



A.S. Research IIc Grovetown, GA 30813 PO#:

PRODUCT NAME: AS Research T Drive 60 capsules

PRODUCT CODE: ASRTdrive60caps

DATE OF MANUFACTURE: 12/31/2014

LOT NUMBER: PB0106714

BATCH YIELD (Units): 2,750

EXPIRATION DATE: 12/31/2017

PRODUCT SPECIFICATIONS

POWDER APPEARANCE: Tanish Powder

AVG. EMPTY CAPSULE WEIGHT (mg.): 120

THEORETICAL POWDER FILLWEIGHT (mg.): 800.00

WEIGHT VARIATION (mg.): 828.00 - 1,012.00 FOR EACH

90% - 110%

OF 30 CAPSULES

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CONFORMS

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AVERAGE: 843.00 mg.

RANGE: 829.00 - 860.00 mg.

SPECIFICATION	RESULT	METHOD OF REF
1.00 each	1.00 each	BY INPUT
3.00 gram	3.00 gram	BY INPUT
1.00 each	1.00 each	BY INPUT
0.75 gram	0.75 gram	BY INPUT
1.00 each	1.00 each	BY INPUT
6.00 gram	6.00 gram	BY INPUT
3.00 gram	3.00 gram	BY INPUT
2.25 gram	2.25 gram	BY INPUT
16.50 gram	16.50 gram	BY INPUT
3.00 gram	3.00 gram	BY INPUT
3.00 gram	3.00 gram	BY INPUT
1.50 gram	1.50 gram	BY INPUT
9.00 gram	9.00 gram	BY INPUT
60.00 each	60.00 each	BY INPUT
	1.00 each 3.00 gram 1.00 each 0.75 gram 1.00 each 6.00 gram 3.00 gram 2.25 gram 16.50 gram 3.00 gram 3.00 gram 9.00 gram	1.00 each 1.00 each 3.00 gram 3.00 gram 1.00 each 1.00 each 0.75 gram 0.75 gram 1.00 each 1.00 each 6.00 gram 6.00 gram 3.00 gram 3.00 gram 2.25 gram 16.50 gram 3.00 gram 3.00 gram 3.00 gram 3.00 gram 1.50 gram 1.50 gram 9.00 gram 9.00 gram



DOCID: 7406

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White Button Mushroom Extract 10:1 (Agaricus bisporus) 3.00 gram (Fruit Body)

3.00 gram

BY INPUT

All Batch Formulations are based on a validated Master Manufacturing Record in accordance with subpart H of Title 21 CFR 111. Validition is performed by creating an FTIR signature baseline of 3 unique batches/lots of the formula; all future formulas are tested via FTIR Spectroscopy for conformity to the MMR batch signatures. All baselines are revalidated annually. Copies of the FTIR Signature and Batch Production Record are available upon request.

I certify that the above referenced product has been manufactured by LLC in accordance with FDA mandated Title 21 CFR Part 111 -current Good Manufacturing Practices in Manufacturing, Packaging, Labeling and Holding Operations for Dietary Supplements; Final Rule.

QC Supervisor Comments:

RELEASED BY:

12/31/2014

DATE