

1. Product Information

| Product name | pBMY-AAV6 | |
|---------------------|---|--|
| Catalog Nr. | 506-01-GMP / 506-05-GMP / 506-10-GMP | |
| Quality grade | Good Manufacturing Practice (GMP) | |
| Description | DNA plasmid encoding Rep2/Cap6 genes of adeno-associated virus (AAV), intended for preparation of AAV particles of serotype 6 | |
| Plasmid Size | 6 107 bp | |
| Molecular weight | 3 774 kDa | |
| Concentration | 1.0 ± 0.2 mg/mL solution in buffer (frozen) | |
| Storage buffer | 10 mM Tris-HCl, 1 mM EDTA, pH 8.0 (TE buffer) | |
| Storage temperature | -20 ± 5 °C | |
| Lot. Nr. | Specified on product label | |
| Use by date | Specified on product label | |
| Manufacturer | Biomay AG, Ada-Lovelace-Straße 2; 1220 Vienna, Austria; www.biomay.com; info@biomay.com | |

2. Description

AAV Rep/Cap plasmid encoding p5 promoter and Rep genes of AAV serotype 2 (NCBI Reference: NC 001401.2) and Cap gene from AAV serotype 6 (NCBI Reference: AF028704.1). The plasmid was propagated by cultivation in Escherichia coli, purified by chromatographic methods, 0.2 µm filtered and filled as low bioburden product. Sequence information as .gb or .fasta-file can be provided upon request.

3. Intended Use / Application

GMP grade: product is a DNA plasmid that has been manufactured and quality-controlled under the conditions of Good Manufacturing Practice (GMP). It has been certified and released by a Qualified Person (QP) under EMA law (EMA directive 2001/83/EC).

It was designed and is intended to be used in conjunction with an adenoviral (Ad) helper plasmid and an AAV transgene plasmid to transfect HEK293 cells, in order to manufacture adeno-associated virus (AAV) particles.

The quality of this product is considered appropriate to serve as a starting plasmid for the GMP manufacturing of clinical-grade AAV virus particles which may be used as gene transfer vector for human clinical trials.

4. Safety information

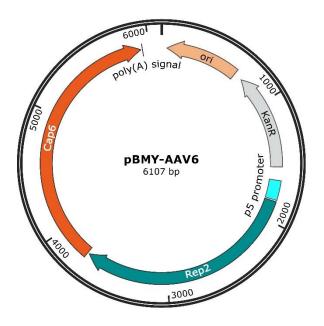
Material is considered non-infectious, non-toxic and non-pathogenic under the conditions of the intended applications. General safety procedures should still be followed to maintain a safe working environment. Always wear appropriate personal protective equipment (PPE), including lab coats, gloves, and safety glasses, to avoid contamination and accidental exposure. Work in a clean, organized space, and handle reagents with care, avoiding direct contact. Dispose of all waste materials, including gloves and pipette

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tips, in designated biohazard containers, even though the plasmid DNA is non-hazardous, to prevent cross-contamination. Always wash hands after handling any biological material and before leaving the lab.

5. Plasmid Map



6. Testing Specifications

| Parameter | Method | Specification GMP-grade |
|--|--|--|
| Appearance | Visual inspection | Clear and colourless solution; free from visible particulates |
| pH value | pH measurement | 8.0 ± 0.5 |
| pDNA homogeneity (% supercoiled) | Anion exchange HPLC | > 85 % supercoiled |
| DNA concentration | UV spectrophotometry (A ₂₆₀) | 1.0 ± 0.2 mg / mL |
| Purity based on A ₂₆₀ /A ₂₈₀ ratio | UV spectrophotometry (ratio A ₂₆₀ /A ₂₈₀) | 1.8 – 2.1 |
| Identity / Integrity | Restriction digest & agarose gel electrophoresis | Major bands conform to theoretical fragments (expected band pattern) |
| pDNA sequence | DNA sequencing (complete sequence) | Conforms to reference sequence |
| Residual host cell DNA | qPCR | < 1.0 % (< 10 μg / mg DNA) |
| Residual host cell RNA | Fluorometry | < 2.0 % (< 20 μg / mg DNA) |
| Residual host cell protein | Colorimetry | < 1.0 % (< 10 μg / mg DNA) |
| Endotoxin | Kinetic chromogenic LAL test (Ph. Eur. 2.6.14 / USP <85>) | ≤ 5.0 EU / mg DNA |
| Bioburden | Testing for bacteria, yeasts & moulds (Ph.Eur.2.6.12/USP <61>) | < 1 cfu / mL |

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