Designation: **DAUDI**

CLS order number: Cryovial: 302009
gDNA: 302009GD05

Origin and General Characteristics	
Organism:	*Homo sapiens* (human)
Ethnicity:	African
Age:	16 years of age
Gender:	Male
Tissue:	Hematopoietic system
Morphology:	B-Lymphoblast
Growth Properties:	Suspension
Description:	Established 1967 from a 16-year-old African boy with Burkitt's Lymphoma. Daudi is reported to be EBV-positive. The cell line is deficient of HLA-ABC expression on the cell surface due to the absence of beta-2-microglobulin, responsible for correct intracellular folding and processing of the MHC class I molecule in the endoplasmic reticulum.
References:	Klein E, Klein G, Nadkami JS, Wigzell H, Clifford P. Surface Igm-kappa specificity on a Burkitt lymphoma cell in vivo and derived culture lines. Cancer Res. 28:1300-1310 (1968) Further information for this cell line see. Cellosaurus, Accession CVCL_0008 A list of further relevant literature can be requested at CLS.
Culture Conditions and Handling	
Culture Medium:	RPMI 1640 with 10% FBS (MG-70, CLS order number 820700).
Sub-culturing:	Maintain culture between 3 to 9 x10^5 cells/ml; A maximum density of 2 x10^6 cells/ml is possible. Incubate at 5% CO_2, 37°C.
Seeding density:	3x10^5/ml
Fluid Renewal:	2 times weekly
Freeze Medium:	CM-1 (CLS order number: 800125, 25ml, 800150, 50ml)
Freezing recovery:	Fast (48 hours)
Sterility:	Mycoplasma specific PCR: negative Cell-based Plasmotest: negative
Biosafety Level:	1 according to TRBA 468
Shipping requirements:	No specific safety requirements for shipping
Safety precautions:	If the cryovial is planned to be stored in liquid nitrogen and to be thawed in the future, special safety precautions should be followed: Protective gloves and clothing should be used and a facemask or safety goggles must be worn when transferring frozen samples into or removing from the liquid nitrogen tank. The removal of a cryovial from liquid nitrogen may result in the explosion of the frozen vial creating flying fragments. Caputo, J.L. Biosafety procedures in cell culture. J. Tissue Cult. Methods 11:223-227, 1988. ATCC Quality Control Methods for Cell Lines, 2nd edition, 1992.
Special Features of the Cell Line	
Karyotype:	46, almost diploid

Surface antigens:	CD10+, CD19+, CD20+, CD21+, CD22+, CD23-, CD24-, CD32+, CD37+, CD38+, CD39-,CD40+, CD54+, CD72+, CD73-, CD75+, CD77+, CD81+, CD82+, CD83-, CD84+, CD86+	
HLA-typing (by NS-seq):	Class I HLA-A: A*01:02,*66:01:01 HLA-B: B*58:01:01,*58:02:01 HLA-C: C*03:02:02,*06:02:01 HLA-E: E*01:03:02,*01:03:05	Class II HLA-DR: DRB1*13:01:01,*13:02:01 HLA-DQ: DQA1*01:02:01,*01:03:01 HLA-DQ: DQB1*06:02:01,*06:04:01 HLA-DP: DPB1*02:01:02,*106 :01
DNA Profile (STR):	Amelogenin: X,Y CSF1PO: 12 D13S317: 11,12 D16S539: 10,12 D5S818: 8,13 D7S820: 8,10 THO1: 6,7 TPOX: 8,11 D19S433: 12,14	vWA: 15,17 D3S1358: 16,18 D21S11: 34,35 D18S51: 16,18 Penta E: 7,9 Penta D: 10,12 D8S1179: 14 FGA: 21,26
Possible applications:	Analysis of B cell surface antigens, testing of cytotoxic drugs, mutational analysis, analysis of apoptotic mechanisms, assay development.	

Certificate of Analysis:	The Certificate of Analysis for each batch can be requested by e-mail at service@clsgmbh.de.

Recommendations for handling of cells growing in suspension following delivery	
Cryopreserved cells	The cells come deep-frozen shipped on dry ice. Please make sure that the vial is still frozen. If immediate culturing is not intended, the cryovial(s) must be stored below -150°C after arrival. If immediate culturing is intended, please follow these instructions: Quickly thaw by rapid agitation in a 37°C water bath within 40-60 seconds. The water bath should have clean water containing an antimicrobial agent. As soon as the sample has thawed, remove the cryovial from the water bath. Note: A small ice clump should still remain and the vial should still be cold. From now on, all operations should be carried out under aseptic conditions. Transfer the cryovial to a sterile flow cabinet and wipe with 70% alcohol. Carefully open the vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of culture medium (room temperature). Resuspend the cells carefully. Centrifuge at 300xg for 3 min and discard the supernatant. The centrifugation step may be omitted, but in this case the remains of the freeze medium have to be removed 24 hours later. Resuspend the cells carefully in 10ml fresh cell culture medium and transfer them into one T25 cell culture flask. All further steps are described in the Subculture section.

Warranty:	CLS warrants for a high cell viability and culture performance only if the product(s) is (are) stored and cultured according to the information described above. Using cell culture media and supplements other than the ones recommended in this product information may result in satisfactory proliferation and viabilities. CLS, however, does not warrant for cell recovery, proliferation and function if differing formulations are employed.
Disclaimer:	The customer shall not be entitled to employ this product for purposes other than research. Commercial utilization shall not be permitted; in particular, the cell line, its components or materials made therefrom shall not be sold or transferred to any third party. In addition, the term ‘Commercial use’ shall mean any activity by a party for consideration and may include, but is not limited to, use of the product or its components

	in manufacturing, for providing services, e.g. fee for service testing, in quality control or assurance processes within the manufacturing of products for sale, for therapeutic, diagnostic or prophylactic purposes, or for resale.