Form Approved: OMB No. 0910-0045. Expiration Date: December 31, 2007. See OMB Statement on Reverse.

DEPARTMENT OF HEALTH AND HUMANS SERVICES
FOOD AND DRUG ADMINISTRATION

DRUG PRODUCT LISTING

(In accordance with Public Law 92-387)

NAME AND ADDRESS OF FIRM
Physician Therapeutics
2980 Beverly Glen Circle, Suite 301
Los Angeles, CA 90077

LABELING REVISON
CHANGE OF:
☐ RTE OF ADMIN ☐ INDICATION
☐ NAME / DOSE / STR / INGR
☐ OTHER *(Specify)*

FOR FDA USE

CONTROL NO. (1–5)	RECORD ID (6–11)	(12–15)

SEC (16–17)	S (18)	U (19)	PRODUCT TRADE NAME OR CATALOG NAME (20–83)	NATIONAL DRUG CODE LABELER (84–89)	PRODUCT (90–93)
01			HYPERTENSA	06840	5100 7

FDA APPLICATION NO. (94–99)	REPORT DATE MO (100–101)	DA (102–103)	YR (104–105)	TYPE RESET (106)	TYPES OF BUSINESS (107–111)	PRODUCT TYPE (112–116)	BNDD (117)	LEGAL STATUS (118)	SCHED (119)	PROF USE (120)	PRODUCT DISCONTINUED (121–125)
	05	28	08	M	OTHER *(Specify)* Manufacturer	OTHER *(Specify)* Medical Food		R		P	

PRODUCT DISCONTINUED — OTHER *(Specify)*

BASIS OF CONCENTRATION

WHOLE NUMBERS (126–133)	DECIMAL (134–137)	UNIT (138–140)
600		

DOSAGE FORM (141–143)	ROUTES OF ADMINISTRATION 1 (144–147)	2 (148–151)	3 (152–155)	OTHER (156)	PT (157–158)	SEC (16–17)	S (18)	U (19)	SMP (20)	PKG CODE (21–22)	PACKAGE SIZE (23–47)	PACKAGE TYPE (48–72)
600	001					03				*3	90	BOT
						03						
						03						
						03						
						03						

INITIAL MARKETING DATE MO (159–160)	YEAR (161–164)	MOST RECENT MARKETING DATE MO (165–166)	YEAR (167–170)	DISCONTINUED DATE MO (171–172)	YEAR (173–176)
04	2004	08	2007		

NOTICE: This report is required by law (21 C.F.R. 207.20). Failure to report can result in imprisonment for not more than one year or a fine of not more than $1,000, or both (FDA&C Act, Section 303).

SEC (16–17)	S (18)	U (19)	TYPE (20)	PT (21–22)	ESTABLISHED NAME OF PRODUCT AND / OR INGREDIENT(S) OR BIOLOGIC PROPER NAME, TEST OBJECTIVE / EQUIPMENT / REAGENT NAME, ETC. (44–107)	FDA USE ONLY INGREDIENT NO. (23–28)	AMOUNT WHOLE NUMBER (29–36)	DECIMAL (37–40)	UNIT (41–43)
05			A		L-ARGININE		63		MG
05			A		CHOLINE BITARTRATE		47		MG
05			A		WHEY PROTEIN HYDROLYSATE		37		MG
05			A		COCOA EXTRACT		20		MG
05			A		CINNAMON		20		MG
05			A		GINSENG		16		MG
05			A		L-LEUCINE		16		MG
05			A		L-GLUTAMINE		16		MG
05			A		L-HISTIDINE		16		MG
05			A		CAFFEINE		14		MG

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[illegible] 2007
DRLS

SEC (16–17)	S (18)	U (19)	SITE OR FIRM ESTABLISHMENT REGISTRATION NUMBER (20–26)	ACTUAL MANUFACTURING SITE OF THE ABOVE DRUG PRODUCT (27–66)	STATE (67–68)	FOREIGN COUNTRY (69–78)	NDC LABELER CODE (79–84)	SHORT NAME (85–99)
07			0825991	ARIZONA NUTRITIONAL CHANDLER	AZ		~~068405~~ 068223	
07								
07								