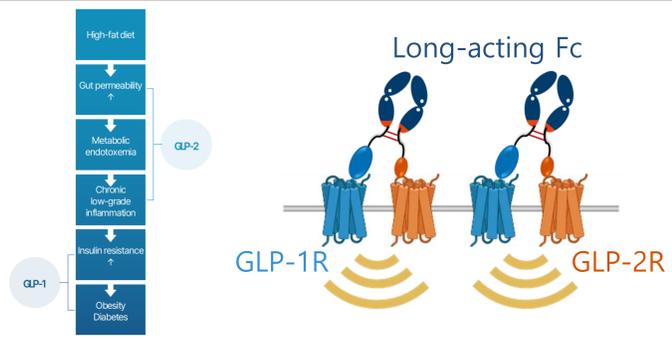
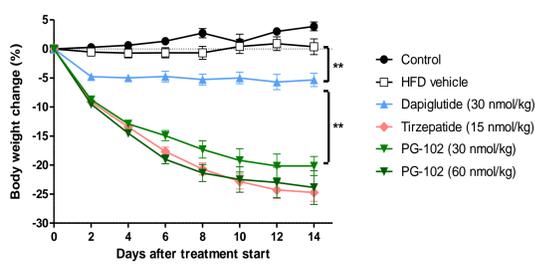
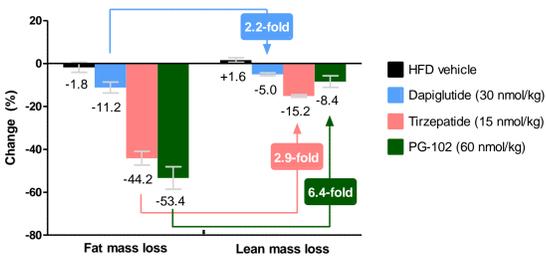
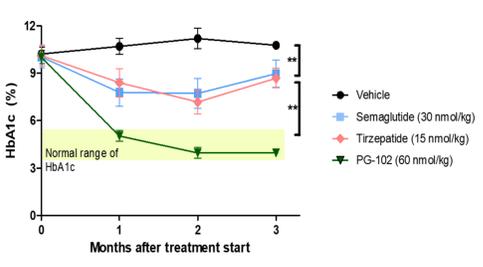
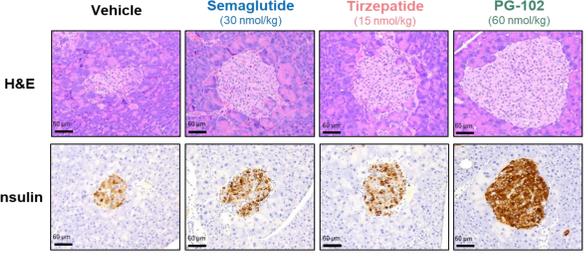
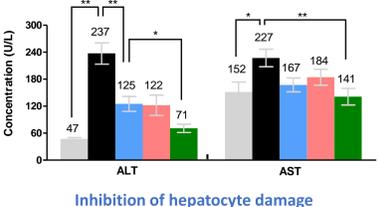
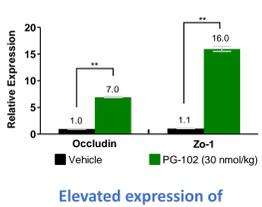
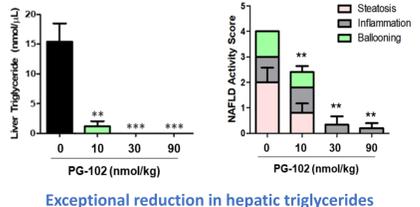
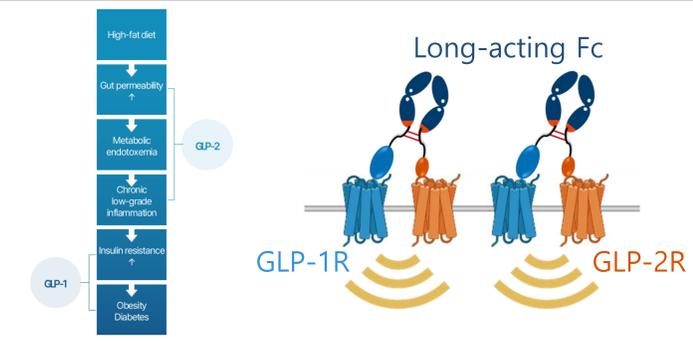


Phase 1 clinical development of PG-102, a next-generation GLP-1/GLP-2 dual agonist for treating obesity and diabetes

ProGen Co., Ltd.

Disease area	Metabolic diseases									
Product Type	Bispecific fusion protein									
Indication	Obesity, Type 2 Diabetes									
Target	GLP-1R and GLP-2R									
Mechanism of Action										
Competitiveness	<table border="1"> <tr> <td>Efficacy</td> <td> <ul style="list-style-type: none"> Superior quality of body weight loss compared to Tirzepatide Fat mass loss ↑ + lean mass loss ↓ Breakthrough glycemic control compared to Semaglutide and Tirzepatide Glucose uptake ↑ + β-cell protection Comorbidity prevention in obesity/diabetes </td> <td>Data 1 Data 2 Data 3</td> </tr> <tr> <td>Safety</td> <td> <ul style="list-style-type: none"> Improved safety & tolerability Gastrointestinal tract inflammation ↓ </td> <td>Data 3</td> </tr> <tr> <td>Convenience</td> <td> <ul style="list-style-type: none"> Extended dosing interval allowing for at least biweekly to monthly administration </td> <td></td> </tr> </table>	Efficacy	<ul style="list-style-type: none"> Superior quality of body weight loss compared to Tirzepatide Fat mass loss ↑ + lean mass loss ↓ Breakthrough glycemic control compared to Semaglutide and Tirzepatide Glucose uptake ↑ + β-cell protection Comorbidity prevention in obesity/diabetes 	Data 1 Data 2 Data 3	Safety	<ul style="list-style-type: none"> Improved safety & tolerability Gastrointestinal tract inflammation ↓ 	Data 3	Convenience	<ul style="list-style-type: none"> Extended dosing interval allowing for at least biweekly to monthly administration 	
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Safety	<ul style="list-style-type: none"> Improved safety & tolerability Gastrointestinal tract inflammation ↓ 	Data 3								
Convenience	<ul style="list-style-type: none"> Extended dosing interval allowing for at least biweekly to monthly administration 									
Development Stage	Phase 1b									
Route of Administration	Subcutaneous injection									
Key Data	<ol style="list-style-type: none"> PG-102 achieves comprehensive weight loss and enhanced fat-to-lean loss ratio compared to Dapigliptide and Tirzepatide   PG-102 excels in glycemic control, achieves normoglycemia, and enhances β-cell protection compared to Semaglutide and Tirzepatide   PG-102 demonstrates effective prevention of comorbidities through superior systemic inflammation control    									
IP	<p>KR Registered (KR10-2349717, KR10-2349718) Under review in Key countries (including US, EP, CN, ID, BR)</p>									

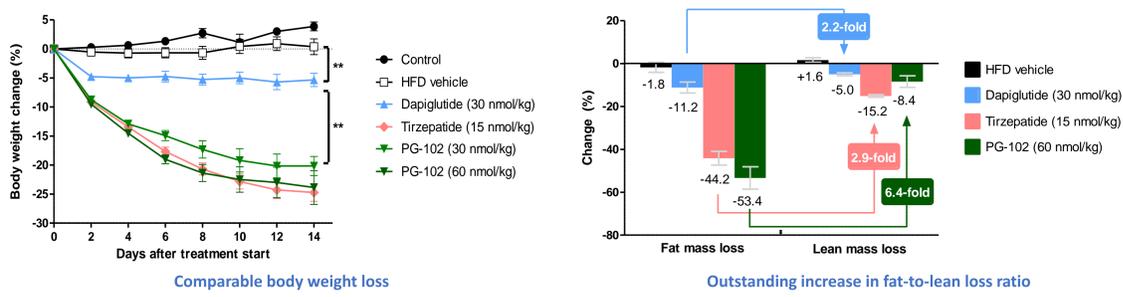


A biased GLP-1R/GLP-2R dual agonist with extended PK properties

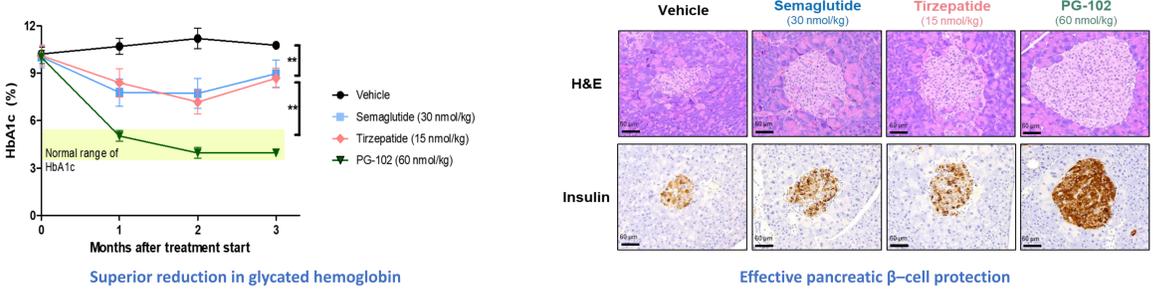
- Relative potency of PG-102 against GLP-1 receptor and GLP-2 receptor is sophisticatedly optimized for simultaneously activation of both receptors.
- PG-102 also features NTIG[®], a long-acting Fc platform technology, for a prolonged action.

Efficacy	<ul style="list-style-type: none"> Superior quality of body weight loss compared to Tirzepatide Fat mass loss ↑ + lean mass loss ↓ Breakthrough glycemic control compared to Semaglutide and Tirzepatide Glucose uptake ↑ + β-cell protection Comorbidity prevention in obesity/diabetes 	Data 1 Data 2 Data 3
Safety	<ul style="list-style-type: none"> Improved safety & tolerability Gastrointestinal tract inflammation ↓ 	Data 3
Convenience	<ul style="list-style-type: none"> Extended dosing interval allowing for at least biweekly to monthly administration 	

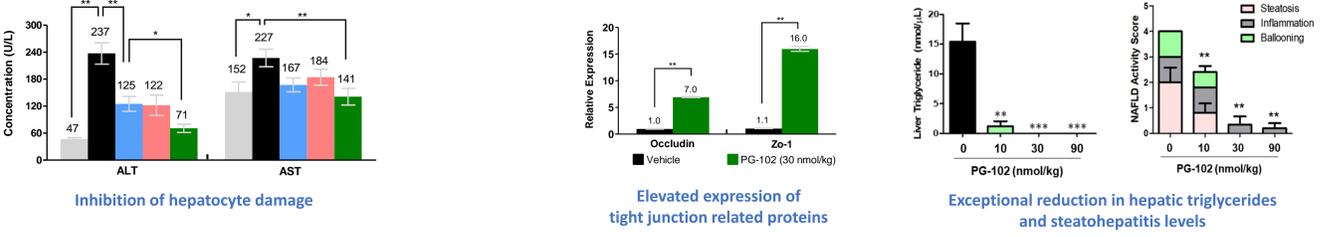
1. PG-102 achieves comprehensive weight loss and enhanced fat-to-lean loss ratio compared to Dapigliptide and Tirzepatide



2. PG-102 excels in glycemic control, achieves normoglycemia, and enhances β-cell protection compared to Semaglutide and Tirzepatide



3. PG-102 demonstrates effective prevention of comorbidities through superior systemic inflammation control



KR Registered (KR10-2349717, KR10-2349718)
Under review in Key countries (including US, EP, CN, ID, BR)