Product Sheet



87723[™]

Description

Clone type: Vector Shipping information: DNA supplied by depositor

Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.

BSL1

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories (BMBL)*, U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies and procedures as well as any other applicable regulations as enforced by your local or national agencies.

Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

Vector Information



pCu413CUP1

87723

Construct size (kb): 5.4520001411438 Intact vector size: 5.452 Vector name: pCu413CUP1 (phagemid) Type of vector: phagemid Construction: pRS413, CUP1 promoter Host range: Saccharomyces cerevisiae; Candida robusta; Escherichia coli Centromere: CEN6 Cloning sites: Spel; BamHI; Smal; EcoRI; EcoRV; ClaI; SalI; XhoI Markers: HIS3; ampR MCS: Spel...XhoI, -> Polylinker sites: Spel; BamHI; SmaI; PstI; EcoRI; EcoRV; HindIII; ClaI; SalI; Xho Promoters: Expression: CUP1 Replicon: f1, ←; pMB1; ARSH4 Terminator: CYC1, <-

Notes

Restriction digests of the clone give the following sizes (kb): SacI--5.4, SacI/SpeI--5.2, 0.26. - ATCC staff

Gene CUP1 is the most potent for copper induction. CUP1 promoter can be used to create a series of versatile vectors to allow its utilization to control the expression of heterologous genes in S.cerevisiae.

- personal communication

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pCu413CUP1 (ATCC 87723)



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References

References and other information relating to this material are available at www.atcc.org.

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Revision

This information on this document was last updated on 2024-10-25

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