

PRODUCT SPECIFICATIONS

Revision N.: 04 (September 2021)

ABG10+[®]



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|----------------------|---|
| Scientific name: | <i>Allium sativum</i> |
| Product code: | GAR0213066 |
| Country of origin: | Spain |
| Plant part used: | Bulb |
| Composition: | Native extract and dextrin |
| Extraction solvents: | Water |
| Appearance: | Natural brown powder with characteristic odor and taste |

| ASSAY | METHOD | SPECIFICATION |
|--|--|-----------------------------------|
| Identification ⁽¹⁾ | HPLC-DAD / TLC | Positive |
| S-Allyl Cysteine ⁽¹⁾ | HPLC-DAD | > 0.1% |
| Bulk density ⁽¹⁾ | Eur. Pharm. (2.9.34) / USP <616> | > 0.3 g/mL |
| Ash ⁽¹⁾ | Eur. Pharm. (2.4.16) / USP 37 <281> | < 10% |
| Loss on drying ⁽¹⁾ | Eur. Pharm. (2.8.17) / USP 37 <731> | < 8% |
| Particle size ⁽¹⁾ | Eur. Pharm. (2.9.12) / USP 34 <786> | 240 µm / 60 mesh |
| MICROBIOLOGY | | EC 2073/2005 & Eur. Pharm. 5.1.8* |
| TAMC ⁽¹⁾ | | < 10 000 cfu/g |
| TYMC ⁽¹⁾ | | < 100 cfu/g |
| Enterobacteria ⁽²⁾ | Eur. Pharm. (2.6.12 / 2.6.13 / 2.6.31) | < 100 cfu/g |
| <i>L. monocytogenes</i> ⁽²⁾ | USP 37 <61 / 62> | Absent/25g |
| <i>E. coli</i> ⁽²⁾ | | Absent/1g |
| <i>S. aureus</i> ⁽²⁾ | | Absent/1g |
| <i>Salmonella</i> spp ⁽²⁾ | | Absent/25g |
| HEAVY METALS | | EC 1881/2006 |
| Lead (Pb) ⁽²⁾ | | < 3.0 ppm |
| Arsenic (As) ⁽²⁾ | Eur. Pharm. (2.4.27) / USP <233> | < 1.0 ppm |
| Cadmium (Cd) ⁽²⁾ | | < 1.0 ppm |
| Mercury (Hg) ⁽²⁾ | | < 0.1 ppm |
| PAHs | | EC 1881/2006 |
| BaP ⁽²⁾ | | < 10 ppb |
| Sum BaP, BaA, BbF, CHR ⁽²⁾ | GC-MS/MS | < 50 ppb |
| AFLATOXINS | | EC 1881/2006 |
| B1 ⁽²⁾ | | < 5 ppb |
| Sum B1, B2, G1, G2 ⁽²⁾ | HPLC-MS | < 10 ppb |
| PESTICIDES | | EC 396/2005 |
| Pesticide residues ⁽²⁾ | LC & GC - MS/MS | According to regulation |
| OTHER CONTAMINANTS | | EC 1881/2006 |
| Melamine ⁽²⁾ | LC-MS/MS | < 2.5 ppm |

- Packaging: food grade LDPE or PA/PE bags.
- Shelf life: 24 months if stored closed in the original container at room temperature (<25°C), sheltered from light and moisture (<60% RH).
- Control plan frequency: ⁽¹⁾ analyzed on each production batch, ⁽²⁾ analyzed externally once a year, on the ingredient or the raw materials.
- *According to Eur. Pharm., products must meet the acceptance criteria or at least the maximum acceptable count tolerated described on this monograph.
- Natural variations in the raw material may lead to color variations from batch to batch but without affecting the quality and efficacy of the product.