APPENDIX A

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NUCLEIC ACID CODES, AMINO ACID CODES, AND GENETIC CODES

Table 6. Nucleic acid	d codes	~ (i)	Side
202	Code	The of	Description
Q	А		Adenine
II	G		Guanine
	С		Cytosine
	Т		Thymine
	U	6062	Uracil
	R		Purine (A or G)
	Y	TINIT	Pyrimidine (C or T)
	N	UNI	Any nucleotide
	W		Weak (A or T)
	S		Strong (G or C)
ລີບສຶກຣົນ	M	Sner	Amino (A or C) Keto (G or T)
CIUCIIIUU	K	JIO	Keto (G or T)
Copyright [©]	B	Chian	Not A (G or C or T) Not G (A or C or T)
All ri	B V	ts	Not C (A or G or T) Not T (A or G or C)

Table 7. Amino acid cod	des 9121%	Ø 91-
1-letter cod	le 3-letter code	Description
A	Ala	Alanine
R	Arg	Arginine
N N	Asn	Asparagine
D	Asp	Aspartic acid
C	Cys	Cysteine
Q Sold R	Gln (?)	Glutamine
E	Glu	Glutamic acid
G	Gly	Glycine
Н	His	Histidine
Ι	Ile	Isoleucine
L	Leu	Leucine
K	Lys	Lysine
М	Met	Methionine
F	Phe	Phenylalanine
Р	Pro	Proline
S	Ser	Serine
Т	Thr	Threonine
W	Trp	Tryptophan
ขสิทธิ์มูห	Tyr	Tyrosine Valine
	Asx	Asn or Asp
opyngnu z	Oy Gix Ian	g Gln or Glu UNIVERSI
J	Xle	Leu or Ile
	Sec	Selenocysteine (UGA)
0	Pyl	Pyrrolysine (UAG)
X	Unk	Unknown

Table 8.	Standard	genetic	code
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		0	181	29	HØ	9			
Table 8.	Standard	d geneti	c code		0		526		
1 st				2 nd pc	osition				3 rd
position	ť	J	C	Ċ,			(3	position
	UUU	Phe	UCU	Ser	UAU	Try	UGU	Cys	U
	UUC	Phe	UCC	Ser	UAC	Try	UGC	Cys	С
	UUA	Leu	UCA	Ser	UAA	Stop	UGA	Stop	A
000	UUG	Leu	UCG	Ser	UAG	Stop	UGG	Trp	G
	CUU	Leu	CCU	Pro	CAU	His	CGU	Arg	U
	CUC	Leu	CCC	Pro	CAC	His	CGC	Arg	C
С	CUA	Leu	CCA	Pro	CAA	Gln	CGA	Arg	А
	CUG	Leu	CCG	Pro	CAG	Gln	CGG	Arg	G
	AUU	Ile	ACU	Thr	AAU	Asn	AGU	Ser	U
	AUC	Ile	ACC	Thr	AAC	Asn	AGC	Ser	С
А	AUA	Ile	ACA	Thr	AAA	Lys	AGA	Arg	А
	AUG	Met	ACG	Thr	AAG	Lys	AGG	Arg	G
	GUU	Val	GCU	Ala	GAU	Asp	GGU	Gly	U
	GUC	Val	GCC	Ala	GAC	Asp	GGC	Gly	C
G	GUA	Val	GCA	Ala	GAA	Glu	GGA	Gly	A
	GUG	Val	GCG	Ala	GAG	Glu	GGG	Gly	G
Copyri	ght [@]	\mathcal{O}	by C	Chia	Ing	Mai	Ur	nive	rsity
	ri	g	h t	S	ľ	e s	e	r v	e d

APPENDIX B

กมยนต์

ANTIRETROVIRAL DRUGS FOR TREATING PREGNANT WOMEN AND PREVENTING HIV INFECTION IN INFANTS: TOWARDS UNIVERSAL ACCESS

RECOMMENDATIONS FOR A PUBLIC HEALTH APPROACH

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Ranking	Time of administration			Advantages	Disadvantages
	Pregnancy	Labour	Postpartum	140 91	
Recommended	AZT (≥28 weeks gestation	Sd-NVP ^a + AZT/3TC	Mother: AZT/3TC x 7 days ^a Infant: Sd- NVP + AZT x 7 days ^b	 Highly effective regimen Substantially reduces <i>in utero</i> and intrapartum transmission The AZT/3TC tail given to the mother reduces the development of her becoming resistant to NVP AZT given to infants reduces the risk of resistance to NVP in those who become infected 	- Longer and more complex than other regimens
Alternative	AZT (>28 weeks gestation	Sd-NVP	Infant: Sd- NVP + AZT x 7 days ^b	 Highly effective regimen Substantially reduces <i>in utero</i> and intrapartum transmission AZT given to infants reduces the risk of resistance to NVP in those who become infected 	 High risk of resistance to NVP Probable sub-optimal viral response if NNRTI-ART is initiated in women within 6 months of childbirth
	r	ig	hts	rese	erved

Table 9. Different approaches to the use of ARV prophylaxis to prevent HIV infection in infants (WHO, 2006 [459])

Ranking	Tim	Time of administration		Advantages	Disadvantages	
	Pregnancy	Labour	Postpartum			
Minimum	6	Sd-NVP + AZT/3TC	Mother: AZT/3TC x 7 days Infant: Sd- NVP	 Effective in reducing MTCT The AZT/3TC tail given to the mother reduces the development of her becoming resistant to NVP 	 Less effective than recommended regimen Does not reduce <i>in utero</i> transmission More complex to deliver than Sd-NVP alone 	
Minimum	<u>.</u>	Sd-NVP	Infant: Sd- NVP	 Effective in reducing MTCT Simplest regimen to administer 	 Less effective than recommended regimen Does not reduce <i>in utero</i> 	
	E.				 transmission High risk of resistance to NVP 	
		G M.		NIVERSI	- Probable sub-optimal viral response if NNRTI- ART is initiated in women within 6 months of childbirth	

^a If the women receives at least four weeks of AZT during pregnancy, omission of the NVP does for mothers may be considered. In this case the NVP does must be given to the infant immediately after birth, AZT is recommended for four weeks instead of one week, and the mother will not require 3TC during labour as well as AZT and 3TC postpartum.

^b If the mother receives less than four weeks of AZT during pregnancy, AZT is recommended for four weeks instead of one week.

 Table 10. ARV prophylaxis regimens for PMTCT among pregnant women living with HIV who have not received antepartum therapy or prophylaxis (WHO, 2006 [459]).

Ranking	Time of administration		Advantages	Disadvantages
	Labour	Postpartum		630
Recommended	Sd-NVP +	Mother:	- Sd-NVP is effective in reducing	- More complex to deliver than Sd-
	AZT/3TC	AZT/3TC x 7	MTCT	NVP alone
	D's	days	- The AZT/3TC tail given to the	-3026
5		Infant: Sd-	mother reduces the development of	
Ű		NVP + AZT x	her becoming resistant to NVP	
	21	4 weeks ^a	- In breastfeeding women, NVP-	5
			based regimen may be	5
			advantageous in reducing early postpartum transmission	\rightarrow
		M	- Consistent with recommended	
		A	regimen for PMTCT when mother	
			receives antepartum prophylaxis	
ຄິບສິ	ทธิ์เ	มหา	- AZT given to infants reduces the risk of resistance to NVP in those	ชียอใหม่
Сору	right ⁽	© by	who become infected	University
	l i	i g h	ts res	erved

Ranking	Time of adr	Time of administration Advantages		Disadvantages
	Labour	Postpartum		
Alternative	AZT/3TC	Mother: AZT/3TC x 7 days Infant: AZT/3TC x 7 days	 Equivalent effecacy to Sd-NVP alone intrapartum/postpartum No risk of resistance to NVP in women or infants should they become infected 	- More complex to deliver than Sd- NVP alone
Minimum	Sd-NVP + AZT/3TC	Mother: AZT/3TC x 7 days Infant: Sd- NVP ^a	 Sd-NVP is effective in reducing MTCT The AZT/3TC tail given to the mother reduces the development of resistance to NVP 	- More complex to deliver than Sd- NVP alone
Minimum	Sd-NVP	Infant: Sd- NVP	 Single-dose NVP is effective in reducing MTCT Simplest regimen to administer 	- High risk of resistance to NVP, with sub-optimal viral response if NNRTI-based ART is initiated in women within 6 months of childbirth

^a Data on added efficacy of four weeks of infant AZT in this situation are limited.

183

Table 11. ARV prophylaxis regimens for infants born to women living with HIV who have not received antepartum or intrapartum therapy or prophylaxis (WHO, 2006 [459]).

Ranking	Time of administration	Advantages	Disadvantages
Recommended	Infant:	- Sd-NVP + AZT given to the infant is more	0.0.
	Sd-NVP immediately	effective in reducing MTCT than just Sd-NVP	than Sd-NVP alone
	after birth + AZT \times 4	- Consistent with recommended regimen for PMTCT	
	weeks ^a	when mother receives antepartum or intrapartum	
, ÇZ		 prophylaxis AZT given to the infant reduces his/her risk of becoming resistant to NVP 	
Alternative	Infant:	- Clinical trial data demonstrate that Sd-NVP + AZT	- More complex to deliver
	Sd-NVP immediately	for one week given to the infant is more effective	than Sd-NVP alone
	after birth + AZT \times 1	in reducing MTCT than just Sd-NVP	
	week	- AZT given to the infant reduces his/her risk of	
		becoming resistant to NVP	
Minimum	Infant:	- Prophylaxis with Sd-NVP for the infant is	- Risk of resistance to NVP in
	Sd-NVP immediately	equivalent to six weeks of AZT	infants who become infected
	after birth	- Simplest regimen to administer	despite NVP prophylaxis

^a NVP administered immediately after birth, if possible within 12 hours after delivery, is likely to result in a larger reduction in transmission than starting it later. Data on added efficacy of four weeks of AZT for infants in this situation are limited.

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188

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