## UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

## Form 6-K

Report of Foreign Private Issuer Pursuant to Rule 13a-16 or 15d-16 under the Securities Exchange Act of 1934

For the month of October 2020

Commission File Number 001-37626

# **Mesoblast Limited**

(Exact name of Registrant as specified in its charter)

Not Applicable

(Translation of Registrant's name into English)

Australia

(Jurisdiction of incorporation or organization)

Silviu Itescu

Chief Executive Officer and Executive Director Level 38 55 Collins Street Melbourne 3000

Australia

(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F:

Form 20-F  $\square$  Form 40-F  $\square$ 

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Yes 🗆 No 🗹

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): Yes  $\Box$  No  $\Box$ 

### INFORMATION CONTAINED ON THIS REPORT ON FORM 6-K

On October 22, 2020, Mesoblast Limited filed with the Australian Securities Exchange a new release announcement, which is attached hereto as <u>Exhibit 99.1</u>, and is incorporated herein by reference.

### SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly organized.

Mesoblast Limited

/s/ Charlie Harrison

Charlie Harrison Company Secretary

Dated: October 23, 2020

99.1 Press release of Mesoblast Ltd, dated October 22, 2020.

## asx announcement



## RANDOMIZED CONTROLLED STUDY USING DIRECT INJECTION OF REMESTEMCEL-L INTO INFLAMED GUT OF PATIENTS WITH CROHN'S DISEASE AND ULCERATIVE COLITIS

Melbourne, Australia; October 22, 2020 and New York, USA; October 21, 2020: Mesoblast Limited (ASX:MSB; Nasdaq:MESO), global leader in allogeneic cellular medicines for inflammatory diseases, today announced that a randomized, controlled study of remestemcel-L delivered by an endoscope directly to the areas of inflammation and tissue injury in up to 48 patients with medically refractory Crohn's disease and ulcerative colitis has commenced at Cleveland Clinic.

Mesoblast Chief Medical Officer Dr Fred Grossman said: "Inflammation of the gut in Crohn's disease and ulcerative colitis closely resembles the most severe manifestation of advanced-stage, lifethreatening acute graft versus host disease (aGVHD). Mesoblast's objective is to confirm the potential for remestencel-L to induce luminal healing and early remission in a wider spectrum of diseases with severe inflammation of the gut, in addition to steroid-refractory aGVHD."

Mesenchymal stem cells (MSCs) promote healing of inflamed gut tissue by downregulating gut mucosal effector T-cell activity and promoting regulatory T-cell formation.<sup>1</sup> MSCs have been tested in clinical trials of Crohn's disease using two different modalities: intravenous infusions of MSCs to treat the primary inflammation of Crohn's disease and local injections of MSCs to treat fistulae complicating Crohn's disease.

A third modality, endoscopic delivery of MSCs, has been successful in preclinical experimental models of colitis, reducing the excessive cytokine storm in the inflamed gut and resulting in tissue healing.<sup>2-3</sup> The study at Cleveland Clinic will be the first in humans using local delivery of MSCs in the gut, and will enable Mesoblast to compare clinical outcomes using this delivery method with results from an ongoing randomized, placebo-controlled trial in patients with biologic-refractory Crohn's disease where remestemcel-L was administered intravenously.

The study's lead investigator Dr Amy L. Lightner, Associate Professor of Surgery in the Department of Colon and Rectal Surgery at Cleveland Clinic, stated: "We are aiming to establish a new treatment paradigm by administering remestemcel-L at one of two escalating doses, or placebo, directly to inflamed gut tissue in patients with medically refractory Crohn's disease and ulcerative colitis, both highly debilitating conditions with significant, unmet medical needs."

According to recent estimates, more than three million people (1.3%) in the US alone have inflammatory bowel disease, with more than 33,000 new cases of Crohn's disease and 38,000 new cases of ulcerative colitis diagnosed every year.<sup>4-6</sup> Despite recent advances, approximately 30% of patients are primarily unresponsive to anti-TNFα agents and even among responders, up to 10% will lose their response to the drug every year. Up to 80% of patients with medically-refractory Crohn's disease eventually require surgical treatment of their disease,<sup>7</sup> which can have a devastating impact on quality of life.

### References

1.Mayne C and Williams C. Induced and natural regulatory T cells in the development of inflammatory bowel disease. *Inflamm Bowel Dis* 2013; 19: 1772–1788. 2.Molendijk I et al. Intraluminal Injection of Mesenchymal Stromal Cells in Spheroids Attenuates Experimental Colitis. Journal of Crohn's and Colitis, 2016, 953–964

2. Robins and Endoscopic Transplantation of Mesenchwal Stem Cell Sheets in Experimental Colitis in Rats. Scientific Reports (2018) 811314 | DOI:10.1038/s41598-018-29617

4.CDC Facts and Figures 2015

5.Globaldata Pharmapoint 2018

6.Dahlhamer JM, MMWR Morb Mortal Wkly Rep. 2016;65(42):1166-1169.

7. Crohn's and Colitis Foundation

### About Remestemcel-L

Mesoblast's lead product candidate, remestemcel-L, is an investigational therapy comprising culture-expanded mesenchymal stem cells derived from the bone marrow of an unrelated donor. It is administered to patients in a series of intravenous infusions. Remestemcel-L is thought to have immunomodulatory properties to counteract severe inflammatory processes by down-regulating the production of pro-inflammatory cytokines, increasing production of anti-inflammatory cytokines, and enabling recruitment of naturally occurring anti-inflammatory cells to involved tissues.

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#### About Mesoblast

Mesoblast Limited (ASX:MSB; Nasdaq:MESO) is a world leader in developing allogeneic (off-the-shelf) cellular medicines. The Company has leveraged its proprietary mesenchymal lineage cell therapy technology platform to establish a broad portfolio of commercial products and late-stage product candidates. Mesoblast has a strong and extensive global intellectual property (IP) portfolio with protection extending through to at least 2040 in all major markets. The Company's proprietary manufacturing processes yield industrial-scale, cryopreserved, off-the-shelf, cellular medicines. These cell therapies, with defined pharmaceutical release criteria, are planned to be readily available to patients worldwide.

Remestemcel-L is being developed for inflammatory diseases in children and adults including steroid-refractory acute graft versus host disease and moderate to severe acute respiratory distress syndrome. Mesoblast is completing Phase 3 trials for its product candidates for advanced heart failure and chronic low back pain. Two products have been commercialized in Japan and Europe by Mesoblast's licensees, and the Company has established commercial partnerships in Europe and China for certain Phase 3 assets.

Mesoblast has locations in Australia, the United States and Singapore and is listed on the Australian Securities Exchange (MSB) and on the Nasdaq (MESO). For more information, please see <a href="http://www.mesoblast.com">www.mesoblast.com</a>, LinkedIn: Mesoblast Limited and Twitter: @Mesoblast

#### Forward-Looking Statements

This announcement includes forward-looking statements that relate to future events or our future financial performance and involve known and unknown risks, uncertainties and other factors that may cause our actual results, levels of activity, performance or achievements or achievements to differ materially from any future results, levels of activity, performance or achievements expressed or implied by these forward-looking statements. All statements other than statements of historical fact are forward-looking statements, which are often indicated by terms such as "anticipate," "could," "estimate," "expect," "goal," "intend," "likely," "look forward to," "may," "plan," "potential," "predict," "project," "should," "will," "would" and similar expressions and variations thereof. We make such forward-looking statements pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995 and other federal securities laws. Forward-looking statements should not be read as a guarantee of future performance or results, and actual results may differ from the results anticipated in these forward-looking statements, and the differences may be material and adverse. The risks, uncertainties and other factors that may impact our forward-looking statements include, but are not limited to: statements about the initiation, timing, progress and results of Mesoblast and its collaborators' clinical studies; in approved; the potential benefits of strategic collaboration agreements and Mesoblast's ability to maintain established strategic collaborations; Mesoblast's ability to establish and maintain intellectual property on its product candidates and Mesoblast's ability to successfully defend these in cases of alleged infringement. You should read this press release together with our risk factors, in our most recently filed reports with the SEC or on our website. Uncertainties and risks that may cause forward-looking statements to be materially different from those which may be expressed or implied by such st

Release authorized by the Chief Executive.

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