

## **Certificate of Analysis**

COA No: CA\_XBN-0006

Version: 10

## dATP 100mM

Suitable for Research and further Manufacturing Use

| Catalog No:         | MDX046        |
|---------------------|---------------|
| Lot No:             | NU060-B122540 |
| Storage Conditions: | -20°C         |
| Component Lot No:   | DA-223111A    |
| Expiry date:        | December 2025 |

## **Quality Control Parameters**

2'-deoxyadenosine-5'-triphosphate  $C_{10}H_{12}N_5O_{12}P_3Li_4$  MW = 514.916 g/mol

Certified <1% deoxynucleoside monophosphates and deoxynucleoside diphosphates

| Characteristics  | Specification             | Result        |
|--|---------------------------|---------------|
| Concentration (at $\lambda$ max, pH 7.0, $\epsilon$ = 15.4 E x mmol <sup>-1</sup> x cm <sup>-1</sup> ) | 100 mM ± 5%               | 101.22 mM     |
| pH of Solution(at 20°C)  | 7.5 – 8.0                 | 7.56 @ 20.3°C |
| λmax (at pH 7.0)   | 259 ± 1 nm                | 259.5 nm      |
| A250/A260  | 0.78 ± 0.03               | 0.77          |
| A280/A260  | 0.15 ± 0.02               | 0.15          |
| Purity dATP (HPLC Area % at λmax)  | ≥99%                      | >99.9 %       |
| dNDP + dNMP (HPLC Area % at λmax)  | <1%                       | Passed        |
| Appearance   | Clear colourless solution | Passed        |



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| Analysis          | Specification   | Result |
|-------------------|---|--------|
| Functional        | A 3Kb Lambda DNA fragment is amplified with a dilution series of dATP, using standard conditions and 30 cycles. Single distinct bands were observed with agarose gel electrophoresis (ethidium stained).  | Passed |
| DNA contamination | Quantitative PCR analysis with no template. Presence of <i>E. coli</i> and mouse genomic DNA checked. Test sample must amplify in line with a reference sample.   | Passed |
| DNase             | Incubation of a 1Kb double stranded DNA fragment. Incubation for 4 hours at 37°C with dilution series of DNase I. Analysed by agarose gel electrophoresis. Test sample must show less degradation than the limit of detection 2.5 x 10 <sup>-3</sup> U DNase. | Passed |
| RNase             | Quantitative PCR analysis with high and low RNase standards. Test sample must show less RNase activity than the limit of detection 9.7x10 <sup>-3</sup> ng/µL RNase.  | Passed |
| Nicking Activity  | Incubation of dATP with supercoiled control plasmid. Analysed by agarose gel electrophoresis. Test sample does not show an increase of linearized or relaxed plasmid.   | Passed |

QA / QC Representative:



Alberta Newton

Date: 9<sup>th</sup> November 2023