homo; **Partitioning Methods:** approx of all possible partitions that can be considered and assign obsv in way that makes clusters as homo as possible; K-Means: get seeds of clusters, measure distance of each osbv to centroid, closest are assigned, repeat until no longer assigned to new clusters.