



CONFIRMATION OF FDA REGISTRATION

Registration No.:3012677127

Dear Official Correspondent:

This document provides notification of the registration number assigned to your establishment:

Establishment:

Address:

Listing Number

See Appendix

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Validity: Dec.31,2020

Conclusion:

This certificate makes no other representations or warranties, nor does it make any representations and warranties to any person or entity other than the named certificate holder. Cytelch assumes no liability to any person or entity in connection with the foregoing. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Cytelch is not affiliated with the U.S. Food and Drug Administration.

Please noted that: FDA registration means the manufacturer registered the factory and certain products with FDA, which does not mean that their products meet certain quality specifications unless the manufacturer shows you the quality certificates.



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Appendix:

Listing Number	Premarket Submission Number/Type	Product Code(s)	Device Name(s)	Activities
D276022	Exempt	HQG	Lens, spectacle, non-custom (prescription)	Manufacturer
D276021	Exempt	HOI	Spectacle, magnifying	Manufacturer
D276020	Exempt	HQY	Sunglasses (non-prescription including photosensitive)	Manufacturer
D276023	Exempt	HQZ	Frame, spectacle	Manufacturer
D391697	Exempt	HOY	Shield, eye, ophthalmic (including sunlamp protective eyewear and post-mydratic eyewear)	Manufacturer
D391699	Enforcement Discretion	QKR	Face mask (except N95 respirator) for general public/healthcare personnel per IIE guidance	Manufacturer

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