

Safety Data Sheet

SECTION 1: Identification

Contact information

General



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Chemtrec: 1-800-424-9300 (Availability: 24 hours/day; 7 days/week; 365 days/year)

Product identifier	mRNA-1283
Synonyms	SARS-CoV2 Vaccine, SARS-CoV2 Vaccine Drug Product, SARS-CoV2 Vaccine Drug Substance
Trade name	mNEXSPIKE®
Chemical family	Mixture - contains ribonucleotides
Recommended uses and restrictions	Formulated pharmaceutical product.
Note	This SDS is written to address potential health and safety issues associated with the handling of the drug substance and/or drug product. Personnel manufacturing this product should consult the SDS of each hazardous ingredient for hazard information and handling recommendations. This SDS will be revisited if more data become available.

SECTION 2: Hazard(s) identification

Classification of the substance or mixture	The classification and labeling listed below is for bulk drug substance and drug product.
Label elements	Not classified
GHS Hazard pictograms	Not applicable
GHS Signal word	Not applicable
GHS Hazard statements	Not applicable
GHS Precautionary statements	Not applicable
Other hazards	mRNA-1283 is a lipid nanoparticle (LNP) based delivery system that is non-viral and non-infectious and does not include the possibility of DNA integration. The LNP is comprised of several lipids, including a novel lipid. Commonly observed adverse effects with parenteral administration of similar mRNA vaccines include reversible injection site inflammation, fatigue, feverishness, myalgia and nausea. Little to no systemic absorption is expected to occur during routine handling of intact containers or through incidental contact with intact skin. The mRNA component is expected to rapidly degrade in the digestive tract following accidental ingestion. Aerosol-generating operations, contact with compromised skin, or accidental inject may present different exposure potential and should be addressed through task-specific controls.
Note	This mixture does not meet criteria for classification under GHS as implemented by Regulation EC No 1272/2008 (EU CLP), WHMIS 2015 (Health Canada), EC No 1272/2008 (UK CLP), Australia WHS Regulations and Hazard Communication Standard No. 1910.1200 (US OSHA). Nevertheless, it should be handled with caution as it has not yet been fully tested and is pharmacologically active.

SECTION 3: Composition/Information on ingredients

Ingredient	CAS number	EINECS/ELINCS#	Amount	GHS classification
Water	7732-18-5	231-791-2	85 – 95 %	Not classified
Sucrose	57-50-1	200-334-9	5 – 10 %	Not classified
Lipids	N/A	N/A	0.5 – 5 %	Not classified
mRNA-1283	N/A	N/A	0.01 – 1 %	Not classified

Note	The components of this product are non-hazardous and/or present at amounts below reportable limits. Amounts are listed as ranges; the exact percentage of composition is withheld as a trade secret. Sucrose is listed because it has an OEL. See Section 16 for full text of GHS classifications.
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SECTION 4: First-aid measures

Description of first aid measures

Immediate medical attention and special treatment, if necessary	No.
Inhalation	Immediately move exposed subject to fresh air. If not breathing, give artificial respiration. If breathing is labored, administer oxygen. Immediately notify medical personnel and supervisor.
Skin contact	Wash exposed area with soap and water and remove contaminated clothing/shoes. If irritation occurs or persists, notify medical personnel and supervisor.
Eye contact	If easy to do, remove contact lenses, if worn. Immediately flush eyes with copious quantities of water for at least 15 minutes. If irritation occurs or persists, notify medical personnel and supervisor.
Ingestion	Do not induce vomiting unless directed by medical personnel. Do not give anything to drink unless directed by medical personnel. Never give anything by mouth to an unconscious person. Notify medical personnel and supervisor. If swallowed, call a physician immediately.
Most Important Symptoms/Effects	Medical conditions aggravated by exposure: None known or reported. Treat symptomatically and supportively.
Expected Symptoms/Effects, Acute and Delayed	See Sections 2 and 11

SECTION 5: Fire-fighting measures

Suitable (and unsuitable) extinguishing media

Suitable extinguishing media	Use water spray (fog), foam, dry powder, or carbon dioxide, as appropriate for surrounding fire and materials.
Specific hazards arising from the chemical	No information identified. May emit carbon monoxide, carbon dioxide, oxides of nitrogen and other nitrogen-containing compounds.
Fire hazard	As product is an aqueous solution, it is not expected to be flammable.
Explosion hazard	As product is an aqueous solution, it is not expected to be explosive.
Special protective equipment and precautions for fire-fighters	
Firefighting instructions	In case of fire in the surroundings: use the appropriate extinguishing agent. Wear full protective clothing and an approved, positive pressure, self-contained breathing apparatus. Decontaminate all equipment after use.

SECTION 6: Accidental release measures

Personal precautions, protective equipment and emergency procedures

Protective equipment	If product is released or spilled, take proper precautions to minimize exposure by using appropriate personal protective equipment (see Section 8). Area should be adequately ventilated.
Emergency procedures	Do not breathe vapors/mist/spray.
Environmental precautions	Do not empty into drains. Avoid release to the environment.
Methods and material for containment and cleaning up	
Methods for cleaning up	DO NOT CAUSE MATERIAL TO BECOME AIRBORNE. For small spills, soak up material with absorbent, e.g. paper towels. For large spills, cordon off spill area and minimize the spreading of spilled material. Soak up material with absorbent. Collect spilled material, absorbent, and rinse water into suitable containers for proper disposal in accordance with applicable waste disposal regulations (see Section 13). Clean the affected area in accordance with an approved site cleaning or decontamination procedure
Other information	Dispose of materials or solid residues at an authorized site.
Reference to other sections	See Sections 8 and 13 for more information.

SECTION 7: Handling and storage

Precautions for safe handling	Follow recommendations for handling pharmaceutical agents (i.e. use of engineering controls and/or other personal protective equipment if needed). Avoid contact with eyes, skin, and other mucous membranes. Wash thoroughly after handling. Do not breathe vapor/mist/spray.
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Conditions for safe storage, including any incompatibilities

Storage conditions	Protect from light. Store in a dry, cool and well-ventilated place. Keep/Store away from incompatible materials.
Storage temperature	Refer to Summary of Analysis Information
Specific end use(s)	Pharmaceuticals.

SECTION 8: Exposure controls/personal protection

Note	Dispose of broken vials in a sharps container.
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Control parameters/Occupational Exposure Limits

Name	Issuer	Value
Water	No data available	No data available
mRNA	No data available	No data available
Sucrose	BE - Limit value (mg/m ³)	10 mg/m ³
	ES - VLA-ED (mg/m ³)	10 mg/m ³
	FR - VME (mg/m ³)	10 mg/m ³
	IE - OEL (8 hours ref) (mg/m ³)	10 mg/m ³
	IE - OEL (15 min ref) (mg/m ³)	20 mg/m ³
	LT - IPRV (mg/m ³)	10 mg/m ³
	PT - OEL TWA (mg/m ³)	10 mg/m ³
	GB - WEL TWA (mg/m ³)	10 mg/m ³
	GB - WEL STEL (mg/m ³)	20 mg/m ³
	ACGIH TWA (mg/m ³)	10 mg/m ³
	NIOSH REL (TWA) (mg/m ³)	10 mg/m ³ (total dust)
	OSHA PEL (TWA) (mg/m ³)	15 mg/m ³ (total dust)
Lipids	No data available	No data available

Appropriate engineering controls	None required for normal handling of packaged product. If handling bulk product and/or vials or syringes are open/crushed/broken: selection and use of containment devices and personal protective equipment should be based on a risk assessment of exposure potential. Use local exhaust and/or enclosure at aerosol/mist-generating points. Use engineered local exhaust ventilation (LEV) and/or enclosure for procedures where aerosolization may occur such as opened transfers, pumping, and spraying. Solutions can be handled outside a containment system or without LEV during procedures with no potential for aerosolization. All containers for solutions and slurries must be covered while being transferred.
Respiratory protection	None required for normal handling of packaged product. If handling bulk product and/or vials or syringes are open/crushed/broken: choice of respiratory protection should be appropriate to the task and the level of existing engineering controls. For bulk manufacturing operations, a tight-fitting full-face respirator with HEPA filters may be required if performing aerosol generating operations. For liquid spill clean-up using gentle application of sorbent materials, standard PPE should be used.
Hand protection	None required for the normal handling of packaged product. If handling bulk product and/or vials or syringes are open/crushed/broken: wear nitrile or other impervious gloves if skin contact is possible. When the material is diluted in an organic solvent, wear gloves that provide protection against the solvent.
Eye protection	None required for normal handling of packaged product. If handling bulk product and/or vials or syringes are open/crushed/broken: wear safety glasses with side shields, chemical splash goggles, or full face shield, if necessary. Base the choice of protection on the job activity and potential for contact with eyes or face. An emergency eye wash station should be available.
Skin and body protection	None required for normal handling of packaged product. If handling bulk product and/or vials or syringes are open/crushed/broken: wear disposable coveralls appropriate to the task, booties, and safety glasses with side shields. Ensure gloves are protective against solvents in use. Protective garments (coveralls, disposable coveralls, lab coats) are not to be worn in common areas (e.g., cafeterias) or out-of-doors. Employees must be trained in proper gowning and degowning practices
Other protective measures	Wash hands in the event of contact with this substance, especially before eating, drinking or smoking. Protective equipment is not to be worn outside the work area (e.g., in common areas or out-of-doors).
Environmental exposure controls	Avoid release to the environment and operate within closed systems wherever practicable. Air and liquid emissions should be directed to appropriate pollution control devices. In case of spill, do not release to drains. Implement appropriate and effective emergency response procedures to prevent release or spread of contamination and to prevent inadvertent contact by personnel.

SECTION 9: Physical and chemical properties

Physical state	Solid (frozen in shipment); Liquid (in use)
Appearance	Frozen or liquid, packaged in glass vial or pre-filled syringe
Formula	N/A (codon-optimized mRNA)
Molecular mass	Not applicable (Mixture)
Color	Frozen - White to off-white; Thawed - Colorless
Odor	No data available

pH	7.0 - 8.0
Melting point	Not applicable
Freezing point	No data available
Boiling point	No data available
Flash point	No data available
Relative evaporation rate (butyl acetate=1)	No data available
Flammability (solid, gas)	Not applicable
Vapor pressure	No data available
Relative vapor density at 20 °C	No data available
Relative density	No data available
Solubility	Soluble in water
Log Kow	No data available
Auto-ignition temperature	No data available
Decomposition temperature	No data available
Viscosity, kinematic	No data available
Viscosity, dynamic	Not applicable
Explosion limits	Not applicable
Explosive properties	As product is an aqueous solution, it is not expected to be explosive
Oxidizing properties	No oxidizing properties

SECTION 10: Stability and reactivity

Reactivity	The product is non-reactive under normal conditions of use, storage and transport.
Chemical stability	Stable under normal conditions.
Possibility of hazardous reactions	No dangerous reactions known under normal conditions of use.
Conditions to avoid	(See section 7: Handling and Storage).
Incompatible materials	No data available
Hazardous decomposition products	Under normal conditions of storage and use, hazardous decomposition products should not be produced.

SECTION 11: Toxicological information

Likely routes of exposure	mRNA-1283 is expected to degrade in gastric fluids, and consequently the drug is not expected to be absorbed by ingestion. As a large molecular weight compound, mRNA-1283 is not likely to be systemically absorbed through inhalation and skin contact.
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Toxicological information

Acute toxicity

Component	Type	Dose
Water	LD50 oral rat	> 90 ml/kg
	LC50 inhalation rat (Dust/Mist - mg/l/4h)	> 6 mg/l/4h
mRNA	No data available	No data available
Sucrose	LD50 oral rat	29700 mg/kg
Lipids	No data available	No data available

Serious eye damage/irritation	No data available
Skin corrosion/irritation	No data available
Sensitization	No data available
STOT-single exposure	No data available
STOT-repeated exposure	No data available for mRNA-1283 For another chemically similar, same platform mRNA vaccine: Rat (4 doses), IM LOAEL: $\geq 9 \mu\text{g/dose}$ Effects: Hematology changes, reversible effects on lymph node, spleen, liver, and bone marrow, and inflammation at the site of injection. No data available for mRNA-1283 For another, chemically similar, same platform mRNA vaccine: Rat (4 doses) IM NOAEL: $150 \mu\text{g/dose}$ (highest dose tested)
Reproductive toxicity	No data available for mRNA-1283 For another, chemically similar, same platform mRNA vaccine: Rat (4 doses) IM Fetal NOAEL: $150 \mu\text{g/dose}$ (highest dose tested)
Developmental toxicity	Maternal IM LOAEL: $15 \mu\text{g/dose}$ Effects: Injection site reactions, abnormal gait, piloerection, transient effects on body weight
Genotoxicity	No data identified for mRNA-1283. As a ribonucleotide it is not expected to be genotoxic. For another chemically similar, same platform mRNA vaccine: <i>In vitro</i>: Bacterial reverse mutation assay (e.g. Ames test): negative Micronucleus test (peripheral human lymphocytes): negative For another chemically similar, same platform mRNA vaccine: <i>In vivo</i>: IV dosed up to 2.6 mg/kg (female) and 5.2 mg/kg (male) rat micronucleus test: positive Effects: Results were not dose dependent and were associated with minimal bone marrow toxicity in the animals
Carcinogenicity	No data available for mRNA-1283. None of the components of this mixture present at levels greater than or equal to 0.1% are listed by NTP, IARC, ACGIH or OSHA as a carcinogen.
Aspiration hazard	No data available
Experience with humans	See "Section 2 - Other Hazards".

SECTION 12: Ecological information

Component	Type	Concentration
Water	No data available	No data available
mRNA	No data available	No data available
Sucrose	No data available	No data available
Lipids	No data available	No data available
Persistence and degradability	Ribonucleotides in water is expected to degrade rapidly in environment.	
Bioaccumulative potential	Not expected to bioaccumulate.	
Mobility in soil	No data available	
Results of PBT assessment	No data available	
Other adverse effects	No data available	
Note	No special precautions are necessary.	

SECTION 13: Disposal considerations

Waste treatment methods	Used product should be disposed of according to local, state, and federal regulations. All wastes containing the material should be properly labeled. Dispose of wastes in accordance to prescribed federal, state, and local guidelines, e.g, appropriately permitted chemical waste incinerator. Rinse waters resulting from spill cleanups should be discharged in an environmentally safe manner, e.g, appropriately permitted municipal or on-site wastewater treatment facility.
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SECTION 14: Transport information

Transport	Based on the available data, this mixture is not regulated as a hazardous material/dangerous good under EU ADR/RID, US DOT, Canada TDG, IATA, or IMDG.
UN number	None assigned.
UN proper shipping name	None assigned.
Transport hazard class(es) (DOT)	None assigned.
Packing group	None assigned.
Marine pollutant	Based on the available data, this mixture is not regulated as an environmental hazard or a marine pollutant.
Special transport precautions	Avoid release to the environment.
Transport in bulk according to Annex II of Marpol and the IBC Code	Not applicable

SECTION 15: Regulatory information

Safety, health and environmental regulations/legislation specific for the substance or mixture	This SDS generally complies with the requirements listed under current guidelines in the US, EU, UK, Australia and Canada. Consult your local or regional authorities for more information.
Chemical safety assessment	No chemical safety assessment has been carried out
TSCA	Drugs are exempt from TSCA.
SARA Section 313 - Emission Reporting	This substance or mixture is not known to contain a toxic chemical or chemicals in excess of the applicable de minimis concentration as specified in 40 CFR §372.38(a) subject to the reporting requirements of section 313 of Title III of the Superfund Amendments and Reauthorization Act of 1986 and 40 CFR Part 372.
California Proposition 65	California Proposition 65 - This product does not contain any substances known to the state of California to cause cancer, developmental and/or reproductive harm
Additional information	No additional information available

SECTION 16: Other information

Data sources	Information from published literature and internal company data.
Abbreviations and acronyms	ACGIH - American Conference of Governmental Industrial Hygienists; ADR/RID - European Agreement Concerning the International Carriage of Dangerous Goods by Road/Rail; AIHA - American Industrial Hygiene Association; CAS# - Chemical Abstract Services Number; CLP - Classification, Labeling, and Packaging of Substances and Mixtures; DNEL - Derived No Effect Level; DOT - Department of Transportation; EINECS - European Inventory of New and Existing Chemical Substances; ELINCS - European List of Notified Chemical Substances; EU - European Union; GHS - Globally Harmonized System of Classification and Labeling of Chemicals; IARC - International Agency for Research on Cancer; IDLH - Immediately Dangerous to Life or Health; IATA - International Air Transport Association; IMDG - International Maritime Dangerous Goods; LOEL - Lowest Observed Effect Level; LOAEL - Lowest Observed Adverse Effect Level; NIOSH - The National Institute for Occupational Safety and Health; NOEL - No Observed Effect Level; NOAEL - No Observed Adverse Effect Level; NTP - National Toxicology Program; OEL - Occupational Exposure Limit; OSHA - Occupational Safety and Health Administration; PBT - Persistent, Bio-accumulative, and Toxic; PNEC - Predicted No Effect Concentration; SARA - Superfund Amendments and Reauthorization Act; STOT - Specific Target Organ Toxicity; STEL - Short Term Exposure Limit; TDG - Transportation of Dangerous Goods; TSCA - Toxic Substances Control Act; TWA - Time Weighted Average; vPvB - Very Persistent and Very Bio-accumulative; WHMIS - Workplace Hazardous Materials Information System
Issue date	5 August 2026
Current revision	4.0
Indication of Changes	Updated scope of SDS to reflect both drug substance and drug product.
Disclaimer	The above information is based on data available to us and is believed to be correct. Since the information may be applied under conditions beyond our control and with which we may be unfamiliar, we do not assume any responsibility for the results of its use and all persons receiving it must make their own determination of the effects, properties and protections which pertain to their particular conditions. No representation, warranty, or guarantee, express or implied (including a warranty of fitness or merchantability for a particular purpose), is made with respect to the materials, the accuracy of this information, the results to be obtained from the use thereof, or the hazards connected with the use of the material. Caution should be used in the handling and use of the material because it is a pharmaceutical product. The above information is offered in good faith and with the belief that it is accurate. As of the date of issuance, we are providing all information relevant to the foreseeable handling of the material. However, in the event of an adverse incident associated with this product, this Safety Data Sheet is not, and is not intended to be, a substitute for consultation with appropriately trained personnel.