# Lecture 3: Resemblance and relatedness

Bruce Walsh lecture notes Introduction to Quantitative Genetics SISG (Module 9), Seattle 15 – 17 July 2019

# Heritability

- Central concept in quantitative genetics
- Fraction of phenotypic variance due to additive genetic values (Breeding values)
  - $-h^2 = V_A/V_P$
  - This is called the narrow-sense heritability
  - Phenotypes (and hence V<sub>P</sub>) can be directly measured
  - Breeding values (and hence V<sub>A</sub>) must be estimated
- Estimates of V<sub>A</sub> require known collections of relatives

# Broad-sense heritability

- Narrow-sense heritability h<sup>2</sup> applies when outcrossing,
  - $h^2 = Var(A)/Var(P)$
  - = the fraction of all trait variation due to variation in breeding (additive genetic) values
- Broad-sense heritability H<sup>2</sup> applies when selecting among a series of pure lines
  - $H^2 = Var(G)/Var(P)$
  - = the fraction of all trait variation due to variation in Genotypic values

## Defining H<sup>2</sup> for Plant Populations

Plant breeders often do not measure individual plants (especially with pure lines), but instead often measure a plot or a block of individuals.

This replication can result in inconsistent measures of H<sup>2</sup> even for otherwise identical populations.

Let  $z_{ijkl}$  denote the value of the l-th replicate in plot k of genotype in environment j. We can decompose this value as

$$z_{ijkl} = G_i + E_j + GE_{ij} + p_{ijk} + e_{ijkl}$$
 deviations of individual plants within this plot

Suppose we replicate the genotype over e environments, with r plots (replicates) per environment, and n individuals per plot.

If we set our unit of measurement as the average over all plots, the phenotypic variance for the mean of line i becomes

$$\sigma^{2}(\bar{z_{i}}) = \sigma_{G}^{2} + \sigma_{E}^{2} + \frac{\sigma_{GE}^{2}}{e} + \frac{\sigma_{p}^{2}}{er} + \frac{\sigma_{e}^{2}}{ern}$$

Thus,  $V_P$ , and  $H^2 = V_G/V_P$ , depend on our choice of e, r, and n

In order to compare board-sense heritabilities we need to use a consistent design (same values of e, r, and n)

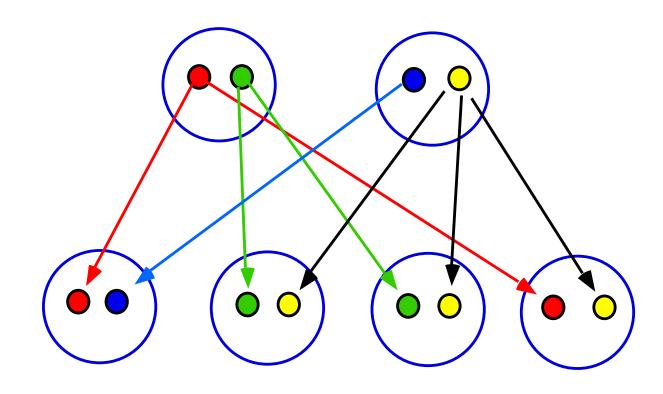
# Key observations

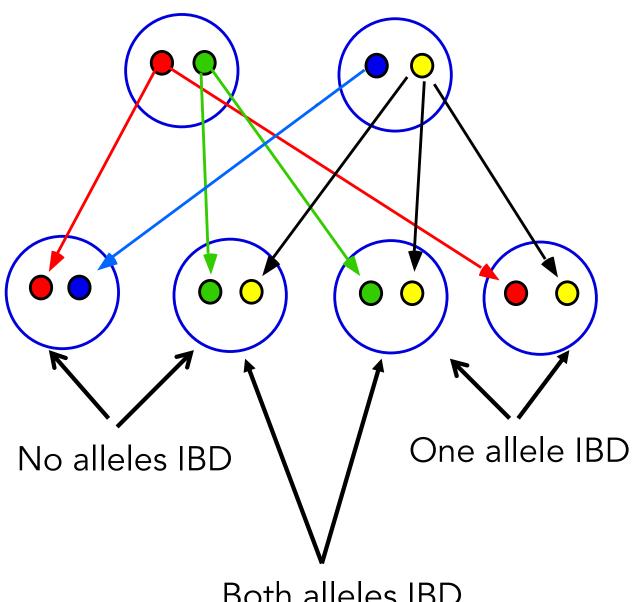
- The amount of phenotypic resemblance among relatives for the trait provides an indication of the amount of genetic variation for the trait.
- If trait variation has a significant genetic basis, the closer the relatives, the more similar their appearance
- The covariance between the phenotypic value of relatives measures the strength of this similarity, with larger Cov = more similarity

#### Genetic Covariance between relatives

Sharing alleles means having alleles that are identical by descent (IBD): both copies can be traced back to a single copy in a recent common ancestor.

Genetic covariances arise because two related individuals are more likely to share alleles than are two unrelated individuals.





Both alleles IBD

# Resemblance between relatives and variance components

- The phenotypic variance between relatives can be expressed in terms of genetic variance components
  - $-\operatorname{Cov}(z_{x},z_{y}) = a_{xy}V_{A} + b_{xy}V_{D}.$
  - The weights a and b depend on the nature of the relatives x and y, and are measures of how often they are expected to share alleles identical by descent
  - These are critical in predicting selection response

### Parent-offspring genetic covariance

 $Cov(G_p, G_o)$  --- Parents and offspring share EXACTLY one allele IBD

Denote this common allele by A<sub>1</sub>

$$G_{p} = A_{p} + D_{p} = \alpha_{1} + \alpha_{x} + D_{1x}$$

$$G_{o} = A_{o} + D_{o} \neq \alpha_{1} + \alpha_{y} + D_{1y}$$
IBD allele

Non-IBD alleles

$$Cov(G_{o},G_{p}) = Cov(\alpha_{1} + \alpha_{x} + D_{1x}, \alpha_{1} + \alpha_{y} + D_{1y})$$

$$= Cov(\alpha_{1}, \alpha_{1}) + Cov(\alpha_{1}, \alpha_{y}) + Cov(\alpha_{1}, D_{1y})$$

$$+ Cov(\alpha_{x}, \alpha_{1}) + Cov(\alpha_{x}, \alpha_{y}) + Cov(\alpha_{x}, D_{1y})$$

$$+ Cov(D_{1x}, \alpha_{1}) + Cov(D_{1x}, \alpha_{y}) + Cov(D_{1x}, D_{1y})$$

All blue covariance terms are zero.

- By construction,  $\alpha$  and D are uncorrelated
  - By construction,  $\alpha$  from non-IBD alleles are uncorrelated
  - By construction, D values are uncorrelated unless both alleles are IBD

$$Cov(\alpha_x, \alpha_y) = \begin{cases} 0 & \text{if } x \neq y, \text{ i.e., not IBD} \\ Var(A)/2 & \text{if } x = y, \text{ i.e., IBD} \end{cases}$$

$$Var(A) = Var(\alpha_1 + \alpha_2) = 2Var(\alpha_1)$$

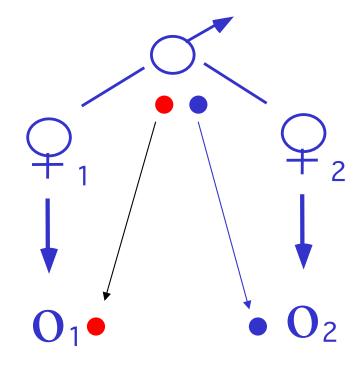
so that

$$Var(\alpha_1) = Cov(\alpha_1, \alpha_1) = Var(A)/2$$

Hence, relatives sharing one allele IBD have a genetic covariance of Var(A)/2

The resulting parent-offspring genetic covariance becomes  $Cov(G_p,G_o) = Var(A)/2$ 

### Half-sibs



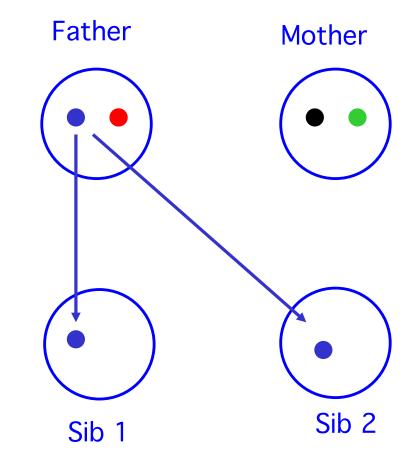
Each sib gets exactly one allele from common father, different alleles from the different mothers

The half-sibs share no alleles IBD

occurs with probability 1/2

Hence, the genetic covariance of half-sibs is just (1/2)Var(A)/2 = Var(A)/4

### Full-sibs



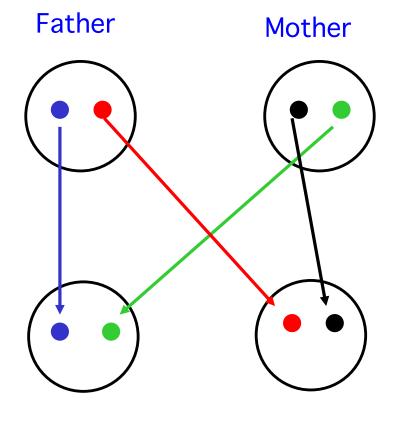
Each sib gets exact one allele from each parent

Prob(Allele from father IBD) = 1/2. Given the allele in parent one, prob = 1/2 that sib 2 gets same allele

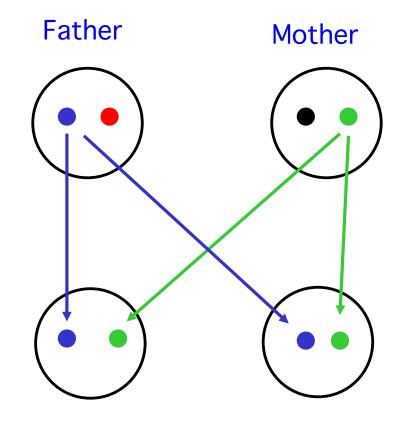
Prob(Allele from father not IBD) = 1/2. Given the allele in parent one, prob = 1/2 that sib 2 gets different allele

### Full-sibs

Each sib gets exact one allele from each parent



Paternal allele not IBD [ Prob = 1/2 ] Maternal allele not IBD [ Prob = 1/2 ] Prob(sibs share 0 alleles IBD) = 1/2\*1/2 = 1/4



Each sib gets exact one allele from each parent

> Paternal allele IBD [ Prob = 1/2 ] Maternal allele IBD [ Prob = 1/2 ] Prob(sibs share 2 alleles IBD) = 1/2\*1/2 = 1/4

> Prob(share 1 allele IBD) = 1-Pr(0) - Pr(2) = 1/2

#### Resulting Genetic Covariance between full-sibs

I BD alleles	Probability	Cantr ibution			
0	1/4	0			
1	1/2	Var(A)/2			
2	1/4	Var(A) + Var(D)			
Cov(Full-sibs) = Var(A)/2 + Var(D)/4					

#### Genetic Covariances for General Relatives

```
Let r = (1/2)Prob(1 \text{ allele IBD}) + Prob(2 \text{ alleles IBD})
Let u = Prob(both \text{ alleles IBD})
```

General genetic covariance between relatives Cov(G) = rVar(A) + uVar(D)

When epistasis is present, additional terms appear r<sup>2</sup>Var(AA) + ruVar(AD) + u<sup>2</sup>Var(DD) + r<sup>3</sup>Var(AAA) +

# More general relationships

- To obtain the expected covariance for any set of relatives, we normally need only compute r and u for that set of relatives
- With general inbreeding, becomes more complex (as three other terms, in addition to  $V_{\rm A}$  and  $V_{\rm D}$  arise)
- With crosses involving inbred and/or related parents, values for r and u are different from those presented above.

# Coefficients of Coancestry

Suppose we pick a single allele each at random from two relatives. The probability that these are IBD is called  $\Theta$ , the coefficient of coancestry. In terms of our previous notation,  $2\Theta = r =$ the coeff on Var(A)

 $\Theta_{xy}$  denotes the coefficient for relatives x and y

Consider an offspring z from a (hypothetical) cross of x and y.  $\Theta_{xy} = f_z$ , the inbreeding coefficient of z. Why? Because the offspring of x and y each get a randomly-chosen allele from each parent. The probability  $f_z$  that both alleles are IBD (the probability of inbreeding) is thus just  $\Theta_{xy}$ .

# $\theta$ and the coefficient on $V_A$

- The coefficient on the additive variance for the relatives x and y is just  $2\theta_{xy}$ .
- To see this,
  - let  $A_iA_j$  denote the two alleles in x and  $A_kA_l$  those in y.
  - Cov(breeding values) =  $Pr(A_i \text{ ibd } A_k) \text{ cov}(\alpha_i, \alpha_k) + Pr(A_i \text{ ibd } A_l) \text{ cov}(\alpha_i, \alpha_l) + Pr(A_j \text{ ibd } A_k) \text{ cov}(\alpha_j, \alpha_k) + Pr(A_i \text{ ibd } A_l) \text{ cov}(\alpha_j, \alpha_l) = 4 \theta_{xy} Var(\alpha)$
  - Since  $Var(A) = 2Var(\alpha)$ ,  $Cov = 2 \theta_{xv}Var(A)$

# $\Theta_{xx}$ : The Coancestry of an individual with itself

Self x, what is the inbreeding coefficient of its offspring?

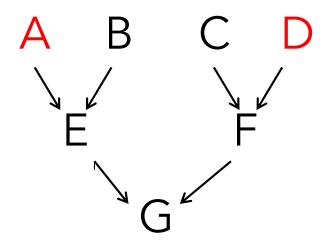
To compute  $\Theta_{xx}$ , denote the two alleles in x by  $A_1$  and  $A_2$ 

Hence, for a non-inbred individual,  $\Theta_{xx} = 2/4 = 1/2$ 

If x is inbred,  $f_x = \text{prob } A_1 \text{ and } A_2 \text{ IBD}$ ,

$$\Theta_{xx} = (1 + f_x)/2$$

# Example

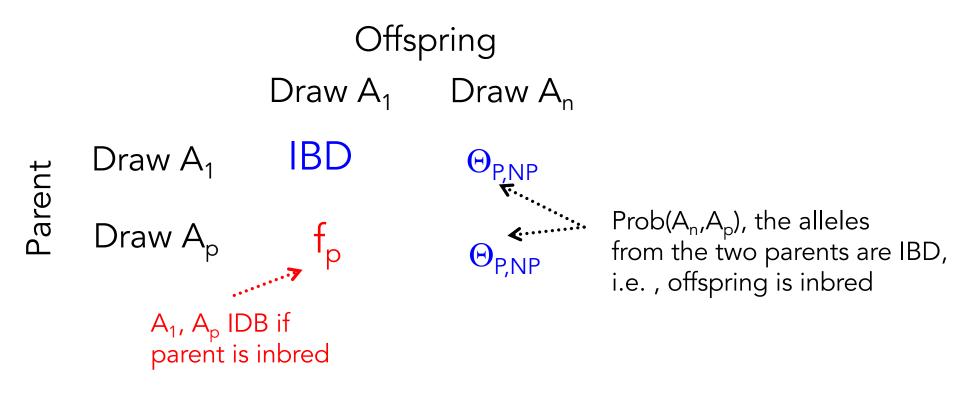


Consider the following pedigree Suppose A and D are fully-inbred, and related, lines with  $\theta_{AD} = 0.5$ . Further, B and C are unrelated and outcrossed individuals

Individual	А	В	С	О
F <sub>x</sub>	1	0	0	1
$\theta_{xx} = (1 + F_x)/2$	1	1/2	1/2	1

## The Parent-offspring Coancestry

Let A<sub>1</sub>, A<sub>n</sub> denote the two alleles in the offspring, where  $A_n$  is the allele from the nonfocal parent (NP), while  $A_1,A_p$  are the two alleles in the focal parent (P)



For a non-inbred individual,  $\Theta_{P0} = 1/4$ 

General: 
$$\Theta_{PO} = (1 + f_p + 2\Theta_{P,NP})/4 = (1 + f_p + 2f_o)/4$$
 24

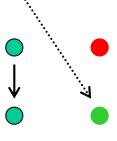
## $\Theta_{op}$ = Parent & Offspring

Parent inbred

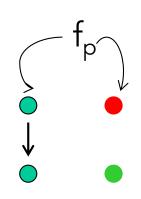
Paternal allele

Mother

Offspring

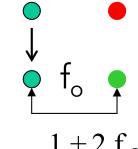


$$\theta_{po} = \frac{1}{4}$$



$$\theta_{po} = \frac{1 + f_p}{4}$$

Offspring inbred



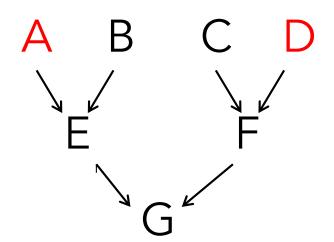
$$\theta_{po} = \frac{1 + 2 \cdot 1_0}{4}$$

1/2 = Prob random offspring allele from father. Prob =  $\theta_{mf}$  =  $f_o$  that this allele is IBD to mother giving a contribution of  $f_o/2$ 

$$\theta_{po} = \frac{1}{4} (1 + f_p + 2 \theta_{mf})$$

 $\dot{}^{\cdot}$ . This is just  $2f_0$ 

#### From before



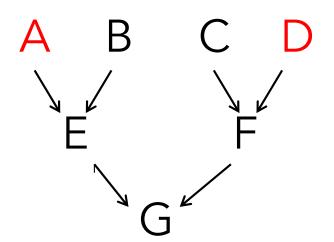
$$\theta_{AA} = \theta_{DD} = 1$$
;  $\theta_{BB} = \theta_{CC} = 1/2$ ;  
 $\theta_{AD} = 1/2$ ,  
 $\theta_{AB} = \theta_{AC} = \theta_{BC} = \theta_{BD} = \theta_{CD} = 0$ 

Consider A - E (inbred parent - offspring)  $\theta_{AE} = (1+f_A)/4 = (1+1)/4 = 1/2$ . Same value for  $\theta_{DF}$ 

Consider B - E (outbred parent - offspring)  $\theta_{BE} = (1+f_B)/4 = (1+0)/4 = 1/4$ . Same value for  $\theta_{CF}$ 

Consider E - G (outbred parent - offspring)  $\theta_{EG} = (1+f_E)/4 = (1+0)/4 = 1/4$ . Same value for  $\theta_{FG}$ 

#### From before



$$\theta_{AA} = \theta_{DD} = 1; \ \theta_{BB} = \theta_{CC} = 1/2;$$
  
 $\theta_{AD} = 1/2,$   
 $\theta_{AB} = \theta_{AC} = \theta_{BC} = \theta_{BD} = \theta_{CD} = 0$ 

#### What about $\theta_{EF}$ ?

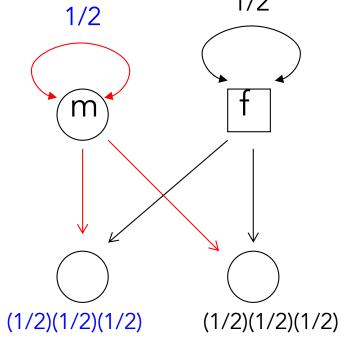
The randomly-chosen allele from E has equal chance of being from A or B. Likewise for F (from C or D)

Of these four possible combinations (A&C, A&D, B&C, B&D), only an allele from A and an allele from D have a chance of being IBD, which is  $\theta_{AD} = 1/2$ .

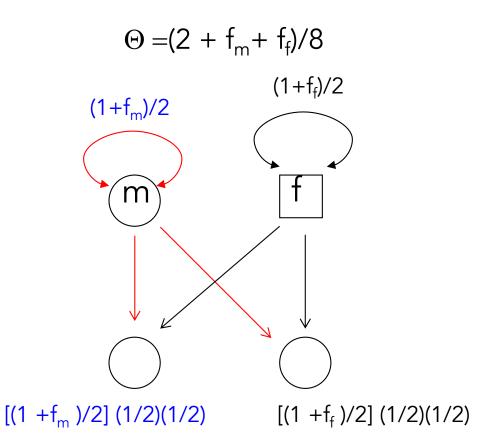
Hence, 
$$\theta_{EF} = \theta_{AD}/4 = 1/8$$

### Full sibs (x and y) from parents m and f

$$\Theta = 1/8 + 1/8 = 1/4$$

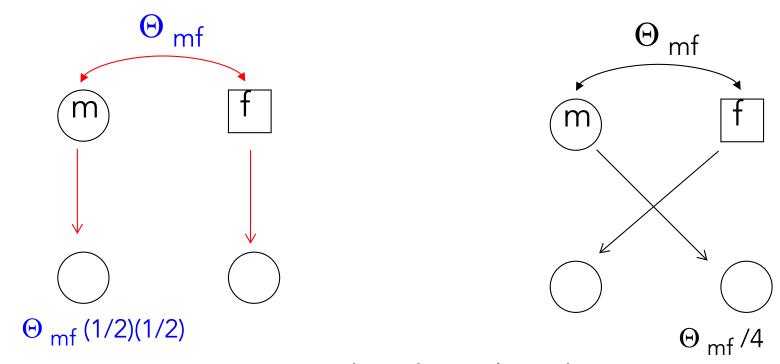


Unrelated, non-inbred parents



Unrelated, inbred parents

### Full sibs (x and y) from parents m and f

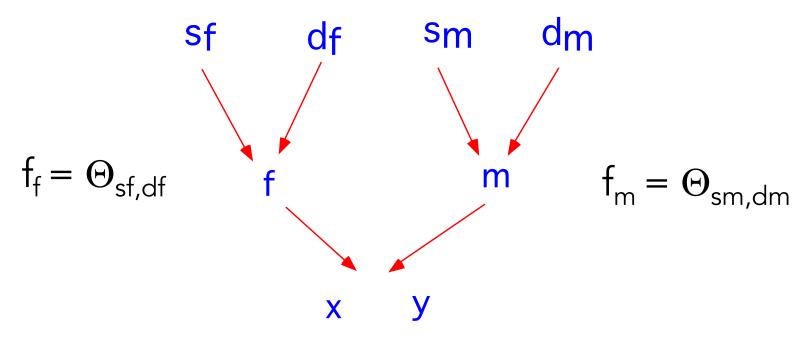


Parents inbred & related. Two additional paths to add to  $\Theta = (2+f_m+f_f)/8$ 

This gives 
$$\Theta = (2+f_m+f_f+4\Theta_{mf})/8$$

### Full sibs (x and y) from parents m and f

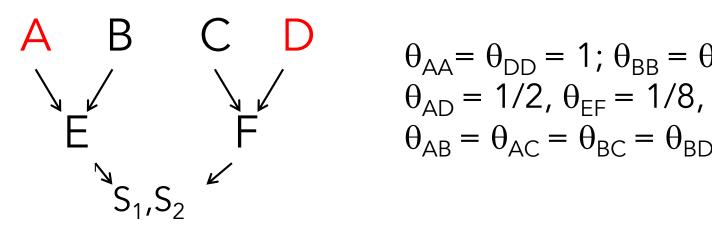
$$\Theta_{xy} = (2 + f_m + f_f + 4\Theta_{mf})/8$$



Putting all this together gives

$$\Theta_{xy} = (2 + \Theta_{sm,dm} + \Theta_{sf,df} + 4\Theta_{mf})/8$$

# Example



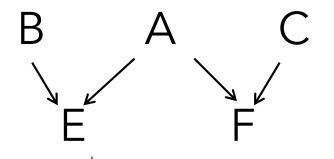
#### From before

$$\theta_{AA} = \theta_{DD} = 1$$
;  $\theta_{BB} = \theta_{CC} = 1/2$ ;  
 $\theta_{AD} = 1/2$ ,  $\theta_{EF} = 1/8$ ,  
 $\theta_{AB} = \theta_{AC} = \theta_{BC} = \theta_{BD} = \theta_{CD} = 0$ 

$$\Theta_{xy} = (2 + \Theta_{AB} + \Theta_{CD} + 4\Theta_{EF})/8$$

$$\theta_{S1S2} = (2 + 0 + 0 + 4[1/8])/8 = (4 + 1)/16 = 5/16$$

### Half-sibs



A is the common parent

Using the same arguments as above,

$$\theta_{EF} = (\theta_{AA} + \theta_{AB} + \theta_{AC} + \theta_{BC})/4$$
$$= ([1 + f_A]/2 + \theta_{AB} + \theta_{AC} + \theta_{BC})/4$$

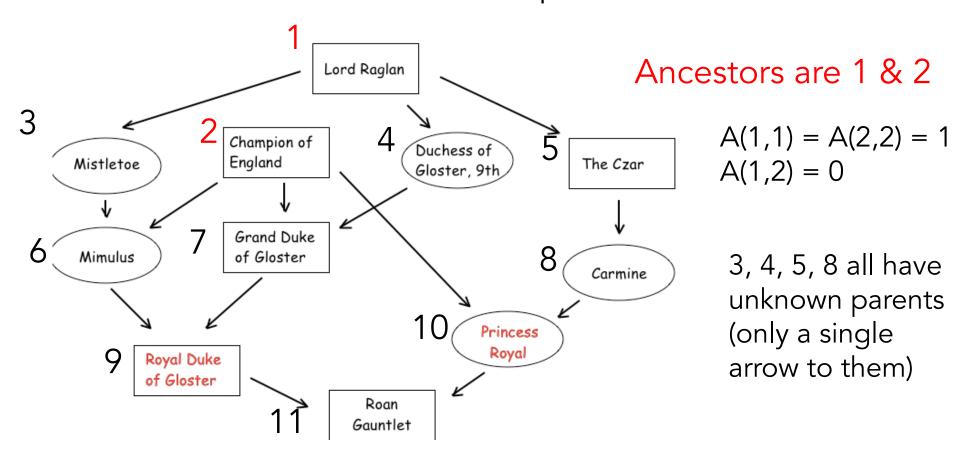
Hence, if B and C unrelated,

$$\theta_{EF} = (1 + f_A)/8$$

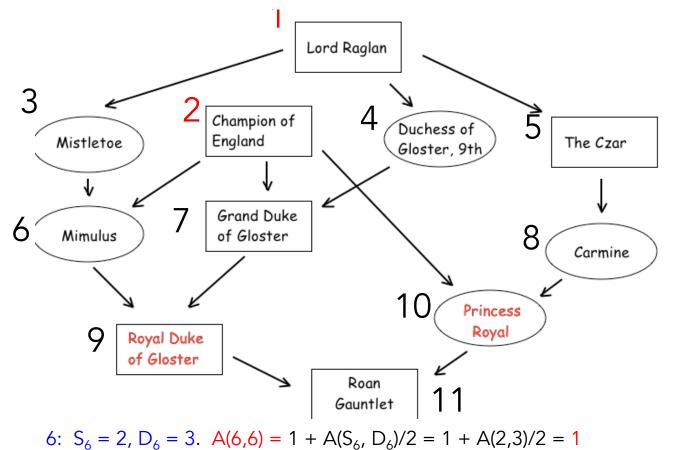
### Computing $\theta_{xy}$ -- The Recursive Method

- There is a simple recursive method for generating the elements  $A_{ij}$  = 2  $\theta_{ij}$  of a relationship matrix (used for BLUP selection). For ease of reading, we use the notation  $A(i,j) = A_{ij}$ 
  - Basic idea is that the founding individuals of the pedigree are assumed to be unrelated and not inbred (although this can also be accommodated). These founders are assigned values of A(i,i) = 1.
  - Likewise, any unknown parent of any future individual is assumed to be unrelated to all others in the pedigree and not inbred, and they are also assigned a value of A(i,i) = 1.
  - Let  $S_i$  and  $D_i$  denote the sire and dam (father and mother) of individual i. For this offspring  $A(i,i) = 1 + A(S_i, D_i)/2$
  - $\ \ A(i,j) = A(j,i) = [A(j,S_i) + A(j,D_i)]/2 = [A(i,S_i) + A(i,D_i)]/2$
  - The <u>recursive</u> (or <u>tabular</u>) method starts with the founding parents and then proceeds down the pedigree in a recursive fashion to fill out A for the desired pedigree.

#### Example



3:  $S_3 = 1$ ,  $D_3 = Unknown$ ,  $A(3,3) = 1 + A(S_3,D_3)/2 = 1 + A(1,unk)/2 = 1$   $A(1,3) = [A(1,S_3) + A(1,D_3)]/2 = [A(1,1) + A(1,unk)]/2 = 1/2$ . Note also that A(1,4) = A(1,5) = 1/2, A(4,4) = A(5,5) = 1.  $A(3,4) = [A(3,S_4) + A(3,D_4)]/2 = [A(3,1) + A(3,unk)]/2 = (1/2+0)/2 = 1/4$ . Same for A(3,5) = 1/4. 2 is unrelated to 3, 4, 5, giving A(2,3) = A(2,4) = A(2,5) = 0.



#### So far

```
6: S_6 = 2, D_6 = 3. A(6,6) = 1 + A(S_6, D_6)/2 = 1 + A(2,3)/2 = 1

A(6,1) = [A(1, S_6) + A(1, D_6)]/2 = [A(1,2) + A(1,3)]/2 = [0 + 1/2]/2 = 1/4

A(6,2) = [A(2, S_6) + A(2, D_6)]/2 = [A(2,2) + A(2,3)]/2 = [1 + 0]/2 = 1/2

A(6,3) = [A(3, S_6) + A(3, D_6)]/2 = [A(3,2) + A(3,3)]/2 = [0 + 1]/2 = 1/2

A(6,4) = [A(4, S_6) + A(4, D_6)]/2 = [A(4,2) + A(4,3)]/2 = [0 + 1/4]/2 = 1/8

A(6,5) = [A(5, S_6) + A(5, D_6)]/2 = [A(5,2) + A(5,3)]/2 = (0 + 1/4)/2 = 1/8

7: S_7 = 2, D_7 = 4. A(7,7) = 1 + A(S_7, D_7)/2 = 1 + A(2,4)/2 = 1 + 0/2 = 1

A(6,7) = [A(6, S_7) + A(6, D_7)]/2 = [A(6, 2) + A(6, 4)]/2 = (1/2 + 1/8)/2 = 5/16

8: S_8 = 5, D_8 = \text{unk}. A(8,8) = 1 + A(S_8, D_8)/2 = 1 + A(5,\text{unk})/2 = 1.

A(6,8) = [A(6, S_8) + A(6, D_8)]/2 = [A(6, 5) + A(6,\text{unk})]/2 = (1/8)/2 = 1/16

9: S_9 = 7, D_9 = 6. A(9,9) = 1 + A(S_9, D_9)/2 = 1 + A(6,7)/2 = 1 + 5/32 = 1.156 < - \text{inbred!}
```

# Actual relatedness versus expected values from pedigrees

Values for the coefficient of coancestry ( $\theta$ ) and the coefficient of fraternity ( $\Delta$ ) obtained from pedigrees are <u>expected values</u>. Due to random segregation of genes from parents, The actual value (or realization) can be different.

For example, we expect  $2\theta$  to be ½ for full subs. However, one pair of sibs may actually be more similar (0.6) and another less similar (say 0.35). On average,  $2\theta$  is ½ for pairs of full sibs, but if we knew the <u>actual value</u> of  $\theta$ , we have more information. With sufficient dense genetic markers, we can estimate these relationships directly.

Genomic selection uses this extra information.

#### What about coefficient of coancestry $\theta$ ?

Genotype of $j$	Genotype of $i$				
	11	10	00		
11	1	0.5	0		
10	0.5	0.5	0.5		
00	0	0.5	1		

One computes the coefficient of coancestry for each SNP, taking the average value over all loci as the coefficient of coancestry for that pair of individuals. Toro et al. (2002) refer to this as **molecular coancestry**. Note that we can compare an individual with itself (i = j), which returns 1 for each homozygous locus and 1/2 for each heterozygous loci.

					Genot	ype of $i$				
	Genoty	pe of $j$		11		10		00		
		11		1		0.5		0		
		10		0.5		0.5		0.5		
		00		0	(	0.5		1		
Indiv x:	00	00	10	10	00	10	11	00	11	00
Indiv y:	10	00	11	11	10	11	11	10	11	10
Locus-specific $\theta$	0.5	1.0	0.5	0.5	0.5	0.5	1.0	0.5	1.0	0.5

Estimated  $\theta$  is the average over all ten loci, = 0.65

#### The coefficient of fraternity

- While (twice) the coefficient of coancestry gives the weight on the additive variance for two relatives, a related measure of IDB status among relatives gives the weight on the dominance variance
- The probability that the two alleles in individual x are IBD to two alleles in individual y is denoted  $\Delta_{xy}$ , and is called the coefficient of fraternity.
- This can be expressed as a function of the coefficients of coancestry for the parents of (mx and fx) of x and the parents (my and fy) of y.

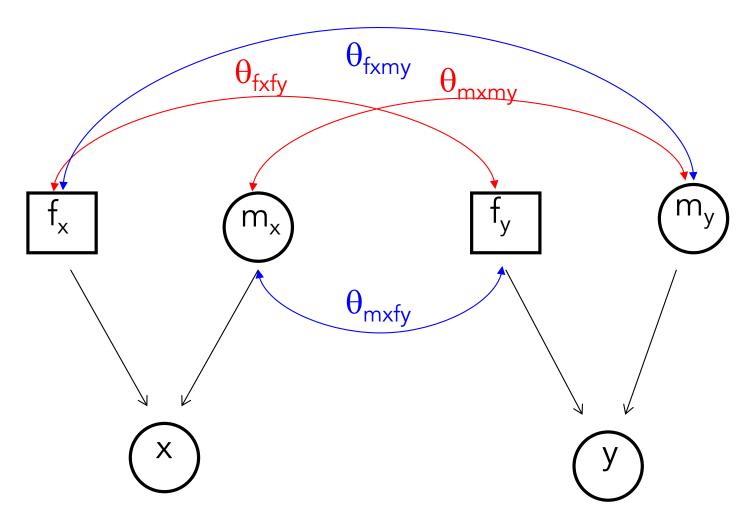
$$- \Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy}$$

#### The coefficient of fraternity (cont)

- x and y can have both alleles IBD if
  - The allele from the father (fx) of x and the father (fy) of y are IDB (probability  $\theta_{fxfy}$ ) AND the allele from the mother (mx) of x and the mother (my) of y are IDB (probability  $\theta_{mxmy}$ ), or  $\theta_{fxfy}$   $\theta_{mxmy}$
  - OR the allele from the mother (mx) of x and the father (fy) of y are IDB (probability  $\theta_{mxfy}$ ) AND the allele from the father (fx) of x and the mother (my) of y are IDB (probability  $\theta_{fxmy}$ ), or  $\theta_{mxfy}$   $\theta_{fxmy}$
  - Putting these together gives
    - $\Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy}$

#### $\Delta_{xy}$ , The Coefficient of Fraternity

 $\Delta_{xy}$  = Prob(both alleles in x & y IBD)



$$\Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy}$$

## Examples of $\Delta_{xy}$ : Full sibs

• Full sibs share same mon, dad

- 
$$m_x = m_y = m$$
,  $f_x = f_y = f$   
-  $\Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy} = \theta_{mm}\theta_{ff} + \theta_{mf}^2$   
-  $\Delta_{xy} = (1+f_m)(1+f_f)/4 + \theta_{mf}^2$ 

- If parents unrelated,  $\theta_{fm} = 0$ , giving
  - $-\Delta_{xy} = (1+f_m)(1+f_f)/4$
- If parents are unrelated and not inbred,

$$-\Delta_{xy} = 1/4$$

### Examples of $\Delta_{xy}$ : Half sibs

 Paternal half sibs share same dad, different moms

$$- f_x = f_y = f; m_x \text{ and } m_y$$

$$- \Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy} = \theta_{mxmy}\theta_{ff} + \theta_{mxf}\theta_{myf}$$

$$- \Delta_{xy} = \theta_{mxmy} (1+f_m)/2 + \theta_{mxf}\theta_{myf}$$

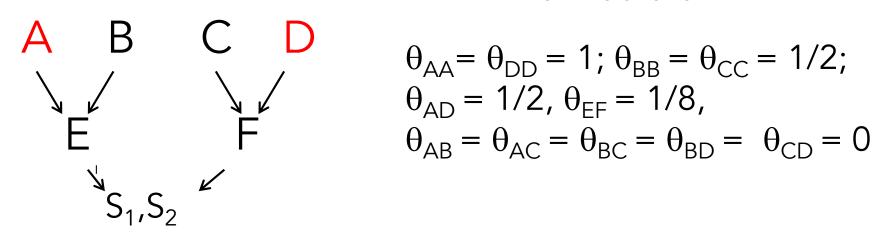
• If mothers are unrelated to each other and to the common father,  $\theta_{mxmy} = \theta_{mxf} = \theta_{myf} = 0$ , giving

$$-\Delta_{xy}=0$$

#### When is $\Delta$ non-zero?

- Since  $\Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy}$
- ullet A nonzero value for  $\Delta$  requires either
  - That the fathers of both x and y are related AND the mothers of both x and y are related
  - OR that the father of x is related to the mother of y AND the mother of x is related to the father of y

#### From before



What is  $\Delta$  for the full sibs (S<sub>1</sub> and S<sub>2</sub>)?

$$\Delta_{xy} = \theta_{mxmy}\theta_{fxfy} + \theta_{mxfy}\theta_{fxmy} = \theta_{EE}\theta_{FF} + \theta_{EF}^{2}$$
Giving  $\Delta_{xy} = \theta_{EE}\theta_{FF} + \theta_{EF}^{2}$ 

$$= (1/2)(1/2) + (1/8)^{2}$$

$$= 1/4 + 1/64 = 17/64 = 0.266$$

#### $\Delta_{xy}$ and the coefficient on $V_D$

- The coefficient on the dominance variance for the relatives x and y is just  $\Delta_{xv}$ .
- To see this,
  - let  $A_iA_j$  denote the two alleles in x and  $A_kA_l$  those in y.
  - Suppose that alleles i and k come from the mothers of these two relatives and alleles j and l from their fathers.
  - Cov(dominance values) =  $Pr(A_i \text{ ibd } A_k, A_j \text{ ibd } A_l)$  $cov(\delta_{ij}, \delta_{kl}) + Pr(A_i \text{ ibd } A_l, A_j \text{ ibd } A_k)cov(\delta_{ij}, \delta_{kl})$
  - $= (\theta_{fxfy}\theta_{mxmy} + \theta_{mxfy}\theta_{jxmy}) Var(D) = \Delta_{xy} Var(D)$

# Estimating relationships using molecular data

With SNP data, treat identity in state (also called alike in state, AIS) as IBD

Suppose the genotypes of two individual at 10 SNPs are

3/10 loci have  $\Delta_{xy} = 1$ , so average  $\Delta_{xy}$  over all loci is  $0.3*\ 1 = 0.3$ 

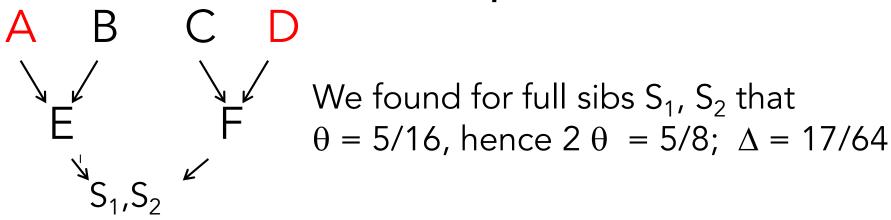
## General Resemblance between relatives

$$2\theta_{xy} = r_{xy}, \qquad u_{xy} = \Delta_{xy}$$

$$Cov(G_x, G_y) = 2\theta_{xy}V_A + \Delta_{xy}V_D$$

$$Cov(G_x, G_y) = 2\theta_{xy}V_A + \Delta_{xy}V_D + (2\theta_{xy})^2V_{AA} + 2\theta_{xy}\Delta_{xy}V_{AD} + \Delta_{xy}^2V_{DD} + \cdots$$

#### Example



Expected genetic covariance between this sibs is

$$(5/8)$$
Var(A) +  $(17/64)$ Var(D) +  $(5/8)^2$ Var(AA) +  $(5/8)(17/64)$ Var(AD) +  $(17/64)^2$ Var(DD) + ...

#### Autotetraploids

- Peanut, Potato, alfalfa, soybeans all examples of crops with at least some autotetraploid lines
- With autotetraploid, four alleles per locus, with a parent passing along two alleles to an offspring
- As a result, a parent can pass along the dominance contribution in G to an offspring
- Further, now there are four variance components assocated with each locus

# Genetic variances for autotetraploids

- G = A + D + T + Q
  - A (additive) and D (dominance, or digenic effects) as with diploids
  - T (trigenic effects) are the three-way interactions among alleles at a locus
  - Q (quadrigenic effects) are the four-way interactions at a locus
- Total genetic variance becomes

$$-V_G = V_A + V_D + V_T + V_Q$$

# Resemblance between autotetraploid relatives

Relatives	V <sub>A</sub>	$V_{D}$	$V_{T}$	VQ
Half-sibs	1/4	1/36		
Full-sibs	1/2	2/9	1/12	1/36
Parent- offspring	1/2	1/6		

Assumes unrelated, non-inbred parents