iLite® Assay Ready Cells containing cryoprotective medium from Lonza
BM3060, BM3071, BM4012, BM4023, BM4050

SECTION 1: IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND OF THE COMPANY/UNDERTAKING

1.1 Product identifier
Product name: iLite® FGF21 Assay Ready Cells
iLite® GM-CSF Assay Ready Cells
iLite® IL-12 Assay Ready Cells
iLite® IL-23 Assay Ready Cells
iLite® Insulin Assay Ready Cells

Product description: iLite Assay Ready Cells containing cryoprotective medium from Lonza (cat no 12-132A)

Product code: BM3060, BM3071, BM4012, BM4023, BM4050

1.2 Relevant identified uses of the substance or mixture and uses advised against
Use of the product: Laboratory chemicals. For research use only.

1.3 Details of the supplier of the safety data sheet
Company: Svar Life Science AB
Address: Lundvägen 151
Zip code/Place: SE-212 24 Malmö, Sweden
Telephone: +46 40 53 76 00
Website: www.svarlifescience.com
E-mail: info@svarlifescience.com

1.4 Emergency telephone number
Emergency telephone number: (Sweden) Acute: 112 – Ask for “Giftinformation”. If less acute call: +46 010 4566700. Other countries: Please contact local emergency telephone number.

SECTION 2: HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture
Classification according to the Regulation (EC) No. 1272/2008 (CLP):
The mixture is not to be classified according to CLP.
The mixture is covered by Directive 2009/41/EC on the contained use of genetically modified micro-organisms and classified as a Class 1 Genetically Modified Microorganism.
The mixture is covered by Directive 2000/54/EC of 18 September 2000 on the protection of workers from risks related to exposure to biological agents at work.

2.2 Label elements
None

2.3 Other hazards
Other hazards which do not result in classification: Contain Fetal Bovine serum, which is derived from cattle. The Certificate of Analysis for FBS show that the substance has been analyzed for Bluetongue Virus, Bovine Adenovirus, Bovine Parvovirus, Rabies Virus, Reovirus, BRSV Fluorescent Antibody, Cytopathogenic agents and Hemadsorbing agents with a negative result. The products are considered to be biological agents in group 1 (ie. a biological agent that is unlikely to cause human infection). As a precaution, it is recommended that the work is carried out under measures similar to Group 2 in Council Directive 2000/54/EC.

Substance meets the criteria for PBT/ vPvB under Regulation EC No. 1907/2006, appendix XIII: PBT/ vPvB: No
SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

3.2 Mixtures
Assay Ready Cells suspended in RPMI 1640 medium with:

<table>
<thead>
<tr>
<th>No</th>
<th>Product/ingredient name</th>
<th>EC-number</th>
<th>CAS-number</th>
<th>REACH registration number</th>
<th>Conc. (%w/w)</th>
<th>Classification Regulation (EC) No. 1272/2008 (CLP)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fetal Bovine Serum (Heat inactivated FBS)</td>
<td>--</td>
<td>--</td>
<td>--</td>
<td>20</td>
<td>None</td>
<td></td>
</tr>
</tbody>
</table>

Mixed 1:1 with cryoprotective medium from Lonza (cat no 12-132A) containing the following substance:

| Dimethyl Sulfoxid (DMSO) | 200-664-3 | 67-68-5 | -- | 15 | None |

SECTION 4: FIRST-AID MEASURES

4.1 Description of first aid measures
On suspicion of possible infection from biological agents – seek medical advice!

Inhalation: Move to fresh air. Keep at rest and under surveillance. If needed: seek medical advice.
Skin contact: Remove contaminated clothing at once. Flush skin and wash thoroughly with soap and water.
Eye contact: Keep eyelids well apart. Rinse with water for a couple of minutes, remember to remove contact lenses if any. If irritation persists: Seek medical advice.
Ingestion: Rinse mouth and drink plenty of water. If needed or if larger amounts has been swallowed: Seek medical advice.

4.2 Most important symptoms and effects, both acute and delayed
Skin contact: May cause irritation of skin.
Eye contact: May cause irritation of eyes.
Inhalation: Prolonged or frequent exposure to vapours of volatile organic compounds may result in damage on liver, kidneys, blood or central nervous system (including brain damage).

4.3 Indication of any immediate medical attention and special treatment needed
Show this safety data sheet to a physician or emergency ward

SECTION 5: FIREFIGHTING MEASURES

5.1 Extinguishing media
Suitable extinguishing media
Use water spray, carbon dioxide, dry chemical or foam.

Unsuitable extinguishing media
Waterjet

5.2 Special hazards arising from the substance or mixture
Hazardous thermal decomposition products
Decomposition products may include the following materials: oxides of carbon and sulphur.

5.3 Advice for firefighters
Special protective actions for fire-fighters
Promptly isolate the scene by removing all persons from the vicinity of the incident if there is a fire. No action shall be taken involving any personal risk or without suitable training.

Special protective equipment for fire-fighters
Fire-fighters should wear appropriate protective equipment and self-contained breathing apparatus (SCBA) with a full face-piece operated in positive pressure mode. Clothing for fire-fighters (including helmets, protective boots and gloves) conforming to European standard EN 469 will provide a basic level of protection for chemical incidents.

Further information
Not applicable
SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

For non-emergency personnel
Use personal protective equipment – see section 8. No action shall be taken involving any personal risk or without suitable training. Evacuate surrounding areas. Keep unnecessary and unprotected personnel from entering. Do not touch or walk through spilt material. The employees or the company’s occupational health and safety organization must be informed immediately of any accident or incident that may have resulted in the release of biological agents, which may cause disease in humans.

For emergency responders
If specialized clothing is required to deal with the spillage, take note of any information in Section 8 on suitable and unsuitable materials. See also Section 8 for additional information on hygiene measures.

6.2 Environmental precautions
Do not empty into drains – see section 12. Inform appropriate authorities in accordance with local regulations.

6.3 Methods and material for containment and cleaning up

Small spill
Stop leak if without risk. Move containers from spill area. Wipe up spillage etc. with paper towels. Use wet towels to finish cleaning up. Follow the laboratory’s general decontamination procedure for infectious waste. Flush area of decontamination with water. Further handling of spillage – see section 13.

Large spill
Stop leak if without risk. Move containers from spill area. Prevent entry into sewers, water courses, basements or confined areas. Contain and collect spillage with absorbent material as vermiculite. Further handling of spillage – see section 13.

6.4 Reference to other sections
See Section 8 for information on appropriate personal protective equipment. See Section 13 for additional waste treatment information.

SECTION 7: HANDLING AND STORAGE

7.1 Precautions for safe handling

Protective measures
Before use, a workplace assessment of the safety and health conditions at the working place must be carried out according to the general specifications mentioned in the EU Directive 2000/54/EC on biological agents and work. Work must be planned, organized and carried out so that the influence of biological agents is avoided as much as possible. Use laboratory facilities, which generally qualify for handling of biological agents. No tool or used material should after end use be placed on tables or similar but collected immediately in special sealed containers. Recycling of tools should only take place after proper disinfecting and purification. If appliances are contaminated, washing must be made with appropriate disinfectant before further use.

Advice on general occupational hygiene
Eating, drinking and smoking should be prohibited in areas where this material is handled. Avoid contact with skin, eyes and clothing. Always wash hands with soap and water after completing work, and when leaving the laboratory (e.g. before going to the toilet and at the end of the workday). Do not pipette by mouth pipetting. See also Section 8 for additional information on hygiene measures.

7.2 Conditions for safe storage, including any incompatibilities

Storage:
Upon receipt confirm that adequate dry-ice is present and the cells are frozen. Immediately transfer to -80 °C (do not store at any other temperature). Cells should be used within 30 min of thawing and should be diluted immediately after thawing.

Further information:
Not applicable

7.3 Specific end use(s)
Laboratory chemicals for research use only.
SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

Occupational exposure limits

European Union: None

UK: None

Sweden:

<table>
<thead>
<tr>
<th>NGV</th>
<th>KGV</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>50 ppm = 150 g/cm³</td>
<td>150 ppm = 500 g/cm³</td>
<td>H,V</td>
</tr>
</tbody>
</table>

H: Skin permeable
V: Indicative short-term exposure limit

Germany, MAK: 50 ppm = 160 mg/m³

Denmark: 50 ppm = 160 mg/m³

Finland: 50 ppm (8h)

Austria: 50 ppm = 160 mg/m³

Switzerland: 50 ppm = 160 mg/m³

Recommended monitoring procedure

Not relevant

Derived effect levels

Product/ingredient name

<table>
<thead>
<tr>
<th>Type</th>
<th>Exposure Value</th>
<th>Population</th>
<th>Effects</th>
</tr>
</thead>
</table>

Predicted effect concentrations

Not available

PNEC Summary

Not available – No CSR

8.2 Exposure controls

Appropriate engineering controls

Sufficient ventilation.

Hygiene measures

Wash hands thoroughly after handling chemical products, before eating, drinking, smoking and using the lavatory and at the end of the working period. Appropriate techniques should be used to remove potentially contaminated clothing. Wash contaminated clothing before reusing. Ensure that eyewash stations and safety showers are close to the workstation location.

Respiratory protection

Not relevant during normal condition.

Eye/face protection

Use safety glasses (according to EN166) when there is risk of splashes.

Hand protection

Wear protective gloves (according to EN374) of butyl rubber or nitrile rubber.

Body protection

Wear suitable protective clothing.

Environmental exposure controls

Not applicable
SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physical state</td>
<td>Aquaous solution</td>
</tr>
<tr>
<td>Colour</td>
<td>Light red</td>
</tr>
<tr>
<td>Odour threshold</td>
<td>n.d</td>
</tr>
<tr>
<td>Solubility(ies)</td>
<td>Soluble in water</td>
</tr>
<tr>
<td>pH (product)</td>
<td>n.d</td>
</tr>
<tr>
<td>Melting point/freezing point</td>
<td>n.d</td>
</tr>
<tr>
<td>Initial boiling point and boiling range</td>
<td>n.d</td>
</tr>
<tr>
<td>Flash point</td>
<td>n.a</td>
</tr>
<tr>
<td>Evaporation rate (butyl acetate = 1)</td>
<td>n.d</td>
</tr>
<tr>
<td>Flammability (solid, gas)</td>
<td>n.a</td>
</tr>
<tr>
<td>Upper/lower flammability or explosive limits</td>
<td>n.d</td>
</tr>
<tr>
<td>Combustion rate</td>
<td>n.d</td>
</tr>
<tr>
<td>Vapour pressure (at 20°C)</td>
<td>n.d</td>
</tr>
<tr>
<td>Vapour density</td>
<td>n.d</td>
</tr>
<tr>
<td>Relative density (Water = 1)</td>
<td>n.d</td>
</tr>
<tr>
<td>Partition coefficient:</td>
<td>n.d</td>
</tr>
<tr>
<td>n-octanol/water</td>
<td>n.d</td>
</tr>
<tr>
<td>Autoignition temperature</td>
<td>n.d</td>
</tr>
<tr>
<td>Decomposition temperature</td>
<td>n.d</td>
</tr>
<tr>
<td>Viscosity</td>
<td>n.d</td>
</tr>
<tr>
<td>Explosive properties</td>
<td>n.d</td>
</tr>
<tr>
<td>Oxidising properties</td>
<td>n.d</td>
</tr>
</tbody>
</table>

n.d = not determined
n.a = not applicable

9.2 Other information
Not applicable

SECTION 10: STABILITY AND REACTIVITY

10.1 Reactivity
No available information

10.2 Chemical stability
Stable at recommended storage conditions – see section 7.

10.3 Possibility of hazardous reactions
No available information.

10.4 Condition to avoid
No available information.

10.5 Incompatible materials
Strong oxidizing agents

10.6 Hazardous decomposition products
When heated to high temperatures (decomposition) toxic fumes are emitted: Oxides of carbon and sulphur.
11.1 Information on toxicological effects

<table>
<thead>
<tr>
<th>Hazard class</th>
<th>Data (DMSO)</th>
<th>Test</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute toxicity: Inhalation</td>
<td>LD₅₀ (rat) &gt; 2 mg/l/4h</td>
<td>No information</td>
<td>IUCLID</td>
</tr>
<tr>
<td>Dermal</td>
<td>LD₅₀ (rat) &gt; 40000 mg/kg</td>
<td>No information</td>
<td>IUCLID</td>
</tr>
<tr>
<td>Oral</td>
<td>LD₅₀ (rat) = 14500 mg/kg</td>
<td>No information</td>
<td>IUCLID</td>
</tr>
<tr>
<td>Corrosion/irritation</td>
<td>Mild eye and skin irritation, rabbit</td>
<td>OECD 404, EU Method B.5</td>
<td>ECHA</td>
</tr>
<tr>
<td>Sensitization</td>
<td>No skin sensitization, guinea pig</td>
<td>Buehler</td>
<td>IUCLID</td>
</tr>
<tr>
<td>CMR</td>
<td>No mutagenicity, carcinogenicity, genotoxicity</td>
<td>Several</td>
<td>Merck/IUCLID</td>
</tr>
</tbody>
</table>

**Acute toxicity**
Assessment for other reagents than DMSO: No data available.

**Irritation/Corrosion**
Assessment for other reagents than DMSO: No data available.

**Sensitization by inhalation/skin contact**
Assessment for other reagents than DMSO: No data available.

**Germ cell mutagenicity**
Assessment for other reagents than DMSO: The chemical structure of the different reagents doesn’t indicate any mutagenic effects.

**Carcinogenicity**
Assessment for other reagents than DMSO: The chemical structure of the different reagents doesn’t indicate any carcinogenic effects.

**Reproduction toxicity**
Assessment for other reagents than DMSO: The chemical structure of the different reagents doesn’t indicate any reproduction toxic effects.

**Developmental toxicity**
Assessment for other reagents than DMSO: The chemical structure of the different reagents doesn’t indicate any teratogenic effects.

**Specific target organ toxicity (single exposure)**
STOT assessment single dose toxicity: No data available.

**Repeated dose toxicity and specific organ toxicity (repeated exposure)**
Prolonged or frequent exposure to vapours of volatile organic compounds may result in damage on liver, kidneys, blood or central nervous system (including brain damage).
SECTION 12: ECOLOGICAL INFORMATION

12.1 Toxicity

12.1.1 Acute toxicity in the aquatic environment for DMSO

<table>
<thead>
<tr>
<th>Test</th>
<th>Value/unit (mg/l)</th>
<th>Test method</th>
<th>Exp. time (h)</th>
<th>Species</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fish LC₅₀</td>
<td>32000</td>
<td>Static (FW)</td>
<td>96</td>
<td>Oncorhynchus mykiss</td>
</tr>
<tr>
<td>Daphnia EC₅₀</td>
<td>7000</td>
<td>No info. (FW)</td>
<td>24</td>
<td>Daphnia sp.</td>
</tr>
<tr>
<td>Algae EC₅₀</td>
<td>12350-25500</td>
<td>No info. (SW)</td>
<td>96</td>
<td>Skeletonema costatum</td>
</tr>
</tbody>
</table>

12.1.2 Acute toxicity in the aquatic environment other reagents than DMSO

No data available.

12.1.3 Ecotoxicity

No data available.

12.2 Persistence and degradability

Conclusion/Summary DMSO is not readily degradable (3.1% after 14 days in OECD 301C test).

12.3 Bioaccumulative potential

Conclusion/Summary DMSO: Log Kₐw -1.35 – No significant bioaccumulation.

12.4 Mobility in soil

Soil/water partition coefficient (KOC) DMSO: KOC (calculated) < 10 – Very high mobility expected in soil environments.

Mobility No available data

12.5 Results of PBT and vPvB assessment

PBT The substance is not considered PBT according to criteria in Annex XIII.

vPvB The substance is not considered vPvB according to criteria in Annex XIII.

12.6 Other adverse effects

None known

SECTION 13: DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods

Method of disposal Biological agents are considered hazardous waste. Disposal should be according to local, state or national legislation.

Note! Waste containers containing biological material must be labeled with: (black symbol on yellow background). The generation of waste should be avoided or minimized wherever possible. This material and its container must be disposed of in a safe way by incineration.

Hazardous waste Within the present knowledge of the supplier, this product is regarded as hazardous waste, as defined by EU Directive 2008/98/EC.

European Waste Catalogue (EWC)

<table>
<thead>
<tr>
<th>EWC Waste Code</th>
<th>Type of waste</th>
</tr>
</thead>
<tbody>
<tr>
<td>18 01 03</td>
<td>Wastes whose collection and disposal is subject to special requirements in order to prevent infection</td>
</tr>
<tr>
<td>15 02 02</td>
<td>Absorbent material containing residues of or contaminated by dangerous substances</td>
</tr>
</tbody>
</table>

Packaging

Method of disposal Incineration.

Special precautions None.

SECTION 14: TRANSPORT INFORMATION
Product is not classified as dangerous goods.

### 14.1 UN number

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>--</td>
<td>--</td>
<td>--</td>
<td>--</td>
</tr>
</tbody>
</table>

### 14.2 UN proper shipping name

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>--</td>
<td>--</td>
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<td>--</td>
</tr>
</tbody>
</table>

### 14.3 Transport hazard class(es)

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>--</td>
<td>--</td>
<td>--</td>
<td>--</td>
</tr>
</tbody>
</table>

### 14.4 Packing Group

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>--</td>
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<td>--</td>
</tr>
</tbody>
</table>

### 14.5 Environmental hazards

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>--</td>
<td>--</td>
<td>--</td>
<td>--</td>
</tr>
</tbody>
</table>

### 14.6 Special precautions for user

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
</tr>
</tbody>
</table>

### 14.7 Transport in bulk according to Annex II of MARPOL and the IBC Code

<table>
<thead>
<tr>
<th>ADR/RID</th>
<th>ADN/ADNR</th>
<th>IMDG</th>
<th>IATA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Not applicable</td>
<td>Not applicable</td>
<td>Not applicable</td>
<td>Not applicable</td>
</tr>
</tbody>
</table>

### Additional information

Waste containing used biological agents may be considered as dangerous goods;
UN 3291, CLINICAL WASTE, UNSPECIFIED: N.O.S., or (BIO) MEDICAL WASTE N.O.S. or REGULATED MEDICAL WASTE, N.O.S. Class 6.2 Packing Group II

### SECTION 15: REGULATORY INFORMATION

#### 15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture

Must not be used by persons under 18 years of age (Directive 94/33/EC). The employer shall assess the working conditions and, if there is any risk to the safety or health and any effects on the pregnancy or breastfeeding of workers, take the necessary measures to adjust the working conditions (Directive 92/85/EEC).

The mixture is covered by:
- Directive 2009/41/EC on the contained use of genetically modified micro-organisms
- Directive 2000/54/EC – biological agents at work

**EU Regulation (EC) No. 1907/2006 (REACH)**
- Annex XIV – List of substances subject to authorization
- Substances of very high concern
  None of the components are listed.

- Annex XVII – Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles
  Not applicable

#### 15.2 Chemical Safety Assessment

No CSR.

### Other information

<table>
<thead>
<tr>
<th>Tariff Code harmonized system</th>
<th>Not applicable</th>
</tr>
</thead>
<tbody>
<tr>
<td>The EU Seveso Directive</td>
<td>Not applicable</td>
</tr>
</tbody>
</table>
**International regulations**

<table>
<thead>
<tr>
<th>Chemical Weapons Convention List</th>
<th>Chemical Weapons Convention List</th>
<th>Chemical Weapons Convention List</th>
</tr>
</thead>
<tbody>
<tr>
<td>Schedule I Chemicals</td>
<td>Schedule II Chemicals</td>
<td>Schedule III Chemicals</td>
</tr>
<tr>
<td>Not regulated</td>
<td>Not regulated</td>
<td>Not regulated</td>
</tr>
</tbody>
</table>

**SECTION 16: OTHER INFORMATION**

Conforms to Regulation (EC) No. 1907/2006 (REACH), Annex II.

**LIST OF HAZARD STATEMENTS MENTIONED UNDER SECTION 3:** None

**Abbreviations:**
- BRSV = Bovine Respiratory Syncytial Virus
- CMR = Carcinogenicity, Mutagenicity, and Reproduction toxicity
- CSR = Chemical Safety Report
- DNEL = Derived No-Effect Level
- EC50 = Half maximal effective concentration
- FW = Fresh Water (Färskvatten)
- KGV = Korttidsvärde (Swedish for short term exposure limit)
- LC50 = Lethal Concentration 50 %
- LD50 = Lethal Dose 50 %
- MAK = Maximale Arbeitsplatzkonzentrationen (German for maximum workplace concentration)
- NGV = Nivågränsvärde (Swedish for exposure limit)
- PBT = Persistent, Bioaccumulative, Toxic
- PNEC = Predicted No-Effect Concentration
- vPvB = very Persistent, very Bioaccumulative

**Literature:**
- Merck (Safety Data Sheet)
- IUCLID = International Uniform Chemical Information Database
- ECHA = European Chemicals Agency

**Other information**

No special training is required. However, the user should be well instructed in the execution of his/her task, be familiar with this Safety Data Sheet and have normal training in the use of personal protective equipment.

**Revisions**

<table>
<thead>
<tr>
<th>Version</th>
<th>Valid from (date)</th>
<th>Changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.0</td>
<td>14-Mar-2019</td>
<td>New document</td>
</tr>
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</table>